PROTOZOOLOGY
PROTOZOOLOGY

By

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With three hundred and seventy-six illustrations

Fourth Edition

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"The revelations of the Microscope are perhaps not excelled in importance by those of the telescope. While exciting our curiosity, our wonder and admiration, they have proved of infinite service in advancing our knowledge of things around us."

Leidy
Preface

The fourth edition of Protozoology maintains its original aim in setting forth "introductory information on the common and representative genera of all groups of both free-living and parasitic Protozoa" for seniors and graduates in zoology in colleges and universities. It has been noted in recent years that students frequently wished to obtain a fuller knowledge on certain topics, organisms, processes, etc., than that which was found in the former edition. In order to meet this need without too great an expansion, references have been given to various items in the text and a list of a much larger number of literature has been appended to each chapter. Furthermore, this enlargement of references increases the usefulness of this work to advanced students, teachers of biology, field workers in various areas of biological science, veterinarians, physicians, public health workers, laboratory diagnosticians and technicians, etc.

While the chapter arrangement remains the same as before, a thorough revision has been carried on throughout the text in the light of many recently published contributions to protozoology. Good illustrations are indispensable in this kind of work, since they are far more easily comprehended than lengthy statements. Therefore, old illustrations were replaced by more suitable ones and many new illustrations have been added, bringing up the total number of the text figures now to 376. Except diagrams, all figures are accompanied by the scales of magnification. For illustrations that have been adopted from published papers, the indebtedness of the author is expressed by mentioning the authors' names.

R. R. Kudo

Urbana, Illinois
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PROTOZOOLOGY
PART I: GENERAL BIOLOGY
Chapter 1

Introduction

PROTOZOA are unicellular animals. The body of a protozoan is morphologically a single cell and manifests all characteristics common to the living thing. The various activities which make up the phenomena of life are carried on by parts within the body or cell. These parts are comparable with the organs of a metazoan which are composed of a large number of cells grouped into tissues and are called organellae or cell-organs. Thus the one-celled protozoan is a complete organism somewhat unlike the cell of a metazoan, each of which is dependent upon other cells and cannot live independently. From this viewpoint, certain students of protozoology maintain that the Protozoa are non-cellular, and not unicellular, organisms. Dobell (1911), for example, pointed out that the term “cell” is employed to designate (1) the whole protozoan body, (2) a part of a metazoan organism, and (3) a potential whole organism (a fertilized egg) which consequently resulted in a confused state of knowledge regarding living things, and, therefore, proposed to define a cell as a mass of protoplasm composing part of an organism, and further considered that the protozoan is a non-cellular but complete organism, differently organized as compared with cellular organisms, the Metazoa and Metaphyta. Although some writers (Hyman, 1940; Lwoff, 1951) follow this view, the great majority of protozoologists continue to consider the Protozoa as unicellular animals. Through the processes of organic evolution, they have undergone cytological differentiation and the Metazoa histological differentiation.

In being unicellular, the Protozoa and the Protophyta are alike. The majority of Protozoa may be distinguished from the majority of Protophyta on the basis of dimensions, methods of nutrition, direction of division-plane, etc. While many Protophyta possess nuclear material, it is not easy to detect it in many forms; on the other hand, all Protozoa contain at least one easily observable nucleus. The binary fission of Protozoa and Protophyta is longitudinal and transverse respectively. Most of Ciliata, however, multiply by transverse division. In general the nutrition of Protozoa is holozoic and of Protophyta, holophytic or saprophytic; but there are large numbers of Protozoa which nourish themselves by the latter methods. Thus an absolute and clean-cut separation of the two groups of unicellular organisms is not possible. Haeckel (1866) coined the name Protista to include these organisms in a single group, but this is not generally
adopted, since it includes undoubted animals and plants, thus creating an equal amount of confusion between it and the animal or the plant. Calkins (1933) excluded chromatophore-bearing Mastigophora from his treatment of Protozoa, thus placing organisms similar in every way, except the presence or absence of chromatophores, in two different (animal and plant) groups. This intermingling of characteristics between the two groups of microorganisms shows clearly their close interrelationship and suggests strongly their common ancestry.

Although the majority of Protozoa are solitary and the body is composed of a single cell, there are several forms in which the organism is made up of more than one cell. These forms, which are called colonial Protozoa (p. 173), are well represented by the members of Phytomastigina, in which the individuals are either joined by cytoplasmic threads or embedded in a common matrix. These cells are alike both in structure and in function, although in a few forms there may be a differentiation of the individuals into reproductive and vegetative cells. Unlike the cells in a metazoan which form tissues, these vegetative cells of colonial Protozoa are not so dependent upon other cells as are the cells in Metazoa; therefore, they do not form any true tissue. The reproductive cells produce zygotes through sexual fusion, which subsequently undergo repeated division and may produce a stage comparable with the blastula stage of a metazoan, but never reaching the gastrula stage. Thus, colonial Protozoa are only cell-aggregates without histological differentiation and may thus be distinguished from the Metazoa.

An enormous number of species of Protozoa are known to man. From comparatively simple forms such as Amoeba, up to highly complicated organisms as represented by numerous ciliates, the Protozoa vary exceedingly in their body organization, morphological characteristics, behavior, habitat, etc., which necessitates a taxonomic arrangement for proper consideration as set forth in detail in Chapters 8 to 44.

Relationship of protozoology to other fields of biological science

A brief consideration of the relationship of Protozoology to other fields of biology and its possible applications may not be out of place here. Since the Protozoa are single-celled animals manifesting the characteristics common to all living things, they have been studied by numerous investigators with a view to discovering the nature and mechanism of various phenomena, the
INTRODUCTION

sum-total of which is known collectively as life. Though the investigators generally have been disappointed in the results, inasmuch as the assumed simplicity of unicellular organisms has proved to be offset by the complexity of their cell-structure, nevertheless discussion of any biological principles today must take into account the information obtained from studies of Protozoa. It is now commonly recognized that adequate information on various types of Protozoa is a prerequisite to a thorough comprehension of biology and to proper application of biological principles.

Practically all students agree in assuming that the higher types of animals have been derived from organisms which existed in the remote past and which probably were somewhat similar to the primitive Protozoa of the present day. Since there is no sharp distinction between the Protozoa and the Protophyta or between the Protozoa and the Metazoa, and since there are intermediate forms between the major classes of the Protozoa themselves, progress in protozoology contributes toward the advancement of our knowledge on the probable steps by which living things in general evolved.

Geneticists have undertaken studies on heredity and variation among Protozoa. "Unicellular animals," wrote Jennings (1909), "present all the problems of heredity and variation in miniature. The struggle for existence in a fauna of untold thousands showing as much variety of form and function as any higher group, works itself out, with ultimate survival of the fittest, in a few days under our eyes, in a finger bowl. For studying heredity and variation we get a generation a day, and we may keep unlimited numbers of pedigreed stock in a watch glass that can be placed under the microscope." Morphological and physiological variations are encountered commonly in all forms. Whether variation is due to germinal or environmental conditions, is often difficult to determine. Studies on conjugation in Paramecium by utilizing the mating types first noted by Sonneborn (1937, 1938) not only brought to light a wealth of important information regarding the genetics of Protozoa, but also are revealing a close insight concerning the relationship between the nuclear and cytoplasmic factors of heredity in the animal.

Parasitic Protozoa are confined to one or more specific hosts. Through studies of the forms belonging to one and the same genus or species, the phylogenetic relation among the host animals may be established or verified. The mosquitoes belonging to the genera Culex and Anopheles, for instance, are known to transmit avian and human Plasmodium respectively. They are further infected by specific microsporidian parasites. For instance, *Thelohania legeri*
has been found widely only in many species of anopheline mosquitoes; *T. opacita* has, on the other hand, been found exclusively in culicine mosquitoes, although the larvae of the species belonging to these two genera live frequently in the same body of water (Kudo, 1924, 1925). By observing certain intestinal Protozoa in some monkeys, Hegner (1928) obtained evidence on the probable phylogenetic relationship between them and other higher mammals. The relation of various Protozoa of the wood-roach to those of the termite, as revealed by Cleveland and his associates (1934), gives further proof that the Blattidae and the Isoptera are closely related.

Study of a particular group of parasitic Protozoa and their hosts may throw light on the geographic condition of the earth which existed in the remote past. The members of the genus *Zelleriella* are usually found in the colon of the frogs belonging to the family Leptodactylidae. Through an extensive study of these amphibians from South America and Australia, Metcalf (1920, 1929) found that the species of *Zelleriella* occurring in the frogs of the two continents are almost identical. He finds it more difficult to conceive of convergent or parallel evolution of both the hosts and the parasites, than to assume that there once existed between Patagonia and Australia a land connection over which frogs, containing *Zelleriella*, migrated.

Experimental studies of large Protozoa have thrown light on the relation between the nucleus and the cytoplasm, and have furnished a basis for an understanding of regeneration in animals. In Protozoa we find various types of nuclear divisions ranging from a simple amitotic division to a complex process comparable in every detail with the typical metazoan mitosis. A part of our knowledge in cytology is based upon studies of Protozoa.

Through the efforts of various investigators in the past fifty years, it has now become known that some 25 species of Protozoa occur in man. *Entamoeba histolytica*, *Balantidium coli*, and four species of *Plasmodium*, all of which are pathogenic to man, are widely distributed throughout the world. In certain restricted areas are found other pathogenic forms, such as Trypanosoma and Leishmania. Since all parasitic Protozoa presumably have originated in free-living forms and since our knowledge of the morphology, physiology, and reproduction of the parasitic forms has largely been obtained in conjunction with the studies of the free-living organisms, a general knowledge of the entire phylum is necessary to understand these parasitic forms.

Recent studies have further revealed that almost all domestic animals are hosts to numerous parasitic Protozoa, many of which
are responsible for serious infectious diseases. Some of the forms found in domestic animals are morphologically indistinguishable from those occurring in man. Balantidium coli is considered as a parasite of swine, and man is its secondary host. Knowledge of protozoan parasites is useful to medical practitioners, just as it is essential to veterinarians inasmuch as certain diseases of animals, such as southern cattle fever, dourine, nagana, blackhead, coccidiosis, etc., are caused by Protozoa.

Sanitary betterment and improvement are fundamental requirements in the modern civilized world. One of man’s necessities is safe drinking water. The majority of Protozoa live freely in various bodies of water and some of them are responsible, if present in sufficiently large numbers, for giving certain odors to the waters of reservoirs or ponds (p. 114). But these Protozoa which are occasionally harmful are relatively small in number compared with those which are beneficial to man. It is generally understood that bacteria live on various waste materials present in the polluted water, but that upon reaching a certain population, they would cease to multiply and would allow the excess organic substances to undergo decomposition. Numerous holozoic Protozoa, however, feed on the bacteria and prevent them from reaching the saturation population. Protozoa thus seem to help indirectly in the purification of the water. Protozoology therefore must be considered as part of modern sanitary science.

Young fish feed extensively on small aquatic organisms, such as larvae of insects, small crustaceans, annelids, etc., all of which depend largely upon Protozoa and Protophyta as sources of food supply. Thus the fish are indirectly dependent upon Protozoa as food material. On the other hand, there are numbers of Protozoa which live at the expense of fish. The Myxosporidia are almost exclusively parasites of fish and sometimes cause death to large numbers of commercially important fishes (Kudo, 1920) (p. 648). Success in fish-culture, therefore, requires among other things a thorough knowledge of Protozoa.

Since Russel and Hutchinson (1909) suggested some forty years ago that Protozoa are probably a cause of limitation of the numbers, and therefore the activities of bacteria in the soil and thus tend to decrease the amount of nitrogen which is given to the soil by the nitrifying bacteria, several investigators have brought out the fact that in the soils of temperate climate various sarcodinans, flagellates and less frequently ciliates, are present and active throughout the year. The exact relation between specific Protozoa and bacteria in
the soil is not yet clear in spite of the numerous experiments and observations. All soil investigators should be acquainted with the biology and taxonomy of free-living Protozoa.

It is a matter of common knowledge that the silkworm and the honey bee suffer from microsporidian infections (p. 670). Sericulture in south-western Europe suffered great damages in the middle of the nineteenth century because of the "pebrine" disease, caused by the microsporidian, Nosema bombycis. During the first decade of the present century, another microsporidian, Nosema apis, was found to infect a large number of honey bees. Methods of control have been developed and put into practice so that these microsporidian infections are at present not serious, even though they still occur. On the other hand, other Microsporidia are now known to infect certain insects, such as mosquitoes and lepidopterous pests, which, when heavily infected, die sooner or later. Methods of destruction of these insects by means of chemicals are more and more used, but attention should also be given to biological control of them by means of Protozoa and Protophyta.

While the majority of Protozoa lack permanent skeletal structures and their fossil forms are little known, there are at least two large groups in the Sarcodina which possess conspicuous shells and which are found as fossils. They are Foraminifera and Radiolaria. From early palaeozoic era down to the present day, the carbonate of lime which makes up the skeletons of numerous Foraminifera has been left embedded in various rock strata. Although there is no distinctive foraminiferan fauna characteristic of a given geologic period, there are certain peculiarities of fossil Foraminifera which distinguish one formation from the other. From this fact one can understand that knowledge of foraminiferous rocks is highly useful in checking up logs in well drilling. The skeletons of the Radiolaria are the main constituent of the ooze of littoral and deep-sea regions. They have been found abundantly in siliceous rocks of the palaeozoic and the mesozoic eras, and are also identified with the clays and other formations of the miocene period. Thus knowledge of these two orders of Sarcodina, at least, is essential for the student of geology and paleontology.

The history of protozoology

Aside from a comparatively small number of large forms, Protozoa are unobservable with the naked eye, so that one can easily understand why they were unknown prior to the invention of the microscope. Antony van Leeuwenhoek (1632-1723) is commonly recog-
nized as the father of protozoology. Grinding lenses himself, Leeuwenhoek made more than 400 simple lenses, including one which, it is said, had a magnification of 270 times (Harting). Among the many things he discovered were various Protozoa. According to Dobell (1932), Leeuwenhoek saw in 1674 for the first time free-living fresh-water Protozoa. Between 1674 and 1716, he observed many Protozoa which he reported to the Royal Society of London and which, as Dobell interpreted, were Euglena ("green in the middle, and before and behind white"), Vorticella, Stylonychia, Carchesium, Volvox, Coleps, Kerona, Anthophysis, Elphidium, etc. Huygens gave in 1678 "unmistakable descriptions of Chilodon(-ella), Paramecium, Astasia and Vorticella, all found in infusions" (Dobell).

Colpoda was seen by Buonanni (1691) and Harris (1696) rediscovered Euglena. In 1718 there appeared the first treatise on microscopic organisms, particularly of Protozoa, by Joblot who emphasized the non-existence of abiogenesis by using boiled hay-infusions in which no Infusoria developed without exposure to the atmosphere. This experiment confirmed that of Redi who, some 40 years before, had made his well-known experiments by excluding flies from meat. Joblot illustrated, according to Woodruff (1937), Paramecium, the slipper animalcule, with the first identifiable figure. Trembley (1744) studied division in some ciliates, including probably Paramecium, which generic name was coined by Hill in 1752. Noctiluca was first described by Baker (1753).

Rösel von Rosenhof (1755) observed an organism, which he called "der kleine Proteus," and also Vorticella, Stentor, and Volvox. The "Proteus" which Linnaeus named Volvox chaos (1758) and later renamed Chaos protheus (1767), cannot be identified with any of the known amoeboid organisms (Kudo, 1946). Wrisberg (1764) coined the term "Infusoria" (Dujardin; Woodruff). By using the juice of geranium, Ellis (1769) caused the extrusion of the "fins" (trichocysts) in Paramecium. Eichhorn (1783) observed the heliozoan, Actinosphaerium, which now bears his name. O. F. Müller described Ceratium a little later and published two works on the Infusoria (1773, 1786) although he included unavoidably some Metazoa and Protophyta in his monographs, some of his descriptions and figures of Ciliata were so well done that they are of value even at the present time. Lamareck (1816) named Folliculina.

At the beginning of the nineteenth century the cylcosis in Paramecium was brought to light by Gruithuisen. Goldfuss (1817) coined the term Protozoa, including in it the coelenterates. Nine years later there appeared d’Orbigny’s systematic study of the Foramini-
fera, which he considered "microscopical cephalopods." In 1828
Ehrenberg began publishing his observations on Protozoa and in
1838 he summarized his contributions in Die Infusionsthierchen als
vollkommene Organismen, in which he diagnosed genera and species
so well that many of them still hold good. Ehrenberg excluded Rotato-
toria and Cercaria from Infusoria. Through the studies of Ehrenberg
the number of known Protozoa increased greatly; he, however, pro-
posed the term "Polygastrichia," under which he placed Mastigo-
phora, Rhizopoda, Ciliata, Suctoria, desmids, etc., since he believed
that the food vacuoles present in them were stomachs. This hypothe-
sis became immediately the center of controversy, which incidentally,
together with the then-propounded cell theory and improvements in
microscopy, stimulated researches on Protozoa.

Dujardin (1835) took pains in studying the protoplasm of various
Protozoa and found it alike in all. He named it sarcode. In 1841 he
published an extensive monograph of various Protozoa which came
under his observations. The term Rhizopoda was coined by this
investigator. The commonly used term protoplasm was employed by
Purkinje (1840) in the same sense as it is used today. The Protozoa
was given a distinct definition by Siebold in 1845, as follows: "Die
Thiere, in welchen die verschiedenen Systeme der Organe nicht
scharf ausgeschieden sind, und deren unregelmässige Form und ein-
fache Organization sich auf eine Zelle reduzieren lassen." Siebold
subdivided Protozoa into Infusoria and Rhizopoda. The sharp differ-
entiation of Protozoa as a group certainly inspired numerous micros-
copists. As a result, several students brought forward various group
names, such as Radiolaria (J. Müller, 1858), Ciliata (Perty, 1852),
Flagellata (Cohn, 1853), Suctoria (Claparède and Laehmann, 1858),
Heliozoa, Protista (Haeckel, 1862, 1866), Mastigophora (Diesing,
1865), etc. Of Suctoria, Stein failed to see the real nature (1849), but
his two monographs on Ciliata and Mastigophora (1854, 1859–1883)
contain concise descriptions and excellent illustrations of numerous
species. Haeckel who went a step further than Siebold by distinguishing
between Protozoa and Metazoa, devoted 10 years to his study of
Radiolaria, especially those of the Challenger collection, and de-
scribed in his celebrated monographs more than 4000 species.

In 1879 the first comprehensive monograph on the Protozoa of
North America was put forward by Leidy under the title of Fresh-
water Rhizopods of North America, which showed the wide distribu-
tion of many known forms of Europe and revealed a number of new
and interesting forms. This work was followed by Stokes' The Fresh-
water Infusoria of the United States, which appeared in 1888.
INTRODUCTION

Bütschli (1880–1889) established Sarcodina and made an excellent contribution to the taxonomy of the then-known species of Protozoa, which is still considered as one of the most important works on general protozoology. The painstaking researches by Maupas, on the conjugation of ciliates, corrected erroneous interpretation of the phenomenon observed by Balbiani some 30 years before and gave impetus to a renewed cytological study of Protozoa. The variety in form and structure of the protozoan nuclei became the subject of intensive studies by several cytologists. Weismann put into words the immortality of the Protozoa. Schaudinn contributed much toward the cytological and developmental studies of Protozoa.

In the first year of the present century, Calkins in the United States and Doflein in Germany wrote modern textbooks of protozoology dealing with the biology as well as the taxonomy. Jennings devoted his time for nearly 40 years to the study of genetics of Protozoa. Recent development of bacteria-free culture technique in certain flagellates and ciliates, has brought to light important information regarding the nutritional requirements and metabolism of these organisms.

Today the Protozoa are more and more intensively and extensively studied from both the biological and the parasitological sides, and important contributions appear continuously. Since all parasitic Protozoa appear to have originated in free-living forms, the comprehension of the morphology, physiology, and development of the latter group is obviously fundamentally important for a thorough understanding of the former group.

Compared with the advancement of our knowledge on free-living Protozoa, that on parasitic forms has been very slow. This is to be expected, of course, since the vast majority of them are so minute that the discovery of their presence has been made possible only through improvements in the microscope and in technique.

Here again Leeuwenhoek seems to have been the first to observe a parasitic protozoan, for he observed, according to Dobell (1932), in the fall of 1674, the oocysts of the coccidian Eimeria stiedae, in the contents of the gall bladder of an old rabbit; in 1681, Giardia intestinalis in his own diarrhoeic stools; and in 1683, Opalina and Nyctotherus in the gut contents of frogs. The oral Trichomonas of man was observed by O. F. Müller (1773) who named it Cercaria tenax (Dobell, 1939). There is no record of anyone having seen Protozoa living in other organisms, until 1828, when Dufour’s account of the gregarine from the intestine of coleopterous insects appeared. Some ten years later, Hake rediscovered the oocysts of Eimeria stiedae. A
flagellate was observed in the blood of salmon by Valentín in 1841, and the frog trypanosome was discovered by Gluge (1842) and Gruby (1843), the latter author creating the genus Trypanosoma for it.

The gregarines were a little later given attention by Kölliker (1848) and Stein (1848). The year 1849 marks the first record of an amoeba being found in man, for Gros then observed Entamoeba gingivalis in the human mouth. Five years later, Davaine found in the stools of cholera patients two flagellates (Trichomonas and Chilomastix). Kloss in 1855 observed the coccidian, Klossia helicina, in the excretory organ of Helix; and Eimer (1870) made an extensive study of Coccidia occurring in various animals. Balantidium coli was discovered by Malmsten in 1857. Lewis in 1870 observed Entamoeba coli in India, and Lösch in 1875 found Entamoeba histolytica in Russia. During the early part of the last century, an epidemic disease, pébrine, of the silkworm appeared in Italy and France, and a number of biologists became engaged in its investigation. Foremost of all, Pasteur (1870) made an extensive report on the nature of the causative organism, now known as Nosema bombycis, and also on the method of control and prevention. Perhaps this is the first scientific study of a parasitic protozoan which resulted in an effective practical method of control of its infection.

Lewis observed in 1878 an organism which is since known as Trypanosoma lewisi in the blood of rats. In 1879 Leuckart created the group Sporozoa, including in it the gregarines and coccidians. Other groups under Sporozoa were soon definitely designated. They are Myxosporidia (Bütschli, 1881), Microsporidia and Sarcosporidia (Balbiani, 1882).

Parasitic protozoology received a far-reaching stimulus when Laveran (November, 1880) discovered the microgamete formation (“flagellation”) of a malaria parasite in the human blood. Smith and Kilborne (1893) demonstrated that Babesia of the Texas fever of cattle in the southern United States was transmitted by the cattle tick from host to host, and thus revealed for the first time the close relationship which exists between an arthropod and a parasitic protozoan. Two years later Bruce discovered Trypanosoma brucei in the blood of domestic animals suffering from “nagana” disease in Africa and later (1897) demonstrated by experiments that the tsetse fly transmits the trypanosome. Studies of malaria organisms continued and several important contributions appeared. Golgi (1886, 1889) studied the schizogony and its relation to the occurrence of fever, and was able to distinguish the types of fever. MacCallum (1897)
INTRODUCTION

observed the microgamete formation in Haemoproteus of birds and suggested that the "flagella" observed by Laveran were microgametes of Plasmodium. In fact, he later observed the formation of the zygote through fusion of a microgamete and a macrogamete of *Plasmodium falciparum*. Almost at the same time, Schaudinn and Siedleke (1897) showed that anisogamy results in the production of zygotes in Coccidia. The latter author published later further observations on the life-cycle of Coccidia (1898, 1899).

Ross (1898, 1898a) revealed the development of *Plasmodium relictum* (*P. praecox*) in *Culex fatigans* and established the fact that the host birds become infected by this protozoan through the bites of the infected mosquitoes. Since that time, investigators too numerous to mention here (p. 600), studied the biology and development of the malarial organisms. Among the more recent findings is the exo-erythrocytic development, fuller information on which is now being sought. In 1902, Dutton found that the sleeping sickness in equatorial Africa was caused by an infection by *Trypanosoma gambiense*. In 1903, Leishman and Donovan discovered simultaneously *Leishmania donovani*, the causative organism of "kala-azar" in India.

Artificial cultivation of bacteria had contributed toward a very rapid advancement in bacteriology, and it was natural, as the number of known parasitic Protozoa rapidly increased, that attempts to cultivate them in vitro should be made. Musgrave and Clegg (1904) cultivated, on bouillon-agar, small free-living amoebae from old faecal matter. In 1905 Novy and MacNeal cultivated successfully the trypanosome of birds in blood-agar medium, which remained free from bacterial contamination and in which the organisms underwent multiplication. Almost all species of Trypanosoma and Leishmania have since been cultivated in a similar manner. This serves for detection of a mild infection and also identification of the species involved. It was found, further, that the changes which these organisms underwent in the culture media were imitative of those that took place in the invertebrate host, thus contributing toward the life-cycle studies of them.

During and since World War I, it became known that numerous intestinal Protozoa of man are widely present throughout the tropical, subtropical and temperate zones. Taxonomic, morphological and developmental studies on these forms have therefore appeared in an enormous number. Cutler (1918) seems to have succeeded in cultivating *Entamoeba histolytica*, though his experiment was not repeated by others. Barret and Yarborough (1921) culti-
vated *Balantium coli* and Boeck (1921) cultivated *Chilomastix mesnili*. Boeck and Drbohlav (1925) succeeded in cultivating *Entamoeba histolytica*, and their work was repeated and improved upon by many investigators. While the in-vitro cultivation has not thrown much light on metabolic activities of this and other parasitic amoebae, as no one of them would grow in culture without some other organisms, it has increased our knowledge on the biology of these parasites.

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Chapter 2

Ecology

With regard to their habitats, the Protozoa may be divided into free-living forms and those living on or in other organisms. Mastigophora, Sarcodina, Ciliata, and Suctoria include both free-living and parasitic Protozoa, but Sporozoa are exclusively parasitic.

Free-living Protozoa

The vegetative or trophic stages of free-living Protozoa have been found in every type of fresh and salt water, soil and decaying organic matter. Even in the circumpolar regions or at extremely high altitudes, certain protozoa occur at times in fairly large numbers. The factors, which influence their distribution in a given body of water, are temperature, light, chemical composition, acidity, kind and amount of food, and degree of adaptability of the individual protozoans to various environmental changes. Their early appearance as living organisms, their adaptability to various habitats, and their capacity to remain viable in the encysted condition, probably account for the wide distribution of the Protozoa throughout the world. The common free-living amoebae, numerous testaceans and others, to mention a few, of fresh waters, have been observed in innumerable places of the world.

Temperature. The majority of Protozoa are able to live only within a small range of temperature variation, although in the encysted state they can withstand a far greater temperature fluctuation. The lower limit of the temperature is marked by the freezing of the protoplasm, and the upper limit by the destructive chemical change within the body protoplasm. The temperature toleration seems to vary among different species of Protozoa; and even in the same species under different conditions. For example, Chalkley (1930) placed Paramecium caudatum in 4 culture media (balanced saline, saline with potassium excess, saline with calcium excess, and saline with sodium excess), all with pH from 5.8 or 6 to 8.4 or 8.6, at 40°C. for 2–16 minutes and found that (1) the resistance varies with the hydrogen-ion concentration, maxima appearing in the alkaline and acid ranges, and a minimum at or near about 7.0; (2) in a balanced saline, and in saline with an excess of sodium or potassium, the alkaline maximum is the higher, while in saline with an excess of calcium, the acid maximum is the higher; (3) in general, acidity decreases and alkalinity increases resistance; and (4) between pH 6.6
and 7.6, excess of potassium decreases resistance and excess of calcium increases resistance. Glaser and Coria (1933) cultivated *Paramecium caudatum* on dead yeast free from living organisms at 20–28°C. (optimum 25°C.) and noted that at 30°C. the organisms were killed. Doudoroff (1936), on the other hand, found that in *P. multimicronucleatum* its resistance to raised temperature was low in the presence of food, but rose to a maximum when the food was exhausted, and there was no appreciable difference in the resistance between single and conjugating individuals.

The thermal waters of hot springs have been known to contain living organisms including Protozoa. Glaser and Coria (1935) obtained from the thermal springs, of Virginia, several species of Mastigophora, Ciliata, and an amoeba which were living in the water, the temperature of which was 34–36°C., but did not notice any protozoan in the water which showed 39–41°C. Uyemura (1936, 1937) made a series of studies on Protozoa living in various thermal waters of Japan, and reported that many species lived at unexpectedly high temperatures. Some of the Protozoa observed and the temperatures of the water in which they were found are as follows: *Amoeba* sp., *Vahlkampfia limax*, *A. radios*, 30–51°C.; *Amoeba verrucosa*, *Chilodonella* sp., *Lionotus fasciola*, *Paramecium caudatum*, 36–40°C.; *Oxytricha fallax*, 30–56°C.

Under experimental conditions, it has been shown repeatedly that many protozoans become accustomed to a very high temperature if the change be made gradually. Dallinger (1887) showed a long time ago that *Tetramitus rostratus* and two other species of flagellates became gradually acclimatized up to 70°C. in several years. In nature, however, the thermal death point of most of the free-living Protozoa appears to lie between 36° and 40°C. and the optimum temperature, between 16° and 25°C.

On the other hand, the low temperature seems to be less detrimental to Protozoa than the higher one. Many protozoans have been found to live in water under ice, and several haematochrome-bearing Phytomastigina undergo vigorous multiplication on snow in high altitudes, producing the so-called "red snow." Klebs (1893) subjected the trophozoites of Euglena to repeated freezing without apparent injury and Jahn (1933) found no harmful effect when Euglena cultures were kept without freezing at −0.2°C. for one hour, but when kept at −4°C. for one hour the majority were killed. Gaylord (1908) exposed *Trypanosoma gambiense* to liquid air for 20 minutes without apparent injury, but the organisms were killed after 40 minutes' immersion.
Kühne (1864) observed that Amoeba and Actinophrys suffered no ill effects when kept at 0°C. for several hours as long as the culture medium did not freeze, but were killed when the latter froze. Molisch (1897) likewise noticed that Amoeba dies as soon as the ice forms in its interior or immediate vicinity. Chambers and Hale (1932) demonstrated that internal freezing could be induced in an amoeba by inserting an ice-tipped pipette at −0.6°C., the ice spreading in the form of fine featherly crystals from the point touched by the pipette. They found that the internal freezing kills the amoebae, although if the ice is prevented from forming, a temperature as low as −5°C. brings about no visible damage to the organism. At 0°C., Deschiens (1934) found the trophozoites of Entamoeba histolytica remained alive, though immobile, for 56 hours, but were destroyed in a short time when the medium froze at −5°C.

According to Greeley (1902), when Stentor coerulescens was slowly subjected to low temperatures, the cilia kept on beating at 0°C. for 1–3 hours, then cilia and gullet were absorbed, the ectoplasm was thrown off, and the body became spherical. When the temperature was raised, this spherical body is said to have undergone a reverse process and resumed its normal activity. If the lowering of temperature is rapid and the medium becomes solidly frozen, Stentor perishes. Efimoff (1924) observed that Paramecium multiplied once in about 13 days at 0°C., withstood freezing at −1°C. for 30 minutes but died when kept for 50–60 minutes at the same temperature. He further stated that Paramecium caudatum, Colpidium colpoda, and Spirostomum ambiguum, perished in less than 30 minutes, when exposed below −4°C., and that quick and short cooling (not lower than −9°C.) produced no injury, but if it is prolonged, Paramecium became spherical and swollen to 4–5 times normal size, while Colpidium and Spirostomum shrunk. Wolfson (1935) studied Paramecium sp. in gradually descending subzero-temperature, and observed that as the temperature decreases the organism often swims backward, its bodily movements cease at −14.2°C., but the cilia continue to beat for some time. While Paramecium recover completely from a momentary exposure to −16°C., long cooling at this temperature brings about degeneration. When the water in which the organisms are kept freezes, no survival was noted. Plasmodium knowlesi and P. inui in the blood of Macacus rhesus remain viable, according to Coggeshall (1939), for as long as 70 days at −76°C., if frozen and thawed rapidly. Low temperature on Protozoa (Luyet and Gehenio, 1940).

Light. In the Phytomastigina which include chromatophore-bear-
ing flagellates, the sun light is essential to photosynthesis (p. 107). The sun light further plays an important rôle in those protozoans which are dependent upon chromatophore-possessing organisms as chief source of food supply. Hence the light is another factor concerned with the distribution of free-living Protozoa.

**Chemical composition of water.** The chemical nature of the water is another important factor which influences the very existence of Protozoa in a given body of water. Protozoa differ from one another in morphological as well as physiological characteristics. Individual protozoan species requires a certain chemical composition of the water in which it can be cultivated under experimental conditions, although this may be more or less variable among different forms (Needham *et al.*, 1937).

In their "biological analysis of water" Kolkwitz and Marsson (1908, 1909) distinguished four types of habitats for many aquatic plant, and a few animal, organisms, which were based upon the kind and amount of inorganic and organic matter and amount of oxygen present in the water: namely, katharobic, oligosaprobic, mesosaprobic, and polysaprobic. **Katharobic** protozoans are those which live in mountain springs, brooks, or ponds, the water of which is rich in oxygen, but free from organic matter. **Oligosaprobic** forms are those that inhabit waters which are rich in mineral matter, but in which no purification processes are taking place. Many Phytomastigina, various testaceans and many ciliates, such as Frontonia, Laerymaria, Oxytricha, Stylonychia, Vorticella, etc. inhabit such waters. **Mesosaprobic** protozoans live in waters in which active oxidation and decomposition of organic matter are taking place. The majority of freshwater protozoans belong to this group: namely, numerous Phytomastigina, Heliozoa, Zoomastigina, and all orders of Ciliata. Finally **polysaprobic** forms are capable of living in waters which, because of dominance of reduction and cleavage processes of organic matter, contain at most a very small amount of oxygen and are rich in carbonic acid gas and nitrogenous decomposition products. The black bottom slime contains usually an abundance of ferrous sulphide and other sulphurous substances. Lauterborn (1901) called this *sapropelic*. Examples of polysaprobic protozoans are *Pelomyxa palustris*, *Euglypha alveolata*, *Pamphaqus armatus*, *Mastigamoeba*, *Trepomonas agilis*, *Hexamita inflata*, *Rhynchosomonas nasuta*, *Heteronema acus*, Bodo, Cercomonas, Daetylochlamys, Ctenostomata, etc. The so-called "sewage organisms" abound in such habitat (Lackey, 1925).

Certain free-living Protozoa which inhabit waters rich in decom-
Protozoologists have observed that organic matter is frequently found in the faecal matter of various animals. Their cysts either pass through the alimentary canal of the animal unharmed or are introduced after the faeces are voided, and undergo development and multiplication in the faecal infusion. Such forms are collectively called coprozoic Protozoa. The coprozoic protozoans grow easily in suspension of old faecal matter which is rich in decomposed organic matter and thus show a strikingly strong capacity of adapting themselves to conditions different from those of the water in which they normally live. Some of the Protozoa which have been referred to as coprozoic and which are mentioned in the present work are, as follows: Scytononas pusilla, Rhynchomonas nasuta, Cercomonas longicauda, C. crassicauda, Trepononas agilis, Naegleria gruberi, Acanthamoeba hyalina, Chlamydothrys stercorea and Tillina magna.

As a rule, the presence of sodium chloride in the sea water prevents the occurrence of numerous species of fresh-water inhabitants. Certain species, however, have been known to live in both fresh and brackish or salt water. Among the species mentioned in the present work, the following species have been reported to occur in both fresh and salt waters: Mastigophora: Amphilidium lacustre, Ceratium hirundinella; Sarcodina: Lieberkühnia wagneri; Ciliata: Mesodinium pulex, Prorodon discolor, Lacrymaria olor, Amphileptus claparedei, Lionotus fasciola, Nassula aurea, Trochilioides recta, Chilodonella cucullulcus, Trimyema compressum, Paramecium calkinsi, Colpidium campylum, Platynematum sociale, Cinetochilum margaritaceum, Pleuronema coronatum, Caenomorpha medusula, Spirostomum minus, S. teres, Climacostomum virens, and Thuricola folliculata; Suctoria: Metacineta mystacina, Endosphaera engelmanni.

It seems probable that many other protozoans are able to live in both fresh and salt water, judging from the observations such as that made by Finley (1930) who subjected some fifty species of freshwater Protozoa of Wisconsin to various concentrations of sea water, either by direct transfer or by gradual addition of the sea water. He found that Bodo uncinnatus, Uronema marinum, Pleuronema jaculans and Colpoda aspera are able to live and reproduce even when directly transferred to sea water, that Amoeba verrucosa, Euglena, Phacus, Monas, Cyclidium, Euplotes, Lionotus, Paramecium, Stylonychia, etc., tolerate only a low salinity when directly transferred, but, if the salinity is gradually increased, they live in 100 per cent sea water, and that Arcella, Cyphoderia, Aspidiscus, Blepharisma, Colpoda cucullus, Halteria, etc. could not tolerate 10 per cent sea water even when the change was gradual. Finley noted no
morphological changes in the experimental protozoans which might be attributed to the presence of the salt in the water, except *Amoeba verrucosa*, in which certain structural and physiological changes were observed as follows: as the salinity increased, the pulsation of the contractile vacuole became slower. The body activity continued up to 44 per cent sea water and the vacuole pulsed only once in 40 minutes, and after systole, it did not reappear for 10-15 minutes. The organism became less active above this concentration and in 84 per cent sea water the vacuole disappeared, but there was still a tendency to form the characteristic ridges, even in 91 per cent sea water, in which the organism was less fan-shaped and the cytoplasm seemed to be more viscous. Yocom (1934) found that *Euplotes patella* was able to live normally and multiply up to 66 per cent of sea water; above that concentration no division was noticed, though the organism lived for a few days in up to 100 per cent salt water, and *Paramaecium caudatum* and *Spirostomum ambiguum* were less adaptive to salt water, rarely living in 60 per cent sea water. Frisch (1939) found that no freshwater Protozoa lived above 40 per cent sea water and that *Paramaecium caudatum* and *P. multimicronucleatum* died in 33-52 per cent sea water. Hardin (1942) reports that *Oikomonas termo* will grow when transferred directly to a glycerol-peptone culture medium, in up to 45 per cent sea water, and cultures contaminated with bacteria and growing in a dilute glycerol-peptone medium will grow in 100 per cent sea water.

**Hydrogen-ion concentration.** Closely related to the chemical composition is the hydrogen-ion concentration (pH) of the water. Some Protozoa appear to tolerate a wide range of pH. The interesting protoctymbxan, *Leptomyxa reticulata*, occurs in soil ranging in pH 4.3 to 7.8, and grows very well in non-nutrient agar between pH 4.2 and 8.7, provided a suitable bacterial strain is supplied as food (Singh, 1948); and according to Loefer and Guido (1950), a strain of *Euglena gracilis* (var. bacillaris) grows between pH 3.2 and 8.3. However, the majority of Protozoa seem to prefer a certain range of pH for the maximum metabolic activity.

The hydrogen-ion concentration of freshwater bodies varies a great deal between highly acid bog waters in which various testaceans may frequently be present, to highly alkaline water in which such forms as Acanthocystis, Hyalobryon, etc., occur. In standing deep fresh water, the bottom region is often acid because of the decomposing organic matter, while the surface water is less acid or slightly alkaline due to the photosynthesis of green plants which utilize carbon dioxide. In some cases different pH may bring about morpho-
logical differences. For example, in bacteria-free cultures of *Paramecium bursaria* in a tryptone medium, Loefer (1938) found that at pH 7.6–8.0 the length averaged 86 or 87μ, but at 6.0–6.3 the length was about 129μ. The greatest variation took place at pH 4.6 in which no growth occurred. The shortest animals at the acid and alkaline extremes of growth were the widest, while the narrowest forms (about 44μ wide) were found in culture at pH 5.7–7.4. Many workers have made observations on the pH range of the water or medium in which certain protozoans live, grow, and multiply, some of which data are collected in Table 1.

**Table 1.—Protozoa and hydrogen-ion concentration**

<table>
<thead>
<tr>
<th>Protozoa</th>
<th>pH range of medium in which growth occurs</th>
<th>Optimum range</th>
<th>Observers</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>A. In bacteria-free cultures</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Euglena gracilis</em></td>
<td>3.5–9.0</td>
<td>6.7</td>
<td>Dusi</td>
</tr>
<tr>
<td></td>
<td>3.0–7.7</td>
<td>3.9–9.9</td>
<td>Alexander</td>
</tr>
<tr>
<td></td>
<td>—</td>
<td>5.0–6.5</td>
<td>Jahn</td>
</tr>
<tr>
<td><em>E. deses</em></td>
<td>6.5–8.0</td>
<td>5.3–8.0</td>
<td>Loefer</td>
</tr>
<tr>
<td><em>E. pisciformis</em></td>
<td>6.0–8.0</td>
<td>6.5–7.5</td>
<td>Dusi</td>
</tr>
<tr>
<td></td>
<td>5.4–7.5</td>
<td>6.8</td>
<td>Hall</td>
</tr>
<tr>
<td><em>E. viridis</em></td>
<td>—</td>
<td>5.0</td>
<td>Schoenborn</td>
</tr>
<tr>
<td><em>Chilomonas paramecium</em></td>
<td>4.8–8.0</td>
<td>4.9;7.0</td>
<td>Mast and Pace</td>
</tr>
<tr>
<td><em>Chlorogonium euthilorum</em></td>
<td>4.8–8.7</td>
<td>7.1–7.5</td>
<td>&quot;</td>
</tr>
<tr>
<td><em>C. elongatum</em></td>
<td>4.8–8.7</td>
<td>7.1–7.5</td>
<td>&quot;</td>
</tr>
<tr>
<td><em>C. teragamum</em></td>
<td>4.2–8.6</td>
<td>6.7–8.3</td>
<td>&quot;</td>
</tr>
<tr>
<td><em>Colpidium campylum</em></td>
<td>—</td>
<td>5.4</td>
<td>Kidder</td>
</tr>
<tr>
<td><em>Glaucoma scintillans</em></td>
<td>—</td>
<td>5.6–6.8</td>
<td>&quot;</td>
</tr>
<tr>
<td><em>G. ficara</em></td>
<td>4.0–9.5</td>
<td>5.1;6.7</td>
<td>Johnson</td>
</tr>
<tr>
<td><em>Tetrahymena pyriformis</em></td>
<td>—</td>
<td>5.6–8.0</td>
<td>Kidder</td>
</tr>
<tr>
<td><em>T. vorax</em></td>
<td>—</td>
<td>6.2–7.6</td>
<td>&quot;</td>
</tr>
<tr>
<td><em>Paramecium bursaria</em></td>
<td>4.9–8.0</td>
<td>6.7–6.8</td>
<td>Loefer</td>
</tr>
<tr>
<td><strong>B. In cultures containing bacteria</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Carteria obtusa</em></td>
<td>—</td>
<td>3.5–4.5</td>
<td>Wermel</td>
</tr>
<tr>
<td><em>Trichomonas vaginalis</em></td>
<td>6.4–8.4</td>
<td>—</td>
<td>Bland <em>et al.</em></td>
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<tr>
<td><em>Actinosphaerium eichhorni</em></td>
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<td><em>Acanthocyctis aculeata</em></td>
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<td></td>
<td>6.0–9.5</td>
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<td>Morea</td>
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<tr>
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<td>5.7–7.8</td>
<td>6.7</td>
<td>Morea</td>
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<td>Protozoa</td>
<td>pH range of medium in which growth occurs</td>
<td>Optimum range</td>
<td>Observers</td>
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<td><em>P. polycaryum</em></td>
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<td><em>P. calkinsi</em></td>
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<td><em>P. woodrufi</em></td>
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<td>7.5</td>
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<td>S. sp.</td>
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<td>Moore</td>
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<td>Pruthi</td>
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<td>Gastrostyla sp.</td>
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<td>6.7; 8.0</td>
<td>Darby</td>
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**Food.** The kind and amount of food available in a given body of water also controls the distribution of Protozoa. The food is ordinarily one of the deciding factors of the number of Protozoa in a natural habitat. Species of *Paramaecium* and many other holozoic protozoans cannot live in waters in which bacteria or minute protozoans do not occur. If other conditions are favorable, then the greater the number of food bacteria, the greater the number of protozoa. Noland (1925) studied more than 65 species of fresh-water ciliates with respect to various factors and came to the conclusion that the nature and amount of available food has more to do with the distribution of these organisms than any other one factor. *Didinium nasutum* feeds almost exclusively on paramecia; therefore, it cannot live in the absence of the latter ciliate. As a rule, euryphagous Protozoa which feed on a variety of food organisms are widely distributed, while stenophagous forms that feed on a few species of food organisms are limited in their distribution.

In nature, Protozoa live in association with diverse organisms. The interrelationships which exist among them are not understood in most cases. For example, the relationship between *Entamoeba histolytica* and certain bacteria in successful in-vitro cultivation has
not yet been comprehended. Certain strains of bacteria were found by Hardin (1944) to be toxic for *Paramecium multimicronucleatum*, but if *Oikomonas termo* was present in the culture, the ciliate was maintained indefinitely. This worker suggested that the flagellate may be able to "detoxify" the metabolic products produced by the bacteria. Food relation in ciliates (Fauré-Fremiet, 1950, 1951a).

The adaptability of Protozoa to varied environmental conditions influences their distribution. The degree of adaptability varies a great deal, not only among different species, but also among the individuals of the same species. *Stentor coeruleus* which grows ordinarily under nearly anaerobic conditions, is obviously not influenced by alkalinity, pH, temperature or free carbon dioxide in the water (Sprugel, 1951).

Some protozoans inhabit soil of various types and localities. Under ordinary circumstances, they occur near the surface, their maximum abundance being found at a depth of about 10–12 cm. (Sandon, 1927). It is said that a very few protozoans occur in the subsoil. Here also one notices a very wide geographical distribution of apparently one and the same species. For example, Sandon found *Amoeba proteus* in samples of soil collected from Greenland, Tristan da Cunha, Gough Island, England, Mauritius, Africa, India, and Argentina. This amoeba is known to occur in various parts of North America, Europe, Japan, and Australia. The majority of Testacea inhabit moist soil in abundance. Sandon observed *Trinema enchelys* in the soils of Spitzbergen, Greenland, England, Japan, Australia, St. Helena, Barbados, Mauritius, Africa, and Argentina.

**Parasitic Protozoa**

Some Protozoa belonging to all groups live on or in other organisms. The Sporozoa are made up exclusively of parasites. The relationships between the host and the protozoan differ in various ways, which make the basis for distinguishing the associations into three types as follows: commensalism, symbiosis, and parasitism.

**Commensalism** is an association in which an organism, the commensal, is benefited, while the host is neither injured nor benefited. Depending upon the location of the commensal in the host body, the term ectocommensalism or endocommensalism is used. Ectocommensalism is often represented by Protozoa which may attach themselves to any aquatic animals that inhabit the same body of water, as shown by various species of Chonotricha, Peritricha, and Suctoria. In other cases, there is a definite relationship between the commensal and the host. For example, *Kerona polyporum* is found
on various species of Hydra, and many ciliates placed in Thigmo-
tricha (p. 774) are inseparably associated with certain species of
mussels.

Endocommensalism is often difficult to distinguish from endo-
parasitism, since the effect of the presence of a commensal upon the
host cannot be easily understood. On the whole, the protozoans
which live in the lumen of the alimentary canal may be looked upon
as endocommensals. These protozoans undoubtedly use part of the
food material which could be used by the host, but they do not in-
vade the host tissue. As examples of endocommensals may be men-
tioned: Endamoeba blattae, Lophomonas blattarum, L. striata,
Nyctotherus ovalis, etc., of the cockroach; Entamoeba coli, Iodamoeba
butschlii, Endolimax nana, Dientamoeba fragilis, Chilomastix mes-
nili, etc., of the human intestine; numerous species of Protoctiliata
of Anura, etc. Because of the difficulties mentioned above, the term
parasitic Protozoa, in its broad sense, includes the commensals also.

Symbiosis on the other hand is an association of two species of
organisms, which is of mutual benefit. The cryptomonads belonging
to Chrysidella ("Zooxanthellae") containing yellow or brown chromo-
atophores, which live in Foraminifera and Radiolaria, and certain
algae belonging to Chlorella ("Zoochlorella") containing green
chromatophores, which occur in some freshwater protozoans, such as
Paramecium bursaria, Stentor amethystinus, etc., are looked upon
as holding symbiotic relationship with the respective protozoan host.
Several species of the highly interesting Hypermastigina, which are
present commonly and abundantly in various species of termites and
the woodroach Cryptocercus, have been demonstrated by Cleveland
to digest the cellulose material which makes up the bulk of wood-
chips the host insects take in and to transform it into glycogenous
substances that are used partly by the host insects. If deprived of
these flagellates by being subjected to oxygen under pressure or to
a high temperature, the termites die, even though the intestine is
filled with wood-chips. If removed from the gut of the termite, the
flagellates perish (Cleveland, 1924, 1925). Recently, Cleveland
(1949–1950c) found that the molting hormone produced by Crypto-
cercus induces sexual reproduction in several flagellates inhabiting
its hind-gut (p. 185). Thus the association here may be said to be an
absolute symbiosis.

Parasitism is an association in which one organism (the parasite)
lives at the expense of the other (the host). Here also ectoparasitism
and endoparasitism occur, although the former is not commonly
found. Hydramoeba hydroxena (p. 464) feeds on the body cells of
Hydra which, according to Reynolds and Looper (1928), die on an average in 6.8 days as a result of the infection and the amoebae disappear in from 4 to 10 days if removed from a host Hydra. *Costia necatrix* (p. 372) often occurs in an enormous number, attached to various freshwater fishes especially in an aquarium, by piercing through the epidermal cells and appears to disturb the normal functions of the host tissue. *Ichthyophthirius multifiliis* (p. 709), another ectoparasite of freshwater and marine fishes, goes further by completely burying themselves in the epidermis and feeds on the host’s tissue cells and, not infrequently, contributes toward the cause of the death of the host fishes.

The endoparasites absorb by osmosis the vital body fluid, feed on the host cells or cell-fragments by pseudopodia or cytostome, or enter the host tissues or cells themselves, living on the cytoplasm or in some cases on the nucleus. Consequently they bring about abnormal or pathological conditions upon the host which often succumbs to the infection. Endoparasitic Protozoa of man are *Entamoeba histolytica*, *Balantidium coli*, species of *Plasmodium* and *Leishmania*, *Trypanosoma gambiense*, etc. The Sporozoa, as was stated before, are without exception coelozoic, histozoic, or cytozoic parasites.

Because of their modes of living, the endoparasitic Protozoa cause certain morphological changes in the cells, tissues, or organs of the host. The active growth of *Entamoeba histolytica* in the glands of the colon of the victim, produces first slightly raised nodules which develop into abscesses and the ulcers formed by the rupture of abscesses, may reach 2 cm. or more in diameter, completely destroying the tissues of the colon wall. Similar pathological changes may also occur in the case of infection by *Balantidium coli*. In *Leishmania donovani*, the victim shows an increase in number of the large macrophages and mononuclears and also an extreme enlargement of the spleen. *Trypanosoma cruzi* brings about the degeneration of the infected host cells and an abundance of leucocytes in the infected tissues, followed by an increase of fibrous tissue. *T. gambiense*, the causative organism of African sleeping sickness, causes enlargement of lymphatic glands and spleen, followed by changes in meninges and an increase of cerebro-spinal fluid. Its most characteristic changes are the thickening of the arterial coat and the round-celled infiltration around the blood vessels of the central nervous system.

Malarial infection is invariably accompanied by an enormous enlargement of the spleen ("spleen index"); the blood becomes watery; the erythrocytes decrease in number; the leucocytes, subnormal; but mononuclear cells increase in number; pigment granules
which are set free in the blood plasma at the time of merozoite-liberation are engulfed by leucocytes; and enlarged spleen contains large amount of pigments which are lodged in leucocytes and endothelial cells. In *Plasmodium falciparum*, the blood capillaries of brain, spleen and other viscera may completely be blocked by infected erythrocytes.

In Myxosporidia which are either histozoic or coelozoic parasites of fishes, the tissue cells that are in direct contact with highly enlarging parasites, undergo various morphological changes. For exam-

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**Fig. 1.** Histological changes in host fish caused by myxosporidian infection, ×1920 (Kudo). a, portion of a cyst of *Myxobolus intestinalis*, surrounded by peri-intestinal muscle of the black crappie; b, part of a cyst of *Thelohanellus notatus*, enveloped by the connective tissue of the blunt-nosed minnow.

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...ple, the circular muscle fibers of the small intestine of *Pomoxis sparooides*, which surround *Myxobolus intestinalis*, a myxosporidian, become modified a great deal and turn about 90° from the original direction, due undoubtedly to the stimulation exercised by the myxosporidian parasite (Fig. 1, a). In the case of another myxosporidian, *Thelohanellus notatus*, the connective tissue cells of the host fish surrounding the protozoan body, transform themselves into "epithelial cells" (Fig. 1, b), a state comparable to the formation of the ciliated epithelium from a layer of fibroblasts lining a cyst formed around a piece of ovary implanted into the adductor muscle of Pecten as observed by Drew (1911).
Practically all Microsporidia are cytozoic, and the infected cells become hypertrophied enormously, producing in one genus the so-called Glugea cysts (Figs. 287, 290). In many cases, the hypertrophy of the nucleus of the infected cell is far more conspicuous than that of the cytoplasm (Figs. 287, 291) (Kudo, 1924).

When the gonads are parasitized heavily, the germ cells of the host animal often do not develop, thus resulting in parasitic castration. For example, the ciliate, *Orchitophrya stellatum*, a parasite in the male reproductive organ of *Asterias rubens*, was found by Vevers (1951) to break down completely all germinal tissues of the testes in the majority of the host starfish. In other cases, the protozoan does not invade the gonads, but there is no development of the germ cells. The microsporidian, *Nosema apis*, attacks solely the gut epithelium of the honey bee, but the ovary of an infected queen bee degenerates to varying degrees (Hassanein, 1951). Still in other instances, the Protozoa invade developing ova of the host, but do not hinder their development, though the parasites multiply, as in *Nosema bombycis* in the silkworm (Stempell, 1909) and *Babesia bigemina* in the cattle tick (Dennis, 1932).

For the great majority of parasitic Protozoa, there exists a definite host-parasite relationship and animals other than the specific hosts possess a natural immunity against an infection by a particular parasitic protozoan. Immunity involved in diseases caused by Protozoa has been most intensively studied on haemozoic forms, especially *Plasmodium* and *Trypanosoma*, since they are the causative organisms of important diseases. Development of these organisms in hosts depends on various factors such as the species and strains of the parasites, the species and strains of vectors, and immunity of the host. Boyd and co-workers showed that reinoculation of persons who have recovered from an infection with *Plasmodium vivax* or *P. falciparum* with the same strain of the parasites, will not result in a second clinical attack, because of the development of homologous immunity, but with a different strain of the same species or different species, a definite clinical attack occurs, thus there being no heterologous tolerance. The homologous immunity was found to continue for at least three years and in one case for about seven years in *P. vivax*, and for at least four months in *P. falciparum* after apparent eradication of the infection. In the case of leishmaniasis, recovery from a natural or induced infection apparently develops a lasting immunity against reinfection with the same species of Leishmania.

It has been shown that in infections with avian, monkey and human *Plasmodium* or *Trypanosoma lewisi*, a considerable number of
the parasites are destroyed during the developmental phase of the infection and that after a variable length of time, resistance to the parasites often develops in the host, as the parasites disappear from the peripheral blood and symptoms subside, though the host still harbors the organisms. In malarious countries, the adults and children show usually a low and a high rate of malaria infection respectively, but the latter frequently do not show symptoms of infection, even though the parasites are detectable in the blood. Apparently repeated infection produces tolerance which can keep, as long as the host remains healthy, the parasites under control. There seems to be also racial difference in the degree of immunity against Plasmodium and Trypanosoma.

As to the mechanism of immunity, the destruction of the parasites by phagocytosis of the endothelial cells of the spleen, bone marrow and liver and continued regenerative process to replace the destroyed blood cells, are the two important phases in the cellular defense mechanism. Besides, there are indications that humoral defense mechanism through the production of antibodies is in active operation in infections by Plasmodium knowlesi and trypanosomes (Taliaferro, 1926; Maegraith, 1948; Culbertson, 1951). Immunity (Taliaferro, 1941).

With regard to the origin of parasitic Protozoa, it is generally agreed among biologists that the parasite in general evolved from the free-living form. The protozoan association with other organisms was begun when various protozoans which lived attached to, or by crawling on, submerged objects happened to transfer themselves to various invertebrates which occur in the same water. These Protozoa benefit by change in location as the host animal moves about, and thus enlarging the opportunity to obtain a continued supply of food material. Such ectocommensals are found abundantly; for example, the peritrichous ciliates attached to the body and appendages of various aquatic animals such as larval insects and microcrustaceans. Ectocommensalism may next lead to ectoparasitism as in the case of Costia or Hydramoeba, and then again instead of confining themselves to the body surface, the Protozoa may bore into the body wall from outside and actually acquire the habit of feeding on tissue cells of the attached animals as in the case of Ichthyophthirius.

The next step in the evolution of parasitism must have been reached when Protozoa, accidentally or passively, were taken into the digestive system of the Metazoa. Such a sudden change in habitat appears to be fatal to most protozoans. But certain others
possess extraordinary capacity to adapt themselves to an entirely different environment. For example, Dobell (1918) observed in the tadpole gut, a typical free-living limax amoeba, with characteristic nucleus, contractile vacuoles, etc., which was found in numbers in the water containing the faecal matter of the tadpole. Glaucoma (Tetrahymena) pyriformis, a free-living ciliate, was found to occur in the body cavity of the larvae of Theobaldia annulata (after MacArthur) and in the larvae of Chironomus plumosus (after Treillard and Lwoff). Lwoff successfully inoculated this ciliate into the larvae of Galleria mellonella which died later from the infection. Janda and Jirovec (1937) injected bacteria-free culture of this ciliate into annelids, molluscs, crustaceans, insects, fishes, and amphibians, and found that only insects—all of 14 species (both larvae and adults)—became infected by this ciliate. In a few days after injection the haemocoele became filled with the ciliates. Of various organs, the ciliates were most abundantly found in the adipose tissue. The organisms were much larger than those present in the original culture. The insects, into which the ciliates were injected, died from the infection in a few days. The course of development of the ciliate within an experimental insect depended not only on the amount of the culture injected, but also on the temperature. At 1–4°C the development was much slower than at 26°C.; but if an infected insect was kept at 32–36°C. for 0.5–3 hours, the ciliates were apparently killed and the insect continued to live. When Glaucoma taken from Dixippus morosus were placed in ordinary water, they continued to live and underwent multiplication. The ciliate showed a remarkable power of withstanding the artificial digestion; namely, at 18°C. they lived 4 days in artificial gastric juice with pH 4.2; 2–3 days in a juice with pH 3.6; and a few hours in a juice with pH 1.0. Cleveland (1928) observed Tritrichomonas fecalis in faeces of a single human subject for three years which grew well in faeces diluted with tap water, in hay infusions with or without free-living protozoans or in tap water with tissues at −3° to 37°C., and which, when fed per os, was able to live indefinitely in the gut of frogs and tadpoles. Reynolds (1936) found that Colpoda steini, a free-living ciliate of fresh water, occurs naturally in the intestine and other viscera of the land slug, Agriolimax agrestis, the slug forms being much larger than the free-living individuals.

It may be further speculated that Vahlkampfia, Hydramoeba, Schizamoeba, and Endamoeba, are the different stages of the course the intestinal amoebae might have taken during their evolution. Obviously endocommensalism in the alimentary canal was the initial phase of endoparasitism. When these endocommensals began
to consume an excessive amount of food or to feed on the tissue cells of the host gut, they became the true endoparasites. Destroying or penetrating through the intestinal wall, they became first established in the body or organ cavities and then invaded tissues, cells or even nuclei, thus developing into pathogenic Protozoa. The endoparasites developing in invertebrates which feed upon the blood of vertebrates as source of food supply, will have opportunities to establish themselves in the higher animals.

**Hyperparasitism.** Certain parasitic Protozoa have been found to parasitize other protozoan or metazoan parasites. This association is named hyperparasitism. The microsporidian *Nosema notabilis* (p. 672) is an exclusive parasite of the myxosporidian *Sphaerospora polymorpha*, which is a very common inhabitant of the urinary bladder of the toad fish along the Atlantic and Gulf coasts. A heavy infection of the microsporidian results in the degeneration and death of the host myxosporidian trophozoite (Kudo, 1944). Thus *Nosema notabilis* is a hyperparasite. Organisms living on and in Protozoa (Duboscq and Grassé, 1927, 1929; Georgévitch, 1936; Grassé, 1936; Kirby, 1932, 1938, 1941, 1941a, 1942, 1942a, 1942b, 1944, 1946)

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Chapter 3
Morphology

Protozoa range in size from submicroscopic to macroscopic, though they are on the whole minute microscopic animals. The parasitic forms, especially cytotic parasites, are often extremely small, while free-living protozoans are usually of much larger dimensions. Noctiluca, Foraminiferida, Radiolaria, many ciliates such as Stentor, Bursaria, etc., represent larger forms. Colonial protozoans such as Carchesium, Zoothamnium, Ophrydium, etc., are even greater than the solitary forms. On the other hand, Plasmodium, Leishmania, and microsporidian spores may be mentioned as examples of the smallest forms. The unit of measurement employed in protozoology is, as in general microscopy, 1 micron (\( \mu \)) which is equal to 0.001 mm.

The body form of Protozoa is even more varied, and because of its extreme plasticity it frequently does not remain constant. Furthermore the form and size of a given species may vary according to the kind and amount of food as is discussed elsewhere (p. 109). From a small simple spheroidal mass up to large highly complex forms, all possible body forms occur. Although the great majority are without symmetry, there are some which possess a definite symmetry. Thus bilateral symmetry is noted in all members of Diplomonadina (p. 392); radial symmetry in Gonim, Cyclonexis, etc.; and universal symmetry, in certain Heliozoa, Volvox, etc.

The fundamental component of the protozoan body is the protoplasm which is without exception differentiated into the nucleus and the cytoplasm. Haeckel's (1868, 1870) monera are now considered as nonexistent, since improved microscopic technique has failed in recent years to reveal any anucleated protozoans. The nucleus and the cytoplasm are inseparably important to the well-being of a protozoan, as has been shown by numerous investigators since Verworn's pioneer work. In all cases, successful regeneration of the body is accomplished only by the nucleus-bearing portions and nucleate parts degenerate sooner or later. On the other hand, when the nucleus is taken out of a protozoan, both the nucleus and cytoplasm degenerate, which indicates their intimate association in carrying on the activities of the body. It appears certain that the nucleus controls the assimilative phase of metabolism which takes place in the cytoplasm in normal animals, while the cytoplasm is capable of carrying on the catabolic phase of the metabolism. Aside from the importance
as the controlling center of metabolism, evidences point to the conclusion that the nucleus contains the genes or hereditary factors which characterize each species of Protozoa from generation to generation, as in the cells of multicellular animals and plants.

The nucleus

Because of a great variety of the body form and organization, the protozoan nuclei are of various forms, sizes and structures. At one extreme there is a small nucleus and, at the other, a large voluminous one and, between these extremes, is found almost every conceivable variety of form and structure. The majority of Protozoa contain a single nucleus, though many may possess two or more throughout the greater part of their life-cycle. In several species, each individual possesses two similar nuclei, as in Diplomonadina, Protoopalina and Zelleriella. In Euciliata and Suctoria, two dissimilar nuclei, a macronucleus and a micronucleus, are typically present. The macronucleus is always larger than the micronucleus, and controls the trophic activities of the organism, while the micronucleus is concerned with the reproductive activity. Certain Protozoa possess numerous nuclei of similar structure, as for example, in Pelomyxa, Mycetozoa, Actinosphaerium, Opalina, Cepedea, Myxosporidia, Microsporidia, etc.

The essential morphological components of the protozoan nucleus are the nuclear membrane, chromatin, plastin and nucleoplasm or nuclear sap. Their interrelationship varies sometimes from one developmental stage to another, and vastly among different species. Structurally, they fall in general into one of the two types: vesicular and compact.

The vesicular nucleus (Fig. 2, a, c, e) consists of a nuclear membrane which is sometimes very delicate but distinct, nucleoplasm, achromatin and chromatin. Besides there is an intranuclear body which is, as a rule, more or less spherical and which appears to be of different make-ups as judged by its staining reactions among different nuclei. It may be composed of chromatin, of plastin, or of a mixture of both. The first type is sometimes called karyosome and the second, nucleolus or plasmosome. Absolute distinction between these two terms cannot be made as they are based solely upon the difference in affinity to nuclear stains which cannot be standardized and hence do not give uniformly the same result. Following Minchin (1912), the term endosome is advocated here to designate one or more conspicuous bodies other than the chromatin granules, present within the nuclear membrane (Fig. 2, b, d).
Fig. 2. a–f, vesicular nuclei; g–j, compact nuclei, ×980. a, b, nuclei of *Entamoeba invadens* (a, in life; b, in stained organism); c, d, nuclei of *Amoeba spumosa* (c, in life, showing a large endosome; d, stained); e, f, nuclei of *A. proteus* (e, in life; f, a nucleus subjected to Feulgen's nucleal reaction); g, h, nuclei of *Paramecium aurelia* (g, in life under phase microscope, showing two vesicular micronuclei and compact macronucleus; h, Feulgen-stained nuclei); i, j, nuclei of *Frontonia leucas*, showing a micronucleus and macronucleus, both of which are compact (i, in life, showing many endosomes imbedded among the granules; j, nuclei stained with acidified methyl green),
When viewed in life, the nucleoplasm is ordinarily homogeneous and structureless. But, upon fixation, there appear invariably achromatic strands or networks which seem to connect the endosome and the nuclear membrane (Fig. 2, b, d). Some investigators hold that these strands or networks exist naturally in life, but due to the similarity of refractive indices of the strands and of the nucleoplasm, they are not visible and that, when fixed, they become readily recognizable because of a change in these indices. In some nuclei, however, certain strands have been observed in life, as for example in the nucleus of the species of Barbulanymphia (Fig. 174, c), according to Cleveland and his associates (1934). Others maintain that the achromatic structures prominent in fixed vesicular nuclei are mere artifacts brought about by fixation and do not exist in life and that the nucleoplasm is a homogeneous liquid matrix of the nucleus in which the chromatin is usually distributed as small granules. Frequently larger granules of various sizes and forms may occur along the inner surface of the nuclear membrane. These so-called peripheral granules that occur in Amoeba, Entamoeba, Pelomyxa, etc., are apparently not chromatinic (Fig. 2, a, e). The vesicular nucleus is most commonly present in various orders of Sarcomina and Mastigophora.

The compact nucleus (Fig. 2, g–j), on the other hand, contains a large amount of chromatin substance and a comparatively small amount of nucleoplasm, and is thus massive. The macronucleus of the Ciliophora is almost always of this kind. The variety of forms of the compact nuclei is indeed remarkable. It may be spherical, ovate, cylindrical, club-shaped, band-form, moniliform, horseshoe-form, filamentous, or dendritic. The nuclear membrane is always distinct, and the chromatin substance is usually of spheroidal form, varying in size among different species and often even in the same species. In the majority of species, the chromatin granules are small and compact (Fig. 2, h, i), though in some forms, such as Nyctotherus ovalis (Fig. 3), they may reach 20μ or more in diameter in some individuals and while the smaller chromatin granules seem to be homogeneous, larger forms contain alveoli of different sizes in which smaller chromatin granules are suspended (Kudo, 1936).

Precise knowledge of chromatin (thymo- or desoxyribose-nucleic acid) is still lacking. At present the determination of the chromatin depends upon the following tests: (1) artificial digestion which does not destroy this substance, while non-chromatinic parts of the nucleus are completely dissolved; (2) acidified methyl green which stains the chromatin bright green; (3) 10 per cent sodium chloride solution which dissolves, or causes swelling of, chromatin granules,
while nuclear membrane and achromatic substances remain unattacked; and (4) in the fixed condition Feulgen's nucleal reaction (p. 897). Action of methyl green (Pollister and Leuchtenberger, 1949).

There is no sharp demarcation between the vesicular and compact nuclei, since there are numerous nuclei the structures of which are intermediate between the two. Moreover what appears to be a vesicular nucleus in life, may approach a compact nucleus when fixed and stained as in the case of Euglenoidina. Several experimental observations show that the number, size, and structure of the endosome in the vesicular nucleus, and the amount and arrangement of the chromatin in the compact nucleus, vary according to the physiological state of the whole organism. The macronucleus may be

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**Fig. 3.** Parts of four macronuclei of *Nyctotherus ovalis*, showing chromatin spherules of different sizes, ×650 (Kudo).
divided into two or more parts with or without connections among
them and in Dileptus anser into more than 200 small nuclei, each of
which is "composed of a plastin core and a chromatin cortex" (Cal-
kins; Hayes).

In a compact nucleus, the chromatin granules or spherules fill, as
a rule, the intranuclear space compactly, in which one or more endo-
somes (Fig. 2, i) may occur. In many nuclei these chromatin granules
appear to be suspended freely, while in others a reticulum appears to
make the background. The chromatin of compact nuclei gives a
strong positive Feulgen's nuclear reaction. The macronuclear and
micronuclear chromatin substances respond differently to Feulgen's
nuclear reaction or to the so-called nuclear stains, as judged by the
difference in the intensity or tone of color. In Paramecium caudatum,
P. aurelia, Chilodonella, Nyctotherus ovalis, etc., the macronuclear
chromatin is colored more deeply than the micronuclear chromatin,
while in Colpoda, Urostyla, Euplotes, Stylonychia, and others, the
reverse seems to be the case, which may support the validity of the
assumption by Heidenhain that the two types of the nuclei of
Euciliata and Suctoria are made up of different chromatin sub-
stances—idiochromatin in the micronucleus and trophochromatin
in the macronucleus—and in other classes of Protozoa, the two kinds
of chromatin are present together in a single nucleus. The macro-
nucleus and the micronucleus of vegetative Paramecium caudatum
were found by Moses (1950) to possess a similar nucleic acid-protein
composition; namely, similar concentrations of total protein, non-
histone protein, desoxyribose nucleic acid and ribose nucleic acid.
Of the two latter nucleic acids, ribose nucleic acid is said to be pres-
ent in a larger amount than desoxyribose nucleic acid in both nuclei.
It may be considered that the two nucleic acids occur in different
proportions in the two nuclei.

Chromidia. Since the detection of chromatin had solely depended
on its affinity to certain nuclear stains, several investigators found
extranuclear chromatin granules in many protozoans. Finding such
granules in the cytoplasm of Actinosphaerium eichhorni, Arcella vul-
garis, and others, Hertwig (1902) called them chromidia, and main-
tained that under certain circumstances, such as lack of food ma-
terial, the nuclei disappear and the chromatin granules become scat-
tered throughout the cytoplasm. In the case of Arcella vulgaris, the
two nuclei break down completely to produce a chromidial-net
which later reforms into smaller secondary nuclei. It has, however,
been found by Bélař that the lack of food caused the encystment
rather than chromidia-formation in Actinosphaerium and, according
to Reichenow, Jollos observed that in Arcella the nuclei persisted, but were thickly covered by chromidial-net which could be cleared away by artificial digestion to reveal the two nuclei. In Diffugia, the chromidial-net is vacuolated or alveolated in the fall and in each alveolus appear glycogen granules which seem to serve as reserve food material for the reproduction that takes place during that season (Zuelzer), and the chromidia occurring in Actinosphaerium appear to be of a combination of a carbohydrate and a protein (Rumjantzew and Wermel, 1925). Apparently the widely distributed volutin (p. 114), and many inclusions or cytozoic parasites, such as Sphaerita volutin (p. 893), which occur occasionally in different Sarcodina, have in some cases been called chromidia. By using Feulgen’s nucleal reaction, Reichenow (1928) obtained a diffused violet-stained zone in Chlamydomonas and held them to be dissolved volutin. Calkins (1933) found the chromidia of Arcella vulgaris negative to the nucleal reaction, but by omitting acid-hydrolysis and treating with fuchsinsulphurous acid for 8–14 hours, the chromidia and the secondary nuclei were found to show a typical positive reaction and believed that the chromidia were chromatin. Thus at present the real nature of chromidia is still not clearly known, although many protozoologists are inclined to think that the substance is not chromatinic, but, in some way, is connected with the metabolism of the protozoan.

The cytoplasm

The extranuclear part of the protozoan body is the cytoplasm. It is composed of a colloidal system, which may be homogeneous, granulated, vacuolated, reticulated, or fibrillar in optical texture, and is almost always colorless. The chromatophore-bearing Protozoa are variously colored, and those with symbiotic algae or cryptomonads are also greenish or brownish in color. Furthermore, pigment or crystals which are produced in the body may give protozoans various colorations. In several forms pigments are diffused throughout the cytoplasm. For example, many dinoflagellates are beautifully colored, which, according to Kofoid and Swezy, is due to a thorough diffusion of pigment in the cytoplasm.

Stentor coeruleus is beautifully blue-colored. This coloration is due to the presence of pigment stentorin (Lankester, 1873) which occurs as granules in the ectoplasm (Fig. 14). The pigment is highly resistant to various solvents such as acids and alkalis, and the sunlight does not affect its nature. It is destroyed by bleaching with chlorine gas or with potassium permanganate, followed by immersion in 5 per cent oxalic acid (Weisz, 1948). Several species of Blepha-
risma are rose- or purple-colored. The color is due to the presence of 
zoopurpurin (Arciehovskij, 1905) which is lodged in numerous gran-
ules present in the ectoplasm. This pigment is soluble in alcohol, 
ether or acetone, and is destroyed by strong light (Giese, 1938). 
Weisz (1950) maintains that both pigment granules are chondri-
somes, and in Stentor, cytochrome oxidase appears to be localized in 
the pigment granules.

The extent and nature of the cytoplasmic differentiation differ 
greatly among various groups. In the majority of Protozoa, the 
cytoplasm is differentiated into the ectoplasm and the endoplasm. 
The ectoplasm is the cortical zone which is hyaline and homogeneous 
in Sarcodina and Sporozoa. In the Ciliophora it is a permanent and 
distinct part of the body and contains several organelles. The endo-
plasm is more voluminous and fluid. It is granulated or alveolated 
and contains various organelles. While the alveolated cytoplasm is 
normal in forms such as the members of Heliozoa and Radiolaria, in 
other cases the alveolation of normally granulated or vacuolated 
cytoplasm indicates invariably the beginning of degeneration of the 
protozoan body. In Amoeba and other Sarcodina, the "hyaline cap" 
and "layer" (Mast) make up the ectoplasm, and the "plasmasol" 
and "plamagel" (Mast) compose the endoplasm (Fig. 46).

In numerous Sarcodina and certain Mastigophora, the body 
surface is naked and not protected by any form-giving organella. 
However, the surface layer is not only elastic, but solid, and there-
fore the name plasma-membrane may be applied to it. Such forms 
are capable of undergoing amoeboid movement by formation of 
pseudopodia and by continuous change of form due to the movement 
of the cytoplasm which is more fluid. However, the majority of 
Protozoa possess a characteristic and constant body form due to the 
development of a special envelope, the pellicle. In Amoeba striata, 
A. verrucosa (Howland, 1924), Pelomyza carolinensis, P. illinoisensis 
(Kudo, 1946, 1951), etc., there is a distinct pellicle. The same is true 
with some flagellates, such as certain species of Euglena, Peranema, 
and Astasia, in which it is elastic and expansible so that the organ-
isms show a great deal of plasticity.

The pellicle of a ciliate is much thicker and more definite, and 
often variously ridged or sculptured. In many, linear furrows and 
ridges run longitudinally, obliquely, or spirally; and, in others, the 
ridges are combined with hexagonal or rectangular depressed areas. 
Still in others, such as Coleps, elevated platelets are arranged paral-
lel to the longitudinal axis of the body. In certain peritrichious 
ciliates, such as Vorticella monilata, Carchesium granulatum, etc.,
the pellicle may possess nodular thickenings arranged in more or less parallel rows at right angles to the body axis.

While the pellicle always covers the protozoan body closely, there are other kinds of protective envelopes produced by Protozoa which may cover the body rather loosely. These are the shell, test, lorica or envelope. The shell of various Phytomastigina is usually made up of cellulose, a carbohydrate, which is widely distributed in the plant kingdom. It may be composed of a single or several layers, and may possess ridges or markings of various patterns on it. In addition to the shell, gelatinous substance may in many forms be produced to surround the shelled body or in the members of Volvocidae to form the matrix of the entire colony in which the individuals are embedded. In the dinoflagellates, the shell is highly developed and often composed of numerous plates which are variously sculptured.

In other Protozoa, the shell is made up of chitin or pseudo-chitin (te tin). Common examples are found in the testaceans; for example, in Arcella and allied forms, the shell is made up of chitinous material constructed in particular ways which characterize the different genera. Newly formed shell is colorless, but older ones become brownish, because of the presence of iron oxide. Diffugia and related genera form shells by gluing together small sand-grains, diatom-shells, debris, etc., with chitinous or pseudochitinous substances which they secrete. Many foraminiferans seem to possess a remarkable selective power in the use of foreign materials, for the construction of their shells. According to Cushman (1933) Psammosphaera fusca uses sand-grains of uniform color but of different sizes, while P. parva uses grains of more or less uniform size but adds, as a rule, a single large acerose sponge spicule which is built into the test and which extends out both ways considerably. Cushman thinks that this is not accidental, since the specimens without the spicules are few and those with a short or broken spicules are not found. P. bowmanni, on the other hand, uses only mica flakes which are found in a comparatively small amount, and P. rustica uses acerose sponge spicules for the framework of the shell, skilfully fitting smaller broken pieces into polygonal areas. Other foraminiferans combine chitinous secretion with calcium carbonate and produce beautifully constructed shells (Fig. 4) with one or numerous pores. In the Coccolithidae, variously shaped platelets of calcium carbonate ornament the shell.

The silica is present in the shells of various Protozoa. In Euglypha and related testaceans, siliceous scales or platelets are produced in the endoplasm and compose a new shell at the time of fission or of
encystment together with the chitinous secretion. In many heliozoans, siliceous substance forms spicules, platelets, or combination of both which are embedded in the mucilaginous envelope that surrounds the body and, in some cases, a special clathrate shell composed of silica, is to be found. In some Radiolaria, isolated siliceous spicules occur as in Heliozoa, while in others the lateral development of the spines results in production of highly complex and the most beautiful shells with various ornamentations or incorporation of foreign materials. Many pelagic radiolarians possess numerous conspicuous radiating spines in connection with the skeleton, which apparently aid the organisms in maintaining their existence in the open sea.

Certain Protomonadina possess a funnel-like collar in the flagellated end and in some in addition a chitinous lorica surrounds the body. The lorica found in the Ciliophora is mostly composed of chitinous substance alone, especially in Peritricha, although others produce a house made up of gelatinous secretion containing foreign materials as in Stentor (p. 806). In the Tintinnidae, the loricae are either solely chitinous in numerous marine forms not mentioned in the present work or composed of sand-grains or coccoliths cemented together by chitinous secretion, which are found in freshwater forms.
Locomotor organellae

Closely associated with the body surface are the organellae of locomotion: pseudopodia, flagella, and cilia. These organellae are not confined to Protozoa alone and occur in various cells of Metazoa. All protoplasmic masses are capable of movement which may result in change of their forms.

**Pseudopodia.** A pseudopodium is a temporary projection of part of the cytoplasm of those protozoans which do not possess a definite pellicle. Pseudopodia are therefore a characteristic organella of Sarcodina, though many Mastigophora and certain Sporozoa, which lack a pellicle, are also able to produce them. According to their form and structure, four kinds of pseudopodia are distinguished.

1). **Lobopodium** is formed by an extension of the ectoplasm, accompanied by a flow of endoplasm as is commonly found in *Amoeba proteus* (Figs. 46; 184). It is finger- or tongue-like, sometimes branched, and its distal end is typically rounded. It is quickly formed and equally quickly retracted. In many cases, there are many pseudopodia formed from the entire body surface, in which the largest one will counteract the smaller ones and the organism will move in one direction; while in others, there may be a single pseudopodium formed, as in *Amoeba striata*, *A. guttula*, *Pelomyxa carolinensis* (Fig. 186, b), etc., in which case it is a broadly tongue-like extension of the body in one direction and the progressive movement of the organisms is comparatively rapid. The lobopodia may occasionally be conical in general shape, as in *Amoeba spumosa* (Fig. 185, a). Although ordinarily the formation of lobopodia is by a general flow of the cytoplasm, in some it is sudden and "eruptive," as in *Endamoeba blattae* or *Entamoeba histolytica* in which the flow of the endoplasm presses against the inner zone of the ectoplasm and the accumulated pressure finally causes a break through the zone, resulting in a sudden extension of the endoplasmic flow at that point.

2). **Filopodium** is a more or less filamentous projection composed almost exclusively of the ectoplasm. It may sometimes be branched, but the branches do not anastomose. Many testaceans, such as Lecythium, Boderia, Plagiophrys, Pamphagus, Euglypha, etc., form this type of pseudopodia. The pseudopodia of *Amoeba radiosa* may be considered as approaching this type rather than the lobopodia.

3). **Rhizopodium** is also filamentous, but branching and anastomosing. It is found in numerous Foraminifera, such as Elphidium (Fig. 5), Peneroplis, etc., and in certain testaceans, such
as Lieberkühnia, Myxotheca, etc. The abundantly branching and anastomosing rhizopodia often produce a large network which serves almost exclusively for capturing prey.

4). Axopodium is, unlike the other three types, a more or less semi-permanent structure and composed of axial rod and cytoplasmic envelope. Axopodia are found in many Heliozoa, such as Actinophrys, Actinosphaerium, Camptonema, Sphaerastrum, and Acan-
thocystis. The axial rod, which is composed of a number of fibrils (Doeflein; Roskin, 1925; Rumjantzew and Wermel, 1925), arises from the central body or the nucleus located in the approximate center of the body, from each of the nuclei in multinucleate forms, or from the zone between the ectoplasm and endoplasm (Fig. 6). Although semipermanent in structure, the axial rod is easily absorbed and reformed. In the genera of Heliozoa not mentioned above and in numerous radiolarians, the radiating filamentous pseudopodia are so extremely delicate that it is difficult to determine whether an axial rod exists in each or not, although they resemble axopodia in general appearance.

There is no sharp demarcation between the four types of pseudopodia, as there are transitional pseudopodia between any two of them. For example, the pseudopodia formed by Arcella, Lesquer-eusia, Hyalosphaenia, etc., resemble more lobopodia than filopodia, though composed of the ectoplasm only. The pseudopodia of Actinomonas, Elaeorhanis, Clathrulina, etc., may be looked upon as transitional between rhizopodia and axopodia.

While the pseudopodia formed by an individual are usually of characteristic form and appearance, they may show an entirely different appearance under different circumstances. According to
the often-quoted experiment of Verworn, a limax amoeba changed into a radiosa amoeba upon addition of potassium hydroxide to the water (Fig. 7). Mast has recently shown that when Amoeba proteus or A. dubia was transferred from a salt medium into pure water, the amoeba produced radiating pseudopodia, and when transferred back to a salt medium, it changed into monopodal form, which change he was inclined to attribute to the difference in the water contents of the amoeba. In some cases during and after certain internal changes, an amoeba may show conspicuous differences in pseudopodia (Neresheimer). As was stated before, pseudopodia occur widely in forms which are placed under classes other than Sarcodina during a part of their life-cycle. Care, therefore, should be exercised in using them for taxonomic consideration of the Protozoa.

Flagella. The flagellum is a filamentous extension of the cytoplasm and is ordinarily extremely fine and highly vibratile, so that it is difficult to recognize it distinctly in life under the microscope. It is most clearly observed under a darkfield or phase microscope. Lugol’s solution usually makes it more easily visible, though the organism is killed. In a small number of species, the flagellum can be seen in life under an ordinary microscope as a long filament, as for example in
Peranema. As a rule, the number of flagella present in an individual is small, varying from one to eight and most commonly one or two; but in Hypermastigina there occur numerous flagella.

A flagellum appears to be composed of two parts: an elastic axial filament or axoneme, made up of one to several fibrils and the contractile cytoplasmic sheath surrounding the axoneme (Fig. 8, a, b). In some flagella, both components extend the entire length and terminate in a bluntly rounded point, while in others the distal portion of the axoneme is apparently very thinly sheathed (Fig. 8, c).

Fig. 8. Diagrams of flagella. a, flagellum of Euglena (Bütschli); b, flagellum of Trachelomonas (Plenge); c, flagella of Polytoma welia; d, flagella of Monas socialis (Vlk).

In some flagellates, stained flagella show numerous lateral fibrils (Fig. 8, d) (Fischer, 1894; Dellinger, 1909; Mainx, 1929; Petersen, 1929; etc.). These flagella or ciliary flagella have also been noticed by several observers in unstained organisms under darkfield microscope (Vlk, 1938; Pitelka, 1949). In recent years, the electron microscope has been used by some to observe the flagellar structure (Schmitt, Hall and Jakus, 1943; Brown, 1945; Pitelka, 1949; Chen, 1950), but in all cases, the organisms were air-dried on collodion films for examination so that the flagella disintegrated more or less completely at the time of observation.

Pitelka (1949) studied flagella of euglenoid organisms under light and electron microscopes. She found that the flagellum of *Euglena*
gracilis, Astasia longa and Rhabdomonas incurva, consists of an axoneme, composed of about 9 fibrils, 350–600 Å in diameter, arranged in two compact, parallel bundles, and a sheath which is made up of fibrillar elements, a probably semi-fluid matrix and a limiting membrane. Under conditions always associated with death of the organism, the fibrils of the sheath fray out on one or more sides of the flagellum into fine lateral filaments or mastigonemes. The electron micrographs obtained by various investigators on supposedly one and the same flagellate present a varied appearance of the structure. Compare, for example, the micrographs of the frayed flagellum of Euglena gracilis by Brown (1945), Pitelka (1949) and Houwink (1951). The anterior flagellum of Peranema trichophorum frays out into three strands during the course of disintegration as first observed by Dellinger (1909) and by several recent observers. It can be easily demonstrated by treating the organism with reagents such as acidified methyl green. Under electron microscope, Petelka noted no frayed mastigonemes in the flagellum of Peranema, while Chen (1950) observed numerous mastigonemes extending out from all sides like a brush, except the basal portion of the flagellum.

The electron micrographs of the flagellum of trypanosomes reveal that it also consists of an axoneme and a sheath of cytoplasm. The axoneme is composed of a number of long parallel fibrils, 8 in Trypanosoma lewisi, each with estimated diameters of 0.055–0.06 μ (Kleinschmidt and Kinder, 1950), and up to 9 in T. evansi, with estimated diameters of 0.04–0.05 μ (Kraneveld, Houwink and Keidel, 1951). The cytoplasmic sheath of the latter species was said to be cross-striated at about 0.05 μ intervals. No mastigonemes occur in these flagella.

The frayed condition of a flagellum which had become detached from the organism or which is still attached to a moribund individual, as revealed by the darkfield microscope, may also indicate a phase in disintegration of the flagellum. It is reasonable to assume that different flagella may have structural differences as revealed by the electron microscope, but evidence for the occurrence of mastigonemes on an active flagellum of a normally living organism appears not to be on hand.

A flagellum takes its origin in a blepharoplast of kinetosome imbedded in the cytoplasm. The blepharoplast is a small compact granule, but in certain parasitic flagellates, it may be comparatively large and ovoid or short rod-shaped, surrounded often by a halo. Whether this is due to the presence of a delicate cortical structure enveloping the compact body or to desiccation or fixation is un-
known. In such forms, the flagellum appears to arise from the outer edge of the halo. Certain observers such as Woodcock (1906), Minchin (1912), etc., used the term kinetoucleus. It has since been found that the blepharoplast of certain trypanosomes often gives a positive Feulgen’s reaction (Bresslau and Scremin, 1924).

The blepharoplast and centriole are considered synonymous by some, since prior to the division of nucleus, it divides and initiates the division of the latter. A new flagellum arises from one of the daughter blepharoplasts. While the blepharoplast is inseparably connected with the flagellum and its activity, it is exceedingly small or absent in *Trypanosoma equinum* and in some strains of *T. evansi*. Furthermore, this condition may be produced by exposure of normal individuals to certain chemical substances (Jírovec, 1929; Piekarski, 1949) or spontaneously (p. 228) without decrease in flagellar activity.

The flagellum is most frequently inserted near the anterior end of the body and directed forward, its movement pulling the organism forward. Combined with this, there may be a trailing flagellum which is directed posteriorly and serves to steer the course of movement or to push the body forward to a certain extent. In a comparatively small number of flagellates, the flagellum is inserted near the posterior end of the body and would push the body forward by its vibration. Under favorable conditions, flagellates regenerate lost flagella. For example, *Peranema trichophorum* from which its anterior flagellum was cut off, regenerated a new one in two hours (Chen, 1950).

In certain parasitic Mastigophora, such as Trypanosoma (Fig. 9), Trichomonas, etc., there is a very delicate membrane extending out from the side of the body, a flagellum bordering its outer margin. When this membrane vibrates, it shows a characteristic undulating movement, as will easily be seen in *Trypanosoma rotatorium* of the frog, and is called the undulating membrane. In many of the dinoflagellates, the transverse flagellum seems to be similarly constructed (Kofoid and Swezy) (Fig. 127, d, f).

**Cilia.** The cilia are the organella of locomotion found in the Ciliophora. They aid in the ingestion of food and serve often as a tactile organella. The cilia are fine and more or less short processes of ectoplasm and occur in large numbers in the majority of the Holotricha. They may be uniformly long, as in Protociliata, or may be of different lengths, being longer at the extremities, on certain areas, in peristome or in circumoral areas. Ordinarily the cilia are arranged in longitudinal, oblique, or spiral rows, being inserted either on the ridges or in the furrows. A cilium originates in a kinetosome embedded
in the ectoplasm. In well-studied ciliates, there occurs a fine fibril, *kinetodesma* (Chatton and Lwoff, 1935), a short distance to the right of the kinetosome (Fig. 23). The ciliary row or *kinety* (Chatton and Lwoff) consists of the kinetosomes and kinetodesma (Fig. 23, a). In forms such as Suctoria in which cilia occur only in the swimming stage, the kinetosomes appear to be present as *infraciliature* (Chatton, Lwoff and Lwoff, 1929).

![Diagram of a trypanosome](image)

**Fig. 9.** A diagram showing the structure of a trypanosome (Kühn).

As to its structure, a cilium appears to be made up of an axoneme and contractile sheath (Fig. 10, a). Gelei observed in flagella and cilia, lipoid substance in granular or rod-like forms which differed even among different individuals of the same species; and Klein (1929) found in many cilia of *Colpidium colpoda*, an argentophilous substance in granular form much resembling the lipoid structure of Gelei and called them "cross striation" of the contractile component (Fig. 10, b, c). In electron micrographs of a dried cilium of *Paramecium*, Jakus and Hall (1946) found that it consisted of a bundle of about 11 fibrils extending the full length (Fig. 10, d). These fibrils were about 300–500 Å in diameter. As there was no visible sheath, the two observers remarked that if a sheath exists, it must be very fragile and easily ruptured.

The cilia are often present more densely in a certain area than in other parts of body and, consequently, such an area stands out conspicuously, and is sometimes referred to as a *ciliary field*. If this area is in the form of a zone, it may be called a *ciliary zone*. Some authors use *pectinellae* for short longitudinal rows or transverse
bands of close-set cilia. In a number of forms, such as Coleps, Stentor, etc., there occur, mingled among the vibratile cilia, immobile stiff cilia which are apparently solely tactile in function.

![Diagram](image.png)

**Fig. 10.** a, cilia of Coleps; b, cilium of Cyclidium glaucoma; c, basal portion of a cilium of Colpidium colpoda, all in silver preparations (Klein); d, electronmicrograph of a dried cilium of Paramecium, shadow-cast with chromium, ×11,000 (Jakus and Hall).

In the Hypotricha, the cilia are largely replaced by cirri, although in some species both may occur. A cirrus is composed of a number of cilia arranged in 2 to 3 rows that fused into one structure completely (Figs. 11, a; 12, a), which was demonstrated by Taylor. Klein also showed by desiccation that each marginal cirrus of Stylonychia
was composed of 7 to 8 cilia. In some instances, the distal portion of a cirrus may show two or more branches. The cirri are confined to the ventral surface in Hypotricha, and called frontal, ventral, anal, caudal, and marginal cirri, according to their location (Fig. 11, b). Unlike cilia, the cirri may move in any direction so that the organisms bearing them show various types of locomotion. Oxytricha,
Stylonychia, etc., “walk” on frontals, ventrals, and analns, while swimming movement by other species is of different types.

In all euciliates except Holotricha, there are adoral membranellae. A **membranella** is composed of a double ciliary lamella, fused completely into a plate (Fig. 12, b). A number of these membranellae occur on a margin of the peristome, forming the **adoral zone of membranellae**, which serves for bringing the food particles to the cytostome as well as for locomotion. The frontal portion of the zone, the so-called **frontal membrane** appears to serve for locomotion and Kahl considers that it is probably made up of three lamellae. The oral membranes which are often found in Holotricha and Heterotricha, are transparent thin membranous structures composed of one or two rows of cilia, which are more or less strongly fused. The membranes, located in the lower end of the peristome, are sometimes called perioral membranes, and those in the cytopharynx, undulating membranes.

In Suctoria, cilia are present only during the developmental stages, and, as the organisms become mature, tentacles develop in their stead. The **tentacles** are concerned with food-capturing, and

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**Fig. 12.** Diagrams of cirrus and membranella of *Euplotes eurystomus*, X1450 (Taylor). a, anal cirrus in side view; b, a membranella (cpg, coagulated protoplasmic granules; cr, ciliary root; fp, fiber plate; k, kinetosome).
are either prehensile or usually suctorial. The prehensile tentacle appears to be essentially similar in structure to the axopodium (Roskin, 1925). The suctorial tentacles are tubular and this type is interpreted by Collin as possibly derived from cytostome and cytopharynx of the ciliate (Fig. 13).

Although the vast majority of Protozoa possess only one of the three organelles of locomotion mentioned above, a few may possess pseudopodia in one stage and flagella in another during their development. Among several examples may be mentioned Naegleriidae (Fig. 183), Tetramitus rostratus (Fig. 155), etc. Furthermore, there are some Protozoa which possess two types of organellae at the same time. Flagellum or flagella and pseudopodia occur in many Phytomastigina and Rhizomastigina, and a flagellum and cilia are present in Ileonema (Fig. 306, b, c).

In the cytoplasm of Protozoa there occur various organellae, each of which will be considered here briefly.

**Fibrillar structures**

One of the fundamental characteristics of the protoplasm is its contractility. If a fully expanded *Amoeba proteus* is subjected to a mechanical pressure, it retracts its pseudopodia and contracts into a more or less spherical form. In this response there is no special organella, and the whole body reacts. But in certain other Protozoa, there are special organellae of contraction. Many Ciliophora are able to contract instantaneously when subjected to mechanical pressure, as will easily be noticed by following the movement of Stentor, Spirostomum, Trachelocerca, Vorticella, etc., under a dissecting microscope. The earliest observer of the contractile elements of Protozoa appears to be Lieberkühn (1857) who noted the "muscle

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**Fig. 13.** Diagrams showing the possible development of a suctorian tentacle from a cytostome and cytopharynx of a ciliate (Collin).
fibers” in the ectoplasm of Stentor which were later named myonemes (Haeckel) or neurophanes (Neresheimer).

The myonemes of Stentor have been studied by several investigators. According to Schröder (1906), there is a canal between each two longitudinal striae and in it occurs a long banded myoneme which measures in cross-section 3-7μ high by about 1μ wide and which appears cross-striated (Fig. 14). Roskin (1923) considers that

![Fig. 14. Myonemes in Stentor coeruleus (Schröder). a, cross-section of the ectoplasm; b, surface view of three myonemes; c, two isolated myonemes (cl, cilium; gis, granules between striae; k, kinetosome; m, myoneme; mc, myoneme canal).](image)

the myoneme is a homogeneous cytoplasm (kinoplasm) and the wall of the canal is highly elastic and counteracts the contraction of the myonemes. All observers agree that the myoneme is a highly contractile organella.

Many stalked peritrichous ciliates have well-developed myonemes not only in the body proper, but also in the stalk. Koltzoff’s (1911) studies show that the stalk is a pseudochitinous tube, enclosing an inner tube filled with granulated thecoplasms, which surrounds a central rod, composed of kinoplasm, on the surface of which are ar-
ranged skeletal fibrils (Fig. 15). The contraction of the stalk is brought about by the action of kinoplasm and walls, while elastic rods will lead to extension of the stalk. Myonemes present in the ciliates aid in the contraction of body, but those which occur in many Gregarinida aid apparently in locomotion, being arranged longitudinally, transversely and probably spirally (Roskin and Levinsohn, 1929) (Fig. 15, c). In certain Radiolaria, such as Acantho-

Fig. 15. a, b, fibrillar structures of the stalk of Zoothamnium (Koltzoff); c, myonemes in Gregarina (Schneider). ef, elastic fiber; ie, inner envelope; k, kinoplasm; oe, outer envelope; t, thecoplas.

metron elasticum (Fig. 219, c), etc., each axial spine is connected with 10–30 myonemes (myophrisks) originating in the body surface. When these myonemes contract, the body volume is increased, thus in this case functioning as a hydrostatic organella.

In Isotricha prostoma and I. intestinalis, Schuberg (1888) observed that the nucleus is suspended by ectoplasmic fibrils and called the apparatus karyophore. In some forms these fibrils are replaced by ectoplasmic membranes as in Nyctotherus ovalis (Zulueta; Kudo). ten Kate (1927, 1928) studied fibrillar systems in Opalina, Nycto-
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therus, Ichthyophthirius, Didinium, and Balantidium, and found that there are numerous fibrils, each of which originates in the kinetosome of a cilium and takes a transverse or oblique course through the endoplasm, ending in a kinetosome located on the other side of the body. He further noted that the cytopharynx and nucleus are also connected with these fibrils. ten Kate suggested morphonemes for them, since he believed that the majority were form-retaining fibrils.

The well-coordinated movement of cilia in the ciliate has long been recognized, but it was Sharp (1914) who definitely showed that this ciliary coordination is made possible by a certain fibrillar system which he discovered in Epidinium (Diplodinium) ecaudatum (Fig. 16). Sharp recognized in this ciliate a complicated fibrillar system connecting all the motor organellae of the cytostomal region, and thinking that it was "probably nervous in function," as its size, arrangement and location did not suggest supporting or contractile function, he gave the name neuromotor apparatus to the whole system. This apparatus consists of a central motor mass, the motorium (which is stained red with Zenker fixation and modified Mallory's connective tissue staining), located in the ectoplasm just above the base of the left skeletal area, from which definite strands radiate: namely, one to the roots of the dorsal membranellae (a dorsal motor strand); one to the roots of the adoral membranellae (a ventral motor strand); one to the cytopharynx (a circum-oesophageal ring and oesophageal fibers); and several strands into the ectoplasm of the operculum (opercular fibers). A similar apparatus has since been observed in many other ciliates: Euplotes (Yocom; Taylor), Balantidium (McDonald), Paramecium (Rees; Brown; Lund), Tintinnopsis (Campbell), Boveria (Pickard), Dileptus (Visscher), Chlamydomon (MacDougall), Entorhipidium and Lechriopyla (Lynch), Eupoterion (MacLennan and Connell), Metopus (Lucas), Troglodytella (Swezey), Oxytricha (Lund), Ancistruma and Couchophthirius (Kidder), etc. Ciliate fibrillar systems (Taylor, 1941).

Euplotes, a common free-living hypotrichous ciliate, has been known for nearly 60 years to possess definite fibrils connecting the anal cirri with the anterior part of the body. Engelmann suggested that their function was more or less nervelike, while others maintained that they were supporting or contracting in function. Yocom (1918) traced the fibrils to the motorium, a very small bilobed body (about 8μ by 2μ) located close to the right anterior corner of the triangular cytostome (Fig. 17, m). Joining with its left end are five
Fig. 16. A composite drawing from three median sagittal sections of *Epidinium ecaudatum*, fixed in Zenker and stained with Mallory's connective tissue stain, ×1200 (Sharp). am, adoral membranellae; c, cytostome; cp, cytopharynx; cpg, cytopyge; cpr, circumpharyngeal ring; dd, dorsal disk; dm, dorsal membrane; ec, ectoplasm; en, endoplasm; m, motorium; oc, oral cilia; od, oral disk; oef, oesophageal fibers; of, opercular fibers; p, pellicle; prs, pharyngeal retractor strands; sl, skeletal laminae; vs, ventral skeletal area.
long fibers (acf) from the anal cirri which converge and appear to unite with the motorium as a single strand. From the right end of the motorium extends the membranella-fiber anteriorly and then to left along the proximal border of the oral lip and the bases of all membranelles. Yocom further noticed that within the lip there is a latticework structure whose bases very closely approximate the cyto-stomal fiber. Taylor (1920) recognized two additional groups of fibrils in the same organism: (1) membranella fiber plates, each of which is contiguous with a membranella basal plate, and is attached at one end to the membranella fiber; (2) dissociated fiber plates contiguous with the basal plates of the frontal, ventral and marginal cirri, to each of which are attached the dissociated fibers (rf). By means of microdissection needles, Taylor demonstrated that these
fibers have nothing to do with the maintenance of the body form, since there results no deformity when Euplotes is cut fully two-thirds its width, thus cutting the fibers, and that when the motorium is destroyed or its attached fibers are cut, there is no coordination in the movements of the adoral membranellae and anal cirri. Hammond (1937) and Hammond and Kofoid (1937) find the neuromotor system continuous throughout the stages during asexual reproduction and conjugation so that functional activity is maintained at all times.

A striking feature common to all neuromotor systems, is that there seems to be a central motorium from which radiate fibers to different ciliary structures and that, at the bases of such motor organelae, are found the kinetosomes or basal plates to which the "nerve" fibers from the motorium are attached.

Independent of the studies on the neuromotor system of American investigators, Klein (1926) introduced the silver-impregnation method which had first been used by Golgi in 1873 to demonstrate various fibrillar structures of metazoan cells, to Protozoa in order to demonstrate the cortical fibers present in ciliates, by dry-fixation and impregnating with silver nitrate. Klein (1926–1942) subjected ciliates of numerous genera and species to this method, and observed that there was a fibrillar system in the ectoplasm at the level of the kinetosomes which could not be demonstrated by other methods. Klein (1927) named the fibers silver lines and the whole complex, the silverline system, which vary among different species (Figs. 18–20). Gelei, Chatton and Lwoff, Jírovec, Lynch, Jacobson, Kidder, Lund, Burt, and others, applied the silver-impregnation method to many other ciliates and confirmed Klein's observations. Chatton and Lwoff (1935) found in Apostomea, the system remains even after the embryonic cilia have entirely disappeared and considered it infraciliature.

The question whether the neuromotor apparatus and the silverline system are independent structures or different aspects of the same structure has been raised frequently. Turner (1933) found that in Euplotes patella (E. eurystomus) the silverline system is a regular latticework on the dorsal surface and a more irregular network on the ventral surface. These lines are associated with rows of rosettes from which bristles extend. These bristles are held to be sensory in function and the network, a sensory conductor system, which is connected with the neuromotor system. Turner maintains that the neuromotor apparatus in Euplotes is augmented by a distinct but connected external network of sensory fibrils. He however finds no motorium in this protozoan.
Lund (1933) also made a comparative study of the two systems in *Paramecium multimicronucleatum*, and observed that the silverline system of this ciliate consists of two parts. One portion is made up of a series of closely-set polygons, usually hexagons, but flattened into rhomboids or other quadrilaterals in the regions of the cytostome, cytopyge, and suture. This system of lines stains if the organisms are well dried. Usually the lines appear solid, but frequently they are interrupted to appear double at the vertices of the polygons which Klein called "indirectly connected" (pellicular) conductile system. In the middle of the anterior and posterior sides of the hexagons is found one granule or a cluster of 2–4 granules, which marks the outer end of the trichocyst. The second part which Klein called "directly connected" (subpellicular) conductile system consists essentially of the longitudinal lines connecting all kinetosomes in a longitudinal row of hexagons and of delicate transverse fibrils connecting granules of adjacent rows especially in the cytostomal region (Fig. 19).

By using Sharp’s technique, Lund found the neuromotor system

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**Fig. 18.** The silverline system of *Ancistruma mytili*, ×1000 (Kidder). a, ventral view; b, dorsal view.
of *Paramecium multimicronucleatum* constructed as follows: The subpellicular portion of the system is the longitudinal fibrils which connect the kinetosomes. In the cytostomal region, the fibrils of right and left sides curve inward forming complete circuits (the circular cytostomal fibrils) (Fig. 20). The postoral suture is separated at the point where the cytopyge is situated. Usually 40–50 fibrils radiate outward from the cytostome (the radial cytostomal fibrils).

The pharyngeal portion is more complex and consists of (1) the oesophageal network, (2) the motorium and associated fibrils, (3) penniculus which is composed of 8 rows of kinetosomes, thus forming a heavy band of cilia in the cytopharynx, (4) oesophageal process, (5) paraoesophageal fibrils, (6) posterior neuromotor chain, and (7) postoesophageal fibrils. Lund concludes that the so-called silverline system includes three structures: namely, the peculiarly ridged pellicle; trichocysts which have no fibrillar connections among them or with fibrils, hence not conductile; and the subpellicular system, the last of which is that part of the neuromotor system that concerns with the body cilia. ten Kate (1927) suggested that senso-motor apparatus is a better term than the neuromotor apparatus. Silverline system (Klein, 1926–1942; Gelei, 1932); fibrils in ciliates

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**Fig. 19.** Diagram of the cortical region of *Paramecium multimicronucleatum*, showing various organellae (Lund). c, cilia; et, tip of trichocyst; k, kinetosome; lf, longitudinal fibril; p, pellicle; t, trichocyst; tf, transverse fibril.
Fig. 20. The neuromotor system of Paramecium multimicronucleatum (Lund). a, oral network; b, motorium, X1670. aep, anterior end of penniculus; c, cytopyge; ccf, circular cytostomal fibril; cof, circular oesophageal fibril; cpf, circular pharyngeal fibril; ef, endoplasmic fibrils; lbf, longitudinal body fibril; lof, longitudinal oesophageal fibrils; lpf, longitudinal pharyngeal fibril; m, motorium; oo, opening of oesophagus; op, oesophageal process; paf, paraoesophageal fibrils; pep, posterior end of penniculus; pnc, posterior neuromotor chain; pof, postoesophageal fibrils; rcf, radial cytostomal fibril; s, suture.
Protective or supportive organellae

The external structures as found among various Protozoa which serve for body protection, have already been considered (p. 47). Here certain internal structures will be discussed. The greater part of the shell of Foraminifera is to be looked upon as endoskeleton and thus supportive in function. In Radiolaria, there is a membranous structure, the central capsule, which divides the body into a central region and a peripheral zone. The intracapsular portion contains the nucleus or nuclei, and is the seat of reproductive processes, and thus the capsule is to be considered as a protective organella. The skeletal structures of Radiolaria vary in chemical composition and forms, and are arranged with a remarkable regularity (p. 517).

In some of the astomatous euciliates, there are certain structures which seem to serve for attaching the body to the host's organ, but which seem to be supportive to a certain extent also. The peculiar organella furcula, observed by Lynch in Lechriopyla (p. 741) is said to be concerned with either the neuromotor system or protection. The members of the family Ophryoscolecidae (p. 816), which are common commensals in the stomach of ruminants, have conspicuous endoskeletal plates which arise in the oral region and extend posteriorly. Dogiel (1923) believed that the skeletal plates of Cycloposthium and Ophryoscolecidae are made up of hemicellulose, "ophryoscolecin," which was also observed by Strelkow (1929). MacLennan found that the skeletal plates of Polyplostron multivesiculatum were composed of small, roughly prismatic blocks of paraglycogen, each possessing a central granule.

In certain Polymastigina and Hypermastigina, there occurs a flexible structure known as the axostyle, which varies from a filamentous structure as in several Trichomonas, to a very conspicuous rod-like structure occurring in Parajoenia, Gigantomonas, etc. The anterior end of the axostyle is very close to the anterior tip of the body, and it extends lengthwise through the cytoplasm, ending near the posterior end or extending beyond the body surface. In other cases, the axostyle is replaced by a bundle of axostylar filaments that are connected with the flagella (Lophomonas). The axostyle appears to be supportive in function, but in forms such as Saccinobaculus, it undulates and aids in locomotion (p. 379).

In trichomonad flagellates there is often present along the line of
attachment of the undulating membrane, a rod-like structure which has been known as **costa** (Kunstler) and which, according to Kirby’s extensive study, appears to be most highly developed in Pseudo-

trypanosoma and Trichomonas. The staining reaction indicates that its chemical composition is different from that of flagella, blepharoplast, parabasal body, or chromatin.

In the gymnostomatous ciliates, the cytopharynx is often sur-

rounded by rod-like bodies, and the entire apparatus is often called **oral** or **pharyngeal basket**, which is considered as supportive in function. These rods are arranged to form the wall of the cytopharynx in a characteristic way. For example, the oral basket of *Chilodonella cucullulus* (Fig. 312, c, d) is made up of 12 long rods which are so completely fused in part that it appears to be a smooth tube; in other forms, the rods are evidently similar to the tubular trichocysts or trichites mentioned below.

In numerous holotrichs, there occur unique organelles, **trichocysts**, imbedded in the ectoplasm, and usually arranged at right angles to the body surface, though in forms such as Cyclogramma, they are arranged obliquely. Under certain stimulations, the trichocysts “explode” and form long filaments which extend out into the surrounding medium. The shape of the trichocyst varies somewhat among different ciliates, being pyriform, fusiform or cylindrical (Penard, 1922; Krüger, 1936). They appear as homogeneous refractile bodies. The extrusion of the trichocyst is easily brought about by means of mechanical pressure or of chemical (acid or alkaline) stimulation.

In forms such as Paramecium, Frontonia, etc., the trichocyst is elongate pyriform or fusiform. It is supposed that within an expansible membrane, there is a layer of swelling body which is responsible for the remarkable longitudinal extension of the membrane (Krüger) (Fig. 21, a). In other forms such as Prorodon, Didinium, etc., the tubular trichocyst or **trichites** are cylindrical in shape and the membrane is a thick capsule with a coiled thread, and when stimulated, the extrusion of the thread takes place. The trichites of *Prorodon teres* measure about 10–11 μ long (Fig. 21, d) and when extruded, the whole measures about 20 μ; those of *Didinium nasutum* are 15–

20 μ long and after extrusion, measure about 40 μ in length (Fig. 21, e, f). In *Spathidium spathula* (Fig. 21, c), trichites are imbedded like a paling in the thickened rim of the anterior end. They are also distributed throughout the endoplasm and, according to Woodruff and Spencer, “some of these are apparently newly formed and being transported to the oral region, while others may well be trichites which have been torn away during the process of prey ingestion.”
Fig. 21. a, a schematic drawing of the trichocyst of *Paramecium caudatum* (Krüger) (b, base of the tip; c, cap; m, membrane; mt, membrane of extruded trichocyst; s, swelling body; t, tip); b, an extruded trichocyst, viewed under phase dark contrast, ×1800; c, trichites in *Spathidium spathula*, ×300 (Woodruff and Spencer); d, a diagram of the trichocyst of *Prorodon teres* (Krüger) (cg, capsule-granule; e, end-piece of filament; f, filament; w, capsule wall); e, f, normal and extruded trichocysts of *Didinium nasulum* (Krüger).
Whether the numerous 12–20μ long needle-like structures which Kahl observed in Remanella (p. 727) are modified trichites or not, is not known.

*Dileptus anser* feeds on various ciliates through the cytostome, located at the base of the proboscis, which possesses a band of long trichocysts on its ventral side. When food organisms come in contact with the ventral side of the proboscis, they give a violent jerk, and remain motionless. Visscher saw no formed elements discharged from the trichocysts, and, therefore, considered that these trichocysts contained a toxic fluid and named them toxicysts. But Krüger and Hayes (1938) found that the extruded trichocysts can be recognized.

Perhaps the most frequently studied trichocysts are those of Paramecium. They are elongate pyriform, with a fine tip at the broad end facing the body surface. The tip is connected with the pellicle (Fig. 19, t). Krüger found this tip is covered by a cap (Fig. 21, a) which can be seen under darkfield or phase microscope and which was demonstrated by Jakus (1945) in an electron micrograph (Fig. 22, a). When extruded violently, the entire structure is to be found outside the body of Paramecium. The extruded trichocyst is composed of two parts: the tip and the main body (Fig. 21, b). The tip is a small inverted tack, and may be straight, curved or bent. The main body or shaft is a straight rod, tapering gradually into a sharp point at the end opposite the tip. Extruded trichocysts measure 20–40μ or more in length, and do not show any visible structures, except a highly refractile granule present at the base of the tuck-shaped tip (Fig. 21, b). The electron microscope studies of the extruded trichocysts by Jakus (1945), Jakus and Hall (1946) and Wohlfarth-Bottermann (1950), show the shaft to be cross-striated (Fig. 22). Jakus considers that the main component of the trichocyst is a thin cylindrical membrane formed by close packing of longitudinal fibrils characterized by a periodic pattern (somewhat resembling that of collagen), and as the fibrils are in phase with respect to this pattern, the membrane appears cross-striated.

As to the mechanism of the extrusion, no precise information is available, though all observers agree that the contents of the trichocyst suddenly increase in volume. Krüger maintains that the trichocyst cap is first lifted and the swelling body increases enormously in volume by absorbing water and lengthwise extension takes place, while Jakus is inclined to think that the membrane itself extends by the sudden uptake of water.
How are these organelles formed? Tönniges (1914) believes that the trichocysts of *Frontonia leucas* originate in the endosomes of the macronucleus and development takes place during their migration to the ectoplasm. Brodsky (1924) holds that the trichocyst is composed of colloidal excretory substances and is first formed in the vicinity of the macronucleus. Chatton and Lwoff (1935) find how-

**Fig. 22.** Electronmicrographs of extruded trichocysts of Paramecium. a, dried and stained with phosphotungstic acid, ×11,000 (Jakus); b, a similarly treated one, ×15,000 (Jakus); c, shadow-cast with chromium, ×16,000 (Jakus and Hall).

ever in Gymnodinioides the trichocysts are formed only in tomite stage and each trichocyst arises from a *trichocystosome*, a granule formed by division of a kinetosome (Fig. 23, a–c). In Polyspira, the trichocyst formation is not confined to one phase, each kinetosome is said to give rise to two granules, one of which may detach itself, migrate into other part of the body and develops into a trichocyst (d). In Foettingeria, the kinetosomes divide in young trophont stage into *trichitosomes* which develop into trichites (e). The two authors note that normally cilia-producing kinetosomes may give rise to
trichocysts or trichites, depending upon their position (or environment) and the phase of development of the organism.

Although the trichocyst was first discovered by Ellis (1769) and so named by Allman (1855), nothing concrete is yet known as to their function. Ordinarily the trichocysts are considered as a defensive organella as in the case of the oft-quoted example Paramecium, but, as Mast demonstrated, the extruded trichocysts of this ciliate do not have any effect upon Didinium other than forming a viscid mass about the former to hamper the latter. On the other hand, the trichocysts and trichites are clearly an offensive organelle in capturing food organisms in organisms such as Dileptus, Didinium, Spathidium, etc. Saunders (1925) considered that the extruded trichocysts of Paramecium serve for attachment of the body to other objects. But Wohlfarth-Bottermann (1950) saw Paramecium caudatum extruding up to 300 trichocysts without any apparent external stimulation and trichocyst-less individuals were able to adhere to foreign objects. This worker suggested that the trichocyst secretes calcium salt and probably also sodium and potassium, and thus may serve an osmoregulatory function. Some years ago Penard (1922) considered that some trichocysts may be secretory organellae to produce material for loricae or envelope, with which view Kahl concurs, as granular to rod-shaped trichocysts occur in Metopus, Amphilep-

Fig. 23. Diagrams showing the formation of trichocysts in Gymnodinioides (a-c) and in Polyspira (d) and of trichites in Foettingeria (e) (Chatton and Lwoff). a, a ciliary row, composed of kinetosomes, large satellite corpuscles and kinetodesma (a solid line); b, each kinetosome divides into two, producing trichocystosome; c, transformation of trichocystosomes into trichocysts; d, formation of trichocyst from one of the two division products of kinetosome; e, formation of trichites from the division products of kinetosomes.
tus, etc. Klein has called these ectoplasmic granules **protrichocysts**, and in Prorodon, Krüger observed, besides typical tubular trichocysts, torpedo-like forms to which he applied the same name. To this group may belong the trichocysts recognized by Kidder in *Conchophthirus mytili*. The trichocysts present in certain Cryptomonadina (Chilomonas and Cyathomonas) are probably homologous with the protrichocysts (Krüger, 1934; Hollande, 1942; Dragesco, 1951).

**Hold-fast organellae**

In the Mastigophora, Ciliophora, and a few Sarcodina, there are forms which possess a **stalk** supporting the body or the lorica. With the stalk the organism is attached to a solid surface. In some cases, as in Anthophysis, Maryna, etc., the dendritic stalks are made up of gelatinous substances rich in iron, which gives to them a reddish brown color. In parasitic Protozoa, there are special organelles developed for attachment. Many genera of cephaline gregarines are provided with an **epimerite** of different structures (Figs. 235–237), by which the organisms are able to attach themselves to the gut epithelium of the host. In Astomata, such as Intoshellina, Maupasella, Lachmannella, etc., simple or complex protrusible chitinaceous structures are often present in the anterior region; or a certain area of the body may be concave and serves for adhesion to the host, as in Rhizocaryum, Perezella, etc.; or, again, there may be a distinctive sucker-like organella near the anterior extremity of the body, as in Haptophyra, Steinella, etc. A sucker is also present on the antero-ventral part of *Giardia intestinalis*.

In the Myxosporidia and Actinomyxidia, there appear, during the development of spore, 1–4 special cells which develop into **polar capsules**, each, when fully formed, enclosing a more or less long spirally coiled delicate thread, the **polar filament** (Figs. 279, 286). The polar filament is considered as a temporary anchoring organelle of the spore at the time of its germination after it gained entrance into the alimentary canal of a suitable host. In the Microsporidia, the filament may or may not be enclosed within a capsule (Figs. 288; 289). The **nematocysts** (Fig. 132, b) of certain dinoflagellates belonging to Nematoidium and Polykrikos, are almost identical in structure with those found in the coelenterates. They are distributed through the cytoplasm, and various developmental stages were noticed by Chatton, and Kofoid and Swezy, which indicates that they are characteristic structures of these dinoflagellates and not foreign in origin as had been held by some. The function of the nematocysts in these protozoans is not understood.
Parabasal apparatus

In the cytoplasm of many parasitic flagellates, there is frequently present a conspicuous structure known as the **parabasal apparatus** (Janicki, 1911), consisting of the parabasal body and often thread (Cleveland), which latter may be absent in some cases. This structure varies greatly among different genera and species in appearance, structure and position within the body. It is usually connected with the blepharoplast and located very close to the nucleus, though not directly connected with it. It may be single, double, or multiple, and may be pyriform, straight or curved rod-like, bandform, spirally coiled or collar-like (Fig. 24). Kofoid and Swezy considered that the parabasal body is derived from the nuclear chromatin, varies in size according to the metabolic demands of the organism, and is a "kinetic reservoir." On the other hand, Duboscq and Grassé (1933) maintain that this body is the Golgi apparatus, since (1) acetic acid destroys both the parabasal body and the Golgi apparatus; (2) both are demonstrable with the same technique; (3) the parabasal body

**Fig. 24.** Parabasal apparatus in: a, *Lophomonas blattarum* (Kudo); b, *Metadevescovina debilis*; c, *Devescovina* sp. (Kirby). af, axosty lar filaments; bl, blepharoplasts; f, food particles; fl, flagella; n, nucleus; pa, parabasal apparatus.
is made up of chromophile and chromophobe parts as is the Golgi apparatus; and (4) there is a strong evidence that the parabasal body is secretory in function. According to Kirby (1931), who has made an extensive study of this organella, the parabasal body could be stained with Delafield’s haematoxylin or Mallory’s triple stain after fixation with acetic acid-containing fixatives and the body does not show any evidence to indicate that it is a secretory organella. Moreover the parabasal body is discarded or absorbed at the time of division of the body and two new ones are formed.

The parabasal body of *Lophomonas blattarum* is discarded when the organism divides and two new ones are reformed from the centriole or blepharoplast (Fig. 65), and its function appears to be supportive. Possibly not all so-called parabasal bodies are homologous or analogous. A fuller comprehension of the structure and function of the organella rests on further investigations.

Golgi apparatus

With the discovery of a wide distribution of the so-called Golgi apparatus in metazoan cells, a number of protozoologists also reported a homologous structure from many protozoans. It seems impossible at present to indicate just exactly what the Golgi apparatus is, since the so-called Golgi techniques, the important ones of which are based upon the assumption that the Golgi material is osmiophile and argentophile, and possesses a strong affinity to neutral red, are not specific and the results obtained by using the same method often vary a great deal. Some of the examples of the Golgi apparatus reported from Protozoa are summarized in Table 2.

It appears thus that the Golgi bodies occurring in Protozoa are small osmiophilic granules or larger spherules which are composed of osmiophile cortical and osmiophobe central substances. Frequently the cortical layer is of unequal thickness, and, therefore, crescentic forms appear. Ringform apparatus was noted in *Chilodonella* and *Dogielella* by Nassonov (1925) and network-like forms were observed by Brown in *Pyrsonympha* and *Dinenympha*. The Golgi apparatus of Protozoa as well as of Metazoa appears to be composed of a lipoidal material in combination with protein substance.

In line with the suggestion made for the metazoan cell, the Golgi apparatus of Protozoa is considered as having something to do with secretion or excretion. Nassonov (1924) considers that osmiophilic lipoidal substance, which he observed in the vicinity of the walls of the contractile vacuole and its collecting canals in many ciliates and
flagellates, is homologous with the metazoan Golgi apparatus and secretes the fluid waste material into the vacuole from which it is excreted to the exterior. According to Brown, there is no blackening by osmic impregnation of the contractile vacuole in *Amoeba proteus*, (Fig. 25), but fusion of minute vacuoles associated with crescentic Golgi bodies produces the vacuole and Park (1929) noted osmiophile knob-like elevations on the surface of the macronucleus of Stentor and Leucophrys, while the contractile vacuole system did not blacken.

Duboscq and Grassé (1933) maintain that this body is a source of energy which is utilized by motor organelles. Joyet-Laverghne points out that in certain Sporozoa, the Golgi body is composed of granules and may be the center of enzyme production. Similar to Golgi material, the so-called vacuome, which consists of neutral red-staining and osmiophile globules, has been reported to occur in many Proto-

<table>
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<td>Hall</td>
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<tr>
<td>Chilomonas</td>
<td>Granules, vacuoles</td>
<td>Hall</td>
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<td>Euglenoidina</td>
<td>Stigma</td>
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<td><em>Euglena gracilis</em></td>
<td>Spherical, discoidal with dark rim, tend to group around or near nucleus</td>
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<tr>
<td>Pyrsonymphma, <em>Dinynymphma</em></td>
<td>Rings, crescents, spherules; granules break down to form network near posterior end</td>
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<td>Nassonov</td>
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**Table 2.---*Golgi apparatus in Protozoa***
zoa (Hall, 1931; Hall and Nigrelli, 1937). The exact morphological and physiological significance of these organellae and the relation between them must be looked for in future investigations. Golgi apparatus in Protozoa (Alexeieff, 1928; MacLennan, 1941; Grasse, 1952).

**Chondriosomes**

Widely distributed in many metazoan cells, the chondriosomes have also been recognized in various Protozoa. The chondriosomes possess a low refractive index, and are composed of substances easily soluble in alcohol, acetic acid, etc. Osmium tetroxide blackens the chondriosomes, but the color bleaches faster than in the Golgi bodies. Janus green B stains them even in 1:500,000 solution, but stains also other inclusions, such as the Golgi bodies (in some cases) and certain bacteria. According to Horning (1926), janus red is said to be a more exclusive chondriosome stain, as it does not stain bacteria. The chemical composition of the chondriosome seems to be somewhat similar to that of the Golgi body; namely, it is a protein compounded with a lipoidal substance. If the protein is small in amount, it is said to be unstable and easily attacked by reagents; on the other hand, if the protein is relatively abundant, it is more stable and resistant to reagents.

The chondriosomes occur as small spherical to oval granules, rod-

Fig. 25. The Golgi bodies in *Amoeba proteus* (Brown).
like or filamentous bodies, and show a tendency to adhere to or remain near protoplasmic surfaces. In many cases they are distributed without any definite order; in others, as in Paramecium or Opalina, they are regularly arranged between the kinetosomes of cilia (Horning). In *Tillina canaliculata*, Turner (1940) noticed that the endoplasmic chondriosomes are evenly distributed throughout the cytoplasm (Fig. 26, *b*), while the ectoplasmic chondriosomes are arranged in regular cross rows, one in the center of each square formed by four cilia (Fig. 26, *a*). In *Peranema trichophorum*, Hall (1929) observed peripheral chondriosomes located along the spiral striae, which Chadefaud (1938) considered as mucus bodies. Weisz (1949, 1950) finds that stentorin and zoopurpurin already mentioned (p. 45) are chondriosomes.

In certain Protozoa, the chondriosomes are not always demonstrable. For example, Horning states in Monocystis the chondriosomes present throughout the asexual life-cycle as rod-shaped bodies, but at the beginning of the spore formation they decrease in size and number, and in the spore none exists. The chondriosomes appear as soon as the sporozoites are set free. Thus it would appear that the
chondriosomes are reformed de novo. On the other hand, Fauré-Fremiet, the first student of the chondriosomes in Protozoa, maintained that they reproduce by division, which has since been confirmed by many observers. As a matter of fact, Horning found in Opalina, the chondriosomes are twisted filamentous structures and undergo multiple longitudinal fission in asexual division phase. Before encystment, the chondriosomes divide repeatedly transversely and become spherical bodies which persist during encystment and in the gametes. In zygotes, these spherical bodies fuse to produce longer forms which break up into elongate filamentous structures. Richardson and Horning further succeeded in bringing about division of the chondriosomes in Opalina by changing pH of the medium.

As to the function of chondriosomes, opinions vary. A number of observers hold that they are concerned with the digestive process. After studying the relationship between the chondriosomes and food vacuoles of Amoeba and Paramecium, Horning suggested that the chondriosomes are the seat of enzyme activity and it is even probable that they actually give up their own substance for this purpose. Mast (1926) described "beta granules" in Amoeba proteus which are more abundantly found around the contractile vacuole. Mast and Doyle (1935, 1935a) noted that these spherical to rod-like beta granules are plastic and stain like chondriosomes and that there is a direct relation between the number of beta granules in the cytoplasm and the frequency of contraction of the contractile vacuole. They maintained that these granules "probably function in transferring substances from place to place in the cytoplasm." Similar granules are recognizable in the species of Pelomyxa (Andresen, 1942; Wilber, 1942; Kudo, 1951).

The view that the chondriosomes may have something to do with the cell-respiration expressed by Kingsbury was further elaborated by Joyet-Laverne through his studies on certain Sporozoa. That the chondriosomes are actively concerned with the development of the gametes of the Metazoa is well known. Zweibaum's observation, showing an increase in the amount of fatty acid in Paramecium just prior to conjugation, appears to suggest this function. On the other hand, Calkins found that in Uroleptus, the chondriosomes became abundant in exconjugants, due to transformation of the macronuclear material into the chondriosomes. The author agrees with McBride and Hewer who wrote: "it is a remarkable thing that so little is known positively about one of the 'best known' protoplasmic inclusions" (Piney, 1931). Condriosomes in Protozoa (MacLennan, 1941; Grassé, 1952).
Numerous minute granules, less than 1µ in diameter, occur usually abundantly suspended in the cytoplasm. They can most clearly be noted under phase microscope. Mast named those found in Amoeba "alpha granules."

**Contractile and other vacuoles**

The majority of Protozoa possess one or more vacuoles known as pulsating or **contractile vacuoles**. They occur regularly in all freshwater-inhabiting Sarcodina, Mastigophora and Ciliophora. Marine or parasitic Sarcodina and Mastigophora do not ordinarily have a contractile vacuole. This organelle is present with a few exceptions in all marine and parasitic Ciliophora, while it is wholly absent in Sporozoa.

In various species of free-living amoebae, the contractile vacuole is formed by accumulation of water in one or more droplets which finally fuse into one. It enlarges itself continuously until it reaches a maximum size (diastole) and suddenly bursts through the thin cytoplasmic layer above it (systole), discharging its content to outside. The location of the vacuole is not definite in such forms and, therefore, it moves about with the cytoplasmic movements; and, as a rule, it is confined to the temporary posterior region of the body. Although almost spherical in form, it may occasionally be irregular in shape, as in *Amoeba striata* (Fig. 184, f). In many testaceans and heliozoans, the contractile vacuoles which are variable in number, are formed in the ectoplasm and the body surface bulges out above the vacuoles at diastole. In Mastigophora, the contractile vacuole appears to be located in the anterior region.

In the Ciliophora, except Protociliata, there occur one to many contractile vacuoles, which seem to be located in the deepest part of the ectoplasm and therefore constant in position. Directly above each vacuole is found a pore in the pellicle, through which the content of the vacuole is discharged to outside. In the species of Conchophthirius, Kidder (1934) observed a narrow slit in the pellicle just posterior to the vacuole on the dorsal surface (Fig. 27). The margin of the slit is thickened and highly refractile. During diastole, the slit is nearly closed and, at systole, the wall of the contractile vacuole appears to break and the slit opens suddenly, the vacuolar content pouring out slowly. When there is only one contractile vacuole, it is usually located either near the cytopharynx or, more often, in the posterior part of the body. When several to many vacuoles are present, they may be distributed without apparent order, in linear series, or along the body outline. When the contrac-
tile vacuoles are deeply seated, there is a delicate duct which connects the vacuole with the pore on the pellicle as in Paramecium woodrufi or in Ophryoscolecidae. In Balantidium, Nyctotherus, etc., the contractile vacuole is formed very close to the permanent cytopharynx located at the posterior extremity, through which it empties its content.

In a number of ciliates there occur radiating or collecting canals besides the main contractile vacuole. These canals radiate from the central vacuole in Paramecium, Frontonia, Disematostoma, etc. But when the vacuole is terminal, the collecting canals of course do not radiate, in which case the number of the canals varies among different species: one in Spirostomum, Stentor, etc., 2 in Climacostomum, Eschaneustyla, etc., and several in Tillina. In Peritricha, the contractile vacuole occurs near the posterior region of the cytopharynx and its content is discharged through a canal into the vestibule and in Ophrydium ectatum, the contractile vacuole empties its content into the cytopharynx through a long duct (Mast).

Of numerous observations concerning the operation of the contractile vacuole, that of King (1935) on Paramecium multimicronucleatum (Figs. 28, 29) may be quoted here. In this ciliate, there are 2 to 7 contractile vacuoles which are located below the ectoplasm on the aboral side. There is a permanent pore above each vacuole. Leading to the pore is a short tube-like invagination of the pellicle, with inner end of which the temporary membrane of the vacuole is in contact (Fig. 28, a). Each vacuole has 5–10 long collecting canals with strongly osmiophilic walls (Fig. 29), in which Gelei (1939) demonstrated longitudinal fibrils, and each canal is made up of terminal portion, a proximal injection canal, and an ampulla between them. Surrounding the distal portion, there is osmiophilic cytoplasm which may be granulated or finely reticulated, and

![Diagrams showing the contractile vacuole, the accessory vacuoles and the aperture, during diastole and systole in Conchophthirus (Kidder).](image_url)
which Nassonov (1924) interpreted as homologous with the Golgi apparatus of the metazoan cell. The injection canal extends up to the pore. The ampulla becomes distended first with fluid transported discontinuously down the canal and the fluid next moves into the injection canal. The fluid now is expelled into the cytoplasm just beneath the pore as a vesicle, the membrane of which is derived from that which closed the end of the injection canal. These fluid vesicles coalesce presently to form the contractile vacuole in full diastole and the fluid is discharged to exterior through the pore, which becomes closed by the remains of the membrane of the discharged vacuole.

In *Haptophrya michiganensis*, MacLennan (1944) observed that accessory vacuoles appear in the wall of the contractile canal which extends along the dorsal side from the sucker to the posterior end, as the canal contracts (Fig. 30). The canal wall expands and enlarging accessory vacuoles fuse with one another, followed by a full expansion of the canal. Through several excretory pores with short ducts the content of the contractile canal is excreted to the exterior. The function of the contractile vacuole is considered in the following

\[
\begin{array}{c}
\text{a} \\
\text{b} \\
\text{c} \\
\text{d} \\
\text{e} \\
\text{f} \\
\text{g} \\
\text{h}
\end{array}
\]

Fig. 28. Diagrams showing the successive stages in the formation of the contractile vacuole in *Paramecium multimicronucleatum* (King); upper figures are side views; lower figures front views; solid lines indicate permanent structures; dotted lines temporary structures. a, full diastole; b–d, stages of systole; e, content of ampulla passing into injection canal; f, formation of vesicles from injection canals; g, fusion of vesicles to form contractile vacuole; h, full diastole.
Fig. 29. Contractile vacuoles of *Paramecium multimicronucleatum*, X1200 (King). a, early systole, side view; b, diastole, front view; c, complete systole, front view; d, systole, side view.
chapter (p. 118). Comparative study of contractile vacuoles (Haye, 1930; Weatherby, 1941).

Various other vacuoles or vesicles occur in different Protozoa. In the ciliates belonging to Loxodidae, there are variable numbers of Müller's vesicles or bodies, arranged in 1–2 rows along the aboral surface. These vesicles (Fig. 31, a–c) vary in diameter from 5 to 8.5μ

Fig. 30. Excretory canal of *Haptophrya michiganensis* (MacLennan). a, an individual in side view, showing a contraction wave passing down the canal; b, successive views of the same region of the contractile canal during a full pulsatory cycle (a–c, systole; d–g, diastole); c, diagram showing a contractile wave passing from left to right between two adjacent excretory pores.

and contain a clear fluid in which one large spherule or several small highly refractile spherules are suspended. In some, there is a filamentous connection between the spherules and the wall of the vesicle. Penard maintains that these bodies are balancing cell-organs and called the vesicle, the statocyst, and the spherules, the statoliths.

Another vacuole, known as concrement vacuole, is a characteristic organella in Bütschliidae and Paraisotrichidae. As a rule, there is a single vacuole present in an individual in the anterior third of body. It is spherical to oval and its structure appears to be highly
complex. According to Dogiel (1929), the vacuole is composed of a pellicular cap, a permanent vacuolar wall, concrement grains and two fibrillar systems (Fig. 31, d). When the organism divides, the anterior daughter individual retains it, and the posterior individual develops a new one from the pellicle into which concrement grains enter after first appearing in the endoplasm. This vacuole shows no external pore. Dogiel believes that its function is sensory and has named the vacuole, the statocyst, and the enclosed grains, the statoliths.

**Food vacuoles** are conspicuously present in the holozoic Protozoa which take in whole or parts of other organisms as food. The food vacuole is a space in the cytoplasm, containing the fluid medium which surrounds the protozoans and in which are suspended the food matter, such as various Protophyta, other Protozoa or small Metazoa. In the Sarcodina and the Mastigophora which do not possess a cytostome, the food vacuoles assume the shape of the food materials and, when these particles are large, it is difficult to make out the thin film of water which surrounds them. When minute food

![Diagram](image_url)
particles are taken through a cytostome, as is the case with the majority of euciliates, the food vacuoles are usually spherical and of approximately the same size within a single protozoan. In the saprozoic Protozoa, which absorb fluid substances through the body surface, food vacuoles containing solid food, of course, do not occur.

**Chromatophore and associated organellae**

In the Phytomastigina and certain other forms which are green-colored, one to many chromatophores (Fig. 32) containing chlorophyll occur in the cytoplasm. The chromatophores vary in form among different species; namely, discoidal, ovoid, band-form, rod-like, cup-like, fusiform, network or irregularly diffused. The color of the chromatophore depends upon the amount and kinds of pigment which envelops the underlying chlorophyll substance. Thus the chromatophores of Chrysomonadina are brown or orange, as they contain one or more accessory pigments, including phycochrysin, and those of Cryptomonadina are of various types of brown with

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**Fig. 32.** a, *Trachelomonas hispida*, ×530 (Doflein); b, c, living and stained reproductive cells of *Pleodorina illinoisensis*, ×1000 (Merton); d–f, terminal cells of *Hydrurus foetidus*, showing division of chromatophore and pyrenoid (Geitler); g–i, *Chlamydomonas* sp., showing the division of pyrenoid (Geitler).
very diverse pigmentation. In Chloromonadina, the chromatophores are bright green, containing an excess of xanthophyll. In dinoflagellates, they are dark yellow or brown, because of the presence of pigments: carotin, phylloxyanthin, and peridinin (Kylin, 1927), the last of which is said to give the brown coloration. A few species of Gymnodinium contain blue-green chromatophores for which phyco-cyanin is held to be responsible. The chromatophores of Phytomonadina and Euglenoidina are free from any pigmentation, and therefore green. Aside from various pigments associated with the chromatophores, there are carotinoid pigments which occur often outside the chromatophores, and are collectively known as haematochrome. The haematochrome occurs in *Haematococcus pluvialis*, *Euglena sanguinea*, *E. rubra*, Chlamydomonas, etc. In Haematococcus, it increases in volume and in intensity when there is a deficiency in phosphorus and especially in nitrogen; and when nitrogen and phosphorus are present sufficiently in the culture medium, the haematochrome loses its color completely (Reichenow, 1909; Pringsheim, 1914). Steinecke also noticed that the frequent yellow coloration of phytomonads in moorland pools is due to a development of carotin in the chromatophores as a result of deficiency in nitrogen. Johnson (1939) noted that the haematochrome granules of *Euglena rubra* become collected in the central portion instead of being scattered throughout the body when sunlight becomes weaker. Thus this Euglena appears green in a weak light and red in a strong light. The chromatophores undergo division at the time when the organism which contains them, divides, and therefore the number of chromatophores appears to remain about the same through different generations (Fig. 32).

In association with the chromatophores are found the pyrenoids (Fig. 32) which are usually embedded in them. The pyrenoid is a viscous structureless mass of protein (Czurda), and may or may not be covered by tightly fitting starch-envelope, composed of several pieces or grains which appear to grow by apposition of new material on the external surface. A pyrenoid divides when it reaches a certain size, and also at the time of the division of the organism in which it occurs. As to its function, it is generally agreed that the pyrenoid is concerned with the formation of the starch and allied anabolic products of photosynthesis. Pyrenoid (Geitler, 1926).

Chromatophore-bearing Protozoa usually possess also a stigma (Fig. 32) or eye-spot. The stigma may occur in exceptional cases in colorless forms, as in Khawkinea, Polytomella, etc. It is ordinarily situated in the anterior region and appears as a reddish or
brownish red dot or short rod, embedded in the cortical layer of the cytoplasm. The color of the stigma is due to the presence of droplets of haematochrome in a cytoplasmic network. The stigma is incapable of division and a new one is formed de novo at the time of cell division. In many species, the stigma possesses no accessory parts, but, according to Mast (1928), the pigment mass in Chlamydomonas, Pandorina, Eudorina, Euglena, Trachelomonas, etc., is in cup-form, the concavity being deeper in the colonial than in solitary forms. There is a colorless mass in the concavity, which appears to function as a lens. In certain dinoflagellates, there is an ocellus (Fig. 127, c, d, g, h) which is composed of amyloid lens and a dark pigment mass (melanosome) that is sometimes capable of amoeboid change of form. The stigma is, in general, regarded as an organella for the perception of light intensity. Mast (1926) considers that the stigma in the Volvocidae is an organella which determines the direction of the movement.

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Chapter 4

Physiology

THE morphological consideration which has been given in the last chapter, is, though necessarily brief, indicative of the occurrence of various and often complex organellae in Protozoa. The physiological activity of the whole protozoan is the sum-total of all the functions which are carried on by numerous minute parts or organellae of the cell body, unlike the condition found in a metazoan. Indeed, as Calkins (1933) stated, “physiological problems (of Protozoa) for the most part begin where similar problems of the Metazoa leave off, namely the ultimate processes of the single cell. Here the functional activities have to do with the action and interaction of different substances which enter into the make-up of protoplasm and, for the most part, these are beyond our powers of analysis.” A full discussion of various physiological problems pertaining to Protozoa is out of question in the present work and, therefore, a general consideration on protozoan physiology will suffice for our purpose.

Nutrition

Protozoa obtain nourishment in manifold ways. Information on the nutrition of the Protozoa is undergoing an accelerated progress through improvements in technique in experimental cultivation. In many Phytomastigina (Pringsheim, 1937a; Hall, 1939), a few ciliates (Kidder and Dewey, 1951) and many blood-inhabiting flagellates (Lwoff, 1951) which have been cultivated in vitro free from other organisms, a much clearer information is becoming available. But for the majority of Protozoa a thorough comprehension of the nutrition is to be sought in future (Doyle, 1943; Lwoff, 1951; Most, 1951; Kidder, 1951).

Holozoic (zootrophic, heterotrophic) nutrition. This is the method by which all higher animals obtain their nourishment; namely, the protozoan uses other animals or plants as sources of food. It involves the food-capture and ingestion, digestion and assimilation, and rejection of indigestible portions.

The methods of food-capture vary among different forms. In the Sarcodina, the food organisms are captured and taken into the body at any point. The methods however vary. According to Rhumbler’s (1910) oft-quoted observations, four methods of food-ingestion occur in amoebae (Fig. 33); namely, (1) by “import,” in which the food is taken into the body upon contact, with very little movement on
the part of the amoeba (a); (2) by “circumfluence,” in which the cytoplasm flows around the food organism as soon as it comes in contact with it on all sides and engulfs it (b); (3) by “circumvallation,” in which the amoeba without contact with the food, forms pseudopodia which surround the food on all sides and ingest it (c);

(4) by “invagination,” in which the amoeba touches and adheres to the food, and the ectoplasm in contact with it is invaginated into the endoplasm as a tube, the cytoplasmic membrane later disappears (d–h). In a species of Hartmannella, Ray (1951) reports an agglutination of large numbers of motile bacteria over the body surface, which later form a large mass and are taken into a food cup.

In certain testaceans, such as Gromia, several rhizopodia cooperate in engulfing the prey and, in Lieberkühnia (Fig. 34), Verworn noted ciliates are captured by and digested in rhizopodia. Similar
observation was made by Schaudinn in the heliozoan Camptonema in which several axopodia anastomose to capture a prey (Fig. 214, d). In the holozoic Mastigophora, such as Hypermastigina, which do not possess cytostome, the food-ingestion is by import or invagination as noted in Trichonympha campanula (Cleveland, 1925a; Emik, 1941) (Fig. 35, a) and Lophomonas blattarum (Kudo, 1926).

The food particles become attached to the pseudopodium and are held there on account of the viscid nature of the pseudopod. The sudden immobility of active organisms upon coming in contact with pseudopodia of certain forms, such as Actinophrys, Actinosphaerium, Gromia, Elphidium, etc., suggests, however, probable discharge of poisonous substances. In the Suctoria which lack a cytostome, the tentacles serve as food-capturing organellae. The suctorial tentacle bears on its distal end a rounded knob which, when it comes in contact with an actively swimming ciliate, stops the latter immediately (Parapodophrya typha, Fig. 369, a). The prehensile tentacles of Ephelotidae are said to be similar in structure to the axopodia, in that each possesses a bundle of axial filaments around a cytoplasmic core (Roskin, 1925). These tentacles are capable of piercing through the body of a prey. In some suctorians, such as Choanophrya (Fig. 374, a), the tubular tentacles are clearly observable, and both solid and liquid food materials are sucked in through the cavity. The rapidity with which tentacles of a suctorian stop a very actively swimming ciliate is attributed to a certain substance secreted by the tentacles, which paralyses the prey.

In the cytostome-bearing Mastigophora, the lashing of flagella will aid in bringing about the food particles to the cytostome, where

Fig. 34. Rhizopodium of Lieberkühnia, capturing and digesting Colpidium colpoda (Verworn).
it is taken into the endoplasm. Chen (1950) observed Peranema feeding on immobile organisms. When the tip of the anterior flagellum comes in contact with an immobile Euglena, the whole flagellum

beats actively and the body contracts, followed by elongation. The process is repeated several times until the body touches Euglena. Then the cytostome stretches open, the oral rods move up, protrude from the body and become attached to Euglena. Peranema advances

Fig. 35. a, eight outline sketches of a *Trichonympha campanula*, ingesting a large particle of food, ×150 (Emik); b, four outline sketches of a *Peranema trichophorum* feeding on an immobile Euglena (Chen).
toward the prey and the whole Euglena is engulfed in 2 to 15 minutes (Fig. 35, b).

In the ciliates, there are many types of cytostome and associated organelles, but the food-capturing seems to be in general of two kinds. When the cytostome is permanently open, the organism ingests continuously food particles that are small enough to pass the cytostome and cytopharynx, as in the case of Paramecium. The other type is carried on by organisms bearing cytostome which is ordinarily closed such as seen in Coleps, Didinium, Perispira (Dewey and Kidder, 1940), but which expands to often an extraordinary size when the ingestion of prey takes place. Cannibalism in Protozoa (Dawson, 1919; Lapage, 1922; Gelei, 1925a; Tanabe and Komada, 1932; Giese and Alden, 1938; Chen, 1950).

The ingested food particles are usually surrounded by a film of fluid which envelops the organism and the whole is known as the food vacuole (p. 88). The quantity of fluid taken in with the food varies greatly and, generally speaking, it seems to be inversely proportional to the size, but proportional to the activity, of the food organisms. Food vacuoles composed entirely of surrounding liquid medium have occasionally been observed. Edwards (1925) noticed ingestion of fluid medium by an amoeba by forming food-cups under changed chemical composition. Brug (1928) reports seeing Entamoeba histolytica engulf liquid culture medium by formation of lip-like elevation of the ectoplasm and Kirby (1932) figures ingestion of the brine containing no visible organisms by the cytostome of Rhopalophrya salina (Fig. 36). Mast and Doyle (1934) state that if Amoeba proteus, A. dubia, A. doeleini, or A. radios a is placed in an albumin solution, a hypertonic balanced salt solution, or a hypertonic solution of calcium gluconate it rapidly decreases in volume, and forms numerous tubes filled with fluid, which disintegrate sooner or later and release their fluid content in the cytoplasm. At times 50 or more such tubes may be present, which indicates that the organism ingests considerable quantities of fluid in this way. The two authors consider that it is "a biological adaptation which serves to compensate for the rapid loss of water."

The food vacuoles finally reach the endoplasm and in forms such as Amoebina the vacuoles are carried about by the moving endoplasm. In the ciliates, the fluid endoplasm shows often a definite rotation movement. In Paramecium, the general direction is along the aboral side to the anterior region and down the other side, with a short cyclossis in the posterior half of the body.

Some observers maintain that in ciliates there is a definite "diges-
tive tubule” beginning with the cytostome and ending in the cytopyge, and the food vacuoles travel through it. Cosmovici (1931, 1932) saw such a canal in soluble starch-fed Colpidium colpoda upon staining with iodine, but Hall and Alvey (1933) could not detect such a structure in the same organism. Kitching (1938b) observed no such tubule in the peritrichous ciliates he studied, and concluded that the food vacuoles are propelled over the determined part of the course by the contraction of surrounding cytoplasm. In Vorticella sp., food vacuoles are formed one by one at the end of cytopharynx, migrate through different parts of the cytoplasm without order and food material is digested (Fig. 37, a). Old food vacuoles are defecated through a small papilla on the lower wall of the cytopharynx and thence to the outside (Hall and Dunihue, 1931) (Fig. 37, b–d).

![Fig. 36. Ingestion of brine by Rhopalophrya salina (Kirby).](image)

As stated above, in a number of species the food organisms are paralyzed or killed upon contact with pseudopodia, tentacles or exploded trichocysts. In numerous other cases, the captured organism is taken into the food vacuole alive, as will easily be noted by observing Chilomonas taken in by Amoeba proteus or actively moving bacteria ingested by Paramecium. But the prey ceases to move in a very short time. It is generally believed that some substances are secreted into the food vacuole by the protoplasm of the organisms to stop the activity of the prey within the food vacuole. Engelmann (1878) demonstrated that the granules of blue litmus, when ingested by Paramecium or Amoeba, became red in a few minutes. Brandt (1881) examined the staining reactions of amoebae by means of haematoxylin, and found that the watery vacuoles contained an acid. Metschnikoff (1889) also showed that there appears an acid secretion around the ingested litmus grains in Mycetozoa. Greenwood and Saunders (1894) found in Carchesium that ingestion of
food particles stimulated the cytoplasm to secrete a mineral acid. According to Nirenstein (1925), the food vacuole in Paramecium undergoes change in reaction which can be grouped in two periods. The first is acid reaction and the second alkaline reaction, in which albumin digestion takes place. On the other hand, Khainsky (1910) observed that the food vacuole of ciliates, such as Paramecium, is

acid during the entire period of protein digestion, and becomes neutral to finally alkaline when the solution of the food substance is ended. Metalnikoff (1912) found that in the food vacuoles of Paramecium, besides acid-alkaline reaction change, some vacuoles never show acid reaction and others occasionally show sustained acid reaction. Shapiro (1927) studied the reaction change of the food vacuoles in Paramecium caudatum by using phenol red, neutral red, Congo red, and litmus, and found that when the organism is kept in a medium with pH 7, its food vacuoles are first alkaline (pH 7.6), soon reach a maximum acidity (pH 4.0), while still in the posterior
half of the body. Later, the vacuoles show a decreased acidity, finally reaching pH 7.0. In Vorticella sp. and Stylonychia pustulata, the range of pH observed in the food vacuoles was said to be 4.5–7.0 and 4.8–7.0 respectively. The food vacuoles of Actinosphaerium, according to Howland (1928), possess at the beginning pH 6.0–7.0 for 5 to 10 minutes, but this soon changes to acid (pH 4.3) in which digestion appears to be carried on. In older food vacuoles which are of less acid (pH 5.4–5.6), the digestion appears to be at an end. In the species of Bresslaua, Claff, Dewey and Kidder (1941) noted that a Colpoda taken into the food vacuole is instantly killed with a sudden release of an acid which shows pH 3.0–4.2. During digestion the protoplasm of the prey becomes alkaline and the undigested residue becomes acid before extrusion.

Mast's observations (1942) on the food vacuoles in Amoeba proteus and A. dubia containing Chilomonas or Colpidium, indicate: (1) the fluid in the vacuoles becomes first acid and then alkaline; (2) the increase in the acidity of the fluid in the vacuole is not due to cytoplasmic secretion, but is probably due to respiration in the ingested organisms, chemical changes associated with their death, etc.; and (3) the death of the organisms taken in the food vacuoles is probably caused by the decrease in oxygen in the vacuoles, owing to the respiration of the organisms in them. De La Arena (1941, 1942) found the maximum acidity of the fluid of food vacuoles in Pelomyxa carolinensis containing Colpidium striatum was pH 5.8 and was not fatal for the ciliate, but considered the possibility of the existence in the food vacuole of "some lethal agent" which kills the prey.

Just exactly what processes take place in the food vacuole have been observed only in a few cases. Nirenstein (1925) noticed the appearance of numerous neutral red-stainable granules around the food vacuole which pass into the interior of the vacuole, and regarded them as carriers of a trypic ferment, while Roskin and Levinsohn (1926) demonstrated the oxidase reaction in these granules. Hopkins and Warner (1946) believe that the digestion of food in Entamoeba histolytica is brought about by enzymes carried to the food vacuoles by "digestive spherules" which arise at the periphery of the nucleus, apparently due to the action of the substances diffusing from the nucleus into the cytoplasm.

As to the localization or distribution of enzymes within protozoan body, definite information is not yet available. In centrifuged Amoeba proteus, Holter and Kopac (1937) found the peptidase activity independent of all cytoplasmic inclusions that were stratified by centrifugal forces. Holter and Løvtrup (1949) found peptidase in
centrifuged *Pelomyxa carolinensis* comparatively evenly distributed after centrifugation, possibly with a tendency to be concentrated in the lighter half, while proteinase was largely localized in the heavier half in which cytoplasmic granules were accumulated, and concluded that these two enzymes are bound, at least in part, to different cytoplasmic components. A number of enzymes have been reported to occur in Protozoa, some of which are listed in Table 3.

These findings suffice to indicate that the digestion in Protozoa is carried on also by enzymes and its course appears to vary among different Protozoa. The albuminous substances are digested and decomposed into simpler compounds by enzymes and absorbed by the surrounding cytoplasm. The power to digest starch into soluble sugars is widely found among various Protozoa. It has been reported in Mycetozoa, Foraminifera, Pelomyxa, Amoeba, Entamoeba, Ophryoscolecidae and other ciliates by several investigators.

The members of Vampyrella (p. 420) are known to dissolve the cellulose wall of algae, especially Spirogyra in order to feed on their contents. Pelomyxa (Stole), Foraminifera (Schaudinn), Amoeba (Rhumbler), Hypermastigina, Polymastigina (Cleveland), etc., have also been known for possessing the power of cellulose digestion. Many of the Hypermastigina and Polymastigina which lead symbiotic life in the intestine of the termite and of the wood roach, as demonstrated by Cleveland and his co-workers, digest by enzymes the cellulose which the host insect ingests. The assimilation products produced by an enormous number of these flagellates are seemingly sufficient to support the protozoans as well as the host. The ciliate commensals inhabiting the stomach of ruminants also apparently digest the cellulose, since the faecal matter as a rule does not contain this substance (Becker et al., 1930; Weineck, 1934).

Dawson and Belkin (1928) injected oils into *Amoeba dubia* and found 1.4 to 8.3 per cent digested. Mast (1938) reported that the neutral fat globules of Colpidium are digested by *Amoeba proteus* and transformed into fatty acid and glycerine which unite and form neutral fat. Chen (1950) found that when *Peranema trichophorum* was fed on almond oil (stained dark blue with Sudan black), Sudan III-stainable droplets gradually increased in number in five to 10 hours, while ingested oil-droplets decreased in size, and considered that the droplets were "fat-substances" resynthesized from products of digestion of almond oil by this flagellate. The digestion of rice starch is followed by the appearance of increasing number of ovoid paramylon granules, and the digestion of casein results in the formation of oil droplets and paramylon bodies.
### Table 3.—Enzymes in Protozoa

<table>
<thead>
<tr>
<th>Protozoa</th>
<th>Enzymes</th>
<th>Observers</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Amoeba proteus</em></td>
<td>Peptidase</td>
<td>Holter and Kopac (1937); Holter and Doyle (1938); Andreasen and Holter (1949); Holter and Løvtrup (1949)</td>
</tr>
<tr>
<td></td>
<td>Proteinase</td>
<td>Andreasen and Holter (1949); Holter and Løvtrup (1949)</td>
</tr>
<tr>
<td></td>
<td>Amylase</td>
<td>Holter and Doyle (1938a)</td>
</tr>
<tr>
<td><em>A. dubia</em></td>
<td>Lipolytic substance</td>
<td>Dawson and Belkin (1928)</td>
</tr>
<tr>
<td><em>Pelomyxa palustris</em></td>
<td>Diastatic enzyme</td>
<td>Hartog and Dixon (1893); Stolc (1900)</td>
</tr>
<tr>
<td></td>
<td>Pepsin-like enzyme</td>
<td>Hartog and Dixon (1893)</td>
</tr>
<tr>
<td></td>
<td>Peptidase</td>
<td>Andreasen and Holter (1949)</td>
</tr>
<tr>
<td></td>
<td>Proteinase</td>
<td>&quot;</td>
</tr>
<tr>
<td><em>P. carolinensis</em></td>
<td>Peptidase</td>
<td>&quot;</td>
</tr>
<tr>
<td></td>
<td>Protease</td>
<td>&quot;</td>
</tr>
<tr>
<td></td>
<td>Succinic dehydrogenase</td>
<td>Andreasen, Engel and Holter (1951)</td>
</tr>
<tr>
<td></td>
<td>Lipase</td>
<td>Wilber (1946)</td>
</tr>
<tr>
<td>Soil amoeba</td>
<td>&quot;Amoeboidiastase,&quot; and a trypsine-like enzyme</td>
<td>Mouton (1902)</td>
</tr>
<tr>
<td><em>Aethalium septicum</em></td>
<td>Pepsin-like enzyme</td>
<td>Krukenberg (1886)</td>
</tr>
<tr>
<td><em>Englena gracilis</em></td>
<td>Proteolytic enzyme</td>
<td>Jahn (1931)</td>
</tr>
<tr>
<td><em>Xylophagous Poly- and Hyper-mastigina</em></td>
<td>Cellulase</td>
<td>Trager (1932)</td>
</tr>
<tr>
<td></td>
<td>Cellobiase</td>
<td>Cleveland <em>et al.</em> (1934)</td>
</tr>
<tr>
<td><em>Didinium nasutum</em></td>
<td>Dipeptidase</td>
<td>Doyle and Patterson (1942)</td>
</tr>
<tr>
<td><em>Tetrahymena pyriformis</em></td>
<td>Proteolytic enzyme</td>
<td>Lwoff (1932); Llawrie (1937)</td>
</tr>
<tr>
<td></td>
<td>Peptidases</td>
<td>Kidder and Dewey (1951)</td>
</tr>
<tr>
<td></td>
<td>Acetylocholinesterase</td>
<td>Seaman and Houlihan (1951)</td>
</tr>
<tr>
<td><em>Colpidium striatum</em></td>
<td>Proteolytic enzyme</td>
<td>Elliott (1933)</td>
</tr>
<tr>
<td><em>Paramecium caudatum</em></td>
<td>Peptidase</td>
<td>Holter and Doyle (1938)</td>
</tr>
<tr>
<td></td>
<td>Amylase</td>
<td>&quot;</td>
</tr>
<tr>
<td><em>P. multimicronucleatum</em></td>
<td>Dipeptidase</td>
<td>Doyle and Patterson (1942)</td>
</tr>
<tr>
<td><em>Frontonia sp.</em></td>
<td>Peptidase</td>
<td>Holter and Doyle (1938)</td>
</tr>
<tr>
<td></td>
<td>Amylase</td>
<td>&quot;</td>
</tr>
<tr>
<td><em>Balantidium coli</em></td>
<td>Diastase</td>
<td>Glaessner (1908)</td>
</tr>
</tbody>
</table>

In certain Sarcodina such as Amoeba and Pelomyxa, refringent bodies occur conspicuously in the cytoplasm. They were first noticed in *Pelomyxa palustris* by Greeff (1874) who called them "Glanzkörper." Stolc (1900) and Leiner (1924) considered them as glycogen enclosed within a membrane and associated intimately with the
carbohydrate metabolism of the organism, since their number was proportionate to the amount of food obtained by the organism. Veley (1905) on the other hand found them albuminoid in nature. Studies of the refringent bodies in *Amoeba proteus* led Mast and Doyle (1935, 1935a) to conclude that the outer layer is composed of a protein stroma impregnated with lipid containing fatty acid, which gives positive reaction for Golgi substance; the envelope is made up of a carbohydrate which is neither starch nor glycogen; and the refringent bodies function as reserve food, since they disintegrate during starvation. The same function was assigned to those occurring in *Pelomyxa carolinensis* by Wilber (1945, 1945a), but Andresen and Holter (1945) do not agree with this view, as they observed the number of the refringent bodies (“heavy spherical bodies”) remains the same in starvation. Thus a full comprehension of the nature and function of the refringent body must depend on future observations.

The indigestible residue of the food is extruded from the body. The extrusion may take place at any point on the surface in many Sarcodina by a reverse process of the ingestion of food. But in pellicle-bearing forms, the defecation takes place either through the cytopyge located in the posterior region of the body or through an aperture to the vestibule (Fig. 37, b–d). Permanent cytopyge is lacking in some forms. In *Fabrea salina*, Kirby (1934) noticed that a large opening is formed at the posterior end, the contents of food vacuoles are discharged, and the opening closes over. At first the margin of the body is left uneven, but soon the evenly rounded outline is restored. The same seems to be the case with *Spirostomum* (Fig. 38), *Blepharisma*, etc. *Cytopyge* (Klein, 1939).

**Holophytic** (autotrophic, phytotrophic) nutrition. This is the type of nutrition in which the Protozoa are able to decompose carbon dioxide by means of chlorophyll contained in chromatophores (p. 89) in the presence of the sunlight, liberating the oxygen and combining the carbon with other elements derived from water and inorganic salts (photosynthesis). Aside from the Phytomastigina, chromatophores were definitely observed in a ciliate *Cyclotrichium meunieri* (Figs. 300, o; 301) (Powers, 1932; Bary and Stuckey, 1950). In a number of other cases, the organism itself is without chromatophores, but is apparently not holozoic, because of the presence of chlorophyll-bearing organisms within it. For example, in the testacean *Paulinella* (Fig. 206, c) in which occur no food vacuoles, chromatophores of peculiar shape are always present. The latter appear to be a species of alga which holds a symbiotic relationship with the testacean, and perhaps acts for the sarcodinan as the chromatophores
of the Phytomastigina. A similar relationship seems to exist between *Paramecium bursaria*, *Stentor polymorphus*, etc. and zoochlorellae; *Paraeuplotes tortugensis* and a zooxanthella and others (p. 29). Pringsheim (1928) showed that organic matters from zoochlorellae are passed on to their host, *Paramecium bursaria*, to be used as food. Through studies of relationships between zooxanthellae and invertebrates, Yonge observed that the zooxanthellae utilize carbon dioxide, nitrogen and phosphorus which are the catabolic products of the host and supply in return oxygen, fats and carbohydrates to the host. Photosynthesis in Phytomastigina (Hutner and Provasoli, 1951).

**Saprozoic** (saprophytic) nutrition. In this nutrition, the Protozoa obtain nourishment by diffusion through the body surface. This is accomplished without any special organellae. Perhaps the only in-

![Fig. 38. Outline sketches showing the defecation process in *Spirostomum ambiguum* (Blättner).](image)

stance in which the saprozoic nutrition is accomplished through a special organella is the pusules (Figs. 127, 129) in marine dinoflagellates which, according to Kofoid and Swezy (1921), appear to contain decomposed organic matter and aid the organisms in carrying on this process.

The dissolved food matters are simpler compounds which originate in animal or vegetable matter due to the decomposing activities of bacterial organisms. Numerous free-living flagellates nourish themselves with this method. Recently a number of investigators found that saprozoic Protozoa could be cultivated in bacteria-free media of known compositions. For example, Pringsheim (1937) observed in *Polytoma uvea* (Fig. 113, h) that sodium acetate is needed from which the starch among others is produced and carbohydrates have no direct bearing upon the nutrition, but fatty acids derived from them participate in the metabolism.

The Protozoa which live within the body of another organism are
able to nourish themselves by absorbing the digested or decomposed substances of the host and could be considered as saprozoic, though the term parasitic has sometimes been used. Coelozoic Protozoa belong to this group, as for example, Protociliata, astomatous ciliates, Trypanosomatidae, etc. In the case of cytozoic or certain histozoic forms, such as Cnidosporidia, the host cytoplasm is apparently liquefied or hydrolyzed by enzymes before being absorbed by them. The parasitic Protozoa, which actually feed on host tissue cells, such as Entamoeba histolytica, Balantidium coli, etc., or endocommensals, (Endamoeba blattae, Entamoeba coli, etc.) employ, of course, the holozoic nutrition.

Many Protozoa nourish themselves by more than one method at the same or different times, subject to a change in external conditions. This is sometimes referred to as mixotrophic nutrition (Pfeiffer). For example, Euglena gracilis, according to Zumstein (1900), Lwoff (1932) and Pringsheim and Hovasse (1948), loses its green coloration in the darkness or even in the light when the culture medium is very abundant in decomposed organic substances, which may indicate that this organism is capable of carrying on both holophytic and saprozoic nutrition.

With the introduction of bacteria-free culture technique in recent years, it has now become well established that a protozoan species exhibits conspicuous differences in form, size and structure, which are exclusively due to differences in the kind and amount of food material. For example, Kidder, Lilly and Claff (1940) noted in Tetrahymena vorax (Fig. 39), bacteria-feeders are tailed (50-75 μ long), saprozoic forms are fusiform to ovoid (30-70 μ long), forms feeding on sterile dead ciliates are fusiform (60-80 μ long), and carnivores and cannibals are irregularly ovoid (100-250 μ long), in the latter form of which a large "preparatory vacuole" becomes developed. In Chilomonas paramecium, Mast (1939) observed the individuals grown in sterile glucose-peptone solution were much smaller than those cultured in acetate-ammonium solution and moreover the former contained many small starch grains, but no fat, while the latter showed many larger starch grains and a little fat. Amoeba proteus when fed exclusively on Colpidium, became very large and extremely "fat" and sluggish, growing and multiplying slowly, but indefinitely; when fed on Chilomonas only, they grew and multiplied for several days, then decreased in number and soon died, but lived longer on Chilomonas cultured in the glucose-peptone. It is well known that Protozoa as any other organism, show atypical or abnormal morphological and physiological peculiarities. In the
case of carnivorous forms, the condition of food organisms may produce abnormalities in them, as was shown by Beers (1933) in Didinium fed on starved paramecia (Fig. 40).

Some thirty years ago, Robertson (1921–1927) reported that when two ciliates, Enchelys and Colpoda, are placed in a small amount of fresh culture medium, the rate of reproduction following a "lag pe-

Fig. 39. Form and size variation in Tetrahymena vorax, due to differences in kind and amount of food material, as seen in life, ×400 (Kidder, Lilly and Claff). a, bacteria-feeder; b, c, saprozoic forms; d, individual which has fed on killed Colpidium campylum; e, starved individual from a killed-Colpidium culture; f–i, progressive form and size changes of saprozoic form in the presence of living Colpidium; j, a young carnivore which has been removed to a culture with living yeast.
period” is more than twice (up to ten times) that of a single animal in the same amount of the medium. He assumed that this acceleration was due to a certain agent or substance produced within the animal, which diffused into the culture medium. When more than one animal is confined in a limited amount of culture fluid, this substance is present in a higher concentration than with one animal, and an increased rate of division is the result. Robertson called this “allelo-catalytic result,” and the phenomenon, “allelocatalysis.”

Fig. 40. Didinium nasutum, ×265 (Beers). a, normal fully grown animal; b–e, abnormal organisms which were fed on starved Paramecium.
Soon a large number of observers came forward with varying results—some confirmatory, others contradictory. The vast majority of these observations including Robertson's own, were carried on ciliates which were grown in association with various bacteria, and naturally, the results lacked agreement. For a review of these observations too numerous to mention here, the reader is referred to Allee (1931, 1934), Mast and Pace (1938) and Richards (1941). When bacteria-free cultivation became possible for some Protozoa, it was hoped that this problem might be solved under controlled conditions. However, the results still lack agreement. For example, Phelps (1935) reported that in Tetrahymena (Glaucoma), the growth rate and the maximum yield were the same between two cultures: one started with 0.014 organism and the other, with 1600 organisms per ml. Thus there was no allelocatalysis. On the other hand, Mast and Pace (1938) noted a significant acceleration of the growth rate in Chilomonas when up to 50 organisms were inoculated into 0.4 cc. of culture fluid as compared to the growth rate in cultures with one or more Chilomonas inocula, and furthermore, a single Chilomonas showed an increased rate of reproduction as the volume of the culture fluid was reduced.

Various aspects of metabolic processes in Protozoa such as inorganic requirements, carbon and nitrogen metabolism, growth factors, vitamins, etc., have recently been studied by a number of investigators. For information, the reader is referred to Hall (1941) and Lwoff (1951).

Reserve food matter

The anabolic activities of Protozoa result in the growth and increase in the volume of the organism, and also in the formation and storage of reserve food-substances which are deposited in the cytoplasm to be utilized later for growth or reproduction. The reserve food stuff is ordinarily glycogen or glycogenous substances, which seem to be present widely. Thus, in saprozoic Gregarinida, there occur in the cytoplasm numerous refractile bodies which stain brown to brownish-violet in Lugol's solution; are insoluble in cold water, alcohol, and ether; become swollen and later dissolved in boiling water; and are reduced to a sugar by boiling in dilute sulphuric acid. This substance which composes the refractile bodies is called paraglycogen (Bütschli) or zooamylon. Göhre (1943) considers it a stabilized polymerization product of glycogen.

Rumnjantzw and Wermel (1925) demonstrated glycogen in Actinosphaerium. In the cysts of Iodamoeba, glycogen body is con-
spicuously present and is looked upon as a characteristic feature of the organism. The iodinophile vacuole of the spores of Myxobolidae is a well-defined vacuole containing glycogenous substance and is also considered as possessing a taxonomic value. In many ciliates, both free-living (Paramecium, Glaucoma, Vorticella, Stentor, etc.) and parasitic (Ophryoscolecidae, Nyctotherus, Balantidium (Faure-Fremiet and Thaureaux, 1944)), glycogenous bodies are always present. According to MacLennan (1936), the development of the paraglycogen in Ichthyophthirius is associated with the chondriosomes. In Eimeria tenella, glycogenous substance does apparently not occur in the schizonts, merozoites, or microgametocytes; but becomes apparent first in the macrogametocyte, and increases in amount with its development, a small amount being demonstrable in the sporozoites (Edgar et al., 1944).

![Diagram of paramylon in Euglena gracilis](image)

The anabolic products of the holophytic nutrition are starch, paramylon, oil and fats. The paramylon bodies are of various forms among different species, but appear to maintain a certain characteristic form within a species and can be used to a certain extent in taxonomic consideration. According to Heidt (1937), the paramylon of Euglena sanguinea (Fig. 41) is spirally coiled which confirms Bütschli's observation. The paramylon appears to be a polysaccharide which is insoluble in boiling water, but dissolves in concentrated sulphuric acid, potassium hydroxide, and slowly in formaldehyde. It does not stain with either iodine or chlor-zinc-iodide and when treated with a dilute potassium hydroxide, the paramylon bodies become enlarged and frequently exhibit a concentric stratification.

In the Chrysomonadina, the reserve food material is in the form of refractile spheroid bodies which are known as leucosin, probably a carbohydrate which when boiled in water stains with iodine. Oil
droplets occur in various Protozoa and when there is a large number of oil-producing forms in a body of water, the water may develop various odors as indicated in Table 4.

**Table 4.—Protozoa and odors of water**

<table>
<thead>
<tr>
<th>Protozoa</th>
<th>Odor produced by them</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cryptomonas</td>
<td>candied violets</td>
</tr>
<tr>
<td>Mallomonas</td>
<td>aromatic, violets, fishy</td>
</tr>
<tr>
<td>Synura</td>
<td>ripe cucumber, muskmelon, bitter and spicy taste</td>
</tr>
<tr>
<td>Uroglenopsis</td>
<td>fishy, cod-liver oil-like</td>
</tr>
<tr>
<td>Dinobryon</td>
<td>fishy, like rockweed</td>
</tr>
<tr>
<td>Chlamydomonas</td>
<td>fishy, unpleasant or aromatic</td>
</tr>
<tr>
<td>Eudorina</td>
<td>faintly fishy</td>
</tr>
<tr>
<td>Pandorina</td>
<td>faintly fishy</td>
</tr>
<tr>
<td>Volvox</td>
<td>fishy</td>
</tr>
<tr>
<td>Ceratium</td>
<td>vile stench</td>
</tr>
<tr>
<td>Glenodinium</td>
<td>fishy</td>
</tr>
<tr>
<td>Peridinium</td>
<td>fishy, like clam-shells</td>
</tr>
<tr>
<td>Bursaria</td>
<td>Irish moss, salt marsh, fishy (Whipple, 1927)</td>
</tr>
<tr>
<td>Pelomyxa</td>
<td>ripe cucumber (Schaeffer, 1937)</td>
</tr>
</tbody>
</table>

**Fats** occur widely in Protozoa. They appear usually as small refractile globules. Zingher (1934) found that in the Sarcomeda and Ciliata he studied, each species showed morphological characteristics of the fatty substance it contained. Fat globules occur abundantly in Amoeba and Pelomyxa which are easily seen by staining with Sudan III. In *Tillina canalisfera*, fat droplets, 1–2 μ in diameter, are present especially in the region to the right of the cytopharynx (Turner, 1940). According to Panzer (1913), the fat content of *Eimeria gadi* was 3.55 per cent and Pratje (1921) reports that 12 per cent of the dry matter of *Noctiluca scintillans* appeared to be the fatty substance present in the form of granules and is said to give luminescence upon mechanical or chemical stimulation. But the chemical nature of these “photogenic” granules is still unknown at present (Harvey, 1952). A number of other dinoflagellates, such as Peridinium, Ceratium, Gonyaulax, Gymnodinium, etc., also emit luminescence. In other forms the fat may be hydrostatic in function, as is the case with a number of pelagic Radiolaria, many of which are also luminous. Luminescence in Protozoa (Harvey, 1952).

Another reserve food-stuff which occurs widely in Protozoa, excepting Ciliophora, is the so-called **volutin** or metachromatic granule. It is apparently equally widely present in Protophyta. In fact it was first discovered in the protophytyan *Spirillum volutans*. Meyer
coined the name and held it to be made up of a nucleic acid. It stains deeply with nuclear dyes. Reichenow (1909) demonstrated that if *Haematococcus plurialis* (Fig. 42) is cultivated in a phosphorus-free medium, the volutin is quickly used up and does not reappear. If however, the organisms are cultivated in a medium rich in phosphorus, the volutin increases greatly in volume and, as the culture becomes old, it gradually breaks down. In *Polytomella agilis* (Fig. 14, c, d), Doflein (1918) showed that an addition of sodium phosphate resulted in an increase of volutin. Reichenow, Schumacher, and others, hold that the volutin appears to be a free nucleic acid, and is a special reserve food material for the nuclear substance. Sassuchin (1935) studied the volutin in *Spirillum volutans* and *Sarcina flava* and found that the volutin appears during the period of strong growth, nourishment and multiplication, disappears in unfavorable condition of nourishment and gives a series of characteristic carbohydrate reactions. Sassuchin considers that the volutin is not related to the nucleus, but is a reserve food material of the cell, and is composed of glycoprotein. Volutin (Jirovec, 1926).

**Starvation.** As in all living things, when deprived of food, Protozoa perish sooner or later. The changes noticeable under the microscope are: gradual loss of cytoplasmic movement, increasing number of vacuoles and their coalescence, and finally the disintegration of the body. In starved *Pelomyxa carolinensis*, Andresen and Holter (1945) noticed the following changes: the animals disintegrate in 10–25 days at 22°C.: body volume decreases particularly during the early days of starvation and is about 20–30 per cent of the initial volume at the time of death; food vacuoles are extruded from the body in 24 to 48 hours; the cytoplasm becomes less viscous and many fluid vacuoles make their appearance; crystals and refringent bodies enclosed within vacuoles, form large groups as the vacuoles coalesce, some of which are extruded from the body; crystals and refringent bodies remain approximately constant during starvation and there

![Fig. 42. *Haematococcus plurialis*, showing the development of volutin in the medium rich in phosphorus and its disintegration in an exhausted medium, ×570 (Reichenow). a, second day; b, third day; c, fourth day; d, e, sixth day; f, eighth day.](image-url)
is no indication that they are utilized as food reserves. The ratio of reduced weight and volume and the specific gravity remain reasonably constant during starvation (Zeuthen, 1948). Andresen (1945) found starved *Amoeba proteus* to show a similar change on the whole, except that the number of chondriosomes decreased and in some cases dissolution of crystals occurred just before disintegration.

**Respiration**

In order to carry on various vital activities, the Protozoa, like all other organisms, must transform the potential energy stored in highly complex chemical compounds present in the cytoplasm, into various forms of active energy by oxidation. The oxygen involved in this process appears to be brought into contact with the substances in two ways in Protozoa. The great majority of free-living, and certain parasitic forms absorb free molecular oxygen from the surrounding media. The absorption of oxygen appears to be carried on by the permeable body surface, since there is no special organella for this purpose. The polysaprobic Protozoa are known to live in water containing no free oxygen. For example, Noland (1927) observed *Metopus es* in a pool, 6 feet in diameter and 18 inches deep, filled with dead leaves which gave a strong odor of hydrogen sulphide. The water in it showed pH 7.2 at 14°C., and contained no dissolved oxygen, 14.9 c.c. per liter of free carbon dioxide, and 78.7 c.c. per liter of fixed carbon dioxide. The parasitic Protozoa of metazoan digestive systems live also in a medium containing no molecular oxygen. All these forms appear to possess capacity of splitting complex oxygen-bearing substances present in the body to produce necessary oxygen.

Several investigators studied the influence of abundance or lack of oxygen upon different Protozoa. For example, Pütter (1905) demonstrated that several ciliates reacted differently when subjected to anaerobic condition, some perishing rapidly, others living for a considerable length of time. Death is said by Löhner to be brought about by a volume-increase due to accumulation of the waste products. When first starved for a few days and then placed in anaerobic environment, Paramecium and Colpidium died much more rapidly than unstarved individuals. Pütter, therefore, supposed that the difference in longevity of aerobic Protozoa in anaerobic conditions was correlated with that of the amount of reserve food material such as protein, glycogen and paraglycogen present in the body. Pütter further noticed that Paramecium is less affected by anaerobic condition than Spirostomum in a small amount of water, and maintained that
the smaller the size of body and the more elaborate the contractile vacuole system, the organisms suffer the less the lack of oxygen in the water, since the removal of catabolic products depends upon these factors.

The variety of habitats and results of artificial cultivations of various Protozoa indicate clearly that the oxygen requirements vary a great deal among different forms. Attempts were made in recent years to determine the oxygen requirement of Protozoa. The results of the observations are not always convincing. The oxygen consumption of Paramecium is said, according to Lund (1918) and Amberson (1928), to be fairly constant over a wide range of oxygen concentration. Specht (1934) found the measurements of the oxygen consumption and carbon dioxide production in Spirostomum ambiguum vary because of the presence of a base produced by the organism. Soule (1925) observed in the cultural tubes of Trypanosoma lewisi and Leishmania tropica, the oxygen contained in about 100 c.c. of air of the test tube is used up in about 12 and 6 days respectively. A single Paramecium caudatum is said to consume in one hour at 21°C, from 0.0052 c.c. (Kalmus) to 0.00049 c.c. (Howland and Bernstein) of oxygen. The oxygen consumption of this ciliate in heavy suspensions (3×10^3 to 301×10^3 in 3 c.c.) and associated bacteria, ranged, according to Gremsbergen and Reynaerts-De Pont (1952), from 1000 to 4000 nM^2 per hour per million individuals at 23.5°C. The two observers considered that P. caudatum possesses a typical cytochrome-oxidase system. Amoeba proteus, according to Hulpiue (1930), succumbs slowly when the amount of oxygen in water is less than 0.005 per cent and also in excess, which latter confirms Pütter's observation on Spirostomum. According to Clark (1942), a normal Amoeba proteus consumes 1.4×10^{-3} mm^3 of oxygen per hour, while an enucleated amoeba only 0.2×10^{-3} mm^3. He suggests that "the oxygen-carriers concerned with 70 per cent of the normal respiration of an amoeba are related in some way to the presence of the nucleus." In Pelomyxa carolinensis, the rate of oxygen consumption at 25°C. was found by Pace and Belda (1944) to be 0.244 ± 0.028 mm^3 per hour per mm^3 cell substance and does not differ greatly from that of Amoeba proteus and Actinosphaerium eichhorni. The temperature coefficient for the rate of respiration is nearly the same as that in Paramecium, varying from 1.7 at 15–25°C. to 2.1 at 25–35°C. Pace and Kimura (1946) further note in Pelomyxa carolinensis that carbohydrate metabolism is greater at higher than at lower temperature and that a cytochrome-cytochrome oxidase system is the mechanism chiefly involved in oxidation of carbohydrate.
The Hypermastigina of termites are killed, according to Cleve-
dland (1925), when the host animals are kept in an excess of oxygen.
Jahn found that Chilomonas paramecium in bacteria-free cultures in
heavily buffered peptone-phosphate media at pH 6.0, required for
rapid growth carbon dioxide which apparently brings about a favor-
able intracellular hydrogen-ion concentration. Respiratory metabo-
lism (Meldrum, 1934; Jahn, 1941).

Excretion and secretion

The catabolic waste material composed of water, carbon dioxide,
and nitrogenous compounds, all of which are soluble, pass out of the
body by diffusion through the surface or by means of the contractile
vacuole (p. 83). The protoplasm of the Protozoa is generally con-
sidered to possess a molecular make-up which appears to be similar
among those living in various habitats. In the freshwater Protozoa
the body of which is hypertonic to surrounding water, the water
diffuses through the body surface and so increases the water content
of the body protoplasm as to interfere with its normal function. The
contractile vacuole, which is invariably present in all freshwater
forms, is the means of getting rid of this excess water from the body.
On the other hand, marine or parasitic Protozoa live in nearly iso-
tonic media and there is no excess of water entering the body, hence
the contractile vacuoles are not found in them. Just exactly why
nearly all euciliates and suctorians possess the contractile vacuole
regardless of habitat, has not fully been explained. It is assumed that
the pellicle of the ciliate is impermeable to salts and slowly permeable
to water (Kitching, 1936) or impermeable to water, salts and prob-
bly gases (Frisch, 1937). If this is the case with all ciliates, it is not
difficult to understand the universal occurrence of the contractile
vacuole in the ciliates and suctorians.

That the elimination of excess amount of water from the body
is one of the functions of the contractile vacuole appears to be be-
yond doubt judging from the observations of Zuelzer (1907), Finley
(1930) and others, on Amoeba verrucosa which lost gradually its con-
tractile vacuole as sodium chloride was added to the water, losing
the organella completely in the seawater concentration and of Yo-
com (1934) on Paramecium caudatum and Euplotes patella, the con-
tractile vacuoles of which nearly ceased functioning when the ani-
mal's were placed in 10 per cent sea water. Furthermore, marine
amoebae develop contractile vacuoles de novo when they are trans-
planted to fresh water as in the case of Vahlkampfia calkinsi (Hogue,
1923) and Amoeba biddulphiae (Zuelzer, 1927). Herfs (1922) studied
the pulsation of the contractile vacuoles of *Paramecium caudatum* in fresh water as well as in salt water and obtained the following measurements:

<table>
<thead>
<tr>
<th>Per cent NaCl in water</th>
<th>0</th>
<th>0.25</th>
<th>0.5</th>
<th>0.75</th>
<th>1.00</th>
</tr>
</thead>
<tbody>
<tr>
<td>Contraction period in second</td>
<td>6.2</td>
<td>9.3</td>
<td>18.4</td>
<td>24.8</td>
<td>163.0</td>
</tr>
<tr>
<td>Excretion per hour in body volumes</td>
<td>4.8</td>
<td>2.82</td>
<td>1.38</td>
<td>1.08</td>
<td>0.16</td>
</tr>
</tbody>
</table>

The number of the contractile vacuoles present in a species is constant under normal conditions. The contraction period varies from a few seconds to several minutes in freshwater inhabitants, and is, as a rule, considerably longer in marine Protozoa. Kitching (1938a) estimated that a quantity of water equivalent to the body volume is eliminated by freshwater Protozoa in four to 45 minutes and by marine forms in about three to four hours. The size of contractile vacuole in diastole may vary. Botsford (1926) reported that the contractile vacuole in *Amoeba proteus* varied considerably within a short period of time in size and rate of contraction under seemingly identical conditions. The rate of contraction is subject to change with the temperature, physiological state of the organism, amount of food substances, etc. For example, Rossbach noted in the three ciliates listed below, the contraction was accelerated first rapidly and then more slowly with rise of the temperature:

| Time in seconds between two systoles at different temperature (°C) |
|------------------------|---|---|---|---|---|---|
| Euplotes charon         | 61 | 48 | 31 | 28 | 22 | 23 |
| Stylonychia pustulata   | 18 | 14 | 10-11 | 6-8 | 5-6 | 4 |
| Chilodonella cucullulus| 9  | 7  | 5  | 4  | 4  | — |

How much water enters through the body surface of Protozoa is not known, but it appears to be the major portion that is excreted through contractile vacuoles. Water also enters the protozoan body in food vacuoles. In *Vampyrella lateritia* which feeds on the cell contents of Spirogyra in a single feeding, many contractile vacuoles appear within the cytoplasm and evacuate the water that has come in with the food (Lloyd, 1926) and the members of Ophryoscolecidae show an increased number and activity of contractile vacuoles while feeding (MacLennan, 1933). The amount of water contained in food vacuoles seems, however, to be far smaller than the amount evacuated by contractile vacuoles (Gelei, 1925; Eisenberg, 1925). Other evidences such as the contractile vacuole continues to pulsate when cytosome-bearing Protozoa are not feeding and its occurrence in astomatous ciliates, would indicate also that the water entering
through this avenue is not of a large quantity. How much water is produced during the metabolic activity of the organisms is unknown, but it is considered to be a very small amount (Kitching, 1938). The mechanism by which the difference in osmotic pressure can be maintained at the body surface is unknown. It may be, as suggested by Kitching (1934), that the contractile vacuole extrudes water but retains the solutes or some osmotically active substances must be continuously produced within the body.

Attempts to detect catabolic products in the contractile vacuole, in the body protoplasm or in the culture fluid, were unsuccessful, because of technical difficulties. Weatherby (1927) detected in the spring water in which he kept a number of thoroughly washed Paramecium, urea and ammonia after 30–36 hours and supposed that the urea excreted by the organisms gave rise to ammonia. He found also urea in similar experiments with Spirostomum and Didinium (Weatherby, 1929). Doyle and Harding (1937) found Glaucoma excreting ammonia, and not urea. Carbon dioxide is obviously excreted by the body surface as well as the contractile vacuole. At present the composition of the fluid in the contractile vacuole is not known. General reference (Weatherby, 1941); permeability of water in Protozoa (Belda, 1942; Løvtrup and Pigón, 1951); physiology of contractile vacuole (Stempell, 1924; Fortner, 1926; Gaw, 1936; Kitching, 1938a).

Aside from the soluble forms, there often occur in the protozoan body insoluble substances in the forms of crystals and granules of various kinds. Schewiakoff (1894) first noticed that Paramecium often contained crystals (Fig. 43) composed of calcium phosphate, which disappeared completely in 1–2 days when the organisms were starved, and reappeared when food was given. Schewiakoff did not see the extrusion of these crystals, but considered that these crystals

![Fig. 43. Examples of crystals present in Protozoa. a–e, in Paramecium caudatum (Schewiakoff), (a–d, ×1000, e, ×2600); f, in Amoeba proteus; g, in A. discoides; h–l, in A. dubia (Schaeffer).](image-url)
were first dissolved and excreted by the contractile vacuoles, as they were seen collected around the vacuoles. When exposed to X-irradiation, the symbiotic Chlorella of Paramecium bursaria disappear gradually and crystals appear and persist in the cytoplasm of the ciliate (Wichterman, 1948a). These crystals varying in size from a few to 12μ, are found mainly in the posterior region of the body. Wichterman notes that the appearance or disappearance of crystals seems to be correlated with the absence or presence of symbiotic Chlorella and with the holozoic or holophytic (by the alga) nutrition of the organism.

In Amoeba proteus, Schubotz (1905) noted crystals of calcium phosphate which were bipyramidal or rhombic in form, were doubly refractile and measured about 2–5μ in length. In three species of Amoeba, Schaeffer (1920) points out the different shape, number and dimensions of the crystals. Thus in Amoeba proteus, they are truncate bipyramids, rarely flat plates, up to 4.5μ long; in A. discoides, abundant, truncate bipyramids, up to 2.5μ long; and in A. dubia, variously shaped (4 kinds), few, but large, up to 10μ, 12μ, 30μ long (Fig. 43). Bipyramidal or plate-like crystals are especially abundant in Pelomyxa illinoisensis at all times (Kudo, 1951); the crystals of P. carolinensis remain the same during the starvation of the organism (Andresen and Holter, 1945; Holter, 1950).

The crystals present in Protozoa appear to be of varied chemical nature. Luce and Pohl (1935) noticed that at certain times amoeboae in culture are clear and contain relatively a few crystals but, as the culture grows older and the water becomes more neutral, the crystals become abundant and the organisms become opaque in transmitted light. These crystals are tubular and six-sided, and vary in length from 0.5 to 3.5μ. They considered the crystals were composed of calcium chlorophosphate. Mast and Doyle (1935), on the other hand, noted in Amoeba proteus two kinds of crystals, plate-like and bipyramidal, which vary in size up to 7μ in length and which are suspended in alkaline fluid to viscid vacuoles. These two authors believed that the plate-like crystals are probably leucine, while the bipyramidal crystals consist of a magnesium salt of a substituted glycine. Other crystals are said to be composed of urate, carbonate, oxalate, etc.

Another catabolic product is the haemozoin (melanin) grains which occur in many haemosporidians and which appear to be composed of a derivative of the haemoglobin of the infected erythrocyte (p. 605). In certain Radiolaria, there occurs a brownish amorphous mass which is considered as catabolic waste material and, in Foram-
Protozoa move about by means of the pseudopodia, flagella, or cilia, which may be combined with internal contractile organelles.

**Movement by pseudopodia.** Amoeboid movements have long been studied by numerous observers. The first attempt to explain the movement was made by Berthold (1886), who held that the difference in the surface tension was the cause of amoeboid movements, which view was supported by the observations and experiments of Bütschli (1894) and Rhumbler (1898). According to this view, when an amoeba forms a pseudopodium, there probably occurs a diminution of the surface tension of the cytoplasm at that point, due to certain internal changes which are continuously going on within the body and possibly due to external causes, and the internal pressure of the cytoplasm will then cause the streaming of the cytoplasm. This results in the formation of a pseudopodium which becomes attached to the substratum and an increase in tension of the plasma-membrane draws up the posterior end of the amoeba, thus bringing about the movement of the whole body.

Jennings (1904) found that the movement of *Amoeba verrucosa* (Fig. 44, a) could not be explained by the surface tension theory, since he observed "in an advancing amoeba substance flows forward on the upper surface, rolls over at the anterior edge, coming in contact with the substratum, then remains quiet until the body of the amoeba has passed over it. It then moves upward at the posterior end, and forward again on the upper surface, continuing in rotation as long as the amoeba continues to progress." Thus *Amoeba verrucosa* may be compared with an elastic sac filled with fluid. Dellinger (1906) studied the movement of *Amoeba proteus, A. verrucosa* and *Difflugia spiralis*. Studying in side view, he found that the amoeba (Fig. 45) extends a pseudopod, "swings it about,
brings it into the line of advance, and attaches it” to the substratum and that there is then a concentration of the substance back of this point and a flow of the substance toward the anterior end. Dellinger held thus that “the movements of amoebae are due to the presence

![Diagram](image1)

**Fig. 44.** a, diagram showing the movement of *Amoeba verrucosa* in side view (Jennings); b, a marine limax-amoeba in locomotion (Pantin from Reichenow). ac, area of conversion; cet, contracting ectoplasmic tube; fe, fluid ectoplasm; ge, gelated ectoplasm.

of a contractile substance,” which was said to be located in the endoplasm as a coarse reticulum. Wilber (1946) pointed out that *Pelmomyxa carolinensis* carries on a similar movement at times.

![Diagram](image2)

**Fig. 45.** Outline sketches of photomicrographs of *Amoeba proteus* during locomotion, as viewed from side (Dellinger).

In the face of advancement of our knowledge on the nature of protoplasm, Rhumbler (1910) realized the difficulties of the surface tension theory and later suggested that the conversion of the ectoplasm to endoplasm and vice versa were the cause of the cytoplasmic
movements, which was much extended by Hyman (1917). Hyman considered that: (1) a gradient in susceptibility to potassium cyanide exists in each pseudopodium, being the greatest at the distal end, and the most recent pseudopodium, the most susceptible; (2) the susceptibility gradient (or metabolic gradient) arises in the amoebae before the pseudopodium appears and hence the metabolic change which produces increased susceptibility, is the primary cause of pseudopodium formation; and (3) since the surface is in a state of gelation, amoeboid movement must be due to alterations of the colloidal state. Solution, which is brought about by the metabolic change, is regarded as the cause of the extension of a pseudopodium, and gelation, of the withdrawal of pseudopodia and of active contraction. Schaeffer (1920) mentioned the importance of the surface layer which is a true surface tension film, the ectoplasm, and the streaming of endoplasm in the amoeboid movement.

Pantin (1923) studied a marine limax-type amoeba (Fig. 44, b) and came to recognize acid secretion and absorption of water at the place where the pseudopodium was formed. This results in swelling of the cytoplasm and the pseudopodium is formed. Because of the acidity, the surface tension increases and to lower or reduce this, concentration of substances in the "wall" of the pseudopodium follows. This leads to the formation of a gelatinous ectoplasmic tube which, as the pseudopodium extends, moves toward the posterior region where the acid condition is lost, gives up water and contracts finally becoming transformed into endoplasm near the posterior end. The contraction of the ectoplasmic tube forces the endoplasmic streaming to the front.

This observation is in agreement with that of Mast (1923, 1926, 1931) who after a series of carefully conducted observations on Amoeba proteus came to hold that the amoeboid movement is brought about by "four primary processes; namely, attachment to the substratum, gelation of plasmasol at the anterior end, solution of plasmagel at the posterior end and the contraction of the plasmagel at the posterior end" (Fig. 46). As to how these processes work, Mast states: "The gelation of the plasmasol at the anterior end extends ordinarily the plasmagel tube forward as rapidly as it is broken down at the posterior end by solation and the contraction of the plasmagel tube at the posterior end drives the plasmasol forward. The plasmagel tube is sometimes open at the anterior end and the plasmasol extends forward and comes in contact with the plasma-lemma at this end (Fig. 47, a), but at other times it is closed by a thin sheet of gel which prevents the plasmasol from reaching the
Fig. 46. Diagram of *Amoeba proteus*, showing the solution and gelation of the cytoplasm during ameboid movement (Mast). c, crystal; cv, contractile vacuole; f, food vacuole; hc, hyaline cap; n, nucleus; pg, plasmagel; pgs, plasmagel sheet; pl, plasmalemma; ps, plasmasol.
anterior end (b). This gel sheet at times persists intact for considerable periods, being built up by gelation as rapidly as it is broken down by stretching, owing to the pressure of the plasmagel against it. Usually it breaks periodically at various places. Sometimes the breaks are small and only a few granules of plasmasol pass through and these gelate immediately and close the openings (d). At other times the breaks are large and plasmasol streams through, filling the hyaline cap (c), after which the sol adjoining the plasmalemma gelates forming a new gel sheet. An amoeba is a turgid system, and the plasmagel is under continuous tension. The plasmagel is elastic and, consequently, is pushed out at the region where its elasticity is weakest and this results in pseudopodial formation. When an amoeba is elongated and undergoing movement, the elastic strength of the plasmagel is the highest at its sides, lowest at the anterior end and intermediate at the posterior end, which results in continuity of the elongated form and in extension of the anterior end. If pressure is brought against the anterior end, the direction of streaming of plasmasol is immediately reversed, and a new hyaline cap is formed at the posterior end which is thus changed into a new anterior end."

The rate of amoeboid locomotion appears to be influenced by environmental factors such as pH, osmotic pressure, salt concentration, substratum, temperature, etc. (Mast and Prosser, 1932).

Flagellar movement. The flagellar movement is in a few instances observable as in Peranema, but in most cases it is very difficult to observe in life. Since there is difference in the number, location, size, and probably structure (p. 53) of flagella occurring in Protozoa, it is supposed that there are varieties of flagellar movements. The first explanation was advanced by Bütschli, who observed that the flagel-

![Fig. 47. Diagrams of varied cytoplasmic movements at the tip of a pseudopodium in *Amoeba proteus* (Mast). g, plasmagel; he, hyaline cap; hl, hyaline layer; pl, plasmalemma; s, plasmasol.](image-url)
lum undergoes a series of lateral movements and, in so doing, a pressure is exerted on the water at right angles to its surface. This pressure can be resolved into two forces: one directed parallel, and the other at right angles, to the main body axis. The former will drive the organism forward, while the latter will tend to rotate the animal on its own axis.

Gray (1928), who gave an excellent account of the movement of flagella, points out that “in order to produce propulsion there must be a force which is always applied to the water in the same direction and which is independent of the phase of lateral movement. There can be little doubt that this condition is satisfied in flagellated organisms not because each particle of the flagellum is moving laterally to and fro, but by the transmission of the waves from one end of the flagellum to the other, and because the direction of the transmission is always the same. A stationary wave, as apparently contemplated by Bütschli, could not effect propulsion since the forces acting on the water are equal and opposite during the two phases of the movement. If however the waves are being transmitted in one direction only, definite propulsive forces are present which always act in a direction opposite to that of the waves.”

Because of the nature of the flagellar movement, the actual process has often not been observed. Verworn observed long ago that in Peranema trichophorum the undulation of the distal portion of flagellum is accompanied by a slow forward movement, while undulation along the entire length is followed by a rapid forward movement. Krijgsman (1925) studied the movements of the long flagellum of Monas sp. (Fig. 48) which he found in soil cultures, under the dark-field microscope and stated: (1) when the organism moves forward with the maximum speed, the flagellum starting from c 1, with the wave beginning at the base, stretches back (c 1-6), and then waves back (d, e), which brings about the forward movement. Another type is one in which the flagellum bends back beginning at its base (f) until it coincides with the body axis, and in its effective stroke waves back as a more or less rigid structure (g); (2) when the organism moves forward with moderate speed, the tip of the flagellum passes through 45° or less (h-j); (3) when the animal moves backward, the flagellum undergoes undulation which begins at its base (k-o); (4) when the animal moves to one side, the flagellum becomes bent at right angles to the body and undulation passes along it from its base to tip (p); and (5) when the organism undergoes a slight lateral movement, only the distal end of the flagellum undulates (q).

Ciliary movement. The cilia are the locomotor organella present
permanently in the ciliates and vary in size and distribution among different species. Just as flagellates show various types of movements, so do the ciliates, though nearly all free-swimming forms swim in a spiral path (Bullington, 1925, 1930). Individual cilium on a

Fig. 48. Diagrams illustrating flagellar movements of Monas sp. (Krijgsman). a–g, rapid forward movement (a, b, optical image of the movement in front and side view; c, preparatory and d, e, effective stroke; f, preparatory and g, effective stroke); h–j, moderate forward movement (h, optical image; i, preparatory and j, effective stroke); k–o, undulatory movement of the flagellum in backward movement; p, lateral movement; q, turning movement.
progressing ciliate bends throughout its length and strikes the water so that the organism tends to move in a direction opposite to that of the effective beat, while the water moves in the direction of the beat (Fig. 49, a–d). In the Protociliata and the majority of holotrichous and heterotrichous ciliates, the cilia are arranged in longitudinal, or oblique rows and it is clearly noticeable that the cilia are not beating in the same phase, although they are moving at the same rate. A

![Diagram](https://via.placeholder.com/150)

**Fig. 49.** Diagrams illustrating ciliary movements (Verworn). a–d, movement of a marginal cilium of *Urostyla grandis* (a, preparatory and b, effective stroke, resulting in rapid movement; c, preparatory, and d, effective stroke, bringing about moderate speed); e, metachronous movements of cilia in a longitudinal row.

cilium (Fig. 49, e) in a single row is slightly in advance of the cilium behind it and slightly behind the one just in front of it, thus the cilia on the same longitudinal row beat metachronously. On the other hand, the cilia on the same transverse row beat synchronously, the condition clearly being recognizable on *Opalina* among others, which is much like the waves passing over a wheat field on a windy day. The organized movements of cilia, cirri, membranellae and undulating membranes are probably controlled by the neuromotor system (p. 63) which appears to be conductile as judged by the results of micro-dissection experiments of Taylor (p. 65). Ciliary movement (Gray, 1928); spiral movement of ciliates (Bullington, 1925, 1930); movement of *Paramecium* (Dembowski, 1923, 1929a) and of *Spirostomum* (Blättner, 1926).

The Protozoa which possess myonemes are able to move by con-
traction of the body or of the stalk, and others combine this with the secretion of mucous substance as is found in Haemogregarina and Gregarinida.

**Irritability**

Under natural conditions, the Protozoa do not behave always in the same manner, because several stimuli act upon them usually in combination and predominating stimulus or stimuli vary under different circumstances. Many investigators have, up to the present time, studied the reactions of various Protozoa to external stimulations, full discussion of which is beyond the scope of the present work. Here one or two examples in connection with the reactions to each of the various stimuli only will be mentioned. Of various responses expressed by a protozoan against a stimulus such as changes in body form, movement, structure, behavior, etc., the movement is the most clearly recognizable one and, therefore, free-swimming forms, particularly ciliates, have been the favorite objects of study. We consider the reaction to a stimulus in protozoans as the movement response, and this appears in one of the two directions: namely, toward, or away from, the source of the stimulus. Here we speak of positive or negative reaction. In forms such as Amoeba, the external stimulation is first received by the body surface and then by the whole protoplasmic body. In flagellated or ciliated Protozoa, the flagella or cilia act in part sensory; in fact in a number of ciliates are found non-vibratile cilia which appear to be sensory in function. In a comparatively small number of forms, there are sensory organellae such as stigma, ocellus, statocysts, concretion vacuoles, etc.

In general, the reaction of a protozoan to any external stimulus depends upon its intensity so that a certain chemical substance may bring about entirely opposite reactions on the part of the protozoans in different concentrations and, even under identical conditions, different individuals of a given species may react differently. Irritability (Jennings, 1906; Mast, 1941); in Spirostomum (Blättner, 1926).

**Reaction to mechanical stimuli.** One of the most common stimuli a protozoan would encounter in the natural habitat is that which comes from contact with a solid object. When an amoeba which Jennings observed, came in contact with the end of a dead algal filament at the middle of its anterior surface (Fig. 50, a), the amoeboid movements proceeded on both sides of the filament (b), but soon motion ceased on one side, while it continued on the other, and
the organism avoided the obstacle by reversing a part of the current and flowing in another direction (c). When an amoeba is stimulated mechanically by the tip of a glass rod (d), it turns away from the side touched, by changing endoplasmic streaming and forming new pseudopodia (e). Positive reactions are also often noted, when a suspended amoeba (f) comes in contact with a solid surface with the tip of a pseudopodium, the latter adheres to it by spreading out (g). Streaming of the cytoplasm follows and it becomes a creeping form (h). Positive reactions toward solid bodies account of course for the ingestion of food particles.

In Paramecium, according to Jennings, the anterior end is more sensitive than any other parts, and while swimming, if it comes in contact with a solid object, the response may be either negative or positive. In the former case, avoiding movement (Fig. 51, c) follows and in the latter case, the organism rests with its anterior end or the whole side in direct contact with the object, in which position it ingests food particles through the cytostome.

**Reaction to gravity.** The reaction to gravity varies among different Protozoa, according to body organization, locomotor organelles, etc. Amoebae, Testacea and others which are usually found attached to the bottom of the container, react as a rule positively
toward gravity, while others manifest negative reaction as in the case of Paramecium (Jensen; Jennings), which explains in part why Paramecium in a culture jar are found just below the surface film in mass, although the vertical movement of *P. caudatum* is undoubtedly influenced by various factors (Koehler, 1922, 1930; Dembowski, 1923, 1929, 1929a; Merton, 1935).

**Reaction to current.** Free-swimming Protozoa appear to move or orientate themselves against the current of water. In the case of

![Diagram](image)

**Fig. 51. Reactions of Paramecium (Jennings).** a, collecting in a drop of 0.02% acetic acid; b, ring-formation around a drop of a stronger solution of the acid; c, avoiding reaction.

Paramecium, Jennings observed the majority place themselves in line with the current, with anterior end upstream. The mycetozoa is said to exhibit also a well-marked positive reaction.

**Reaction to chemical stimuli.** When methylgreen, methylene blue, or sodium chloride is brought in contact with an advancing amoeba, the latter organism reacts negatively (Jennings). Jennings further observed various reactions of Paramecium against chemical stimulation. This ciliate shows positive reaction to weak solutions of many acids and negative reactions above certain concentrations. For example, Paramecium enters and stays within the
area of a drop of 0.02 per cent acetic acid introduced to the preparation (Fig. 51, a); and if stronger acid is used, the organisms collect about its periphery where the acid is diluted by the surrounding water (b). The reaction to chemical stimuli is probably of the greatest importance for the existence of Protozoa, since it leads them to proper food substances, the ingestion of which is the foundation of metabolic activities. In the case of parasitic Protozoa, possibly the reaction to chemical stimuli results in their finding specific host animals and their distribution in different organs and tissues within the host body. Recent investigations tend to indicate that chemotaxis plays an important rôle in the sexual reproduction in Protozoa. Chemotaxis in Peranema (Chen, 1950).

**Reaction to light stimuli.** Most Protozoa seem to be indifferent to the ordinary light, but when the light intensity is suddenly increased, there is usually a negative reaction. Verworn saw the direction of movements of an amoeba reversed when its anterior end was subjected to a sudden illumination; Rhumbler observed that an amoeba, which was in the act of feeding, stopped feeding when it was subjected to strong light. According to Mast, *Amoeba proteus* ceases to move when suddenly strongly illuminated, but continues to move if the increase in intensity is gradual and if the illumination remains constant, the amoeba begins to move. *Pelomyxa carolinensis* reacts negatively to light (Kudo, 1946).

The positive reaction to light is most clearly shown in stigma-bearing Mastigophora, as is well observable in a jar containing Euglena, Phacus, etc., in which the organisms collect at the place where the light is strongest. If the light is excluded completely, the organisms become scattered throughout the container, inactive and sometimes encyst, although the mixotrophic forms would continue activities by saprozoic method. The positive reaction to light by chromatophore-bearing forms enables them to find places in the water where photosynthesis can be carried on to the maximum degree.

All Protozoa seem to be more sensitive to ultraviolet rays. Inman found that amoeba shows a greater reaction to the rays than others and Hertel observed that Paramecium which was indifferent to an ordinary light, showed an immediate response (negative reaction) to the rays. MacDougall brought about mutations in Chilodonella by means of these rays (p. 229). Horváth (1950) exposed *Kahlia simplex* to ultraviolet rays and destroyed the micronucleus. The emicronucleate individuals lived and showed a greater vitality than normal individuals, as judged by the division rate at 34°C. Mazia and
Hirshfield (1951) subjected *Amoeba proteus* to ultraviolet radiation and noticed that irradiation of the whole and nucleated half amoebae delays division immediately following exposure; later progeny of the irradiated amoebae have a normal division rate; amputation of half of the cytoplasm greatly increases the radiation sensitivity as measured by delayed division or by the dose required for permanent inhibition of division (sterilization dose); individuals that have received this dose may survive for 20–30 days; and the survival time of an enucleate fragment is very much reduced by small (200–500 ergs/sq. mm) doses. The two workers consider that the overall radiation effect may have both nuclear and cytoplasmic components. By exposing *Pelomyxa carolinensis* to 2537 Å ultraviolet irradiation, Wilber and Slane (1951) found the effects variable; however, all recovered from a two minutes’ exposure, none survived a 10-minute exposure, and 70 per cent of fat were released after two minutes’ exposure.

Zuelzer (1905) found the effect of radium rays upon various Protozoa vary; in all cases, a long exposure was fatal to Protozoa, the first effect of exposure being shown by accelerated movement. Halberstaedter and Luntz (1929, 1930) studied injuries and death of *Eudorina elegans* by exposure to radium rays. *Entamoeba histolytica* in culture when subjected to radium rays, Nasset and Kofoid (1928) noticed the following changes: the division rate rose two to four times by the exposure, which effect continued for not more than 24 hours after the removal of the radium and was followed by a retardation of the rate; radium exposure produced changes in nuclear structure, increase in size, enucleation or autotomy, which were more striking when a larger amount of radium was used for a short time than a smaller amount acting on for a long time; and the effects persisted for four to six days after the removal of the radium and then the culture gradually returned to normalcy. Halberstaedter (1914) reported that when exposed to Beta rays, *Trypanosoma brucei* lost its infectivity, though remained alive.

Halberstaedter (1938) exposed *Trypanosoma gambiense* to X-rays and found that 12,000r rendered the organisms not infectious for mice, while 600,000r was needed to kill the flagellates. Emmett (1950) exposed *T. cruzi* to X-rays and noticed that dosages between 51,000r and 100,000r were necessary to destroy the infectivity of this trypanosome; the cultures, after exposure to 100,000r, appeared to be thriving up to three months; and the effects of exposure were not passed on to new generations.

When *Paramecium bursaria* were exposed to X-rays, Wichterman
(1948) noted: dosages higher than 100,000r retard the locomotion of the ciliate; none survives 700,000r; the symbiotic Chlorella is destroyed by exposure to 300,000–600,000r; irradiation inhibits division temporarily, but the animals recover normal division rate after certain length of time; and mating types are not destroyed, though minor changes occur. In Pelomyxa carolinensis, Daniels (1951) observed: the median lethal dose of X-rays is 96,000r; with dosages 15,000–140,000r, the first plasmotomy is greatly delayed and the second plasmotomy is also somewhat delayed, but later plasmotomies show complete recovery; X-irradiation does not change the type of plasmotomy; and in individuals formed by plasmogamy of X-irradiated halves to non-irradiated halves, the nuclei divide simultaneously as in a normal individual.

Reaction to temperature stimuli. As was stated before, there seems to be an optimum temperature range for each protozoan, although it can withstand temperatures which are lower or higher than that range. As a general rule, the higher the temperature, the greater the metabolic activities, and the latter condition results in turn in a more rapid growth and more frequent reproduction. It has been suggested that change to different phases in the life-cycle of a protozoan in association with the seasonal change may be largely due to temperature changes of the environment. In the case of parasitic Protozoa which inhabit two hosts: warm-blooded and cold-blooded animals, such as Plasmodium and Leishmania, the difference in body temperature of host animals may bring about specific stages in their development.

Reaction to electrical stimuli. Since Verworn’s experiments, several investigators studied the effects of electric current which is passed through Protozoa in water. Amoeba shows negative reaction to the anode and moves toward the cathode either by reversing the cytoplasmic streaming (Verworn) or by turning around the body (Jennings). The free-swimming ciliates move mostly toward the cathode, but a few may take a transverse position (Spirostomum) or swim to the anode (Paramecium, Stentor, etc.). Of flagellates, Verworn noticed that Trachelomonas and Peridinium moved to the cathode, while Chilomonas, Cryptomonas, and Polytomella, swam to the anode. When Paramecium caudatum was exposed to a high-frequency electrostatic or electromagnetic field, Kahler, Chalkley and Voegtltn (1929) found the effect was primarily caused by a temperature increase in the organism. By subjecting Pelomyxa carolinensis to a direct current electric field, Daniel and May (1950) noted that the time required for the rupture of the body in a given current
density is directly correlated with the size of the organism and that calcium increases the time required for rupture at a fixed body size and current density, but does not alter the size effect. Galvanotaxis of Oxytricha (Luntz, 1935), of Arcella (Miller, 1932).

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Chapter 5

Reproduction

The mode of reproduction in Protozoa is highly variable among different groups, although it is primarily a cell division. The reproduction is initiated by the nuclear division in nearly all cases, which will therefore be considered first.

Nuclear division

Between a simple direct division on the one hand and a complicated indirect division which is comparable with the typical metazoan mitosis on the other hand, all types of nuclear division occur.

Direct nuclear division. Although not so widely found as it was thought to be in former years, amitosis occurs normally and regularly in many forms. While the micronuclear division of the Ciliophora is mitotic (p. 165), the macronuclear division is invariably amitosis. The sole exception to this general statement appears to be the so-called promitosis reported by Ivanić (1938) in the macronucleus in the "Vermehrungsruhe" stage of Chilodonella uncinata in which chromosomes and spindle-fibers were observed. In Paramecium caudatum (Fig. 52), the micronucleus initiates the division by mitosis and the macronucleus elongates itself without any visible changes in its internal structure. The elongated nucleus becomes constricted through the middle and two daughter nuclei are produced.

It is assumed that the nuclear components undergo solution during division, since the formed particles of nucleus which are stationary in the resting stage manifest a very active Brownian movement. Furthermore, in some cases the nuclear components may undergo phase reversal, that is to say, the chromatin granules which are dispersed phase in the non-staining fluid dispersion medium in the resting nucleus, become dispersion medium in which the latter is suspended as dispersed phase. By using Feulgen's nucleal reaction, Reichenow (1928) demonstrated this reversal phenomenon in the division of the macronucleus of Chilodonella cucullulus (Fig. 53).

The macronucleus becomes at the time of its division somewhat enlarged and its chromatin granules are more deeply stained than before. But chromosomes which characterize the mitotic division are entirely absent, although in a few forms in which mating types occur, the type difference and certain other characters, according to
Sonneborn and Kimball, appear to be under control of genic constituents of the macronucleus. Since the number of chromatin granules appear approximately the same in the macronuclei of different generations of a given species, the reduced number of chromatin granules must be restored sometime before the next division takes place. Calkins (1926) is of the opinion that "each granule elongates and divides into two parts, thus doubling the number of chromomerces." Reichenow (1928) found that in Chilodonella cucullulus the lightly Feulgen positive endosome appeared to form chromatin granules and Kudo (1936) maintained that the large chromatin spherules of

Fig. 52. Nuclear and cytoplasmic division of Paramecium caudatum as seen in stained smears, ×260 (Kudo).
the macronucleus of *Nyctotherus ovalis* probably produce smaller spherules in their alveoli (Fig. 3).

When the macronucleus is elongated as in Spirostomum, Stentor, Euplotes, etc., the nucleus becomes condensed into a rounded form prior to its division. During the "shortening period" of the elongated macronuclei prior to division, there appear 1–3 characteristic zones which have been called by various names, such as nuclear clefts, reconstruction bands, reorganization bands, etc. In *Euplotes patella* (E. *eurystomus*), Turner (1930) observed prior to division of the macronucleus a reorganization band consisting of a faintly staining zone ("reconstruction plane") and a deeply staining zone ("solution plane"), appears at each end of the nucleus (Fig. 54, a) and as each moves toward the center, a more chromatinic area is left behind (b–d). The two bands finally meet in the center and the nucleus assumes an ovoid form. This is followed by a simple division into two. In the T-shaped macronucleus of *E. woodruffii*, according to Pierson (1943), a reorganization band appears first in the right arm and the posterior tip of the stem of the nucleus. When the anterior band reaches the junction of the arm and stem, it splits into two, one part

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**Fig. 53.** The solution of chromatin during the macronuclear division of *Chilodonella cucullulus*, as demonstrated by Feulgen's nucleal reaction, ×1800 (Reichenow).
moving along the left arm to its tip, and the other entering and passing down the stem to join the posterior band. According to Summers (1935) a process similar to that of *E. eurystomus* occurs in *Diophrys appendiculata* and *Stylonychia pustulata*; but in *Aspidisca lynceus* (Fig. 55) a reorganization band appears first near the middle region of the macronucleus (*b*), divides into two and each moves toward an end, leaving between them a greater chromatinic content of the reticulum (*c–i*). Summers suggested that "the reorganization bands are local regions of karyolysis and resynthesis of macronuclear materials with the possibility of an elimination of physically or possibly chemically modified nonstaining substances into the cytoplasm." Weisz (1950a) finds that the nodes of the moniliform macronucleus of *Stentor coeruleus* contain different concentration of thymonucleic acid which is correlated with morphogenetic activity of individual nodes, and that fusion of ill-staining nodes results in a return of strong affinity to methyl green. It appears, therefore, concentration of bandform or moniliform macronucleus prior to division may serve to recover morphogenetic potential prior to division.

In a small number of ciliates, the macronucleus is distributed as small bodies throughout the cytoplasm. In *Urostyla grandis*, the macronuclear material is lodged in 100 or more small bodies scattered in the cytoplasm. Prior to fission, all macronuclear bodies fuse with one another and form one macronucleus which then divides three times into eight and the latter are evenly distributed between the two daughter individuals, followed by divisions until the number reaches 100 or more (Raabe, 1947). On the other hand, in *Dileptus*

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Fig. 54. Macronuclear reorganization before division in *Euplotes eurystomus*, ×240 (Turner). a, reorganization band appearing at a tip of the macronucleus; b–d, later stages.
anser (Fig. 310, c), "each granule divides where it happens to be and with the majority of granules both halves remain in one daughter cell after division" (Calkins). Hayes noticed a similar division, but at the time of simultaneous division prior to cell division, each macronucleus becomes elongated and breaks into several small nuclei.

Fig. 55. Macronuclear reorganization prior to division in *Aspidisca lyncus*, ×1400 (Summers). a, resting nucleus; b–i, successive stages in reorganization process; j, a daughter macronucleus shortly after division.

The extrusion of a certain portion of the macronuclear material during division has been observed in a number of species. In *Uroleptus halseyi*, Calkins actually noticed each of the eight macronuclei is "purified" by discarding a reorganization band and an "x-body" into the cytoplasm before fusing into a single macronucleus which then divides into two nuclei. In the more or less rounded macronucleus that is commonly found in many ciliates, no reorganization band has been recognized. A number of observers have however noted
that during the nuclear division there appears and persists a small body within the nuclear figure, located at the division plane as in the case of Loxocephalus (Behrend), Eupoterion (MacLennan and Connell) and even in the widely different protozoan, *Endamoeba blatiae* (Kudo, 1926). Kidder (1933) observed that during the division of the macronucleus of *Conchophthirus mytili* (Fig. 56), the nucleus "casts out a part of its chromatin at every vegetative division," which "is broken down and disappears in the cytoplasm of either daughter organism." A similar phenomenon has since been found further in *C. anodontae*, *C. curtus*, *C. magna* (Kidder), *Urocentrum turbo*, *Colpidium colpoda*, *C. campylum*, *Glaucoma scintillans* (Kidder and Diller), *Allosphaerium convexa* (Kidder and Summers), *Colpoda inflata*, *C. maupasi*, *Tillina canalisfera*, *Breslaua vorax*, etc. (Burt et al., 1941). Beers (1946) noted chromatin extrusion from the macronucleus during division and in permanent cysts in *Tillina magna*. What is the significance of this phenomenon? Kidder and his associates believe that the process is probably elimination of waste substances of the prolonged cell-division, since chromatin extrusion does not take place during a few divisions subsequent to reorganization.
after conjugation in *Conchophthirus mytili* and since in Colpidium and *Glaucoma*, the chromatin elimination appears to be followed by a high division rate and infrequency of conjugation. Dass (1950) noticed a dark body between two daughter macronuclei of a ciliate designated by him as *Glaucoma pyriformis* and considered it as surplus desoxyribonucleic acid about to be converted by the cytoplasm to ribonucleic acid necessary for active growth.

In *Paramecium aurelia*, Woodruff and Erdmann (1914) reported the occurrence of "endomixis." At regular intervals of about 30 days, the old macronucleus breaks down and disappears, while each of the two micronuclei divides twice, forming eight nuclei. Of these, six disintegrate. The animal then divides into two, each daughter individual receiving one micronucleus. This nucleus soon divides twice into four, two of which develop into two macronuclei, while the other two divide once more. Here the organism divides again into two individuals, each bearing one macronucleus and two micronuclei. This process, they maintained, is "a complete periodic nuclear reorganization without cell fusion in a pedigreed race of Paramecium."

The so-called endomixis has since been reported to occur in many ciliates. However, as pointed out by Wilson (1928), Diller (1936), Sonneborn (1947) and others, there are several difficulties in holding that endomixis is a valid process. Diller considers that endomixis may have been based upon partial observations on hemixis (p. 206) and autogamy (p. 203). Sonneborn could not find any indication that this process occurs in numerous stocks and varieties of *Paramecium aurelia*, including the progeny of the strains studied by Woodruff, and maintained that endomixis does not occur in this species of Paramecium.

As has been stated already, two types of nuclei: macronucleus and micronucleus, occur in Euciliata and Suctoria. The macronucleus is the center of the whole metabolic activity of the organism and in the absence of this nucleus, the animal perishes. The waste substances which become accumulated in the macronucleus through its manifold activities, are apparently eliminated at the time of division, as has been cited above in many species. On the other hand, it is also probable that under certain circumstances, the macronucleus becomes impregnated with waste materials which cannot be eliminated through this process. Prior to and during conjugation (p. 188) and autogamy (p. 203), the macronucleus becomes transformed, in many species, into irregularly coiled thread-like structure (Fig. 85) which undergoes segmentation into pieces and finally is absorbed by the cytoplasm. New macronuclei are produced from
some of the division-products of micronuclei by probably incorporating the old macronuclear material. In most cases this supposition is not demonstrable. However, Kidder (1938) has shown in

Fig. 57. Diagram showing the macronuclear regeneration in *Paramecium aurelia* (Sonneborn). a, an individual before the first division after conjugation or autogamy, containing two macronuclear (stippled) anlagen, two micronuclei (rings) and about 30 disintegrating (solid black) masses of the old macronucleus; b, two individuals formed by the first division, each containing one macronuclear anlage, two micronuclei and macronuclear masses; c, two individuals produced by the second division: one (above) with the new macronucleus, two micronuclei and macronuclear masses, and the other without new macronucleus; d–f, binary fissions in which the two micronuclei divide, but old macronuclear masses are distributed equally between the two daughters until there is one large regenerated macronucleus and two micronuclei; g, division following f, goes on in an ordinary manner.

the encysted *Paraclevelandia simplex*, an endocommensal of the colon of certain wood-feeding roaches, this is actually the case; namely, one of the divided micronuclei fuses directly with a part of macronucleus to form a macronuclear anlage which then develops into a macronucleus after passing through "ball-of-yarn" stage similar to that which appears in an exconjugant of Nyctotherus (Fig. 85).
Since the macronucleus originates in a micronucleus, it must contain all structures which characterize the micronucleus. Why then does it not divide mitotically as does the micronucleus? During conjugation or autogamy in a ciliate, the macronucleus degenerates, disintegrates and finally becomes absorbed in the cytoplasm. In Paramaecium aurelia, Sonneborn (1940, 1942, 1947) (Fig. 57) observed that when the animal in conjugation is exposed to 38°C. from the time of the synkaryon-formation until before the second postzygotic nuclear division (a–c), the development of the two newly formed macronuclei is retarded and do not divide as usual with the result that one of the individuals formed by the second postzygotic division receives the newly formed macronucleus, while the other lacks this (c). In the latter, however, division continues, during which some of the original 20–40 pieces of the old macronucleus that have been present in the cytoplasm segregate in approximately equal number at each division (d, e) until there is only one in the animal (f). Thereafter the macronucleus divides at each division (g). Sonneborn found this “macronuclear regeneration” in the varieties 1 and 4, but considered that it occurs in all stocks. Thus the macronucleus in this ciliate appears to be a compound structure with its 20–40 component parts, each containing all that is needed for development into a complete macronucleus. From these observations, Sonneborn concludes that the macronucleus in P. aurelia appears to undergo amitosis, since it is a compound nucleus composed of many “subnuclei” and since at fission all that is necessary to bring about genetically equivalent functional macronuclei is to segregate these multiple subnuclei into two random groups.

While the macronuclear division usually follows the micronuclear division, it takes place in the absence of the latter as seen in amicronucleate individuals of ciliates which possess normally a micronucleus. Amicronucleate ciliates have been found to occur naturally or produced experimentally in the following species: Didinium nasutum (Thon, 1905; Patten, 1921), Oxytricha hymenostoma (Dawson, 1919), O. fallax, Urostyla grandis (Woodruff, 1921), Paramaecium caudatum (Landis, 1920; Woodruff, 1921), etc. Amicronucleate Oxytricha fallax which were kept under observation by Reynolds (1932) for 29 months, showed the same course of regeneration as the normal individuals. Beers (1946b) saw no difference in vegetative activity between amicronucleate and normal individuals of Tillina magna. In Euplotes patella, amicronucleates arise from “double” form (p. 229) with a single micronucleus, and Kimball (1941a) found that the micronucleus is not essential for continued life in at least some
clones, though its absence results in a marked decrease in vigor. The bi-micronucleate *Paramecium bursaria* which Woodruff (1931) isolated, developed in the course of 7 years of cultivation, unimicronucleate and finally amicronucleate forms, in which no marked variation in the vitality of the race was observed. These data indicate that amicronucleates are capable of carrying on vegetative activity and multiplication, but are unable to conjugate or if cell-pairing occurs, the result is abortive, though Chen (1940c) reported conjugation between normal and amicronucleate individuals of *P. bursaria* (p. 189). Horváth (1950) succeeded in destroying the micronucleus in *Kahlia simplex* (p. 133) and found the emicronucleates as vigorous as the normal forms, judged by the division rate, but were killed within 15 days by proactinomycin, while normal individuals resisted by encystment. This worker reasons that the emicronucleates are easily destroyed by unfavorable conditions and, therefore, ciliates without a micronucleus occur rarely in nature.

![Fig. 58. Amitosis of the vegetative nucleus in the trophozoite of *Myxosoma catostomi*, ×2250 (Kudo).](image)

Other examples of amitosis are found in the vegetative nuclei in the trophozoite of Myxosporidia, as for example, *Myxosoma catostomi* (Fig. 58), *Thelohanellus notatus* (Debaisieux), etc., in which the endosome divides first, followed by the nuclear constriction. In *Streblomastix strix*, the compact elongated nucleus was found to undergo a simple division by Kofoid and Swezy. **Indirect nuclear division.** The indirect division which occurs in the protozoan nuclei is of manifold types as compared with the mitosis in the metazoan cell, in which, aside from minor variations, the change is of a uniform pattern. Chatton, Alexeieff and others, have proposed several terms to designate the various types of indirect nuclear division, but no one of these types is sharply defined. For our purpose, mentioning of a few examples will suffice.

A veritable mitosis was noted by Dobell in the heliozoan *Oxnerella maritima* (Fig. 59), which possesses an eccentrically situated nucleus containing a large endosome and a central centriole, from which radiate many axopodia (a). The first sign of the nuclear division is
the slight enlargement, and migration toward the centriole, of the nucleus (b). The centriole first divides into two (c, d) and the nucleus becomes located between the two centrioles (e). Presently spindle fibers are formed and the nuclear membrane disappears (f, g). After

Fig. 59. Nuclear and cytoplasmic division in Oxnerella maritima, × about 1000 (Dobell). a, a living individual; b, stained specimen; c–g, prophase; h, metaphase; i, anaphase; j, k, telophase; l, division completed.

passing through an equatorial-plate stage, the two groups of 24 chromosomes move toward the opposite poles (g–i). As the spindle fibers become indistinct, radiation around the centrioles becomes conspicuous and the two daughter nuclei are completely reconstructed to assume the resting phase (j–l). The mitosis of another heliozoan Acanthocystis aculeata is, according to Schaudinn and
Stern, very similar to the above. Aside from these two species, the centriole has been reported in many others, such as Hartmannella (Arndt), Euglypha, Monocystis (Bélař), Aggregata (Dobell; Bélař;

Fig. 60. Mitosis in Trichonympha campanula, ×800 (Kofoid and Swezy). a, resting nucleus; b–g, prophase; h, metaphase; i, j, anaphase; k, telophase; l, a daughter nucleus being reconstructed.
Naville), various Hypermastigina (Kofoid; Duboseq and Grassé; Kirby; Cleveland and his associates).

In numerous species the division of the centriole (or blepharoplast) and a connecting strand between them, which has been called desmose (centrodesmose or paradesmose), have been observed. According to Kofoid and Swezy (1919), in *Trichonympha campanula* (Fig. 60), the prophase begins early, during which 52 chromosomes are formed and become split. The nucleus moves nearer the anterior end where the centriole divides into two, between which develops a desmose. From the posterior end of each centriole, astral rays extend out and the split chromosomes form loops and pass through "tangled skein" stage. In the metaphase, the equatorial plate is made up of V-shaped chromosomes as each of the split chromosomes is still connected at one end, which finally becomes separate in anaphase, followed by reformation of two daughter nuclei.

As to the origin and development of the achromatic figure, various observations and interpretations have been advanced. Certain Hypermastigina possess very large filiform centrioles and a large rounded nucleus. In *Barbulanympha* (Fig. 61), Cleveland (1938a) found that the centrioles vary from 15 to 30μ in length in the four species of the genus which he studied. They can be seen, according to Cleveland, in life as made up of a dense hyaline protoplasm. When stained, it becomes apparent that the two centrioles are joined at their anterior ends by a desmose and their distal ends 20 to 30μ apart, each of which is surrounded by a special centrosome (a). In the resting stage no fibers extend from either centriole, but in the prophase, astral rays begin to grow out from the distal end of each centriole (b). As the rays grow longer (c), the two sets soon meet and the individual rays or fibers join, grow along one another and overlap to form the central spindle (d). In the resting nucleus, there are large irregular chromatin granules which are connected by fibrils with one another and also with the nuclear membrane. As the achromatic figure is formed and approaches the nucleus, the chromatin becomes arranged in a single spireme imbedded in matrix. The spireme soon divides longitudinally and the double spireme presently breaks up transversely into paired chromosomes. The central spindle begins to compress the nuclear membrane and the chromosomes become shorter and move apart. The intra- and extra-nuclear fibrils unite as the process goes on (e), the central spindle now assumes an axial position, and two groups of V-shaped chromosomes are drawn to opposite poles. In the telophase, the chromosomes elongate and become branched, thus assuming conditions seen in the resting nucleus.
Fig. 61. Development of spindle and astral rays during the mitosis in Barbulanympha, ×930 (Cleveland). a, interphase centrioles and centrosomes; b, prophase centrioles with astral rays developing from their distal ends through the centrosomes; c, meeting of astral rays from two centrioles; d, astral rays developing into the early central spindle; e, a later stage showing the entire mitotic figure.

In Holomastigotoides tusitala (Fig. 172, a, b), Cleveland (1949) brought to light the formation of the achromatic figure, and the minute structure and change in chromosomes (Fig. 62). In the late telophase, after cytoplasmic division, the centrioles follow the flagellar bands 4 and 5 for 1.5 turns (a). The two chromosomes are anchored to the old centriole. When the new centriole has become as
Fig. 62. Mitosis in *Holomastigotoides tusitala* (Cleveland). a, anterior region showing flagellar bands, centrioles, centromeres and chromosomes. b–h, telophase; i, j, prophase; k, metaphase; l, anaphase; m, telophase. b, c, new and old centrioles forming achromatic figure; d, one chromosome has shifted its connection from old to new centriole; e, f, flattening out of centrioles and achromatic figure; g, h, beginning of chromosomal twisting; i, chromosomes duplicated, producing many gyres of close-together relational coiling of chromatics, and centromeres duplicated; j, chromatids losing their relational coiling by unwinding; k, relational coiling disappeared, achromatic figure elongating and separating sister chromatids; l, central spindle bent, chromatids in two groups; m, central spindle pulled apart.
long as the old one, the centrioles begin to produce astral rays (b) which soon meet and form the central spindle (c). An astral ray from the new centriole becomes connected with the centromere of one of the chromosomes (d). The spindle grows in length and enters resting stage (e–j), later the spindle fibers lengthen (k, l) and pull apart (m).

The chromosome is composed of the matrix and chromonema (Fig. 63), of which the former disintegrates in the telophase and re-appears in the early prophase of each chromosome generation, while the latter remains throughout. From late prophase to mid-telophase, minor coils are incorporated in major coils (a–e); from mid-telophase to late telophase, they are in very loose majors (d); and after the majors have disappeared completely, they become free (e). Soon after cytoplasmic division, the majors become looser and irregular and finally disappear, while minors and twisting remain. Each chromosome presently divides into 2 chromatids (f) and a new matrix is formed for each. As the matrix contracts the chromatids lose their relational coiling and the minors become bent and thus the new generation of major coils makes its appearance (g). With the further concentration of the matrix, the majors become more conspicuous (h), the minors being incorporated into them. When most of the relational coiling has been lost and majors are close together, the chromosomal changes cease for days or weeks. This is the late prophase. After the resting stage, the achromatic figure commences to grow again (i, j) and the two groups of chromatids are carried to the poles, followed by transverse cytoplasmic division (Fig. 64). The coils remain nearly the same during metaphase to early telophase. Thus Cleveland showed the continuity of chromosomes from generation to generation. He finds that the resting stage of chromosomes varies in different types of cells: some chromosomes rest in interphase, some in early prophase and others in telophase, and that the centromere is an important structure associated with the movement of chromatids and in the reduction of chromosomes in meiosis. For fuller information the reader is referred to the profusely illustrated original paper (Cleveland, 1949).

In Lophomonas blattarum, the nuclear division (Fig. 65) is initiated by the migration of the nucleus out of the calyx. On the nuclear membrane is attached the centriole which probably originates in the blepharoplast ring; the centriole divides and the desmose which grows, now stains very deeply, the centrioles becoming more conspicuous in the anaphase when new flagella develop from them. Chromatin granules become larger and form a spireme, from which
Fig. 63. Chromosomal changes in *Holomastigoloides tusitala*, ×1050 (Cleveland). a, telophase shortly after cytoplasmic division, new fifth band and new centriole are growing out and chromosomes are twisted; b, c, the same chromosome showing major and minor coils respectively; d, later telophase, showing minor coils; e, matrix completely disintegrated, showing minor coils; f, a prophase nucleus, showing division of chromosomes into two chromatids; g, later prophase, in which majors are developing with minors; h, later prophase; i, metaphase in which distal halves of the chromatids have not yet separated, showing minor coils; j, anaphase, showing major and minor coils of chromonemata.
Fig. 64. Cytoplasmic division in *Holomastigotoides tusitala*, X about 430 (Cleveland). a, fifth flagellar band has separated from others; b, one nucleus and fifth band moving toward posterior end; c, the movement of the band and nucleus has been completed; d, e, anterior and posterior daughter individuals, produced by transverse division.
6–8 chromosomes are produced. Two groups of chromosomes move toward the opposite poles, and when the division is completed, each centriole becomes the center of formation of all motor organellae.

In some forms, such as Noctiluca (Calkins), Actinophrys (Bělař), etc., there may appear at each pole, a structureless mass of cytoplasm (centrosphere), but in a very large number of species there appear no special structures at poles and the spindle fibers become stretched seemingly between the two extremities of the elongating nuclear membrane. Such is the condition found in Pelomyxa (Kudo) (Fig. 66), Cryptomonas (Bělař), Rhizochrysis (Doflein), Aulacantha (Borgert), and in micronuclear division of the majority of Euciliata and Suctoria.

The behavior of the endosome during the mitosis differs among different species as are probably their functions. In Eimeria schubergi (Schaudinn), Euglena viridis (Tschenzoff), Oxyrrhis marina (Hall),
Colacium vesiculosum (Johnson), Haplosporidium limnodrili (Granata), etc., the conspicuously staining endosome divides by elongation and constriction along with other chromatic elements, but in many other cases, it disappears during the early part of division and reappears when the daughter nuclei are reconstructed as observed in Monocystis, Dimorpha, Euglypha, Pamphagus (Bélař), Acanthocystis (Stern), Chilomonas (Doflein), Dinenympha (Kirby), etc.

In the vegetative division of the micronucleus of Conchophthirus anodontae, Kidder (1934) found that prior to division the micronucleus moves out of the pocket in the macronucleus and the chromatin becomes irregularly disposed in a reticulum; swelling continues and the chromatin condenses into a twisted band, a spireme, which breaks into many small segments, each composed of large chromatin granules. With the rapid development of the spindle fibers, the twelve bands become arranged in the equatorial plane and condense. Each chromosome now splits longitudinally and two groups of 12 daughter chromosomes move to opposite poles and transform them-
selves into two compact daughter nuclei. A detailed study of micronuclear division (Fig. 67) of Urostyla grandis was made by Raabe (1946). The micronucleus is a compact body in the interphase (a),

Fig. 67. Micronuclear division of Urostyla grandis, ×2100 (H. Raabe). a, resting stage; b–j, prophase (b–e, stages in the formation of spireme; f, g, spireme ribbon; h, i, twelve segments of ribbon arranged in the direction of the elongating nuclear axis; j, a polar view of the same); k, l, metaphase, condensation of the segments; m–o, anaphase; p, late anaphase; q, a daughter nucleus in telophase; r–t, reconstruction stages; u, a resting daughter nucleus.
but increases in size and the chromatin becomes grouped into small masses (b, c), which become associated into a spiral ribbon (d–g). The latter then breaks up into 12 segments that are arranged parallel to the axis of the elongating nucleus (h–i). Each segment condenses into a chromosome which splits longitudinally into two (k) and the two groups of chromosomes move to opposite poles (l–P). In Zelleriella elliptica (Fig. 295) and four other species of the genus inhabiting the colon of Bufo valliceps, Chen (1936, 1948) observed the formation of 24 chromosomes, each of which is connected with a fiber of the intranuclear spindle and splits lengthwise in the metaphase.

While in the majority of protozoan mitosis, the chromosomes split longitudinally, there are observations which suggest a transverse division. As examples may be mentioned the chromosomal divisions in Astasia laevis (Bélař), Entosiphon sulcatum (Lackey), and a number of ciliates. In a small number of species observations vary within a species, as, for example, in Peranema trichophorum in which the chromosomes were observed to divide transversely (Hartmann and Chagas) as well as longitudinally (Hall and Powell; Brown). It is inconceivable that the division of the chromosome in a single species of organism is haphazard. The apparent transverse division might be explained by assuming, as Hall (1937) showed in Euglena gracilis, that the splitting is not completed at once and the pulling force acting upon them soon after division, brings forth the long chromosomes still connected at one end. Thus the chromosomes remain together before the anaphase begins.

In the instances considered on the preceding pages, the so-called chromosomes found in them, appear to be essentially similar in structure and behavior to typical metazoan chromosomes. In many other cases, the so-called chromosomes or "pseudochromosomes" are slightly enlarged chromatin granules which differ from the ordinary chromatin granules in their time of appearance and movement only. In these cases it is of course not possible at present to determine how and when their division occurs before separating to the respective division pole. In Table 5 are listed the number of the "chromosomes" which have been reported by various investigators in the Protozoa that are mentioned in the present work.

**Cytoplasmic division**

The division of the nucleus is accompanied by division of extranuclear organelles such as chromatophores, pyrenoids, etc. The blepharoplast of the flagellates and kinetosomes of the ciliates undergo di-
<table>
<thead>
<tr>
<th>Protozoa</th>
<th>Number of chromosomes</th>
<th>Observers</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Rhizochrysis scherffelii</em></td>
<td>22</td>
<td>Dofflein</td>
</tr>
<tr>
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</tr>
<tr>
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<td>Dofflein</td>
</tr>
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<td>Moewus</td>
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<td><em>Euglena psciformis</em></td>
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<td>Dangeard</td>
</tr>
<tr>
<td><em>E. viridis</em></td>
<td>30 or more</td>
<td>Dangeard</td>
</tr>
<tr>
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<td>30–40</td>
<td>Dangeard</td>
</tr>
<tr>
<td><em>Rhodomonas incurva</em></td>
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<td>Hall</td>
</tr>
<tr>
<td><em>Vacuolaria virescens</em></td>
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<td>Fott</td>
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<td>Chatton</td>
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<tr>
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</tr>
<tr>
<td><em>Cercomonas longicauda</em></td>
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</tr>
<tr>
<td><em>C. coli</em></td>
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<td>Bélař</td>
</tr>
<tr>
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<td>Boeck and Tanabe</td>
</tr>
<tr>
<td><em>Eutrichomastix serpentis</em></td>
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<td>Kofoid and Swezy</td>
</tr>
<tr>
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<td>Kirby</td>
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<td>Light</td>
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<tr>
<td><em>T. hominis</em></td>
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<td>Bishop</td>
</tr>
<tr>
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</tr>
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<tr>
<td></td>
<td>6</td>
<td>Samuels</td>
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</tr>
<tr>
<td><em>G. muris</em></td>
<td>4</td>
<td>Kofoid and Christiansen</td>
</tr>
<tr>
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</tr>
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<td>Cup</td>
</tr>
<tr>
<td><em>S. bispira</em></td>
<td>2 doubles</td>
<td>Cleveland</td>
</tr>
<tr>
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<td>Janicki</td>
</tr>
<tr>
<td></td>
<td>8 or 6</td>
<td>Kudo</td>
</tr>
<tr>
<td></td>
<td>12 or 6 double</td>
<td>Bélař</td>
</tr>
<tr>
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<td>Cleveland</td>
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<td><em>Trichonympha campanula</em></td>
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<td>Kofoid and Swezy</td>
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<td>Protozoa</td>
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<td><em>M. serotinum</em></td>
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<td><em>Sphaeromyxa sabrazesi</em></td>
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<td><em>Myxobolus pfeifferi</em></td>
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<td>Naville</td>
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### Table 5.—Continued

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<th>Protozoa</th>
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<td>Chilodonella uncinata</td>
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<td>C. uncinata (tetraploid)</td>
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<td>MacDougall</td>
</tr>
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<td>Kidder</td>
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<tr>
<td>E. eurystomus</td>
<td>8 (diploid)</td>
<td>Turner</td>
</tr>
<tr>
<td>Vorticella microstoma</td>
<td>4</td>
<td>Finley</td>
</tr>
<tr>
<td>Carchesium polypinum</td>
<td>16 (diploid)</td>
<td>Popoff</td>
</tr>
<tr>
<td>Trichodina sp.</td>
<td>4–6</td>
<td>Diller</td>
</tr>
</tbody>
</table>

Vision, giving rise to daughter blepharoplasts and kinetosomes that become organized into characteristic locomotor organelles. Morphogenesis in the apostomes (Chatton and Lwoff, 1935; Lwoff, 1950); mechanism of morphogenesis in ciliates (Fauré-Fremiet, 1948; Guiller, 1950; Weisz, 1951, 1951a).

**Binary fission.** As in metazoan cells, the binary fission occurs very widely among the Protozoa. It is a division of the body through middle of the extended long axis into two nearly equal daughter individuals. In *Amoeba proteus*, Chalkley and Daniel found that there is a definite correlation between the stages of nuclear division and external morphological changes (Fig. 68). During the prophase, the organism is rounded, studded with fine pseudopodia and exhibits under reflected light a clearly defined hyaline area near its center (a), which disappears in the metaphase (b, c). During the anaphase the pseudopodia rapidly become coarser; in the telophase the elongation of body, cleft formation, and return to normal pseudopodia, take place.

In Testacea, one of the daughter individuals remains, as a rule, within the old test, while the other moves into a newly formed one,
as in Arcella, Pyxidicula, Euglypha, etc. According to Doflein, the
division plane coincides with the axis of body in Cochliopodium,
Pseudodifflugia, etc., and the delicate homogeneous test also divides
into two parts. In the majority of the Mastigophora, the division is
longitudinal, as is shown by that of *Rhabdomonas incurva* (Fig. 69).
In certain dinoflagellates, such as Ceratium, Cochliodinium, etc.,
the division plane is oblique, while in forms such as Oxyrrhis (Dunk-
ery; Hall), the fission is transverse. In *Streblomastix strix* (Kofoid
and Swezy, 1919), *Lophomonas striata* (Kudo, 1926b), *Spirotrichon-
nympha bispira* (Cleveland, 1938), *Holomastigotoides tusitala* (Fig.
64) and others (Cleveland, 1947), and *Strombidium clavellinae* (Bud-
denbrock, 1922), the division takes place transversely but the polar-
ity of the posterior individual is reversed so that the posterior end
of the parent organism becomes the anterior end of the posterior
daughter individual. In the ciliate Bursaria, Lund (1917), observed
reversal of polarity in one of the daughter organisms at the time of
division of normal individuals and also in those which regenerated
after being cut into one-half the normal size.

**Fig. 68.** External morphological changes during division of *Amoeba proteus*, as viewed in life in reflected light, × about 20 (Chalkley and Daniel). a, shortly before the formation of the division sphere; b, a later stage; c, prior to elongation; d, further elongation; e, division almost completed.
In the Ciliophora the division is as a rule transverse (Fig. 52), in which the body without any enlargement or elongation divides by constriction through the middle so that the two daughter individuals are about half as large at the end of division. Both individuals usually retain their polarity.

**Multiple division.** In multiple division the body divides into a number of daughter individuals, with or without residual cytoplasmic masses of the parent body. In this process the nucleus may undergo either simultaneous multiple division, as in Aggregata, or more commonly, repeated binary fission, as in Plasmodium (Fig. 256) to produce large numbers of nuclei, each of which becomes the center of a new individual. The number of daughter individuals often varies, not only among the different species, but also within one and the same species. Multiple division occurs commonly in the Foraminifera (Fig. 208); the Radiolaria (Fig. 218), and various groups of Sporozoa in which the trophozoite multiplies abundantly by this method.

**Budding.** Multiplication by budding which occurs in the Protozoa is the formation of one or more smaller individuals from the

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**Fig. 69.** Nuclear and cytoplasmic division in *Rhabdomonas incurva*, × about 1400 (Hall). a, resting stage; b, c, prophase; d, equatorial plate; e, f, anaphase; g, telophase.
parent organism. It is either exogenous or endogenous, depending upon the location of the developing buds or gemmules. Exogenous budding has been reported in Acanthocystis, Noctiluca (Fig. 127), Myxosporidia (Fig. 70, b), astomatous ciliates (Fig. 298), Chonotricha, Suctoria (Fig. 371, k), etc. Endogenous budding has been found in Testacea, Gregarinida, Myxosporidia (Figs. 279, e; 281, j), and other Sporozoa as well as Suctoria (Fig. 371, h). Collin observed a unique budding in *Tokophrya cyclopum* in which the entire body, excepting the stalk and pellicle, transforms itself into a young ciliated bud and leaves sooner or later the parent pellicle.

**Plasmotomy.** Occasionally the multinucleate body of a protozoan divides into two or more small, multinucleate individuals, the cytoplasmic division taking place independently of nuclear division. This has been called plasmotomy by Doflein. It has been observed in the

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Fig. 70. a, b, budding in *Myxidium lieberkühni*; c, d, plasmotomy in *Chloromyxum leydi*; e, plasmotomy in *Sphaeromyxa balbianii*. 
trophozoites of several coelozoic myxosporidians, such as *Chloromyxum leydigi*, *Sphaeromyxa balbianii* (Fig. 70), etc. It occurs further in certain Sarcodina such as Mycetozoa (Fig. 179) and Pelomyxa (Fig. 71), and Protociliata.

Fig. 71. Eight individuals of *Pelomyxa carolinensis*, seen undisturbed in culture dishes, in which mitotic stages occurred as follows, ×40 (Kudo): a, early prophase; b, c, later prophase; d, metaphase; e, f, early and late anaphase; g, h, late telophase to resting nuclei (g, plasmotomy into two individuals; h, plasmotomy into three daughters).

**Colony formation**

When the division is repeated without a complete separation of the daughter individuals, a colonial form is produced. The compon-
ent individuals of a colony may either have protoplasmic connections among them or be grouped within a gelatinous envelope if completely separated. Or, in the case of loricate or stalked forms, these exoskeletal structures may become attached to one another. Although varied in appearance, the arrangement and relationship of the component individuals are constant, and this makes the basis for distinguishing the types of protozoan colonies, as follows:

**Catenoid or linear colony.** The daughter individuals are attached endwise, forming a chain of several individuals. It is of comparatively uncommon occurrence. Examples: Astomatous ciliates such as Radiophrya (Fig. 298), Protoradiophrya (Fig. 298) and dinoflagellates such as Ceratium, Haplozoon (Fig. 130) and Polykrikos (Fig. 132).

**Arboroid or dendritic colony.** The individuals remain connected with one another in a tree-form. The attachment may be by means of the lorica, stalk, or gelatinous secretions. It is a very common colony found in different groups. Examples: Dinobryon (Fig. 108), Hyalobryon (Fig. 108), etc. (connection by lorica); Colacium (Fig. 121), many Peritricha (Figs. 362; 364), etc. (by stalk); Poteriodendron (Fig. 139), Stylobryon (Fig. 151), etc. (by lorica and stalk); Hydrurus (Fig. 109), Spongomonas (Fig. 150), Cladomonas (Fig. 150) and Anthophysis (Fig. 151) (by gelatinous secretions).

**Discoid colony.** A small number of individuals are arranged in a single plane and grouped together by a gelatinous substance. Examples: Cyclonexis (Fig. 108), Gonium (Fig. 116), Platydorina (Fig. 117), Protospongia (Fig. 138), Bicosoeca (Fig. 139), etc.

**Spheroid colony.** The individuals are grouped in a spherical form. Usually enveloped by a distinct gelatinous mass, the component individuals may possess protoplasmic connections among them. Examples: Uroglena (Fig. 108, e), Uroglenopsis (Fig. 108, d), Volvox (Fig. 115), Pandorina (Fig. 117, f), Eudorina (Fig. 117, k), etc. Such forms as Stephanoon (Fig. 117, a) appear to be intermediate between this and the discoid type. The component cells of some spheroid colonies show a distinct differentiation into somatic and reproductive individuals, the latter developing from certain somatic cells during the course of development.

The gregaroid colony, which is sometimes spoken of, is a loose group of individuals of one species, usually of Sarcodina, which become attached to one another by means of pseudopodia in an irregular form.
Asexual reproduction

The Protozoa nourish themselves by certain methods, grow and multiply, by the methods described in the preceding pages. This phase of the life-cycle of a protozoan is the vegetative stage or the trophozoite. The trophozoite repeats its asexual reproduction process under favorable circumstances. Generally speaking, the Sporozoa increase to a much greater number by multiple division or schizogony and the trophozoites are called schizonts.

Under certain conditions, the trophozoite undergoes encystment (Fig. 72). Prior to encystment, the trophozoites cease to ingest, and extrude remains of, food particles, resulting in somewhat smaller forms which are usually rounded and less active. This phase is some-

![Fig. 72. Encystment of Lophomonas blattarum, x1150 (Kudo).](image)

times called the precystic stage. The whole organism becomes de-differentiated; namely, various cell organs such as cilia, cirri, flagella, axostyle, peristome, etc., become usually absorbed. Finally the organism secretes substances which become solidified into a resistant wall, and thus the cyst is formed. In this condition, the protozoan is apparently able to maintain its vitality for a certain length of time under unfavorable conditions.

Protozoa appear to encyst under various conditions. Low temperature (Schmähl, 1926), evaporation (Bělař, 1921; Bodine, 1923; Garnjobst, 1928), change in pH (Koffman, 1924; Darby, 1929), low or high oxygen content (Brand, 1923; Rosenberg, 1938), accumulation of metabolic products (Bělař, 1921; Mast and Ibara, 1923; Beers, 1926) or of associated bacteria (Mouton, 1902; Bělař, 1921) and over-population (Barker and Taylor, 1931) in the water in which Protozoa live, have been reported to bring about encystment. While
lack of food in the culture has been noted by many observers (Oehler, 1916; Claff, Dewey and Kidder, 1941; Singh, 1941; Beers, 1948; etc.) as a cause of encystment in a number of Protozoa such as Blepharisma (Stolte, 1922), Polytomella (Kater and Burroughs, 1926), Didinium (Mast and Ibara, 1931), Uroleptus (Calkins, 1933), etc., an abundance of food and adequate nourishment seem to be prerequisite for encystment. Particular food was found in some instances to induce encystment. For example, Singh (1948) employed for culture of Leptomyxa reticulata, 40 strains of bacteria, of which 15 led to the production of a large number of cysts in this sarcodinian. Encystment of Entamoeba histolytica is easily obtained by adding starch to the culture (Dobell and Laidlow, 1926; Balamuth, 1951).

The age of culture, if kept under favorable conditions, does not influence encystment. Didinium after 750 generations, according to Beers (1927), showed practically the same encystment rate as those which had passed through 10 or 20 generations since the last encystment. When Leptomyxa mentioned above is cultured for more than a year, no encystment occurred, but young cultures when supplied with certain bacteria encysted (Singh, 1948).

In some cases, the organisms encyst temporarily in order to undergo nuclear reorganization and multiplication as in Colpoda (Fig. 73) (Kidder and Claff, 1938; Stuart, Kidder and Griffin, 1939), Tiliina (Beers, 1946), etc. In Ichthyophthirius, the organism encysts after leaving the host fish and upon coming in contact with a solid object, and multiplies into numerous "ciliospores" (MacLennan, 1937). Pelomyxa carolinensis (Illinois stock) has not encysted since its discovery in 1944, although the cultures were subjected to various environmental changes, but P. illinoensis has been found to encyst and excyst frequently in flourishing cultures (Kudo, 1951). Thus it may be assumed that some unknown internal factors play as great a part as do the external factors in the phenomenon of encystment (Ivanic, 1934; Cutler and Crump, 1935).

The cyst is covered by one to three membranes. Though generally homogeneous, the wall of cyst may contain siliceous scales as in Euglypha (Fig. 74). While chitinous substance is the common material of which the cyst wall is composed, cellulose makes up the cyst membrane of many Phytomastigina. Entz (1925) found the cysts of various species of Ceratium less variable in size as compared with the vegetative form, and found in all, glycogen, oil and volutin.

The capacity of Protozoa to produce cyst is probably one of the
reasons why they are so widely distributed over the surface of the globe. The minute protozoan cysts are easily carried from place to place by wind, attached to soil particles, debris, etc., by the flowing water of rivers or the current in oceans or by insects, birds, other animals to which they become readily attached. The cyst is capable of remaining viable for a long period of time: eight years in *Haematococcus pluvialis* (Reichenow, 1929), four years in *Spathidium spatula* and *Oxytricha* sp. (Dawson and Mitchell, 1929), five years in *Colpoda cucullus* (Dawson and Hewitt, 1931), 10 years in *Didinium nasutum* (Beers, 1937), etc.

![Diagram](image-url)

**Fig. 73.** Diagram showing the life cycle of *Colpoda cucullus* (Kidder and Claff). a–j, normal reproductive activity repeated (j–b) under favorable cultural conditions; k–o, resistant cyst (k–n, nuclear reorganization and chromatin elimination).
When a cyst encounters a proper environment, redifferentiation takes place within the cyst. Various organellae which characterize the organism, are regenerated and reformed, and the young trophozoite excysts. The emerged organism returns once more to its trophic phase of existence. Experimental data indicate that excystment takes place under conditions such as addition of fresh culture medium (Kühn, 1915; Rosenberg, 1938), hypertonic solution (Ilowaisky, 1926), distilled water (Johnson and Evans, 1941), organic infusion (Mast, 1917; Beers, 1926; Barker and Taylor, 1933), and bacterial infusion (Singh, 1941; Beers, 1946a) to the culture medium. Change in pH (Koffman, 1924), lowering the temperature (Johnson and Evans, 1941) and increase in oxygen content (Brand, 1923; Finley, 1936) of the medium have also been reported as bringing about excystment. Excystment in Colpoda cucullus is said to be due to specific inducing substances present in plant infusion (Thimann and Barker, 1934; Haagen-Smit and Thimann, 1938). Experimenting with two soil amoebae, “species 4 and Z,” Crump (1950) found that the excystment in species Z took place without the presence of bacteria and regardless of the age of the cysts, but species 4 excysted only in the presence of certain bacteria (Aerobacter sp. or “4036”) and the excystment diminished with the age of cysts. Crump suggested that the two strains of bacteria appeared to produce some material which induced excystment in Amoeba species 4. In Tillina magna, Beers (1945) found, however, the primary excystment-inducing factor to be of an osmotic nature and inducing substances, a secondary one.

As to how an aperture or apertures are formed in the cyst wall prior to the emergence of the content, precise information is not yet on hand, though there are many observations. In the excystment in Didinium and Tillina, Beers (1935, 1945, 1945a) notes that

Fig. 74. Encystment of Euglypha acanthophora, ×320 (Kühn).
an increased internal pressure due to the imbibition of water, results in the rupture of the cyst wall which had lost its rigidity and resistance (Fig. 75). Apertures in the cyst wall of *Pelomyxa illinoisensis* are apparently produced by pseudopodial pressure (Kudo, 1951). Seeing a similar aperture formation in the cyst of *Entamoeba histolytica*, Dobell (1928) "imagined that the amoeba secretes a ferment which dissolves the cyst wall."

Fig. 75. Excystment in *Didinium nasutum*, as seen in a single individual, ×250 (Beers). a, resting cyst; b, appearance of "excystment" vacuole; c, rupture of the cyst membrane, the vacuole is becoming enlarged; d, e, emergence of the cyst content, the vacuole increasing in size; f, the empty outer cyst membrane; g, the free organism with the inner membrane; h, organism after discharge of vacuole; i, j, later stages of emergence of the ciliate.

Although encystment seems to be an essential phase in the life cycle of Protozoa in general, there are certain Protozoa including such common and widely distributed forms as the species of *Paramecium* in which this phenomenon has not been definitely observed (p. 744). In some Sporozoa, encystment is followed by production of large numbers of spores, while in others there is no encystment. Here at the end of active multiplication of trophozoite, sexual re-
production usually initiates the production of the spores (Fig. 76). The spores which are protected by a resistant membrane are capable of remaining viable for a long period of time outside the host body.

![Diagram](image)

**Fig. 76.** Diagram illustrating the life-cycle of *Thelohania legeri* (Kudo). 

*Fig. 76.* Diagram illustrating the life-cycle of *Thelohania legeri* (Kudo). a, extrusion of the polar filament in gut of anopheline larva; b, emerged amoebula; c–f, schizogony in fat body; g–m, sporont-formation; m–x, stages in spore-formation.

**Sexual reproduction and life-cycles**

Besides reproducing by the asexual method, numerous Protozoa reproduce themselves in a manner comparable with the sexual reproduction which occurs universally in the Metazoa. Various types of sexual reproduction have been reported in literature, of which a few will be considered here. The sexual fusion or **syngamy** which is a complete union of two gametes, has been reported from various groups, while the conjugation which is a temporary union of two individuals for the purpose of exchanging the nuclear material, is found almost exclusively in the Ciliophora.

**Sexual fusion.** The **gametes** which develop from trophozoites, may be morphologically alike (**isogametes**) or unlike (**anisogametes**),
both of which are, in well-studied forms, physiologically different as judged by their behavior toward each other. If a gamete does not meet with another one, it perishes. Anisogametes are called microgametes and macrogametes. Difference between them is comparable in many instances (Figs. 77, 256) with that which exists between the spermatozoa and the ova of Metazoa. The microgametes are motile, relatively small and usually numerous, while the macrogametes are usually not motile, much more voluminous and fewer in number. Therefore, they have sometimes been referred to as male and female gametes (Fig. 77).

While morphological differences between the gametes have long been known and studied by many workers, whatever information we possess on physiological differences between them is of recent origin. Since 1933, Moewus and his co-workers have published a series of papers based upon their extended studies of bacteria-free cultures of many species (and strains) of Chlamydomonas (p. 276) which throw some light on the gamete differentiation among these phytomonadins. The gametes in Chlamydomonas are mostly isogamous, except in a few forms. Sexual fusion takes place in the majority of species and strains between the gametes produced in different clones, and there is no gametic fusion within a single clone. Moewus obtained "sex substances" from some of the cultures and showed that these are chemotactic substances. Each gamete secretes substances that attract the other and each reacts to the substances secreted by the other. Kühn, Moewus and Wendt (1939) recognized "hormones," and named them, termones (sex-determining hormones), andrortermone (male-determining hormone) and gynoter­mone (female-determining hormone).

In a few strains or species of Chlamydomonas, sexual fusion is found to take place among the gametes that develop within a single clone. Moewus considers in these cases there exist two types of gametes in a clone. However, Pascher, Pringsheim, and others ob-
tained results which seem to indicate that there is no physiological or sex differentiation between the fusing gametes. In the much-studied Sporozoa, for example, Plasmodium, the two gametes are both morphologically and physiologically differentiated, and sexual fusion always takes place between two anisogametes.

Fig. 78. Sexual fusion in Copromonas subtilis, ×1300 (Dobell).

Fig. 79. Sexual fusion in Trinema linearis, ×960 (Dunkerly). a, an organism in life, with the resting nucleus and two contractile vacuoles; b, union of two individuals; c, fusion of the organisms in one test, surrounded by cyst membrane; d, older cyst; e, still older cyst with a single nucleus.
The isogamy is typically represented by the flagellate *Copro-
monas subtilis* (Fig. 78), in which there occurs, according to Dobell,
a complete nuclear and cytoplasmic fusion between two isogametes. Each nucleus, after casting off a portion of its nuclear material, fuses with the other, thus forming a zygote containing a synkaryon. In *Trinema lineare* (Fig. 79), Dunkerly (1923) saw isogamy in which
Fig. 81. Sexual reproduction in Trichonympha of Cryptocercus (Cleveland). a, vegetative individual; b, gametocyte in early stage of encystment; c, anterior end of the same organism (chromosomes have been duplicated, nuclear sleeve is opening at seams and granules are flowing into the cytoplasm); d, further separation of the male and female chromosomes; e, the nuclear division has been completed, few old flagella remain and new post rostral flagella are growing; f, the cytoplasmic division has begun at the anterior end; g, the gametes just before encystment, the female showing the developing ring of fertilization granules; h, a female gamete; i, a female gamete with a fertilization ring. a, X350; b, X320; c, X600; d-i, X280.
two individuals undergo a complete fusion within one test and encyst. In *Stephanosphaera plurialis* (Fig. 80), both asexual and sexual reproductions occur, according to Hieronymus. Each individual multiplies and develops into numerous biflagellate gametes, all of which are alike. Isogamy between two gametes results in formation of numerous zygotes which later develop into trophozoites.

Anisogamy has been observed in certain Foraminifera. It perhaps occurs in the Radiolarians also, although positive evidence has yet to be presented. Anisogamy seems to be more widely distributed. In *Pandorina morum*, Pringsheim observed that each cell develops asexually into a young colony or into anisogametes which undergo sexual fusion and encyst. The organism emerges from the cyst and develops into a young trophozoite. A similar life-cycle was found by Goebel in *Eudorina elegans*.

The wood-roach inhabiting flagellates belonging to Trichonympha, Oxymonas, Saccinobaeulus, Notila and Eucomonympha, were found by Cleveland (1949a–1951a) to undergo sexual reproduction when the host insect molts. It has been observed that the gamete-formation is induced by the molting hormone produced by the prothoracic glands of the host insect. The sexual reproduction of Trichonympha, possessing 24 chromosomes, as observed and described by Cleveland, is briefly as follows (Figs. 81, 82): About three days before its host molts, the haploid nucleus in the flagellate divides, in which two types of daughter chromosomes (or chromatids) become separated from each other: the dark-staining male gamete nucleus and light-staining female gamete nucleus (Fig. 81, b–d); in the meantime, a membrane is formed to envelop the organism (b, d). When the cytoplasmic division is completed (e–g), the two gametes “encyst” and become free in the host gut (h; Fig. 82, b). In the female gamete, there appear “fertilization granules” (Fig. 81, h), which gather at the posterior extremity (i), through which a fluid-filled vesicle (“fertilization cone”) protrudes (Fig. 82, a). A male gamete (b) comes in touch with a female gamete only at this point (c), and enters the latter (d–f). The two gamete nuclei fuse into a diploid synkaryon (g, h). The zygote and its nucleus begin immediately to increase in size, and undergo two meiotic divisions (i–k), finally giving rise to vegetative individuals (Fig. 81, a).

Among the Sporozoans, anisogamy is of common occurrence. In Coccidia, the process was well studied in *Eimeria schubergi* (Fig. 213), *Aggregata eberthi* (Fig. 246), *Adelaea ovata* (Fig. 253), etc., and the resulting products are the *oocysts* (zygotes) in which the spores or sporozoites develop. Similarly in Haemosporidia such as *Plasmo-
Fig. 82. Sexual reproduction in Trichonympha of Cryptocercus (Cleveland). a, a female gamete with a fertilization ring and cone; b, a male gamete; c–g, stages in fusion and fertilization; h, a zygote; i, telophase of the first meiotic division of the zygote nucleus; j, k, prophase and anaphase of the second meiotic division. a–g, ×280; h, ×215; i–k, ×600.
diem vivax (Fig. 256), anisogamy results in the formation of the ookinetes or motile zygotes which give rise to a large number of sporozoites. Among Myxosporidia, a complete information as to how the initiation of sporogony is associated with sexual reproduction, is still lacking. Naville, however, states that in the trophozoite of Sphaeromyxa sabrazesi (Fig. 277), micro- and macro-gametes develop, each with a haploid nucleus. Anisogamy, however, is peculiar in that the two nuclei remain independent. The microgametic nucleus divides once and the two nuclei remain as the vegetative nuclei of the pansporoblast, while the macrogamete nucleus multiplies repeatedly and develop into two spores. Anisogamy has been suggested to occur in some members of Amoebina, particularly in Endamoeba blattae (Mercier, 1909). Cultural studies of various parasitic amoebae in recent years show, however, no evidence of sexual reproduction. Among the Ciliophora, the sexual fusion occurs only in Protociliata (Fig. 294).

Conjugation. The conjugation is a temporary union of two individuals of one and the same species for the purpose of exchanging part of the nuclear material and occurs almost exclusively in the Eugiliata and Suctoria. The two individuals which participate in this process may be either isogamous or anisogamous. In Paramecium caudatum (Fig. 83), the process of conjugation has been studied by many workers, including Bütschli (1876), Maupas (1889), Calkins and Cull (1907), and others. Briefly the process is as follows: Two similar individuals come in contact on their oral surface (a). The micronucleus in each conjugant divides twice (b–e), forming four micronuclei, three of which degenerate and do not take active part during further changes (f–h). The remaining micronucleus divides once more, producing a wandering pronucleus and a stationary pronucleus (f, g). The wandering pronucleus in each of the conjugants enters the other individual and fuses with its stationary pronucleus (h, r). The two conjugants now separate from each other and become exconjugants. In each exconjugant, the synkaryon divides three times in succession (i–m) and produces eight nuclei (n), four of which remain as micronuclei, while the other four develop into new macronuclei (o). Cytoplasmic fission follows then, producing first, two individuals with four nuclei (p) and then, four small individuals, each containing a micronucleus and a macronucleus (a). Jennings maintained that of the four smaller nuclei formed in the exconjugant (o), only one remains active and the other three degenerate. This active nucleus divides prior to the cytoplasmic divi-
Fig. 83. Diagram illustrating the conjugation of *Paramecium caudatum*. a-q, × about 130 (Calkins); r, a synkaryon formation as in h, ×1200 (Dehorne).
sion so that in the next stage (p), there are two developing macronuclei and one micronucleus which divides once more before the second and last cytoplasmic division (q). During these changes, the original macronucleus disintegrates, degenerates, and finally becomes absorbed in the cytoplasm.

Although this is the general course of events in the conjugation of this ciliate, recent observations revealed a number of different nuclear behavior. For example, there may not be pronuclear exchange between the conjugants (cytogamy, p. 204), thus resulting in self fertilization (Diller, 1950a). In a number of races, Diller (1950) found that one of the two nuclei produced by the first division of the synkaryon degenerates, while the other nucleus divides three times, forming 8 nuclei, and furthermore, an exconjugant may conjugate occasionally with another individual before the reorganization has been completed.

The conjugation of P. bursaria has also received attention of many workers. According to Chen (1946a), the first micronuclear division is a long process. One daughter nucleus degenerates and the other undergoes a second division. Here again one nucleus degenerates, while the other divides once more, giving rise to a wandering and a stationary pronucleus. Exchange of the wandering pronuclei is followed by the fusion of the two pronuclei in each conjugant. The synkaryon then divides. One of the two nuclei formed by this division degenerates, while the other gives rise to four nuclei by two divisions. The latter presently become differentiated into two micronuclei and two macronuclei, followed by a cytoplasmic division. The time two conjugants remain paired is said to be 20–38 or more hours (Chen, 1946c). In this Paramecium also, various nuclear activities have been reported. Chen (1940a, e) found that conjugation between a micronucleate and an amicronucleate can sometimes occur. In such a case, the micronucleus in the normal individual divides three times, and one of the pronuclei migrates into the amicronucleate in which there is naturally no nuclear division. The single haploid nucleus ("hemicaryon") in each individual divides three times as mentioned above and four nuclei are produced. Thus amicronucleate becomes micronucleated. Conjugating pairs sometimes separate from each other in a few hours. Chen (1946c) found that when such pairs are kept in a depression slide, temporary pairing recurs daily for many days, though there is seemingly no nuclear change. Chen (1940) further observed that the micronucleus in this species is subject to variation in size and
in the quantity of chromatin it contains, which gives rise to different (about 80 to several hundred) chromosome numbers during conjugation in different races, and that polyploidy is not uncommon in this ciliate. This investigator considers that polyploidy is a result of fusion of more than two pronuclei which he observed on several occasions. The increased number of pronuclei in a conjugant may be due to: (1) the failure of one of the two nuclei produced by the first or second division to degenerate; (2) the conjugation between a unimicronucleate and a bimicronucleate, or (3) the failure of the wandering pronucleus to enter the other conjugant; with this latter view Wichterman (1946) agrees. Apparently polyploidy occurs in other species also; for example, in *P. caudatum* (Calkins and Cull, 1907; Penn, 1937).

In *P. trichium*, Diller (1948) reported that the usual process of conjugation is the sequence of three micronuclear divisions, producing the pronuclei (during which degeneration of nuclei may occur at the end of both the first and second divisions), cross- or self-fertilization and three divisions of the synkarya. Ordinarily four of the eight nuclei become macronuclei, one remains as the micronucleus and the other three degenerate. The micronucleus divides at each of the two cytoplasmic divisions. Exchange of strands of the macronuclear skein may take place between the conjugants. Diller found a number of variations such as omission of the third prefertilization division, autogamous development, etc., and remarked that heteroploidy is pronounced and common.

In *P. aurelia* possessing typically two micronuclei, the process of conjugation was studied by Maupas (1889), Hertwig (1889), Diller (1936), Sonneborn (1947), etc., and is as follows: Soon after bi-association begins, the two micronuclei in each conjugant divide twice and produce eight nuclei, seven of which degenerate, while the remaining one divides into two gametic nuclei (Maupas, Woodruff, Sonneborn) Diller notes that two or more of the eight nuclei divide for the third time, but all but two degenerate; the two gametic nuclei may or may not be sister nuclei. All agree that there are two functional pronuclei in each conjugant. As in other species of *Paramaecium* already noted, there is a nuclear exchange which results in the formation of a synkaryon in each conjugant. The synkaryon divides twice and the conjugants separate from each other at about this time. Two nuclei develop into macronuclei and the other two into micronuclei. Prior to the first cytoplasmic division of the exconjugant, the micronuclei divide once, but the macronucleus does not divide, so that each of the two daughters receives one macronucleus
and two micronuclei. The original macronucleus in the conjugant becomes transformed into a skein which breaks up into 20 to 40 small masses. These are resorbed in the cytoplasm as in other species. As to when these nuclear fragments are absorbed, depends upon the nutritive condition of the organism (Sonneborn); namely, under a poor nutritional condition the resorption begins and is completed early, but under a better condition this resorption takes place after several divisions.

During conjugation reciprocal migration of a pronucleus thus occurs in all cases. During biassociation and even in autogamy (p. 203), there develops a conical elevation ("paroral cone") and the nuclear migration takes place through this region. Although there is ordinarily no cytoplasmic exchange between the conjugants, this may occur in some cases as observed by Sonneborn (1943a, 1944). _P. aurelia_ of variety 4, according to Sonneborn, do occasionally not separate after fertilization, but remain united by a thin strand in the region of the paroral cones. In some pairs, the strand enlarges into a broad band through which cytoplasm flows from one individual to the other. The first division gives off a normal single animal from each of the "parabiotic twins" and the two clones derived from the two individuals belong to the same mating type (p. 192).

Conjugation between different species of _Paramecium_ has been attempted by several workers. Müller (1932) succeeded in producing a few pairings between normal _P. caudatum_ and exconjugant _P. multimicronucleatum_. The nuclear process ran normally in caudatum, which led Müller to believe that crossing might be possible, but without success. De Garis (1935) mixed "double animals" (p. 228) of _P. caudatum_ and conjugating population of _P. aurelia_. Pairing between them occurred readily, in which the aurelia mates remained attached to caudatum for five to 12 hours. Four pairs remained together, but aurelia underwent cytolysis on the second day. The separated aurelia from other pairs died after showing "cloudy swelling" on the second or third day after biassociation. The caudatum double-animals on the other hand lived for two to 12 (average six) days during which there was neither growth nor division and finally perished after "hyaline degeneration." No information on nuclear behavior in these animals is available. Apparently, the different species of _Paramecium_ are incompatible with one another.

In 1937, Sonneborn discovered that in certain races of _P. aurelia_, there are two classes of individuals with respect to "sexual" differentiation and that the members of different classes conjugate with each other, while the members of each class do not. The members of
a class or caryonide (Sonneborn, 1939) are progeny of one of the two individuals formed by the first division of an exconjugant and thus possess the same macronuclear constitution. These classes were designated by Sonneborn (1938) as mating types. Soon a similar phenomenon was found by several workers in other species of Para-

![Fig. 84. Mating behavior of Paramecium bursaria (Jennings).](image)

...
large masses break down into small masses (c) and still later, conjugants appear in pairs (d). The only other ciliate in which mating types are definitely known to occur is *Euplotes patella* in which, according to Kimball (1939), there occurs no agglutination mating reaction.

How widely mating types occur is not known at present. But as was pointed out by Jennings, the mating types may be of general occurrence among ciliates; for example, Maupas (1889) observed that in *Lionotus* (*Loxophyllum*) *fasciola*, *Leucophrys patula*, *Stylonychia pustulata*, and *Onychodromus grandis*, conjugation took place between the members of two clones of different origin, and not among the members of a single clone. Precise information on the occurrence of mating types among different ciliates depends on future research.

In *Paramecium aurelia*, Sonneborn distinguishes seven varieties which possess the same morphological characteristics of the species, but which differ in addition to mating types, also in size, division rate, conditions of temperature and light under which mating reaction may occur, etc. (Sonneborn, 1947). There occurs ordinarily no conjugation between the clones of different varieties. Within each of six varieties, there are two mating types, while there is only one type in the seventh variety. Animals belonging to the same variety, but to different mating types, only conjugate when put together (Table 6).

Under optimum breeding conditions two mating types of the same variety give 95 per cent immediate agglutination and conjugation. But exceptions occur. Sonneborn and Dipell (1946) place the 7 varieties of *aurelia* under two groups: A (varieties 1, 3, 5 and 7) and B (varieties 2, 4 and 6) on the basis of their conjugational reactions. Mating types in group A do not conjugate with those of group B; no mating type of group B is known to conjugate with any type of other varieties in this group; but a number of combinations of mating types belonging to different varieties of group A conjugate with each other. For example, varieties 1 and 5 conjugate (namely, type I with type X and type II with type IX); however these intervarietal mating reactions are (1) always less intense than intravarietal reaction, (2) dependent upon the degree of reactivity of the culture, and (3) different from the intravarietal reaction with respect to the conditions for optimum reaction. Furthermore in most cases, the progeny of intervarietal matings are not viable. In the varieties of group A, the mating types appear to be of a more general sort. Therefore, Sonneborn (1947) designated even- and odd-numbered types as + and — respectively.
In *P. bursaria*, Jennings (1938, 1939) found three varieties. Varieties 1 and 3 contain 4 mating types each, while variety 2, eight mating types. Jennings and Opitz (1944) further found variety 4 (Russian), composed of two mating types and variety 5 under which several Russian clones were placed. Chen (1946a) added variety 6 (originating in Europe) containing four mating types. Thus in this species of Paramecium, there are now six varieties, containing 23 mating types (Table 7), and mating reaction occurs even among enucleate fragments of animals of different mating types of the same variety (Tartar and Chen, 1941). In *Euplotes patella*, Kimball (1939) observed six mating types which he designated as type I to type VI (Table 8).

Though the members of a clone are of the same mating type and therefore do not conjugate, a clone may undergo at very long intervals (some 2000 culture days), "self-differentiation" into two mating types which then conjugate (Jennings, 1941). Furthermore, Jennings

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Table 7.—Varieties and mating types in Paramecium bursaria
(Jennings; Jennings and Opitz; Chen)

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and Opitz (1944) found that mating type R (variety 4) conjugated with E, K, L or M (variety 2), but all conjugants or exconjugants perished without multiplication. Chen (1946a) made a cytological study of them and observed that the nuclear changes which are

Table 8.—Mating types in Euplotes patella (Kimball)

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seemingly normal during the first 16 hours, become abnormal suddenly after that time, and the micronuclei divide only once and there is no nuclear exchange. The death of conjugants or exconjugants is possibly due to physiological incompatibility between the varieties upon coming in contact or probably due to "something that diffuses from one conjugant to the other."

Studies of mating types have revealed much information regarding conjugation. Conjugation usually does not occur in well-fed or extremely starved animals, and appears to take place shortly after the depletion of food. Temperature also plays a rôle in conjugation, as it takes place within a certain range of temperature which varies even in a single species among different varieties (Sonneborn). Light seems to have different effects on conjugation in different varieties of P. aurelia. The time between two conjugations also varies in different species and varieties. In P. bursaria, Jennings found that in some races the second conjugation would not take place for many months after the first, while in others such an "immature" period may be only a few weeks. In P. aurelia, in some varieties there is no "immature" period, while in others there is 6 to 10 days' "immaturity."

Very little is known about the physiological state of conjugants as compared with vegetative individuals. Several investigators observed that animals which participate in conjugation show much viscous body surface. Boell and Woodruff (1941) found that the mating individuals of Paramecium calcinski show a lower respiratory rate than not-mating individuals. Neither is the mechanism of conjugation understood at present. Kimball (1942) discovered in Euplotes patella, the fluid taken from cultures of animals of one type induces conjugation among the animals of other types (p. 235). Presumably certain substances are secreted by the organisms and become diffused in the culture fluid. In Paramecium aurelia, Sonneborn (1943) found that of the four races of variety 4, race 51 was a "killer," while the other three races, "sensitive." Fluid in which the killer race grew, kills the individuals of the sensitive races. As has been mentioned already, P. bursaria designated as type T (variety 5) (Table 7) conjugates with none. But Chen (1945) found that its culture fluid induces conjugation among a small number of the individuals of one mating type of varieties 2, 3, 4 and 6, in which nuclear changes proceed as in normal conjugation. Furthermore, this fluid is capable of inducing autogamy in single animals. Other visible influences of the fluid on organisms are sluggishness of movement and darker coloration and distortion of the body.
Boell and Woodruff (1941) noticed that in *P. calkinsi*, living individuals of one mating type will agglutinate with dead ones of the complementary mating type. A similar phenomenon was also observed by Metz (1946, 1947, 1948) who employed various methods of killing the animals. The pairs composed of living and formaldehyde-killed animals, behave much like normal conjugating pairs; there is of course no cross-fertilization, but the living member of the pair undergoes autogamy. While the "mating type substances" can be destroyed by exposure to 52°C, for five minutes; by X-irradiation; by exposure of formaldehyde-killed reactive animals to specific antisera or to 100°C, etc., Metz demonstrated that animals may be killed by many reagents which do not destroy these substances. Furthermore, all mating activities disappear when the animals are thoroughly broken up, which suggests that Paramecium might release some mating substance inhibitory agent. This agent was later found in this Paramecium (Metz and Butterfield, 1950). Metz (1948) points out that the mating reaction involves substances present on the surfaces of the cilia, and supposes that the interaction between two mating-type substances initiates a chain of reactions leading up to the process of conjugation and autogamy. Hiwataishi (1949a, 1950) using four groups (each composed of two mating types) of *P. caudatum*, confirmed Metz's observation. Metz and Butterfield (1951) more recently report that non-proteolytic enzymes (lecithinase, hyaluronidase, lysozyme, ptyalin, ribonuclease) have no detectable effect on the mating reactivity of *P. calkinsi*; but proteolytic enzymes such as trypsin and chymotrypsin destroy the mating reactivity, and mating substance activity was not found in the digest of enzyme-treated organisms. The two observers believe that the mating reactivity is dependent upon protein integrity.

When the ciliate possesses more than one micronucleus, the first division ordinarily occurs in all and the second may or may not take place in all, varying apparently even among individuals of the same species. This seems to be the case with the majority, although more than one micronucleus may divide for the third time to produce several pronuclei, for example, two in *Euplotes patella*, *Stylonychia pustulata*; two to three in *Oxytricha fallax* and two to four in *Uroleptus mobilis*. This third division is often characterized by long extended nuclear membrane stretched between the division products.

Ordinarily the individuals which undergo conjugation appear to be morphologically similar to those that are engaged in the trophic activity, but in some species, the organism divides just prior to
Fig. 85. The life-cycle of *Nyctotherus cordiformis* in *Hyla versicolor* (Wichterman). a, a cyst; b, excystment in tadpole; c, d, division is repeated until host metamorphoses; e, smaller preconjugant; f–j, conjugation; k, exconjugant; l, amphinucleus divides into 2 nuclei, one micronucleus and the other passes through the “spireme ball” stage before developing into a macronucleus; k–n, exconjugants found nearly exclusively in recently transformed host; o, mature trophozoite; p–s, binary fission stages; t, precystic stage.
conjugation. According to Wichterman (1936), conjugation in *Nyciototherus cordiformis* (Fig. 85) takes place only among those which live in the tadpoles undergoing metamorphosis (f–j). The conjugants are said to be much smaller than the ordinary trophozoites, because of the preconjugation fission (d–e). The micronuclear divisions are similar to those that have been described for *Paramaecium caudatum* and finally two pronuclei are formed in each conjugant. Exchange and fusion of pronuclei follow. In each exconjugant, the synkaryon divides once to form the micronucleus and the macronuclear anlage (k–l) which develops into the “spireme ball” and finally into the macronucleus (m–o).

A sexual process which is somewhat intermediate between the sexual fusion and conjugation, is noted in several instances. According to Maupas’ (1888) classical work on *Vorticella nebulifera*, the ordinary vegetative form divides twice, forming four small individuals, which become detached from one another and swim about independently. Presently each becomes attached to one side of a stalked individual. In it, the micronucleus divides three times and produces eight nuclei, of which seven degenerate; and the remaining nucleus divides once more. In the stalked form the micronucleus divides twice, forming four nuclei, of which three degenerate, and the other dividing into two. During these changes the two conjugants fuse completely. The wandering nucleus of the smaller conjugant unites with the stationary nucleus of the larger conjugant, the other two pronuclei degenerating. The synkaryon divides several times to form a number of nuclei, from some of which macronuclei are differentiated and exconjugant undergoes multiplication. In *Vorticella microstoma* (Fig. 86), Finley (1943) notes that a vegetative individual undergoes unequal division except the micronucleus which divides equally (a), and forms a large stalked macroconjugant and a small free microconjugant (b). The conjugation which requires 18–24 hours for completion, begins when a microconjugant attaches itself to the lower third of a macroconjugant. The protoplasm of the microconjugant enters the macroconjugant (c). The micronucleus of the microconjugant divides three times, the last one of which being reductional (d, e), while that of the macroconjugant divides twice (one mitotic and one meiotic). Fusion of one of each produces a synkaryon (f) which divides three times. One of the division products becomes a micronucleus and the other seven macronuclear anlagen (g, h) which are distributed among the progeny (i, j).

Another example of this type has been observed in *Metopus es*
(Fig. 87). According to Noland (1927), the conjugants fuse along the anterior end (a), and the micronucleus in each individual divides in the same way as was observed in *Paramecium caudatum* (b–e). But the cytoplasm and both pronuclei of one conjugant pass into the other (f), leaving the degenerating macronucleus and a small

Fig. 86. Sexual reproduction in *Vorticella microstoma*, ×800 (Finley). a, preconjugation division which forms a macroconjugant and a microconjugant; b, a macroconjugant with three microconjugants; c, a microconjugant fusing with a macroconjugant; d, the micronucleus of the microconjugant divided into four nuclei; e, with 12 nuclei formed by divisions of the two micronuclei of conjugants; f, synkaryon; g, eight nuclei after three divisions of synkaryon; h, seven enlarging macronuclear anlagen and a micronucleus in division; i, first division; j, a daughter individual with a micronucleus, four macronuclear anlagen, and old macronuclear fragments.
amount of cytoplasm behind in the shrunken pellicle of the smaller conjugant which then separates from the other (j). In the larger exconjugant, two pronuclei fuse, and the other two degenerate and disappear (g, h). The synkaryon divides into two nuclei, one of which condenses into the micronucleus and the other grows into the macronucleus (i, k–m). This is followed by the loss of cilia and encystment.

While ordinarily two individuals participate in conjugation, three

Fig. 87. Conjugation of *Metopus es* (Noland). a, early stage; b, first micronuclear division; c, d, second micronuclear division; e, third micronuclear division; f, migration of pronuclei from one conjugant into the other; g, large conjugant with two pronuclei ready to fuse; h, large conjugant with the synkaryon, degenerating pronuclei and macronucleus; i, large exconjugant with newly formed micronucleus and macronucleus; j, small exconjugant with degenerating macronucleus; k–m, development of two nuclei. a, ×290; b–j, ×250, k–m, ×590.
or four individuals are occasionally involved. For example, conjugation of three animals was observed in *P. caudatum* by Stein (1867), Jickeli (1884), Maupas (1889) and in *Blepharisma undulans* by Giese (1938) and Weisz (1950). Chen (1940b, 1948) made a careful study of such a conjugation which he found in *Paramecium burs-*

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**Fig. 88.** Conjugation of three individuals in *Paramecium bursaria*, ×365 (Chen). a, late prophase of the first nuclear division (the individual on right is a member of a race with "several hundred chromosomes," while the other two belong to another race with "about 80 chromosomes"); b, anaphase of the third division (each individual contains 2 degenerating nuclei); c, beginning of pronuclear exchange between two anterior animals; d, e, synkaryon formation; f, after the first division of synkaryon, one daughter nucleus undergoing degeneration in all animals.
saria (Fig. 88). He found that the usual manner of association is conjugation between a pair with the third conjugant attached to the posterior part of one of them (a). Nuclear changes occur in all three individuals, and in each, two pronuclei are formed by three divisions (c). But the exchange of the pronuclei takes place only between two anterior conjugants (c-e) and autogamy (see below) occurs in the third individual.

![Diagram illustrating autogamy in Paramecium aurelia (Diller).](image)

Fig. 89. Diagram illustrating autogamy in *Paramecium aurelia* (Diller). a, normal animal; b, first micronuclear division; c, second micronuclear division; d, individual with 8 micronuclei and macronucleus preparing for skein formation; e, two micronuclei dividing for the third time; f, two gamete-nuclei formed by the third division in the paroral cone; g, fusion of the nuclei, producing synkaryon; h, i, first and second division of synkaryon; j, with 4 nuclei, 2 becoming macronuclei and the other 2 remaining as micronuclei; k, macronuclei developing, micronuclei dividing; l, one of the daughter individuals produced by fission.

**Automixis.** In certain Protozoa, the fusion occurs between two nuclei which originate in a single nucleus of an individual. This process has been called automixis by Hartmann, in contrast to the amphimixis (Weismann) which is the complete fusion of two nuclei originating in two individuals, as was discussed in the preceding pages. If the two nuclei which undergo a complete fusion are present in a single cell, the process is called **autogamy**, but, if they are in two
different cells, then **paedogamy**. The autogamy is of common occurrence in the myxosporidian spores. The young sporoplasm contains two nuclei which fuse together prior to or during the process of germination in the alimentary canal of a specific host fish, as for example in *Sphaeromyxa sabrazesi* (Figs. 276; 277) and *Myxosoma catostomi* (Fig. 275). In the Microsporidia, autogamy appears to initiate the spore-formation at the end of schizogonic activity of individuals as in *Theclohania legeri* (Fig. 76).

Diller (1936) observed in solitary *Paramecium aurelia* (Fig. 89), certain micronuclear changes similar to those which occur in conjugating individuals. The two micronuclei divide twice, forming eight nuclei (a–d), some of which divide for the third time (e), producing two functional and several degenerating nuclei (f). The two functional nuclei then fuse in the "paroral cone" and form the synkaryon (g, h) which divides twice into four (i, j). The original macronucleus undergoes fragmentation and becomes absorbed in the cytoplasm. Of the four micronuclei, two transform into the new macronuclei and two remain as micronuclei (k) each dividing into two after the body divided into two (l).

Another sexual process appears to have been observed by Diller (1934) in conjugating *Paramecium trichium* in which there was no nuclear exchange between the two conjugants. Wichterman (1940) observed a similar process in *P. caudatum* and named it **cytogamy**. Two small (about 200μ long) individuals of *P. caudatum* fuse on their oral surfaces. There occur three micronuclear divisions as in the case of conjugation, but there is no nuclear exchange between the members of the pair. The two gametic nuclei in each individual are said to fuse and form a synkaryon as in autogamy. Sonneborn (1941) finds the frequency of cytogamy in *P. aurelia* to be correlated with temperature. At 17°C., conjugation occurs in about 95 per cent of the pairs and cytogamy in about 5 per cent; but at 10° and 27°C., cytogamy takes place in 47 and 60 per cent respectively. In addition, there is some indication that sodium decreases and calcium increases the frequency of occurrence of cytogamy.

The paedogamy occurs in at least two species of Myxosporidia, namely, *Leptotheca ohlmacheri* (Fig. 279) and *Unicapsula muscularis* (Fig. 280). The spores of these myxosporidians contain two uninucleate sporplasms which are independent at first, but prior to emergence from the spore, they undergo a complete fusion to metamorphose into a uninucleate amoeoba. Perhaps the classical example of the paedogamy is that which was found by Hertwig (1898) in *Actinosphaerium eichhorni*. The organism encysts and the body di-
vides into numerous uninucleate secondary cysts. Each secondary cyst divides into two and remains together within a common cyst-wall. In each the nucleus divides twice, and forms four nuclei, one of which remains functional, the remaining three degenerating. The paedogamy results in formation of a zygote in place of a secondary cyst. Bělař (1923) observed a similar process in Actinophrys sol (Fig. 90). This heliozoan withdraws its axopodia and divides into two uninucleate bodies which become surrounded by a common gelatinous envelope. Both nuclei divide twice and produce four nuclei, three of which degenerate. The two daughter cells, each with one haploid nucleus, undergo paedogamy and the resulting individual now contains a diploid nucleus.

In Paramecium aurelia, Diller (1936) found simple fragmentation of the macronucleus which was not correlated with any special micronuclear activity and which could not be stages in conjugation or autogamy. Diller suggests that if conjugation or autogamy is to create a new nuclear complex, as is generally held, it is conceivable that somewhat the same result might be achieved by "purification act" (through fragmentation) on the part of the macronucleus itself,
without involving micronuclei. He coined the term **hemixis** for this reorganization.

**Meiosis.** In the foregoing sections, references have been made to the divisions which the nuclei undergo prior to sexual fusion or conjugation. In all Metazoa, during the development of the gametes, the gametocytes undergo reduction division or meiosis, by which the number of chromosomes is halved; that is to say, each fully mature gamete possesses half (haploid) number of chromosomes typical of the species (diploid). In the zygote, the diploid number is reestablished. In the Protozoa in which sexual reproduction occurs during their life-cycle, meiosis presumably takes place to maintain the constancy of chromosome-number, but the process is understood only in a small number of species.

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**Fig. 91.** Mitotic and meiotic micronuclear divisions in conjugating *Didinium nasutum.* (Prandtl, modified). *a,* normal micronucleus; *b,* equatorial plate in the first (mitotic) division; *c,* anaphase in the first division; *d,* equatorial plate in the second division; *e,* anaphase in the second (meiotic) division.

In conjugation, the meiosis seems to take place in the second micronuclear division, although in some, for example, *Oxytricha fallax,* according to Gregory, the actual reduction occurs during the first division. Prandtl (1906) was the first to note a reduction in number of chromosomes in the Protozoa. In conjugating *Didinium nasutum* (Fig. 91), he observed 16 chromosomes in each of the daughter micronuclei during the first division, but only 8 in the second division. Since that time, the fact that meiosis occurs during the second micronuclear division has been observed in *Chilodonella uncinata* (Enrique; MacDougall), *Carchesium polypinum* (Popoff), *Uroleptus halseyi* (Calkins), etc. (note the ciliates in Table 5 on p. 168). In various species of Paramecium and many other forms, the number of chromosomes appears to be too great to allow a precise counting, but the observations of Sonneborn, as quoted elsewhere (p. 234) and of Jennings (1942) on *P. aurelia and P. bursaria* respec-
tively, indicate clearly the occurrence of meiosis prior to nuclear exchange during conjugation.

Information on the meiosis involved in the complete fusion of gametes is even more scanty and fragmentary. In *Monocystis rostrata* (Fig. 92), a parasite of the earthworm, Mulsow (1911) noticed that

Fig. 92. Mitosis and meiosis in *Monocystis rostrata* (Mulsow). a–g, mitosis; h–j, meiosis. a, a resting nucleus in the gametocyte; b, development of chromosomes; c, polar view of equatorial plate; d, longitudinal splitting of eight chromosomes; e, separation of chromosomes in two groups; f, late anaphase; g, two daughter nuclei; h, i, polar view of the equatorial plate in the last division; j, anaphase, the gamete nucleus is now haploid (4). a–c, ×1840; d–g, ×1400; h–j, ×3000.

the nuclei of two gametocytes which encyst together, multiply by mitosis in which eight chromosomes are constantly present (a–g), but in the last division in gamete formation, each daughter nucleus receives only 4 chromosomes (h–j). In another species of *Monocystis*, Calkins and Bowling (1926) observed that the diploid number of chromosomes was 10 and that haploid condition is established in the last gametic division thus confirming Mulsow’s finding.

In the paedogamy of *Actinophrys sol* (Fig. 90), Bělař (1923) finds 44 chromosomes in the first nuclear division, but after two meiotic divisions, the remaining functional nucleus contains only 22 chromosomes so that when paedogamy is completed the diploid number is restored. In *Polytoma wella*, Moewus finds each of the two gametes is haploid (8 chromosomes) and the zygotes are diploid. The syn-karyon divides twice, and during the first division reduction division takes place.
In the coccidian, *Aggregata eberthi* (Fig. 246), according to Dobell (1925), Naville (1925) and Bélař (1926) and in the gregarine, *Diplonychis schneideri*, according to Jameson (1920), there is no reduction in the number of chromosomes during the gamete-formation, but the first zygotic division is meiotic, 12 to 6 and 6 to 3, respectively. A similar reduction takes place also in *Actinocephalus parvus* (8 to 4, after Weschenfelder, 1938), *Gregarina blattarum* (6 to 3, after Sprague, 1941), *Adelina deronis* (20 to 10, after Hauschka, 1943), etc. *Trichonympha* and other flagellates (p. 185) of woodroach, *Polytoma*

![Diagram](image_url)

**Fig. 93. Degeneration or aging in *Stylonychia pustulata*. ×340 (Maupas, modified).** a, Beginning stage with reduction in size and completely atrophied micronucleus; b, c, advanced stages in which disappearance of the frontal zone, reduction in size, and fragmentation of the macronucleus occurred; d, final stage before disintegration.

and *Chlamydomonas* (p. 276) also undergo postzygotic meiosis. Thus in these organisms, the zygote is the only stage in which the nucleus is diploid.

Some seventy years ago Weismann pointed out that a protozoan grows and multiplies by binary fission or budding into two equal or unequal individuals without loss of any protoplasmic part and these in turn grow and divide, and that thus in Protozoa there is neither senescence nor natural death which occur invariably in Metazoa in which germ and soma cells are differentiated. Since that time, the problem of potential immortality of Protozoa has been a matter which attracted the attention of numerous investigators. Because of large dimensions, rapid growth and reproduction, and ease with
which they can be cultivated in the laboratory, the majority of Protozoa used in the study of the problem have been free-living freshwater ciliates that feed on bacteria and other microorganisms.

The very first extended study was made by Maupas (1888) who isolated *Stylonychia pustulata* on February 27, 1886, and observed 316 binary fissions until July 10. During this period, there was noted a gradual decrease in size and increasing abnormality in form and structure, until the animals could no longer divide and died (Fig. 93). A large number of isolation culture experiments have since been carried on numerous species of ciliates by many investigators. The results obtained are not in agreement. However, the bulk of obtained data indicates that the vitality of animals decreases with the passing of generations until finally the organisms suffer inevitable death, and that in the species in which conjugation or other sexual reproduction occurs, the declining vitality often becomes restored. Perhaps the most thorough experiment was carried on by Calkins (1919, 1933) with *Uroleptus mobilis*. Starting with an exconjugant on November 17, 1917, a series of pure-line cultures was established by the daily isolation method. It was found that no series lived longer than a year, but when two of the progeny of a series were allowed to conjugate after the first 75 generations, the exconjugants repeated the history of the parent series, and did not die when the parent series died. In this way, lines of the same organism have lived for more than 12 years, passing through numerous series. In a series, the average division for the first 60 days was 15.4 divisions per 10 days, but the rate gradually declined until death. Woodruff and Spencer (1924) also found the isolation cultures of *Spathidium spathula* (fed on *Colpidium colpoda*) died after a gradual decline in the division rate, but were inclined to think that improper environmental conditions rather than internal factors were responsible for the decline.

On the other hand, Woodruff (1932) found that 5071 generations produced by binary fission from a single individual of *Paramecium aurelia* between May 1, 1907 and May 1, 1915, did not manifest any decrease in vitality after eight years of continued asexual reproduction. Other examples of longevity of ciliates without conjugation are: Glaucoma for 2701 generations (Enriques, 1916), *Paramecium caudatum* for 3967 generations (Metalnikov, 1922), *Spathidium spathula* for 1080 generations (Woodruff and Moore, 1924), *Didinium nasutum* for 1384 generations (Beers, 1929), etc. With *Actinophrys sol*, Bělař (1924) carried on isolation cultures for 1244 generations for a period of 32 months and noticed no decline in the division rate.
Hartmann (1921) made a similar observation on *Eudorina elegans*. It would appear that in these forms, the life continues indefinitely without apparent decrease in vital activity.

As has been noted in the beginning part of the chapter, the macronucleus in the ciliates undergoes, at the time of binary fission, a reorganization process before dividing into two parts and undoubted- edly, there occurs at the same time extensive cytoplasmic reorganiza- tion as judged by the degeneration and absorption of the old, and formation of the new, organellae. It is reasonable to suppose that this reorganization of the whole body structure at the time of divi- sion is an elimination process of waste material accumulated by the organism during the various phases of vital activities as was con- sidered by Kidder and others (p. 150) and that this elimination, though not complete, enables the protoplasm of the products of divi- sion to carry on their metabolic functions more actively.

As the generations are multiplied, the general decline in vitality is manifest not only in the decreased division-rate, slow growth, abnormal form and function of certain organellae, etc., but also in inability to complete the process involved in conjugation. Jennings (1944) distinguished four successive periods in various clone cultures of *Paramecium bursaria*; namely, (1) a period of sexual immaturity during which neither sexual reaction nor conjugation occurs; (2) a period of transition during which weak sexual reactions appear in a few individuals; (3) a period of maturity in which conjugation takes place readily when proper mating types are brought together; and (4) a period of decline, ending in death. The length of the first two periods depends on the cultural conditions. Exconjugant clones that are kept in condition under which the animals multiply rapidly, reach maturity in three to five months, while those subjected to de- pressing condition require 10 to 14 months to reach maturity. The third period lasts for several years and is followed by the fourth period during which fission becomes slower, abnormalities appear, many individuals die and the clones die out completely.

Does conjugation affect the longevity of clones in *Paramecium bursaria*? A comparative study of the fate of exconjugants and non- conjugants led Jennings (1944a) to conclude that (1) conjugation results in production of one of the following four types: (a) excon- jugants perish without division, (b) exconjugants divide one to four times and then die, (c) exconjugants produce weak abnormal clones which may become numerous, and (d) exconjugants multiply vigor- ously and later undergo conjugation again; at times the latter are
more vigorous than the parent clones, thus showing rejuvenescence through conjugation; (2) conjugation of young clones results in little or no mortality, while that of old clones results in high (often 100 per cent) mortality; (3) conjugation between a young and an old clone, results in the death of most or all of the exconjugants; (4) the two members of a conjugating pair have the same fate; and (5) what other causes besides age bring about the death, weakness or abnormality of the exconjugants, are not known.

It is probable that the process of replacing old macronuclei by micronuclear material which are derived from the products of fusion of two micronuclei of either the same (autogamy) or two different animals (conjugation), would perhaps result in a complete elimination of waste substances from the newly formed macronuclei, and divisions which follow this fusion may result in shifting the waste substances unequally among different daughter individuals. Thus in some individuals there may be a complete elimination of waste material and consequently a restored high vitality, while in others the influence of waste substances present in the cytoplasm may offset or handicap the activity of new macronuclei, giving rise to stocks of low vitality which will perish sooner or later. In addition in conjugation, the union of two haploid micronuclei produces diverse genetic constitutions which would be manifest in progeny in manifold ways. Experimental evidences indicate clearly such is actually the case.

In many ciliates, the elimination of waste substances at the time of binary fission and sexual reproduction (conjugation, and autogamy), seemingly allow the organisms continued existence through a long chain of generations indefinitely. Jennings (1929, 1942) who reviewed the whole problem states: "Some Protozoa are so constituted that they are predestined to decline and death after a number of generations. Some are so constituted that decline occurs, but this is checked or reversed by substitution of reserve parts for those that are exhausted; they can live indefinitely, but are dependent on this substitution. In some the constitution is such that life and multiplication can continue indefinitely without visible substitution of a reserve nucleus for an exhausted one; but whether this is due to the continued substitution, on a minute scale, of reserve parts for those that are outworn cannot now be positively stated. This perfected condition, in which living itself includes continuously the necessary processes of repair and elimination, is found in some free cells, but not in all."
Regeneration

The capacity of regenerating the lost parts, though variable among different species, is characteristic of all Protozoa from simple forms to those with highly complex organizations, as shown by observations of numerous investigators. It is now a well established fact that when a protozoan is cut into two parts and the parts are kept under proper environmental conditions, the enucleate portion is able to carry on catabolic activities, but unable to undertake anabolic activities, and consequently degenerates sooner or later. Brandt (1877) studied regeneration in Actinosphaerium eichhorni and found that only nucleate portions containing at least one nucleus regenerated and enucleate portions or isolated nuclei degenerated. Similarly Gruber (1886) found in Amoeba proteus the nucleate portion regenerated completely, while enucleate part became rounded and perished in a few days. The parts which do not contain nuclear material may continue to show certain metabolic activities such as locomotion, contraction of contractile vacuoles, etc., for some time; for example, Grosse-Allermann (1909) saw enucleate portions of Amoeba verrucosa alive for 20 to 25 days, while Stole (1910) found enucleate Amoeba proteus living for 30 days. Clark (1942, 1943) showed that Amoeba proteus lives for about seven days after it has been deprived of its nucleus. Enucleated individuals show a 70 per cent depression of respiration and are unable to digest food due to the failure of zymogens to be activated in the dedifferentiating cytoplasm. According to Brachet (1950), the enucleated half of an amoeba shows a steady decrease in ribonucleic acid content, while the nucleated half retains a much larger amount of this substance. Thus it appears that the synthesis of the cytoplasmic particles containing ribonucleic acid is under the control of the nucleus.

In Arcella (Martini; Hegner) and Diffugia (Verworn; Penard), when the tests are partially destroyed, the broken tests remain unchanged. Verworn considered that in these testaceans test-forming activity of the nucleus is limited to the time of asexual reproduction of the organisms. On the other hand several observers report in Foraminifera the broken shell is completely regenerated at all times. Verworn pointed out that this indicates that here the nucleus controls the formation of shell at all times. In a radiolarian, Thalassicolla nucleata, the central capsule, if dissected out from the rest of body, will regenerate into a complete organism (Schneider). A few regeneration studies on Sporozoa have not given any results to be considered here, because of the difficulties in finding suitable media for cultivation in vitro.
An enormous number of regeneration experiments have been conducted on more than 50 ciliates by numerous investigators. Here also the general conclusion is that the nucleus is necessary for regeneration. In many cases, the macronucleus seems to be the only essential nucleus for regeneration, as judged by the continued division on record of several amicronucleate ciliates and by experiments such as Schwartz's in which there was no regeneration in Stentor coeruleus from which the whole macronucleus had been removed.

A remarkably small part of a protozoan is known to be able to regenerate completely if nuclear material is included. For example, Sokoloff found 1/53–1/69 of Spirostomum ambiguum and 1/70–1/75 of Dileptus anser regenerated and Phelps showed portions down to 1/80 of an amoeba were able to regenerate. In Stentor coeruleus, pieces as small as 1/27 (Lilly) or 1/64 (Morgan) of the original specimens or about 70μ in diameter (Weisz) regenerate. Burnside cut 27 specimens of this ciliate belonging to a single clone, into two or more parts in such a way that some of the pieces contained a large portion of the nucleus while others a small portion. These fragments regenerated and multiplied, giving rise to 268 individuals. No dimensional differences resulted from the different amounts of nuclear material present in the cut specimens. Apparently regulatory processes took place and in all cases normal size was restored, regardless of the amount of the nuclear material in ancestral pieces. Thus biotypes of diverse sizes are not produced by causing inequalities in the proportions of nuclear material in different individuals.

In addition to these restorative regenerations, there are physiological regenerations in which as in the case of asexual and sexual reproduction, various organellae such as cilia, flagella, cytostome, contractile vacuoles, etc., are completely regenerated. Information is now available on the process of morphogenesis in regeneration and reorganization in certain ciliates (Chatton and Lwoff, 1935; Balamuth, 1940; Summers, 1941; Fauré-Fremiet, 1948; Weisz, 1948, 1951).

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Chapter 6

Variation and heredity

IT IS generally recognized that individuals of all species of organism vary in morphological and physiological characteristics. Protozoa are no exception, and manifest a wide variation in size, form, structure, and physiological characters among the members of a single species. The different groups in a species are spoken of as the races, varieties, strains, etc. It is well known that dinoflagellates show a great morphological variation in different localities. Wesenberg-Lund (1908) noticed a definite seasonal morphological variation in Cerctium hirundinella in Danish lakes, while Schröder (1914) found at least nine varieties of this organism (Fig. 94) occurring in various bodies of water in Europe, and List (1913) reported that the organisms living in shallow ponds possess a marked morphological difference from those living in deep ponds. Cyphoderia ampulla is said to vary in size among those inhabiting the same deep lakes; namely, individuals from the deep water may reach 200μ in length, while those from the surface layer measure only about 100μ long.

In many species of Foraminifera, the shell varies in thickness according to the part of ocean in which the organisms live. Thus the strains which live floating in surface water have a much thinner shell than those that dwell on the bottom. For example, according to Rhumbler, Orbulina universa inhabiting surface water has a comparatively thin shell, 1.28–18μ thick, while individuals living on the bottom have a thick shell, up to 24μ in thickness. According to Uyemura, a species of Amoeba living in thermal waters, showed a distinct dimensional difference in different springs. It measured 10–40μ in diameter in sulphurous water and 45–80μ in ferrous water; in both types of water the amoebae were larger at 36–40°C. than at 51°C.

Such differences or varieties appear to be due to the influence of diverse environmental conditions, and will continue to exist under these conditions; but when the organisms of different varieties are subjected to a similar environment, the strain differences usually disappear sooner or later. That the differences in kind and amount of foods bring about extremely diverse individuals in Tetrahymena vorax and Chilomonas paramecium in bacteria-free cultures has already been mentioned (p. 109). Chlamydomonas debaryana are represented by many races differing in form, size, and structure, in various localities as well as under different laboratory conditions. Moewus
(1934) distinguished 12 such varieties and showed that any variety could be changed into another by using different culture media. This transformation, however, did not occur at the same rate among different races. It was found that the longer a strain has remained under

![Varieties of Ceratium hirundinella from various European waters (Schröder). a, furcoides-type (130–300μ by 30–45μ); b, brachyceroides-type (130–145μ by 30–45μ); c, silesiacum-type (148–280μ by 28–34μ); d, carinthiacum-type (120–145μ by 45–60μ); e, gracile-type (140–200μ by 60–75μ); f, austriacum-type (120–160μ by 45–60μ); g, robustum-type (270–310μ by 45–55μ); h, scotticum-type (160–210μ by 50–60μ); i, piburgense-type (180–260μ by 50–60μ).](image)

conditions producing a given type, the greater the time and the number of generations needed to change it to a new type under a new condition, as is shown in Table 9.

While in many species, the races or varieties have apparently been brought about into being under the influence of environmental conditions, in others the inherited characters persist for a long period, and still in others the biotype may show different inherited char-
Table 9.—Relation between the number of days cultivated in peptone medium and the number of days cultivated in salt-sugar medium needed to change from type 1 to type 5 in Chlamydomonas debaryana (Moewus).

<table>
<thead>
<tr>
<th>Days in peptone medium as type 1</th>
<th>Days in salt-sugar medium needed to change to type 5</th>
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</thead>
<tbody>
<tr>
<td>28</td>
<td>28</td>
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<tr>
<td>140</td>
<td>49</td>
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<td>273</td>
<td>133</td>
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<td>441</td>
<td>175</td>
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<td>567</td>
<td>231</td>
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<td>609</td>
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<td>672</td>
<td>531</td>
</tr>
<tr>
<td>690</td>
<td>534</td>
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</tbody>
</table>

acters. To the last-mentioned category belongs perhaps a strain of Tetrahymena pyriformis in which, according to Furgason (1940), a pure-line bacteria-free culture derived from a single individual was found to be composed of individuals differing in shape and size which became more marked in older cultures.

The first comprehensive study dealing with the variation in size and its inheritance in asexual reproduction of Protozoa was conducted by Jennings (1909). From a "wild" lot of Paramecium caudatum, eight races or biotypes with the relative mean lengths of 206, 200, 194, 176, 142, 125, 100, and 45μ were isolated. It was found that within each clone derived from a single parent, the size of individuals varies greatly (which is attributable to growth, amount of food, and other environmental conditions), any one of which may give rise to progeny of the same mean size. Thus selection within the pure race has no effect on the size, and the differences brought about merely by environment are not inherited. Jennings (1916) examined the inheritance of the size and number of spines, size of shell, diameter of mouth, and size and number of teeth of the testacean Diffugia corona, and showed that "a population consists of many hereditarily diverse stocks, and a single stock, derived from a single progenitor, gradually differentiates into such hereditarily diverse stocks, so that by selection marked results are produced." Root (1918) with Centropyxis aculeata, Hegner (1919) with Arcella dentata, and Reynolds (1924) with A. polypora, obtained similar results. Jennings (1937) studied the inheritance of teeth in Diffugia corona in normal fission and by altering through operation, and found that operated mouth or teeth were restored to
normal form in 3 or 4 generations and that three factors appeared to
determine the character and number of teeth: namely, the size of the
mouth, the number and arrangement of teeth in the parent, and
“something in the constitution of the clone (its genotype) which
tends toward the production of a mouth of a certain size, with teeth
of a certain form, arrangement, and number.”

Races or strains have been recognized in almost all intensively
studied Protozoa. For example, Ujihara (1914) and Dobell and Jepps
(1918) noticed five races in *Entamoeba histolytica* on the basis of dif-
ferences in the size of cysts. Spector (1936) distinguished two races in
the trophozoite of this amoeba. The large strain was found to be
pathogenic to kittens, but the small strain was not. Melene
Y and Frye (1933, 1935) and Frye and Meleney (1939) also hold that there
is a small race in *Entamoeba histolytica* which has a weak capacity for
invading the intestinal wall and not pathogenic to man. Sapiro,
Hakansson and Louttit (1942) similarly notice two races which can be
distinguished by the diameters of cysts, the division line being
10\(\mu\) and 9\(\mu\) in living and balsam-mounted specimens respectively.
The race with large cysts gives rise to trophozoites which are more
actively motile, ingest erythrocytes, and culture easily, is patho-
genic to man and kitten, while the race with small cysts develops
into less actively motile amoebae which do not ingest erythrocytes
and are difficult to culture, is not pathogenic to hosts, thus not being
histozoic. It is interesting to note, however, that Cleveland and
Sanders (1930) found the diameter of the cysts produced in a pure-
line culture of this sarcodinan, which had originated in a single cyst,
varied from 7 to 23\(\mu\). Furthermore, the small race of Frye and
Meleney mentioned above was later found by Meleney and Zucker-
man (1948) to give rise to larger forms in culture, which led the last
two observers to consider that the size range of the strains of this
amoeba is a characteristic which may change from small to large or
*vice versa* under different environmental conditions.

Investigations by Boyd and his co-workers and others show that
the species of *Plasmodium* appear to be composed of many strains
which vary in diverse physiological characters. In an extended study
on *Trypanosoma lewisi*, Taliaferro (1921–1926) found that this flagel-
late multiplies only during the first ten days in the blood of a rat after
inoculation, after which the organisms do not reproduce. In the adult
trypanosomes, the variability for total length in a population is about
3 per cent. Inoculation of the same pure line into different rats some-
times brings about small but significant differences in the mean size
and passage through a rat-flea generally results in a significant vari-
ability of the pure line. It is considered that some differences in dimensions among strains are apparently due to environment (host), but others cannot be considered as due to this cause, since they persist when several strains showing such differences are inoculated into the same host. The two strains of *T. cruzi* isolated from human hosts and maintained for 28 and 41 months by Hauschka (1949), showed well defined and constant strain-specific levels of virulence, different degrees of affinity for certain host tissues, unequal susceptibility to the quinoline-derivative Bayer 7602, and a difference in response to environmental temperature. The five strains of *Trichomonas gallinae* studied by Stabler (1948) were found to possess a marked variation in virulence to its hosts.

According to Kidder and his associates, the six strains (H, E, T, T–P, W, GHH) of *Tetrahymena pyriformis* and the two strains (V, PP) of *T. vorax* differ in biochemical reactions. They found the appearance of a biochemical variation between a parent strain (T) and a daughter strain (T–P) during a few years of separation and a greater difference in the reactions between the two species than that between the strains of each species. These strains show further differences in antigenic relationships. Five strains of *pyriformis* contain qualitatively identical antigens, but differ quantitatively with respect to amount, concentration or distribution of antigenic materials. The sixth strain (T) contains all the antigens of the other five strains and additional antigens. The two strains of *vorax* are said to be nearly identical antigenically. The antigenic differences between the two species were marked, since there is no cross-reaction within the standard testing time. In these cases, thus, some aspects of the physiological difference among different strains are understood.

Jollos (1921) subjected *Paramecium caudatum* to various environmental influences such as temperature and chemicals, and found that the animals develop tolerance which is inherited through many generations even after removal to the original environment. For example, one of the clones which tolerated only 1.1% of standard solution of arsenic acid, was cultivated in gradually increasing concentrations for four months, at the end of which the tolerance for this chemical was raised to 5%. After being removed to water without arsenic acid, the tolerance changed as follows: 22 days, 5%; 46 days, 4.5%; 151 days, 4%; 166 days, 3%; 183 days, 2.5%; 198 days, 1.25% and 255 days, 1%. As the organisms reproduced about once a day, the acquired increased tolerance to arsenic was inherited for about 250 generations.

There are also known inherited changes in form and structure
which are produced under the influence of certain environmental conditions. Jollos designated these changes long-lasting modifications (Dauermodifikationen) and maintained that a change in environmental conditions, if applied gradually, brings about a change, not in the nucleus, but in the cytoplasm, of the organism which when transferred to the original environment, is inherited for a number of generations. These modifications are lost usually during sexual processes at which time the whole organism is reorganized.

The long-lasting morphological and physiological modifications induced by chemical substances have long been known in parasitic Protozoa. Werbitzki (1910) discovered that Trypanosoma brucei loses its blepharoplast when inoculated into mice which have been treated with pyronin, acridin, oxazin and allied dyes, and Piekarski (1949) showed that trypanflavin and organic metal compounds which act as nuclear poisons and interfere with nuclear division, also bring about the loss of blepharoplast in this trypanosome. Laveran and Roudsky (1911) found that the dyes mentioned above have a special affinity for, and bring about the destruction by auto-oxidation of, the blepharoplast. Such trypanosomes lacking a blepharoplast behave normally and remain in that condition during many passages through mice. When subjected to small doses of certain drugs repeatedly, species of Trypanosoma often develop into drug-fast or drug-resistant strains which resist doses of the drug greater than those used for the treatment of the disease for which they are responsible. These modifications may also persist for several hundred passages through host animals and invertebrate vectors, but are eventually lost.

Long-lasting modifications have also been produced by several investigators by subjecting Protozoa to various environmental influences during the nuclear reorganization at the time of fission, conjugation, or autogamy. In Stentor (Popoff) and Glaucoma (Chatton), long-lasting modifications appeared during asexual divisions. Calkins (1924) observed a double-type Uroleptus mobilis (Fig. 95, b) which was formed by a complete fusion of two conjugants. This abnormal animal underwent fission 367 times for 405 days, but finally reverted back to normal forms, without reversion to double form. The double animal of Euplotes patella (d) is, according to Kimball (1941) and Powers (1943), said to be formed by incomplete division and rarely through conjugation. De Garis (1930) produced double animals in Paramecium caudatum through inhibition of division by exposing the animals to cyanide vapor or to low temperatures.
Jennings (1941) outlined five types of long-lasting inherited changes during vegetative reproduction, as follows: (1) changes that occur in the course of normal life history, immaturity to sexual maturity which involves many generations; (2) degenerative changes resulting from existence under unfavorable conditions; (3) adaptive changes or inherited acclimitization or immunity; (4) changes which are neither adaptive nor degenerative, occurring under specific environmental conditions; and (5) changes in form, size, and other characters, which are apparently not due to environment.

Whatever exact mechanism by which the long-lasting modifications are brought about may be, they are difficult to distinguish from permanent modification or mutation, since they persist for hundreds of generations, and cases of mutation have in most instances not been followed by sufficiently long enough pure-line cultures to definitely establish them as such (Jollos, 1934; Moewus, 1934; Sonneborn, 1947).

Jollos observed that if Paramecium were subjected to environmental change during late stages of conjugation, certain individuals, if not all, become permanently changed. Possibly the recombining and reorganizing nuclear materials are affected in such a way that the hereditary constitution or genotype becomes altered. MacDougall subjected Chilodonella uncinata to ultraviolet rays and produced many changes which were placed in three groups: (1) abnormalities which caused the death of the organism; (2) temporary variations which disappeared by the third generation; and (3) variations which
were inherited through successive generations and hence considered as mutations. The mutants were triploid, tetraploid, and tailed diploid forms (Fig. 96), which bred true for a variable length of time in pure-line cultures, either being lost or dying off finally. The tailed form differed from the normal form in the body shape, in the number of ciliary rows and contractile vacuoles, and in the mode of movement, but during conjugation it showed the diploid number of chromosomes as in the typical form. The tailed mutant remained true and underwent 20 conjugations during ten months.

![Fig. 96. Chilodonella uncinata (MacDougall). a, b, ventral and side view of normal individual; c, d, ventral and side view of the tailed mutant.](image)

Kimball (1950) exposed Paramecium aurelia to beta particles from plaques containing P³² and obtained many clones which multiplied more slowly than normal animals or died, which conditions were interpreted by him to be due to mutational changes induced in the micronuclei by the radiation. Kimball found that the radiation was less effective if given just before the cytoplasmic division than if given at other times during the division interval and that exposure of the organisms to ultraviolet ray of wave length 2537 Å inactivates the Kappa (p. 239).

The loss of the blepharoplast in trypanosomes mentioned above occurs also spontaneously in nature. A strain of Trypanosoma evansi which had been maintained in laboratory animals for five years, suddenly lost the blepharoplast (Wenyon, 1928) which condition remained for 12½ years (Hoare, 1940). Hoare and Bennett (1937) found five camels out of 100 they examined infected by the same species of trypanosome that was without a blepharoplast. One strain inoculated into laboratory animals has retained this peculiarity for nearly
three years. Nothing is known as to how such strains arise, though some workers suggest mutational change.

In sexual reproduction, the nuclei of two individuals participate in producing new combinations which would naturally bring about diverse genetic constitutions. The new combination is accomplished either by sexual fusion in Sarcodina, Mastigophora, and Sporozoa, or by conjugation in Euciliata and Suctoria.

The genetics of sexual fusion is only known in a few forms. Perhaps the most complete information was obtained by Moewus through his extended studies of certain Phytomonadina. In Polytoma (p. 281), Chlamydomonas (p. 276), and allied forms, the motile individuals are usually haploid. Two such individuals (gametes) fuse with each other and produce a diploid zygote which encysts. The zygote later undergoes at least two divisions within the cyst wall, in the first division of which chromosome reduction takes place. These swarmers when set free become trophozoites and multiply asexually by division for many generations, the descendants of each swarmer giving rise to a clone.

Moewus (1935) demonstrated the segregation and independent assortment of factors by hybridization of Polytoma. He used two varieties each of two species: P. uvella and P. pascheri, both of which possess 8 haploid chromosomes. Their constitutions were as follows:

**P. uvella**

Form A: Oval (F), without papilla (p), with stigma (S), large (D) (Fig. 97, a).
Form B: Oval (F), without papilla (p), without stigma (s), large (D) (Fig. 97, b).

**P. pascheri**

Form C: Pyriform (f), with papilla (P), without stigma (s), large (D) (Fig. 97, c).
Form D: Pyriform (f), with papilla (P), without stigma (s), small (d) (Fig. 97, d).

Thus six different crosses were possible from the four pairs of characters. When A (FpSD) and B (FpsD) fuse, the zygote divides into four swarmers, two swarmers have stigma (S), and the other two lack this cell organ, which indicates the occurrence of segregation of the two characters (S, s) during the reduction division. When B (FpsD) is crossed with C (fPsD), thus differing in two pairs of characters, two swarmers possess one combination or type and the other two another combination. Different pairs of combinations are
of course found. It was found that about half the zygotes gives rise to the two parental combinations (Fig. 97, b, c), while the other half gives rise to FpsD (e) and fpsD (f).

When B (FpsD) is crossed with D (fPsd) or A (fPsd) is crossed with D (fPsD), only two types of swarmers are also formed from each zygote, and in the case of BxD, eight different combinations are produced, while in the case of AxD, sixteen different combinations, which appear in about equal numbers, are formed. Thus these four factors or characters show independent assortment during divisions of the zygote.

Furthermore, Moewus noticed that certain other characters appeared to be linked with some of the four characters mentioned above. For example, the length of flagella, if it is under control of a factor, is linked on the same chromosome with the size-controlling factors (D, d), for large individuals have invariably long flagella and small individuals short flagella. During the experiments to determine this linkage, it was found that crossing over occurs between two entire chromosomes that are undergoing synapsis.

In certain races of *Polytoma pascheri* and *Chlamydomonas eugametos*, the sexual fusion takes place between members of different clones only. The zygote gives rise as was stated before to four swarmers by two divisions, which are evenly divided between the two sexes, which shows that the sex-determining factors are lodged in a single chromosome pair. In a cross between *Chlamydomonas paradox* and *C. pseudoparadoxa*, both of which produce only one type of gamete in a clone, the majority of the zygotes yield four clones, two
producing male gametes and the other two female gametes; but a small number of zygotes gives rise to four clones which contain both gametes. It is considered that this is due to crossing-over that brought the two sex factors (P and M) together into one chromosome, and hence the "mixed" condition, while the other chromosome which is devoid of the sex factors gives rise to individuals that soon perish.

In crosses between Chlamydomonas eugametos which possesses a stigma and 10 haploid chromosomes and C. paupera which lacks a stigma and 10 haploid chromosomes, 12 pairs of factors including sex factor are distinguishable. Consequently at least two chromosomes must have two factors in them. Thus adaptation to acid or alkaline culture media was found to be linked with differences in the number of divisions in zygote. That there occurs a sex-linked inheritance in Chlamydomonas was demonstrated by crossing stigma-bearing C. eugametos of one sex with stigma-lacking C. paupera of the opposite sex. The progeny that were of the same sex as C. eugametos parent possessed stigma, while those that were of the same sex as C. paupera parent lacked stigma. Thus it is seen that the sex factor and stigma factor are located in the same chromosome.

The genetics of conjugation which takes place between two diploid conjugants has been studied by various investigators. Pure-line cultures of exconjugants show that conjugation brings about diverse inherited constitutions in the clones characterized by difference in size, form, division-rate, mortality-rate, vigor, resistance, etc. The discovery of mating types in Paramecium and in Euplotes, and intensive studies of conjugation and related phenomena, are bringing to light hitherto unknown information on some of the fundamental problems in genetics.

Sonneborn (1939) has made extended studies of variety 1 of Paramecium aurelia (p. 194) and found that genetically diverse materials show different types of inheritance, as follows:

(1) Stocks containing two mating types. When types I and II conjugate, among a set of exconjugants some produce all of one mating type, others all of the other mating type and still others both types (one of one type and the other of the other type). In the last mentioned exconjugants, the types segregate usually at the first division, since of the two individuals produced by the first division, one and all its progeny, are of one mating type, and the other and all its progeny are of the other mating type. A similar change was also found to take place at autogamy. Sonneborn therefore considers that the mating types are determined by macronuclei, as
judged by segregation at first or sometimes second division in exconjugants and by the influence of temperature during conjugation and the first division.

(2) Stocks containing only one mating type. No conjugation occurs in such stocks. Autogamy does not produce any change in type which is always type I. Stocks that contain type II only have not yet been found.

(3) Hybrids between stocks containing one and two mating types. When the members of the stock containing both types I and II (two-type condition) conjugate with those of the stock containing one type (one-type condition), all the descendants of the hybrid exconjugants show two-type condition, which shows the dominancy of two-type condition over one-type condition. The factor for the two-type condition may be designated A and that for the one-type condition a. The parent stocks are AA and aa, and all F₁ hybrids Aa. When the hybrids (Aa) are backcrossed to recessive parent (aa) (158 conjugating pairs in one experiment), approximately one-half (81) of the pairs give rise to two-type condition (Aa) and the remaining one-half (77) of the pairs to one-type condition (aa), thus showing a typical Mendelian result. When F₁ hybrids (Aa) were interbred by 120 conjugating pairs, each exconjugant in 88 of the pairs gave rise to two-type condition and each exconjugant in 32 pairs produced one-type condition, thus approximating an expected Mendelian ratio of 3 dominants to 1 recessive. That the F₂ dominants are composed of two-thirds heterozygotes (Aa) and one-third homozygotes (AA) was confirmed by the results obtained by allowing F₂ dominants to conjugate with the recessive parent stock (aa). Of 19 pairs of conjugants, 6 pairs gave rise to only dominant progeny, which shows that they were homozygous (AA) and their progeny heterozygous (Aa), while 13 pairs produced one-half dominants and one-half recessives, which indicates that they were heterozygous (Aa) and their progeny half homozygous (aa) and half heterozygous (Aa). Thus the genic agreement between two conjugants of a pair and the relative frequency of various gene combinations as shown in these experiments confirm definitely the occurrence of meiosis and chromosomal exchange during conjugation which have hitherto been considered only on cytological ground.

In Euplotes patella, Kimball (1942) made various matings with respect to the inheritance of the mating type. The results obtained can be explained if it is assumed that mating types I, II, and V, are determined by different heterozygous combinations of three allelic genes which if homozygous determine mating types III, IV, and VI.
Upon this supposition, type I has one allele in common with type II, and this allele is homozygous in type IV. It has one allele in common with type V, and this allele is homozygous in type VI. Type II has one allele in common with type V and this is homozygous in type III. These alleles were designated by Kimball, mt\(^1\), mt\(^2\), and mt\(^3\).

The genotypes of the six mating types may be indicated as follows: mt\(^1\)mt\(^2\) (I), mt\(^2\)mt\(^3\) (II), mt\(^3\)mt\(^1\) (III), mt\(^1\)mt\(^1\) (IV), mt\(^2\)mt\(^3\) (V), and mt\(^2\)mt\(^2\) (VI).

There is no dominance among these alleles, the three homozygous combinations determining three mating types being different from one another and from the three determined by homozygous combination. Kimball (1939, 1941) had shown that the fluid obtained free of Euplotes from a culture of one mating type will induce conjugation among animals of certain other mating types. When all possible combinations of fluids and animals are made, it was found that the fluid from any of the heterozygous types induces conjugation among animals of any types other than its own and the fluid from any of the homozygous types induces conjugation only among animals of the types which do not have the same allele as the type from which the fluid came. These reactions may be explained by an assumption that each of the mating type alleles is responsible for the production by the animal of a specific conjugation-inducing substance. Thus the two alleles in a heterozygote act independently of each other; each brings about the production by the animal of a substance of its own. Thus heterozygous animals are induced to conjugate only by the fluids from individuals which possess an allele not present in the heterozygotes.

The double animals of *Euplotes patella* (p. 228) conjugate with double animals or with single animals in appropriate mixtures and at times a double animal gives rise by binary fission to a double and two single animals instead of two animals (Fig. 98). Powers (1943) obtained doubles of various genotypes for mating types which were determined by observing the mating type of each of the two singles that arose from the doubles. Doubles of type IV (mt\(^1\)mt\(^1\)) with a single micronucleus (Fig. 98, a) were mated with singles of type VI (mt\(^3\)mt\(^2\)) (b). The double exconjugants (d) were “split” into their component singles belonging to mating types IV and VI (g), while the doubles were type I (f). Thus it was found that the phenotype of a double animal with separate nuclei was the same as though the alleles present in the nuclei were located within one nucleus. The fact that loss of one micronucleus had no effect on the type of doubles, tends to show that the micronucleus has no direct effect on
mating types. Sonneborn's view (p. 233) that the macronucleus is the determinant of the mating types in *Paramecium aurelia* appears to hold true in *Euplotes* also.

The relation between the cytoplasm and nucleus in respect to inheritance has become better known in recent years in some ciliates. Sonneborn (1934) crossed two clones of *Paramecium aurelia* differing markedly in size and division rate, and found the difference persisted.

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**Fig. 98.** Diagram showing conjugation between a double (type IV) and a single (type VI) of *Euplotes patella* (Powers). a, a double organism with one micronucleus (genotype mt'mt'); b, a normal single with a micronucleus (genotype mt2mt2); c, conjugation of the single with the amicronucleate half of the double (one of the pronuclei produced in the single migrates into the double, while the two pronuclei of the double undergo autogamy); d, the exconjugant double is shown to be type I (mt'mt2); e, exconjugant single remains type VI; f, the double divides into two type I doubles; g, occasionally the anterior half of the double is widely "split," and division produces a double and two singles, the latter testing as type IV and type VI; h, line of exconjugant single. Newly formed macronuclei are stippled.
for a time between the two F₁ clones produced from the two members of each hybrid pair of exconjugants, but later both clones became practically identical in size and division rate (Sonneborn, 1947). De Garis (1935) succeeded in bringing about conjugation in Paramecium caudatum, between the members of a large clone (198μ long) (Fig. 99, a) and of a small clone (73μ long) (b). The exconjugants of a pair are different only in the cytoplasm as the nuclei are alike through exchange of a haploid set of chromosomes. The two exconjugants divide and give rise to progeny which grow to size characteristic of each parent clone, division continuing at the rate of once or twice a day. However, as division is repeated, the descend-

Fig. 99. Diagram showing the size changes in two clones derived from a pair of conjugants of Paramecium caudatum, differing in size (a, b). Gradual change in dimensions in each clone during 22 days resulted in intermediate size (Jennings).

ants of the large clone become gradually smaller after successive fissions, while the descendants of the small clone become gradually larger, until at the end of 22 days (in one experiment) both clones produced individuals of intermediate size (about 135μ long) which remained in the generations that followed. Since the exconjugants differed in the cytoplasm only, it must be considered probable that at first the cytoplasmic character was inherited through several vegetative divisions, but ultimately the influence of the new nucleus gradually changed the cytoplasmic character. The ultimate size between the two clones is however not always midway between the mean sizes of the two parent clones, and is apparently dependent upon the nuclear combinations brought about by conjugation. It has also become known that different pairs of conjugants between the same two clones give rise to diverse progeny, similar to those of sexual reproduction in Metazoa, which indicates that clones of Paramecium caudatum are probably not true clonal lines without genetic mixing.
mecium caudatum are in many cases heterozygous for size factors and recombination of factors occurs at the time of conjugation.

In *P. aurelia*, Kimball (1939) observed that there occasionally occurs a change of one mating type into another following autogamy. When the change is from type II to type I, not all animals change type immediately. Following the first few divisions of the product of the first division after autogamy there are present still some type II animals, although ultimately all become transformed into type I. Here also the cytoplasmic influence persists and is inherited through vegetative divisions. Jennings (1941) in his excellent review writes: “The primary source of diversities in inherited characters lies in the nucleus. But the nucleus by known material interchanges impresses its constitution on the cytoplasm. The cytoplasm retains the constitution so impressed for a considerable length of time, during which it assimilates and reproduces true to its impressed character. It may do this after removal from contact with the nucleus to which its present constitution is due, and even for a time in the presence of another nucleus of different constitution. During this period, cytoplasmic inheritance may occur in vegetative reproduction. The new cells produced show the characteristics due to this cytoplasmic constitution impressed earlier by a nucleus that is no longer present. But in time the new nucleus asserts itself, impressing its own constitution on the cytoplasm. Such cycles are repeated as often as the nucleus is changed by conjugation.”

Since the first demonstration some forty years ago of “cytoplasmic inheritance” in higher plants, many cytoplasmic factors have been observed in various plants (Michaelis and Michaelis, 1948). Information on similar phenomena in Metazoa and Protozoa is of recent origin.

As was already mentioned (p. 196), Sonneborn found in four races of variety 4 of *Paramecium aurelia* a pair of characters which he designated as “killer” and “sensitive.” The killers liberate paramecin, a desoxyribonucleoprotein (Wagtendonk and Zill, 1947), into the culture fluid, to which they are resistant. When the sensitive races are exposed to paramecin in the fluid in which the killer race 51 lived, they show after hours a hump on the oral surface toward the posterior end which becomes enlarged, while the anterior part of the body gradually wastes away. The body becomes smaller and rounded; finally the organisms perish (Fig. 100). Sensitives can be mated to the killers, however, without injury if proper precaution is taken, since paramecin does not affect them during conjugation. The two exconjugants obtain identical genotypes, but their progeny
are different; that is, one is a killer and the other is a sensitive culture. F\(_2\) progeny obtained by selfing show no segregation. Therefore, the difference between the killer and the sensitive is due to a cytoplasmic difference and not to a genic difference.

The same observer noted that the thin cytoplasmic paroral strand which appears between conjugating pair that ordinarily breaks off within a minute, occasionally may remain for a long time, and if the strand persists as long as 30 minutes, there occurs an interchange of cytoplasm between the pair (Fig. 101). When this happens, both exconjugants produce killer clones. In F\(_2\) no segregation takes place. Thus killers can introduce the killer trait to sensitives through a cytoplasmic connection between them. Sonneborn supposed that the killers contain a cytoplasmic genic factor or a \textit{plasmagene} which de-

![Fig. 100. Paramecium aurelia. The changes leading up to death when the sensitives are exposed to the killer stock 51 (variety 4) (Sonneborn).](image)

termines the killer trait and called it \textit{kappa}. Preer (1948) demonstrated that this kappa is a particle which can be recognized in Giemsa-stained specimens (Fig. 102). It was further found that killers can be irreversibly transformed into hereditary sensitives by eliminating kappa particles by exposure to high temperature (Sonneborn, 1946), x-irradiation (Preer, 1948b) or nitrogen mustard (Geckler, 1949) and that sensitives can be transformed to hereditary killers by placing them in concentrated suspensions of broken bodies of killers (Sonneborn, 1948a). Therefore, it became clear that kappa is a self-multiplying cytoplasmic body which is produced when some are already present.

Killer races of variety 2 differ from each other and from that of variety 4 mentioned above, in the effects produced on sensitives before the latter are killed. These sensitives possess a gene different from that of the killers and cannot be changed into killers by immersing it to kappa suspensions of broken bodies of killers. When this sensitive is mated with a killer, F\(_2\) generation produced by self-
ing among the killer F\textsubscript{1} clones, shows segregation of sensitives and killers in the ratio of a single gene difference. In the presence of dominant gene K, kappa is maintained, but in recessive k homozygotes, kappa cannot be maintained and any kappa carried over from killers is rapidly lost. Thus it is evident, Sonneborn points out, that the plasmagene kappa is dependent on gene K.

Dipell (1948, 1950) found a number of killer mutants in variety 4.

She showed through breeding analysis that these mutations have brought about no change in any gene affecting kappa or the killer trait, but have been in every case due to changes in kappa. In a mutant which was capable of producing two types of killing, there were two kinds of kappa which she succeeded in separating in different animals and their progeny. Thus it became apparent that kappa can undergo mutation, that various mutant kappas can multiply in animals with the original genome, and that the kappas are determined by themselves and not by nuclear genes.

According to Preer (1948), the kappa particles (Fig. 102) in the killer race G are about 0.4\textmu long, and those in a mutant Gml only about 0.2–0.3\textmu long, while in other strains they measure as much as
0.8μ in length. Preer (1948a, 1950) further observed that the kappa particles contain desoxyribonucleic acid and vary in form (rod-like or spherical), size and number in different races of killers, and that an increase, reduction or destruction of the kappas, as determined by indirect methods, was correlated with the observed number of the stained particles. As to the suggestion that the kappa particles may be viruses, symbionts (Altenberg, 1948), etc., the reader is referred to Sonneborn (1946, 1950).

The application of antigen-antibody reactions to free-living Protozoa began some forty years ago. Bernheimer and Harrison (1940, 1941) pointed out the antigenic dissimilarity of three species of Paramecium in which the members of a clone differ widely in their
susceptibility to the immobilizing action of a given serum. Strains of *Tetrahymena pyriformis* differ in antigenic reactions, as has already been mentioned (p. 227). Sonneborn and his co-workers have studied serological reactions in *Paramecium aurelia* (Sonneborn, 1950).

When a rabbit is inoculated intraperitoneally with a large number of a strain of *P. aurelia*, its serum immobilizes in a high dilution, the organisms of the same strain, but not of other strains. Such a serologically distinct strain is called a *serotype* or antigenic type. It was found that a clone originating in a homozygous individual gives rise to a series of various serotypes. Race 51 gave rise to eight serotypes: A, B, C, D, E, G, H and J, and race 29, to seven serotypes: A, B, C, D, F, H and J. When a serotype is exposed to its antiserum, it changes into other types, which course Sonneborn was able to control by temperature and other conditions. For example, serotype D (stock 29) may be changed by its antiserum to type B at 32°C. and to type H at 20°C., types B, F and H are convertible one into the other and all other types can be transformed to any of the three; and serotypes A and B (stock 51) are convertible one into the other, and other types can be changed to A or B. The antigenic types are inherited, if the cultures are kept at 26°–27°C. with food enough to allow one division a day. When induced or spontaneous changes of serotype occur, crosses made among different serotypes of the same strain reveal no effective gene differences among them; thus all serotypes of a strain possess apparently an identical genetic constitution. Sonneborn finds serotype A of stock 29 is not exactly the same as the type A of stock 51. When these are crossed, it is found that the difference between two antigens is controlled by a pair of allelic genes of which the 51A-gene is dominant over the 29A-gene. On the basis of these observations, it has been concluded that nuclear genes control the specificity of the physical basis of cytoplasmic inheritance in these antigenic traits, and hereditary transformations of serotype are cytoplasmic "mutations" of hitherto unknown type.

In the inheritance of the killer trait and of serotype, both traits are cytoplasmically determined and inherited; hereditary changes are brought about by environmental conditions; and the traits are dependent for their maintenance upon nuclear genes. However, the specific type of killer trait is controlled by the kind of kappa present, not by the genes, while the specific type of A antigen is determined by the nuclear genes. The transformation of the killer to the sensitive is made irreversible, but that of serotypes is not. The various types of killer character are not mutually exclusive, as different kinds of kappa can coexist in the same organism and
its progeny, each kind of kappa controlling production of its corresponding kind of paramecin, while in serotype, two kinds of antigen substances cannot coexist, thus being mutually exclusive. The physical basis of the killer trait lies in the visible Feulgen-positive kappa particles, while no such particles have so far been found in association with the serotype.

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PART II: TAXONOMY AND SPECIAL BIOLOGY
Chapter 7

Major groups and phylogeny of Protozoa

THE Protozoa are grouped into two subphyla: Plasmodroma (p. 254) and Ciliophora (p. 683). The Plasmodroma are more primitive Protozoa and subdivided into three classes: Mastigophora (p. 254), Sarcodina (p. 417), and Sporozoa (p. 526). The Ciliophora possess more complex body organizations, and are divided into two classes: Ciliata (p. 683) and Suctoria (p. 863).

In classifying Protozoa, the natural system would be one which is based upon the phylogenetic relationships among them in conformity with the doctrine that the present day organisms have descended from primitive ancestral forms through organic evolution. Unlike Metazoa, the great majority of Protozoa now existing do not possess skeletal structures, which condition also seemingly prevailed among their ancestors, and when they die, they disintegrate and leave nothing behind. The exceptions are Foraminifera (p. 493) and Radiolaria (p. 516) which produce multiform varieties of skeletal structures composed of inorganic substances and which are found abundantly preserved as fossils in the earliest fossiliferous strata. These fossils show clearly that the two classes of Sarcodina were already well-differentiated groups at the time of fossilization. The sole information the palaeontological record reveals for our reference is that the differentiation of the major groups of Protozoa must have occurred in an extremely remote period of the earth history. Therefore, consideration of phylogeny of Protozoa had to depend exclusively upon the data obtained through morphological, physiological, and developmental observations of the present-day forms.

The older concept which found its advocates until the beginning of the present century, holds that the Sarcodina are the most primitive of Protozoa. It was supposed that at the very beginning of the living world, there came into being undifferentiated mass of protoplasm which later became differentiated into the nucleus and the cytoplasm. The Sarcodina represented by amoebae and allied forms do not have any further differentiation and lack a definite body wall, they are, therefore, able to change body form by forming pseudopodia. These pseudopodia are temporary cytoplasmic processes and formed or withdrawn freely, even in the more or less permanent axopodia. On the other hand, flagella and cilia are permanent cell-organs possessing definite structural plans. Thus from the morphological viewpoint, the advocates of this concept main-
tained that the Sarcodina are the Protozoa which were most closely related to ancestral forms and which gave rise to Mastigophora, Ciliata, and Sporozoa.

This concept is however difficult to follow, since it does not agree with the general belief that the plant came into existence before the animal; namely, holophytic organisms living on inorganic substances anteceded holozoic organisms living on organic substances. Therefore, from the physiological standpoint the Mastigophora which include a vast number of chlorophyll-bearing forms, must be considered as more primitive than the holozoic Sarcodina. The class Mastigophora is composed of Phytomastigina (chromatophore-bearing flagellates and closely related colorless forms) and Zoomastigina (colorless flagellates). Of the former, Chrysomonadina (p. 256) are mostly naked, and are characterized by possession of 1–2 flagella, 1–2 yellow chromatophores and leucosin. Though holophytic nutrition is general, many are also able to carry on holozoic nutrition. Numerous chrysomonads produce pseudopodia of different types; some possess both flagellum and pseudopodia; others such as Chrysoamoeba (Fig. 105) may show flagellate and ameboid forms (Klebs; Scherffel); still others, for example, members of Rhizochrysidina (p. 267), may lack flagella completely, though retaining the characteristics of Chrysomonadina. When individuals of Rhizochrysis (p. 267) divide, Scherffel (1901) noticed unequal distribution of the chromatophore resulted in the formation of colorless and colored individuals (Fig. 110, a, b). Pascher (1917) also observed that in the colonial chrysomonad, Chryssarachnion (p. 267), the division of component individuals produces many in which the chromatophore is entirely lacking (Fig. 110, c, d). Thus these chrysomonads which lack chromatophores, resemble Sarcodina rather than the parent Chrysomonadina.

Throughout all groups of Phytomastigina, there occur forms which are morphologically alike except the presence or absence of chromatophores. For example, Cryptomonas (p. 273) and Chilomonas (p. 273), the two genera of Cryptomonadina, are so morphologically alike that had it not been for the chromatophore, the former can hardly be distinguished from the latter. Other examples are Euglena, Astasia, and Khawkinea; Chlorogonium and Hyalogonium; Chlamydomonas and Polytoma; etc.

The chromatophores of various Phytomastigina degenerate readily under experimental conditions. For instance, Zumstein (1900) and recently Pringsheim and Hovasse (1948) showed that *Euglena gracilis* loses its green coloration even in light if cultured in fluids
rich in organic substances; in a culture fluid with a small amount of organic substances, the organisms retain green color in light, lose it in darkness; and when cultured in a pure inorganic culture fluid, the flagellates remain green even in darkness. Therefore, it would appear reasonable to consider that the morphologically similar forms with or without chromatophores such as are cited above, are closely related to each other phylogenetically, that they should be grouped together in any scheme of classification, and that the apparent heterogeneity among Phytomastigina is due to the natural course of events. The newer concept which is at present followed widely is that the Mastigophora are the most primitive unicellular animal organisms.

Of Mastigophora, Phytomastigina are to be considered on the same ground more primitive than Zoomastigina. According to the studies of Pascher, Scherffel and others, Chrysomonadina appear to be the nearest to ancestral forms from which other groups of Phytomastigina arose. Among Zoomastigina, Rhizomastigina possibly gave rise to Protomonadina, from which Polymastigina and Hypermastigina later arose. The last-mentioned group is the most highly advanced one of Mastigophora in which an increased number of flagella is an outstanding characteristic.

As to the origin of Sarcodina, many arose undoubtedly from various Zoomastigina, but there are indications that they may have evolved directly from Phytomastigina. As was stated already, Rhizochrysidina possess no flagella and the chromatophore often degenerates or is lost through unequal distribution during division, apparently being able to nourish themselves by methods other than holophytic nutrition. Such forms may have given rise to Amoebina. Some chrysomonads such as Cyrtophora (p. 260) and Palatinella, have axopodia, and it may be considered that they are closer to the ancestral forms from which Heliozoa arose through stages such as shown by Actinomonas (p. 335), Dimorpha (p. 335), and Pteridomonas (p. 335) than any other forms. Another chrysomonad, Porochrysis (p. 260), possesses a striking resemblance to Testacea. The interesting marine chrysomonad, Chrysothylakion (p. 267) that produces a brownish calcareous test from which extrudes astomosing rhizopodial network, resembling a monothalamous foraminiferan, and forms such as Distephanus (p. 267) with siliceous skeletons, may depict the ancestral forms of Foraminifera and Radiolaria respectively. The flagellate origin of these two groups of Sarcodina is also seen in the appearance of flagellated swarmers during their development. The Mycetozoa show also flagellated phase
during their life cycle, which perhaps suggests their origin in flagellated organisms. In fact, in the chrysonad Myxochrysis (p. 261), Pascher (1917) finds a multinucleate and chromatophore-bearing organism (Fig. 105, e–j) that stands intermediate between Chrysonadina and Mycetozoa. Thus there are a number of morphological, developmental, and physiological observations which suggest the flagellate origin of various Sarcodina.

The Sporozoa appear to be equally polyphyletic. The Telosporidida contain three groups in which flagellated microgametes occur, which suggests their derivation from flagellated organisms. Léger and Duboseq even considered them to have arisen from Bodonidae (p. 362) on the basis of flagellar arrangement. Obviously Gregarinida are the most primitive of the three groups. The occurrence of such a form as Selenococcidium (p. 572), would indicate the gregarine-origin of the Coccidia and the members of Haemogregarinidae (p. 592) suggest the probable origin of the Haemosporidia in the Coccidida. The Nidosporidida are characterized by multinucleate trophozoites and by the spore in which at least one polar capsule with a coiled filament occurs. Some consider them as having evolved from Mycetozoa-like organisms, because of the similarity in multinucleate trophozoites, while others compare the polar filament with the flagellum. It is interesting to note here that the nematocyst, similar to the polar capsule, occurs in certain Dinoflagellata (p. 310) independent of flagella. The life cycle of Acnidosporidida is still incompletely known, but the group may have differentiated from such Sarcodina as Mycetozoa.

The Ciliata and Suctoria are distinctly separated from the other groups. They possess the most complex body organization seen among Protozoa. All ciliates possess cilia or cirri which differ from flagella essentially only in size. Apparently Protociliata and Euciliata have different origins, as judged by their morphological and physiological differences. It is probable that Protociliata arose from forms which gave rise to Hypermastigina. Among Euciliata, one finds such forms as Coleps, Urotricha, Plagiocampa, Microregma, Trinyema, Anophrys, etc., which have, in addition to numerous cilia, a long flagellum-like process at the posterior end, and Ileonema that possesses an anterior vibratile flagellum and numerous cilia, which also indicates flagellated organisms as their ancestors. It is reasonable to assume that Holotricha are the most primitive ciliates from which Spirotrichia, Chonotrichia, and Peritrichia evolved. The Suctoria are obviously very closely related to Ciliata and most probably arose from ciliated ancestors by loss of cilia during adult stage
and by developing tentacles in some forms from cytostomes as was suggested by Collin (Fig. 13). General reference (Franz, 1919; Lwoff, 1951).

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THE Plasmodroma possess pseudopodia which are used for locomotion and food-getting or flagella that serve for cell-organs of locomotion. In Sporozoa, the adult stage does not possess any cell-organs of locomotion. The body structure is less complicated than that of Ciliophora. In some groups, are found various endo- and exo-skeletons. The nucleus is of one kind, but may vary in number. All types of nutrition occur. Sexual reproduction is exclusively by sexual fusion or automixis; asexual reproduction is by binary or multiple fission or budding. The majority are free-living, but numerous parasitic forms occur, Sporozoa being all parasitic.

The Plasmodroma are subdivided into three classes as follows:

- Trophozoite with flagellum. Class 1 Mastigophora
- Trophozoite with pseudopodium. Class 2 Sarcodina (p. 417)
- Without cell-organs of locomotion; producing spores; all parasitic. Class 3 Sporozoa (p. 526)

Class 1 Mastigophora Diesing

The Mastigophora includes those Protozoa which possess one to several flagella. Aside from this common characteristic, this class makes a very heterogeneous assemblage and seems to prevent a sharp distinction between the Protozoa and the Protophyta, as it includes Phytomastigina which are often dealt with by botanists.

In the majority of Mastigophora, each individual possesses 1–4 flagella during the vegetative stage, although species of Polymastigina may possess up to 8 or more flagella and of Hypermastigina a greater number of flagella. The palmella stage (Fig. 103) is common among the Phytomastigina and the organism is capable in this stage not only of metabolic activity and growth, but also of reproduction. In this respect, this group shows also a close relationship to algae.

All three types of nutrition, carried on separately or in combination, are to be found among the members of Mastigophora. In holophytic forms, the chlorophyll is contained in the chromatophores which are of various forms among different species and which differ in colors, from green to red. The difference in color appears to be due to the pigments which envelop the chlorophyll body (p. 89). Many forms adapt their mode of nutrition to changed environmental conditions; for instance, from holophytic to saprozoic in the absence of the sunlight. Holozoic, saprozoic and holophytic nutrition are said
to be combined in such a form as Ochromonas. In association with chromatophores, there occurs refractile granules or bodies, the pyrenoids, which are connected with starch-formation. Reserve food substances are starch, oil, etc. (p. 113).

In less complicated forms, the body is naked except for a slight cortical differentiation of the ectoplasm to delimit the body surface and is capable of forming pseudopodia. In others, there occurs a thin plastic pellicle produced by the cytoplasm, which covers the body surface closely. In still others, the body form is constant, being encased in a shell, test, or lorica, which is composed of chitin, pseudo-chitin, or cellulose. Not infrequently a gelatinous secretion envelopes the body. In three families of Protomonadina there is a collar-like structure located at the anterior end, through which the flagellum protrudes.

The great majority of Mastigophora possess a single nucleus, and only a few are multinucleated. The nucleus is vesicular and contains a conspicuous endosome. Contractile vacuoles are always present in the forms inhabiting fresh water. In simple forms, the contents of the vacuoles are discharged directly through the body surface to the exterior; in others there occurs a single contractile vacuole near a reservoir which opens to the exterior through the so-called cytopharynx. In the Dinoflagellata, there are apparently no contractile vacuoles, but non-contractile pusules (p. 310) occur in some forms. In chromatophore-bearing forms, there occurs usually a stigma which is located near the base of the flagellum and seems to be the center of phototactic activity of the organism which possesses it.

Asexual reproduction is, as a rule, by longitudinal fission, but in some forms multiple fission also takes place under certain circumstances, and in others budding may take place. Colony-formation (p. 174), due to incomplete separation of daughter individuals, is widely found among this group. Sexual reproduction has been reported in a number of species.

The Mastigophora are free-living or parasitic. The free-living forms are found in fresh and salt waters of every description; many are free-swimming, others creep over the surface of submerged objects, and still others are sessile. Together with algae, the Mastigophora compose a major portion of plankton life which makes the foundation for the existence of all higher aquatic organisms. The parasitic forms are ecto- or endo-parasitic, and the latter inhabit either the digestive tract or the circulatory system of the host animal. Trypanosoma, a representative genus of the latter group, includes important disease-causing parasites of man and of domestic animals.
The Mastigophora are divided into two subclasses as follows:

With chromatophores ................. Subclass 1 Phytomastigina
Without chromatophores .......... Subclass 2 Zoomastigina (p. 333)

Subclass 1 Phytomastigina Doflein

The Phytomastigina possess the chromatophores and their usual method of nutrition is holophytic, though some are holozoic, saprozoic or mixotrophic; the majority are conspicuously colored; some that lack chromatophores are included in this group, since their structure and development resemble closely those of typical Phytomastigina.

Some observers consider the types of flagella as one of the characters in taxonomic consideration (Petersen, 1929; Vlk, 1933; Owen, 1949; etc.). Owen found, for example, "lash flagella" (with a terminal filament) in some species of Phytomonadina, Rhizomastigina, Protomonadina and Polymastigina and simple flagella in the forms included in Chrysomonadina, Cryptomonadina, Euglenoidina and Dinoflagellata; and simple flagellum and flagella on Oikomonas and Monas. He advocated the transfer of the latter two genera from Protomonadina to Chrysomonadina.

1-4 flagella, either directed anteriorly or trailing
Chromatophores yellow, brown or orange
Anabolic products fat, leucosin ........ Order 1 Chrysomonadina
Anabolic products starch or similar carbohydrates .......... Order 2 Cryptomonadina (p. 272)

Chromatophores green
Anabolic products starch and oil. Order 3 Phytomonadina (p. 276)
Anabolic products paramylon ......... Order 4 Euglenoidina (p. 293)
Anabolic products oil ............. Order 5 Chloromonadina (p. 306)

2 flagella, one of which transverse ...... Order 6 Dinoflagellata (p. 310)

Order 1 Chrysomonadina Stein

The chrysomonads are minute organisms and are plastic, since the majority lack a definite cell-wall. Chromatophores are yellow to brown and usually discoid, though sometimes reticulated, in form. Metabolic products are leucosin and fats. 1-2 flagella are inserted at or near the anterior end of body where a stigma is present.

Many chrysomonads are able to form pseudopodia for obtaining food materials which vary among different species. Nutrition, though chiefly holophytic, is also holozoic or saprozoic. Contractile vacuoles are invariably found in freshwater forms, and are ordinarily of simple structure.

Under conditions not fully understood, the chrysomonads lose
their flagella and undergo division with development of mucilaginous envelope and thus transform themselves often into large bodies known as the palmella phase and undertake metabolic activities as well as multiplication (Fig. 103). Asexual reproduction is, as a rule, by longitudinal division during either the motile or the palmella stage. Incomplete separation of the daughter individuals followed by repeated fission, results in numerous colonial forms mentioned elsewhere (p. 174). Some resemble higher algae very closely. Sexual reproduction is unknown in this group. Encystment occurs commonly; the cyst is often enveloped by a silicious wall possessing an opening with a plug. Taxonomy (Doflein, 1923; Schiller, 1925a; Pascher, 1926; Conrad, 1926; Scherffel, 1926; Hollande, 1952).

The chrysomonads inhabit both fresh and salt waters, often occurring abundantly in plankton.

Motile stage dominant .......... Suborder 1 Euchrysomonadina
Palmella stage dominant
Sarcodina-like; flagellate stage unknown .. Suborder 2 Rhizochrysidina (p. 267)
With flagellate phase .......... Suborder 3 Chryscapsina (p. 269)

Suborder 1 Euchrysomonadina Pascher

With or without simple shell
One flagellum .......... Family 1 Chromulinidae (p. 258)
2 flagella
Flagella equally long .......... Family 2 Syneyryptididae (p. 262)
Flagella unequally long ....... Family 3 Ochromonadidae (p. 264)
With calcareous or silicious shell
Bearing calcareous discs and rods .... Family 4 Coccolithidae (p. 266)
Bearing silicious skeleton ....... Family 5 Silicoflagellidae (p. 267)
Family 1 Chromulinidae Engler

Minute forms, naked or with sculptured shell; with a single flagellum; often with rhizopodia; a few colonial; free-swimming or attached.

Genus Chromulina Cienkowski. Oval; round in cross-section; amoeboid; 1–2 chromatophores; palmella stage often large; in fresh water. Numerous species. The presence of a large number of these organisms gives a golden-brown color to the surface of the water. Development (Doflein, 1923); species (Doflein, 1921, 1922; Schiller, 1929; Pascher, 1929; Conrad, 1930).

C. pascheri Hofeneder (Fig. 104, a, b). 15–20 μ in diameter.

Genus Pseudochromulina Doflein. Spheroid body amoeboid; cytoplasm granulated; two contractile vacuoles anterior; a single flagellum about the body length; a yellow tray-like chromatophore with upturned edge; stigma and pyrenoid absent; nucleus central; cyst ovoid, with asymmetrical siliceous wall with an aperture tube (Doflein, 1921).

P. asymmetrica D. Body 3–4 μ in diameter; cytoplasm with fat and probably leucosin; cyst 4 μ by 3 μ; aperture tube about 1 μ; fresh water (Doflein, 1921).

Genus Chrysamoeba Klebs. Body naked; flagellate stage ovoid, with 2 chromatophores, sometimes slender pseudopodia at the same time; flagellum may be lost and the organism becomes amoeboid, resembling Rhizochrysis (p. 267); standing fresh water.

C. radians K. (Fig. 105, a, b). Flagellated form measures 8 μ by 3.5 μ; amoeboid stage about 8–10 μ by 3–4 μ, with 10–20 μ long radiating pseudopodia; cyst 7 μ in diameter (Doflein, 1922).

Genus Chrysapsis Pascher. Solitary; plastic or rigid; chromatophore diffused or branching; with stigma; amoeboid movement; holophytic, holozoic; fresh water. Several species.

C. sagene P. (Fig. 104, c). Anterior region actively plastic; stigma small; 8–14 μ long; flagellum about 30 μ long.

Genus Chrysococcus Klebs. Shell spheroidal or ovoidal, smooth or sculptured and often brown-colored; through an opening a flagellum protrudes; 1–2 chromatophores; one of the daughter individuals formed by binary fission leaves the parent shell and forms a new one; fresh water. Lackey (1938) found several species in Scioto River, Ohio.

C. ornatus Pascher (Fig. 104, d). 14–16 μ by 7–10 μ.

Genus Mallomonas Perty (Pseudomallomonas Chodat). Body elongated; with silicious scales and often spines; 2 chromatophores
rod-shaped; fresh water. Numerous species (Pascher, 1921; Conrad, 1927, 1930).

M. litomosa Stokes (Fig. 104, e). Scales very delicate, needle-like projections at both ends; flagellum as long as body; 24–32μ by 8μ.

Fig. 104. a, b, Chromulina pascheri, ×670 (Hofeneder); e, Chrysapsis sagene, ×1000 (Pascher); d, Chrysococcus ornatus, ×600 (Pascher); e, Mallomonas litomosa, ×400 (Stokes); f, Pyramidochrysis modesta, ×670 (Pascher); g, Sphaleromantis ochracea, ×600 (Pascher); h, Kephyrion ovum, ×1600 (Pascher); i, Chrysopyxis cyathus, ×600 (Pascher); j, Cyrtophora pedicellata, ×400 (Pascher); k, Palatinella cyrtophora, ×400 (Lauterborn); l, Chrysosphaerella longispina, ×600 (Lauterborn).
Genus **Microglena** Ehrenberg. Body ovoid to cylindrical; with a firm envelope in the surface of which are embedded many lenticular masses of silica (Conrad, 1928); a single flagellum at anterior end; a reservoir around which four to eight contractile vacuoles occur; a sheet-like chromatophore; stigma; leucosin; fresh water.

*M. ovum* Conrad (Fig. 106, a). 31–38μ by 18–25μ (Conrad, 1928).

Genus **Pyramidochrysis** Pascher. Body form constant; pyriform with 3 longitudinal ridges; flagellate end drawn out; a single chromatophore; 2 contractile vacuoles; fresh water.

*P. modesta* P. (Fig. 104, j). 11–13μ long.

Genus **Sphaleromantis** Pascher. Triangular or heart-shaped; highly flattened; slightly plastic; 2 chromatophores; 2 contractile vacuoles; stigma large; long flagellum; fresh water.

*S. ochracea* P. (Fig. 104, g). 6–13μ long.

Genus **Kephyrion** Pascher. With oval or fusiform lorica; body fills posterior half of lorica; one chromatophore; a single short flagellum; small; fresh water. Species (Conrad, 1930).

*K. ovum* P. (Fig. 104, h). Lorica up to 7μ by 4μ.

Genus **Chrysopyxis** Stein. With lorica of various forms, more or less flattened; 1–2 chromatophores; a flagellum; attached to algae in fresh water.

*C. cyathus* Pascher (Fig. 104, i). One chromatophore; flagellum twice body length; lorica 20–25μ by 12–15μ.

Genus **Cyrtophora** Pascher. Body inverted pyramid with 6–8 axopodia and a single flagellum; with a contractile stalk; a single chromatophore; a contractile vacuole; fresh water.

*C. pedicellata* P. (Fig. 104, j). Body 18–22μ long; axopodia 40–60μ long; stalk 50–80μ long.

Genus **Palatinella** Lauterborn. Lorica tubular; body heart-shaped; anterior border with 16–20 axopodia; a single flagellum; a chromatophore; several contractile vacuoles; fresh water.

*P. cyrtophora* L. (Fig. 104, k). Lorica 80–150μ long; body 20–25μ by 18–25μ; axopodia 50μ long.

Genus **Chrysochromella** Lauterborn. In spherical colony, individual cell, oval or pyriform, with 2 chromatophores; imbedded in gelatinous mass; fresh water.

*C. longispina* L. (Fig. 104, l). Individuals up to 15μ by 9μ; colony up to 250μ in diameter; in standing water rich in vegetation.

Genus **Porochrysis** Pascher. Shell with several pores through which rhizopodia are extended; a flagellum passes through an apical pore; a single small chromatophore; leucosin; a contractile vacuole; fresh water.
**P. aspergillus** P. (Fig. 105, c, d). Shell about 35μ long by 25μ wide; chromatophore very small; a large leucosin grain; fresh water.

Genus *Myxochrysis* Pascher. Body multinucleate, amoeboid; with yellowish moniliform chromatophores, many leucosin granules and contractile vacuoles; holozoic; surrounded by a brownish envelop which conforms with body form; flagellated swarmers develop into

![Diagrams](image)

Fig. 105. a, flagellate and b, amoeboid phase of *Chrysamoeba radians*, ×670 (Klebs); c, surface view and d, optical section of *Porochrysis aspergillus*, ×400 (Pascher); e–j, *Myxochrysis paradoxa* (Pascher). e, a medium large plasmodium with characteristic envelop; the large food vacuole contains protophytan, Scenedesmus, ×830; f, diagrammatic side view of a plasmodium, engulfing a diatom; moniliform bodies are yellowish chromatophores, ×1000; g–i, development of swarmer into plasmodium (stippled bodies are chromatophores), ×1200.

multinucleate plasmodium; plasmotomy and plasmogamy; fresh water (Pascher, 1916a).

*M. paradoxa* P. (Fig. 105, e–j). Plasmodium 15–18μ or more in diameter; in standing water.

Genus *Angulochrysis* Lackey. Body ovoid: colorless, thin lorica rounded anteriorly and flattened posteriorly into “wings”; a single flagellum long; no cytostome; two bright yellow-brown chromatophores; no stigma; swims with a slow rotation; marine (Lackey, 1940).
A. erratica L. (Fig. 106, b, c). Body up to 12μ long; lorica up to 30μ high; flagellum about four times the body length; Woods Hole.

Genus Stylochromonas L. Body ovoid, sessile with a stiff stalk; with a large collar at anterior end; a single flagellum; two golden brown chromatophores; no stigma; marine (Lackey, 1940).

S. minuta L. (Fig. 106, d). Body 5–8μ long; collar about 6μ high; flagellum about twice the body length.

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Fig. 106. a, Microglena ovum, ×680 (Conrad); b, c, two views of Angulochrysis erratica, ×900 (Lackey); d, Stylochromonas minuta, ×1200 (Lackey).

Family 2 Syncryptidae Poche

Solitary or colonial chrysomonads with 2 equal flagella; with or without pellicle (when present, often sculptured); some possess stalk.

Genus Syncrypta Ehrenberg. Spherical colonies; individuals with 2 lateral chromatophores, embedded in a gelatinous mass; 2 contractile vacuoles; without stigma; cysts unknown; fresh water.

S. volvox E. (Fig. 107, a). 8–14μ by 7–12μ; colony 20–70μ in diameter; in standing water.

Genus Synura Ehrenberg (Synuropsis Schiller). Spherical or ellipsoidal colony composed of 2–50 ovoid individuals arranged radially; body usually covered by short bristles; 2 chromatophores lateral; no stigma; asexual reproduction of individuals is by longitudinal division; that of colony by bipartition; cysts spherical; fresh water. Species (Korshikov, 1929).

S. uvella E. (Fig. 107, b). Cells oval; bristles conspicuous; 20–40μ by 8–17μ; colony 100–400μ in diameter; if present in large numbers,
the organism is said to be responsible for an odor of the water resembling that of ripe cucumber.

*S. adamsi* Smith (Fig., 107 c). Spherical colony with individuals radiating; individuals long spindle, 42–47μ by 6.5–7μ; 2 flagella up to 17μ long; in fresh water pond.

Genus *Hymenomonas* Stein. Solitary; ellipsoid to cylindrical; membrane brownish, often sculptured; 2 chromatophores; without stigma; a contractile vacuole anterior; fresh water.

*H. roseola* S. (Fig. 107, d). 17–50μ by 10–20μ.

Genus *Derepyxis* Stokes. With cellulose lorica, with or without a short stalk; body ellipsoid to spherical with 1–2 chromatophores; 2 equal flagella; fresh water.

*D. amphora* S. (Fig. 107, e). Lorica 25–30μ by 9–18μ; on algae in standing water.

*D. ollula* S. (Fig. 107, f). Lorica 20–25μ by 15μ.

Genus *Stylochrysalis* Stein. Body fusiform; with a gelatinous stalk attached to Volvocidae; 2 equal flagella; 2 chromatophores; without stigma; fresh water.

*S. parasitica* S. (Fig. 107, g). Body 9–11μ long; stalk about 15μ long; on phytomonads.
Family 3 Ochromonadidae Pascher

With 2 unequal flagella; no pellicle and plastic; contractile vacuoles simple; with or without a delicate test; solitary or colonial; free-swimming or attached.

Genus Ochromonas Wyssotzki. Solitary or colonial; body surface delicate; posterior end often drawn out for attachment; 1–2 chromatophores; usually with a stigma; encystment; fresh water. Many species (Doflein, 1921, 1923).

O. mutabilis Klebs (Fig. 108, a). Ovoid to spherical; plastic, 15–30μ by 8–22μ.

O. ludibunda Pascher (Fig. 108, b). Not plastic; 12–17μ by 6–12μ.

O. granularis Doflein. No stigma; 5–12μ long (Doflein, 1922).

Fig. 108. a, Ochromonas mutabilis, ×670 (Senn); b, O. ludibunda, ×540 (Pascher); c, Uroglena volvox, ×430 (Stein); d, Uroglenopsis americana, ×470 (Lemmernann); e, Cyclonexis annularis, ×540 (Stokes); f, Dinobryon sertularia, ×670 (Scherffel); g, Hyalobryon ramosum, ×540 (Lauterborn); h, Stylopyxis mucicola, ×470 (Bolochonzew).
Genus **Uroglena** Ehrenberg. Spherical or ovoidal colony, composed of ovoid or ellipsoidal individuals arranged on periphery of a gelatinous mass; all individuals connected with one another by gelatinous processes running inward and meeting at a point; with a stigma and a plate-like chromatophore; asexual reproduction of individuals by longitudinal fission, that of colony by bipartition; cysts spherical with spinous projections, and a long tubular process; fresh water. One species.

*U. volvox* E. (Fig. 108, c). Cells 12–20μ by 8–13μ; colony 40–400μ in diameter; in standing water.

Genus **Uroglenopsis** Lemmermann. Similar to *Uroglena*, but individuals without inner connecting processes.

*U. americana* (Calkins) (Fig. 108, d). Each cell with one chromatophore; 5–8μ long; flagellum up to 32μ long; colony up to 300μ in diameter; when present in abundance, the organism gives an offensive odor to the water (Calkins). Morphology, development (Troitzkaja, 1924).

*U. europaea* Pascher. Similar to the last-named species; but chromatophores 2; cells up to 7μ long; colony 150–300μ in diameter.

Genus **Cyclonexis** Stokes. Wheel-like colony, composed of 10–20 wedge-shaped individuals; young colony funnel-shaped; chromatophores 2, lateral; no stigma; reproduction and encystment unknown; fresh water.

*C. annularis* S. (Fig. 108, e). Cells 11–14μ long; colony 25–30μ in diameter; in marshy water with sphagnum.

Genus **Dinobryon** Ehrenberg. Solitary or colonial; individuals with vase-like, hyaline, but sometimes, yellowish cellulose test, drawn out at its base; elongated and attached to the base of test with its attenuated posterior tip; 1–2 lateral chromatophores; usually with a stigma; asexual reproduction by binary fission; one of the daughter individuals leaving test as a swarmer, to form a new one; in colonial forms daughter individuals remain attached to the inner margin of aperture of parent tests and there secrete new tests; encystment common; the spherical cysts possess a short process; Ahlstrom (1937) studied variability of North American species and found the organisms occur more commonly in alkaline regions than elsewhere; fresh water. Numerous species.

*D. serrulata*ria E. (Fig. 108, f). 23–43μ by 10–14μ.

*D. divergens* Imhof. 26–65μ long; great variation in different localities.

Genus **Hyalobryon** Lauterborn. Solitary or colonial; individual body structure similar to that of *Dinobryon*; lorica in some cases
tubular, and those of young individuals are attached to the exterior of parent tests; fresh water.

*H. ramosum* L. (Fig. 108, g). Lorica 50–70μ long by 5–9μ in diameter; body up to 30μ by 5μ; on vegetation in standing fresh water.

Genus *Stylopyxis* Bolchoziew. Solitary; body located at bottom of a delicate stalked lorica with a wide aperture; 2 lateral chromatophores; fresh water.

*S. mucicola* B. (Fig. 108, h). Lorica 17–18μ long; stalk about 33μ long; body 9–11μ long; fresh water.

Family 4 Coccolithidae Lohmann

The members of this family occur, with a few exceptions, in salt water only; with perforate (tremalith) or imperforate (discolith) discs, composed of calcium carbonate; 1–2 flagella; 2 yellowish

![Fig. 109. a, Pontosphaerahaecckeli, ×1070 (Kühn); b, Discosphaeratubifer, ×670 (Kühn); c, Distephanus speculum, ×530 (Kühn); d, Rhizo- 
chrysischerffeli, ×670 (Doflein); e–g, Hydrurusfoetidus (e, entire 
colony; f, portion; g, cyst), e (Berthold), f, ×330, g, ×800 (Klebs); h, i, Chrysocapsapaludosa, ×530 (West); j, k, Phaeosphaeragelatinosa (j, part 
of a mass, ×70; k, three cells, ×330) (West).](image-url)
chromatophores; a single nucleus; oil drops and leucosin; holophytic. Taxonomy and phylogeny (Schiller, 1925, 1926; Conrad, 1928a; Kamptner, 1928; Deflandre, 1952a).

Examples:

*Pontosphaera haeckeli* Lohmann (Fig. 109, a).

*Discosphaera tubifer* Murray and Blackman (Fig. 109, b).

**Family 5 Silicoflagellidae** Borgert

Exclusively marine planktons; with siliceous skeleton which envelops the body. Example: *Distephanus speculum* (Müller) (Fig. 109, c) (Deflandre, 1952).

**Suborder 2 Rhizochrysidina** Pascher

No flagellate stage is known to occur; the organism possesses pseudopodia; highly provisional group, based wholly upon the absence of flagella; naked or with test; various forms; in some species chromatophores are entirely lacking, so that the organisms resemble some members of the Sarcodina. Several genera.

Genus **Rhizochrysis** Pascher. Body naked and amoeboïd; with 1–2 chromatophores: fresh water.

*R. scherffeli* P. (Figs. 109, d; 110, a, b). 10–14μ in diameter; 1–2 chromatophores: branching rhizopods; fresh water.

Genus **Chrysidiastrum** Lauterborn. Naked; spherical; often several in linear association by pseudopodia; one yellow-brown chromatophore; fresh water.

*C. catenatum* L. Cells 12–14μ in diameter (Pascher, 1916a).

Genus **Chrysoarachnion** Pascher. Ameboid organism; with a chromatophore, leucosin grain and contractile vacuole; many individuals arranged in a plane and connected by extremely fine rhizopods, the whole forming a cobweb network. Small animals are trapped by the net; chromatophores are small; nutrition both holophytic and holozoic; during division the chromatophore is often unevenly distributed so that many individuals without any chromatophore are produced; fresh water (Pascher, 1916a).

*C. insidians* P. (Fig. 110, c, d). Highly amœboïd individuals 3–4μ in diameter; chromatophore pale yellowish brown, but becomes bluish green upon death of organisms; a leucosin grain and a contractile vacuole; colony made up of 200 or more individuals.

Genus **Chrysothylakion** Pascher. With retort-shaped calcareous shell with a bent neck and an opening; shell reddish brown (with
Fig. 110. a, b, *Rhizochysis scherffeli*, ×500 (Scherffel). a, 4 chromatophore-bearing individuals and an individual without chromatophore; b, the last-mentioned individual after 7 hours. c, d, *Chrysarachnion insidians* (Pascher). c, part of a colony composed of individuals with and without chromatophore, ×1270; d, products of division, one individual lacks chromatophore, but with a leucosin body, ×2530. e, f, *Chrysothylakion vorax* (Pascher). e, an individual with anastomosing rhizopodia and "excretion granules," ×870; f, optical section of an individual; the cytoplasm contains two fusiform brownish chromatophores, a spheroid nucleus, a large leucosin body and contractile vacuole, × about 1200.
iron) in old individuals; through the aperture are extruded extremely fine anastomosing rhizopods; protoplasm which fills the shell is colorless; a single nucleus, two spindle-form brown chromatophores, several contractile vacuoles and leucosin body; marine water.

*C. vorax* P. (Fig. 110, e, f). The shell measures 14–18μ long, 7–10μ broad, and 5–6μ high; on marine algae.

Suborder 3 *Chrysocapsina* Pascher

Palmella stage prominent; flagellate forms transient; colonial; individuals enclosed in a gelatinous mass; 1–2 flagella, one chromatophore, and a contractile vacuole; one group of relatively minute forms and the other of large organisms.

Genus *Hydrurus* Agardh. In a large (1–30 cm. long) branching gelatinous cylindrical mass; cells yellowish brown; spherical to ellipsoidal; with a chromatophore; individuals arranged loosely in gelatinous matrix; apical growth resembles much higher algae; multiplication of individuals results in formation of pyrimidal forms with a flagellum, a chromatophore, and a leucosin mass; eyst may show a wing-like rim; cold freshwater streams.

*H. foetidus* Kirschner (Figs. 32, d–f; 109, e–g). Olive-green, feathery tufts, 1–30 cm. long, develops an offensive odor; sticky to touch; occasionally encrusted with calcium carbonate; in running fresh water.

Genus *Chrysocapsa* Pascher. In a spherical to ellipsoidal gelatinous mass; cells spherical to ellipsoid; 1–2 chromatophores; with or without stigma; freshwater.

*C. paludosa* P. (Fig. 109, h, i). Spherical or ellipsoidal with cells distributed without order; with a stigma; 2 chromatophores; swarmer pyriform with 2 flagella; cells 11μ long; colony up to 100μ in diameter.

Genus *Phaeosphaera* West and West. In a simple or branching cylindrical gelatinous mass; cells spherical with a single chromatophore; fresh water.

*P. gelatinosa* W. and W. (Fig. 109, j, k). Cells 14–17.5μ in diameter.

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Chapter 9

Order 2 Cryptomonadina Stein

The cryptomonads differ from the chrysomonads in having a constant body form. Pseudopodia are very rarely formed, as the body is covered by a pellicle. The majority show dorso-ventral differentiation, with an oblique longitudinal furrow. 1–2 unequal flagella arise from the furrow or from the cytopharynx. In case 2 flagella are present, both may be directed anteriorly or one posteriorly. These organisms are free-swimming or creeping.

One or two chromatophores are usually present. They are discoid or band-form. The color of chromatophores varies: yellow, brown, red, olive-green; the nature of the pigment is not well understood, but it is said to be similar to that which is found in the Dinoflagellata (Pascher). One or more spherical pyrenoids which are enclosed within a starch envelope appear to occur outside the chromatophores. Nutrition is mostly holophytic; a few are saprozoic or holozoic. Assimilation products are solid discoid carbohydrates which stain blue with iodine in Cryptomonas or which stain reddish violet by iodine in Cryptochrysis; fat and starch are produced in holozoic forms which feed upon bacteria and small Protozoa. The stigma is usually located near the insertion point of the flagella. Contractile vacuoles, one to several, are simple and are situated near the cytopharynx. A single vesicular nucleus is ordinarily located near the middle of the body.

Asexual reproduction, by longitudinal fission, takes place in either the active or the non-motile stage. Sexual reproduction is unknown. Some cryptomonads form palmella stage and others gelatinous aggregates. In the suborder Phaeocapsina, the palmella stage is permanent. Cysts are spherical, and the cyst wall is composed of cellulose. The Cryptomonadina occur in fresh or sea water, living also often as symbionts in marine organisms.

Flagellate forms predominant............Suborder 1 Eucryptomonadina
Palmella stage permanent..............Suborder 2 Phaeocapsina (p. 275)

Suborder 1 Eucryptomonadina Pascher

Truncate anteriorly; 2 anterior flagella; with an oblique furrow near anterior end.................Family 1 Cryptomonadidae (p. 273)
Reniform; with 2 lateral flagella; furrow equatorial.....................
.........................................................Family 2 Nephroselmidae (p. 274)
Family 1 Cryptomonadidae Stein

Genus Cryptomonas Ehrenberg. Elliptical body with a firm pellicle; anterior end truncate, with 2 flagella; dorsal side convex, ventral side slightly so or flat; nucleus posterior; "cytopharynx" with granules, considered trichocysts by some observers (Hollande, 1942, 1952); 2 lateral chromatophores vary in color from green to blue-green, brown or rarely red; holophytic; with small starch-like bodies which stain blue in iodine; 1–3 contractile vacuoles anterior; fresh water. Several species. Morphology and taxonomy (Hollande, 1942, 1952).

Fig. 111. a, Cryptomonas ovata, ×800 (Pascher); b, Chilomonas paramecium, ×1330 (Bütschli); c, d, Chrysidella schaudinni, ×1330 (Winter); e, Cyathomonas truncata, ×670 (Uehla); f, Cryptochrysis commutata, ×670 (Pascher); g, Rhodomonas lens, ×1330 (Ruttner); h, Nephroselmis olvacea, ×670 (Pascher); i, Protochrysis phaeophycearum, ×800 (Pascher); j, k, Phaeothamnion confervicolum, ×600 (Kühn).

C. ovata E. (Fig. 111, a). 30–60 μ by 20–25 μ; among vegetation.

Genus Chilomonas Ehrenberg. Similar to Cryptomonas in general body form and structure, but colorless because of the absence of chromatophores; without pyrenoid; "cytopharynx" deep, lower half surrounded by granules, considered by Hollande (1942) and Dragesco (1951) as trichocysts; one contractile vacuole anterior; nucleus in posterior half; endoplasm usually filled with polygonal starch grains; saprozoic fresh water.

C. paramecium E. (Fig. 111, b). Posteriorly narrowed, slightly bent "dorsally"; 30–40 μ by 10–15 μ; saprozoic; widely distributed in stag-
nant water. Cytology (Mast and Doyle, 1935; Hollande, 1942; Dragesco, 1951); bacteria-free culture (Mast and Pace, 1933); metabolism (Mast and Pace, 1929; Pace, 1941); growth substances (Pace, 1944, 1947; Mast and Pace, 1946); effects of vitamins (Pace, 1947).

*C. oblonga* Pascher. Oblong; posterior end broadly rounded; 20–50μ long.

Genus *Chrysidella* Pascher. Somewhat similar to *Cryptomonas*-but much smaller; yellow chromatophores much shorter; those occurring in Foraminifera or Radiolaria as symbionts are known as *Zooxanthellae*. Several species.

*C. schaudinii* (Winter) (Fig. 111, c, d). Body less than 10μ long; in the foraminiferan *Peneroplis pertusus*.

Genus *Cyathomonas* Fromentel. Body small, somewhat oval; without chromatophores; much compressed; anterior end obliquely truncate; with 2 equal or subequal anterior flagella; colorless; nucleus central; anabolic products, stained red or reddish violet by iodine; contractile vacuole usually anterior; a row of refractile granules, prorichocysts, close and parallel to anterior margin of body; asexual reproduction by longitudinal fission; holozoic; in stagnant water and infusion. One species.

*C. truncata* Ehrenberg (Fig. 111, e). 15–25μ by 10–15μ.

Genus *Cryptochrysis* Pascher. Furrow indistinctly granulated; 2 or more chromatophores brownish, olive-green, or dark green, rarely red; pyrenoid central; 2 equal flagella; some lose flagella and may assume amoeboid form; fresh water.

*C. commutata* P. (Fig. 111, f). Bean-shaped; 2 chromatophores; 19μ by 10μ.

Genus *Rhodomonas* Karsten. Furrow granulated; chromatophore one, red (upon degeneration the coloring matter becomes dissolved in water); pyrenoid central; fresh water.

*R. lens* Pascher and Ruttner (Fig. 111, g). Spindle-form; about 16μ long; in fresh water.

Family 2 *Nephroselmidae* Pascher

Body reniform; with lateral equatorial furrow; 2 flagella arising from furrow, one directed anteriorly and the other posteriorly.

Genus *Nephroselmis* Stein. Reniform; flattened; furrow and cytopharynx distinct; no stigma; 1–2 chromatophores, discoid, brownish green; nucleus dorsal; a central pyrenoid; 2 contractile vacuoles; with reddish globules; fresh water.

*N. olvacea* S. (Fig. 111, h). 20–25μ by 15μ.
Genus *Protochrysis* Pascher. Reniform; not flattened; with a distinct furrow, but without cytopharynx; a stigma at base of flagella; 1–2 chromatophores, brownish yellow; pyrenoid central; 2 contractile vacuoles; fission seems to take place during the resting stage; fresh water.

*P. phaeophycearum* P. (Fig. 111, i). 15–17μ by 7–9μ.

**Suborder 2 Phaeocapsina** Pascher

Palmella stage predominant; perhaps border-line forms between brown algae and cryptomonads. Example: *Phaeothamnion confervicolum* Lagerheim (Fig. 111, j, k) which is less than 10μ long.

**References**


Order 3 Phytomonadina Blochmann

The phytomonads are small, more or less rounded, green flagellates, with a close resemblance to the algae. They show a definite body form, and most of them possess a cellulose membrane, which is thick in some and thin in others. There is a distinct opening in the membrane at the anterior end, through which 1–2 (or 4 or more) flagella protrude. The majority possess numerous grass-green chromatophores, each of which contains one or more pyrenoids. The method of nutrition is mostly holophytic or mixotrophic; some colorless forms are, however, saprozoic. The metabolic products are usually starch and oils. Some phytomonads are stained red, owing to the presence of haematochrome. The contractile vacuoles may be located in the anterior part or scattered throughout the body. The nucleus is ordinarily centrally located, and its division seems to be mitotic, chromosomes having been definitely noted in several species.

Asexual reproduction is by longitudinal fission, and the daughter individuals remain within the parent membrane for some time. Sexual reproduction seems to occur widely. Colony formation also occurs, especially in the family Volvocidae. Encystment and formation of the palmella stage are common among many forms. The phytomonads have a much wider distribution in fresh than in salt water.

Solitary
Membrane a single piece; rarely indistinct
   2 flagella ........................................... Family 1 Chlamydomonadidae
   3 flagella ........................................... Family 2 Trichlorididae (p. 281)
   4 flagella ........................................... Family 3 Carteriidae (p. 281)
   5 flagella ........................................... Family 4 Chlorasteridae (p. 283)
   6 or more flagella ................................. Family 5 Polyblepharididae (p. 284)
Membrane bivalve .................................. Family 6 Phacotidae (p. 284)
Colonial, of 4 or more individuals; 2 (1 or 4) flagella .........................
   ................................................. Family 7 Volvocidae (p. 285)

Family 1 Chlamydomonadidae Bütschli

Solitary; spheroid, oval, or ellipsoid; with a cellulose membrane; 2 flagella; chromatophores, stigma, and pyrenoids usually present. Cytology (Hollande, 1942).

Genus Chlamydomonas Ehrenberg. Spherical, ovoid or elongated; sometimes flattened; 2 flagella; membrane often thickened at anterior end; a large chromatophore, containing one or more pyrenoids;
stigma; a single nucleus; 2 contractile vacuoles anterior; asexual reproduction and palmella formation; sexual reproduction isogamy or anisogamy; fresh water. Numerous species (Pascher, 1921, 1925, 1929, 1930, 1932; Skvortzow, 1929; Pringsheim, 1930; Pascher and Jahoda, 1928; Moewus, 1932, 1933; Gerloff, 1940); variation (Moewus, 1933); sexual development (Moewus, 1933a); variation (p. 223); genetics (p. 231).

C. monadina Stein (Fig. 112, a-c). 15–30 μ long; fresh water; Landacre noted that the organisms obstructed the sand filters used in connection with a septic tank, together with the diatom Navicula.

C. angulosa Dill. About 20 μ by 12–15 μ; fresh water.

C. epiphytica Smith (Fig. 112, d). 8–9 μ by 7–8 μ; in freshwater lakes.

C. globosa Snow (Fig. 112, e). Spheroid or ellipsoid; 5–7 μ in diameter; in freshwater lakes.

C. gracilis S. (Fig. 112, f). 10–13 μ by 5–7 μ; fresh water.

Genus Haematococcus Agardh (Sphaerella Sommerfeldt). Spheroidal or ovoid with a gelatinous envelope; chromatophore peripheral and reticulate, with 2–8 scattered pyrenoids; several contractile vacuoles; haematochrome frequently abundant in both motile and encysted stages; asexual reproduction in motile form; sexual reproduction isogamy; fresh water.

H. pluvialis (Flotow) (Figs. 42; 112, g). Spherical; with numerous radial cytoplasmic processes; chromatophore U-shape in optical section; body 8–50 μ, stigma fusiform, lateral; fresh water. Reichenow (1909) noticed the disappearance of haematochrome if the culture medium was rich in nitrogen and phosphorus. In bacteria-free cultures, Elliott (1934) observed 4 types of cells: large and small flagellates, palmella stage and haemato cysts. Large flagellates predominate in liquid cultures, but when conditions become unfavorable, palmella stage and then haemato cysts develop. When the cysts are placed in a favorable environment after exposure to freezing, desiccation, etc., they give rise to small flagellates which grow into palmella stage or large flagellates. No syngamy of small flagellates was noticed. Haematochrome appears during certain phases in sunlight and its appearance is accelerated by sodium acetate under sunlight. Sexuality (Schulze, 1927).

Genus Sphaerellopsis Korschikoff (Chlamydococcus Stein). With gelatinous envelope which is usually ellipsoid with rounded ends; body elongate fusiform or pyriform, no protoplasmic processes to envelope; 2 equally long flagella; chromatophore large; a pyrenoid; with or without stigma; nucleus in anterior half; 2 contractile vacuoles; fresh water.
S. fluviatilis (Stein) (Fig. 112, h). 14–30μ by 10–20μ; fresh water.
Genus Brachiomonas Bohlin. Lobate; with horn-like processes, all directed posteriorly; contractile vacuoles; ill-defined chromatophore; pyrenoids; with or without stigma; sexual and asexual reproduction; fresh, brackish or salt water.

Fig. 112. a–c, Chlamydomonas monadina, ×470 (Goroschankin) (a, typical organism; b, anisogamy; c, palmella stage); d, C. epiphytica, ×1030 (Smith); e, C. globosa, ×2000 (Snow); f, C. gracilis, ×770 (Snow); g, Haematococcus pluvialis, ×500 (Reichenow); h, Sphaerellopsis fluviatilis, ×490 (Korschikoff); i, Brachiomonas westiana, ×960 (West); j, Lobomonas rostrata, ×1335 (Hazen); k, Diplostauron pentagonium, ×1110 (Hazen); l, Gigantochloris permoxima, ×370 (Pascher); m, Gloeomonas ovalis, ×330 (Pascher); n, Scourfieldia complanata, ×1540 (West); o, Thorakomonas sabulosa, ×670 (Korschikoff).
B. westiana Pascher (Fig. 112, i). 15–24\(\mu\) by 13–23\(\mu\); brackish water.

Genus Lobomonas Dangeard. Ovoid or irregularly angular; chromatophore cup-shaped; pyrenoid; stigma; a contractile vacuole, fresh water.

L. rostrata Hazen (Fig. 112, j). 5–12\(\mu\) by 4–8\(\mu\).

Genus Diplostauron Korschikoff. Rectangular with raised corners; 2 equally long flagella; chromatophore; one pyrenoid; stigma; 2 contractile vacuoles anterior; fresh water.

D. pentagonium (Hazen) (Fig. 112, k). 10–13\(\mu\) by 9–10\(\mu\).

Genus Gigantochloris Pascher. Unusually large form, equalling in size a colony of Eudorina; flattened; oval in front view; elongate ellipsoid in profile; membrane radially striated; 2 flagella widely apart, less than body length; chromatophore in network; numerous pyrenoids; often without stigma; in woodland pools.

G. permaxima P. (Fig. 112, l). 70–150\(\mu\) by 40–80\(\mu\) by 25–50\(\mu\).

Genus Gloeomonas Klebs. Broadly ovoid, nearly subspherical; with a delicate membrane and a thin gelatinous envelope; 2 flagella widely apart; chromatophores numerous, circular or oval discs; pyrenoids (?); stigma; 2 contractile vacuoles anterior; fresh water.

G. ovalis K. (Fig. 112, m). 38–42\(\mu\) by 23–33\(\mu\); gelatinous envelope over 2\(\mu\) thick.

Genus Scourfieldia West. Whole body flattened; ovoid in front view; membrane delicate; 2 flagella 2–5 times body length; a chromatophore; without pyrenoid or stigma; contractile vacuole anterior; nucleus central; fresh water.

S. complanata W. (Fig. 112, n). 5.2–5.7\(\mu\) by 4.4–4.6\(\mu\); fresh water.

Genus Thorakomonas Korschikoff. Flattened; somewhat irregularly shaped or ellipsoid in front view; membrane thick, enclustered with iron-bearing material, deep brown to black in color; protoplasmic body similar to that of Chlamydomonas; a chromatophore with a pyrenoid; 2 contractile vacuoles; standing fresh water.

T. sabulosa K. (Fig. 112, o). Up to 16\(\mu\) by 14\(\mu\).

Genus Coccomonas Stein. Shell smooth; globular; body not filling intracapsular space; stigma; contractile vacuole; asexual reproduction into 4 individuals; fresh water. Species (Conrad 1930).

C. orbicularis S. (Fig. 113, a). 18–25\(\mu\) in diameter; fresh water.

Genus Chlorogonium Ehrenberg. Fusiform; membrane thin and adheres closely to protoplasmic body; plate-like chromatophores usually present, sometimes ill-contoured; one or more pyrenoids; numerous scattered contractile vacuoles; usually a stigma; a central nucleus; asexual reproduction by 2 successive transverse fissions
during the motile phase; isogamy reported; fresh water.
during the motile phase; isogamy reported; fresh water. Sexuality
(Schulze, 1927); nutrition (Loefer, 1935).

*C. euchlorum* E. (Fig. 113, b). 25–70µ by 4–15µ; in stagnant water.

Genus **Phyllomonas** Korschikoff. Extremely flattened; membrane
delicate; 2 flagella; chromatophore often faded or indistinct; numer-
ous pyrenoids; with or without stigma; many contractile vacuoles;
fresh water.

![Fig. 113. a, Coccomonas orbicularis, ×500 (Stein); b, Chlorogonium
euchlorum, ×430 (Jacobsen); c, Phyllomonas phacoides, ×200 (Kor-
schikoff); d, Sphaenochloris printzi, ×600 (Printz); e, Korschikoffia
guttula, ×1670 (Pascher); f, Furcilla lobosa, ×670 (Stokes); g, Hyalo-
gonium klebsi, ×470 (Klebs); h, Polytoma uvella, ×670 (Dangeard);
i, Parapolytoma satura, ×1600 (Jameson); j, Trichloris paradoxa, ×990
(Pascher).]*

*P. phacoides* K. (Fig. 113, c). Leaf-like; rotation movement; up to
100µ long; in standing fresh water.

Genus **Sphaenochloris** Pascher. Body truncate or concave at flagel-
late end in front view; sharply pointed in profile; 2 flagella widely
apart; chromatophore large; pyrenoid; stigma; contractile vacuole
anterior; fresh water.

*S. printzi* P. (Fig. 113, d). Up to 18µ by 9µ.

Genus **Korschikoffia** Pascher. Elongate pyriform with an undu-
lating outline; anterior end narrow, posterior end more bluntly
rounded; plastic; chromatophores in posterior half; stigma absent;
contractile vacuole anterior; 2 equally long flagella; nucleus nearly
central; salt water.
PHYTOMONADINA

*K. guttula* P. (Fig. 113, e). 6–10μ by 5μ; brackish water.

Genus *Furcilla* Stokes. U-shape, with 2 posterior processes; in side view somewhat flattened; anterior end with a papilla; 2 flagella equally long; 1–2 contractile vacuoles anterior; oil droplets; fresh water.

*F. lobosa* S. (Fig. 113, f). 11–14μ long; fresh water.

Genus *Hyalogonium* Pascher. Elongate spindle-form; anterior end bluntly rounded; posterior end more pointed; 2 flagella; protoplasm colorless; with starch granules; a stigma; asexual reproduction results in up to 8 daughter cells; fresh water.

*H. klebsi* P. (Fig. 113, g). 30–80μ by up to 10μ; stagnant water.

Genus *Polytoma* Ehrenberg (*Chlamydoblepharis* Francé; *Tussetia* Pascher). Ovoid; no chromatophores; membrane yellowish to brown; pyrenoid unobserved; 2 contractile vacuoles; 2 flagella about body length; stigma if present, red or pale-colored; many starch bodies and oil droplets in posterior half of body; asexual reproduction in motile stage; isogamy (Dogiel, 1935); saprozoic; in stagnant fresh water. Genetics (p. 231).

*P. uvella* E. (Figs. 8, c; 97, a, b; 113, h). Oval to pyriform; stigma may be absent; 15–30μ by 9–20μ. Cytology (Entz, 1918; Hollande, 1942).

Genus *Parapolytoma* Jameson. Anterior margin obliquely truncate, resembling a cryptomonad, but without chromatophores; without stigma and starch; division into 4 individuals within envelope; fresh water.

*P. sutara* J. (Fig. 113, i). About 15μ by 10μ; fresh water.

Family 2 Trichlorididae

Genus *Trichloris* Scherffel and Pascher. Bean-shape; flagellate side flattened or concave; opposite side convex; chromatophore large, covering convex side; 2 pyrenoids surrounded by starch granules; a stigma near posterior end of chromatophore; nucleus central; numerous contractile vacuoles scattered; 3 flagella near anterior end.

*T. paradoxa* S and P. (Fig. 113, j). 12–15μ broad by 10–12μ high; flagella up to 30μ long.

Family 3 Carteriidae

Genus *Carteria* Diesing (*Corbierea, Pithiscus* Dangeard). Ovoid, chromatophore cup-shaped; pyrenoid; stigma; 2 contractile vacuoles; fresh water. Numerous species (Pascher, 1925, 1932; Schiller, 1925).
**C. cordiformis** (Carter) (Fig. 114, a). Heart-shaped in front view; ovoid in profile; chromatophore large; 18–23μ by 16–20μ.

**C. ellipsoidalis** Bold. Ellipsoid; chromatophore; a small stigma; division into 2, 4, or 8 individuals in encysted stage; 6–24μ long; fresh water, Maryland (Bold, 1938).

Genus **Pyramimonas** Schmarda (**Pyramidomonas** Stein). Small pyramidal or heart-shaped body; with bluntly drawn-out posterior end; usually 4 ridges in anterior region; 4 flagella; green chromatophore cup-shaped; with or without stigma; a large pyrenoid in the posterior part; 2 contractile vacuoles in the anterior portion; encystment; fresh water. Several species (Geitler, 1925).

**P. tetrarhynchus** S. (Fig. 114, b). 20–28μ by 12–18μ; fresh water; Wisconsin (Smith, 1933).

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**Fig. 114.** a, *Carteria cordiformis*, ×600 (Dill); b, *Pyramimonas tetrarhynchus*, ×400 (Dill); c, d, *Polytomella agilis*, ×1000 (Doflein) (d, a cyst); e, *Spirogonium chlorogonioides*, ×670 (Pascher); f, *Tetrablepharis multifilis*, ×670 (Pascher); g, *Spermatozopsis exultans*, ×1630 (Pascher); h, *Chloraster gyrota*, ×670 (Stein); i, *Polyblepharides singularis*, ×870 (Dangeard); j, k, *Pocillomonas flos aquae*, ×920 (Steinecke); l, m, *Phacotus lenticularis*, ×430 (Stein); n, *Pteromonas angulosa*, ×670 (West); o, p, *Dysmorphococcus variabilis*, ×1000 (Bold).
P. montana Geitler. Bluntly conical; anterior end 4-lobed or truncate; posterior end narrowly rounded; plastic; pyriform nucleus anterior, closely associated with 4 flagella; stigma; 2 contractile vacuoles anterior; chromatophore cup-shaped, granular, with scattered starch grains and oil droplets; a pyrenoid with a ring of small starch grains; 17–22.5μ long (Geitler, 1925); 12–20μ by 8–16μ (Bold); flagella about body length; fresh water, Maryland (Bold, 1938).

Genus Polytomella Aragão. Ellipsoid, or oval, with a small papilla at anterior end, where 4 equally long flagella arise; with or without stigma; starch; fresh water (Aragão, 1910; Doflein, 1916).

P. agilis A. (Fig. 114, c, d). Numerous starch grains; 8–18μ by 5–9μ; flagella 12–17μ long; fresh water; hay infusion.

P. caeca Pringsheim. Ovoid with bluntly pointed posterior end; 12–20μ by 10–12μ; membrane delicate; a small papilla at anterior end; no stigma; two contractile vacuoles below papilla; cytoplasm ordinarily filled with starch grains; fresh water (Pringsheim, 1937).

Genus Medusochloris Pascher. Hollowed hemisphere with 4 processes, each bearing a flagellum at its lower edge; a lobed plate-shaped chromatophore; without pyrenoid. One species.

M. phiale P. In salt water pools with decaying algae in the Baltic.

Genus Spirogonium Pascher. Body spindle-form; membrane delicate; flagella a little longer than body; chromatophore conspicuous; a pyrenoid; stigma anterior; 2 contractile vacuoles; fresh water. One species.

S. chlorogonioides (P.). (Fig. 114, e). Body up to 25μ by 15μ.

Genus Tetrablepharis Senn. Ellipsoid to ovoid; pyrenoid present; fresh water.

T. multifilis (Klebs) (Fig. 114, f). 12–20μ by 8–15μ; stagnant water.

Genus Spermatozopsis Korschikoff. Sickle-form; bent easily, occasionally plastic; chromatophore mostly on convex side; a distinct stigma at more rounded anterior end; flagella equally long; 2 contractile vacuoles anterior; fresh water infusion.

S. exultans K. (Fig. 114, g). 7–9μ long; also biflagellate; in fresh water with algae, leaves, etc.

Family 4 Chlorasteridae

Genus Chloraster Ehrenberg. Similar to Pyramimonas, but anterior half with a conical envelope drawn out at four corners; with 5 flagella; fresh or salt water.

C. gyrans E. (Fig. 114, h). Up to 18μ long; standing water; also reported from salt water.
Family 5 Polyblepharididae Dangeard

Genus Polyblepharides Dangeard. Ellipsoid or ovoid; flagella 6–8, shorter than body length; chromatophore; a pyrenoid; a central nucleus; 2 contractile vacuoles anterior; cysts; a questionable genus; fresh water.

*P. singularis* D. (Fig. 114, i). 10–14µ by 8–9µ.

Genus Pocillomonas Steinecke. Ovoid with broadly concave anterior end; covered with gelatinous substance with numerous small projections; 6 flagella; chromatophores disc-shaped; 2 contractile vacuoles anterior; nucleus central; starch bodies; without pyrenoid.

*P. flos aquae* S. (Fig. 114, j, k). 13µ by 10µ; fresh water pools.

Family 6 Phacotidae Poche

The shell typically composed of 2 valves; 2 flagella protrude from anterior end; with stigma and chromatophores; asexual reproduction within the shell; valves may become separated from each other owing to an increase in gelatinous contents.

Genus Phacotus Perty. Oval to circular in front view; lenticular in profile; protoplasmic body does not fill dark-colored shell completely; flagella protrude through a foramen; asexual reproduction into 2 to 8 individuals; fresh water.

*P. lenticularis* (Ehrenberg) (Fig. 114, l, m). 13–20µ in diameter; in stagnant water.

Genus Pteromonas Seligo. Body broadly winged in plane of suture of 2 valves; protoplasmic body fills shell; chromatophore cup-shaped; one or more pyrenoids; stigma; 2 contractile vacuoles; asexual reproduction into 2–4 individuals; sexual reproduction by isogamy; zygotes usually brown; fresh water. Several species.

*P. angulosa* (Lemmermann) (Fig. 114, n). With a rounded wing and 4 protoplasmic projections in profile; 13–17µ by 9–20µ; fresh water.

Genus Dysmorphococcus Takeda. Circular in front view; anterior region narrowed; posterior end broad; shell distinctly flattened posteriorly, ornamented by numerous pores; sutural ridge without pores; 2 flagella; 2 contractile vacuoles; stigma, pyrenoid, cup-shaped chromatophore; nucleus; multiplication by binary fission; fresh water.

*D. variabilis* T. (Fig. 114, o, p). Shell 14–19µ by 13–17µ; older shells dark brown; fresh water; Maryland (Bold, 1938).
PHYTOMONADINA

Family 7 Volvocidae Ehrenberg

An interesting group of colonial flagellates; individual similar to Chlamydomonadidae, with 2 equally long flagella (one in Mastigosphaera; 4 in Spondylomorum), green chromatophores, pyrenoids, stigma, and contractile vacuoles; body covered by a cellulose membrane and not plastic; colony or coenobium is discoid or spherical; exclusively freshwater inhabitants.

Genus Volvox Linnaeus. Often large spherical or subspherical colonies, consisting of a large number of cells which are differentiated into somatic and reproductive cells; somatic cells numerous, embedded in gelatinous matrix, and contains a chromatophore, one or more pyrenoids, a stigma, 2 flagella and several contractile vacuoles; in some species cytoplasmic connection occurs between adjacent cells; generative cells few and large. Reproduction is by parthenogenesis or true sexual fusion. In parthenogenetic colonies, the gametes are larger in size and fewer in number as compared with the macrogametes of the female colonies. Sexual fusion is anisogamy (Fig. 77) and sexual colonies may be monoecious or dioecious. Zygotes are usually yellowish to brownish red in color and covered by a smooth, ridged or spinous wall. Fresh water. Many species. Smith (1944) made a comprehensive study of 18 species on which the following species descriptions are based.

V. globator L. (Fig. 115, a, b). Monoecious. Sexual colonies 350–500μ in diameter; 5000–15,000 cells, with cytoplasmic connections; 3–7 microgametocytes, each of which develops into over 250 microgametes; 10–40 macrogametes; zygotes 35–45μ in diameter, covered with many spines with rounded tip. Parthenogenetic colonies 400–600μ in diameter; 4–10 gametes, 10–13μ in diameter; young colonies up to 250μ. Europe and North America.

V. aureus Ehrenberg (Figs. 77; 115, c–e). Dioecious. Male colonies 300–350μ in diameter; 1000–1500 cells, with cytoplasmic connections; numerous microgametocytes; clusters of some 32 microgametes, 15–18μ in diameter. Female colonies 300–400μ; 2000–3000 cells; 10–14 macrogametes; zygotes 40–60μ with smooth surface. Parthenogenetic colonies up to 500μ; 4–12 gametes; young colonies 150μ in diameter. Europe and North America. Sexual differentiation (Mainx, 1929).

V. tertius Meyer. Dioecious. Male colonies up to 170μ in diameter; 180–500 cells, without cytoplasmic connections; about 50 microgametocytes. Female colonies up to 500μ; 500–2000 cells; 2–12 macrogametes; zygotes 60–65μ with smooth wall. Parthenogenetic
Fig. 115. Species of Volvox (Smith). a, b, *Volvox globator* (a, a female colony, ×150; b, a zygote, ×370); c–e, *V. aureus* (c, a young parthenogenetic colony; d, a mature male colony, ×125; e, a zygote, ×370); f–h, *V. spermatosphaera*: f, a parthenogenetic colony, ×185; g, a mature male colony, ×370; h, a zygote, ×370); i, a zygote of *V. weismannia*, ×370; j, k, *V. perglobator* (j, a male colony, ×150; k, a zygote, ×370).
colonies up to 600µ in diameter; 500–2000 cells; 2–12 gametes. Europe and North America.

*V. spermatosphaera* Powers (Fig. 115, f–h). Dioecious. Male colonies up to 100µ in diameter; cells, without connection, up to 128 microgametocytes. Female colonies up to 500µ in diameter; 6–16 macrogametes; zygotes 35–45µ, with smooth membrane. Parthenogenetic colonies up to 650µ in diameter; 8–10 gametes; young colonies ellipsoid, up to 100µ in diameter. North America (Powers, 1908).

*V. weismannia* P. (Fig. 115, i). Male colonies 100–150µ in diameter; 250–500 cells; 6–50 microgametocytes; clusters of microgametes (up to 128) discoid, 12–15µ in diameter. Female colonies up to 400µ; 2000–3000 cells; 8–24 macrogametes; zygotes 30–50µ in diameter, with reticulate ridges on shell. Parthenogenetic colonies up to 400µ; 1500–3000 cells; 8 or 10 gametes; 40–60µ in diameter; young colonies 100–200µ in diameter. North America (Powers, 1908).

*V. perglobator* P. (Fig. 115, j, k). Dioecious. Male colonies 300–450µ in diameter 5000–10,000 cells, with delicate cytoplasmic connections; 60–80 microgametocytes. Female colonies 300–550µ in diameter; 9000–13,000 cells; 50–120 macrogametes; zygotes 30–34µ, covered with bluntly pointed spines. Parthenogenetic colonies as large as 1.1 mm; three to nine gametes; young colonies 250–275µ in diameter. North America.

Genus *Gonium* Müller. 4 or 16 individuals arranged in one plane; cell ovoid or slightly polygonal; with 2 flagella arranged in the plane of coenobium; with or without a gelatinous envelope; protoplasmic connections among individuals occur occasionally; asexual reproduction through simultaneous divisions of component cells; sexual reproduction isogamy; zygotes reddish; fresh water. Colony formation (Hartmann, 1924).

*G. sociale* (Dujardin) (Fig. 116, a). 4 individuals form a discoid colony; cells 10–22µ by 6–16µ wide; in open waters of ponds and lakes.

*G. pectorale* M. (Fig. 116, b). 16 (rarely 4 or 8) individuals form a colony; 4 cells in center; 12 peripheral, closely arranged; cells 5–14µ by 10µ; colony up to 90µ in diameter; fresh water.

*G. formosum* Pascher. 16 cells in a colony further apart; peripheral gelatinous envelope reduced; cells similar in size to those of *G. sociale* but colony somewhat larger; freshwater lakes.

Genus *Stephanoon* Schewiakoff. Spherical or ellipsoidal colony, surrounded by gelatinous envelope, and composed of 8 or 16 bi-
flagellate cells, arranged in 2 alternating rows on equatorial plane; fresh water.

*S. askenasii* S. (Fig. 117, a). 16 individuals in ellipsoidal colony; cells 9μ in diameter; flagella up to 30μ long; colony 78μ by 60μ.

Genus *Platydorina* Kofoid. 32 cells arranged in a slightly twisted plane; flagella directed alternately to both sides; dioecious; fresh water.

*P. caudata* K. (Fig. 117, b). Individual cells 10–15μ long; colony up to 165μ long by 145μ wide, and 25μ thick; dioecious; anisogamy; macrogametes escape from female colonies and remain attached to them or swim about until fertilized by microgametes; zygotes become thickly walled (Taft, 1940).

Genus *Spondylomorum* Ehrenberg. 16 cells in a compact group in 4 transverse rings; each with 4 flagella; asexual reproduction by simultaneous division of component cells; fresh water. One species.

*S. quaternarium* E. (Fig. 117, c). Cells 12–26μ by 8–15μ; colony up to 60μ long.

Genus *Chlamydoebotrys* Korschikoff. Colony composed of 8 or 16 individuals; cells with 2 flagella; chromatophore; stigma; no pyrenoid; fresh water. Species (Pascher, 1925); culture (Schulze, 1927).

*C. stellata* K. (Fig. 117, d). Colony composed of 8 individuals arranged in 2 rings; individuals 14–15μ long; colony 30–40μ in diameter; Maryland (Bold, 1933).
Genus **Stephanosphaera** Cohn. Spherical or subspherical colony, with 8 (rarely 4 or 16) cells arranged in a ring; cells pyriform, but with several processes; 2 flagella on one face; asexual reproduction and isogamy (p. 183); fresh water.

![Diagram](image)

**Fig. 117.** a, *Stephanon askenasii*, ×440 (Schewiakoff); b, *Platydorina caudata*, ×280 (Kofoid); c, *Spondylomorum quaternarium*, ×330 (Stein); d, *Chlamydothrix stellata*, ×450 (Korschikoff); e, *Stephanosphaera pluvialis*, ×250 (Hieronymus); f, *Pandorina morum*, ×300 (Smith); g, *Mastigosphaera gobii*, ×520 (Schewiakoff); h, *Eudorina elegans*, ×310 (Goebel); i, *Pleodorina illinoisensis*, ×200 (Kofoid).

*S. pluvialis* C. (Figs. 80; 117, e). Cells 7–13μ long; colony 30–60μ in diameter. Culture and sexuality (Schulze, 1927).

Genus **Pandorina** Bory. Spherical or subspherical colony of usually 16 (sometimes 8 or 32) biflagellate individuals, closely packed within a gelatinous, but firm and thick matrix; individuals often angular; with stigma and chromatophores; asexual reproduction
through simultaneous division of component individuals; anisogamy; zygotes colored and covered by a smooth wall; fresh water. One species.

*P. morum* (Müller) (Figs. 117, f). Cells 8–17μ long; colony 20–40μ, up to 250μ in diameter; ponds and ditches.

Genus *Mastigosphera* Schewiakoff. Similar to *Pandorina*; but individuals with a single flagellum which is 3.5 times the body length; fresh water.

*M. gobii* S. (Fig. 117, g). Individual 9μ long; colony 30–33μ.

Genus *Eudorina* Ehrenberg. Spherical or ellipsoidal colony of usually 32 or sometimes 16 spherical cells; asexual reproduction similar to that of *Pandorina*; sexual reproduction with 32–64 spherical green macrogametes and numerous clustered microgametes which when mature, unite with the macrogametes within the colony; reddish zygotes with a smooth wall; fresh water. Colony formation (Hartmann, 1924).

*E. elegans* E. (Fig. 117, h). Cells 10–24μ in diameter; colony 40–150μ in diameter; in ponds, ditches and lakes. Culture and morphology (Hartmann, 1921); response to light (Luntz, 1935).

Genus *Pleodorina* Shaw. Somewhat similar to *Eudorina*, being composed of 32, 64, or 128 ovoid or spherical cells of 2 types: small somatic and large generative, located within a gelatinous matrix; Sexual reproduction similar to that of *Eudorina*; fresh water.

*P. illinoisensis* Kofoid (Figs. 32, b, c; 117, i). 32 cells in ellipsoid colony, 4 vegetative and 28 reproductive individuals; arranged in 5 circles, 4 in each polar circle, 8 at equator and 8 on either side of equator; 4 small vegetative cells at anterior pole; vegetative cells 10–16μ in diameter; reproductive cells 19–25μ in diameter; colony up to 160μ by 130μ.

*P. californica* S. Spherical colony with 64 or 128 cells, of which 1/2–2/3 are reproductive cells; vegetative cells 13–15μ; reproductive cells up to 27μ; colony up to 450μ, both in diameter. Variation (Tiffany, 1935); in Ukraine (Swirenko, 1926).

**References**


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——— (1933a) Untersuchungen über die Sexualität und Entwick- lungen von Chlorophyceen. Ibid., 80:469.


——— (1925) Neue oder wenig bekannte Protisten. XVII. Ibid., 51: 549.
XVIII. Ibid., 52:566.


(1930) Neue Volvocalen. Ibid., 69:103.

(1932) Zur Kenntnis der einzelligen Volvocalen. Ibid., 76:1.


(1908) Further studies in Volvox with descriptions of three new species. Ibid., 28:141.


(1937) Zur Kenntnis saprotropher Algen und Flagellaten. II. Ibid., 88:151.


Chapter 11
Order 4 Euglenoidina Blochmann

The body is as a rule elongated; some are plastic, others have a definite body form with a well-developed, striated or variously sculptured pellicle. At the anterior end, there is an opening through which a flagellum protrudes. In holophytic forms the so-called cyto-stome and cytopharynx, if present, are apparently not concerned with the food-taking, but seem to give a passage-way for the flagellum and also to excrete the waste fluid matters which become collected in one or more contractile vacuoles located near the reservoir. In holozoic forms, a well-developed cytostome and cytopharynx are present. Ordinarily there is only one flagellum, but some possess two or three. Chromatophores are present in the majority of the Euglenidae, but absent in two families. They are green, vary in shape, such as spheroidal, band-form, cup-form, discoidal, or fusiform, and usually possess pyrenoids. Some forms may contain haematochrome. A small but conspicuous stigma is invariably present near the anterior end of the body in chromatophore-bearing forms.

Reserve food material is the paramylon body, fat, and oil, the presence of which depends naturally on the metabolic condition of the organism. The paramylon body assumes diverse forms in different species, but is, as a rule, constant in each species, and this facilitates specific identification to a certain extent. Nutrition is holophytic in chromatophore-possessing forms, which, however, may be saprozoic, depending on the amount of light and organic substances present in the water. The holozoic forms feed upon bacteria, algae, and smaller Protozoa.

The nucleus is, as a rule, large and distinct and contains almost always a large endosome. Asexual reproduction is by longitudinal fission; sexual reproduction has been observed in a few species. Encystment is common. The majority inhabit fresh water, but some live in brackish or salt water, and a few are parasitic in animals. Taxonomy (Mainx, 1928; Hollande, 1942, 1952a); Jahn, 1946; Pringsheim, 1950.

With stigma .................................. Family 1 Euglenidae (p. 294)
Without stigma
   With 1 flagellum ......................... Family 2 Astasiidae (p. 302)
   With 2 flagella ....................... Family 3 Anisonemidae (p. 303)
Family 1 Euglenidae Stein

Body plastic ("euglenoid"), but, as a rule, more or less spindle-form during locomotion. The flagellum arises from a blepharoplast located in the cytoplasm at the posterior margin of the reservoir. Between the blepharoplast and the "cytostome," the flagellum shows a swelling which appears to be photosensitive (Mast, 1938). Many observers consider that the basal portion of the flagellum is bifurcated and ends in two blepharoplasts, but Hollande (1942), Pringsheim (1948) and others, hold that in addition to a long flagellum arising from a blepharoplast, there is present a short flagellum which does not extend beyond the neck of the reservoir and often adheres to the long flagellum, producing the appearance of bifurcation. Culture and physiology (Mainx, 1928); cytology (Günther, 1928; Hollande, 1942).

Genus Euglena Ehrenberg. Short or elongated spindle, cylindrical, or band-form; pellicle usually marked by longitudinal or spiral striae; some with a thin pellicle highly plastic; others regularly spirally twisted; stigma usually anterior; chromatophores numerous and discoid, band-form, or fusiform; pyrenoids may or may not be surrounded by starch envelope; paramylon bodies which may be two in number, one being located on either side of nucleus, and rod-like to ovoid in shape or numerous and scattered throughout; contractile vacuole small, near reservoir; asexual reproduction by longitudinal fission; sexual reproduction reported in Euglena sanguinea; common in stagnant water, especially where algae occur; when present in large numbers, the active organisms may form a green film on the surface of water and resting or encysted stages may produce conspicuous green spots on the bottom of pond or pool; in fresh water. Numerous species (Pascher, 1925; Johnson, 1944; Gojdics, 1953).

E. pisciformis Klebs (Fig. 118, a). 20–35μ by 5–10μ; spindle-form with bluntly pointed anterior and sharply attenuated posterior end; slightly plastic; a body-length flagellum, active; 2–3 chromatophores; division into two or four individuals in encysted stage (Johnson, 1944).

E. viridis Ehrenberg (Fig. 118, b). 40–65μ by 14–20μ; anterior end rounded, posterior end pointed; fusiform during locomotion; highly plastic when stationary; flagellum as long as the body; pellicle obliquely striated; chromatophores more or less bandform, radially arranged; nucleus posterior; nutrition holophytic, but also saprozoic. Multiplication in thin-walled cysts (Johnson).

E. acus E. (Fig. 118, c). 50–175μ by 8–18μ; body long spindle or
cylinder, with a sharply pointed posterior end; flagellum short, about \( \frac{1}{4} \) the body length; spiral striation of pellicle very delicate; numerous discoid chromatophores; several paramylon bodies, rod-form and 12–20\( \mu \) long; nucleus central; stigma distinct; movement sluggish.

**E. spirogyra** E. (Fig. 118, d). 80–125\( \mu \) by 10–35\( \mu \); cylindrical; anterior end a little narrowed and rounded, posterior end drawn out; spiral striae, made up of small knobs, conspicuous; many discoid chromatophores; two ovoidal paramylon bodies, 18–45\( \mu \) by 10–18\( \mu \), one on either side of centrally located nucleus; flagellum about \( \frac{1}{4} \) the body length; stigma prominent; sluggish.

**E. oxyuris** Schmarda (Fig. 118, e). 150–500\( \mu \) by 20–40\( \mu \); cylindri-
cal; almost always twisted, somewhat flattened; anterior end round-
ed, posterior end pointed; pellicle with spiral striae; numerous dis-
coid chromatophores; two ovoid paramylon bodies, 20–40μ long, one
on either side of nucleus, and also small bodies; stigma large; flagel-
num short; sluggish.

E. sanguinea E. (Fig. 118, f). 80–170μ by 25–45μ; posterior end
bluntly rounded; flagellum about the body length; pellicle striated;
elongate chromatophores lie parallel to the striae; haematochrome
granules scattered in sun light and collected in the central area in
darkness.

E. deses E. (Fig. 118, g). 85–170μ by 10–20μ; elongate; highly
plastic; faint striae; stigma distinct; nucleus central; chromatop-
hores discoid with pyrenoid; several small rod-shaped paramylon
scattered; flagellum less than ¼ the body length.

E. gracilis Klebs (Fig. 118, h). 35–55μ by 6–25μ; cylindrical to
elongate oval; highly plastic; flagellum about the body length; fus-
iform chromatophores 10–20; nucleus central; pyrenoids.

E. tripteris Dujardin (Fig. 118, i). 70–120μ by 12–16μ; elongate;
three-ridged, rounded anteriorly and drawn out posteriorly; pellicle
longitudinally striated; only slightly plastic; stigma prominent; dis-
coid chromatophores numerous; two paramylon bodies, rod-shaped
and one on either side of the nucleus; flagellum about ¾ the body
length.

E. ehrenbergi Klebs (Fig. 118, j). 170–400μ by 15–40μ; cylindrical
and flattened, posterior end rounded; plastic, often twisted; spiral
striation; numerous small discoid chromatophores; stigma conspicu-
ous; 2 paramylon bodies elongate, up to over 100μ long; flagellum
about ¼ the body length or less.

E. terricola Dangeard (Fig. 118, k). 65–95μ by 8–18μ; pellicle thin
and highly plastic; nucleus central; chromatophores long (20–30μ)
rods; paramylon bodies small and annular; flagellum about ½ the
body length.

E. sociabilis D. (Fig. 118, l). 65–112μ by 15–30μ; cylindrical; deli-
cate pellicle; highly plastic; numerous elongate chromatophores;
paramylon bodies discoid; flagellum slightly longer than body.

E. klebsi Mainx (Fig. 118, m). 45–85μ by 5–10μ; form highly
plastic; chromatophores discoid; paramylon bodies rod-shaped, up
to several; flagellum short.

E. rubra Hardy (Fig. 118, n). 70–170μ by 25–36μ; cylindrical;
rounded anteriorly and drawn out posteriorly; spiral striation; nu-
cleus posterior; flagellum longer than body; stigma about 7μ in di-
ameter; many fusiform chromatophores aligned with the body striae;
numerous haematochrome granules, 0.3–0.5\(\mu\) in diameter; ovoid paramylon bodies; reproductive and temporary cysts and protective cysts, 34–47\(\mu\) in diameter, with a gelatinous envelope.

Johnson (1939) found that the color of this Euglena was red in the morning and dull green in the late afternoon, due to the difference in the distribution of haematochrome within the body. When haematochrome granules are distributed throughout the body, the organism is bright-red, but when they are condensed in the center of the body, the organism is dull green. When part of the area of the pond was shaded with a board early in the morning, shortly after sunrise all the scum became red except the shaded area. When the board was removed, the red color appeared in 11 minutes while the temperature of the water remained 21°C. In the evening the change was reversed. Johnson and Jahn (1942) later found that green-red color change could be induced by raising the temperature of the water to 30–40°C. and by irradiation with infrared rays or visible light. The two workers hold that the function of haematochrome may be protective, since it migrates to a position which shields the chromatophores from very bright light. If this is true, it is easy to find the species thriving in hot weather in shallow ponds where temperature of the water rises to 35–45°C. In colder weather, it is supposed that this Euglena is less abundant and it exists in a green phase, containing a few haematochrome granules.

Genus Khawkinea Jahn and McKibben. Similar to Genus Euglena, but without chromatophores and thus permanently colorless; fresh water.

K. halli J. and M. 30–65\(\mu\) by 12–14\(\mu\); fusiform; pellicle spirally striated; plastic; flagellum slightly longer than body; stigma 2–3\(\mu\) in diameter, yellow-orange to reddish-orange, composed of many granules; numerous (25–100) paramylon bodies elliptical or polyhedral; cysts 20–30\(\mu\) in diameter; putrid leaf infusion; saprozoic (Jahn and McKibben, 1937).

K. ocellata (Khawkine). Similar to above; flagellum 1.5–2 times body length; fresh water.

Genus Phacus Dujardin. Highly flattened; asymmetrical; pellicle firm; body form constant; prominent longitudinal or oblique stria- tion; flagellum and a stigma; chromatophores without pyrenoid (Pringsheim) are discoid and green; holophytic; fresh water. Numerous species (Skvortzov, 1937; Pochmann, 1942; Conrad, 1943; Allegre and Jahn, 1943); Morphology and cytology (Krichenbauer, 1937; Conrad, 1943).

P. pleuronectes (Müller) (Fig. 119, a). 45–100\(\mu\) by 30–70\(\mu\); short
posterior prolongation slightly curved; a prominent ridge on the convex side, extending to posterior end; longitudinally striated; usually one circular paramylon body near center; flagellum as long as body.

*P. longicauda* (Ehrenberg) (Fig. 119, b). 120–170 µ by 45–70 µ; usually slightly twisted; a long caudal prolongation; flagellum about

\[ \frac{1}{2} \text{ the body length}; \text{ stigma prominent; numerous chromatophores; one discoidal paramylon body central; pellicle longitudinally striated.} \]

*P. pyrum* (E.) (Fig. 119, c). About 30–50 µ by 10–20 µ; circular in cross-section; with a medium long caudal prolongation; pellicle obliquely ridged; stigma inconspicuous; two discoid paramylon bodies; flagellum as long as the body.

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**Fig. 119.** Species of Phacus (Allegre and Jahn). a, *Phacus pleuronectes* and an end view, ×800; b, *P. longicauda*, ×500; c, *P. pyrum* and an end view, ×880; d, *P. acuminata* and an end view, ×1300; e, *P. monilata*, ×800; f, *P. torta*, and an end view, ×800; g, *P. oscillans*, ×1400.
P. acuminata Stokes (Fig. 119, d). About 30–40 μ by 20–30 μ; nearly circular in outline; longitudinally striated; usually one small paramylon body; flagellum as long as the body.

P. monilata (S) (Fig. 119, e). 40–55 μ by 32–40 μ; a short caudal projection; pellicle with minute knobs arranged in longitudinal rows; discoid chromatophores; flagellum about the body length.

P. torta Lemmermann (Fig. 119, f). 80–100 μ by 40–45 μ; body twisted, with a long caudal prolongation; longitudinal striae on pellicle; chromatophores discoid; one large circular paramylon body; flagellum about 1/2 the body length.

P. oscillans Klebs (Fig. 119, g). 15–35 μ by 7–10 μ; rounded anteriorly and bluntly pointed posteriorly; striation oblique; 1 or 2 paramylon bodies; flagellum about as long as the body.

Genus Lepocinclis Perty (Crumenula Dujardin). Body more or less ovo-cylindrical; rigid with spirally striated pellicle; often with a short posterior spinous projection; stigma sometimes present; discoidal chromatophores numerous and marginal; paramylon bodies usually large and ring-shaped, laterally disposed; without pyrenoids; fresh water. Many species (Pascher, 1925, 1929: Conrad, 1934; Skvortzov, 1937).

L. ovum (Ehrenberg) (Fig. 120, a). Body 20–40 μ long.

Genus Trachelomonas Ehrenberg. With a lorica which often possesses numerous spines; sometimes yellowish to dark brown, composed of ferric hydroxide impregnated with a brown manganic compound (Pringsheim, 1948); a single long flagellum protrudes from the anterior aperture, the rim of which is frequently thickened to form a collar; chromatophores either two curved plates or numerous discs; paramylon bodies small grains; a stigma and pyrenoid; multiplication by fission, one daughter individual retains the lorica and flagellum, while the other escapes and forms a new one; cysts common; fresh water. Numerous species (Palmer, 1902, 1905, 1925a; Pascher, 1924, 1925, 1925a, 1926, 1929; Gordienko, 1929; Conrad, 1932; Skvortzov, 1937; Balech, 1944).

T. hispida (Perty) (Figs. 32, a; 120, b). Lorica oval, with numerous minute spines; brownish; 8–10 chromatophores; 20–42 μ by 15–26 μ; many varieties.

T. urceolata Stokes (Fig. 120, c). Lorica vasiform, smooth with a short neck; about 45 μ long.

T. piscatoris (Fisher) (Fig. 120, d). Lorica cylindrical with a short neck and with numerous short, conical spines; 25–40 μ long; flagellum 1–2 times body length.
T. verrucosa Stokes (Fig. 120, e). Lorica spherical, with numerous knob-like attachments; no neck; 24-25μ in diameter.

T. vermiculosa Palmer (Fig. 120, f). Lorica spherical; with many sausage-form markings; 23μ in diameter.

Genus Cryptoglena Ehrenberg. Body rigid, flattened; 2 band-form chromatophores lateral; a single flagellum; nucleus posterior; among freshwater algae. One species.

C. pigra E. (Fig. 120, g). Ovoid, pointed posteriorly; flagellum short; stigma prominent; 10-15μ by 6-10μ; standing water.

![Fig. 120. a, Lepocinclus ovum, ×430 (Stein); b, Trachelomonas hispida, ×430 (Stein); c, T. urecolata, ×430 (Stokes); d, T. piscatoris, ×520 (Fisher); e, T. verrucosa, ×550 (Stokes); f, T. vermiculosa, ×800 (Palmer); g, Cryptoglena pigra, ×430 (Stein); h, Ascoglena vaginicola, ×390 (Stein); i, Eutreptia viridis, ×270 (Klebs); j, E. marina, ×670 (da Cunha); k, Euglenamorpha hegneri, ×730 (Wenrich).](image)

Genus Ascoglena Stein. Encased in a flexible, colorless to brown lorica, attached with its base to foreign object; solitary; without stalk; body ovoidal, plastic; attached to test with its posterior end; a single flagellum; a stigma; numerous chromatophores discoid; with or without pyrenoids; reproduction as in Trachelomonas; fresh water.

A. vaginicola S. (Fig. 120, h). Lorica about 43μ by 15μ.

Genus Colacium Ehrenberg. Stalked individuals form colony; frequently attached to animals such as copepods, rotifers, etc.; stalk mucilaginous; individual cells pyriform, ellipsoidal or cylindrical; without flagellum; a single flagellum only in free-swimming stage; discoidal chromatophores numerous; with pyrenoids; multiplication
by longitudinal fission; also by swarmers, possessing a flagellum and a stigma; fresh water. Several species.

*C. vesiculosum* E. (Fig. 121). Solitary or colonial, made up of two to eight individuals; flagellate form ovoid to spindle; 22μ by 12μ; seven to ten elongate chromatophores along the periphery; flagellum

![Diagram of Colacium vesiculosum](image)

one to two times the body length; a stigma; many paramylon bodies; palmella stage conspicuous; stalked form (Johnson, 1934).

Genus *Eutreptia* Perty (*Eutreptiella* da Cunha). With 2 flagella at anterior end; pellicle distinctly striated; plastic; spindle-shaped during movement; stigma; numerous discoid chromatophores; pyrenoids absent; paramylon bodies spherical or subcylindrical; multiplication as in Euglena; cyst with a thick stratified wall; fresh or salt water.

*E. viridis* P. (Fig. 120, i). 50–70μ by 5–13μ; in fresh water; a variety was reported from brackish water ponds.

*E. marina* (da Cunha) (Fig. 120, j). Flagella unequal in length;
longer one as long as body, shorter one \( \frac{3}{4} \); body 40–50\( \mu \) by 8–10\( \mu \); salt water.

Genus *Euglenamorpha* Wenrich. Body form and structure similar to those of *Euglena*, but with 3 flagella; in gut of frog tadpoles. One species.

*E. hegneri* W. (Fig. 120, k). 40–50\( \mu \) long (Wenrich, 1924).

Family 2 *Astasiidae* Bütschli

Similar to Euglenidae in body form and general structure, but without chromatophores; body highly plastic, although usually elongate spindle.

Genus *Astasia* Dujardin. Body plastic, although ordinarily elongate; fresh water or parasitic (?) in microcrustaceans. Many species (Pringsheim, 1942). Bacteria-free cultivation (Schoenborn, 1946).

*A. klebsi* Lemmermann (Fig. 122, a). Spindle-form; posterior portion drawn out; flagellum as long as body; plastic; paramylon bodies oval; 40–50\( \mu \) by 13–20\( \mu \); stagnant water.

Genus *Urceolus* Mereschkowsky (*Phialonema* Stein). Body colorless; plastic; flask-shaped; striated; a funnel-like neck; posterior region stout; a single flagellum protrudes from funnel and reaches inward the posterior third of body; fresh or salt water.

*U. cyclostomus* (Stein) (Fig. 122, b). 25–50\( \mu \) long; fresh water.

*U. sabulosus* (Stokes) (Fig. 122, c). Spindle-form; covered with minute sand-grains; about 58\( \mu \) long; fresh water.

Genus *Petalomonas* Stein. Oval or pyriform; not plastic; pellicle often with straight or spiral furrows; a single flagellum; paramylon

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**Fig. 122.** a, *Astasia klebsi*, \( \times 500 \) (Klebs); b, *Urceolus cyclostomus*, \( \times 430 \) (Stein); c, *U. sabulosus*, \( \times 430 \) (Stokes); d, *Petalomonas mediocanellata*, \( \times 1000 \) (Klebs); e, *Rhabdomonas incurva*, \( \times 1400 \) (Hall); f, *Scytononas pusilla*, \( \times 430 \) (Stein).
bodies; a nucleus; holozoic or saprozoic. Many species in fresh water and a few in salt water. Species (Shawhan and Jahn, 1947).

*P. mediocanellata* S. (Fig. 122, *d*). Ovoid with longitudinal furrows on two sides; flagellum about as long as the body; 21–26 µ long.

**Genus Rhabdomonas** Fresenius. Rigid body, cylindrical and not flattened, more or less arched; pellicle longitudinally ridged; a flagellum through aperture at the anterior tip; fresh water (Pringsheim, 1942). Species (Pascher, 1925); relation to *Menoidium* (Pringsheim, 1942).

*R. incurva* F. (Figs. 69, 122, *e*). Banana-shaped; longitudinal ridges conspicuous; flagellum as long as the body; 15–25 µ by 7–8 µ (Hall, 1923); 13–15 µ by 5–7 µ (Hollande, 1952a); common in standing water.

**Genus Scytomonas** Stein. Oval or pyriform, with a delicate pellicle; a single flagellum; a contractile vacuole with a reservoir; holozoic on bacteria; longitudinal fission in motile stage; stagnant water and coprozoic.

*S. pusilla* S. (Fig. 122, *f*). About 15 µ long. Cytology (Schüessler, 1917).

**Genus Copromonas** Dobell. Elongate ovoid; with a single flagellum; a small cytostome at anterior end; holozoic on bacteria; permanent fusion followed by encystment (p. 183); coprozoic in faecal matters of frog, toad, and man; several authors hold that this genus is probably identical with *Scytomonas* which was incompletely described by Stein.

*C. subtilis* D. (Fig. 78). 7–20 µ long. Golgi body (Gatenby and Singh, 1938).

**Family 3 Anisonemidae** Schewiakoff

Colorless body plastic or rigid with a variously marked pellicle; 2 flagella, one directed anteriorly and the other usually posteriorly; contractile vacuoles and reservoir; stigma absent; paramylon bodies usually present; free-swimming or creeping.

**Genus Anisonema** Dujardin. Generally ovoid; more or less flattened; asymmetrical; plastic or rigid; a slit-like ventral furrow; flagella at anterior end; cytopharynx long; contractile vacuole anterior; nucleus posterior; in fresh water. Several species.

*A. acinus* D. (Fig. 123, *a*). Rigid; oval; somewhat flattened; pellicle slightly striated; 25–40 µ by 16–22 µ.

*A. truncatum* Stein (Fig. 123, *b*). Rigid; elongate ovoid: 60 µ by 20 µ.

*A. emarginatum* Stokes (Fig. 123, *c*). Rigid; 14 µ long; flagella long.
Genus **Peranema** Dujardin. Elongate, with a broad rounded or truncate posterior end during locomotion; highly plastic when stationary; delicate pellicle shows a fine striation; expansible cytostome with a thickened ridge and two oral rods at anterior end; aperture through which the flagella protrude is also at anterior end; a free flagellum, long and conspicuous, tapers toward free end; a second flagellum adheres to the pellicle; nucleus central; a contractile vacuole, anterior, close to the reservoir; holozoic; fresh water.

*P. trichophorum* (Ehrenberg) (Fig. 123, d). 40–70μ long; body ordinarily filled with paramylon or starch grains derived from Astasia, Rhabdomonas, Euglena, etc., which coinhabit the culture; holozoic; very common in stagnant water. Cell inclusion (Hall, 1929); structure and behavior (Chen, 1950); development (Lackey, 1929); flagellar apparatus (Lackey, 1933; Pitelka, 1945); food intake (Hall, 1933; Hollande, 1942; Hyman, 1936; Chen, 1950).

*P. granulifera* Penard. Much smaller in size. 8–15μ long; elongate, but plastic; pellicle granulated; standing water.

Genus **Heteronema** Dujardin. Plastic; rounded or elongate; flagella arise from anterior end, one directed forward and the other trailing; cytostome near base of flagella; holozoic; fresh water. Several species.

*H. acus* (Ehrenberg) (Fig. 123, e). Extended body tapers towards both ends; anterior flagellum as long as body, trailing one about 1/2; contractile vacuole anterior; nucleus central; 45–50μ long; fresh water. Morphology, reproduction (Loefer, 1931).

*H. mutabile* (Stokes) (Fig. 123, f). Elongate; highly plastic; longitudinally striated; about 254μ long; in cypress swamp.

Genus **Tropidoscyphus** Stein. Slightly plastic; pellicle with 8 longitudinal ridges; 2 unequal flagella at anterior pole; holozoic or saprozoic; fresh or salt water.

*T. octocostatus* S. (Fig. 123, g). 35–63μ long; fresh water, rich in vegetation.

Genus **Distigma** Ehrenberg. Plastic; elongate when extended; body surface without any marking; 2 flagella unequal in length, directed forward; cytostome and cytopharynx located at anterior end; endoplasm usually transparent; holozoic. Several species (Pringsheim, 1942).

*D. proteus* E. (Fig. 123, h). 50–110μ long when extended; nucleus central; stagnant water; infusion. Cytology (Hollande, 1937).

Genus **Entosiphon** Stein. Oval, flattened; more or less rigid; flagella arise from a cytostome, one flagellum trailing; protrusible cytopharynx a long conical tubule almost reaching posterior end;
EUGLENOIDINA

nucleus centro-lateral; fresh water.

*E. sulcatum* (Dujardin) (Fig. 123, i). About 20μ long (Lackey, 1929, 1929a).

*E. ovatum* Stokes. Anterior end rounded; 10–12 longitudinal striae; about 25–28μ long.

Genus **N**otosolenus Stokes. Free-swimming; rigid oval; ventral

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Fig. 123. a, Anisonema acinus, ×400 (Klebs); b, *A. truncatum*, ×430 (Stein); c, *A. emerginatum*, ×530 (Stokes); d, *Peranema trichophorum*, ×670; e, *Heteronema acus*, ×430 (Stein); f, *H. mutabile*, ×120 (Stokes); g, *Tropidocycphus octocostatus*, ×290 (Lemmermann); h, *Distigma proteus*, ×430 (Stein); i, *Entosiphon sulcatum*, ×430 (Stein); j, *Notosolenus apocampus*, ×120 (Stokes); k, *N. sinuatus*, ×600 (Stokes); l, m, front and side views of *Triangulomonas rigida*, ×935 (Lackey); n, *Marsupiogaster striata*, ×590 (Schewiakoff); o, *M. picta* (Faria, da Cunha and Pinto).
surface convex, dorsal surface with a broad longitudinal groove; flagella anterior; one long, directed anteriorly and vibratile; the other shorter and trailing; fresh water with vegetation.

_N. apocamptus_ S. (Fig. 123, _j_). Oval with broad posterior end; 6–11µ long.

_N. sinuatus_ S. (Fig. 123, _k_). Posterior end truncate or concave; about 22µ long.

Genus **Triangulomonas** Lackey. Rigid body, triangular, with convex sides; one surface flat, the other elevated near the anterior end; pellicle brownish; a mouth at anterior end with cytopharynx and reservoir: two flagella, one trailing; salt water.

_T. rigida_ L. (Fig. 123, _l, m_). Body 18µ by 15µ; anterior flagellum as long as the body; posterior flagellum 1.5 times the body length; Woods Hole (Lackey, 1940).

Genus **Marsupiogaster** Schewiakoff. Oval; flattened; asymmetrical; cytostome occupies entire anterior end; cytopharynx conspicuous, 1/2 body length; body longitudinally striated; 2 flagella, one directed anteriorly, the other posteriorly; spherical nucleus; contractile vacuole anterior; fresh or salt water.

_M. striata_ Schewiakoff (Fig. 123, _n_). About 27µ by 15µ; fresh water; Hawaii.

_M. picta_ Faria, da Cunha and Pinto (Fig. 123, _o_). In salt water; Rio de Janeiro.

**Order 5 Chloromonadina** Klebs

The chloromonads are of rare occurrence and consequently not well known. The majority possess small discoidal grass-green chromatophores with a large amount of xanthophyll which on addition of an acid become blue-green. No pyrenoids occur. The metabolic products are fatty oil. Starch or allied carbohydrates are absent. Stigma is also not present. Genera (Poisson and Hollande, 1943; Hollande, 1952).

Genus **Gonyostomum** Diesing (_Rhaphidomonas_ Stein). With a single flagellum: chromatophores grass-green; highly refractile trichocyst-like bodies in cytoplasm; fresh water. A few species.

_G. semen_ D. (Fig. 124, _a_). Sluggish animal; about 45–60µ long; among decaying vegetation.

Genus **Vacuolaria** Cienkowski (_Coelomonas_ Stein). Highly plastic; without trichocyst-like structures; anterior end narrow; two flagella; cyst with a gelatinous envelope. One species.

_V. virescens_ C. (Fig. 124, _b_). 50–70µ by 18–25µ; fresh water. Cytology (Fott, 1935; Poisson and Hollande, 1943).
Genus Trentonia Stokes. Bi-flagellate as in the last genus; but flattened; anterior margin slightly bilobed. One species.

*T. flagellata* S. (Fig. 124, c). Slow-moving organism; encystment followed by binary fission; about 60μ long; fresh water.

Genus Thaumatomastix Lauterborn. Colorless; pseudopodia formed; 2 flagella, one extended anteriorly, the other trailing; holo-

zoic; perhaps a transitional form between the Mastigophora and the Sarcodina. One species.

*T. setifera* L. (Fig. 104, d). About 20–35μ by 15–28μ; fresh water.

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Chapter 12
Order 6 Dinoflagellata Bütschli

THE dinoflagellates make one of the most distinct groups of the Mastigophora, inhabiting mostly marine water, and to a lesser extent fresh water. In the general appearance, the arrangement of the two flagella, the characteristic furrows, and the possession of brown chromatophores, they are closely related to the Cryptomonadina.

The body is covered by an envelope composed of cellulose which may be a simple smooth piece, or may be composed of two valves or of numerous plates, that are variously sculptured and possess manifold projections. Differences in the position and course of the furrows and in the projections of the envelope produce numerous asymmetrical forms. The furrows, or grooves, are a transverse annulus and a longitudinal sulcus. The annulus is a girdle around the middle or toward one end of the body. It may be a complete, incomplete or sometimes spiral ring. While the majority show a single transverse furrow, a few may possess several. The part of the shell anterior to the annulus is called the epitheca and that posterior to the annulus the hypotheca. In case the envelope is not developed, the terms epicone and hypocone are used (Fig. 105). The sulcus may run from end to end or from one end to the annulus. The two flagella arise typically from the furrows, one being transverse and the other longitudinal.

The transverse flagellum which is often band-form, encircles the body and undergoes undulating movements, which in former years were looked upon as ciliary movements (hence the name Cilioflagellata). In the suborder Prorocentrinae, this flagellum vibrates freely in a circle near the anterior end. The longitudinal flagellum often projects beyond the body and vibrates. Combination of the movements of these flagella produces whirling movements characteristic of the organisms.

The majority of dinoflagellates possess a single somewhat massive nucleus with evenly scattered chromatin, and usually several endosomes. There are two kinds of vacuoles. One is often surrounded by a ring of smaller vacuoles, while the other is large, contains pink-colored fluid and connected with the exterior by a canal opening into a flagellar pore. The latter is known as the pusule which functions as a digestive organella (Kofoid and Swezy). In many freshwater forms a stigma is present, and in Pouchetiidae there is an ocellus composed of an amyloid lens and a dark pigment-ball. The majority
of planktonic forms possess a large number of small chromatophores which are usually dark yellow, brown or sometimes slightly greenish and are located in the periphery of the body, while bottom-dwelling and parasitic forms are, as a rule, colorless, because of the absence of chromatophores. A few forms contain haematochrome. The method of nutrition is holophytic, holozoic, saprozoic, or mixotrophic. In holophytic forms, anabolic products are starch, oil, or fats.

Asexual reproduction is by binary or multiple fission or budding in either the active or the resting stage and differs among different groups. Encystment is of common occurrence. In some forms the cyst wall is formed within the test. The cysts remain alive for many years; for example, Ceratium cysts were found to retain their vitality in one instance for six and one-half years. Conjugation and sexual fusion have been reported in certain forms, but definite knowledge on sexual reproduction awaits further investigation.

The dinoflagellates are abundant in the plankton of the sea and play an important part in the economy of marine life as a whole. A number of parasitic forms are also known. Their hosts include various diatoms, copepods and several pelagic animals.

Some dinoflagellates inhabiting various seas multiply suddenly in enormous numbers within certain areas, and bring about distinct discolorations of water, often referred to as "red tide" or "red water." Occasionally the red water causes the death of a large number of fishes and of various invertebrates. According to Galtsoff (1948, 1949), the red water which appeared on the west coast of Florida in 1946 and 1947, was due to the presence of an enormous number of Gymnodinium breve and this dinoflagellate seemed in some manner to have been closely correlated with the fatal effect on animals entering the discolored water. Ketchum and Keen (1948) found the total phosphorus content of the water containing dense Gymnodinium populations to be 2.5 to 10 times the maximum expected in
the sea, and the substance associated with Gymnodinium and other dinoflagellates causes nose and throat irritations in man. Woodcock (1948) observed that similar irritations can be produced by breathing air artificially laden with small drops of the red water containing $50 \times 10^6$ dinoflagellates per liter. The irritant substance passed through a fine bacterial filter, and was found to be very stable, remaining active in stored red water for several weeks. Distribution and taxonomy (Kofoid, 1906, 1907, 1909, 1931; Kofoid and Swezy, 1921; Prescott, 1928; Eddy, 1930; Playfair, 1919; Wailes, 1934; Thompson, 1947, 1950; Balech, 1944, 1949, 1951; Rampi, 1950; Chatton, 1952); locomotion (Peters, 1929).

The Dinoflagellata are subdivided into three major groups:

Bivalve shell without furrows. . . . . . . . . Suborder 1 Prorocentrinae
Naked or with shell showing furrows. . . Suborder 2 Peridiniinea (p. 313)
Naked; without furrows; no transverse flagellum. . . . . . . . . . . . . . . . . . . . . Suborder 3 Cystoflagellata (p. 329)

Suborder 1 Prorocentrinae Poche

Test bivalve; without any groove; with yellow chromatophores; 2 flagella anterior, one directed anteriorly, the other vibrates in a circle; fresh or salt water.

Family Prorocentridae Kofoid

Genus Prorocentrum Ehrenberg. Elongate oval; anterior end bluntly pointed, with a spinous projection at pole; chromatophores small, yellowish brown; salt water. Species (Schiller, 1918, 1928).

$P. \text{micans}$ E. (Fig. 126, a). 36–52\(\mu\) long; a cause of “red water.”

$P. \text{triangulatum}$ Martin. Triangular with rounded posterior end; shell-valves flattened; one valve with a delicate tooth; surface covered with minute pores; margin striated; chromatophores yellow-brown, irregular, broken up in small masses; 17–22\(\mu\). Martin (1929) found it extremely abundant in brackish water in New Jersey.

Genus Exuviaella Cienkowski. Subspherical or oval; no anterior projection, except 2 flagella; 2 lateral chromatophores, large, brown, each with a pyrenoid and a starch body; nucleus posterior; salt and fresh water. Several species (Schiller, 1918, 1928).

$E. \text{marina}$ C. (Fig. 126, b, c). 36–50\(\mu\) long.

$E. \text{apora}$ Schiller. Compressed, oval; striae on margin of valves; chromatophores numerous yellow-brown, irregular in form; 30–32\(\mu\) by 21–26\(\mu\) (Schiller); 17–22\(\mu\) by 14–19\(\mu\) (Lebour; Martin); common in brackish water, New Jersey.

$E. \text{compressa}$ (Stein). Flattened ellipsoid test; anterior end with a
depression through which two flagella emerge; two chromatophores pale or deep green, each with a pyrenoid; nucleus posterior; no stigma; 22–26 μ by 15–18 μ by 11–12 μ; fresh and salt water (Thompson, 1950).

Suborder 2 Peridiniinea Poche

Typical dinoflagellates with one to many transverse annuli and a sulcus; 2 flagella, one of which undergoes a typical undulating movement, while the other usually directed posteriorly. According to Kofoid and Swezy, this suborder is divided into two tribes.

Body naked or covered by a thin shell ........ Tribe 1 Gymnodinioidae
Body covered by a thick shell ............. Tribe 2 Peridinioidae (p. 324)

Tribe 1 Gymnodinioidae Poche

Naked or covered by a single piece cellulose membrane with annulus and sulcus, and 2 flagella; chromatophores abundant, yellow or greenish platelets or bands; stigma sometimes present; asexual reproduction, binary or multiple division; holophytic, holozoic, or
saprozoic; the majority are deep-sea forms; a few coastal or fresh water forms also occur.

With a cellulose membrane..........................Family 1 Cystodiniidae
Without shell
   Furrows rudimentary..............................Family 2 Pronoictilucidae
   Annulus and sulcus distinct
      Solitary
         With ocellus..............................Family 3 Pouchetiidae (p. 316)
         Without ocellus
           With tentacles.............................Family 4 Noctilucidae (p. 316)
           Without tentacles
              Free-living............................Family 5 Gymnodiniidae (p. 318)
              Parasitic.............................Family 6 Blastodiniidae (p. 321)
              Permanently colonial.....................Family 7 Polykrikidae (p. 324)

Family 1 Cystodiniidae Kofoi and Swezy

Genus Cystodinium Klebs. In swimming phase, oval, with extremely delicate envelope; annulus somewhat acyclic; cyst-membrane drawn out into 2 horns. Species (Pascher, 1928; Thompson, 1949).

   C. steini K. (Fig. 126, d, e). Stigma beneath sulcus; chromatophores brown; swarmer about 45μ long; freshwater ponds.

Genus Glenodinium Ehrenberg. (Glenodiniopsis, Stasizecella Woloszynska). Spherical; ellipsoidal or reniform in end-view; annulus a circle; several discoidal, yellow to brown chromatophores; horseshoe- or rod-shaped stigma in some; often with gelatinous envelope; fresh water. Many species (Thompson, 1950).

   G. cinctum E. (Fig. 126, f). Spherical to ovoid; annulus equatorial; stigma horseshoe-shaped; 43μ by 40μ. Morphology and reproduction (Lindemman, 1929).

Family 2 Pronoictilucidae Lebour

Genus Pronoictiluca Fabre-Domergue. Body with an antero-ventral tentacle and sulcus; annulus poorly marked; salt water.

   P. tentaculatum (Kofoi and Swezy) (Fig. 127, a). About 54μ long; off California coast.

Genus Oxyrrhis Dujardin. Subovoidal, asymmetrical posteriorly; annulus incomplete; salt water.

   O. marina D. (Fig. 127, b). 10–37μ long. Division (Dunkerly, 1921; Hall, 1925).
Fig. 127. a, Pronoctiluca tentaculatum, ×730 (Kofoid and Swezy); b, Oxyrrhis marina, ×840 (Senn); c. Pouchetia fusus, ×340 (Schütt); d, P. maxima, ×330 (Kofoid and Swezy); e, Protopsis ochrea, ×340 (Wright); f, Nematodinium partitum, ×560 (Kofoid and Swezy); g, Proterothropsis crassicaudata, ×740 (Kofoid and Swezy); h, Erythropsis cornuta, ×340 (Kofoid and Swezy); i, j, Noctiluca scintillans (i, side view; j, budding process), ×140 (Robin).
Family 3 Pouchetiidae Kofoid and Swezy

Ocellus consists of lens and melanosome (pigment mass); sulcus and annulus somewhat twisted; pusules usually present; cytoplasm colored; salt water (pelagic).

Genus Pouchetia Schütt. Nucleus anterior to ocellus; ocellus with red or black pigment mass with a red, brown, yellow, or colorless central core; lens hyaline; body surface usually smooth; holozoic; encystment common; salt water. Many species (Schiller, 1928a).

P. fusus S. (Fig. 127, c). About 94/ by 41/x; ocellus 27m long.

P. maxima Kofoid and Swezy (Fig. 127, d). 145/ by 92/*; ocellus 20/*; off California coast.

Genus Protopsis Kofoid and Swezy. Annulus and sulcus similar to those of Gymnodinium or Gyrodinium; with a simple or compound ocellus; no tentacles; body not twisted; salt water. A few species.

P. ochrea (Wright) (Fig. 127, e). 55/* by 45/z; ocellus 22/*; Nova Scotia.

Genus Nematodinium Kofoid and Swezy. With nematoceysts; girdle more than 1 turn; ocellus distributed or concentrated, posterior; holozoic; salt water.

N. partitum K. and S. (Fig. 127, f). 91/* long; off California coast.

Genus Proterythropsis Kofoid and Swezy. Annulus median; ocellus posterior; a stout rudimentary tentacle; salt water. One species.

P. crassicaudata K. and S. (Fig. 127, g). 70/* long; off California.

Genus Erythropsis Hertwig. Epicone flattened, less than 1/4 hypocone; ocellus very large, composed of one or several hyaline lenses attached to or imbedded in a red, brownish or black pigment body with a red, brown or yellow core, located at left of sulcus; sulcus expands posteriorly into ventro-posterior tentacle; salt water. Several species.

E. cornuta (Schütt) (Fig. 127, h). 104/* long; off California coast (Kofoid and Swezy).

Family 4 Noctilucidae Kent

Contractile tentacle arises from sulcal area and extends posteriorly; a flagellum; this group has formerly been included in the Cystoflagellata; studies by recent investigators, particularly by Kofoid, show its affinity with the present suborder; holozoic; salt water.

Genus Noctiluca Suriray. Spherical, bilaterally symmetrical; peristome marks the median line of body; cytostome at the bottom of peristome; with a conspicuous tentacle and a short flagellum; cytoplasm greatly vacuolated, and cytoplasmic strands connect the central mass with periphery; specific gravity is less than that of sea wa-
ter, due to the presence of an osmotically active substance with a lower specific gravity than sodium chloride, which appears to be ammonium chloride (Goethard and Heinsius); certain granules are luminescent (Fig. 128); cytoplasm colorless or blue-green; sometimes tinged with yellow coloration in center; swarmers formed by budding, and each possesses one flagellum, annulus, and tentale; widely distributed in salt water; holozoic. One species.

_N. scintillans_ (Macartney) (_N. miliaris_ S.) (Figs. 127, _i, j_; 128). Usually 500–1000µ in diameter, with extremes of 200µ and 3 mm. Gross (1934) observed that complete fusion of two swarmers (isogametes) results in cyst formation from which trophozoites develop. Acid content of the body fluid is said to be about pH 3. Nuclear di-

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**Fig. 128. Noctiluca scintillans,** as seen under darkfield microscope (Pratje). _a,_ an active individual; _b,_ a so-called “resting stage,” with fat droplets in the central cytoplasm, prior to either division or swarmer formation; _c,_ _d,_ appearance of luminescent individuals (F, fat-droplets; K, nucleus; P, peristome; T, tentacle; V, food body; Z, central protoplasm).
vision (Calkins, 1898); morphology and physiology (Goor, 1918; Kofoid, 1920; Pratje, 1921); feeding (Hofker, 1930); luminescence (Harvey, 1952).

Genus Pavillardia Kofoid and Swezy. Annulus and sulcus similar to those of Gymnodinium; longitudinal flagellum absent; stout finger-like mobile tentacle directed posteriorly; salt water. One species.

*P. tentaculifera* K. and S. 58μ by 27μ; pale yellow; off California.

Family 5 Gymnodiniidae Kofoid

Naked forms with simple but distinct 1/2–4 turns of annulus; with or without chromatophores; fresh or salt water.

Genus Gymnodinium Stein. Pellicle delicate; subcircular; bilaterally symmetrical; numerous discoid chromatophores varicolored (yellow to deep brown, green, or blue) or sometimes absent; stigma present in few; many with mucilaginous envelope; salt, brackish, or fresh water. Numerous species (Schiller, 1928a); cultivation and development (Lindemann, 1929).

*G. aeruginosum* S. (Fig. 129, a). Green chromatophores; 20–32μ by 13–25μ (Thompson, 1950); ponds and lakes.

*G. rotundatum* Klebs (Fig. 129, b). 32–35μ by 22–25μ; fresh water.

*G. palustre* Schilling (Fig. 129, c). 45μ by 38μ; fresh water.

*G. agile* Kofoid and Swezy (Fig. 129, d). About 28μ long; along sandy beaches.

Genus Hemidinium Stein. Asymmetrical; oval; annulus about half a turn, only on left half. One species.

*H. nasutum* S. (Fig. 129, e). Sulcus posterior; chromatophores yellow to brown; with a reddish brown oil drop; nucleus posterior; transverse fission; 24–28μ by 16–17μ; fresh water.

Genus Amphidinium Claparède and Lachmann. Form variable; epicone small; annulus anterior; sulcus straight on hypocone or also on part of epicone; with or without chromatophores; mainly holophytic, some holozoic; coastal or fresh water. Numerous species (Schiller, 1928a).

*A. lacustre* Stein (Fig. 129, f). 30μ by 18μ; in fresh and salt (?) water.

*A. scissum* Kofoid and Swezy (Fig. 129, g). 50–60μ long; along sandy beaches.

*A. fusiforme* Martin. Fusiform, twice as long as broad: circular in cross-section; epicone rounded conical; annulus anterior; hypocone 2–2.5 times as long as epicone; sulcus obscure; body filled with
Fig. 129. a, Gymnodinium aeruginosum, ×500 (Schilling); b, G. rotundatum, ×360 (Klebs); c, G. palustre, ×360 (Schilling); d, G. agile, ×740 (Kofoid and Swezy); e, Hemidinium nasutum, ×670 (Stein); f, Amphidinium lacustre, ×440 (Stein); g, A. scissum, ×880 (Kofoid and Swezy); h, Gyrodinium biconicum, ×340 (Kofoid and Swezy); i, G. hyalinum, ×670 (Kofoid and Swezy); j, Cochlodinium atromaculatum, ×340 (Kofoid and Swezy); k, Torodinium robustum, ×670 (Kofoid and Swezy); l, Massartia nieuportensis, ×670 (Conrad); m, Chilodinium cruciatum, ×900 (Conrad); n, o, Trochodinium prismaticum, ×1270 (Conrad); p, Ceratodinium asymmetricum, ×670 (Conrad).
yellowish green chromatophores except at posterior end; stigma dull orange, below girdle; nucleus ellipsoid, posterior to annulus; pellicle delicate; 17–22\(\mu\) by 8–11\(\mu\) in diameter. Martin (1929) found that it was extremely abundant in parts of Delaware Bay and gave rise to red coloration of the water (“Red water”).

Genus **Gyrodinium** Kofoid and Swezy. Annulus descending left spiral; sulcus extending from end to end; nucleus central; pusules; surface smooth or striated; chromatophores rarely present; cytoplasm colored; holozoic; salt or fresh water. Many species (Schiller, 1928a).

*G. biconicum* K. and S. (Fig. 129, h). 68\(\mu\) long; salt water; off California.

*G. hyalinum* (Schilling) (Fig. 129, i). About 24\(\mu\) long; fresh water.

Genus **Cochlodinium** Schütt. Twisted at least 1.5 turns; annulus descending left spiral; pusules; cytoplasm colorless to highly colored; chromatophores rarely present; holozoic; surface smooth or striated; salt water. Numerous species (Schiller, 1928a).

*C. atromaculatum* Kofoid and Swezy (Fig. 129, j). 183–185\(\mu\) by 72\(\mu\); longitudinal flagellum 45\(\mu\) long; off California.

Genus **Torodinium** Kofoid and Swezy. Ellongate; epicone several times longer than hypocone; annulus and hypocone form augur-shaped cone; sulcus long; nucleus greatly elongate; salt water. 2 species (Schiller, 1928).

*T. robustum* K. and S. (Fig. 129, k). 67–75\(\mu\) long; off California.

Genus **Massartia** Conrad. Cylindrical; epicone larger (9–10 times longer and 3 times wider) than hypocone; no sulcus; with or without yellowish discoid chromatophore (Thompson, 1950).

*M. nieuportensis* C. (Fig. 129, l). 28–37\(\mu\) long; brackish water.

Genus **Chilodinium** Conrad. Ellipsoid; posterior end broadly rounded, anterior end narrowed and drawn out into a digitiform process closely adhering to body; sulcus, apex to 1/5 from posterior end; annulus oblique, in anterior 1/3 (Conrad, 1926).

*C. cruciatum* C. (Fig. 129, m). 40–50\(\mu\) by 30–40\(\mu\); with trichocysts; brackish water.

Genus **Trochodinium** Conrad. Somewhat similar to *Amphidiniun*; epicone small, button-like; hypocone with 4 longitudinal rounded ridges; stigma; without chromatophores.

*T. prismaticum* C. (Fig. 129, n, o). 18–22\(\mu\) by 9–12\(\mu\); epicone 5–7\(\mu\) in diameter; brackish water (Conrad, 1926).

Genus **Ceratodinium** Conrad. Cuneiform; asymmetrical, colorless, more or less flattened; annulus complete, oblique; sulcus on half of epicone and full length of hypocone; stigma.
C. asymmetricum C. (Fig. 129, p). 68–80μ by about 10μ; brackish water (Conrad, 1926).

Family 6 Blastodiniidae Kofoid and Swezy

All parasitic in or on plants and animals; in colony forming genera, there occur trophocyte (Chatton) by which organism is attached to host and more or less numerous gonocytes (Chatton). Taxonomy (Chatton, 1920; Reichenow, 1930).

Genus Blastodinium Chatton. In the gut of copepods; spindle-shaped, arched, ends attenuated; envelope (not cellulose) often with 2 spiral rows of bristles; young forms binucleate; when present, chromatophores in yellowish brown network; swarmers similar to those of Gymnodinium; in salt water. Many species.

B. spinulosum C. (Fig. 130, a). About 235μ by 33–39μ; swarmers 5–10μ; in Palacalanus parvus, Clausocalanus arcuicornis and C. furcatus.

Genus Oodinium Chatton. Spherical or pyriform; with a short stalk; nucleus large; often with yellowish pigment; on Salpa, Annelida, Siphonophora, marine fishes, etc.

O. poucheti (Lemmermann) (Fig. 130, b, c). Fully grown individuals up to 170μ long; bright yellow ochre; mature forms become detached and free, dividing into numerous gymnodinium-like swarmers; on the tunicate. Oikopleura dioica.

O. ocellatum Brown (Fig. 131, a, b). Attached to the gill filaments of marine fish by means of cytoplasmic processes; oval in form; 12μ by 10μ to 104μ by 80μ, average 60μ by 50μ; nucleus spherical; many chromatophores and starch grains; a stigma. When grown, the organism drops off the gill and becomes enlarged to as much as 150μ in diameter. Soon the cytoplasmic processes and the broad flagellum are retracted and the aperture of shell closes by secretion of cellulose substance. The body divides up to 128 cells, which become flagellated and each divides once more. These flagellates, 12μ by 5μ, reach the gills of fish and become attached (Brown, 1931; Nigrelli, 1936).

O. limneticum Jacobs (Fig. 131, c, d). Pyriform; 12μ by 7.5μ to 20μ by 13μ; light green chromatophores variable in size and shape; no stigma; without flagella; filopodia straight or branched; the organism grows into about 60μ long in three days at 25°C.; observed maximum, 96μ by 80μ; starch becomes abundant; fission takes place in cyst; flagellate forms measure about 15μ long; ectoparasitic on the integument of freshwater fishes in aquaria (Jacobs, 1946).
Genus Apodinium Chatton. Young individuals elongate, spherical or pyriform; binucleate; adult colorless; formation of numerous swarmers in adult stage is peculiar in that lower of the 2 individuals formed at each division secretes a new envelope, and delays its

Fig. 130. a, Blastodinium spinulosum, ×240 (Chatton); b, c, Oo- 
dinium poucheti (c, a swarmer) (Chatton); d, e, Apodinium mycetoides (d, swarmer-formation, ×450; e, a younger stage, ×640) (Chatton); f, Chytriodinium parasiticum in a copepod egg (Dogiel); g, Trypanodinium oricola, ×1070 (Chatton); h, Duboscqella tintinnicola (Duboscq and Collin); i, j, Haplozoon clymenellae (i, mature colony, ×300; j, a swarmer, ×1340) (Shumway); k, Syndinium turbo, ×1340 (Chatton); l, Paradi- 
nium poucheti, ×800 (Chatton); m, Ellobiopsis chattoni on Calanus fin- 
marchicus (Caullery); n, Paraellobiopsis coutieri (Collin).
further division until the upper one has divided for the second time, leaving several open cups; on tunicates.

A. *mycetoides* C. (Fig. 130, d, e). On gill-slits of *Fritillaria pel- lucida*.

Genus *Chytriodinium* Chatton. In eggs of planktonic copepods; young individuals grow at the expense of host egg and when fully formed, body divides into many parts, each producing 4 swimmers. Several species.

*C. parasiticum* (Dogiel) (Fig. 130, f). In copepod eggs; Naples.

Genus *Trypanodinium* Chatton. In copepod eggs; swarmer-stage only known.

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Fig. 131. *Oo dinium ocellatum*, recently detached from host gill; b, a free living flagellate form, ×760 (Nigrelli); c, d, *O. limneticum*, ×800 (Jacobs).

*T. ovicola* C. (Fig. 130, g). Swarmers biflagellate; about 15μ long.

Genus *Duboscqella* Chatton. Rounded cell with a large nucleus; parasitic in Tintinnidae. One species.

*D. tintinnicola* (Lohmann) (Fig. 130, h). Intracellular stage oval, about 100μ in diameter with a large nucleus; swarmers biflagellate.

Genus *Haplozoon* Dogiel. In gut of polychaetes; mature forms composed of variable number of cells arranged in line or in pyramid; salt water. Many species.

*H. clymenellae* (Calkins) (*Microtaeniella clymenellae* C.) (Fig. 130, i, j). In the intestine of *Clymenella torquata*; colonial forms consist of 250 or more cells; Woods Hole (Shumway, 1924).

Genus *Syndinium* Chatton. In gut and body cavity of marine copepods; multinucleate round cysts in gut considered as young
forms; multinucleate body in host body cavity with numerous needle-like inclusions.

*S. turbo* C. (Fig. 130, k). In *Paracalanus parvus*, *Corycaeus venus*, *Calanus finmarchicus*; swarmers about 15μ long.

Genus **Paradinium** Chatton. In body-cavity of copepods; multinucleate body without inclusions; swarmers formed outside the host body.

*P. poucheti* C. (Fig. 130, l). In the copepod, *Acartia clausi*; swarmers about 25μ long, amoeboid.

Genus **Ellobiopsis** Caullery. Pyriform; with stalk; often a septum near stalked end; attached to anterior appendages of marine copepods.

*E. chattoni* C. (Fig. 130, m). Up to 700μ long; on antennae and oral appendages of *Calanus finmarchicus*, *Pseudocalanus elongatus* and *Acartia clausi*. Development (Steuer, 1928).

Genus **Paraellobiopsis** Collin. Young forms stalkless; spherical; mature individuals in chain-form; on Malacostraca.

*P. coutieri* C. (Fig. 130, n). On appendages of *Nebalia bipes*.

**Family 7 Polykrikidae** Kofoid and Swezey

Two, 4, 8, or 16 individuals permanently joined; individuals similar to *Gymnodinium*; sulcus however extending entire body length; with nematocysts (Fig. 132, b); greenish to pink; nuclei about 1/2 the number of individuals; holozoic; salt water. Nematocysts (Hovasse, 1951).

Genus **Polykrikos** Bötschli. With the above-mentioned characters; salt or brackish water. Species (Schiller, 1928).

*P. kofoidi* (Chatton) (Fig. 132, a, b). Greenish grey to rose; composed of 2, 4, 8, or 16 individuals; with nematocysts; each nematocyst possesses presumably a hollow thread, and discharges under suitable stimulation its content; a binucleate colony composed of 4 individuals about 110μ long; off California.

*P. barnegatensis* Martin. Ovate, nearly circular in cross-section, slightly concave ventrally; composed of 2 individuals; constriction slight; beaded nucleus in center; annuli descending left spiral, displaced twice their width; sulcus ends near anterior end; cytoplasm colorless, with numerous oval, yellow-brown chromatophores; nematocysts absent; 46μ by 31.5μ; in brackish water of Barnegat Bay.

**Tribe 2 Peridinioidae** Poche

The shell composed of epitheca, annulus and hypotheca, which may be divided into numerous plates; body form variable.
With annulus and sulcus
Shell composed of plates; but no suture... Family 1 Peridiniidae (p. 326)
Breast plate divided by sagittal suture. Family 2 Dinophysidae (p. 328)
Without annulus or sulcus.............. Family 3 Phytodiniidae (p. 329)

**Fig. 132.** a, b, *Polykrikos kofoidi* (a, colony of four individuals, ×340; b, a nematocyst, ×1040) (Kofoid and Swezy); c, *Peridinium tabulatum*, ×460 (Schilling); d, *P. divergens*, ×340 (Calkins); e, *Ceratium hirundinella*, ×540 (Stein); f, *C. longipes*, ×100 (Wailes); g, *C. tripos*, ×140 (Wailes); h, *C. fusus*, ×100 (Wailes); i, *Heterodinium scrippsi*, ×570 (Kofoid and Adamson).
Family 1 Peridiniidae Kent

Shell composed of numerous plates; annulus usually at equator, covered by a plate known as cingulum; variously sculptured and finely perforated plates vary in shape and number among different species; in many species certain plates drawn out into various processes, varying greatly in different seasons and localities even among one and the same species; these processes seem to retard descending movement of organisms from upper to lower level in water when flagellar activity ceases; chromatophores numerous small platelets, yellow or green; some deep-sea forms without chromatophores; chain formation in some forms; mostly surface and pelagic inhabitants in fresh or salt water.

Genus Peridinium Ehrenberg. Subspherical to ovoid; reniform in cross-section; annulus slightly spiral with projecting rims; hypotheca often with short horns and epitheca drawn out; colorless, green, or brown; stigma usually present; cysts spherical; salt or fresh water. Numerous species. Species and variation (Böhm, 1933; Diwald, 1939); Chromatophore and pyrenoid (Geitler, 1926).

P. tabulatum Claparède and Lachmann (Fig. 132, c). 48μ by 44μ; fresh water. P. divergens (E.) (Fig. 132, d). About 45μ in diameter; yellowish, salt water.

Genus Ceratium Schrank. Body flattened; with one anterior and 1-4 posterior horn-like processes; often large; chromatophores yellow, brown, or greenish; color variation conspicuous; fission is said to take place at night and in the early morning; fresh or salt water. Numerous species; specific identification is difficult due to a great variation (p. 223). Biology and morphology (Entz, 1927); encystment (Entz, 1925).

C. hirundinella (Müller) (Figs. 94; 132, e). 1 apical and 2-3 antapical horns; seasonal and geographical variations (p. 223); chain-formation frequent; 95-700μ long; fresh and salt water. Numerous varieties. Reproduction (Entz, 1921, 1931; Hall, 1925a; Borgert, 1935); holozoic nutrition (Hofeneder, 1930).

C. longipes (Bailey) (Fig. 132, f). About 210μ by 51-57μ; salt water.

C. tripus (Müller) (Fig. 132, g). About 225μ by 75μ; salt water. Wailes (1928) observed var. atlantica in British Columbia; Martin (1929) in Barnegat Inlet, New Jersey. Nuclear division (Schneider, 1924).

C. fusus (Ehrenberg) (Fig. 132, h). 300-600μ by 15-30μ; salt water; widely distributed; British Columbia (Wailes), New Jersey (Martin), etc.
Genus *Heterodinium* Kofoid. Flattened or spheroidal; 2 large antapical horns; annulus submedian; with post-cingular ridge; sulcus short, narrow; shell hyaline, reticulate, porulate; salt water. Numerous species.

*H. scrippsi* K. (Fig. 132, i). 130–155\(\mu\) long; Pacific and Atlantic (tropical).

Genus *Dolichodinium* Kofoid and Adamson. Subconical, elongate; without apical or antapical horns; sulcus only 1/2 the length of hypotheca; plate porulate; salt water.

*D. lineatum* (Kofoid and Michener) (Fig. 133, a). 58\(\mu\) long; eastern tropical Pacific.

Genus *Goniodoma* Stein. Polyhedral with a deep annulus; epitheca and hypotheca slightly unequal in size, composed of regularly arranged armored plates; chromatophores small brown platelets; fresh or salt water.

*G. acuminata* (Ehrenberg) (Fig. 133, b). About 50\(\mu\) long; salt water.

Genus *Gonyaulax* Diesing. Spherical, polyhedral, fusiform, elongated with stout apical and antapical prolongations, or dorso-ventrally flattened; apex never sharply attenuated; annulus equatorial; sulcus from apex to antapex, broadened posteriorly; plates 1–6 apical, 0–3 anterior intercalaries, 6 precingulars, 6 annular plates, 6 postcingulars, 1 posterior intercalary and 1 antapical; porulate; chromatophores yellow to dark brown, often dense; without stigma; fresh, brackish or salt water. Numerous species (Kofoid, 1911; Whedon and Kofoid, 1936).

*G. polyedra* Stein (Fig. 133, c). Angular, polyhedral; ridges along sutures, annulus displaced 1–2 annulus widths, regularly pitted; salt water. “Very abundant in the San Diego region in the summer plankton, July–September, when it causes local outbreaks of ‘red water,’ which extend along the coast of southern and lower California” (Kofoid, 1911; Allen, 1946). The organisms occurred also in abundance (85 per cent of plankton) in pools of sea water off the beach of Areia Branca, Portugal, and caused “red water” during the day and an extreme luminescence when agitated at night (Santos-Pinto, 1949).

*G. apiculata* (Penard) (Fig. 133, d). Ovate, chromatophores yellowish brown; 30–60\(\mu\) long; fresh water.

Genus *Spiraulax* Kofoid. Biconical; apices pointed; sulcus not reaching apex; no ventral pore; surface heavily pitted; salt water.

*S. jolliffi* (Murray and Whitting) (Fig. 133, e). 132\(\mu\) by 92\(\mu\); California (Kofoid, 1911a).

**Family 2 Dinophysidae Kofoid**

Genus *Dinophysis* Ehrenberg. Highly compressed; annulus widened, funnel-like, surrounding small epitheca; chromatophores yellow; salt water. Several species (Schiller, 1928). Morphology and taxonomy (Tai and Skogsberg, 1934).

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Fig. 133. a, *Dolichodinium lineatum*, ×670 (Kofoid and Adamson); b, *Coniodoma acuminata*, ×340 (Stein); c, *Gonyaulax polyedra*, ×670 (Kofoid); d, *G. apiculata*, ×670 (Lindemann); e, *Spiraulax jolliffei*, right side of theca, ×340 (Kofoid); f, *Dinophysis acuta*, ×580 (Schütt); g, h, *Oxyphysis oxytoxoides*, ×780 (Kofoid); i, *Phytodinium simplex*, ×340 (Klebs); j, k, *Dissodinium lunula*: j, primary cyst (Dogiel); k, secondary cyst with 4 swarmers (Wailes), ×220.
D. acuta E. (Fig. 133, f). Oval; attenuated posteriorly; 54-94μ long; widely distributed; British Columbia (Wailes).

Genus Oxyphysis Kofoid. Epitheca developed; sulcus short; sulcal lists feebly developed; sagittal suture conspicuous; annulus impressed; salt water (Kofoid, 1926).

O. oxytoxoides K. (Fig. 133, g, h). 63-68μ by 15μ; off Alaska.

Family 3 Phytodiniidae Klebs

Genus Phytodinium Klebs. Spherical or ellipsoidal; without furrows; chromatophores discoidal, yellowish brown.

P. simplex K. (Fig. 133, i). Spherical or oval; 42-50μ by 30-45μ fresh water.

Genus Dissodinium Klebs (Pyrocystis Paulsen). Primary cyst, spherical, uninucleate; contents divide into 8-16 crescentic secondary cysts which become set free; in them are formed 2, 4, 6, or 8 Gymnodinium-like swarmers; salt water.

D. lunula (Schütz) (Fig. 133, j, k). Primary cysts 80-155μ in diameter; secondary cysts 104-130μ long; swarmers 22μ long; widely distributed; British Columbia (Wailes).

Suborder 3 Cystoflagellata Haeckel

Since Noctiluca which had for many years been placed in this suborder, has been removed, according to Kofoid, to the second suborder, the Cystoflagellata becomes a highly ill-defined group and includes two peculiar marine forms: Leptodiscus medusoides Hertwig (Fig. 134, a), and Craspedotella pileolus Kofoid (Fig. 134, b), both of which are medusoid in general body form.

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Chapter 13
Subclass 2 Zoomastigina Doflein

The Zoomastigina lack chromatophores and their body organizations vary greatly from a simple to a very complex type. The majority possess a single nucleus which is, as a rule, vesicular in structure. Characteristic organelles such as parabasal body, axostyle, etc., are present in numerous forms and myonemes are found in some species. Nutrition is holozoic or saprozoic (parasitic). Asexual reproduction is by longitudinal fission; sexual reproduction is unknown. Encystment occurs commonly. The Zoomastigina are free-living or parasitic in various animals.

With pseudopodia besides flagella.............Order 1 Rhizomastigina
With flagella only
   With 1–2 flagella.........................Order 2 Protomonadina (p. 339)
   With 3–8 flagella.........................Order 3 Polymastigina (p. 369)
   With more than 8 flagella.............Order 4 Hypermastigina (p. 404)

Order 1 Rhizomastigina Bütschli

A number of borderline forms between the Sarcodina and the Mastigophora are placed here. Flagella vary in number from one to several and pseudopods also vary greatly in number and in appearance.

With many flagella..........................Family 1 Multiciliidae
With 1–3 rarely 4 flagella..................Family 2 Mastigamoebidae

Family 1 Multiciliidae Poche

Genus Multicilia Cienkowski. Generally spheroidal, but amoeboid; with 40–50 flagella, long and evenly distributed; one or more nuclei; holozoic; food obtained by means of pseudopodia; multiplication by fission; fresh or salt water.

M. marina C. (Fig. 135, a). 20–30μ in diameter; uninucleate; salt water.

M. lacustris Lauterborn (Fig. 135, b). Multinucleate; 30–40μ in diameter; fresh water.

Family 2 Mastigamoebidae

With 1–3 or rarely 4 flagella and axopodia or lobopodia; uninucleate; flagellum arises from a basal granule which is connected with the nucleus by a rhizoplast; binary fission in both trophic and encysted stages; sexual reproduction has been reported in one species; holozoic or saprozoic; the majority are free-living, though a few parasitic.
Genus *Mastigamoeba* Schulze (*Mastigina* Frenzel). Monomastigote, uninucleate, with finger-like pseudopodia; flagellum long and connected with nucleus; fresh water, soil or endocommensal. Species (Klug, 1936).

*M. aspera* S. (Fig. 135, c). Subspherical or oval; during locomotion elongate and narrowed anteriorly, while posterior end rounded or lobed; numerous pseudopods slender, straight; nucleus near flagellate end; 2 contractile vacuoles; 150–200μ by about 50μ; in ooze of pond.

*M. longifilum* Stokes (Fig. 135, d). Elongate, transparent; flagellum twice body length; pseudopods few, short; contractile vacuole anterior; body 28μ long when extended, contracted about 10μ; stagnant water.
M. setosa (Goldschmidt) (Fig. 135, e). Up to 140μ long.

M. hylae (Frenzel) (Fig. 136, a). In the hind-gut of the tadpoles of frogs and toads; 80–135μ by 21–31μ; flagellum about 10μ long (Becker, 1925). Development (Ivanic, 1936).

Genus Mastigella Frenzel. Flagellum apparently not connected with nucleus; pseudopods numerous, digitate; body form changes actively and continuously; contractile vacuole.

M. vitrea Goldschmidt (Fig. 135, f). 150μ long; sexual reproduction (Goldschmidt).

Genus Actinomonas Kent. Generally spheroidal, with a single flagellum and radiating pseudopods; ordinarily attached to foreign object with a cytoplasmic process, but swims freely by withdrawing it; nucleus central; several contractile vacuoles; holozoic.

A. mirabilis K. (Fig. 136, b). Numerous simple filopodia; about 10μ in diameter; flagellum 20μ long; fresh water.

Genus Dimorpha Gruber. Ovoid or subspherical; with 2 flagella and radiating axopodia, all arising from an eccentric centriole; nucleus eccentric; pseudopods sometimes withdrawn; fresh water. Species (Pascher, 1925).

D. mutans G. (Fig. 136, c). 15–20μ in diameter; flagella about 20–30μ long.

Genus Tetradimorpha Hsiung. Spherical with radiating axopodia; four flagella originate in a slightly depressed area; nucleus central. When disturbed, all axopodia turn away from the flagellated pole and are withdrawn into body, and the organism undergoes swimming movement; freshwater ponds.

T. radiata H. (Fig. 136, d, e). Body 27–38μ in diameter; axopodia 27–65μ long; flagella 38–57μ long (Hsiung, 1927).

Genus Pteridomonas Penard. Small, heart-shaped; usually attached with a long cytoplasmic process; from opposite pole there arises a single flagellum, around which occurs a ring of extremely fine filopods; nucleus central; a contractile vacuole; holozoic; fresh water.

P. pulex P. (Fig. 136, f). 6–12μ broad.

Genus Histomonas Tyzzer. Actively amoeboid; mostly rounded, sometimes elongate; a single nucleus; an extremely fine flagellum arises from a blepharoplast, located close to nucleus; axostyle (?) sometimes present; in domestic fowls. One species.

H. meleagridis (Smith) (Amoeba meleagridis S.) (Fig. 137). Actively amoeboid organism; usually rounded; 8–21μ (average 10–14μ) in the largest diameter; nucleus circular or pyriform with usually a large endosome; a fine flagellum; food vacuoles contain bacteria, starch grains and erythrocytes; binary fission; during division flagel-
lum is discarded; cysts unobserved; in young turkeys, chicks, grouse, and quail. Bayon and Bishop (1937) successfully cultured the organism from hen’s liver. Morphology of the cultured forms (Bishop, 1938).

This organism is the cause of enterohepatitis known as “blackhead,” an infectious disease, in young turkeys and also in other fowls, in which it is often fatal. Smith (1895) discovered the organism and considered it an amoeba (1910). It invades and destroys the mucosa of the intestine and caeca as well as the liver tissues. Tropho-

Fig. 136. a, Mastigamoeba hylae, X690 (Becker); b, Actinomonas mirabilis, X1140 (Griessmann); c, Dimorpha mutans, X940 (Bloehmann); d, e, Tetradmorpha radiata (Hsiung) (d, a typical specimen, X430; e, swimming individual, X300); f, Pteridomonas pulex, X540 (Penard); g, Rhizomastix gracilis, X1340 (Mackinnon).
zoites voided in faeces by infected birds may become the source of new infection when taken in by young birds with drink or food. Tyzzer (1920) found the organism to possess a flagellate stage and established the genus *Histomonas* for it. Tyzzer and Fabyan (1922) and Tyzzer (1934) demonstrated that the organism is transmissible from bird to bird in the eggs of the nematode *Heterakis gallinae*, which method appears to be a convenient and reliable one for producing *Histomonas* infection in turkeys (McKay and Morehouse, 1948). Desowitz (1950) noticed in a *Heterakis* two enlarged gut cells filled with amoebulae which he suggested might be a stage of this protozoan. Niimi (1937) reported that the organism enters through the mouth of the nematode and invades its eggs. Dobell (1940) points out the similarity between this flagellate and *Dientamoeba fragilis* (p. 462). Wenrich (1943) made a comparative study of forms found in

Fig. 137. *Histomonas meleagridis*. a–d, from host animals (Wenrich); e–h, from cultures (Bishop). a, b, organisms in caecum of chicken (in a Tyzzer slide); c, an individual from pheasant showing "ingestion tube" with a bacterial rod; d, a large individual from the same source, all \(\times 1765\); e, an amoeboid form; f, a rounded form with axostyle (?); g, h, stages in nuclear division, \(\times 2200\).
the caecal smears of wild ring-neck pheasants and of chicks. The organisms measured 5–30μ in diameter and possessed 1–4 flagella, though often there were no flagella.

Genus *Rhizomastix* Alexeieff. Body amoeboid; nucleus central; blepharoplast located between nucleus and posterior end; a long fiber runs from it to anterior end and continues into the flagellum; without contractile vacuole; division in spherical cyst.

*R. gracilis* A. (Fig. 136, g). 8–14μ long; flagellum 20μ long; in intestine of axolotles and tipulid larvae.

References


——— (1920) The flagellate character and reclassification of the parasite producing “blackhead” in turkey, etc. J. Parasit., 6:124.


CHAPTER 14

Order 2 Protomonadina Blochmann

THE protomonads possess one or two flagella and are composed of a heterogeneous lot of Protozoa, mostly parasitic, whose affinities to one another are very incompletely known. The body is in many cases plastic, having no definite pellicle, and in some forms amoeboid. The method of nutrition is holozoic, or saprozoic (parasitic). Reproduction is, as a rule, by longitudinal fission, although budding or multiple fission has also been known to occur, while sexual reproduction, though reported in some forms, has not been confirmed.

With 1 flagellum
  With collar
    Collar enclosed in jelly..................Family 1 Phalansteriidae
    Collar not enclosed in jelly
      Without lorica..........................Family 2 Codosigidae
      With lorica...........................Family 3 Bicosoecidae (p. 341)
  Without collar
    Free-living.............................Family 4 Oikomondaidae (p. 343)
    Parasitic..............................Family 5 Trypanosomatidae (p. 344)

With 2 flagella
  With undulating membrane..............Family 6 Cryptobiidae (p. 357)
  Without undulating membrane
    Flagella equally long..................Family 7 Amphimonadidae (p. 358)
    Flagella unequally long
      No trailing flagellum.................Family 8 Monadidae (p. 360)
      One flagellum trailing...............Family 9 Bodonidae (p. 362)

Family 1 Phalansteriidae Kent

Genus Phalansterium Cienkowski. Small, ovoid; one flagellum and a small collar; numerous individuals are embedded in gelatinous substance, with protruding flagella; fresh water.

P. digitatum Stein (Fig. 138, a). Cells about 17μ long; oval; colony dendritic; fresh water among vegetation.

Family 2 Codosigidae Kent

Small flagellates; delicate collar surrounds flagellum; ordinarily sedentary forms; if temporarily free, organisms swim with flagellum directed backward; holozoic on bacteria or saprozoic; often colonial; free-living in fresh water. Feeding process (Lapage, 1925).

Genus Codosiga Kent (Codonocladium Stein; Astrosiga Kent). Individuals clustered at end of a simple or branching stalk; fresh water.
C. utriculus Stokes (Fig. 138, b). About 11μ long; attached to freshwater plants.

C. disjuncta (Fromentel) (Fig. 138, c). In stellate clusters; cells about 15μ long; fresh water.

Fig. 138. a, Phalansterium digitatum, ×540 (Stein); b, Codosiga utriculus, ×1340 (Stokes); c, C. disjuncta, ×400 (Kent); d, Monosiga ovata, ×800 (Kent); e, M. robusta, ×770 (Stokes); f, Desmarella moniliformis, ×800 (Kent); g, Protospongia haeckeli, ×400 (Lemmermann); h, an individual of Sphaeroeca volvox, ×890 (Lemmermann); i, Diplosiga francei, ×400 (Lemmermann); j, D. socialis, ×670 (Francé).

Genus Monosiga Kent. Solitary; with or without stalk; occasionally with short pseudopodia; attached to freshwater plants. Several species.

M. ovata K. (Fig. 138, d). 5–15μ long; with a short stalk.

M. robusta Stokes (Fig. 138, e). 13μ long; stalk very long.

Genus Desmarella Kent. Cells united laterally to one another; fresh water.

D. moniliformis K. (Fig. 138, f). Cells about 6μ long; cluster composed of 2–12 individuals; standing fresh water.

D. irregularis Stokes. Cluster of individuals irregularly branching, composed of more than 50 cells; cells 7–11μ long; pond water.
Genus *Proterospongia* Kent. Stalkless individuals embedded irregularly in a jelly mass, collars protruding; fresh water.

*P. haeckeli* K. (Fig. 138, g). Body oval; 8μ long; flagellum 24–32μ long; 6–60 cells in a colony.

Genus *Sphaeroeca* Lauterborn. Somewhat similar to the last genus; but individuals with stalks and radiating; gelatinous mass spheroidal; fresh water.

*S. volvox* L. (Fig. 138, h). Cells ovoid, 8–12μ long; stalk about twice as long; flagellum long; contractile vacuole posterior; colony 82–200μ in diameter; fresh water.

Genus *Diplosiga* Frenzel (*Codonosigopsis* Senn). With 2 collars; without lorica; a contractile vacuole; solitary or clustered (up to 4); fresh water.

*D. francei* Lemmermann (Fig. 138, i). With a short pedicel; 12μ long; flagellum as long as body.

*D. socialis* F. (Fig. 138, j). Body about 15μ long; usually 4 clustered at one end of stalk (15μ long).

**Family 3 Bicosoecidae** Poche

Small monomastigote; with lorica; solitary or colonial; collar may be rudimentary; holozoic; fresh water. Taxonomy and morphology (Grasse and Deflandre, 1952).

Genus *Bicosoeca* James-Clark. With vase-like lorica; body small, ovoid with rudimentary collar, a flagellum extending through it; protoplasmic body anchored to base by a contractile filament (flagellum?); a nucleus and a contractile vacuole; attached or free-swimming.

*B. socialis* Lauterborn (Fig. 139, a). Lorica cylindrical, 23μ by 12μ; body about 10μ long; often in groups; free-swimming in fresh water.

*B. kepneri* Reynolds. Body pyriform: 10μ by 6μ; lorica about 1.5 times the body length; flagellum about 30μ long (Reynolds, 1927).

Genus *Salpingoeca* James-Clark. With a vase-like chitinous lorica to which stalked or stalkless organism is attached; fresh or salt water. Numerous species (Pascher, 1925, 1929). Morphology (Hof-neider, 1925).

*S. fusiformis* Kent (Fig. 139, b). Lorica short vase-like, about 15–16μ long; body filling lorica; flagellum as long as body; fresh water.

Genus *Diplosigopsis* Francé. Similar to *Diplosiga* but with lorica; solitary; fresh water on algae.

*D. affinis* Lemmermann (Fig. 139, c). Chitinous lorica, spindle-form, about 15μ long; body not filling lorica; fresh water.
Genus **Histiona** Voigt. With lorica; but body without attaching filament; anterior end with lips and sail-like projection; fresh water. Morphology (Pascher, 1943).

*H. zachariasi* V. (Fig. 139, d). Lorica cup-like; without stalk; about 13μ long; oval body 13μ long; flagellum long; standing fresh water.

Genus **Poteriodendron** Stein. Similar to *Bicosoeca*; but colonial;

Genus **Codonoeca** James-Clark. With a stalked lorica; a single flagellum; 1–2 contractile vacuoles; fresh or salt water.

*C. inclinata* Kent (Fig. 139, f). Lorica oval; aperture truncate; about 23μ long; stalk twice as long; body oval, about 17μ long; flagellum 1.5 times as long as body; contractile vacuole posterior; standing fresh water.

Genus **Lagenoea** Kent. Resembles somewhat *Salpingoeca*; with lorica; but without any pedicel between body and lorica; solitary; free-swimming; fresh water.

*L. ovata* Lemmermann (Fig. 139, g). Lorica oval, 15μ long; body loosely filling lorica; flagellum 1.5 times body length; fresh water.

Genus **Stelexomonas** Lackey. A single collar longer than body;
vesicular nucleus median; a contractile vacuole terminal; individuals are enclosed in arboroid, dichotomously branching tubes; fresh water.

*S. dichotoma* L. (Fig. 140, a). Body ovoid, 10\(\mu\) by 8\(\mu\); flagellum up to 25\(\mu\) long; collar 12\(\mu\) long; the dichotomous tube infolded and wrinkled where branched; organisms are not attached to the tube (Lackey, 1942).

![Diagram of Stelexomonas dichotoma](image)

Fig. 140. a, *Stelexomonas dichotoma*, \(\times 1000\) (Lackey); b, *Oikomonas termo*, \(\times 1330\) (Lemmermann); c, *Thylacomonas compressa*, \(\times 640\) (Lemmermann); d, *Ancyromonas contorta*, \(\times 2000\) (Lemmermann); e, *Platytheca microspora*, \(\times 650\) (Stein); f, *Aulomonas purdyi*, \(\times 1000\) (Lackey); g, *Caviomonas mobilis*, \(\times 2400\) (Nie).

**Family 4 Oikomonadidae** Hartog

**Genus Oikomonas** Kent. A rounded monomastigote; uninucleate; encystment common; stagnant water, soil and exposed faecal matter. Several workers note the affinity of the members of this genus with Chrysomonadina, on the basis of general structure, cyst, etc., though lacking chromatophores. Owen (1949) points out the flagellum of Oikomonas is a simple one, typical of Chrysomonadina.

*O. termo* (Ehrenberg) (Fig. 140, b). Spherical or oval; anterior end lip-like; flagellum about twice body length; a contractile vacuole; 5–20\(\mu\) in diameter; stagnant water. Bacteria-free culture (Hardin, 1942); bacterial food (Hardin, 1944, 1944a).

**Genus Thylacomonas** Schewiakoff. Pellicle distinct; cytostome
anterior; one flagellum; contractile vacuole anterior; rare.

_T. compressa_ S. (Fig. 140, c). 22μ by 18μ; flagellum body length; fresh water.

Genus _Ancyromonas_ Kent. Ovate to triangular; free-swimming or adherent; flagellum trailing, adhesive or anchorate at its distal end, vibratile throughout remainder of its length; nucleus central; a contractile vacuole; fresh or salt water.

_A. contorta_ (Klebs) (Fig. 140, d). Triangular, flattened; posterior end pointed; 6–7μ by 5–6μ; flagellum short; a contractile vacuole; standing fresh water.

Genus _Platytheca_ Stein. With a flattened pyriform lorica, with a small aperture; 1 or more contractile vacuoles; fresh water.

_P. microspora_ S. (Fig. 140, e). Lorica yellowish brown, with a small aperture; 12–18μ long; flagellum short; among roots of _Lemna_.

Genus _Aulomonas_ Lackey. Solitary and colorless; enclosed in, but not attached to, a thin hyaline cylindrical tube, which expands like a funnel at one end and broken at the other end; fresh water.

_A. purdyi_ L. (Fig. 140, f). Ovoid, 6–8μ by 4–5μ; flagellum 10–16μ long; nucleus median; one contractile vacuole at each end of the body (Lackey, 1942).

Genus _Caviomonas_ Nie. Elongate pyriform; a single flagellum from the rounded anterior end where a vesicular nucleus is located; a band-like “peristyle” runs along the body; without cytostome; parasitic. One species (Nie, 1949).

_C. mobilis_ N. (Fig. 140, g). Body 2.2–6.6μ by 2–3μ; average 4μ by 3μ; in addition to the peristyle, a short, fine spinous strand occurs; in the caecal contents of guinea-pig, _Cavia porcella_.

Family 5 _Trypanosomatidae_ Doflein

Body characteristically leaf-like, though changeable to a certain extent; a single nucleus and a blepharoplast from which a flagellum arises (Figs. 9; 141); basal portion of the flagellum forms the outer margin of undulating membrane which extends along one side of body; exclusively parasitic; a number of important parasitic Protozoa which are responsible for serious diseases of man and domestic animals in various parts of the world are included in it. Morphology and taxonomy (Grassé, 1952).

Genus _Trypanosoma_ Gruby. Parasitic in the circulatory system of vertebrates; highly flattened, pointed at flagellate end, and bluntly rounded, or pointed, at other; polymorphism due to differences in development common; nucleus central; near aflagellate end, there is a blepharoplast from which the flagellum arises and runs toward
opposite end, marking the outer boundary of the undulating membrane; in most cases flagellum extends freely beyond body; many with myonemes; multiplication by binary or multiple fission. The organism is carried from host to host by blood-sucking invertebrates and undergoes a series of changes in the digestive system of the latter (Fig. 142). A number of forms are pathogenic to their hosts and the diseased condition is termed *trypanosomiasis* in general.

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Fig. 141. Diagram illustrating the morphological differences among the genera of Trypanosomatidae (Wenyon)

*T. gambiense* Dutton (Fig. 143, a–d). The trypanosome, as it occurs in the blood, lymph or cerebro-spinal fluid of man, is extremely active; body elongate, tapering towards both ends and sinuous; 15–30μ by 1–3μ; the small blepharoplast is located near the posterior end; flagellum arises from the blepharoplast and runs forward along the outer border of somewhat spiral undulating membrane, extending freely; binary fission; between long (dividing) and short (recently divided) forms, various intermediates occur; in man in central Africa.

No other stages are found in the human host. When a “tse-tse” fly, *Glossina palpalis* or *G. tachinoides*, sucks the blood of an infected person, the trypanosomes remain in its stomach for a few days and undergo multiplication which produces flagellates of diverse size and form until the 7th to 10th days when the organisms show a very wide range of forms. From 10th to 12th days on, long
slender forms appear in great numbers and these migrate back gradually towards proventriculus in which they become predominant forms. They further migrate to the salivary glands and attach them-
selves to the duct-wall in crithidia form. Here the development continues for 2–5 days and the flagellates finally transform themselves into small trypanosomes which are now infective. These metacyclic trypanosomes pass down through the ducts and hypopharynx. When the fly bites a person, the trypanosomes enter the victim. In addition to this so-called cyclic transmission, mechanical transmission may take place.

*Trypanosoma gambiense* is a pathogenic protozoan which causes Gambian or Central African sleeping sickness. The disease occurs in,

![Diagram](https://via.placeholder.com/150)

Fig. 143. a–d, *Trypanosoma gambiense*; e–h, *T. rhodesiense*, in stained blood smears of experimental rats, ×2300. An erythrocyte of rat is shown for comparison. a, b, typical forms; c, d, division stages; e, f, typical forms; g, h, post-nuclear forms.

and confined to, central Africa within a zone on both sides of the equator where the vectors, *Glossina palpalis* and *G. tachinoides* (on the west coastal region) live. Many wild animals have been found naturally infected by the organisms and are considered to be reservoir hosts. Among the domestic animals, the pigs appear to be one of the most significant, as they themselves are said not to suffer from infection.

The chief lesions of infection are in the lymphatic glands and in the central nervous system. In all cases, there is an extensive small-
cell infiltration of the perivascular lymphatic tissue throughout the central nervous system.

*T. rhodesiense* Stephens and Fantham (Fig. 143, e–h). Morphologically similar to *T. gambiense*, but when inoculated into rats, the position of the nucleus shifts in certain proportion (usually less than 5%) of individuals toward the posterior end, near or behind the blepharoplast, together with the shortening of body. Some consider this trypanosome as a virulent race of *T. gambiense* or one transmitted by a different vector, others consider it a human strain of *T. brucei*.

The disease caused by this trypanosome appears to be more virulent and runs a course of only a few months. It is known as Rhodesian or East African sleeping sickness. The organism is confined to south-eastern coastal areas of Africa and transmitted by *Glossina morsitans*.

*T. cruzi* Chagas (*Schizotrypanum cruzi* C). (Fig. 144). A small

![Fig. 144. *Trypanosoma cruzi* in experimental rats. a–c, flagellate forms in blood; d, e, cytozoic forms, all ×2300; f, a portion of infected cardiac muscle, ×900.](image)

curved (C or U) form about 20μ long; nucleus central; blepharoplast conspicuously large, located close to sharply pointed non-flagellate end; multiplication takes place in the cells of nearly every organ of the host body; upon entering a host cell, the trypanosome loses its flagellum and undulating membrane, and assumes a leishmania form which measures 2 to 5μ in diameter; this form undergoes repeated binary fission, and a large number of daughter individuals are produced; they develop sooner or later into trypanosomes which, through rupture of host cells, become liberated into blood stream. Life cycle (Elkeles, 1951).

This trypanosome is the causative organism of Chagas’ disease or South American trypanosomiasis which is mainly a children's disease, and is widely distributed in South and Central America and as
far north as Mexico in North America. In the infected person, the heart and skeletal muscles show minute cyst-like bodies.

The transmission of the organism is carried on apparently by numerous species of reduviid bugs, bed bugs and certain ticks, though the first named bugs belonging to genus Triatoma (cone-nosed or kissing bugs) especially *T. megista* (*Panstrongylus megistus*), are the chief vectors. When *P. megistus* (nymph or adult) ingests the infected blood, the organisms undergo division in the stomach and intestine, and become transformed into crithidia forms which continue to multiply. In eight to 10 days the metacyclic or infective trypanosomes make their appearance in the rectal region and pass out in the faeces of the bug at the time of feeding on host. The parasites gain entrance to the circulatory system when the victim scratches the bite-site or through the mucous membrane of the eye (Brumpt, 1912; Denecke and von Haller, 1939; Weinstein and Pratt, 1948).

Cats, dogs, opossums, monkeys, armadillos, bats, foxes, squirrels, wood rats, etc., have been found to be naturally infected by *T. cruzi*, and are considered as reservoir hosts. Vectors are also numerous.

No cases of Chagas' disease have been reported from the United States, but Wood (1934) found a San Diego wood rat (*Neotoma fuscipes macrotis*) in the vicinity of San Diego, California, infected by *Trypanosoma cruzi* and Packchanian (1942) observed in Texas, 1 nine-banded armadillo (*Dasypis novemcinctus*), 8 opossums (*Didelphys virginiana*), 2 house mice (*Mus musculus*), and 32 wood rats (*Neotoma micropus microps*), naturally infected by *Trypanosoma cruzi*. It has now become known through the studies of Kofoid, Wood, and others that *Triatoma protracta* (California, New Mexico), *T. rubida* (Arizona, Texas), *T. gerstaeckeri* (Texas), *T. heidemanni* (Texas), *T. longipes* (Arizona), etc., are naturally infected by *T. cruzi*. Wood and Wood (1941) consider it probable that human cases of Chagas' disease may exist in southwestern United States. In fact, the organisms from a naturally infected *Triatoma heidemanni* were shown by Packchanian (1943) to give rise to a typical Chagas' disease in a volunteer. Reduviid bugs (Usinger, 1944); Chagas' disease in the United States (Packchanian, 1950); in central Brazil (Dias, 1949).

*T. brucei* Plimmer and Bradford (Fig. 145, a). Polymorphic; 15–30μ long (average 20μ); transmitted by various species of tsetse flies, Glossina; the most virulent of all trypanosomes; the cause of the fatal disease known as "nagana" among mules, donkeys, horses, camels, cattle, swine, dogs, etc., which terminates in the death of
the host animal in from two weeks to a few months; wild animals are equally susceptible; the disease occurs, of course, only in the region in Africa where the tsetse flies live.

*T. theileri* Laveran (Fig. 145, b). Large trypanosome which occurs in blood of cattle; sharply pointed at both ends; 60–70µ long; myonemes are well developed. Cytology (Hartmann and Nöller, 1918).

*T. americanum* Crawley. In American cattle; 17–25µ or longer; only crithidia forms develop in culture. Crawley (1909, 1912) found it in 74 per cent and Glaser (1922a) in 25 per cent of cattle they examined. The latter worker considered that this organism was an intermediate form between Trypanosoma and Crithidia.

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**Fig. 145.** a, *Trypanosoma brucei*; b, *T. theileri*; c, *T. melophagium*; d, *T. evansi*; e, *T. equinum*; f, *T. equiperdum*; g, *T. lewisi*; all ×1330 (several authors).

*T. melophagium* (Flu) (Fig. 145, c). A trypanosome of the sheep; 50–60µ long with attenuated ends; transmitted by *Melophagus ovinus*.

*T. evansi* (Steel) (Fig. 145, d). In horses, mules, donkeys, cattle, dogs, camels, elephants, etc.; infection in horses seems to be usually fatal and known as “surra”; about 25µ long; monomorphic; transmitted by tabanid flies; widely distributed. Transmission (Nieschulz, 1928).

*T. equinum* Vages (Fig. 145, e). In horses in South America, causing an acute disease known as “mal de Caderas”; other domestic animals do not suffer as much as do horses; 20–25µ long; without blepharoplast.
**T. equiperdum** Doflein (Fig. 145, f). In horses and donkeys; causes "dourine," a chronic disease; widely distributed; 25-30μ long; no intermediate host; transmission takes place directly from host to host during sexual act. Nuclear division (Roskin and Schisch., 1928).

**T. hippocum** Darling. In horses and mules in Panama; the cause of "murrina" or "derrengadera"; 16-18μ long; posterior end obtuse; mechanically transmitted by flies; experimentally various domestic and wild animals are susceptible, but calf is refractory (Darling, 1910, 1911). Serological tests (Taliaferro and Taliaferro, 1934).

**T. lewisi** (Kent) (Figs. 142; 145, g). In the blood of rats; widely distributed; about 30μ long; body slender with a long flagellum; transmitted by the flea *Ceratophyllus fasciatus*, in which the organism undergoes multiplication and form change (Fig. 142); when a rat swallows freshly voided faecal matter of infected fleas containing the metacyclic organisms, it becomes infected. Many laboratory animals are refractory to this trypanosome, but guinea pigs are susceptible (Laveran and Mesnil, 1901; Coventry, 1929). Variation and inheritance of size (Taliaferro, 1921, 1921a, 1923); reproduction-inhibiting reaction product (Taliaferro, 1924, 1932); nuclear division (Wolcott, 1952).

**T. neotomae** Wood (? *T. triatomae* Kofoid and McCulloch). In wood rats, *Neotoma fuscipes annectens* and *N. f. macrotis*; resembles *T. lewisi*; about 29μ long; blepharoplast large, rod-form; free flagellum relatively short; the development in the vector flea *Orchopeas W. wickhami*, similar to that of *T. lewisi*; experimentally Norway rats are refractory (and wood rats are refractory to *T. lewisi* (Fae D. Wood, 1936)); comparative morphology of trypanosomes which occur in California rodents and shrews (Davis, 1952).

**T. duttoni** Thiroux. In the mouse; similar to *T. lewisi*, but rats are said not to be susceptible, hence considered as a distinct species; transmission by fleas. Antibodies (Taliaferro, 1938).

**T. peromysci** Watson. Similar to *T. lewisi*; in Canadian deer mice, *Peromyscus maniculatus* and others.

**T. nabiasi** Railliet. Similar to *T. lewisi*; in rabbits, *Lepus domesticus* and *L. cuniculus*.

**T. paddae** Laveran and Mesnil. In Java sparrow, *Munia oryzivora*.

**T. noctuae** (Schaudinn). In the owl *Athene noctua*.

Numerous other species occur in birds (Novy and MacNeal, 1905; Laveran and Mesnil, 1912; Wenyon, 1926). Crocodiles, snakes and turtles are also hosts for trypanosomes (Roudabush and Coatney,
Transmission is by blood-sucking arthropods or leeches.

*T. rotatorium* (Mayer) (Fig. 146, a). In tadpoles and adults of various species of frog; between a slender form with a long projecting flagellum measuring about 35μ long and a very broad one without free portion of flagellum, various intermediate forms are to be noted in a single host; blood vessels of internal organs, such as kidneys, contain more individuals than the peripheral vessels; nucleus central, hard to stain; blepharoplast small; undulating membrane highly developed; myonemes prominent; multiplication by longitudinal fission; the leech, *Placobdella marginata*, has been found to be the transmitter in some localities

*T. inopinatum* Sergent and Sergent (Fig. 146, b). In blood of various frogs; slender; 12–20μ long; larger forms 30–35μ long; blepharoplast comparatively large; transmitted by leeches.

Numerous species of Trypanosoma have been reported from the frog, but specific identification is difficult; it is better and safer to hold that they belong to one of the 2 species mentioned above until their development and transmission become known.

*T. diemyctyli* Tobey (Fig. 146, c). In blood of the newt, *Triturus viridescens*; a comparatively large form; slender; about 50μ by 2–5μ; flagellum 20–25μ long; with well developed undulating membrane.
Both fresh and salt water fish are hosts to different species of trypanosomes; what effect these parasites exercise upon the host fish is not understood; as a rule, only a few individuals are observed in the peripheral blood of the host. Transmission (Robertson, 1911); species (Laveran and Mesnil, 1912; Wenyon, 1926; Laird, 1951).

*T. granulosum* Laveran and Mesnil (Fig. 146, e). In the eel, *Anguilla vulgaris*; 70–80μ long.

*T. giganteum* Neumann (Fig. 146, d). In *Raja oxyrhynchus*; 125–130μ long.

*T. remaki* Laveran and Mesnil (Fig. 146, f). In *Esox lucius, E. reticulatus* and probably other species; 24–33μ long. (Kudo, 1921).

*T. percae* Brumpt (Fig. 146, g). In *Percia fluviatilis*; 45–50μ long.

*T. danilewskyi* Laveran and Mesnil (Fig. 146, h). In carp and goldfish; widely distributed; 40μ long.

*T. raja* Laveran and Mesnil (Fig. 146, i). In various species of *Raja*; 30–35μ long (Kudo, 1923).

Genus *Crithidia* Léger. Parasitic in arthropods and other invertebrates; blepharoplast located between central nucleus and flagellum-bearing end (Fig. 141); undulating membrane not so well developed as in *Trypanosoma*; it may lose the flagellum and form a leptomonas or rounded leishmania stage which leaves host intestine with faecal matter and becomes the source of infection in other host animals.

*C. euryophthalmi* McCulloch (Fig. 147, a–c). In gut of *Euryophthalmus convivus*; California coast.

*C. gerridis* Patton (Fig. 147, d). In intestine of water bugs, *Gerris* and Microvelia; 22–45μ long. Becker (1923) saw this in *Gerris remigis*.

*C. hyalommae* O'Farrell (Fig. 147, e, f). In body cavity of the cattle tick, *Hyalomma aegyptium* in Egypt; the flagellate through its invasion of ova is said to be capable of infecting the offspring while it is still in the body of the parent tick.

Genus *Leptomonas* Kent. Exclusively parasitic in invertebrates; blepharoplast very close to flagellate end; without undulating membrane (Fig. 141); non-flagellate phase resembles *Leishmania*.

*L. ctenocephali* Fantham (Fig. 147, g, h). In hindgut of the dog flea, *Ctenocephalus canis*; widely distributed. Morphology (Yamasaki, 1924).

Genus *Phytomonas* Donovan. Morphologically similar to *Leptomonas* (Fig. 141); in the latex of plants belonging to the families Euphorbiaceae, Asclepiadaceae, Apocynaceae, Sapotaceae and Utricaceae; transmitted by hemipterous insects; often found in
enormous numbers in localized areas in host plant; infection spreads from part to part; infected latex is a clear fluid, owing to the absence of starch grains and other particles, and this results in degeneration of the infected part of the plant. Several species.

*P. davidi* (Lafranc). 15–20μ by about 1.5μ; posterior portion of body often twisted two or three times; multiplication by longitudinal fission; widely distributed; in various species of Euphorbia.

*P. elmassiani* (Migone) (Fig. 147, *i, j*). In various species of milk-weeds; 9–20μ long; suspected transmitter, *Oncopeltus fasciatus* (Holmes, 1924); in South and North America.

Genus *Herpetomonas* Kent. Ill-defined genus (Fig. 141); exclusively invertebrate parasites; *Trypanosoma-, Crithidia-, Leptomonas-, and Leishmania-*forms occur during development. Several species. Species in insects (Drbohlav, 1925).

*H. muscarum* (Leidy) (*H. muscae-domesticae* Burnett) (Fig. 147, *k*). In the digestive tube of flies belonging to the genera: *Musca, Calliphora, Cochliomyia, Sarcophaga, Lucilia, Phormia*, etc.; up to 30μ by 2–3μ. Effect on experimental animals (Glaser, 1922); comparative study (Becker, 1923a).

*H. drosophilae* (Chatton and Alilaire) (Fig. 147, *l–n*). In intestine of *Drosophila confusa*; large leptomonal forms 21–25μ long, flagellum body-length; forms attached to rectum 4–5μ long.
Genus *Leishmania* Ross. In man or dog, the organism is an ovoid body with a nucleus and a blepharoplast; 2–5μ in diameter; with vacuoles and sometimes a rhizoplast near the blepharoplast; intracellular parasite in the cells of reticulo-endothelial system; multiplication by binary fission. In the intestine of blood-sucking insects or in blood-agar cultures, the organism develops into leptomonal form (Fig. 148, d–f) which multiplies by longitudinal fission. Nuclear division (Roskin and Romanowa, 1928).

There are known at present three “species” of *Leishmania* which are morphologically alike. They do not show any distinct differential characteristics either by animal inoculation experiments or by culture method or agglutination test.

Species of *Phlebotomus* (sand-flies) have long been suspected as vectors of *Leishmania*. When a *Phlebotomus* feeds on kala-azar patient, the leishmania bodies become flagellated and undergo multiplication so that by the third day after the feeding, there are large numbers of *Leptomonas* flagellates in the mid-gut. These flagellates migrate forward to the pharynx and mouth cavity on the 4th or 5th day. On the 7th to 9th days (after the fly is fed a second time), the organisms may be found in the proboscis. But the great majority of the attempts to infect animals and man by the bite of infected *Phlebotomus* have failed, although in a number of cases small numbers of positive infection have been reported. Adler and Ber (1941) have finally succeeded in producing cutaneous leishmaniasis in 5 out of 9 human volunteers on the site of bites by laboratory-bred *P. papatasii* which were fed on the flagellates of *Leishmania tropica* suspended in 3 parts 2.7% saline and 1 part defibrinated blood and kept at a temperature of 30°C. Swaminath, Shortt and Anderson (1942) also succeeded in producing cutaneous infections in 3 out of 5 volunteers through the bites of infected *P. argentipes*.

*L. donovani* (Laveran and Mesnil) (*L. infantum* Nicolle) (Fig. 148). As seen in stained spleen puncture smears, the organism is rounded (1–3μ) or ovoid (2–4μ by 1.5–2.5μ); cytoplasm homogeneous, but often with minute vacuoles; nucleus comparatively large, often spread out and of varied shapes; blepharoplast stains more deeply and small; number of parasites in a host cell varies from a few to over 100.

This is the cause of kala-azar or visceral leishmaniasis which is widely distributed in Europe (Portugal, Spain, Italy, Malta, Greece, and southern Russia), in Africa (Morocco, Algeria, Tunisia, Libya, Abyssinia, Sudan, northern Kenya and Nigeria), in Asia (India,
China, Turkestan, etc.), and in South America. The parasite is most abundantly found in the macrophages, mononuclear leucocytes, and polymorphonuclears of the reticulo-endothelial system of various organs such as spleen, liver, bone marrow, intestinal mucosa, lymphatic glands, etc. The most characteristic histological change appears to be an increase in number of large macrophages and mononuclears. The spleen and liver become enlarged due in part to increased fibrous tissue and macrophages.

The organism is easily cultivated in blood-agar media (p. 886). After two days, it becomes larger and elongate until it measures 14–20μ by 2μ. A flagellum as long as the body develops from the blepharoplast and it thus assumes leptomonad form (Fig. 148, f) which repeats longitudinal division. Dogs are naturally infected with *L. donovani* and may be looked upon as a reservoir host. Vectors are *Phlebotomus argentipes* and other species of Phlebotomus.

*L. tropica* (Wright). This is the causative organism of the Oriental sore or cutaneous leishmaniasis. It has been reported from Africa (mainly regions bordering the Mediterranean Sea), Europe (Spain, Italy, France, and Greece), Asia (Syria, Palestine, Armenia, Southern Russia, Iraq, Iran, Arabia, Turkestan, India, Indo-China, and China), and Australia (northern Queensland). The organisms are present in the endothelial cells in and around the cutaneous lesions, located on hands, feet, legs, face, etc.

*L. tropica* is morphologically indistinguishable from *L. donovani*, but some believe that it shows a wider range of form and size than the latter. In addition to rounded or ovoid forms, elongate forms are
often found, and even leptomonad forms have been reported from the scrapings of lesions. The insect vectors are *Phlebotomus papatasii* (p. 355), *P. sergenti* and others. Direct transmission through wounds in the skin also takes place. The lesion appears first as a small papula on skin; it increases in size and later becomes ulcerated. Microscopically an infiltration of corium and its papillae by lymphocytes and macrophages is noticed; in ulcerated lesions leishmania bodies are found in the peripheral zone and below the floor of the ulcers.

*L. brasiliensis* Vianna. This organism causes Espundia, Bubos, or American or naso-oral leishmaniasis, which appears to be confined to South and Central America. It has been reported from Brazil, Peru, Paraguay, Argentina, Uruguay, Bolivia, Venezuela, Ecuador, Colombia, Panama, Costa Rica, and Mexico.

Its morphological characteristics are identical with those of *L. tropica*, and a number of investigators combine the two species into one. However, *L. brasiliensis* produces lesions in the mucous membrane of the nose and mouth. Vectors appear to be *Phlebotomus intermedius*, *P. panamensis* and other species of the genus. Direct transmission through wounds is also possible. Fuller and Geiman (1942) find *Citellus tridecemlineatus* a suitable experimental animal.

**Family 6 Cryptobiidae** Poche

Biflagellate trypanosome-like protomonads; 1 flagellum free, the other marks outer margin of undulating membrane; blepharoplast an elongated rod-like structure, often referred to as the parabasal body; all parasitic.

**Genus Cryptobia** Leidy (*Trypanoplasma* Laveran and Mesnil). Parasitic in the reproductive organ of molluscs (Leidy, 1846) and other invertebrates; also in the blood of fishes.

*C. helicis* L. (Fig. 149, a–c). In the reproductive organ of various species of pulmonate snails: *Triodopsis albolabris*, *T. tridentata*, *Anguispira alternata* (Leidy, 1846), *Helix aspersa*, and *Monadenia fidelis* (Kozloff, 1948); 16–26.5μ by 1.5–3.3μ. Morphology and culture (Schindera, 1922).

*C. borreli* (Laveran and Mesnil) (Fig. 149, d, e). In blood of various freshwater fishes such as *Catostomus*, *Cyprinus*, etc.; 20–25μ long (Mavor, 1915).

*C. cyprini* (Plehn) (Fig. 149, f). In blood of carp and goldfish; 10–30μ long; rare.

*C. grobbeni* (Keysselitz). In coelenteric cavity of Siphonophora; about 65μ by 4μ.
Family 7 Amphimonadidae Kent

Body naked or with a gelatinous envelope; 2 equally long anterior flagella; often colonial; 1–2 contractile vacuoles; free-swimming or attached; mainly fresh water.

Genus Amphimonas Dujardin. Small oval or rounded amoeboid; flagella at anterior end; free-swimming or attached by an elongated stalk-like posterior process; fresh or salt water.

Fig. 149. a, a neutral red stained and b, a fixed and stained Cryptobia helicis, X2200 (Kozloff); c, stained specimen of the same organism, X1690 (Bélař); d, a living and e, stained C. borreli, X1730 (Mavor); f, C. cyprini, X600 (Plehn).

A. globosa Kent (Fig. 150, a). Spherical; about 13µ in diameter; stalk long, delicate; fresh water.

Genus Spongomonas Stein. Individuals in granulated gelatinous masses; 2 flagella; one contractile vacuole; colonial; with pointed pseudopodia in motile stage; fresh water.

S. uvella S. (Fig. 150, b). Oval; 8–12µ long; flagella 2–3 times as long; colony about 50µ high; fresh water.

Genus Cladomonas Stein. Individuals are embedded in dichotomous dendritic gelatinous tubes which are united laterally; fresh water.

C. fruticulosa S. (Fig. 150, c). Oval; about 8µ long; colony up to 85µ high.

Genus Rhipidodendron Stein. Similar to Cladomonas, but tubes are fused lengthwise; fresh water.
**R. splendidum** S. (Fig. 150, d, e). Oval; about 13μ long; flagella about 2–3 times body length; fully grown colony 350μ high.

Genus **Spiromonas** Perty. Elongate; without gelatinous covering; spirally twisted; 2 flagella anterior; solitary; fresh water.

*S. augusta* (Dujardin) (Fig. 150, f). Spindle-form; about 10μ long; stagnant water.

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![Fig. 150. a, Amphimonas globosa, ×540 (Kent); b, Spongomonas wella, ×440 (Stein); c, Cladomonas fruticulosa, ×440 (Stein); d, e, Rhipidodendron splendidum (d, a young colony, ×440; e, a freeliving individual, ×770) (Stein); f, Spiromonas augusta, ×1000 (Kent); g, Diplomita socialis, ×1000 (Kent); h, Streptomonas cordata, ×890 (Lemmermann); i, Dinomomas vorax, ×800 (Kent).](image-url)

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Genus **Diplomita** Kent. With transparent lorica; body attached to bottom of lorica by a retractile filamentous process; a rudimentary stigma (?); fresh water.

*D. socialis* K. (Fig. 150, g). Oval flagellum about 2–3 times the body length; lorica yellowish or pale brown; broadly spindle in form; about 15μ long; pond water.

Genus **Streptomonas** Klebs. Free-swimming; naked; distinctly keeled; fresh water.

*S. cordata* (Perty) (Fig. 150, h). Heart-shaped; 15μ by 13μ; rotation movement.
Genus **Dinomonas** Kent. Ovate or pyriform, plastic, free-swimming; 2 flagella, equal or sub-equal, inserted at anterior extremity, where large oral aperture, visible only at time of food ingestion, is also located, feeding on other flagellates; in infusions.  

*D. vorax* K. (Fig. 150, *i*). Ovoid, anterior end pointed; 15–16μ long; flagella longer than body; hay infusion and stagnant water.

**Family 8 Monadidae** Stein

Two unequal flagella; one primary and the other secondary; swimming or attached; 1–2 contractile vacuoles; colony formation frequent; free-living.

Genus **Monas** Müller (*Physomonas* Kent). Active and plastic; often attached to foreign objects; small, up to 20μ long; fresh and salt water. Some authors hold that this genus should be placed in *Chrysomonadina* on the same ground mentioned for *Oikomonas* (p. 343). Flagellar movement (Krijgsman, 1925); cyst (Scherffel, 1924); morphology and taxonomy (Reynolds, 1934).

*M. guttula* Ehrenberg (Fig. 151, *a*). Spherical to ovoid; 14–16μ long; free-swimming or attached; longer flagellum about 1–2 times body length; cysts 12μ in diameter; stagnant water.

*M. elongata* (Stokes) (Fig. 151, *b*). Elongate; about 11μ long; free-swimming or attached; anterior end obliquely truncate; fresh water.

*M. socialis* (Kent) (Figs. 8, *d*; 151, *c*). Spherical; 5–10μ long; among decaying vegetation in fresh water.

*M. vestita* (Stokes) (Fig. 151, *d*). Spherical; about 13.5μ in diameter; stalk about 40μ long; pond water. Reynolds (1934) made a careful study of the organism.

*M. sociabilis* Meyer. Body 8–10μ long by 5μ; two unequal flagella; the longer one is as long as the body and the shorter one about one-fourth; 20–50 individuals form a spheroid colony, resembling a detached colony of *Anthophysis*; polysaprobie.

Genus **Stokesiella** Lemmermann. Body attached by a fine cytoplasmic thread to a delicate and stalked vase-like lorica; 2 contractile vacuoles; fresh water.

*S. dissimilis* (Stokes) (Fig. 151, *c*). Solitary; lorica about 28μ long.

*S. leptostoma* (S.) (Fig. 151, *f*). Lorica about 17μ long; often in groups; on vegetation.

Genus **Stylobryon** Fromentel. Similar to *Stokesiella*; but colonial; on algae in fresh water.

*S. abbotti* Stokes (Fig. 151, *g*). Lorica campanulate; about 17μ long; main stalk about 100μ high; body oval or spheroidal; flagella short.
Genus **Dendromonas** Stein. Colonial; individuals without lorica, located at end of branched stalks; fresh water among vegetation.

*D. virgaria* (Weisse) (Fig. 151, *h*). About 8μ long; colony 200μ high; pond water.

**Fig. 151.** a, *Monas guttula*, ×620 (Fisch); b, *M. elongata*, ×670 (Stokes); c, *M. socialis*, ×670 (Kent); d, *M. vestita*, ×570 (Stokes); e, *Stokesiella dissimilis*, ×500 (Stokes); f, *S. leptostoma*, ×840 (Stokes); g, *Stylobryon abbotti*, ×480 (Stokes); h, *Dendromonas virgaria*, a young colony of, ×670 (Stein); i, *Cephalothamnium cyclopum*, ×440 (Stein); j, k, *Anthophysis vegetans* (*j*, part of a colony, ×230; k, an individual, ×770) (Stein).

Genus **Cephalothamnium** Stein. Colonial; without lorica, but individuals clustered at the end of a stalk which is colorless and rigid; fresh water.

*C. cyclopum* S. (Fig. 151, *i*). Ovoid; 5–10μ long; attached to body of Cyclops and also among plankton.

Genus **Anthophysis** Bory (*Anthophysa*). Colonial forms, somewhat similar to *Cephalothamnium*; stalks yellow or brownish and usually bent; detached individuals amoeboid with pointed pseudopodia.

*A. vegetans* (Müller) (Fig. 151, *j, k*). About 5–6μ long; common in stagnant water and infusion.
Family 9 **Bodinidae** Bütschli

With 2 flagella; one directed anteriorly and the other posteriorly and trailing; flagella originate in anterior end which is drawn out to a varying degree; one to several contractile vacuoles; asexual reproduction by binary fission; holozoic or saprozoic (parasitic). Morphology and taxonomy (Hollande, 1942, 1952).

Genus **Bodo** Ehrenberg (*Prowazekia* Hartman and Chagas). Small, ovoid, but plastic; cytostome anterior; nucleus central or anterior; flagella connected with 2 blepharoplasts in some species; encystment common; in stagnant water and coprozoic. Numerous species. Cytology (Bélař, 1920; Hollande, 1936).

**B. caudatus** (Dujardin) (Fig. 152, *a, b*). Highly flattened, usually tapering posteriorly; 11–22μ by 5–10μ; anterior flagellum about body length, trailing flagellum longer; blepharoplast; cysts spherical; stagnant water.

**B. edax** Klebs (Fig. 152, *c*). Pyriform with bluntly pointed ends; 11–15μ by 5–7μ; stagnant water.

Genus **Pleuromonas** Perty. Naked, somewhat amoeboid; usually attached with trailing flagellum; active cytoplasmic movement; fresh water.

**P. jaculans** P. (Fig. 152, *d*). Body 6–10μ by about 5μ; flagellum 2–3 times body length; 4–8 young individuals are said to emerge from a spherical cyst; stagnant water.

Genus **Rhynchomonas** Klebs (*Cruzella* Faria, da Cunha and Pinto). Similar to *Bodo*, but there is an anterior extension of body, in which one of the flagella is embedded, while the other flagellum trails; a single nucleus; minute forms; fresh or salt water; also sometimes coprozoic.

**R. nasuta** (Stokes) (Fig. 152, *e*). Oval, flattened; 5–6μ by 2–3μ; fresh water and coprozoic.

**R. marina** (F., C. and P.). In salt water.

Genus **Proteromonas** Kunstler (*Prowazekella* Alexeieff). Elongated pyriform; 2 flagella from anterior end, one directed anteriorly and the other, posteriorly; nucleus anterior; encysted stage is remarkable in that it is capable of increasing in size to a marked degree; exclusively parasitic; in gut of various species of lizards. Species (Grasse, 1926, 1952).

**P. lacertae** (Grassi) (Fig. 152, *f*). Elongate, pyriform; 10–30μ long, gut of lizards belonging to the genera Lacerta, Tarentola, etc.

Genus **Retortamonas** Grassi (*Embadomonas* Mackinnon). Body plastic, usually pyriform or fusiform, drawn out posteriorly; a large
cytostome toward anterior end; nucleus anterior; 2 flagella; cysts pyriform or ovoid; parasitic in the intestines of various animals. Taxonomy (Wenrich, 1932; Kirby and Honigberg, 1950).

*R. gryllotalpae* G. (Fig. 152, g). About 7–14μ (average 10μ) long; in intestine of the mole cricket, *Gryllotalpa gryllotalpa*.

![Figure 152](image-url)
with an endosome near anterior end; anterior flagellum as long as the body; posterior flagellum shorter, but thicker, in or near cytostome; cysts pyriform; 4.5–7μ long; a single nucleus and an oblong area surrounded by fibril; commensal in the lumen of human intestine; trophozoites and also cysts occur in diarrhoeic faeces; of comparatively rare occurrence. Varieties (Hogue, 1933, 1936).

*R. caviae* (Hegner and Schumaker, 1928). In the caecum of guinea-pigs; stained trophozoites 4–7μ by 2.4–3.2μ (H. and S.), 4.4–7.7μ by 4–4.3μ (Nie, 1950); stained cysts 3.4–5.2μ by 3.3–3.6μ (H. and S.), 4.5–5.7μ by 3.4–3.7μ (Nie).

Genus *Phyllomitus* Stein. Oval; highly plastic; cytostome large and conspicuous; 2 unequal flagella, each originates in a blepharoplast; fresh water or coprozoic.

*P. undulans* S. (Fig. 152, j), Ovoid; 21–27μ long; trailing flagellum much longer than anterior one; stagnant water.

Genus *Colponema* Stein. Body small; rigid; ventral furrow conspicuous, wide at anterior end; one flagellum arises from anterior end and the other from middle of body; fresh water.

*C. loxodes* S. (Fig. 152, k). 18–30μ by 14μ cytoplasm with refractile globules.

Genus *Cercomonas* Dujardin. Biflagellate, both flagella arising from anterior end of body; one directed anteriorly and the other runs backward over body surface, becoming a trailing flagellum; plastic; pyriform nucleus connected with the blepharoplast of flagella; spherical cysts uninucleate; fresh water or coprozoic.
C. longicauda D. (Fig. 152, l). Pyriform or ovoid; posterior end drawn out; 18–36μ by 9–14μ; flagella as long as body; pseudopodia; fresh water and coprozoic.

C. crassicauda D. (Fig. 152, m). 10–16μ by 7–10μ; fresh water and coprozoic.

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Chapter 15

Order 3 Polymastigina Blochmann

The Zoomastigina placed in this group possess 3–8 (in one family up to a dozen or more) flagella and generally speaking, are minute forms with varied characters and structures. Many possess a cytostome and one to many nuclei and the body is covered by a thin pellicle which allows the organism to change form, although each species shows a typical form. The cytoplasm does not show any special cortical differentiation; in many, there is an axial structure known as *axostyle* or axostylar filaments (p. 70). In Trichomonadidae, there is usually a rod-like structure, known as *costa* (Kunstler), along the base of the undulating membrane and in Devescovinidae, there is a subtriangular body, the *cresta*, directly below the basal portion of the trailing flagellum, which in some species is very large and capable of movement. At the time of division, the old costa is retained and a new one is formed; the cresta however is resorbed and two new ones are produced (Kirby). Parabasal bodies of various form and structure occur in many species.

The majority of Polymastigina inhabit the digestive tract of animals and nutrition is holozoic or saprozoic (parasitic). Many xylophagous forms hold symbiotic relationship with the host termites. Asexual reproduction is binary or multiple fission. Encystment is common. Sexual reproduction has been recognized in a few species. Taxonomy of species living in termites (Kirby, 1926).

With 1 nucleus.......................... Suborder 1 Monomondina
With 2 nuclei.......................... Suborder 2 Diplomondina (p. 392)
With more than 2 nuclei........... Suborder 3 Polymondina (p. 396)

Suborder I Monomondina

Without axial organella
With 3 flagella.......................... Family 1 Trimastigidae (p. 370)
With 4 flagella
  None undulates on body surface
  Without cell-organ of attachment. Family 2 Tetramitidae (p. 371)
  With rostellum................. Family 3 Streblomastigidae (p. 374)
  One undulates on body surface. Family 4 Chilomastigidae (p. 374)
With more than 4 flagella........... Family 5 Callimastigidae (p. 375)
With axial organella
Without undulating membrane
Without cresta
  Flagella not adhering to body
    Without rostellum........... Family 6 Polymastigidae (p. 376)
    With rostellum.............. Family 7 Oxymonadidae (p. 378)
Flagellar cords on body surface ...........................................
                        ...................................................... Family 8 Dinenymphidae (p. 379)
With cresta ...................... Family 9 Devescovinidae (p. 380)
With undulating membrane . . Family 10 Trichomonadidae (p. 385)

Family 1 Trimastigidae Kent

Genus Trimastix Kent. Oval or pyriform; naked; free-swimming; with a laterally produced membranous border; 3 flagella (1 anterior flagellum vibrating, 2 trailing); salt water. Species (Grasse, 1952a).

*T. marina* K. (Fig. 154, a). About 18μ long; salt water.

![Fig. 154. a, Trimastix marina, x1250 (Kent); b, Dallingeria drysdali, x2000 (Kent); c, Macromastix lapsa, x1500 (Stokes).](image)

*T. convexa* Grassé (*Coelotrichomastix convexa* Hollande) (Fig. 167, a). In life 10–22μ by 8–10μ; dorsal side strongly convex, ventral side concave; three free flagella nearly equally long, fourth flagellum borders the undulating membrane, present on the concave side and becomes free beyond the posterior end of body; spherical nucleus voluminous, with a large endosome; free-living and coprozoic (Hollande, 1939; Grassé, 1952a).

Genus Dallingeria Kent. Free-Swimming or attached; with trailing flagella; body small; with drawn-out anterior end; fresh water with decomposed organic matter.

*D. drysdali* K. (Fig. 154, b). Small; elongate oval; less than 6μ long; stagnant water.

Genus Macromastix Stokes. Free-swimming, somewhat like
Dallingeria, but anterior region not constricted; 3 flagella from anterior end; one contractile vacuole; fresh water.

M. lapsa S. (Fig. 154, c). Ovoid; 5.5μ long; anterior flagellum 1/2 and trailing flagella 2–3 times body length; pond water.

Genus Mixotricha Sutherland. Large; elongate; anterior tip spirally twisted and motile; body surface with a coat of flagella in closely packed transverse bands (insertion and movement entirely different from those of Trichonympha) except posterior end; 3 short flagella at anterior end; nucleus, 20μ by 2μ, connected with blepharoplasts by prolonged tube which encloses nucleus itself; cytoplasm with scattered wood chips; in termite gut. One species. Taxonomic position undetermined.

M. paradoxa S. About 340μ long, 200μ broad and 25μ thick; in gut of Mastotermes darwiniensis; Australia (Sutherland).

Family 2 Tetramitidae Bütschli

Genus Tetramitus Perty. Ellipsoidal or pyriform; free-swimming; cytostome at anterior end; 4 flagella unequal in length; a contractile vacuole; holozoic; fresh or salt water or parasitic. Species (Klug, 1936).

T. rostratus P. (Fig. 156, a). Body form variable, usually ovoid and narrowed posteriorly: 18–30μ by 8–11μ; stagnant water. Bunting (1922, 1926) observed an interesting life cycle of what appeared to be this organism which she had found in cultures of the caecal content of rats (Fig. 155). Nuclear division (Bunting and Wenrich, 1929).

T. pyriformis Klebs (Fig. 156, b). Pyriform, with pointed posterior end; 11–13μ by 10–12μ; stagnant water.

T. salinus (Entz) (Fig. 156, c). 2 anterior flagella, 2 long trailing flagella; nucleus anterior; cytostome anterior to nucleus; a groove to posterior end; cytopharynx temporary and length variable; 20–30μ long (Entz); 15–19μ long (Kirby). Kirby observed it in a pool with a high salinity at Marina, California.

Genus Collodictyon Carter. Body highly plastic; with longitudinal furrows; posterior end bluntly narrowed or lobed; no apparent cytostome; 4 flagella; a contractile vacuole anterior; fresh water.

C. triciliatum C. (Fig. 156, d). Spherical, ovoid or heart-shaped; 27–60μ long; flagella as long as the body; pond water. Cytology (Rhodes, 1919); food ingestion (Bělař, 1921).

Genus Costia Leclerque. Ovoid in front view, pyriform in profile; toward the right side, there is a shallow depression which leads into cytostome (?) and from which extend two long and two short flagella (only two flagella (Andai, 1933)); contractile vacuole posterior; encystment; ectoparasitic in freshwater fishes.
Fig. 155. Diagram illustrating the life-cycle of Tetramitus rostratus (Bunting). a, cyst; b, vegetative amoeba; c, division; d, after division; e, f, stages in transformation to flagellate form; g, fully formed flagellate; h, flagellate prior to division; i, flagellate after division; j–l, transformation stages to amoeba.

C. necatrix (Henneguy) (Fig. 156, e–j). 10–20 μ by 5–10 μ (Henneguy), 5–18 μ by 2.5–7 μ (Tavolga and Nigrelli, 1947); nucleus central; uninucleate cyst, spherical, 7–10 μ in diameter; when present in large numbers, the epidermis of the fish appears to be covered by a whitish coat. Davis (1943) found a similar organism which measured 9–14 μ by 5–8 μ, on trout, Salmo irideus and Salvelinus fontinalis, and named it Costia pyriformis.

Genus Enteromonas da Fonseca (Tricercomonas Wenyon and O’Connor). Spherical or pyriform, though plastic; 3 anterior flagella; the fourth flagellum runs along the flattened body surface and extends a little freely at the posterior tip of body; nucleus anterior; no cytostome; cyst ovoid and with 4 nuclei when mature; parasitic
in mammals. da Fonseca (1915) originally observed only 3 flagella and no cysts; 4 flagella and encysted forms were noticed in Tricercomonas by Wenyon and O'Connor (1917); in da Fonseca's original preparations, Dobell (1935) observed 4 flagella as well as cysts and concluded that Enteromonas and Tricercomonas are one and the same flagellate.

**Fig. 156.** a, *Tetramitus rostratus,* ×620 (Lemmermann); b, *T. pyriformis,* ×670 (Klebs); c, *T. salinus,* ×1630 (Kirby); d, *Collodictyon triciliatum,* ×400 (Carter); e–j, *Costia necatrix* (e, f, ×800 (Weltner); g–i, ×1400 (Moroff); j, two individuals attached to host integument ×500 (Kudo)); k, *Enteromonas hominis,* ×1730 (Wenyon and O'Connor); l, *Copromastix prowazeki,* ×1070 (Aragão).

*E. hominis* da F. (*T. intestinalis* W. and O) (Figs. 156, k; 157, a–d). Trophozoites 4–10μ by 3–6μ; nucleus circular or pyriform, with a large endosome, near anterior end; 4 flagella take their origins in blepharoplasts located close to nucleus; cytoplasm vacuolated or reticulated, contains bacteria; cysts ovoid, 6–8μ by 4–6μ; with 1, 2, or 4 nuclei; commensal in the lumen of human intestine; found in diarrhoeic stools. Widely distributed.

*E. caviae* Lynch. Similar to the species mentioned above, but slightly smaller; in the caecum of guinea-pigs (Lynch, 1922). Cytology (Nie, 1950).
Genus Copromastix Aragão. Four anterior flagella equally long; body triangular or pyramidal; coprozoic.

C. prowazeki A. (Fig. 156, l). About 16–18μ long; in human and rat faeces.

Genus Karotomorpha Travis (Tetramastix Alexeieff). Elongate pyriform; body more or less rigid; four unequal flagella at the anterior end, in two groups; nucleus anterior; without cytostome; parasitic in the intestine of Amphibia. Species (Travis, 1934).

K. bufonis (Dobell) (Fig. 157, e). Spindle in shape; 12–16μ by 2–6μ; in the intestine of frogs and toads. Cytology (Grasse, 1926).

Family 3 Streblomastigidae Kofoid and Swezy

Genus Streblomastix Kofoid and S. Spindle-form; with a rostellum, the anterior tip of which is enlarged into a sucker-like cup; below the cup are inserted 4 (Kidder) or 6 (Kofoid and Swezy) equally long flagella; extremely elongate nucleus below rostellum; body surface with 4 or more spiral ridges; in termite gut. One species.

S. strix K. and S. (Fig. 157, f, g). 15–52μ by 2–15μ; 4–8 spiral ridges; blepharoplast in rostellum; in Termopsis angusticollis.

Family 4 Chilomastigidae Wenyon

Four flagella, one of which undulates in the cytostome.

Genus Chilomastix Alexeieff. Pyriform; with a large cytostomal cleft at anterior end; nucleus anterior; 3 anteriorly directed flagella; short fourth flagellum undulates within the cleft; cysts common; in intestine of vertebrates. Several species.

C. mesnili (Wenyon) (Fig. 157, h–k). The trophozoite is oval or pyriform; 5–20 (10–15)μ long; jerky movements; a large cytostomal cleft near anterior end; nucleus, vesicular, often without endosome; 3 anterior flagella about 7–10μ long; the fourth flagellum short, undulates in the cleft which ridge is marked by 2 fibrils. The cyst pyriform; 7–10μ long; a single nucleus; 2 cytostomal fibrils and a short flagellum; commensal in the caecum and colon (some consider also in small intestine) of man. Both trophozoites and cysts occur in diarrhoeic faeces. It is widely distributed and very common. Cytology (Kofoid and Swezy, 1920); cultivation (Boeck, 1921).

C. intestinalis Kuczynski. In guinea-pigs; 13–27μ by 5–11μ (Geiman, 1935); 8.8–28μ by 6.6–11μ (Nie, 1950).

C. bettencourtii da Fonseca. In rats and mice.

C. cuniculi da F. In rabbits.

C. caprae d. F. In goat.
Fig. 157. a–d, Enteromonas hominis, ×1730 (Wenyon and O’Connor) (a, b, living and c, stained trophozoites; d, a stained cyst); e, Karotomorpha bufonis, ×2000 (Grassé); f, Streblomastix strix, ×1030; g, anterior end of the organism, showing the rostellum, blepharoplast, sucking cup and flagella (Kidder); h–k, Chilomastix mesnili, ×1530 (h, living and i, stained trophozoites; j, a fresh cyst; k, a stained cyst); l, a stained trophozoite, and m, a stained cyst of C. gallinarum, ×1330 (Boeck and Tanabe); n, Callimastix frontalis, ×1500 (Braune); o, C. equi, ×1100 (Hsiung).

C. gallinarum Martin and Robertson (Fig. 157, l, m). 11–20μ by 5–6μ; in the caeca of turkeys and chicks. Morphology (Boeck and Tanabe, 1926).

Family 5 Callimastigidae da Fonseca

Flagella 12 or more; in stomach of ruminants or in caecum and colon of horse.

Genus Callimastix Weissenberg. Ovoid; compact nucleus central or anterior; 12–15 long flagella near anterior end, vibrate in unison. Weissenberg (1912) considered this genus to be related to Lophomonas (p. 407), but organism lacks axial organellae; in Cyclops and alimentary canal of ruminants and horse.

C. cyclopis W. In body-cavity of Cyclops sp.

C. frontalis Braune (Fig. 157, n). 12 flagella; about 12μ long; flagella 30μ long; in cattle, sheep and goats.
C. equi Hsiung (Fig. 157, o). 12–15 flagella; 12–18μ by 7–10μ; nucleus central; in caecum and colon of horse.

Family 6 Polymastigidae Bützchli

Genus Polymastix Bützchli. Pyriform; four flagella arise from two blepharoplasts located at anterior end; cytostome and axostyle inconspicuous; body often covered by a protophyтан; commensals in insects. Species (Grassé, 1926, 1952).

P. melolonthae (Grassi) (Fig. 158, a). 10–15μ by 4–8μ; body covered by Fusiformis melolonthae (Grassé, 1926): in the intestine of Melolontha, Oryctes, Cetonia, Rhizotrogus, Tipula, etc.

Genus Eutrichomastix Kofoid and Swezy (Trichomastix Blochmann). Pyriform; anterior end rounded; cytostome and nucleus anterior; 3 flagella of equal length arise from anterior end, the fourth trailing; axostyle projects beyond posterior end of body; all endocommensals.

E. serpents (Dobell) (Fig. 158, b). About 10–25μ long; in intestine
of snakes: Pituophis, Eutaenia, and Python (Kofoid and Swezy, 1915).

_E_. _batrachorum_ (Dobell) (Fig. 158, c). Ovoid; 6–20μ long; in intestine of _Rana fusca_ (Dobell, 1909).

_E_. _axostylis_ Kirby (Fig. 158, d). Elongate, ellipsoid, or pyriform; axostyle projecting; 5–10.5μ by 2–3.5μ; 3 anterior flagella 5–10μ long; in gut of _Nasutitermes kirbyi_ (Kirby, 1931).

Genus _Chilomitus_ da Fonseca. Elongate oval; pellicle well developed; aboral surface convex; cytostome near anterior end, through which four flagella originating in a bi-lobed blepharoplast, protrude; rudimentary axostyle; nucleus and parabasal body below the cytostome (da Fonseca, 1915).

_C_. _caviae_ da F. (Fig. 158, e). In the caecum of guinea-pigs; stained trophozoites 6–14μ by 3.1–4.6μ; cytoplasm contains siderophilic bodies of unknown nature (Nie, 1950).

Genus _Hexamastix_ Alexeieff. Body similar to _Eutrichomastix_, but with 6 flagella, of which one trails; axostyle conspicuous; parabasal body prominent.

_H_. _termopsidis_ Kirby (Fig. 158, f). Ovoidal or pyriform; 5–11μ long; flagella 15–25μ long; in gut of _Zootermopsis angusticollis_ and _Z. nevadensis_; California (Kirby, 1930).

_H_. _caviae_ and _H_. _robustus_ were described by Nie (1950) from the caecum of guinea-pigs.

_H_. _batrachorum_ Alexeieff (Fig. 158, g). Oval or spindle form; 8–14μ by 4–8μ; flagella about body length; in gut of _Triton taeniatus_.

Genus _Protrichomonas_ Alexeieff. 3 anterior flagella of equal length, arising from a blepharoplast located at anterior end; parasitic.

_P_. _legeri_ A. (Fig. 158, h). In oesophagus of the marine fish, _Box boops_.

Genus _Monocercomonoides_ Travis (_Monocercomonas_ Grassi). Small; 4 flagella inserted in pairs in two places; two directed anteriorly and the other two posteriorly; axostyle filamentous; parasitic. Taxonomy (Travis, 1932).

_M_. _melolonthae_ (Grassi) (Fig. 158, i). Ovoid; 4–15μ long; in the larvae of _Melolontha melolontha_, etc.

Genus _Cochlosoma_ Kotlán. Body small, oval; sucker in the anterior half; 6 flagella; axostyle filamentous; parasitic (Kotlán, 1923).

_C_. _rostratum_ Kimura (Fig. 158, j). In the colon of domestic ducks, _Anas platyrhynchos_ and _Carina moschata_; 6–10μ by 4–6.5μ (Kimura, 1934). McNeil and Hinshaw (1942) observed this organism in the intestine of young poults and in the region of caecal tonsil in adults.
Family 7 Oxymonadidae Kirby

Genus Oxymonas Janicki. Attached phase with a conspicuous rostellum, the anterior end of which forms a sucking-cup for attachment; pyriform. In motile phase, rostellum is less conspicuous; 2 blepharoplasts located near the anterior extremity of axostyle, give rise to 2 flagella each; axostyle conspicuous; xylophagous; in termite and woodroach; sexual reproduction in some (Cleveland, 1950).

O. dimorpha Connell (Fig. 159, a, b). Subovoid; delicate pellicle; axostyle slightly protruding; a pair of long anterior flagella from 2 blepharoplasts, connected by rhizoplast; nucleus anterior. When attached to intestine, rostellum elongate, flagella disappear; 17µ by 14µ to 195µ by 165µ; in Neotermes simplicicornis; California and Arizona (Connell, 1930).

O. grandis Cleveland (Fig. 159, c). Body 76µ by 31µ to 183µ by 79µ; rostellum varies 30–200µ in length; nucleus without an endo-
some, anterior, about 20–23μ in diameter; axostyle consists of a staining part and a non-staining part; in the intestine of Neotermes dalbergiae and N. tectonae (Cleveland, 1935).

Genus Proboscidiella Kofoid and Swezy (Microrhopalodina Grassi and Foà; Kirbyella Zeliff). Attached and motile forms similar to Oxy-

omonas; but multinucleate; 4 flagella from each karyomastigont (p. 315); rostellum with filaments which extend posteriorly as axo-

styles; in termite gut (Kofoid and Swezy, 1926; Zeliff, 1930a).

P. kofoidi Kirby (Fig. 159, d). Average size 66μ by 46μ; rostellum as long as, or longer than, the body; karyomastigonts 2–19 or more (average 8); each mastigont with 2 blepharoplasts from which extend 4 flagella; in Cryptotermes dudleyi (Kirby, 1928).

Family 8 Dinenymphidae Grassi and Foà

Genus Dinenympha Leidy. Medium large; spindle form; 4–8 flagellar cords adhering to body which are spirally twisted about one turn; the flagella free at the posterior end; axostyle varies from cord to band; pyriform nucleus, anterior, with a large endosome; in termite gut. Species (Koidzumi, 1921).

D. gracilis L. (Fig. 160, a). 24–50μ by 6–12μ; body flattened and twisted; ends attenuated; with adhering prophytes; in Reticulitermes flavipes.

D. fimbriata Kirby (Fig. 140, b). 52–64μ by 8–18μ; 4–8 flagellar cords; with adherent prophytes; axostyle varies in width; in Reticulitermes hesperus (Kirby, 1924).

Genus Pyrsonympha Leidy. Large; club-shaped, the posterior end is rounded; body surface with 4–8 flagellar cords which are arranged lengthwise or slightly spirally; flagella extend freely posteriorly; blepharoplast at the anterior tip, often with a short process for attachment; axostyle a narrow band, may be divided into parts; large pyriform nucleus anterior; in termite gut. Species (Koidzumi, 1921); nuclear division (Cleveland, 1938).

P. vertens L. (Fig. 160, c). About 100–150μ long; 4–8 flagellar cords; in Reticulitermes flavipes. Cytology (Duboscq and Grassé, 1925).

P. granulata Powell (Fig. 160, d). 40–120μ by 5–35μ; 4–8 flagellar cords; in Reticulitermes hesperus (Powell, 1928).

Genus Saccinobaculus Cleveland. Elongate to spherical; 4, 8, or 12 flagella adhere to the body, and project out freely; axostyle is an extremely large paddle-like body and undulates, serving as cell-organ of locomotion; posterior end of axostyle enclosed in a sheath; in woodroach gut.
S. ambloaxostylus C. (Fig. 160, e–g). 65–110\(\mu\) by 18–26\(\mu\); in Cryptocercus punctulatus. Sexual reproduction (Cleveland, 1950a).

Genus Notila Cleveland. Body elongate, plastic; four flagella, the attached portion of which shows attached granules (Fig. 160, i); axostyle large, paddle-like, much broader than that of Pyrsonympha;

no axostyler sheath at posterior end, but with large granules or spherules embedded in it; in Cryptocercus punctulatus.

N. proteus C. (Fig. 160, h–j). Size not given; gametogenesis and sexual fusion, induced by the molting hormone of the host; diploid number of chromosomes 28 (Cleveland, 1950b).

Family 9 Devescovinidae Doflein

Usually 3 anterior flagella and a trailing stout flagellum; near base of trailing flagellum an elongated cresta (becoming a large
internal membrane in some species) (Fig. 161); trailing flagellum lightly adheres to body surface along edge of cresta; axostyle; parabasal body of various forms; single nucleus anterior; without undulating membrane; generally xylophagous. Cytology and morphogenesis (Kirby, 1944).

Fig. 161. A diagrammatic view of the anterior part of *Devescovina lemniscata*, showing the cresta and other organellae (Kirby).

Genus *Devescovina* Foà. Elongate body, usually pointed posteriorly; 3 anterior flagella about the body length; trailing flagellum, slender to band-form, about 1–1.5 times the body length; cresta; parabasal body spiraled around axostyle or nucleus; in termite intestine. Many species (Kirby, 1941, 1949).

*D. lemniscata* Kirby (Figs. 161; 162, a). 21–51µ by 9–17µ; trailing flagellum a band; cresta long, 7–9µ; in *Cryptotermes hermsi* and many species of the genus; species of Neotermes, Glyptotermes and Kalotermes (Kirby, 1926a).

Genus *Parajoenia* Janicki. Medium large; with rounded extremities; 3 anterior flagella and trailing flagellum long; cresta of moderate size; parabasal body well developed with its anterior end close to blepharoplast; stout axostyle expanded anteriorly into leaf-like capitulum, bearing a longitudinal keel; in intestine of termites.

*P. grassii* J. (Fig. 162, b). 29–59µ by 12–33µ; trailing flagellum
Fig. 162. a, Devescovina lemniscata, ×1600; b, Parajoenia grassii, with attached spirochaetes, ×1150; c, Foaina nana, ×1150; d, Macrotrichomonas pulchra, ×1600 (all after Kirby); e, Metadevescovina debilis, ×1130 (Light, modified).
stout, cordlike; cresta about 9\(\mu\) long; in *Neotermes connexus* (Kirby, 1937, 1942a).

Genus *Foaina* Janicki (*Janickiella* Duboscq and Grasse; *Para-devescovina*, *Crucinympha* Kirby). Small to medium large; 3 anterior flagella; trailing flagellum about twice the body length; cresta slender, 2.5–17\(\mu\) long; parabasal body single, in some with rami; in intestine of termites. Many species (Kirby, 1942a, 1949).

*F. nana* Kirby (Fig. 162, c). 6–18\(\mu\) by 4.5–8.5\(\mu\); trailing flagellum a moderately stout cord, 2–3 times the body length; cresta slender, 8.5\(\mu\) long; filament part of the parabasal body reaching the middle of body; in *Cryptotermes hermsi* and many species of the genus; also species of Glyptotermes, Iugitermes, and Procryptotermes (Kirby, 1942a).

Genus *Macrotrichomonas* Grassi. Large; 3 anterior flagella; trailing flagellum well developed, 1–1.5 times the body length; cresta a broad internal membrane, 21–86\(\mu\) long; parabasal body coiled around the axostyle, 1–13 times; in termite gut. Several species (Kirby, 1942, 1949).

*M. pulchra* G. (Fig. 162, d). 44–91\(\mu\) by 21–41\(\mu\); trailing flagellum band-form; cresta large; parabasal body coiled closely 4–10 times; in *Glyptotermes parvulus*, and many other species of the genus (Kirby, 1912).

Genus *Metadevescovina* Light. Moderately large; 3 anterior flagella; a short trailing flagellum; cresta small; parabasal body loosely coiled around axostyle; anterior end of axostyle in a loop; in termite gut. Many species (Light, 1926; Kirby, 1945).

*M. debilis* L. (Fig. 162, e). 30–70\(\mu\) by 15–30\(\mu\); in *Kalotermes hubbardi*.

Genus *Caduceia* França. Large; 3 long anterior flagella; trailing flagellum slender, shorter than body; cresta relatively small, 1–12\(\mu\) long; parabasal body coiled around axostyle 2–20 times; nucleus relatively large; axostyle terminates in filament; in termites. Several species (Kirby, 1942, 1949).

*C. bugnioni* Kirby (Fig. 163, a). 48–80\(\mu\) by 18–40\(\mu\); in *Neotermes greeni* (Kirby, 1942).

Genus *Hyperdevescovina* Kirby. Similar to *Caduceia*; but cresta very small; stout axostyle projects from the body; in Proglyptotermes, Neotermes; New Zealand and South Africa. Many species (Kirby, 1949).


Genus *Pseudodevescovina* Sutherland. Large; 3 short anterior
flagella; one short trailing flagellum; axostyle stout; cresta of moderate size; parabasal body large, divided into a number of attached cords; in termite gut. Several species (Kirby, 1945).

*P. uniflagellata* S. (Fig. 163, b). 52–95μ by 26–60μ; 3 delicate flagella, 30μ long; trailing flagellum a little stouter; cresta 11–20μ long; main parabasal body C-shaped, with 7–19 attached cords; in *Kalotermes insularis* (Kirby, 1936, 1945).

Fig. 163. a, *Caduceia bugnioni*, ×930; b, *Pseudoderescorina uniflagellata*, ×1190; c, *Bullanympha silvestrii*, ×780 (all after Kirby); d, e, *Gigantomonas herculea* (Dogiel) (d, ×530; e, amoeboid phase (Myxomonas), ×400).
Genus *Bulanympha* Kirby. Flagella and cresta similar to those in *Pseudodevescovina*; axostyle similar to that in *Caduccia*; proximal part of parabasal body bent in U-form around the nucleus and attached voluminous distal portion coiled around the axostyle; in termite gut (Kirby, 1938, 1949).

*B. silvestrii* K. (Fig. 163, c). 50-138 by 35-100; cresta about 5.8 μ long; distal portion of parabasal body coils around axostyle about twice; in *Neotermes erythraeus*.

Genus *Gigantomonas* Dogiel (*Myxomonas* D.). Medium large; 3 anterior flagella; a long and stout trailing flagellum; cresta conspicuously large; large axostyle; in termite gut. According to Kirby (1946), the so-called undulating membrane is a large cresta; in aflagellate phase (*Myxomonas*) the nuclear division takes place.

*G. herculea* D. (*M. polymorpha* D.) (Fig. 163, d, e). 60-75 by 30-35; in the intestine of *Hodotermes mossambicus* (Kirby, 1946).

Family 10 **Trichomonadidae** Wenyon

Kirby (1947) considers that *Trichomonas* and allied genera should be grouped in a new order *Trichomonadina*. He proposes four families: Monocercomonadidae, *Devescovinidae*, Calonymphidae and Trichomonadidae to be placed under it. Morphology and taxonomy (Grasse, 1952a).

Genus *Trichomonas* Donné. Pyriform; typically with four free anterior flagella; fifth flagellum along the outer margin of the undulating membrane; costa at the base of the membrane; axostyle developed, often protruding beyond the posterior end of the body; encystment has not been definitely observed; all parasitic. Numerous species (Wenrich, 1944). Cytology and morphogenesis (Kirby, 1944); division process (Kuczyński, 1918).

*T. hominis* (Davaine) (Fig. 164, a). Active flagellate, undergoing a jerky or spinning movement; highly plastic, but usually ovoid or pyriform; 5-20 μ long; cytostome near anterior end; 4 anterior flagella equally long; fifth flagellum borders undulating membrane which is seen in life; in degenerating individuals the membrane may undulate, even after loss of flagella, simulating amoeboid movement; axostyle straight along the median line; vacuolated cytoplasm with bacteria; commensal in the colon and ileum of man; found in diarrhoeic stools. Wenrich (1944) states that in all 20 cases which he studied, some or most of the individuals showed five anterior flagella and two unequal blepharooplasts.

Since encysted forms have not yet been found, transmission is assumed to be carried on by trophozoites. According to Dobell (1934),
he became infected by an intestinal Trichomonas of a monkey (Macacus nemestrinus) by swallowing “a rich two-day culture” plus bacteria which were mixed with 10 cc. of sterilized milk on an empty stomach. The presence of Trichomonas in his stools was established on the 6th day by culture and on the 13th day by microscopical examination after taking in the cultures. The infection which lasted for about four and a half years, did not cause any ill effects upon

![Diagrams](image1)

**Fig. 164.** Diagrams showing the species of Trichomonas which live in man, ×2500 (modified after Wenrich). a, *Trichomonas hominis*; b, *T. tenax*; c, *T. vaginalis*.

him. The organism is killed after five minutes' exposure to N/20 HCl at 37°C., but at 15–22°C., is able to survive, though in small numbers, up to 15 minutes after exposure to the acid (Bishop, 1930). This flagellate is widely distributed and of common occurrence, especially in tropical and subtropical regions.

*T. tenax* (Müller) (*T. elongata* Steinberg; *T. buccalis* Goodey) (Fig. 164, b). Similar to the last mentioned species; commensal in the tartar and gum of human mouth. Nomenclature (Dobell, 1939).

*T. vaginalis* Donné (Fig. 164, c). Broadly pyriform; 10–30μ by 10–20μ; cytoplasm contains many granules and bacteria; cytostome inconspicuous; nutrition parasitic and holozoic; parasitic in human reproductive organ. Although the organism does not enter the vaginal tissues, many observers believe it to be responsible for certain diseases of the vagina. Trussell and Johnson (1945) maintain that it
is capable of inciting an inflammatory reaction in the vaginal mucous membrane and according to Hogue (1943), this flagellate produces a substance which injures the cells in tissue culture. It occurs also in the male urethra (Feo, 1944). Morphology (Reuling, 1921; Wenrich, 1939, 1944, 1944a, 1947); taxonomy, structure and division (Hawes, 1947); comprehensive monograph (Trussell, 1947).

Because of the morphological similarity of these three species of human Trichomonas, a number of workers maintain that they may be one and the same species. Dobell (1934) inoculated a rich culture of Trichomonas obtained from his stools into the vagina of a monkey (Macacus rhesus) and obtained a positive infection which was easily proven by culture, but unsatisfactorily by microscopical examination of smears. The infection thus produced lasted over three years and did not bring about any ill effect on the monkey. He considers that T. vaginalis and T. hominis are synonyms and that there occur diverse strains different in minor morphological characters and physiological properties. Andrews (1929) noted the organism obtained from vaginal secretion was larger than T. hominis and its undulating membrane extended for 1/2 or 2/3 the body length, but when cultured in vitro, the organisms became smaller in size and the undulating membrane protruded beyond the body as a free flagellum. On the other hand, Stabler and his co-workers (1941, 1942) failed to obtain infections in volunteers by inoculating intravaginally with cul-

Fig. 165. a, Trichomonas microti, ×2000 (Wenrich and Saxe); b–d, T. gallinae, ×1765 (Stabler) (b, from domestic pigeon; c, from turkey; d, from red-tailed hawk); e, T. linearis, ×2000 (Kirby).
tures of *T. hominis*. Wenrich (1944) who made comparative studies of human *Trichomonas*, considers that there exist distinctly recognizable morphological differences among the three human species of *Trichomonas*, as shown in Fig. 164.

*T. macacovaginæ* Hegner and Ratcliffe. In the vagina of *Macacus rhesus*. Dobell (1934) held that this is identical with *T. vaginalis* and *T. hominis*.

*T. microti* Wenrich and Saxe (Fig. 165, a). In the caecum of rodents, *Microtus pennsylvanicus, Peromyscus leucopus, Rattus norvegicus, Mesocricetus auratus*; 4–9μ long; four free flagella; a blepharoplast; undulating membrane medium long; axostyle conspicuous.

*T. gallinae* (Rivolta) (*T. columbae* Rivolta and Delprato) (Fig. 165, b–d). Pyriform; 6–19μ by 2–9μ; ovoid nucleus anterior together with a blepharoplast and parabalidal body; axostyle protrudes a little; cytoplasmic granules; four anterior flagella 8–13μ long; autotomy; in the upper digestive tract of pigeon and also turkey, chicken, and dove. Experimentally it is transferable to quail, bob-white, hawk, canary, etc., and often fatal to hosts. Species (Travis, 1932a). Morphology (Stabler, 1941); pathology (Levine and Brandly, 1940); transmission (Levine *et al.*, 1941); distribution (Barnes, 1951; Stabler, 1951).

*T. linearis* Kirby (Fig. 165, e). Elongate spindle in form; 9–24μ by 3–8μ; in the intestine of *Orthognathotermes wheeleri*; Panama. Other species in termites (Kirby, 1931).

*T. limacis* (Dujardin). In the intestine and liver-tubules of slugs, *Deroceras agreste* (Dujardin, 1841) and *Limax flavus* (Kozloff, 1945); subspherical to ellipsoidal; 11–17μ by 8–13μ; four anterior flagella; undulating membrane extends to posterior end, with free flagellum (Kozloff).

**Genus Tritrichomonas** Kofoid. Similar to *Trichomonas* in appearance, behavior and structure, but with only three anterior flagella; parasitic. Many species.

*T. foetus* (Riedmüller) (Fig. 166, a, b). In the genitalia of cattle; pathogenic; 10–15μ long; transmission by sexual act, from cow to bull or bull to cow and also by “natural contamination” (Andrews and Miller, 1936) from cow to cow. Infection brings about permanent or temporary suspension of the conception or the death of foetus. Sheep is susceptible (Andrews and Rees, 1936). Morphology (Wenrich and Emmerson, 1933; Morgan and Noland, 1943; Kirby, 1951); effect on tissue culture (Hogue, 1938); effect on reproductibility of cow (Bartlett, 1947, 1948).

*T. fecalis* Cleveland. 5μ by 4μ to 12μ by 6μ; average dimensions
8.5μ by 5.7μ; axostyle long, protruding 1/3–1/2 the body length from the posterior end; of 3 flagella, one is longer and less active than the other two; in the faeces of man. Its remarkable adaptability observed by Cleveland was noted elsewhere (p. 34).

Fig. 166. a, *Trichomonas foetus* in life, ×1330 (Morgan and Noland); b, a stained *T. foetus*, ×1765 (Wenrich and Emmerson); c, d, *T. muris*, ×2000 (Wenrich); e, *T. batracorum*, ×1465 (Bishop); f, g, *T. augusta*, ×1455 (Samuels); h, *T. brevicollis*, ×2000 (Kirby); i, j, *Pseudotrichomonas keilini*, ×2200 (Bishop).

*T. muris* (Grassi) (Fig. 166, c, d). Fusiform; 10–16μ by 5–10μ; 3 anterior flagella short, posterior flagellum extends beyond body; axostyle large, its tip protruding; in the caecum and colon of mice (Mus, Peromyscus) (Wenrich, 1921) and ground squirrel (*Citellus lateralis chrysodeirus*) (Kirby and Honigberg, 1949). The organism
has been found within nematodes which coinhabit the host intestine. For example, Theiler and Farber (1932) found the flagellate in the chyle-stomach of Aspicularis tetraptera and Syphacia obvelata, and Becker (1933) noted two active individuals of this flagellate within the egg shell of the last-named nematode. Morphology and division (Kofoid and Swezy, 1915; Wenrich, 1921).

\[ T. caviae \] (Davaine). Ovoid or pyriform; 5–22μ long; undulating membrane long; axostyle protrudes; spherical cysts about 7μ in diameter (Galli-Valerio, 1903; Wenyon, 1926). Cytology and reproduction (Grasse and Faure, 1939).

\[ T. batrachorum \] (Perty) (Fig. 166, e). Ovoid; 14–18μ by 6–10μ (Alexeieff); in culture, 7–22μ by 4–7μ (Bishop, 1931); axostyle without granules; in the colon of frogs and toads. Bishop (1934) succeeded in infecting the tadpoles of Rana temporaria and Bufo vulgaris by feeding them on cultures free from cysts.

\[ T. augusta \] Alexeieff (Fig. 166, f, g). Elongate spindle; 15–27μ by 5–13μ; thick axostyle protrudes, and contains dark-staining granules; in the colon of frogs and toads. Morphology and division (Kofoid and Swezy, 1915; Samuels, 1941); viability (Rosenberg, 1936); in frog liver lesions (Stabler and Pennypacker, 1939).

\[ T. brevicollis \] Kirby (Fig. 166, h). Ovoid, undulating membrane curved around end; 10–17μ by 4–8μ; in the intestine of Kalotermes brevicollis; Panama.

Genus \textit{Pseudotrichomonas} Bishop. Body form, structure and movement, are exactly like those of \textit{Tritrichomonas}, but free-living in freshwater pond (Bishop, 1939).

\[ P. keilini \] B. (Fig. 166, i, j). When alive 7–11μ by 3–6μ; highly plastic; young cultures contain more globular forms, while old cultures more elongated organisms; three unequally long anterior flagella; undulating membrane short, does not extend more than 1/2 the body and without a free flagellum; cytostome; holozoic, feeding on bacteria; nucleus anterior; axostyle filamentous, invisible in life; no cysts; in a pond in Lincolnshire, England. Bishop (1935) cultivated this flagellate in serum-saline medium, in hay infusion and in pond or rain water with boiled wheat grains at 4–31°C. (Bishop, 1936, 1939).

Genus \textit{Tricercomitus} Kirby. Small; 3 anterior flagella; a long trailing flagellum, adhering to body; nucleus anterior, without endosome; blepharoplast large, with a parabasal body and an axial filament; parasitic.

\[ T. termopsidis \] K. (Fig. 167, b). 4–12μ by 2–3μ; anterior flagella 6–20μ long; trailing flagellum 19–65μ long; in gut of Zootermopsis
angusticollis, Z. nevadensis and Z. laticeps; California and Arizona. Culture and encystment (Trager, 1934).

Genus Pentatrichomonas Mesnil. Similar to Trichomonas, but with 5 free anterior flagella.

P. bengalensis Chatterjee. 9–20μ by 7–14μ; in human intestine. Kirby (1943, 1945a) observed that of the five flagella, four arise from

the end of a columnar (1–2μ long) extension, while the fifth flagellum is a little shorter and takes its origin about 1μ behind the extension.

Genus Pentatrichomonoides Kirby. Five anterior flagella and the undulating membrane; axostyle very slightly developed; fusiform parabasal body; nucleus separated from the anterior blepharoplast; in termite gut.

P. scroa K. (Fig. 167, c). 14–45μ by 6–15μ; in Cryptotermes dudleyi and Lobitermes longicollis.

Genus Pseudotrypanosoma Grassi. Large, elongate; 3 anterior flagella; undulating membrane; slender axostyle; band-like structure between nucleus and blepharoplast; parabasal body long, narrow; in termite gut.

P. giganteum G. (Fig. 167, d). 55–111μ long (Grassi); 145–205μ by
20–40μ; anterior flagella about 30μ long (Kirby); in gut of *Porotermes adamsoni* and *P. grandis*.

**Suborder 2 Diplomonadina**

The suborder consists of a number of binucleate flagellates possessing bilateral symmetry.

**Family Hexamitidae Kent**

Genus *Hexamita* Dujardin (*Octomitus Prowazek*). Pyriform; 2 nuclei near anterior end; 6 anterior and 2 posterior flagella; 2 axo-

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**Fig. 168.** a, *Hexamita inflata*, ×600 (Klebs); b, c, trophozoite and cyst of *H. intestinalis*, ×1600 (Alexeieff); d, *H. salmonis*, ×2100 (Davis); e, *H. cryptocerci*, ×1600 (Cleveland); f, *Trepomonas agilis*, ×1070 (Klebs); g, *T. rotans*, ×710 (Lemmermann); h, *Gyromonas ambulans*, ×530 (Seligo); i, *Trigonomonas compressa*, ×490 (Klebs); j, *Urophagus rostratus*, ×800 (Klebs).
styles; 1–2 contractile vacuoles in free-living forms; cytostome obscure; endoplasm with refractile granules; encystment; in stagnant water or parasitic.

_H. inflata_ D. (Fig. 168, a). Broadly oval; posterior end truncate; 13–25 μ by 9–15 μ; in stagnant water.

_H. intestinalis_ D. (Fig. 168, b, c). 10–16 μ long; in intestine of frogs, also in midgut of _Trutta fario_ and in rectum of _Motella tricirrata_ and _M. mustela_ in European waters. Morphology (Schmidt, 1920).

_H. salmonis_ (Moore) (Fig. 168, d). 10–12 μ by 6–8 μ; in intestine of various species of trout and salmon; schizogony in epithelium of pyloric caeca and intestine; cysts; pathogenic to young host fish (Moore, 1922, 1923; Davis, 1925).

_H. periplanetae_ (Bélař). 5–8 μ long; in intestine of cockroaches.

_H. cryptocerci_ Cleveland (Fig. 168, e). 8–13 μ by 4–5.5 μ; in _Crypto cercus punctulatus._

_H. meleagridis_ McNeil, Hinshaw and Kofoid (Fig. 169, a). Body 6–12 μ by 2–5 μ. It causes a severe catarrhal enteritis in young turkeys. Experimentally it is transmitted to young quail, chicks, and duckling (McNeil, Hinshaw and Kofoid, 1941).

_H. sp._ Hunminen and Wichterman (1938) (Fig. 169, b). Average dimensions 10 μ by 5.5 μ; found in the reproductive organs of the trematode, _Dero pristis inflata_, parasitic in the eel; heavily infected eggs are said not to develop.

**Genus Giardia** Kunstler (_Lamblia_ Blanchard). Pyriform to ellipsoid; anterior end broadly rounded, posterior end drawn out; bilaterally symmetrical; dorsal side convex, ventral side concave or flat, with a sucking disc in anterior half; 2 nuclei; 2 axostyles; 8 flagella in 4 pairs; cysts oval to ellipsoid; with 2 or 4 nuclei and fibrils; in the intestine of various vertebrates. Many species. Criteria for species differentiation (Simon, 1921; Hegner, 1922); cytology and taxonomy (Filice, 1952).

_G. intestinalis_ (Lambl) (G. _enterica_ Grassi; _G. lamblia_ Stiles (Fig. 169, c–q). When the flagella lash actively, the organism shows a slight forward movement with a sidewise rocking motion. The trophozoite is broadly pyriform, not plastic; 9–20 μ by 5–10 μ; sucking disc acts as attachment organella; cytoplasm hyaline; 2 needle-like axostyles; 2 vesicular nuclei near anterior margin; 8 flagella in 4 pairs; two flagella originate near the anterior end of axostyles, cross each other and follow the anterolateral margin of the disc, becoming free; two originating in anterior part of axostyles, leave the body about 1/3 from the posterior tip; two (ventral) which are thicker than others, originate in axostyles at nuclear level and remain free; two (caudal)
flagella arise from the posterior tips of axostyles; a deeply staining body may be found in cytoplasm.

The cysts are ovoid and refractile; 8–14 μ by 6–10 μ; cyst wall thin; contents do not fill the wall; 2 or 4 nuclei, axostyles, fibrils and flagella are visible in stained specimens.

This flagellate inhabits the lumen of the duodenum and other

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**Fig. 169.** a, *Hexamita meleagridis*, ×1875 (McNeal et al.); b, an egg of *Deropristis inflata* containing *Hexamita*, ×770 (Hunninen and Wichterman); c–g, *Giardia intestinalis*, ×2300 (c, front and d, side view of living organisms; e, stained trophozoite; f, fresh and g, stained mature cysts).
parts of small intestine and colon of man. Both trophozoites and cysts are ordinarily found in diarrhoeic faeces. In severe cases of infection, an enormous number of the organisms attach themselves to the mucous membrane of the intestine which may result in abnormal functions of the host tissues. In some cases, the flagellate has been reported from the gall bladder. The stools often contain unusual amount of mucus. Although there is no evidence that *G. intestinalis* attacks the intestinal epithelium, experimental observations point to its pathogenicity (Tsuchiya and Andrews, 1930). Cytology (Kofoid and Swezy, 1922).

*G. duodenalis* (Davaine). In the intestine of rabbits; 13–19 μ by 8–11 μ (Hegner, 1922).

*G. canis* Hegner. In dogs; 12–17 μ by 7.6–10 μ; cysts oval, 9–13 μ by 7–9 μ (Hegner, 1922).

*G. muris* (Grassi). In rats and mice; 7–13 μ by 5–10 μ (Simon, 1922).

*G. simoni* Lavier. In the small intestine of rats; 14–19 μ by 7–10.5 μ (Lavier, 1924); 11–16 μ by 5–8 μ (Nieschulz and Krijgsman, 1925).

*G. ondatrae* Travis. In the intestine of the muskrat, *Ondatra zibethica*; 13 μ by 7 μ (Travis, 1939); 10 μ by 5.5 μ (Waters et al.).

*G. caviae* Hegner. In the intestine of guinea-pigs; 8–14 μ by 5.5–10 μ (Hegner, 1923).

Genus *Trepomonas* Dujardin. Free-swimming; flattened; more or less rounded; cytostomal grooves on posterior half, one on each side; 8 flagella (one long and 3 short flagella on each side) arise from anterior margin of groove; near anterior margin there is a horseshoe-form structure, in which two nuclei are located; fresh water, parasitic, or coprozoic.

*T. agilis* D. (Fig. 168, f). More or less ovoid; 7–30 μ long; 1 long and 3 short flagella on each side; rotation movement; stagnant water; also reported from intestine of amphibians.

*T. rotans* Klebs (Fig. 168, g). Broadly oval; posterior half highly flattened; 2 long and 2 short flagella on each of 2 cytostomes; stagnant water.

Genus *Gyromonas* Seligo. Free-swimming; small; form constant, flattened; slightly spirally coiled; 4 flagella at anterior end; cytostome not observed; fresh water.

*G. ambulans* S. (Fig. 168, h). Rounded; 8–15 μ long; standing water.

Genus *Trigonomonas* Klebs. Free-swimming; pyriform, plastic; cytostome on either side, from anterior margin of which arise 3 flagella; flagella 6 in all; 2 nuclei situated near anterior end; movement rotation; holozoic; fresh water.
T. compressa K. (Fig. 168, i). 24–33μ by 10–16μ; flagella of different lengths; standing water (Klug, 1936).

Genus Urophagus Klebs. Somewhat similar to Hexamita; but a single cytostome; 2 moveable posterior processes; holozoic; stagnant water.

U. rostratus (Stein) (Fig. 168, j). Spindle-form; 16–25μ by 6–12μ.

Suborder 3 Polymonadina

The polymonads are multinucleate. Each nucleus is associated with a blepharoplast (from which a flagellum extends), a parabasal

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**Fig. 170.** a, Calonympha grassii, ×900 (Janicki); b, Stephanonympha nelumbium, ×400 (Kirby); c, Coronympha clevelandi, ×1000 (Kirby); d, Metacoronympha senta, ×485 (Kirby); e, Snyderella tabogae, ×350 (Kirby).
body, and an axial filament. Janicki called this complex karyomastigont (Fig. 170, a) and the complex which does not contain a nucleus, akaryomastigont (Fig. 170, e). This group includes the forms which inhabit the gut of various species of termites, most probably as symbionts.

Genus Calonympha Foà. Body rounded; large; numerous long flagella arise from anterior region; numerous nuclei; karyomastigonts and akaryomastigonts; axial filaments form a bundle; in termite gut (Foà, 1905).

C. grassii F. (Fig. 170, a). 69–90μ long; in Cryptotermes grassii.

Genus Stephanonympha Janicki. Oval, but plastic; numerous nuclei spirally arranged in the anterior half; karyomastigonts; axial filaments form a bundle; in termite gut (Janicki, 1911).

S. nelumbium Kirby (Fig. 170, b). 45μ by 27μ; in Cryptotermes hermsi.

Genus Coronympha Kirby. Pyriform with 8 or 16 nuclei, arranged in a single circle in anterior region; 8 or 16 karyomastigonts; axostyles distributed; in termite gut (Kirby, 1929, 1939).

C. clevelandi K. (Fig. 170, c). 25–53μ by 18–46μ; in Kalotermes clevelandi.

Genus Metacoronympha Kirby. Pyriform; one hundred or more karyomastigonts arranged in spiral rows meeting at the anterior end; each karyomastigont is composed of nucleus, blepharoplast, cresta, 3 anterior flagella, a trailing flagellum, and an axostyle; axostyle as in the last genus; in termite gut (Kirby, 1939).

M. senta K. (Fig. 170, d). 22–92μ by 15–67μ; karyomastigonts about 66–345 (average 150) in usually 6 spiral rows; in Kalotermes emersoni and four other species of the genus.

Genus Snyderella Kirby. Numerous nuclei scattered through the cytoplasm; akaryomastigonts close together and extend through the greater part of peripheral region; axial filaments in a bundle; in termite gut (Kirby, 1929).

S. tabogae K. (Fig. 170, e). Pyriform; rounded posteriorly; bluntly conical anteriorly; 77–172μ by 53–97μ; in Cryptotermes longicollis.

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Chapter 16
Order 4 Hypermastigina Grassi and Foà

All members of this order are inhabitants of the alimentary canal of termites, cockroaches, and woodroaches. The cytoplasmic organization is of high complexity, although there is only a single nucleus. Flagella are numerous and have their origin in the blepharoplasts located in the anterior region of body. In many species which are xylophagous, there exists a true symbiotic relationship between the host termite and the protozoans (p. 29). Method of nutrition is either holozoic or saprozoic (parasitic). Bits of wood, starch grains, and other food material are taken in by means of pseudopodia (p. 99).

Asexual reproduction is by binary fission; multiple division has also been noted in some species under certain conditions, while sexual reproduction has been observed in a few species. Encystment occurs in some genera of Lophomonadidae and certain species inhabiting woodroaches in which moulting of the host insect leads to encystment and sexual reproduction. The protozoan fauna of the colon is lost at the time of molting of the host insect, but newly molted individuals regain the fauna by proctodeal feeding (Andrews, 1930).

The number of Protozoa present in the colon of the termite is usually very enormous. The total weight of all Protozoa present in a termite worker has been estimated to be from about 1/7–1/4 (Hun- gate, 1939) or 1/3 (Katzin and Kirby, 1939) to as much as 1/2 (Cleveland, 1925) of the body weight of the host. The correlation between the termite and its intestinal flagellate fauna, has been studied by several observers. Kirby (1937) notes that certain groups of flagellates occur only in certain groups of termites, while others are widely distributed. Flagellates of one host termite introduced into individuals of another species survive for a limited time only (Light and Sanford, 1928; Cleveland, Hall et al., 1934; Dropkin, 1941, 1946). Taxonomy (Koidzumi, 1921; Kirby, 1926; Bernstein, 1928).

Body without segmented appearance
Flagella in spiral rows.............Family 1 Holomastigotidae (p. 405)
Flagella not arranged in spiral rows
Flagella in one or more anterior tufts
1 tuft of flagella.............Family 2 Lophomonadidae (p. 407)
2 tufts of flagella.............Family 3 Hoplonymphidae (p. 410)
4 tufts of flagella.............Family 4 Staurojoeninidae (p. 412)
Several tufts (loriculae).............Family 5 Kofoidiidae (p. 412)
Flagella not arranged in tufts
Posterior part without flagella.

Family 6 Trichonymphidae (p. 412)
Flagella over entire body...

Family 7 Eucomonymphidae (p. 414)
Body with segmented appearance.

Family 8 Teratologymphpidae (p. 414)

Family 1 Holomastigotidae Janicki

Genus Holomastigotes Grassi. Body small; spindle-shaped; few spiral rows reach from anterior to posterior end; nucleus anterior, surrounded by a mass of dense cytoplasm; saprozoic; in the termite gut.

*H. elongatum* G. (Fig. 171, a). In gut of *Reticulitermes lucifugus*, *R. speratus*, *R. flaviceps*, and *Macrohodoterme massambicus*; up to 70μ by 24μ (Grassi, 1892).

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**Fig. 171.** a, *Holomastigotes elongatum*, ×700 (Koidzumi); b, *Holomastigotoides hartmanni*, ×250 (Koidzumi); c, *Spirotrichonympha leidyi*, ×400 (Koidzumi); d, *S. pulchella*, ×900 (Brown); e, *Microspirotrichonympha porteri*, ×250 (Koidzumi); f, *M. ovalis*, ×600 (Brown); g, *Macrospironympha xylopletha*, ×300 (Cleveland et al.); h, *Leptospironympha eupora*, ×1050 (Cleveland et al.).
Genus **Holomastigotoides** Grassi and Foà. Large; pyriform; spiral rows of flagella as in the last genus, but more numerous (12–40 rows); a mass of dense cytoplasm surrounds ovoid nucleus near the anterior end; in termite gut (Grassi and Foà, 1911). Cytology (Cleveland, 1949).

**H. hartmanni** Koidzumi (Fig. 171, b). 50–140μ long; in Coptotermes formosanus.

**H. tusitala** Cleveland (Figs. 62; 63; 64; 172, a, b). In the hindgut of Prorhinoter mes simplex; largest species in this host; elongate pyriform; five flagellar bands, arise at the anterior end and spiral the body 5½ times; dimorphic with respect to chromosome numbers, 2 and 3; 130–200μ long. Cleveland’s observation on its chromosome cycle has been mentioned elsewhere (p. 158).

Genus **Spirotrichonympha** Grassi and Foà (1911). Moderately large; elongate pyriform; flagella deeply embedded in cytoplasm in anterior region, arising from 1 to several spiral bands; mass of dense cytoplasm conical and its base indistinct; nucleus spherical; in termite gut. Development (Duboscq and Grassé, 1928).

**S. leidyi** Koidzumi (Fig. 171, c). In Coptotermes formosanus; 15–50μ by 8–30μ.

**S. pulchella** Brown (Fig. 171, d). 36–42μ by 14–16μ; in Reticulitermes hageni.

**S. bispira** Cleveland. In Kalotermes simplicicornis; 59–102μ by 32–48μ; two flagellar bands in 34 spiral turns; resting nucleus with two chromosomes; the cytoplasmic division is unique in that portion of the anterior end shifts its position to the posterior end, where a new flagellar band develops; thus the division is longitudinal (Cleveland, 1938).

Genus **Spirotrichonymphella** Grassi. Small; without spiral ridges; flagella long; saprozoic, not wood-feeding; in termite gut.

**S. pudibunda** G. In Porotermes adamsoni; Australia. Multiple fusion (Sutherland).

Genus **Microspirotrichonympha** Koidzumi (Spironympha Koidzumi). Small, surface not ridged; spiral rows of flagella only on anterior half; a tubular structure between nucleus and anterior extremity; a mass of dense cytoplasm surrounds nucleus; with or without axial rod; in termite gut (Koidzumi, 1917, 1921).

**M. porteri** K. (Fig. 171, e). In Reticulitermes flaviceps; 20–55μ by 20–40μ.

**M. ovalis** (Brown) (Fig. 171, f). 36–48μ by about 40μ; in Reticulitermes hesperus (Brown, 1931).

Genus **Spirotrichosoma** Sutherland. Pyriform or elongate; below
operculum, two deeply staining rods from which flagella arise and which extend posteriorly into 2 spiral flagellar bands; without axostyle; nucleus anterior, median; wood chips always present, but method of feeding unknown; in Stolotermes victoriensis; Australia.

*S. capitata* S. 87μ by 38μ; flagellar bands closely spiral, reach posterior end.

Genus *Macrospironympha* Cleveland *et al.* Broadly conical: flagella on 2 broad flagellar bands which make 10–12 spiral turns, 2 inner bands; axostyles 36–50 or more; during mitosis nucleus migrates posteriorly; encystment, in which only nucleus and centrioles are retained, takes place at each ecdysis of host; in Cryptocercio punctulatus.

*M. xylopletha* C. *et al.* (Fig. 171, g). 112–154μ by 72–127μ.

Genus *Leptospironympha* Cleveland *et al.* Cylindrical; small; flagella on 2 bands winding spirally along body axis; axostyle single, hyaline; in Cryptocercio punctulatus. Several species. Sexual reproduction (Cleveland, 1951).

*L. eupora* C. *et al.* (Fig. 171, h). 30–38μ by 18–21μ.

Genus *Rostronympha* Duboscq, Grassé and Rose. Form variable, ovoid to medusoid; with or without a long contractile attaching organelle like a trunk, constricted in three places and of annulated surface; spiral ridges from which flagella arise, do not reach the posterior half; posterior half with attached spirochaetes; xylophagous; in the intestine of Anacanthotermes in Algier.

*R. magna* D., G. and R. (Fig. 172, c–e). Large individuals, 135–180μ by 110–135μ, with the trunk-like extension reaching a length of 180μ; the body proper is divided into two parts; the posterior portion may be drawn out like the manubrium of a medusa; axostyle conspicuous; in the gut of Anacanthotermes ochraceus of Algier (Duboscq and Grassé, 1943).

**Family 2 Lophomonadidae** Kent

Genus *Lophomonas* Stein. Ovoid or elongate; small: a vesicular nucleus anterior; axostyle composed of many filaments; cysts common; in colon of cockroaches.

*L. blattarum* S. (Figs. 24, a; 65; 72; 173, a–e). Small pyriform, plastic; bundle of axostylar filaments may project beyond the posterior end; active movements; binary or multiple fission; 25–30μ long; encystment; holozoic; in the colon of cockroaches, Blatta orientalis in particular; widely distributed (Kudo, 1926). Cytology (Janciki, 1910; Bělař, 1926; Kudo, 1926).

*L. striata* Bütschli (Fig. 173, f–h). Elongate spindle; body with
obliquely arranged needle-like structures which some investigators believe to be a protophyta (to which Grassé gave the name, \textit{Fusiformis lophomonadis}); bundle of axial filaments short, never protruding; movement sluggish; cyst spherical with needle-like structures; in same habitat as the last species. Cytology (Kudo, 1926a).

![Fig. 172. a, b, Holomastigotoides tusitala (Cleveland) (a, surface view; b, flagellar bands, parabasal bodies, thin axostyles); c–e, Rostronympha magna (Duboscq and Grassé) (c, a large individual with the completely extended trunk, with axostyle, ×500; d, a small medusoid form, ×1000; e, a young individual with posteriorly attached spirochaetes, ×500); f, anterior end of \textit{Joenia annectens} (Duboscq and Grassé).]

Genus \textbf{Eulophomonas} Grassi and Foà. Similar to \textit{Lophomonas}, but flagella vary from 5–15 or a little more in number; in termite gut.

\textit{E. kalotermitis} Grassi. In \textit{Kalotermes flavicollis}; this flagellate has not been observed by other workers.

Genus \textbf{Prolophomonas} Cleveland \textit{et al.} Similar to \textit{Eulophomonas}; established since Eulophomonas had not been seen by recent observers; it would become a synonym “if Eulophomonas can be found in \textit{K. flavicollis}” (Cleveland \textit{et al.}).
**P. tocopola** C. et al. (Fig. 173, i). 14–19μ by 12–15μ; in *Cryptocercus punctulatus*.

Genus **Joenia** Grassi. Ellipsoidal; anterior portion capable of forming pseudopodia; flagellar tufts in part directed posteriorly; surface covered by numerous immobile short filamentous processes,

![Diagram of various protozoa](image)

Fig. 173. a–e, *Lophomonas blattarum* (a, b, in life, ×320; c, a stained specimen; d, a trophozoite in which the nucleus is dividing; e, a stained cyst, ×1150) (Kudo); f–h, *L. striata* (f, in life, ×320; g, a stained dividing individual; h, a stained cyst, ×1150) (Kudo); i, *Prolophomonas tocopola*, ×1200 (Cleveland et al.); j, *Joenia annectens* (Grassi and Foà); k, *Microjoenia pyriformis*, ×920 (Brown); l, *Torquenympha octopus*, ×920 (Brown).

nucleus spherical, anterior; posterior to it a conspicuous axostyle composed of numerous axial filaments, a parabasal apparatus surrounding it; xylophagous; in termite gut (Grassi, 1885).

**J. annectens** G. (Figs. 172, f; 173, j). In *Kalotermes flavicollis*. Parabasal apparatus (Duboscq and Grassé, 1928a).
Genus *Joenina* Grassi. More complex in structure than that of *Joenia*; flagella inserted at anterior end in a semi-circle; parabasal bodies 2 elongated curved rods; xylophagous (Grassi, 1917).

*J. pulchella* G. In *Porotermes adamsoni*.

Genus *Joenopsis* Cutler. Oval; large; a horseshoe-shaped pillar at anterior end, flagella arising from it; some directed anteriorly, others posteriorly; parabasal bodies long rods; a strong axostyle; xylophagous; in the termite gut (Cutler, 1920).

*J. polytricha* C. In *Archotermopsis wroughtoni*; 95-129 μ long.

Genus *Microjoenia* Grassi. Small, pyriform; anterior end flat-topped; flagella arranged in longitudinal rows; axostyle; parabasal body simple; in termite gut (Grassi, 1892).

*M. pyriformis* Brown (Fig. 173, k). 44–52 μ by 24–30 μ; in *Reticulitermes hageni* (Brown, 1930).

Genus *Mesojoenia* Grassi and Foà. Large; flagellar tuft spreads over a wide area; distinct axostyle, bent at posterior end; 2 parabasal bodies; in termite gut (Grassi and Foà, 1911).

*M. decipiens* G. In *Kalotermes flavicollis*.

Genus *Torquenympha* Brown. Small; pyriform or top-form; axostyle; radially symmetrical; 8 radially arranged parabasal bodies; nucleus anterior; in termite gut (Brown, 1930).

*T. octoplus* B. (Fig. 173, l). 15–26 μ by 9–13 μ; in *Reticulitermes hesperus*.

Family 3 *Hoplonymphidae* Light

Genus *Hoplonympha* Light. Slender fusiform, covered with thick, rigid pellicular armor; each of the two flagellar tufts arises from a plate connected with blepharoplast at anterior end; nucleus near anterior extremity, more or less triangular in form; in termite gut (Light, 1926).

*H. natator* L. (Fig. 174, a, b). 60–120 μ by 5–12 μ; in *Kalotermes simplicicornis*.

Genus *Barbulanympha* Cleveland *et al.* Acorn-shaped: small, narrow, nuclear sleeve between centrioles; number of rows of flagella greater at base; large chromatin granules; numerous (80–350) parabasals; axostylar filaments 80–350; flagella 1500–13,000; different species show different number of chromosomes during mitosis; in gut of *Cryptocercus punctulatus*. Four species.

*B. ufulula* C. *et al.* (Figs. 61; 174, c). 250–340 μ by 175–275 μ; 50 chromosomes; flagellated area 36–41 μ long; centriole 28–35 μ long.

*B. laurabuda* C. *et al.* 180–240 μ by 135–170 μ; 40 chromosomes; flagellated area 29–33 μ long; centriole 24–28 μ long.
Genus *Rhynchonympha* Cleveland *et al.* Elongate; number of flagellar rows same throughout; axial filaments somewhat larger and longer, about 30; 30 parabasals: 2400 flagella: in *Cryptocercus punctulatus*. Sexual cycle (Cleveland, 1952).

*R. tarda* C. *et al.* (Fig. 175, f). 130–215\(\mu\) by 30–70\(\mu\).

Genus *Urinympha* Cleveland *et al.* Narrow, slender; flagellated area, smaller than that of the two genera mentioned above; flagella move as a unit; about 24 axial filaments; 24 parabasals; 600 flagella;

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**Fig. 174.** a, b, *Hoplonympha natator*, \(\times 450\) (Light); c, *Barbulanympha uvalula*, \(\times 210\) (Cleveland *et al.*); d, *Urinympha talea*, \(\times 350\) (Cleveland *et al.*); e, *Staurojoenina assimilis*, \(\times 200\) (Kirby); f, *Idionympha perissa*, \(\times 250\) (Cleveland *et al.*); g, *Teratonympha mirabilis*, \(\times 200\) (Dogiel).
in gut of *Cryptocercus punctulatus* (Cleveland, 1951a).

*U. talea* C. (Fig. 174, d). 75–300\(\mu\) by 15–50\(\mu\); sexual reproduction (Cleveland, 1951a).

**Family 4 Staurojoeninidae Grassi**

Genus *Staurojoenina* Grassi. Pyriform to cylindrical; anterior region conical; nucleus spherical, central; 4 flagellar tufts from anterior end; ingest wood fragments; in termite gut (Grassi, 1917).

*S. assimilis* Kirby (Fig. 174, e). 105–190\(\mu\) long; in *Kalotermes minor* (Kirby, 1926).

Genus *Idionympha* Cleveland et al. Acorn-shaped; axostyles 8–18; fine parabasals grouped in 4 areas; pellicle non-striated; nucleus nearer anterior end than that of *Staurojoenina*; flagellated areas smaller; in gut of *Cryptocercus punctulatus*.

*I. perissa* C. et. al (Fig. 174, f). 169–275\(\mu\) by 98–155\(\mu\).

**Family 5 Kofoidiidae Light**

Genus *Kofoidia* Light. Spherical; flagellar tufts composed of 8–16 *loriculae* (permanently fused bundles of flagella); without either axostyle or parabasal body; between oval nucleus and bases of flagellar tufts, there occurs a chromatin collar; in termite gut (Light, 1927).

*K. loriculata* L. (Fig. 175, a, b). 60–140\(\mu\) in diameter; in *Kalotermes simplicicorinis*.

**Family 6 Trichonymphidae Kent**

Genus *Trichonympha* Leidy (*Leidyonella* Frenzel; *Gymnonympha* Dobell; ? *Leidyopsis* Kofoid and Swezy). Anterior portion consists of nipple and bell, both of which are composed of 2 layers; a distinct axial core; nucleus central; flagella located in longitudinal rows on bell; xylophagous; in the intestine of termites and woodroach. Many species. The species inhabiting the woodroach undergo sexual reproduction at the time of molting of the host (Cleveland, 1949a) (p. 185). Species (Leidy, 1877; Kirby, 1932, 1944); nomenclature (Cleveland, 1938; Dobell, 1939); mineral ash (MacLennan and Murer, 1934).

*T. campanula* Kofoid and Swezy (Figs. 60; 175, c). 144–313\(\mu\) by 57–144\(\mu\); wood particles are taken in by posterior region of the body (Fig. 35, a); in *Zootermopsis angusticollis, Z. nevadensis* and *Z. laticeps* (Kofoid and Swezy, 1919).

*T. agilis* Leidy (Fig. 175, d). 55–115\(\mu\) by 22–45\(\mu\); in *Reticulitermes flavipes, R. lucifugus, R. speratus, R. flaviceps, R. hesperus, R. tibialis*. (Leidy, 1877).
HYPERMASTIGINA

T. grandis Cleveland et al. 190–205µ by 79–88µ; in Cryptocercus punctulatus.

Genus Pseudotrichonympha Grassi and Foà. 2 parts in anterior end as in Trichonympha; head organ with a spherical body at its tip and surrounded by a single layer of ectoplasm; bell covered by 2 layers of ectoplasm; nucleus lies freely; body covered by slightly

Fig. 175. a, b, Kofoidia loriculata, ×175, ×300 (Light); c, Trichonympha campanula, ×150 (Kofoid and Swezy); d, T. agilis, ×410 (Kirby); e, Eucomonympha imla, ×350 (Cleveland et al.); f, Rhynchonympha tarda, ×350 (Cleveland et al.).
oblique rows of short flagella; in termite gut (Grassi and Foà, 1911).

P. grassii Koidzumi. In Coptotermes formosanus; spindle-form; 200–300\(\mu\) by 50–120\(\mu\) (Koidzumi, 1921).

Genus Deltotrichonympha Sutherland. Triangular; with a small dome-shaped "head"; composed of 2 layers; head and neck with long active flagella; body flagella short, arranged in 5 longitudinal rows; flagella absent along posterior margin; nucleus large oval, located in anterior third; cytoplasm with wood chips; in termite gut. One species.

D. operculata S. Up to 230\(\mu\) long, 164\(\mu\) wide, and about 50\(\mu\) thick; in gut of Mastotermes darwiniensis; Australia.

Family 7 Eucomonymphidae Cleveland et al.

Genus Eucomonympha Cleveland et al. Body covered with flagella arranged in 2 (longer rostral and shorter post-rostral) zones; rostral tube very broad, filled with hyaline material; nucleus at base of rostrum; in gut of Cryptocercus punctulatus.

E. imla C. et al. (Fig. 175, e). 100–165\(\mu\) by 48–160\(\mu\); attached forms more elongate than free individuals; sexual reproduction (Cleveland, 1950).

Family 8 Teratonymphidae Koidzumi

Genus Teratonympha Koidzumi (Teranympha K.; Cyclonympha Dogiel). Large and elongate; transversely ridged, and presents a metameric appearance; each ridge with a single row of flagella; anterior end complex, containing a nucleus; reproduction by longitudinal fission; in termite gut (Koidzumi, 1917, 1921; Dogiel, 1917).

T. mirabilis K. (Fig. 174, g). 200–300\(\mu\) or longer by 40–50\(\mu\); in Reticulitermes speratus. Mitosis (Cleveland, 1938a).

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— (1937) Host-parasite relations in the distribution of Protozoa in termites. Ibid., 41:189.
— (1919a) IV. Ibid., 20:99.
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— and Sanford, Mary F.: (1928) Experimental transfaunation of termites. Ibid., 31:269.
THE members of this class do not possess any thick pellicle and, therefore, are capable of forming pseudopodia (p. 49). The term 'amoeboid' is often used to describe their appearance. The pseudopodia serve ordinarily for both locomotion and food-capturing. The peripheral portion of the body shows no structural differentiation in Amoebina, Proteomyxa, and Mycetozoa. Internal and external skeletal structures are variously developed in other orders. Thus, in Testacea and Foraminifera, there is a well-developed test or shell that usually has an aperture, through which the pseudopodia are extruded; in Heliozoa and Radiolaria, skeletons of various forms and materials are developed.

The cytoplasm is, as a rule, differentiated into the ectoplasm and the endoplasm, but this differentiation is not constant. In Radiolaria, there is a perforated membranous central capsule which marks the border line between the two cytoplasmic regions. The endoplasm contains the nucleus, food vacuoles and various granules. The majority of Sarcodina are uninucleate, but species of Foraminifera and Mycetozoa are multinucleate in certain phases during their development. In the family Paramoebidae, there occurs a peculiar secondary nucleus.

The Sarcodina are typically holozoic. Their food organisms are Protozoa, small Metazoa and Protophyta, which present themselves conspicuously in the cytoplasm. The methods of ingestion have already been considered (p. 97). One or more contractile vacuoles are invariably present in forms inhabiting the fresh water, but absent in parasitic forms or in those which live in the salt water.

Asexual reproduction is usually by binary (or rarely multiple) fission, budding, or plasmotomy. Definite proof of sexual reproduction has been noted in a comparatively small number of species. Encystment is common in the majority of Sarcodina, but is unknown in a few species. The life-cycle has been worked out in some forms and seems to vary among different groups. The young stages are either amoeboid or flagellate, and on this account, it is sometimes very difficult to distinguish the Sarcodina and the Mastigophora. In some forms the mature trophic stage may show an amoeboid or flagellate phase, owing to differences in environmental conditions.

The Sarcodina are divided into two subclasses as follows:

With lobopodia, rhizopodia, or filopodia... Subclass 1 Rhizopoda (p. 418)
With axopodia.......................... Subclass 2 Actinopoda (p. 505)
Subclass 1 Rhizopoda Siebold

The name Rhizopoda has often been used to designate the entire class, but it is used here for one of the subclasses, which is further subdivided into five orders, as follows:

Without test or shell
   With radiating pseudopodia ...................... Order 1 Proteomyxa
   With rhizopodia; forming plasmodium . . Order 2 Mycetozoa (p. 427)
   With lobopodia ................................ Order 3 Amoebina (p. 435)

With test or shell
   Test single-chambered; chitinous .......... Order 4 Testacea (p. 472)
   Test 1- to many-chambered; calcareous . Order 5 Foraminifera (p. 493)

Order 1 Proteomyxa Lankester

A number of incompletely known Rhizopods are placed in this group. The pseudopodia are filopodia which often branch or anastomose with one another. In this respect the Proteomyxa show affinity to the Mycetozoa. Flagellate swarmer and encystment occur commonly. The majority of Proteomyxa lead parasitic life in algae or higher plants in fresh or salt water. Taxonomy (Valkanov, 1940).

Pseudoplasmodium-formation .................... Family 1 Labyrinthulidae
   Solitary and Heliozoa-like
      With flagellate swarmer ................. Family 2 Pseudosporidae (p. 420)
      Without flagellate swarmer .......... Family 3 Vampyrellidae (p. 420)

Family 1 Labyrinthulidae Haeckel

Small fusiform protoplasmic masses are grouped in network of sparingly branched and anastomosing filopodia; individuals encyst independently; with or without flagellate stages.

Genus Labyrinthula Cienkowski. Minute forms feeding on various species of algae in fresh or salt water; often brightly colored due to carotin. Jepps (1931) found these organisms common in marine aquaria. Young (1943) considers the six known species as actually three species and two varieties, while Watson (1951) holds that only one species, L. macrocystis, should be recognized.

L. cienkowskii Zopf (Fig. 176, a). Attacks Vaucheria in fresh water.
L. macrocystis Cienkowski. Renn (1934, 1936) found a species in the diseased leaf-tissue of the 'spotting and darkening' eel-grass, Zostera marina, along the Atlantic coast of the United States. Young (1943) identified the organism which he studied as L. macrocystis, and noted that its hosts included various algae and three genera of Naiadaceae: Zostera, Ruppia and Zannichellia.

The 'net-plasmodium' contains fusiform cells which average in size
18μ by 4μ and which multiply by binary fission; many cells encyst together within a tough, opaque membrane. The growth is best at 14–24°C. and at 12–22 per cent chlorinity (Young). Watson and Ordal (1951) cultivated the organism on agar and sea water with various bacteria, and found that the organism is fusiform in young cultures; highly motile; filamentous projections are formed from the flat mucoid lamellae, secreted by the organism, and expand to form passways over which the organism travels; holozoic, saprozoic.

Genus Labyrinthomyxa Duboscq. Body fusiform; amoeboid and flagellate phases, variable in size; flagellate stage penetrates the host cell membrane; in plants.

Fig. 176. a, Labyrinthula cienkowskii, ×200 (Doflein); b–e, Labyrinthomyxa sauvageau (b, c, flagellate forms, ×100; d, e, amoeboid forms, ×500) (Duboscq); f, g, Pseudospora volvocis, ×670 (Robertson); h–j, Protomonas amyli (Zopf); k, l, Vampyrella lateritia, ×530 (k (Leidy), l (Doflein)); m, n, Nuclearia delicatula, ×300 (Cash).
L. sauvageau D. (Fig. 176, b–e). Fusiform body 7–11μ long; pseudoplasmomium-formation; ameboid stage 2.5–14μ long; flagellate stage 7–18μ long; parasitic in Laminaria lejolii at Roscoff, France.

Family 2 Pseudosporidae Berlese

Genus Pseudospora Cienkowski. Body minute; parasitic in algae and Mastigophora (including Volvocidae); organism nourishes itself on host protoplasm, grows and multiplies into a number of smaller individuals, by repeated division; the latter biflagellate, seek a new host, and transform themselves into ameboid stage; encystment common. Morphology and development (Schussnig, 1929).

P. volvocis C. (Fig. 176, f, g). Heliozoan form about 12–30μ in diameter; pseudopodia radiating; cysts about 25μ in diameter; in species of Volvox. Morphology (Roskin, 1927).

P. parasitica C. Attacks Spirogyra and allied algae.

P. eudorini Roskin. Heliozoan forms 10–12μ in diameter; radiating pseudopodia 2–3 times longer; ameboid within host colony; cysts 15μ in diameter; in Eudorina elegans.

Genus Protomonas Cienkowski. Body irregularly rounded with radiating filopodia; food consists of starch grains; division into bi-flagellate organisms which become ameboid and unite to form pseudoplasmomium; fresh or salt water.

P. amyli C. (Fig. 176, h–j). In fresh water.

Family 3 Vampyrellidae Doflein

Filopodia radiate from all sides or formed from a limited area; flagellate forms do not occur; the organism is able to bore through the cellulose membrane of various algae and feeds on protoplasmic contents; body often reddish because of the presence of carotin; multinucleate; multiplication in encysted stage into uni- or multi-nucleate bodies; cysts often also reddish.

Genus Vampyrella Cienkowski. Heliozoa-like; endoplasm vacuolated or granulated, with carotin granules; numerous vesicular nuclei and contractile vacuoles; multinucleate cysts, sometimes with stalk; 50–700μ in diameter. Several species.

V. lateritia (Fresenius) (Fig. 176, k, l). Spherical; orange-red except the hyaline ectoplasm; feeds on Spirogyra and other algae in fresh water. On coming in contact with an alga, it often travels along it and sometimes breaks it at joints, or pierces individual cell and extracts chlorophyll bodies by means of pseudopodia; multiplication in encysted condition; 30–40μ in diameter. Behavior (Lloyd, 1926, 1929).
Genus *Nuclearia* Cienkowski. Subspherical, with sharply pointed fine radiating pseudopodia; actively moving forms vary in shape; with or without a mucous envelope; with one or many nuclei; fresh water.

Fig. 177. a, *Arachnula impatiens*, ×670 (Dobell); b, c, *Chalmydomyxa montana*: b, ×270 (Cash); c, ×530 (Penard); d, *Rhizoplasma kaiseri*, (Verworn); e, *Biomyxa vagans*, ×200 (Cash); f, *Penardia mutabilis*, ×200 (Cash); g, *Hyalodiscus rubicundus*, ×370 (Penard).
N. delicatula C. (Fig. 176, m, n). Multinucleate; bacteria often adhering to gelatinous envelope; up to 60μ in diameter.

N. simplex C. Uninucleate; 30μ in diameter.

Genus Arachnula Cienkowski. Body irregularly chain-form with filopodia extending from ends of branches; numerous nuclei and contractile vacuoles; feeds on diatoms and other microorganisms.

A. impatiens C. (Fig. 177, a). 40–350μ in diameter.

Genus Chlamydomyxa Archer. Body spheroidal; ectoplasm and endoplasm well differentiated; endoplasm often green-colored due to the presence of green spherules; numerous vesicular nuclei; 1–2 contractile vacuoles; secretion of an envelope around the body is followed by multiplication into numerous secondary cysts; cyst wall cellulose; in sphagnum swamp.

C. montana Lankester (Fig. 177, b, c). Rounded or ovoid; cytoplasm colored; about 50μ in diameter; when moving, elongate with extremely fine pseudopodia which are straight or slightly curved and which are capable of movement from side to side; non-contractile vacuoles at bases of grouped pseudopods; in active individual there is a constant movement of minute fusiform bodies (function?); when extended 100–150μ long; total length 300μ or more; fresh water among vegetation.

Genus Rhizoplasma Verworn. Spherical or sausage-shaped; with anastomosing filopodia; orange-red; with a few nuclei.

R. kaiseri V. (Fig. 177, d). Contracted form 0.5–1 mm. in diameter; with 1–3 nuclei; pseudopodia up to 3 cm. long; extended body up to 10 mm. long; originally described from Red Sea.

Genus Chondropus Greeff. Spherical to oval; peripheral portion transparent but often yellowish; endoplasm filled with green, yellow, brown bodies; neither nucleus nor contractile vacuoles observed; pseudopods straight, fine, often branched; small pearl-like bodies on body surface and pseudopodia.

C. viridis G. Average diameter 35–45μ; fresh water among algae.

Genus Biomyxa Leidy (Gymnophrys Cienkowski). Body form inconstant; initial form spherical; cytoplasm colorless, finely granulated, capable of expanding and extending in any direction, with many filopodia which freely branch and anastomose; cytoplasmic movement active throughout; numerous small contractile vacuoles in body and pseudopodia; with one or more nuclei.

B. vagans L. (Fig. 177, e). Main part of body, of various forms; size varies greatly; in sphagnous swamps, bog-water, etc.

B. cometa (C.). Subspherical or irregularly ellipsoidal; pseudopodia small in number, formed from 2 or more points; body 35–40μ, or up
SARCODINA, PROTEOMYXA

to 80μ or more; pseudopodia 400μ long or longer. Cienkowski maintained that this was a moneran.

Genus *Penardia* Cash. When inactive, rounded or ovoid; at other times expanded; exceedingly mobile; endoplasm chlorophyll-green with a pale marginal zone; filopodia, branching and anastomosing, colorless; nucleus inconspicuous; one or more contractile vacuoles, small; fresh water.

*P. mutabilis* C. (Fig. 177, f). Resting form 90–100μ in diameter; extended forms (including pseudopodia) 300–400μ long.

Genus *Hyalodiscus* Hertwig and Lesser. Discoid, though outline varies; endoplasm reddish, often vacuolated and sometimes shows filamentous projections reaching body surface; a single nucleus; ectoplasmic band of varying width surrounds the body completely; closely allied to *Vampyrella*; fresh water.

*H. rubicundus* H. and L. (Fig. 177, g). 50–80μ by about 30μ; polymorphic; when its progress during movement is interrupted by an object, the body doubles back upon itself, and moves on in some other direction; fresh water ponds among surface vegetation.

Genus *Leptomyxa* Goodey. Multinucleate, thin, amoeboid organisms; multinucleate cysts formed by condensation of protoplasm; free-living in soil (Goodey, 1915).

*L. reticulata* G. (Fig. 178, a–c). Body composed of a thin transparent protoplasm; when fully extended, 3 mm. or more in length; superficially resembles an endosporous mycetozoan, but no reversible cytoplasmic movement; multinucleate with eight to 20 to several hundred nuclei; nuclei, 5–6μ in diameter, with a large endosome; nuclear division simultaneous, but not synchronous; plasmotomy; plasmogamy; cysts multinucleate, by local condensation of protoplasm; widely distributed in British soil (Singh, 1948, 1948a). McLennan (1930) found a similar organism in and on the root of diseased hops in Tasmania.

Genus *Megamoebomyxa* Nyholm. Extremely large amoeboid organism; when contracted, lobulate, with adhering detritus; when cultured at 8–10°C. on debris, filopodia are formed and form-change occurs; lobate during locomotion; “nutrient chiefly detritus”; Marine. One species (Nyholm, 1950).

*M. argillobia* N. (Fig. 178, d). An opaque white organism; up to 25 mm. long; polymorphic; in marine sediment, rich in debris at the depth of 45–70 m.; Gullmar Fjord, Sweden.

Genus *Reticulomyxa* Nauss. Highly polymorphic, multinucleate amoeboid organism; rhizopodia radiating from a central mass of undifferentiated granular protoplasm with many non-contractile vacu-
Fig. 178. a–c, *Leptomyxa reticulata*, ×73 (Singh) (a, a trophozoite; b, cyst-formation; c, a cyst); d, an individual of *Megamoebomyxa argillobia*, showing the changes of body form, ×2/3 (Nyholm); e, a young trophozoite of *Reticulomyxa filosa*, ×3 (Nauss).

oles; plasmotomy usually into three, after discarding extraneous particles and migrating to new site; when transferred to fresh dish of water, "spore-like" bodies are dispersed; fresh water among decaying leaves. Nauss (1949) points out its affinity to Proteomyxa, Mycetozoa and Foraminifera.

*R. filosa* N. (Fig. 178, e). On moist blotting paper the central mass is an elevated body, but in water it spreads into a broad sheet, 4–6 mm. in diameter; pseudopodia may be up to 10 times the diameter of the central white mass; encystment occurs when subjected to
lower temperature or when cultured with algae; food consists of “worms,” rotifers and organic debris.

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Chapter 18
Order 2 Mycetozoa de Bary

The Mycetozoa had been considered to be closely related to the fungi, being known as Myxomycetes, or Myxogasteres, the 'slime molds.' Through extended studies of their development, de Bary showed that they are more closely related to the Protozoa than to the Protophyta, although they stand undoubtedly on the border-line between these two groups of microorganisms. The Mycetozoa occur on dead wood or decaying vegetable matter of various kinds.

The most conspicuous part of a mycetozoan is its plasmodium which is formed by fusion of many myxamoebae, thus producing a large multinucleate body (Fig. 179, a). The greater part of the cytoplasm is granulated, although there is a thin layer of hyaline and homogeneous cytoplasm surrounding the whole body. The numerous vesicular nuclei are distributed throughout the granular cytoplasm. Many small contractile vacuoles are present in the peripheral portion of the plasmodium. The nuclei increase in number by division as the body grows; the division seems to be amitotic during the growth period of the plasmodium, but is mitotic prior to the spore-formation. The granulation of the cytoplasm is due to the presence of enormous numbers of granules which in some forms are made up of carbonate of lime. The plasmodium is usually colorless, but sometimes yellow, green, or reddish, because of the numerous droplets of fluid pigment present in the cytoplasm.

The food of Mycetozoa varies among different species. The great majority feed on decaying vegetable matter, but some, such as Badhamia, devour living fungi. Thus the Mycetozoa are holozoic or saprozoic in their mode of nutrition. Pepsin has been found in the plasmodium of Fuligo and is perhaps secreted into the food vacuoles, into which protein materials are taken. The plasmodium of Badhamia is said to possess the power of cellulose digestion.

When exposed to unfavorable conditions, such as desiccation, the protoplasmic movement ceases gradually, foreign bodies are extruded, and the whole plasmodium becomes divided into numerous sclerotia or cysts, each containing 10–20 nuclei and being surrounded by a resistant wall (b). These cysts may live as long as three years. Upon return of favorable conditions, the contents of the sclerotia germinate, fuse together, and thus again produce plasmodia (c–e).

When lack of food material occurs, the plasmodium undergoes
changes and develops sporangia. The first indication of this process is the appearance of lobular masses of protoplasm in various parts of the body (f, g). These masses are at first connected with the streaming protoplasmic thickenings, but later become completely segregated into young sporangia. During the course of sporangium-formation, foreign bodies are thrown out of the body, and around each sporangium there is secreted a wall which, when mature, possesses a wrinkled appearance (h). The wall continues down to the substratum as a slender stalk of varying length, and in many genera the end of a stalk spreads into a network over the substratum, which forms the base, hypothallus, for the stalk. With these changes the interior

Fig. 179. The life-cycle of the endosporous mycetozoan (de Bary, Lister, and others). a, plasmodium-formation by fusion of numerous myxamoebae; b, c, formation of sclerotium; d, e, germination of sclerotium and formation of plasmodium; f, portion of a plasmodium showing streaming protoplasmic thickenings; g, h, formation of sporangia; i, a sporangium opened, showing capillitium; j, a spore; k, germination of spore; l, myxamoeba; m, n, myxoflagellates; o-q, multiplication of myxoflagellate; r, microcyst; s, myxamoeba. Variously magnified.
of the sporangium becomes penetrated by an anastomosing network, *capillitium*, of flat bands which are continuous with the outer covering (j). Soon after the differentiation of these protective and supporting structures, the nuclei divide simultaneously by mitosis and the cytoplasm breaks up into many small bodies. These uninucleate bodies are the *spores* which measure 3–20μ in diameter and which soon become covered by a more or less thick cellulose membrane (j), variously colored in different species.

The mature sporangium breaks open sooner or later and the spores are carried, and scattered, by the wind. When a spore falls in water, its membrane ruptures, and the protoplasmic contents emerge as an amoebula (k, l). The amoebula possesses a single vesicular nucleus and contractile vacuoles, and undergoes a typical amoeboid movement. It presently assumes an elongate form and one flagellum or two unequally long flagella (Elliott, 1948) develop from the nucleated end, thus forming a *myxoflagellate* (m, n) which undergoes a peculiar dancing movement and is able to form short, pointed pseudopodia from the posterior end. It feeds on bacteria, grows and multiplies by binary fission (o–q). After a series of division, the myxoflagellate may encyst and becomes a *microcyst* (r). When the microcyst germinates, the content develops into a myxamoeba (s) which, through fusion with many others, produces the plasmodium mentioned above. This is the life-cycle of a typical endosporous mycetozoon.

In the genus Ceratiomyxa in which spores are formed on the surface of *sporophores*, the development is briefly as follows: the plasmodium lives on or in decayed wood and presents a horn-like appearance. The body is covered by a gelatinous hyaline substance, within which the protoplasmic movements may be noted. The protoplasm soon leaves the interior and accumulates at the surface of the mass; at first as a close-set reticulum and then into a mosaic of polygonal cells, each containing a single nucleus. Each of these cells moves outward at right angles to the surface, still enveloped by the thin hyaline layer, which forms a stalk below. These cells are spores which become ellipsoid and covered by a membrane when fully formed. The spore is uninucleate at first, but soon becomes tetranucleate. When a spore reaches the water, its content emerges as an amoebula which divides three times, forming 8 small bodies, each of which develops a flagellum and becomes a myxoflagellate. The remaining part of the development is presumably similar to that of the endosporous form. Morphology (de Bary, 1864, 1884; MacBride, 1922; Jahn, 1928; MacBride and Martin, 1934).
A large number of mycetozoan genera and species are known (Hagelstein, 1944). The order is divided here into two suborders. Spore develops into myxoflagellate; myxamoebae fuse completely and form plasmodium. Suborder 1 Eumycetozoa No flagellate stage; myxamoebae grouped prior to spore-formation, but do not fuse to form a true plasmodium. Suborder 2 Sorophora (p. 433)

Suborder 1 Eumycetozoa Zopf

Spores develop within sporangia
Spores violet or violet-brown
Sporangia with lime
Lime in small granular form.

Family 1 Physaridae

Fig. 180. a, b, Badhamia utricularis Berkeley (a, cluster of sporangia, X4; b, part of capillitium and spore-cluster, X140) (Lister); c, d, Fuligo septica Gmelin (c, a group of sporangia, X3/4; d, part of capillitium and two spores, X120) (Lister); e, f, Didymium effusum Link (e, sporangium, X12; f, portion of capillitium and wall of sporangium showing the crystals of calcium carbonate and two spores, X200) (Lister); g, h, Stemonitis splendens Rostafinski (g, three sporangia, X2; h, columella and capillitium, X42) (Lister).

Genus Badhamia Berkeley (Fig. 180, a, b)
Capillitium, a course network with lime throughout.

Genus Fuligo Haller (Fig. 180, c, d)
Capillitium, a delicate network of threads with vesicular expansions filled with granules of lime.
Lime in crystalline form.

Family 2 Didymiidae
Genus Didymium Schrader (Fig. 180, e, f)
Lime crystals stellate, distributed over the wall of sporangium.
Sporangia without lime
Sporangia stalked..........................Family 3 Stemonitidae

Genus Stemonitis Gleditsch (Fig. 180, g, h)
Sporangium-wall evanescent; capillitium arising from all parts of
columella to form a network.
Sporangium combined into aethalium..........................
..........................Family 4 Amaurochaetidae

Genus Amaurochaete Rostafinski (Fig. 181, a, b)
With irregularly branching thread-like capillitium.
Spores variously colored, except violet
Capillitium absent or not forming a system of uniform threads.
Sporangium-wall membranous; with minute round granules......
..........................Family 5 Cribrariidae

Fig. 181. a, b, Amaurochaete fuliginosa MacBride (a, group of sporangia, \( \times \frac{1}{4} \); b, capillitium, \( \times 10 \) (Lister); c, empty sporangium of Cribraria aurantiaca Schrader, \( \times 20 \) (Lister); d, sporangium of Orcadella operculata Wingate, \( \times 80 \) (Lister); e, cluster of sporangia of Tubulina fragiformis Persoon, \( \times 3 \) (Lister); f, aethalium of Reticularia lycoperdon Bull., \( \times 1 \) (Lister); g, aethalium of Lycogala miniatum Persoon \( \times 1 \) (Lister); h-j, Trichia affinis de Bary (h, group of sporangia, \( \times 2 \); i, elater, \( \times 250 \); j, spore, \( \times 400 \) (Lister); k, l, Arcyria punicea Persoon (k, four sporangia, \( \times 2 \); l, part of capillitium, \( \times 250 \) and a spore, \( \times 560 \) (Lister); m, n, Ceratiomyxa fruticulosa MacBride (m, sporophore, \( \times 40 \); n, part of mature sporophore, showing two spores, \( \times 480 \) (Lister).
Genus *Cribraria* Persoon (Fig. 181, c)
Sporangia stalked; wall thickened and forms a delicate persistent network expanded at the nodes.

Sporangia solitary; stalked .......................... Family 6 Liceidae

Genus *Orcadella* Wingate (Fig. 181, d)
Sporangia stalked, furnished with a lid of thinner substance.

Sporangium-wall membranous without granular deposits ........

.......................... Family 7 Tubulinidae

Genus *Tubulina* Persoon (Fig. 181, e)
Sporangia without tubular extensions.

Many sporangia more or less closely fused to form large bodies (aethalia); sporangium-wall incomplete and perforated  ......

.......................... Family 8 Reticulariidae

Genus *Reticularia* Bulliard (Fig. 181, f)
Walls of convoluted sporangia incomplete, forming tubes and folds with numerous anastomosing threads.

Sporangia forming aethalium ............... Family 9 Lycogalidae

Genus *Lycogala* Micheli (Fig. 181, g)
Capillitium a system of uniform threads
Capillitium threads with spiral or annular thickenings ........

.......................... Family 10 Trichiidae

Genus *Trichia* Haller (Fig. 181, h–j)
Capillitium abundant, consisting of free elasters with spiral thickenings.

Capillitium combined into an elastic network with thickenings in forms of cogs, half-rings, spines, or warts. Family 11 Areyriidae

Genus *Arcyria* Wiggers (Fig. 181, k, l)
Sporangia stalked; sporangium-wall evanescent above, persistent and membranous in the lower third.

Capillitium abundant; sporangia normally sessile ..........

.......................... Family 12 Margaritidae

Genus *Margarita* Lister
Capillitium profuse, long, coiled hair-like.

Spores develop on the surface of sporophores
Spores white; borne singly on filiform stalk ..................

.......................... Family 13 Ceratiomyxidae
Genus Ceratiomyxa Schröter (Fig. 181, m, n)
Suborder 2 Sorophora Lister
Pseudoplasmodium incomplete; myxamoeba of limax-form.

Family 1 Guttuliniidae
Pseudoplasmodium complete; myxamoeba with short pointed pseudopodia.

Family 2 Dictyosteliidae

The Proteomyxa and the Mycetozoa as outlined above, are not distinctly defined groups. In reality, there are a number of forms which stand on the border line between them. Development of Dictyostelium discoideum (Raper, 1940); food habits and distribution of Dictyostelium (Singh, 1947, 1947a).

Phytomyxinea Poche

These organisms which possess a large multinucleate amoeboid body, are parasitic in various plants and also in a few animals. Taxonomy (Palm and Burk, 1933; Cook, 1933).

Genus Plasmodiophora Woronin. Parasitic in the roots of cabbage and other cruciferous plants. The organism produces knotty enlargements, sometimes known as "root-hernia," or "fingers and toes" (Fig. 182, a). The small (haploid) spore (b) gives rise to a myxoflagellate (c–f) which penetrates the host cell. The organism grows in size and multiplies (g, h). The plasmodium divides into sporangia. Flagellated gametes that develop from them fuse in pairs, giving rise to diploid zygotes. These zygotes develop further into plasmodia in which haploid spores are produced. Morphology (Jones, 1928); cytology (Milovidov, 1931).

P. brassicae W. (Fig. 182). In Brassica spp.
Genus **Sorosphaera** Schröter. Parasitic in *Veronica* spp.
Genus **Tetramyxa** Goebel. In Rupia, Zannichellia, etc.
Genus **Octomyxa** Couch, Leitner and Whiffen. In *Achlya glomerata*.
Genus **Sorodiscus** Lagerheim and Winge. In Chara, Callitriche, etc.
Genus **Polymyxa** Ledingham. In Triticum, etc.
Genus **Membranosorus** Ostenfeld and Petersen. In *Heteranthera dubia*.
Genus **Spongospora** Brunchorst. Parasitic in Solanum; the diseased condition of potatoes is known as powdery or corky scab.
Genus **Ligniera** Maire and Tison. In Alisma, Juncus, etc.

**References**

—— (1884) Vergleichende Morphologie und Biologie der Pilze, Mycetozoen, und Bacterien. Leipzig.
—— (1928) Myxomycetenstudien. XII. Ibid., p. 80.
Palm, B. T. and Burk, Myrle: (1933) The taxonomy of the Plasmodiophoraceae. Ibid., 79:262.
—— (1947a) II. Ibid., 1:361.
THE Amoebina show a very little cortical differentiation. There is no thick pellicle or test, surrounding the body, although in some a delicate pellicle occurs. The cytoplasm is more or less distinctly differentiated into the ectoplasm and the endoplasm. The ectoplasm is hyaline and homogeneous, and appears tougher than the endoplasm. In the endoplasm, which is granulated or vacuolated, are found one or more nuclei, various food vacuoles, crystals, and other inclusions. In the freshwater forms, there is at least one distinctly visible contractile vacuole. The pseudopodia are lobopodia, and ordinarily both the ectoplasm and endoplasm are found in them. They are formed by streaming or fountain movement of the cytoplasm. In some members of this order, the formation of pseudopodia is eruptive or explosive, since the granules present in the endoplasm break through the border line between the two cytoplasmic layers and suddenly flow into the pseudopodia. Asexual reproduction is ordinarily by binary fission, although multiple fission may occasionally take place. Encystment is of common occurrence. Sexual reproduction, which has been reported in a few species, has not been confirmed.

The Amoebina inhabit all sorts of fresh, brackish, and salt waters. They are also found in moist soil and on ground covered with decaying leaves. Many are inhabitants of the digestive tract of various animals, and some are pathogenic to the hosts.

The taxonomic status of the group is highly uncertain and confusing, since their life-histories are mostly unknown and since numerous protozoans other than the members of this group often possess amoeboid stages.

The order is subdivided into four families as follows:

With amoeboid and flagellate stages ........................................... Family 1 Naegleriidae

Amoeboid stage only

With one or more nuclei of one kind

Free-living ........................................... Family 2 Amoebidae (p. 437)
Parasitic ........................................... Family 3 Endamoebidae (p. 443)
With a secondary nucleus ............... Family 4 Paramoebidae (p. 405)

Family 1 Naegleriidae

The members of the two genera placed in this family possess both amoeboid and flagellate phases (diphyasic). In the former, the organ-
ism undergoes amoeboid movement by means of lobopodia and in the latter the body is more or less elongated. Binary fission seems to take place during the amoeboid phase only. Thus these are diphasic amoebae, in which the amoeboid stage predominates over the flagellate. The amoeboid phase is often a 'limax' form; under natural circumstances, it is often exceedingly difficult by observing the amoeboid stage only, to determine whether they belong to this family or the family Amoebidae.

Genus Naegleria Alexeieff. Minute flagellate stage with 2 flagella; amoeboid stage resembles Vahlkampfia (p. 442), with lobopodia; cytoplasm differentiated; vesicular nucleus with a large endosome; contractile vacuole conspicuous; food vacuoles contain bacteria; cysts uninucleate; free-living in stagnant water and often coprozoic. Taxonomy and cytology (Rafalko, 1947; Singh, 1952).

Fig. 183. a–c, trophozoite, flagellate phase and cyst (all stained) of Naegleria gruberi, ×750 (Alexeieff); d–f, similar stages of N. bistadialis, ×750 (Kühn); g–j, trophozoite, flagellate phase, cyst, and excystment of Trimastigamoeba philippinensis, ×950 (Whitmore).

N. gruberi (Schrading) (Fig. 183, a–c). Amoeboid stage 10–36μ by 8–18μ; cyst uninucleate; cyst wall with several apertures; flagellate stage 18μ by 8μ; stagnant water and often coprozoic.

N. bistadialis (Puschkarew) (Fig. 183, d–f). Similar in size; but cyst with a smooth wall.

Genus Trimastigamoeba Whitmore. Flagellate stage bears 3 flagella of nearly equal length; vesicular nucleus with a large endosome; amoeboid stage small, less than 20μ in diameter; uninucleate cysts with smooth wall; stagnant water.

T. philippinensis W. (Fig. 183, g–j). Amoeboid stage 16–18μ in diameter; oval cysts 13–14μ by 8–12μ; flagellate stage 16–22μ by 6–8μ.
Family 2 Amoebidae Bronn

These amoebae do not have flagellate stage and are exclusively amoeboid (*monophasic*). They are free-living in fresh or salt water, in damp soil, moss, etc., and a few parasitic; 1, 2, or many nuclei; contractile vacuoles in freshwater forms; multiplication by binary or multiple fission or plasmotomy; encystment common. Genera (Leidy, 1879; Penard, 1902; Singh, 1952).

Genus *Amoeba* Ehrenberg (*Proteus* Müller; *Amiba* Bory). Amoeboid; a vesicular nucleus, either with many spherical granules or with a conspicuous endosome; usually one contractile vacuole; pseudopodia are lobopodia, never anastomosing with one another; holozoic; in fresh, brackish or salt water. Numerous species. Nomenclature (SchaefTer, 1926; Mast and Johnson, 1931; Kudo, 1952).

*A. proteus* (Pallas) (Figs. 2, e, f; 25; 33, b, c; 43, f; 45–47; 68; 184, a, b). Up to 600μ or longer in largest diameter; creeping with a few large lobopodia, showing longitudinal ridges; ectoplasm and endoplasm usually distinctly differentiated; typically uninucleate; nucleus discoidal but polymorphic; endoplasmic crystals truncate bipyramidal, up to 4.5μ long (SchaefTer, 1916); nuclear and cytosome divisions show a distinct correlation (p. 169); fresh water. Cytology (Mast, 1926; Mast and Doyle, 1935, 1935a); nuclear division (Chalkley, 1936; Liesche, 1938).

*A. discoides* Schaeffer (Figs. 43, g; 184, c). About 400μ long during locomotion; a few blunt, smooth pseudopodia; crystals abundant, truncate bipyramidal, about 2.5μ long (Schaeffer); endoplasm with numerous coarse granules; fresh water.

*A. dubia* S. (Figs. 43, h–l; 184, d). About 400μ long; numerous pseudopodia flattened and with smooth surface; crystals, few, large, up to 30μ long and of various forms among which at least 4 types are said to be distinct (Schaeffer); contractile vacuole one or more; fresh water. Nuclear division (Dawson *et al*., 1935); viscosity (Angerer, 1942); contractile vacuole (Dawson, 1945).

*A. verrucosa* Ehrenberg (Figs. 33, a, d–k; 44, a; 184, c). Ovoid in general outline with wart-like expansions; body surface usually wrinkled, with a definite pellicle; pseudopodia short, broad and blunt, very slowly formed; nucleus ovoid, vesicular, with a large endosome; contractile vacuole; up to 200μ in diameter; fresh water among algae.

*A. striata* Penard (Fig. 184, f). Somewhat similar to *A. verrucosa*, but small; body flattened; ovoid, narrowed and rounded posteriorly; nucleus vesicular; contractile vacuole comparatively large and often
not spherical; extremely delicate pellicle shows 3 or 4 fine longitudinal lines which appear and disappear with the movement of the body; 25–45μ by 20–35μ; fresh water among vegetation.

Fig. 184. a, b, *Amoeba proteus* (a, ×130 (Schaeffer), b, cyst (Doflein)); c, *A. discoides*, ×130 (Schaeffer); d, *A. dubia*, ×130 (Schaeffer); e, *A. verrucosa*, ×200 (Cash); f, *A. striata*, ×400 (Penard); g, *A. guttula*, ×800 (Penard); h, *A. limicola*, ×530 (Penard).

*A. guttula* Dujardin (Fig. 184, g). Ovoid during locomotion, narrowed posteriorly and often with a few minute, nipple-like dentations; movement by wave-like expansions of ectoplasm; endoplasm granulated, with crystals; nucleus vesicular; a single contractile vacuole; 30–35μ by 20–25μ; fresh water in vegetation.

*A. limicola* Rhumbler (Fig. 184, h). Somewhat similar to *A. gut-
tula; body more rounded; locomotion by eruption of cytoplasm through the body surface; 45–55μ by 35μ; nucleus vesicular; fresh water among vegetation.

A. spumosa Gruber (Fig. 2, c, d; 185, a). Somewhat fan-shaped; flattened; during locomotion broad pseudopodia with pointed end; temporary posterior region with nipple-like projections; a small
number of striae become visible during movement, showing there is a very thin pellicle; endoplasm always vacuolated, the vacuoles varying in size (up to 30μ in diameter); vesicular nucleus with an endosome; 50–125μ long during locomotion; fresh water.

_A. vespertilio_ Penard (Fig. 185, b, c.) Pseudopodia conical, comparatively short, connected at base by web-like expansions of ectoplasm; endoplasm colorless, with numerous granules and food particles; a single vesicular nucleus with a large endosome; contractile vacuoles; 60–100μ long; fresh water. Cannibalism (Lapage, 1922); contractile vacuole (Hyman, 1936); morphology and biology (Raabe, 1951).

_A. gorgonia_ P. (Fig. 185, d–f). Body globular when inactive with a variable number of radiating “arms,” formed on all sides; when in locomotion, clavate; nucleus vesicular, with a large endosome; rounded forms 40–50μ in diameter; clavate individuals up to 100μ; fresh water among vegetation.

_A. radiosa_ Ehrenberg (Fig. 185, g). Small, usually inactive; globular or oval in outline; with 3–10 radiating slender pseudopodia which vary in length and degree of rigidity; when pseudopods are withdrawn, the organism may be similar to _A. proteus_ in general appearance; pseudopods straight, curved or spirally coiled; size varies, usually about 30μ in diameter, up to 120μ or more; fresh water.

Genus *Dinamoeba* Leidy. Essentially Amoeba, but the temporary posterior region of body with retractile papillae; body surface including pseudopods and papillae, bristling with minute spicules or motionless cils; often surrounded by a thick layer of delicate hyaline jelly, even during locomotion; fresh water.

_D. mirabilis_ L. (Fig. 185, h). Oval to limaciform; spheroid when floating; pseudopodia numerous, conical; ectoplasm clear, usually with cils; endoplasm with food vacuoles, oil (?) spherules and large clear globules; nucleus and contractile vacuole obscure; spherical forms 64–160μ in diameter; creeping forms 152–340μ by 60–220μ; cyst about 160μ in diameter (Groot, 1936); in sphagnous swamp.

Genus *Pelomyxa* Greeff. Large amoeboid organisms, ranging from 0.5 to 4 or 5 mm. in length when clavate and moving progressively; nuclei numerous, less than 100 to 1000 or more; many small contractile vacuoles; refringent bodies (“Glanzkörper”) of various dimension and number; with or without bacterial inclusions (which Penard and others consider as symbiotic); holozoic on plant or animal organisms or detritus; plasmotomy simple or multiple; in fresh water. Several species (Kudo, 1946). Nomenclature (Schaeffer, 1926; Mast and Johnson, 1931; Rice, 1945; Kudo, 1946, 1952; Wilber, 1947).
P. palustris G. (P. villosa Leidy) (Fig. 186, a). Large; 2–3 mm. or larger in diameter; sluggish, with usually one broad pseudopodium; undifferentiated cytoplasm with many nuclei and various inclusions such as fragments of plant bodies, numerous small sand particles, etc., which brings about opacity and dark coloration of body; in addition bacteria (Cladothrix pelomyxae Veley, Myxococcus pelomyxae Keller and Bacterium parapelomyxae Keller) occur in the cytoplasm.

Fig. 186. a, Pelomyxa palustris, ×160 (Leidy); b, c, P. carolinensis, ×45 (Kudo) (b, an individual in locomotion; c, feeding form); d, e, P. illinoensis, ×40 (Kudo) (d, an individual in locomotion; e, a more or less stationary animal); f, Vahlkampfia patuxent, ×660 (Hogue); g, h, Acanthamoeba castellanii, ×1270 (Hewitt); i, j, A. hyalina, ×840 (Dobell).

which some observers consider as symbionts; cyst with two to three envelopes (Stolc, see Kudo, 1951); feeds on plant and inorganic debris; polysaprobic in still stagnant water, buried in mud. Central Europe, Great Britain and North America. Morphology (Greeff, 1874; Hollande, 1945); locomotion (Okada, 1930a; Mast, 1934); plasmogamy (Okada, 1930); laboratory cultivation (Hollande, 1945).
P. carolinensis Wilson (Figs. 66; 71; 186, b, c). Monopodal forms 1–5 mm. long; polypodal forms 1–2 mm. in diameter; locomotion active; nuclei up to 1000 or more, circular in front view, about 20μ in diameter and ellipsoid in profile; fluid and food vacuoles, crystals, many contractile vacuoles; feeds on various Protozoa and invertebrates; easily cultivated in laboratory; plasmotomy into two to six individuals; nuclear division simultaneous and synchronous; experimental plasmogamy; no encystment in the Illinois stock, but New Jersey stock is said to encyst (Musacchia, 1950); North America. Distribution (Kudo, 1946); morphology (Wilson, 1900; Andersen, 1942; Kudo, 1946); plasmotomy (Schaeffer, 1938; Kudo, 1949); nuclear division (Kudo, 1947); locomotion (Wilber, 1946); permeability (Belda, 1942-1943); effect of x-irradiation (Daniels, 1951, 1952, 1952a).

P. illinoensis Kudo (Fig. 186, d, e). The organism resembles the last-named species, but much smaller in size; 500–1000μ in length; clavate forms seldom exceed 1.5 mm.; several hundred nuclei, spherical, 14–16μ in diameter; peripheral granules of the nuclei are large and often discoid, irregularly distributed; crystals occur abundantly in all physiological conditions; chalky white in reflected light; plasmotomy into two to five daughters; encystment and excystment take place freely in cultures; cysts measure 250–350μ in diameter with usually two membranes, a multinucleate amoeba emerges from a cyst after several weeks (Kudo, 1950, 1951). Other species of Pelomyxa (Kudo, 1951).

Genus Vahlkampfia Chatton and Lalung-Bonnaire. Small amoebae; vesicular nucleus with a large endosome and peripheral chromat; with polar caps during nuclear division; snail-like movement, with one broad pseudopodium; cysts with a perforated wall; fresh water or parasitic. Nuclear division (Jollos, 1917).

V. limax (Dujardin). 30–40μ long; fresh water.

V. patuxent Hogue (Fig. 186, f). In the alimentary canal of the oyster; about 20μ long during the first few days of artificial cultivation, but later reaching as long as 140μ in diameter; ordinarily one large broad fan-shaped pseudopodium composed of the ectoplasm; in culture, pseudopodium-formation eruptive; holozoic on bacteria; multiplication by fission or budding; encystment rare; cysts uninucleate.

Genus Hartmannella Alexeieff. Small amoebae, with moderately or well-developed ectoplasm; vesicular nucleus with a large endosome; mitotic figure ellipsoidal or cylindrical, without polar caps. Cysts rounded; wall smooth or slightly wrinkled in one species.
Several species. Volkonsky (1933) distinguishes four groups. Species and morphology (Singh, 1952); nuclear division (Jollos, 1917).

*H. hyalina* (Dangeard). 20–25μ in diameter; ectoplasm well developed; endoplasm vacuolated; slender pseudopodia extend in different directions; Hartmann and Chagas observed a centriole in the endosome.

Genus *Acanthamoeba* Volkonsky. Small amoebae similar to *Hartmannella*; ectoplasm is not well developed; mitotic figure at the end of metaphase, a straight or concave spindle with sharply pointed poles. Cysts enveloped by two membranes, the outer envelope being highly wrinkled and mammillated. Several species.

A. *castellanii* (Douglas) (Fig. 186, g, h). In association with fungi and certain bacteria; Hewitt obtained the organism from agar cultures of sample soil taken from among the roots of white clover; co-existing with yeast-like fungi, *Flavobacterium trifolium* and *Rhizobium* sp.; 12–30μ in diameter; some cysts are said to remain viable at 37°C, for 6 days.

A. *hyalina* (Dobell and O’Connor) (Fig. 186, i, j). According to Volkonsky, the organism described by Dobell and O’Connor as *Hartmannella hyalina*, is transferred to this genus. Small amoeba; 9–17μ in diameter when rounded; a single contractile vacuole; binary fission; mitotic figure a sharply pointed spindle. Cysts spherical; 10–15μ in diameter; with a smooth inner and a much wrinkled outer wall; easily cultivated from old faeces of man and animals; also in soil and fresh water.

Genus *Sappinia* Dangeard. With two closely associated nuclei.

*S. diploidea* (Hartmann and Nägler). Coprozoic in the faeces of different animals; pseudopodia short, broad, and few; highly vacuolated endoplasm with 2 nuclei, food vacuoles, and a contractile vacuole; surface sometimes wrinkled; the nuclei divide simultaneously; during encystment, two individuals come together and secrete a common cyst wall; 2 nuclei fuse so that each individual possesses a single nucleus; finally cytoplasmic masses unite into one; each nucleus gives off reduction bodies (?) which degenerate; 2 nuclei now come in contact without fusion, thus producing a binucleate cyst (Hartmann and Nägler).

**Family 3 Endamoebidae** Calkins

Exclusively parasitic amoebae; the vegetative form is relatively small and occurs mostly in the alimentary canal of the hosts; contractile vacuoles absent, except in *Hydramoeba*; multiplication by binary fission; encystment common. The generic differentiation is
based upon the morphological characteristics of the nucleus. Summary No. 99 of 'Opinions Rendered' by the International Commission of Zoological Nomenclature (1928) holds that Entamoeba is a synonym of Endamoeba; in the present work, however, Endamoeba and Entamoeba are separated, since the two groups of species placed under them possess different nuclear characteristics (Fig. 187). Nomenclature (Dobell, 1919, 1938; Kirby, 1945; Hemming, 1951).

Genus **Endamoeba** Leidy (1879). Nucleus spheroidal to ovoid; membrane thick; in life, filled with numerous granules of uniform dimensions along its peripheral region; upon fixation, a fine chromatic network becomes noticeable in their stead; central portion

coarsely reticulated; with several endosomes between the two zones (Fig. 187, a); in some, cytoplasm becomes prominently striated during locomotion; in the intestine of invertebrates.

**E. blattae** (Bütschli) (Fig. 188). In the colon of cockroaches; 10–150μ in diameter; rounded individuals with broad pseudopodia, show a distinct differentiation of cytoplasm; elongated forms with a few pseudopodia, show ectoplasm only at the extremities of the pseudopods; endoplasm of actively motile trophozoites shows a distinct striation, a condition not seen in other amoebae; fluid-filled vacuoles occur in large numbers; amoebae feed on starch grains, yeast cells, and bacteria, all of which coexist in the host organ; cysts, 20–50μ in diameter, commonly seen in the colon contents, with often more than 60 nuclei. The life-cycle of this amoeba is still unknown. Mercier (1909) held that when the multinucleate cysts gain entrance to the host intestine through its mouth, each of the cyst-nuclei becomes the center of a gamete; when the cyst-membrane ruptures, the gametes are set free and anisogamy takes place, resulting in forma-

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**Fig. 187.** Diagram showing the stained nuclei of the trophozoites of five genera of parasitic amoebae. a, Endamoeba; b, Entamoeba; c, Iodamoeba; d, Endolimax; e, Dientamoeba.
tion of numerous zygotes which develop into the habitual trophozoites. Morphology (Leidy, 1879; Kudo, 1926; Morris, 1936; Meglitsch, 1940).

*E. thomsoni* Lucas. In the colon of cockroaches; 7–30μ in diameter; very adhesive; 1–3 large peripheral granules on the nuclear membrane; cysts 8–16μ in diameter, with 1–4 nuclei (Lucas, 1927).

*E. disparata* Kirby. In colon of *Microtermes hispaniolae*; 20–40μ long; active; xylophagous (Kirby, 1927).

![Fig. 188. *Endamoeba blattae.* a–c, trophozoites in life, X530; d, a stained binucleate amoeba; e, f, stained and fresh cysts, X700 (Kudo).](image)

*E. majestas* K. (Fig. 189, a). In the same habitat; 65–165μ in diameter; many short pseudopodia; cytoplasm filled with food particles (Kirby, 1927).

*E. simulans* K. (Fig. 189, b). In the gut of *Microtermes pana-maensis*; 50–150μ in diameter.

*E. sabulosa* K. In the same habitat; small 19–35μ in diameter.

*E. pellucida, E. granosa, E. lutea and E. suggrandis* were described from the colon of *Cubitermes* sp. of Africa (Henderson, 1941).
Genus *Entamoeba* Casagrandi and Barbagallo (1895). Nucleus vesicular, with a comparatively small endosome, located in or near the center and with varying number of peripheral nonchromatinic granules attached to the nuclear membrane (Fig. 187, b); chromatin in the endosome and in peri-endosomal region. The genus was established by the two Italian authors who were unaware of the existence of the genus *Endamoeba* (p. 444). Numerous species in vertebrates and invertebrates; one species in Protozoa.

![Fig. 189. a, Endamoeba majestus, ×420 (Kirby); b, E. simulans, ×420 (Kirby); c, Entamoeba paulista in Zelleriella, ×290 (Stabler and Chen).](image)

*E. histolytica* Schaudinn (1903) (Figs. 190, 191). The trophozoite is an active amoeba and measures 7–35 (9–20) μ in diameter; cytoplasm usually well differentiated; eruptive formation of large lobopodia, composed largely of ectoplasm; when fresh, active monopodal progressive movement; the vesicular nucleus appears in life as a ring, difficult to recognize; food vacuoles contain erythrocytes, tissue cell fragments, leucocytes, etc.; stained nucleus shows a membrane, comparatively small peripheral granules, a centrally located small endosome and an indistinct network with a few scattered chromatin granules. The trophozoite multiplies by binary fission. The amoeba lives in the lumen and in the tissues of the wall of the colon, and brings about characteristic ulceration of the colon which is typically accompanied by symptoms of *amoebic dysentery*. Through the portal vein, the amoeba may invade the liver in which it produces abscess, and other organs such as lung, brain, testis, etc. The infection in these organs is referred to as *amoebiasis*.

Under certain circumstances not well understood, the amoebae remain small after division. Such amoebae are sluggish and known
as the precystic forms. The precystic amoeba secretes presently a resistant wall and becomes encysted. The highly refractile cyst is spherical and measures 5–20\(\mu\) in diameter. At first it contains a single nucleus which divides twice. The mature cyst contains four nuclei. In addition the cyst contains diffused glycogen and elongated refractile rod-like bodies with rounded extremities which stain deeply with haematoxylin (hence called chromatoïd bodies). These inclusions are absorbed and disappear as the cyst matures. No further changes

![Figure 190](image)

**Fig. 190. Entamoeba histolytica, \(\times1150\) (Kudo).** 1, a living trophozoite; 2–4, stained trophozoites; 5, a fresh cyst; 6–9, stained cysts.

take place in the cyst as long as it remains outside the host’s intestine. The trophozoites are found in dysenteric or diarrhoeic faeces, but formed faeces usually contain cysts. The cyst is the stage by which the organism begins its life in a new host.

The life-cycle of *Entamoeba histolytica* in human host is unknown. The amoeba has, however, been cultivated in vitro by numerous investigators since the first successful cultivation by Boeck and Drbohlav (1924) (p. 887). The excystment of cysts and metacystic development have also been observed and studied especially by Dobell (1928) and Cleveland and Sanders (1930) in cultures. Snyder
and Meleney (1941) found that bacteria-free cysts usually excyst when suspended in various media with living bacteria and in the absence of bacteria, excystment was observed only in the presence of the reducing agents, cysteine or neutralized thioglycollic acid or under conditions of reduced oxygen tension. According to Dobell, in the process of excystation, a single tetranucleate amoeba emerges from a cyst through a minute pore in the cyst wall. The tetranucleate metacystic amoeba produces a new generation of trophozoites by a diverse series of nuclear and cytoplasmic divisions (Fig. 191) which result in production of eight uninucleate amoebulae. These amoebulae are young trophozoites which grow into larger ones. No sexual phenomena have been observed during these changes. It is supposed that when viable cysts reach the lower portion of the small intestine or the colon, the changes stated above take place in the lumen and the young uninucleate amoebulae initiate an infection.

Fig. 191. Diagram showing excystment and a common way by which a metacystic amoeba of *Entamoeba histolytica* divides into 8 uninucleate amoebulae (Dobell).
While the description of *Entamoeba histolytica* given above applies in general, diversities in dimensions of trophozoites and cysts, and in pathogenicity in human host as well as in experimental animals have been reported. A number of observers are inclined to think that there are several varieties or races of this amoeba, as has already been mentioned (p. 226).

*Entamoeba histolytica*, commonly known as "the dysentery amoeba," was first definitely recognized by L"osch in Russia in 1875. It is now known to be widely distributed in tropical, subtropical and temperate regions alike, although it is more prevalent in warmer regions. The incidence of infection depends mainly on the sanitary conditions of the community, since the cysts of the organism are voided from host in faeces. Faecal examinations which have been carried on by numerous investigators in different countries of the world, reveal that the incidence of infection is as high as over 50 per cent in some areas. According to Craig (1934), 49,336 examinations made by many observers in various parts of the United States show that the infection rate varied from 0.2 to 53 per cent, averaging 11.6 per cent, which justifies Craig's (1926) earlier estimate that about 10 per cent of the general population harbor this protozoan. An acute infection by *E. histolytica* is accompanied by dysentery, while in chronic cases or in convalescence, the host may void infectious cysts without suffering from the infection himself. Such a person is known as a cyst-carrier or -passer.

The trophozoite if voided in faeces perish in a comparatively short time. The dissemination of infection is thus exclusively carried on by the cyst. Viable cysts may be transmitted (1) by contamination of food through contact with contaminated water or through unsanitary habit of food handlers who are cyst-carriers; (2) by droppings of flies and cockroaches which, as noted below, contain viable cysts for a comparatively long time after feeding on faeces containing cysts and by soiled appendages of these insects which may directly transfer the cysts to food by walking on it; and (3) by contaminated water in which the cysts live considerably longer than in faeces (p. 450).

The seriousness of water-borne infection in crowded areas is easily realized when one recalls the outbreak (some 1400 cases) of amoebic dysentery and amoebiasis which originated in Chicago in 1933, where defective plumbing in certain establishments contaminated the water system with the cysts of *Entamoeba histolytica* (Bundesen et al., 1936) and the development of some 100 cases of amoebic dysentery among firemen who drank contaminated water in connection with the 1934 fire of the Union Stockyards in Chicago (Hardy and Spector), although in the latter instance, some workers believe that se-
vere amoebic infections may have resulted from already existing dormant infections aided by the newly formed association with bacteria.

The cysts remain viable for a considerable length of time outside the human intestine, if environmental conditions are favorable. Since information regarding the viability and longevity of the cyst is highly important from the epidemiological standpoint, many papers have dealt with it. In testing the viability of the cyst, the following two tests have been used by the majority of investigators.

(a) Eosin-staining test. Kuenen and Swellengrebel (1913) first used a dilute solution of eosin (1:1000). It has since been used by Wenyon and O'Connor, Root, Boeck, and many others. Solutions used vary from 1:10,000 (Root) to 1:100 (Boeck). A small amount of fresh cyst-containing material and a drop of eosin solution are mixed on a slide, then dead cysts will appear stained reddish under the microscope, while living cysts remain unstained. Whether or not unstained cysts might be dead or uninfectious is unknown. But as Wenyon and O'Connor wrote, "if we accept the eosin test as a criterion and regard all unstained cysts as living, the error in judgment will be on the safe side." Root found neutral red in 1:10,000 dilution to give a slightly larger proportion of stained cysts than eosin. Frye and Meleney's (1936) comparative study leads one to look upon this method as a fairly dependable one.

(b) Cultivation test. Improved cultural technique now brings about easily excystment of viable cysts in a proper culture medium. For example, Yorke and Adams (1926) obtained in 24 hours "a plentiful growth of vegetative forms" from cysts in Locke-egg-serum medium (p. 887). Snyder and Meleney (1941) note recently that the excystation does not take place in various culture media unless living bacteria were added or oxygen concentration of the media was decreased. Animal infection method has not been used much, as experimental animals (cats) show individual difference in susceptibility. Some of the published results are summarized below. The testing method used is indicated by: $a$ for eosin test or $b$ for cultivation test and is given after the name of the investigators.

1. Cysts in faeces kept in a covered container. All cysts disappeared in 3 days at 37°C.; at 27–30°C. half of the cysts found dead by the 4th and all dead by the 9th day (Kuenen and Swellengrebel; $a$). Alive for 3 weeks (Thomson and Thomson; $a$). Remain unchanged for several weeks if kept "cool and moist" (Dobell). All dead within 10 days at 16–20°C or 0°C. (Yorke and Adams; $b$).

2. Cysts kept in water emulsion. All alive on the 9th, but almost
all dead on the 13th day (Kuenen and Swellengrebel; a). Viable for 25 days (Thomson and Thomson; a). Cysts in running water for 15 days, excysted in pancreatic juice (Penfold, Woodcock and Drew). Viable for 30 days (Wenyon and O’Connor; a); for 5 weeks (Dobell); for 153 days (Boeck; a). Alive for 10 and 17 days at 16-20° and 0°C. respectively (Yorke and Adams; b); for 3, 10, 30, and 90 days at 30°, 20°, 10° and 0°C. respectively (Chang and Fair; b).

3. Cysts in relation to high temperatures. Cysts are killed at 68°C. in 5 minutes (Boeck; a); at 50°C. in 5 minutes (Yorke and Adams; b). Dipping in boiling water for 5-10 seconds kills the cysts (Kessel; a).

4. Cysts in relation to desiccation. Desiccation kills cysts instantly (Kuenen and Swellengrebel; Wenyon and O’Connor, Dobell, etc.). Therefore, the cysts carried in dust are most probably not viable under ordinary circumstances.

5. Cysts in relation to chemicals.

\( \text{HgCl}_2 \). 0.1% solution kills cysts in 4 hours (Kuenen and Swellengrebel; a); kills readily (Lin; b). 1:2500 solution kills cysts in 30 minutes at 20–25°C. (Yorke and Adams; b).

Creolin. 1:250 solution kills cysts in 5–10 minutes (Kuenen and Swellengrebel; a).

Alcohol. 50% alcohol kills cysts immediately (Kuenen and Swellengrebel; a); in one hour (Kessel; a).

Formaldehyde. Cysts treated in 1% solution for 4 hours were apparently dead, though not stained with eosin (Wenyon and O’Connor). 0.5% solution kills cysts in 30 minutes at 20–25° or 37°C. (Yorke and Adams; b).

Cresol. 1:20, 1:30, and 1:100, killed the cysts immediately, in one minute and in 30 minutes respectively (Wenyon and O’Connor; a).

Phenol. 1:40 and 1:100 killed cysts in 15 minutes and 7 hours respectively (Wenyon and O’Connor; a). 1% solution of phenol or lysol kills cysts in 30 minutes at 20–25° or 37°C. (Yorke and Adams; b).

HCl. 7.5% solution at 20–25°C. and 5% at 37°C. kill the cysts in 30 minutes (Yorke and Adams; b).

NaOH. 2.5% solution kills cysts in 30 minutes at 20–25° or 37°C. (Yorke and Adams; b).

Chlorine. 1:10,000 solution did not have any effect on cysts after several hours (Wenyon and O’Connor; a). 0.2% and 0.5% solutions kill the cysts in 7 days and 72 hours respectively (Kessel; a). 0.5% and 1% solutions kill the cysts in
36–48 and 12–24 hours respectively (Lin; b). 1/64 of a saturated solution of chlorine (about 0.7 weight %) at 20–25°C. and 1/320 solution at 37°C. killed the cysts in 30 minutes (Yorke and Adams; b). Exposure to the residual chlorine 5, 8 and even 10 parts per million for 30 minutes allowed cysts to remain viable (Becker et al.). Thus the cysts of *E. histolytica* are resistant to chlorinated water far above the concentration which is used ordinarily in water treatment.

Potassium permanganate. 2% solution kills the cysts in 3 days (Kessel; a). 1:500 solution kills cysts in 24–48 hours (Lin; b). 1% solution does not kill cysts at 20–25° or 37°C. in 30 minutes (Yorke and Adams; b).

Emetin hydrochloride and yetren. 5% solutions of the two drugs did not have any effects upon cysts at 20–25° or 37°C. in 30 minutes (Yorke and Adams; b).

**Antibiotics.** The majority of antibiotics appear to inhibit the growth of bacteria, which results in the death of the amoeba in culture. Prodigiosin, however, according to Balamuth and Brent (1950), kills the amoebae when added in the dilution of 1:400,000, while bacterial flora, oxidation-reduction potentials and pH are not affected by it.

6. Cysts in relation to passage through the intestine of insects. Wenyon and O’Connor found that the cysts of *E. histolytica* survived as long as 24 hours in the intestines of flies, *Musca domestica*, Calliphora, and Lucilia, and living cysts were voided for 16 hours after feeding on faecal material containing cysts. Roubaud using *Musca domestica*, found also unaltered cysts for over 24 hours (but rarely after 40 hours) after taking the cysts in its gut, and if a fly drowned in water, the cysts remained viable for about a week. Root (1921) using *Musca domestica*, Calliphora erythrocephala (and *Fannia canicularis*, Lucilia caesar, and *Chrysomyia macellaria*) found that about half the cysts were dead after 15 hours and last living cysts were found after 49 hours in the intestines of these flies after feeding on cyst-containing material, and that when the flies which ingested cysts were drowned in water, about half the cysts were found dead in 3 days and last living cysts were noticed on the 7th day. Frye and Meleney (1932) found cysts in the intestines of flies which were caught in 4 of 12 houses where infected subjects lived.

Macfie (1922) reported that the cysts of *Entamoeba histolytica* he observed in the intestine of *Periplaneta americana* appeared un-
harmed. Tejera (1926) reports successful experimental infection in two kittens that were fed on the droppings of cockroaches (sp.? caught in a kitchen, which contained cysts resembling those of *E. histolytica*. Frye and Melaney (1936) observed that the cysts passed through the intestine of *Periplaneta americana* in as early as 10–12 hours and remained in the intestine for as long as 72 hours, after feeding on experimental material. Cysts which stayed in the cockroach intestine for 48 hours gave good cultures of trophozoites in egg-horse-serum-Ringer medium.

In addition to *E. histolytica*, there are now known four other intestinal amoebae living in man. They are *E. coli*, *Endolimax nana*, *Iodamoeba bütschlii* and *Dientamoeba fragilis*. In Table 10 are given the characteristics necessary for distinguishing *E. histolytica* from the other four intestinal amoebae.

*E. coli* (Grassi) (Fig. 192). The trophozoite measures 15–40μ in diameter; average individuals 20–35μ; cytoplasm not well differentiated; movement sluggish; endoplasm granulated, contains microorganisms and faecal debris of various sizes in food vacuoles; erythrocytes are not ingested, though in a few cases (Tyzzer and Geiman)
Table 10.—Differential diagnosis of the intestinal amoebae of man

<table>
<thead>
<tr>
<th>Trophezoite</th>
<th>Entamoeba histolytica</th>
<th>Entamoeba coli</th>
<th>Iodamoeba bütschlii</th>
<th>Endolimax nana</th>
<th>Dientamoeba fragilis</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Living specimens</td>
<td>7–35µ</td>
<td>10–40µ</td>
<td>6–25µ</td>
<td>6–18µ</td>
<td>4–18µ</td>
</tr>
<tr>
<td>a. Diameter</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>b. Movement</td>
<td>Active progressive movement; eruptive formation of pseudopodia</td>
<td>Less active amoeboid movement</td>
<td>Less active amoeboid movement</td>
<td>Progressive movement</td>
<td>Progressive movement</td>
</tr>
<tr>
<td>c. Cytoplasm</td>
<td>Hyaline; erythrocytes, tissue cells, taken in as food</td>
<td>Granulated; bacteria, yeasts, faecal debris in food vacuoles</td>
<td>Granulated; bacteria in food vacuoles</td>
<td>Hyaline; bacteria in food vacuoles</td>
<td>Hyaline; bacteria in food vacuoles</td>
</tr>
<tr>
<td>d. Nucleus</td>
<td>Faintly visible ring</td>
<td>Ring of coarse granules</td>
<td>Faintly seen</td>
<td>Rarely seen</td>
<td>Faintly seen</td>
</tr>
<tr>
<td>2. Stained specimens</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>a. Nucleus</td>
<td>Fig. 190, 2–4</td>
<td>Fig. 192, 2–5</td>
<td>Fig. 195, 2–5</td>
<td>Fig. 196, b</td>
<td>Fig. 197, c, d</td>
</tr>
<tr>
<td>b. Inclusions</td>
<td>Erythrocytes, fragments of tissue cells</td>
<td>Bacteria, faecal debris, etc.</td>
<td>Bacteria</td>
<td>Bacteria</td>
<td></td>
</tr>
<tr>
<td>Cyst</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1. Living specimens</td>
<td>Spherical; circular in outline</td>
<td>Circular, often oval</td>
<td>Of various forms</td>
<td>Often oval to ellipsoid</td>
<td></td>
</tr>
<tr>
<td>a. Form</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>b. Diameter</td>
<td>5–20µ</td>
<td>10–30µ</td>
<td>6–15µ</td>
<td>5–12µ</td>
<td></td>
</tr>
<tr>
<td>2. Lugol-treated specimens</td>
<td>Cytoplasm greenish yellow; glycogen diffused; 1, 2, or 4 nuclei</td>
<td>Cytoplasm yellowish brown; glycogen body often big; 1, 2, 4, or 8 nuclei</td>
<td>Cytoplasm yellowish; large glycogen body sharply outlined; 1 nucleus</td>
<td>Cytoplasm greenish yellow; glycogen scanty, diffused; 1, 2, or 4 nuclei</td>
<td></td>
</tr>
<tr>
<td>3. Stained specimens</td>
<td>1, 2, or 4 nuclei; chromatoid bodies with rounded ends</td>
<td>1, 2, 4, or 8 nuclei; chromatoid bodies few, acicular or irregular with pointed ends</td>
<td>One nucleus; conspicuous glycogen vacuole; no chromatoid body</td>
<td>1, 2, or 4 nuclei; chromatoid bodies very small if present</td>
<td></td>
</tr>
</tbody>
</table>
and in culture (Dobell, etc.), they may be taken in as food particles (see below); nucleus, 5–8μ in diameter, seen in vivo; compared with *E. histolytica*, the endosome is somewhat large (about 1μ in diameter) and located eccentrically; peripheral granules more conspicuous. The precystic form, 10–30μ in diameter, resembles that of *E. histolytica*. Separation of the two species of amoebae by this stage is ordinarily impossible.

The cyst is spherical or often ovoid, highly refractile; 10–30μ in diameter; immature cyst contains 1, 2 or 4 nuclei, one or more large glycogen bodies with distinct outlines, but comparatively small number of acicular, filamentous or irregular chromatoid bodies with sharply pointed extremities; when mature the cyst contains 8 nuclei and a few or no chromatoid bodies. The trophozoites and small number of cysts occur in diarrhoeic or semiformed faeces and the formed faeces contain mostly cysts.

This amoeba lives in the lumen of the colon and does not enter the tissues of the wall. As noted above, it has been observed in a few instances to ingest erythrocytes, but there is no evidence to show that it takes them in from living tissues. This amoeba is therefore considered a commensal. The abundant occurrence of the trophozoite in diarrhoeic faeces is to be looked upon as a result and not the cause of the intestinal disturbance. This amoeba is of common occurrence and widely distributed throughout the world.

Nothing is known about its life-cycle in the human intestine. Cultivation of cysts in vitro indicates, according to Dobell (1938), the following changes: The cyst content usually emerges as a single multinucleate amoeba through a large opening in the cyst wall. Prior to or during the emergence, the amoeba may divide. Normal mature cysts "frequently lose" 1–4 of their original 8 nuclei before germination, thus becoming "infranucleate" (with 4–7 nuclei). Unlike in *E. histolytica*, there is no nuclear division in the metacystic stages. By a series of binary divisions with random nuclear distribution, uninucleate amoebulae are finally produced. These are young amoebae which develop into large trophozoites. Here also, there is no sexual phenomenon in the life-cycle. Nomenclature and morphology (Dobell, 1919, 1938).

*E. gingivalis* (Gros) (*E. buccalis* Prowazek) (Fig. 193). This amoeba lives in carious teeth, in tartar and debris accumulated around the roots of teeth, and in abscesses of gums, tonsils, etc. The trophozoite is as active as that of *E. histolytica*; 8–30μ (average 10–20μ) in diameter; cytoplasm well differentiated; monopodal progressive movement in some individuals; endoplasm hyaline, but
vacuolated, and contains ordinarily a large number of pale greenish bodies (which are probably nuclei of leucocytes, pus cells or other degenerating host cells) and bacteria in food vacuoles; nucleus, 2–4μ in diameter, appears as a ring; when stained it shows a small central endosome and small peripheral granules closely attached to the membrane. Stabler (1940) observed 5 chromosomes during binary fission. Encysted forms have not been observed in this amoeba. Kofoid and Johnstone (1930) reported having seen the same organism in the mouth of monkeys (Rhesus and Cynomolgus) from southeast Asia.

*E. gingivalis* is the very first parasitic amoeba that has become known to man. Gros (1849) found it in Russia in the tartar on the surface of the teeth. Some observers maintain that this amoeba is the cause of pyorrhoea alveolaris, but evidence for such an assumption seems to be still lacking. It has been found in the healthy gums and even in false teeth (Lynch). Therefore, it is generally considered as a commensal. It is widely distributed and of common occurrence.

In the absence of the encysted stage, it is supposed that the organism is transmitted in trophic forms. According to Koch (1927) who studied the effects of desiccation and temperatures upon the amoeba in culture, the amoeba is killed at 0°C. in 18 hours, at 5°C. in 24 hours, at 10°C. in 48 hours, at 45°C. in 20 minutes, at 50°C. in 15 minutes, and at 55°C. in 2 minutes. At 40°C., the survival is said to be for an indefinite length of time. Complete desiccation of the culture medium or immersion in water at 60°C. kills the amoeba. She
considered that *E. gingivalis* may be disseminated both by direct contact and by intermediate contaminated articles. Nuclear division (Stabler, 1940; Noble, 1947).

*E. gedoelsti* Hsiung (*E. intestinalis* (Gedoelst)). In the colon and caecum of horse; 6–13μ by 6–11μ; endosome eccentric; bacteria-feeder.

*E. equi* Fantham. 40–50μ by 23–29μ; nucleus oval; cysts tetranucleate, 15–24μ in diameter; seen in the faeces of horse; Fantham reports that the endoplasm contained erythrocytes.

*E. gedoelsli* Hsiung (*E. intestinalis* (Gedoelst)). In the colon and caecum of horse; 6–13μ by 6–11μ; endosome eccentric; bacteria-feeder.

*E. equi* Fantham. 40–50μ by 23–29μ; nucleus oval; cysts tetranucleate, 15–24μ in diameter; seen in the faeces of horse; Fantham reports that the endoplasm contained erythrocytes.


*E. ovis* Swellengrebel. Cyst uninucleate; in the intestine of sheep.

*E. caprae* Fantham. In goat intestine.

*E. polecki* (Prowazek). In the colon of pigs; 10–12μ in diameter; cyst uninucleate, 5–11μ in diameter.

*E. debliecki* Nieschulz (Fig. 194, a). 5–10μ in diameter; cysts uninucleate; in the intestine of pigs and goats. Two races (Hoare, 1940); morphology (Nieschulz, 1924); Entamoebae of domestic animals (Noble and Noble, 1952).

*E. venaticum* Darling. In the colon of dog; similar to *E. histolytica*; since the dog is experimentally infected with the latter, this amoeba discovered from spontaneous amoebic dysentery cases of dogs, in one of which were noted abscesses of liver, is probably *E. histolytica*.

*E. cuniculi* Brug. Similar to *E. coli* in both trophic and encysted stages; in the intestine of rabbits.

*E. cobayae* Walker (*E. caviae* Chatton). Similar to *E. coli*; in the intestine of guinea-pigs (Nie, 1950).

*E. muris* (Grassi) (Fig. 194, b, c). In the caecum of rats and mice; trophozoite 8–30μ; cytoplasm with rod-shaped or fusiform bacteria and flagellates cohabiting the host’s organ; nucleus 3–9μ in diameter and resembles closely that of *E. coli*; cysts 9–20μ in diameter, with eight nuclei when mature. Nuclear division (Wenrich, 1940); food habits (Wenrich, 1941).

*E. citelli* Becker (Fig. 194, d, e). In the caecum and colon of the striped ground squirrel, *Citellus tridecemlineatus*; rounded trophozoites 10–25μ in diameter; nucleus 4–6μ in diameter, with a comparatively large endosome which varies in position from central to peripheral; cysts with eight nuclei, about 15μ in diameter.

*E. gallinarum* Tyzzer. In the caeca of chicken, turkeys and possibly other fowls; trophozoites 9–25 (16–18)μ; cysts octonucleate, 15μ by 12μ.
E. testudinis Hartmann. In intestine of turtles, Testudo graeca, T. argentina, T. calcarata and Terrapene carolina.

E. barreti (Taliaferro and Holmes) (Fig. 194, f). In the colon of snapping turtle, Chelydra serpentina; trophozoites 14–23 (18)μ long. Cultivation (Barret and Smith, 1924).

E. terrapinae Sanders and Cleveland (Fig. 194, g, h). Trophozoites 10–15μ long; cysts 8–14μ in diameter, tetranucleate when mature;

![Figure 194](image)

upon excystment, the cyst content divides into four uninucleate amoebulae; in the colon of Chrysemys elegans (Sanders and Cleveland, 1930).

E. invadens Rodhain (Figs. 2, a, b; 194, i, j). Resembles E. histolytica. Trophozoites measure 15.9μ in average diameter (9.2–38.6μ by 9–30μ); active locomotion; feed on leucocytes, liver cells, epithelial cell debris, bacteria, etc.; nucleus similar to that of E. histolytica. Cysts 13.9μ (11–20μ) in diameter; 1–4 nuclei; glycogen vacuole; chromatoid bodies acicular, rod-like or cylindrical.

Hosts include various reptiles: Varanus salvator, V. varius, Tiliqua scincoides, Pseudoboa clelia, Lampropeltis getulus, Ancis-
amoebina

trodon mokasen, *Natrix rhombifer*, *N. sipedon* *sipedon*, *N. cyclopion*, *Python sebae*, *Rachidelus brazili*, etc. Zoological Gardens in Philadelphia (Geiman and Ratcliffe) and Antwerp (Rodhain).

The amoeba produces lesions in the stomach, duodenum, ileum, colon and liver in host animals. Time for excystation in host's intestine (jejunum and ileum) five to 14 hours; time for metacystic development in host's intestine seven-24 hours; the excysted amoeba with four nuclei, each of which divides once, divides finally into eight amoebulae; optimum temperature for culture 20-30°C. (Geiman and Ratcliffe, 1936).

Ratcliffe and Geiman (1938) observed spontaneous and experimental amoebiasis in 32 reptiles. *E. ranarum* (Grassi). In colon of various species of frogs; resembles *E. histolytica*; 10–50μ in diameter; cysts are usually tetranucleate, but some contain as many as 16 nuclei; amoebic abscess of the liver was reported in one frog. Comparison with *E. histolytica* (Dobell, 1918); life cycle (Sanders, 1931).

*E. (?) phallusiae* Mackinnon and Ray. In the intestine of the ascidian, *Phallusia mamillata*; 15–30μ by 10–15μ; nucleus about 5μ in diameter, structure not well defined; cysts uninucleate, about 20μ in diameter; parasitic nutrition.

*E. minchini* Mackinnon. In gut of tipulid larvae; 5–30μ in diameter; cyst nuclei up to 10 in number.

*E. apis* Fantham and Porter. In *Apis mellifica*; similar to *E. coli*.

*E. thomsoni* Lucas. In the colon of cockroaches; when rounded 7–30 (15–25)μ in diameter; usually attached to debris by a knob-like process, highly adhesive; cytoplasm poorly differentiated; vesicular nucleus with peripheral granules; endosome variable, with loosely aggregated granules and a central dot; cysts 8–16μ in diameter, with one to four nuclei (Lucas, 1927).

*E. aulastomi* Nöller. In the gut of the horse-leech, *Haemopis sanguisuga*; cysts with four nuclei. Morphology and development (Bishop, 1932).

*E. paulista* (Carini) (*Brumptina paulista* C.) (Fig. 189, c). In the cytoplasm of many species of Protociliata; trophozoites 5.3–14.3μ in diameter; cysts about 9.4μ in diameter, uninucleate; no effect upon host ciliates even in case of heavy infection (Stabler and Chen, 1936; Chen and Stabler, 1936). Carini and Reichenow (1935); trophozoites 8–14μ in diameter; cysts 8–12μ; either identical with *E. ranarum* or a race derived from it.

Genus *Iodamoeba* Dobell. Vesicular nucleus, with a large endosome rich in chromatin, a layer of globules which surrounds the endosome and do not stain deeply, and achromatic strands between
the endosome and membrane (Fig. 187, c); cysts ordinarily uninucleate, contain a large glycogenous vacuole which stains conspicuously with iodine; in intestine of man and mammals (Dobell, 1919).

*I. bütschlii* (Prowazek) (*I. williamsi* P.) (Fig. 195). The trophozoite is 6–25μ (average 8–15μ) in diameter; fairly active with progressive movement, when fresh; cytoplasm not well differentiated; endoplasm granulated, contains bacteria and yeasts in food vacuoles; the nucleus (3–4μ in diameter) visible in vivo; the large endosome about ½ the diameter of nucleus, surrounded by small spherules.

Fig. 195. Iodamoeba bütschlii, ×1150 (Kudo). 1, a living amoeba; 2–5, stained trophozoites; 4, 5, somewhat degenerating trophozoites; 6, a fresh cyst; 7–10, stained cysts.

The cysts are spherical, ovoid, ellipsoid, triangular, pyriform or square; rounded cysts measure about 6–15μ in the largest diameter; a large glycogen body which becomes conspicuously stained with Lugol’s solution (hence formerly called “iodine cysts”) persists; nucleus with a large, usually eccentric endosome.

The trophozoites and cysts are ordinarily present in diarrhoeic faeces, while the formed faeces contain cysts only. This amoeba apparently lives in the lumen of the colon and does not seem to attack host’s tissues and is, therefore, considered to be a commensal. Nomenclature (Dobell, 1919); nuclear structure (Wenrich, 1937a).

*I. suis* O’Connor. In colon of pig; widely distributed; indistinguishable from *I. bütschlii*; it is considered by some that pigs are probably reservoir host of *I. bütschlii*.

Genus *Endolimax* Kuenen and Swellengrebel. Small; vesicular nucleus with a comparatively large irregularly shaped endosome,
composed of chromatin granules embedded in an achromatic ground mass and several achromatic threads connecting the endosome with membrane (Fig. 187, d); commensal in hindgut in man and animals. Several species.

*E. nana* (Wenyon and O'Connor) (Fig. 196, a–d). The trophozoite measures 6–18 μ in diameter; fairly active monopodal movement by forming a broad pseudopodium; when stationary pseudopodia are formed at different points; endoplasm is granulated and contains bacteria as food particles; the vesicular nucleus, 1.5–3 μ in diameter, is composed of a delicate membrane with a few chromatin granules and a large irregularly shaped endosome.

The cyst is usually ovoid; young cyst contains 1 or 2 nuclei; mature cyst with 4 nuclei; indistinctly outlined glycogen body may be present while immature; dimensions 5–12 μ (majority 7–10 μ) in diameter.

The trophozoites are found in diarrhoeic or semifluid faeces together with the cysts, and formed faeces contain cysts. This amoeba is coelozoic in the lumen of the upper portion of colon and is considered to be a commensal. Cytology and life-history (Dobell, 1943).

E. gregariniformis (Tyzzer). In the caeca of fowls; 4–12μ in diameter; cysts uninucleate (Tyzzer, 1920).

E. clevelandi Gutierrez-Ballesteros and Wenrich (Fig. 196, e, f). In the rectal contents of Pseudemys floridana mobilensis; trophozoites 5–14μ in diameter; cysts tetranucleate, 4.5–10μ large.

E. ranarum Epstein and Illovaisky. In the colon of frogs; cysts octonucleate, up to 25μ in diameter.

E. blattae Lucas. In the colon of cockroaches; trophozoites 3–15μ long; cysts, 7–11μ in diameter and with one to three nuclei (Lucas, 1927).

Genus Dientamoeba Jepps and Dobell. Small amoeba; number of binucleate trophozoites often greater than that of uninucleate forms; nuclear membrane delicate; endosome consists of several chromatin granules embedded in plasmosomic substances and connected with the membrane by delicate strands (Fig. 187, e); in colon of man (Jepps and Dobell, 1918).

![Fig. 197. Dientamoeba fragilis, ×2300 (Kudo). a, b, living bi- and uni-nucleate trophozoites; c, d, stained uni- and bi-nucleate trophozoites.](image)

D. fragilis J. and D. (Fig. 197). The trophozoite is actively amoeboid; 4–18μ (average 5–12μ) in diameter; progressive movement; cytoplasm well differentiated; endoplasm granulated contains bacteria in food vacuoles; nucleus only faintly visible; 1 or 2 nuclei, the ratio is variable; in some material binucleate forms may be 80% or more (Jepps and Dobell), while in others uninucleate forms may predominate (Kudo, 1926a; Wenrich, 1937); nucleus is made up of a delicate membrane and a large endosome (more than one-half the diameter of nucleus) in which are embedded 4–8 chromatin granules along the periphery. According to Dobell (1940), the binucleate condition represents an arrested telophase stage of mitosis and the chromatin granules are in reality chromosomes, probably 6 in number. Comparison with Histomonas meleagridis (p. 335) led this author to think that this amoeba may be an aberrant flagellate closely related to Histomonas.
Encysted stage has not been observed. Degenerating trophozoites often develop vacuoles which coalesce into a large one and the organisms may then resemble *Blastocystis hominis* (p. 893) which is very common in faeces. Transmission may be carried on by trophozoites. According to Wenrich (1940), this amoeba if left in the faeces remains alive up to 48 hours at room temperature, but disappears probably by disintegration in 2 hours at 3.5°C. Since all attempts to bring about experimental infection by mouth or by rectum failed, Dobell considered that the amoeba may be transmitted from host to host in the eggs of nematodes such as *Trichuris* or *Ascaris*, as in the case of *Histomonas* (p. 335).

The amoeba inhabits the lumen of the colon. There is no indication that it is histozoic or cytozoic. Some workers attribute certain intestinal disturbances to this amoeba, but no definite evidence for its pathogenicity is available at present. It seems to be widely distributed, but not as common as the other intestinal amoebae mentioned above, although in some areas it appears to be common. Nuclear division (Wenrich, 1936, 1939, 1944a; Dobell, 1940).

Genus *Martinezia* Hegner and Hewitt. The nucleus consists of a wrinkled membrane, a large compact or granular endosome and heavy peripheral beads; cysts unknown; parasitic.

*M. baezi* H. and H. (Fig. 196, g, h). In the intestine of iguanas, *Ctenosaura acaenthura*; 8–21 μ by 6.5–16 μ; nucleus about 4 μ in diameter; two nuclei in about 3 per cent of the organisms; cysts not seen.

Genus *Dobellina* Bishop and Tate. Trophozoite: small amoeba; ectoplasm and endoplasm differentiated; usually monopodal; nucleus one to many; nucleus with a large central endosome and an achromatic nuclear membrane; nuclear divisions mitotic and simultaneous; no solid food vacuoles; no contractile vacuole; with refringent granules. Cysts: spherical; thin-walled; devoid of glycogen and of chromatoid bodies; 2 or more nuclei; parasitic (Bishop and Tate, 1939).

*D. mesnili* (Keilin) (Fig. 198, a–c). Uninucleate amoebae as small as 3.6 μ in diameter; multinucleate forms 20–25 μ by 10–15 μ; cysts 8–11 μ in diameter; in the space between the peritrophic membrane and the epithelium of the gut in the larvae of *Trichocera hiemalis*, *T. annulata*, and *T. regelationis* (winter gnats).

Genus *Schizamoeba* Davis. Nucleus vesicular, without endosome, but with large discoid granules arranged along nuclear membrane; 1 to many nuclei; cyst-nuclei formed by fragmentation of those of the trophozoite and possess a large rounded chromatic endosome, connected at one side with the nuclear membrane by achromatic
strands to which chromatin granules are attached; in stomach of salmonid fish. One species (Davis, 1926).

*Salmonis* D. (Fig. 198, d, e). Sluggish amoeba; 10–25μ in diameter; 1 to several nuclei; multiplication by binary fission; nuclear division amitotic. Cyst are said to be more abundant than trophozoites and their appearance seems to be correlated with the amount of available food; cysts spherical, 15–35μ in diameter; cyst-membrane thin and nuclei vary from 3 to many; during encystment, chromatin bodies of trophozoite become collected in several masses which then break up and each chromatin grain becomes the endosome of newly formed nucleus; cyst contents divide sooner or later into 4–11 multinucleate bodies and the whole increases in size; finally cyst-membrane disintegrates and the multinucleate bodies become set free. Trophozoites are said to occur in the mucous covering of stomach of host fish; cysts occur in both stomach and intestine. Aside from the loss of certain amount of available food, no pathogenic effect of the amoeba upon the host fish was noticed (Davis).

Genus *Hydramoeba* Reynolds and Looper. Nucleus vesicular with a large central endosome composed of a centriole (?) and chromatin granules embedded in an achromatic mass, achromatic strands radiating from endosome to membrane; a ring made up of numerous rod-shaped chromatin bodies in the nuclear-sap zone; 1 or more contractile vacuoles; apparently the most primitive parasitic amoeba; parasitic on Hydra.

*Hydrozoena* (Entz) (Fig. 198, f–l). Parasitic in various species of Hydra; first observed by Entz; Wermel found 90 per cent of Hydra he studied in Russia were infected by the amoeba; Reynolds and Looper (1928) stated that infected Hydra die on an average in 6.8 days and that the amoebae disappear in 4–10 days if removed from a host hydra. More or less spheroidal, with blunt pseudopods; 60–380μ in diameter; nucleus shows some 20 refractile peripheral granules in life; contractile vacuoles; food vacuoles contain host cells; multiplication by binary fission.

Ito (1949) found this organism in *Hydra japonica, H. magnipapillata, Palmatycnora robusta*, etc. in Japan. The trophozoites measured 26–210μ long with a nucleus, 10–12μ in diameter. Early infection occurs on the tip of tentacles and spreads to the body proper (Fig. 198, i–l). Since the tentacles remain contracted, the host hydra cannot feed on food organisms and becomes "depressed." The amoebae finally enter the coelenteric cavity and feed on the endoderm cells. The host hydra becomes spherical. At 25°C. death of the hydra may
occur in one week. Encystment takes place soon after the death of the host or occasionally when the organisms become detached from the host; cysts are spherical, measure 27.5–29μ, and contain one or more nuclei, nematocysts and a large vacuole (h). Nuclear division (Reynolds and Threlkeld, 1929).

Fig. 198. a–c, Dobellina mesnili (Bishop and Tate) (a, b, stained uni- and multi-nucleate trophozoites, ×2200; c, a stained cyst with six nu- clei, ×1760); d, e, stained trophozoite and cyst of Schizamoeba salmonis, ×1070 (Davis); f–l, Hydramoeba hydroxena (f, h–l, Ito; g, Reynolds and Looper) (f, a trophozoite in life, ×330; g, a trophozoite feeding on ecto- dermal cells of a Hydra in section, ×470; h, a living cyst, ×530; i–l, stages of infection in Hydra, ×6.5); m, Paramoeba pigmentifera with its nucleus in center, ×800 (Janicki).

Family 4 Paramoebidae Poche

Genus Paramoeba Schaudinn. The amoeba possesses a nucleus and nucleus-like secondary cytoplasmic structure, both of which multiply by division simultaneously; free-living or parasitic.

P. pigmentifera (Grassi) (Fig. 198, m). About 30μ long; sluggish;
cytoplasm distinctly differentiated; secondary body larger than the nucleus; flagellated swarvers are said to occur; parasitic in coelom of Chaetognatha such as Sagitta claparedi, Spadella bipunctata, S. inflata, and S. serratodentata. Cytology (Jauicki, 1928, 1932).

P. schaudinni Faria, da Cunha and Pinto. About 7–22μ in diameter; in salt water; Rio de Janeiro, Brazil.

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AMOEBINA

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Chapter 20
Order 4 Testacea Schultze

The Testacea or Thecamoeba comprise those amoeboid organisms which are enveloped by a simple shell or test, within which the body can be completely withdrawn. The shell has usually a single aperture through which pseudopodia protrude, and varies in shape and structure, although a chitinous or pseudochitinous membrane forms the basis of all. It may be thickened, as in Arcella and others, or composed of foreign bodies cemented together as in Diffugia, while in Euglypha siliceous platelets or scales are formed in the endoplasm and deposited in the shell.

The cytoplasm is ordinarily differentiated into the ectoplasm and endoplasm. The ectoplasm is conspicuously observable at the aperture of the shell where filopodia or slender ectoplasmic lobopodia are produced. The endoplasm is granulated or vacuolated and contains food vacuoles, contractile vacuoles and nuclei. In some forms there are present regularly in the cytoplasm numerous basophilic granules which are known as 'chromidia' (p. 44).

Asexual reproduction is either by longitudinal fission in the forms with thin tests, or by transverse division or budding, while in others multiple division occurs. Encystment is common. Sexual reproduction by amoeboid or flagellate gametes has been reported in some species.

The testaceans are mostly inhabitants of fresh water, but some live in salt water and others are semi-terrestrial, being found in moss or moist soil, especially peaty soil. Biology of soil-inhabiting forms (Volz, 1929); ecology (Hoogenraad, 1935).

Shell simple and membranous
Filopodia, in some anastomosing
Pseudopodia filose, simply branched

Family 1 Gromiidae
Eimer and Fickert

These forms are frequently included in the Foraminifera by other authors.

Genus Gromia Dujardin (Allogromia, Rhynchogromia, Diplogromia Rhumbler). Thin test rigid or flexible, smooth or slightly coated with foreign bodies; spherical to elongate ellipsoid; aperture
terminal; 1 or more nuclei; contractile vacuoles; many filopodia, branching and anastomosing; cytoplasm with numerous motile granules; fresh or salt water. Many species.

_G. fluvialis_ D. (Fig. 199, a). Test spherical to subspherical; smooth or sparsely covered with siliceous particles; yellowish cytoplasm fills the test; aperture not seen; a large nucleus and numerous contractile vacuoles; filopodia long, often enveloping test; 90–250μ long; on aquatic plants, in moss or soil.

_G. ovoidea_ (Rhumbler) (Fig. 199, b). In salt water.

_G. nigricans_ (Penard) (Fig. 199, c). Test large, circular in cross-section; a single nucleus; 220–400μ long; in pond water among vegetation.

Genus _Microgromia_ Hertwig and Lesser. Test small, hyaline, spherical or pyriform, not compressed; aperture terminal, circular; filopodia long straight or anastomosing, arising from a peduncle; a single nucleus and contractile vacuole; solitary or grouped. Morphology (Valkanov, 1930).

_M. socialis_ (Archer) (Fig. 199, d). Cytoplasm bluish; contractile vacuole near aperture; filopodia arise from a peduncle, attenuate, branching, anastomosing; often numerous individuals are grouped; multiplication by fission and also by swarmers; 25–35μ in diameter; among vegetation in fresh water.

Genus _Microcometes_ Cienkowski. Body globular, enclosed within a transparent, delicate, light yellowish and pliable envelope with 3–5 apertures, through which long branching filopodia extend; body protoplasm occupies about 1/2 the space of envelope; 1–2 contractile vacuoles; fresh water.

_M. paludosa_ C. (Fig. 199, e). About 16–17μ in diameter; fresh water among algae (Valkanov, 1931; Jepps, 1934).

Genus _Artodiscus_ Penard. Body globular, plastic; covered by envelope containing small grains of various kinds; nucleus eccentric; a few pseudopodia extend through pores of the envelope; movement very rapid; fresh water.

_A. saltans_ P. (Fig. 199, f). 18–23μ in diameter; fresh water.

Genus _Lieberkühnia_ Claparède and Lachmann. Test ovoidal or spherical, with or without attached foreign particles; aperture usually single, lateral or subterminal; one or more nuclei; many contractile vacuoles; pseudopodia formed from a long peduncle, reticulate, often enveloping test; fresh or salt water.

_L. wagneri_ C. and L. (Fig. 200, a). Spheroidal; aperture subterminal, oblique, flexible; cytoplasm slightly yellowish, fills the test; 80–150 vesicular nuclei; nuclei 6μ in diameter; many contractile vac-
Fig. 199. a, *Gromia fluvialis*, ×120 (Dujardin); b, *G. ovoidea*, ×50 (Schultze); c, *G. nigricans*, ×200 (Cash and Wailes); d, *Microgromia socialis*, ×170 (Cash); e, *Microcometes paludosa*, ×670? (Penard); f, *Artodiscus saltans*, ×670 (Penard); g, *Schultzella diffluens*, ×120 (Rhumbler).
uoles; pseudopodia long, anastomosing; 60–160μ long; among algae in fresh and salt water.

Genus **Diplophrys** Barker. Test thin, spherical; 2 apertures, one at each pole; cytoplasm colorless; a single nucleus; several contractile vacuoles; filopodia radiating. One species.

*D. archeri* B. (Fig. 200, b). With 1–3 colored oil droplets; pseudopodia highly attenuate, radiating, straight or branched; multiplication into 2 or 4 daughter individuals; solitary or in groups; diameter 8–20μ; on submerged plants in fresh water.

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**Fig. 200.** a, *Lieberkühnia wagneri*, ×160 (Verworn); b, *Diplophrys archeri*, ×930 (Hertwig and Lesser); c, *Lecythium hyalinum*, ×330 (Cash and Wailes); d, *Myxotheca arenilega*, ×70 (Schaudinn); e, *Dactylosaccus vermiformis*, ×15 (Rhumbler); f, *Boderia turneri* (Wright).

Genus **Lecythium** Hertwig and Lesser. Test thin, flexible, colorless; aperture elastic, terminal; colorless cytoplasm fills the test; large nucleus posterior; numerous filopodia long, branching, not anastomosing; fresh water.

*L. hyalinum* (Ehrenberg) (Fig. 200, c). Spheroidal; aperture circular with a short flexible neck; a single contractile vacuole; diameter 20–45μ; in submerged vegetation.
Genus *Schultzella* Rhumbler. Test thin, delicate, difficult to recognize in life, easily broken at any point for formation of pseudopodia which branch and anastomose; irregularly rounded; without foreign material; salt water.

*S. diffusus* (Grubler) (Fig. 199, g). Cytoplasm finely granulated; opaque, colorless; with oil droplets, vacuoles and numerous small nuclei; up to 220μ in diameter.

Genus *Myxotheca* Schaudinn. Amoeboid; spherical or hemispherical, being flattened on the attached surface; a thin pseudochitinous test with foreign bodies, especially sand grains; pseudopodia anastomosing; salt water. Nucleus (Foyn, 1936).

*M. arenilega* S. (Fig. 200, d). Test yellow, with loosely attached foreign bodies; cytoplasm bright red due to the presence of highly refractile granules; 1–2 nuclei, 39–75μ in diameter; body diameter 160–560μ.

Genus *Dactylosaccus* Rhumbler. Test sausage-shape and variously twisted; pseudopodia filiform, anastomosing; salt water.

*D. vermiformis* R. (Fig. 200, e). Test smooth; pseudopodia rise from small finger-like projections; 1–2 nuclei; body 4 mm. by 340μ; salt water.

Genus *Boderia* Wright. Body form changeable; often spherical, but usually flattened and angular; filopodia long; test extremely delicate, colorless; salt water.

*B. turneri* W. (Fig. 200, f). Body brown to orange; active cytoplasmic movement; 1–10 nuclei; multiple division(?); 1.56–6.25 mm. in diameter; in shallow water.

**Family 2 Arcellidae** Schultze

Genus *Arcella* Ehrenberg. Test transparent, chitinous, densely punctated; colorless to brown (when old); in front view circular, angular, or stellate; in profile plano-convex or semicircular; variously ornamented; aperture circular, central, inverted like a funnel; protoplasmic body does not fill the test and connected with the latter by many ectoplasmic strands; slender lobopodia, few, digitate, simple or branched; 2 or more nuclei; several contractile vacuoles; fresh water. Numerous species. Taxonomy and morphology (Deflandre, 1928); variation and heredity (Jollos, 1924).

*A. vulgaris* E. (Fig. 201, a, b). Height of test about 1/2 the diameter; dome of hemispherical test evenly convex; aperture circular, central; colorless, yellow, or brown; protoplasmic body conforms with the shape of, but does not fill, the test; lobopodia hyaline; 2 vesicular nuclei; several contractile vacuoles; test 30–100μ in dia-
meter; in the ooze and vegetation in stagnant water and also in soil. Of several varieties, two may be mentioned; var. *angulosa* (Perty), test smaller, 30–40μ in diameter, faceted, forming a 5- to 8-sided figure, with obtuse angles; var. *gibbosa* (Penard), test gibbous, surface pitted with circular depressions of uniform dimensions; 45–50μ up to 100μ in diameter.

![Fig. 201. a, b, Arcella vulgaris, ×170; ×230 (Leidy); c, A. discoides, ×170 (Leidy); d, A. mitrata, ×140 (Leidy); e, f, A. catinus. ×170 (Cash); g–i, A. dentata, ×170 (Leidy); j, k, A. artocrea, ×170 (Leidy).](image)

*A. discoides* E. (Fig. 201, c). Test circular in front view, plano-convex in profile; diameter about 3–4 times the height; test coloration and body structure similar to those of *A. vulgaris*; test 70–260μ in diameter; in fresh water.

*A. mitrata* Leidy (Fig. 201, d). Test balloon-shaped or polyhedral; height exceeds diameter of base; aperture circular, crenulated and usually everted within inverted funnel; protoplasmic body spheroidal, with ‘neck’ to aperture and cytoplasmic strands to test; 6 or more slender lobopodia; test 100–145μ high, 100–152μ in diameter; in fresh water among vegetation.

*A. catinus* Penard (Fig. 201, e, f). Test oval or quadrate, not circular, in front view; aperture oval; dome compressed; lateral margin with 6 or 8 facets; test 100–120μ in diameter and about 45μ high; fresh water among vegetation.

*A. dentata* Ehrenberg (Fig. 201, g–i). Test circular and dentate
in front view, crown-like in profile; diameter more than twice the height; aperture circular, large; colorless to brown; about 95\(\mu\) in diameter, aperture 30\(\mu\) in diameter; 15–17 spines; in the ooze of freshwater ponds.

_A. artocrea_ Leidy (Fig. 201, _j, k_). Height of test 1/4–1/2 the diameter; dome convex; surface mammillated or pitted; border of test everted and rising 1/4–1/2 the height of test; about 175\(\mu\) in diameter; fresh water.

![Illustration](image)

Genus _Pyxidicula_ Ehrenberg. Test patelliform; rigid, transparent, punctate; aperture circular, almost the entire diameter of test; cytoplasm similar to that of _Arcella_; a single nucleus; 1 or more contractile vacuoles; fresh water.

_P. operculata_ (Agardh) (Fig. 202, _a, b_). Test smooth, colorless to brown; a single vesicular nucleus; pseudopodia short, lobose or digitate; 20\(\mu\) in diameter; on vegetation.
Genus *Pseudochlamys* Claparède and Lachmann. Test discoid, flexible when young; body with a central nucleus and several contractile vacuoles.

*P. patella* C. and L. (Fig. 202, c). Young test hyaline, older one rigid and brown; often rolled up like a scroll; a short finger-like pseudopodium between folds; 40–45μ in diameter; in fresh water among vegetation, in moss and soil.

Genus *Difflugiella* Cash. Test ovoid, not compressed, flexible and transparent membrane; colorless cytoplasm fills the test, usually with chlorophyllous food material; median pseudopodia lobate or digitate with aciculate ends, while lateral pseudopods long, straight, and fine, tapering to a point; fresh water. One species.

*D. apiculata* C. (Fig. 202, d, e). About 40μ by 28μ; among vegetation.

Genus *Cryptodiffugia* Penard. Small test yellowish to brownish; Diffugia-like in general appearance, compressed; with or without foreign bodies; pseudopodia long, acutely pointed; fresh water.

*C. oviformis* P. (Fig. 202, f). Test ovoid; without foreign bodies; crown hemispherical; aperture truncate; cytoplasm with chlorophyllous food particles; 16–20μ by 12–15μ; in marshy soil.

Genus *Lesquereusia* Schlumberger. Test compressed, oval or globular in profile, narrowed at bent back; semispiral in appearance; with curved or comma-shaped rods or with sand-grains (in one species); body does not fill up the test; pseudopodia simple or branched; fresh water.

*L. spiralis* (Ehrenberg) (Fig. 202, g). Aperture circular; border distinct; cytoplasm appears pale yellow; a single nucleus; 96–188μ by 68–114μ; in marsh water.

Genus *Hyalosphenia* Stein. Test ovoid or pyriform; aperture end convex; homogeneous and hyaline, mostly compressed; crown uniformly arched; protoplasm partly filling the test; several blunt pseudopodia simple or digitate. Several species.

*H. papilio* Leidy (Fig. 202, h). Test yellowish; transparent; pyriform or oblong in front view; a minute pore on each side of crown and sometimes one also in center; aperture convex; in narrow lateral view, elongate pyriform, aperture a shallow notch; with chlorophyllous particles and oil globules; 110–140μ long; in fresh water among vegetation.

Genus *Corycia* Dujardin. Envelope extremely pliable, open at base, but when closed, sack-like; envelope changes its shape with movement and contraction of body; with or without spinous projections.
C. coronata Penard (Fig. 202, i). 6-12 spines; 140μ in diameter; in moss.
Genus Pamphagus Bailey. Test hyaline membranous, flexible; aperture small; body fills the envelope completely; spherical nucleus large; contractile vacuoles; filopodia long, delicate, branching, but not anastomosing; fresh water. Species (Hoogenraad, 1936).

P. mutabilis B. (Fig. 202, j). Envelope 40-100μ by 28-68μ.
Genus Plagiophrys Claparède and Lachmann. Envelope thin, hyaline, changeable with body form; usually elongate-oval with rounded posterior end; narrowed at other half; envelope finely punctated with a few small plates; aperture round; cytoplasm clear; nucleus large; pseudopods straight filopodia, sometimes branching; fresh water.

P. parvipunctata Penard (Fig. 202, k). Envelope 50μ long.
Genus Leptochlamys West. Test ovoid, thin transparent chitinous membrane, circular in optical section; aperture end slightly expanded with a short neck; aperture circular, often oblique; body fills test; without vacuoles; pseudopodium short, broadly expanded and sometimes cordate; fresh water.

L. ampullacea W. (Fig. 203, a). Nucleus large, posterior; with green or brown food particles; test 45-55μ by 36-40μ in diameter; aperture 15-17μ; among algae.
Genus Chlamydophrys Cienkowski. Test rigid, circular in cross-section; aperture often on drawn-out neck; body fills the test; zonal differentiation of cytoplasm distinct; nucleus vesicular; refractile waste granules; pseudopodia branching; fresh water or coprozoic. Species (Bélař, 1926); plasmogamy and division (Bélař, 1926).

C. stercorea C. (Fig. 203, k). Test 18-20μ by 12-15μ; mature cysts yellowish brown, 12-15μ in diameter; multiplication by budding; coprozoic and fresh water.
Genus Cochliopodium Hertwig and Lesser. Test thin, flexible, expansible and contractile; with or without extremely fine hair-like processes; pseudopodia blunt or pointed, but not acicular. Several species.

C. bilimbosum (Auerbach) (Fig. 203, b). Test hemispherical; pseudopodia conical with pointed ends; test 24-56μ in diameter; fresh water among algae.
Genus Amphizonella Greeff. Test membranous with a double marginal contour; inner membrane smooth, well-defined; outer serrulate; aperture inverted; a single nucleus; pseudopodia blunt, digitate, and divergent.
A. violacea G. (Fig. 203, c). Test patelliform, violet-tinted; with
chlorophyllous corpuscles and grains; sluggish; average diameter 160µ; fresh water.

Genus *Zonomyxa* Nüsslin. Test rounded pyriform, flexible, chitinous, violet-colored; endoplasm vacuolated, with chlorophyllous particles; several nuclei; pseudopodia simple, not digitate; fresh water.

*Z. violacea* N. (Fig. 203, d). A single lobular pseudopodium with acuminate end; 4 nuclei; diameter 140–160µ; actively motile forms 250µ or longer; among sphagnum.

Genus *Microcorycia* Cockerell. Test discoidal or hemispherical,
flexible, with a diaphanous continuation or fringe around periphery, being folded together or completely closed; crown of test with circular or radial ridges; body does not fill the test; 1-2 nuclei; pseudopodia lobular or digitate; fresh water. A few species.

*M. flava* (Greeff) (Fig. 203, *e, f*). Test yellowish brown; crown with few small foreign bodies; endoplasm with yellowish brown granules; 2 nuclei; contractile vacuoles; diameter 80-100μ; young individuals as small as 20μ; in moss.

Genus *Parmulina* Penard. Test ovoid, chitinoid with foreign bodies; aperture may be closed; a single nucleus; 1 or more contractile vacuoles; fresh water. A few species.

*P. cyathus* P. (Fig. 203, *g, h*). Test small, flexible; ovoid in aperture view, semicircular in profile; aperture a long, narrow slit when test is closed, but circular or elliptical when opened; 40-55μ long; in moss.

Genus *Capsellina* Penard. Test hyaline, ovoid, membranous; with or without a second outer covering; aperture long slit; a single nucleus; 1 or more contractile vacuoles; filose pseudopodia; fresh water.

*C. timida* Brown (Fig. 203, *j*). Small, ovoid; elliptical in cross-section; with many oil (?) globules; filopodium; 34μ by 25μ; in moss.

Genus *Diplochlamys* Greeff. Test hemispherical or cup-shaped, flexible with a double envelope; inner envelope a membranous sack with an elastic aperture; outer envelope with loosely attached foreign bodies; aperture large; nuclei up to 100; pseudopodia few, short, digitate or pointed; fresh water. Several species.

*D. leidyi* G. (Fig. 203, *i*). Test dark gray; inner envelope projecting beyond outer aperture; nuclei up to 20 in number; diameter 80-100μ.

Family 3 *Diffugiidae* Taránek

Genus *Diffugia* Leclerc. Test variable in shape, but generally circular in cross-section; composed of cemented quartz-sand, diatoms, and other foreign bodies; aperture terminal; often with zoochlorellae; cytoplasmic body almost fills the test; a single nucleus; many contractile vacuoles; pseudopodia cylindrical, simple or branching; end rounded or pointed; fresh water, woodland soil, etc.

*D. oblonga* Ehrenberg (*D. pyriformis* Perty) (Fig. 204, *a*). Test pyriform, flask-shaped, or ovoid; neck variable in length; fundus rounded, with occasionally 1-3 conical processes; aperture terminal, typically circular; test composed of angular sand-grains, diatoms; bright green with chlorophyllous bodies; 60-580μ by 40-240μ; in
the ooze of fresh water ponds, ditches and bogs; also in moist soil. Several varieties.

*D. urceolata* Carter (Fig. 204, b). A large ovoid, rotund test, with a short neck and a rim around aperture; 200–230μ by 150–200μ: in ditches, ponds, sphagnous swamps, etc.

![Diagram of *Difflugia oblonga*, *D. urceolata*, *D. arcula*, *D. lobostoma*, *D. constricta*, *Centropyxis aculeata*, *Campuscus cornutus*, *Cucurbitella mespiliformis*]

Fig. 204. a, *Difflugia oblonga*, ×130 (Cash); b, *D. urceolata*, ×130 (Leidy); c, d, *D. arcula*, ×170 (Leidy); e, *D. lobostoma*, ×130 (Leidy); f, *D. constricta*, ×200 (Cash); g, *Centropyxis aculeata*, ×200 (Cash); h, *Campuscus cornutus*, ×170 (Leidy); i, *Cucurbitella mespiliformis*, ×200 (Wailes).

*D. arcula* Leidy (Fig. 204, c, d). Test hemispherical, base slightly concave, but not invaginated; aperture triangular, central, trilobed; test yellowish with scattered sand-grains or diatoms; diameter 100–140μ; in sphagnous swamp, moss, soil, etc.

*D. lobostoma* L. (Fig. 204, e). Test ovoid to subspherical; aperture terminal; with 3–6 lobes; test usually composed of sand-grains, rarely with diatoms; endoplasm colorless or greenish; diameter 80–120μ; in fresh water. Sexual fusion and life cycle (Goette, 1916).

*D. constricta* (Ehrenberg) (Fig. 204, f). Test laterally ovoid, fundus more or less prolonged obliquely upward, rounded, and simple or provided with spines; soil forms generally spineless; aperture antero-inferior, large, circular or oval and its edge inverted; test composed of quartz grains; colorless to brown; cytoplasm colorless; 80–340μ long; in the ooze of ponds and in soil.
D. corona Wallich. Test ovoid to spheroid, circular in cross-section; crown broadly rounded, with a variable number of spines, aperture more or less convex in profile, central and its border multidentate or multilobate; test with fine sand-grains, opaque; cytoplasm colorless; pseudopodia numerous, long, branching or bifurcating; 180-230μ by about 150μ; in fresh water. Genetics (Jennings, 1916, 1937).

Genus Centropyxis Stein. Test circular, ovoid, or discoid; aperture eccentric, circular or ovoidal, often with a lobate border; with or without spines; cytoplasm colorless; pseudopodia digitate; fresh water. Species (Deflandre, 1929).

C. aculeata S. (Fig. 204, g). Test variable in contour and size; with 4–6 spines; opaque or semitransparent; with fine sand-grains or diatom shells; pseudopodia sometimes knotted or branching; when encysted, the body assumes a spherical form in wider part of test; granulated, colorless or with green globules; diameter 100–150μ; aperture 50–60μ in diameter.

Genus Campascus Leidy. Test retort-shaped with curved neck, rounded triangular in cross-section; aperture circular, oblique, with a thin transparent discoid collar; nucleus large; 1 or more contractile vacuoles; body does not fill the test; fresh water.

C. cornutus L. (Fig. 204, h). Test pale-yellow, retort-form; with a covering of small sand particles; triangular in cross-section; a single nucleus and contractile vacuole; filopodia straight; 110–140μ long; aperture 24–28μ in diameter; in the ooze of mountain lakes.

Genus Cucurbitella Penard. Test ovoid with sand-grains, not compressed; aperture terminal, circular, surrounded by a 4-lobed annular collar; cytoplasm grayish, with zoochlorellae; nucleus large; 1 to many contractile vacuoles; pseudopodia numerous, digitate; fresh water.

C. mespiliformis P. (Fig. 204, i). 115–140μ long; diameter 80–105μ; in the ooze or on vegetation in ponds and ditches.

Genus Plagiopyxis Penard. Test subcircular in front view; ovoid in profile; aperture linear or lunate; cytoplasm gray, with a single nucleus and a contractile vacuole; fresh water.

P. callida P. (Fig. 205, a). Test gray, yellowish, or brown; large nucleus vesicular; pseudopodia numerous, radiating, short, pointed or palmate; diameter 55–135μ; in vegetation.

Genus Pontigulasia Rhumbler. Test similar to that of Diffugia, but with a constriction of neck and internally a diaphragm made of the same substances as those of the test.

P. vas (Leidy) (Fig. 205, b). Round or ovoid test; constriction
deep and well-marked; with sand-grains and other particles; aperture terminal; 125-170µ long; fresh water ponds. Stump (1943) made a study of the nuclear division of the organism. During metaphase 8-12 "chromosomes" form a well-defined equatorial plate; average time for completion of the division was found to be 80 minutes.

Genus Phryganella Penard. Test spheroidal or ovoid, with sand-grains and minute diatom shells; aperture terminal, round; pseudopodia drawn out to a point; fresh water.

*P. acropodia* (Hertwig and Lesser) (Fig. 205, c). Test circular in

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Fig. 205. a, Plagiopyxis callida, ×200 (Wailes); b, Pontigulasia vas ×200 (Cash); c, Phryganella acropodia, ×190 (Cash); d, Bullinula indica, ×130 (Wailes); e, f, Heleopera petricola, ×190 (Cash); g, Nadinella tenella, ×400 (Penard); h, Frenzelina reniformis, ×600 (Penard); i, Amphitrema flavum, ×360 (Cash and Wailes); j, Pseudodifflugia gracilis, ×330 (Cash); k, Diaphoropodon mobile, ×270 (Cash and Wailes); l, m, Clypeolina marginata, ×330 (Cash and Wailes).
aperture view; hemispherical in profile; yellowish or brownish, semi-transparent, and covered with sand-grains and scales; in front view sharply pointed pseudopodia radiating; colorless endoplasm usually with chlorophyllous bodies; 30–50μ in diameter.

Genus Bullinula Penard. Test ellipsoidal, flattened on one face, with silicious plates; on the flattened surface, \( \infty \)-shaped aperture; a single nucleus; pseudopodia digitate or spatulate, simple or branched; fresh water. 

\( B. \text{ indica} \) P. (Fig. 205, d). Test dark brown; 120–250μ in diameter. Distribution and morphology (Hoogenraad, 1933).

Genus Heleopera Leidy. Test variously colored; fundus hemispherical, with sand-grains; surface covered with amorphous scales, often overlapping; aperture truncate, narrow, elliptic notched in narrow lateral view; a single nucleus; pseudopodia variable in number, thin digitate or branching; fresh water. Several species.

\( H. \text{ petricola} \) L. (Fig. 205, e, f). Test variable in size and color, strongly compressed; fundus rough with sand-grains of various sizes; aperture linear or elliptic, convex in front view; pseudopodia slender, branching; 80–100μ long; in boggy places.

Genus Averintzia Schouteden. Test similar to that of Heleopera, but small aperture elliptical; test thickened around aperture; fresh water. 

\( A. \text{ cyclostoma} \) (Penard). Test dark violet, with sand-grains of different sizes; elliptical in cross-section; pseudopodia unobserved; 135–180μ long; in sphagnum and aquatic plants.

Genus Nadinella Penard. Test chitinous, thin, hyaline, with foreign bodies and collar around aperture; filopodia; fresh water. 

\( N. \text{ tenella} \) P. (Fig. 205, g). 50–55μ long; fresh water lakes.

Genus Frenzelina Penard. Two envelopes, outer envelope hemispherical, thin, rigid, covered with siliceous particles; inner envelope round or ovoid, drawn out at aperture, thin, hyaline and covering the body closely; aperture round, through which a part of body with its often branching straight filopods extends; cytoplasm with diatoms, etc.; a nucleus and a contractile vacuole; fresh water. 

\( F. \text{ reniformis} \) P. (Fig. 205, h). Outer envelope 26–30μ in diameter; fresh water lakes.

Genus Amphitrema Archer. Test ovoid, symmetrical, compressed; composed of a transparent membrane, with or without adherent foreign bodies; 2 apertures at opposite poles; with zoochlorellae; nucleus central; 1 to several contractile vacuoles; straight filopodia, sparsely branched, radiating; fresh water. Several species.

\( A. \text{ flavum} \) A. (Fig. 205, i). Test brown, cylindrical with equally
rounded ends in front view; elliptical in profile; ovoid with a small central oval aperture in end view; 45–77μ by 23–45μ; in sphagnum.

Genus **Pseudodifflugia** Schlumberger. Test ovoid, usually rigid, with foreign bodies; circular or elliptical in cross-section; aperture terminal; granulated cytoplasm colorless or greyish; nucleus posterior; a contractile vacuole; filopodia long, straight or branching; fresh water. Several species.

*P. gracilis* S. (Fig. 205, j). Test yellowish or brownish; subspherical, with sand-grains; aperture without neck; 20–65μ long.

Genus **Diaphoropodon** Archer. Test ovoid, flexible, with minute foreign bodies and a thick covering of hyaline hair-like projections; pseudopodia long, filose, branching; fresh water.

*D. mobile* A. (Fig. 205, k). Test brown; of various shapes; aperture terminal; body does not fill the test; nucleus large; 1–2 contractile vacuoles; 60–120μ long; projections 8–10μ long; in vegetation.

Genus **Clypeolina** Penard. Test ovoid, compressed, formed of a double envelope; outer envelope composed of 2 valves with scales and particles; inner envelope a membranous sack; long filopodia, often branching; fresh water.

*C. marginata* P. (Fig. 205, l, m). Outer test-valves yellow to dark brown; lenticular in cross-section; wide terminal aperture; endoplasm with many small globules; a single nucleus and contractile vacuole; 80–150μ long.

Family **Euglyphidae** Wallich

Genus **Euglypha** Dujardin (*Pareuglypha* Penard). Test hyaline, ovoid, composed of circular, oval, or scutiform siliceous imbricated scales, arranged in longitudinal rows; aperture bordered with regularly arranged denticulate scales; usually with spines; 1–2 nuclei large, placed centrally; filopodia dichotomously branched; contractile vacuoles; fresh water. Numerous species. Division and encystment (Ivanić, 1934).

*E. acanthophora* (Ehrenberg) (*E. alveolata* D.) (Fig. 74). Test ovoid, or slightly elongate; 3–7 scales protruding around the circular aperture; scales elliptical; body almost fills the test; 50–100μ long.

*E. cristata* Leidy (Fig. 206, a). Test small, elongate with a long neck, fundus with 3–8 spines; scales oval; aperture circular, bordered by a single row of 5–6 denticulate scales; cytoplasm colorless; nucleus posterior; reserve scales are said to be collected around the exterior of aperture, unlike other species in which they are kept within the cytoplasm; 30–70μ long; 12–23μ in diameter; aperture 6–12μ; scales 4.5–9.5μ by 2.5–6.5μ; spines 10–15μ long.
E. mucronata L. (Fig. 206, b). Test large; fundus conical, with 1–2 terminal spines (12–44μ long); aperture circular, bordered by a single row of 6–8 denticulate scales; 100–150μ long, diameter 30–60μ; aperture 15–20μ in diameter.

Fig. 206. a, Euglypha cristata, ×330 (Wailes); b, E. mucronata, ×330 (Wailes); c, Paulinella chromatophora, ×1000 (Wailes); d, Cyphoderia ampulla, ×200 (Cash); e, f, Corythion pulchellum, ×350 (Wailes).

Genus Paulinella Lauterborn. Test small ovoid, not compressed; with siliceous scales in alternating transverse rows; aperture terminal; body does not fill the test completely; nucleus posterior; among vegetation in fresh or brackish water.

P. chromatophora L. (Fig. 206, c). Scales arranged in 11–12 rows; with 1–2 curved algal symbionts; no food particles; a single contractile vacuole; 20–32μ long; 14–23μ in diameter.

Genus Cyphoderia Schlumberger. Test retort-shaped; colorless to yellow; made up of a thin chitinous membrane, covered with discs or scales; aperture terminal, oblique, circular; body does not fill the test completely; nucleus large, posterior; pseudopodia, few, long filose, simple or branched; fresh water (Husnot, 1943).

C. ampulla (Ehrenberg) (Fig. 206, d). Test usually yellow, translucent, composed of discs, arranged in diagonal rows; circular in
cross-section; aperture circular; cytoplasm gray, with many granules and food particles; 2 contractile vacuoles; 60–200μ long; diameter 30–70μ. Several varieties.

Genus Trinema Dujardin. Test small, hyaline, ovoid, compressed anteriorly, with circular siliceous scales; aperture circular, oblique, invaginate; nucleus posterior; filopodia not branched; fresh water in vegetation.

T. enchelys (Ehrenberg) (Fig. 207, a). 1–2 contractile vacuoles;

Fig. 207. a, Trinema enchelys, ×330 (Wailes); b, Placocista spinosa, ×200 (Wailes); c, Assulina seminulum, ×400 (Wailes); d, Nebela collaris, ×200 (Cash); e, Quadrula symmetrica, ×200 (Cash); f, Sphenoderia lenta, ×330 (Leidy).

pseudopodia attenuate, radiating; 30–100μ long; 15–60μ wide; scales 4–12μ in diameter.

T. lineare Penard (Fig. 79). Test transparent; scales indistinct; about 35μ by 17μ; filopodia. Sexual fusion (Dunkerly, 1923) (p. 183).

Genus Corythion Taránek. Test small, hyaline, composed of small oval siliceous plates; compressed; elliptical in cross-section; aperture subterminal, ventral or oblique, and circular or oval; numerous filopodia; fresh water.

C. pulchellum Penard (Fig. 206, e, f). Aperture lenticular; cytoplasm colorless; 2–3 contractile vacuoles; 25–35μ by 15–20μ; aperture 7–10μ by 3–4μ.

Genus Placocista Leidy. Test ovoid, hyaline, compressed; lenticular in cross-section; with oval or subcircular siliceous scales; aperture wide, linear, with flexible undulate borders; nucleus large,
posterior; often with zoochlorellae; filopodia branching and many, generally arising from a protruded portion of cytoplasm; fresh water.

_P. spinosa_ (Carter) (Fig. 207, _b_). Margin of test with spines, either singly or in pairs; 116–174μ by 70–100μ; in sphagnum.

Genus _Assulina_ Ehrenberg. Test colorless or brown; ovoid; with elliptical scales, arranged in diagonal rows; aperture oval, terminal bordered by a thin chitinous dentate membrane; nucleus posterior; contractile vacuoles; filopodia divergent, sometimes branching; fresh water.

_A. seminulum_ (E.) (Fig. 207, _c_). Body does not fill the test; with numerous food particles; pseudopodia few, straight, divergent, slender, seldom branched; 60–150μ by 50–75μ; in sphagnum.

Genus _Nebela_ Leidy. Test thin, ovate or pyriform; with circular or oval platelets of uniform or various sizes; highly irregular; endoplasm with oil globules; nucleus posterior; body does not fill the test, and is connected with the latter by many ectoplasmic strands at fundus end; pseudopodia blunt, rarely branched; fresh water. Numerous species. Taxonomy (Jung, 1942a).

_N. collaris_ (Ehrenberg) (Fig. 207, _d_). Test pyriform, fundus obtuse in profile; aperture without any notch; endoplasm with chlorophyllous food particles; pseudopodia digitate, short, usually 3–6 in number: about 130μ by 85–90μ; in marshes among sphagnum. Feeding habit, binary fission and plasmogamy (MacKinlay, 1936).

Genus _Quadrula_ Schulze. Test pyriform, hemispherical, or discoidal; with quadrangular siliceous or calcareous platelets, arranged generally in oblique series, not overlapping; a single nucleus; body and pseudopodia similar to those of _Difflugia_; fresh water.

_Q. symmetrica_ (Wallich) (Fig. 207, _e_). Compressed, smaller platelets near aperture; cytoplasm very clear, with chlorophyllous granules; 3–5 pseudopodia digitate; nucleus posterior; 80–140μ by 40–96μ; in sphagnum.

Genus _Sphenoderia_ Schlumberger. Test globular or oval, sometimes slightly compressed; hyaline, membranous, with a short broad neck, and a wide elliptical aperture; scales circular, oval, or hexagonal, arranged in alternating series; cytoplasm colorless; 1–2 contractile vacuoles; filopodia, fine, branching; fresh water.

_S. lenta_ S. (Fig. 207, _f_). Hyaline test ovoid or globular; scales circular or broadly oval; aperture terminal, surrounded by a thin chitinous collar, one side inclined inwards; nucleus large; cytoplasm colorless; 2 contractile vacuoles; 30–64μ by 20–46μ; aperture 10–22μ in diameter.
References


Chapter 21
Order 5 Foraminifera d’Orbigny

The Foraminifera are comparatively large Protozoa, living almost exclusively in the sea. They were very abundant in geologic times and the fossil forms are important in applied geology (p. 10). The majority live on ocean bottom, moving about sluggishly over the mud and ooze by means of their pseudopodia. Some are attached to various objects on the ocean floor, while others are pelagic.

The cytoplasm is ordinarily not differentiated into the two zones and streams out through the apertures, and in perforated forms through the numerous pores, of the shell, forming rhizopodia which are fine and often very long and which anastomose with one another to present a characteristic appearance (Fig. 5). The streaming movement of the cytoplasm in the pseudopodia are quite striking; the granules move toward the end of a pseudopodium and stream back along its periphery. The body cytoplasm is often loaded with brown granules which are apparently waste matter and in some forms such as Peneroplis pertusus these masses are extruded from the body from time to time, especially prior to the formation of a new chamber. Contractile vacuoles are usually not found in the Foraminifera.

The test of the Foraminifera varies greatly in form and structure. It may show various colorations—orange, red, brown, etc. The majority measure less than one millimeter, although larger forms may frequently reach several millimeters. The test may be siliceous or calcareous and in some forms, various foreign materials, such as sand-grains, sponge-spicules, etc. which are more or less abundantly found where these organisms live, are loosely or compactly cemented together by pseudochitinious or gelatinous substances. Certain forms show a specific tendency in the selection of foreign materials for the test (p. 47). Siliceous tests are comparatively rare, being found in some species of Miliolidae inhabiting either the brackish water or deep sea. Calcareous tests are sometimes imperforated, but even in such cases those of the young are always perforated. By far the majority of the Foraminifera possess perforated calcareous tests. The thickness of the shell varies considerably, as do also the size and number of apertures, among different species. Frequently the perforations are very small in the young and later become large and coarse, while in others the reverse may be the case.

The form of the shell varies greatly. In some there is only one chamber composed of a central body and radiating arms which repre-
sent the material collected around the pseudopodia, as in Rhabdam-
mina (Fig. 209, a), or of a tubular body alone, as in Hyperammina (Fig.
209, d). The polythalamous forms possess shells of various spirals.
The first chamber is called the proloculum which may be formed
either by the union of two swarmer or by asexual reproduction. The
former is ordinarily small and known as the microspheric proloculum,
while the latter, which is usually large, is called the megalospheric
proloculum. To the proloculum are added many chambers which
may be closely or loosely coiled or not coiled at all. These chambers
are ordinarily undivided, but in many higher forms they are divided
into chamberlets. The chambers are delimited by the suture on the
exterior of the shell. The septa which divide the chambers are per-
forated by one or more foramina known as stolon canals, through
which the protoplasm extends throughout the chambers. The last
chamber has one or more apertures of variable sizes, through which
the cytoplasm extends to the exterior as pseudopodia. The food of
Foraminifera consists mostly of diatoms and algae, though pelagic
forms are known to capture other Protozoa and microcrustaceans.

All species of Foraminifera manifest a more or less distinct tend-
ency toward a dimorphism: the megalospheric form has a large pro-
loculum, is uninucleate and is relatively small in size; while the micro-
spheric form possesses a small proloculum, is multinucleate, and is
large. In addition, there is a difference in the direction of rotation of
spiral chambers of tests in some species (Myers). For example, in
Discorbis opercularis, the microspheric form has clockwise rotation
of the chambers, and the megalospheric form shows counterclock-
wise rotation. The megalospheric forms are said to be much more
numerous than the microspheric forms, especially in pelagic species.
It is possible that, as Myers (1938) pointed out, the flagellate gam-
etes are set free in open water and have a minimum of opportunity
for syngamy.

Lister (1895) observed the development of the megalospheric
form in Elphidium by asexual reproduction from the microspheric
form. He noticed flagellated swarmer in megalospheric tests and
considered them as gametes which through syngamy gave rise to
microspheric individuals. Recent studies by Myers (1935–1940)
confirm the correctness of this view, except that in some species the
gametes are amoeboid. In Spirillina vivipara (Fig. 208, A, 1–5) the
mature microspheric form (1) which measures 125–152 μ in diameter,
becomes surrounded by an envelope composed of substrate debris
and viscous substance. Within the “multiple fission cyst,” nuclear
and cytoplasmic fissions form numerous small uninucleate megalо-
spheric individuals which produce tests and emerge from the cyst (2). They grow into mature megalospheric forms which measure 60–72μ in diameter. Two to four such individuals become associated

Fig. 208. Developmental cycles of Foraminifera (Myers). A, Spirillina vivipara; B, Discorbis patelliformis; C. Elphidium crispa. 1, microospheric forms; 2, megalospheric forms, a–c, enlarged views of young megalospheric forms; 3, beginning of sexual reproduction; 4, gamete and zygote formation, a–c, gametes; 5, young microospheric forms, a–c, enlarged views of one in each species.
and transform into “fertilization cyst.” (5). The nucleus in each individual divides twice or occasionally three times and thus formed multinucleate bodies escape from the tests within the cyst envelope where many gametocytes are produced by multiple fissions. Each gametocyte which contains 12 chromosomes divides into two amoeboid haploid gametes by meiosis. Gametes developed from different parents presumably undergo fusion in pairs and zygotes are produced (4). Each zygote becomes proloculum in which the nucleus divides twice and when the coiled tubular chamber of test grows to about three-quarters of a whorl, young microspheric individuals escape from the cyst and lead independent existence (5). Myers reports the development of *Patellina corrugata* is similar to that of *Spirillina*, except the amoeboid gametes possess 12 haploid number of chromosomes.

In *Discorbis patelliformis* (Fig. 208, B, 1–5), the same investigator noticed no fertilization cyst during the sexual reproduction, but two megalospheric individuals come in contact and flagellate gametes are produced in them. The zygotes develop within the space formed by the dissolution of septa between chambers and tests; the zygote nucleus divides repeatedly within each zygote and forms about 40 nuclei before a test is secreted. In *Elphidium crispa* (Fig. 208, C, 1–5), there is no direct association of megalospheric individuals during sexual reproduction. The flagellated gametes produced in each, are set free in the water and the fusion of the gametes depends entirely upon the chance meeting.

In *Patellina corrugata* and *Discorbis vilardeboanus*, Calvez (1950) finds that the postzygotic divisions of the nucleus are mitotic and the trophozoite nucleus is diploid, but meiosis occurs in the trophozoite just before multiple division.

More than 300 genera of extinct and living Foraminifera are now known. Cushman distinguished 45 families. The present work follows Cushman in recognizing and differentiating 44 families, and lists one genus as an example for each, but places Gromia and allied genera in the order Testacea (p. 472). Taxonomy (Cushman, 1948); ecology (Phleger and Walton, 1950; Phleger and Parker, 1951); distribution (Post, 1951, Illing, 1952).

Test entirely or in part arenaceous

Test single-chambered or rarely an irregular group of similar chambers loosely attached

Test with a central chamber, 2 or more arms; fossil and recent

............................................Family 1 Astrorhizidae
Genus **Rhabdammina** Sars (Fig. 209, a)

Test without a central chamber, elongate, open at both ends; fossil and recent.........................Family 2 Rhizamminidae

Genus **Rhizammina** Brady (Fig. 209, b)

Test a chamber or rarely series of similar chambers loosely attached, with normally a single opening; fossil and recent...............Family 3 Saccamminidae

Genus **Saccammina** Sars (Fig. 209, c)

Test 2-chambered, a proloculum and long undivided tubular second chamber

Fig. 209. a, Rhabdammina abyssorum, ×5 (Kühn); b, Rhizammina algaeformis, fragment of, ×14 (Cushman); c, Saccammina sphaerica, ×8 (Rhumbler); d, Hyperammina subnodosa, ×4 (Brady); e, Ammodiscus incertus, ×20 (Kühn); f, Silicina limitata, ×13 (Cushman); g, Reophax nodulosus, ×3 (Brady).

Test with the second chamber, simple or branching, not coiled; mostly recent and also fossil........Family 4 Hyperamminidae

Genus **Hyperammina** Brady (Fig. 209, d)

Test with the second chamber usually coiled at least in young

Test of arenaceous material with much cement, usually yellowish or reddish brown; fossil and recent. Family 5 Ammodiscidae

Genus **Ammodiscus** Reuss (Fig. 209, e)

Test of siliceous material, second chamber partially divided; fossils only.........................Family 6 Silicinidae

Genus **Silicina** Bornemann (Fig. 209, f)

Test typically many-chambered

Test with all chambers in a rectilinear series; fossil and recent........Family 7 Reophacidae
Genus **Reophax** Montfort (Fig. 209, g)
Test planispirally coiled at least in young
Axis of coil, short; many uncoiled forms; fossil and recent

.................................Family 8 Lituolidae

Genus **Lituola** Lamarck (Fig. 210, a)
Axis of coil usually long, all close-coiled
Interior not labyrinthic; fossil only

Family 9 Fusulinidae

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Fig. 210. a, *Lituola nautiloidea* (Cushman); b, section through a Fusulina (Carpenter); c. *Textularia agglutinans*, ×90 (Rhumbler); d, *Verneuilina propinqua*, ×8 (Brady); e, *Valvulina triangularis*, (d'Orbigny); f, *Trochammina inflata*, ×32 (Brady); g, *Placopsilina cenomana* (Reuss); h, *Tetrataxis palaeotrochus*, ×15 (Brady); i, *Spiroloculina limbata*, ×20 (Brady); j, *Triloculina trigonula*, ×15 (Brady); k, *Fischerina helix*, ×32 (Heron-Allen and Earland); l, *Vertebralina striata*, ×40 (Kühn); m, *Alveolinella mello*, ×35 (Brady).

Genus **Fusulina** Fisher (Fig. 210, b)
Interior labyrinthic; fossil only

Family 10 Loftusiidae

Genus **Loftusia** Brady
Test typically biserial at least in young of microspheric form; fossil and recent

Family 11 Textulariidae

Genus **Textularia** Defrance (Fig. 210, c)
Test typically triserial at least in young of microspheric form
Aperture usually without a tooth, test becoming simpler in higher forms; fossil and recent

Family 12 Verneuilinidae
Genus *Verneuilina* d'Orbigny (Fig. 210, d)
Aperture typically with a tooth, test becoming conical in higher forms; fossil and recent. ...............Family 13 Valvulinidae

Genus *Valvulina* d'Orbigny (Fig. 210, e)
Test with whole body labyrinthic, large, flattened, or cylindrical; recent..........................Family 14 Neusinidae

Genus *Neusina* Goës
Test trochoid at least while young
Mostly free, typically trochoid throughout; fossil and recent...
........................................Family 15 Trochamminidae

Genus *Trochammina* Parker and Jones (Fig. 210, f)
Attached; young trochoid, later stages variously formed; fossil and recent. .................Family 16 Placopsilinidae

Genus *Placopsilina* d'Orbigny (Fig. 210, g)
Free; conical, mostly of large size; fossil only............................Family 17 Orbitolinidae

Genus *Tetrataxis* Ehrenberg (Fig. 210, h)
Test coiled in varying planes, wall imperforate, with arenaceous portion only on the exterior; fossil and recent. .........Family 18 Miliolidae (in part)

Genus *Spiroloculina* d'Orbigny (Fig. 210, i)
Test calcareous, imperforate, porcellaneous
Test with chambers coiled in varying planes, at least in young; aperture large, toothed; fossil and recent. ....Family 18 Miliolidae (in part)

Genus *Triloculina* d'Orbigny (Fig. 210, j)
Test trochoid; fossil and recent.............Family 19 Fischerinidae

Genus *Fischerina* Terquem (Fig. 210, k)
Test planispiral at least in young
Axis very short, chambers usually simple; fossil and recent.................Family 20 Ophthalmidiidae

Genus *Vertebralina* d'Orbigny (Fig. 210, l)
Axis short, test typically compressed and often discoid, chambers mostly with many chamberlets; fossil and recent...............
........................................Family 21 Peneroplidae

Genus *Peneroplis* Montfort (Figs. 4; 211)
Axis typically elongate, chamberlets developed; mainly fossil..........
........................................Family 22 Alveolinellidae
Fig. 211. Diagram illustrating the life-cycle of *Peneroplis pertusus* (Winter). a–f, megalospheric generation; g, gamete formation; h–k, isogamy; l–n, microspheric generation; o, multiple division.

Genus *Alveolinella* Douvillé (Fig. 210, m)

Test globular, aperture small, not toothed; recent only. ............... Family 23 Keramosphaeridae

Genus *Keramosphaera* Brady

Test calcareous, perforate
Test vitreous with a glassy lustre, aperture typically radiate, not trochoid
Test planispirally coiled or becoming straight, or single-chambered; fossil and recent. Family 24 Lagenidae

Genus Lagena Walker and Jacob (Fig. 212, a)
Test biserial or elongate spiral; fossil and recent. Family 25 Polymorphinidae

Genus Polymorphina d’Orbigny
Test not vitreous; aperture not radiating
Test planispiral, occasionally trochoid, then usually with processes along the suture lines, septa single, no canal system; fossil and recent. Family 26 Nonionidae

Genus Elphidium Montfort (Figs. 5; 208, C; 212, b)
(Polystomella Lamarck)
Test planispiral, at least in young, generally lenticular, septa double, canal system in higher forms; fossil and recent. Family 27 Camerinidae

Genus Operculina d’Orbigny (Fig. 212, c)
Test generally biserial in at least microspheric form, aperture usually large, without teeth; fossil and recent. Family 28 Heterohelicidae
Genus **Pavonina** d’Orbigny (Fig. 212, d)
Test planispiral, bi- or tri-serial with elongate spines and lobed aperture; fossil and recent. .......... Family 29 Hantkeninidae

Genus **Hantkenina** Cushman (Fig. 212, e)
Test typically with an internal tube, elongate
Aperture generally loop-shaped or cribrate; fossil and recent...
................................................. Family 30 Buliminidae

Genus **Bolivina** d’Orbigny (Fig. 212, f)
Aperture narrow, curved, with an overhanging portion; mostly fossil, also recent............ Family 31 Ellipsoidinidae

Genus **Ellipsoidina** Seguenza
Test trochoid, at least in young of microspheric form, usually coarsely perforate; when lenticular, with equatorial and lateral chambers
Test trochoid throughout, simple; aperture ventral
No alternating supplementary chambers on ventral side; fossil and recent....................... Family 32 Rotaliidae

Genus **Rotalia** Lamarck (Fig. 212, g)

Genus **Spirillina** Ehrenberg (Fig. 208, A)

Genus **Patellina** Williamson.

Genus **Discorbis** Lamarck (Fig. 208, B)
Alternating supplementary chambers on ventral side; fossil and recent......................... Family 33 Amphisteginidae

Genus **Asterigerina** d’Orbigny (Fig. 212, h)
Test trochoid and aperture ventral in young
With supplementary material and large spines, independent of chambers; fossil and recent...... Family 34 Calcarinidae

Genus **Calcarina** d’Orbigny (Fig. 213, a)
With later chambers in annular series or globose with multiple apertures, but not covering earlier ones; fossil and recent....
................................................. Family 35 Halkyardiiidae

Genus **Halkyardia** Heron-Allen and Earland (Fig. 213, b)
With later chambers somewhat biserial; aperture elongate in the axis of coil; fossil and recent.. Family 36 Cassidulinidae
Genus Cassidulina d'Orbigny (Fig. 213, c)
With later chambers becoming involute, very few making up the exterior in adult; aperture typically elongate, semicircular; in a few species circular; fossil and recent. ......................... Family 37 Chilostomellidae

Genus Allomorphina Reuss (Fig. 213, d)
With chambers mostly finely spinose and wall cancellated, adapted, for pelagic life, globular forms with the last chamber completely involute; aperture umbilicate or along the suture; fossil and recent. ................. Family 38 Globigerinidae

Fig. 213. a, Calcarina defrancei, ×25 (Brady); b, Halkyardia radiata, ×15 (Cushman); c, Cassidulina laevigata, ×25 (Brady); d, Allomorphina trigona, ×40 (Brady); e, Globigerina bulloides, ×30 (Kühn); f, Anomalina punctulata (d'Orbigny); g, Rupertia stabilis, ×50 (Brady).

Genus Globigerina d'Orbigny (Fig. 213, e)
Early chambers globigerine, later ones spreading and compressed; fossil and recent. ................. Family 39 Globorotaliidae

Genus Globorotalia Cushman
Test trochoid at least in young, aperture peripheral or becoming dorsal
Mostly attached, dorsal side usually flattened; fossil and recent. .................................. Family 40 Anomaliniidae

Genus Anomalina d'Orbigny (Fig. 213, f)
Later chambers in annular series; fossil and recent. .................................. Family 41 Planorbulinidae
Genus **Planorbulina** d’Orbigny

Test trochoid in very young, later growing upward
Later chambers in loose spiral; fossil and recent............... Family 42 Rupertiidae

Genus **Rupertia** Wallich (Fig. 213, g)

Later chambers in masses or branching, highly colored; mostly recent, also fossil............ Family 43 Homotremidae

Genus **Homotrema** Hickson

Test trochoid in the very young of microspheric form, chambers becoming annular later, with definite equatorial and lateral chambers, often with pillars; fossil only................... Family 44 Orbitoididae

Genus **Orbitoides** d’Orbigny

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Chapter 22

Subclass 2 Actinopoda Calkins

The Actinopoda are divided into two orders as follows:
Without central capsule.........................Order 1 Heliozoa
With central capsule.........................Order 2 Radiolaria (p. 516)

Order 1 Heliozoa Haeckel

The Heliozoa are, as a rule, spherical in form with many radiating axopodia. The cytoplasm is differentiated, distinctly in Actinosphaerium, or indistinctly in other species, into the coarsely vacuolated ectoplasm and the less transparent and vacuolated endoplasm. The food of Heliozoa consists of living Protozoa or Protophyta; thus their mode of obtaining nourishment is holozoic. A large organism may sometimes be captured by a group of Heliozoa which gather around the prey. When an active ciliate or a small rotifer comes in contact with an axopodium, it seems to become suddenly paralyzed and, therefore, it has been suggested that the pseudopodia contain some poisonous substances. The axial filaments of the axopodia disappear and the pseudopodia become enlarged and surround the food completely. Then the food matter is carried into the main part of the body and is digested. The ectoplasm contains several contractile vacuoles and numerous refractile granules which are scattered throughout. The endoplasm is denser and usually devoid of granules. In the axopodium, the cytoplasm undergoes streaming movements. The hyaline and homogeneous axial filament runs straight through both the ectoplasm and the endoplasm, and terminates in a point just outside the nuclear membrane. When the pseudopodium is withdrawn, its axial filament disappears completely, though the latter sometimes disappears without the withdrawal of the pseudopodium itself. In Acanthocystis the nucleus is eccentric (Fig. 216, b), but there is a central granule, or centroplast, in the center of the body from which radiate the axial filaments of the axopodia. In multinucleate Actinosphaerium, the axilia filaments terminate at the periphery of the endoplasm. In Camptonema, an axial filament arises from each of the nuclei (Fig. 214, d).

The skeletal structure of the Heliozoa varies among different species. The body may be naked, covered by a gelatinous mantle, or provided with a lattice-test with or without spicules. The spicules are variable in form and location and may be used for specific differentiation. In some forms there occur colored bodies bearing chromatophores, which are considered as holophytic Mastigophora.
(p. 29) living in the heliozoans as symbionts.

The Heliozoa multiply by binary fission or budding. Incomplete division may result in the formation of colonies, as in Rhaphidio-phys. In Actinosphaerium, nuclear phenomena have been studied by several investigators (p. 204). In Acanthocystis and Oxnerella (Fig. 59), the central granule behaves somewhat like the centriole in a metazoan mitosis. Budding has been known in numerous species. In Acanthocystis the nucleus undergoes amitosis several times, thus forming several nuclei, one of which remains in place while the other migrates toward the body surface. Each peripheral nucleus becomes surrounded by a protruding cytoplasmic body which becomes covered by spicules and which is set free in the water as a bud. These small individuals are supposed to grow into larger forms, the central granules being produced from the nucleus during the growth. Formation of swarmers is known in a few genera and sexual reproduction occurs in some forms. The Heliozoa live chiefly in fresh water, although some inhabit the sea. Taxonomy and morphology (Penard, 1905, 1905a; Cash and Wailes, 1921; Roskin, 1929, Valkanov, 1940).

Without gelatinous envelope
Without flagella
Pseudopodia arise from thick basal parts, branching. .............
.................................................Family 1 Actinocomidae
Pseudopodia not branching, cytoplasm highly vacuolated..........
.................................................Family 2 Actinophryidae (p. 507)
With 1–2 flagella...............................Family 3 Ciliophryidae (p. 508)

With gelatinous envelope; with or without skeleton
Without flagella
Without chitinous capsule
Without definite skeleton........Family 4 Lithocollidae (p. 508)
With chitinous or siliceous spicules or scales
With chitinous spicules....Family 5 Heterophryidae (p. 510)
With siliceous skeleton
Cup-like plates over body; 2–3 pseudopodia often grouped
.................................................Family 6 Clathrellidae (p. 511)
Scales flattened, not cup-like
.................................................Family 7 Acanthocystidae (p. 511)
With chitinous retiform capsule....Family 8 Clathulinidae (p. 513)
With numerous flagella, among axopodia; siliceous scales
.................................................Family 9 Myriophryidae (p. 514)

Family 1 Actinocomidae Poche

Genus Actinocoma Penard. Body spherical; one or more contractile vacuoles; nucleus with a thick membrane, central; filopodia, not axopodia, simple or in brush-like groups; fresh water.

A. ramosa P. (Fig. 214, a). Average diameter 14–26μ.
Family 2 Actinophyridae Claus

Genus Actinophrys Ehrenberg. Spheroidal; cytoplasm highly vacuolated, especially ectoplasm; with often symbiotic zoochlorellae; nucleus central; 1 to many contractile vacuoles; axopodia straight,

![Diagram](img)

Fig. 214. a, Actinocoma ramosa, ×630 (Penard); b, Actinophrys sol, ×400 (Kudo); c, Actinosphaerium eichhorni, ×45 (Kudo); d, Camptonema nutans, ×350 (Schaudinn).

numerous, axial filaments terminate at surface of the nucleus; “sun animalcules”; fresh water.

A. sol E. (Figs. 90; 214, b). Spherical; ectoplasm vacuolated; endoplasm granulated with numerous small vacuoles; a large central nucleus; solitary but may be colonial when young; diameter variable, average being 40–50μ; among plants in still fresh water. Reproduction, morphology and physiology (Bělař, 1923, 1924); food habit (Looper, 1928).
A. vesiculata Penard. Ectoplasm with saccate secondary vesicles, extending out of body surface between axopodia; nucleus central, with many endosomes; 25–30μ in average diameter; fresh water.

Genus Actinosphaerium Stein. Spherical; ectoplasm consists almost entirely of large vacuoles in one or several layers; endoplasm with numerous small vacuoles; numerous nuclei; axopodia end in the inner zone of ectoplasm (Fig. 6). 2 species.

A. eichhorni Ehrenberg (Figs. 6; 214, c). Numerous nuclei scattered in the periphery of endoplasm; 2 or more contractile vacuoles, large; axial filaments arise from a narrow zone of dense cytoplasm at the border line between endoplasm and ectoplasm; body large, diameter 200–300μ, sometimes up to 1 mm.; nuclei 12–20μ in diameter; among vegetation in freshwater bodies. Nuclear change (Speeth, 1919); morphology (Rumjantzew and Wermel, 1925); transplantation (Okada, 1930).

A. arachnoideum Penard. Ectoplasm irregularly vacuolated; no distinct endoplasmic differentiation; nuclei smaller in number; pseudopodia of 2 kinds; one straight, very long and the other filiform, and anastomosing; 70–80μ in diameter; fresh water.

Genus Camptonema Schaudinn. Spheroidal; axial filaments of axopodia end in nuclei about 50 in number; vacuoles numerous and small in size; salt water.

C. nutans S. (Fig. 214, d). About 150μ in diameter.

Genus Oxnerella Dobell. Spherical; cytoplasm indistinctly differentiated; eccentric nucleus with a large endosome; axial filaments take their origin in the central granule; no contractile vacuole; nuclear division typical mitosis (Fig. 59).

O. maritima D. (Fig. 59). Small, 10–22μ in diameter; solitary, floating or creeping; salt water.

Family 3 Ciliophryidae Poche

Genus Ciliophrys Cienkowski. Spherical with extremely fine radiating filopodia, giving the appearance of a typical heliozoan, with a single flagellum which is difficult to distinguish from the numerous filopodia, but which becomes conspicuous when the pseudopodia are withdrawn; fresh or salt water.

C. infusionum C. (Fig. 215, a). 25–30μ long; freshwater infusion.  
C. marina Caullery. About 10μ in diameter; salt water.

Family 4 Lithocollidae Poche

Genus Lithocolla Schulze. Spherical body; outer envelope with usually one layer of sand-grains, diatoms, etc.; nucleus eccentric.
L. globosa S. (Fig. 215, b). Body reddish with numerous small colored granules; nucleus large; central granule unknown; envelope 35–50μ in diameter; in lakes, ponds, and rivers; also in brackish water.

Fig. 215. a, Ciliophrys infusionum, ×400 (Bütschli); b, Lithocolla globosa, ×250 (Penard); c, Astrodisculus radians, ×600 (Penard); d, Actinolophus pedunculatus, ×400 (Schultze); e, Elaeorhanis cincta, ×300 (Penard); f, Sphaerastrum fockei, ×300 (Stubenrauch); g, Heterophrys myriopoda, ×270 (Penard).
Genus *Astrodisculus* Greeff. Spherical with gelatinous envelope, free from inclusions, sometimes absent; no demarcation between 2 regions of the cytoplasm; pseudopodia fine without granules; fresh water.

*A. radians* G. (Fig. 215, c). Outer surface usually with adherent foreign bodies and bacteria; cytoplasm often loaded with green, yellow, or brown granules; nucleus eccentric; a contractile vacuole; diameter 25–30μ including envelope; in pools and ditches.

Genus *Actinolophus* Schulze. Body pyriform, enveloped in a gelatinous mantle; stalked; stalk apparently hollow; axopodia long, numerous; nucleus eccentric; salt water.

*A. pedunculatus* S. (Fig. 215, d). Diameter about 30μ; stalk about 100μ long.

Genus *Elaeorhanis* Greeff. Spherical; mucilaginous envelope with sand-grains and diatoms; cytoplasm with a large oil globule; nucleus eccentric; 1 or more contractile vacuoles; pseudopodia not granulated, sometimes forked; fresh water.

*E. cineta* G. (Fig. 215, e). Bluish with a large yellow oil globule; without any food particles; no central granule; pseudopodia rigid, but apparently without axial filaments, sometimes forked; young forms colonial; solitary when mature; outer diameter 50–60μ; body itself 25–30μ; in lakes and pools.

Genus *Sphaerastrum* Greeff. Somewhat flattened; greater part of axopodia and body covered by a thick gelatinous mantle; a central granule and an eccentric nucleus; fresh water.

*S. fockei* G. (Fig. 215, f). Diameter about 30μ; often colonial; in swamps.

Family 5 *Heterophryidae* Poche

Genus *Heterophrys* Archer. Spherical; mucilaginous envelope thick, with numerous radial, chitinous spicules which project beyond periphery; nucleus eccentric; axial filaments originate in a central granule; fresh or salt water.

*H. myriopoda* A. (Fig. 215, g). Nucleus eccentric; cytoplasm loaded with spherical algae, living probably as symbionts; contractile vacuoles indistinct; 50–80μ in diameter; in pools and marshes; and also among marine algae.

*H. glabrescens* Penard. Spherical; gelatinous envelope poorly developed; chitinous needles indistinct; pseudopodia very long; 11–15μ in diameter; fresh water.
Family 6 Clathrellidae Poche

Genus *Clathrella* Penard. Envelope distinct, polygonal; surface with uniform alveoli with interalveolar portion extending out; envelope appears to be continuous, but in reality formed by a series of cup-like bodies; contractile vacuole large; voluminous nucleus eccentric; filopodia straight, some bifurcated, arising between "cups."

*C. foerti* P. (Fig. 216, a). Envelope about 40–55µ in diameter; fresh water.

Family 7 Acanthocystidae Claus

Genus *Acanthocystis* Carter. Spherical; siliceous scales, arranged tangentially and radiating siliceous spines with pointed or bifurcated ends; nucleus eccentric; a distinct central granule in which the axial filaments terminate. Several species.

*A. aculeata* Hertwig and Lesser (Fig. 216, b). Tangential scales stout and pointed; spines curved and nail-headed; cytoplasm greyish; a single contractile vacuole; diameter 35–40µ; spines about 1/3 the body diameter; in fresh water. Morphology and reproduction (Stern, 1924).

Genus *Pompholyxophrys* Archer. Spherical; outer mucilaginous envelope with minute colorless spherical granules arranged in concentric layers; nucleus eccentric; contractile vacuoles; pseudopodia long, straight, acicular; fresh water.

*P. punicea* A. (Fig. 216, c). Body colorless or reddish, with usually many colored granules and green or brown food particles; nucleus large, eccentric; solitary, active; diameter 25–35µ; outer envelope 5–10µ larger; in pools.

Genus *Raphidiophrys* Archer. Spherical; mucilaginous envelope with spindle-shaped or discoidal spicules which extend normally outwards along pseudopodia; nucleus and endoplasm eccentric; solitary or colonial; fresh water. Several species.

*R. pallida* Schulze (Fig. 216, d). Outer gelatinous envelope crowded with curved lenticular spicules, forming accumulations around pseudopodia; ectoplasm granulated; nucleus eccentric; contractile vacuoles; axial filaments arise from the central granule; solitary; diameter 50–60µ; nucleus 12–15µ in diameter; spicules 20µ long; among vegetation in still fresh water.

Genus *Raphidocystis* Penard. Spicules of various forms, but unlike those found in the last genus.

*R. tubifera* P. (Fig. 216, e). Spicules tubular with enlarged extrem-
ity; diameter about 18\(\mu\); envelope 25\(\mu\); fresh water.

*R. infestans* Wetzel. Body 20–40\(\mu\) in diameter; thin axopodia twice the body diameter; without radial spicules; feeds on ciliates (Wetzel, 1925).

Genus *Wagnerella* Mereschkowsky. Spherical, supported by a

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**Fig. 216.** a, *Clathrella foreli*, ×250 (Penard); b, *Acanthocystis aculeata*, ×300 (Stern); c, *Pompholyxophrys punicea*, ×260 (West); d, *Raphidiophrys pallida*, ×300 (Penard); e, *Raphidocystis tubifera*, ×500 (Penard); f, *Wagnerella borealis*, ×75 (Kühn); g, *Pinaciophora fluviatilis*, ×250 (Penard).
cylindrical stalk with an enlarged base; small siliceous spicules; nucleus in the base of stalk; multiplication by budding.

*W. borealis* M. (Fig. 216, f). About 180μ in diameter; stalk often up to 1.1 mm. long; salt water.

Genus **Pinaciophora** Greeff. Spherical; outer envelope composed of circular discs, each being perforated with 19 minute pores; cytoplasm reddish; fresh water.

*P. fluviatilis* G. (Fig. 216, g). Diameter 45–50μ, but somewhat variable; in freshwater ponds.

Family 8 **Clathrulinidae** Claus

Genus **Clathrulina** Cienkowski. Envelope spherical, homogeneous, with numerous regularly arranged openings; with a stalk; protoplasm central, not filling the capsule; nucleus central; pseudopodia numerous, straight or forked, granulated; fresh water.

*C. elegans* C. (Fig. 217, a). Envelope colorless to brown, perforated by numerous comparatively large circular or polygonal openings; 1 or more contractile vacuoles; nucleus central; diameter 60–90μ, openings 6–10μ; length of stalk 2–4 times the diameter of envelope, 3–4μ wide; solitary or colonial; among vegetation in ponds. Taxonomy and stalk formation (Valkanov, 1928).

Genus **Hedriocystis** Hertwig and Lesser. Envelope spherical, openings minute, surrounded by polyhedral facets or ridges; with stalk; solitary or colonial; fresh water.

*H. reticulata* Penard (Fig. 217, b). Envelope colorless or pale yellow, facets regularly polygonal with raised borders; stalk solid, nucleus central; 1 contractile vacuole; each pseudopodium arises from a pore located in the center of a facet; solitary; capsule about 25μ in diameter; body about 12μ in diameter; stalk about 70μ by 1.5μ; in marshy pools.

Genus **Elaster** Grimm. Envelope spherical, delicate, penetrated by numerous more or less large pores; without stalk; pseudopodia many, straight filose.

*E. greeffi* G. (Fig. 217, c). Diameter of envelope 20μ; envelope delicate, colorless; many pseudopodia; in peaty soil.

Genus **Choanocystis** Penard. Spherical envelope with perforations which possess conical borders; openings of cones provided with funnel-like expansions, edges of which nearly touch one another; fresh water.

*C. lepidula* P. (Fig. 217, d). Diameter 10–13μ; envelope delicate; 1 or more contractile vacuoles; pseudopodia very long.
Family 9 Myriophryidae Poche

Genus Myriophrys Penard. Spherical or ovoid, covered with a protoplasmic envelope containing scales (?), surrounded by numer-

Fig. 217. a, Clathrulina elegans, ×250 (Leidy); b, Hedriocystis reticulata, ×500 (Brown); c, Elaster greefi, ×680 (Penard); d, Choanocystis lepidula, ×690 (Penard); e, Myriophrys paradoxa, ×300 (Penard).
ous fine processes; endoplasm vesicular; a large nucleus eccentric; a large contractile vacuole; long pseudopodia granulated and attenuated toward ends.

M. paradoxa P. (Fig. 217, ε). Average diameter 40μ; in fresh-water swamps.

References

Valkanov, A.: (1928) Protistenstudien. III. Ibid., 64: 446.
Chapter 23

Order 2 Radiolaria Müller

The Radiolaria are pelagic in various oceans. A vast area of the ocean floor is known to be covered with the ooze made up chiefly of radiolarian skeletons. They seem to have been equally abundant during former geologic ages, since rocks composed of their skeletons occur in various geological formations. Thus this group is the second group of Protozoa important to geologists.

The body is generally spherical, although radially or bilaterally symmetrical forms are also encountered. The cytoplasm is divided distinctly into two regions which are sharply delimited by a membranous structure known as the central capsule. This is a single or double perforated membrane of pseudochitinous or mucinoid nature. Although its thickness varies a great deal, the capsule is ordinarily very thin and only made visible after addition of reagents. Its shape varies according to the form of the organism; thus in spherical forms it is spherical, in discoidal or lenticular forms it is more or less ellipsoidal, while in few cases it shows a number of protruding processes. The capsule is capable of extension as the organism grows and of dissolution at the time of multiplication. The cytoplasm on either side of the capsule communicates with the other side through pores which may be large and few or small and numerous. The intracapsular portion of the body is the seat of reproduction, while the extracapsular region is nutritive and hydrostatic in function. The intracapsular cytoplasm is granulated, often greatly vacuolated, and is stratified either radially or concentrically. It contains one or more nuclei, pigments, oil droplets, fat globules, and crystals. The nucleus is usually of vesicular type, but its form, size, and structure, vary among different species and also at different stages of development even in one and the same species.

A thin assimilative layer, or matrix, surrounds the central capsule. In Tripylea, waste material forms a brownish mass known as phaeodarium, around the chief aperture (astropyile) of the capsule. Then there is a highly alveolated region, termed calymma, in which the alveoli are apparently filled with a mucilaginous secretion of the cytoplasm. Brandt showed that the vertical movement of some Radiolaria is due to the formation and expulsion of a fluid which consists of water saturated with carbon dioxide. Under ordinary weather and temperature conditions, the interchange between the alveoli and the exterior is gradual and there is a balance of loss and gain of the fluid, so that the organisms float on the surface of the sea. Under
rough weather conditions or at extraordinary high temperatures, the pseudopodia are withdrawn, the alveoli burst, and the organisms descend into deeper water, where the alveoli are reformed.

The Radiolaria feed on microplankton such as copepods, diatoms, and various Protozoa. The food is taken in through pseudopodia and passed down into the deeper region of calymma where it is digested in food vacuoles. The Radiolaria can, however, live under experimental conditions without solid food if kept under light. This is ordinarily attributed to the action of the yellow corpuscles which are present in various parts of the body, although they are, as a rule, located in the calymma. In Actipylea they are found only in intracapsular cytoplasm, and in Tripylea they are absent altogether. They are spherical bodies, about 15μ in diameter, with a cellulose wall, 2 chromatophores, a pyrenoid, starch, and a single nucleus. They appear to multiply by fission. These bodies are considered as zooxanthellae (p. 274). In the absence of organic food material, the Radiolaria live probably by utilizing the products of holophytic nutrition of these symbiotic organisms.

The axopodia arise from either the extracapsular or the intracapsular portion and radiate in spherical forms in all directions, as in Heliozoa. In Actipylea, myonemes are present in certain pseudopodia and produce circular groups of short, rod-like bodies clustered around each of the radial spines (Fig. 219, c). They connect the peripheral portion of the body with the pseudopodial covering of the spicule and possess a great contractile power, supposedly with hydrostatic function (p. 62).

The skeletal structure of Radiolaria varies considerably from simple to complex and has a taxonomic value. The chemical nature of the skeleton is used in distinguishing the major subdivisions of the order. In the Actipylea it seems to be made up of strontium sulphate, while in the three other groups, Peripylea, Monopylea, and Tripylea, it consists fundamentally of siliceous substances. The skeleton of the Actipylea is sharply marked from others in form and structure. The majority of this group possess 20 rods radiating from center. The rod-shaped skeletons emerge from the body in most cases along five circles, which are comparable to the equatorial, two tropical and two circumpolar circles of the globe, which arrangement is known as Müller's law, since J. Müller first noticed it in 1858.

The life-cycle of the Radiolaria is very incompletely known (Fig. 218). Binary or multiple fission or budding has been seen in some Peripylea, Actipylea, and Tripylea. Multiple division is also known to occur in Thalassophysidae in which it is the sole known means of
reproduction. The central capsule becomes very irregular in its outline and the nucleus breaks up into numerous chromatin globules. Finally the capsule and the intrcapsular cytoplasm become trans-

Fig. 218. Diagram illustrating the probable life-cycle of *Actipylea* (Kühn). a, mature individual; b, c, binary fission; d, e, multiplication by budding; f, mature individual similar to a; g, formation of swarmers; h–j, supposed, but not observed, union of two swarmers producing a zygote; k, l, young individuals.

formed into numerous small bodies, each containing several nuclei. Further changes are unknown. Swarmer-formation is known in some forms. In *Thalassicolla*, the central capsule becomes separated from the remaining part of the body and the nuclei divide into a number of small nuclei, around each of which condenses a small ovoidal mass of cytoplasm. They soon develop flagellum. In the meantime the capsule descends to a depth of several hundred meters, where its
wall bursts and the flagellates are liberated (q). Both isoswarers and anisoswarers occur. The former often contain a crystal and a few fat globules. Of the latter, the macroswarers possess a nucleus and refringent spherules in the cytoplasm. Some forms possess 2 flagella, one of which is coiled around the groove of the body, which makes them resemble certain dinoflagellates. Further development is unknown; it is supposed that the anisoswarers are sexual and isoswarers asexual generations. Nuclear relationship (Hertwig, 1930).

Enormous numbers of species of Radiolaria are known. An outline of the classification is given below, together with a few examples, of the genera.

Skeleton composed of strontium sulphate .......... Suborder 1 Actipylea
Skeleton composed of other substances

Central capsule uniformly perforated, skeleton either tangential to the capsule or radiating without reaching the intracapsular region .......... Suborder 2 Peripylea (p. 520)

Central capsule not uniformly perforated

Capsule monaxonic, bears at one pole a perforated plate forming the base of an inward-directed cone .......... Suborder 3 Monopylea (p. 522)

Capsule with 3 openings: 1 astropyple and 2 parapyles .......... Suborder 4 Tripylea (p. 523)

Suborder 1 Actipylea Hertwig
Radial spines, 10–200, not arranged according to Müller's law.
Spines radiate from a common center, ancestral forms (Haeckel) ......

Genus Actinelius (Fig. 219, a)
10–16 spines irregularly set .......... Family 1 Actineliidae

Genus Acanthociasma (Fig. 219, b)
Radial spines, few, arranged according to Müller's law
Without tangential skeletons
Spines more or less uniform in size
Spicules circular in cross-section .... Family 3 Acanthometridae

Genus Acanthometron (Fig. 219, c)
Spicules cruciform in cross-section .... Family 4 Acanthoniidae

Genus Acanthonia (Fig. 219, d)
2 opposite spines much larger .......... Family 5 Amphilonchidae
Fig. 219. a, Actinellus primordialis, ×25 (Haeckel); b, Acanthocissos planum, ×65 (Mielek); c, Acanthometron elasticum (Hertwig); d, Acan- thonia tetracopa, ×40 (Schewiakoff); e, Amphilonche hydrometrica, ×130 (Haeckel); f, Hexaconus serratus, ×100 (Haeckel).

Genus Amphilonche (Fig. 219, e)

With tangential skeletons
20 radial spines of equal size, shell composed of small plates, each with one pore. Family 6 Sphaerocapsidae

Genus Sphaerocapsa

2 or 6 larger spines
2 enormously large conical sheathed spines. Family 7 Diploconidae

Genus Diploconus

6 large spines. Family 8 Hexalaspidae

Genus Hexaconus (Fig. 219, f)

Suborder 2 Peripylea Hertwig

Solitary, skeleton wanting or simple spicules; mostly spherical
Nucleus spherical with smooth membrane
Vacuoles intracapsular. Family 1 Physematiidae

Genus Lampoxanthium (Fig. 220, a)

Vacuoles extracapsular. Family 2 Thalassicollidae

Genus Thalassicolla (Fig. 220, b)

Nuclear membrane not smoothly contoured
Nuclear wall branching out into pouches, structure similar to the last. Family 3 Thalassophysidae
Radialaria

Fig. 220 a, *Lampoxanthium pandora*, ×20 (Haeckel); b, *Thalassicolla nucleata*, ×15 (Huth).

**Genus Thalassophysa**

Nuclear wall crenate

Huge double spicule ............... Family 4 Thalassothamnidae

**Genus Thalassothamnus**

A latticed skeleton, with branching and thorny spines .............

................................. Family 5 Orosphaeridae

**Genus Orosphaera**

Solitary, skeleton complex, often concentric

Central capsule and skeleton spherical .... Family 6 Sphaeridae

**Genus Hexacontium** (Fig. 221, a)

Central capsule and skeleton elliptical or cylindrical ..............

................................. Family 7 Prunoidae

Fig. 221. a, *Hexacontium asteracanthion*, ×130; b, *Pipetta tuba*, ×100;

*Staurocyclia phacostaurus*, ×130; d, *Cenolarus primordialis*, ×100;

e, *Sphaerodium ovodimare*, ×30 (Haeckel).
Genus *Pipetta* (Fig. 221, *b*)
Central capsule and skeleton discoidal or lenticular.......................... Family 8 Discoidae

Genus *Staurocyclia* (Fig. 221, *c*)
Similar to the above, but flattened.............. Family 9 Larcoidae

Genus *Cenolarus* (Fig. 221, *d*)
Colonial, individuals with anastomosing extracapsular cytoplasm, embedded in a jelly mass
Without latticed skeleton, but with siliceous spicules arranged tangentially to central capsule.............. Family 10 Sphaerozoidae

Genus *Sphaerozoum* (Fig. 221, *e*)
Central capsule of each individual enclosed in a latticed skeleton..... ........................................... Family 11 Collossphaeridae

Genus *Collossphaera*
Suborder 3 Monopylea Hertwig
Without any skeleton.......................... Family 1 Nassoidae

Genus *Cystidium* (Fig. 222, *a*)
With skeleton
Without a complete latticed skeleton
Skeleton a basal tripod....................... Family 2 Plectoidae

Fig. 222. *a*, *Cystidium princeps*, ×120; *b*, *Triplagia primordialis*, ×25; *c*, *Lithocircus magnificus*, ×100; *d*, *Dictyophimus hertwigi*, ×80 (Haeckel).

Genus *Triplagia* (Fig. 222, *b*)
Skeleton a simple or multiple sagittal ring... Family 3 Stephoidae

Genus *Lithocircus* (Fig. 222, *c*)
With a complete latticed skeleton
Lattice skeleton single, without constriction... Family 4 Cyrtoidae

Genus *Dictyophimus* (Fig. 222, *d*)
Lattice skeleton multiple...................... Family 5 Botryoidae
Genus **Phormobothrys**

Suborder 4 **Triplylea** Hertwig

Without skeleton; with isolated spicules
Skeleton consists of radial hollow rods and fine tangential needles

Family 1 **Aulacanthidae**

Genus **Aulacantha** (Fig. 223, a)

With foreign skeletons covering body surface

Family 2 **Caementellidae**

Genus **Caementella** (Fig. 223, b)

With skeleton
1–2 (concentric) usually spherical skeletons
Outer lattice skeleton with triangular or areolar meshes

Family 3 **Sagosphaeridae**

Genus **Sagenoscene**

One lattice skeleton with hollow radial bars

Family 4 **Aulosphaeridae**

Genus **Aulosphaera** (Fig. 223, c)

2 concentric lattice skeletons connected by radial bars

Family 5 **Cannosphaeridae**

Genus **Cannosphaera**

One skeleton, simple, but variable in shape; bilaterally symmetrical
Skeleton with fine diatomaceous graining

Family 6 **Challengeridae**

Genus **Challengeron** (Fig. 224, a)

Skeleton smooth or with small spines

Family 7 **Medusettidae**

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Fig. 223. a, *Aulacantha scolymantha*, ×30 (Kühn); b, *Caementella stapedia*, ×65 (Haeckel); c, *Aulosphaera labradoriensis*, ×10 (Haecker).
Genus **Medusetta** (Fig. 224, b)

One skeleton; spherical or polyhedral, with an opening and with radiating spines.

Skeleton spherical or polyhedral, with uniformly large round pores

Family 8 Castanellidae

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**Fig. 224.** a, *Challengeron wyvillei*, ×105 (Haeckel); b, *Medusetta ansata*, ×230 (Borgert); c, *Castanidium murrayi*, ×25 (Haecker); d, *Circoporus octahedrus*, ×65 (Haeckel); e, *Tuscarora murrayi*, ×7 (Haecker); f, *Coelodendrum ramosissimum*, ×10 (Haecker).

Genus **Castanidium** (Fig. 224, c)

Skeleton similar to the last, but the base of each radial spine surrounded by pores.

Family 9 Circoporidae

Genus **Circoporus** (Fig. 224, d)

Skeleton flask-shaped with 1–2 groups of spines.

Family 10 Tuscaroridae

Genus **Tuscarora** (Fig. 224, e)

Central portion of skeleton consists of 2 valves

Valves thin, each with a conical process which divides into branched tubes.

Family 11 Coelodendridae

Genus **Coelodendrum** (Fig. 224, f)

**References**

——— (1904) II. Ibid., 19:733.
——— (1913) II. Ibid., 3:539.
——— (1887) II.
THE Sporozoa are without exception parasitic and bear spores. Their hosts are widely distributed in the animal kingdom, from Protozoa to Chordata. As a rule, they are incapable of locomotion, but some when immature may move about by pseudopodia or myonemes. They possess neither cilia nor flagella, except in the gamete stage. In the forms that are confined to one host, the spore is usually enveloped by a resistant membrane which would enable it to withstand unfavorable conditions while outside the host body, but in those having two host animals, as in Plasmodium, the sporozoite is naked. The method of nutrition is saprozoic or parasitic, the food being dissolved cytoplasm, tissue fluid, body fluid, or dissolved food material of the host.

Both asexual and sexual reproductions are well known in many species. Asexual reproduction is by repeated binary or multiple fission or budding of intracellular trophozoites. The multiple division in a host cell produces far greater number of individuals than that of protozoans belonging to other classes and often is referred to as schizogony. The sexual reproduction is by isogamous or anisogamous fusion or autogamy and marks in many cases the beginning of sporogony or spore-formation.

Schaudinn (1900) divided the Sporozoa into two groups, Telosporidia and Neosporidia, and this scheme has been followed by several authors. Some recent writers consider these two groups as separate classes. This, however, seems to be improper, as the basis of distinction between them is entirely different from that which is used for distinguishing the other four classes: Sarcodina, Mastigophora, Ciliata, and Suctoria. For this reason, the Sporozoa are placed in a single class and divided into three subclasses as follows:

Spore simple; without polar filament
Spore with or without membrane; with 1–many sporozoites
Subclass 1 Telosporidia
Spore with membrane; with one sporozoite
Subclass 2 Acnidosporidia (p. 635)
Spore with polar filament
Subclass 3 Cnidosporidia (p. 643)

Subclass 1 Telosporidia Schaudinn

The spore which contains neither a polar capsule nor a polar filament possesses one to several sporozoites and is formed at the end of the trophic life of the individual. In the forms which invade two host
animals to complete their development, there occur naked sporozoites instead of spores.

The infection of a new host begins with the entrance of mature spores through mouth, or with the introduction of the sporozoites by blood-sucking invertebrates directly into the blood stream. The sporozoites enter specific host cells and there grow at the expense of the latter. In the Coccidia and the Haemosporidia, the trophozoite continues its intracellular existence, but in the Gregarinida it leaves the host cell and grows in an organ cavity. Except Eugregarinina, the vegetative form undergoes schizogony and produces a large number of daughter individuals which invade new host cells, thus spreading the infection within the host body. The sporozoites finally develop into gametocytes. In the Coccidia and the Haemosporidia, anisogametes are, as a rule, produced. Each macrogametocyte develops into a single macrogamete and each microgametocyte, into several microgametes. Fusion of the gametes in pairs results in formation of a large number of zygotes, each of which develops either into one to many spores or into a number of naked sporozoites. In the Gregarinida, two fully mature trophozoites (or gametocytes) encyst together and the nucleus in each multiplies repeatedly to form numerous gametes, which fuse in pairs with those produced in the other individual within the common envelope. The zygotes develop into spores, each containing variable number of sporozoites. When these spores enter a new host, the changes outlined above are repeated. The Telosporidia are parasitic in vertebrates and higher invertebrates.

Three orders are distinguished in this subclass:

Mature trophozoite extracellular, large; zygote not motile; sporozoites enveloped...........................................Order 1 Gregarinida
Mature trophozoite intracellular, small
Zygote not motile; sporozoites enveloped...Order 2 Coccidia (p. 570)
Zygote motile; sporozoites naked....Order 3 Haemosporidia (p. 599)

Order 1 Gregarinida Lankester

The gregarines are chiefly coelozoic parasites in invertebrates, especially arthropods and annelids. They obtain their nourishment from the host organ-cavity through osmosis. The vast majority of gregarines do not undergo schizogony and an increase in number is carried on solely by sporogony. In a small group, however, schizogony takes place and this is used as the basis for grouping these protozoans into two suborders as follows:

No schizogony..............................Suborder 1 Eugregarinina (p. 528)
Schizogony occurs.......................Suborder 2 Schizogregarinina (p. 560)
Suborder 1 Eugregarinina Doflein

This suborder includes the majority of the so-called gregarines which are common parasites of arthropods. When the spore gains entrance into a suitable host, it germinates and the sporozoites emerge and enter the epithelial cells of the digestive tract. There they grow at the expense of the host cells which they leave soon and to which they become attached by various organellae of attachment (Fig. 235). These trophozoites become detached later from the host cells and move about in the lumen of the gut. This stage, sporadin, is ordinarily most frequently recognized. It is usually large and vermiform. The body is covered by a definite pellicle and its cytoplasm is clearly differentiated into the ectoplasm and endoplasm. The former contains myonemes (p. 62) which enable the organisms to undergo gliding movements (Watson, 1916).

In one group, Acephalina, the body is of a single compartment, but in the other group, Cephalina, the body is divided into two compartments by an ectoplasmic septum. The smaller anterior part is the protomerite and the larger posterior part, the deutomerite, contains a single nucleus. In Pileocephalus (Fig. 236, S) the nucleus is said to be located in the protomerite and according to Goodrich (1938) both the protomerite and deutomerite of Nina gracilis contain a nucleus. The endoplasm contains numerous spherical or ovoidal bodies which are called zooamylon or paraglycogen grains and which are apparently reserve food material (p. 112). The protomerite may possess an attaching process with hooks or other structures at its anterior border; this is called the epimerite. The epimerite is usually not found on detached sporadins. Goodrich observed recently that in Nina the protomerite is a knob-like part of the gregarine when contracted, but expands freely and used as a mobile sucker for attachment to the gut epithelium of the host Scolependra. Presently multiple filiform epimerite grows at the free edge of the sucker and penetrates between the host cells. Epimerite bearing trophozoites are called cephalins. Cytology (Göhre, 1943).

Many gregarines are solitary, others are often found in an endwise association of two or more sporadins. This association is called syzygy. The anterior individual is known as the primite and the posterior, the satellite. What differences exist between the two individuals that become associated is not well known. But Mühl (1921) reported in Gregarina cuneata, the granules in the primite and the satellite stained differently with neutral red. Sporadins usually encyst in pairs and become gametocytes. This process following bisassociation was observed in a number of species; for example, in
Leidyana erratica (Watson, 1916), Gregarina blattarum (Sprague, 1941) (Fig. 226), etc. Within the cyst-membrane, the nucleus in each individual undergoes repeated division, forming a large number of small nuclei which by a process of budding transform themselves into numerous gametes. The gametes may be isogamous or anisogamous. Each of the gametes in one gametocyte appears to unite with one formed in the other, so that a large number of zygotes are produced. In some species such as Nina gracilis the microgametes enter the individual in which macrogametes develop, and the development of zygotes takes place, thus producing the so-called pseudocyst. The zygote becomes surrounded by a resistant membrane and its content

Fig. 225. Diagram illustrating the developmental cycle of Lankesteria culicis (Wenyon). a, entrance of sporozoite into the mid-gut epithelium and growth of trophozoites; b, mature trophozoite found in the lumen of gut; c, association of two gametocytes prior to encystment; d–f, gamete formation; g, zygote formation; h, development of spores from zygotes; i, a spore; j, emergence of eight sporozoites from a spore in a new host gut.
develops into the sporozoites, thus developing into a spore. The spores germinate when taken into the alimentary canal of a host animal and the life-cycle is repeated.

According to Wenyon, in a typical Eugregarinina, *Lankesteria culicis* (Fig. 225) of *Aedes aegypti*, the development in a new host begins when a larva of the latter ingests the spores which had been set free by infected adult mosquitoes in the water. From each spore are liberated 8 sporozoites (*j*), which enter the epithelial cells of the stomach and grow (*a*). These vegetative forms leave the host cells later and become mingled with the food material present in the stomach lumen of the host (*b*). When the larva pupates, the sporadins enter the Malpighian tubules, where they encyst (*c*). The repeated nuclear division is followed by formation of large numbers of gametes (*d–f*) which unite in pairs (*g*). The zygotes thus formed develop into spores, each possessing 8 sporozoites (*h*). Meanwhile the host pupa emerges as an adult mosquito, and the spores which become set free in the lumen of the tubules pass into the intestine, from which they are discharged into water. Larvae swallow the spores and acquire infection.

Eugregarinina are divided into 2 tribes:

Trophozoite not septate.............................. Tribe 1 Acephalina (p. 531)
Trophozoite septate................................. Tribe 2 Cephalina (p. 541)

Fig. 226. Encystment in *Gregarina blattarum*, ×60 (Sprague). a, a trophozoite with epimerite and 3 pairs of syzygy; b, association of three individuals; c–h, encystment as seen in a single pair in about one hour.
Tribe 1 Acephalina Kölliker

The acephalines are mainly found in the body cavity and organs associated with it. The infection begins by the ingestion of mature spores by a host, in the digestive tract of which the sporozoites are set free and undergo development or make their way through the gut wall and reach the coelom or various organs such as seminal vesicles. Young trophozoites are intracellular, while more mature forms are either intracellular or extracellular. Acephaline gregarines (Berlin, 1924; Bhatia and Chatterjee, 1925; Bhatia and Setna, 1926; Bhatia, 1929; Troisi, 1933).

Spores with similar ends
  Spores biconical
    Sporadins solitary
      Anterior end not differentiated .............. Family 1 Monocystidae
      Anterior end conical or cylindro-conical ............ Family 2 Rhynchocystidae (p. 534)
    Sporadins in syzygy
      Spores with thickenings at ends. Family 3 Zygocystidae (p. 534)
      Spores without thickenings. Family 4 Aikinetocystidae (p. 535)
  Spores not biconical
    Spores navicular ................. Family 5 Stomatophoridae (p. 536)
    Spores round or oval
      No encystment ............ Family 6 Schaudinnellidae (p. 537)
      2 sporadins encyst together .... Family 7 Diplocystidae (p. 538)
  Spores with dissimilar ends
    Spores with epispore ............. Family 8 Urosporidae (p. 538)
    Spores without epispore ........ Family 9 Allantocystidae (p. 540)
  Spores unobserved; grown trophozoites with cup-like depression at posterior end for syzygy ........ Family 10 Ganymedidae (p. 541)

Family 1 Monocystidae Bütschli

Trophozoites spheroidal to cylindrical; anterior end not differentiated; solitary; spores biconical, without any spines, with 8 sporozoites.

Genus Monocystis Stein. Trophozoites variable in form; motile; incomplete sporulation in cyst; spore biconical, symmetrical; in coelom or seminal vesicles of oligochaetes. Numerous species (Berlin, 1924).

*M. ventrosa* Berlin (Fig. 227, a-e). Sporadins 109–183μ by 72–135μ; nucleus up to 43μ by 20μ; cysts 185–223μ by 154–182μ; spores 17–25μ by 8–19μ; in *Lumbricus rubellus*, *L. castaneus* and *Helodrilus foetidus*.

*M. lumbrici* Henle (Fig. 227, d, e). Sporadins about 200μ by 60–70μ; cysts about 162μ in diameter; in *Lumbricus terrestris*, *L. rubellus*, and *L. castaneus* (Berlin, 1924).
M. rostrata Mulsow (Figs. 92, 228). Elongate oval; average dimensions 450\(\mu\) by 220\(\mu\); anterior end often drawn out into a process; pellicle thick, longitudinally striated; cysts about 750\(\mu\) in diameter; spores 23\(\mu\) by 9\(\mu\); in the seminal vesicles of Lumbricus terrestris. Mulsow (1911) found vegetative stages in autumn and winter and sporogony in spring. Meiosis in the last pre-gametic division (p. 207).

Fig. 227. a–c, Monocystis ventrosa (a, \(\times260\); b, \(\times150\); c, \(\times830\)) (Berlin); d, e, M. lumbrici, \(\times280\) (Berlin); f. Apolocystis gigantea, \(\times90\) (Troisi); g, A. minuta, with attached phagocytes, \(\times770\) (Troisi); h, Nematocystis vermicularis, \(\times80\) (Hesse); i, j, Rhabdocystis claviformis (i, \(\times220\); j, \(\times270\)) (Boldt); k, l, Enterocystis ensis (k, \(\times140\)) (Zwetkow).

Genus Apolocystis Martii. Trophozoites spherical; without principal axis marked by presence of any special peripheral organ; solitary; spore biconical; in seminal vesicles or coelom of various oligochaetes. Many species.

A. gigantea Troisi (Fig. 227, f). In seminal vesicles of Helodrilus foetidus and Lumbricus rubellus; late October to March only; fully
grown trophozoites 250–800\(\mu\) in diameter; whitish to naked eyes; pellicle thickly covered by 10–15\(\mu\) long 'hairs'; endoplasm packed with spherical paraglycogen grains (3\(\mu\) in diameter); nucleus 35–43\(\mu\) in diameter; cysts 400–800\(\mu\) in diameter; spores 19\(\mu\) by 8.6\(\mu\) (Troisi, 1933).

A. minuta Troisi (Fig. 227, g). In seminal vesicles of *Lumbricus terrestris*, *L. castaneus* and *L. rubellus*; mature trophozoites 40–46\(\mu\) in diameter; endoplasm yellowish brown, packed with spherical paraglycogen grains (5.3–7\(\mu\) in diameter); nucleus 10\(\mu\) in diameter; cysts 68–74\(\mu\) by 55–65\(\mu\); spores of 3 sizes, 11\(\mu\) by 5.5\(\mu\), 18.8\(\mu\) by 7\(\mu\) and 21.6\(\mu\) by 9.8\(\mu\).

Genus Nematocystis Hesse. Trophozoites elongate, cylindrical and shaped like a nematode; solitary. Many species (Bhatia and Chatterjee, 1925).

![Fig. 228. Monocystis rostrata (Mulsow). a–c, trophozoites, \(\times90\); d, spore, \(\times850\).](image)

*N. vermicularis* H. (Fig. 227, h). In seminal vesicles of *Lumbricus terrestris*, *L. rubellus*, *Helodrilus longus*, *Pheretima barbadensis*; trophozoites 1 mm. by 100\(\mu\); cylindrical, both ends with projections; nucleus oval; endoplasm alveolated, with paraglycogen grains; sporadins become paired lengthwise; cysts and spores unknown.

Genus Rhabdocystis Boldt. Trophozoites elongate, gently curved; anterior end swollen, club-shaped; posterior end attenuated; spores with sharply pointed ends. One species.

*R. claviformis* B. (Fig. 227, i, j). In seminal vesicles of *Octolasium complanatum*; sporadins extended, up to 300\(\mu\) by 30\(\mu\); pellicle distinctly longitudinally striated; zooamylon bodies 2–4\(\mu\) in diameter; cysts biscuit-form, 110\(\mu\) by 70\(\mu\); spores 16\(\mu\) by 8\(\mu\).

Genus Enterocystis Zwetkow. Early stages of trophozoites in syzygy; sporadins in association ensiform; cysts spherical without ducts; spores elongate ovoid, with 8 sporozoites; in gut of ephemerid larvae. Species (Noble, 1938a).
E. ensis Z. (Fig. 227, k, l). Sporadins in syzygy 200–510\(\mu\) long; cysts 200–350\(\mu\) in diameter; spores elongate ovoid; in gut of larvae of Caenis sp.

Genus Echinocystis Bhatia and Chatterjee. Body nearly spherical with two spine-like structures extending out from the body surface; solitary; spores biconical with equally truncated ends; in the seminal vesicles of earthworms (Bhatia and Chatterjee, 1925).

E. globosa B. and C. Body 740\(\mu\) by 65\(\mu\); spines sometimes unequally long; observations on spores incomplete; in the sperm sacs of Pheretima heterochaeta.

Family 2 Rhynchocystidae Bhatia

Trophozoites ovoid, spherical or elongate, with a conical or cylindro-conical trunk at anterior end; solitary; spore biconical, with 8 sporozoites.

Genus Rhynchocystis Hesse. Trophozoites ovoid or cylindrical; plastic epimerite, conical or cylindro-conical trunk; in seminal vesicles of oligochaetes. Many species (Bhatia and Chatterjee, 1925; Troisi, 1933).

R. pilosa Cuénot (Fig. 229, a). In seminal vesicles of Lumbricus terrestris, L. castaneus and Helodrilus foetidus; 217\(\mu\) by 25.5\(\mu\); pellicle with close, longitudinal ridges from which arise 'hairs' up to 40\(\mu\) in length; endoplasm viscous, packed with oval (3\(\mu\) by 2\(\mu\)) paraglycogen bodies; cysts ovoid, 95\(\mu\) by 84\(\mu\); spores 13.3\(\mu\) by 5\(\mu\) (Troisi, 1933).

R. porrecta Schmidt (Fig. 229, b, c). In seminal vesicles of Lumbricus rubellus and Helodrilus foetidus; extremely long with an enlarged head; up to 2.5 mm. by 32–36\(\mu\); sluggish; endoplasm granulated, filled with oval (4\(\mu\) by 2–3\(\mu\)) paraglycogen grains; nucleus 17–25\(\mu\) in diameter; spores 27.7–28\(\mu\) by 12\(\mu\); sporozoites 13–18\(\mu\) by 3–5\(\mu\) (Troisi, 1933).

Family 3 Zygocystidae Bhatia

Trophozoites in association; spores biconical, with peculiar thickenings at extremities; with 8 sporozoites; in seminal vesicles or coelom of oligochaetes.

Genus Zygocystis Stein. Sporadins pyriform, 2–3 in syzygy; in seminal vesicles or coelom of oligochaetes. Several species.

Z. wenrichi Troisi (Fig. 229, d, e). In seminal vesicles of Lumbricus rubellus and Helodrilus foetidus; sporadins up to 1.5 mm. by 250\(\mu\) in diameter; pellicle with longitudinal ridges which become free and form a ‘tuft of hairs’ at the posterior end; cysts 500–800\(\mu\) by 300–500\(\mu\); spores 28\(\mu\) by 13\(\mu\).
Genus **Pleurocystis** Hesse. Trophozoites in longitudinal or lateral association; spores biconical. One species.

*P. cuenoti* H. (Fig. 229, f). In the ciliated seminal horn of *Helodrilus longus* and *H. caliginosus*; 2 mm. by 300μ; pellicle striated longitudinally, obliquely near the posterior end; cysts 1.5–2 mm. in diameter; spores 28.5μ by 12μ (Hesse, 1909).

**Family 4 Aikinetocystidae** Bhatia

Trophozoites solitary or in syzygy; branching dichotomously, branches with sucker-like organellae of attachment; spores biconical.

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**Fig. 229** a, *Rhynchocystis pilosa*, ×200 (Hesse); b, c, *R. porrecta*: b, ×170 (Hesse); c, spore, ×1330 (Troisi); d, e, *Zygocystis wenrichi* (d, ×45; e, ×450) (Troisi); f, *Pleurocystis cuenoti*, ×190 (Hesse); g, h, *Aikinetocystis singularis* (h, ×320) (Gates); i–k, *Stomatophora coronata* (i, j, ×430; k, ×870) (Hesse); l, *Astrocystella lobosa*, ×120; m, *Craterocystis papua*, ×65; n, *Choanocystella tentaculata*, ×570; o, *Choanocystoides costaricensis*, ×470 (Martii).
Genus *Aikinetocystis* Gates. Trophozoites cylindrical or columnar, with a characteristic, regular dichotomous branching at attached end, with sucker-like bodies borne on ultimate branches; solitary or 2 (3–8) individuals in association; spores biconical.

*A. singularis* G. (Fig. 229, g, h). In coelom of *Eutypheoeus foveatus. E. rarus, E. peguanus* and *E. spinulosus* (of Burma); trophozoites up to 4 mm. long; number of branches 8 or 16, each with an irregular sucker; ovoid nucleus near rounded end; spores of two sizes, 20–23 μ long and 7–8 μ long; a few cysts found, ovoid and about 600 μ long.

Family 5 *Stomatophoridae* Bhatia

Trophozoites spherical to cylindrical or cup-shaped; with a sucker-like epimerite; solitary; spores navicular, ends truncate; 8 sporozoites; in seminal vesicles of *Pheretima* (Oligochaeta).

Genus *Stomatophora* Drzewiecki. Trophozoites spherical or ovoid: anterior end with a sucker-like epimeritic organella with a central spine; spores navicular. Several species.

*S. coronata* (Hesse) (Fig. 229, i–k). In seminal vesicles of *Pheretima rodericensis, P. hawayana* and *P. barbadensis*; trophozoites spherical, ovoid or elliptical, about 180 μ by 130 μ; endoplasm with ovoid paraglycogen grains; cysts ellipsoid or fusiform, 70–80 μ by 50–60 μ; spores in 2 sizes, 11 μ by 6 μ and 7 μ by 3 μ and in chain.

Genus *Astrocystella* Martii. Trophozoites solitary; stellate with 5–9 lobes radiating from central part containing nucleus; anterior surface with a depression. One species.

*A. lobosa* M. (Fig. 229, l). In seminal vesicles of *Pheretima beaufortii* (New Guinea); diameter about 200 μ; spores fusiform.

Genus *Craterocystis* M. Trophozoites solitary; rounded; a sucker-like depression on anterior end; myonemes well developed, running from concave to convex side. One species.

*C. papua* M. (Fig. 229, m). In prostate and lymphatic glands of *Pheretima wendessiana* (New Guinea); trophozoites about 360–390 μ in diameter.

Genus *Choanocystella* M. (*Choanocystis* M.). Trophozoites solitary; rounded or ovate; anterior end with a mobile sucker and a tentacle bearing cytoplasmic hairs; myonemes. One species.

*C. tentaculata* M. (Fig. 229, n). In seminal vesicles of *Pheretima beaufortii* (New Guinea); trophozoites 50 μ by 36 μ.

Genus *Choanocystoides* M. Trophozoites solitary, rounded or cup-shaped; anterior end with a mobile sucker, bordered by cytoplasmic filaments. One species.

*C. costaricensis* M. (Fig. 229, o). In seminal vesicles of *Pheretima*
heterochaeta (Costa Rica); trophozoites 40–45μ in diameter; nucleus ovoid, large, 12μ in diameter.

Genus Beccaricystis M. Mature trophozoites elongate, cylindrical, with a sucker-like depression at anterior end; nucleus at its bottom. One species.

*B. loriai* M. (Fig. 230, a). In seminal vesicles of *Pheretima sermowaiana*; trophozoites cylindrical, with wart-like growths, myo-

nemes run lengthwise with radially arranged transverse fibrils; about 100μ long.

Genus Albertisella M. Mature trophozoites cup-shaped, with anterior sucker with a smooth wall; nucleus at its bottom. One species.

*A. crater* C. In seminal vesicles of *Pheretima sermowaiana*.

Family 6 Schaudinnellidae Poche

Parasitic in the digestive system of oligochaetes; spores spherical; trophozoites do not encyst; male trophozoites producing microgam-
etes and female, macrogametes; zygotes or amphionts (spores) rounded.

Genus **Schaudinnella** Nusbaum. Trophozoites elongate spindle, free in lumen or attached to gut wall; sporadins male or female; spherical macrogametes and fusiform microgametes; zygotes or amphionts encapsulated, passed out of host or enter gut epithelium, dividing to produce many sporozoites (autoinfection).

*S. heneleae* N. (Fig. 230, b, c). In gut of *Henlea leptodera*; mature trophozoites about 70 µ by 9 µ; attached trophozoite with a clear wart-like epimerite; female and male sporadins; macrogametes, 5–7.5 µ in diameter; microgametes, spindle-form, 1–1.25 µ long; sporozoites rounded oval, 2.5–3 µ in diameter.

**Family 7 Diplocystidae** Bhatia

Coelomic or gut parasites of insects; trophozoites solitary or associated early in pairs; spores round or oval, with 8 sporozoites.

Genus **Diplocystis** Kunstler. Trophozoites spherical to oval; association of 2 individuals begin early in spherical form; spores round or oval, with 8 sporozoites; in the intestine and coelom of insects.

*D. schneideri* K. (Fig. 230, d, e). In the body cavity of *Periplaneta americana*; young stages in gut epithelium; cysts up to 2 mm. in diameter; spores 7–8 µ in diameter; sporozoites 8 µ long. Meiosis (p. 208).

Genus **Lankesteria** Mingazzini. Trophozoites more or less spatulate; spherical cyst formed by 2 laterally associated sporadins in rotation; spores oval, with flattened ends, with 8 sporozoites; in the gut of tunicates, flatworms and insects. Several species.

*L. culicis* (Ross) (Fig. 225). In gut and Malpighian tubules of *Aedes aegypti* and *A. albopictus*; mature trophozoites about 150–200 µ by 31–41 µ; cysts spherical, in Malpighian tubules of host, about 30 µ in diameter; spores 10 µ by 6 µ.

**Family 8 Urosporidae** Woodcock

Coelomonic parasites in various invertebrates; sporadins associative; spores with unequal ends; with or without epispores of various forms, with 8 sporozoites.

Genus **Urospora** Schneider. Large; frequently in lengthwise association of 2 individuals of unequal sizes, spores oval, with a filamentous process at one end; in body cavity or blood vessel of Tubifex, Nemertinea, Sipunculus, Synapta, and Chiridota. Several species.

*U. chiridotae* (Dogiel) (Fig. 230, f). In blood vessel of *Chiridota laevis* (in Canada); paired trophozoites up to about 1 mm. long; with stiff ‘hairs’ (Goodrich, 1925).
**U. hardyi** Goodrich. In the coelom of *Sipunculus nudus*; spores about 16\(\mu\) long, process 4–6\(\mu\) long, with eight sporozoites; thin-walled cysts 0.5–2 mm. in diameter; active phagocytosis by host cells of cysts and some trophozoites, producing brownish masses, 5 by 2 mm. or more in diameter, which are crowded together in the posterior region of the host.

Genus **Gonospora** Schneider. Trophozoites polymorphic, oval, pyriform or vermiform; cysts spherical; spore with a funnel at one end, rounded at the other; in gut, coelom or ova of polychaetes.

*G. minchini* Goodrich and Pixell-Goodrich (Figs. 230, g–i; 231, g). In coelom of *Arenicola ecaudata*; young trophozoites live in host eggs which float in the coelomic fluid; fully grown trophozoites leave eggs in which they grow up to 200\(\mu\) long, and encyst together in pairs; spores without well-developed funnel, 8–10\(\mu\) long (Goodrich and Goodrich, 1920).

Genus **Lithocystis** Giard. Trophozoites large, ovoid or cylindrical; attached for a long period to host tissue; pellicle with hairlike processes; endoplasm with calcium oxalate crystals; spores ovoid, with a long process at one end; in coelom of echinids.

*L. brachycercus* Goodrich (Fig. 231, a, b). In the coelom of *Chiridota laevis*; fully grown spherical trophozoites up to 200\(\mu\) in diameter; spores with a short flattened tail; in Canada (Goodrich, 1925).

*L. lankesteri* G. In the coelom of *Sipunculus nudus*; trophozoites covered with spinous structures; biassociative; spores 12–14\(\mu\) by 6–8\(\mu\); the long ribbon-like tail 50–60\(\mu\) long.

Genus **Pterospora** Racovitza and Labbé. Sporadins associative or solitary; free end drawn out into 4 bifurcated processes; cysts spherical or oval; spores with epispore drawn out into 3 lateral processes; in coelom of polychaetes.

*P. maldaneorum* R. and L. (Fig. 231, c, d). In the coelom of *Liocephalus liopygue*; trophozoites about 140\(\mu\) long; cysts 288\(\mu\), by 214\(\mu\); epispore 24\(\mu\) in diameter; endospore 10–14\(\mu\) by 3–4\(\mu\).

Genus **Ceratospora** Léger. Sporadins elongate conical, head to head association; without encystment; spores oval with a small collar at one end and 2 divergent elongate filaments at other. One species.

*C. mirabilis* L. (Fig. 231, e, f). Sporadins 500–600\(\mu\) long; spore 12\(\mu\) by 8\(\mu\), filaments 34\(\mu\) long; in general body cavity of *Glycera* sp.

Genus **Cystobia** Mingazzini. Trophozoites, large, irregular; fully grown forms always with 2 nuclei, due to early union of 2 individuals; spores oval, membrane drawn out and truncate at one end; in blood vessels and coelom of Holothuria.
**C. irregularis** (Minchin) (Fig. 231, h, i). Trophozoites irregular in form; up to 500μ long; endoplasm opaque, granulated; cysts in connective tissue of vessels; spore ovoid, epispore bottle-like, 25μ long; in blood vessel of *Holothuria nigra*.

**Family 9 Allantocystidae** Bhatia

Trophozoites elongate cylindrical; cysts elongate, sausage-like; spores fusiform, sides slightly dissimilar.

Genus *Allantocystis* Keilin. Sporadins, head to head association; cysts sausage-like; in dipterous insect. One species.

*A. dasyhelei* K. (Fig. 231, j–m). In gut of larval *Dasyhelea obscura*;
full-grown sporadins \( 65-75\mu \) by \( 20-22\mu \); cysts \( 140-150\mu \) by \( 20\mu \); spores \( 18\mu \) by \( 6.5\mu \) (Keilin, 1920).

**Family 10 Ganymedidae** Huxley

Trophozoites only known; mature individuals biassociative; posterior end of primate with a cup-like depression to which the epimeritic organella of satellite fits; cysts spherical; spores unknown.

**Genus Ganymedes** Huxley. Characters of the family; Huxley considers it as an intermediate form between Acephalina and Cephalina.

*G. anaspides* H. (Fig. 231, n). In gut and liver-tube of the crustacean, *Anaspides tasmaniae* (of Tasmania); trophozoites in association, \( 70-300\mu \) by \( 60-130\mu \); cysts \( 85-115\mu \) in diameter.

**Tribe 2 Cephalina** Delage

The body of a trophozoite is divided into the protomerite and deutomerite by an ectoplasmic septum; inhabitants of the alimentary canal of invertebrates, especially arthropods. Taxonomy and distribution (Watson, 1916; Pinto, 1919; Kamm, 1922, 1922a).

One host species involved

None-septate; epimerite a knob....... Family 1 Lecudinidae (p. 542)

Development intracellular

Sporadins associative.......Family 2 Cephaloidophoridae (p. 543)

Sporadins solitary.............Family 3 Stenophoridae (p. 544)

Development extracellular

Sporadins associative

Satellite non-septate.............Family 4 Didymophyidae (p. 544)

Satellite septate...............Family 5 Gregarinidae (p. 544)

Sporadins solitary

Epimerite simple knob-like

Cysts with several ducts.......Family 6 Leidyanidae (p. 547)

Cysts without or with one duct...........................................

............................................Family 7 Monoductidae (p. 548)

Epimerite not simple knob-like

Epimerite cup-shaped or digitate

Epimerite cup-shaped......Family 8 Menosporidae (p. 549)

Epimerite digitate.......Family 9 Daetylophoridae (p. 550)

Epimerite otherwise

Spore hat-shaped.......Family 10 Stylocephalidae (p. 552)

Spore of other shapes

Spore with spines..Family 11 Acanthosporidae (p. 554)

Spore without spines...........................

............................................Family 12 Actinocephalidae (p. 554)

Two host species involved............Family 13 Porosporidae (p. 558)
Family 1 Lecudinidae Kamm

Epimerite simple, symmetrical; non-septate; spores ovoidal, thickened at one pole; solitary; in gut of polychaetes and termites. Undoubtedly intermediate forms between Acephalina and Cephalina.

Genus Lecudina Mingazzini. Epimerite simple, knob-like; in polychaetes. Species (Kamm, 1922).

Fig. 232. a, Lecudina pellucida (Kölliker); b, Polyrhabdina spionis, $\times 800$ (Reichenow); c, Sycia inspinata (Léger); d, e, Zygosoma globosum (d, $\times 60$; e, $\times 1260$) (Noble); f, Cephaloidophora olivia, $\times 190$ (Kamm); g, Stenophora larvata, $\times 50$ (Leidy); h, S. robusta, $\times 130$ (Ellis); i, j, Fonsecaia polymorpha (i, $\times 220$; j, $\times 430$) (Pinto); k, Gregarina blattarum, $\times 55$ (Kudo); l, G. locustae, $\times 65$ (Leidy); m, G. oviceps, $\times 30$ (Crawley); n, Protomagalhaesia serpentula, $\times 35$ (Pinto); o, Gamocystis tenax (Schneider).
L. pellucida (Kölliker) (Fig. 232, a). In Nereis cultrifera and N. beaucourdrayi; trophozoites ellipsoid; spores 7 \( \mu \) by 5 \( \mu \) (Ellis, 1913).

Genus Polyrhabdina Mingazzini. Trophozoites flattened, ovoidal; epimerite with a corona of processes with split ends, deeply stainable; in polychaetes (Spionidae).

P. spionis (Kölliker) (Fig. 232, b). In Scololepis fuligionosa; 100 \( \mu \) by 35 \( \mu \); epimerite with a corona of 8–10 processes; cysts unknown. Mackinnon and Ray (1931) report var. bifurcata, the epimerite of which is a "knob-shaped structure with a circket of 14 to 16 minute teeth at its base, and at its crown, two much larger, diverging, claw-like processes."

Genus Kofoïdina Henry. Epimerite rudimentary; development intracellular; 2–14 sporadins in association; cysts and spores unknown (Henry, 1933).

K. ovata H. In midgut of Zootermopsis angusticollis and Z. nevadensis; syzygy 153–672 \( \mu \) long; sporadins 41–105 \( \mu \) long.

Genus Sycia Léger. Epimerite knobbed, bordered by a thick ring; protomerite subspherical; deutomerite conical, with navicular inclusions; in marine annelids (Léger, 1892).

S. inspinata L. (Fig. 232, c). In Audouinia lamarcki.

Genus Zygosoma Labbé. Trophozoites with wart-like projections; epimerite a simple knob; spores oval; in gut of marine annelids.

Z. globosum Noble (Fig. 232, d, e). Trophozoites 250–500 \( \mu \) by 200–380 \( \mu \); epimerite a large globule; cysts 400 \( \mu \) by 360 \( \mu \), without ducts; spores oval, with 4 sporozoites, 9 \( \mu \) by 7 \( \mu \); reduction zygotice, 12 to 6 chromosomes; in gut of Urechis caupo in California.

Genus Ulivina Mingazzini. Elongate ellipsoid; epimerite simple; spores unknown; in gut of polychaetes.

U. rhynchoboli (Crawley). Sporadins up to 700 \( \mu \) long; in the gut of Rhynchobolus americanus (Crawley, 1903).

Family 2 Cephaloidophoridae Kamm

Development intracellular; early association; cysts without sporoducts; spores ovoidal, with equatorial line; in gut of Crustacea.

Genus Cephaloidophora Mawrodiadi. Sporadins biassociative, early; epimerite rudimentary; cysts without sporoducts; spores in chain, ovoidal.

C. olivia (Watson) (Fig. 232 f,). Biasssociated sporadins up to 218 \( \mu \) long; individuals up to 118 \( \mu \) by 36 \( \mu \); cysts spheroidal, 60 \( \mu \) in diameter; spores (?); in gut of Libinia dubia; Long Island.

C. nigrofusca (Watson). Sporadins, ovoid to rectangular, up to 125 \( \mu \) by 75 \( \mu \); cysts and spores (?); in gut of Uca pugnax and U. pugilator.
Family 3 *Stenophoridae* Léger and Duboscq

Development intracellular; sporadins solitary; with a simple epimerite or none; cysts open by rupture; spores ovoid, with or without equatorial line, not extruded in chain; in Diplopoda.

Genus *Stenophora* Labbé. With or without simple epimerite; spores ovoid with equatorial line, not in chain. Species (Watson, 1916; Pinto, 1919).

*S. larvata* (Leidy) (Fig. 232, g). Sporadins up to 800μ by 23μ; protomerite small; in gut of *Spirobolus spinigerus*.

*S. robusta* Ellis (Fig. 232, h). Sporadins 140–180μ by 67μ; cysts and spores both unobserved; in gut of *Parajulus venustus, Orthomorpha gracilis* and *O*. sp.; Colorado.

Genus *Fonsecaia* Pinto. Spores elongate ovoid; without equatorial line.

*F. polymorpha* Pinto (Fig. 232, i, j). Sporadins 170μ long; spores 18μ by 8μ; in gut of *Orthomorpha gracilis*; Brazil.

Family 4 *Didymophyidae* Léger

Two to three sporadins in association; satellite without septum.

Genus *Didymophyes* Stein. Epimerite a small pointed papilla; cysts spherical, open by rupture; spores ellipsoidal.

*D. gigantea* S. Sporadins slender, 1 cm. by 80–100μ; 2 deutomerites; cysts spherical, 600–700μ in diameter; spores oval, 6.5μ by 6μ; in gut of larvae of *Oryctes nasicornis, O*. sp., and *Phyllognathus* sp. (Léger, 1892).

Family 5 *Gregarinidae* Labbé

Sporadins in association; epimerite simple, symmetrical; cysts with or without ducts; spores symmetrical.

Genus *Gregarina* Dufour. Sporadins biassociative; epimerite small, globular or cylindrical; spores dolioform to cylindrical; cysts open by sporoducts; in the gut of arthropods. Numerous species (Watson, 1916). Morphology and physiology (Mühl, 1921).

*G. blattarum* Siebold (Figs. 226; 232, k). Sporadins in syzygy, 500–1100μ; by 160–400μ; cysts spherical or ovoidal; eight to 10 sporoducts; spores cylindrical to dolioform, truncate at ends, 8–8.5μ by 3.5–4μ; in the midgut of cockroaches, especially *Blatta orientalis*. Reproduction (Schiffmann, 1919; Sprague, 1941).

*G. locustae* Lankester (Fig. 232, l). Sporadins 150–350μ long; in *Dissosteria carolina*.

*G. oviceps* Diesing (Fig. 232, m). Sporadins up to 500μ by 225μ; in syzygy; spherical cysts 250μ in diameter; two to five sporoducts up
SPOROZOA, GREGARINIDA

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to 1 mm. long; spores dolioform, 4.5 μ by 2.25 μ; in Gryllus abbreviatus and G. americanus (Leidy, 1853).

G. polymorpha (Hammerschmidt). Cylindrical sporadins up to 350 μ by 100 μ; in syzygy; protomerite dome-shaped; deutomerite cylindrical, rounded posteriorly; a small nucleus with an endosome; in the intestine of larvae and adults of Tenebrio molitor (“meal worm”).

G. rigida (Hall). Sporadins 28 μ by 20 μ up to 424 μ by 196 μ; syzygy; spherical cysts 212–505 μ in diameter; in the species of Melanoplus (grasshoppers) (Kararo, 1920; Allegre, 1948).

Genus Protomagalhaesia Pinto. Sporadins cylindrical; in syzygy, protomerite of satellite draws in the posterior end of primite; cysts without ducts; spores dolioform, with spines at ends.

P. serpentula (Magalhães) (Fig. 232, n). Sporadins up to 1.2 mm. by 180 μ; in gut and coelom of Blatta orientalis.

Genus Gamocystis Schneider. Septate only in trophozoites; sporadins non-septate; in syzygy; spore formation partial; with sporoducts; spores cylindrical. A few species.

G. tenax S. (Fig. 232, o). Association head to head; spherical cysts with 15 or more ducts; spore cylindrical, with rounded ends; in gut of Blattella lapponica (Schneider, 1875).

Genus Hyalospora Schneider. Sporadins in syzygy; cytoplasm yellowish orange; epimerite a simple knob; cysts open by rupture; spores fusiform.

H. affinis S. Trophozoites 300 μ long; cysts, yellow, 60 μ in diameter; spores 8.7 μ by 6 μ; in gut of Machilis cylindrica (Labbé, 1899).

Genus Tettigonospora Smith. Similar to Hyalospora, but cytoplasm opaque white; spores spherical. One species (Smith, 1930).

T. stenopelmati S. Sporadins 225–542 μ by 118–225 μ; spherical cysts 434–551 μ in diameter, wall 17–66 μ thick; spores 4.8–5 μ in diameter; in the midgut of Stenopelmatus fuscus and S. pictus (“Jerusalem crickets”).

Genus Hirmocystis Labbé. Sporadins associative, 2–12 or more; with a small cylindrical papilla-like epimerite; cysts without ducts; spores ovoidal.

H. harpali Watson (Fig. 233, a). Total length of association up to 1060 μ; sporadins up to 560 μ by 80 μ; cysts unknown; in gut of Harpalus pennsylvanicus erythropus (Watson, 1916).

H. termitis (Leidy) (Fig. 233, b). Association 614–803 μ long; epimerite simple sphere; cysts rare; spores (?); in Zootermopsis angusticollis, Z. nevadensis, etc. (Henry, 1933).

Genus Uradiophora Mercier. Sporadins in syzygy; deutomerite
with small process; epimerite an elongate papilla; cysts oval without ducts; spores spherical, in chains (Mercier, 1911).

U. cuenoti M. (Fig. 234, a). 2–4 sporadins in syzygy; individuals up to 700μ long; cysts ovoid, 44μ long; spores 4μ in diameter; in gut of Atyaephyra desmaresti.

Genus Pyxinioides Trégouboff. Sporadins biassociative; epimerite with 16 longitudinal furrows, small cone at end.

P. balani (Kölliker). Primites up to 130μ; satellite 60μ long; in gut of Balanus amphitrite and B. eburneus.

Genus Anisolobus Vincent. Sporadins in syzygy; epimerite lacking; protomerite of primites expanded to form sucker-like organella.
SPOROZOA, GREGARINIDA

Cysts ellipsoid, with thick envelope; with 6–8 sporoducts; spores barrel-shaped. One species.

*A. dacnecola* V. (Fig. 233, c). In the midgut of the coleopteran *Dacne rufifrons*; 2 sporadins in syzygy 100–300 μ by 20–50 μ; cysts without envelope, 130–150 μ by 80–90 μ; sporoducts 40–50 μ long; spores in chain, dolioform, 6 μ by 4 μ (Vincent, 1924).

Genus *Carcinoecetes* Ball. Sporadins in syzygy of 2 or more individuals; epimerite rudimentary; cysts without sporoducts; spores round to oval, not in chain; in gut of Crustacea (Ball, 1938).

*C. hesperus* B. (Fig. 233, d, e). 2–6 sporadins in association; sporadins up to 320 μ by 9 μ; cysts about 140 μ by 123 μ, attached to the wall of hindgut; spores 8.6 μ by 7.7 μ, with 8 radially arranged sporozoites; in gut of the striped shore crab, *Pachygrapsus crassipes*; in California.

*C. bermudensis* B. In the mid- and hind-gut of *Pachygrapsus transversus*; in Bermuda (Ball, 1951).

*C. mithraxi* B. In the gut of *Mithrax forceps*; in Bermuda.

*C. calappae* B. In the gut of *Calappaflammca*; in Bermuda.

Genus *Heliospora* Goodrich. Elongated, septate; spores more or less spherical, with equatorial ray-like processes (Goodrich, 1949).

*H. longissima* (Siebold) (Fig. 234, b–e). Trophozoites elongate filiform, up to 228 μ long; no intracellular stage; epimerite small, and is retained until the sporadins roll up for encystment; spherical cyst thinly walled and ruptures easily; microgametes flagellated; spores 7–8 μ in diameter, with eight sporozoites and bear six long ray-like processes at the equator; in the gut of *Gammarus pulex*.

Genus *Rotundula* Goodrich. Rotund; button-like epimerite; precocious association; cyst without duct; spores, small, spherical or subspherical (Goodrich, 1949).

*R. gammari* (Diesing) (Fig. 234, f). Cysts 40–50 μ; microgametes flagellate, 4 μ in diameter; spores spherical, 5–6 μ in diameter; in the gut of *Gammarus pulex*.

Family 6 *Leidyanae*

Similar to the last two families; but sporadins are solitary and epimerite simple knob-like; cysts with several sporoducts.

Genus *Leidyana* Watson. Solitary; epimerite a simple globular sessile knob; cysts with ducts; spores dolioform (Watson, 1915).

*L. erratica* (Crawley) (Fig. 233, f). Sporadins up to 500 μ by 160 μ; cysts about 350 μ in diameter; membrane about 30 μ thick; 1–12 sporoducts; spores extruded in chains, 6 μ by 3 μ; in gut of *Gryllus abbreviatus* and *G. pennsylvanicus*. 
Family 7 Monoductidae Ray and Chakravatry

As in the last family solitary; but cyst with a single sporoduct or none; spore with 8 sporozoites.

Genus Monoductus R. and C. Sporadins solitary; epimerite a small elevation with prongs attached to its base; anisogamy; cyst with a single sporoduct; spores flattened fusiform, with dissimilar ends, each with 8 sporozoites. One species.

*M. lunatus* R. and C. (Fig. 233, *m-o*). Cephalins 225–445 μ by 33–47 μ; epimerite with about 16 prongs; nucleus parachute-shaped, with myonemes attached at posterior margin; sporadins develop posterior pseudopodial processes before association; cysts spherical, 225–230 μ in diameter, voided by host; development completed in 3–4 days outside the host body, with one duct; spores 10.25 μ by 4 μ, truncate at one end, attenuated at other and discharged in a single chain; in gut of *Diplopoda* sp.

Genus Sphaerocystis Léger. Sporadins solitary; without protomerite; spherical.

*S. simplex* L. Sporadins 100–140 μ in diameter; protomerite in young trophozoites; spherical cysts in which individuals are not associative, 100 μ in diameter; spores ovoid, 10.5 μ by 7.5 μ; in gut of *Cyphon pallidulus*.

Genus Lepismatophila Adams and Travis. Epimerite a simple knob; cysts without ducts; spores ellipsoidal, smooth, in chain. One species (Adams and Travis, 1935).

*L. thermobiae* A. and T. (Fig. 233, *g–i*). Sporadins 67–390 μ by 30–174 μ; cysts white to black, ellipsoidal to subspherical, 244–378 μ by 171–262 μ; spores brown, 13.6 μ by 6.8 μ; in the ventriculus of the firebrat, *Thermobia domestica*.

Genus Colepismatophila Adams and Travis. Similar to the last genus; but larger; spores in wavy chains, hat-shaped, with 2 curved filamentous processes attached at opposite ends. One species.

*C. watsonae* A. and T. (Fig. 233, *j–l*). Sporadins 92–562 μ by 55–189 μ; cysts 226–464 μ by 158–336 μ; spores 16.5 μ by 9.7 μ, processes 21 μ long; in ventriculus of *Thermobia domestica* (Adams and Travis, 1935).

Genus Hyalosporina Chakravarty. Sporont solitary; epimerite small, tongue-like; anisogametes; cyst without ducts; spores oval, with a hyaline membrane. One species (Chakravarty, 1935, 1936).

*H. cambolopsisae* C. (Fig. 234, *g–j*). Trophozoites 247–1111 μ by 37–111 μ; cysts oval, 292–390 μ by 263–375 μ; spores 8 μ; by 6 μ; in the gut of the milliped, *Cambolopsis* sp.
Fig. 234. a, Uradiophora cuenoti in syzygy, ×65 (Mercier); b–e, Heliospora longissima (Goodrich) (b, a pair in syzygy, ×330; c, microgamete; d, zygote; e, a spore with 4 nuclei, ×2665); f, Rotundula gamnari in syzygy, ×330 (Goodrich); g–j, Hyalosporina cambolopsisae (Chakravarty) (g, intracellular trophozoite, ×1110; h, a mature individual with fibrils tethering the nucleus, ×120; i, anterior part of an attached organism, ×2330; j, a spore, ×1110); k, the digestive tube of Nepa cinerea with eight trophozoites attached to the stomach (opened) epithelium and three cysts of Coleorhynchus heros (Poisson).

Family 8 Menosporidae Léger

Sporadins solitary; epimerite a large cup, bordered with hooks, with a long neck; cysts without sporoducts; spores crescentic, smooth.

Genus Menospora Léger. With the characters of the family.

M. polyacantha L. (Fig. 235, a, b). Sporadins 600–700μ long; cysts 200μ in diameter; spores 15μ by 4μ; in gut of Agrion puella.
Family 9 Dactylophoridae Léger

Sporadins solitary; epimerite complex, digitate; cysts dehiscence by pseudocyst; spores cylindrical; in gut of chilopods.

Genus Dactylophorus Balbiani. Protomerite wide, bordered by digitiform processes; spores cylindrical.

*D. robustus* Léger (Fig. 235, c, d). Sporadins 700–800μ long; cysts spherical, 200μ in diameter; spores 11μ by 4.3μ; in gut of Cryptops hortensis.

Genus Echinomera Labbé. Epimerite an eccentric cone with 8 or more digitiform processes; cysts without sporoducts; spores cylindrical.

*E. magalhaesi* (Pinto) (Fig. 235, e). Sporadins up to 300μ by 70μ; in gut of Scolopendra sp.

Genus Rhopalonia Léger. Epimerite spherical, with 10 or more digitiform processes; pseudocysts; spores cylindrical.

*R. hispida* (Schneider) (Fig. 235, f, g). Endoplasm yellowish orange; cysts 200–250μ in diameter; spores 16μ by 6.5μ; in gut of Geophiles sp. and Stigmatogaster gracilis.

Genus Dendrorynchus Keilin. Elongate; epimerite a disc, surrounded by numerous ramified papillae; transverse fibrils conspicuous; cysts elliptical; spores fusiform.

*D. system K.* (Fig. 235, h). Sporadins 255μ by 18.5–20μ; spores 18–19μ by 7μ; in midgut of larvae of Systenus sp., a dolichopodid fly, found in decomposed sap of elm tree.

Genus Trichorhynchus Schneider. Protomerite prolonged anteriorly into a long neck, dilated at tip; pseudocyst; spores cylindrical to ellipsoidal.

*T. pulcher* S. (Fig. 235, i). Cysts 303–316μ in diameter; spores 9.7μ by 5.8μ; in gut of Scutigera sp. and S. forceps (Watson, 1916).

Genus Nina Grebnecki (Pterocephalus Schneider). Protomerite made up of 2 long narrow horizontal lobes fused and upturned spirally at one end, peripheral portion with many teeth, from which project long filaments; spores in chain; in gut of myriapods. Species (Watson, 1916).

*N. gracilis* G. (Fig. 235, j, k). 1.5–5 mm. long; cyst spherical; spores ellipsoidal; in the gut of Scolopendra cingulata and S. subspinipes (Goodrich, 1938).

Genus Seticephalus Kamm. Protomerite with closely set brush-like bristles.

*S. elegans* (Pinto) (Fig. 235, l). Sporadins up to 75μ by 35μ; cysts and spores unknown; in gut of Scolopendra sp.

Genus Acutispora Crawley. Solitary; pseudocyst; spore biconical,
with a thick blunt endosporal rod at each end. One species (Crawley, 1903).

*A. macrocephala* C. (Fig. 235, *m*). Sporadins up to 600μ long; cysts spherical, 410μ in diameter; spores navicular, slightly curved, 19μ by 4μ; in gut of *Lithobius forficatus*.

![Diagram of Sporozoa, Gregarinida](image)

FIG. 235. *a, b, Menospora polyacantha* (Léger); *c, d, Dactylophorus robustus* (*c, X130; d, X900*) (Léger); *e, Echinomera magalhaesi, X130* (Pinto); *f, g, Rhopalonia hispida* (*g, X830*) (Léger); *h, Dendrorhynchus systemi, X770* (Keilin); *i, Trichorhynchus pulcher* (Schneider); *j, k, Nina gracilis* (*j, X10*) (Schneider); *l, Seticephalus elegans, X450* (Pinto); *m, Acutispora macrocephala, X65* (Crawley); *n, Metamera schubergi, X270* (Duke); *o, p, Hentschelia thalassemae* (*o, X230; p, X620*) (Mackinnon and Ray); *q, r, Lecythion thalassemiae* (*q, X270; r, X930*) (Mackinnon and Ray).

*M. schubergi* D. (Fig. 235, n). Sporadins 150μ by 45μ; spores 9μ by 7μ; in gut of the leeches, *Glossosiphonia complanata* and *Placobdella marginata*.

*M. reynoldsi* Jones. Sporadins with epimerite measure 280μ by 50μ; cysts spherical; dehiscence by rupture; spore biconical, 5μ by 3μ, with 8 sporozoites; in the stomach diverticula and intestine of *Glossosiphonia complanata*.

Genus *Hentschelia* Mackinnon and Ray. Epimerite with a short neck, umbrella-like with its margin divided into 4–5 lobes, each fluted on anterior surface; 2 sporadins encyst together; gametes anisogamous; flagellate and non-flagellate; zygote gives rise to a spherical spore with 8 sporozoites. One species.

*H. thalassemae* M. and R. (Fig. 235, o, p). Cephalins 75–98μ by 30–45μ; in gut of *Thalassemia neptuni* (Mackinnon and Ray, 1931).

Genus *Lecythion* Mackinnon and Ray. Epimerite a low cone, surrounded by 14–15 petal-shaped lobes, with a neck; cysts and spores unknown.

*L. thalassemae* M. and R. (Fig. 235, q, r). Cephalins 135μ by 52μ; epimerite about 27μ long; in gut of *Thalassemia neptuni*.

Family 10 *Stylocephalidae* Ellis

Sporadins solitary; epimerite varied; pseudocysts; hat-shaped spores in chains.

Genus *Stylocephalus* Ellis. Epimerite nipple-like; cysts covered with papillae; in arthropods and molluscs.

*S. giganteus* E. (Fig. 236, a). Sporadins 1.2–1.8 mm. long; cysts spherical, 450μ in diameter; spores subspherical black, 11μ by 7μ; in *Eleodes* sp., *Asida opaca*, *A*. sp., and *Eusattus* sp. (Coleoptera) (Ellis, 1912).


*B. elongatus* W. (Fig. 236, b). Sporadins up to 1.6 mm. by 50μ; nucleus diagonal; cysts and spores unknown; in gut of Cucujus larva (a coleopteran).

Genus *Sphaerorhynchus* Labbé. Epimerite a small sphere at end of a long neck.

*S. ophioides* (Schneider). Cephalins 1.3 mm. long; epimerite 220μ long; terminal part 8.5μ; sporadins 3–4 mm. long; in gut of *Acis* sp.
Genus *Cystocephalus* Schneider. Epimerite a large lance-shaped papilla with a short neck; spore hat-shaped.

*C. algerianus* S. (Fig. 236, c, d). Sporadins 3–4 mm. long; spores 10–10.5μ long; in gut of *Pimelia* sp. (Labbé, 1899).

Genus *Lophocephalus* Labbé. Epimerite sessile crateriform disc with crenulate periphery, surrounded by digitiform processes.

*L. insignis* (Schneider) (Fig. 236, e). Sporadins 1 mm. long; cysts rounded; 430μ by 330μ; pseudocysts; spores 10μ long; in gut of *Helops striatus*.
Family 11 Acanthosporidae Léger

Sporadins solitary; epimerite complex; cysts without sporoducts; spores with equatorial and polar spines.

Genus Acanthospora Léger. Epimerite simple conical knob; spores with spines.

*A. polymorpha* L. (Fig. 236, f). Sporadins polymorphic; up to 1 mm. long; protomerite cylindro-conical; deutomerite ovoidal; endoplasm yellowish brown; cyst 500–700μ in diameter; spore with 4 spines at each pole and 6 at equatorial plane, 8μ by 4.4μ; in gut of *Hydrous ceraboides*.

Genus Corycella Léger. Epimerite globular, with 8 hooks; spores biconical, with one row of polar spines (Léger, 1892).

*C. armata* L. (Fig. 236, g, h). Sporadins 250–300μ long; cysts spherical, 250μ in diameter; spores 13μ by 6.5μ; in gut of larva of *Gyrinus natator*.

Genus Prismatospora Ellis. Epimerite subglobular with 8 lateral hooks; spores hexagonal, prismatic with one row of spines at each pole.

*P. evansi* E. (Fig. 236, i). Sporadins broadly conical, 400μ long; cysts 370μ in diameter; without sporoducts; spores with 6 long spines at each pole, 11μ by 5.8μ; in gut of *Tramea lacerta* and *Sympetrum rubicundulum*; Michigan.

Genus Ancyrophora Léger. Epimerite globular with 5–10 digitiform processes directed posteriorly; spores biconical, with spines.

*A. gracilis* L. (Fig. 236, j, k). Sporadins 200μ–2 mm. long; cysts spherical, 200μ in diameter; spores hexagonal in optical section, with 4 polar and 6 equatorial spines, 8.5μ by 5μ; in gut of larvae and adults of *Carabus auratus*, *C. violaceus*, *C. sp.*, and of larvae of *Silpha thoracica* (Coleoptera) (Léger, 1892).

Genus Cometoides Labbé. Epimerite globular with 6–15 long filaments; spores with polar spines and 2 rows of equatorial spines.

*C. capitatus* (Léger) (Fig. 236, l, m). Sporadins up to 2 mm. long, active; epimerite with 12–15 filaments, 32–35μ long; cysts 300μ in diameter; spores 5.1μ by 2.5μ; in gut of larvae of *Hydrous* sp. (Coleoptera) (Watson, 1916).

Family 12 Actinocephalidae Léger

Sporadins solitary; epimerite variously formed; cysts without sporoducts; spores irregular, biconical or cylindro-biconical; in gut of insects.

Genus Actinocephalus Stein. Epimerite sessile or with a short
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neck, with 8–10 simple digitiform processes at its apex; spores biconical.

A. acutispora Léger (Fig. 236, n, o). Sporadins 1–1.5 mm. long; cysts ovoid, 550–600μ by 280μ; spores, acutely pointed, of 2 sizes, 4.5μ by 2.8μ and 6.4μ by 3.6μ; in gut of the coleopteran Silpha laevigata.

A. parvus Wellmer. Sporadins 180μ by 50μ; cysts rounded, 62–112μ in diameter; spores spindle-form, 6–7.5μ by 3–3.8μ; 8 diploid chromosomes; the first division in the zygote is meiotic; in the gut of larvae of dog-flea, Ctenocephalus canis. Development (Weschenfelder, 1938).

Genus Amphoroides Labbé. Epimerite a globular sessile papilla; protomerite cup-shaped; spores curved; in myriapods.

A. calverti (Crawley) (Fig. 236, p). Sporadins up to 1670μ by 120μ; cysts spherical, 380μ in diameter; spores unknown; in gut of Callipus lactarius.

Genus Asterophora Léger. Epimerite a thick horizontal disc with a milled border and a stout style projecting from center; spore cylindrobiconical; in Neuroptera and Coleoptera.

A. philica (Leidy) (Fig. 236, q). Sporadins 300μ–2 mm. long; cysts and spores unknown; in gut of Nyctobates pennsylvanica (Crawley, 1903).

Genus Steinina Léger and Duboscq. Solitary; epimerite a short motile digitiform process, changing into a flattened structure; spore biconical; in Coleoptera (Léger and Duboscq, 1904).

S. rotunda Watson (Fig. 236, r). Sporadins 180–250μ long; in gut of Amara augustata (Coleoptera) (Watson, 1915).

Genus Pileocephalus Schneider. Epimerite lance-shaped, with a short neck.

P. striatus Léger and Duboscq (Fig. 236, s). Sporadins 150μ long; nucleus in protomerite; cysts spherical; in gut of larvae of Ptychoptera contaminata.

Genus Stylocystis Léger. Epimerite a sharply pointed, curved process; spores biconical (Léger, 1899).

S. praecox L. (Fig. 236, t). Sporadins up to 500μ long; cysts ovoidal, 200μ long; spores 8μ by 5μ in gut of larval Tanypus sp.

Genus Discorhynchus Labbé. Epimerite a large spheroidal papilla with collar and short neck; spores biconical, slightly curved.

D. truncatus (Léger) (Fig. 237, a, b). Sporadins 300μ long; cysts spherical, 140μ in diameter; in gut of larvae of Sericostoma sp.

Genus Anthorhynchus Labbé. Epimerite a large flattened fluted disc; spores biconical, chained laterally.
A. sophiae (Schneider) (Fig. 237, c, d). Cephalins up to 2 mm. long, with 200\(\mu\) long epimerite; protomerite 150\(\mu\) long; endoplasm opaque; spores 7\(\mu\) by 5\(\mu\); in gut of Phalangium opilio.

Genus Sciadiophora Labbé. Epimerite a large sessile disc with crenulate border; protomerite with numerous vertical laminations; spores biconical.

**Fig. 237.** a, b, Discorhynchus truncatus (a, ×130) (Léger); c, d, Anthorhynchus sophiae (c, ×15; d, ×1330) (Schneider); e–g, Sciadiophora phalangii (g, spore, ×1040) (Léger); h, Amphorocephalus amphorellus (Ellis); i, Pyzinia bulbifera (Watson); j, Schneideria mucronata, ×75 (Léger); k, Beloides firmus (Léger); l, Taeniocystis mira, ×85 (Léger); m, n, Stictospora provincialis (Léger); o, Bothriopsis histrio (Léger); p–r, Coleorhynchus heros (p, ×14) (Schneider); s, Legeria agilis (Schneider); t–v, Phialoides ornata (t, ×45; v, ×930) (Léger); w, Geneiorhynchus aeschnae, ×60 (Crawley).

S. phalangii (Léger) (Fig. 237, e–g). Sporadins 2–2.5 mm. long; protomerite with 15–16 plates; cysts 500\(\mu\) in diameter; spores 9\(\mu\) by 5\(\mu\); in gut of Phalangium crassum and P. cornutum (Arachnida).

Genus Amphorocephalus Ellis. Epimerite a sessile peripherally fluted disc set upon a short neck; protomerite constricted superficially; spores unknown (Ellis, 1913).
A. amphorellus E. (Fig. 237, h). Sporadins 500–970μ long; in gut of Scolependra heros.

Genus *Pyxinia* Hammerschmidt. Epimerite a crenulate crateriform disc; with a style in center; spores biconical. Species (Vincent, 1922).

*P. bulbifera* Watson (Fig. 237, i). Sporadins up to 850μ by 260μ; in gut of *Dermestes lardarius* (Watson, 1916a).

Genus *Schneideria* Léger. Epimerite sessile, a thick horizontal disc with milled border; a style arising from center; sporadins without protomerite; spores biconical (Léger, 1892).

*S. mucronata* L. (Fig. 237, j). Sporadins 700–800μ long; agile; polymorphic; cysts 270μ by 190μ; spores fusiform, 15μ by 9μ; in intestinal caeca of larvae of *Bibio marci*.

Genus *Beloides* Labbé. Epimerite bordered by pointed lateral processes and apical style; spores biconical (Labbé, 1899).

*B. firmus* (Léger) (Fig. 237, k). Style 80μ long; cysts 180–200μ in diameter; spores 14.5μ by 6μ; in gut of larvae of *Dermestes lardarius*.

Genus *Taeniocystis* Léger. Epimerite sessile or with a short neck; 8–10 digitiform processes at its apex; deutomerite divided by septa into many chambers; spores biconical.

*T. mira* L. (Fig. 237, l). Sporadins tapeworm-like; 400–500μ long; epimerite with 6–8 curved hooks; cysts spherical, 130μ in diameter; spores 7μ by 3μ; in gut of larval *Ceratopogon solstitialis*.

Genus *Stictospora* Léger. Epimerite with a short neck, a spherical crateriform ball with 12 posteriorly-directed laminations set close to neck; cysts with a gelatinous envelope; without ducts; spores biconical, slightly curved (Léger, 1893).

*S. provincialis* L. (Fig. 237, m, n). Sporadins 1–2 mm. long; cysts 800μ in diameter; in gut of larvae of *Melolontha* sp. and *Rhizotrogus* sp.

Genus *Bothriopsis* Schneider. Epimerite sessile, small, oval, with 6 or more filamentous processes directed upward; spores biconical; cysts spherical (Schneider, 1875).

*B. histrio* S. (Fig. 237, o). Epimerite with 6 filaments, 80–90μ long; cysts 400–500μ long; spores 7.2μ by 5μ; in gut of *Hydaticus* sp.

Genus *Coleorhynchus* Labbé. Epimerite discoid, lower border over deutomerite; spores biconical.

*C. heros* (Schneider) (Figs. 234, k; 237, p–r). Sporadins 2–3 mm. long; in gut of *Nepa cinerea*. Development (Poisson, 1939).

Genus *Legeria* Labbé. Protomerite wider than deutomerite; epi-
merite unknown; cysts without duct; spores cylindro-biconical (Labbé, 1899).

*L. agilis* (Schneider) (Fig. 237, s). In gut of the larvae of *Colymbetidae* sp.

Genus *Phialoides* Labbé. Epimerite a cushion set peripherally with stout teeth, surrounded by a wider collar; with a long neck; cysts spherical, without ducts; spores biconical.

*P. ornatia* (Léger) (Fig. 237, i–v). Sporadins 500μ long; cysts 300–400μ in diameter; spores 10.5μ by 6.7μ; in gut of larvae of *Hydrophilus piceus*.

Genus *Geneiorhynchus* Schneider. Epimerite a tuft of short bristles at end of neck; spores cylindrical.

*G. aeschnae* Crawley (Fig. 237, w). Sporadins 420μ long; cysts and spores unknown; in *Aeschna constricta*.

Family 13 Porosporidae Labbé

When naked or well-protected sporozoites enter the stomach and midgut of a specific crustacean host, they develop into typical cephaline gregarines; 1, 2, or more sporadins become associated and encyst. Repeated nuclear and cytoplasmic division results in formation of an enormous number of gymnospores in hindgut. Some observers consider this change as schizogony, and hence include the family in the suborder Schizogregarina. When the gymnospores are voided in the faeces of crustaceans and come in contact with molluscan host, they enter, or are taken in by phagocytosis of, the epithelial cells of the gills, mantle or digestive system. These gymnospores are found especially in abundance in the lacunae of the gills. Presently they become paired and fuse (Hatt); the zygotes develop into naked or encapsulated sporozoites within the phagocytes of the molluscan host, which when taken in by a crustacean host, develop into cephaline gregarines.

Genus *Porospora* Schneider. Sporozoites formed in molluscan phagocytes without any protective envelope (Hatt, 1931).

*P. gigantea* (van Beneden) (Fig. 238, a–f). Sporadins in *Hymenocera gammarus*, up to 10 mm. long; cysts 3–4 mm. in diameter; gymnospores spherical, 8μ in diameter (Hatt), containing some 1500 merozoites; in molluscan hosts, *Mytilus minimus* and *Trochocochlea mutabilis*, they develop into naked sporozoites (17μ long) which are usually grouped within phagocytes.

Genus *Nematopsis* Schneider. Development similar to that of *Porospora* (Hatt); but each sporozoite in a double envelope.

*N. legeri* (de Beauchamp) (*Porospora galloprovincialis* Léger and
Fig. 238. a–f, Porospora gigantea (Hatt). a, a cephalin attached to Homarus gut, $\times 1250$; b, gymnospores; c, d, developing sporozoites in mollusc; e, sporozoites enveloped by phagocyte; f, a sporozoite, $\times 2250$. g–n, Nematopsis legeri (Hatt). g, h, trophozoites in Eriphia; i, associated trophozoites attached to gut-epithelium, $\times 1250$; j, gymnospores; k, gymnospores after entering molluscan body; l, a young sporozoite, $\times 2250$; m, cyst in mollusc with six spores; n, germination of a spore in Eriphia gut, $\times 1250$.

Duboscq) (Fig. 238, g–n). Sporadins in a crustacean, Eriphia spinifrons, in linear or bifurcated syzygy 75–750\(\mu\) long; cysts about 80\(\mu\) in diameter; gymnospores 7\(\mu\) in diameter, composed of fewer, but larger merozoites; permanent spores with a distinct one-piece shell (endospore) and a less conspicuous epispore, about 14–15\(\mu\) long and circular in cross-section, develop in numerous species of molluscan

*N. ostraeorum* Prytherch. Sporadins in syzygy in the mud crabs, *Panopeus herbsti* and *Eurypanopeus depressus*, 220–342 μ; cysts 80–190 μ in diameter; gymnospores 4 μ in diameter; spores produced in the oyster, *Ostrea virginica*, 16 μ by 11–12 μ (Prytherch, 1940). Landau and Galtsoff (1951) showed that the organism is widely distributed among the oysters along the Atlantic and Gulf coasts, but found no evidence to suppose that the organism is destructive to the host mollusc.

*N. panopei* Ball. Sporadins up to 210 μ by 14 μ; protomerite about 1/15 the body length; epimerite on young individuals only; syzygy often multiple, as in other species; cysts 88 μ by 74 μ, free in the lumen or attached to the wall of the hind-gut; gymnospores about 6.5 μ in diameter; in the gut of *Panopeus herbsti* and *P. occidentalis*; in Bermuda. Molluscan host unknown (Ball, 1951).

Suborder 2 Schizogregarinina Léger

The schizogregarines are intestinal parasites of arthropods, annelids, and tunicates. When the spore gains entrance to the digestive tract of a specific host through mouth, it germinates and the sporozoites are set free (Fig. 239). These sporozoites develop into trophozoites either in the gut-lumen or within the host cells, and undergo schizogony (c), which may be binary or multiple fission or budding. The fully grown trophozoites become paired as in Eugregarinina and encyst, in which condition they undergo sexual reproduction. Each individual which is now a gametocyte produces gametes (d–e). Fusion of two gametes follows (f). The zygote develops into a spore containing 1–8 sporozoites (g, a).

One spore from 2 gametocytes.................Family 1 Ophryocystidae
Two or more spores from 2 gametocytes.......................Family 2 Schizocystidae (p. 562)

Family 1 Ophryocystidae Léger and Duboscq

Two gametocytes produce one spore; in Malpighian tubules of Coleoptera, gut of Ascidia and coelom of Oligochaeta.

Genus *Ophryocystis* Schneider. Multiplication by binary or multiple division; extracellular; trophozoites conical, attached to host cells by pseudopods; a single spore in a pair of spheroidal gameto-
Fig. 239. The life-cycle of *Schizocystis gregarinoides*, ×1000 (Léger). a, germinating spore; b, growth of schizonts; c, schizogony; d, two gametocytes and their association; e, stages in gamete formation, f, zygote formation, g, cyst containing zygotes, each of which develops into a spore shown in a.

cytes; spore with 8 sporozoites; in Malpighian tubules of Coleoptera. Several species.

*O. mesnili* Léger (Fig. 240, a–e). In *Tenebrio molitor*; schizonts 1–4 nuclei; gametocytes 11μ in diameter; pairs 16–17μ by 11μ; spores biconical, 11μ by 7μ.

Genus *Merogregarina* Porter. Schizogony intracellular; trophozoites attached to gut epithelium by a proboscidiform organel-la; 2 gametocytes giving rise to one spore containing 8 sporozoites.

*M. amaroucii* P. (Fig. 240, f, g). In gut of the ascidian, *Amaroucium* sp.; extracellular; trophozoites with epimerite, 27–31μ long; spore about 14μ long.
Fig. 240. a–e, Ophryocystis mesnili (a, trophozoite attached to Malpighian tubule; b–e, sporogony), ×1330 (Léger); f, g, Merogregarina amaroucii, ×1000 (Porter); h, i, Spirocytis nidula (h, ×770; i, ×500) (Léger and Duboscq); j, k, Caulleryella pipiens (j, gut of Culex pipiens with trophozoites, ×200; k, a spore, ×1200) (Buschkiel).

Genus Spirocytis Léger and Duboscq. Schizogony intracellular; schizonts curved, one end highly narrowed; mature schizonts snail-like, with numerous nuclei; repeated schizogony (?) gametes in chloragogen cells, somatic and visceral peritonium; association of 2 gametes produces a spore. One species.

S. nidula L. and D. (Fig. 240, h, i). In coelom and gut epithelium of Lumbricus variegatus; multinucleate schizont about 35μ long; microgametes fusiform or ovoid, 7μ by 3μ; macrogametes ovoid or spherical, 11μ in diameter; fusion of 2 gametes produces one spore which is thick-walled, 35μ long and contains one sporozoite, up to 40μ long.

Family 2 Schizocystidae Léger and Duboscq

Two or more spores are produced in a pair of gametocytes.

Genus Schizocystis Léger. Mature trophozoite multinucleate; ovoid or cylindrical with differentiated anterior end; schizogony by multiple division; trophozoites become associated, encyst, and produce numerous (up to 30) spores, each with 8 sporozoites; in Diptera, Annelida, and Sipunculoida (Léger, 1909).

S. gregarinoides L. (Fig. 239). In gut of larvae of Ceratopogon
sporozoa, Gregarinida, 563

solstitialis; mature schizonts up to 400μ by 15μ; curved or spirally coiled; gametocytes 30–50μ long; cysts ovoid, 16–32μ long; spores biconical, 8μ by 4μ.

Genus Syncystis Schneider. Schizogony and sporogony extracellular; young trophozoites elongate, amoeboid; mature schizonts more or less spheroidal, producing some 150 merozoites; cysts spherical, producing about 150 spores. One species.

S. mirabilis S. (Fig. 241, k, l). In coelomic fluid and fat bodies of Nepa cinerea; merozoites, 7μ long; cysts spherical; spores navicular, 3–4 spines at ends, 10μ by 6μ, with 8 sporozoites.


M. dispora N. (Fig. 241, m). In adipose tissue cells of larvae of the flour moth, Ephestia kuhniella and Plodia interpunctella (pupa and adult also); schizonts 2.5–12μ long; cyst 8–12μ in diameter, with 2 spores, each with 8 sporozoites; spores 14μ by 7.5μ (Naville, 1938); 11μ by 6μ (maximum 13.5μ by 8μ) (Musgrave and Mackinnon). Highly pathogenic according to Musgrave and Mackinnon.

Genus Caulleryella Keilin. Multiplication extracellular; each gametocyte gives rise to 8 gametes, a pair forming 8 zygotes or spores; spore with 8 sporozoites; in gut of dipterous larvae. Several species.

C. pipientis Buschkiel (Fig. 240, j, k). Average trophozoites 50–60μ by 23–26μ; with paraglycogen grains; schizogony produces 30–38 merozoites; in gut of larvae of Culex pipiens.

Genus Lipotropha Keilin. Schizogony and sporogony intracellular; cyst contains 16 spores, each with 8 sporozoites; in fat body of Systenus larvae. One species.

L. macrospora K. (Fig. 241, n). Spores about 13.5μ by 3μ.

Genus Lipocystis Grell. Schizogony and sporogony intracellular; gamete formation on the surface of cytomeres; isogamy; cyst produces 100–200 spores, each with eight sporozoites. One species (Grell, 1938).

L. polyspora G. (Fig. 242, a). Spores elongate ellipsoid, about 10μ by 4μ; in the fat body of Panorpa communis.

Genus Selenidium Giard. Schizogony intracellular; many spores produced by a pair of extracellular gametocytes; spore with 4 or more sporozoites; in gut of annelids. Generic status (Mackinnon and Ray, 1933).

S. potamillae Mackinnon and Ray (Fig. 241, a–c). Trophozoites euglenoid, average size 40μ by 15μ; longitudinal striae; cysts ob-
long, producing many spores; spore, spherical with 4 (up to 10) sporozoites; in gut of the polychaete, *Potamilla reniformis* (Mackinnon and Ray, 1933).

Genus **Meroselenidium** Mackinnon and Ray. Schizogony intracellular, initiated by formation of small masses which give rise

to merozoites; about 20 spores from a pair of gametocytes; spores with numerous sporozoites. One species (Mackinnon and Ray, 1933).

*M. keilini* M. and R. (Fig. 241, d–f). Large schizonts about 150μ by 30μ; sporadins free in gut 200–300μ by 40–70μ; paired gametocytes 85μ by 40μ; spores 26–28μ by 14–16μ, bivalve (?), transverse ridges, with many sporozoites; in gut of *Potomilla reniformis*.

Genus **Selenocystis** Dibb. Sporadins leaf-like with a median ridge; biassociation with posterior ends, forming an elongated cyst, at-

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**Fig. 241.** a–c, *Selenidium potamillae* (a, ×420; b, cyst with spores, ×330; c, spore) (Mackinnon and Ray); d–f, *Meroselenidium keilini* (d, sporadin, ×670; e, f, different views of spore, ×930) (Mackinnon and Ray); g–j, *Machadoella triatoma* (g, a schizont, ×1420; h, i, a single and associated sporadins, ×710; j, spore, ×1920) (Reichenow); k, l, *Syncystis mirabilis*: k, a cyst, ×470 (Steopoe); l, spore (Schneider); m, *Mattesia dispora*, ×1480 (Naville); n, *Lipotropha macrospora*, ×800 (Keilin).
tached to the host epithelium by a foot-like organelle; isogametes with a short flagellum; spores with four or eight sporozoites. One species (Dibb, 1938).

*S. foliata* (Ray) (Fig. 242, b–f). Trophozoites 30–250μ long; pellicle with 16–24 striae; the broader end with which the organism is attached to the host epithelium depressed; surrounding this depression, a number of about 8μ long refringent filaments occur; while one

![Fig. 242. a, two views of a spore of Lipocystis polyspora, ×1485 (Grell); b–f, Selenocystis foliata (b, a mature trophozoite, ×665 (Ray); c, migration of nuclei of gametocytes to the surface of cyst, ×565; d, gamete in life; e, f, spores with four and eight sporozoites, ×1130 (Dibb)).](image)

organism is still attached, biassociation by posterior ends takes place; 26–226μ by 9–34μ; isogametes; subspherical spores about 8.5μ in diameter, with four or eight sporozoites; in the gut of the polychaete, *Scolelepis fuliginosa*.

Genus *Machadoella* Reichenow. Nematode-like, rigid; simple rounded anterior end; thick pellicle, longitudinally striated; schizogony in vermiform stage; head to head association of gametocytes; cysts with 3–6 spores, each with 8 sporozoites.

*M. triatomae* R. (Fig. 241, g–j). Schizonts about 55μ long; gametocytes 100–120μ long; schizogony into 6–8 merozoites; cysts with 3–6 spores; spore 10–11μ by 7–7.5μ; in Malpighian tubules of *Triatoma dimidiata* (of Guatemala) (Reichenow, 1935).

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Chapter 25

Order 2 Coccidia Leuckart

The Coccidia show a wide zoological distribution, attacking all vertebrates and higher invertebrates alike. The majority are parasites of the epithelium of the digestive tract and its associated glands. Asexual reproduction is by schizogony and sexual reproduction by anisogamy in the majority of species. Both kinds of reproduction take place in one and the same host body, with the exception of such forms as Aggregata in which alternation of generations and of hosts occurs. Taxonomy (Léger, 1911).

Gametocytes similar; independent; a microgametocyte developing into many microgametes.......................Suborder 1 Eimeridea
Gametocytes dissimilar; association begins during the late trophic life; a few microgametes..................Suborder 2 Adeleidea (p. 590)

Suborder 1 Eimeridia Léger

These coccidians are, as a rule, intracellular parasites of the gut epithelium. Both asexual (schizogonic) and sexual (sporogonic) generations occur in one host, although in some there is also alternation of hosts. The life-cycle of Eimeria schubergi, a gut parasite of the centipede, Lithobius forficatus, as observed by Schaudinn, is as follows (Fig. 243). The infection begins when the mature oocysts of the coccidian gain entrance into the host through the mouth. The sporozoites escape from the spores and make their way through the micropyle of the oocyst into the gut lumen (p). By active movement they reach and enter the epithelial cells (a). These schizonts grow into large rounded bodies and their nuclei multiply in number. The newly formed nuclei move to the body surface, and each becomes surrounded by a small mass of cytoplasm, forming a merozoite. When the host cells rupture, the merozoites are set free in the gut lumen, make their way into new host cells and repeat the development (b). Instead of growing into schizonts, some merozoites transform themselves into macro- or micro-gametocytes (c). Each macrogametocyte contains refractile bodies, and becomes a mature macrogamete, after extruding a part of its nuclear material (d, e). In the microgametocyte, the nucleus divides several times and each division-product assumes a compact appearance (f–h). The biflagellate comma-shaped microgametes thus produced, show activity when freed from the host cells (i). A microgamete and a macrogamete unite to form a zygote which secretes a membrane around itself (j). This stage is
Fig. 243. The life-cycle of *Eimeria schubergi*, ×400 (Schaudinn) a, entrance of a sporozoite in the gut epithelial cell of host and growth of schizont; b, schizogony; c, macro- and micro-gametocyte; d, e, formation of macrogamete; f–h, formation of microgametes; i, mature gametes prior to fusion, j, k, fertilization; l–n, spore-formation; o, oocyst containing four mature spores, each with two sporozoites; p, germination of spores in host's gut.

known as the oocyst. The nucleus divides twice and produces four nuclei (*k–m*). Each of the four nuclei becomes the center of a sporoblast which secretes a membrane and transforms itself into a spore (*n*). Its nucleus, in the meantime, undergoes a division, and two sporozoites develop in the spore (*o*). Oocysts leave the host in the faecal matter and become the source of infection.

Body veriform; schizogony in motile stage.......................... Family 1 Selenococcidiidae (p. 572)

Body not veriform
 Alteration of generations and of hosts.. Family 2 Aggregatidae (p. 572)
 Only one host
 Gametocytes become associated early; many microgametes..........
 .......................................................... Family 3 Dobelliidae (p. 576)
 Gametocytes independent.............. Family 4 Eimeriidae (p. 576)
Family 1 Selenococcidiidae Poche

Vermiform body and gametic differentiation place this family on the borderline between Coccidia and Gregarinida.

Genus Selenococcidium Léger and Duboscq. Nucleus of vermiciform trophozoite divides 3 times, producing 8 nuclei; trophozoite becomes rounded after entering gut-epithelium and divides into 8 schizonts; this is apparently repeated; schizonts develop into gametocytes; microgametocyte produces numerous microgametes; gametic union and sporogony (?). One species.

Fig. 244. Selenococcidium intermedium, ×550 (Léger and Duboscq). a, schizont in host gut; b, c, schizogony; d, microgametocyte; e, microgametes; f, macrogametocyte; g, macrogamete; h, zygote (oocyst).

S. intermedium L. and D. (Fig. 244). Octonucleate vermiciform schizont 60–100μ long, and divides into vermicular merozoites in gut cells; parasitic in gut lumen of European lobster.

Genus Ovivora Mackinnon and Ray. Trophozoites large and vermiciform (Fig. 245, a); gametocytes spherical (c); large macrogametocytes; small microgametocytes, giving rise to numerous biflagellate microgametes (d); oocyst membrane delicate or lacking; ovoid spores contain variable (averaging 12?) number of sporozoites; schizogony produces many merozoites; one host. One species (Mackinnon and Ray, 1937).

O. thalassaeae (Lankester) (Fig. 245). In the egg of the echiurid worm, Thalassema neptuni; merozoites about 10μ long (b); macrogametocytes (c) 40–75μ in diameter; microgametocytes (c) 23–65μ; chromosome reduction, 14 to 7, in the zygote; spores (f) 15.5μ by 13.5μ (Mackinnon and Ray, 1937).

Family 2 Aggregatidae Labbé

Anisogamy results in production of zygotes which become transformed into many spores, each with 2–30 sporozoites; in schizogony
cytomeres first appear and then merozoites; alternation of generations and of hosts which are marine annelids, molluscs and crustaceans.

Genus *Aggregata* Frenzel. Schizogony in a crustacean and sporogony in a cephalopod; zygote produces many spores, each with 3 sporozoites. Many species. Cytology (Moroff, 1908).

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**Fig. 245. Ovivora thalassemæ** (Mackinnon and Ray). a, two mature organisms in host egg, seen in reflected light, ×250; b, schizonts in sectioned egg; c, micro- and macro-gametocytes in an egg, ×500; d, two maturing microgametocytes still attached to cytoplasmic residuum, ×1075; e, cyst with zygotes in some of which nuclei are dividing, ×500; f, a spore with 10 nuclei, ×900.

*A. eberthi* (Labbé) (Fig. 246). Schizogony in *Portunus depurator* and sporogony in *Sepia officinalis*. Spores (a) germinate in the crab gut, each liberating 3 sporozoites (b) which grow and produce merozoites (10μ by 2μ) by schizogony in peri-intestinal connective tissue cells (6 chromosomes) (c–f); when host crab is eaten by a cuttlefish, merozoites penetrate gut wall and develop into micro- and macro-gametocytes (h, k), and further into gametes (j–l); anisogamy (m) produces zygotes; zygote nucleus contains 12 chromosomes which become divided into 2 groups of 6 in the first division (n, o); repeated nuclear division (p) forms many sporoblasts (q), each transforms itself into a spherical spore with 3 sporozoites (Dobell, 1925; Naville, 1925; Bělař, 1926).
Fig. 246. The life-cycle of *Aggregata eberthi* (Dobell). a, a mature spore; b, germination of spore; c–f, schizogony; g, a merozoite, swallowed by Sepia; h–j, development of microgametes; k–l, development of macrogamete; m, fertilization; n, o, first zygotic division, chromosomes reduced in number from 12 to 6; p, q, development of sporoblasts, each of which develops into a spore with three sporozoites.

Genus *Merocystis* Dakin. Sporogony in the kidney of the whelk, Buccinum; schizogony unknown, in another host (possibly a crab); microgametocytes produce first cytomeres which in turn form microgametes; anisogamy gives rise to zygotes, zygote forms many sporoblasts, each developing into a spore; spore spherical, with 2 sporozoites. One species.

*M. kathae* D. (Fig. 247, a, b). In the kidney of *Buccinum undatum*; spores spherical, about 14µ in diameter. Patten (1935)
studied its life cycle and found that during microgametogenesis and sporogony, 6 chromosomes occur. She added that meiosis occurs in the zygote which is the only diploid stage as in *Aggregata eberthi*.

Genus *Pseudoklossia* Léger and Duboseq. Anisogamy and sporogony in the kidney of marine mussels; oocyst or zygote produces numerous spores; spore with 2 sporozoites; no residual body; schizogony unknown, in another host (Léger and Duboseq, 1915, 1917).

*P. pectinis* L. and D. (Fig. 247, c). In kidney of *Pecten maximus* in France; association of 2 sporozoites which are 3.5μ in diameter.

Genus *Caryotropha* Siedlecki. Both schizogony and sporogony take place in a host. One species.

*C. mesnili* S. In coelom (in floating bundles of spermatogonia) of the polychaete, *Polymnia nebula;* schizogony in bundle of spermatogonia, in which cytomeres with 10–16 nuclei and then merozoites are formed; schizogony repeated; gametocytes undergo development also in the same host cells; microgametes become set free in coelom, where union with macrogametes takes place; each oocyst forms about 16 spores; spore with usually 12 sporozoites; cysts are extruded with the reproductive cells of the host worm.
Genus **Myriospora** Lermantoff. Anisogamy and sporogony in marine snails; schizogony unknown; oocyst forms numerous spores each with 2 sporozoites. One species.

*M. trophonææ* L. In the polychaete, *Trophonia plumosa*; macrogametes, vermiform, up to 800μ long, later ovoid; microgametocyte forms first about 100 cytomeres, each with some 20 nuclei; microgametes comma-shaped; anisogamy; oocyst with several hundred spores, each with about 24 sporozoites. One species.

Genus **Hyaloklossia** Labbé. Schizogony unknown; sporogony in the kidney of marine mussels; oocyst in the organ-cavity; spherical spores of 2 kinds: smaller one with 2 spirally coiled sporozoites and the other with 4–6 sporozoites. One species.

*H. pelseneeri* Léger. Spherical oocysts 75–80μ in diameter; spores 8μ and 11–12μ in diameter; in kidney of *Tellina* sp. and *Donax* sp.

Genus **Angeiocystis** Brasil. Schizogony unknown; sporogony in polychaetes; oocyst forms 4 spores; spore oval, with about 30 sporozoites and residual body at a pole. One species.

*A. audouiniaæ* B. In the cardiac body of *Audouinia tentaculata*; macrogametes vermiform, up to 65μ long.

**Family 3 Dobelliidae** Ikeda

Numerous microgametes develop from each microgametocyte; the union of gametocytes begins early.

Genus **Dobellia** Ikeda. Schizonts sexually differentiated: microschizonts and macroschizonts; young schizonts binucleate; association of 2 gametocytes begins early as in *Adeleidea* (p. 590), but many microgametes are formed in each microgametocyte. One species (Ikeda, 1914).

*D. binucleata* I. In the gut of *Petalostoma minutum*; mature oocyst 20–25μ in diameter, with a thin wall, contains some 100 sporozoites without any spore membrane around them.

**Family 4 Eimeriidae** Léger

Macro- and micro-gametocytes develop independently; microgametocyte produces many gametes; an oocyst from a pair of anisogametes; oocyst with variable number of spores containing 1–many sporozoites, which condition is used as basis of generic differentiation. Oocysts found in the faeces of hosts are usually immature; time needed for completion of spore formation depends upon the species, temperature, moisture, etc. Becker (1934) recommends the following bactericidal solutions in which oocysts develop to maturity: 1% formaldehyde, 1% chromic acid of 2–4% potassium dichromate.
Genus *Eimeria* Schneider (*Coccidium* Leuckart). Zygote or oocyst produces four spores, each with two sporozoites. Numerous species (Levine and Becker, 1933; Boughton and Volk, 1938; Hardcastle, 1943); host specificity (Becker, 1933).

*E. schubergi* (Schaudinn) (Fig. 243). In the gut of *Lithobius forficatus*; oocysts spherical, 22–25 μ in diameter.

*E. stiedae* (Lindemann) (*Coccidium oviforme* Leuckart) (Fig. 247, d–k). In the epithelium of the bile-duct and liver (with white nodules) of wild and domestic rabbits; schizonts ovoid or spherical, 15–18 μ in diameter; merozoites 8–10 μ long; oocysts ovoid to ellipsoid, often yellowish, micropylar end flattened; mature oocysts 28–40 μ by 16–25 μ; sporulation in 60–70 hours; heavy infection is believed to be fatal to young animals, which may occur in an epidemic form. Transmission and comparison with *E. perforans* (Uhlhorn, 1926).

*E. perforans* (Leuckart) (Fig. 247, l, m). In the small intestine of rabbits; oocysts with equally rounded ends, 24–30 μ by 14–20 μ; sporulation in 48 hours at 33°C; the thermal death point of immature oocysts 51°C. (Becker and Crouch, 1931); pathogenic. Other species (Péard, 1925; Becker, 1934). Lund (1950) found 17 per cent of coccidian infection among 1200 faecal specimens collected from 23 commercial rabbitries in southern California.

*E. zürnii* (Rivolta). In the gut of cattle; oocystss pherical to ellipsoidal, 12–28 μ by 10–20 μ; sporulation in 48–72 hours; said to cause diarrhoea.

*E. bovis* (Züblin) (*E. smithi* Yakimoff and Galouzo). In the gut of cattle; oocysts 23–34 μ by 17–23 μ; sporulation in three to five days in shallow dishes, and two weeks in deep dishes (Becker). Development (Hammond et al., 1946).

*E. ellipsoidalis* Becker and Frye (Fig. 248, a). In the faeces of calf; oocysts ellipsoidal, 20–26 μ by 13–17 μ; sporulation in 18 days (Becker and Frye, 1929).

*E. cylindrica* Wilson. In the faeces of cattle; oocysts cylindrical, 19–27 μ by 12–15 μ; sporulation in two to 10 days.

*E. wyomingensis* Huizinga and Winger. In the faeces of cattle; oocysts pyriform, 37–45 μ by 26–31 μ; spores 19 μ by 3 μ (Huizinga and Winger, 1942).

*E. faurei* Moussu and Marotel (Fig. 247, r). In the gut of sheep and goat; oocysts ovoid, 20–40 μ by 17–26 μ; sporulation in 24–48 hours.

*E. arloingi* Marotel. In the gut of sheep and goat; oocysts with a cap, ovoid, 25–35 μ by 18–25 μ; sporulation in three days.

*E. intricata* Spiegel. In the gut of sheep and goat; oocysts with
Fig. 248. Oocysts of Eimeria. a, *Eimeria ellipsoidalis*, ×1500 (Becker and Frye); b, *E. debliecki*, ×1070 (Wenyon); c, *E. canis*, ×650 (Wenyon); d, *E. falciformis*, ×730 (Wenyon); e, *E. separata*; f, *E. miyairii*, ×2000 (Becker, Hall and Hager); g, *E. mephitidis*, ×1000 (Andrews); h, *E. cynomysis*, ×1000 (Andrews); i, *E. citelli*, ×1360 (Kartchner and Becker), j, *E. monacis*, ×1630 (Fish); k, *E. tenella*, ×600 (Tyzzer); l, *E. mitis*, ×430 (Tyzzer); m, *E. acervulina*, ×430 (Tyzzer); n, *E. maxima*, ×470 (Tyzzer); o, *E. ranarum*, ×670; p, *E. prevoti*, ×670 (Laveran and Mesnil); q, *E. ranae*, ×670 (Dobell); r, *E. sardinae*, ×600, s, *E. clupearum*, ×600 (Thomson and Robertson); t, *E. brevortiana* (Hardcastle).
thick wall, with or without cap, ellipsoidal, 42–60 μ by 30–36 μ; sporulation in about 9 days. Species in North American sheep (Christensen, 1938), in Rocky Mountain Bighorn sheep (Honess, 1942).

*E. debliecki* Douwes (Fig. 248, b). In the gut of pigs; 30–82 per cent infection in California (Henry); oocysts 12–29 μ by 12–20 μ; sporulation in seven to nine days. Development (Nöller and Frenz, 1922).

*E. scabra* Henry. In the caecal contents of pigs; oocysts, brown, ellipsoidal, 22–36 μ by 16–26 μ. Henry (1931) recognized 2 other species in California swine.

*E. caviae* Sheather. In the gut of guinea pigs; oocysts subspherical to ellipsoid, 13–26 μ by 13–22 μ (Sheather, 1924). Morphology and development (Lapage, 1940).

*E. canis* Wenyon (Fig. 248, c). In the gut of dogs; oocysts, ellipsoidal, 18–45 μ by 11–28 μ; spores 9.5 μ by 2.5 μ; sporulation in 24 hours.

*E. felina* Nieschulz. In the gut of cat; oocysts 21–26 μ by 13–17 μ.

*E. falciformis* (Eimer) (Fig. 248, d). In the gut of mice; oocysts spherical to ovoid, 16–21 μ by 11–17 μ; sporulation in 3 days.

*E. nieschulzi* Dieben. In the small intestine of rats; oocysts 16–26.4 μ by 13–21 μ; sporulation in 65–72 hours. Growth-promoting potency of feeding stuffs (Becker, 1941; Becker, Manresa and Smith, 1943).

*E. separata* Becker and Hall (Fig. 248, e). In the caecum and colon of rats; oocysts 13–19.5 μ by 11–17 μ; sporulation in 27–36 hours.

*E. miyairii* Ohira (Fig. 248, f). In the small intestine of rats; oocysts 16.5–29 μ by 16–26 μ; sporulation in 96–120 hours. Unsporulated oocysts perish in 15 seconds at 53°C. and in 24 hours at 41°C.; sporulated oocysts are killed in two minutes at 52°C. (Reinhardt and Becker, 1933). Structure of oocyst wall (Henry, 1932); Eimeria in rodents (Fish, 1930; Henry, 1932a; Roudabush, 1937a).

*E. mephitidis* Andrews (Fig. 248, g). In the faeces of the common skunk; oocysts oval to spherical, 17–25 μ by 16–22 μ; wall 1 μ thick; a circular micropyle; spores with a rostrum, 10–12 μ by 7–9 μ; extended sporozoites 10–14 μ by 4–5 μ; other stages unknown (Andrews, 1928).

*E. cynomysis* A. (Fig. 248, h). In the faeces of the prairie dog; oocysts oval, 33–37 μ by 28–32 μ; a double fibrous wall, 1.5–2.5 μ thick; the inner wall slightly orange-yellow; micropyle 5–6 μ in diameter; spores, broad pyriform, 13–17 μ by 8–12 μ.

*E. citelli* Kartchner and Becker (Fig. 248, i). In the caecal contents of the striped ground squirrel, *Citellus tridecemlineatus*; subspherical to ellipsoidal oocysts 15–23 μ by 14–19 μ.
Fig. 249. Diagram illustrating the development of *Eimeria tenella* in the caecal glands of chick (Tyzzer). The numbers below indicate the days of infection. *ma*, macrogamete; *me*, merozoite (*me*¹, *me*², *me*³, generation 1, 2, 3 merozoites respectively); *mi*, microgametocyte; *oo*, oocyst; *ret. oo* and *ret. sch*, oocysts and schizonts which failed to escape; *sch*¹, *sch*², schizonts of generation 1 and 2; *tr*, young growing trophozoites. (Continue to upper left of Fig. 250.)

*E. monacis* Fish (Fig. 248, j). In the intestine of the woodchuck, *Marmota monax*; spherical to subspherical oocysts 20µ by 18µ (Fish, 1930), 14–20µ in diameter (Crouch and Becker, 1931); wall comparatively thick; sporulation completed in 60–64 hours in 2 per cent potassium bichromate at room temperature. Crouch and Becker found two other species: *E. perforoides* and *E. os*, in the woodchuck in Iowa. *Eimeria* in lemming (Levine, 1952).

*E. tenella* (Railliet and Lucet) (Figs. 248, k; 249; 250). In the caeca, colon and lower small intestine of chicken; a cause of acute coccidiosis characterized by haemorrhage (Tyzzer); in the caecal contents of California quail (Henry); oocysts 19.5–26µ by 16.5–23µ; sporulation in 48 hours. Tyzzer’s observation on experimental infection in
Fig. 250. Continuation of the diagram shown in Fig. 249 (Tyzzer). From the right end of the upper figure continue to the left of the lower figure; for explanation see Figure 249.

chicken is as follows (Figs. 249 and 250): When a large number of
Oocysts are fed to chickens, the sporozoites emerge from the oocysts and spores, in as early as 20 hours and are found on the surface of the caecal mucosa. Toward the end of the second day, growing trophozoites are found in the gland epithelial cells; they undergo schizogony (Fig. 249, sch1) by the middle of the third day. A single first generation schizont is estimated to produce about 900 pyriform merozoites which measure 2–4μ by 1–1.5μ and occur in the gland lumen (me1). As these merozoites invade the epithelial cells of the fundi of the glands and become trophozoites, the infected host cells increase in size, become rounded and no longer form a continuous layer (tr). These trophozoites (Fig. 250, sch) grow to much greater dimensions (up to as much as 45μ in diameter) than those of the first generation and multiply into merozoites (me2) by the fifth day. These merozoites are much larger and more elongated than those of the first generation and measure 16μ by 2μ. The haemorrhage in the affected mucosa which begins usually with the growth of the second generation trophozoites, increases in volume so that by the fifth day after infection, a great portion of the mucosa sloughs off, which coincides with the liberation of the merozoites. The merozoites formed in the host cells located in the deeper part of the mucosa are unable to become free and appear to grow into multinucleate forms (ret sch). When the liberated merozoites enter epithelial cells, most of them develop into macrogametocytes (ma) and microgametocytes (mi), while comparatively small numbers become trophozoites and form by budding a few, large third generation merozoites (me3). Mature oocysts (oo) are found on seven to eight days after infection. Eimeria species in chicken (Tyzzer, 1929, 1932; Henry, 1931a); economic importance (Foster, 1949; Brackett and Bliznick, 1950); pathological changes (Tyzzer, 1929, 1932; Mayhew, 1937); statistical study of infections (Fish, 1931); mortality of hosts (Mayhew, 1933); killing oocysts (Fish, 1931a); control measures (Andrews and Tsuchiya, 1931; Andrews, 1933); comparative oocyst production (Brackett and Bliznick, 1950); in wild fowls (Haase, 1939).

E. mitis Tyzzer (Fig. 248, l). In the anterior small intestine of chicken; oocysts subspherical; 16.2μ by 15.5μ; sporulation in 48 hours (Tyzzer, 1929).

E. acervulina T. (Fig. 248, m). In the anterior small intestine of chicken, and in California quail (Henry); oocysts oval, 17.7–20.2μ by 13.7–16.3μ; sporulation in 20 hours; associated with serious chronic coccidiosis (Tyzzer, 1929). Effect on host (Moynihan, 1950).

E. maxima T. (Fig. 248, n). In the small intestine of chicken; oocysts oval, 21.5–42.5μ by 16.5–29.8μ (Tyzzer, 1929).
E. necatrix Johnson. In the small intestine (schizonts) and caeca (oocysts) of chicken; a cause of chronic coccidiosis; oocysts obovate, 13-23μ by 11-18μ; sporulation in 48 hours (Tyzzer, 1932).

E. praecox J. In the upper third of the small intestine of chicken; oocysts ovoid, 20-25μ by 15.5-20μ; sporulation in 48 hours.

E. meleagrimitis T. In the lower small intestine of turkey; somewhat similar to E. mitis; oocysts, 16.5-20.5μ by 13.2-17.2μ (Tyzzer, 1929).

E. adenocides Moore and Brown. In the ileum, caeca and rectum of turkeys; oocysts about 25.6μ by 16.5μ; highly pathogenic to young turkeys (Moore and Brown, 1950).

E. truncata (Railliet and Lucet). In the kidney of geese; oocysts truncate at one pole, ovoid, 14-23μ by 13-18μ; some observers find this coccidian fatal to young geese.

E. anseris Kotlan. In the intestine of geese; oocysts spherical or pyriform, 11-16μ in diameter. Coccidia in Canada goose (Levine, 1952a).

E. labbeana Pinto. In the gut of domestic pigeon; oocysts sometimes light brown, 15-26μ by 14-24μ.

E. dispersa Tyzzer. In the small intestine of bob-white quail and pheasant; oocysts ovate, 18.8-22.8μ (quail), smaller in pheasant, without polar inclusion; sporulation in about 24 hours.

E. amydae Roudabush. In the intestine of Amyda spinifera; oocysts oval with a thin wall, 17-24μ by 12-17μ; elliptical spores about 11-16μ long (Roudabush, 1937).

E. chrysemydis Deeds and Jahn. In the intestine of Chrysemys marginata; oval oocysts 21-27μ by 13-18μ; fusiform spores 12-14μ by 5-8μ (Deeds and Jahn, 1939). Other reptilian species (Roudabush, 1937)

E. ranarum (Labbé) (Fig. 248, o). In the gut epithelium (nuclei) of frogs; oocysts about 17μ by 12μ.

E. prevoti (Laveran and Mesnil) (Fig. 248, p). In the gut epithelium of frogs; oocysts about 17μ by 12μ.

E. ranae Dobell (Fig. 248, q). In the gut of frogs; oocysts 22μ by 18μ.

Species of Eimeria are often parasitic in fishes used for human consumption, and thus may appear in faecal matter. A few examples will be mentioned here.

E. sardinae (Thélohan) (E. oxyspora Dobell) (Fig. 248, r). In the
testis of sardine; spherical oocyst 30–50µ (Thélohan, 1890; Dobell, 1919).

E. clupearum (Thélohan) (E. wenyonii Dobell) (Fig. 248, s). In the liver of herring, mackerel, and sprat; spherical oocysts 18–33µ in diameter (Thélohan, 1894; Dobell, 1919). Taxonomy (Thomson and Robertson, 1926).

E. gadi Fiebiger. In the swim-bladder of Gadus virens, G. morrhua, and G. aeglefinus; schizogony and sporogony; germination of spores takes place in the bladder of the same host individual, bringing about a very heavy infection; oocysts 26–28µ; pathogenic (Fiebiger, 1913).

E. brevoortiana Hardcastle (Fig. 248, t). Schizogony in the epithelium of the pyloric caeca and sporogony in the testis of the menhaden, Brevoortiana tyrannus; mature oocysts, spherical, 17.5–30µ in diameter or ovoid, 21–30µ by 15–27.5µ (Hardcastle, 1944).

Genus Jarrina Léger and Hesse. Oocysts ovoid, one end rounded and the other drawn out into a short neck; 4 spores, each with 2 sporozoites (Léger and Hesse, 1922).

J. paludosa L. and H. (Fig. 251, a, b). In the gut of Fulica atra and Gallinula chloropus; oocysts 15µ by 11µ; sporulation in 15 days.

Genus Wenyonella Hoare. Oocysts with 4 spores, each with 4 sporozoites. Three species.

W. africana H. (Fig. 251, c, d). In the small intestine of Boaedon lineatus (“brown snake”) in Uganda; oocysts ovoid or subspherical, 18.5–19.2µ by 16–17.6µ; spores ovoid, 9.6µ by 8µ; sporulation in 5–6 days.

W. gallinae Ray. In the epithelium of the lower intestine of chick-
en; oval oocysts, 29.5–33.5 μ by 20–23 μ; spores 18.8 μ by 8 μ; sporozoites club-shaped; sporulation in four to six days at 28°C. (Ray, 1945).

Genus Isospora Schneider. Oocyst produces two spores, each containing four sporozoites. Avian Isospora (Boughton, Boughton and Volk, 1938).

I. hominis (Rivolta) (I. bellii Wenyon) (Fig. 251, e, f). This is the sole coccidian parasite of man known up to the present time. Its life cycle is unknown, but most probably the schizogony, gametogenesis and sexual fusion occur in the intestinal epithelium. Oocysts have only been seen in the stools of infected persons.

The oocyst is asymmetrically fusiform; 20–33 μ by 10–16 μ; wall is made up of two membranes which are highly resistant to chemicals; when voided in faeces, the contents either fill up the oocyst or appear as a spherical mass, composed of refractile granules of various sizes; nucleus appears as a clear circular area; when the faecal specimen is kept in a covered container at the room temperature, the protoplasmic mass divides into 2 spherical sporoblasts in about 24 hours each sporoblast develops in another 24 hours into a spore (10–16 μ by 7–10 μ) containing 4 sporozoites. Further changes take place when the oocyst finds its way into the human intestine in contaminated food or water.

I. hominis has been observed in widely separated regions, but appears not to be of common occurrence. As to its effect on the human host, very little is known. Connal (1922) described the course of an accidental oral infection by viable mature oocysts, as follows: The incubation period was about six days, the onset sudden, and the duration over a month. The cure was spontaneous. The symptoms were diarrhoea, abdominal discomfort, flatulence, lassitude, and loss of weight. During the first three weeks of the illness no oocysts were found, but then oocysts appeared in the stools for nine days. On the 10th day they were not seen, but reappeared on the 11th and 12th days, after which they were not found again. The acute signs of illness abated within one week of the finding of the oocysts. The faeces contained a large amount of undigested material, particularly fat which gave it a thick oily consistency, showing signs of slow gaseous formation.

Matsubayashi and Nozawa (1948) found six cases of infection in Japan. A volunteer ingested some 3000 oocysts. Eight days later diarrhoea developed, followed by a rise of temperature above 39°C., which lasted for 10 days. On the following day, the diarrhoea subsided, but later returned and was especially pronounced on the 17th
day, after which it disappeared completely. Oocysts were discharged regularly since the 9th day for 32 days. About a month after the cessation of oocyst-production, the person ingested again some 2500 cysts, but no infection resulted, which the two authors attributed to the immunity produced during the first infection. Another volunteer showed a similar course of infection. The symptoms disappeared without medication after the termination of oocyst discharge. Thus, the coccidiosis of man appears to be a self-limited one. Attempts to infect common laboratory animals with this coccidian have so far failed (Foner, 1939; Herrlich and Liebmann, 1944; Rita and Vida, 1949). History (Dobell, 1919); human species (Dobell, 1926); incidence (Magath, 1935; Barksdale and Routh, 1948).

*I. bigemina* (Stiles) (Fig. 251, g). In the gut of cat and dog; oocysts 10–14μ by 7–9μ.

*I. rivolta* (Grassi) (Fig. 251, h). In the gut of cat and dogs; oocysts 20–25μ by 15–20μ.

*I. felis* Wenyon (Fig. 252, a). In cat and dog; oocysts 39–48μ by 26–37μ.

*I. suis* Biester. In swine faeces; oocysts subspherical, about 22.5μ by 19.4μ; sporulation in 4 days.

*I. lacazii* Labbé. In the small intestine of passerine birds (sparrows, blackbirds, finches, etc); oocysts subspherical or ovoidal, 18.5–30μ by 18–29.2μ; spores, 16.5–18.5μ by 10.3–12.4μ; heavily infected sparrows show definite symptoms of infection; sporulation in 24 hours (Henry, 1932b). Sparrows and other common small birds have been known to be free from Eimeria infection, while the barnyard fowls are seldom infected by Isospora (Boughton, 1929). Significance of size variation in oocysts (Boughton, 1930; Henry, 1932b); development (Chakravarty and Kar, 1944).

*I. buteonis* Henry. In the duodenal contents of several species of hawks: *Buteo borealis*, *B. swainsoni*, *Accipiter cooperii*, and *Asio flammeus*; oocysts irregular in form with a thin wall, 16–19.2μ by 12.8–16μ; spores 9.6–13μ by 8–10.4μ (Henry, 1932b).

*I. lieberkühni* (Labbé) (Fig. 252, b). Oocyst about 40μ long; in the kidney of frogs. Development (Nöller, 1923).

Genus *Cyclospora* Schneider. Development similar to that of *Eimeria*; oocyst with 2 spores, each with 2 sporozoites and covered by a bi-valve shell.

*C. caryolytica* Schaudinn (Fig. 252, c). In the gut of the mole; sporozoites enter and develop in the nuclei of gut epithelial cells; oocyst oval, about 15μ by 11.5μ. Development (Tanabe, 1938).

Genus *Dorisiella* Ray. Zygote develops (without becoming oocyst)
into 2 spores, each with 8 sporozoites; macrogametocytes migratory.

*D. scolelepidis* R. (Fig. 252, d). In the gut of the polychaete, *Scolelepis fuliginosa*; zygote contents divide into 2 oval spores, 12–16μ by 6–10μ; spore with 8 sporozoites (Ray, 1930).

Genus *Caryospora* Léger. Oocyst develops into a single spore with 8 sporozoites and a residual mass; membrane thick and yellow. One species.

*C. simplex* L. (Fig. 252, e,f). In the gut-epithelium of *Vipera aspis*; oocyst thick-walled, 10–15μ in diameter.

Genus *Cryptosporidium* Tyzzer. Lumen-dwelling minute organisms; oocyst with 4 sporozoites.

*C. muris* T. (Fig. 252, g, i). In the peptic glands of the mouse; both schizogony and sporogony in the mucoid material on surface of the epithelium: oocysts 7μ by 5μ; 4 sporozoites, 12–14μ long (Tyzzer, 1910).
C. parvum T. In the glands of small intestine of the mouse; oocysts with 4 sporozoites, 4.5μ in diameter (Tyzzer, 1912).

Genus Pfeifferinella Wasielewski. Macrogamete with a “reception tubule” by which microgamete enters; oocyst produces directly 8 sporozoites.

P. ellipsoides W. (Fig. 252, j). In the liver of Planorbis corneus; oocysts oval, 13–15μ long.

P. impudica Léger and Hollande (Fig. 252, k). In the liver of Limax marginatus; oocysts ovoid, 20μ by 10μ.

Genus Lankesterella Labbé. Oocyst produces 32 or more sporozoites directly without spore-formation; in endothelial cells of cold-blooded vertebrates; mature sporozoites enter erythrocytes in which they are transmitted to a new host individual by bloodsucking invertebrates.

L. minima (Chaussat) (Fig. 252, l). In frogs; transmitted by the leech (Placobdella marginata); frog acquires infection through introduction of sporozoites by a leech; sporozoites make their way into the blood capillaries of various organs; there they enter endothelial cells; schizogony produces numerous merozoites which bring about infection of many host cells; finally macro- and micro-gametocytes are formed; anisogamy produces zygotes which transform into oocysts, in which a number of sporozoites develop; these sporozoites are set free upon disintegration of cyst wall in the blood plasma and enter erythrocytes (Nöller); oocyst oval, about 33μ by 23μ.

Genus Schellackia Reichenow (Tyzzeria Allen). Oocyst spherical with 8 sporozoites, without spore membrane; in the intestine of birds and lizards.

S. bolivari R. In the mid-gut of the lizards, Acanthodactylus vulgaris and Psammodromus hispanicus; development somewhat similar to that of Eimeria schubergi (Fig. 243); oocysts spherical, 15–19μ in diameter, with 8 sporozoites (Reichenow, 1919).

S. perniciosa (Allen). In the small intestine of Anas domesticus; oocysts 10–13.3μ by 9–10.8μ; highly pathogenic.

Genus Barrouxia Schneider. Oocyst with numerous spores, each with a single sporozoite; spore membrane uni- or bi-valve, with or without caudal prolongation. Development (Schellack and Reichenow, 1913).

B. ornata S. (Fig. 252, m). In gut of Nepa cinerea; oocysts spherical, 34–37μ in diameter, with many spores; spore with one sporozoite and bivalve shell, 17–20μ by 7–10μ.

Genus Echinospora Léger. Oocyst with 4–8 spores, each with a sporozoite; endospore with many small spinous projections.
Coccidia

E. labbei L. (Fig. 252, n). In the gut of Lithobius mutabilis; oocysts spherical, 30-40μ in diameter; spores, 11μ by 9.4μ, with bi-valve shell, sporulation completed in about 20 days.

Fig. 253. The life-cycle of Adelea ovata, X600 (Schellack and Reidleman). a, schizont entering the gut epithelium of the host centipede; b-d, macrogametocytes (left); e, association of macrogametocytes; f, fertilization; g, zygote; h, nuclear division in zygote; i, mature oocyst with many spores.
Suborder 2 Adeleidea Léger

The Adeleidea are on the whole similar to Eimeridea in their habitat and development, but the micro- and macro-gametocytes become attached to each other in pairs during the course of development into gametes (Fig. 253), and each microgametocyte produces a few microgametes. The zygote becomes oocyst which produces numerous sporoblasts, each of which develops into a spore with 2 or 4 sporozoites.

In epithelium of gut and its appended glands of chiefly invertebrates...

...Family 1 Adeleidae

In cells of circulatory system of vertebrates...

...Family 2 Haemogregarinidae (p. 592)

Family 1 Adeleidae Léger

Genus Adelea Schneider. Zygote develops into a thinly walled oocyst with numerous flattened spores, each with 2 sporozoites; in arthropods.

A. ovata S. (Fig. 253). In the gut of Lithobius forficatus; merozoites 17–22μ long; oocysts elongate oval, 40–50μ by 30–40μ; 17–33 or more spores; spores circular, flattened, 20μ by 4μ (Hesse, 1910a). Life cycle (Schellack and Reichenow, 1913, 1915).

Genus Adelina Hesse. Oocyst thick-walled; spores spherical, comparatively small in number; in the gut or coelom of arthropods and oligochaetes (Hesse, 1910, 1910a).

A. dimidiata (Schneider) (Fig. 254, a). In the gut of Scolopendra cingulata and other myriapods; oocysts with 3–17 spores (Schellack, 1913).

A. octospora H. (Fig. 254, b). Spherical oocyst contains spores; in the coelom of Slavina appendiculata (Hesse, 1910a).

A. deronis Hauschka and Pennypacker. In peritoneum of Dero limosa; oocyst contains 12 (10–14) spores; meiosis at the first zygotic nuclear division; haploid chromosome number 10; the life cycle is completed in 18 days at room temperature (Hauschka, 1943).

Genus Klossia Schneider. Oocyst with numerous spherical spores, each with 3–10 sporozoites. Several species. Life cycle (Nabih, 1938).

K. helicina S. In the kidneys of various land-snails, belonging to genera Helix, Succinea, and Vitrina; oocyst with a double envelope 120–180μ in diameter; spores 12μ in diameter, with 5–6 sporozoites (Debaisieux, 1911). Cytology and development (Naville, 1927).

Genus Orcheobius Schuberg and Kunze. Macrogametes vermiciform; oocyst with 25–30 spores, each with 4 (or 6) sporozoites.
Fig. 254. a, Adelina dimidiata, a spore, ×1000 (Schellack); b, A. octospora, oocyst, ×1000 (Hesse); c, Orcheobius herpobdellae, ×550 (Kunze); d, e, Klossiella muris (d, renal cell of host with 14 sporoblasts; e, spore), ×280 (Smith and Johnson); f, Legerella hydropori, oocyst, ×1000 (Vincent); g, h, Haemogregarina of frog, ×1400 (Kudo); i–m, H. simondi, in the blood of the sole, Solea vulgaris, ×1300 (Laveran and Mesnil); n, Hepatozoon muris, spore, ×420 (Miller); o, Karyolysus lacertae, ×700 (Reichenow).

O. herpobdellae S. and K. (Fig. 254, c). In the testis of Herpobdella atomaria; mature macrogametes 180μ by 30μ; microgametes 50μ by 12μ; schizogony in April and May; sporogony in June and July.

Genus Klossiella Smith and Johnson. Microgametocyte produces 2 microgametes; oocyst with many spores, each with numerous sporozoites; in the kidney of mammals (Smith and Johnson, 1902).

K. muris S. and J. (Fig. 254, d, e). Oocyst with 12–14 spherical spores; about 30–34 sporozoites in a spore, 16μ by 13μ; spores discharged in the host’s urine; in the epithelium of the tubules and glomeruli in the kidney of the mouse, Mus musculus.

K. cobayae Seidelin. Oocyst with 8–20 spores; spore with about 30 sporozoites; in the kidney of guinea pig.

Genus Legerella Mesnil. Oocyst contains numerous sporozoites; spores entirely lacking; in arthropods (Mesnil, 1900).

L. hydropori Vincent (Fig. 254, f). In the epithelium of Malpighian
tubules of *Hydroporus palustris*; oocysts ovoid, 20–25μ long, with 16 sporozoites which measure 17μ by 3μ (Vincent, 1927).

Genus **Chagassella** Machado. Oocyst with 3 spores, each with 4 or 6 (or more) sporozoites; in hemipterous insects. 

*C. hartmanni* (Chagas). In the gut of *Dysdercus ruficollis*; oocysts with 3 spores, each with 4 or 6 (or more) sporozoites, about 35μ by 15μ (Machado, 1911).

Genus **Ithania** Ludwig. Microgametocyte produces four microgametes; oocyst with one to four spores, each with nine to 33 sporozoites. One species (Ludwig, 1947).

*I. weririchi* L. In the epithelial cells of the gastric caeca and mid-gut of the larvae of the crane-fly, *Tipula abdominalis*; oocysts 34–63μ by 22–50μ.

**Family 2 Haemogregarinidae** Léger

With 2 hosts: vertebrates (circulatory system) and invertebrates (digestive system).

Genus **Haemogregarina** Danilewsky. Schizogony takes place in blood cells of vertebrates; when gametocytes are taken into gut of leech or other blood-sucking invertebrates, sexual reproduction takes place; microgametocyte develops 2 or 4 microgametes; sporozoites formed without production of spores.

*H. stepanowi* D. (Fig. 255). Schizogony in *Emys orbicularis* and sexual reproduction in *Placobdella catenigera*; sporozoites introduced into blood of the chelonian host by leech (a), and enter erythrocytes in which they grow (d–g); schizogony in bone-marrow, each schizont producing 12–24 merozoites (h); schizogony repeated (i); some merozoites produce only 6 merozoites (j, k) which become gametocytes (l–o); gametogony occurs in leech; 4 microgametes formed from each microgametocyte and become associated with macrogamocytes in gut of leech (p–r); zygote (s) divides three times, and develops into 8 sporozoites (t–w).

Haemogregarines are found commonly in various birds (Aragão, 1911), reptiles, amphibians (Fig. 254, g, h) (Roudabush and Coatney, 1937) and fishes (Fig. 254, i–m).

Genus **Hepatozoon** Miller. Schizogony in the cells of liver, spleen, and other organs of vertebrates; merozoites enter erythrocytes or leucocytes and develop into gametocytes; in blood-sucking arthropods (ticks, mites), micro- and macro-gametes develop and unite in pairs; zygotes become oocysts which increase in size and produce sporoblasts, spores, and sporozoites.

*H. muris* (Balfour) (Fig. 254, n). In various species of rat; several
Fig. 255. The life-cycle of Haemogregarina stepanowi, ×1200 (Reichenow). a, sporozoite; b–i, schizogony; j–k, gametocyte-formation, l, m, microgametocytes; n, o, macrogametocytes; p, q, association of gametocytes; r, fertilization; s–w, division of the zygote nucleus to form eight sporozoites.
specific names were proposed on the basis of difference in host, locality, and effect on the host, but they are so indistinctly defined that specific separation appears to be impossible. Schizogony in the liver of rat; young gametocytes invade mononuclear leucocytes and appear as haemogregarines; when blood is taken in by the mite, *Laelaps echidninus*, union of 2 gametes produces vermicular body which penetrates gut epithelium and reaches peri-intestinal tissues and grows; becoming surrounded by a cyst-membrane, cyst content breaks up into a number of sporoblasts and then into spores, each of which contains a number of sporozoites; when a rat devours infected mites, it becomes infected.

Genus *Karyolysus* Labbé. Sporoblasts formed in the oocysts in gut epithelium of a mite, vermiform sporokinete, enter host ova and become mature; when young mites hatch, spores in gut epithelium are cast off and discharged in faeces; a lizard swallows spores; liberated sporozoites enter endothelial cells in which schizogony takes place; merozoites enter erythrocytes as gametocytes which when taken in by a mite complete development in its gut.

*K. lacertae* (Danilewsky) (Fig. 254, o). In *Lacerta muralis*; sexual reproduction in *Liponyssus saurarum*; sporokinetes 40–50μ long; spores 20–25μ in diameter (Reichenow, 1913, 1921).

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Chapter 26
Order 3 Haemopiridia Danilewsky

The development of the Haemopiridia is, on the whole, similar to that of the Coccidia in that they undergo asexual reproduction or schizogony, and also sexual reproduction resulting in sporozoite-formation; but the former takes place in the blood of vertebrates and the latter in the alimentary canal of some blood-sucking invertebrates. Thus one sees that the Haemopiridia remain always within the body of one of the two hosts; hence, the sporozoites do not possess any protective envelope.

The Haemopiridia are minute intracorpuscular parasites of vertebrates. The malarial parasites of man are typical members of this order. The development of Plasmodium vivax is briefly as follows (Fig. 256). An infected female anopheline mosquito introduces sporozoites into human blood when it feeds on it through skin (a). The sporozoites are fusiform and 6–15 μ long. They are capable of slight vibratory and gliding movement when seen under the microscope after removal from mosquitoes. After about 7–10 days of exo-erythrocytic development (p. 602), the organisms are found in erythrocytes (c, d) and are called schizonts. At the beginning the schizonts are small rings. They grow and finally divide into 12–24 or more merozoites (e, f) which are presently set free in the blood plasma (g). This schizogony requires 48 hours. The freed merozoites will, if not ingested by leucocytes, enter and repeat schizogony in the erythrocytes. After repeated and simultaneous schizogony in geometric progression, large numbers of infected erythrocytes will be destroyed at intervals of 48 hours, apparently setting free ever-increasing amounts of toxic substances into the blood. This is the cause of the regular occurrence of a characteristic paroxysm on every third day.

In the meanwhile, some of the merozoites develop into gametocytes instead of undergoing schizogony (h–k). When fully formed they are differentiated into macro- and micro-gametocytes, but remain as such while in the human blood. When a female anopheline mosquito takes in the blood containing gametocytes, the microgametocyte develops into 4–8 microgametes (k, l), and the macrogametocyte into a macrogamete (i, m) in its stomach. An ookinete (zygote) is formed when a microgamete fuses with a macrogamete (m, n). The ookinetes are motile. As they come in contact with the stomach epithelium, they enter it and become rounded into oocysts which lie between the base of the epithelium and the outer membrane of the stomach (o). Within the oocysts, repeated nuclear division
produces numerous sporozoites (p). When fully mature, the oocyst ruptures and the sporozoites are set free in the haemolymph through which they migrate to the salivary glands (q, r). The sporozoites make their way through the gland epithelium and finally to the duct of hypopharynx. They are ready to infect a human victim when the mosquito pierces with its proboscis the skin for another blood meal. Thus the sexual reproduction occurs in the mosquito (primary host) and the asexual reproduction, in man (secondary host).

The Haemosporidia are divided into three families:

With pigment granules
Schizogony in peripheral blood of vertebrates..Family 1 Plasmodiidae
Gametocytes in peripheral blood; schizogony elsewhere..................Family 2 Haemoproteidae (p. 618)
Without pigment granules; minute parasites of erythrocytes..........
..........................................................Family 3 Babesiidae (p. 622)

Family 1 Plasmodiidae Mesnil

Genus Plasmodium Marchiafava and Celli. Schizogony in erythrocytes and also probably in endothelial cells of man, mammals, birds, and reptiles; sexual reproduction in blood-sucking insects; widely distributed. Numerous species.

In all species, the infection in a vertebrate host begins under natural condition with the inoculation of the sporozoites by a vector mosquito. The form, size and structure of the sporozoites vary widely within a species so that identification of the species in this stage appears to be impossible (Boyd, 1935). Until some 20 years ago, it had been generally believed that the sporozoites upon entering the blood, penetrate and enter immediately the erythrocyte and begin intracorporeal development, which process Schaudinn (1902) reported to have seen in life. In this the eminent pioneer protozoologist was in error, since no one has up to the present time been able to confirm his observation. Et. and Edm. Sergent (1922) were the first to find that quinine given in large doses to the canaries on the day the birds were bitten by Culex mosquitoes infected with Plasmodium relictum, did not prevent infection in the birds. During the course of studies on P. vivax in cases of general paresis, Yorke and MacFie (1924) discovered that if quinine was given before the inoculation of infected blood, no infection resulted, but if the sporozoites were inoculated, quinine did not prevent infection. Similar observations were made on other species of malarial organisms. James (1931) suggested the possibility that the sporozoites are carried away from peripheral to visceral circulation and develop in the cells of the reticulo-endothelial system.
Fig. 256. The life-cycle of *Plasmodium vivax* (Kudo). a, sporozoite entering human blood; b, exoerythrocytic stage; c, the initiation of the erythrocytic development; d, a young schizont (“ring form”); e-g, schizogony; h, i, macrogametocytes; j, k, microgametocytes; l, microgamete-formation in the stomach of a mosquito; m, union of the gametes; n, zygote or ookinete; o, rounding up of an ookinete in the stomach wall; p, oocyst in which sporozoites are developing; q, mature oocyst ruptured and sporozoites are set free in the haemolymph; r, sporozoites entering the salivary gland cells.
Boyd and Stratman-Thomas (1934) found that the peripheral blood of a person who had been subjected to the bites of 15 anopheline mosquitoes infected by *Plasmodium vivax*, did not become infectious to other persons by subinoculation until the 9th day and that the parasites were not observed before the 11th day in the stained films of the peripheral blood. Warren and Coggeshall (1937) observed that when suspensions of the sporozoites of *P. catheemerium* obtained from infected *Culex pipiens*, were inoculated into canaries, the blood was not infectious for 72 hours, but emulsions made from the spleen, liver and bone marrow contained infectious parasites which brought about infection by subinoculations in other birds. These and many similar observations cannot be satisfactorily explained if one follows Schaudinn’s view. The fact that *P. elongatum* is capable of undergoing schizogony in the leucocytes and reticuloendothelial cells in addition to erythrocytes of host birds had been observed by Raffaele (1934) and Huff and Bloom (1935).

As to the nature of development of *Plasmodium* during the pre-patent period, James and Tate (1938) showed that there occur schizonts and schizogonic stages in the endothelial cells of the spleen, heart, liver, lung, and brain of the birds infected by *P. gallinaceum* (Fig. 257). They suggested the term *exoerythrocytic* to this schizogony in contrast to the well known *erythrocytic schizogony*. Huff and his co-workers made a series of detailed studies of pre-erythrocytic stages of this avian species. According to Huff and Coulston (1944), the sporozoites that are inoculated into the skin of chickens, are engulfed by phagocytes in 0.5–6 hours. In heterophile leucocytes, the sporozoites are apparently killed, but in the cells of lymphoid-macrophage system they develop into *cryptozoites* (Huff, Coulston and Cantrell, 1943) by assuming a spheroid shape and increasing in size for the first 36 hours, during which time there is a rapid repeated division of the nucleus. The schizogony is completed in 36 to 48 hours, each giving rise to 75–150 merozoites. These merozoites enter new lymphoid-macrophage and endothelial cells and become *metacryptozoites* which undergo schizogony similar to that of the cryptozoite. After three or four generations, the merozoites enter erythrocytes, and thus the erythrocytic stages appear in five to 10 days. Porter (1942) distinguishes two types of exoerythrocytic development in avian Plasmodium; namely, gallinaceum-type just quoted and elongatum-type.

Exoerythrocytic or *E.-E.* stages were further discovered in saurian *Plasmodium* (Thompson and Huff, 1944; Garnham, 1950) and in mammalian malaria organisms (Shortt and Garnham, 1948). In *P.*
Fig. 257. Exoerythrocytic schizogony in avian Plasmodium. a–f, *P. gallinaceum* in smears from chicks (James and Tate). a, monocyte from lung, infected by 2 young schizonts; b, monocyte from liver, with a growing trinucleate schizont; c, monocyte from lung, with a large multinucleate schizont; d, large mature schizont containing many mature merozoites, free in lung; e, portion of broken schizont from lung, showing the attached developing merozoites. (×1660). f, a capillary of brain blocked by 3 large schizonts (×740). g, h, *P. cathelemrium* in sections of organs of canaries (Porter; ×1900). g, capillary in the brain, showing an endothelial cell infected with a uninucleate and a multinucleate schizont; h, a multinucleate schizont and a group of merozoites found in a capillary of heart muscle.

cynomolgi, Shortt and Garnham report that the E.-E. stages occur in the parenchymatous cells of the liver of host monkeys and are inclined to think that there is one generation only. The earliest forms were seen on the fifth day after the inoculation of the sporozoites. They are rounded bodies, about 10 μ in diameter and contain about 50 chromatin granules of irregular shape. They grow in size to about 35 μ in diameter, and divide in eight to nine days into some 1000 merozoites, each measuring about 1 μ. These merozoites presumably invade the erythrocyte. In *P. vivax*, the E.-E. stages develop in the parenchymatous cells of the liver also and resemble those of *P. cynomolgi*. The forms found on the seventh day after sporozoite-inoculation were slightly larger (about 42 μ in diameter) than those
of *P. cynomolgi*, and when mature, give rise to 800–1000 merozoites.

Thus exoerythrocytic stages and development have definitely been demonstrated for *Plasmodium* in various host groups, although morphological and developmental details, distinction between them and other little known organisms such as *Toxoplasma* (p. 625) and interrelationship between them and erythrocytic stages, had to be looked for in future investigations (Fig. 258). General review of E.-E. development (Huff, 1947, 1948; Garnham, 1948).

![Diagram](Image)

**Fig. 258.** Diagrammatical life-cycle of an avian *Plasmodium* (Several authors). Well established phases are connected by solid lines, while undetermined and recently suggested phases are indicated by broken lines. a, sporozoite injected into host bird by a mosquito; b–e, exoerythrocytic schizonts and schizogony in monocytes; f–i, commonly seen schizogony in erythrocytes; j, macrogametocyte; k, microgametocyte.

The incubation period of *Plasmodium* infections in man varies due to various factors such as the strain, vitality and number of the sporozoites injected by the mosquitoes, the varied susceptibility on the part of host, etc. Boyd and co-workers found that the incubation periods for the three species of human *Plasmodium* which they studied were, as follows: In *P. vivax*, 8–21 days (the majority 11–14 days)
after the bites of infected mosquitoes, but in one case as long as 304 days; in *P. malariae*, 4–5 weeks, with the onset of fever lagging 3–12 days behind; and in two strains of *P. falciparum*, one, 6–25 days and the other, 9–13 days; in another observation, *P. falciparum* was observable in the peripheral blood in 5–9 days and the onset of fever in 7–12 days.

The paroxysm of malaria is usually divisible into three stages: chill or rigor stage, high temperature or febrile stage (104°F or over) and sweating or defervescent stage. The time of paroxysm corresponds, as was stated already, with the time of liberation of merozoites from erythrocytes, and is believed to be due to extrusion of certain substance into the blood plasma. The nature of this material is however unknown at present. In the grown schizonts as well as in gametocytes of *Plasmodium*, are found invariably yellowish brown to black pigment granules which vary in form, size and number among different species. They are usually called haemozoin granules and are apparently the catabolic products formed within the parasites. The pigment of *P. gallinaceum* and *P. cynomolgi* has been identified with haematin (ferri protoporphyrin) (Rimington and Fulton, 1947). The pigment possesses certain taxonomic significance, as will be described below. The infected erythrocytes, if stained deeply, may show a punctate appearance. These dots are small and numerous in the erythrocytes infected by *P. vivax* and *P. ovale*, and are known as Schüffner’s (1899) dots, while those in the host cells infected by *P. falciparum* are few and coarse and are referred to as Maurer’s (1902) dots. No dots occur in the erythrocytes infected by *P. malariae*. Pathology (Maegraith, 1948); splenomegaly (Darling, 1924, 1926; Russell, 1935, 1952a; Hackett, 1944); histopathology (Taliaferro and Mulligan, 1937); character of paroxysm (Kitchen and Putnam, 1946); blood proteins during infection (Boyd and Proske, 1941); stippling of erythrocytes (Thomson, 1928).

The condition which brings about the formation of gametocytes is not known at present. The gametocytes appear in the peripheral blood at various intervals after onset of fever, and remain inactive while in the human blood. The assumption that the macrogametocytes undergo parthenogenesis under certain conditions and develop into schizonts as advocated by Grassi, Schaudinn and others, does not seem to be supported by factual evidence. The initiation of further development appears to be correlated with a lower temperature and also a change in pH of the medium (Manwell). If living mature microgametocytes of human *Plasmodium* taken from an infected person are examined microscopically under a sealed cover glass
at room temperature (18–22°C.), development takes place in a short while and motile microgametes are produced ("exflagellation"). Similar changes take place when the gametocytes are taken into the stomach of mosquitoes belonging to genera other than Anopheles, but no sexual fusion between gametes occurs in them and all degenerate sooner or later. In the stomach of an anopheline mosquito, however, the sexual reproduction of human Plasmodium continues, as has been stated before.

All species are transmitted by adult female mosquitoes. The males are not concerned, since they do not take blood meal. The species of Plasmodium which attack man are transmitted only by the mosquitoes placed in genus Anopheles, while the majority of the avian species of Plasmodium are transmitted by those which belong to genera Culex, Aedes, and Theobaldia. The chief vectors of the human malarial parasites in North America are *A. quadrir maculatus* (eastern, southern and middle-western States), *A. punctipennis* (widely distributed), *A. crucians* (southern and south-eastern coastal area), *A. walkeri* (eastern area), and *A. maculipennis freeborni* (Pacific coast). Boyd and coworkers observed that (1) *A. quadrir maculatus* and *A. punctipennis* were about equally susceptible to *Plasmodium vivax*; (2) *A. quadrir maculatus* was susceptible to several strains of *P. falci parum*, while *A. punctipennis* varied from highly susceptible to refractory to the same strains; (3) *A. quadrir maculatus* was more susceptible to all three species of Plasmodium than coastal or inland *A. crucians*. Thus *A. quadrir maculatus* is the most dangerous malaria vector in the United States as it shows high susceptibility to all human Plasmodium. *A. pseudopunctipennis* distributed from south-western United States to Argentina and *A. albimanus* occurring in Central America, are but a few out of many anopheline vectors of human Plasmodium in the areas indicated. Host-parasite relation (Boyd and Coggeshall, 1938); malaria vectors of the world (Komp, 1948); susceptibility of Anopheles to malaria (King, 1916; Boyd and Kitchen, 1936); epidemiology in North America (Boyd, 1941), in Brazil (Boyd, 1926), in Jamaica (Boyd and Aris, 1929), in Cuba (Carr and Hill, 1942), in Trinidad and British West Indies (Downs, Gillette and Shannon, 1943), in Porto Rico (Earle, 1930, 1939), in Haiti (Paul and Bellerive, 1947), in Philippine Islands (Russell, 1934, 1935a), in India (Russell and Jacob, 1942) and in Liberia; general picture (Russell, 1952, 1952a); mosquito control (Russell, 1952a)

The time required for completion of sexual reproduction of Plasmodium in mosquitoes varies according to various conditions such as species and strain differences in both Plasmodium and Anopheles,
temperature, etc. Boyd and co-workers showed that when the anophelines which fed on patients infected by \textit{P. vivax} were allowed to feed on other persons, their infectivity was as follows: 1–10 days after infective feeding, 87.2\%; 11–20 days, 93.8\%; 21–30 days, 78\%; 31–40 days, 66\%; 41–50 days, 20\%; and over 50 days, none. In a similar experiment with \textit{P. falciparum}, during the first 10 days the infection rate was 84\%, but thereafter the infectivity rapidly diminished until there was no infection after 40 days. It is generally known that the development of the parasites in mosquitoes depends greatly on temperature. Although the organisms may survive freezing temperature in mosquitoes (Coggeshall), sporozoite-formation is said not to take place at temperatures below 16° C. or above 35° C. (James). According to Stratman-Thomas (1940), the development of \textit{Plasmodium vivax} in \textit{Anopheles quadrimaculatus} is completed within the temperature range of 15–17° to 30° C. It varies from 8 to 38 days after infective feeding. The optimum temperature is said to be 28° C. at which the development is completed in the shortest time. A period of 24 hours at 37.5° C. will sterilize all but a very small per cent of \textit{Anopheles quadrimaculatus} of their \textit{Plasmodium vivax} infection. This has a bearing on the transmission of \textit{Plasmodium vivax} in summer months. In certain localities oocysts may survive the winter and complete their development in the following spring. Duration of infection in \textit{Anopheles} (Boyd and St.-Thomas, 1943a; Boyd, St.-Thomas and Kitchen, 1936).

There are three long-recognized species of human \textit{Plasmodium}. They are \textit{P. vivax}, \textit{P. falciparum} and \textit{P. malariae}. To these \textit{P. ovale} is here added. Each species appears to be represented by numerous strains or races as judged by the differences in virulence, immunological responses, incubation period, susceptibility to quinine, etc. (Boyd, 1934, 1940, 1940a; Boyd and Kitchen, 1948).

Malaria has been, and still is, perhaps the most important protozoan disease of man. In India alone, malaria fever is held to be the direct cause of over a million deaths annually among nearly 100 million persons who suffer from it (Sinton, 1936). In the United States, the disease had been prevalent in places in south-eastern States. But since 1945, cases of malaria have rapidly declined and there is prospect of the disappearance of endemic malaria from the United States (Andrews, Quinby and Langmuir, 1950; Andrews, 1951). In malarious countries, the disease is a serious economic and social problem, since it affects the majority of population and brings about a large number of persistent sickness, the loss of man power and retardation of both mental and physical development among
children. History of malaria (Ross, 1928; Boyd, 1941; Russell, 1943); general reference (Boyd, 1949; Russell, West and Manwell, 1946); antimalarial drugs (Russell, 1952a).

It must be added here that human ingenuity has been for nearly 30 years utilizing the malarial organisms in combating another disease; namely, naturally induced malaria therapy has been successfully used in the treatment of patients suffering from general paresis

![Fig. 259. Plasmodium vivax, ×1535 (Original). a, young ring-form; b, c, growing schizonts; d, two schizonts in an erythrocyte; e, f, large schizonts; g-i, schizogonic stages; j, fully developed merozoites; k, macrogametocyte; l, microgametocyte.](image)

and other forms of neuro-syphilis. Technique (Boyd and Stratman-Thomas, 1933; Boyd, St.-Thomas and Kitchen, 1936a; Boyd, St.-Thomas, Kitchen and Kupper, 1938; Mayne and Young, 1941).

*P. vivax* (Grassi and Feletti) (Fig. 259). The benign tertian malaria parasite; schizogony completed in 48 hours and paroxysm every third day. *Ring forms*: About 1/4–1/3 the diameter of erythrocytes; unevenly narrow cytoplasmic ring is stained light blue (in Giemsa) and encloses a vacuole; nucleus stained dark-red, conspicuous. *Growth period*: Irregular amoeboid forms; host cell slightly enlarged; Schüffner's dots begin to appear. *Grown schizonts*: In about 26 hours after paroxysm; occupy about 2/3 of the enlarged erythrocytes, up to 12μ in diameter, which are distinctly paler than uninfected ones; Schüffner's dots more numerous; brownish haemoglobin granules; a large nucleus. *Schizogonic stages*: Repeated nuclear division produces
HAEMOSPORIDIA

12–24 or more merozoites; multinucleate schizonts about 8–9μ in diameter; haemozoin granules in loose masses; merozoites about 1.5μ long. 

**Gametocytes:** Time required for development of ringform into a mature gametocyte is estimated to be about four days; smoothly rounded body, occupying almost whole of the enlarged erythrocytes; brown haemozoin granules numerous. Macrogametocytes are about 9–10μ in diameter, stain more deeply and contain a small compact nucleus; microgametocytes are a little smaller (7–8μ in diameter), stain less deeply and contain a less deeply staining large nucleus. This species is said to invade reticulocytes rather than erythrocytes (Kitchen, 1938). Boyd (1953a) distinguished five series of erythro-

cytic organisms on the basis of nuclear and cytoplasmic characteristics. The organisms of series A give rise by schizogony to organisms of series B or D which in turn produce series C (microgametocytes) or series E (macrogametocytes). Onset of infection is said to occur usually when the parasite density is less than 100 per mm³ (Boyd, 1944). Incubation period (Boyd and Stratman-Thomas, 1933c, 1934); concentration of organisms (Ferrebee and Geiman, 1946); immunity (Boyd and Stratman-Thomas, 1933a, b; Boyd and Kitchen, 1936a; Boyd, 1947); susceptibility (Boyd and Stratman-Thomas, 1933c, 1934).

The benign malaria fever parasite is the commonest and the most widely distributed species in the tropical and subtropical regions as well as in the temperate zone. It has been reported as far north as the Great Lakes region in North America; England, southern Sweden and northern Russia in Europe; and as far south as Argentina, Australia, and Natal in the southern hemisphere. Generally speaking this spe-

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**Fig. 260.** *Plasmodium falciparum,* ×1535 (Original). a, three ring-forms in an erythrocyte; b, a somewhat grown schizont in an erythrocyte with Maurer's dots; c–f, growing and schizogonic stages, g, h; merozoite formation; i, macrogametocyte; j, microgametocyte.
cies predominates in the spring and early summer over the other species.

*P. falciparum* (Welch) (*Laverania malariae* Grassi and Feletti; *P. tenue* Stepens) (Fig. 260). The subtertian, malignant tertian or aestivo-autumnal fever parasite; schizogonic cycle is somewhat irregular, though generally about 48 hours. **Ring forms:** Much smaller than those of *P. vivax*; about 1μ in diameter; marginal forms and multiple (2–6) infection common; nucleus often rod-form or divided into two granules; in about 12 hours after paroxysm, all schizonts disappear from the peripheral blood. **Growth and schizogonic stages:** These are almost exclusively found in the capillaries of internal organs; as schizonts mature, Maurer’s dots appear in the infected erythrocytes; when about 5μ in diameter, nucleus divides repeatedly and 8–24 or more small merozoites are produced; haemozoin granules dark brown or black and usually in a compact mass; infected erythrocytes are not enlarged. **Gametocytes:** Mature forms sausage-shaped ("crescent"), about 10–12μ by 2–3μ; appear in the peripheral blood. Macrogametocytes stain blue and contain a compact nucleus and coarser granules, grouped around nucleus; microgametocytes stain less deeply blue or reddish, and contain a large lightly staining nucleus and scattered smaller haemozoin granules. The organism invades both mature and immature erythrocytes (Kitchen, 1939). Cytological study of microgametocytes and microgametes (MacDougall, 1947); different strains (Kitchen and Putnam, 1943); induced infection (Boyd and Kitchen, 1937); incubation period (Boyd and Kitchen, 1937b; Boyd and Matthews, 1939); immunity (Boyd and Kitchen, 1945).

The subtertian fever parasite is widely distributed in the tropics. In the subtropical region, it is more prevalent in late summer or early autumn. It is relatively uncommon in the temperate zone. The malignancy of the fever brought about by this parasite is attributed in part to decreased elasticity of the infected erythrocytes which become clumped together into masses and which adhere to the walls of the capillaries of internal organs especially brain, thus preventing the circulation of blood through these capillaries.

*P. malariae* (Laveran) (Fig. 261). The quartan malaria parasite; schizogony in 72 hours and paroxysm every fourth day. **Ring forms:** Similar to those of *P. vivax*. **Growth period:** Less amoeboid, rounded; in about 6–10 hours haemozoin granules begin to appear; granules are dark brown; in 24 hours, schizonts are about 1/2 the diameter of erythrocytes which remain normal in size; schizonts often stretched into "band-form" across the erythrocytes; no dots comparable with Schüffner’s or Maurer’s dots. **Mature and segmenting**
schizonts: In about 48 hours, schizonts nearly fill the host cells; rounded; haemoglobin granules begin to collect into a mass; nuclear divisions produce 6–12 merozoites which are the largest of the three species and may often be arranged in a circle around a haemoglobin mass. Gametocytes: Circular; with haemoglobin granules. Macrogametocytes stain more deeply and contain a small, more deeply staining nucleus and coarser granules; microgametocytes stain less deeply and contain a larger lightly stained nucleus and finer and numerous granules. The organism invades most frequently mature red corpuscles (Kitchen, 1939).

The quartan fever parasite is distributed in the tropics and subtropics, though it is the least common of the three species. As a rule, in an area where the three species of Plasmodium occur, this species seems to appear later in the year than the other two.

*P. ovale* Stevens (Fig. 262). The Ovale or mild tertian fever parasite; schizogony in about 48 hours; its morphological characters resemble both *P. vivax* and *P. malariae*. Ring forms: Similar to those of the two species just mentioned; Schüffner's dots appear early. Growth period: Infected erythrocytes are more or less oval with irregular fimbriated margin; slightly enlarged; not actively amoeboid, sometimes in band-form; with dark brown haemoglobin.
granules; Schüffner’s dots abundant. Schizogonic stages: 6–12 merozoites. Gametocytes: Resemble closely those of P. malariae; host cells with Schüffner’s dots and slightly enlarged.

This organism appears to be confined to Africa and Asia (Philippine Islands and India). Several malariologists doubt the validity of the species.

The malarial parasites are ordinarily studied in stained blood films (p. 899). Table 11 will serve for differential diagnosis of the three common species.

Several species of Plasmodium have been observed in primates and monkeys, some of which resemble strikingly the human species. Here a few species will be mentioned. Other species (Aberle, 1945).

P. kochi (Laveran) (Fig. 263, a–f). In the monkeys belonging to the genera: Callicebus, Cercocebus, Cercopithecus, Erythrocebus, and Papio; schizogony in 48 hours; organism resembles P. vivax; infected erythrocytes become enlarged and sometimes stippling like Schüffner’s dots occurs; eight to 14 merozoites; gametocytes large and spheroid.

P. brasilianum Gonder and Berenberg-Gossler (Fig. 263, g–l). In New World monkeys belonging to the genera: Alouatta, Ateles, Cacajao and Cebus; schizogony in 72 hours; it resembles P. malariae; no enlargement of infected erythrocytes; band-form schizonts; number of merozoites vary according to the difference in hosts, averaging eight to 10; gametocytes rounded, comparatively small in number (Taliaferro and Taliaferro, 1934). Haematology (Taliaferro and Klüver, 1940).

P. cynomolgi Mayer (Fig. 263, m–r). In Macaca irus (Macacus
Table 11.—Differential diagnosis of three species of human *Plasmodium*

<table>
<thead>
<tr>
<th></th>
<th><em>P. vivax</em></th>
<th><em>P. falciparum</em></th>
<th><em>P. malariae</em></th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Ring forms</strong></td>
<td>About $\frac{1}{4}$–$\frac{1}{2}$ the diameter of erythrocytes; a single granular nucleus.</td>
<td>About $\frac{1}{4}$–$\frac{1}{2}$ the diameter of erythrocytes; marginal forms and multiple (2–6) infection common.</td>
<td>Similar to those of <em>P. vivax</em>; cytoplasm slightly denser.</td>
</tr>
<tr>
<td><strong>Infected erythrocytes</strong></td>
<td>Much enlarged, up to 12μ in diameter, paler than normal (7.5μ in diameter) erythrocytes; Schüffer’s dots.</td>
<td>Normal, some are distorted or contracted in later schizogonic period; Maurer’s dots.</td>
<td>Not enlarged; sometimes slightly smaller than uninfected ones; no dots.</td>
</tr>
<tr>
<td><strong>Growing schizonts</strong></td>
<td>Irregularly ameboid; vacuolated; paler; small yellowish brown haemoglobin granules.</td>
<td>Partly grown ring forms often with rod-shaped or 2 granular nuclei; further development not seen in peripheral blood.</td>
<td>Not ameboid; oval, rounded, band-form, rarely irregular; less vacuolated cytoplasm deeper blue; dark brown granules.</td>
</tr>
<tr>
<td><strong>Fully grown schizonts</strong></td>
<td>Irregular in form; about $\frac{1}{3}$ the enlarged erythrocytes; vacuolated; brown haemoglobin granules.</td>
<td>Only in internal organs; $\frac{1}{2}$–$\frac{2}{3}$ of erythrocytes; dark haemoglobin in compact mass.</td>
<td>Nearly filling erythrocytes; rounded; cytoplasm deeper blue; dark brown pigment granules.</td>
</tr>
<tr>
<td><strong>Schizogonic stages</strong></td>
<td>12–24 or more merozoites; irregularly arranged in much enlarged host cells.</td>
<td>Only in internal organs; 8–24 or more small merozoites; irregularly arranged; dark pigment.</td>
<td>6–12 merozoites which are the largest of all, typically arranged in a circle.</td>
</tr>
<tr>
<td><strong>Gametocytes</strong></td>
<td>Almost filling enlarged erythrocytes; rounded or oval; with brown pigment granules.</td>
<td>Sausage-shaped; haemoglobin dark brown; in the peripheral blood.</td>
<td>Filling normal-sized erythrocytes; round or ovoid, much smaller than those of <em>P. vivax</em>; dark brown pigment.</td>
</tr>
</tbody>
</table>

*cynomolagus*); schizogony in 48 hours; eight to 22 merozoites; infected erythrocytes slightly enlarged and stippled; vectors are Anopheles. Schizogony (Wolfson and Winter, 1946; Taliaferro and Mulligan, 1937); morphology (Mulligan, 1935); cellular changes in host (Taliaferro and Mulligan, 1937).
Fig. 263. Plasmodium of monkeys. Column 1, ring forms; 2, 3, growing trophozoites; 4, segmenting schizonts; 5, macrogametocytes; 6, microgametocytes. a–f, *Plasmodium kochi*, ×1665 (Gonder and Berenberg-Gossler); g–l, *P. brasilianum*, ×1665; m–r, *P. cynomolgi*, ×2000; s–x, *P. knowlesi*, ×2000 (Taliaferro and Taliaferro).

*P. knowlesi* Sinton and Mulligan (Fig. 263, s–x). In *Macaca irus*; experimentally man is susceptible; schizogonic cycle in 24 hours; six to 16 merozoites; infected erythrocytes are somewhat distorted. Morphology and development (Brug, 1934; Mulligan, 1935; Taliaferro and Taliaferro, 1949); infections in man (Milam and Coggshall, 1938).


Many species of *Plasmodium* have been reported from numerous species of birds in which are observed clinical symptoms and pathological changes similar to those which exist in man with malaria infection. In recent years the exoerythrocytic stages have been intensively studied in these forms. According to Hegner and co-
workers the erythrocytes into which merozoites enter are often the most immature erythrocytes (polychromatophilic erythroblasts). The species of avian Plasmodium are transmitted by adult female mosquitoes belonging to Culex, Aedes or Theobaldia. Some of the common species are briefly mentioned here. Avian Plasmodium (Manwell, 1933a; Hewitt, 1940b); avian hosts (Wolfson, 1941); distribution (Manwell and Herman, 1935; Herman, 1938; Hewitt, 1940a; Wood and Herman, 1943).

*P. relictum* Grassi and Feletti (*P. praecox* G. and F.; *P. inconstans* Hartman) (Fig. 264, *a*). In English sparrow (*Passer domesticus*) and other passerine birds, also in mourning doves and pigeons (Coatney 1938); schizogony varies in different strains, in 12, 24, 30 or 36 hours; 8–15 or 16–32 merozoites from a schizont; gametocytes rounded, with small pigment granules; host-cell nucleus displaced; canaries (*Serinus canaria*) susceptible; many strains; transmitted by Culex, Aedes and Theobaldia; widely distributed. Duration of infection (Manwell, 1934; Bishop, Tate and Thorpe, 1938); variety (Manwell, 1940); in *Culex pipiens* (Huff, 1934); development in birds (Mudrow and Reichenow, 1944); relationship of E.-E. and erythrocytic stages (Sergent, 1949).

*P. vaughani* Novy and McNeal (Fig. 264, *b*). In robin (*Turdus m. migratorius*) and starling (*Sturnus v. vulgaris*); 4–8 (usually 4) merozoites from a schizont, ordinarily with 2 pigment granules; schizogony in about 24 hours; gametocytes elongate; host-cell nucleus not displaced.

*P. cathemerium* Hartman (Fig. 264, *c*). In English sparrow, cowbird, red-winged blackbird, and other birds; schizogony in 24 hours, segmentation occurs at 6–10 p.m.; 6–24 merozoites from a schizont; mature schizonts and gametocytes about 7–8μ in diameter; gametocytes rounded; haemozoin granules in microgametocytes longer and more pointed than those present in macrogametocytes; canaries susceptible; numerous strains; common; transmitted by many species of Culex and Aedes (Hartman, 1927). Relapse (Manwell, 1929); acquired immunity (Cannon and Taliaferro, 1931); in ducks (Hegner and West, 1941); cultivation (Hewitt, 1939); effect of plasmochin (Wampler, 1930).

*P. rouxi* Sergent, Sergent and Catanei (Fig. 264, *d*). In English sparrow in Algeria; similar to *P. vaughani*; schizogony in 24 hours; 4 merozoites from a schizont; transmitted by Culex.

*P. elongatum* Huff (Fig. 264, *e*). In English sparrow; schizogony occurs mainly in the bone marrow, and completed in 24 hours; 8–12 merozoites from a schizont; gametocytes elongate, found in periph-
eral blood; transmitted by Culex (Huff, 1930). Canaries and ducks are susceptible. Study of nucleus (Chen, 1944).

*P. circumflexum* Kikuth (Fig. 264, f). In the red-winged blackbird, cowbird and several other birds, including the ruffed grouse (Fallis, 1946); growing schizonts and gametocytes form broken rings around

![Fig. 264](https://example.com/fig264.png)

*Fig. 264.* a, *Plasmodium relictum*; b, *P. vaughani*; c, *P. cathemerium*; d, *P. rouxi*; e, *P. elongatum*; f, *P. circumflexum*; g, *P. polare*; h, *P. nucleophilum*; i, *P. gallinaceum*; j, *P. hexamertium*; k, *P. oti*; l, *P. lophurae*. Columns 1, ring-forms; 2, growing schizonts; 3, segmenting schizonts; 4, macrogametocytes; and 5, microgametocytes. X about 1400 (Several authors; from Hewitt, modified).
the host-cell nucleus; schizogony in 48 hours; 13–30 merozoites; gametocytes originate, with a few haemozoin granules; transmission by Theobaldia (Herman, 1938b).

*P. polare* Manwell (Fig. 264, g). In cliff swallow (*Petrochelidon l. lunifrons*); grown schizonts at one of the poles of host erythrocytes; 8–14 merozoites from a schizont; few in peripheral blood; gametocytes originate (Manwell, 1935a).

*P. nucleophilum* M. (Fig. 264, h). In catbird (*Dumatella carolinensis*); schizogony in 24 hours; 3–10 merozoites from a schizont; mature schizonts usually not seen in the peripheral blood; gametocytes originate, often seen closely applied to the host-cell nucleus; haemozoin granules at one end (Manwell, 1935a).

*P. gallinaceum* Brumpt (Fig. 264, i). In domestic fowl (*Gallus domesticus*) in India; schizogony in 36 hours; 20–36 merozoites from a schizont; mature schizonts often displaced; pheasants, geese, partridges and peacocks are susceptible, but canaries, ducks, guinea fowls, etc., are refractory; transmitted by *Aedes* (Brumpt, 1935). E.-E. development (p. 602); vectors (Russell and Mohan, 1942); phosphorus 32 in study (Clarke, 1952); nucleic acids (Lewert, 1952).

*P. hexamerium* Huff (Fig. 264, j). In bluebird (*Sialis s. sialis*) and Maryland yellow-throats; schizogony in 48 or 72 hours; grown schizonts often elongate; 6 merozoites from a schizont; gametocytes elongate (Huff, 1935).

*P. oti* Wolfson (Fig. 264, k). In eastern screech owl (*Otus asio naevius*); 8 merozoites from a schizont; body outlines irregular, rough; gametocytes originate. Manwell (1949) considers this species identical with *P. hexamerium*.

*P. lophurae* Coggeshall (Fig. 264, l). In fire-back pheasant (*Lophura i. igniti*) from Borneo, examined at New York Zoological Park; 8–18 merozoites from a schizont; gametocytes large, elongate; host-cell nucleus not displaced; canaries are refractory, but chicks and especially ducks are highly susceptible (Coggeshall, 1938, 1941; Wolfson, 1940); young ducklings succumb less readily to its infection than older ducks (Becker, 1950). Experimentally *Aedes aegypti*, *A. albopictus* and *Anopheles quadrimaculatus* serve as vectors, but not *Culex pipiens* (Jeffery, 1944). Characteristics (Terzian, 1941); cultivation (Trager, 1950).

A number of lizards have recently been found to be infected by *Plasmodium*. A few species are described here briefly. Species (Thompson and Huff, 1944a; Laird, 1951).

*P. mexicanum* Thompson and Huff (Fig. 265). In *Sceloporus fer-
rariperezi of Mexico; experimentally *S. olivaceus*, *S. undulatus*, *Crotaphytus collaris*, *Phrynosoma cornutum* and *P. asio*, become infected; in erythrocytes and normoblasts, and in all types of circulating cells; host cells not hypertrophied; schizonts round to elongate; 10–40 merozoites; gametocytes 12–16µ by 6–7.7µ, only in haemoglobin-containing cells which become enlarged and distorted (Thompson and Huff, 1944); a mite, *Hirstella* sp., was considered to be a possible vector (Peláez, Reyes and Barrera, 1948).

**Fig. 265.** *Plasmodium mexicanum, X1780* (Peláez *et al.*). a, b, young and growing trophozoites in host’s erythrocyte; c, segmenting schizont; d, macrogametocyte; e, microgametocyte.

*P. rhadinurum* T. and H. In the erythrocytes of *Iguana iguana rhinolopha* in Mexico; schizonts extremely polymorphic with one or two long processes; 4–5 merozoites; gametocytes 6.5–7.7µ; vector unknown (Thompson and Huff, 1944a).

*P. floridense* T. and H. In the erythrocytes of *Sceloporus undulatus* in Florida; young trophozoites pyriform; 6–21(12) merozoites; gametocytes 7.5–8.0µ in diameter; vector unknown (Thompson and Huff, 1944a).


**Family 2 Haemoproteidae** Doflein

Schizogony occurs in the endothelial cells of vertebrates; merozoites enter circulating blood cells and develop into gametocytes; if blood is taken up by specific blood-sucking insects, gametocytes develop into gametes which unite to form zygotes that undergo changes similar to those stated above for the family Plasmodiidae.

**Genus Haemoproteus** Kruse. Gametocytes in erythrocytes, with pigment granules, halter-shaped when fully formed (hence *Halteridium* Labbé); schizogony in endothelial cells of viscera of vertebrate reptiles. Species (Cerny, 1933; Coatney and Roudabush, 1937); transmission experiments (Nöller, 1920).
H. columbae Celli and Sanfelice (Fig. 266). In pigeons (*Columba livia*), etc.; widely distributed; young schizonts, minute and uninucleate, are in the endothelial cells of lungs and other organs, grow into large multinucleate bodies which divide into 15 or more uninucleate cytomeres (Aragão). Each cytomere now grows and its nucleus divides repeatedly. The host cell in which many cytomeres undergo enlargement, becomes highly hypertrophied and finally ruptures. The multinucleate cytomeres break up into numerous merozoites, some of which possibly repeat the schizogony by invading endothelial cells, while others enter erythrocytes and develop into gametocytes which are seen in the peripheral blood; sexual reproduction

**Fig. 266.** The life-cycle of *Haemoproteus columbae*. (Several authors). a, a sporozoite entering an endothelial cell of the pigeon; b, growth of a schizont; c, segmentation of multinucleate schizont into uninucleate cytomeres; d–i, development of cytomeres to produce merozoites; j–m, development of microgametes; n–p, development of macrogamete; q, fertilization; r, s, ookinetes; t, a young oocyst in the stomach wall of a fly; u, a ruptured mature oocyst with sporozoites. a–k, n, o, in the pigeon, l, m, p–u, in *Pseudolynchia mauro*. 
in, and transmitted by, the flies: *Lynchia brunea, L. lividicolor, L. capensis, Pseudolynchia mauroa,* and *Microlynchia fusilla.* Nomenclature and relapse (Coatney, 1933).

*H. lophortyx* O’Roke. In California valley quail, Gambel quail, and Catalina Island quail (*Lophortyx*); gametocytes in erythrocytes, also occasionally in leucocytes; young gametocytes, spherical to elongate, about 1μ long; more developed forms, cylindrical, about 8μ by 2μ, with 2–10 pigment granules; mature gametocytes, halter-shaped, encircling the nucleus of the host erythrocyte, 18μ by 1.5–2.5μ; numerous pigment granules; 4–8 microgametes, about 13.5μ long, from each microgametocyte; on slide in one instance, gamete-formation, fertilization and ookinet formation, completed in 52 minutes at room temperature; in nature sexual reproduction takes place in the fly, *Lynchia hirsuta*; sporozoites enter salivary glands and fill central tubules; schizonts present in lungs, liver and spleen of quail after infected flies sucked blood from the bird; merozoites found in endothelial cells of capillaries of lungs, in epithelial cells of liver and rarely in peripheral blood cells; how merozoites enter blood cells is unknown; schizonts seldom seen in circulating blood; infected birds show pigment deposits in spleen and lungs (O’Roke, 1934). Duration of infection (Herman and Bischoff, 1949).


Genus *Leucocytozoon* Danilewsky. Schizogony in the endothelial cells as well as visceral cells of vertebrates; sexual reproduction in blood-sucking insects; gametocytes in spindle-shaped host cells. Several species (Cerny, 1933; Coatney and Roudabush, 1937).

*L. simondi* Mathis and Léger (*L. anatis* Wickware) (Fig. 267). Mathis and Léger (1910) described this species from the teal duck (*Querquedula crecca*) in Tonkin, China. Wickware (1915) saw *L. anatis* in ducks in Canada. O’Roke (1934) carried on experimental studies on the developmental cycle with the form which he found in wild and domestic ducks in Michigan. Herman (1938) observed the organism in common black ducks (*Anas rubripes tristis*), red-breasted merganser (*Mergus serrator*), and blue-winged teal (*Querquedula discors*) and considered *L. anatis* as identical with *L. simondi.* Huff (1942) studied the schizogony and gametocytes, and maintained the species he studied in mallard ducks (*Anas p. platyrhynchos*) and domestic ducks from Wisconsin, to be *L. simondi.*

According to O’Roke, the vector is the black fly, *Simulium venustum,* in which the sexual reproduction takes place. Gametocytes de-
velop into mature gametes in 1–2 minutes after blood is obtained from an infected duck; macrogametes about 8μ in diameter; 4–8 microgametes, 15.7–24.1μ long, from a single microgametocyte; zygotes are found in stomach contents of fly in 10–20 minutes after sucking

Fig. 267. The life-cycle of *Leucocytozoon simondi* (Brumpt, modified). a–c, development of macrogamete; d–f, development of microgametes; g, fertilization; h, ookinete; i, j, ookinete piercing through the stomach wall; k–m, development of sporozoites; n, sporozoites entering endothelial cells; o–r, schizogony.

in the infected blood of bird; motile ookinetes abundant after 5 hours, measure 33.3μ by 3–4.6μ; 22 hours after sucking duck blood, oocysts are found on outer wall of stomach; sporozoites mature probably in 24–48 hours; 5 days after a duck has been bitten by infected black flies, schizogonic stages are noticed in endothelial cells of capillaries of lungs, liver, spleen; on about 7th day gametocytes appear in blood; liver and spleen become hypertrophied; the infection among duck-
lings is said to be highly fatal and appears often suddenly. In addition to the Simulium mentioned above, *Simulium parnassum* appears to be a vector (Fallis, Davies and Vickers, 1951).

Mathis and Léger: Macrogametocytes, oval; 14–15μ by 4.5–5.5μ; several vacuoles in darkly stained cytoplasm. Microgametocytes, oval; slightly smaller; cytoplasm stains less deeply. Infected host cells about 48μ long; nucleus elongate.

Huff found that (1) young schizonts are in macrophages of, and also extracellularly in, the spleen and liver; (2) two types of schizonts occur: one, “hepatic schizonts” in hepatic cells which cause no distortion or alteration of the host cell, and the other, “megaloschizonts” in the blood vessels of, or extravascularly in, the heart, spleen, liver and intestine; (3) megaloschizonts become divided into many cytomeres which give rise to numerous merozoites; (4) young gametocytes occur in lymphocytes, monocytes, myelocytes and late polychromatophilic erythroblasts; (5) the cells in which fully grown gametocytes occur, appear to be macrophages. Life history and effect on the blood of host birds (Fallis, Davies and Vickers, 1951); development in ducklings (Chernin, 1952).

Other reported species: *L. smithi* Laveran and Lucet (1905) in turkey; *L. bonasae* Clarke (1935) in ruffed grouse; *L. andrewsi* Atchley (1951) in chicken, etc.

**Family 3 Babesiidae Poche**

Minute non-pigmented parasites of the erythrocytes of various mammals; transmission by ticks.

Genus *Babesia* Starcovici (*Piroplasma* Patton). In erythrocytes of cattle; pear-shaped, arranged in couples; sexual reproduction in female ticks in which developing ova, hence young ticks, become infected with ookinete, producing sporozoites which enter salivary glands (Dennis). Taxonomy (Toit, 1918).

*B. bigemina* (Smith and Kilborne) (Figs. 268; 269, a–d). The causative organism of the haemoglobinuric fever, Texas fever or red-water fever of cattle; the very first demonstration that an arthropod plays an important rôle in the transmission of a protozoan parasite; the infected cattle contain in their erythrocytes oval or pyriform bodies with a compact nucleus and vacuolated cytoplasm; the division is peculiar in that it appears as a budding process at the beginning. We owe Dennis (1932) for our knowledge of the development of the organism.

Sexual reproduction followed by sporozoite formation occurs in the tick, *Boophilus* (*Margaropus*) *annulatus*; when a tick takes in
infected blood into gut lumen, isogametes, 5.5–6μ long, are produced; isogamy results in motile club-shaped ookinete, 7–12μ long, which pass through gut wall and invade larger ova (1–2, in one case about 50, ookinete per egg); each ookinete rounds itself up into a sporont 7.5–12μ in diameter, which grows in size and whose nucleus divides repeatedly; thus are produced multinucleated (4–32 nuclei) amoeboi'd sporokinetes, up to 15μ long, which now migrate throughout

Fig. 268. The life-cycle of Babesia bigemina (Dennis). a–f, division in erythrocytes of cattle; g, h, gametocytes; i, isogametes; j, fertilization; k, zygote; l, ookinete penetrating through the gut wall; m, ookinete in host egg; n–p, sporoblast-formation; q, sporokinetes in a large embryonic cell; r, sporozoites in salivary gland.
embryonic tissue cells of tick, many of which cells develop into salivary gland cells; sporokinetes develop into sporozoites before or after hatching of host tick; sporozoites bring about an infection to cattle when they are inoculated by tick at the time of feeding. Texas fever once caused a considerable amount of damage to the cattle industry in the southern United States to which region the distribution of the tick is limited. Rees (1934) maintains that there is in addition a somewhat smaller species, *B. argentina* Lignières.

Fig. 269. a–d, *Babesia bigemina*, ×3000 (Nuttall); e–h, *B. bovis*, ×3000 (Nuttall); i–l, *Theileria parva*, ×3000 (Nuttall); m–s, *Dactylosoma ranarum* (m–q, schizogony; r, s, gametocytes), ×2700 (Noller).

*B. bovis* Starcovici (Fig. 269, e–h). In European cattle; amoeboïd form usually rounded, though sometimes stretched; 1–1.5μ in diameter; paired pyriform bodies make a larger angle, 1.5–2μ long; transmitted by *Ixodes ricinus*.

*B. canis* (Piana and Galli-Valerio). Pyriform bodies 4.5–5μ long; the organism causes malignant jaundice in dogs; widely distributed; transmitted by the ticks: *Haemaphysalis leachi*, *Rhipicephalus sanguineus*, and *Dermacentor reticulatus* (Regendanz and Reichenow, 1933).
Species of Babesia occur also in sheep, goats, pigs and horses.

Genus Theileria Bettencourt, França and Borges. Schizogony takes place in endothelial cells of capillaries of viscera of mammals; certain forms thus produced enter erythrocytes and appear in the peripheral circulation.

*T. parva* (Theiler) (Fig. 269, i–l). In the cattle in Africa, cause of African coast fever; intracorpusecular forms 1–2μ in diameter; transmitted by the tick, *Rhipicephalus evertsi* and *R. appendiculatus* (Reichenow, 1937).

Genus Dactylosoma Labbé. In blood of reptiles and amphibians; schizogony and gametocytes in erythrocytes; invertebrate hosts unknown.

*D. ranarum* (Kruse) (Fig. 269, m–s). In European frogs; schizonts 4–9μ in diameter; 4–16 merozoites, 2–3μ by 1–1.5μ; gametocytes 5–8μ by 1.5–3μ.

Genus Toxoplasma Nicolle and Manceaux. Minute intracellular parasites in leucocytes and endothelial cells of various mammals, birds and reptiles; round or ovoid; usually not common in peripheral blood, though infective through inoculation; ordinarily abundant in the liver, spleen, bone marrow, lung, brain, etc.; multiplication by binary fission (Nicolle and Manceaux, 1909). Several species were designated by observers on the basis of the difference in host species. Taxonomy (Chatton and Blanc, 1917); morphology (Arantes 1914); relation to Plasmodium (Hegner and Wolfson, 1938; Manwell, 1939, 1941).

Fig. 270. *Toxoplasma gondii*. × about 1750. (Chatton and Blanc) a, isolated organisms; b, 2 trophozoites; c, organisms undergoing binary fission; d, a host cell with many organisms which developed by repeated binary fission.

*T. gondii* N. and M. (Fig. 270). In *Ctenodactylus gundi*, a rodent in North Africa; a variety of experimental animals susceptible to it; crescentic; 4–6μ by 2–3μ; division occurs intra- or extra-cellularly.
Since the subcutaneous tissues of experimentally infected animals such as rats, pigeons and chicks do not harbor parasites in the absence of parasitemia and the organisms are apparently present in the blood, transmission may be carried on by blood-sucking arthropods (Jacobs and Jones, 1950).

Toxoplasma appears to be common in birds. For example, in a survey on the blood parasites of birds on Cape Cod, Herman (1938) found the organism in 11 species of birds examined by him. In the past ten years a considerable amount of information has accumulated on the organisms which attack and produce a disease (toxoplasmosis) in man. References (Sabin, 1942; Schwarz, Rose and Fry, 1948; Mantz, Sailey and Grocott, 1949; Hogan, 1951; Weinman, 1952).

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HAEMOSPORIDIA


Saurian malarial parasites of the United States and Mexico. Ibid., 74:68.


THE sporozoa which are grouped here are mostly incompletely known, although some of them are widely distributed. They produce spores which are simple in structure, being composed of a spore membrane and a sporoplasm.

Order 1 Haplosporidia Caullery and Mesnil

This order includes those sporozoans which produce simple spores. In some species the spores may resemble superficially those of Microsporidia, but do not possess any polar filament. In this regard, Haplosporidia may be considered a more primitive group than Cnidosporidia (p. 643).

The Haplosporidia are cytozoic, histozoic, or coelozoic parasites of invertebrates and lower vertebrates. The spore is spherical or ellipsoidal in form and covered by a resistant membrane which may possess ridges or may be prolonged into a more or less long tail-like projection. In a few species the spore membrane possesses a lid which, when opened, will enable the sporoplasm to emerge as an amoebula. The sporoplasm is uninucleate and fills the intrasporal cavity.

The development of a haplosporidian, Ichthyosporidium giganteum, as worked out by Swarczewsky, is as follows (Fig. 271): The spores germinate in the alimentary canal of the host fish and the emerged amoebulae make their way to the connective tissue of various organs (a). These amoebulae grow and their nuclei multiply in number, thus forming plasmodia. The plasmodia divide into smaller bodies, while the nuclei continue to divide (b–e). Presently the nuclei become paired (f, g) and the nuclear membranes disappear (h). The plasmodia now break up into numerous small bodies, each of which contains one set of the paired nuclei (i, j). This is the sporont (j) which develops into 2 spores by further differentiation (k–o).

Genus Haplosporidium Caullery and Mesnil. After growing into a large form, plasmodium divides into uninucleate bodies, each of which develops into a spore; spore truncate with a lid at one end; envelope sometimes prolonged into processes; in aquatic annelids and mollusces.

H. chitonis (Lankester) (Fig. 272, a, b). In liver and connective tissue of Craspidochilus cinereus; spores oval, 10μ by 6μ; envelope with 2 prolonged projections.
H. limnodrili Granata (Fig. 272, c). In gut epithelium of Limnodrilus udekemianus; spores 10–12μ by 8–10μ.

H. nemertis Debaïsieux (Fig. 272, d). In connective tissue of Lineus bilineatus; spores oval with a flat operculum, but without any projections of envelope, 7μ by 4μ.

H. heterocirri C. and M. (Fig. 272, e). In gut epithelium of Het-

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Fig. 271. The development of Ichthyosporidium giganteum (Swarzewsky). a–e, schizogony; f–n, sporogony; o, stained spore, X about 1280.

erocirrus viridis; mature organisms 50–60μ by 30–40μ; spores 6.5μ by 4μ.

H. scolopli C. and M. (Fig. 272, f). In Scoloplos mülleri; fully grown form 100–150μ by 20–30μ; spores 10μ by 6.5μ.

H. vejdovskii C. and M. (Fig. 272, g). In a freshwater oligochaete, Mesenchytraeus flavus; spores 10–12μ long.

Genus Urosporidium Caullery and Mesnil. Similar to Haplo-

sporidium, but spherical spore with a long projection.
**ACNIDOSPORIDIA, HAPLOSPORIDIA**

*U. fuliginosum* C. and M. (Fig. 272, h, i). In the coelom of the polychaete, *Syllis gracilis*; rare.

Genus **Anurosporidium** Caullery and Chappellier. Similar to *Haplosporidium*, but operculate spore spherical.

*A. pelseneeri* C. and C. In sporocyst of a trematode parasitic in *Donax trunculus*; schizogony intracellular; cysts extracellular, with up to 200 spores; spores about 5μ long.

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**Fig. 272.** a, b, *Haplosporidium chitonis*, ×1000 (Pixell-Goodrich); e, *H. limnodrilli*, ×1000 (Granata); d, *H. nemertis*, ×1000 (Debaisieux); e, *H. heterocirri*, ×1000 (Caullery and Mesnil); f, *H. scolopi*, ×1000 (Caullery and Mesnil); g, *H. vejovskii*, ×1000 (Caullery and Mesnil); h, i, *Urosporidium fuliginosum*, ×1000 (Caullery and Mesnil); j, k, *Bertramia asperospora* (j, cyst with spores; k, empty cyst), ×1040 (Minchin); l, m, *Coelosporidium periplanetae* (l, trophozoite with spores and chromatoid bodies), ×2540 (Sprague).

Genus **Bertramia** Caullery and Mesnil. Parasitic in aquatic worms and rotifers; sausage-shaped bodies in coelom of host; spherical spores which develop in them, possess a uninucleate sporoplasm and a well-developed membrane.

*B. asperospora* (Fritsch) (Fig. 272, j, k). In body cavity of rotifers: Brachionus, Asplanchna, Synchaeta, Hydatina, etc.; fully grown vermicular body 70–90μ with 80–150 spores.
B. capitellae C. and M. In the annelid Capitella capitata; spores 2.5μ in diameter.

B. euchlanis Konsuloff. In coelom of rotifers belonging to the genus Euchlanis.

Genus Ichthyosporidium Caullery and Mesnil. In fish; often looked upon as Microsporidia, as the organism develops into large bodies in body muscles, connective tissue, or gills, which appear as conspicuous "cysts," that are surrounded by a thick wall and contain numerous spores.

I. giganteum (Thélohan) (Fig. 271). In various organs of Crenilabrus melops and C. ocellatus; cysts 30μ–2 mm. in diameter; spores 5–8μ long.

I. hertwigi Swarczewsky. In Crenilabrus paro; cysts 3–4 mm. in diameter in gills; spores 6μ long.

Genus Coelosporidium Mesnil and Marchoux. In coelom of Cladocera or Malpighian tubules of cockroach; body small, forming cysts; spores resemble microsporidian spores; but without a polar filament.

C. periplanetae (Lutz and Splendore) (C. blattellae Crawley) (Fig. 272, l, m). In lumen of Malpighian tubules of cockroaches; common; spores 5.5–7.5μ by 3–4μ. Cytology (Sprague, 1940).

Order 2 Sarcosporidia Balbiani

These organisms are muscle parasites of mammals, birds and reptiles. The infected host muscles are characterized by the presence of opaque white bodies (Miescher’s tubes) (Fig. 273) which vary from microscopic to several centimeters in length, and are cylindrical, ellipsoid or ovoid, with a somewhat lobulated surface. When mature, the parasite becomes filled with the “spores” or Rainey’s corpuscles which are crescentic or banana-shaped. They contain a nucleus and many granules, surrounded by a very delicate membrane (Fig. 274).

The morphological peculiarity and lack of information concerning their transmission and development have characterized these organ-

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Fig. 273. a, Sarcocystis tenella in the oesophagus of sheep; b, S. miescheriana in the muscle of pig; ×1 (Schneidemühl from Doflein).
isms for many years. Spindler and Zimmerman (1945) placed aseptically ruptured cysts of *Sarcocystis miescheriana* (p. 640) of pigs in sterile dextrose solution and kept the preparations at 37°C. for 24 hours and then at room temperature. In from a few days to two weeks, the "spores" budded off minute, coccoid bodies which developed into septate mycelia with vertical hyphae bearing spores, a typical feature of the development of a fungus belonging to Aspergillus. When the conidia from the cultures were injected into or fed to 50 young pigs, 25 showed at necropsy four to six months after the injection or ingestion of the conidia, typical Sarcocystis cysts in the muscles, while the controls remained free from infection. Cultures made from the mature cysts in these pigs, developed a fungus like that which had been injected. Pigs, rats and mice which fed on the cysts, passed faeces and urine containing yeast-like bodies which developed in cultures into a fungus like that which was originally cultured. Spindler's (1947) further study revealed that in the sarcosporidian cysts of sheep and duck, the strands present within the cysts were none other than the connective tissues of the host and the compartments contained a network of jointed hypha-like structures, and the spores appeared to be exogenous growths on the jointed hypha-like structures; and each spore was capable of budding out another spore from its free end. Spindler concludes from these observations that Sarcocystis of pigs, sheep and ducks are fungi, related to Aspergillus. This view will explain reasonably well the difficulties encountered in relation to Sarcosporidia; namely, the unknown life cycle, lack of a protective membrane of the "spore," the absence of a vector, and the common occurrence among herbivorous animals.

Genus *Sarcocystis* Lankester. In the muscles of higher vertebrates. Many species have been reported by various workers from mammals, birds and reptiles on the basis of difference in host species. Species (Babudieri, 1932).

*S. lindemanni* (Rivolta). A few cases of Sarcocystis infection have been reported from man in muscle cells of larynx (Baraban and St. Remy), of biceps and tongue (Darling), of heart (Manifold), of breast (Vasudevan), etc. There seem to be dimensional discrepancies of organisms observed by different investigators. The dimensions of parasitic masses and of spores are as follows: Parasites 1.6 mm. by 170µ and banana-shaped spores 8–9µ long (Baraban and St. Remy); parasites 84µ by 27µ and spores 4.25µ by 1.75µ (Darling); parasites spherical, 500µ in diameter and spores over 10µ long (Manifold); parasites 5.3 cm. by 320µ and spores 8.33µ by 1.6µ (Vasudevan). The parasitic masses are oval to spindle in form and
imbedded in the muscle cells which are distended, and may appear white-streaked to naked eye. Seen in sections, the body is divided into compartments. Gilmore, Kean and Posey (1942) have recently found three bodies in sectioned heart muscles of an eleven year old child who died from an unknown cause, and considered them as sarcosporidian bodies. They measured 25µ by 19µ, 57µ by 30µ, and 41µ by 25µ in cross sections; there were no septa within the bodies; minute bodies present in the masses were mostly rounded and about 1µ in diameter, though a few were crescentic. The questions such as what species infect man, how man becomes infected, etc., are unanswered at present.

![Diagram of muscle layer, connective tissue layer, fibrous zone, external, median, internal cyst membrane, spores, and sporocests.](image)

Fig. 274. Portion of a cyst of *Sarcocystis tenella* in sheep, × about 1000 (Alexeieff).

*S. tenella* Railleit (Figs. 273, a; 274). In the muscles of tongue, pharynx, oesophagus, larynx, neck, heart, etc., of sheep; large parasites 40µ–2 cm. long with a thin membrane; spores sickleform (Alexeieff, 1913; Scott, 1943).

*S. miescheriana* (Kühn) (Fig. 273, b). In the muscles of pig; cysts up to 3–4 mm. by 3 mm.; envelope striated; "spores" reniform. Musfeldt (1950) found 15 of 264 pig diaphragms examined were infected by a Sarcocystis. The pigs were all garbage-fed animals.
Sarcocystis infections were also noticed in the rats from one of the piggeries from which infected pigs were obtained. Fungus nature of the organism (p. 639); effect on host (Spindler, Zimmerman and Jaquette, 1946).

S. bertrami Doflein. In the muscles of horse; similar to S. miescheriana; parasitic mass up to 9–10 mm.; envelope striated.

S. muris Blanchard. In body muscles of rats and mice; parasitic masses up to 3 cm. long; spores 13–15μ by 2.5–3μ; transmissible to guinea pig (Negri) which shows experimental infection in muscles in 50–100 days after feeding on infected muscles.

S. rileyi Stiles. In muscles of various species of ducks; parasites in muscle, opaque white in color and measure up to 5 mm. by 2 mm.; spores are sausage-shaped and 8–10μ by about 3μ.

References


THE members of this subclass possess without exception resistant spores which are of unique structure. Each spore possesses 1-4 polar filaments and one to many sporoplasms. The membrane which envelops these structures may be a single-piece or bi- or trivalved. The polar filament is typically coiled within a polar capsule.

In the order Myxosporidia and Actinomyxidia, there appear several cells during the process of sporulation. These cells give rise to one to many sporoplasms or generative cells, capsulogenous cells, and spore membrane. This condition is not observed in other groups of Protozoa and for this reason some writers recognize a close affinity between these two orders and the Mesozoa. The method of multiplication in the Cnidosporidia is schizogonic and sporogonic. The division is repeated binary or multiple fission, budding, or plasmotomy. The nuclear division varies from amitosis to mitosis. Isogamous, anisogamous, and autogamous reproduction have been reported in a number of species. In many forms, the zygote is the sporont, in which one to many spores become differentiated.

No secondary or intermediate host has been found for any of the Cnidosporidia. They are exclusively parasites of the lower vertebrates and invertebrates. Since cnidosporidian infections occur frequently in epidemic forms among such economically important animals as the silkworm, honey bees, and commercial fishes, these organisms possess considerable practical significance. History and economic importance (Auerbach, 1910; Kudo, 1920, 1924).

The Cnidosporidia are divided into the following four orders:

Spores comparatively large
- Shell bivalve; 1 to 4 polar capsules.................Order 1 Myxosporidia
- Shell trivalve; 3 polar capsules........Order 2 Actinomyxidia (p. 660)

Spores comparatively small
- Shell one-piece; 1 (or 2) polar filament..Order 3 Microsporidia (p. 668)
- Barrel-shaped; a thick filament coiled beneath the shell; 3 sporoplasms

Order 4 Helicosporidia (p. 678)

**Order 1 Myxosporidia Bütschli**

The spore of a myxosporidian is of various shapes and dimensions. It is covered by a bivalve chitinous spore membrane (Kudo, 1921), the two valves meeting in a sutural plane which is either twisted (in three genera) or more or less straight. The membrane may possess various markings or processes. The polar capsule, with
its short coiled filament, varies in number from one to four (Fig. 275). Except in the family Myxidiidae, in which one polar capsule is situated near each of the poles of the spore, the polar capsules are always grouped at one end which is ordinarily designated as the anterior end of the spore. Below or between (in Myxidiidae) the polar capsules, there is almost always a sporoplasm. Ordinarily a

![Diagram of spore stages](image)

**Fig. 275.** Sporogony in *Myxosoma catostomi*, ×2130 (Kudo). a, sporont or pansporoblast; b–h, development of two sporoblasts within the sporont; i, a nearly mature spore; j–l, views of spore.

young spore possesses two sporoplasm nuclei which fuse into one (autogamy) when the spore becomes mature. In Myxobolidae there is a glycogenous substance in a vacuole which stains mahogany red with iodine and is known as the iodinophilous (iodophile) vacuole.

The Myxosporidia are exclusively parasites of lower vertebrates, especially fishes. Both fresh and salt water fishes have been found to harbor, or to be infected by, Myxosporidia in various regions of the world. A few occur in Amphibia and Reptilia, but no species has been found to occur in either birds or mammals. When a spore gains entrance into the digestive tract of a specific host fish, the sporoplasm leaves the spore as an amoebula which penetrates through the gut-epithelium and, after a period of migration, enters the tissues of certain organs, where it grows into a trophozoite at the
expense of the host tissue cells, and the nucleus divides repeatedly. Some nuclei become surrounded by masses of dense cytoplasm and become the sporonts (Fig. 275). The sporonts grow and their nuclei divide several times, forming 6–18 daughter nuclei, each with a small mass of cytoplasm. The number of the nuclei thus produced depends upon the structure of the mature spore, and also upon whether 1 or 2 spores develop in a sporont. When the sporont develops into a single spore, it is called a monosporoblastic sporont, and if two spores are formed within a sporont, which is usually the case, the sporont is called disporoblastic, or pansporoblast. The spore-formation begins usually in the central area of the large trophozoite, which continues to grow. The surrounding host tissue becomes degenerated or modified and forms an envelope that is often large enough to be visible to the naked eye (Figs. 278, 280). This is ordinarily referred to as a myxosporidian cyst. If the site of infection is near the body surface, the large cyst breaks and the mature spores become set free in the water. In case the infection is confined to internal organs, the spores will not be set free while the host fish lives. Upon its death and disintegration of the body, however, the liberated spores become the source of new infection.

The more primitive Myxosporidia are coelozoic in the host's organs, such as the gall bladder, uriniferous tubules of the kidney, urinary bladder, etc. In these forms, the liberated amoebulae make their way into the specific organ and there grow into multinucleate amoeboid trophozoites which are capable of forming pseudopodia of various types. They multiply by exogenous or endogenous budding or plasmotomy. One to several spores are developed in the trophozoite.

Almost all observers agree in maintaining the view that the 2 nuclei of the sporoplasm or 2 uninucleate sporplasms fuse into one (autogamy or paedogamy), but as to the nuclear as well as cytoplasmic changes prior to, and during, spore-formation, there is a diversity of opinions. For illustration, the development of Sphaeromyxa sabrazesi (p. 656) as studied by two investigators may be taken as an example. Debaisieux's (1924) observation is in brief as follows (Fig. 276): Sporoplasms after finding their way into the gall bladder of host fish develop into large trophozoites containing many nuclei ($a, b$) 2 vegetative nuclei become surrounded by a cytoplasmic mass ($c$) and this develops into a primary propagative cell ($d$) which divides (3 chromosomes are noted) ($e$) and forms secondary propagative cells ($f$). A binucleate sporocyte is formed from the latter by unequal nuclear division ($g–i$) and 2 sporocytes unite to form a tetranucleate
pansporoblast (j) which develops into 2 spores (k, l). Sporoplasm shows first 2 nuclei (l), but later 4 (m), of which 2 degenerate (n) and the other 2 fuse into one nucleus (o). On the other hand, according to Naville (1930) a uninucleate amoebula (Fig. 277, a) enters the gall bladder and develops into multinucleate trophozoite in which nuclear

Fig. 276. The development of Sphaeromyxa sabrazesi (Debaisieux). a, vegetative nuclei; b, association of two vegetative nuclei; c, the same within a cell; d, primary propagative cell; e, its division; f, secondary propagative cells; g, their division; h, formation of sporocyte; i, two sporocytes; j, formation of pansporoblast; k, pansporoblast at later stages; l, pansporoblast with two spores, the sporoplasm of which contains two nuclei; m, four nuclei in sporoplasm; n, two nuclei remain functional, the other two degenerate, o, fusion of the two nuclei.
Fig. 277. The development of *Sphaeromyxa sabrazesi* (Naville). a, uninucleate amoebula enters the gall bladder; b, young multinucleate trophozoite; c, development of macrogametes; d, development of microgametes; e, f, plasmogamy; g–m, development of pansporoblast; n, fusion of the two nuclei in the sporoplasm.
division reveals 4 chromosomes (b); within the trophozoite macro-
gametes and microgametes are independently formed, during which
process, chromosome number is reduced into half (2) (c, d); plasog-
amy between a macrogamete and a microgamete results in produc-
tion of a binucleate pansporoblast (e, f), from which repeated nuclear
division (g–l) forms 2 spores (m); each of the 2 nuclei of the sporo-
plasm is haploid and the diploid number is restored when the 2 nuclei
fuse into one (n).

The site of infection by Myxosporidia varies among different
species. They have been found in almost all kinds of tissues and
organs of host fish, although each myxosporidian has its special site
of infection in one to several species of fish. The gills and gall bladder
are most frequently parasitized by Myxosporidia in freshwater
fishes, while the gall bladder and urinary bladder of marine fishes
harbor one or more species of Myxosporidia. When the infection is
concentrated in the fins or integument, the resulting changes are
quite conspicuous (Fig. 278). The infection in the gills is usually

![Fig. 278. A channel cat, heavily infected with *Henneguya exilis*,
×½ (Kudo).](image)

manifest by whitish pustules which can be frequently detected with
the unaided eye. When the wall of the alimentary canal, mesentery,
liver, and other organs are attacked, one sees considerable changes
in them. Heavy myxosporidian infection of the gall bladder or uri-
nary bladder of the host fish may cause abnormal appearance and
coloration or unusual enlargement of the organ, but under ordinary
circumstances the infection is detected only by a microscopical ex-
amination of its contents. Certain histological changes in the host
fish have been mentioned elsewhere (p. 31).

Severe epidemic diseases of fishes are frequently found to be due to
myxosporidian infections. According to Davis (1924), the “wormy”
halibut of the Pacific coast of North America is due to the myxo-
sporidian, *Unicapsula muscalaris* (Fig. 280), which invades the mus-
cular tissue of the host fish. The “boil disease” of the barbel, *Barbus
barbus* and others, of European waters, is caused by *Myxobolus
pfeifferi* (Keysselitz, 1908). *Myxosoma cerebralis*, which attacks the
supporting tissues of salmonid fish, is known to be responsible for the so-called "twist disease" (Plehn, 1904), which is often fatal especially to young fishes and occurs in an epidemic form. *Henneguya salminicola* invades the body muscles of various species of Pacific salmon and produces opaque white cysts, 3–6 mm in diameter; it is thus responsible for the so-called "tapioca disease" of salmon (Fish, 1939). *Kudoa thyrsites* (p. 655) attacks the body muscle fibers of the barracouta in which the infected muscles become liquefied. This condition is known as "milky barracouta" or "pap snoek" and may affect as much as 5 per cent of the commercial catches (Willis, 1949).

The Myxosporidia are divided into three suborders:

Largest diameter of spore at right angles to sutural plane; with 1 polar capsule on each side; sporoplasm without iodinophilous vacuole.

Spores laterally expanded; coelozoic in marine fish, except one species.

Spores less laterally expanded; in freshwater fish; histozoic or coelozoic.

**Suborder 1 Eurysporea Kudo**

Family 1 Ceratomyxidae

Spores are laterally prolonged and therefore sutural diameter is smaller than width; 2 polar capsules at anterior margin; one on each side of sutural plane; in one genus the spores contain three polar capsules and the spore membrane is composed of three shell-valves.

Genus **Ceratomyxa** Thélohan. Shell-valves conical and hollow,
attached on bases; sporoplasm usually not filling intrasporal cavity; Numerous species in the gall-bladder of marine fish, except *C. shasta* (Noble, 1950) which was found “widely distributed in viscera” of fingerling rainbow trout (*Salmo gairdneri*).

![Fig. 279. a, Ceratomyxa mesospora, ×1000 (Davis); b, c, C. hopkinsi, ×1000 (Jameson); d–j, Leptotheca ohlmacheri (d, section of a uriniferous tubule of *Rana pipiens*, with trophozoites and spores, ×800; e, a trophozoite with a bud; f–h, disporous trophozoites; i, a spore with extruded polar filaments; j, surface view of spore, ×1500) (Kudo).](image)
C. mesospora Davis (Fig. 279, a). In the gall-bladder of Cestracion zygaena; spores 8 μ in sutural diameter and 50–65 μ wide.

C. hopkinsi Jameson (Fig. 279, b, c). In the gall-bladder of Parophrys vetulus, Microstomus pacificus and Citharichthys xanthostigmus; trophozoites disporous; spores 5.7–7.5 μ in sutural diameter and 28.8–39 μ wide.

Genus Leptotheca Thelohan. Shell-valves hemispherical; in gall-bladder or urinary bladder of marine fish and one in amphibians. Numerous species.

L. ohlmacheri (Gurley) (Fig. 279, d–j). In the uriniferous tubules of kidney of frogs and toads; spores 9.5–12 μ in sutural diameter and 13–14.5 μ wide; with 2 uninucleate sporoplasms (Kudo 1922).

Genus Myxoproteus Doflein. Spores pyramidal with or without distinct processes at base of pyramid; in urinary bladder of marine fish. 3 species.

M. cordiformis Davis (Fig. 280, a). In the urinary bladder of Chaeotodipterus faber; spores 12 μ by 10–11 μ.

Genus Trilospora Noble. Spores triangular with concave sides in anterior end-view; profile ellipsoid; three polar causles and three shell-valves; in the gall-bladder of marine fish. One species.

T. californica N. Spores 7.2 μ in sutural diameter by 16 μ wide; polar capsules 3 μ by 1.5 μ, often four instead of three in number; in the gall-bladder of Typhlogobius californiensis and Gibbonsia elegans elegans (Noble, 1939).

Family 2 Wardiidae Kudo

Genus Wardia Kudo. Spores isoseceles triangle with 2 convex sides; oval in profile; 2 large polar capsules; tissue parasites of freshwater fish. 2 species.

W. ovinocua K. (Fig. 280, b). In the ovary of Lepomis humilis; spores 9–11 μ in sutural diameter and 10–12 μ wide.

Genus Mitraspora Fujita. Spores circular or ovoidal in front view; somewhat flattened in profile; 2 polar capsules; shell striated; with or without posterior filaments; in kidneys of freshwater fishes. This genus apparently includes border-line forms between this and other suborders. 3 species.

M. elongata Kudo. In the kidney of Apomotis cyanellus; spores 15–17 μ by 5–6 μ.

Suborder 2 Sphaerosporea Kudo

Spore with 1 polar capsule .......... Family 1 Unicapsulidae (p. 652)
Spore with 2 polar capsules .......... Family 2 Sphaerosporidae (p. 653)
Spore with 4 polar capsules .......... Family 3 Chloromyxidae (p. 654)
Family 1 Unicapsulidae Kudo

Genus Unicapsula Davis. Spherical spore with 1 polar capsule; shell-valves asymmetrical; sutural line sinuous; histozoic in marine fish. One species.

Fig. 280. a, Myxoproteus cordiformis, ×1000 (Davis); b, Wardia ovinocua, ×1330 (Kudo); c, Sphaerospora polymorpha, ×1000 (Davis); d–i, S. tincae (d, external appearance of a heavily infected young tench; e, internal appearance, ×3; f, mature pansporoblast; g, h, two spores; i, germination of spore, ×1000) (Léger); j, k, Sinuolinea dimorpha (j, trophozoite with three gemmules, ×420; k, a spore, ×930) (Davis); l, m, Chloromyxum leydigi (l, ×500; m, ×1000) (Thélohan); n, C. trijugum, ×1130 (Kudo).

U. muscularis D. (Fig. 281). Spore about 6μ in diameter; 2 uninucleate sporoplasm; in muscle fibers of halibut; Pacific coast of North America; the cause of the “wormy” halibut (Davis, 1924).
Family 2 Sphaerosporidae Davis

Genus Sphaerospora Thélohan. Spore spherical or subspherical; sutural line straight; 2 polar capsules at anterior end; coelozoic or histozoic in marine or freshwater fishes.

S. polymorpha Davis (Figs. 280, c; 282, a–e). In the urinary bladder of toadfish, Opsanus tau and O. beta. Trophozoites amoeboid with conical pseudopodia; up to 100μ long, the majority being 20–50μ long; plasmodotory; disporoblastic; disporous or polysporous. Spores spheroidal; shell-valves finely striated; polar capsules divergent; fresh spores measure 7.5–9.5μ by 7–8μ. The trophozoites suffer frequently infection by Nosema notabilis (p. 672). Development and hyperparasitism (Kudo, 1944).
S. tincae Plehn (S. pernicialis Léger) (Fig. 280, d–i). In the kidney and other viscera of Tinca tinca in France and Germany; cause of epidemic disease among young tench; disease is manifest by great distension of anterior portion of abdomen and up-turned mouth: in-

![Diagram](image)

Fig. 282. a–e, *Sphaerospora polymorpha* (Kudo) (a, a trophozoite in life, ×1530; b, stage in simple plasmotomy, ×700; c, d, front and anterior end views of fresh spores; e, a spore with the extruded polar filaments, ×1415); f–h, *Myxidium serotinum* (Kudo) (f, a stained young trophozoite, ×1530; g, h, two views of fresh spores, showing the ridges on the membrane, ×915); i–l, *Kudoa clupeidae* (Meglitsch) (i, j, two views of unstained spores, ×1240; k, l, stained spores, ×1430); m–p, *K. thyrsites* (Willis) (m–o, preserved spores; p, a spore from section).

Infection fatal through rupture of abdominal wall; spores 7–8.75μ in diameter (Léger, 1929).

Genus *Sinuolinea* Davis. Spherical or subspherical spores; sutural line sinuous; with or without lateral processes; 2 spherical polar capsules; in urinary bladder of marine fish.

*S. dimorpha* D. (Fig. 280, j, k). In Cynoscion regalis; spores 15μ in diameter (Davis, 1917).

Family 3 Chloromyxidae Thélonean

Genus *Chloromyxum* Mingazzini. Spore with 4 polar capsules, grouped at anterior end; shell surface often striated or ridged;
histozoic or coelozoic in freshwater or marine fish and also in amphibians. Numerous species.

*C. leydigi* M. (Figs. 70, c, d; 280, l, m). In the gall-bladder of various species of *Raja, Torpedo* and *Cestracion*; spores 6–9µ by 5–6µ; widely distributed. Structure and development (Erdmann, 1917; Naville, 1927).

*C. trijugum* Kudo (Fig. 280, n). In the gall-bladder of *Xenotis megalotis* and *Pomoxis sparoides*; spores 8–10µ by 5–7µ.

Genus *Kudoa* Meglitsch. Resembles *Chloromyxum*; but spores stellate or quadrate in anterior end-view; spore membrane delicate and the sutures indistinct; four shell-valves (?); histozoic (Meglitsch, 1947a). Several species.

*K. clupeidae* (Hahn) (Fig. 282, i–l). In the body muscles of *Clupea harengus, Brevoortia tyrannus*, etc.; spores 5.1µ by 6.4µ; polar capsules 1.5µ by 1µ (Meglitsch, 1947). Nigrelli (1946) found this species in the ocean pout (*Macrozoares americanus*).

*K. thyrsites* (Gilchrist) (Fig. 282, m–p). In the body muscles of the barracouta, *Thyrsites atun*, in Australia and Africa; pyramidal spores 6–7µ high and 12–17µ wide; two uninucleate sporoplasms; polar capsules homogeneous in appearance (Willis, 1949). Effect on host (p. 649).

Suborder 3 **Platysporea** Kudo

Without iodonphilous vacuole

2 polar capsules, one at each pole. ............ Family 1 Myxidiidae

1 polar capsule. ......................... Family 2 Coccomyxidae (p. 658)

2 or 4 polar capsules grouped. ............ Family 3 Myxosomatidae (p. 658)

With an iodonphilous vacuole. ............ Family 4 Myxobolidae (p. 658)

**Family 1 Myxidiidae** Thélohan

Genus *Myxidium* Bütschli (*Cystodiscus Lutz*). Spores fusiform with pointed or rounded ends; polar filament comparatively long, fine; coelozoic or histozoic in fishes, also in amphibians and reptiles. Numerous species.

*M. lieberkühni* Bütschli (Figs. 70, a, b; 284, a–d). In urinary bladder of *Esox* spp.; spores 18–20µ by 5–6µ; widely distributed. Development (Cohn, 1896; Debaisieux, 1916); division (Kudo, 1921a; Bremer, 1922).

*M. immersum* (Lutz) (*Cystodiscus immersus* Lutz; *M. lindoyense* Carini). (Fig. 284, e, f). In the gall bladder of species of *Bufo, Leptodactylus, Atelopus*, etc.; in Brazil and Uruguay. Trophozoites circular to oval, and very thin; up to 4 mm. in diameter; disporoblastic; polysporous. Spores 11.8–13.3µ by 7.5–8.6µ; shell-valves
marked with 1 longitudinal and 7–9 transverse ridges (Cordero, 1919; Kudo and Sprague, 1940).

*M. serotinum* Kudo and Sprague (Figs. 282, f–h; 283). In the gall bladder of *Bufo terrestris, Rana pipiens, R. clamitans* and *R. sphencephala*; in the United States. Trophozoites up to 6.5 by 1.8 mm., extremely thin; cytoplasm highly alveolated; endogenous budding; disporoblastic; polysporous. Spores 16–18μ by 9μ; shell-valve with 2–4 longitudinal and 10–13 transverse ridges (Kudo, 1943).

Fig. 283. Scattered spores, young and sporulating trophozoites of *Myxidium serotinum*, as seen in the bile of a frog in life, X64 (Kudo).

*M. kudoi* Meglitsch. In gall-bladder of *Ictalurus furcatus*; trophozoites large disc-like up to 1 mm. in diameter; spores 8.5–12μ long by 4–6μ (Meglitsch, 1937).

Genus *Sphaeromyxa* Thélohan. Spore fusiform, but ends usually truncate; polar filament short, thick; trophozoites large, discoid; coelozoic in marine fish. Several species.

*S. balbianii* T. (Figs. 70, e; 284, g–i). In gall-bladder of Motella and other marine fish in Europe and of *Siphostoma* in the United States; spores 15–20μ by 5–6μ (Naville, 1930).

*S. sabrazesi* Laveran and Mesnil (Figs. 276; 277; 284, j–l). In gall-
bladder of Hippocampus, Motella, etc.; spores 22–28 µ by 3–4 µ (Debaisieux, 1925; Naville, 1930).

Genus Zschokkella Auerbach. Spore semi-circular in front view; fusiform in profile; circular in cross-section; ends pointed obliquely;

polar capsules large, spherical; sutural line usually in S-form, coelozoic in fish or amphibians. A few species.

Z. hildae A. (Fig. 284, m, n). In urinary bladder of Gadus spp.; spores 16–29 µ by 13–18 µ (Auerbach, 1910).
Family 2 Coccomyxidae Léger and Hesse

Spore ellipsoidal; one polar capsule at one end; circular in cross-section; undoubtedly a border-line form between Myxosporidia and Microsporidia.

Genus Coccomyxa Léger and Hesse. Polar filament long, fine; coelozoic parasite in marine fish (Léger and Hesse, 1907).

*C. morovi* L. and H. (Fig. 284, o-t). In the gall-bladder of *Clupea pilchardus*; spores 14\(\mu\) by 5–6\(\mu\) (Georgévitch, 1926).

Family 3 Myxosomatidae Poche

Two or 4 polar capsules at anterior end; sporoplasm without any iodinophilous vacuoles.

Genus Myxosoma Thélohan (*Lentospora* Plehn). Spore circular, oval or ellipsoid in front view, lenticular in profile; 2 polar capsules at anterior end; histozoic in marine or fresh water fish. Several species.

*M. catostomi* Kudo (Figs. 58; 275). In the muscle and connective tissue of *Catostomus commersonii*; spores 13–15\(\mu\) by 10–11.5\(\mu\) (Kudo, 1926).

*M. cerebralis* (Hofer) (Fig. 285, a). In the cartilage and perichondrium of salmonid fish; young fish are especially affected by infection, the disease being known as the "twist-disease" (Drehkrankheit); spores 6–10\(\mu\) in diameter. (p. 648).

*M. funduli* Kudo. In the gills of *Fundulus*; spherical cysts up to 360\(\mu\) by 264\(\mu\); spores pyriform, 14\(\mu\) by 8\(\mu\) by 6\(\mu\); polar capsules 8\(\mu\) by 2\(\mu\) (Kudo, 1918). Other species (Bond, 1938–1939).

Genus Agarella Dunkerly. Spore elongate oval; 4 polar capsules at anterior end; shell prolonged posteriorly into long processes. One species.

*A. gracilis* D. (Fig. 285, b). In the testis of South American lungfish, *Lepidosiren paradoxa* (Dunkerly, 1915, 1925).

Family 4 Myxobolidae Thélohan

One, 2, or 4 polar capsules grouped at anterior end; sporoplasm with an iodinophilous vacuole.

Genus Myxobolus Büttschli. Spores ovoidal or ellipsoidal, flattened; 2 polar capsules at anterior end; sporoplasm with an iodinophilous vacuole; sometimes with a posterior prolongation of shell; exclusively histozoic in freshwater fish or amphibians. Numerous species.

*M. pfeifferi* Thélohan (Fig. 285, e, f). In the muscle and connective
tissue of body and various organs of Barbus barbus, B. fluviatilis, and B. plebejus; tumor up to a diameter of 7 cm; most of infected fish die from the effect (Keysselitz); spores 12–12.5μ by 10–10.5μ.

M. orbiculatus Kudo (Fig. 285, g–i). In muscle of Notropis gilberti; spores 9–10μ in diameter by 6.5–7μ thick.

M. conspicuus K. (Fig. 285, j, k). In corium of head of Moxostoma breviceps; tumors 1/2–4 mm.; spores 9–11.5μ by 6.5–8μ (Kudo, 1929).

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**Fig. 285.** a, Myxosoma cerebralis, showing two views of spore, ×800 (Plehn); b, a spore of Agarella gracilis, ×1660 (Dunkerly); c, d, front and side views of fresh spores of Thelohanellus notatus, ×1530 (Kudo); e, f, Myxobolus pfeifferi (Keysselitz) (e, Part of section of a cyst; f, a spore treated with iodine solution, ×1780); g–i, M. orbiculatus (Kudo) (g, infected host's muscle, ×600; h, a fresh spore; i, Lugol-treated spore, ×1000); j, k, views of fresh spores of M. conspicuus, ×1530 (Kudo); l–o, M. squamosus (l, a cyst under a scale, ×6.5; m–o, views of fresh spores, ×1530); p–r, spores of Henneguya exilis, ×1530; s–u, spores of Unicauda clavicauda, ×1530 (s, t, fresh spores; u, a stained spore without the process (Kudo)).
M. intestinalis K. (Fig. 1, a). In the intestinal wall of *Pomoxis sparoides*; (fixed unstained) spores, 12–13μ by 10–12.5μ; the histological changes brought about by this protozoan have been mentioned elsewhere (p. 27) (Kudo, 1929).

M. squamosus K. (Fig. 285, l-o). In connective tissue below scales of *Hybopsis kentuckiensis*; spore circular in front view, 8–9μ in diameter, 4.5–5μ thick.

Genus Thelohanellus Kudo. Pyriform spores, each with one polar capsule; sporoplasm with an iodinophilous vacuole; histozoic in freshwater fish. 11 species (Kudo, 1933).

*T. notatus* (Mavor) (Figs. 1, b; 285, c, d). In subdermal connective tissue of *Pimephales notatus, Cliola vigilax, Notropis cornutus, N. blennius*, and *Leuciscus rutilus*; tumor up to 7 mm. in diameter; spores 17–18μ by 7.5–10μ; host tissue surrounding the organism becomes so greatly changed that it appears as an epithelium (p. 31) (Debaisieux, 1925; Kudo, 1929, 1934).

Genus Henneguya Thelohan (Myxobilatus Davis). Spore circular or ovoidal in front view; flattened; 2 polar capsules at anterior end; each shell-valve prolonged posteriorly into a long process; sporoplasm with an iodinophilous vacuole; mostly histozoic in freshwater fish. Numerous species.

*H. exiles* Kudo (Figs. 278; 285, p-r). In gills and integument of *Ictalurus punctatus*; cysts up to 3 mm. in diameter, conspicuous; spores, total length 60–70μ, spore proper 18–20μ long by 4–5μ wide by 3–3.5μ thick (Kudo, 1929, 1934).

*H. mictospora* Kudo. In the urinary bladder of *Lepomis* spp. and *Micropterus salmoides*; spores 13.5–15μ long, 8–9μ wide, 6–7.5μ thick; caudal prolongation 30–40μ long.

Genus Unicauda Davis. The spore is similar to that of *Henneguya*, but the single caudal appendage is not an extension of the shell-valves. Several species (Davis, 1944).

*U. clavicauda* (Kudo) (Fig. 285, s-u). In the subdermal connective tissue of the minnow, *Notropis blennius*; oblong or ellipsoid cysts, 1–1.5 mm. in the longest diameter; spores 10.5–11.5μ by 8.5–9.5μ by 6μ; appendage 20–30μ by 3–6.5μ (Kudo, 1934).

Order 2 Actinomyxidia Stolic

The Cnidosporidia placed in this order have been less frequently studied and, therefore, not so well known as the Myxosporidia. The spore is enveloped by a membrane, or shell composed of 3 valves which are sometimes drawn out into simple or bifurcated processes. There are also 3 polar capsules in the spore and the polar filaments
are plainly visible in vivo. One to many sporoplasms occur in each spore. In the fully grown stage, the body is covered by a membrane and contains eight sporoblasts which develop in turn into eight spores. Whether the pansporoblast is formed by union of two cells or not, is unknown. The nuclei and cytoplasm divide and isogamy takes place. The zygote thus formed is the sporont in which a single spore is produced by repeated nuclear division combined with cytoplasmic differentiation.

The Actinomyxidia inhabit the body cavity or the gut-epithelium of fresh or salt water annelids. Taxonomy, morphology and development (Granata, 1925).

Spore with a double membrane; inner membrane a single piece, the outer trivalve; a single binucleate sporoplasm.................. Family 1 Tetractinomyxidae
Spore membrane a single trivalve shell; a single octonucleate sporoplasm or 8 uninucleate sporoplasms..........

Family 1 Tetractinomyxidae Poche

Genus Tetractinomyxon Ikeda. In the coelom of the sipunculid Petalostoma minutum; spores tetrahedron, without processes; trophozoite a rounded body, when mature; pansporoblast develops 8 spores. Seemingly borderline forms between the Myxosporidia and the Actinomyxidia.

T. intermedium I. (Fig. 286, a). Spherical pansporoblasts 20–25μ in diameter; spores 7–8μ in diameter; in coelom of the sipunculid, Petalostoma minutum (Ikeda, 1912).

Family 2 Triactinomyxidae

Genus Triactinomyxon Stolc. Each of 3 shell-valves drawn out into a long process, the whole anchor-like; spore with 8 or more uninucleate sporoplasms; in the gut-epithelium of oligochaetes.

T. ignotum S. (Fig. 286, d). Spore with 8 sporoplasms; in Tubifex tubifex.

T. magnum Granata. Spore with 16 sporoplasms; in Limnodrilus udekemianus.

T. legeri Mackinnon and Adams. Spore with 24 sporoplasms; in Tubifex tubifex.


Genus Sphaeractinomyxon Caullery and Mesnil. In the coelom of oligochaetes; spores rounded, without any processes; in early stage
of development, there are 2 uninucleate bodies surrounded by a binucleate envelope; 2 inner cells multiply into 16 cells which unite in pairs; nucleus of zygote of sporont divides first into 2; 1 of the nuclei divides into 6 which form 3 shell-valves and 3 polar capsules, while the other nucleus together with a portion of cytoplasm remains outside the envelope, and undergoes multiplication; multinucleate sporoplasm migrates into spore; sporoplasm later divides into a large number of uninucleate sporoplasms which, when spores gain entrance into a new host, begin development.

S. stolci C. and M. (Fig. 286, b). Spore spherical; in Clitellis arenarius and Hemitubifex benedii.
S. gigas Granata (Fig. 286, c). In the coelom of Limnodrilus hoffmeisteri (Granata, 1925).

Genus Hexactinomyxon Stolc. Each of 3 shell-valves prolonged into 2 processes; spore appears as a 6-armed anchor.

H. psammoryctis S. (Fig. 286, c). In the gut-epithelium of Psammoryctes barbatus; sporoplasm multinucleate.

Genus Synactinomyxon Stolc. Spore with 2 prolonged shell-valves and 1 conical valve.

S. tubificis S. (Fig. 286, f, g). In the gut-epithelium of Tubifex tubifex.

Genus Neoactinomyxum Granata. 3 shell-valves without any process, distended to hemisphere.

N. globosum G. (Fig. 286, h). In the gut-epithelium of Limnodrilus udekemianus; spore with numerous sporoplasts (Granata, 1925; Jirovec, 1940).

Genus Guyenotia Naville. Pansporoblast with 8 spores; spore spherical with 3 shell-valves, each drawn out posteriorly into digitiform process, longer than diameter of spore; sporoplasm with 32 nuclei.

G. sphaerulosa N. (Fig. 286, i). In the gut-epithelium of Tubifex tubifex; spores 15μ in diameter; appendages of mature spore 40μ long.

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Chapter 29
Order 3 Microsporidia Balbiani

The Microsporidia are far more widely distributed as parasites among various animal phyla than the Myxosporidia. They are however typically parasites of arthropods and fishes. All Microsporidia invade and undergo asexual division and sporogony within the host cell. These infected cells may show frequently an enormous hypertrophy of both the cytoplasmic body and nuclei (Figs. 287, f, g; 290, a–c), a characteristic feature of the host reaction toward this particular group of protozoan parasites.

The microsporidian spore is on the whole relatively small as compared with that of Myxosporidia. In the vast majority it measures 3–6μ in the largest diameter. The chitinous spore membrane which is apparently of a single piece except in a few species, envelops the

Fig. 287. Effects of microsporidian infections upon host animals. a, the central nervous system of Lophius piscatoris infected by Nosema lophii (Doflein); b, a smelt infected by Glugea hertwigi, X3 (Schrader); c, a Culex larva infected by Thelohania opacita, X14 (Kudo); d, a Simulium larva infected by T. multispora, X10 (Strickland); e, portion of testis of Barbus barbus infected by Plistophora longifilis, X1.4 (Schuberg); f, g, normal and hypertrophied nuclei of the adipose tissue cells of larval Culex pipiens, the latter due to a heavy infection by Stempellia magna, X1330 (Kudo).
sporoplasm and the polar filament, a very long delicate filament. The latter may be enclosed within a polar capsule as in a myxosporidian spore. Structure of microsporidian spores (Léger and Hesse, 1916a; Kudo, 1920, 1921, 1924b; Kohler, 1921).

When such spores are taken into the digestive tract of a specific host (Fig. 288), the polar filaments are extruded and perhaps anchor the spores to the gut-epithelium (a). The sporoplasts emerge as amoebulae through the opening after the filaments become completely detached (b). By amoeboid movements they penetrate through the intestinal epithelium and enter the blood stream or body cavity and reach the specific site of infection (c). They then enter the

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**Fig. 288.** The life-cycle of *Stempellia magna*, ×800 (Kudo). a, b, germination of spore in the mid-gut of culicine larva; c–k, division stages; l–p, sporont formation; q–t, formation of 1, 2, 4, and 8 sporoblasts; u, sporoblast; v–x, development of sporoblast into spore.

host cells and undergo multiplication at the expense of the latter (d–n). The trophozoites become sporonts (o), each of which produces a number of spores (p–x) characteristic of each genus. Some spores seem to be capable of germinating in the same host body, and thus the number of infected cells increases. When heavily infected, the host animal dies as a result of the degeneration of enormous numbers of cells thus attacked. Such fatal infections may occur in an epidemic form, as is well known in the case of the pèbrine disease of silkworms.
PROTOZOOLOGY

(Pasteur, 1870; Stempell, 1909; Kudo, 1916; Hutchinson, 1920; Jameson, 1922), Nosema-disease of honey bees (Zander, 1911; White, 1919; Farrar, 1947), microsporidiosis of mosquitoes (Kudo, 1921-1930), etc. Taxonomy (Léger and Hesse, 1922; Kudo, 1924b; Jiřívec, 1936; Weiser, 1947); the polar filament (Kudo, 1913, 1918, 1924b; Morgenthaler, 1922; Ohshima, 1927, 1937).

The Microsporidia are subdivided into two suborders:

Spore with a single polar filament... Suborder 1 Monocnidea (p. 670)
Spore with 2 polar filaments................. Suborder 2 Dienidea (p. 678)

Suborder 1 Monocnidea Léger and Hesse

Spore oval, ovoid, or pyriform, if subcylindrical length less than 4 times breadth..................... Family 1 Nosematidae
Spore spherical or subspherical......... Family 2 Coccosporidiae (p. 676)
Spore tubular or cylindrical, width less than 1/5 length, straight or curved.......................... Family 3 Mrazekiidae (p. 676)

Family 1 Nosematidae Labbé

The majority of Microsporidia belong to this family.


N. bombycis N. (Fig. 289, a, b). In all tissues of embryo, larva, pupa and adult of Bombyx mori; spores 3-4μ by 1.5-2μ, polar filament 57-72μ long when extruded; advanced infection is characterized by numerous minute brownish-black spots scattered over the body surface, which gave rise to such names as pébrine disease (France), Fleckenkrankheit (Germany), Biriushi-Bio (Japan), Cota (India), etc. (Fig. 289, b) to the disease; heavily infected larvae cannot spin cocoon and perish; the organisms invade, and develop in, ova so that newly hatched larvae are already infected with this microsporidian. Viable spores introduced per os bring about infections in Arctia caja (Stempell, 1909), Margarnia pyloalis, Chilo simplex (Ohshima, 1935), and Hyphantria cunea (Kudo and DeCoursey, 1940). Morphology and Development (Stempell, 1909; Kudo, 1924b).

N. bryozoides (Korotneff) (Fig. 289, c, d). In the germ cells and cavity of the bryozoans, Plumatella fungosa and P. repens; spores 7-10μ by 5-6μ (Braem, 1911; Schröder, 1914).

N. apis Zander (Fig. 289, e-g). In the mid-gut of honey bees; spores 4-6μ by 2-4μ; the extruded filament shows often 2 sections of different undulations (Fig. 289, g) (Kudo, 1921a). The infection is confined to the digestive system, but the ovary of an infected queen bee undergoes various degrees of degeneration depending on the extent of the gut infection (Fyg, 1945; Farrar, 1947; Hassanein, 1951),
though the eggs are free from the parasites, which condition may be
looked upon as a parasitic castration. Morphology and development
(Zander, 1909; Fantham and Porter, 1912).

N. cyclopis Kudo (Fig. 289, h, i). In Cyclops fuscus; spores 4.5μ by
3μ (Kudo, 1921b).

N. anophelis K. (Fig. 289, j, k). In the larvae of Anopheles quadri-
maculatus; spores 5–6μ by 2–3μ (Kudo, 1925). It was also found in A.
maculipennis (Missirol, 1928).

Fig. 289. a, b, Nosema bombycis (Kudo) (a, fresh spores, ×1500; b, a
heavily infected silkworm larva showing characteristic dots on integu-
ment, ×1); c, d, N. bryozoides (c, infected funiculus, ×270 (Braem); d, a
stained spore, ×1200 (Schröder)); e–g, N. apis (Kudo) (e, a fresh spore; f,
a stained spore, ×1560; g, a spore with the extruded polar filament as seen
in dark field, ×800); h, i, views of fresh spores of N. cyclopis, ×1560
(Kudo); j, k, fresh spores of N. anophelis, ×1600 (Kudo); l, m, preserved
and stained spores of N. aedis, ×1530 (Kudo); n, Frenzelina conformis, a
gregarine, infected by schizonts and spores of Nosema frenzeliniae (Léger
and Duboscq); o–q, Nosema notabilis, ×1400 (Kudo) (o, a stained tropho-
zoite of Sphaerospora polymorpha, a myxosporidian, infected by six
trophozoites of Nosema notabilis; p, another host trophozoite in which
nine spores and two trophozoites of N. notabilis occur; q, six fresh spores
of N. notabilis).
N. aedis K. (Fig. 289, l, m). In the adipose tissue of a larval *Aedes aegypti*; spores broadly pyriform and measure 7.5–9μ by 4–5μ; polar capsule large; uninucleate sporoplasm posterior (Kudo, 1930).

N. frenzelinae Léger and Duboscq (Fig. 289, n). In the cytoplasm of the cephaline gregarine, *Frenzelina conformis*, parasitic in the gastric caeca and intestine of *Pachygrapsus marmoratus*; spores about 2.8μ long; extruded polar filament up to 25μ long (Léger and Duboscq, 1909).

N. notabilis Kudo (Fig. 289, o–q). In the trophozoite of the myxosporidian, *Sphaerospora polymorpha* (p. 653) which inhabits the urinary bladder of *Opsanus tau* and *O. beta*. The host fish remain free from the microsporidian infection. The entire development takes place in the cytoplasm of the host trophozoites. Trophozoites small binucleate, multiply by binary fission. Spores ovoid to ellipsoid; sporoplasm binucleate; fresh spores 2.9–4μ by 1.4–2.5μ; extruded polar filament 45–62μ. When heavily infected, the host myxosporidian trophozoites degenerate and disintegrate. A unique example of hyperparasitism in which two cnidosporidians are involved (Kudo, 1944).

Genus Glugea Thélohan. Each sporont develops into 2 spores; the infected host cells become extremely hypertrophied, and transform themselves into the so-called Glugea cysts (Figs. 287, b; 290, e). Many species (Kudo, 1924b).

*G. anomala* (Moniez) (Fig. 290, a–f). In *Gasterosteus aculeatus*, *G. pungitus* (sticklebacks) and *Gobius minutus*; cysts conspicuous, up to about 5 mm. in diameter; host cells are extremely hypertrophied; spores 4–6μ by 2–3μ. Morphology and sporogony (Stempell, 1904; Weissenberg, 1913; Debaisieux, 1920).

*G. müllerii* Pfeiffer. In the muscles of *Gammarus pulex* and *G. locusta*; spores 5–6μ by 2–3μ (Debaisieux, 1919).

*G. hertwigi* Weissenberg (Figs. 287, b; 290, g, h). In the smelt, *Osmerus mordax* and *O. eperlanus*. Schrader (1921) found the intestine the primary site of infection, the cysts varying in size, up to 3 mm. in diameter; as the cysts grow in the mucosa, they come to lie immediately under the peritoneum. Spores measure 4–5.5μ by 2–2.5μ. Fantham, Porter and Richardson (1941) found the cysts in the serous membrane of the hind gut; as the spores were 3.5–4.6μ by 1.5–2μ, they named the organism Glugea hertwigi var. canadensis. Morphology and spore-formation (Weissenberg, 1911, 1913; Schrader, 1921).

Genus Perezia Léger and Duboscq. Each sporont produces 2
spores as in Glugea, but infected host cells are not hypertrophied. A few species.

*P. mesnili* Paillot (Fig. 290, i). In cells of silk glands and Malpi-

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**Fig. 290.** a–f, *Glugea anomala* (a, a young trophozoite in a connective tissue cell of the intestine of a young host fish, seven days after feeding on spores; b, c, more advanced stages; d, a later stage, the host cell being multinucleated and 41 μ in diameter, X1000 (Weissenberg); e, section of an infected *Gasterosteus aculeatus*, showing two large cysts (Thélohan); f, a fresh spore, X1500 (Stempell)); g, h, *G. hertwigi* (Schrader) (g, cross-section of the infected intestine of a smelt, X14; h, 2 spores); i, stained spores of *Perezia mesnili*, X2265 (Paillot); j, section of *Lankesteria ascidiae*, a gregarine, infected by *P. lankesteriae*, X900 (Léger and Duboscq); k–o, *Gurleya tetraspora* (k, infected hypodermal cells of Moina, X660 (Jirovec)); l, a mature sporont; m, a fresh spore (Dofflein); n, stained spores; o, spores with extruded polar filaments (Jirovec)); p, q, a sporont and a spore with the extruded filament of *Gurleya richardi*, X1200 (Cépède).
ghian tubules of larvae of *Pieris brassicae*; spores 3.4 μ by 1.5–2 μ (Paillot, 1918, 1929).

*P. lankesteriae* Léger and Duboscq (Fig. 290, f). In the cytoplasm of the gregarine, *Lankesteria ascidiae*, parasitic in the intestine of the tunicate, *Ciona intestinalis*. It attacks only the gregarine which are free in the lumen of the gut; the host nucleus does not undergo hypertrophy; ovoid spores 2.5 μ long.

Genus *Gurleya* Doflein. Each sporont develops into four sporoblasts and finally into four spores. A few species.

*G. tetraspora* D. (Fig. 290, k–o). In the hypodermal cells of *Daphnia maxima* and *Moina rectirostris*; spores pyriform, 2.8–3.4 μ by 1.4–1.6 μ (Jírovec, 1942). The infected host appears opaque white.

*G. richardi* Cépède (Fig. 290, p, q). In *Diaptomus castor*; spores 4–6 μ by 2.8 μ.

Genus *Thelohania* Henneguy. Each sporont develops into 8 sporoblasts and ultimately into 8 spores; sporont membrane may degenerate at different times during spore formation. Numerous species.

*T. legeri* Hesse (*T. illinoisensis* Kudo) (Figs. 76; 291, a–e). In the fat bodies of the larvae of several species of Anopheles; spores 4–6 μ by 3–4 μ; heavily infected larvae die without pupation; widely distributed. Spore-formation (Kudo, 1924).

*T. opacita* Kudo (Figs. 287, c; 291, f, g). In the adipose tissue of the larvae of *Culex* mosquitoes; spores 5.5–6 μ by 3.5–4 μ (Kudo, 1922, 1924a).

*T. reniformis* Kudo and Hetherington (Fig. 291, h). In the gut cells of the nematode, *Protospirura muris*, in mice; reniform spores 3–4 μ by 1.5–1.8 μ (Kudo and Hetherington, 1922).

Genus *Stempellia* Léger and Hesse. Each sporont produces 1, 2, 4, or 8 sporoblasts and finally 1, 2, 4, or 8 spores. 2 species.

*S. magna* Kudo (Figs. 287, f, g; 288; 291, i–l). In fat-bodies of various culicine larvae; spores 12.5–16.5 μ by 4–5 μ; polar capsule visible in life; polar filament when extruded under mechanical pressure measures up to 350–400 μ long (Kudo, 1925a).

Genus *Duboscquia* Pérez. Sporont develops into 16 sporoblasts and finally 16 spores. Host-cell nuclei extremely hypertrophied. One species.

*D. legeri* P. (Fig. 291, m–o). In the fat-body cells of *Reticulitermes lucifugus* and *R. flavipes*. Trophozoites invade the peri-midintestinal adipose tissue cells which become enlarged into “cysts,” up to 660 μ by 300 μ, because of active multiplication of the organisms; each binucleate schizont becomes a sporont which grows and produces 16 spores. Spores ovoid to ellipsoid; fresh spores are 4.3–5.9 μ by 2.2–3 μ;
Fig. 291. a–e, *Thelohania legeri*, ×1570 (Kudo) (a, b, stained sporogenic stages; c, d, mature sporonts; e, a fresh spore); f, g, mature octosporous and tetrasporous sporonts of *T. opacita*, ×1570 (Kudo); h, gut epithelial cells of *Protospirura* infected by *T. reniformis*, ×1040 (Kudo and Hetherington); i–l, *Stempellia magna*, ×1570 (Kudo) (i, j, fresh spores; k, slightly pressed spore in Lugol; l, a spore with the nearly completely extruded polar filament, stained after Fontana); m–o, *Duboscqia legeri* (Kudo) (m, the mid-gut of *Reticulitermes flavipes* with an enlarged and two uninfected fat bodies, ×57; n, portion of an infected and two uninfected fat body cells of the termite in section; o, mature sporont in life, ×1530); p, q, *Trichoduboscqia epeori* (Léger) (p, a mature sporont, ×1330; q, a fresh spore, ×2670); r, s, stained spores of *Plistophora longifilis*, ×1280 (Schuberg).
sporoplasm uninucleate; extruded polar filament 80–95μ long (Pérez, 1908; Kudo, 1942).

Genus *Trichoduboscqia* Léger. Similar to *Duboscqia* in number of spores produced in each sporont; but sporont with 4 (or 3) rigid transparent prolongations, difficult to see in life. One species.

*T. epeori* L. (Fig. 291, p, q). In fat-bodies of nymphs of the mayflies, *Epeorus torrentium* and *Rhithrogena semicolorata*; sporonts spherical, 9–10μ in diameter, with usually 16 spores; prolongations of membrane in sporont, 20–22μ long; spores pyriform, 3.5–4μ long (Léger, 1926).

Genus *Plistophora* Gurley. Sporont develops into variable number (often more than 16) of sporoblasts, each of which becomes a spore. Several species.

*P. longifilis* Schuberg (Figs. 287, e; 291, r, s). In the testis of *Barbus fluviatilis*; spores 3μ by 2μ to 12μ by 6μ; extruded polar filament up to 510μ long.

*P. kudoi* Sprague and Ramsey. In the epithelial cells of the midgut of *Blatta orientalis*; fresh spores about 3.2μ by 1.75μ; polar filament 25–50μ long.

Genus *Pyrotheca* Hesse. Schizogony and sporogony unknown; spores elongate pyriform, anterior end attenuated, posterior end rounded, slightly curved; sporoplasm in posterior region, with 1–2 nuclei; polar capsule large. One species (Hesse, 1935).

*P. incurvata* H. (Fig. 292, a, b). In fat-bodies and haemocoel of *Megacylcops viridis*; spores 14μ by 3μ; polar filament 130μ long.

Family 2 *Coccosporidae* Kudo

Genus *Coccospora* Kudo (*Cocconema* Léger and Hesse). Spore spherical or subspherical. Several species (Léger and Hesse, 1921, 1922; Kudo, 1925b).

*C. slavinae* (L. and H.) (Fig. 292, c, d). In gut-epithelium of *Slavina appendiculata*; spores about 3μ in diameter.

Family 3 *Mrazekiidae* Léger and Hesse


*M. caudata* L. and H. (Fig. 292, e, f). In the lymphocytes of *Tubifex tubifex*; spore cylindrical, 16–18μ by 1.3–1.4μ, with a long process.

Genus *Bacillidium* Janda. Spore cylindrical, but without any process; one end narrowed in a few species (Janda, 1928). Several species (Jírovec, 1936a).
**MICROSPORIDIA**

*B. criodrili* J. (Fig. 292, g). In the lymphocytes in the posterior portion of the body cavity and nephridia of *Criodrilus lacuum*; infected lymphocytes become hypertrophied from 15μ to 200–400μ in diameter; the infected part of the body appears yellowish; spores 20–22μ by 1μ (Janda); 15.5–17μ by 1.2–1.4μ up to 24–25μ by 1.6μ (commonly 18–20μ by 1.4–1.5μ) (Jirovec).

![Diagram](image-url)

Fig. 292. a, b, stained spores of *Pyrotheca incurvata*, ×1330 (Hesse); c, d, spores of *Coccospora slavinae*, the latter with extruded filament, ×1330 (Léger and Hesse); e, f, *Mrazekia caudata* (e, an infected host cell, ×465 (Mrazek); f, a spore, ×1165 (Léger and Hesse)); g, *Criodrilus lacuum*, infected by *Bacillidium criodrili*, showing the enlarged posterior region, ×1/3 (Janda); h, i, *B. limnodrili* (Jirovec) (h, trophozoites and spores of the microsporidian in a host lymphocyte, ×600; i, a stained spore, ×930); j, k, stained spores of *Cougourdella magna*, ×1330 (Hesse); l, a spore of *Octospora muscae-domesticae*, ×1430 (Chatton and Krempf); m, n, spores of *Spiroglugea octospora* (Léger and Hesse) (m, ×665; n, ×2000); o, p, spores of *Toxoglugea vibrio* (Léger and Hesse) (o, ×665; p, ×2000); q, stained spores of *T. gerridis*, ×2000 (Poisson); r, s, a fresh and a stained spore of *Telomyxa glugeiformis*, ×2000 (Léger and Hesse).

*B. limnodrili* Jirovec (Fig. 292, h, i). In lymphocytes within gonads of *Limnodrilus claparedeanus*; spores 22–24μ by 1.5μ (Jirovec, 1936a).

Genus *Cougourdella* Hesse. Spore cylindrical, with an enlarged extremity, resembling the fruit of *Lagenaria cougourda*. 3 species (Hesse, 1935).

*C. magna* H. (Fig. 292, j, k). In haemocoele and fat body of Mega-
cyclops viridis; spores 18μ by 3μ; polar filament 110μ long; sporoplasm with 1–2 nuclei or 2 uninucleate sporoplasms.

Genus Octosporea Flu. Spore cylindrical; more or less curved; ends similar. 6 species (Jírovee, 1936a).

O. muscae-domesticae F. (Fig. 292, l). In gut and germ cells of Musca and Drosophila; spores 5–8μ long (Chatton and Krempf, 1911).

Genus Spiroglugea Léger and Hesse. Spore tubular and spirally curved; polar capsule large. One species.

S. octospora L. and H. (Fig. 292, m, n). In fat body of larvae of Ceratopogon sp.; spores 8–8.5μ by 1μ.

Genus Toxoglugea (Toxonema) Léger and Hesse. Minute spore curved or arched in semi-circle. 4 species (Poisson, 1941).

T. vibrio L. and H. (Fig. 292, o, p). In the fat body of Ceratopogon sp.; spores 3.5μ by less than 0.3μ.

T. gerridis Poisson (Fig. 292, q). In the fat body of the bug, Aquarius najas; sporont gives rise to eight sporoblasts and then to eight spores; also monosporous; microspores 4.5μ by 0.8μ, the polar filament 40–50μ long; macrospores 7–8μ long.

Suborder 2 Dicnidea Léger and Hesse

Family Telomyxidae Léger and Hesse

Genus Telomyxa Léger and Hesse. Spore with 2 polar capsules; sporont develops into 8, 16, or more sporoblasts and finally 8, 16, or more spores (Léger and Hesse, 1910). Four species (Poisson, 1941).

T. glugciformis L. and H. (Fig. 292, r, s). In the fat body of the larva of Ephemera vulgata; spores 6.5μ by 4μ.

Order 4 Helicosporidia Kudo

This order has been created to include the interesting organism, Helicosporidium, observed by Keilin. Although quite peculiar in the structure of its spore, the organism seems to be best placed in the Cnidosporidia.

The minute spore is composed of a thin membrane of one piece and of three uninucleate sporoplasms, around which is coiled a long thick filament. Young trophozoites are found in the host tissues or body cavity. They undergo schizogony, at the end of which uninucleate sporonts become differentiated. A sporont divides apparently twice and thus forms four small cells which develop into a spore. The complete life-history is still unknown.

Genus Helicosporidium Keilin. Parasitic in arthropods; schizog-
Fig. 293. Diagram illustrating the probable development of Helicosporidia, × about 1600 (Keilin). a–c, schizont and schizogony; d, sporont(?); e, three stages in formation of four-celled stage; f, hypothetical stage; g, young spore before the spiral filament is formed; h, mature spore; i, j, opening of spore and liberation of sporoplasms. a–h, in living host larva; i, j, in dead host body.

ony and sporogony; spore with central sporoplasms and a single thick coiled filament. One species (Keilin, 1921).

_H. parasiticum_ K. (Fig. 293). In body cavity, fat body, and nervous tissue of larvae of _Dasyhelea obscura_ and _Mycetobia pallipes_ (Diptera), and _Hericia hericia_ (Acarina), all of which inhabit wounds of elm and horse-chestnut trees; schizonts minute; spores 5–6μ in diameter; extruded filament 60–65μ by 1μ thick.

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Chapter 30

Subphylum 2 Ciliophora Doflein

The Ciliophora possess cilia which serve as cell-organs of locomotion. In Suctoria the cilia are present only during early developmental stages. The members of this subphylum possess a unique organization not seen in the Plasmodromae; namely, except Proto-ciliata, the Ciliophora contain two kinds of nuclei: the macronucleus and the micronucleus. The former is large and massive, and controls the metabolic activities of the organism, while the latter is minute and usually vesicular or less compact, and is concerned with the reproductive processes. Nutrition is holozoic or parasitic; holophytic in Cyclo-trichium meunieri (p. 706). Sexual reproduction is mainly by conjugation, and asexual reproduction is by binary fission or budding. The majority are free-living, but a number of parasitic forms also occur.

The Ciliophora are divided into two classes:

Cilia present throughout trophic life ....................... Class 1 Ciliata
Adult with tentacles; cilia only while young.. Class 2 Suctoria (p. 863)

Class 1 Ciliata Perty

The class Ciliata includes Protozoa of various habitats and body structures, though all possess cilia or cirri during the trophic stage. They inhabit all sorts of fresh and salt water bodies by free-swimming, creeping, or being attached to other objects; some are parasitic in other animals. Free-swimming forms are usually spherical to elliptical, while the creeping forms are, as a rule, flattened or compressed.

The cilia are extremely fine, comparatively short, and as a rule arranged in rows (p. 55). In some forms they diminish in number and are replaced by cirri (p. 57). The cilia are primarily cell-organs of locomotion, but secondarily through their movements bring the food matter into the cytostome. Moreover, certain cilia appear to be tactile organellae. The food of free-living ciliates consists of small plant and animal organisms which ordinarily abound in the water; thus their nutrition is holozoic. The ciliates vary in size from less than 10μ up to 2 mm. in large forms (as in an extended Spirostomum or Stentor). The cytoplasm is distinctly differentiated into the ectoplasm and the endoplasm. The ectoplasm gives rise to the cilia and trichocysts and is covered by a pellicle. The endoplasm contains nuclei, food vacuoles, contractile vacuoles, pigment granules, crystals, etc. 683
In the majority of ciliates, the anterior and posterior extremities are permanent and distinct; in all cytostome-possessing forms, the oral and aboral surfaces are distinguishable, while in numerous creeping forms the dorsal and ventral sides are differentiated.

The body is covered by a very thin yet definite membrane, the pellicle, which is ordinarily uniformly thin and covers the entire body surface so closely that it is not recognizable in life. In some forms, such as Coleps, it develops into numerous platelets and in others, such as Trichodina, into hook-like processes. The outer half of the ectoplasm may show alveolar structure which, in section, exhibits radiating and parallel lines. In this portion the myonemes (p. 61) are lodged. The deeper layer of the ectoplasm is structureless and free from granules. In the ectoplasm are embedded the kinetosomes of cilia, which are arranged in longitudinal, oblique, or spiral rows. In recent years complex fibrillar systems have been recognized in many ciliates (p. 63-70). The cilia may fuse to form cirri, membranelae, and undulating membranes (p. 59) which occur in certain groups. In many euciliates contractile vacuoles with one to several collecting canals are one of the prominent structures. The endoplasm is more fluid and the ground substance is finely granulated or reticulated; it undergoes rotation movement or cyclosis.

Two types of nuclei are present in all euciliates. The massive macronucleus is of various forms. The chromatin granules which may reach 20µ in diameter (p. 42) fill compactly the intranuclear space. The macronucleus multiplies by amitosis. The micronucleus is ordinarily so minute that it is difficult to see in a living specimen. It is vesicular in structure, although in some it appears to be compact, and consists of an endosome, the chromatin, the nucleoplasm, and the membrane. The number of micronuclei present in an individual varies among different species. At the time of reproduction it increases in size and divides mitotically; during conjugation it undergoes a characteristic meiotic division (p. 206).

The protociliates possess from two to many nuclei of a uniformly same structure and numerous ovoid or spindle-shaped bodies, endospherules, the nature of which is open to speculation. Some authors think that they are nuclei (micronuclei (after Hickson, 1903) or macronuclei (after Konsuloff, 1922, 1930)); others consider them as reserve food materials (Patten). Metcalf (1909) considers that each nucleus possesses both metabolic chromatin and reproductive chromatin, the former being seen as large flattened peripheral masses and the latter, as smaller spheroidal granules.

In all except protociliates and a comparatively small number of
CILIOPHORA, PROTOCILIATA

astomatous euciliates, there is a cytostome which in its simplest form is represented by a small opening on the pellicle, and may or may not be closed when the animal is not feeding. The cytostome opens into the cytopharynx (or gullet), a tubule which ends in the deeper portion of the endoplasm. In the cytopharynx there may be present one or more undulating membranes to facilitate intaking of the food. Occasionally the cytostome is surrounded by trichites or trichocysts (p. 71). When the cytostome is not at the anterior region as, for instance, in Paramecium, there is a peristome (or oral groove) which starts at or near the anterior end and runs posteriorly. The peristome is ciliated so that food particles are thrown down along it and ultimately into the cytostome which is located at its posterior end. Solid waste particles are extruded from the cytopyge, or cell-anus, which is usually noticeable only at the time of actual defecation (p. 108).

Cytology (Konsuloff, 1922; Wetzel, 1925). Following Metcalf, Ciliata are here divided into 2 subclasses:

Two to many nuclei of one kind; sexual reproduction permanent fusion...

.................................................................................Subclass 1 Protociliata

Macronucleus and micronucleus; sexual reproduction conjugation......

.................................................................................Subclass 2 Euciliata (p. 690)

Subclass 1 Protociliata Metcalf

The protociliates are almost exclusively inhabitants of the large intestine of Salientia; only a few species have been reported from urodeles, reptiles, and fish (Metcalf, 1923, 1940). The body is covered uniformly by cilia of equal length. There is no cytostome and the nutrition is parasitic (saprozoic). The number of nuclei varies from two to many, all of which are of one type. Asexual reproduction is by binary fission or plasmotomy. In a number of species sexual fusion of 2 gametes has been observed (Metcalf, 1909; Konsuloff, 1922) (Fig. 294, f–i). Grassé (1952) proposed recently to transfer these organisms to "Rhizoflagellata" from Ciliata, since they differ from the ciliates in (1) having nuclei of the same kind, (2) undergoing sexual fusion and not conjugation, and (3) having longitudinal, and not transverse, division or plasmotomy. Taxonomy (Metcalf, 1920a, 1923, 1940); geographical distribution (Metcalf, 1920, 1929, 1940); cytology and development (van Overbeek de Meyer, 1929); species (Bhatia and Gulati, 1927; Carini, 1938–1942; Beltran, 1941, 1941a).

Family Opalinidae Claus

Genus Opalina Purkinje and Valentin. Highly flattened; multinucleate; in amphibians. Numerous species (Metcalf, 1923, 1940).
Growth and nuclear division (Hegner and Wu, 1921); cytology (ten Kate, 1927).

*O. hylaxena* Metcalf (Fig. 294, a). In *Hyla versicolor*; larger in individuals about $420 \mu$ long, $125 \mu$ wide, $28 \mu$ thick. Several subspecies (Metcalf).

*O. obtrigonoidea* M. (Fig. 294, b–d). $400-840 \mu$ long, $175-180 \mu$ wide, $20-25 \mu$ thick; in various species of frogs and toads (*Rana, Hyla, Bufo, Gastrophryne*, etc.), North America. Numerous subspecies (Metcalf).

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Fig. 294. a–i, l, Metcalf; j, k, Léger and Duboscq. a, two individuals of *Opalina hylaxena*, $\times 78$; b–d, three individuals of *O. obtrigonoidea*, $\times 78$ (b, from *Bufo fowleri*; e, from *Rana p. pipiens*; d, from *R. p. palustris*); e, four individuals of *Cepedea cantabrigensis*, $\times 78$; f–i, stages in sexual reproduction in *Protoopalina intestinalis*; j, k, *P. saturnalis*, $\times 500$; l, *P. mitotica*, $\times 240$. 
O. carolinensis M. 90–400\(\mu\) by 32–170\(\mu\); in Rana pipiens sphencephala.
O. pickeringii M. 200–333\(\mu\) by 68–100\(\mu\); in Hyla pickeringii.
O. oregonensis M. 526\(\mu\) by 123\(\mu\); in Hyla regilla.
O. spiralis M. 300–355\(\mu\) long, 130–140\(\mu\) wide, 25–42\(\mu\) thick; in Bufo compactilis.
O. chorophili M. About 470\(\mu\) by 100\(\mu\); in Chorophilus triseriatus.
O. kennicottii M. About 240\(\mu\) by 85\(\mu\); in Rana areolata.

Genus Cepedea Metcalf. Cylindrical or pyriform; circular in cross-section; multinucleate; all in Amphibia. Numerous species. Cytology (Fernandez, 1947).

C. cantabrigensis M. (Fig. 294, e). About 350\(\mu\) by 84\(\mu\); in Rana cantabrigensis.
C. hawaiensis M. 170–200\(\mu\) by 43–60\(\mu\); in Rana catesbeiana; Hawaii.
C. obovoidea M. About 315\(\mu\) by 98\(\mu\); in Bufo lentiginosus.

Fig. 295. Zelleriella elliptica, stained specimens, \(\times 440\) (Chen). a, a typical vegetative individual; b, an individual which is nearly completely divided, the nuclei being at early metaphase.
C. floridensis M. About 230μ by 89μ; in Scaphiopus albus.

Genus Protoopalina Metcalf. Cylindrical or spindle-shaped, circular in cross-section; 2 nuclei; in the colon of various species of Amphibia with one exception. Numerous species.

P. intestinalis (Stein) (Fig. 294, f-i). About 330μ by 68μ; in Bombina bombina, and B. pachypa; Europe.

P. saturnalis Léger and Duboscq (Fig. 294, j, k). In the marine fish, Box boops; 100–152μ by 22–60μ.

P. mitotica (M) (Fig. 294, l). 300μ by 37μ; in Ambystoma tigrinum.


Z. scaphiopodos M. In Scaphiopus solitarius; about 150μ long, 90μ thick.

Z. antilliensis (M). About 180μ long, 113μ wide, 32μ thick; in Bufo marinus.

Z. hirsula M. About 113μ long, 60μ wide, 22μ thick; in Bufo cognatus.

Z. elliptica Chen (Fig. 295). In Bufo valliceps; average dimensions 184μ by 91μ. Chen (1948) distinguishes four other species from the same host, all of which possess 24 chromosomes.

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THE most conspicuous group of Protozoa containing 2 nuclei; macronucleus and micronucleus. Sexual reproduction is through conjugation. We owe Kahl a great deal for his series of comprehensive taxonomic studies of free-living ciliates. The euciliates are grouped under the following four orders:

Without adoral zone of membranellae.................Order 1 Holotricha
With adoral zone of membranellae
  Adoral zone winds clockwise to cytostome
  Peristome not extending beyond general body surface.............Order 2 Spirotricha (p. 796)
  Peristome extending out like funnel....Order 3 Chonotricha (p. 847)
  Adoral zone winds counter-clockwise to cytostome..............Order 4 Peritricha (p. 850)

For a brief, but concise view on the classification of the ciliates, the reader is referred to Faure-Fremiet (1950).

Order 1 Holotricha Stein

The members of this order show uniform ciliation over the entire body surface. Adoral zone does not occur. The majority possess a cytostome which varies among different forms. Nutrition is holozoic or saprozoic. Asexual reproduction is usually by transverse fission and sexual reproduction by conjugation. Encystment is common. The holotrichous ciliates are conspicuous free-living forms in all sorts of fresh, brackish, and salt waters, though some are parasitic.

The order is here divided into 6 suborders:

Without cytostome..........................Suborder 1 Astomata (p. 691)
With cytostome

  Cytostome not rosette-like
  Without special thigmotaetic ciliated field
    Cytostome on body surface or in peristome, without strong cilia...
    Suborder 2 Gymnostomata (p. 700)
  Cytostome in peristome, bearing special cilia or membranes
    Peristome lined with rows of free cilia.
    Suborder 3 Trichostomata (p. 737)
    Peristome with membrane; with or without free cilia.
    Suborder 4 Hymenostomata (p. 758)
  With well-developed thigmotactic ciliated field; commensals in mussels.
    Suborder 5 Thigmotricha (p. 774)
  Cytostome small rosette-like aperture or obscure; parasitic
    Suborder 6 Apostomata (p. 789)
The ciliates placed in this suborder possess no cytostome, although there may occur a slit-like organella which has been looked upon as a vestigial cytostome. The body ciliation is usually uniform. Asexual division is carried on by transverse fission and often by budding which results in chain formation. Sexual reproduction is conjugation and in some encystment is known. These organisms are parasitic in various invertebrates living in fresh or salt water. Taxonomy (Cépède, 1910, 1923; Cheissin, 1930; Heidenreich, 1935; Delphy, 1936); skeletal structures (Rossolimo and Perzewa, 1929); Argyrome (Puytorac, 1951).

Without attaching organellae or skeletal structures
Macronucleus round to elongate. Family 1 Anoplophryidae
Macronucleus irregular network. Family 2 Opalinopsidae (p. 694)
With attaching organellae or skeletal structures
Contractile vacuole, a long dorsal canal; usually with a sucking organella. Family 3 Haptophryidae (p. 694)
Contractile vacuoles not canal-like; with various attaching organellae or skeletal structures. Family 4 Intoshellinidae (p. 696)

Family 1 Anoplophryidae Cépède

Genus Anoplophrya Stein (Collinia Cépède). Oval, elongate, ellipsoid or cylindrical; macronucleus ovoid to cylindrical; micronucleus small; one to several contractile vacuoles; ciliation dense and uniform; in coelom and gut of Annelida and Crustacea. Numerous species (Rossolimo, 1926).

A. marylandensis Conklin (Fig. 296, a). 36–72µ by 16–42µ; in the intestine of Lumbricus terrestris and Helodrilus caliginosus; Baltimore, Maryland (Conklin, 1930).

A. orchestii Summers and Kidder (Fig. 296, b). Polymorphic according to size; pyriform to broadly ovoid; 7–45 ciliary rows meridional, unequally spaced, and more on one surface; macronucleus voluminous, a compact micronucleus; body 6–68µ long; in the sand flea, Orchestia agilis; Woods Hole, Massachusetts (Summers and Kidder, 1936).

Genus Rhizocaryum Caullery and Mesnil. With hollowed ventral surface which serves for attachment; macronucleus drawn out like a tree-root. One species.

R. concavum C. and M. (Fig. 296, c). In the gut of Polydora caeca and P. flava (polychaetes).

Genus Metaphrya Ikeda. Pyriform, anterior end bent slightly to one side; 12 longitudinal ciliary furrows; below ectoplasm, a
layer of refringent materials; endoplasm sparse; macronucleus basket-like, large, with a spacious hollow; a micronucleus; no contractile vacuoles. One species.

*M. sagittae* I. (Fig. 296, d). About 250µ by 130µ; in the body cavity of *Sagitta* sp.

Genus *Perezella* Cépède. Ovoid; ventral surface concave, serves for attachment; macronucleus ellipsoid; contractile vacuole ter-

Fig. 296. a, *Anoplophrya marylandensis*, ×500 (Conklin); b, *A. orchestii*, ×500 (Summers and Kidder); c, *Rhizocaryum concavum*, ×670 (Cépède); d, *Metaphrya sagittae*, ×120 (Ikeda); e, *Perezella pelagica*, ×340 (Cépède); f, *Dogielella sphaerii*, ×470 (Poljansky); g, *D. minuta*, ×670 (Poljansky); h, *D. virginia*, ×670 (Kepner and Carroll); i, *Orchitophrya stellarum*, ×870; j, *Kofoidella eleutheriae*, ×270; k, *Bütschliella opheliae*, ×350 (Cépède).
minal; longitudinally, uniformly, ciliated. A few species.

P. pelagica C. (Fig. 296, e). In the coelom of copepods (Ascartia, Clausia, Paracalanus); about 48μ long.

Genus Dogielella Poljansky. Pyriform; longitudinal ciliary rows; contractile vacuole terminal; macronucleus spherical, with a spherical or elliptical micronucleus; in the parenchyma of flatworms or molluscs. 4 species (Poljansky, 1925).

D. sphaerii P. (Fig. 296, f). 40–100μ by 25–54μ: in Sphaerium corneum. Conjugation (Poljansky, 1926).

D. minuta P. (Fig. 296, g). 12–28μ by up to 20μ; in Stenostomum leucops (Platyhelminthes).

D. virginia (Kepner and Carroll) (Fig. 296, h). 40–50μ long; in the same host animal; Virginia.

D. renalis Kay. Elongate pyriform, but extremely plastic; 61–184μ by 27–82μ; spherical macronucleus in the middle of body; one micronucleus; a contractile vacuole anterior; in the renal organ of Physella sp. (Kay, 1946).

Genus Orchitophrya Cépède. Elongate pyriform; ciliary rows oblique; macronucleus spherical, central. One species.

O. stellatum C. (Fig. 296, i). In gonads of the echinoderm, Asteracanthion (Asterias) rubens; 35–65μ long.

Genus Kofoidella Cépède. Pyriform; macronucleus broadly oval; contractile vacuole, subterminal. One species.

K. eleutheriae C. (Fig. 296, j). In gastrovascular cavity of the medusa, Eleutheria dichotoma; 30–80μ long.

Genus Herpetophrya Siedlecki. Ovoid; with a pointed, mobile, tactile, non-ciliated cone; macronucleus globular; without contractile vacuole. One species.

H. astomata S. In coelom of Polymnia (annelid).

Genus Biitschliella Awerinzew. Elongate with pointed anterior end, with non-ciliated retractile anterior cap; cilia in about 10 slightly spiral rows; macronucleus band-form; several contractile vacuoles in a longitudinal row. Several species.

B. opheliae A. (Fig. 296, k). In Ophelia limacina; 280–360μ by 35–50μ.

B. chaetogastri Penard. Elongate lanceolate, slightly flattened; longitudinal rows of long cilia; cytoplasm colorless; macronucleus elongate; micronucleus voluminous, vesicular; without contractile vacuole; 60–120μ long; in the oesophagus of Chaetogaster sp.

Genus Spirobutschliella Hovasse (1950). Elongate fusiform with rounded extremities; ciliation uniform and in spiral rows; anterior tip not ciliated; pellicle thick; macronucleus, a long spindle reaching
the both ends of the body; a median micronucleus; in the intestine of Annelida.

*S. chattoni* H. In the mid-gut of *Potamoceros triqueter*, a common annelid in the vicinity of Banyuls; 180–550μ by 50μ; micronucleus fusiform, 6–10μ long; often infected by a microsporidian, *Gurleya nova* H.

Genus *Protanoplophrya* Miyashita. Similar to *Anoplophrya*; but with rudimentary oral apparatus, a long slit, an undulating membrane and cytopharynx in anterior region of body; macronucleus elongate band; numerous contractile vacuoles. One species.

*P. stomata* Miyashita (Fig. 297, a). Cylindrical; up to 1.5 mm. by about 70μ; in hind-gut of *Viviparus japonicus* and *V. malleatus*.

**Family 2 Opalinopsidae** Hartog

Genus *Opalinopsis* Foettinger. Oval or ellipsoid; macronucleus fragmented; ciliation uniform and close; parasite in the liver of cephalopods. A few species.

*O. sepiolae* F. (Fig. 297, b). 40–80μ long; in the liver of *Sepiola rondeletii* and *Octopus tetracirrhus*.

Genus *Chromidina* Gonder (*Benedenia* Foettinger). Elongate; anterior region broader, end pointed; uniform ciliation; macronucleus in irregular network distributed throughout body; micronucleus obscure; budding and encystment; Cheissin holds that this is identical with *Opalinopsis*. One species.

*C. elegans* (Foettinger) (Fig. 297, c, d). 500–1500μ by about 30–60μ in kidney and gonad of cephalopods: Sepia, Loligo, Illex and Spirula (Jepps, 1931). Morphology (Wermel, 1928).

**Family 3 Haptophryidae** Cépède

Genus *Haptophrya* Stein. Elongate; uniformly ciliated; anterior end with a neck-like constriction; a circular sucker surrounded by 1–2 rows of cilia. A few species.

*H. michiganensis* Woodhead (Fig. 297, e). 1.1–1.6 mm. long; in the gut of the four-toed salamander, *Hemidactylyum scutatum*; Michigan. Cytology (Bush, 1933); contractile canal (MacLennan, 1944).

*H. virginiensis* Meyer. 354μ by 95μ; macronucleus about one-third of the body length; in the intestine of *Rana palustris*.

Genus *Steinella* Cépède. Anterior end broad; sucker-like depression without encircling cilia, but with 2 chitinous hooks. One species.

*S. uncinata* (Schultze). Up to 200μ long; in gastrovascular cavity of *Planaria uvae*, *Gunda segmentata* and *Procercos* sp.

Genus *Lachmannella* Cépède. With a chitinous hook at anterior
end; elongate pyriform, anterior end curved; ciliation longitudinal and dense. One species.

*L. recurva* (Claparède and Lachmann) (Fig. 297, f). In the gastrovascular cavity of *Planaria limacina*; about 200μ long.

Genus *Sieboldiellina* Collin. Vermiform, with neck-like constriction; simple sucker at anterior end. One species.

*S. planaria*rum (Siebold) (Fig. 297, g). Up to 700μ long; in gastro-

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Fig. 297. a, *Protanoplophrya stomata*, ×100 (Miyashita); b, *Opalinopsis sepiolae*, ×670 (Gonder); c, d, *Chromidina elegans* (c, ×330 (Chatton and Lwoff); d, ×220 (Wermel)); e, *Haptophrya michiganensis*, ×35 (Woodhead); f, *Lachmannella recurva*, ×100 (Cépède); g, *Sieboldiellina planaria*rum, ×100 (Cépède); h, i, *Intoshellina poljanskii* (h, ×300; i, attaching organella seen from ventral side, ×870) (Cheissin); j, k, *Monodontophrya kijenskii* (j, ×100; k, anterior end in profile, ×870) (Cheissin).
vascular cavity of various fresh- and salt-water turbellarians, most frequently *Planaria torva*.

Family 4 Intoshellinidae Cépède

Genus *Intoshellina* Cépède. Elongate; ciliary rows slightly spiral; macronucleus voluminous, highly elongate; 5–7 contractile vacuoles scattered in posterior region; a complicated attaching organella at anterior end (Fig. 297, i); vestigial cytopharynx.

*I. poljanskyi* Cheissin (Fig. 297, h, i). 170–280μ long; in the intestine of *Limnodrilus arenarius*.

Genus *Monodontophrya* Vejdowsky. Elongate; anterior end with thick ectoplasm; attaching organella at anterior end, with fibrils; macronucleus elongate; numerous contractile vacuoles in a longitudinal row.

*M. kijenskiji* Cheissin (Fig. 297, j, k). 400–800μ long; in anterior portion of intestine of *Tubifex inflatus*.

Genus *Maupasella* Cépède. Ellipsoid; close longitudinal ciliary rows; with a spinous attaching organella at anterior end, with fibrils; contractile vacuoles in 2 irregular rows; macronucleus elongate. One species.

*M. nova* C. (Fig. 298, a). 70–130μ long; in the intestine of *Allolobophora caliginosa* (annelid). Supplementary chromatic body (Keilin, 1920).

Genus *Schultzellina* Cépède. Similar to *Maupasella*; but with attaching organella set obliquely; macronucleus voluminous, reniform.

*S. mucronata* C. (Fig. 298, b). In the intestine of *Allurus tetraedurus* (annelid).

Genus *Hoplitophrya* Stein. Slender, elongate; elongated macronucleus; a micronucleus; a single longitudinal row of many contractile vacuoles on the dorsal side; a single median spicule with a small pointed tooth at its anterior end; in the intestine of oligochaetes. Several species.

*H. secans* S. Elongated; 160–500μ by 20–35μ; 15–30 contractile vacuoles in a row; spicule 10–15μ long; in the intestine of *Lumbricus variiegatus*.

*H. criodrili* Miyashita (Fig. 298, c). Ellipsoid, slightly flattened; 90–130μ by 45–60μ; periphery of endoplasm highly granulated; attaching organelle about 25μ long; macronucleus bandform; two rows of contractile vacuoles; in the anterior half of the gut of an oligochaete, *Criodrilus* sp.

Genus *Radiophrya* Rossolimo. Elongate, often with satellites; attaching organella composed of an arrowhead, a tooth and ecto-
plasmic fibrils; macronucleus a narrow long band; a single row of many small contractile vacuoles, close to the nucleus. Many species.

*R. hoplites* R. (Fig. 298, d, e). 100–1000μ long; in the intestine of Lamprodrilus, Teleuscoleox, Stylosolech, and other oligochaetes.

Genus *Metaradiophrya* Heidenreich. Ovoid to ellipsoid; with 2 lateral rows of contractile vacuoles; with a hook attached to a long shaft; ectoplasmic fibers supporting the hook; in the intestine of oligochaetes. Several species.

*M. lumbrici* (Dujardin) (Fig. 298, f). 120–140μ by 60–70μ; in the intestine of *Lumbricus terrestris*, *L. rubellus* and *Eisenia foetida*. Morphology (Williams, 1942); argyrome (Puytorac, 1951).
M. asymmetrica Beers. 115–150μ by 55–70μ; hook 10μ long; shaft 25–30μ by 2μ in antero-lateral margin in ectoplasm; 25–30 supporting fibrils; 2 rows of 4 vacuoles each, which do not contract regularly in vitro; in the intestine (middle third) of Eisenia lönnergi (Beers, 1938).

Genus Protoradiophrya Rossolimo. Elongate; near anterior end a shallow depression along which is found a spicule which may be split posteriorly. A few species.

P. fissispicidata Cheissin (Fig. 298, g). 180–350μ long; in the anterior portion of intestine of Styloscolex sp.

Genus Mrazekiella Kijenskij. Elongate; anterior portion broad with sucker-like depression, posterior region cylindrical; anterior end with attaching organella composed of arrowhead and skeletal ribs; macronucleus an elongate band; contractile vacuoles distributed. A few species.

M. intermedia Cheissin (Fig. 298, h). 180–260μ long; in the anterior portion of intestine of Branchiura coccinea.

Genus Mesnilella Cépède. Elongate; with one or more long spicules imbedded in endoplasm; contractile vacuoles in 1–2 rows. Numerous species.

M. rostrata Rossolimo (Fig. 298, i). 100–1200μ long; in the intestine of various oligochaetes (Styloscolex, Teleuscolex, Lamprodrilus, Agriodrilus, etc.).

M. clavata (Leidy) (Fig. 298, j). 100–200μ long; in the intestine of Lumbricus variegatus.

References


KEILIN, D.: (1920) On the occurrence of a supplementary chromatic body in Maupasella nova, etc. Parasitology, 12:92.


SUMMERS, F. M. and KIDDER, G. W.: (1936) Taxonomic and cytological studies on the ciliates associated with the amphipod family Orchestiidae from the Woods Hole district. Ibid., 86:379.


Chapter 32
Order 1 Holotricha Stein (continued)

Suborder 2 Gymnostomata Bütschli

Cytostome at or near anterior end.................. Tribe 1 Prostomata
Cytostome not at or near anterior end
  Cytostome lateral, narrow or round...Tribe 2 Pleurostomata (p. 723)
  Cytostome ventral, in anterior half....Tribe 3 Hypostomata (p. 728)

Tribe 1 Prostomata Schewiakoff

Free-living
Cytostomal region compressed; bearing trichites.................... Family 1 Spathidiidae
Cytostomal region not compressed
  Cytostome opens into anterior receptaculum; with lorica........ Family 2 Metacystidae (p. 703)
  Cytostome at tip of apical cone.....Family 3 Didiniidae (p. 703)
  Cytostome otherwise
    Body covered with regularly arranged, perforated, ectoplasmic plates.................. Family 4 Colepidae (p. 706)
    Body not covered with plates
      With radially arranged tentacles.................................. Family 5 Actinobolinidae (p. 707)
      Without tentacles..........Family 6 Holobolinidae (p. 708)
Parasitic in mammalian gut..............Family 7 Bütschliidae (p. 717)

Family 1 Spathidiidae Kahl

Genus Spathidium Dujardin. Flask- or sack-shaped; compressed; anterior region slightly narrowed into a neck, and truncate; ciliation uniform; cytostome occupies whole anterior end; contractile vacuole posterior; macronucleus elongate; several micronuclei; trichocysts around cytostome and scattered throughout; fresh or salt water. Numerous species.

S. spathula Müller (Figs. 21, c; 299, a, b). Up to 250µ long; fresh water. Morphology and food-capture (Woodruff and Spencer, 1922); conjugation (Woodruff and Spencer, 1924).

Genus Paraspathidium Noland. Form resembles that of Spathidium; but cytostome an elongate slit, bordered on one side by strong cilia and on the other by weaker cilia and a shelf-like, nonundulatory membrane; 2 longer cilia on dorsal edge near anterior tip; anterior 1/3 compressed; posterior 2/3 nearly cylindrical; 2 oval macronuclei, each with a micronucleus; cytoplasm filled with numerous refractile granules; about 70 rows of cilia; contractile vacuole terminal; salt water. One species.

P. trichostomum N. (Fig. 299, c-e). About 220µ long; macronuclei 44µ long each; salt water; Florida (Noland, 1937).
Genus Spathidioides Brodsky (*Spathidiella* Kahl). Somewhat similar to *Spathidium*; but oral ridge highly flattened on ventral side and conspicuously developed into a wart-like swelling on dorsal side; this knob contains trichocysts; sapropelic.

*S. sulcata* B. (Fig. 299, f). 65–85μ long; posterior end pointed,
highly flattened; anterior end elevated at one side where cytostome and cytopharynx with 10 rods are located.

Genus *Enchelydium* Kahl. Somewhat similar to *Spathidium*; but oral ridge forms a swollen ring with trichocysts; the ridge circular or elongated in cross-section; when swimming, the organisms appear as if cytostome is opened; with dorsal bristle; fresh water.

*E. fusidens* K. (Fig. 299, g). Cylindrical, contractile; cilia dense and rather long; macronucleus reniform, often appears as composed of 2 spherical parts; contractile vacuole terminal; oral ring with spindle-like trichocysts; food vacuoles not seen; extended body 110μ long; contracted 75μ; sapropelic.

Genus *Homalozoon* Stokes. Elongate; cilia conspicuous on flattened right side; left side swollen or keeled; fresh water.

*H. vermiculare* (S.) (Fig. 299, h). Extended body 450–850μ long; vermiciform; macronucleus band form; contractile vacuoles about 30 or more in a row; standing fresh water.

Genus *Cranotheridium* Schewiakoff. Spathidium-like organisms; anterior end obliquely truncate, near the extended side of which is located the cytostome; cytopharynx surrounded by a group of trichites; fresh water.

*C. taeniatum* S. (Fig. 299, i). Anterior end flattened; with a group of trichites; macronucleus long band-form; with many micronuclei; contractile vacuole terminal; ciliation and striation close; colorless; movement slow; about 170μ long; fresh water.

Genus *Penardiella* Kahl. Ellipsoid, somewhat compressed; oral ridge slightly oblique; a girdle with trichocysts encircling the body; fresh water.

*P. crassa* (Penard) (Fig. 299, j). Elongate ellipsoid, flattened; trichocysts in posterior portion of girdle are longer and those in the dorsal region are fewer in number and shorter; macronucleus sausage-form; contractile vacuole posterior, in front of the girdle; body 160μ by 50μ; sapropelic.

Genus *Perispira* Stein. Ovoid or cylindrical; oral ridge turns right-spirally down to posterior end.

*P. ovum* S. (Fig. 299, k). Oval; starved individuals 30–60μ by 20–45μ, well-fed forms 65–120μ by 50–110μ; spiral ridge one complete turn; cytostome in the anterior end of the ridge, with a number of delicate trichites; ovoid to elongate macronucleus; a micronucleus; a terminal contractile vacuole; in fresh water (Dewey and Kidder, 1940). The ciliate was cultured bacteria-free by feeding on sterile *Euglena gracilis*.

*P. strephosoma* Stokes (Fig. 299, l). Oval to cylindrical; about 85μ long; standing water with sphagnum.
Genus **Legendrea** Fauré-Fremiet. Ellipsoid or ovoid; a peripheral zone with small tentacular processes bearing trichocysts.

*L. bellerophon* Penard (Fig. 299, *m*). 100–180μ; fresh water.

Genus **Teuthophrys** Chatton and Beauchamp. Body rounded posteriorly, anterior end with 3 radially equidistant, spirally curved arms (counter-clockwise when viewed from posterior end); the depressions between arms form furrows; cytostome apical, at the inner bases of arms; contractile vacuole terminal; ciliation uniform, except the inner surfaces of arms where longer cilia as well as trichocysts are present; with zoochlorellae; macronucleus rope-shaped and wound; micronucleus unobserved. One species.

*T. trisula* C and B. (Fig. 300, *a*). 150–300μ long; length: width 3:1–2:1; ponds in Pennsylvania and California (Wenrich, 1929).

**Family 2 Metacystidae** Kahl

Genus **Metacystis** Cohn. Oblong; ciliation general, except posterior end; ciliary circle around cytostome; usually one caudal cilium; with a large posterior vesicle containing turbid fluid.

*M. truncata* C. (Fig. 300, *b*). Elongate, not much difference in body width at different levels; with about 12 furrow rings; body length up to 30μ; salt water.

Genus **Vasicola** Tatem (*Pelamphora* Lauterborn). Ovoid with caudal cilia; lorica flask-shape, highly ringed; cytostome at anterior end, its lip with 4 rows of long cilia; body surface with shorter cilia; macronucleus round, central, with a micronucleus; contractile vacuole near macronucleus; fresh or salt water.

*V. ciliata* T. (*Pelamphora bütschlii* L.) (Fig. 300, *c*). Body about 100μ long; sapropelic in fresh water.

Genus **Pelatractus** Kahl. Somewhat similar to **Vasicola**; but without lorica or caudal cilia; with a terminal vacuole; without lip of **Vasicola**; sapropelic.

*P. (Vasicola) grandis* (Penard) (Fig. 300, *d*). Free-swimming; elongated fusiform; numerous contractile vacuoles on one side; body 125–220μ long; sapropelic in fresh water.

**Family 3 Didiniidae** Poche

Genus **Didinium** Stein (*Monodinium* Fabre-Domergue). Barrel-shaped; one to several girdles of cilia (pectinellae); expandable cytostome at the tip of a proboscis, supported by a dense layer of long trichites; macronucleus horseshoe-shaped; two to three and occasionally four micronuclei, close to macronucleus; contractile vacuole terminal; fresh or salt water. Several species.
D. nasutum (Müller) (Figs. 21, e, f; 40; 75; 91; 300, e–g). 80–200μ long; endoplasm highly granulated; with two girdles of pectinelles; feeds on Paramecium; spherical cysts (Fig. 75) with three walls, 60–80μ in diameter; fresh water. Morphology (Thon, 1905; Calkins, 1915; Beers, 1935); encystment, food requirement and conjugation (Beers, 1927, 1930, 1933, 1935); longevity of cysts (Beers, 1937);

Fig. 300. a, Teuthophrys trisula, ×330 (Wenrich); b, Metacystis truncata, ×270 (Cohn); c, Vasicola ciliata, ×250 (Kahl); d, Pelattractus grandis, ×170 (Penard); e–g, Didinium nasutum, ×170 (Kudo); h, D. balbianii, ×290 (Butschli); i–k, Mesodinium pulex (i, ×670; j, oral view; k, oral tentacles, ×1330) (Noland); l, m, M. acarus (l, ×670; m, oral tentacles, ×1330) (Noland); n, Askenasia faurei, ×530 (Fauré-Fremiet); o, Cyclotrichium meunieri, ×780 (Powers).
excystment (Beers, 1945, 1946) (Fig. 75); fibrillar structures (ten Kate, 1927); meiosis in conjugation (p. 206) (Prandtl, 1906).

*D. balbianii* (Fabre-Domergue) (Fig. 300, h). 60–100μ long; a single girdle of pectinelles near anterior end; fresh water.

Genus *Mesodinium* Stein. Ovoid; an equatorial furrow marks conical anterior and spherical posterior parts; in the furrow are inserted 2 (or 1) rings of strong cilia; one directed anteriorly and the other posteriorly; with tentacle-like retractile processes around the cytostome; fresh and salt water.

*M. pulex* (Claparède and Lachmann) (Fig. 300, i–k). Oral tentacles with trifurcate tips; body 20–31μ long; salt water; Florida. Noland states that the freshwater forms are 21–38μ long.

*M. acarus* Stein (Fig. 300, l, m). Oral tentacles with capitate tip; 10–16μ long; salt water, Florida (Noland, 1937).

Genus *Askenasia* Blochmann. Resembles *Didinium*; ovoid; with 2 closely arranged rings of long cilia; anterior ring made up of some 60 pectinelles which are directed anteriorly; posterior ring composed of about the same number of long cilia directed posteriorly and arranged parallel to body surface; fresh or salt water.

*A. faurei* Kahl (Fig. 300, n). Body oval, anterior end broadly rounded; posterior region conical; pectinelles about 13μ long; the second band (10μ) of long cilia; an ellipsoid macronucleus; a micronucleus; body about 58–60μ long; fresh water.

Genus *Cyclotrichium* Meunier. Body spheroid to ellipsoid with a large non-ciliated oral field which is surrounded by a pectinelle-ring,
one end dome-like, and the other truncate; macronucleus sausage-shaped; in salt water.

*C. meunieri* Powers (Fig. 300, o; 301). Anterior end broadly rounded; posterior region conical; cytostome obscure; oral funnel at anterior end in a depression; broad viliated band at about middle; ectoplasm with concave chromatophore (covered with haematochrome) plates on surface, below which numerous pyrenoids occur in vacuoles; endoplasm with numerous granules; 25–42μ by 18–34μ; Powers (1932) found that the ‘red water’ in Frenchman Bay in Maine was caused by the swarming of this organism. The same author held later that this ciliate may be the same as *Mesodinium rubrum* as observed by Leegaard (1920).

Bary and Stuckey (1950) found this organism in an extensive area of brownish-maroon water in Wellington Harbour in April and August, 1948. Their description follows: body 22–47μ by 19–41μ; anterior half dome-like, posterior half expanded; posterior end truncate; “cytostome”; greenish-maroon chromatophores close to body surface; no ingested food material.

**Family 4 Colepidae** Claparède and Lachmann

Genus *Coleps* Nitzsch. Body-form constant, barrel-shaped; with regularly arranged ectoplasmic plates; cytostome at anterior end, surrounded by slightly longer cilia; often spinous projections at or near posterior end; 1 or more long caudal cilia, often overlooked; fresh or salt water. Many species (Noland, 1925, 1937; Kahl, 1930).

*C. hirtus* (Müller) (Fig. 302, a). 40–65μ long; 15–20 rows of platelets; 3 posterior processes; fresh water.

*C. elongatus* Ehrenberg (Fig. 302, b). 40–55μ long; slender; about 13 rows (Noland, 1925) or 14–17 rows (Kahl) of platelets; 3 posterior processes; fresh water.

*C. bicuspis* Noland (Fig. 302, c). About 55μ long; 16 rows of platelets; 2 posterior processes; fresh water.

*C. octospinus* N. (Fig. 302, d). 80–110μ long; 8 posterior spines; about 24 rows of platelets; Geiman (1931) found this organism in an acid marsh pond and noted variation in number and location of accessory spines; fresh water.

*C. spiralis* N. (Fig. 302, e). About 23 longitudinal rows of platelets slightly spirally twisted; posterior spines drawn together; a long caudal cillum; about 50μ long; salt water; Florida (Noland, 1937).

*C. heteracanthus* N. (Fig. 302, f). Anterior processes only on one side; posterior spines; caudal cillum; about 90μ by 35μ; salt water; Florida.
Genus Tiarina Bergh. Somewhat similar to Coleps, but posterior end tapering to a point; salt water.

*T. fusus* (Claparède and Lachmann) (Fig. 302, g). 85–135μ long.

Family 5 Actinobolinidae Kent

Genus Actinobolina Strand (*Actinobolus* Stein). Ovate or spheroidal; ciliation uniform; extensible tentacles among cilia; contractile vacuole terminal; macronucleus curved band; fresh water.

A. *vorax* (Wenrich) (Fig. 302, h). Body 100–200μ long; elongate oval to spheroid; yellowish brown in color; cytostome at anterior end; contractile vacuole terminal; macronucleus rope-like; 30–60

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**Fig. 302.** a, *Coleps hirtus*, ×530 (Noland); b, *C. elongatus*, ×530 (Noland); c, *C. bicuspid*, ×530 (Noland); d, *C. octospinus*, ×530 (Noland); e, *C. spiralis*, ×400 (Noland); f, *C. heteracanthus*, ×400 (Noland); g, *Tiarina fusus*, ×530 (Fauré-Fremiet); h, *Actinobolina vorax*, ×300 (Wenrich); i, *Dactylochlamys pisciformis*, ×330 (Kahl); j, *Enchelyomorpha vermicularis*, ×670 (Kahl).
ciliary rows; about 30 tentacles in each ciliary row; tentacles may be extended to twice the diameter of the body or be completely withdrawn; feeds chiefly on rotifers which stop all movements as though completely paralyzed upon coming in contact with the tentacles (Wenrich, 1929a).

Genus *Dactylochlamys* Lauterborn. Body spindle-form, though variable; posterior end drawn out into tail; pellicle with 8–12 undulating spiral ridges on which tentacle-like processes and long cilia are alternately situated; these processes are retractile (Kahl) and similar in structure to those of Suctoria; cytostome has not been detected; possibly allied to Suctoria; fresh water. One species.

*D. pisciformis* L. (Fig. 302, i). Body 80–120 μ long.

Genus *Enchelyomorpha* Kahl. Conical, compressed; posterior end broadly rounded; anterior portion narrow; cilia on ring-furrows; anterior half with unretractile short tentacles; cytostome not noted; macronucleus with a central endosome surrounded by spherules; contractile vacuole terminal, large.

*E. vermicularis* (Smith) (Fig. 302, j). Body 30–45 μ; fresh and brackish water.

Family 6 *Holophryidae* Schouteden

Genus *Holophrya* Ehrenberg. Oval, globose or ellipsoidal; ciliation uniform; sometimes longer cilia at the anterior or posterior region; systostome circular, simple, without any ciliary ring around it; cytopharynx with or without trichites or trichocysts; fresh or salt water. Numerous species.

*H. simplex* Schewiakoff (Fig. 304, a). Ellipsoidal; 18–20 ciliary rows; cilia uniformly long; cytostome small; cytopharynx without trichocysts or trichites; contractile vacuole and cytopyge posterior; macronucleus large, round; 34 μ by 18 μ; fresh water.

Genus *Lagynophrya* Kahl. Resembles *Holophrya*; small elongate ovoid to short cylindrical; one side convex, the other more or less flattened; cytopharynx terminates anteriorly in a small cone-like process which may or may not be distinct; stagnant fresh or salt water. Several species.

*L. mutans* K. (Fig. 304, b). Body plastic; oval to cylindrical; colorless; narrowly striated; oval cone hemispherical without any trichocysts; body about 90 μ long, when contracted about 65 μ in diameter; among decaying leaves in fresh water.

Genus *Ichthyophthirius* Fouquet. Body oval; ciliation uniform; pellicle longitudinally striated; cytostome at anterior end, with a short cytopharynx with cilia; horseshoe-shaped macronucleus;
micronucleus adhering to macronucleus; macronucleus undergoes reorganization by discarding small chromatin masses (Haas, 1934); no division within the host body; multiplication within cyst which is formed after dropping off the fish skin and in which numerous (up to 1000) ciliated bodies (30–45μ in diameter) are produced; conjugation has been reported; parasitic in the integument of freshwater and marine fishes; in aquarium, host fish may suffer death; widely distributed.

*I. multifiliis* F. (Fig. 303). 100–1000μ long; ovoid; produces pustules in the epidermis or gills; cytostome is large, 30–40μ in diameter.
Pearson (1932) and Kudo (1934) reported extensive infections in large open ponds in Indiana and Illinois and Butcher (1941, 1943) noted infections in many yearling trout in hatcheries in 1939 and 1940. MacLennan (1935, 1935a, 1937, 1942) observed that the grown trophozoites leave the host epithelium and encyst on the bottom of aquarium; the cytostome is absorbed; the body protoplasm divides into 100–1000 small spherical ciliated cells, 18–22μ in diameter, which presently metamorphose into elongated forms, measuring about 40μ to 10μ. These young ciliates break through the cyst wall and seek new host fish by active swimming. The young ciliates are able to attack the fish integument for at least 96 hours, though their infectivity decreases markedly after 48 hours.

Sikama (1938) observed a similar organism on 44 species of marine fishes. This ciliate was somewhat smaller in dimensions, measuring up to 452μ by 360μ, and possessed a macronucleus typically constricted into four beads. Fibrillar structures (ten Kate, 1927).

Genus Bursella Schmidt. Oval; anterior end broadly and obliquely truncate where a large ciliated groove-like pit occurs; ridges of pit contractile; cilia short; macronucleus, spherical to ellipsoidal; several micronuclei; endoplasm reticulated; with symbiotic algae; ectoplasm with trichocysts; fresh water.

*B. spumosa* S. 240–560μ long; freshwater pond.

Genus Spasmostoma Kahl. Somewhat similar to *Holophrya*; cytostome with flaps which beat alternately; ciliation uniform.

*S. viride* K. (Fig. 304, c). Spherical or oval; always with green food vacuoles containing Euglena and allied flagellates; cytostome at anterior end; cytopharynx with trichocysts, which are extensible at the time when food is taken in; cilia on about 20 rows, near cytostome somewhat longer; macronucleus round; body 50–75μ long; sapropelic.

Genus Urotricha Claraparède and Lachmann (*Balanitozoon* Stokes). Body oval to ellipsoidal or conical; with 1 or more longer caudal cilia; ciliation uniform, except in posterior region which may be without cilia; cytostome at or near anterior end, surrounded by ring of heavier cilia; contractile vacuole, posterior; macronucleus spherical; fresh water.

*U. agilis* (Stokes) (Fig. 304, d). Body small; about 15–20μ long; swimming as well as leaping movement; standing fresh water with sphagnum.

*U. farcta* C. and L. (Fig. 304, e). Body 20–30μ long; fresh water. Kahl considers *U. parvula* Penard and *Balanitozoon gyrans* Stokes are identical with this species.
Fig. 304. a, Holophrya simplex, ×800 (Roux); b, Lagynophrya mutans, ×380 (Kahl); c, Spasmostoma viride, ×330 (Kahl); d, Urot richa agilis, ×530 (Stokes); e, U. farcta, ×470 (Lieberkühn); f, g, Plagiocampa marina (Noland) (f, ×400; g, anterior end, ×670); h, Chilophrya utahensis, ×840 (Pack); i, C. labiata, ×500 (Edmondson); j, Platyophrya lata, ×280 (Kahl); k, Stephanopogon colpoda, non-ciliate side, ×500 (Kahl); l, Prorodon discolor, ×330 (Biitschli); m, Pseudoprorodon farctus, ×270 (Roux); n, o, Coelosomides marina, ×245 (Fauré-Fremiet) (n, silver-imregnated surface view; o, optical section); p, q, Placus socialis, ×530 (Noland) (p, anterior end view).
Genus *Plagiocampa* Schewiakoff. Ovoid, spindle-form or cylindrical; slightly asymmetrical; cytostome at anterior end in a slit; right ridge thickened and lip-like, with about 8 long cilia; with or without long caudal cilium; fresh or salt water. Several species.

*P. marina* Kahl (Fig. 304, f, g). Cylindrical; oval macronucleus central; contractile vacuole terminal; a caudal cilium; 55–90μ long; salt water; Florida (Noland).

Genus *Chilophrya* Kahl. Ovoid or ellipsoid; cytostome at anterior end, surrounded by protrusible rods; on one side there is a lip-like ectoplasmic projection; fresh or salt water.

*C. (Prorodon) utahensis* (Pack) (Fig. 304, h). Body ellipsoid, somewhat asymmetrical; comparatively small number of furrows; ciliation uniform; a finger-like process in front of cytostome; macronucleus small, central; contractile vacuole terminal; endoplasm with zoochlorellae; encystment common; cysts highly sensitive to light; 50μ long; Great Salt Lake, Utah (Pack).

*C. (Urotricha) labiata* (Edmondson) (Fig. 304, i). Body ovoid; a lip-like process in front of cytostome; macronucleus oblong, central; contractile vacuole terminal; 30μ long; fresh water.

Genus *Platyophrya* Kahl. Compressed; flask-like or elongate ovoid; asymmetrical; dorsal surface convex, ventral surface flat or partly concave; spiral striation; position and direction of cytostome variable; macronucleus round; contractile vacuole terminal; fresh water.

*P. lata* K. (Fig. 304, j). Highly compressed; colorless; many striae; on left edge of cytostome 5–6 cirrus-like projections and on right edge many short bristles; 105μ long; fresh water with sphagnum.

Genus *Stephanopogon* Entz. Somewhat resembles *Platyophrya*; compressed; cytostome at anterior extremity which is drawn out; cytostome surrounded by lobed membranous structures; salt water.

*S. colpoda* E. (Fig. 304, k). Longitudinal striae on ‘neck’ 4–8 in number; 2 contractile vacuoles; 50–70μ long; creeping movement; salt water among algae.

Genus *Prorodon* Ehrenberg (*Rhagadostoma* Kahl). Ovoid to cylindrical; ciliation uniform, with sometimes longer caudal cilia; oral basket made up of double trichites which end deep in ectoplasm, oval in cross-section; contractile vacuole terminal; macronucleus massive, spherical or oval; fresh or salt water. Numerous species.

*P. discolor* (E.) (Fig. 304, l). Ovoidal; 45–55 ciliary rows; macronucleus ellipsoid; micronucleus hemispherical; contractile vacuole terminal; 100–130μ long; fresh water; Kahl (1930) states that it occurs also in brackish water containing 2.5 per cent salt; sapropelic
form in salt water is said to possess often long caudal cilia.

*P. griseus* Claparède and Lachmann. Oblong; 165–200μ long; fresh water.

Genus *Pseudoprorodon* Blochmann. Similar to *Prorodon*; usually flattened; one side convex, the other concave; ectoplasm conspicuously alveolated; trichocysts grouped; 1 or more contractile vacuoles posterior-lateral or distributed, with many pores; macronucleus elongate; cytopharynx with trichites; fresh or salt water.

*P. farctus* (Claparède and Lachmann) (Figs. 304, m). Ellipsoid; cytostome surrounded by long trichites; contractile vacuole posterior, with secondary vacuoles; macronucleus elongate; body 150–200μ long; fresh water.

Genus *Coelosomides* Anigstein (*Coelosoma A*.). General appearance similar to *Prorodon* and *Holophrya*; body cylindrical; ciliation uniform; at anterior end, a conspicuous ciliated vestibule runs down deep; mouth and cytopharynx; endoplasm vacuolated; a macronucleus and a micronucleus; marine.

*C. marina* A. (Fig. 304, n, o). About 200μ long; central endoplasm highly vacuolated; periphery finely reticulated; macronucleus elongate; micronucleus compact (Anigstein, 1911; Fauré-Fremiet, 1950).

Genus *Placus* Cohn (*Spathidiopis* Fabre-Domergue; *Thoraco-phrya* Kahl). Body small; ellipsoid or ovoid; somewhat compressed; pellicle with conspicuous spiral furrows; cytostome a narrow slit at anterior extremity; with strong cilia on right margin of slit; cytopyge a long narrow slit with cilia on both sides; macronucleus ellipsoid to sausage-form; contractile vacuole posterior; salt, brackish or fresh water.

*P. socialis* (Fabre-Domergue) (Fig. 304, p, q). 40–50μ by 28–32μ, about 22μ thick; salt water; Florida (Noland, 1937).

Genus *Lacrymaria* Ehrenberg. Polymorphic; cylindrical, spindle-or flask-shaped; with a long contractile proboscis; cytostome round; ciliary rows meridional or spiral to right; near cytostome a ring-like constriction with a circle of longer cilia; cytopharynx usually distinct; contractile vacuole terminal; fresh or salt water. Numerous species.

*L. olor* (Müller) (Fig. 305, a). Elongate; highly contractile; 2 macronuclei; 2 contractile vacuoles; extended forms 400–500μ up to 1.2 mm. long; when dividing, long neck is formed sidewise so that it appears as oblique division (Penard); fresh and salt water.

*L. lagenula* Claparède and Lachmann (Fig. 305, b). Body flask-shape; neck highly extensible; striation distinct, spiral when contracted; macronucleus short sausage-like or horseshoe-shape; endo-
plasm granulated; body 70\(\mu\) long, up to 150\(\mu\) (Kahl); salt water.

L. coronata C. and L. (Fig. 305, c). Large; neck extensible; body form variable, but usually with bluntly rounded posterior end; endoplasm appears dark; striae spiral; 85–100\(\mu\) long; salt and brackish water.

Genus Enchelys Hill. Flask-shape; anterior end obliquely truncate; cytostome slit-like, rarely round; fresh or salt water. Several species (Fauré-Fremiet, 1944).

E. curvilata (Smith) (Fig. 305, d). Elongate ovoid; posterior end rounded; longitudinal striation; macronucleus band-form; contractile vacuole terminal; endoplasm yellowish, granulated; about 150\(\mu\) long; fresh water among algae.

Genus Crobylura André. Body when extended spindle-form, with truncate ends; when contracted, thimble-form; cilia short and thick; several long caudal cilia; slit-like cytostome at anterior end; no apparent cytopharynx; macronucleus irregularly rounded, hard to stain; micronucleus not observed; contractile vacuole latero-posterior; fresh water. One species.

C. pelagica A. (Fig. 305, e). Body 65–95\(\mu\) long; in freshwater plankton.

Genus Microregma Kahl. Small, ovoid; dorsal side convex; ventral side flat; with a small slit-like cytostome near anterior end; with or without caudal bristle; fresh or salt water.

M. (Enchelys) auduboni (Smith) (Fig. 305, f). Body plastic; coarsely ciliated; caudal bristle thin; cytostome at anterior end, surrounded by longer cilia; cytopharynx small with trichocysts; round macronucleus central; contractile vacuole near posterior end; 40–55\(\mu\); fresh water.

Genus Chaenea Quennerstedt. Elongate; anterior end drawn out into a narrow truncated ‘head’; but without any ring furrow; ‘head’ spirally or longitudinally furrowed; often with longer cilia directed anteriorly; cytostome terminal, not lateral; cytopharynx with trichocysts; body striation meridional, or slightly right spiral; macronucleus often distributed; fresh or salt water.

C. limicola Lauterborn (Fig. 305, g). Anterior half of body broad; posterior end drawn out into a point; contractile; cytopharynx with trichocysts; many trichocysts in endoplasm; contractile vacuoles in a row; 130–150\(\mu\) long; stagnant fresh water.

Genus Pithothorax Kahl. Slender, barrel-shaped; with firm pellicle; a fairly long caudal bristle, contractile vacuole in posterior half; ciliation coarse and not over entire body surface; resembles Coleps; fresh water.
P. ovatus K. (Fig. 305, h). Caudal bristle breaks off easily; body 30μ long; fresh water among decaying vegetation.

Genus Rhopalophrya Kahl. Cylindrical; furrows widely separated; slightly asymmetrical; curved ventrally; dorsal surface convex; ventral surface flat or slightly concave; anterior end with ‘neck’; 2 spherical macronuclei; fresh or salt water; sapropelic.

Genus Enchelyodon Claparède and Lachmann. Elongated; cy-
lindrical, ovoid or flask-shaped; some with head-like prolongation; cytopharynx with trichites; cilia long at anterior end; fresh or salt water. Several species.

*E. californicus* Kahl. 120–130μ long; elongate ovoid to nearly cylindrical; not distinctly flattened; macronucleus horseshoe-like, with a large micronucleus; in mosses; California.

Genus *Trachelophyllum* Claparède and Lachmann. Elongate; flattened; flexible, ribbon-like; anterior end neck-like and tip truncate; cytopharynx narrow, round in cross-section, with trichocysts; ciliary rows widely apart; 2 macronuclei, each with a micronucleus; contractile vacuole terminal; fresh or salt water. Several species.

*T. clavatum* Stokes (Fig. 305, i). About 200μ long; fresh water.

Genus *Ileonema* Stokes (*Monomastix* Roux). Body flattened; flask-shaped; somewhat similar to *Trachelophyllum*, but there is a remarkable flagellum-like process extending from anterior end; cytopharynx with trichocysts; fresh water.

*I. dispar* S. (Fig. 306, b). Highly contractile; anterior flagellum half body length, whose basal portion spirally furrowed; cytostome

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**Fig. 306.** a, *Rhopalophrya salina*, ×1040 (Kirby); b, *Ileonema dispar*, ×190 (Stokes); c, *I. ciliata*, ×800 (Roux); d, e, *Trachelocerca phaenicopterus* (Kahl) (d, whole organism, ×120; e, anterior end, ×260); f, g, *T. subviridis* (Noland) (f, whole organism, ×155; g, the nucleus, ×480).
at base of the flagellum; cytopharynx spindle-form with trichites; 2 contractile vacuoles and cytopyge posterior; ovoid macronucleus; movement slow creeping; about 120μ long; fresh water among algae.

*I. ciliata* (Roux) (Fig. 306, c). 75μ by 14μ; fresh water.

Genus *Trachelocerca* Ehrenberg. Elongate, vermiform or flask-shaped; more or less extensible, with drawn-out anterior end; without any ring-furrow which marks the ‘head’ of Lacrymaria, and when contracted pellicular striae not spiral and no neck as is the case with Chaenea; salt water. Many species.

*T. phoenicopterus* Cohn (Fig. 306, d, e). Elongate; extensible and contractile; neck and tail distinct when contracted; cytostome at anterior end, surrounded by a ridge containing indistinctly visible short trichocysts, cytopharynx with trichocysts; macronuclei made up of 4 radially arranged endosomes suspended in the nucleoplasm (Gruber, Kahl); micronucleus difficult to make out; contractile vacuoles apparently in a row, rarely seen; salt water; Woods Hole (Calkins).

*T. subviridis* Sauerbrey (Fig. 306, f, g). Highly extensible and contractile; nucleus contains peculiar crystal-like bodies; size variable; when extended 320-480μ long; salt water. Noland (1937) observed the organism in a salt spring in Florida.

**Family 7 Büttschliidae Poche**

This family includes species that inhabit the alimentary canal of mammals; circular cytostome at anterior end, cytopyge usually located at posterior end; ciliation uniform or in a few zones; with refractile concrement vacuole (Fig. 31, d) in anterior portion; one or more contractile vacuoles.

Genus *Büttschia* Schuberg. Ovoid, anterior end truncate, posterior end rounded; cytostome at anterior end, surrounded by long cilia; thick ectoplasm at anterior end; macronucleus spherical micronucleus(?); concretion vacuole; ciliation uniform; in stomach of cattle.

*B. parva* S. (Fig. 307, a). 30–50μ by 20–30μ Conjugation (Dogiel, 1928).

Genus *Blepharoprosthium* Bundle. Pyriform, anterior half contractile, ciliated; caudal cilia; macronucleus reniform; in the caecum and colon of horse.

*B. pircum* B. (Fig. 307, b). 54–86μ by 34–52μ (Hsiung, 1930a).

Genus *Didesmis* Fiorentini. Anterior end neck-like, with large cytostome; anterior and posterior ends ciliated; macronucleus ellipsoidal; in the caecum and colon of horse. Species (Hsiung, 1930a).
D. quadrata F. (Fig. 307, c). 50–90\(\mu\) by 33–68\(\mu\); with a deep dorsal groove.

Genus **Blepharosphaera** Bundle. Spherical or ellipsoidal; ciliation uniform except in posterior region; caudal cilia; in the caecum and colon of horse.

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Fig. 307. a, *Büttschlia parva*, \(\times 670\) (Schuberg); b, *Blepharoprosthium pireum*, \(\times 470\) (Hsiung); c, *Didesmis quadrata*, \(\times 270\) (Hsiung); d, *Blepharosphaera intestinalis*, \(\times 600\) (Hsiung); e, *Blepharoconus cervicalis*, \(\times 360\) (Hsiung); f, *Bundleia postciliata*, \(\times 530\) (Hsiung); g, *Blepharozoum zonatum*, \(\times 200\) (Gassovsky).

**B. intestinalis** B. (Fig. 307, d). 38–74\(\mu\) in diameter (Hsiung, 1930a).

Genus **Blepharoconus** Gassovsky. Oval; small cytostome; cilia on anterior 1/3–1/2; caudal cilia; macronucleus ovoid; 3 contractile vacuoles; cytopharynx with rods; in the colon of horse.

**B. cervicalis** Hsiung (Fig. 307, e). 56–83\(\mu\) by 48–70\(\mu\); Iowa (Hsiung, 1930, 1930a).

Genus **Bundleia** da Cunha and Muniz. Ellipsoid; cytostome small; cilia at anterior and posterior ends, posterior cilia much less numerous; in the caecum and colon of horse.

**B. postciliata** (Bundle) (Fig. 307, f). 30–56\(\mu\) by 17–32\(\mu\) (Hsiung, 1930a).
Genus Polymorpha Dogiel. Flask-shaped; ciliation on anterior region, a few caudal cilia; macronucleus disc-shaped; contractile vacuole terminal; in the caecum and colon of horse.

P. ampulla D. (Fig. 308, a). 22–36 μ by 13–21 μ (Hsiung, 1930a).

Genus Holophryoides Gassovsky. Oval, with comparatively large cytostome at anterior end; ciliation uniform; macronucleus small, ellipsoid; contractile vacuole subterminal; in the colon and caecum of horse.

H. ovalis (Fiorentini) (Fig. 308, b). 95–140 μ by 65–90 μ.

Genus Blepharozoum Gassovsky. Ellipsoid, with attenuated posterior end; ciliation uniform; cytostome near anterior tip; 2 contractile vacuoles; macronucleus small, reniform; in caecum of horse.

B. zonatum G. (Fig. 307, g). 230–245 μ by 115–122 μ (Hsiung, 1930a).
Genus Prorodonopsis Gassovsky. Pyriform; ciliation uniform; 3 contractile vacuoles; macronucleus sausage-shaped; in the colon of horse.

P. coli G. (Fig. 308, c). 55–67 µ by 38–45 µ (Hsiung, 1930a).

Genus Paraisotrichopsis Gassovsky. Body uniformly ciliated; spiral groove from anterior to posterior end; in the caecum of horse.

P. composita G. (Fig. 308, d). 43–56 µ by 31–40 µ (Hsiung, 1930a).

Genus Sulcoarcus Hsiung. Ovoid, compressed; a short spiral groove at anterior end; cytostome at ventral end of the groove; cytopyge terminal; concretion vacuole mid-ventral, contractile vacuole posterior to it; cilia on groove, posterior end and mid-ventral region (Hsiung, 1935).

S. pellucidulus H. (Fig. 308, e). 33–56 µ by 30–40 µ; in faeces of mule.

Genus Alloiozona Hsiung. Cilia in 3 (anterior, equatorial and posterior) zones; in the caecum and colon of horse (Hsiung, 1930, 1930a).

A. trizona H. (Fig. 308, f). 50–90 µ by 30–60 µ.

Genus Ampullacula Hsiung. Flask-shaped; posterior half bearing fine, short cilia; neck with longer cilia; in the caecum of horse.

A. ampulla (Fiorentini). About 110 µ by 40 µ (Hsiung, 1930a).

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Chapter 33

Order 1 Holotricha Stein (continued)

Suborder 2 Gymnostomata Bütschli (continued)

Tribe 2 Pleurostomata Schewiakoff

Cytostome on convex ventral surface.

Cytostome a long slit.........................Family 1 Amphileptidae
Cytostome round, at base of trichocyst-bearing neck..................

.................Family 2 Tracheliidae (p. 725)
Cytostome on concave ventral side........Family 3 Loxodidae (p. 727)

Family 1 Amphileptidae Schouteden

Genus Amphileptus Ehrenberg. Flask-shaped; somewhat compressed; ciliation uniform and complete; slit-like cytostome not reaching the middle of body, without trichocyst-borders; many contractile vacuoles; 2 or more macronuclei; fresh or salt water.

A. claparedei Stein (A. meleagris Claparède and Lachmann) (Fig. 309, a). Slightly flattened; broadly flask-shaped; with bluntly pointed posterior and neck-like anterior end; cytostome about 2/5 from ventral margin; trichocysts indistinct; dorsal ciliary rows also not distinct; contractile vacuoles irregularly distributed; 120–150μ long; fresh and salt water, on stalks of Zoothamnium, Carchesium, Epistylis, etc.

A. branchiarum Wenrich (Fig. 309, b). On the integument and gills of frog tadpoles; swimming individuals killed with iodine, 100–135μ by 40–60/x (Wenrich, 1924).

Genus Lionotus Wrzesniowski (Hemiophrys W.). Flask-shape; elongate, flattened; anterior region neck-like; cilia only on right side; without trichocyst-borders; cytostome with trichocysts; 1 (terminal) or many (in 1–2 rows) contractile vacuoles; 2 macronuclei; 1 micronucleus; fresh or salt water.

L. fasciola (Ehrenberg) (Fig. 309, c). Elongate flask in form; hyaline; with flattened neck and tail, both of which are moderately contractile; posterior end bluntly rounded; without trichocysts; neck stout, bent toward the dorsal side; cytostome a long slit; contractile vacuole posterior; 2 spherical macronuclei between which a micronucleus is located; 100μ long; fresh water and probably also in salt water.

Genus Loxophyllum Dujardin (Opisthodon Stein). Generally similar to Lionotus in appearance; but ventral side with a hyaline border, reaching posterior end and bearing trichocysts; dorsal side with
either similar trichocyst-border or with trichocyst-warts; macronucleus a single mass or moniliform; contractile vacuole, one to many; fresh or salt water. Many species.

*L. meleagris* D. (Fig. 309, d). Form and size highly variable; flask-shape to broad leaf-like; broad ventral seam with trichocysts and often undulating; dorsal seam narrow and near its edge, groups of trichocysts in wart-like protuberances; macronucleus moniliform; micronuclei, as many as the beads of the macronucleus (Penard, 1922); contractile vacuole terminal, with a long canal; 300–400\(\mu\) long, up to 700\(\mu\) (Penard); feeds mainly on rotifers; fresh water.

Fig. 309. a, *Amphileptus clapareei*, ×370 (Roux); b, *A. branchiarum*, flattened, ×490 (Wenrich); c, *Lionotus fasciola*, ×540 (Kahl); d, *Loxophyllum meleagris*, ×120 (Penard); e, *L. setigerum*, ×570 (Sauerbrey); f, *Bryophyllum vorax*, ×360 (Stokes); g, h, *Centrophorella fasciolatum* (g, ×50; h, ×110) (Noland).
L. setigerum Quennerstedt (Fig. 309, e). 100–350μ long; average 150μ by 60μ; form variable; 1–4 macronuclei; several contractile vacuoles in a row; salt and brackish water. Morphology (Sauerbrey, 1928).

Genus Bryophyllum Kahl. Similar to Loxophyllum; but uniformly ciliated on both broad surfaces; ventral ridge with closely arranged trichocysts, extends to the posterior extremity and ends there or may continue on to the opposite side for some distance; macronucleus ovoid to coiled bandform; in mosses. Species (Gelei, 1933).

B. vorax (Stokes) (Fig. 309, f). Elongate; trichocyst-bearing ventral ridge turns up a little on dorsal side; contractile vacuole posterior; macronucleus oval; 130μ long; in fresh water among sphagnum.

Genus Centrophorella Kahl (Kentrophoros Sauerbrey). Extremely elongate, nematode-like; anterior end greatly attenuated; posterior end pointed; body surface longitudinally striated; ciliation uniform; 1–3 macronuclei; numerous contractile vacuoles in 2 rows; cytostome not observed.

C. fasciolatum (S.) (Fig. 309, g, h). About 270μ by 38μ. Noland (1937) observed 2 specimens in sediment taken from sandy bottom in Florida; contracted 650μ long; extended 1 mm. long.

C. lanceolata Fauré-Fremiet. Ribbon-like; 460–520μ by 40μ; ventral side ciliated; dorsal side covered with dark sulphur bacteria (Caulobacteria), except the extremities; five to six spherical micronuclei, about 4μ in diameter; on sandy flat of Cape Cod (Fauré-Fremiet, 1951).

Family 2 Tracheliidae Kent

Genus Trachelius Schrank. Oval to spherical; anterior end drawn out into a relatively short finger-like process or a snout; posterior end rounded; round cytostome at base of neck; cytopharynx with trichites; contractile vacuoles many; macronucleus simple or band-form; fresh water.

T. ovum Ehrenberg (Fig. 310, a). Spheroidal to ellipsoid; right side flattened and with a longitudinal groove; left side convex; proboscis about 1/4–1/2 the body length; cilia short and closely set; numerous contractile vacuoles; macronucleus short sausage-form, often divided into spherules; endoplasm penetrated by branching cytoplasmic skeins or bands and often with numerous small brown excretion granules; 200–400μ long; fresh water.

Genus Dileptus Dujardin. Elongate; snout or neck-like prolongation conspicuous; somewhat bent dorsally; along convex ventral side
of neck many rows of trichocysts; a row of strong cilia; dorsal surface with 3 rows of short bristles; cytostome surrounded by a ring; cytospharynx with long trichocysts; posterior end drawn out into a tail; contractile vacuoles, 2 or more; body ciliation uniform; macronu-

Fig. 310. a, Trachelius ovum, ×130 (Roux); b, Dileptus americanus, ×250 (Kahl); c, D. anser, ×310 (Hayes); d, Paradileptus conicus, ×340 (Wenrich); e, P. robustus, ×340 (Wenrich); f, Branchioecetes gammari ×200 (Penard); g, Lozodes vorax, ×190 (Stokes); h, L. magnus, ×80 (Kahl); i, j, Remanella rugosa (i, dorsal side, ×130; j, anterior part showing the endoskeleton) (Kahl).
cleus bandform, moniliform or divided into numerous independent bodies; fresh or salt water. Many species.

D. americanus Kahl (Fig. 310, b). Proboscis bent dorsally sickle-like; macronucleus made up of 2 sausage-shaped or often horseshoe-shaped parts; 2 contractile vacuoles on dorsal side; 200μ long; in mosses.

D. anser (Müller) (Fig. 310, c). Proboscis slightly flattened; macronucleus divided into 100 or more discoid bodies; 16 or more vesicular micronuclei (Jones, 1951); contractile vacuoles in a row on the aboral surface, with 2–3 in proboscis; 250–400μ, sometimes up to 600μ long; in fresh water. Culture, encystment and excystment (Jones, 1951).

Genus Paradileptus Wenrich (Tentaculifera Sokoloff). Body broader at the level of cytostome; with a wide peristomal field that bears the cytostome and is surrounded for 2/3–3/4 its circumference by a raised rim which is continuous anteriorly with the spirally wound proboscis; trichocyst-zone traversing the rim and anterior edge of proboscis; contractile vacuoles small, numerous, distributed; macronucleus segmented; fresh water (Wenrich, 1929a).

P. conicus W. (Fig. 310, d). 100–200μ by 50–100μ.

P. robustus W. (Fig. 310, e). 180–450μ long.

P. estensis Canella. 600–800μ long; feeds on rotifers (Canella, 1951).

Genus Branchioecetes Kahl. Preoral part somewhat like that of Amphileptus, and bent dorsally; ventral side of neck with 2 rows of trichocysts; cytostome at posterior end of neck; cytopharynx with trichocysts; ectocommensals on Asellus or Gammarus.

B. gammari (Penard) (Fig. 310, f). 130–200μ long; on Gammarus.

Family 3 Loxodidae Bütschli

Genus Loxodes Ehrenberg. Lancet-like; strongly compressed; anterior end curved ventrally, and usually pointed; right side slightly convex; uniform ciliation on about 12 longitudinal rows; ectoplasm appears brownish, because of closely arranged brownish protrichocysts; endoplasm reticulated; 2 or more vesicular macronuclei; one or more micronuclei; 5–25 Müller’s vesicles (p. 87; Fig. 31, a, b) in dorsal region; fresh water.

L. vorax Stokes (Fig. 310, g). 125–140μ long; yellowish brown, a row of slightly longer cilia; sapropelic in standing fresh water.

L. magnus S. (Fig. 310, h). Extended about 700μ long; dark brown; 12–20 or more Müller’s vesicles in a row along dorsal border; standing pond water.

Genus Remanella Kahl. Similar to Loxodes in general appearance;
but with endoskeleton consisting of 12–20μ long spindle-form needles lying below broad ciliated surface in 3–5 longitudinal strings connected with fibrils; Müller's vesicles (Fig. 31, c) in some, said to be different from those of Loxodes (Kahl); sandy shore of sea.

*R. rugosa* K. (Fig. 310, i, j). 200–300μ long.

**Tribe 3 Hypostomata** Schewiakoff

Without furrow; free-living; conspicuous oral or pharyngeal basket

Ciliation complete; dorsal cilia usually less dense than those on ventral surface.............................. Family 1 Nassulidae

Ciliation incomplete; dorsal surface without cilia or with a few sensory bristles

Posterior ventral surface with a style. Family 2 Dysteriidae (p. 730)
Without a style......................... Family 3 Chlamydodontidae (p. 731)
Furrow from anterior end of cytostome; parasitic............................... Family 4 Pycnothricidae (p. 733)

**Family 1 Nassulidae** Schouteden

Genus *Nassula* Ehrenberg. Oval to elongate; ventral surface flat, dorsal surface convex; usually brightly colored, due to food material; cytostome 1/3–1/4 from anterior end; body often bent to left near cytostome; opening of oral basket deep, in a vestibule with a membrane; macronucleus spherical or ovoid, central; a single micronucleus; contractile vacuole large, with accessory vacuoles and opens ventrally through a tubule-pore; fresh or salt water. Many species.

*N. aurea* E. (Fig. 311, a). 200–250μ long; fresh and brackish water (Kahl).

Genus *Paranassula* Kahl. Similar in general appearance to *Nassula*; but with preoral and dorsal suture line; longer caudal cilia on dorsal suture; pharyngeal basket not funnel-like, with 16–18 trichites; about 75 ciliary rows; trichocysts especially in anterior region.

*P. microstoma* (Claparède and Lachmann) (Fig. 311, b). Pellicle roughened by a criss-cross of longitudinal and circular furrows; macronucleus elongate oval, posterior; contractile vacuole near middle and right-dorsal; about 80–95μ long; salt water; Florida (Noland).

Genus *Cyclogramma* Perty. Somewhat resembling *Nassula*; but conspicuous oral basket in pyriform depression and opens toward left on ventral surface; depression with a short row of small membranes at its anterior edge; trichocysts usually better developed than in Nassula; fresh water.

*C. trichocystis* (Stokes) (Fig. 311, c). Body colorless or slightly rose-colored; trichocysts thick and obliquely arranged; one contractile vacuole; usually full of blue-green food vacuoles; actively motile; about 60μ long; in fresh water among algae.
Fig. 311. a, *Nassula aurea*, $\times 190$ (Schewiakoff); b, *Paranassula microstoma*, $\times 400$ (Noland); c, *Cyclogramma trichocystis*, $\times 510$ (Stokes); d, *Chilodontopsis vorax*, $\times 200$ (Stokes); e, *Eucamplocerca longa*, $\times 320$ (da Cunha); f, *Orthodon hamatus*, $\times 160$ (Entz); g, *Dysteria calkinsi*, $\times 540$ (Calkins); h, *Trochilia palustris*, $\times 1070$ (Roux); i, *Trochilioides recta*, $\times 740$ (Kahl); j, *Hartmannula entzi*, $\times 220$ (Entz); k, *Chlamydodon mnemosyne*, $\times 520$ (MacDougall); l, *Phascolodon vorticella*, $\times 340$ (Stein).

Genus *Chilodontopsis* Blochmann. Elongate ellipsoid; colorless; ventral surface flattened, dorsal surface slightly convex; both sides ciliated; oral basket without vestibule; cytostome with a membranous ring; usually with a postoral ciliary furrow; fresh water.

*C. vorax* (Stokes) (Fig. 311, d). Elongate ellipsoid; anterior re-
region slightly curved to left; snout fairly distinct; oral basket with about 16 rods; several contractile vacuoles distributed, a large one terminal; macronucleus large, lenticular, granulated; with a closely attached micronucleus; 50–160μ long; fresh water.

Genus *Eucamptocerca* da Cunha. Elongate; posterior part drawn out into a caudal prolongation; dorso-ventrally flattened; ciliation on both sides; round cytostome with oral basket in anterior ventral surface. One species.

*E. longa* da C. (Fig. 311, e). 300μ by 25μ; macronucleus ovoid, with a micronucleus; contractile vacuole(?); in brackish water (salt content 3 per cent); Brazil.

Genus *Orthodon* Gruber. Oval; contractile; colorless; much flattened; anterior region curved toward left; striation on both dorsal and ventral sides; cytostome toward right border; oral basket long; macronucleus oval; contractile vacuole terminal; fresh or salt water.

*O. hamatus* G. (Fig. 311, f). Extended 200–260μ long, contracted 90–150μ long; flask-shaped; oral basket with 16 trichites; salt water.

Family 2 *Dysteriidae* Kent

Genus *Dysteria* Huxley (*Ervilia* Dujardin; *Iduna*, *Aegyria* Claparède and Lachmann; *Cypridium* Kent). Ovate, dorsal surface convex, ventral surface flat or concave; left ventral side with nonciliated ventral plate; postoral ciliation is continuation of preoral to right of cytostome and parallel to right margin; cytostome in a furrow near right side; posterior style or spine conspicuous; macronucleus spheroid or ovoid, central; with a micronucleus; usually 2 contractile vacuoles; fresh or salt water. Numerous species.

*D. calcinsi* Kahl (*D. lanceolata* Calkins) (Fig. 311, g). About 45μ by 27μ; salt water; Woods Hole.

Genus *Trochilia* Dujardin. Similar to *Dysteria*; but ciliation on the ventral side in an arched zone; fresh or salt water. Several species.

*T. palustris* Stein (Fig. 311, h). 25μ long; fresh water.

Genus *Trochilioides* Kahl. Rounded at anterior end, narrowed posteriorly; right side more convex than left; cytostome anterior with cytopharynx and preoral membrane; conspicuous longitudinal bands on right half with longitudinal striae, becoming shorter toward left; fresh or salt water.

*T. recta* K. (Fig. 311, i). 40–50μ long; sapropelic in fresh and brackish water.

Genus *Hartmannula* Poche (*Onychodactylus* Entz). Ventral surface uniformly ciliated; cytopharynx with short rods; in salt water.

*H. entzi* Kahl (Fig. 311, j). 80–140μ long; salt water.
Family 3 Chlamydodontidae Claus

Genus Chlamydodon Ehrenberg. Ellipsoid, reniform, elongate triangular, etc.; cilia only on ventral surface, anterior cilia longer; cytostome elongate oval and covered with a membrane bearing a slit; oral basket made up of closely arranged rods with apical processes; along lateral margin, there is a characteristic striped band which is a canalicule of unknown function; fresh or salt water.

C. mnemosyne E. (Fig. 311, k). Ellipsoid or reniform; right side convex, left side concave; ventral side flat, dorsal side greatly convex; a band of trichites, 'railroad track,' parallel to body outline; oral basket with 8–10 rods; macronucleus oval and covered with a membrane bearing a slit; oral basket made up of closely arranged rods with apical processes; along lateral margin, there is a characteristic striped band which is a canalicule of unknown function; fresh or salt water. MacDougall (1928) observed it in the brackish water at Woods Hole and studied its neuromotor system.

Genus Phascolodon Stein. Ovoid; with broad anterior end and bluntly pointed posterior end; ventral side concave or flat, dorsal side convex; ciliated field on ventral surface narrowed laterally behind cytostome, forming V-shaped ciliated area (about 12 rows); cytostome ellipsoid with oral basket; macronucleus oval with a micronucleus; 2 contractile vacuoles; fresh water.

P. vorticella S. (Fig. 311, l). 80–110μ long; cytostome covered by a slit-bearing membrane; with 2 preoral membranes; macronucleus ovoid; fresh water.

Genus Cryptopharynx Kahl. Ellipsoid, anterior third bent to left; ventral surface flat, dorsal surface with hump; spiral interciliary furrows ridged; oval cytostome at anterior end; no cryptopharynx; dorsal hump yellowish, granulated with gelatinous cover; 2 macronuclei; 1 micronucleus; 2 contractile vacuoles, one posterior and the other toward left side at the bend of body. One species.

C. setigerus K. (Fig. 312, a, b). Elongate ellipsoid; anterior region bent to left; ventral surface flat, dorsal surface with a hump; about 15 ventral ciliary rows; 2 vesicular macronuclei and 1 micronucleus dorso-central; 33–96μ by 21–45μ (Kirby). Kirby (1934) found the organism in salt marsh pools (salinity 1.2–9.7 per cent) with purple bacteria; California.

Genus Chilodonella Strand (Chilodon Ehrenberg). Ovoid; dorso-ventrally flattened; dorsal surface convex, ventral surface flat; ventral surface with ciliary rows; anteriorly flattened dorsal surface with a cross-row of bristles; cytostome round; oral basket conspicuous, protrusible; macronucleus rounded; contractile vacuoles variable in number; fresh or salt water or ectocommensal on fish and amphipods. Many species.
C. cucullulus (Müller) (Chilodon steini Blochmann) (Figs. 53; 312, c–e). 19–20 ventral ciliary rows; oral basket with about 12 rods and with 3 preoral membranes; macronucleus oval, a characteristic concentric structure; micronucleus small; body 100–300μ long, most

often 130–150μ long; fresh and brackish water. Conjugation (Ivanić, 1933).

C. caudata (Stokes) (Fig. 312, f). About 42μ long; standing water.
C. fluvialis (S.) (Fig. 312, g). About 50μ long; fresh water.
C. uncinata (Ehrenberg) (Fig. 96). 50–90μ long; about 11 ventral
ciliary rows; some 7 dorsal bristles; widely distributed in various freshwater bodies; several varieties. Conjugation (MacDougall, 1935).

*C. cyprini* (Moroff) (Fig. 312, h). 50–70 μ by 30–40 μ; in the integument and gills of cyprinoid fishes; the organism, if freed from the host body, dies in 12–24 hours. Ciliation (Krascheninnikow, 1934).

*C. longipharynx* Kidder and Summers. 17–21 μ (average 19 μ) long; cytopharynx long, reaches posterior end; ectocommensal on amphipods, *Talorchestia longicornis* and *Orchestia palustris*; Woods Hole (Kidder and Summers, 1935).

*C. hyalina* K. and S. 40 μ (36–47 μ) long; ectocommensal on *Orchestia agilis*; Woods Hole.

*C. rotunda* K. and S. 29 μ (27–34 μ) long; ectocommensal on *Orchestia agilis*; Woods Hole.

Genus *Allosphaerium* Kidder and Summers. Oval; right side concave, left side more or less flat; body highly flattened; arched dorsal surface devoid of cilia; ventral surface slightly concave with 12–27 ciliary rows; right and left margin of ventral surface with a papillar fold; cytostome anterior-ventral, oval or irregular, surrounded by ridge on posterior border, extending to left margin; 3 groups of ciliary membranes extending out of cytostome; macronucleus oval, central or anterior; a micronucleus; 2 (or 1) contractile vacuoles; a refractile spherule regularly present in posterior portion of endoplasm; ectocommensal on the carapace and gills of amphipods.

*A. palustris* K. and S. (Fig. 312, i). 46–59 μ long; 27 ventral ciliary rows; on *Orchestia palustris* and *Talorchestia longicornis*; Woods Hole.

*A. sulcatum* K. and S. 24–32 μ long; 12 ciliary rows; on the carapace of *Orchestia agilis* and *O. palustris*; Woods Hole.

*A. granulosum* K. and S. 32–42 μ long; rotund; 17 ciliary rows; cytoplasm granulated; on carapace of *Orchestia agilis* and *O. palustris*; Woods Hole.

*A. caudatum* K. and S. Resembles *A. palustris*; 35–45 μ long; 14 ciliary rows; 1 contractile vacuole; ectoplasm at posterior end, drawn out into a shelf; on *Orchestia agilis*; Woods Hole.

*A. convexa* K. and S. 24–36 μ long; 17 ciliary rows; on the carapace and gill lamellae of *Talorchestia longicornis*; Woods Hole.

Family 4 *Pycnothricidae* Poche

Ciliation uniform; ectoplasm thick and conspicuous; a furrow or groove connects the cytostome with the anterior end; parasitic in the alimentary canal of mammals.
Genus *Pycnothrix* Schubotz. Large, elongate; with broadly rounded anterior and narrowed posterior end; somewhat flattened; short thick cilia throughout; ectoplasm thick; macronucleus spherical, in anterior 1/6; micronucleus(?); 2 longitudinal grooves, one beginning on each side near anterior end, united at the notched posterior end; a series of apertures in grooves considered as cytostomes; at posterior 1/3, an aperture gives rise to branching canals running through endoplasm, and is considered as excretory in function. One species.

*P. monocystoides* S. (Fig. 313, a). 300μ—2 mm. long; in the colon of *Procavia capensis* and *P. brucei*.

Genus *Nicollella* Chatton and Pérard. Elongate; a narrow groove extends from the anterior end to cytostome, located at the middle of

![Fig. 313. a, Pycnothrix monocystoides, x50; b, Nicollella ctenodactyli, x170; c, Collinella gundi, x170 (Chatton and Pérard); d, Buxtonella sulcata, x395 (Jameson); e, Taliaferria clarki, x500 (Hegner and Rees).](image-url)
body; bilobed posteriorly; contractile vacuole terminal; macronucleus ellipsoidal, anterior; a micronucleus; ectoplasm thick anteriorly; ciliation uniform (Chatton and Pérand, 1921). One species.

*N. ctenodactyli* C. and P. (Fig. 313, b). 70–550 μ by 40–150 μ; in the colon of *Ctenodactylus gundi*.

Genus **Collinella** Chatton and Pérand. More elongate than *Nicollella*; uniform ciliation; a groove extends from end to end; cytostome at posterior end of the groove; contractile vacuole terminal; macronucleus much elongated, central or posterior (Chatton and Pérand, 1921). One species.

*C. gundi* C. and P. (Fig. 313, c). 550–600 μ by 100 μ; in the colon of *Ctenodactylus gundi*.

Genus **Buxtonella** Jameson. Ovoid; a prominent curved groove bordered by two ridges from end to end; cytostome near anterior end; uniform ciliation; in the caecum of cattle (Jameson, 1926). One species.

*B. sulcata* J. (Fig. 313, d). 55–124 μ by 40–72 μ.

Genus **Taliaferria** Hegner and Rees. Body ovate; circular in cross-section; ectoplasm is two-layered and thick; ciliation uniform; cytostome anterior, subterminal; macronucleus and a closely attached micronucleus near center; two contractile vacuoles; cytopyge (Hegner and Rees, 1933). One species.

*T. clarki* H. and R. (Fig. 313, e). 83–146 μ by 42–83 μ; in the caecum and colon of the red spider monkey (*Ateles geoffroyi*).

**References**


Chapter 34

Order 1 Holotricha Stein (continued)

Suborder 3 Trichostomata Bütschli

With gelatinous lorica; swimming backward ........................................... Family 1 Marynidae (p. 738)

Without lorica

Compressed, armor-like pellicle; ciliation sparse, mainly on flat right side in 2–9 broken rows on semicircular or crescentic keel; cytostome on flattened ventral surface, with an obscure membrane. .................................................. Family 2 Trichopelmidae (p. 739)

Body form and ciliation otherwise

With a long caudal cilium; cilia in 3–4 spiral rows on anterior half, very small forms .................. Family 3 Trimyemidae (p. 739)

Without a caudal cilium; form and ciliation otherwise

With a spiral zone of special cilia, from cytostome to posterior end
Spiral zone extends from anterior right to posterior left........... Family 4 Spirozonidae (p. 740)
Spiral zone extends from anterior left to posterior right........... Family 5 Trichospiridae (p. 740)

Without a spiral zone of special cilia

Ciliated cross-furrow in anterior 1/5 on ventral surface, leads to cytostome........... Family 6 Plagiopylidae (p. 740)

Without ciliated cross-furrow

Cytostome in flat oval groove with heavily ciliated ridge in anterior 1/4............ Family 7 Clathrostomidae (p. 742)
Cytostome funnel-like, deeply situated
Cytostomal funnel with strong cilia; peristome from anterior left to posterior right................................................. Family 8 Parameciidae (p. 742)

Without such a peristome

Free-living; oral funnel deep; cilia at bottom and top.................. Family 9 Colpodidae (p. 745)

Endozoic

Commensal in vertebrates................................................................. Family 10 Entorhipidiidae (p. 748)

Parasitic in vertebrates

Ciliation uniform

With concrement vacuoles................................................................. Family 11 Paraisotrichidae (p. 750)

Without such vacuoles................................................................. Family 12 Isotrichidae (p. 751)

Ciliation not uniform

Cytostome occupies the entire anterior end; cilia only in anterior region................................................. Family 13 Cyathodiriidae (p. 752)
Cytostome not terminal; tufts of cilia above and below cytostome and in posterior region ....... Family 14 Blepharocororidae (p. 752)
Body with infraciliature, but cilia only in free-swimming stage; adult enclosed within a thick pellicle and attached to secretory hair of arthropods. Family 15 Conidophryidae (p. 753).

Family 1 Marynidae Poche

Genus Maryna Gruber. Peristome makes a complete circle, thus the cone is entirely separated from anterior edge of body; cytostome left ventral, elongate slit; ridge also with a slit; gelatinous lorica dichotomous.

*M. socialis* G. (Fig. 314, a, b). About 150μ long; in infusion made from long-dried mud.

Fig. 314. a, b, *Maryna socialis* (a, ×40; b, ×160) (Gruber); c, *Mycterothrix erlangeri*, ×310 (Kahl); d, *Trichopelma sphagnetorum*, ×570 (Kahl); e, f, *Pseudomicrothorax agilis* (e, ×340; f, ×670) (Kahl); g, *Drepanomonas dentata*, ×540 (Penard); h, *Microthorax simulans*, ×620 (Kahl); i, *Trimyema compressum*, ×410 (Lackey); j, *Spirozoa caudata*, ×370 (Kahl); k, *Trichospira inversa*, ×360 (Kahl).
Genus *Mycterothrix* Lauterborn (*Trichorhynchus Balbiani*). Anterior cone continuous on dorsal side with body ridge; hence free edge of body only on ventral side; no ventral slit.

*M. erlangeri* L. (Fig. 314, c). Nearly spherical with zoochlorellae; 50-55μ by 40-50μ; fresh water.

**Family 2 Trichopelmidae Kahl**

Genus *Trichopelma* Levander (*Leptopharynx Mermod*). Compressed; surface with longitudinal furrows, seen as lines in end-view; coarse ciliation throughout; cytostome toward left edge about 1/3 from the anterior end; cytopharynx tubular; macronucleus spheroid, central; 2 contractile vacuoles; fresh water.

*T. sphagnetorum* (L.) (Fig. 314, d). 25-40μ long; in fresh water.

Genus *Pseudomicrothorax* Mermod (*Craspedothorax Sondheim*). More or less compressed; cytostome opens in anterior half toward left side, in a depression surrounded by ciliary rows; body surface marked with a broad longitudinal ridge with cross striation; furrows canal-like; cilia on ventral side; cytopharynx tubular, with elastic rods; fresh water.

*P. agilis* M. (Fig. 314, e, f). Ellipsoid; 48-58μ long; in fresh water.

Genus *Drepanomonas* Fresenius (*Drepanoceras Stein*). Highly flattened; aboral surface convex; oral surface flat or concave; with a few deep longitudinal furrows; ciliation sparse; cytostome and a small cytopharynx simple, near the middle of body; fresh water. Several species.

*D. dentata* F. (Fig. 314, g). With a small process near cytostome; 2 rows of ciliary furrows on both oral and aboral surfaces; cilia on both ends of oral surface; 40-65μ long; in fresh water.

Genus *Microthorax* Engelmann (*Kreyella Kahl*). Small, flattened; with delicate keeled armor which is more or less pointed anteriorly and rounded posteriorly; ventral armor with 3 ciliary rows; oral depression posterior-ventral, with a stiff ectoplasmic lip on right side, below which there is a small membrane, and with a small tooth on left margin; no cytopharynx; macronucleus spherical; 2 contractile vacuoles; in fresh water. Many species.

*M. simulans* Kahl (Fig. 314, h). 30-35μ long; decaying plant infusion, also in moss.

**Family 3 Trimyemidae Kahl**

Genus *Trimyema* Lackey (*Sciadostoma Kahl*). Ovoid, more or less flattened; anterior end bluntly pointed, posterior end similar or rounded; with a long caudal cillum; cilia on 3-4 spiral rows which are
usually located in the anterior half of body; round cytostome near anterior end with a small cytopharynx; spherical macronucleus central with a small micronucleus; one contractile vacuole; active swimmer; fresh or salt water.

*T. compressum* L. (Fig. 314, i). About 65μ by 35μ; Lackey found it in Imhoff tank; fresh and salt water (Kahl). Klein (1930) studied its silverline system.

Family 4 Spirozonidae Kahl

Genus *Spirozona* Kahl. Short spindle-form; anterior end truncate, posterior region drawn out to a rounded end, with a group of longer cilia; spiral ciliation; beginning near right posterior third the central ciliary row runs over ridge to left and then reaches the cytostome; other rows are parallel to it; cytostome in anterior 1/4, with cytopharynx; ellipsoid macronucleus nearly central; contractile vacuole terminal; fresh water, sapropelic.

*S. caudata* K. (Fig. 314, j). 80–100μ long.

Family 5 Trichospiridae Kahl

Genus *Trichospira* Roux. Body cylindrical; posterior end rounded, anterior end conical in profile, where the cytostome surrounded by 2 spiral rows of cilia, is located; a special ciliary band beginning in the cytostomial region runs down on ventral side, turns spirally to left and circles partially posterior region of body; ciliary rows parallel to it; macronucleus oval, with a micronucleus; contractile vacuole posterior; fresh water, sapropelic.

*T. inversa* (Claparède and Lachmann) (Fig. 314, k). 70–100μ long.

Family 6 Plagiopylidae Schewiakoff

Genus *Plagiopyla* Stein. Peristome a broad ventrally opened groove from which body ciliation begins; peristomal cilia short, except a zone of longer cilia at anterior end; cytostome near median line at the end of the peristome; cytopharynx long; a peculiar ‘stripe band’ located on dorsal surface has usually its origin in the peristomal groove, after taking an anterior course for a short distance, curves back and runs down posteriorly near right edge and terminates about 1/3 the body length from posterior end; macronucleus rounded; a micronucleus; contractile vacuole terminal; free-living or endozoic.

*P. nasuta* S. (Fig. 315, a). Ovoid; tapering anteriorly; peristome at right angles or slightly oblique to the edge; trichocysts at right angles to body surface; macronucleus round to irregular in shape; body about 100μ (80–180μ) long; sapropelic in brackish water. Lynch
(1930) observed this ciliate in salt water cultures in California and found it to be 70–114μ by 31–56μ by 22–37μ.

P. minuta Powers (Fig. 315, b). 50–75μ by 36–46μ; in the intestine of Strongylocentrotus droebachiensis; the Bay of Fundy (Powers, 1933).

Genus **Lechriopyla** Lynch. Similar to Plagiopyla; but with a large internal organella, *furcula*, embracing the vestibule from right, and a large crescentic motorium at left end of peristome; in the intestine of sea-urchins.

![Fig. 315. a, Plagiopyla nasuta, ×340 (Kahl); b, P. minuta, ×400 (Powers); c, Lechriopyla mystax, ×340 (Lynch); d, Sonderia pharyngea, ×590 (Kirby); e, S. vorax, ×310 (Kahl); f, Clathrostoma viminal, ×220 (Penard); g, Physalophrya spumosa, ×160 (Penard).](image)

L. mystax L. (Fig. 315, c). 113–174μ long; in the gut of Strongylocentrotus purpuratus and S. franciscanus; California.

Genus **Sonderia** Kahl. Similar to Plagiopyla in general appearance; ellipsoid; flattened; peristome small and varied; body covered by 2–4μ thick gelatinous envelope which regulates osmosis, since no
contractile vacuole occurs (Kahl); with or without a striped band; trichocysts slanting posteriorly; in salt or brackish water. Kirby (1934) showed that several species of the genus are common in the pools and ditches in salt marshes of California, salinities of which range 3.5–10 per cent or even up to 15–20 per cent.

*S. pharyngea* Kirby (Fig. 315, d). Ovoid to ellipsoid; flattened; 84–110μ by 48–65μ; gelatinous layer about 2μ thick, with bacteria; about 60 longitudinal ciliary rows, each with 2 borders; peristome about 35μ long, at anterior end, oblique; with closely set cilia from the opposite inner surfaces; cytopharynx conspicuous; spherical macronucleus anterior, with a micronucleus; trichocysts (7–9μ long) distributed sparsely and unevenly, oblique to body surface; a group of bristle-like cilia at posterior end; often brightly colored because of food material; in salt marsh, California.

*S. vorax* Kahl (Fig. 315, e). Broadly ellipsoid; size variable, 70–180μ long; ventral surface flattened; posterior border of peristomal cavity extending anteriorly; in salt marsh; California (Kirby, 1934).

**Family 7 Clathrostomidae** Kahl

Genus *Clathrostoma* Penard. Ellipsoid; with an oval pit in anterior half of the flattened ventral surface, in which occur 3–5 concentric rows of shorter cilia; cytostome a long slit located at the bottom of this pit; with a basket composed of long fibrils on the outer edge of the pit; in fresh water.

*C. viminale* P. (Fig. 315, f). Resembles a small *Frontonia leucas*; macronucleus short sausage-form; 4 micronuclei in a compact group; endoplasm with excretion crystals; 5 preoral ciliary rows; 130–180μ long; in fresh water.

**Family 8 Parameciidae** Grobben

Genus *Paramecium* Hill (*Paramaccium* Müller). Cigar- or foot-shaped; circular or ellipsoid in cross section; with a single macronucleus and 1 to several vesicular or compact micronuclei; peristome long, broad, and slightly oblique; in fresh or brackish water. Several species. Comparative morphology (Wenrich, 1928a; Wichterman, 1933); ciliary arrangement (Lieberman, 1928); pellicular structure (Gelei, 1939); excretory system (Gelei, 1939a); spiral movement (Bullington, 1930); cultivation (Wichterman, 1949).

*P. caudatum* Ehrenberg (Figs. 21, a, b; 43, a–e; 52; 83; 316, a). 180–300μ long; with a compact micronucleus, a massive macronucleus; 2 contractile vacuoles on aboral surface; posterior end bluntly pointed; in fresh water. The most widely distributed species. Cytol-
ogy and physiology (Müller, 1932); contractile vacuoles (Dimitrowa, 1928); cytopharynx (Gelei, 1934); calcium and iron (Kruszynski, 1939); nuclear variation (Diller, 1940); re-conjugation (Diller, 1942); food vacuoles (Bozler, 1924); conjugation (p. 187).

*P. aurelia* E. (Figs. 2, g, h; 57; 89; 100; 101; 102; 316, b). 120–180 μ long; two small vesicular micronuclei, a massive macronucleus; two contractile vacuoles on aboral surface; posterior end more rounded than *P. caudatum*; in fresh water. Nutrition (Phelps, 1934); autogamy and hemixis (Diller, 1936); conjugation and mating types (p. 190).

Fig. 316. Semi-diagrammatic drawings of nine species of *Paramecium* in oral surface view, showing distinguishing characteristics taken from fresh and stained specimens, ×230 (several authors). a, *P. caudatum*; b, *P. aurelia*; c, *P. multimicronucleatum*; d, *P. bursaria*; e, *P. putrinum*; f, *P. calkinsi*; g, *P. trichium*; h, *P. polycaryum*; i, *P. woodruffi*. 
P. multimicronucleatum Powers and Mitchell (Figs. 19; 20; 28; 29; 316, c). The largest species, 200-330μ long; three to seven contractile vacuoles; four or more vesicular micronuclei; a single macronucleus; in fresh water. Cytology and physiology (Müller, 1932); division and conjugation (Stanghoner, 1932; Köster, 1933); relation to Oikomonas and bacteria in culture (Hardin, 1944).

P. bursaria (Ehrenberg) (Figs. 84; 88; 316, d). Foot-shaped, somewhat compressed; about 100-150μ by 50-60μ; green with zoochlorellae as symbionts; a compact micronucleus; a macronucleus; two contractile vacuoles; in fresh water. Relation between Chlorella and host (Parker, 1926; Pringsheim, 1928); micronuclear variation (Woodruff, 1931); bacteria-free culture (Loefer, 1936); removal of symbionts (Jennings, 1938; Wichterman, 1948); conjugation (p. 189).

P. putrinum Claparède and Lachmann (Fig. 316, e). Similar to P. bursaria, but a single contractile vacuole and an elongated macronucleus; no zoochlorellae; 80-150μ long; in fresh water.

P. calkinsi Woodruff (Fig. 316, f). Foot-shaped; posterior end broadly rounded; 100-150μ by 50μ; 2 vesicular micronuclei; 2 contractile vacuoles; in fresh, brackish and salt water. Ecology, morphology, mating types (Wichterman, 1951).

P. trichium Stokes (Fig. 316, g). Oblong; somewhat compressed; 50-105 (80-90)μ long; a compact micronucleus; two contractile vacuoles deeply situated, each with a convoluted outlet; in fresh water. Structure and division (Wenrich, 1926); conjugation (p. 190) (Diller, 1948, 1949).

P. polycaryum Woodruff and Spencer (Fig. 316, h). Form similar to P. bursaria; 70-110μ long; 2 contractile vacuoles; 3-8 vesicular micronuclei; in fresh water.

P. woodruffii Wenrich (Fig. 316, i). Similar to P. polycaryum; 150-210μ long; 2 contractile vacuoles; 3-4 vesicular micronuclei; brackish water (Wenrich, 1928).

Although Paramecium occurs widely in various freshwater bodies throughout the world and has been studied extensively by numerous investigators by mass or pedigree culture method, there are only a few observations concerning the process of encystment. Bütschli considered that Paramecium was one of the Protozoa in which encystment did not occur. Stages in encystment were however observed in P. bursaria (by Prowazek) and in P. putrinum (by Lindner). In recent years, four observers reported their findings on the encystment of Paramecium. Curtis and Guthrie (1927) give figures in their textbook of zoology, showing the process (in P. caudatum?) (Fig. 317, a-c), while Cleveland (1927) injected Paramecium culture into the rectum
of frogs and observed that the ciliate encysted within a thin membrane. Michelson (1928) found that if *P. caudatum* is kept in Knop-agar medium, the organism becomes ellipsoidal under certain conditions, later spherical to oval, losing all organellae except the nuclei, and develops a thick membrane; the fully formed cyst is elongated and angular, and resembles a sand particle (Fig. 317, f). Michelson considers its resemblance to a sand grain as the chief cause of the cyst having been overlooked by workers. In all these cases, it may however be added that excystment has not been established.

![Fig. 317. a–c, encystment in a species of Paramecium (Curtis and Guthrie); d–f, encystment of *P. caudatum*, ×380 (Michelson).](image)

Genus *Physalophrya* Kahl. Without peristome; but cytostome located near the anterior half of body, resembles much that of *Paramecium*; although there is no membrane, a ciliary row occurs in the left dorsal wall of cytopharynx; in fresh water. Taxonomic status is not clear; but because of its general resemblance to Paramecium, the genus with only one species is mentioned here.

*P. spumosa* (Penard) (Fig. 315, g). Oval to cylindrical; highly plastic; cytoplasm reticulated; numerous contractile vacuoles; 150–320μ long; in fresh water.

Family 9 *Colpodidae* Poche

Genus *Colpoda* Müller. Reniform; compressed; right border semicircular; posterior half of the left border often convex; oral funnel in the middle of flattened ventral side; cytostome is displaced to the
right of the median plane, which leads into peristome cavity and gives rise dorsally to a diagonal groove; right edge of cytostome bears a ciliated area, but no protruding membrane as in *Bryophrya* (p. 747); macronucleus spherical or oval, central; a compact micronucleus; a contractile vacuole terminal; in fresh water. Many species. Burt (1940) made a comparative study of five species, which are mentioned here.

*C. cucullus* M. (Fig. 318, a). 40–110\(\mu\) long; anterior keel with eight
to 10 indentations; 29–34 ciliary grooves; cilia mostly paired; macronucleus with a stellate endosome; trichocysts rod-form; usually with abundant food vacuoles; in fresh water with decaying plants.

*C. inflata* (Stokes) (Fig. 318, b). 35–90\(\mu\) long; anterior keel with 6–8 indentations; number of ciliary grooves (or meridians) 21–24; cilia mostly in pairs; in fresh water among vegetation.

*C. maupasi* Enriques (Fig. 318, c). 35–90\(\mu\) long; cytostome about
one-fourth from the anterior end; anterior keel with five indentations; 16–18 meridians; in fresh water.

*C. aspera* Kahl (Fig. 318, d). 12–42 μ long; cytostome about one-third from the anterior end; 14–16 meridians; anterior keel with five indentations; in fresh water.

*C. steini* Maupas (Fig. 318, e). 15–42 μ; cytostome about two-fifths from the anterior end, and with a bundle of long membranellae; five to six preoral ridges; paired and single cilia; one pair of long caudal cilia; 12 meridians; in fresh water. The organism can live in various organs of the land slug, *Agriolimax agrestis* (Reynolds, 1936).

*C. duodenaria* Taylor and Furgason. 20–40 μ (9–60 μ) long; 12 longitudinal ciliary rows; 3 postoral rows; 2 long cilia at the posterior end; long cilia project out from the cytostome along its posterior margin, forming a “beard”; a contractile vacuole terminal; macronucleus ovoid, with crescentic micronucleus; division into 2–8 individuals in division cyst; but no division in trophozoite stage; bacteria-feeder; fresh water. Encystment (Taylor and Strickland, 1939); identity (Burt, 1940).

Genus *Tillina* Gruber. Similar to *Colpoda* in general appearance and structure; but cytopharynx a long curved, ciliated tube; in fresh water.

*T. magna* G. (Fig. 318, f, g). 180–200 μ long (Gruber), up to 400 μ long (Bresslau); macronucleus oval to rod-shape; micronuclei vesicular, highly variable in number (2–16) (Beers); a contractile vacuole terminal, with six long collecting canals; division cyst produces four individuals; in stagnant water and also coprozoic. Morphology (Gregory, 1909; Beers, 1944, 1945); encystment and excystment (Beers, 1945, 1946, 1946a).

*T. canalifera* Turner (Figs. 26; 318, h). 150–200 μ by 100–150 μ; resembles magna; but macronucleus ellipsoid, about one-third the body length; four to 14 micronuclei, clustered around the macronucleus; a terminal contractile vacuole with seven to nine long permanent collecting canals; cytoplasm with 3–7 μ long refractile rods; in fresh water (Turner, 1937). Cytoplasmic inclusions (Turner, 1940).

Genus *Bresslaua* Kahl. General body form resembles *Colpoda*; but cytopharynx large and occupies the entire anterior half.

*B. vorax* K. (Fig. 318, i). 80–250 μ long; in fresh water.

Genus *Bryophrya* Kahl. Ovoid to ellipsoid; anterior end more or less bent toward left side; cytostome median, about 1/3 from anterior end, its right edge continues in horseshoe form around the posterior end and half of the left edge; anterior portion of left edge of
the cytostome with posteriorly directed membrane; macronucleus oval or spherical; micronuclei; in fresh water.

_B. bavariensis_ K. (Fig. 318, j). 50–120μ long.

Genus _Woodruffia_ Kahl. Form similar to _Chilodonella_ (p. 731); highly flattened snout bent toward left; cytostome, a narrow diagonal slit, its left edge with a membranous structure and its right edge with densely standing short cilia; macronucleus spherical; several (?) micronuclei; contractile vacuole flattened, terminal; in salt water.

_W. rostrata_ K. (Fig. 318, k). 120–180μ long; salt water culture with Oscillatoria.

_W. metabolica_ Johnson and Larson (1938). Pyriform; 85–400μ long; division cysts 85–155μ in diameter; resting cysts 40–62μ in diameter; in freshwater ponds. Johnson and Evans (1939, 1940) find two types of protective cysts in this ciliate: “stable” and “unstable” cysts, formation of both of which depends upon the absence of food. These cysts have three membranes: a thin innermost endocyst, a rigid mesocyst and a gelatinous outer ectocyst. The protoplasmic mass of the stable cyst is smaller, and free from vacuoles, and its ectocyst is thick, while that of the unstable cyst is larger, contains at least one fluid vacuole and its ectocyst is very thin. Crowding, feeding on starved Paramecium, increasing the temperature, and increasing the salt concentration of the medium, are said to influence the formation of unstable cysts. The two authors (1941) further reported that when free-swimming individuals were subjected, in the absence of food, to extremes of temperature, high concentrations of hydrogen-ion, and low oxygen tension, unstable cysts were formed; when the oxygen tension decreased, the tendency to encyst increased, even when ample food was present. The unstable cysts are said to remain viable for six months. Excystment is induced by changing the balanced salt solution, by replacing it with distilled water and by lowering temperature from 30° to 20°C.

Family 10 _Entorhipidiidae_ Madsen

Genus _Entorhipidium_ Lynch. Triangular in general outline; colorless; large, 155–350μ long; flattened; posterior end drawn out, with a bristle; anterior end bent to left; cytostome in depression close to left anterior border, with long cilia; with or without a cross-groove from preoral region; cytopharynx inconspicuous; trichocysts; macronucleus oval to sausage-form; one to several micronuclei; several (excretory) vacuoles left-ventral; in intestine of the starfish, _Strongylocentrotus purpuratus_. Four species.
**E. echini** L. (Fig. 319, a). About 253µ by 125µ; California.

Genus *Entodiscus* Madsen. Broadly or narrowly lancet-like, without narrowed posterior portion; cytostome small on left narrow side, about 2/5 the body length from anterior end; without trichocysts; macronucleus central, with a micronucleus; contractile vacuole sub-terminal; swimming movement rapid without interruption. Two species. Morphology (Powers, 1933, 1933a).

*E. indomitus* M. (Fig. 319, b). 80–117µ by 20–23µ; in the intestine of *Strongylocentrotus droebachiensis*.

*E. borealis* (Hentschel) (Fig. 319, c). Oval; cytostome nearer anterior end; 105–170µ by 60–115µ; in the gut of *Strongylocentrotus*

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Fig. 319. a, *Entorhipidium echini*, ×270 (Lynch); b, *Entodiscus indomitus*, ×380 (Madsen); c, *E. borealis*, ×380 (Powers); d, *Biggaria bermudense*, ×380 (Powers); e, *B. echinometris*, ×380 (Powers); f, *Anophrys elongata*, ×390 (Powers); g, *A. aglycus*, ×390 (Powers).
droebachiensis and Echinus esculentus; Powers (1933) studied this species in the first-named host from Maine, and found a supporting rod which is imbedded in the margin along the right wall of the oral cavity and which he named stomatostyle.

Genus Biggaria Kahl. Scoop-like form; anterior 2/3 thin, posterior region thickened, terminating in a rudder-like style; cilia in longitudinal rows; longer cilia on caudal prolongation; cytostome in the posterior half, opening into a vestibule, into which long cilia project from the roof; aperture to cytopharynx with 2 membranes; contractile vacuole subterminal; in the intestine of sea-urchins.

B. bermudense (Biggar) (Fig. 319, d). 90–185μ by 48–82μ; in Lytechinus variegatus; Bermuda (Biggar), North Carolina (Powers). Powers (1935) found the organism at Tortugas in Lytechinus variegatus, Centrechinus antillarum, Echinometra lucunter, Tripneustes esculentus and Astrophyga magnifica.

B. echinometris (B.) (Fig. 319, e). 80–195μ by 33–70μ; in Echinometris subangularis (Bermuda) and Lytechinus variegatus (North Carolina).

Genus Anophrys Cohn. Cigar-shaped; flexible; longitudinal ciliary rows; peristome begins near the anterior end, parallel to body axis and about 1/3 the body length; a row of free cilia on right edge of peristome; cytostome inconspicuous; spherical macronucleus central; contractile vacuole terminal; in the intestine of sea-urchins.

A. elongata Biggar (Fig. 319, f). About 96μ long (Powers); 166μ long (Biggar); in the gut of Lytechinus variegatus and Echinometris subangularis; Bermuda (Biggar); Powers (1935) found this species also in the hosts mentioned for Biggaria bermudense.

A. aglycus Powers (Fig. 319, g). 56–120μ by 16–35μ; in the gut of Centrechinus antillarum and Echinometra lucunter; Tortugas (Powers, 1935).

Family 11 Paraisotrichidae da Cunha

Genus Paraisotricha Fiorentini. Uniformly ciliated in more or less spiral longitudinal rows; longer cilia at anterior end; cytostome near anterior tip; contractile vacuole posterior; in the caecum and colon of horse.

P. colpoidea F. (Fig. 320, a). 70–100μ by 42–60μ. Conjugation (Dogiel, 1930).

P. beckeri Hsiung (Fig. 320, b). 52–98μ by 30–52μ (Hsiung, 1930, 1930a).
Family 12 Isotrichidae Bützchli

Genus Isotricha Stein. Ovoid; flattened; dense longitudinal ciliary rows; cytostome at or near anterior end; several contractile vacuoles; reniform macronucleus and a micronucleus connected with, and suspended by, fibrils, karyophore; locomotion with posterior end directed forward; in the stomach of cattle and sheep.

Fig. 320. a, Paraisotricha colpoidea, ×270 (Hsiung); b, P. beckeri, ×360 (Hsiung); c, Isotricha prostoma, ×500 (Becker and Talbott); d, I. intestinalis, ×500 (Becker and Talbott); e, Dasytricha ruminantium, ×330 (Becker and Talbott); f, Cyathodinium piriforme, ×1290 (Lucas); g, Blepharocorys uncinata, ×540 (Reichenow); h, B. bovis, ×850 (Dogiel); i, Charon equi, ×570 (Hsiung).

I. prostoma S. (Fig. 320, c). 80–195 μ by 53–85 μ. Cytology (Campbell, 1929).

I. intestinalis S. (Fig. 320, d). 97–130 μ by 68–88 μ.

Genus Dasytricha Schuberg. Oval, flattened; cilia in longitudinal spiral rows; no karyophore; in the stomach of cattle.

D. ruminantium S. (Fig. 320, e). 50–75 μ by 30–40 μ.
Family 13 Cyathodiniidae da Cunha

Genus Cyathodinium da Cunha. Conical or pyriform; broad cytostome occupies the entire anterior end and extends posteriorly 1/4–3/4 the body length; deep with prominent ridges; oral cilia in a single row on left ridge; body cilia comparatively long, confined to anterior half; macronucleus round or ellipsoid; a micronucleus; one to several contractile vacuoles; in the caecum and colon of guinea pigs.

*C. conicum* da C. Inverted cone; 50–80μ by 20–30μ; in the caecum of *Cavia aperea* and *C. porcella*.

*C. piriforme* da C. (Fig. 320, f). Typical form inverted pyriform; second form conical with tapering anterior end; contractile vacuole posterior; 30–40μ by 20–30μ; in the caecum of *Cavia aperea* and *C. porcella*. Occurrence and cytology (Lucas, 1932, 1932a; Nie, 1950).

Family 14 Blepharocoridae Hsiung

Genus Blepharocorys Bundle. Oral groove deep, near anterior end; 3 (oral, dorsal and ventral) ciliary zones at anterior end; a caudal ciliary zone; in the caecum and colon of horse or stomach of cattle. Many species.

Fig. 321. The developmental cycle of *Conidiophrys pilisuctor* (Chatton and Lwoff). a, trophont with two tomites; b, freed tomite; c, tomite becoming attached to host's hair; d, lacrymoid trophont; e, spheroid stage; f, g, cucurbitoid stage.
**B. uncinata** (Fiorentini) (*B. equi* Schumacher) (Fig. 320, g). With a screw-like anterior process; 55–74μ by 22–30μ; in the caecum and colon of horse (Hsiung, 1930a).

**B. bovis** Dogiel (Fig. 320, h). 23–37μ by 10–17μ; in the stomach of cattle (Dogiel, 1926).

Genus **Charon** Jameson. Two caudal ciliary zones; in the colon of horse or in stomach of ruminants.

**C. equi** Hsiung (Fig. 320, i). 30–48μ by 10–14μ; in the colon of horse (Hsiung, 1930, 1930a).

**Family 15 Conidophryidae** Mohr and LeVeque

(*Pilisuctoridae* Chatton and Lwoff)

Genus **Conidophrys** Chatton and Lwoff (Fig. 321). Trophont or the form attached to host's appendages (a), cylindrical, with a thick pellicle; contents divide into two or three (and up to several) smaller bodies which develop into tomites or free-swimming individuals (b); when the latter come in contact with the ends of the secretory hairs

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**Fig. 322. Conidiophrys pilisuctor** (Chatton and Lwoff). a, trophonts of all ages on an appendage of *Corophium acherusicum*; b, a stained mature trophont with two formed and one developing tomites, ×1330; c, a tomite emerging from trophont, ×1330; d, a living tomite, ×2230; e, newly attached lacrymoid trophont, ×1330.
of the host, they become attached through their cytopharynx (c) and lose their cilia; during the development into the cucurbitoid mature stage (f, g), the organism passes through lacrymoid (d) and spheroid (e) stages; on freshwater amphipods and isopods (Chatton and Lwoff, 1934, 1936).

*C. pilisuctor* C. and L. (Fig. 322). Lacrymoid trophont 12–15μ by 6–7μ; cucurbitoid forms 50–60μ long; free-swimming tomites 12–14μ in diameter by 6–7μ high, ciliated and possess a comparatively long cytopharynx; nourishment of trophont through host’s hairs; in amphipods and isopods, especially on *Corophium acherusicum*, France. Mohr and LeVeque (1948) found it on the wood-boring isopods, *Limnoria lignorum* and *Corophium acherusicum* in California.

**References**


— (1948) Nuclear behavior of *Paramecium trichium* during conjugation. Ibid., 82:1.


Chapter 35

Order 1 Holotricha Stein (continued)

Suborder 4 Hymenostomata Delage and Hérouard

Cytostome not connected with peristome .......... Family 1 Frontoniidae
Cytostome at end or bottom of peristome

Peristome sickle-form, ciliated slit; sunk at right angles to body surface

................................. Family 2 Ophryoglenidae (p. 767)

Peristome long, begins at anterior end of body

Peristome with a one-layered membrane which forms a pocket surrounding cytostome on right edge and a row of cilia or membrane on left ............... Family 3 Pleuronematidae (p. 769)

Peristome otherwise

Peristome with 2 one-layered membranes; no distinct ectoplasmic pocket around cytostome. Family 4 Cohnilembidae (p. 771)

Peristome furrow either covered densely with cilia, besides an undulating membrane on right edge, or with only a thick undulating membrane on the right edge. .................. Family 5 Philasteridae (p. 771)

Family 1 Frontoniidae Kahl

Genus Frontonia Ehrenberg. Ovoid to ellipsoid; anterior end more broadly rounded than posterior end; flattened; oral groove lies in anterior third or more or less flattened ventral surface, to right of median line; lancet-like with pointed anterior and truncate posterior end; left edge is more curved than right edge, and posteriorly becomes a prominent ectoplasmic lip; cytostome with a complex organization (on left edge a large undulating membrane composed of 3 layers, each being made up of 4 rows of cilia; on right, semimembranous groups of cilia; 3 outer rows of cilia from the postoral suture; along this suture ectoplasm is discontinuous so that large food matter is taken in; with a small triangular ciliated field posterior to cytostome and left of suture); a long narrow postoral groove which is ordinarily nearly closed; cytopharynx with numerous strong fibrils; ciliary rows close and uniform; ectoplasm with numerous fusiform trichocysts; macronucleus oval; one to several micronuclei; 1–2 contractile vacuoles, with collecting canals and an external pore; in fresh or salt water. Species identification and movement (Bullinghton, 1939); trichocysts (Krüger, 1931).

F. leucas E. (Figs. 2, i, j; 323, a–c). 150–600μ long; feeds on filamentous algae, but may take in Arcella and even large amoebae (Beers, 1933); among algae in fresh water.

F. branchiostoma Codreanu (Fig. 323, d). 75–100μ by 55–95μ; commensal in the branchial cavity of Amphioxus.

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Genus *Disematostoma* Lauterborn. Somewhat similar to *Frontonia*; pyriform; with broadly rounded, truncate or concave anterior end and bluntly pointed narrow posterior end; preoral canal wide; a dorsal ridge in posterior region of body; macronucleus sausage-form; a micronucleus; contractile vacuole in middle of body, with long collecting canals; in fresh water.

Fig. 323. a–c, *Frontonia leucas* (Bullington) (a, aboral view showing a contractile vacuole, collecting canals, macronucleus, four micronuclei and trichocysts, ×220; b, oral view, showing the cytostome with undulating membrane and groove, ×165; c, portion of pellicle with wart-like projections over trichocysts); d, *F. branchiostomae*, ×490 (Codreanu); e, *Disematostoma bütschlii*, ×340 (Kahl); f, *Lembadion bullinum*, ×170 (Kahl); g, *Tetrahymena pyriformis*, ×950 (Furgason).
**D. bütschlii** L. (Fig. 323, e). 135–155μ long; with or without zoochlorellae; in fresh water.

Genus **Lembadion** Perty. Oval; dorsal side convex, ventral side concave; cytostome 3/4–4/5 the body length; on its left with a large membrane composed of many ciliary rows and on its right, numerous narrow rows of short free cilia; an undulating membrane and ciliary rows near posterior end; contractile vacuole in mid-dorsal region with a long tubule opening at posterior-right side; close ciliation uniform; macronucleus ellipsoid, subterminal; a micronucleus; long caudal cilia; in fresh water.

**L. bullinum** P. (Fig. 323, f). 120–200μ long; posterior cilia 40–50μ long.

Genus **Tetrahymena** Furgason (1941). Pyriform; small forms; uniform ciliation; ciliary rows or meridians 17–42; 2 postoral meridians; preoral suture straight; cytostome small, close to anterior end, pyriform; its axis parallel to body axis; inconspicuous ectoplasmic ridge or flange on the left margin of mouth; an undulating membrane on right side and 3 membranellae on left of the cytostome; a single contractile vacuole; macronucleus ovoid; micronucleus absent in some species; in fresh water or parasitic. Corliss (1952, 1952a) made a comparative study of different strains and allied forms.

**T. pyriformis** (Ehrenberg) (*T. geleii* Furgason) (Figs. 323, g; 324, a–c). 59 strains (Corliss, 1952a); 40–60μ long; 17–23 ciliary meridians; pyriform cytostome about 1/10 the body length; with or without micronucleus; bacteria-feeder; in fresh water (Corless, 1952, 1952a). Bacteria-free or pure culture (Kidder, 1941) (p. 884).

**T. vorax** (Kidder, Lilly and Claff) (*Glaucoma vorax* K. L. and C.) (Fig. 39). Form and size vary; bacteria-feeders elongate pyriform, 50–75μ long; saprozoic forms fusiform, 30–70μ long, decreasing in size with the age of culture; sterile particle-feeders, 60–80μ long; carnivores and cannibals broadly pyriform, 100–250μ long; 19–21 ciliary meridians; macronucleus ovoid, central; in carnivores, outline irregular; apparently without micronucleus; pond water.

**T. limacis** (Warren). In the liver and other visceral organs of the gray garden slug, *Deroceras agreste*; 33–68 (55)μ by 18–35(27)μ; those from cultures measure 28–68(44)μ by 17–42(27)μ; the parasitic phase is cucumber-shaped with apiculate anterior end; the free-living organisms are pyriform, somewhat pointed anteriorly; cytostome at about 1/4 from the anterior end, with an undulating membrane and three membranellae; 33–37 ciliary rows (Kozloff, 1946).

Genus **Leucophrys** Ehrenberg. Broadly pyriform; cytostome large, pyriform, with its axis parallel to body axis; ectoplasmic flange along
left margin; undulating membrane on right and 3 membranellae on left of mouth; 5 postoral ciliary meridians; macronucleus ovoid; a micronucleus; fresh water.

$L.\ patula$ E. (Fig. 324, d–f). Broadly pyriform; 80–160$\mu$ long; occasionally small forms occur; cytostome pyriform, about 1/3 the body length; 40–45 ciliary meridians; macronucleus irregularly ovoid; a micronucleus attached to macronucleus; carnivorous, but may be cultured in sterile media (Kidder); fresh water. Morphogenesis (Fauré-Fremiet, 1948).

Genus Glaucoma Ehrenberg (Dallasia Stokes). Ovoid or ellipsoid; cytostome about one-fourth the body length, near anterior end, ellipsoid; cytostome with an inconspicuous undulating membrane

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Fig. 324. a–e, Tetrahymena pyriformis (a, $\times535$ (Kidder); b, c, cytostomal structure (Furgason)); d–f, Leucophrys patula (d, a well-fed animal, $\times280$ (Maupas); e, a diagram, $\times535$ (Kidder); f, cytostome (Furgason)); g, h, Glaucoma scintillans (g, a diagram, $\times535$ (Kidder); h, cytostome (Furgason)); i, j, Colpidium colpoda (i, $\times180$ (Kahl); j, cytostome (Furgason)); k, C. campylum, $\times535$ (Kidder); l, C. echini, $\times385$ (Powers); m, Paraglaucoma rostrata, $\times400$ (Kahl); n, Malacophrys rotans, $\times500$ (Kahl).
on right and 3 membranellae on left; ectoplasmic ridge on right border of mouth; ciliation uniform; 30-40 ciliary meridians; 7 postoral meridians; macronucleus rounded; a micronucleus; a contractile vacuole; with or without 1 or more caudal bristles; fresh water.

*G. scintillans* E. (Fig. 324, g, h). Ovate with rounded ends; 45-75μ long; U-shaped cytostome, about one-fourth the body length, oblique; ectoplasmic flange and 3 membranellae conspicuous; a contractile vacuole in posterior one-third; macronucleus oval, central; a micronucleus; bacteria-feeder; in fresh water. Bacteria-free culture (Kidder, 1941); division (Kidder and Diller, 1934)

Genus **Colpidium** Stein. Elongate reniform; ciliary meridians variable in number, but typically one postoral meridian; small triangular cytostome one-fourth from anterior end toward right side; a small ectoplasmic flange along right border of cytostome which shows an undulating membrane on right and 3 membranellae on left; rounded macronucleus; a micronucleus; a contractile vacuole; fresh or salt water or parasitic.

*C. colpoda* (Ehrenberg) (*Tillina helia* Stokes) (Figs. 10, e; 324, i, j). Elongate reniform; 90-150μ long; cytostome about one-tenth the body length; 55-60 ciliary meridians; preoral suture curved to left; macronucleus oval, central; a micronucleus; fresh water. Bacteria-free culture (Kidder, 1941); division (Kidder and Diller, 1934); effect of food bacteria on division (Burbank, 1942).

*C. campylum* (Stokes) (Fig. 324, k). Elongate reniform; 27-30 ciliary meridians; preoral suture curved to right; 50-70μ long; in fresh and brackish water. Division (Kidder and Diller, 1934).

*C. striatum* S. Similar to the last species; contractile vacuole further posterior; 50μ long; in standing water.

*C. echini* (Russo) (Fig. 324, l). In the intestinal caeca of *Strongylocentrotus lividus*; 37-64(55)μ by 21-28(25)μ; 24 longitudinal ciliary rows; cytostome at anterior third (Powers, 1933).

Genus **Paraglaucoma** Kahl. Somewhat similar to *Glaucoma*; but without perioral ectoplasmic ridge; a membrane on right ridge of the cytostome; anterior end drawn out to a point in profile, posterior end rounded; a stiff posterior bristle; a contractile vacuole; rapid zig-zag movement. One species.

*P. rostrata* K. (Fig. 324, m). 60-80μ long; in fresh water (often in dead rotiferan body); California, Wisconsin (Kahl).

Genus **Malacophrys** Kahl. Ellipsoid or cylindrical; plastic; cilia uniformly close-set in longitudinal rows; slit-like cytostome at anterior extremity; in fresh water.

*M. rotans* K. (Fig. 324, n). Oval; close and dense ciliation; spheri-
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cal macronucleus central; a micronucleus; a single contractile vacuole; body 40–50\(\mu\) long; fresh water.

Genus *Espejoia* Bürger (*Balantiophorus* Penard). Ellipsoid; anterior end obliquely truncate; large cytostome at anterior end; post-oral groove on ventral side, 1/4–1/3 the body length; a conspicuous membrane on the left edge of groove; in gelatinous envelope of eggs of insects and molluses.

*E. musicola* (P.) (Fig. 325, a). Elongate; right side flat, left side convex; 80–100\(\mu\) long (Penard); 70–80\(\mu\) long and dimorphic (Fauré-Fremiet and Mugard, 1949).

Genus *Cryptochilidium* Schouteden (*Cryptochilum* Maupas). Ellipsoid; with rounded anterior end, posterior end pointed in profile; highly compressed; uniform and close ciliation; cytostome near middle; one or more longer cilia at posterior end; contractile vacuole posterior; macronucleus round; a micronucleus; commensal. Several species (Powers, 1933, 1935).

*C. echini* (Maupas) (Fig. 325, b). 70–140\(\mu\) long; in the gut of *Echinus lividus*.

Genus *Eurychilum* André. Elongate ellipsoid; anterior end somewhat narrowed; cilia short; dense ciliation not in rows; contractile vacuole terminal; macronucleus band-form; cytostome about 2/5 from anterior end and toward right, with a strong undulating membrane on left; no cytopharynx; actively swimming. One species.

*E. actiniae* A. (Fig. 325, c). About 155\(\mu\) long; in gastrovascular cavity of *Sagartia parasitica*.

Genus *Monochilum* Schewiakoff. Ovoid to ellipsoid; medium large; uniform and dense ciliation in rows; oblong cytostome left of median line, in about 1/4 the body length from anterior end; short cytopharynx conical, with an undulating membrane; contractile vacuole near middle; in fresh water.

*M. frontatum* S. (Fig. 325, d). Anterior end broader; ventrally flattened, dorsally somewhat convex; macronucleus ellipsoid; a micronucleus; feeds on algae; 80\(\mu\) by 30\(\mu\).

Genus *Dichilum* Schewiakoff. Similar to *Monochilum*; but membrane on both edges of the cytostome; in fresh or salt water.

*D. cuneiforme* S. (Fig. 325, e). Ellipsoid; cytostome about 1/5 the body length from anterior end; right membrane larger than left; small cytopharynx; macronucleus ellipsoid; about 40\(\mu\) by 24\(\mu\); in fresh water.

Genus *Loxocephalus* Eberhard. Ovoid to cylindrical; sometimes compressed; crescentic cytostome on slightly flattened area near anterior end, with 2 membranes; often a zone of cilia around body;
usually 1 (or more) long caudal cilium; endoplasm granulated, yellowish to dark brown; macronucleus ovoid; a single contractile vacuole; in fresh or brackish water. Many species.

*L. plagius* (Stokes) (Fig. 325, f). 50–65μ long; nearly cylindrical; 15–16 ciliary rows; endoplasm usually darkly colored; in fresh water among decaying vegetation.

Genus *Balanonema* Kahl. Similar to *Loxocephalus*; but with plug-like ends; cytostome difficult to see; a caudal cilium; macronucleus oval; contractile-vacuole; ciliation uniform or broken in the middle zone; fresh water.

*B. biceps* (Penard) (Fig. 325, g). Ellipsoid; no cilia in the middle region; contractile vacuole central; macronucleus posterior to it; 42–50μ long.

Genus *Platynematum* Kahl. Ovoid or ellipsoid; highly flattened; with a long caudal cilium; contractile vacuole posterior-right; small
cytostome more or less toward right side, with 2 outer membranes; ciliary furrows horseshoe-shaped; in fresh or salt water.

*P. sociale* (Penard) (Fig. 325, h). Anterior half more flattened; ventral side concave; cytostome in the anterior third; yellowish and granulated; 30–50μ long; sapropelic in fresh and brackish water.

Genus *Saprophilus* Stokes. Ovoid or pyriform; compressed, cytostome in anterior 1/4–1/3 near right edge, with two membranes; macronucleus spherical; contractile vacuole posterior; in fresh water.

*S. agitatus* S. (Fig. 325, i). Ellipsoid; ends bluntly pointed; compressed; plastic; close striation; about 40μ long; in fresh water in decomposing animal matter such as Gammarus.

*S. muscorum* Kahl (Fig. 325, j). Cytostome large, with a large membrane; trichocysts; contractile vacuole with a distinct canal; body about 35μ long; in fresh water.

Genus *Cinetochilum* Perty. Oval to ellipsoid; highly flattened; cilia on flat ventral surface only; cytostome right of median line in posterior half, with a membrane on both edges which form a pocket; oblique non-ciliated postoral field leads to left posterior end; with 3–4 caudal cilia; macronucleus spherical, central; contractile vacuole terminal; in fresh or salt water. Neuroneme system (Gelei, 1940).

*C. margaritaceum* P. (Fig. 325, k). 15–45μ long; in fresh and brackish water.

Genus *Dexiotrichides* Kahl (*Dexiotricha* Stokes). Reniform; compressed; cytostome near middle, with two membranes; long cilia sparse; a special oblique row of cilia; a single caudal cillum; contractile vacuole terminal; spheroidal macronucleus anterior; a micronucleus; in fresh water. One species.

*D. centralis* (Stokes) (Fig. 326, a). About 30–45μ long; in decaying vegetable matter.

Genus *Cyrtolophosis* Stokes. Ovoid or ellipsoid; with a mucilaginous envelope in which it lives, but from which it emerges freely; cytostome near anterior end with a pocket-forming membrane; on right side a short row of special stiff cilia, bent ventrally; sparse ciliation spiral to posterior-left; spherical macronucleus central; a contractile vacuole; in fresh water.

*C. mucicola* S. (Fig. 326, b). 25–28μ long; in infusion of leaves.

Genus *Urocentrum* Nitzsch. Short cocoon-shaped, constricted in the middle; ventral surface flat; 2 broad girdles of cilia; fused cilia at posterior end; with a zone of short cilia in the constricted area; cytopharynx with a stiff ectoplasmic membrane which separates two undulating membranes; macronucleus horseshoe-shaped,
posterior; a micronucleus; contractile vacuole terminal, with eight long collecting canals which reach the middle of body; in fresh water.

*U. turbo* (Müller) (Fig. 326, c). 50–80μ long; unique movement. Fission (Kidder and Diller, 1934).

Genus *Urozona* Schewiakoff. Ovoid, both ends broadly rounded; a distinct constriction in the ciliated middle region; ciliary band composed of 5–6 rows of cilia, directed anteriorly and arranged longitudinally; cytostome with a membrane; rounded macronucleus and a micronucleus posterior; contractile vacuole subterminal; in fresh water.

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Fig. 326. a, *Dexiotrichides centralis*, ×500 (Kahl); b, *Cyrtolophosis mucicola*, ×670 (Kahl); c, *Urocentrum turbo*, ×200 (Bütschli); d, *Urozona bütschlii*, ×440 (Kahl); e, *Uronema marinum*, ×490 (Kahl); f, g, *U. pluricaudatum*, ×940 (Noland); h, *Homologastra setosa*, ×450 (Kahl); i, j, *Stokesia vernalis*, ×340 (Wenrich); k, *Ophyroglena collini*, ×150 (Lichtenstein); l, *O. pyriformis*, ×180 (Rossolimo); m, *O. intestinalis*, ×55 (Rossolimo).
HOLOTRICHA

U. bâttschlii S. (Fig. 326, d). 20–25μ long (Kahl); 30–40μ (Schewiakoff); in stagnant water.

Genus Uronema Dujardin. Oval to elongate ovoid; slightly flattened; anterior region not ciliated; inconspicuous peristome with ciliated right edge; cytostome on the ventral side close to left border in the anterior half, with a small tongue-like membrane; cytopharynx indistinct; macronucleus spherical, central; contractile vacuole terminal; in salt or fresh water. Comparison with Cyclidium (Párducz, 1940).

U. marinum D. (Fig. 326, e). 30–50μ long; in salt water among algae. Structure (Párducz, 1939).

U. pluricaudatum Noland (Fig. 326, f, g). Body appears to be twisted in dorsal view, due to a spiral depression that runs obliquely down toward cytostome; with about 8 caudal cilia; in salt water; Florida (Noland, 1937).

Genus Homalogastra Kahl. Broad fusiform; furrows spiral to left; a long caudal cilium; a group of cilia on right and left side of it; macronucleus spherical, anterior; contractile vacuole posterior; in fresh water.

H. setosa K. (Fig. 326, h). About 30μ long; fresh water.

Genus Stokesia Wenrich. Oblique cone with rounded angles; flat anterior surface uniformly ciliated; with peristome bearing zones of longer cilia, at the bottom of which is located the cytostome; a girdle of longer cilia around the organism in the region of its greatest diameter; pellicle finely striated; with zoochlorellae; trichocysts; free-swimming; in freshwater pond. One species (Wenrich, 1929).

S. vernalis W. (Fig. 326, i, j). 100–160μ in diameter; macronucleus; 2–4 micronuclei; fresh water.

Family 2 Ophryoglenidae Kent

Genus Ophryoglena Ehrenberg. Ellipsoidal to cylindrical; ends rounded or attenuated; preoral depression in form of ‘6’ due to an ectoplasmic membrane extending from the left edge, cilia on the right edge; cytostome deep-seated; 1 (or 2) contractile vacuole with long radiating canals, opens through pores on right ventral side; macronucleus of various forms with several endosomes; a micronucleus; fresh or salt water or parasitic. Many species.

O. collini Lichtenstein (Fig. 326, k). Pyriform; macronucleus horseshoe-shape; 200–300μ by 120–230μ; in the caecum of Baetis larvae.

*O. pyriformis* Rossolimo (Fig. 326, l). Flask-shape; 240–300\(\mu\) long; in the gastrovascular cavity of various Turbellaria.

*O. intestinalis* R. (Fig. 326, m). Up to 1.5 mm. by 450–500\(\mu\); smallest 60\(\mu\) long; in the gastrovascular cavity of *Dicotylus* sp.

*O. atra* Lieberkühn. Oval, posterior end broadly rounded; 300–500\(\mu\) long; grayish; filled with globules; cytostome near anterior end; macronucleus elongated; a contractile vacuole; trichocysts; stagnant fresh water.

Genus *Deltopylum* Fauré-Fremiet and Mugard. Cylindrical; uniform ciliation on about 70 ciliary rows; a triangular cytostome in the anterior fourth, with a paroral undulating membrane on right and three adoral membranes; a contractile vacuole on mid-right side, a pore being located in a depression of pellicle above it; macronucleus

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**Fig. 327.** a, *Deltopylum rhabdoïdes*, \(\times 665\) (Fauré-Fremiet and Mugard); b, *Pleuronema crassum*, \(\times 240\) (Kahl); c, *P. anodontae*, \(\times 290\) (Kahl); d, e, *P. setigerum*, \(\times 540\) (Noland); f, *P. coronatum*, \(\times 540\) (Noland); g, *P. marinum*, \(\times 400\) (Noland); h, *Cyclidium litomesum*, \(\times 300\) (Stokes); i, *Cristigera phoenix*, \(\times 500\) (Penard); j, *C. media*, \(\times 400\) (Kahl).
irregularly ribbon-like; five or six micronuclei; in fresh water (Fauré-Fremiet and Mugard, 1946).

D. rhabdoides F. and M. (Fig. 327, a). Cylindrical; 150–180µ by 40–45µ; anterior end slightly attenuated and curved, posterior end rounded; the organism grows well on the gut of Chironomus larvae in laboratory.

Family 3 Pleuronematidae Kent

Genus Pleuronema Dujardin. Ovoid to ellipsoid; peristome begins at anterior end and extends for 2/3 the body length; a conspicuous membrane at both edges; semicircular swelling to left near oral area; no cytopharynx; close striation longitudinal; one to many posterior sensory stiff cilia; macronucleus round or oval; a micronucleus; a contractile vacuole; trichocysts in some species; fresh or salt water, also commensal in freshwater mussels.

P. crassum D. (Fig. 327, b). 70–120µ long; somewhat compressed; Woods Hole (Calkins).

P. anodontae Kahl (Fig. 327, c). About 55µ long; posterior cilium about 1/2 the body length; in Sphaerium, Anodonta.

P. setigerum Calkins (Fig. 327, d, e). Ellipsoid; flattened; ventral surface slightly concave; about 25 ciliary rows; 38–50µ long (Noland); in salt water; Massachusetts, Florida.

P. coronatum Kent (Fig. 327, f). Elongate ovoid; both ends equally rounded; caudal cilia long; about 40 ciliary rows; 47–75µ long (Noland, 1937); in fresh and salt water; Florida.

P. marinum D. (Fig. 327, g). Elongate ovoid; trichocysts distinct; caudal cilia medium long; about 50 ciliary rows; 51–126µ long (Noland); in salt water; Florida.

Genus Cyclidium Müller. Small, 15–60µ long; ovoid; usually with refractile pellicle; with a caudal cilium; peristome near right side; on its right edge occurs a membrane which forms a pocket around cytostomal groove and on its left edge either free cilia or a membrane which unites with that on right; no semicircular swelling on left of oral region; round macronucleus with a micronucleus; contractile vacuole posterior; fresh or salt water. Numerous species. 3 species in sea urchin (Powers, 1935); comparison with Uronema (Párducz, 1940).

C. litomesum Stokes (Fig. 327, h). About 40µ long; dorsal surface slightly convex with a depression in middle; ventral surface more or less concave; cilia long; in fresh water.

Genus Cristigera Roux. Similar to Cyclidium; much compressed;
with a postoral depression; peristome closer to mid-ventral line; fresh or salt water. Several species.

*C. phoenix* Penard (Fig. 327, i). 35–50μ long; fresh water.

*C. media* Kahl (Fig. 327, j). 45–50μ long; in salt water.

Genus *Ctedoctema* Stokes. Similar to *Cyclidium* in body form; peristome nearer median line, diagonally right to left; right peri-

stomal ridge with a sail-like membrane which surrounds the cytostome at its posterior end; trichocysts throughout; fresh water.

*C. acanthocrypta* S. (Fig. 328, a). Ovoid; anterior end truncate; macronucleus round, anterior; about 35μ long; in fresh water among vegetation.

Genus *Calyptotricha* Phillips. Somewhat resembles *Pleuronema* or *Cyclidium*; but dwelling in a lorica which is opened at both ends; with zoochlorellae; fresh water.

*C. pleuronemoides* P. (Fig. 328, b). Lorica about 85μ high; body
about 50µ long; Kellicott's (1885) form is more elongated; in fresh water.

Genus **Histiobalantium** Stokes. Ovoid; ventral side flattened; ciliation uniform; long stiff cilia distributed over the body surface; peristome deep; both anterior and posterior regions with a well-developed membrane, connected with the undulating membrane; macronucleus in 2 parts; 1–2 micronuclei; several contractile vacuoles distributed; fresh water.

*H. natans* (Claparède and Lachmann) (Fig. 328, c). 70–110µ long.

*H. semisetatum* Noland (Fig. 328, d). Elongate ellipsoid; posterior end bluntly rounded; macronucleus spherical; longer cilia on posterior half only; contractile vacuoles on dorsal side; 126–205µ long; salt water; Florida (Noland, 1937).

Genus **Pleurocoptes** Wallengren. Ovoid, dorsal side hemispherical, ventral side flattened; peristome large, reaching the posterior 1/3; cytopharynx indistinct; longer cilia along peristome; macronucleus spherical; several micronuclei; contractile vacuole terminal; ectocommensal.

*P. hydractiniae* W. (Fig. 328, e). 60–70µ long; on *Hydractinia echinata*.

**Family 4 Cohnilembidae** Kahl

Genus **Cohnilembus** Kahl (*Lembus* Cohn). Slender spindle-form; flexible; peristome from anterior end to the middle of body or longer, curved to right, with 2 membranes on right edge; a caudal cilium or a few longer cilia at posterior end; macronucleus oval, central; in salt or fresh water, some parasitic.

*C. fusiformis* (C.) (Fig. 328, f). Striation spiral; peristome about 1/6 the body length; a few cilia at posterior end; oval macronucleus central; contractile vacuole posterior; about 60µ long; in fresh water.

*C. caeci* Powers (Fig. 328, g). About 32–92µ long; in the intestine of *Tripneustes esculentus* and other echinoids; Tortugas.

**Family 5 Philasteridae** Kahl

Genus **Philaster** Fabre-Domergue (*Philasterides* Kahl). Body cylindrical; peristome about 1/3–2/5 the body length, broader near cytostome and with a series of longer cilia; cytostome with a triangular membrane; cytopharynx (?); ciliation uniform; a caudal cilium; trichocysts; oval macronucleus with a micronucleus, central; contractile vacuole terminal or central; in salt or fresh water.

*P. digitiformis* F–D. (Fig. 328, h). Anterior region bent dorsally; contractile vacuole terminal; 100–150µ long; salt water.
P. armata (K.) (Fig. 328, i). Anterior end more or less straight; peristome difficult to see; contractile vacuole central; 70–80μ long; fresh water.

Genus Helicostoma Cohn. Similar to Philaster in general appearance; preoral side-pouch curved around posterior edge of peristome and separated from it by a refractile curved band; with or without a pigment spot near cytostome; macronucleus oval or band-form; contractile vacuole terminal; in salt water.

H. buddenbrocki Kahl (Fig. 328, j). 130–200μ long; in salt and brackish water.

References


——— and Diller, W. F.: (1934) Observations on the binary fission of four species of common free-living ciliates, etc. Ibid., 67: 201.


THE majority of the ciliates placed in this suborder are parasites or commensals of molluscs. They possess thigmotactic cilia with which they attach themselves to the host body. Though appearing heterogeneous, Chatton and Lwoff (1949) maintain that there is a phylogenetic unity among them, which condition has been brought about by degenerative influence because of similar conditions of habitat. Taxonomy (Jarocki and Raabe, 1932; Chatton and Lwoff, 1949).

Following Chatton and Lwoff (1939), the suborder is here divided into seven families:

- **Family 1 Conchophthiridae**
- **Family 2 Thigmophryidae** (p. 776)
- **Family 3 Hemispeiridae** (p. 776)
- **Family 4 Hysterocinetidae** (p. 779)
- **Family 5 Ancistrocomidae** (p. 780)
- **Family 6 Hypocomidae** (p. 784)
- **Family 7 Sphenophryidae** (p. 785).

**Family 1 Conchophthiridae** Reichenow

Genus *Conchophthirus* Stein. Oval to ellipsoidal; flattened; right margin concave at cytostomal region, left margin convex; ventral surface somewhat flattened, dorsal surface convex; cytostome on right side near middle in a depression with an undulating membrane; macronucleus; micronucleus; contractile vacuole opens through a canal to right side; in the mantle cavity and gills of various mussels. Species (Kidder, 1934, 1934a; Uyemura, 1934, 1935); morphology (Raabe, 1932, 1934; Kidder, 1934).

*C. anodontae* ( Ehrenberg) (Fig. 329, a). Ovoid; cytostome in anterior third, with an overhanging projection in front; cytopharynx, surrounded by circular fibrils, continues down as a fine, distensible tubule, to near the macronucleus; with peristomal basket; ciliary grooves originate in a wide ventral suture near anterior end; anterior region filled with refractile granules; macronucleus posterior; contractile vacuole between nuclei and peristome, with a slit-like aperture (Fig. 27); 65–125μ by 47–86μ; in the mantle cavity, gills and on non-ciliated surface of palps of *Elliptio complanatus*; Woods Hole.
Fig. 329. a, Concophthirus anodontae; b, C. magna, ×300 (Kidder); c, Myxophyllum steenstrupi, ×280 (Raabe); d, Hemispeira asteriasi, ×705 (Wallengren); e, f, Protophrya ovicola (Cépède) (f, a young Littorina rudis with the ciliate); g, h, two views of Ancistruma mytili, ×500 (Kidder); i, A. isseli, ×500 (Kidder); j, A. japonica, ×600 (Uyemura); k, Eupoteron pernix, ×500 (MacLennan and Connell); l, Ancistrina ovata, ×630 (Cheissin).

C. magna Kidder (Fig. 329, b). Much larger; 123–204μ by 63–116μ; closer ciliation; anterior 1/3 filled with smaller granules; irregularly outlined macronucleus, 25–30μ in diameter, central; 2 (or 1) micronuclei; aperture for contractile vacuole large; mantle cavity of Elliptio complanatus; Massachusetts.
C. *mytili* de Morgan (Fig. 56). Reniform; 130–220μ by 76–161μ; peristomal groove on the right side; trichocysts conspicuous along frontal margin; macronucleus oval; 2 micronuclei; on the foot of the common mussel, *Mytilus edulis*. Division and conjugation (Kidder, 1933b, c).

Genus *Myxophyllum* Raabe. Oval or spheroid; pellicle elastic and flexible; peristome on posterior right, without undulating membrane; 7 macronuclei; a micronucleus; ciliation uniform; in the slime covering land pulmonates.

*M. steenstrupi* (Stein) (Fig. 329, c). 120μ by 100–120μ; on *Succinea putris*, etc.

**Family 2 Thigmophryidae** Chatton and Lwoff

Genus *Thigmophrya* Chatton and Lwoff. Elongate; round or oblong in cross section; cytostome in posterior third; contractile vacuole opens in cytopharynx; on the gills or palps of lamellibranchs.

*T. macomae* C. and L. Elongate ovoid; flattened; ventral surface slightly concave; oral funnel opened; contractile vacuole opens at the bottom of cytopharynx; numerous ciliary rows; about 110μ by 40μ; on the gills of *Macoma* (*Tellina*) *balthica* (Chatton and Lwoff, 1923).

**Family 3 Hemispeiridae** König

Genus *Hemispeira* Fabre-Domergue (*Hemispeiropsis* König). Nearly spherical; flattened; longitudinal non-ciliated furrow on ventral surface, which encircles thigmotactic posterior cilia; 4–5 cross-furrows of cilia: a huge adoral membrane at anterior end; macronucleus, micronucleus large; contractile vacuole, anterior-right; commensal.

*H. asteriasi* F.–D (Fig. 329, d). 20–30μ long; ectocommensal on *Asterias glacialis* (Wallengren, 1895).

Genus *Protophrya* Kofoid (*Isselina* Cépède). Ellipsoid to pyriform; spherical macronucleus; cytostome close to the posterior end. Taxonomy (Raabe, 1949); ciliation (Chatton and Lwoff, 1949).

*P. ovicola* K. (Fig. 329, e, f). About 60μ long; in the uterus and brood-sac of the molluscs, *Littorina rudis* and *L. obtusata* (Kofoid, 1903).

Genus *Ancistruma* Strand (*Ancistrum* Maupas). Ovoid, pyriform or somewhat irregular; flattened; right side with more numerous large cilia than the left; peristome on right side; cytostome near posterior extremity; macronucleus round or sausage-shape, central; a micronucleus; contractile vacuole posterior; commensal in the mantle cavity of various marine mussels. Many species. Morphology, reproduction (Kidder, 1933, 1933a).
A. *mytili* (Quennerstedt) (Figs. 18; 329, g, h). Oval; dorsal surface convex, ventral surface concave; dorsal edge of peristome curves around the cytostome; peristomial floor folded and protruding; longitudinal ciliary rows on both surfaces; three rows of long cilia on peristomial edges; macronucleus sausage-form; a compact micronucleus anterior; 52–74μ by 20–38μ. Kidder (1933) found it in abundance in the mantle cavity of *Mytilus edulis* at Woods Hole and New York.

A. *isseli* Kahl (Fig. 329, i). Bluntly pointed at both ends; 70–88μ by 31–54μ. Kidder (1933) observed it abundantly in the mantle cavity of the solitary mussel, *Modiolus modiolus*, Massachusetts and New York, and studied its conjugation and nuclear reorganization.

A. *japonica* Uyemura (Fig. 329, j). Body oval or elongate pyriform; 55–76(67)μ by 14–29(20)μ; subspherical macronucleus conspicuous; a compact micronucleus; usually a single contractile vacuole, posterior; in the mantle cavity of marine mussels; *Meritrix meritrix*, *Paphia philippinarum*, *Cyclina sinensis*, *Mactra veneriformis*, *M. sulcataria*, and *Dosinia bilinulata* (Uyemura, 1937).

Genus *Eupoterion* MacLennan and Connell. Small ovoid; slightly compressed; cilia short, in longitudinal rows; rows of long cilia in peristome on mid-ventral surface and extend posteriorly, making a half turn to left around cytostome; small conical cytostome lies in postero-ventral margin of body; contractile vacuole terminal; large round macronucleus anterior; a micronucleus; commensal.

*E. pernix* M. and C. (Fig. 329, k). 46–48 ciliary rows; 6 rows of heavy peristomial cilia; 38–56μ long; in the intestinal contents of the mask limpet, *Acmaea persona*; California.

Genus *Ancistrina* Cheissin. Ovoid; anterior end attenuated; peristomial field along narrow right side; 15–18 ciliary rows parallel to peristomial ridges; cytostome right-posterior, marked with oral ring, with a membrane and a zone of membranellae; right ridge of peristome marked by two adoral ciliary rows; macronucleus anterior, spheroidal; a micronucleus; commensal.

A. *ovata* C. (Fig. 329, l). 38–48μ by 15–20μ; in the mantle cavity of molluscs: *Benedictia biacalensis*, *B. limneoides* and *Choanomphalus* sp.

Genus *Cochliophilus* Kozloff. Ovoid and compressed; peristome in right-posterior fourth of the body; membrane-like fine cilia overlie a series of thick cilia from the anterior end of the peristome to cytostome; longitudinal rows of cilia; a vesicular micronucleus; an ovoid macronucleus; a contractile vacuole; in molluscs.

*C. depressus* K. (Fig. 330, a). About 93μ by 63μ by 15μ; 52–56
ciliary rows; peristomal membranous cilia motile; macronucleus oblong; in the mantle cavity of the pulmonate snail, *Phytia setifer* in San Francisco Bay (Kozloff, 1945).

Genus Ancistrella Cheissin. Elongate; ends rounded; ventral surface less convex than dorsal surface; 16–17 longitudinal ciliary rows; ciliation uniform, except anterior-dorsal region, bearing bristle-like longer cilia; 2 adoral ciliary rows on right of peristome, curved dorsally behind cytostome; contractile vacuole posterior; macronucleus single or divided into as many as 7 parts; micronucleus; commensal.

*A. choanomphali* C. (Fig. 330, b). 55–90µ by 18–20µ; in the mantle cavity of *Choanomphalus* sp.

![Fig. 330. a, Cochliophilus depressus, X600 (Kozloff); b, Ancistrella choanomphali, X840 (Cheissin); c, Boveria teredinidi, X550 (Pickard); d, Plagiospira crinita, X740 (Issel); e, Hysterocineta eiseniae, X250 (Beers); f, Ptychostomum bacteriophilum, X500 (Miyashita).](image)
Genus *Ancistrospira* Chatton and Lwoff. Ciliation meridional to spiral; peristome right spiral; commensal.

*A. veneris* C. and L. 50–60μ by 22–28μ; ovoid, anterior end pointed; ciliary rows meridional; thigmotactic field on the left side, sharply marked from body ciliation; on the gills of *Venus fasciata*.

Genus *Boveria* Stevens (*Tiarella* Cheissin). Conical; cytostome at posterior end; peristome spiral posteriorly; macronucleus oval, anterior half; a micronucleus; contractile vacuole posterior; ectocommensal on gills of various marine animals such as Teredo, Bankia, Tellina, Capsa and Holothuria. Several species.

*B. teredinidi* Pickard (Fig. 330, c). 27–173μ by 12–31μ; on gills of *Teredo navalis*; California (Pickard, 1927).

Genus *Plagiospira* Issel. Conical; anterior end attenuated; peristome runs spirally from middle of body to cytostome, with long cilia; macronucleus oval, anterior; a micronucleus; contractile vacuole near middle of body; somewhat spirally arranged striae widely apart on right side; commensal.

*P. crinita* I. (Fig. 330, d). 32–58μ by 18–34μ; in *Cardita calyculata* and *Loripes lacteus*.

Family 4 *Hysterocinetidae* Diesing

Inclusion of this family in the present suborder is provisional, since its affinity to other forms is not yet clear. Beers (1938) who placed it in Hymenostomata, in agreement with Cheissin (1931), states that the nutrition is in part saprozoic, and that the organisms are in the process of acquiring the saprozoic and astomatous condition.

Genus *Hysterocineta* Diesing (*Ladopsis* Cheissen). Elongate; flattened; flexible, an inverted V- or U-shaped sucker conspicuously present in antero-ventral margin; ciliation uniform; cytostome and cytopharynx at the posterior end; an undulating membrane along peristome which borders the posterior margin of body; macronucleus elongate; a micronucleus; contractile vacuole posterior; in the intestine of gastropods and oligochaetes. 4 species. Taxonomy (Jarocki, 1934; Beers, 1938; Raabe, 1949).

*H. eiseniae* Beers (Fig. 330, e). 190–210μ by 35–40μ; cytostome not functional; endoplasm with small granules; macronucleus 45–50μ long; sucker inverted V, about 25–30μ long; in the intestine of *Eisenia lönnerbergi* (Beers, 1938).

Genus *Ptychostomum* Stein (*Lada* Vejdovsky). Sucker circular or ovoid; macronucleus ovoid or reniform, not elongate; in oligochaetes. Several species. Taxonomy (Studitsky, 1932; Raabe, 1949).
P. bacteriophilum Miyashita (Fig. 330, f). Elongate oval; 70–130μ by 30–45μ; sucker oval and large, about 50μ in diameter; macronucleus ellipsoid; endoplasm with numerous rods (symbiotic bacteria?); in the freshwater oligochaete, Criodrilus sp.

Family 5 Ancistrocomidae Chatton and Lwoff

Genus Ancistrocoma C. and L. (Parachaenia Kofoid and Bush). Elongate pyriform with attenuated anterior end; somewhat flattened dorso-ventrally; a contractile suctorial tentacle at the anterior tip, which is used for attachment to the epithelium of host, and which continues internally as a long curved canal; longitudinal ciliation on dorso-lateral and ventral sides, beginning at the anterior end; parasitic in the gills and palps of mollusks. Taxonomy (Kozloff, 1946b; Chatton and Lwoff, 1950).

A. pelseneeri C. and L. (Parachaenia myae Kofoid and Bush) (Fig. 331, a). Body 50–83(62)μ by 14–20(16)μ by 11–16(12.5)μ; 14 ciliary rows on dorso-lateral and ventral surfaces; five rows on the ventral side extend only 2/3 from the anterior end; tentacle continues internally for about 2/3 of body, curved; macronucleus sausage-shaped; a single micronucleus; on the gills and palps of mussels: Mya arenaria, M. irus, M. inconspicua, M. nasuta, M. secta, Cryptomya californica (Kozloff, 1946b).

Genus Hypocomagalma Jarocki and Raabe. Ovoid or pyriform with attenuated anterior end; asymmetrical; 22–24 ciliary rows which do not reach the posterior end; a suctorial tentacle at the anterior end; on mollusks.

H. pholadidis Kozloff (Fig. 331, b). 63–89μ by 18–25μ by 16–21μ; anterior end bent ventrally; 24 or 25 ciliary rows; one or more contractile vacuoles; macronucleus sausage-shaped; a single micronucleus; parasitic in the epithelium of the gills and palps of Pholadidea penita (Kozloff, 1946b).

Genus Syringopharynx Collin. Elongate ovoid, narrowed anteriorly; a suctorial tentacle at anterior end; 14 ciliary rows (six dorsal, six ventral and two lateral); on molluscs (Collin, 1914).

S. pterotrachae C. Body 55μ by 25μ; macronucleus elongate band; on the gills of Pterotracha coronata (Chatton and Lwoff, 1950).

Genus Goniocoma Chatton and Lwoff. Ovoid with attenuated anterior end; end of suctorial tentacle extremely slender; 27–29 ciliary rows; of the 14 dorsal rows, the median row is very short and the rows on either side of it are progressively longer; ventral rows pass over the posterior end and terminate on dorsal surface; on the gills of molluscs.
Fig. 331, a, ventral view of a stained Ancistrocoma pelseneeri, $\times 1120$ (Kozloff); b, Hypocomagalma pholadidis, $\times 840$ (Kozloff); c, ciliate as viewed from right side of Holocoma primigenius, $\times 1130$ (Chatton and Lwoff); d, ventral view of Insignicoma venusta, $\times 1245$; e, Raabella botulae, $\times 1245$; f, Crebricoma kozloffi, $\times 755$ (Kozloff).


Genus *Holocoma* Chatton and Lwoff. Cylindrical; ventral surface convex; tentacle at anterior end; 19–23 ciliary rows; 6–10 median dorsal rows relatively short, seven left and six right rows long; on the gills of mollusks.

*H. primigenius* C. and L. (Fig. 331, c). Elongated body 41–59$\mu$ by 15$\mu$; ventral surface convex; elongate macronucleus; on the gills of *Macoma balthica* (Chatton and Lwoff, 1950).
Genus **Insignicoma** Kozloff. Elongate pyriform; a contractile tentacle with internal canalicule; median ciliary rows on anterior half of ventral surface; two right ciliary rows; left rows short and closely set; an inverted V-shaped row of long cilia on left-lateral surface at about the middle of body; on mollusks.

*I. venusta* K. (Fig. 331, d). 42–52 μ by 18–21 μ by 15–18 μ; 15 median, two right, and 16–17 left ciliary rows; macronucleus ovoid; micronucleus spherical; on the gills and palps of *Botula californiensis* (Kozloff, 1946a).

Genus **Raabella** Chatton and Lwoff. Three groups of ciliary rows; eight to 11 short median rows; six to 11 longer rows on left-lateral side; two longer rows on the right side; on mollusks.

*R. botulae* (Kozloff) (Fig. 331, e). 31–39 μ by 14–17 μ by 12–14 μ; 11 median rows; 11 closely set left rows; two longer right rows; macronucleus ovoid to sausage-shaped; spherical micronucleus; on the gills and palps of *Botula californiensis* (Kozloff, 1946a).

Genus **Crebricoma** Kozloff. Pyriform; anterior suctorial tentacle; many ciliary rows, the majority of which are closely set; two long rows on the right side; anterior terminals of the rows make a V-shaped suture; on the gills of mollusks.

*C. kozloffii* Chatton and Lwoff (*C. carinata* K.) (Fig. 331, f). Body 58–71 μ by 27–39 μ by 22–31 μ; two ciliary rows on right side long, about 2/3 the body length; more than 30 rows of closely set cilia (1/2–2/3 the body length and longer toward left); macronucleus ellipsoid; on the gills and palps of *Mytilus edulis* (Kozloff, 1946; Chatton and Lwoff, 1950).

Genus **Hypocomides** Chatton and Lwoff. Elongate; some 23 ciliary rows; about 20 median rows, short; two longer rows on right; a short curved row near the posterior end; on mollusks.

*H. mediolariae* C. and L. (Fig. 332, a). 27–50 μ by 15–27 μ; on the gills of *Mediolaria marmorata* (Chatton and Lwoff, 1922).

Genus **Anisocomides** Chatton and Lwoff. Body ovoid, slightly flattened; 12 ciliary rows; two short median rows with five additional rows which are progressively longer toward left; a short oblique row, posterior to the outermost left row; four right rows much longer; on the gills of mollusks.

*A. zyrphea* (C. and L.) (Fig. 332, b). 19–38 μ by 10–15 μ by 7–10 μ; on the gills of *Pholas (Zyrphea) crispata* (Chatton and Lwoff, 1926).

Genus **Hypocomatidium** Jarocki and Raabe. Similar to **Anisocomides**, but without the short posterior ciliary row; on the gills of mollusks.
Fig. 332, a, Hypocomides mediolariae, ×1000; b, left side view of Anisocomides zyrpheae in life, ×1065; c, Isocomides mytili in life, ×1000 (Chatton and Lwoff); d, Hypocomina tegularum, ×1245; e, Heterocinetopsis goniobasidis, ×1145; f–h, Hypocomella phoronopsis, ×1300 (f, ventral view of a stained specimen; g, h, dorsal and right side views in life); i, Enerthecoma kozloffi, ×1145 (Kozloff).

H. sphaerii J. and R. Ovoid; 30–45 μ by 14–18 μ by 9–12 μ; nine ciliary rows; five rows on left-ventral and four on right; on the gills of Sphaerium corneum and S. rivicola (Jarocki and Raabe, 1932).

Genus Isocomides Chatton and Lwoff. Elongated; 14–18 ciliary rows on anterior 2/3 of the ventral surface; six to seven on right and eight to 11 on left; in addition, there is a short transverse row with a dozen long cilia, posterior to other rows; on mussels.

I. mytili (C. and L.) (Fig. 332, c). 57–64 μ by 20–22 μ; on the gills of Mytilus edulis (Chatton and Lwoff, 1922).

Genus Hypocomina Chatton and Lwoff. Ovoid to pyriform; an
anterior tentacle; eight to 10 ciliary rows about half the body-length and starting a little distance away from the anterior tip; on mollusks.

_H. tegularum_ Kozloff (Fig. 332, d). 26–36μ by 12–17μ by 9–12μ; anterior end bent ventrally; nine ciliary rows, five rows on right being longer than the other four; spherical macronucleus; parasitic on the ctenidium of _Tegula brunnae_ (Kozloff, 1946).

Genus _Heterocinetopsis_ Jarocki. Body elongate, flattened dorso-ventrally; a contractile tentacle, its canalicule extending 1/3–2/3 the body length; 10–12 ciliary rows; the median rows about one-half the body length, the rows toward left being progressively longer; on mollusks (Jarocki, 1935).

_H. goniobasidis_ (Kozloff) (Fig. 332, e). 36–48μ by 15–20μ by 11–14μ; 10 ciliary rows; macronucleus pyriform; ovoid micronucleus inconspicuous; parasitic on the epithelium of the gills and mantle of _Goniobasis plicifera silicula_ (Kozloff, 1946c).

Genus _Hypocomella_ Chatton and Lwoff (Hypocomidium Raabe). Pyriform, asymmetrical, flattened; a long retractile tentacle; seven to 13 ciliary rows on the ventral surface, three rows on left being progressively longer; on mollusks (Chatton and Lwoff, 1922, 1950).

_H. phoronopsidis_ (Kozloff) (Fig. 332, f–h). 26–37μ by 11–16μ by 6.5–11μ; eight ventral ciliary rows; ovoid macronucleus and micronucleus; on the tentacles of _Phoronopsis viridis_ (Kozloff, 1945a).

Genus _Enerthecoma_ Jarocki. Pyriform, symmetrical; 8 ciliary rows on the ventral side; three on left are somewhat separated from five others and closely set; on the gills of mollusks.

_E. kozloffi_ Chatton and Lwoff (Fig. 332, i). 32–56μ by 13–21μ by 10–13μ; eight ciliary rows about 2/3 the body length; macronucleus elongate; micronucleus fusiform; on the gills of _Viviparus fasciatus_ and _V. malleatus_ (Kozloff, 1946c; Chatton and Lwoff, 1950).

Genus _Cepedella_ Poyarkoff. Pyriform with a pointed anterior end; macronucleus globular; without contractile vacuole.

_C. hepatica_ P. Body 16–26μ long; in the liver of _Sphaerium cornueum_.

Family 6 _Hypocomidae_ Bütschli

Genus _Hypocoma_ Grüber. Dorsal side convex; ventral side flattened with a ciliated oval field; a suctorial tentacle at the anterior end; about 13 ciliary rows; an adoral zone, a short row (eight granules) at anterior-left side; on colonial Protozoa.

_H. parasitica_ G. (Fig. 333, a). 30–38μ by 18–20μ by 18μ; 13 ciliary rows on the flattened surface: adoral zone, a short row; 11 general
ciliary rows; macronucleus horseshoe-shape; a large central food vacuole; on solitary or colonial peritrichs such as Vorticella, Zoo-thamnium, etc. (Chatton and Lwoff, 1950).

Genus Heterocoma Chatton and Lwoff. Body ovoid; ventral side flattened; suctorial tentacle antero-ventral; 13 ciliary rows make an ellipsoidal field; an adoral zone, five closely-set rows on left and seven widely separated rows on right; in the branchial cavity of Salpa (Chatton and Lwoff, 1939).

H. hyperparasitica C. and L. (Fig. 333, b). Body ovoid, with bluntly pointed posterior end; about 44 μ long; a large food vacuole in cytoplasm; in the branchial cavity of Salpa mucronata-democratica (Chatton and Lwoff, 1950).

Genus Parahypocoma Chatton and Lwoff. Ellipsoid; highly flattened; anterior end tapers slightly; 29–34 ciliary rows; the adoral zone as in the other two genera; a comparatively short suctorial tentacle at anterior end; macronucleus horseshoe-shaped; a large central food vacuole; parasitic in ascidians.

P. collini C. and L. (Fig. 333, c). In Ascidia mentula and Ciona intestinalis (Chatton and Lwoff, 1950).

Family 7 Sphenophryidae Chatton and Lwoff

Genus Sphenophrya Chatton and Lwoff. Body elongated, “quarter orange-” or banana-shaped; attached to the gills of host mollusks
by a suctorional tentacle; adult stage without cilia; ciliature is reduced to infraciliature of 2 groups; multiplication by budding; embryos are ciliated; on the gills of mollusks (Chatton and Lwoff, 1921).

*S. dosiniae* C. and L. (Fig. 334, a–c). Body 120μ by 15–20μ; young embryo ciliated; on the gills of *Dosinia exoleta, Venus ovata, Corbula gibba*, etc. (France); *Mactra solidissima* (Woods Hole) (Chatton and Lwoff, 1950).

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**Fig. 334, a–c, Sphaenophrya dosiniae** (a, a young embryo; b, a growing individual attached to an epithelial cell of the host by a suctorional tentacle; c, an individual from which a bud is ready to separate); d, a side view of *Pelecyophrya tapetis* in life; e, f, *Gargarius gargarius*, ×1200 (e, in life, showing a macronucleus and a micronucleus; f, diagram showing the ciliature) (Chatton and Lwoff).

Genus *Pelecyophrya* Chatton and Lwoff. Body hatchet-shaped, laterally compressed; posterior end rounded; a large "sucker" at the anterior end; infraciliature in two groups, five on right and four on left; multiplication by budding; on the gills of mollusks (Chatton and Lwoff, 1922).

*P. tapetis* C. and L. (Fig. 334, d). Body 23–25μ by about 10μ; macronucleus spherical; ovoid micronucleus; cytoplasm contains
fragments of host cells including nuclei; conjugation; on the gills of *Tapes aureus* (Chatton and Lwoff, 1950).

Genus **Gargarius** Chatton and Lwoff. Dorso-ventrally flattened; with a "horn" near the anterior end; sucker occupies the entire ventral surface, its margin showing papillous extensions; on the ventral surface there are two groups of ciliature; four rows on each side; on *Mytilus* (Chatton and Lwoff, 1950).

*G. gargarius* C. and L. (Fig. 334, *e, f*). Body about 35μ long; ciliated embryos formed by budding or unequal division; macronucleus elongate; micronucleus spherical; on *Mytilus edulis* (Chatton and Lwoff, 1950).

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Order 1 Holotricha Stein (continued)

Suborder 6 Apostomea Chatton and Lwoff

ASYMMETRICAL forms with a rosette-like cytostome through which liquid or small solid particles are taken into the body; sparse ciliary rows spiral; adoral rows short; macronucleus oval to band-form; a micronucleus; a single contractile vacuole.

The life-cycle of the ciliates grouped here appears to be highly complex and Chatton and Lwoff (1935) distinguished several developmental phases (Fig. 335), as follows: (1) Trophont or vegetative phase: right-spiral ciliary rows; nucleus pushed aside by food bodies; body grows, but does not divide. (2) Protomont: transitory stage between 1 and 3 in which the organism does not nourish itself, but produces "vitelloid" reserve plates; nucleus central, condensed; ciliary rows become straight. (3) Tomont: the body undergoes division usually in encysted condition into more or less a large number of small ciliated individuals. (4) Protomite: a stage in which a renewed torsion begins, and which leads to tomite stage. (5) Tomite: small free-swimming and non-feeding stage, but serves for distribution. (6) Phoront: a stage which is produced by a tomite when it becomes attached to a crustacean and encysts; within the cyst a complete transformation to trophont takes place.

Family Foetingeriidae Chatton

Genus Foetingeria Caullery and Mesnil. Trophonts large, up to 1 mm. long; sublenticular, anterior end attenuated; dorsal surface convex, ventral surface concave; right side less convex than left side; 9 spiral ciliary rows nearly evenly arranged; in gastrovascular cavity of various actinozoans; tomont on outer surface of host body, gives rise to numerous tomites with meridional ciliary rows; each tomite becomes a phoront by encysting on a crustacean, and develops into a trophont when taken into gastrovascular cavity of an actinozoan. One species.

F. actiniarum (Claparède) (Fig. 336, a). Phoronts on Copepoda, Ostracoda, Amphipoda, Isopoda and Decapoda; trophonts in Actinia mesembryanthemum, A. equina, Anemonea sulcata and other actinozoans in European waters; Chatton and Lwoff found Metridium marginatum, Sagartia leucolena and Astrangia danae of Woods Hole free from this ciliate.

Genus Spirophrya Chatton and Lwoff. Trophonts ovoid, pointed anteriorly; 16 uninterrupted ciliary rows of which striae 1 and 2 ap-
Fig. 335. Diagram illustrating the life-cycle of *Spirophrya subparasitica* (Chatton and Lwoff).

proach each other in posterior-dorsal region; phoronts attached to a crustacean; when eaten by Cladonema, trophonts enter the crustacean body and complete growth; protomonts upon leaving the host body encyst and each divides into 4–82 tomites (Fig. 335). One species.

*S. subparasitica* C. and L. (Figs. 335; 336, b). Phoronts attached to *Idyaea furcata*; ovoid trophonts enter the copepod when eaten by *Cladnema radiatum.*
Genus **Gymnodinioides** Minkiewicz (*Physosphaga* Percy; *Oospira* Chatton and Lwoff). Trophonts twisted along equatorial plane; generally 9 ciliary rows, in some a rudimentary row between striae 5 and 6 at anterior end. Many species.

*G. calkinsi* Chatton and Lwoff. Phoronts on gills and trophonts in the moult of *Palaemonetes* sp.; Woods Hole.

Genus **Phoretrophyra** Chatton and Lwoff. Trophonts generally with 9 ciliary rows; striae 1, 2, and 3, close to one another. One species.

Fig. 336. a, *Foettingeria actiniarum*, a trophont; b, *Spirophrya subparasitica*, a trophont, ×1000; c, *Phoretrophyra nebaliae*, ×1180; d, *Synophrya hypertrophica* (Chatton and Lwoff).
P. nebaliæ C. and L. (Fig. 336, c). Phoronts and tomonts on appendages, and trophonts in the moult, of Nebalia geoffroyi.

Genus Synophrya Chatton and Lwoff. Trophonts and tomonts similar to those of Gymnodinioides; but development highly complicated. One species.

S. hypertrophica C. and L. (Fig. 336, d). Phoronts in branchial lamellae, and trophonts in the moult, of Portunus depurator, etc.
Fig. 338. a, Polyspira delagei; b, Calospira minkiewiczi, a trophont, ×1300; c, Vampyrophrya pelagica; d, Traumatiophlora punctata, ×1300 (Chatton and Lwoff).

Genus Ophiurespira Chatton and Lwoff. Trophonts ovoid; 10 ciliary rows; striae 9 and 10 interrupted. One species.

O. weilli C. and L. (Fig. 337, a). Trophonts in the intestine of Ophiothrix fragilis and Amphiura squamata (Ophiuroidea).

Genus Photorophrya Chatton and Lwoff. Trophonts small; ciliation approximately that of Ophiurespira; massive macronucleus; with peculiar trichocysts comparable with the nematocysts of Polykrikos (p. 324); ecto- or endo-parasitic in encysted stages of other apostomes. Several species.
PROTOZOOLOGY

*P. insidiosa* C. and L. (Fig. 337, b). Phoronts, trophonts and tomites in phoronts of *Gymnodinioides*.

Genus *Polyspira* Minkiewicz. Trophonts reniform; 9 rows and several extra rows; striae 1–4 and 5–9 with 2 others in 2 bands.

*P. delagei* M. (Fig. 338, a). Phoronts on gills and trophonts in the moult of *Eupagurus berhardus*.

Genus *Pericaryon* Chatton. Trophonts ellipsoid; 14 ciliary rows.

*P. cesticola* C. (Fig. 337, d). Trophonts in the gastrovascular cavity of the ctenophore, *Cestus veneris*; other stages unknown.

Genus *Calospira* Chatton and Lwoff. Trophonts resemble those of *Spirophrya*; 20 ciliary rows; macro nucleus twisted band-form; a micronucleus.

*C. minkiewiczii* C. and L. (Fig. 338, b). Phoronts attached to integument of *Harpacticus gracilis* (copepod); trophonts in its fresh carcass; tomonts and tomites in water.

Genus *Vampyrophrya* Chatton and Lwoff. Trophonts ovoid; 10 ciliary rows. One species.

*V. pelagica* C. and L. (Fig. 337, c; 338, c). Phoronts on *Paracalanus parous, Clausocalanus furcatus*, etc., and trophonts in their fresh carcasses.

Genus *Traumatiphtora* Chatton and Lwoff. Trophonts oval; 11 ciliary rows. One species.

*T. punctata* C. and L. (Fig. 338, d). Trophonts in fresh carcass of *Acartia clausi*.

Genus *Hyalospira* Miyashita. Trophonts in the moult of a freshwater crustacean, with a contractile vacuole and a long accessory canal, and with a band-shaped macronucleus; protomont encysts in narrow crevices; tomont divides into 2–16 tomites; tomite with a tubular macronucleus, two ciliated grooves on ventral side, and 9 ciliary rows; phoront cysts occur on the body hairs of *Xiphocaridina* to metamorphose into trophont (Miyashita, 1933).

*H. caridinae* M. (Fig. 339 a). Fully grown trophonts 80–120μ long; phoronts and phoront cysts present in fresh moult and body hairs respectively of the freshwater shrimp, *Xiphocaridina compressa*.

Genus *Cyrtocaryum* Fauré-Fremiet and Mugard. Trophont, astomous; external appearance resembles *Anoplophrya* (p. 691); macronucleus reticulate as in *Foettingeria*; liberated in sea water; no encystment, but multiplication in free state; differentiation of an oral ciliary field.

*C. halosydnae* F. and M. (Fig. 339, b–e). Trophont in the lateral caeca of the digestive tube of *Halosydna gelatinosa*; pyriform, 90–120μ by 65–80μ; with about 60 slightly spiral ciliary rows; cilia in
the anterior region strongly thygmotactic. When freed in the sea water, no encystment occurs, but division into eight to 16 sub-spherical individuals in chain, takes place. Tomont 45μ long; tomites 20μ by 16μ, asymmetrical, with a long caudal bristle.

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Chapter 38

Order 2 Spirotricha Bütschli

With free cilia only; exceptionally with small groups of cirrus-like projections in addition to cilia
Uniformly ciliated; in Peritromidae dorsal surface without or with a few cilia; in Lienophoridae cilia only on edge of attaching disk; peristome usually extended; peristomal field mostly ciliated......

Suborder 1 Heterotricha
Ciliation much reduced or none at all
Rounded in cross-section; cilia usually much reduced; adoral zone encloses a non-ciliated peristomal field in spiral form......

Suborder 2 Oligotricha (p. 814)
Compressed; carapaced; peristomal zone reduced to 8 membranellae which lie in an oval hollow. Suborder 3 Ctenostomata (p. 829)
Cirri only, on ventral side; dorsal side usually with rows of short bristles..

Suborder 1 Heterotricha Stein

Body ciliation complete and uniformly the same
Peristome sunk in a funnel-like hollow at anterior end, thus mostly covered. Family 1 Bursariidae (p. 797)
Peristome lies almost completely free, leading to a short and narrow oral funnel (absent in one family)
Peristome in anterior region
A narrow non-ciliated zone on right of adoral zone; usually an undulating membrane or ciliary row to right of this non-ciliated zone and anterior to cytostome; a small peristomal field between the membrane and adoral zone
Adoral zone extends diagonally to posterior-right on ventral surface; highly developed forms, with a long zone twisting spirally around body......Family 2 Metopidae (p. 800)
Adoral zone parallel to body axis on flat ventral surface, turns somewhat to right in front of cytostome; oral funnel distinct; typically an undulating membrane or a double ciliated furrow in front of cytostome..................

Suborder 4 Hypotricha (p. 832)

Family 1 Bursariidae (p. 797)
Peristome lies almost completely free, leading to a short and narrow oral funnel (absent in one family)
Peristome in anterior region
A narrow non-ciliated zone on right of adoral zone; usually an undulating membrane or ciliary row to right of this non-ciliated zone and anterior to cytostome; a small peristomal field between the membrane and adoral zone
Adoral zone extends diagonally to posterior-right on ventral surface; highly developed forms, with a long zone twisting spirally around body......Family 2 Metopidae (p. 800)
Adoral zone parallel to body axis on flat ventral surface, turns somewhat to right in front of cytostome; oral funnel distinct; typically an undulating membrane or a double ciliated furrow in front of cytostome..................

Suborder 5 Stentoridae (p. 806)
Peristomal field drawn out into 2 wings; free-swimming or secretes gelatinous lorica..................

Family 5 Stentoridae (p. 806)
Peristomal field drawn out into 2 wings; with flask-shaped, thin pseudochitinuous lorica. Family 6 Folliculinidae (p. 807)
Peristome at posterior end; cytopharynx directed anteriorly......

Family 7 Clevelandellidae (p. 809)
Body ciliation either confined to ventral side or lacking
Free-living; flattened; cilia only on ventral surface; adoral zone surrounds anterior region of ventral surface; cytostome on left edge near the middle of body......Family 8 Peritromidae (p. 810)
Ectocommensal; extremities discoid; body narrowed; anterior disk surrounded spirally by adoral zone; posterior disk bears membranous cilia.................Family 9 Licnophoridae (p. 810)

Family 1 Bursariidae Perty

Genus Bursaria Müller. Ovoid; anterior end truncate, posterior end broadly rounded; dorsal surface convex, ventral surface flattened; deep peristome begins at anterior end and reaches about

Fig. 340. a, Bursaria truncatella, ×60 (Kahl); b, Thylacidium truncatum, ×440 (Schewiakoff); c, Bursaridium difficile, ×210 (Kahl); d, Balantidium duodeni, ×170 (Stein); e, B. praenucleatum, ×950 (Kudo and Meglichts).

central part of body, where it gives rise to cytostome and cytopharynx, which is bent to left; lengthwise fold divides peristome into 2 chambers; striation longitudinal; ciliation complete and uniform; macronucleus band-form; many micronuclei; many contractile vacuoles distributed along lateral and posterior borders; cysts with a double envelope; fresh water. Cytology and conjugation (Poljansky, 1934); division (Schmäh1, 1926); fibrils (Peschkowsky, 1927).
B. truncatella M. (Fig. 340, a). 500–1000μ long; macronucleus a long rod; 10–34 vesicular micronuclei; fission mostly during night; feeds on various Protozoa; cysts 120–200μ in diameter; macronucleus becomes coiled and intertwined; fresh water (Schmähl, 1926; Beers, 1948).

Genus Thylacidium Schewiakoff. Similar to Bursaria in general appearance; but smaller in size; peristome simple in structure without longitudinal fold; with zoochlorellae; fresh water. One species. T. truncatum S. (Fig. 340, b). 60–100μ long.

Genus Bursaridium Lauterborn. Similar to Bursaria; peristome funnel turns to right; fresh water.

B. difficile Kahl (Fig. 340, c). Anterior end truncate, cytopharynx slanting toward right; about 130μ long.

Fig. 341. Balantidium coli, ×530 (Kudo). a, a living trophozoite; b, a stained trophozoite; c, a fresh cyst; d, a stained cyst.

Genus Balantidium Claparède and Lachmann (Balantidiopsis Bütschli; Balantiodoides Alexeieff). Oval, ellipsoid to subcylindrical; peristome begins at or near anterior end; cytopharynx not well developed; longitudinal ciliation uniform; macronucleus elongated; a micronucleus; contractile vacuole and cytopyge terminal; in the gut of vertebrates and invertebrates. Numerous species (Hegner, 1934; Kudo and Meglitsch, 1938).

B. coli (Malmsten) (Fig. 341). Ovoid; 40–80μ by 30–60μ, but length varies 30–150μ; body covered by many slightly obliquely longitudinal rows of cilia; peristome small near anterior tip, lined with coarser cilia; inconspicuous cytostome and cytopharynx are located at the end of peristome; 2 contractile vacuoles, one terminal, the other near the middle of body; macronucleus sausage-shape and a vesicular micronucleus; cytopyge near the posterior tip; food particles are of various kinds, including erythrocytes and other host
cell fragments, starch grains, faecal debris, etc. The trophozoite multiples by binary fission. Conjugation (Brumpt, 1909; Jameson, 1927; Scott, 1927; Nelson, 1934).

The cysts are circular to ovoid in outline; slightly yellowish or greenish and refractile; 40–60μ in diameter; cyst wall made up of 2 membranes; cytoplasm hyaline; macronucleus and a contractile vacuole are usually seen.

This ciliate lives in the colon and caecum of man and causes balantidiosis or balantidial dysentery. Strong (1904) made the first histological study of the infection. The organisms invade the tissues and blood vessels of the mucosa and submucosa. At the beginning there is hyperaemia with punctiform haemorrhages, and later vascular dilatation, round cell infiltration, eosinophilia, etc., develop in the infected area. Finally deep-seated ulcers are produced. The balantidial dysentery is usually of chronic type. It has a wide geographical distribution. In the United States a few cases of infections have been observed in recent years. In the Philippine Islands, more cases have been noticed than anywhere else.

This ciliate is a very common parasite in the intestine of pigs, and also of chimpanzee and orang-outang. In pigs, the organism ordinarily confines itself to the lumen of the intestine, but according to Ratcliffe (1934), when the host animals become infected by organisms belonging to Salmonella, it invades and ulcerates the intestinal wall. The cysts developing in pigs appear to become the chief source of infection, since balantidial dysentery is more commonly found among those who come in contact with pigs or pig intestine. The cysts remain viable for weeks in pig faeces in moist and dark places, though they are easily killed by desiccation or exposure to sunlight. The cysts may reach human mouth in food or in water contaminated with them, through unclean hands of persons who come in contact with faeces or intestine of pigs, and in some cases perhaps through uncooked sausage.

*B. suis* McDonald. Ellipsoid; 35–120μ by 20–60μ; macronucleus more elongate than that of *B. coli*; in the intestine of pigs (McDonald, 1922). Levine (1940) through a series of culture studies, has come to consider that *B. coli* and *B. suis* are only morphological variations due to the nutritional condition and that *B. suis* is synonymous with *B. coli*. Lamy and Roux (1950) observed both forms in cultures started with single individuals, and considered the elongate *suis* as conjugants and the oval *coli* as vegetative forms.

*B. caviae* Neiva, da Cunha and Travassos. In the caecum of guinea-pig. Morphology (Scott, 1927; Nie, 1950).
Other domestic and wild animals harbor various species of Balantidium.

*B. duodeni* Stein (Fig. 340, d). 70–80 μ by 55–60 μ; in the intestine of the frog.

*B. praenucleatum* Kudo and Meglitsch (Fig. 340, e). 42–127 μ long, 32–102 μ thick, 25–80 μ wide; macronucleus close to anterior end; in the colon of *Blatta orientalis* (Kudo and Meglitsch, 1938).

**Family 2 Metopidae** Kahl

Genus **Metopus** Claparède and Lachmann. Body form changeable; when extended oblong or fusiform; peristome conspicuous, slightly spirally diagonal, beginning at the anterior end and reaching the middle of the body; when contracted, peristome much spirally coiled; cytopharynx short; body ciliation uniform, longitudinal or in some, spiral; longer cilia at ends; conspicuous contractile vacuole terminal; macronucleus ovoid to elongate; fresh or salt water (sapropelic), some parasitic. Numerous species.

*M. es* Müller (*M. sigmoides* C. and L.) (Figs. 87; 342, a). 120–200 μ long; sapropelic. Noland’s (1927) study on its conjugation has been described (p. 161).

*M. striatus* McMurrich (Fig. 342, b). 80–120 μ long; fresh water.

*M. fuscus* Kahl (Fig. 342, c). 180–300 μ long by 60 μ wide and 40 μ thick; fresh water.

*M. circumlabens* Biggar (Fig. 342, d). 70–165 μ by 50–75 μ; in the digestive tract of sea urchins, *Diadema setosum* and *Echinometris subangularis* in Bermuda (Biggar, 1932; Lucas, 1934); in *Centrechinus antillarum*, etc., in Tortugas (Powers, 1935); in *Diadema setosum* and *Echinometra oblonga* in Japan (Uyemura, 1933).

Genus **Spirorhynchus** da Cunha. Fusiform; somewhat flattened; anterior end drawn out and curved toward left; posterior end also drawn out; spiral peristome; cytopharynx small with an undulating membrane; cilia uniformly long; contractile vacuole posterior; longitudinally striated; body surface with closely adhering bacteria (Kirby); three spherical macronuclei; micronucleus (?); in brackish water (da Cunha, 1915).

*S. verrucosus* da C. (Fig. 342, e). 122–140 μ by 20–22 μ. Kirby (1934) observed it in salt marsh with 3 per cent salinity; California.

Genus **Caenomorpha** Perty (*Gyrocoris* Stein). Bell-shaped; carpapaced ectoplasm in some species bears prorichocysts; strong marginal zone of about 8 rows of cilia; 1–2 dorsal rows of longer cilia and a dense spiral field around caudal prolongation; peristome long; cytostome posterior; cytopharynx directed anteriorly; a single
Fig. 342. a, *Metopus es*, ×260 (Kahl); b, *M. striatus*, ×220 (Kahl); c, *M. fuscus*, ×150 (Kahl); d, *M. circumlabens*, ×370 (Powers); e, *Spirohynchus verrucosus*, ×360 (Kirby); f, *Caenomorpha medusula*, ×200 (Blochmann); g, *Blepharisma lateritium*, ×160 (Penard); h, *B. persicinum*, ×290 (Penard); i, *B. steini*, ×340 (Penard); j, *Protocruzia pigerima*, ×390 (Faria, da Cunha and Pinto); k, *Phacodinium metschnicofi*, ×270 (Kahl).

elongate or two spherical macronuclei; a micronucleus; fresh or salt water (sapropelic). Several species.

*C. medusula* P. (Fig. 342, f). 150μ by 130μ; fresh and brackish water. Several varieties.

**Family 3 Spirostomidae** Kent.

**Genus Spirostomum** Ehrenberg. Elongated; cylindrical; somewhat compressed; ectoplasm with highly developed myonemes which are arranged lengthwise independent of ciliary rows, hence highly contractile; yellowish to brown; excretory vacuole terminal large, with a long dorsal canal; macronucleus either ovoid or chain form;
cilia short; rows longitudinal; caudal cilia are thigmotactic, secrete mucous threads (Jennings); peristome closely lined with short mem-
branellae; fresh or salt water. Several species.

Fig. 343. a, Spirostomum ambiguum, ×65 (Kahl); b, S. minus, ×140 (Kahl); c, S. loxodes, ×240 (Stokes); d, S. intermedium, ×140 (Kahl); e, S. teres, ×200 (Kahl); f, S. filum, ×190 (Penard); g, Gruberia calcinsi, ×140 (Bertran); h, Pseudoblepharisma tenuis, ×310 (Kahl); i, Parablepharisma pellitum, ×340 (Kahl).
S. ambiguurn E. (Figs. 38; 343, a). 1–3 mm. long; macronucleus composed of many beads; many micronuclei; peristome 2/3 the body length; fresh water. Regeneration (Seyd, 1936); irritability (Blättner, 1926).

S. minus Roux (Fig. 343, b). 500–800μ long; macronucleus chain-form; in fresh and salt water (Kahl).

S. loxodes Stokes (Fig. 343, c). About 300μ long (length: width, 6–7:1); peristome about 1/3 the body length; oblique striation; longer cilia at ends; macronucleus chain-form; fresh water.

S. intermedium Kahl (Fig. 343, d). Slender; 400–600μ long; macronucleus chain-form; fresh water.

S. teres Claparède and Lachmann (Fig. 343, e). 150–400μ long; macronucleus oval; in fresh water and also reported from salt water.

Genus Gruberia Kahl. Similar to Spirostomum in general appearance; but posterior end drawn out; slightly contractile; contractile vacuole posterior; macronucleus compact or beaded; salt water.

G. calcinski Beltrán (Fig. 343, g). 200–800μ long; peristome 2/3 the body length; many (contractile?) vacuoles distributed; monili-form macronucleus; many micronuclei; Woods Hole (Beltrán, 1933).

Genus Blepharisma Perty. Pyriform, spindle-form or ellipsoid; somewhat narrowed anteriorly; compressed; peristome on the left border, which is twisted to right at posterior end and connected with oral funnel with membrane; in front of cytostome a 2-layered undulating membrane on right edge; ciliary rows longitudinal; ciliation dense; contractile vacuole and cytopyge terminal; macronucleus one or divided into several parts; several species rose-colored; fresh or salt water. Many species.

B. lateritium (Ehrenberg) (Fig. 342, g). 130–200μ long; pyriform; macronucleus oval; a micronucleus; rose-colored; fresh water among decaying leaves.

B. persicinum P. (Fig. 342, h). 80–120μ long; elongate oval; posterior end pointed; left peristomal edge sigmoid; preoral membrane large; macronucleus in 3–7 parts; rose-colored; fresh water among decaying vegetation.

B. steini Kahl (Fig. 342, i). 80–200μ long; macronucleus ovoid; reddish to colorless; fresh water in sphagnum.

B. undulans Stein. 150–300μ long; macronucleus in 2 parts; undulating membrane long; cytopharynx directed posteriorly; fresh water among decaying vegetation. Contractile vacuole (Moore, 1934); influence of light on color (Giese, 1938) (p. 46); morphology
and physiology (Stolte, 1924); macronuclear reorganization (Young, 1939); multiconjugation (Weisz, 1950a); zoopurpurin (Weisz, 1950).

Genus **Protocruzia** Faria, da Cunha and Pinto. Peristome does not turn right, leads directly into cytostome; convex left side not ciliated, but bears bristles; flat right side with 3–5 faintly marked ciliary rows; peristome begins at pointed anterior end and extends 1/4–1/3 the body length; cytopharynx (?); macronucleus simple; contractile vacuole subterminal; salt water.

*P. pigerrima* (Cohn) (Fig. 342, j). About 20μ (da Cunha); 50–60μ long (Kahl); peristome 1/4–1/3 the body length; salt water.

Genus **Phacodinium** Prowazek. Oval; marked grooves on body surface; cilia in cirrus-like fused groups; peristome long on left margin; cytostome posterior; contractile vacuole terminal; macronucleus horseshoe-shape; 5–9 micronuclei; fresh water. One species.

*P. metschnicoffi* (Certes) (Fig. 342, k). About 100μ long.

Genus **Pseudoblepharisma** Kahl. Body form intermediate between *Spirostomum* and *Blepharisma*; right peristomal edge with 2 rows of cilia; fresh water.

*P. tenuis* K. (Fig. 343, h). 100–200μ long.

Genus **Parablepharisma** Kahl. Similar to *Blepharisma*; but peristome-bearing anterior half narrowed neck-like and pointed; ectoplasm covered with gelatinous layer in which symbiotic bacteria are imbedded; salt water.

*P. pellitum* K. (Fig. 343, i). 120–180μ long.

Genus **Nyctotherus** Leidy. Oval or reniform; compressed; peristome begins at anterior end, turns slightly to right and ends in cytostome located midway between the ends; cytopharynx runs dorsally and posteriorly, a long tube with undulating membrane; ciliary rows longitudinal and close-set; massive macronucleus in anterior half with a micronucleus; in some, nuclei are suspended by a karyophore; endoplasm with discoid glycogen bodies, especially in anterior region, hence yellowish to brown; contractile vacuole and cytopyge terminal; in the colon of Amphibia and various invertebrates. Numerous species (Geiman and Wichterman, 1937; Wichterman, 1938; Carini, 1938–1945).

*N. ovalis* L. (Figs. 3; 344, a, b). Ovoid; anterior half compressed; macronucleus elongate, at right angles to dorso-ventral axis at anterior 1/3; micronucleus in front of macronucleus; distinct karyophore; glycogen bodies; 90–185μ by 62–95μ; giant forms up to 360μ by 240μ; cysts 72–106μ by 58–80μ; in the colon of cockroaches. The chromatin spherules of the macronucleus are often very large (p. 42). Fibrillar structure (ten Kate, 1927); nuclei (Kudo, 1936).
**N. cordiformis** (Ehrenberg) (Figs. 85; 344, c). 60–200μ by 40–140μ; ovoid; micronucleus behind macronucleus; no karyophore; in the colon of frogs and toads. Higgins (1929) notes that there are certain differences between American and European forms and that the

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Fig. 344. a, b, *Nyctotherus ovalis*, ×340 (Kudo); c, *N. cordiformis* ×170 (Stein); d, *Condylostoma vorticella*, ×120 (Penard); e, *Stentor coerules*, somewhat contracted, ×70 (Roux); f, *S. polymorphus*, ×60 (Roux); g, *S. mulleri*, ×50 (Kahl); h, *S. roeselt*, ×75 (Roux); i, *S. igneus*, ×160 (Kahl); j, *S. amethystinus*, ×100 (Kahl).
organisms exhibit a great variety of form and size in the tadpoles of various frogs, although those of adult frogs are relatively constant in form. Life cycle (Wichterman, 1936) (p. 198); tactile cilia (Fernandez-Galiano, 1948); fibrillar structure (ten Kate, 1927).

**Family 4 Condylostomidae** Kahl

**Genus Condylostoma** Bory. Ellipsoid; anterior end truncate, posterior end rounded or bluntly pointed; slightly flattened; peristome wide at anterior end and V-shaped, peristomial field not ciliated; a large membrane on right edge and adoral zone on left; macronucleus moniliform; one to several contractile vacuoles often with canal; cytopyge posterior; fresh or salt water. Many species (Spiegel, 1926).

*C. vorticella* (Ehrenberg) (Fig. 344, d). 100–200μ long; fresh water.
*C. patens* (Müller). 250–550μ long; salt water; Woods Hole (Calkins).

**Family 5 Stentoridae** Carus

**Genus Stentor** Oken. When extended, trumpet-shaped or cylindrical; highly contractile; some with mucilaginous lorica; usually oval to pyriform while swimming; conspicuous peristomial field frontal; adoral zone encircles peristome in a spiral form, leaving a narrow gap on ventral side; the zone and field sink toward cytostome and the former continues into cytopharynx; macronucleus round, oval or elongated, in a single mass or moniliform; contractile vacuole anterior-left; free-swimming or attached; fresh water.

*S. coeruleus* Ehrenberg (Figs. 14; 344, e). Fully extended body 1–2 mm. long; anterior end greatly expanded; the beautiful blue color is due to a pigment, stentorin, lodged in interstria tion granules (p. 45); macronucleus moniliform; fresh water. Body and nuclear size (Burnside, 1929); physiology (Dierks, 1926); effect of environment (Stolte, 1922); cytology (Dierks, 1926; Weisz, 1949); regeneration (Schwartz, 1935; Weisz, 1948, 1948a, 1951); vertical distribution (Sprugel, 1951).

*S. striatus* Barraud-Maskell. Dark bluish green; funnel-shaped; peristomal edge irregularly undulating; striation conspicuous; macronucleus beaded; up to 2.2 mm. long.

*S. polymorphus* (Müller) (Fig. 344, f). Colorless; with symbiotic Chlorella 1–2 mm. long when extended; macronucleus beaded; anterior end expanded.

*S. müleri* (Bory) (Fig. 344, g). Colorless; with zoochlorellae; 2–3 mm. long; anterior end expanded; posterior portion drawn out into
stalk, often housed in a gelatinous tube; on body surface 3–4 longer and stiff cilia grouped among cilia; macronucleus moniliform.

*S. roeseli* Ehrenberg (Fig. 344, h). 0.5–1 mm. long; anterior end expanded; body surface with groups of longer cilia; posterior portion drawn out and often housed in a gelatinous tube; macronucleus long band-form.

*S. igneus* E. (Fig. 344, i). Rose-colored or colorless; 200–400μ long; macronucleus oval; ciliation uniform.

*S. niger* (Müller). Yellowish or brown; macronucleus oval; 200–300μ long.

*S. multiformis* (M.) Dark blue to bluish green; anterior end not expanded; 150–200μ long; macronucleus oval.

*S. amethystinus* Leidy (Fig. 344, j). Habitually pyriform (contracted); amethyst-blue; with zoochlorellae; 300–600μ long; macronucleus oval.

*S. pyriformis* Johnson. When extended 500μ long; anterior end 200μ in diameter.

Genus *Fabrea* Henneguy. Pyriform; posterior end broadly rounded, anterior end bluntly pointed; peristome extends down from anterior end 2/5 or more the body length, its posterior portion closely wound; peculiar black spot beneath membranellae in anterior portion of spiral adoral zone, composed of numerous pigment granules; without contractile vacuole; macronucleus, a sausage-shaped body or in 4 parts; in salt water.

*F. salina* H. (Fig. 345, a, b). 120–220μ by 67–125μ (Kirby); 130–450μ by 70–200μ (Henneguy); cysts ovoidal, with gelatinous envelope; 89–111μ by 72–105μ. Kirby (1934) found the organism in ditches and pools in salt marshes, showing salinities 7.5–20.1 per cent in California.

Genus *Climacostomum* Stein. Oval; flattened; right edge of peristome without membrane, left edge, semicircular or spiral with a strong adoral zone; peristomal field ciliated; cytopharynx a long curved tube with a longitudinal row of cilia; macronucleus band-form; contractile vacuole terminal, with two long canals; fresh or brackish water.

*C. virens* (Ehrenberg) (Fig. 345, c). 100–300μ long; with or without zoochlorellae; fresh and brackish water.

Family 6 Folliculinidae Dons

Genus *Folliculina* Lamarck. Horny or chitinous lorica (Fig. 345, d) attached on broad surface; neck of the lorica oblique to perpendicular; sometimes with a collar or spiral ridge; neck uniform in
Fig. 345. a, b, *Fabrea salina* (Kirby) (a, trophozoite, ×170; b, cyst, ×330); c, *Climacostomum vireus*, ×100 (Stein); d, side-view of the loria of a Folliculina, ×150 (Andrews); e, *Folliculina moebiisi*, ×170 (Stein); f, *F. producta*, ×110 (Wright); g, *Pseudofolliculina arctica*, ×50 (Dons); h, *Parafolliculina violacea*, ×230 (Andrews).

diameter; in salt or fresh water. Species (Andrews, 1914, 1921, 1923; Sahrhage, 1916); test secretion (Dewey, 1939).

*F. moebiisi* Kahl (Fig. 345, e). Loria about 500μ long.

*F. producta* (Wright) (Fig. 345, f). Loria yellowish brown; 250μ long; neck often long; Atlantic coast.

*F. boltoni* Kent. Loria about 200μ; loria and body blue green; aperture only slightly enlarged; short neck oblique or upright; in fresh water (Hamilton, 1950, 1952).

Genus **Microfolliculina** Dons. Posterior end or sides of loria with sack-like protuberances.

*M. limnoriae* (Giard). Loria dark blue; pellicle faintly striated; salt water.

Genus **Pseudofolliculina** Dons. Loria attached with its posterior
end; more or less vertical; without ring-furrow in middle; with or without style; salt water.

*P. arctica* D. (Fig. 345, g). Lorica about 430μ high, with spiral ridge; off Norweigian coast 15–28 m. deep.

Genus *Parafolliculina* Dons. Neck of lorica with a basal swelling; attached either with posterior end or on a lateral surface; salt water.

*P. violacea* (Giard) (Fig. 345, h). Total length 225–288μ; widely distributed in salt water (Andrews, 1921, 1942).

Family 7 *Clevelandellidae* Kidder

Genus *Clevelandella* Kidder (*Clevelandia* K.). Elongate pyriform or spear-shaped; posterior region drawn out, at the end of which peristome and cytostome are located; body more or less flexible; completely ciliated; one macronucleus supported by a karyophore; a micronucleus; a contractile vacuole at posterior left, near cytopyge;

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Fig. 346. a, b, ventral and dorsal views of *Clevelandella panesthiae*, X300; c, d, *Paraclelevelandia brevis* (c, ventral view, X760; d, a cyst, X740) (Kidder); e, *Peritromus californicus*, X360 (Kirby); f, *Lichnophora macrofarlandi*, X420; g, *L. conklini*, X340 (Stevens).
endocommensals in the colon of wood-feeding roaches, *Panesthia javanica* and *P. spadica*. Several species.

*C. panesthiae* K. (Fig. 346, *a, b*). Broadly fusiform with bluntly pointed anterior end and truncate posterior end; 87–156(123)μ by 53–78(62)μ; peristomal projection about one-fifth the body length; peristome is nearly enclosed; macronucleus massive; a vesicular micronucleus on its anterior border; karyophore separates the endoplasm into 2 parts: anterior part with glycogenous platelets, posterior part with numerous food particles; often parasitized by *Sphaerita* (p. 893); in the colon of *Panesthia javanica* and *P. spadica* (Kidder, 1937, 1938).

Genus *Paraclevelandia* Kidder. Elongate pyriform; body rigid; posterior end truncated obliquely to left; no peristomal projection; one macronucleus and one micronucleus; at anterior end, there is a sac connected with the karyophore, which is said to be a "macronuclear reservoir"; endocommensals.

*P. brevis* K. (Fig. 346, *c–d*). Conical in shape; 16–38 (38)μ by 9–21 (19)μ; macronucleus spherical to elongate ellipsoid; micronucleus comparatively large, retains nuclear stains longer than macronucleus; anterior sac may sometimes be absent; cysts, 14–19μ long; ovoid; with a spherical macronucleus and a micronucleus; in the colon of *Panesthia javanica* and *P. spadica* (Kidder, 1938).

Family 8 *Peritromidae* Stein

Genus *Peritromus* Stein. Ovoid; ventral surface flattened, dorsal surface with hump of irregular outline bearing a few stiff cilia; ciliary rows only on ventral surface; a small undulating membrane at posterior end of peristome; short marginal spines; 2 macro- and 2 micro-nuclei; salt water.

*P. emmae* S. 90–100μ long; creeping on bottom; Woods Hole.

*P. californicus* Kirby (Fig. 346, *e*). Peristome short; left margin slightly concave; dorsal hump with wart-like protuberances, bearing spines (about 12μ long); 16–19 or more ventral ciliary rows; 2 spherical macronuclei, one anterior right and the other posterior left of hump; micronuclei 4 (2–5); 89–165μ by 60–96μ; salt marsh pools with salinity "1.2–6 per cent" in California (Kirby, 1934).

Family 9 *Licnophoridae* Stevens

Genus *Licnophora* Claparède. Discoid; body roughly divisible into basal disc, neck and oral disc; basal disc for attachment, with several concentric ciliary coronas; neck flattened, contractile narrowed part with or without a ventral furrow and fibril-bundles (both running from oral groove to basal disc); oral disc highly flat-
SPIROTRICHA, HETEROTRICHA

tened, round or ovoid; edge with membrandelle zone which extends to pharyngeal funnel; macronucleus long chain-form; without contractile vacuole; commensal in salt water animals.

*L. macfarlandi* Stevens (Fig. 346, f). Average 90–110μ by 45–60μ; diameter of basal disc 40–45μ; basal disc circular; macronuclei in 25–35 parts in 4 groups; commensal in the respiratory tree of *Stichopus californicus* (Stevens, 1901). Morphology, fission and regeneration (Balmuth, 1941, 1942).

*L. conklini* S. (Fig. 346, g). 100–135μ long; commensal in *Crepidula plana* of Atlantic coast.

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SPIROTRICHA, HETEROTRICHA


Chapter 39

Order 2 Spirotricha Bütschli (continued)

Suborder 2 Oligotricha Bütschli

The cilia are greatly reduced in number in the Oligotricha and the adoral zone encloses a non-ciliated spiral peristomal field.

Free-living

Oral portion of peristome lies free on ventral surface..........................

Adoral zone encloses frontal peristomal field

Without lorica.................. Family 2 Strobilidiidae (p. 815)

With lorica or test'.................. Family 3 Tintinnidae (p. 816)

Parasitic

Adoral and dorsal zones, both directed anteriorly and retractile; no other cilia................. Family 4 Ophryoscolecidae (p. 816)

In addition to adoral and dorsal zones, groups of cirri in posterior half of body, directed posteriorly and nonretractile..................

................................. Family 5 Cycloposthiidae (p. 823)

Family 1 Halteriidae Claparède and Lachmann

Genus Halteria Dujardin. Spherical or broadly fusiform; anterior border bears conspicuous adoral zone; oral part of peristome with a small membrane on right edge and cirri on left; with an equatorial zone of small oblique grooves, each bearing 3 long cirri or bristles; macronucleus oval; a micronucleus; contractile vacuole left of cytostome; fresh water. Several species (Szabó, 1935).

H. grandinella (Müller) (Fig. 347, a). About 7 bristle-bearing grooves; 15 frontal and 7 adoral membranellae; 20-40μ long. Kahl (1932) distinguishes 2 varieties: var. cirrifera (Fig. 347, b), 25-50μ long, with huge cirri instead of fine body cirri; and var. chlorelligera (Fig. 347, c), 40-50μ long, with bristles and large zoochlorellae; fresh water.

Genus Strombidium Claparède and Lachmann. Ovoid to spherical; adoral zone very conspicuous (2-4 conspicuous sickle-form frontal membranellae and adoral membranellae extend down cytopharynx, the first section surrounding an apical process); no body bristles or cirri; trichocysts; macronucleus oval or band-form; a micronucleus; a contractile vacuole; salt or fresh water. Numerous species.

S. calkinsi Fauré-Fremiet (Fig. 347, d). 35-60μ long; brackish and salt water; Calkins (1902) first observed it at Woods Hole.
Genus **Tontonia** Fauré-Fremiet. With well-developed apical collar; a long cytoplasmic (contractile) caudal process; salt water.

*T. gracillima* F.-F. (Fig. 347, e). 48–52μ long; caudal process 250–300μ long; macronucleus moniliform; with zoochlorellae.

![Fig. 347. a, Halteria grandinella, ×490 (Kahl); b, H. g. var. cirrifera, ×370 (Kahl); c, H. g. var. chlorelligera, ×260 (Kahl); d, Strombidium calcinski, ×900 (Calkins); e, Tontonia gracillima, ×540 (Fauré-Fremiet); f, Strobilidium gyrans, ×340 (Kahl); g, Tintinnidium fluviatile, ×140 (Kent); h, i, T. semiciliatum, ×140 (Sterki); j, Strombidinopsis gyrans, ×270 (Kent); k, Tintinnopsis cylindrata, ×440 (Daday); l, T. illinoisensis, ×420 (Hempel); m, Codonella cratera, ×540 (Fauré-Fremiet).](image)

**Family 2 Strobilidiidae** Kahl

Genus **Strobilidium** Schewiakoff. Pyriform or turnip-shaped; cytoplasm at anterior end; without cytopharynx; horseshoe-shaped
macronucleus anterior; a micronucleus; a contractile vacuole; fresh or salt water. Several species (Busch, 1921).

*S. gyrans* (Stokes) (Fig. 347, f). Lateral border with rounded elevation near anterior end, posterior end truncate; 40–70μ long; in standing fresh water.

**Family 3 Tintinnidae** Claparède and Lachmann

Conical or trumpet-like, attached inside a lorica of various forms, composed of gelatinous or pseudochitinous substances; with longitudinal rows of cilia, and 2 (1–4) macro- and a micro-nuclei; mostly pelagic, a few inhabiting fresh or brackish water. Kofoid and Campbell (1929) distinguished more than 300 species and placed them in 12 families and 51 genera, of which 23 genera were created by them. A few genera and species are mentioned here. Taxonomy (Hofker, 1932); species (Campbell, 1942; Balech, 1942–1951; Rampi, 1950; Silva, 1950); factors in evolution (Kofoid, 1930); lorica formation (Busch, 1925).

**Genus Tintinnidium** Stein. Elongated lorica, highly irregular in form; soft; aboral end closed or with a minute opening; wall viscous and freely agglomerates foreign bodies; salt or fresh water.

*T. fluviatile* Stein (Fig. 347, g). Lorica 100–200μ by 45μ; on vegetation in fresh water.

*T. semiciliatum* Sterki (Fig. 347, h, i). 40–60μ long; on plants in fresh water.

**Genus Strombidinopsis** Kent. Lorica often absent; ovate or pyriform; frontal border with numerous long cirrus-like cilia; body covered by fine cilia; contractile vacuole posterior; fresh water.

*S. gyrans* K. (Fig. 347, j). 30–80μ long; fresh water pond.

**Genus Tintinnopsis** Stein. Lorica bowl-shaped; always with a broad aperture; aboral end closed; wall thin and covered with foreign bodies; salt or fresh water. Species (Balech, 1945).

*T. cylindrata* Kofoid and Campbell (Fig. 347, k). Lorica 40–50μ long; in lakes.

*T. illinoisensis* Hempel (Fig. 347, l). Lorica 59μ long; in rivers.

**Genus Codonella** Haeckel. Lorica urn- to pot-shaped; sharply divided externally and internally into a collar and bowl; collar without spiral structure; in fresh water.

*C. cratera* (Leidy) (Fig. 347, m). Lorica 60–70μ by 40μ; a number of varieties are often mentioned.

**Family 4 Ophryoscolecidae** Stein

Elongate oval, asymmetrical; with 1 or 2 (adoral and dorsal) zones of membranellae; in digestive tract of mammals. Sharp (1914)
employed “forma” to distinguish forms in Entodinium with common characteristics, differing in certain others, which scheme was extended to the whole family by Dogiel (1927). It is most probable that many species are varieties of a single species as judged by the work of Poljansky and Strelkow (1934); but since information is still incomplete, the present work ranks various formae with species, in agreement with Kofoid and MacLennan (1930).

The relationship between these oligotrichs and host ruminants has not definitely been determined, but it appears to be commensalism rather than symbiosis (Becker, Schulz and Emmerson, 1930; Mowry and Becker, 1930). Morphology (Bretschnieder, 1934, 1935); contractile vacuoles (MacLennan, 1933); conjugation (Dogiel, 1925); numbers in cattle stomach (Dogiel and Fedorowa, 1929); fauna in African antelopes (Dogiel, 1932); in yaks (Dogiel, 1934); in Indian goat (Das-Gupta, 1935); in Indian ox (Kofoid and MacLennan, 1930, 1932, 1933); in gaur (Kofoid and Christenson, 1934); in sheep, wild sheep and goat (Ferber and Fedorowa, 1929; Bush and Kofoid, 1948).

Genus Ophryoscolex Stein. Ovoid; with adoral and dorsal zones of membranellae; dorsal zone some distance behind anterior end, encircling 3/4 the body circumference at middle, broken on right ventral side; 3 skeletal plates extend over the body length on right-ventral side; 9–15 contractile vacuoles in 2 (anterior and posterior) circles; macronucleus simple, elongate; in the stomach of cattle, sheep, goat and wild sheep (Ovis orientalis cycloceros). Several species (Kofoid and MacLennan, 1933); neuromotor system (Fernandez, 1949).

Dogiel (1927) designated the following species as 3 formae of O. caudatus Eberlein.

O. bicoronatus Dogiel (Fig. 348, a). 120–170μ by 81–90μ; primary spine 38–58μ long; in sheep.

O. caudatus Eberlein (Fig. 348, b). 137–162μ by 80–98μ; preanal spines 47–60μ long; in sheep, goat, and cattle.

O. quadricoronatus Dogiel (Fig. 348, c). 128–180μ by 86–100μ; preanal spines 48–63μ long; in sheep and Ovis orientalis cycloceros.

Genus Caloscolex Dogiel. Ovoid; anterior end truncate, posterior end rounded with or without processes; 2 zones of membranellae; dorsal zone encircles the body completely; 3 skeletal plates variously modified; 7 contractile vacuoles in a single circle; nucleus elongate; in the stomach of Camelus dromedarius. Several species.

C. cuspidatus D. (Fig. 348, d). 130–160μ by 73–90μ.

Genus Entodinium Stein. Without dorsal zone; adoral zone at truncate anterior end; without skeleton; contractile vacuole ante-
rior; macronucleus, cylindrical or sausage-form, dorsal; micronucleus anterior to middle and on left-ventral side of macronucleus; in cattle and sheep. Numerous species (Kofoid and MacLennan, 1930; MacLennan, 1935).

*E. caudatum* S. (Fig. 348, c). 50–80μ long; in cattle and sheep.

*E. bursa* S. (Fig. 348, f). 55–114μ by 37–78μ (Schuberg); 80μ by 60μ (Becker and Talbott); in the stomach of cattle.

Genus *Amphacanthus* Dogiel. Similar to *Entodinium*; but spinous processes at both anterior and posterior ends; in stomach of *Camelus dromedarius*. One species.

*A. ovum-rajae* D. (Fig. 348, g). 46–55μ by 32–48μ.
Genus *Eodinium* Kofoid and MacLennan. Dorsal zone on the same level as adoral zone; without skeleton; macronucleus a straight, rod-like body beneath dorsal surface; 2 contractile vacuoles; in cattle and sheep. Several species.

*E. lobatum* K. and M. (Fig. 349, a). 44–60μ by 29–37μ; in *Bos indicus* (Kofoid and MacLennan, 1932).

Genus *Diplodinium* Schuberg. Adoral and dorsal zones at the

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**Fig. 349.** a, *Eodinium lobatum*, ×540 (Kofoid and MacLennan); b, *Diplodinium dentatum*, ×250 (Kofoid and MacLennan); c, *Eremoplastron bovis*, ×550 (Kofoid and MacLennan); d, *Eudiplodinium maggi*, ×500 (Dogiel); e, *Diploplastron affine*, ×320 (Dogiel); f, *Metadinium medium*, ×320 (Dogiel).
same level; without skeletal plates; macronucleus beneath right side, its anterior third bent ventrally at an angle of 30°–90°; 2 contractile vacuoles; in cattle, antelope, Camelus dromedarius, reindeer, goat. Numerous species (Kofoid and MacLennan, 1932).

D. dentatum (Stein) (Fig. 349, b). 65–82μ by 40–50μ; in cattle (including Bos indicus).

Genus Eremoplastron Kofoid and MacLennan. Adoral and dorsal zones at anterior end; a single narrow skeletal plate beneath right surface; triangular or rod-like macronucleus, anterior end of which is often bent ventrally; 2 contractile vacuoles; in cattle, antelope, sheep, reindeer. Numerous species (Kofoid and MacLennan, 1932).

E. bovis (Dogiel) (Fig. 349, c). 52–100μ by 34–50μ; in cattle and sheep.

Genus Eudiplodinium Dogiel. Adoral and dorsal zones at anterior end; a single, narrow, skeletal plate beneath right surface; rod-like macronucleus with its anterior end enlarged to form a hook opening dorsally; pellicle and ectoplasm thick; 2 contractile vacuoles with heavy membranes and prominent pores; in cattle. Species (Kofoid and MacLennan, 1932).

E. maggi (Fiorentini) (Fig. 349, d). 104–255μ by 63–170μ; in cattle, sheep and reindeer.

Genus Diploplastron Kofoid and MacLennan. Adoral and dorsal zones at anterior end; 2 skeletal plates beneath right surface; macronucleus narrow; rod-like; 2 contractile vacuoles below dorsal surface, separated from macronucleus. One species (Kofoid and MacLennan, 1932).

D. affine (Dogiel and Fedorowa) (Fig. 349, e). 88–120μ by 47–65μ; in the stomach of cattle, sheep, and goat.

Genus Metadinium Awerinzew and Mutafowa. Adoral and dorsal zones at anterior end; 2 skeletal plates beneath right surface sometimes fused posteriorly; macronucleus with 2–3 dorsal lobes; 2 contractile vacuoles; pellicle and ectoplasm thick; conspicuous oesophageal fibrils beneath dorsal and right sides; in the stomach of cattle, sheep, goat, and reindeer (Awerinzew and Mutafowa, 1914).

M. medium A. and M. (Fig. 349, f). 180–272μ by 111–175μ; in cattle.

Genus Polyplastron Dogiel. Adoral and dorsal zones at anterior end; 2 skeletal plates beneath right surface, separate or fused; 3 longitudinal plates beneath left surface, with anterior ends connected by cross bars; contractile vacuoles beneath dorsal surface in a longitudinal row, also with additional vacuoles; in the stomach of cattle and sheep. Species (Kofoid and MacLennan, 1932).
SPIROTRICHA, OLIGOTRICHA

*P. multivesiculatum* (D. and Fedorowa) (Fig. 350, a). 120–190μ by 78–140μ; in cattle and sheep. MacLennan (1934) found that the skeletal plates are made up of small, roughly prismatic blocks of glycogen, each with a central granule.

Genus *Elytroplastron* Kofoid and MacLennan. 2 zones at anterior end, 2 skeletal plates beneath right surface, a small plate beneath ventral surface, and a long plate below left side; pellicle and ectoplasm thick; conspicuous fibrils beneath dorsal–right side. One species.

*E. hegneri* (Becker and Talbott) (Fig. 350, b). 110–160μ by 67–97μ; in cattle, sheep, *Buffelus bubalus* and *Bos indicus* (Becker, 1933).

Genus *Ostracodinium* Dogiel. 2 zones at anterior end; broad skeletal plate beneath right side; 2–6 contractile vacuoles in a dorsal row;
cytopharyngeal fibrils thick, extend to posterior end; in cattle, sheep, antelope, steenbok, and reindeer. Numerous species (Kofoid and MacLennan, 1932).

O. dentatum (Fiorentini) (Fig. 350, c). 52-110μ by 31-68μ; in the stomach of cattle.

Genus Enoploplastron Kofoid and MacLennan. 2 zones near anterior end; 3 skeletal plates beneath right–ventral side, either separate or partly fused; 2 contractile vacuoles; heavy pharyngeal fibrils; in cattle, reindeer and antelope.

E. triloricatum (Dogiel) (Fig. 350, d). Dogiel (1927) mentions size differences among those occurring in different host species, as follows: in cattle, 85–112μ by 51–70μ; in reindeer, 75–103μ by 40–58μ; in antelope (Rhaphicerus sp.), 60–110μ by 37–56μ.

Genus Epidinium Crawley. Elongate; twisted around the main axis; 2 zones; dorsal zone not at anterior end; 3 skeletal plates, with secondary plates; simple macronucleus club-shaped; 2 contractile vacuoles; in cattle, sheep, reindeer, camels, etc. Species (Kofoid and MacLennan, 1932).

E. caudatum (Fiorentini) (Fig. 350, e). 113–151μ by 45–61μ; in cattle, camels, Cervus canadensis and reindeer.

E. (Diplodinium) ecaudatum (F.) (Figs. 16; 350, f). 112–140μ by 40–60μ (Becker and Talbott); in cattle, sheep, and reindeer. The classical observation of Sharp (1914) on its neuromotor system has been described elsewhere (p. 63).

Genus Epiplastron Kofoid and MacLennan. Elongate; 2 zones; dorsal zone not at anterior end; 5 skeletal plates, with secondary plates; macronucleus simple, elongate; 2 contractile vacuoles; in antelopes.

E. africanum (Dogiel) (Fig. 350, g). 90–140μ by 30–55μ; in Rhaphicerus sp.

Genus Ophisthotrichum Buisson. 2 zones; dorsal zone at middle or near posterior end of body; one-piece skeletal plate well developed; 2 contractile vacuoles posterior; conjugation (Dogiel); in many African antelopes. One species.

O. janus (Dogiel) (O. thomasi B.) (Fig. 351, a). 90–150μ by 42–60μ. Conjugation (Dogiel, 1925).

Genus Cunhaia Hasselmann. Cytostome near anterior end, with adoral zone; dorsal zone on 1/3 of anterior-dorsal surface; 2 contractile vacuoles; skeleton (?); in the caecum of guinea pig, Cavia aperea. One species.

C. curvata H. (Fig. 351, b). 60–80μ by 30–40μ; in Brazil.
Family 5 Cycloposthiidae Poche

Pellicle firm and body rigid; zones of membranellae at anterior and posterior ends; more or less compressed; cytopharynx short and wide; macronucleus elongate; a single micronucleus; 2 or more contractile vacuoles; in horse and anthropoid apes.

Genus Cycloposthium Bundle. Large, elongate barrel-shaped; cytostome in center of a retractile conical elevation at anterior end; adoral zone conspicuous; an open ring-zone of membranellae near posterior end on both dorsal and ventral sides; pellicle ridged; skele-

Fig. 351. a, Ophisthotrichum janus, ×370 (Dogiel); b, Cunhaia curvata, ×670 (Hasselmann); c, Cycloposthium bipalatum, ×300 (Bundle); d, C. dentiferum, ×270 (Hsiung), e, Spirodinium equi, ×350 (Davis); f, Triadinium caudatum, ×300 (Hsiung); g, T. minimum, ×440 (Hsiung); h, Tetratoxum unifasciculatum, ×280 (Hsiung).
ton club-shaped; several contractile vacuoles in a row along band-
form macronucleus; in the caecum and colon of horse. Many species
(Hsiung, 1930). Cytology (Strelkow, 1929, 1932).

*C. bipalmatum* (Fiorentini) (Fig. 351, c). 80–127 µ by 35–57 µ. Con-
jugation (Dogiel, 1925).

*C. dentiferum* Gassovsky (Fig. 351, d). 140–222 µ by 80–110 µ.

Genus *Spirodinium* Fiorentini. Elongate, more or less fusiform;
adoral zone at anterior end; anterior ciliary zone encircles the body
at least once; a posterior ciliary arch, only 1/2 spiral; a dorsal cavity
of unknown function (Davis, 1941), lined with stiff rods; in the colon
and caecum of the horse. Species (Hsiung, 1930, 1935).

*S. equi* F. (Fig. 351, e). 82–196 µ by 46–108 µ; widely distributed.

Morphology (Hsiung, 1935a; Davis, 1941); division (Davis, 1941).

Genus *Triadinium* Fiorentini. More or less helmet-shaped; com-
pressed; adoral zone at anterior end; 2 posterior (ventral and dorsal)
zones; with or without a caudal projection; in the caecum and colon
of horse. Species (Hsiung, 1935).

*T. caudatum* F. (Fig. 351, f). 59–86 µ by 50–68 µ.

*T. galea* Gassovsky. 59–78 µ by 50–60 µ.

*T. minimum* G. (Fig. 351, g). 35–58 µ by 30–40 µ.

Genus *Tetratoxum* Gassovsky. Slightly compressed; 2 anterior
and 2 posterior zones of membranellae; in the colon of horse. Species
(Hsiung, 1930).

*T. unifasciculatum* (Fiorentini) (Fig. 351, h). 88–186 µ by 60–108 µ;
widely distributed. Morphology and micronuclear division (Davis,
1941a).

*T. excavatum* Hsiung. 95–135 µ by 55–90 µ.


Genus *Tripalmaria* Gassovsky (*Tricaudalia* Buisson). Adoral zone
at anterior end; 2 dorsal and 1 ventral-posterior zones in tuft-form;
macronucleus inverted U-shape; in the colon of horse. Cytology
(Strelkow, 1932).

*T. dogieli* G. (Fig. 352, a). 77–123 µ by 47–62 µ (Hsiung, 1930).

Genus *Triplumaria* Hoare. Adoral zone; 2 dorsal and 1 ventral
cirrose tufts (caudals); skeleton, composed of polygonal plates ar-
nanged in a single layer, surrounds the body except the dorsal sur-
face; dorsal groove supported by rod-like skeleton; macronucleus
elongate sausage-form, with a micronucleus attached to its dorsal
surface near middle; about 6 contractile vacuoles arranged in line
along dorsal surface of body; in the intestine of Indian rhinoceros
(Hoare, 1937).

*T. hamertoni* H. 129–207 µ long, 65–82 µ thick, 4–39 µ broad; endo-
Fig. 352. a, Tripalmaria dogieli, ×180 (Gassovsky); b, Cochliatoxum periachtum, ×270 (Hsiung); c, Ditoxum funinucleum, ×270 (Hsiung); d–f, Troglodytella abrassarti (d, ×670 (Swezey); e, ventral and f, dorsal view, ×210 (Brumpt and Joyeux)).
commensal in the intestine of *Rhinoceros unicornis* in Zoological Garden in London.

Genus **Cochliatoxum** Gassovsky. Adoral zone near anterior end; 3 additional zones, 1 antero-dorsal, 1 postero-dorsal and 1 postero-ventral; macronucleus with curved anterior end; in the colon of horse. One species.

*C. periachtum* G. (Fig. 352, b). 210–370μ by 130–210μ (Hsiung, 1930).

Genus **Ditoxum** Gassovsky. Large adoral zone near anterior end; 2 dorsal (anterior and posterior) zones; macronucleus curved club-shaped; in the colon of horse (Hsiung, 1935).

*D. funinucleum* G. (Fig. 352, c). 135–203μ by 70–101μ.

Genus **Troglodytella** Brumpt and Joyeux. Ellipsoid; flattened; adoral zone; 3 additional zones (anterior zone continuous or not continuous on ventral surface; posterior zone continuous on dorsal surface; between them a small zone on each side); skeletal plates in anterior region; macronucleus L-form; contractile vacuoles in 2 circles; in the colon of anthropoid apes.

*T. abrassarti* B. and J. (Fig. 352, d–f). About 145–220μ by 120–160μ; in the colon of chimpanzees (Brumpt and Joyeux, 1912). Reichenow (1920) distinguished var. *acuminata* on the basis of the drawn-out posterior end, which was found by Swezey (1932) to be a variant of *T. abrassarti*. Cytology (Swezey, 1934); cultivation (Nelson, 1932; Swezey, 1935).

*T. gorillae* Reichenow. 200–280μ by 120–160μ; in the colon of gorilla; with anterior zone not reaching the right side.

**References**


SPIROTRICHA, Oligotricha


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Chapter 40

Order 2 Spirotricha Bütschli (continued)

Suborder 3 Ctenostomata Lauterborn

The ciliates placed in this group are carapaced and compressed forms with a very sparse ciliation. The adoral zone is also reduced to about 8 membranellae. These organisms are exclusively free living and sapropelic in fresh, brackish, or salt water. Morphology and taxonomy (Kahl).

Posterior half of carapace with 4 ciliated rows on left and at least 2 rows on right; with anterior row of cilia on left side..Family 1 Epalcidae

Posterior half of carapace with cirrus-like groups on left only, none on right; without frontal cilia

Long ciliated band extends over both broad sides.....................

.................................Family 2 Discomorphidae

Short ciliated band ventral, extending equally on both broad sides...

.................................Family 3 Mylestomidae (p. 830)

Family 1 Epalcidae Wetzel

Genus Epalxis Roux. Rounded triangular; anterior end pointed toward ventral surface, posterior end irregularly truncate; dorsal surface more convex; right carapace with 1 dorsal and 1 ventral ciliary row in posterior region; usually 4 (2–3) median teeth; all anal teeth without spine; with comb-like structures posterior to oral aperture; 1–2 oval macronuclei dorsal; contractile vacuole posterior-ventral; sapropelic in fresh or salt water. Many species.

E. mirabilis R. (Fig. 353, a). 38–45μ by 27–30μ; fresh water.

Genus Saprodinium Lauterborn. Similar to Epalxis; but some (left and right) of anal teeth with spines; sapropelic in fresh or salt water. Several species.

S. dentatum L. (Fig. 353, b). 60–80μ long; fresh water (Lackey, 1925).

S. putrinium Lackey (Fig. 353, c). 50μ long, 40μ wide, about 15μ thick; in Imhoff tanks.

Genus Pelodinium Lauterborn. Right carapace with 2 median rows of cilia, its median anal teeth fused into one so that there are only three teeth. One species.

P. reniforme L. (Fig. 353, d). 40–50μ long; sapropelic.

Family 2 Discomorphidae Poche

Genus Discomorpha Levander. Oval; ventrally directed anterior spine long; posterior end without teeth or ridges; ciliated bands on
Fig. 353. a, Epalxis mirabilis, ×1200 (Roux); b, Saprodinium dentatum, ×430 (Kahl); c, S. putrini, ×470 (Lackey); d, Pelodinium reniforme, ×600 (Lauterborn); e, f, Discomorpha pectinata, (e, ×500; f, ×220) (Kahl); g, Mylestoma bipartitum, ×470 (Kahl); h, Atopodinium fibulatum, ×520 (Kahl).

both lateral surfaces; 2 spines on right side; 2 cirrus-like groups on posterior-left; sapropelic. A few species.

D. pectinata L. (Fig. 353, e, f). 70–90μ long; sapropelic.

Family 3 Mylestomidae Kahl

Genus Mylestoma Kahl. Posterior margin without any indentation, though sometimes a small one on right side, but none on left;
3 often long ribbon-like cirri on peristome; fresh or salt water. Several species.

*M. bipartitum* (Gourret and Roesner) (Fig. 353, *g*). 35–50μ long; two caudal processes; salt water.

Genus *Atopodinium* Kahl. Posterior left side with one large, and right side with 2 indentations; macronucleus spherical; sapropelic. *A. fibulatum* K. (Fig. 353, *h*). 40–50μ long.

References


Chapter 41

Order 2 Spirotricha Bütschli (continued)

Suborder 4 Hypotricha Stein

THE members of this suborder are, as a rule, flattened and strong cilia or cirri are restricted to the ventral surface. Except the family Aspidiscidae, the dorsal surface possesses rows of short slightly moveable tactile bristles. The peristome is very large with a well-developed adoral zone. The cirri on the ventral surface are called, according to their location, frontals, ventrals, marginals, anals (transversals), and caudals, as was mentioned before (Fig. 11, b). Asexual reproduction is by binary fission and sexual reproduction by conjugation. Encystment is common. Mostly free-living in fresh, brackish or salt water; a few parasitic.

Adoral zone fully formed
Cirri on ventral surface
Ventrals in rows, though in some reduced; 2 rows of marginals...

.................................Family 1 Oxytrichidae
Ventrals and marginals not in longitudinal rows...

.................................Family 2 Euplotidae (p. 839)
No ventral cirri; caudal cirri......Family 3 Paraeuplotidae (p. 843)
Adoral zone reduced.............Family 4 Aspidiscidae (p. 845)

Family 1 Oxytrichidae Kent

Genus Oxytricha Ehrenberg (Histrio Sterki; Opisthotricha Kent; Steinia Diesing). Ellipsoid; flexible; ventral surface flattened, dorsal surface convex; 8 frontals; 5 ventrals; 5 anals; short caudals; marginals may or may not be continuous along posterior border; macronucleus in 2 parts, rarely single or in 4 parts; fresh or salt water. Numerous species (Horváth, 1933); neuromotor system (Lund, 1935).

O. fallax Stein (Fig. 354, a). Posterior region broadly rounded; about 150μ long; fresh water. Amicronucleate race (Reynolds, 1932).

O. bifaria Stokes (Fig. 354, b). Right side convex; left side flattened; posterior end pointed; about 250μ long; fresh water infusion.

O. ludibunda S. (Fig. 354, c). Ellipsoid; flexible; 100μ long; fresh water among sphagnum.

O. setigera S. (Fig. 354, d). Elongate ellipsoid; 5 frontals; ventrals shifted anteriorly; 50μ long; fresh water.

Genus Tachysoma Stokes (Actinotricha Cohn). Flexible; frontals 8–10, of which anterior three are usually the largest; 5 ventrals
scattered; 5 anals; marginals at some distance from lateral borders, interrupted posteriorly; fresh or salt water.

*T. parvistyla* S. (Fig. 354, e). 10 frontals scattered; about 63μ long; in shallow freshwater pools.

Genus *Urosoma* Kowalewski. Similar to *Oxytricha*; but posterior portion drawn out and much narrowed; fresh water.

*U. caudata* (Stokes) (Fig. 354, f). 200–250μ long; pond water.

Genus *Amphisiella* Gourret and Roeser. With a single row of ventrals and 2 marginal rows; salt or fresh water. Several species.

*A. thiophaga* (Kahl) (Fig. 354, g). 70–100μ long; salt water.

Genus *Eschaneustyla* Stokes. Elliptical or ovate; narrow peri-

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Fig. 354. a, *Oxytricha fallax*, ×230 (Stein); b, *O. bifaria*, ×180 (Stokes); c, *O. ludibunda*, ×400 (Stokes); d, *O. setigera*, ×870 (Stokes); e, *Tachysoma parvistyla*, ×490 (Stokes); f, *Urosoma caudata*, ×250 (Stokes); g, *Amphiisiella thiophaga*, ×380 (Kahl); h, *Eschaneustyla brachytona*, ×240 (Stokes); i, *Gonostomum strenuum*, ×160 (Engelmann); j, *Hemicycliostyla sphagni*, ×100 (Stokes); k, l, *Cladotricha koltzowii* (k, ×170; l, ×300) (Kahl).
stome 1/3 the body length; frontals numerous, about 22 in addition to 2 at anterior margin; ventrals small and numerous in 3 oblique rows; no anals; marginals uninterrupted; contractile vacuole a long canal near left border; fresh water. One species.

*E. brachytona* S. (Fig. 354, h). 170–220μ long.

Genus *Gonostomum* Sterki (*Plagiotricha* Kent). Flexible; 8 or more frontals; 1–2 oblique ventral rows of short cirri; 4 or 5 anals; 2 marginal rows; fresh water.

*G. strenuum* (Engelmann) (Fig. 354, i). Elongate; with caudal bristles; about 150μ long; fresh water.

Genus *Hemicycliostyla* Stokes. Elongate oval; flexible; 8 or more frontals; 1–2 oblique ventral rows of short cirri; 4 or 5 anals; 2 marginal rows; fresh water.

*H. sphagni* S. (Fig. 354, j). About 400–500μ long; marsh water with sphagnum.

Genus *Hypotrichidium* Ilowaisky. Two ventral and marginal rows of cirri spirally arranged; peristome large, extends 1/2 the body length, with a large undulating membrane; 2 macro- and micro-nuclei; contractile vacuole anterior-left; fresh water.

*H. conicum* I. (Fig. 355, a). 90–150μ long.

Genus *Cladotricha* Gajevskaja. Elongate band-form; anterior end rounded, posterior end rounded or attenuated; frontals only 2 feathery cirri; macronucleus spheroidal; micronucleus; without contractile vacuoles; salt water, with 5–20 per cent salt content. One species.

*C. koltzowii* G. (Fig. 354, k, l). Band-form up to about 200μ long; posteriorly attenuated forms up to about 100μ long.

Genus *Psilotricha* Stein. Oval to ellipsoid; frontals and anals undifferentiated; ventrals and marginals long cirri, few; ventrals in 2 rows and a rudimentary row toward left; with or without zoochlorellae; fresh water. A few species.

*P. acuminata* S. (Fig. 355, b). 80–100μ long.

Genus *Kahlia* Horvath. Frontal margin with 3–4 strong cirri; 5–8 ventral longitudinal rows; marginals; sapropelic in fresh water.

*K. acrobates* H. (Fig. 355, c). 100–200μ long; soil infusion.

Genus *Uroleptus* Ehrenberg. Elongate body drawn out into a tail-like portion; 3 frontals; 2–4 rows of ventral cirri; marginals; no anals; sometimes rose- or violet-colored; fresh or salt water. Many species.

*U. limnetis* Stokes (Fig. 355, d). About 200μ long; fresh water among vegetation.
**U. longicaudatus** S. (Fig. 355, e). About 200μ long; marsh water with sphagnum.

**U. halseyi** Calkins (Fig. 355, f). About 160μ by 20μ; peristome 1/6–1/7 the body length; 3 ventrals; macronucleus divided into many (up to 26) parts; 2 (1–3) micronuclei; fresh water (Calkins, 1930).

Genus **Uroleptopsis** Kahl. Ventrals in 2 uninterrupted rows; salt water. A few species.

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**Fig. 355.** a, *Hypotrichidium conicum*, ×200 (Kahl); b, *Psilotricha acuminata*, ×230 (Stein); c, *Kahlia acrobates*, ×240 (Kahl); d, *Uroleptus limnetis*, ×240 (Stokes); e, *U. longicaudatus*, ×240 (Stokes); f, *U. halseyi*, ×470 (Calkins); g, *Uroleptopsis citrina*, ×260 (Kahl); h, *Strongylidium californicum*, ×200 (Kahl); i, *Stichotricha secunda*, ×340 (Kahl); j, *S. intermedia* (Froud); k, *Chaetospora mülleri* (Froud); l, *Urostyla grandis*, ×140 (Stein); m, *U. trichogaster*, ×150 (Kahl).
**U. citrina** K. (Fig. 355, g). Elongate; flexible; ectoplasm with pale-yellow ringed bodies which give the organism yellowish color; marginals discontinuous posteriorly; 2 contractile vacuoles near left border; 150–250μ long; salt water.

Genus **Strongylidium** Sterki. 2–5 ventral rows of cirri; marginals spirally arranged; 3–6 frontals; 2 or more macronuclei; fresh or salt water. Many species.

**S. californicum** Kahl (Fig. 355, h). 4–5 frontals; macronuclei about 30 in number; 4 micronuclei; contractile vacuole with short canals; about 250μ long; fresh water among vegetation.

Genus **Stichotricha** Perty. Slender ovoid or fusiform; peristome-bearing part narrowed; not flexible; usually 4 spiral rows of cirri; sometimes tube-dwelling, and then in groups; fresh or salt water. Many species.

**S. secunda** P. (Fig. 355, i). 130–200μ long; in fresh water.

**S. intermedia** Froud (Fig. 355, j). Solitary; non-loricate; 40–170μ long, 2/5 of which is a bent proboscis; two rows of body cilia; two rows of dorsal cilia, 5μ long; among Lemna in fresh water (Froud, 1949).

Genus **Chaetospira** Lachmann. Similar to **Stichotricha**; but peristome-bearing part flexible; fresh or salt water.

**C. mülleri** L. (Fig. 355, k). Flask-shaped, 60–200μ long, in a lorica; cytostome at the base of proboscis; a single (two or more) micronucleus; macronucleus in two to eight parts; ingested diatoms lose color in 10 minutes; Bodo is immobilized in less than one minute; binary fission; the anterior individual remains in the lorica, while the posterior individual (averaging 46μ long) swims away and sooner or later becomes attached to substrate; cysts pyriform, 35–55μ by 15–20μ; among Lemna in fresh water (Froud, 1949).

Genus **Urostyla** Ehrenberg. Ellipsoid; flexible; ends rounded; flattened ventral surface with 4–10 rows of small cirri and 2 marginal rows; 3 or more frontals; 5–12 anals; macronucleus a single body or in many parts; fresh or salt water. Numerous species.

**U. grandis** E. (Figs. 49; 355, l). 300–400μ long; macronucleus in 100 or more parts; 6–8 micronuclei; fresh water. Nuclei (Raabe, 1946, 1947) (p. 165).

**U. trichogaster** Stokes (Fig. 355, m). 250–330μ long; fresh water.

**U. caudata** S. (Fig. 356, a). Elongate ellipsoid; flexible; narrowed anterior part bent to left; peristome 1/3 the body length; macronucleus in many parts; contractile vacuoles near left margin; about 600μ long; fresh water with sphagnum.
**U. polymicronucleata** Merriman. Elliptical with broadly rounded ends; flexible; 225μ by 65μ; opaque, green or brown because of the ingested diatoms; 3 large and 10 small frontals; four ventral rows of cirri; marginals; macronucleus in two parts; three to 11 micronuclei (Merriman, 1937).

**U. coei** Turner. Elliptical, with a more pointed posterior end; 200μ by 50μ; four rows of ventral cirri, the right row being the longest; five frontals; macronucleus in two masses; four micronuclei (Turner, 1939).

Genus **Kerona** Ehrenberg. Reniform; no caudals; 6 oblique rows of ventral cirri; commensal. One species.

Fig. 356. a, *Urostyla caudata*, ×90 (Stokes); b, *Kerona polyporum*, ×200 (Stein); c, *Keronopsis rubra*, ×270 (Entz); d, *Epiclinites pluvialis*, ×100 (Smith); e, *Holosticha vernalis*, ×220 (Stokes); f, *H. hymenophora*, ×180 (Stokes); g, *Paraholosticha herbicola*, ×200 (Kahl); h, *Trichotaxis stagnatilis*, ×190 (Stokes); i, *Balladyna elongata*, ×800 (Roux); j, *Pleurotricha lanceolata*, ×250 (Stein); k, *Gastrostyla muscorum*, ×200 (Kahl).
**K. polyporum** E. (Fig. 356, b). 120–200μ long; commensal on Hydra.

Genus *Keronopsis* Penard. Two ventral rows of cirri reaching frontal field; caudals variable; macronucleus usually in several (rarely 2) parts; fresh or salt water. Numerous species.

*K. rubra* (Ehrenberg) (Fig. 356, c). Reddish; 200–300μ long; salt water.

Genus *Epiclintes* Stein. Elongate; spoon-shaped; flattened ventral surface with more than 2 rows of cirri; 2 marginal rows; frontals undifferentiated; anals; no caudals; salt or fresh water. A few species.

*E. pluvialis* Smith (Fig. 356, d). About 375μ long; fresh water.

Genus *Holosticha* Wrzesniowski. Three frontals along anterior margin; 2 ventral and 2 marginal rows of cirri; anals; fresh or salt water. Numerous species.

*H. vernalis* Stokes (Fig. 356, e). 7 anals; about 180μ long; shallow pools with algae.

*H. hymenophora* S. (Fig. 356, f). 5 anals; 2 contractile vacuoles; 160–200μ long; shallow pools.

Genus *Paraholosticha* Kahl. Elongate-oval; flexible; ventral cirri in 2 parallel oblique rows; with a row of stiff cirri along frontal margin, posterior to it 2 short rows of cirri; marginals continuous or interrupted at posterior border; fresh water.

*P. herbicola* K. (Fig. 356, g). 150–190μ long; fresh water among algae.

Genus *Trichotaxis* Stokes. Similar to *Holosticha*; but with 3 rows of ventral cirri; fresh or salt water.

*T. stagnatilis* S. (Fig. 356, h). About 160μ long; ellipsoid; in fresh water among decaying vegetation.

Genus *Balladyna* Kowalewski. Ellipsoid; frontals not well developed or lacking; 1 ventral and 2 marginal rows of cirri; long dorsal and lateral stiff cirri; fresh water.

*B. elongata* Roux (Fig. 356, i). 32–35μ by 11–12μ; fresh water among plants and detritus.

Genus *Pleurotricha* Stein. Oblong to ellipsoid; marginals continuous; 8 frontals; 3–4 ventrals; 7 anals of which 2 are more posterior; 2 rows of ventral cirri; between ventrals and marginals 1–3 rows of few coarse cilia; fresh water.

*P. lanceolata* (Ehrenberg) (Fig. 356, j). 100–165μ long; 2 macro- and 2 micro-nuclei. Manwell (1928) studied its conjugation, division, encystment and nuclear variation. Encystment (Penn, 1935).

Genus *Gastrostyla* Engelmann. Frontals distributed except 3 along the frontal margin; ventrals irregular; 5 anals; macronucleus divided
into 2–8 parts; fresh or salt water. Morphology and physiology (Weyer, 1930).

*G. muscorum* Kahl (Fig. 356, k). 130–200μ long; macronucleus in 8 parts; fresh water in vegetation.

Genus *Stylonychia* Ehrenberg. Ovoid to reniform; not flexible; ventral surface flat, dorsal surface convex; 8 frontals; 5 ventrals; 5 anals; marginals; 3 caudals; with short dorsal bristles; fresh or salt water. Many species.

*G. muscorum* Kahl (Fig. 356, k). 130–200μ long; macronucleus in 8 parts; fresh water in vegetation.

Genus *Stylonychia* Ehrenberg. Ovoid to reniform; not flexible; ventral surface flat, dorsal surface convex; 8 frontals; 5 ventrals; 5 anals; marginals; 3 caudals; with short dorsal bristles; fresh or salt water. Many species.

Genus *Stylonychia* Ehrenberg. Ovoid to reniform; not flexible; ventral surface flat, dorsal surface convex; 8 frontals; 5 ventrals; 5 anals; marginals; 3 caudals; with short dorsal bristles; fresh or salt water. Many species.

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Fig. 357. a, Stylonychia mytilus, X200 (Stein); b, S. pustulata, X400 (Roux); c, S. putrina, X200 (Stokes); d, S. notophora, X200 (Stokes); e, Onychodromus grandis, X230 (Stein); f, Onychodromopsis flexilis, X240 (Stokes); g, Euplotes patella, X420 (Pierson); h, E. eurystomus, X330 (Pierson); i, E. woodruffi, X310 (Pierson); j, E. aediculatus, X290 (Pierson).
**E. eurystomus** Wrzesniowski (Fig. 357, h). Elongated ellipsoid; length 100–195μ; average dimensions 138μ by 78μ; 9 frontal-ventrals; no aboral ridges, but 7 rows of bristles; peristome wide, deep; peristomal depression sigmoid; membranellae forming sigmoid curve; end of cytopharynx far to left and anterior to the fifth anal cirrus; post-pharyngeal sac; macronucleus 3-shaped; micronucleus near flattened anterior corner of macronucleus; fresh and brackish water. Division and conjugation (Turner, 1930); neuromotor system (Turner, 1933; Hammond, 1937; Hammond and Kofoi, 1937).

**E. woodruffi** Gaw (Fig. 357, i). Oval; length 120–165μ; average dimensions 140μ by 90μ; 9 frontal-ventrals; aboral surface often with 8 low ridges; peristome wide, with a small peristomal plate; end of cytopharynx almost below the median ridge; 4th ridge between anal cirri often extends to anterior end of body; post-pharyngeal sac; macronucleus consistently T-shaped; micronucleus anterior-right; brackish (with salinity 2.30 parts of salt per 1000) and fresh water (Gaw, 1939).

**E. aediculatus** Pierson (Fig. 357, j). Elliptical; length 110–165μ; average dimensions 132μ by 84μ; 9 frontal-ventrals; aboral surface usually without ridges, but with about 6 rows of bristles; peristome narrow; peristomal plate long triangular, drawn out posteriorly; a niche midway on the right border of peristome; anal cirri often form a straight transverse line; 4th ridge between anals may reach anterior end of body; macronucleus C-shape with a flattened part in the left-anterior region; micronucleus some distance from macronucleus at anterior-left region; post-pharyngeal sac; fresh and brackish (salinity 2.30 parts of salt per 1000) water.

**E. plumipes** Stokes. Similar to **E. eurystomus**. About 125μ long; fresh water.

**E. carinatus** S. (Fig. 358, a). About 70μ by 50μ; fresh water.

**E. charon** (Müller) (Fig. 358, b). 70–90μ long; salt water.

Genus **Euplotidium** Noland. Cylindrical; 9 frontal-ventrals in 2 rows toward right; 5 anals; a groove extends backward from oral region to ventral side, in which the left-most anal cirrus lies; peristome opened widely at anterior end, but covered posteriorly by a transparent, curved, flap-like membrane; adoral zone made up of about 80 membranellae; longitudinal ridges (carinae), 3 dorsal and 2 lateral; a row of protrichocysts under each carina; a broad zone of protrichocysts in antero-dorsal region; cytoplasm densely granulated; salt water. One species (Noland, 1937).

**E. agitatum** N. (Fig. 358, c, d). 65–95μ long; erratic movement rapid; observed in half-dead sponges in Florida.
Fig. 358. a, Euplotes carinatus, ×430 (Stokes); b, E. charon, ×440 (Kahl); c, d, Euplotidium agitatum, ×540 (Noland); e, Certesia quadrinucleata, ×670 (Sauerbrey); f, Diophrys appendiculata, ×570 (Wallengren); g, Uronychia setigera, ×870 (Calkins); h, Aspidisca lyncus, ×300 (Stein); i, A. polystyla, ×290 (Kahl).
Genus *Certesia* Fabre-Domergue. Ellipsoid; flattened; dorsal surface slightly convex, ventral surface flat or concave; 5 frontals at anterior border; 7 ventrals; 5 anals; no caudals; marginals small in number; 4 macronuclei; salt water. One species.

*C. quadrinucleata* F.-D. (Fig. 358, e). 70-100μ by about 45μ. Morphology (Sauerbrey, 1928).

Genus *Diophrys* Dujardin. Peristome relatively large, often reaching anals; 7-9 frontal-ventrals; 5 anals; 3 strong cirri right-dorsal near posterior margin; salt water.

*D. appendiculata* (Ehrenberg) (Fig. 358, f). 60-100μ long; salt water; Woods Hole (Calkins). Division and reorganization (Summers, 1935).

Genus *Uronychia* Stein. Without frontals and ventrals; 5 anals; 3 right-dorsal cirri (as in *Diophrys*); 2 left-ventral cirri near posterior margin; peristome, oval with a large undulating membrane on right edge; salt water. Several species.

*U. setigera* Calkins (Fig. 358, g). 40μ by 25μ; salt water; Woods Hole.

Genus *Gastrocirrhus* Lepsi. Anterior end truncate with a ring of cilia; posterior end bluntly pointed; slightly flattened; a wide peristome leading to cytostome, with undulating membrane on left; 16 cirri on ventral surface arranged on right and posterior margins (Lepsi) or six frontals, five ventrals, five caudals (Bullington); marine. Apparently intermediate between Heterotricha and Hypotricha (Lepsi).

*G. stentoreus* Bullington (Fig. 359, a). About 104μ by 71-81μ; dark granulated cytoplasm; active jumping as well as swimming movement; in Tortugas (Bullington, 1940).

Family 3 *Paraeuplotidae* Wichterman

Genus *Paraeuplotes* Wichterman. Ovoid; ventral surface slightly concave, dorsal surface highly convex and bare, but with one ridge; frontal and adoral zones well developed; ventral surface with a semi-circular ciliary ring on the right half, posterior half of which is marked by a plate and with two ciliary tufts, near the middle of anterior half; 5-6 caudal cirri; macronucleus curved band-form; a terminal contractile vacuole; zooxanthellae, but no food vacuole in cytoplasm; marine, on the coral.

*P. tortugensis* W. (Fig. 359, b, c). Subcircular to ovoid; average individuals 85μ by 75μ; ciliary plate 37μ long, with longer cilia; adoral zone reaches nearly the posterior end; “micronucleus not clearly differentiated” (Wichterman); 5-6 caudal cirri about 13μ
long; zooxanthellae yellowish brown, about 12µ in diameter, fill the body; found on *Eunicea crassa* (coral); Tortugas, Florida.

Genus *Euplotaspis* Chatton and Seguela. Ellipsoid; ventral surface flat or slightly concave; dorsal surface convex; membranelles and cirri with fringed tips; peristome very long; 10 frontal-ventrals; five anals; three or four caudals difficult to see in life; dorsal surface without striae or ciliary processes; macronucleus arched band; a single micronucleus. One species (Chatton and Seguela, 1936).

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**Fig. 359.** a, ventral view of *Gastrocirrhus stentoreus*, ×330 (Bullington) (ac, anterior cirri; cc, caudal cirri; cp, cytopharynx; oc, oral cilia; om, oral membrane; vc, ventral cirri); b, c, dorsal and ventral views of *Paraeuplotes tortugensis*, ×490 (Wichterman) (avc, anterior ventral cilia; cp, ciliary plate; cpc, ciliary plate cilia; cv, contractile vacuole; ds, dorsal swelling; fm, frontal membranelles; om, adoral membranelles; sp, caudal cirri; tc, tufts of cilia; zo, zoothanthellae); d, ventral view of *Euplotaspis cionaecola*, ×1285 (Chatton and Seguela).
E. cionaccola C. and S. (Fig. 359, d). 60–70μ by 45–55μ; in the branchial cavity of the asidian, Ciona intestinalis.

Family 4 Aspidiscidae Claus

Genus Aspidisca Ehrenberg. Small; ovoid; inflexible; right and dorsal side convex, ventral side flattened; dorsal surface conspicuously ridged; adoral zone reduced or rudimentary; 7 frontal-ventrals; 5–12 anals; macronucleus horseshoe-shaped or occasionally in 2 rounded parts; contractile vacuole posterior; fresh or salt water. Numerous species.

A. lyneus E. (Figs. 55; 358, h). 30–50/x long; fresh water. Division and reorganization (Summers, 1935).

A. polystyla Stein (Fig. 358, i). About 50/x long; marine; Woods Hole (Calkins).

References


— (1947) II. Ibid., 1:133.


Chapter 42
Order 3 Chonotricha Wallengren

These ciliates live attached to aquatic animals, especially crustaceans and have developed a peculiar organization. The body is, as a rule, vase-form with an apical peristome, around which extends a more or less complicated ectoplasmic collar or funnel and along which are found ciliary rows that lead to the deeply located cytopharynx. The macronucleus is oval and situated centrally; there is a contractile vacuole usually near the cytopharynx. Asexual reproduction is by lateral budding, and conjugation has been observed in a few species. Taxonomy (Kahl, 1935); distribution (Mohr, 1948).

Family Spirochonidae Stein

Genus Spirochona Stein. Peristome funnel spirally wound; ciliary zone on floor of the spiral furrow; attached to Gammarus in fresh water. Many species (Swarczewsky, 1928).

S. gemmipara S. (Fig. 360, a). 80–120μ long; attached to the gill-plates of Gammarus pulex and other species. Morphology (Guilcher, 1950).

Genus Stylochona Kent. Peristomal funnel with an inner funnel. One species.

S. coronata K. (Fig. 360, b). About 60μ long; on marine Gammarus.

Genus Kentrochona Rompel (Kentrochonopsis Doflein). Peristomal funnel wide, simple, membranous; with or without a few (2) spines.

K. nebaliae R. (Fig. 360, c). About 40μ long; much flattened, with its broad side attached by means of gelatinous substance to epipodite of Nebalia geoffroyi; salt water.

Genus Trichochona Mohr. Elongate; with a long stalk; pellicle thick; a single and simple funnel; two ciliary patches, one parallel to funnel rim and the other diagonal in the deep part of funnel; one macronucleus; one to four micronuclei; budding; marine. One species (Mohr, 1948).

T. lecythoides M. (Fig. 360, d, e). Body 35–86μ by 3–28μ; funnel 8–21.5μ high; stalk 16–51μ long; peristomal funnel with horizontal ciliary lines, up to 32; diagonal lines about 20; on the appendages of the marine crustacean, Amphithoe sp.

Genus Heliochona Plate. Peristomal funnel with numerous needle-like spines. Taxonomy (Wallengren, 1895; Guilcher, 1950).
Fig. 360. a, *Spirochona gemmipara*, ×300 (Hertwig); b, *Stylochona coronata*, ×400 (Kent); c, *Kentrochona nebaliae*, ×970 (Rompel); d, e, *Trichochona lecythoides* (Mohr) (d, a portion of a host’s appendage with 16 attached organisms, ×110; e, an individual, ×405); f, *Heliochona scheuteni*, ×550 (Wallengren); g, *H. sessilis*, ×510 (Wallengren); h, *Chilodochona quennerstedti*, ×400 (Wallengren).

*H. scheuteni* (Stein) (Fig. 360, f). About 80–90μ long; on appendages of *Gammarus locusta*; salt water.

*H. sessilis* P. (Fig. 360, g). About 60μ long; on *Gammarus locusta*; salt water.

Genus *Chilodochona* Wallengren. Peristome drawn out into two lips; with a long stalk.

*C. quennerstedti* W. (Fig. 360, h). 60–115μ long; stalk, 40–160μ; on *Ebalia turnefacta* and *Portunus depurator*; salt water.

References


Mohr, J. L.: (1948) Trichochona lecythroides, a new genus and species, etc. Allan Hancock Found. Publ., Occasional Papers, no. 5.


THE peritrichous ciliates possess a much enlarged disk-like anterior region which is conspicuously ciliated. The adoral zone is counter-clockwise to the eyestome viewed from the anterior end. The body ciliation is more or less limited. The stalked forms produce free-swimming individuals, *telotrochs*. Asexual reproduction is by binary fission; and conjugation occurs commonly. The majority are free-living or attached to various aquatic animals and plants, although a few are parasitic. Taxonomy (Kahl, 1935; Stiller, 1939, 1940; Nenninger, 1948); structure of stalk (Precht, 1935).

Attached to submerged objects; usually no body cilia, though telotroch possesses a posterior ring of cilia........Suborder 1 Sessilia Free-swimming; but with highly developed attaching organellae on aboral end..................Suborder 2 Mobilia (p. 859)

Suborder 1 Sessilia Kahl

Without lorica, although some with a gelatinous or mucilaginous envelope..........................Tribe 1 Aloricata

With definite pseudochitinous lorica ..........Tribe 2 Loricata (p. 857)

Tribe 1 Aloricata Kahl

Posterior end with 1–2 short spines; swimming with peristome-bearing end forward..................Family 1 Astylozooidae

Posterior end, directly or indirectly through stalk, attached to submerged objects

Anterior region a long cylindrical, highly contractile neck; contractile vacuole connected with vestibule by a long canal; reservoir of contractile vacuole distinct; with or without a thin stalk..............

.................................Family 2 Ophryidiidae (p. 852)

Anterior portion not drawn out into a neck

Without stalk.....................Family 3 Scyphidiidae (p. 852)

With stalk

Stalk non-contractile............Family 4 Epistyliidae (p. 853)

Stalk contractile................Family 5 Vorticellidae (p. 855)

Family 1 Astylozooidae Kahl

Genus *Astylozoon* Engelmann (*Geleiella* Stiller). Free-swimming; pyriform or conical; aboral end attenuated, with 1–2 thigmotactic stiff cilia; pellicle smooth or furrowed; with or without gelatinous envelope; in fresh water. A few species.

*A. fallax* E. (Fig. 361, a). 70–100μ; fresh water.

Genus *Hastatella* Erlanger. Free-swimming; body surface with 2–4 rings of long conical ectoplasmic processes; fresh water.
H. aesculacantha Jarocki and Jacubowska (Fig. 361, b). 30–52μ by 24–40μ; in stagnant water.

Genus Opisthonecta Fauré-Fremiet. Conical; ends broadly rounded; a ring of long cilia close to aboral end; adoral zone about 1.1 turns, composed of 2 parallel rows; a papilla with about 12 long cilia, just above the opening into vestibule; macronucleus sausage-

![Fig. 361.](image-url)
form; micronucleus; 3 contractile vacuoles connected with cytopharynx; fresh water. One species.

O. henneguyi F.-F. (Fig. 361, c, d). 148–170μ long; cysts about 57μ in diameter; sometimes infected by a parasitic suctorian, Endosphaera engelmannii (Lynch and Noble, 1931) (p. 873). Conjugation (Rosenberg, 1940); neuromotor system (Kofoid and Rosenberg, 1940); encystment (Rosenberg, 1938).

Family 2 Ophrydiidae Kent

Genus Ophrydium Ehrenberg (Gerda Claparède and Lachmann). Cylindrical with a contractile neck; posterior end pointed or rounded; variable number of individuals in a common mucilaginous mass; pellicle usually cross-striated; fresh water.

O. sessile Kent (Fig. 361, e). Fully extended body up to 300μ long; colorless or slightly brownish; ovoid colony up to 5 mm. by 3 mm.; attached to freshwater plants.

O. vernalis (Stokes) (Fig. 361, f). About 250μ long; highly contractile; in shallow freshwater ponds in early spring (Stokes).

O. ectatum Mast (Fig. 361, g). 225–400μ long; with many zoochlorellae; colony up to 3 mm. in diameter; in fresh water (Mast, 1944).

Family 3 Scyphidiidae Kahl

Genus Scyphidia Dujardin. Cylindrical; posterior end attached to submerged objects or aquatic animals; body usually cross-striated; fresh or salt water. Species (Nenninger, 1948).

S. amphibiārum Nenninger (Fig. 361, h). On tadpoles; about 76μ long.

Genus Paravorticella Kahl. Similar to Scyphidia; but posterior portion is much elongated and contractile; salt water, attached or parasitic.

P. clymenellae (Shumway) (Fig. 361, i). 100μ long; in the colon of the annelid, Clymenella torquata; Woods Hole.

Genus Glossatella Bütschli. With a large adoral membrane; often attached to fish and amphibian larvae.

G. tintinnabulum (Kent) (Fig. 362, a). 30–43μ long; attached to the epidermis and gills of young Triton.

Genus Ellobiophrya Chatton and Lwoff. Posterior end drawn out into 2 arm-like processes by means of which the organism holds fast to the gill bars of the mussel, Donax vittatus. One species.

E. donacis C. and L. (Fig. 362, b). 50μ by 40μ, excluding the processes.
Family 4 Epistylidae Kent

Genus *Epistylis* Ehrenberg. Inverted bell-form; individuals usually on dichotomous non-contractile stalk, forming large colonies; attached to fresh or salt water animals. Numerous species (Nenninger, 1948).

*E. plicatilis* E. (Fig. 362, c). 110–162 μ long (Nenninger); colony often up to 3 mm. high; in fresh water.
E. fugitans Kellicott. 50–60 μ long; attached to Sida in early spring.
E. cambari K. (Fig. 362, d, e). About 50 μ long; attached to the gills of Cambarus.

E. niagarae (Fig. 362, f). Expanded body about 160 μ long; peristomal ring prominent; flat cap makes a slight angle with the ring; bandform macronucleus transverse to long axis, in the anterior third; gullet with ciliated wall; 40–50 in a colony; attached to the antennae and body surface of crayfish (Kellicott, 1883) or to painted and snapping turtles (Bishop and Jahn, 1941).

Genus Rhabdostyla Kent. Similar to Epistylis; but solitary with a non-contractile stalk; attached to aquatic animals in fresh or salt water. Numerous species (Nenninger, 1948).

R. vernalis Stokes (Fig. 362, g). About 50 μ long; attached to Cyclops and Cypris in pools in early spring.

Genus Opisthostyla Stokes. Similar to Rhabdostyla; but stalk long, is bent at its point of attachment to submerged object, and acts like a spring; fresh or salt water (Nenninger, 1948).

O. annulata S. (Fig. 362, h). Body about 23 μ long; fresh water.

Genus Campanella Goldfuss. Similar to Epistylis; but adoral double zone turns 4–6 times; fresh water.

C. umbellaria (Linnaeus) (Fig. 362, i). Colony may reach several millimeters in height; individuals 130–250 μ long (Kent).

Genus Pyxidium Kent. Stalk simple, not branching; peristome even when fully opened, not constricted from the body proper; frontal disk small, oblique, supported by style-like slender process arising from peristome; attached to freshwater animals and in vegetation. Taxonomy (Nenninger).

P. vernale Stokes (Fig. 362, j). Solitary or few together; 70–85 μ long; fresh water among algae.

P. urceolatum S. (Fig. 362, k). About 90 μ long; fresh water on plants.

Genus Opercularia Stein. Individuals similar to Pyxidium; but short stalk dichotomous; peristome border like a band.

O. stenostoma S. (Fig. 362, l). When extended, up to 125 μ long; attached to Asellus aquaticus and others.

O. plicatilis Stokes (Fig. 362, m). About 254 μ long; colony 1.25–2.5 mm. high; pond water.

Genus Operculariella Stammer. Fixed stalk, branched, short and rigid; peristome small, without border, smooth; without disk or frontal cilia; vestibule large (Stammer, 1948).

O. parasitica S. (Fig. 362, n). 100–110 μ long; barrel-shaped; peristome opening only 1/4 the body breadth; macronucleus about 30 μ
long; parasitic in the oesophagus of *Dytiscus marginalis*, *Acilius sulcatus*, *Hydaticus transversalis*, *Graphoderes zonatus* and *G. bilineatus*.

Family 5 *Vorticellidae* Fromental

Genus *Vorticella* Linnaeus. Inverted bell-form; colorless, yellowish, or greenish; peristome more or less outwardly extended; pellicle sometimes annulated; with a contractile stalk, macronucleus band-form; micronucleus; 1–2 contractile vacuoles; solitary; in fresh or salt water, attached to submerged objects and aquatic plants or animals. Numerous species. Taxonomy (Noland and Finley, 1931;

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![Diagram of Vorticella species](image-url)

*Fig. 363. a–e, Vorticella campanula (a, ×400; b, part of stalk, ×800; c, telotroch, ×200); d, e, V. convallaria (d, ×400; e, ×800); f–p, V. microstoma (f, g, ×400; h, ×840; i, telotroch, ×400; j–p, telotroch-formation in vitro, ×270); q, r, V. picta (q, ×400; r, ×800); s, t, V. monilata (s, ×400; t, ×800) (Noland and Finley).*
Kahl, 1935; Nenninger, 1948); movements of food vacuoles (Hall and Dunihue, 1931).

V. *campanula* Ehrenberg (Fig. 363, a–c). Usually in groups; endoplasm filled with refractile reserve granules; vestibule very large with an outer pharyngeal membrane; 50–157μ by 35–99μ; peristome 60–125μ wide; stalk 50–4150μ by 5.6–12μ fresh water.

V. *convallaria* (L.) (Fig. 363, d, e). Resembles the last-named species; but anterior end somewhat narrow; usually without refractile granules in endoplasm; 50–95μ by 35–53μ; peristome 55–75μ wide; stalk 25–460μ by 4–6.5μ; fresh water.

V. *microstoma* Ehrenberg (Figs. 86; 363, f–p). 35–83μ by 22–50μ; peristome 12–25μ wide; stalk 20–385μ by 1.5–4μ; common in fresh-water infusion. Conjugation (Finley, 1943); encystment (von Brand, 1923).

V. *picta* (E.) (Fig. 363, q, r). 41–63μ by 20–37μ; peristome 35–50μ; stalk 205–550μ by 4–7μ; 2 contractile vacuoles; with refractile granules in stalk; fresh water.

V. *monilata* Tatem (Fig. 363, s, t). Body with pellicular tubercles composed of paraglycogen (Fauré-Fremiet and Thaureaux, 1944); 2 contractile vacuoles; 50–78μ by 35–57μ; peristome 36–63μ wide; stalk 50–200μ by 5–6.5μ; fresh water.

**Genus Carchesium** Ehrenberg. Similar to *Vorticella*; but colonial; myonemes in stalk not continuous, and therefore individual stalks contract independently; attached to fresh or salt water animals or plants; occasionally colonies up to 4 mm. high. Several species (Kahl, 1935; Nenninger, 1948).

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**Fig. 364.** a, *Carchesium polypinum*, ×200 (Stein); b, *C. granulatum*, ×220 (Kellicott); c, *Zoothamnium arbuscula*, ×200 (Stein); d, *Z. adamsi*, ×150 (Stokes).
C. *polypinum* (Linnaeus) (Fig. 364, a). 100–125μ long; colony up to 3 mm. long; fresh water.

C. *granulatum* Kellicott (Fig. 364, b). About 100μ long; 2 contractile vacuoles anterior; on Cambarus and aquatic plants.

Genus *Zoothamnium* Bory. Similar to *Carchesium*; but myonemes (Fig. 15) of all stalks of a colony are continuous with one another, so that the entire colony contracts or expands simultaneously; fresh or salt water; colonies sometimes several millimeters high. Numerous species (Kahl, 1935; Nenninger, 1948). Development (Summers, 1938, 1938a).

Z. *arbuscula* Ehrenberg (Fig. 364, c). 40–60μ long; colony up to more than 6 mm. high; fresh water. Morphology and life cycle (Furssenko, 1929).

Z. *adamsi* Stokes (Fig. 364, d). About 60μ long; colony about 250μ high; attached to Cladophora.

**Tribe 2 Loricata** Kahl

Peristomal margin not connected with lorica; body attached only at posterior end, and extends, out of lorica. Family 1 Vaginicolidae

Peristomal margin connected with inner margin of aperture of lorica; stalked disk only extends out of lorica.

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Family 2 Lagenophryidae (p. 691)

Family 1 Vaginicolidae Kent

Genus *Vaginicola* Lamarck. Lorica without stalk, attached to substratum directly with its posterior end; body elongate and cylindrical; fresh or salt water. Numerous species (Swarczewsky, 1930).

V. *leptosoma* Stokes (Fig. 365, a). Lorica about 160μ high; when extended, about 1/3 of body protruding; on algae in pond water.

V. *annulata* S. (Fig. 365, b). Lorica about 120μ high; below middle, a ring-like elevation; anterior 1/3 of body protruding, when extended; pond water.

Genus *Cothurnia* Ehrenberg. Similar to *Vaginicola*; but lorica stands on a short stalk; fresh or salt water. Numerous species (Swarczewsky, 1930).

C. *canthocampti* Stokes (Fig. 365, c). Lorica about 80μ high; on *Canthocamptus minutus*.

C. *annulata* S. (Fig. 365, d). Lorica about 55μ high; fresh water.

Genus *Thuricola* Kent. Body and lorica as in *Vaginicola*; but lorica with a simple or complex valve-like apparatus which closes obliquely after the manner of a door when protoplasmic body contracts; salt or fresh water.
Fig. 365. a, Vaginicola leptosoma, ×130 (Stokes); b, V. annulata, ×170 (Stokes); c, Cothurnia canthocampti, ×150 (Stokes); d, C. annulata, ×340 (Stokes); e, Thuricola folliculata, ×110 (Kahl); f, Thuricolopsis kellicottiana, ×110 (Stokes); g, Caulicola valvata, ×760 (Stokes); h, i, Pyxicola affinis, ×170 (Kent); j, P. socialis, ×170 (Kent); k, Platycola longicollis, ×200 (De Fromentel); l, Lagenophrys vaginicola, ×380 (Penard); m, L. patina, ×150 (Stokes); n, L. labiata, ×340 (Penard).

T. folliculata (Müller) (Fig. 365, e). Lorica 127–170μ high (Kent); 160–200μ high (Kahl); salt and fresh water.

Genus Thuricolopsis Stokes. Lorica with an internal, narrow, flexible valve-rest, adherent to lorica wall and projecting across cavity to receive and support the descended valve; protoplasmic body attached to lorica by a pedicel; on freshwater plants.

T. kellicottiana S. (Fig. 365, f). Lorica about 220μ long.

Genus Caulicola Stokes. Similar to Thuricola; but lorica-lid attached to aperture; fresh or brackish water. 2 species.

C. valvata S. (Fig. 365, g). Lorica about 50μ high; stalk about 1/2; body protrudes about 1/3 when extended; brackish water.

Genus Pyxicola Kent. Body attached posteriorly to a corneous lorica; lorica colorless to brown, erect, on a pedicel; a discoidal corneous operculum developed beneath border of peristome, which closes lorica when organism contracts; fresh or salt water. Many species.

P. affinis K. (Fig. 365, h, i). Lorica about 85μ long; in marsh water.
P. socialis (Gruber) (Fig. 365, j). Lorica about 100μ long; often in
groups; salt water.

Genus Platycola Kent. Body similar to that of Vaginicola; but
lorica always decumbent and attached throughout one side to its
fulcrum of support; fresh or salt water. Many species.
P. longicollis K. (Fig. 365, k). Lorica yellow to brown when older;
about 126μ long; fresh water.

Family 2 Lagenophryidae Bütschli

Genus Lagenophrys Stein. Lorica with flattened adhering surface,
short neck and convex surface; "striped body" connects body with
lorica near aperture; attached to fresh or salt water animals. Many
species (Swarczewsky, 1930). Biology (Awerinzew, 1936).
L. vaginicola S. (Fig. 365, l). Lorica 70μ by 48μ; attached to caudal
bristles and appendages of Cyclops minutus and Canthocamptus sp.
L. patina Stokes (Fig. 365, m). Lorica 55μ by 50μ; on Gammarus.
L. labiata S. (Fig. 365, n). Lorica 60μ by 55μ; on Gammarus.

Suborder 2 Mobilia Kahl

Family Urceolariidae Stein

Genus Urceolaria Lamarck. Peristome more or less obliquely
placed; external ciliary ring difficult to see; horny corona of attach-
ing disk with obliquely arranged simple teeth without radial pro-
tesses; commensal. A few species. Morphology (Wallengren, 1897).
U. mitra (Siebold) (Fig. 366, a). 80–140μ long; on planarians.
U. paradoxa (Claparède and Lachmann) (Fig. 366, b). 70–80μ
in diameter; colonial forms; in the respiratory cavity of Cyclostoma
elegans.
U. karyolobia Hirshfield. 45–50μ in diameter, 20–30μ high; mac-
ronucleus lobate and conspicuous; in the mantle cavity of limpets,
Lottia gigantea and Acmaea spp. (Hirshfield, 1949).
Genus Trichodina Ehrenberg. Low barrel-shaped; with a row of
posterior cilia; horny ring of attaching disk with radially arranged
hooked teeth; commensal on, or parasitic in, aquatic animals. Several
species (Mueller, 1932, 1937). Structure (Wallengren, 1897a); bi-
ometry (Fauré-Fremiet, 1943).
T. pediculus (Müller) (Fig. 366, c). A shallow constriction in mid-
dle of body; 50–70μ in diameter; on fish. Those found on Hydra and
on the gills of Necturus and Triturus larvae are probably this species
(Fulton, 1923). Reproduction (Cavallini, 1931).
T. urinicola Fulton (Fig. 366, d). 50–90μ long; teeth 28–36; in
Fig. 366. a, Urceolaria mitra, $\times 270$ (Wallengren); b, U. paradoxa, $\times 215$ (Claparède and Lachmann); c, Trichodina pediculus, $\times 425$ (James-Clark); d, T. urinicola, $\times 470$ (Fulton); e, T. ranae (Cunha); f, T. sp., $\times 460$ (Diller); g, Cyclochaeta spongillae, $\times 460$ (Jackson); h, i, C. domerguei, $\times 535$ (MacLennan).

urinary bladder of a moribund *Bufo* sp. (Fulton) and in frogs Fauré-Fremiet and Mugard, 1946).

*T. sp.* Diller (Fig. 366, e). 30–40 $\mu$ in diameter; on the skin and gills of frog and toad tadpoles. Division (Diller, 1928).

*T. ranae* da Cunha (Fig. 366, f). 40–50 $\mu$ in diameter, 30–50 $\mu$ high; 23–31 V-shaped teeth on the attaching ring; in the urinary bladder of *Rana ridibunda perezi* (da Cunha, 1950).

Genus *Cyclochaeta* Jackson. Saucer-form; peristomal surface parallel to the basal disc; upper surface with numerous flat wrinkles; basal disc composed of cuticular rings, velum, cirri, and membranel-
lae; commensal on, or parasitic in, fresh or salt water animals. Several species. MacLennan (1939) made a careful study of two species. 

*C. spongillae* J. (Fig. 366, g). About 60µ in diameter; in interstices of *Spongilla fluviatilis.*

*C. domerguei* Wallengren (Fig. 366, h, i). 23–56µ in diameter; about one-fifth high; 18–25 denticles, each with a narrow slightly curved spine; outer cuticular ring more finely striated than inner ring; cirri longer than membranellae (MacLennan, 1939); on fresh water fishes.

**References**


—— and Mugard, Hélène: (1946) Une trichodine parasite endovésicale chez *Rana esculenta.* Ibid., 71:36.


—— (1938a) Form regulation in Zoothamnium alterans. Ibid., 74:130.


Chapter 44

Class 2 Suctoria Claparède and Lachmann

The Suctoria, which have been also known as Acinetaria, Tentaculifera, etc., do not possess any cilia or any other cell-organs of locomotion in the mature stage. The cilia are present only on young individuals which are capable of free-swimming, and lost with the development of a stalk or attaching disk, and of tentacles. Therefore, an adult suctorian is incapable of active movement. The body may be spheroidal, elliptical, or dendritic; and is covered with a pellicle and occasionally possesses a lorica. There is no cytostome, and the food-capturing is carried on exclusively by the tentacles. Tentacles are of two kinds: one is suctorial in function and bears a rounded knob on the extremity and the other is for piercing through the body of a prey and more or less sharply pointed. The tentacles may be confined to limited areas or may be distributed over the entire body surface. The food organisms are usually small ciliates and nutrition is thus holozoic.

Asexual reproduction is by binary fission or by budding. The buds which are formed by either exogenous or endogenous gemmation are ciliated, and swim around actively after leaving the parent individual. Finally becoming attached to a suitable object, the buds metamorphose into adult forms. Sexual reproduction is through a complete fusion of conjugants. Relation to prostomatous ciliates (Kahl, 1931); morphogenesis (Guilcher, 1950).

The Suctoria live attached to animals, plants or non-living matter submerged in fresh or salt water, although a few are parasitic.

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Family 1 Dendrosomidae Bütschli

Genus Dendrosoma Ehrenberg. Dendritic; often large; nucleus band-form, branched; numerous contractile vacuoles; fresh water. Taxonomy and morphology (Gönnert, 1935).

Fig. 367. a, Dendrosoma radians, ×35 (Kent); b, Trichophrya epistyloidis ×250 (Stokes); c, T. salparum, ×170 (Collin); d, T. columbiae, ×200 (Wailes); e, T. micropteri, ×650 (Davis); f, Erastophrya chattoni (Fauré-Fremiet); g, Astrophrya arenaria, ×65 (Awerinzew); h, Lernaeophrya capitata, ×35 (Pérez); i, Dendrosomides paguri, ×200 (Collin).
D. radians E. (Fig. 367, a). Brownish; 1.2–2.5 mm. high; on vegetation. Morphology (Gönnert).

Genus Trichophrya Claparède and Lachmann (Platophrya Gönnert). Body small; rounded or elongate, but variable; without stalk; tentacles in fascicles, not branching; simple or multiple endogenous budding; fresh or salt water.

T. epistylidis C. and L. (T. sinuosa Stokes) (Fig. 367, b). Form irregular; with many fascicles of tentacles; nucleus band-form, curved; numerous vacuoles; up to 240μ long; on Epistylis, etc., in fresh water. Morphology (Gönnert).

T. salparum Entz (Fig. 367, c). On various tunicates such as Molgula manhattensis; 40–60μ long; tentacles in 2 groups; salt water; Woods Hole (Calkins).

T. columbiae Wailes (Fig. 367, d). 60–75μ by 40–48μ in diameter; cylindrical; tentacles at ends; nucleus spherical; in marine plankton; Vancouver (Wailes).

T. micropteri Davis (Fig. 367, e). Body elongate, irregular or rounded; up to 30–40μ long by 10–12μ; fully extended tentacles 10–12μ long; cytoplasm often filled with yellow to orange spherules; a single micronucleus; a single contractile vacuole; attached to the gill of small mouth black bass, Micropterus dolomieu. Davis (1942) states that when abundantly present, the suctorian may cause serious injury to the host.

Genus Erastophrya Fauré-Fremiet. Pyriform; distributed tentacles; posterior end drawn out into two “arms” by means of which the organism grasps the stalk of a peritrich; fresh water (Fauré-Fremiet, 1943). One species.

E. chattoni F.-F. (Fig. 367, f). Body up to 130μ long; macronucleus spherical to sausage form; a single micronucleus; a contractile vacuole; endogenous budding, gemma about 40μ long; a commensal on Glossatella piscicola.

Genus Astrophrya Awerinzew. Stellate; central portion drawn out into 8 elongate processes, each with a fascicle of tentacles; body covered by sand grains and other objects. One species.

A. arenaria A. (Fig. 367, g). 145–188μ in diameter; processes 80–190μ long; in Volga river plankton.

Genus Lernaeophrya Pérez. Body large; with numerous short prolongations, bearing very long multifasciculate tentacles; nucleus branched; brackish water. One species.

L. capitata P. (Fig. 367, h). Attached to the hydrozoan, Cordylophora lacustris in brackish water; 400–500μ long; tentacles 400μ long. Morphology (Gönnert).
Genus **Dendrosomides** Collin. Branched body similar to Dendrosoma, but with a peduncle; reproduction by budding of vermicular form; salt water. One species.

*D. paguri* C. (Fig. 367, i). 200–300μ long; vermicular forms 350μ long; on the crabs, *Eupagurus excavatus* and *E. cuanensis*.

Genus **Rhabdophrya** Chatton and Collin. Elongate, rod-form; with short peduncle, not branched; tentacles distributed over entire sur-

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**Fig. 368.** a, *Rhabdophrya trimorpha*, ×430 (Collin); b, *Staurophrya elegans*, ×200 (Zacharias); c, *Ophryodendron porcellanum*, ×220 (Collin); d, *O. belgicum*, ×270 (Fraipont); e, *Dendrocometes paradoxus*, ×270 (Wrzesnowski); f, *Dendrocometides priscus*, ×220; g, *Discosoma tenella*, ×220; h, *Cometodendron clavatum*, ×220 (Swareczewsky); i, j, *Podophrya fixa* (i, ×400 (Wales); j, ×220 (Collin)); k, *P. elongata*, ×240 (Wailes).
face; macronucleus ellipsoid; micronucleus small; 2–3 contractile vacuoles; salt or brackish water. Several species.

*R. trimorpha* C. and C. (Fig. 368, a). Up to 150μ long; on the copepod, *Cletodes longicaudatus*.

Genus *Staurophrya* Zacharias. Rounded body drawn out into 6 processes.

*S. elegans* Z. (Fig. 368, b). Tentacles not capitate; macronucleus round; 1–2 contractile vacuoles; about 50μ in diameter; in fresh water.

Swarczewsky (1928) established the following genera for the forms he had found in Lake Baikal: *Baikalophrya*, *Stylophrya*, *Baikalodendron* and *Gorgonosoma*.

**Family 2 Ophryodendridae Stein**

Genus *Ophryodendron* Claparède and Lachmann. With one long or 3–6 shorter retractile processes, bearing suctorial tentacles; on Crustacea, Annelida, etc.; salt water. Several species.

*O. porcellanum* Kent (Fig. 368, c). 60–100μ long; on *Porcellana platycheles*, etc.

*O. belgicum* Fraipont (Fig. 368, d). 38–114μ long; vermicular form 100μ; on Bryozoa and hydrozoans; Vancouver (Wailes).

**Family 3 Dendrocometidae Stein**

Genus *Dendrocometes* Stein. Body rounded; with variable number of branched arms; fresh water. Taxonomy (Swarczewsky, 1928a).

*D. paradoxus* S. (Fig. 368, e). Up to 100μ long; on *Gammarus pulex*, *G. puteanus*, etc. Morphology and biology (Pestel, 1932).

Genus *Stylocometes* Stein. Arms not branched; tentacles finger-like; fresh water.

*S. digitatus* (Claparède and Lachmann). Up to 110μ long; on the gills of *Asellus aquaticus* and on *Aphrydium versatile*.

Genus *Dendrocometides* Swarczewsky. Body more or less arched; suctorial tentacles slender, pointed and simple or branched; attached to crustaceans on its broad and circular surface (Swarczewsky, 1928a).

*D. priscus* S. (Fig. 368, f). Diameter 60–65μ, height 18–20μ; on *Acanthogammarus albus*; Lake Baikal.

Genus *Discosoma* S. Discoid; circular in front view; short and pointed tentacles radially arranged, four or six in each row; gemmation, endogenous and simple.

*D. tenella* S. (Fig. 368, g). Diameter 75μ, height 10μ; on *Acanthogammarus victorii*, etc.; Lake Baikal.
Genus Cometodendron S. Body elongate; attached to substrate by a “foot,” well-developed arms; short and pointed tentacles at the ends of arms; simple endogenous gemmation.

C. clavatum S. (Fig. 368, h). 150μ by 40–50μ; the foot 20–22μ; on Acanthogammarus victorii, etc.; Lake Baikal.

Family 4 Podophryidae Büttschli

Genus Podophrya Ehrenberg. Subspherical; normally with a rigid stalk; suckorial tentacles in fascicles or distributed on entire body surface; encystment common; fresh or salt water. Many species.

P. fixa Müller (Fig. 368, i, j). Spherical; tentacles of various lengths; stalked; nucleus spheroid; one contractile vacuole; 10–28μ long; fresh water.

P. collini Root. Ovoid; stalked; 30–60 capitate tentacles, distributed; nucleus spherical; one contractile vacuole; 40–50μ in diameter; in swamp (Root, 1914).

P. elongata Wailes (Fig. 368, k). Elongate; flattened; with a pedicle; tentacles distributed; nucleus cylindrical; 95–105μ long; stalk 65–85μ by 7–9μ; on the marine copepod, Euchaeta japonica; Vancouver.

Genus Parapodophrya Kahl. Spherical; tentacles radiating, a few long, more or less conical at proximal portion; stalk thin; salt water.

P. typha K. (Fig. 369, a). 50–60μ in diameter; salt water (Kahl, 1931).

Genus Sphaerophrya Claparède and Lachmann. Spherical, without stalk; with or without distributed tentacles; multiplication by binary fission or exogenous budding; fresh water, free-living or parasitic.

S. soliformis Lauterborn (Fig. 369, b). Spherical; numerous tentacles about 1/4–1/3 the body diameter; a contractile vacuole; nucleus oval; diameter about 100μ; sapropelic.

S. magna Maupas. Spherical; about 50μ in diameter; numerous tentacles of different length; nucleus spheroid; standing fresh water with decaying vegetation.

S. stentoris M. Parasitic in Stentor; swarmers ciliated on posterior end; the other end with capitate tentacles; nucleus spheroid; 2 contractile vacuoles; about 50μ long.

Genus Paracineta Collin. Spherical to ellipsoidal; tentacles distributed; mostly in salt water, a few in fresh water.

P. limbata (Maupas) (Fig. 369, c, d). With or without gelatinous envelope; 20–50μ in diameter; swarmer with many ciliated bands, contractile; on plants and animals in salt water.
Genus **Metacineta** Bütschli. Lorica funnel-shaped, lower end drawn out for attachment; tentacles grouped at anterior end; nucleus spherical; one contractile vacuole. One species.

*M. mystacina* (Ehrenberg) (Fig. 369, e). Lorica up to 700μ long; in fresh and salt water.

Genus **Urnula** Claparède and Lachmann. Lorica colorless; lower end pointed, attached; aperture narrowed, round or triangular; body
more or less filling lorica; 1–2 (up to 5) long active tentacles; nucleus central, oval; one or more contractile vacuoles; fresh water.

_U. epistylidis_ C. and L. (Fig. 369, f). Up to 80μ long; on Epistylis, Dendrosoma, etc.

Genus _Lecanophrya_ Kahl. Body rounded rectangular in cross section; anterior region bowl-shaped; somewhat rigid tentacles located on the inner surface of bowl; salt water.

_L. drosera_ K. (Fig. 369, g). 40–70μ high; hollow stalk; tentacles in 3–5 indistinct rows; attached to the antennae of the copepod, _Nitocra typica_.

Genus _Ophryocephalus_ Wailes. Spheroidal, stalked; a single long mobile, capitate tentacle; multiplication by multiple exogenous budding from apical region; on _Ephelota gemmipara_ and _E. coronata_ (p. 877); salt water. One species.

_O. capitatum_ W. (Fig. 369, h). About 55μ long; tentacle up to 100μ by 1.5–5μ; Vancouver.

Family 5 _Acinetidae_ Bütschli

Genus _Acineta_ Ehrenberg. Lorica more or less flattened; usually with stalk; tentacles in 2 (1 or 3) fascicles; body completely or partly filling lorica; swarmer with ciliated band or completely ciliated; fresh or salt water. Numerous species (Swarczewsky, 1928a).

_A. tuberosa_ E. (Fig. 370, a). Lorica 50–100μ high; with stalk; salt and brackish water.

_A. cuspidata_ Stokes (Fig. 370, b). Lorica cup-shaped; front end with 2 opposing sharp points; lorica 32–42μ high; on Oedogonium in fresh water.

_A. lacustris_ S. (Fig. 369, i). Lorica elongate ovoid; flattened; 75–185μ high; on Anacharis in pond.

Genus _Tokophrya_ Bütschli. Pyriform or pyramidal; without lorica; tentacles in 1–4 fascicles on anterior surface; stalk not rigid; simple endogenous budding; fresh water. Several species.

_T. infusionum_ (Stein) (Fig. 370, c–e). Inverted pyramid; stalk with or without attaching disk; macronucleus oval; 2 contractile vacuoles; about 60μ long. Relation between contractile vacuole and feeding (Rudzinska and Chambers, 1951); life span (Rudzinska, 1951).

_T. cyclopum_ (Claparède and Lachmann) (Fig. 370, f). Oval or spherical; stalk short; tentacles in 2–5 bundles; macronucleus spherical; 1–2 contractile vacuoles; about 50μ long; on Cyclops, etc.

Genus _Thecacineta_ Collin. Lorica with free margin; body usually attached to bottom of lorica, more or less long; tentacles from anterior end; salt water. Several species (Swarczewsky, 1928).
**Fig. 370.** a, *Acineta tuberosa*, ×670 (Calkins); b, *A. cuspidata*, ×670 (Stokes); c–e, *Tokophrya infusionum* (c, ×400; d, a free-swimming bud; e, a young attached form, ×800) (Collin); f, *T. cyclopum*, a young individual, ×500 (Collin).

*T. cothurnioiides* C. (Fig. 371, a). Lorica about 50μ high; stalk knobbled; on *Cletodes longicaudatus*.

*T. gracilis* (Wailes) (Fig. 371, b). Lorica 110μ by 35μ; stalk 200μ by 4μ; on hydrozoans.

Genus *Periacineta* Collin. Elongate lorica; attached with its drawn-out posterior end; tentacles from the opposite surface in bundles; fresh water.
P. buckei (Kent) (Fig. 371, c). Attached end of lorica with basal plate; 3 contractile vacuoles; up to 125μ long; on Lymnaea stagnalis and Ranatra linearis.

Fig. 371. a, Thecacincta coturnioides, ×400 (Collin); b, T. gracilis, ×270 (Wailes); c, Periacineta buckei, feeding on Chilodonella, ×530 (Collin); d, Hallezia brachypoda, ×200 (Stokes); e, Solenophrya inclusa, ×230 (Stokes); f, S. pera, ×230 (Stokes); g, h, Acinetopsis tentaculata (g, ×130; h, ×230) (Root); i, j, Tachyblaston ephelotensis (i, a young individual in Ephelota, ×260; j, mature form, ×500) (Martin); k, Dactylophrya roscolevita, ×830 (Collin).
Genus Hallezia Sand. Without lorica; with or without a short stalk; tentacles in bundles; fresh water.

*H. brachypoda* (Stokes) (Fig. 371, d). 34–42μ in diameter; in standing water among leaves.

Genus Solenophrya Claparède and Lachmann. Lorica attached directly with its underside; body usually not filling lorica; tentacles in fascicles; fresh water.

*S. inclusa* Stokes (Fig. 371, c). Lorica subspherical; about 44μ in diameter; standing fresh water.

*S. pera* S. (Fig. 371, f). Lorica satchel-form; about 40–45μ high; body about 35μ long; standing fresh water.

Genus Acinetopsis Robin. Lorica in close contact with body on sides; stalked; 1–6 large retractile tentacles and numerous small tentacles from apical end; mainly salt water.

*A. tentaculata* Root (Fig. 371, g, h). Lorica 187μ high; stalk 287μ long; large tentacles up to 500μ long; body about 138μ by 100μ; on *Obelia commissuralis* and *O. geniculata*; Woods Hole (Root, 1922).

Genus Tachyblaston Martin. Lorica with short stalk; tentacles distributed on anterior surface; nucleus oval; salt water. One species.

*T. ephelotensis* M. (Fig. 371, i, j). Lorica 30–95μ high; stalk 20–30μ long; attached to *Ephelota gemmipara*.

Genus Dactylophrya Collin. Cup-like lorica, filled with the protoplasmic body; with a short stalk; 12–15 arm-like tentacles from anterior surface; salt water. One species.

*D. ros covita* C. (Fig. 371, k). About 40μ long excluding stalk; on the hydrozoan, *Diphasia attenuata*.

Genus Pseudogemma Collin. Attached with a short stalk to larger suctorians; without tentacles; endogenous budding; swarmer with 4 ciliary bands; salt water.

*P. pachystyla* C. (Fig. 373, a). About 30μ long; stalk 3–4μ wide; swarmer 15μ by 9μ; on *Acineta tuberosa*.

Genus Endosphaera Engelmann. Spherical without lorica; without tentacles; budding endogenous; swarmer with 3 equatorial ciliary bands; parasitic in Peritricha; fresh and salt water.

*E. engelmanni* Entz (Fig. 373, b). 15–41μ in diameter; imbedded in the host’s cytoplasm; swarmer 13–19μ in diameter; in *Opisthonia henneguyi* (p. 852), and other peritrichs.

Genus Allantosoma Gassovsky. With neither lorica nor stalk; elongate; one or more tentacles at ends; macronucleus oval or spherical; compact micronucleus; a single contractile vacuole; cytoplasm often filled with small spheroidal bodies; development unknown; in mammalian intestine. Species (Hsiung, 1930).
A. intestinalis G. (Fig. 373, c). 33–60μ by 18–37μ; attached to various ciliates living in the caecum and colon of horse.

A. dicorniger Hsiung (Fig. 373, d). 20–33μ by 10–20μ; unattached; in the colon of horse (Hsiung, 1928).

A. brevicorniger H. (Fig. 373, e). 23–36μ by 7–11μ; attached to various ciliates in the caecum and colon of horse.

Fig. 372. a–d, Anarma multiruga, ×about 230; b, budding individual; c, cross-section; d, with an internal ciliated bud; e, f, Squalorophrya macrostyla, ×about 670; f, cross-section; g, Multifasciculatum elegans, ×about 660 (Goodrich and Jahn).

Genus Anarma Goodrich and Jahn. Radially or somewhat bilaterally symmetrical; without stalk or lorica; attached directly or by a short protoplasmic process to substratum; 1–2 fascicles of capitate tentacles; multiplication by external budding near base or by a single internal ciliated bud; conjugation; ectocommensal on Chrysemys picta bellii (Goodrich and Jahn, 1943).

A. multiruga G. and J. (Fig. 372, a–d). Body cylindrical, 70–150μ by 35–70μ; body surface with 7 or 8 longitudinal folds; pellicle thin; cytoplasm granulated; nucleus ribbon-form; 2–6 contractile vacuoles,
each with a permanent canal and a pore; attached directly or indirectly to the carapace and plastron of the turtle.

Genus **Squalorophrya** Goodrich and Jahn. Elongate; radially symmetrical; lorica, rigid, close-fitting, covered with debris; with a stalk; capitate tentacles at distal end; ectocommensal on *Chrysemys picta bellii*.

*S. macrostyla* G. and J. (Fig. 372, e, f). Cylindrical, with 4 longitudinal grooves; body about 90μ by 40μ; striated stalk, short and thick, about 30μ long; lorica highly viscous with debris; nucleus ovoid to elongate, sometimes Y-shaped; 2 contractile vacuoles, each with a permanent canal and a pore; on *Chrysemys picta bellii*.

Genus **Multifasciculatum** Goodrich and Jahn. Radially or bilaterally symmetrical; stalked; without lorica; pellicle thin; several fascicles of tentacles on distal, lateral and proximal regions of body; ectocommensal on *Chrysemys picta bellii*.

*M. elegans* G. and J. (Fig. 372, g). Body ovoid; 50–90μ by 20–50μ; stalk striated, about 150–270μ long; tentacles in 4 groups; nucleus ovoid; 1–3 contractile vacuoles; attached to the plastron of the turtle.

**Family 6. Discophryidae** Collin

Genus **Discophrya** Lachmann. Elongate; a short stout pedicel with a plate; tentacles evenly distributed on anterior surface or in fascicles; contractile vacuoles, each with a canalicule leading to body surface; mainly fresh water. Several species (Swarzewsky, 1928b).

*D. elongata* (Claparède and L.) (Fig. 373, f). Cylindrical; tentacles on anterior end and in 2 posterior fascicles; stalk striated; about 80μ long; on the shell of *Paldina vivipara* in fresh water.

Genus **Thaumatophrya** Collin. Spherical; long stalk; tentacles distributed, tapering toward distal end; salt water. One species.

*T. trold* (Claparède and Lachmann) (Fig. 373, g). About 75μ in diameter.

Genus **Rhynchophrya** Collin. Oblong; bilaterally symmetrical; a short striated stalk; 1 main long and a few shorter tentacles; 6–10 contractile vacuoles, each with a canalicule leading to outside; fresh water. One species.

*R. palpans* C. (Fig. 373, h). 85μ by 50μ; tentacles retractile, 10–200μ long; stalk 20μ by 10μ; on *Hydrophilus piceus*.

Genus **Choanophrya** Hartog. Spheroidal to oval; stalked; 10–12 tentacles; tubular, expansible at distal end to engulf voluminous food particles; macronucleus oval to spherical; a micronucleus; fresh water. One species.

*C. infundibulifera* H. (Fig. 374, a). 65μ by 60μ; fully extended ten-
Fig. 373. a, *Pseudogemma pachystyla*, $\times 400$ (Collin); b, *Endosphaera engelmanni*, $\times 500$ (Lynch and Noble); c, *Allantosoma intestinalis*, $\times 1050$ (Hsiung); d, *A. dicorniger*, $\times 1300$ (Hsiung); e, *A. brevicorniger*, $\times 1400$ (Hsiung); f, *Discophrya elongata*, $\times 440$ (Collin); g, *Thaumatophrya trold*, $\times 1150$ (Claparède and Lachmann); h, *Rhynchophrya palpans*, $\times 440$ (Collin).

Tentacles $200\mu$ long; on *Cyclops ornatus*. Tentacles and feeding (Farkas, 1924).

Genus *Rhyncheta* Zenker. Protoplasmic body attached directly to an aquatic animal; with a long mobile tentacle bearing a sucker at its end.

*R. cyclopum* Z. (Fig. 374, b, c). About $170\mu$ long; on *Cyclops*. 
Family 7 Ephelotidae Sand

Genus Ephelota Wright. Without lorica; stalk stout, often striated; suctorial and prehensile tentacles distributed; macronucleus usually elongate, curved; on hydroids, bryozoans, algae, etc.; salt water. Numerous species.

_E. gemmipara_ Hertwig (Fig. 374, _d_). About 250\(\mu\) by 220\(\mu\); stalk up to 1.5 mm. long; on hydroids, bryozoans, etc.

_E. coronata_ Kent (Fig. 374, _e_). Flattened; 90–200\(\mu\) long; stalk longitudinally striated (Kent); on hydroids, bryozoans, algae, etc.

_E. plana_ Wailes (Fig. 374, _f_). 150–320\(\mu\) by 100–150\(\mu\); stalk 100\(\mu\)–1 mm. long; on bryozoans; Vancouver.
Genus *Podocyathus* Kent. It differs from *Ephelota* in having a conspicuous lorica; salt water. One species.

*P. diadema* K. (Fig. 374, g). Lorica about 42μ long; on bryozoans, hydrozoans, etc.

**References**


—— (1912) II. Ibid., 51:1.


—— (1930) A monograph on the Protozoa of the large intestine of the horse. Ibid., 4:356.


Root, F. M.: (1914) Reproduction and reactions to food in the suctorian, Podophrya collini n. sp. Ibid., 35:164.


Rudzinska, Maria A.: (1951) The effect of overfeeding and starvation on the life span and reproduction of Tokophrya infusionum, etc. J. Gerontol., 6, Suppl. 3:144.


—— (1928a) II. Ibid., 62:41.

—— (1928b) III. Ibid., 63:1.

—— (1928c) IV. Ibid., 63:362.

Chapter 45

Collection, Cultivation, and Observation of Protozoa

Collection

IN THE foregoing chapters it has been pointed out that various species of Protozoa have characteristic habitats and that many of free-living forms are widely distributed in bodies of water: fresh, brackish, and salt; while the parasitic forms are confined to specific host animals. Of free-living Protozoa many species may occur in large numbers within a small area under favorable conditions, but the majority are present in comparatively small numbers. If one who has become acquainted with the representative forms, intends to make collection, it is well to carry a compound microscope in order to avoid bringing back numerous jars containing much water, but few organisms. Submerged plants, decaying leaves, surface scum, ooze, etc., should be examined under the microscope. When desired forms are found, they should be collected together with a quantity of water in which they occur.

When the material is brought into the laboratory, it is often necessary to concentrate the organisms in a relatively small volume of water. For this purpose the water may partly be filtered rapidly through a fine milling cloth and the residue quickly poured back into a suitable container before filtration is completed. The container should be placed in a cool moderately lighted room to allow the organisms to become established in the new environment. Stigma-bearing Phytomastigina will then be collected in a few hours on the side of the container, facing the strongest light, and the members of Sarcodina will be found among the debris on the bottom. Many forms will not only live long, but also multiply in such a container.

For obtaining large freshwater amoebae, fill several finger bowls with the collected material and water, and place one or two rice grains to each. After a few days, examine the bottom surface of the bowls under a binocular dissecting microscope. If amoebae were included in the collection, they will be found particularly around the rice grains. Pipette them off and begin separate cultures (p. 881).

In order to collect parasitic Protozoa, one must, of course, find the host organisms that harbor them. Various species of tadpoles, frogs, cockroaches, termites, etc., which are of common occurrence or easily obtained and which are hosts to numerous species of Protozoa, are useful material for class work.
Intestinal Protozoa of man are usually studied in the faeces of an infected person. Natural movement should be collected. Do not use oily purgatives in obtaining faecal specimens, as they make the microscopical examination difficult by the presence of numerous oil droplets. The receptacle must be thoroughly cleaned and dry, and provided with a cover. Urine or water must be excluded completely. The faeces must be examined as soon as possible, since the active trophozoites degenerate quickly once leaving the human intestine. If dysenteric or diarrhoeic stools are to be examined, they must not be older than one hour or two. In case this is not possible, wrap the container with woolen cloth while transporting, the organisms may live for several hours. Care must however be exercised during the microscopical examination, as there will be present unavoidably a large number of degenerating forms. If the stool is formed and normal, it would contain usually encysted forms and no trophozoites if the host is infected by a protozoan, unless mucus, puss, or blood is present in it. Examination of such faeces can be delayed, as the cysts are quite resistant (p. 450).

Cultivation

For extensive study or for class work, a large number of certain species of Protozoa are frequently needed. Detection and diagnosis of human Protozoa are often more satisfactorily made by culture method than by microscopical examination of the collected material. Success in culturing Protozoa depends upon several factors. First an abundant supply of proper food material must be made available. For example, several species of Paramecium live almost exclusively on bacterial organisms, while Didinium and allied ciliates depend upon Paramecium and other ciliates as sources of food supply. For cultivating chromatophore-bearing forms successfully, good light and proper kinds and amount of inorganic substances are necessary. In the second place, the temperature and chemical constituents of the culture medium must be adjusted to suit individual species. As a rule, lower temperatures seem to be much more favorable for culture than higher temperatures, although this is naturally not the case with those parasitic in homoiothermal animals. Furthermore, proper hydrogen ion concentration of the culture must be maintained. In the third place, both Protozoa and Metazoa which prey upon the forms under cultivation must be excluded from the culture. For instance, it is necessary to remove *Didinium nasutum* in order to obtain a rich culture of Paramecium. For successful culture of *Amoeba proteus*, Aeolosoma, Daphnia, Cyclops, etc., must be excluded from the culture.
Mixed cultures of many free-living Protozoa are easily maintained by adding from time to time a small amount of ripe hay-infusion or dried lettuce powder to the collected water mentioned before. Chilomonas, Peranema, Bodo, Arcella, Amoeba, Paramecium, Colpoda, Stylonychia, Euplotes, etc., often multiply in such cultures. To obtain a large number of a single species, individuals are taken out under a binocular dissecting microscope by means of a finely drawn-out pipette and transferred to a suitable culture medium. Such a culture is called a mass or stock culture. If a culture is started with a single individual, the resulting population makes up a clone or a pure line.

Aside from the cultures of blood-inhabiting Protozoa and of some 100 free-living forms, the protozoan cultures are by no means "pure" cultures in the bacteriological sense, even if only one species of Protozoa is present, since bacteria and other microorganisms are invariably abundantly present in them.

A. Free-living Protozoa

To deal with all the culture media employed by numerous workers for various free-living Protozoa is beyond the scope of the present work. Here only a few examples will be given. For further information, the reader is referred to Bělař (1928), Needham et al. (1937), etc.

Chromatophore-bearing flagellates.—There are a number of culture fluids. Two examples:

(a) Peptone or tryptone 2.0 gm.
    KH₂PO₄ 0.25 gm.
    MgSO₄ 0.25 gm.
    KCl 0.25 gm.
    FeCl₃ trace
    Sodium acetate 2.0 gm.
    Pyrex distilled water 1000 cc.

(b) Peptone or tryptone 2.5 gm.
    KNO₃ 0.5 gm.
    KH₂PO₄ 0.5 gm.
    MgSO₄ 0.1 gm.
    NaCl 0.1 gm.
    Sodium acetate 2.5 gm.
    Dextrose 2.0 gm.
    Glass distilled water 1000 cc.

Peranema, Chilomonas, Astacia and other colorless flagellates.—A number of culture fluids have been advocated. A simple yet satis-
factory one is as follows: Fill a finger bowl with about 150 cc. of glass distilled water and place 4 rice grains on the bottom. Let the dish stand for a few days, and then introduce with a pipette a number of desired flagellates from a mass culture into it. Cover the bowl and keep it at about 20°C.

Mast (1939) used the following media for Chilomonas paramecium.

(a) Glucose-peptone solution:

- Peptone: 8 gm.
- Glucose: 2 gm.
- Water: 1000 cc.

(b) Acetate-ammonium solution:

- Sodium acetate: 1.5 gm.
- Ammonium chloride: 0.46 gm.
- Ammonium sulphate: 0.1 gm.
- Dipotassium hydrogen phosphate: 0.2 gm.
- Magnesium chloride: 0.01 gm.
- Calcium chloride: 0.012 gm.
- Water: 1000 cc.

Amoeba proteus and other freshwater amoebae.—Fill a finger bowl with 200 cc. of glass distilled water, and place 4 rice grains. After a few days seed with amoebae (p. 879), add about 5 cc. of Chilomonas culture, and cover the bowl with a glass cover. In about two weeks a ring of amoebae will be found around each rice grain, and if Chilomonas do not overmultiply, the amoebae will be found abundantly in another two weeks. If properly maintained, subcultures may be made every 4–6 weeks. Chalkley (1930) advocates substitution of the plain water with a salt solution which is composed of

- NaCl: 0.1 gm.
- KCl: 0.004 gm.
- CaCl₂: 0.006 gm.
- Glass distilled water: 1000 cc.

If the culture water becomes turbid, make subcultures or pour off the water and fill with fresh distilled water or the solution. Culture should be kept at 18–22°C.

Hahnert (1932) used the following culture solution:

- KCl: 0.004 gm.
- CaCl₂: 0.004 gm.
- Ca₃(PO₄)₂: 0.002 gm.
- Mg₃(PO₄)₂: 0.002 gm.
Pelomyxa carolinensis.—These amoebae grow well in a finger bowl with 150 cc. of redistilled water to which large numbers of Paramecium are added daily. Pace and Belda (1944) advocate the following solution instead of distilled water:

- K₂HPO₄ 0.08 gm.
- KH₂PO₄ 0.08 gm.
- CaCl₂ 0.104 gm.
- Mg₃(PO₄)₂·4H₂O 0.002 gm.
- Pyrex water 1000 cc.

Small mono- or di-phasic amoebae.—Musgrave and Clegg's medium, modified by Walker, is as follows:

- Agar 2.5 gm.
- NaCl 0.05 gm.
- Liebig's beef-extract 0.05 gm.
- Normal NaOH 2 cc.
- Distilled water 100 cc.

Arcella and other Testacea.—The testaceans commonly multiply in a mixed culture for several weeks after the collection was made. Hegner's method for Arcella: Pond water with weeds is shaken up violently and filtered through eight thicknesses of cheese cloth, which prevents the passage of coarse particles. The filtrate is distributed among Petri dishes, and when suspended particles have settled down to the bottom, specimens of Arcella are introduced. This will serve also for Diffugia and other testaceans. Hay or rice infusion is also a good culture medium for these organisms.

Actinophrys and Actinosphaerium.—Bělař cultivated these heliozoans successfully in Knop's solution:

- Magnesium sulphate 0.25 gm.
- Calcium nitrate 1 gm.
- Potassium phosphate 0.25 gm.
- Potassium chloride 0.12 gm.
- Iron chloride trace
- Distilled water 1000 cc.

Freshwater ciliates.—They are easily cultivated in a weak infusion of hay, bread, cracker, lettuce leaf, etc. The battery jars containing the infusions should be left standing uncovered for a few days to allow a rich bacterial growth in them. Seed them with material such
as submerged leaves or surface scum containing the ciliates. If desired, culture may be started with a single individual in a watch glass. Collection, cultivation and sterilization of Paramecium (Wichertman, 1949).

Pure culture

Many free-living flagellates and certain ciliates have in recent years been successfully cultured free from any other associated organisms. The protozoan to be cultivated must be freed from other Protozoa and bacteria. For this, washing, dilution, migration and bactericidal agents have been used. For information, the reader is referred to Glaser and Coria (1930), Claff (1940), Taylor and Van Wagentock (1941), Kidder (1941), etc.

Free-living Phytomastigina.—Many media are known. See Pringsheim (1926, 1937, 1946), Hall (1937, 1941), Hutner and Provasoli (1951), etc.

Tetrahymena and allied forms.—Kidder, Dewey and Parks use a basal medium as quoted below:

<table>
<thead>
<tr>
<th>Substance</th>
<th>γ per ml</th>
</tr>
</thead>
<tbody>
<tr>
<td>DL-alanine</td>
<td>110</td>
</tr>
<tr>
<td>L-arginine</td>
<td>206</td>
</tr>
<tr>
<td>L-aspartic acid</td>
<td>122</td>
</tr>
<tr>
<td>Glycine</td>
<td>10</td>
</tr>
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<td>L-glutamic acid</td>
<td>233</td>
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<tr>
<td>L-histidine</td>
<td>87</td>
</tr>
<tr>
<td>DL-isoleucine</td>
<td>276</td>
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<tr>
<td>L-leucine</td>
<td>344</td>
</tr>
<tr>
<td>L-lysine</td>
<td>272</td>
</tr>
<tr>
<td>DL-methionine</td>
<td>248</td>
</tr>
<tr>
<td>L-phenylalanine</td>
<td>160</td>
</tr>
<tr>
<td>L-proline</td>
<td>250</td>
</tr>
<tr>
<td>DL-serine</td>
<td>394</td>
</tr>
<tr>
<td>DL-threonine</td>
<td>326</td>
</tr>
<tr>
<td>L-tryptophane</td>
<td>72</td>
</tr>
<tr>
<td>DL-valine</td>
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<tr>
<td>Ca pantothenate</td>
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</tr>
<tr>
<td>Nicotinamide</td>
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</tr>
<tr>
<td>Pyridoxine HCl</td>
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</tr>
<tr>
<td>Pyridoxal HCl</td>
<td>0.10</td>
</tr>
<tr>
<td>Pyridoxamine HCl</td>
<td>0.10</td>
</tr>
<tr>
<td>Riboflavin</td>
<td>0.10</td>
</tr>
<tr>
<td>Pteroylglutamic acid</td>
<td>0.01</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Substance</th>
<th>γ per ml</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thiamine HCl</td>
<td>1.00</td>
</tr>
<tr>
<td>Biotin (free acid)</td>
<td>0.0005</td>
</tr>
<tr>
<td>Choline Cl</td>
<td>1.00</td>
</tr>
<tr>
<td>MgSO₄·7H₂O</td>
<td>100</td>
</tr>
<tr>
<td>Fe(NH₄)₂(SO₄)₂·6H₂O</td>
<td>25</td>
</tr>
<tr>
<td>MnCl₂·4H₂O</td>
<td>0.5</td>
</tr>
<tr>
<td>ZnCl₂</td>
<td>0.05</td>
</tr>
<tr>
<td>CaCl₂·2H₂O</td>
<td>50</td>
</tr>
<tr>
<td>CuCl₂·2H₂O</td>
<td>5</td>
</tr>
<tr>
<td>FeCl₃·6H₂O</td>
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<tr>
<td>K₂HPO₄</td>
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<tr>
<td>KH₂PO₄</td>
<td>1,000</td>
</tr>
<tr>
<td>Guanylic acid</td>
<td>30</td>
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<tr>
<td>Adenylic acid</td>
<td>20</td>
</tr>
<tr>
<td>Cytidyllic acid</td>
<td>25</td>
</tr>
<tr>
<td>Uracil</td>
<td>10</td>
</tr>
<tr>
<td>Dextrose</td>
<td>2,500</td>
</tr>
<tr>
<td>Na acetate</td>
<td>1,000</td>
</tr>
<tr>
<td>Tween 85</td>
<td>700</td>
</tr>
<tr>
<td>Protogen</td>
<td>1 unit</td>
</tr>
</tbody>
</table>

B. Parasitic Protozoa

Intestinal flagellates of man.—There are numerous media which have been used successfully by several investigators.
(a) Ovo-mucoid medium (Hogue, 1921). White of two eggs are broken in a sterile flask with beads. Add 200 cc. of 0.7 % NaCl solution and cook the whole for 30 minutes over a boiling water bath, shaking the mixture constantly. Filter through a coarse cheese cloth and through cotton-wool with the aid of a suction pump. Put 6 cc. of the filtrate in each test tube. Autoclave the tubes for 20 minutes under 15 pounds pressure. After cooling, a small amount of fresh faecal material containing the flagellates is introduced into the tubes. Incubate at 37°C.

(b) Sodium chloride sheep serum water (Hogue, 1922). Composed of 100 cc. of sterile 0.95% NaCl and 10–15 cc. of sterile sheep serum water (dilution 1:3). 15 cc. to each tube. Trichomonas hominis, T. tenax, and Retortamonas intestinalis grow well. Trichomonas vaginalis.—Johnson and Trussell (1943) reported the following mixture the most suitable medium:

| Bacto-peptone | 32 gm. |
| Bacto-agar | 1.6 gm. |
| Cysteine HCl | 2.4 gm. |
| Maltose | 1.6 gm. |
| Difco liver infusion | 320 cc. |
| Ringer's solution | 960 cc. |
| NaOH(N/1) | 11–13 cc. |

Heat the mixture in a water bath to melt the agar; filter through a coarse paper; add 0.7 cc. of 0.5 per cent aqueous methylene blue; adjust pH to 5.8–6.0 with N/1 HCl or NaOH; tube 8 cc.; autoclave. After cooling, add aseptically 2 cc. of sterile (filtered) human serum. Incubate at least four days; store at room temperature for two to three weeks or as long as an amber “anaerobic” zone is apparent.

Termite flagellates.—Trager's (1934) media are as follows:

<table>
<thead>
<tr>
<th></th>
<th>Solution A gm. per liter water</th>
<th>Solution U gm. per liter water</th>
</tr>
</thead>
<tbody>
<tr>
<td>NaCl</td>
<td>1.169</td>
<td>2.164</td>
</tr>
<tr>
<td>NaHCO₃</td>
<td>0.840</td>
<td>0.773</td>
</tr>
<tr>
<td>Na₂C₆H₅O₇·2H₂O (citrate)</td>
<td>2.943</td>
<td>1.509</td>
</tr>
<tr>
<td>NaH₂PO₄·H₂O</td>
<td>0.690</td>
<td>0</td>
</tr>
<tr>
<td>KCl</td>
<td>0.745</td>
<td>0</td>
</tr>
<tr>
<td>KH₂PO₄</td>
<td>0</td>
<td>1.784</td>
</tr>
<tr>
<td>CaCl₂</td>
<td>0.111</td>
<td>0.083</td>
</tr>
<tr>
<td>MgSO₄</td>
<td>0</td>
<td>0.048</td>
</tr>
</tbody>
</table>
In solution A, *Trichomonas* sp. and *Tricercomitus termopsidis* were cultivated. For *Trichomonas termopsidis*, a small amount of Loeffler's blood serum and cellulose were added. All three flagellates were cultured for over three years. In solution U to which 0.01 per cent blood serum, cellulose and charcoal, were added, *Trichonympha sphaerica* (from *Termopsis angusticollis*) grew well and multiplied up to two weeks, although *T. campanula* and *T. collaris* failed to do so. The culture in a test tube was inoculated with the entire hindgut of a termite and kept at room temperature.

*Lophomonas blattarum* and *L. striata.*—A mixture of one sterile egg-white and 100 cc. of sterile Ringer's solution, to which a small amount of yeast cake is added, is an excellent culture medium. Incubation at room temperature; subcultures every 4–6 days.

*Trypanosoma and Leishmania.*—Novy, MacNeal and Nicolle (NNN) medium: 14 gm. of agar and 6 gm. of NaCl are dissolved by heating in 900 cc. of distilled water. When the mixture cools to about 50°C., 50–100 cc. of sterile defibrinated rabbit blood is gently added and carefully mixed so as to prevent the formation of bubbles. The blood agar is now distributed among sterile test tubes to the height of about 3 cm., and the tubes are left slanted until the medium becomes solid. The tubes are then incubated at 37°C. for 24 hours to determine sterility and further to hasten the formation of condensation water (pH 7.6). Sterile blood or splenic puncture containing *Trypanosoma cruzi* or Leishmania is introduced by a sterile pipette to the condensation water in which organisms multiply. Incubation at 37°C. for trypanosomes and at 20–24°C. for Leishmania.

For cultivating *T. gambiense* and *T. rhodesiense*, Tobie, von Brand and Mehlman (1950) used the following medium:

(a) Base. 1.5 gm. Bacto-beef, 2.5 gm. Bacto-peptone, 4 gm. sodium chloride and 7.5 gm. Bacto-agar, are dissolved in 500 cc. distilled water. After adjusting pH to 7.2–7.4 with NaOH, autoclave at 15 lbs pressure for 20 minutes. Cool this to about 45°C., then add whole rabbit blood which had been inactivated at 56°C. for 30 minutes, in the proportion of 25 cc. blood to 75 cc. base, using 0.5 per cent sterile sodium citrate to prevent the coagulation. This base is placed in test tubes (5 cc. each and slanted) or in flasks (25 cc.), and allowed to solidify.

(b) Liquid phase. Sterile Locke's solution. This is added in amounts of 2 cc. (to test tubes) or 10–15 cc. (to flasks), and cotton plugs are applied. The trypanosomes are said to grow well and to reach the peak population in 10–14 days.

*Entamoeba barreti.*—Barret and Smith (1924) used a mixture of
9 parts of 0.5% NaCl and 1 part of human blood serum. Incubation at 10–15°C.

*E. invadens.*—Ratcliffe and Geiman (1938) used a mixture of gastric mucin 0.3 gm., “ground alum” salt 0.5 gm., and distilled water 100 cc. About 2 mg. of sterile rice starch is added to each culture tube at the time of inoculation. Culture at 20–30° C. and subculture every 7 days.

*E. histolytica and other amoebae of man.*—The first successful culture was made by Boeck and Drbohlav (1925) who used the following media.

(a) Locke-egg-serum (LES) medium. The contents of 4 eggs (washed and dipped in alcohol) are mixed with, and broken in, 50 cc. of Locke’s solution in a sterile flask with beads. The solution is made up as follows:

<table>
<thead>
<tr>
<th>Component</th>
<th>Concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>NaCl</td>
<td>9 gm.</td>
</tr>
<tr>
<td>CaCl₂</td>
<td>0.2 gm.</td>
</tr>
<tr>
<td>KCl</td>
<td>0.4 gm.</td>
</tr>
<tr>
<td>NaHCO₃</td>
<td>0.2 gm.</td>
</tr>
<tr>
<td>Glucose</td>
<td>2.5 gm.</td>
</tr>
<tr>
<td>Distilled water</td>
<td>1000 cc.</td>
</tr>
</tbody>
</table>

The emulsion is now tubed so that when coagulated by heat, there is 1–1.5 inches of slant. These tubes are now slanted and heated at 70°C, until the medium becomes solidified. They are then autoclaved for 20 minutes at 15 pounds pressure (temperature must be raised and lowered slowly). After cooling the slant is covered with a mixture of 8 parts of sterile Locke’s solution and 1 part of sterile inactivated human blood serum. The tubes are next incubated to determine sterility. The culture tubes are inoculated with a small amount of faecal matter containing active trophozoites. Incubation at 37°C. Yorke and Adams (1926) obtained rich cultures by inoculating this medium with washed and concentrated cysts of *E. histolytica* in 24 hours.

(b) Locke-egg-albumin (LEA) medium. The serum in LES medium is replaced by 1% solution of crystallized egg albumin in Locke’s solution which has been sterilized by passage through a Berkefeld filter.

Dobell and Laidlaw (1926) used Ringer’s solution instead of Locke’s.

(c) Ringer-egg-serum (RES) or Ringer-egg-albumin (REA) medium. Solid medium is the same as that of (a) or (b), but made up in Ringer’s solution which is composed of
NaCl   9 gm.
KCl   0.2 gm.
CaCl$_2$  0.2 gm.
Distilled water  1000 cc.

The covering liquid is serum-Ringer or egg-albumin. The latter is prepared by breaking one egg white in 250 cc. of Ringer's solution which is passed through a Seitz filter. Before inoculating with amoeboae, a small amount of sterile solid rice-starch (dry-heated at 180°C. for 1 hour) is added to the culture tube.

(d) Horse-serum-serum (HSS) or Horse-serum-egg-albumin (HSA) medium. Whole horse-serum, sterilized by filtration, is tubed and slanted at 80°C. for about 60-70 minutes (do not heat longer). When the slants have cooled, they are covered with diluted serum or egg-albumin given for (c). The tubes are incubated for sterility and sterile rice-starch is added immediately before inoculation. Frye and Meleny (1939) substituted the liquid portion of this medium by 0.5% solution of Lily liver extract No. 343 in 0.85% NaCl.

(e) Liver-agar-serum (LAS) medium. Cleveland and Sanders (1930) used the following medium:

Liver infusion agar 30 gm.
(Difco dehydrated)
Glass distilled water 1000 cc.

The medium is tubed, autoclaved, and slanted. The slants are covered with a 1:6 dilution of sterile fresh horse serum in 0.85% NaCl solution. A 5 mm. loop of sterile rice flour or powdered unpolished rice is added to each tube. In making subculture, remove 2 or 3 drops of the rice flour debris from the bottom with a sterile pipette.

(f) Egg-yolk-saline medium (Balamuth and Sandza, 1944). Two eggs are hard-boiled. Upon cooling, the egg white is discarded and the yolks are crumbled in a beaker containing 125 ml of 0.8 per cent sodium chloride solution. The mixture is boiled for 10 minutes, and after replacement of evaporated water the infusion is filtered by suction pump and restored to 125 ml. The filtrate is autoclaved 20 minutes at 15 pounds pressure. Upon cooling, a slight precipitation of yolk settles, and is removed by simple filtration, after which 125 ml of N/15 phosphate buffer (pH 7.5) is added, making the total salt concentration N/30 phosphate solution in 0.4 per cent sodium chloride. This final mixture is tubed in 5 ml amounts, autoclaved as before, and then is stored under refrigeration until use. Before introducing amoebae a loop of sterile rice starch is added to each tube.
To inhibit bacterial growth in cultures of Entamoeba, various antibiotics have been tried. For example, Spingarn and Edelman (1947) found that when streptomycin was added in the amount of 1000–3000 units per cc. to culture of *E. histolytica*, the survival of the amoebae in culture was prolonged from an average of 8 days to 33.7 days, which effect was apparently due to the inhibition of bacteria.

Encystment of *Entamoeba histolytica* is usually brought about by first cultivating the organisms in starch-free media and then by transferring them into media with starch. Balamuth (1951) recommends a diphase medium of the following composition: 2 gm. of Wilson liver concentrate powder is brought to boiling in 80 ml. distilled water and filtered. Then 6.4 ml. of 0.25 molar Na$_2$PO$_4$, 12 H$_2$O and 7.6 ml. of 1.0 molar potassium phosphate buffer (in the ratio of 4.7 parts K$_2$HPO$_4$ to 0.3 part KH$_2$PO$_4$) are added. By adding distilled water in a volumetric flask bring the mixture to 100 ml. Transfer it to a beaker and add 3 gm. Bacto-agar. Heat gently until agar dissolves; then autoclave for 20 minutes at 15 lbs pressure. The pH should be about 7.2. The overlay is prepared by mixing double-strength egg yolk and normal horse serum (10:1) and rice starch is added last.

*Plasmodium.*—Bass and John's (1912) culture is as follows: 10 cc. of defibrinated human blood containing Plasmodium and 0.1 cc. of 50% sterile dextrose solution are mixed in test tubes and incubated at 37–39°C. In the culture, the organisms develop in the upper layer of erythrocytes. Since that time a number of investigators have undertaken cultivation of different species of Plasmodium. For information the reader is referred to Geiman, Anfinsen *et al.* (1946) and Trager (1950).

*Balantidium coli.*—Barret and Yarbrough (1921) first cultivated this ciliate in a medium consisting of 16 parts of 0.5% NaCl and 1 part of inactivated human blood serum. The medium is tubed. Inoculation of a small amount of the faecal matter containing the trophozoites is made into the bottom of the tubes. Incubation at 37°C. Maximum development is reached in 48–72 hours. Subcultures are made every second day. Rees used a mixture of 16 parts of Ringer's solution and 1 part of Loeffler's dehydrated blood serum.

Atehley (1935) employed a medium composed of 4 parts of Ringer's solution and 1 part of faeces, which is filtered after 24 hours, centrifuged and sterilized by passage through a Seitz filter. Nelson (1940) also used 1 part of caecal contents of pig in 9 parts of Ringer's solution, which mixture is passed through a sieve and then filtered through a thick absorbent cotton. *Balantidium* which shows posi-
tive geotropism, is freed of faecal debris by passage downward through cotton in V-tube. The ciliates are introduced into the culture tubes. Incubation at 37°C. Subcultures are made every 7–22 days. Nelson found that autoclaved medium is unsuitable until a living bacterial population has been established. Balantidium can also be cultivated in the media given for the intestinal amoebae.

Microscopical examination

Protozoa should be studied as far as possible in life. Permanent preparations while indispensable in revealing many intracellular structures, cannot replace fresh preparations. The microscopic slides of standard size, 3" by 1", should be of white glass and preferably thin. The so-called No. 1 slides measure about 0.75 mm. in thickness. For darkfield illumination thin slides are essential. No. 1 coverglasses should be used for both fresh and permanent preparations. They are about 130–170µ thick. The most convenient size of the coverglass is about 7/8 square inch which many prefer to circular ones.

The slides and coverglasses must be thoroughly cleaned before being used. Immerse them in concentrated mineral acids (nitric acid is best fitted) for 10 minutes. Pour off the acid, wash the slides and coverglasses for about 10 minutes in running water, rinse in distilled water, and keep them in 95% alcohol. When needed they are dried one by one with clean cheese cloth. Handle slides and covers with a pair of forceps. If thumb and fingers are used, hold them edgewise.

A. Fresh preparations

In making fresh preparations with large Protozoa care must be exercised to avoid pressure of the coverglass on the organisms as this will cause deformities. If small bits of detritus or debris are included in the preparation, the coverglass will be supported by them and the organisms will not be subjected to any pressure. Although ordinary slides are used most frequently, it is sometimes advisable to use a depression slide especially for prolonged observation. To make a preparation with this slide, a small drop of water containing specimens is placed in the center of a coverglass, and is covered by a small circular coverglass (about 1 cm. in diameter), which in turn is covered by a depression slide with a thin coat of vaseline along the edge of the depression, so as to make an air-tight compartment. In turning over the whole, care must be taken to prevent the smaller circular cover from touching any part of the slide, as this would cause the water to run down into the depression. Nemeczki (1926) seems to have been the first one who used the second coverglass for this prepa-
rations. If the Protozoa to be examined are large and observation can be made under a low power objective, the small cover-glass should be omitted.

As far as possible examine fresh preparations with low power objectives. The lower the magnification, the brighter and the larger the field. The microscopical objects can quickly and easily be measured, if an ocular micrometer division has been calculated in combination with different objectives.

The free-living ciliates swim about so actively as to make their observation difficult. However, an actively swimming ciliate will sooner or later come to stop upon coming in contact with various debris, air bubbles or margin of the coverglass to allow a study of its structure. Various reagents recommended for retardation of swimming movements of ciliates, bring about deformities in the organisms and therefore, must not be used; but a drop of saturated solution of methyl cellulose may be added to a ciliate preparation to retard the active movement of the organism without causing any visible abnormality (Marsland, 1943).

For observation of cilia, flagella, extruded polar filament of Microsporidia, etc., the so-called changeable condenser is useful, since it gives both bright and dark fields under dry objectives. The ordinary dark field condenser is used almost exclusively in conjunction with an oil immersion objective and therefore for very active organisms a great deal of time is often lost before satisfactory observation is made. The phase microscope is highly useful in studying various intracellular structures in life.

When treated with highly diluted solutions of certain dyes, living Protozoa exhibit some of their organellae or inclusions stained without apparent injury to the organisms. These vital stains are usually prepared in absolute alcohol solutions. A small amount is uniformly applied to the slide and allowed to dry, before water containing Protozoa is placed on it. Congo red (1:1,000) is used as an indicator, as its red color of the salt changes blue in weak acids. Janus Green B (1:10,000–20,000) stains chondriosomes. Methylene blue (1:10,000 or more) stains cytoplasmic granules, nucleus, cytoplasmic processes, etc., Neutral red (1:3,000–30,000) is an indicator: yellowish red (alkaline), cherry red (weak acid), and blue (strong acid). It also stains nucleus slightly. Golgi bodies are studied in it, though its specificity for this structure is not clear.

Parasitic Protozoa should be studied in the tissue or body fluids in which they occur. When they are too small in amount to make a suitable preparation, one of the following solutions may be used.
Physiological salt solution. Widely used concentrations of NaCl solutions are 0.5–0.7% for cold-blooded animals and 0.8–0.9% for warm-blooded animals.

Ringer’s solution. The one Dobell advocated has been given already (p. 887). Another frequently used solution consists of

<table>
<thead>
<tr>
<th>Component</th>
<th>Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td>NaCl</td>
<td>0.8 gm.</td>
</tr>
<tr>
<td>KCl</td>
<td>0.02 gm.</td>
</tr>
<tr>
<td>CaCl₂</td>
<td>0.02 gm.</td>
</tr>
<tr>
<td>(NaHCO₃)</td>
<td>0.02 gm.</td>
</tr>
<tr>
<td>Glass distilled water</td>
<td>100 cc.</td>
</tr>
</tbody>
</table>

For demonstrating organellae, the following reagents which kill the Protozoa upon application, may be used on living Protozoa.

Lugol’s solution. This is made up of potassium iodide 1.5 gm., water 25 cc., and iodine 1 gm. The solution deteriorates easily. Flagella and cilia stain clearly. Glycogen bodies stain ordinarily reddish brown. Cysts of intestinal Protozoa are more easily studied in Lugol’s solution.

Sudan III and IV. 2% absolute alcohol solution diluted before use with the same amount of 45% alcohol. Neutral fats are stained red.

Methyl green. 1% solution in 1% acetic acid solution makes an excellent nuclear stain.

Nigrosin. 10% solution if used in smears and air-dried makes the pellicular patterns of flagellates and ciliates stand out clearly.

In the case of faecal examination if the stool is dysenteric, a small portion is placed by a tooth-pick or platinum loop on a slide and covered with a cover glass. Before placing the cover, all large particles must be removed quickly so that the smear will be uniformly thin. Smears of diarrhoeic stools can be made in a similar way. But if the faecal material is formed or semifomed, a small drop of warm (37°C.) 0.85% NaCl solution is first placed on the slide, and a small portion of the faeces, particularly mucus, pus or blood, is emulsified in it. The whole is covered by a coverglass. The faecal smear should not be too thick or too thin for a satisfactory observation. If the smear is too thick, it will be impossible to distinguish objects clearly, and on the other hand, if it is too thin, there will be much time lost in observing widely scattered Protozoa. The optimum thickness of the smear is one through which the print of this page can be read.

The success in faecal examination for intestinal Protozoa depends almost entirely on continued practice, since the faecal matter contains myriads of objects which may resemble Protozoa (Fig. 375,
Aside from certain coprozoic Protozoa (p. 24) which appear in old faeces, Blastocystis hominis (Fig. 375, c–f) occur in almost all faeces. This organism which is considered to be a fungus and harmless to its host, is usually spherical and measures about 5–25μ in diameter. Within a very thin membrane, there is a narrow peripheral cytoplasmic layer in which 1 or 2 nuclei and several refractile granules are present. The cytoplasmic ring encloses a large homogeneous body which is somewhat eosinophile, but not iodinophile. In some the cytoplasm may be more abundant and the inclusion body smaller. Dividing forms appear peanut-shaped. Blastocystis (Grasse, 1926; Reyer, 1939).

Fig. 375. a, Sphaerita in a stained trophozoite of Entamoeba coli; b, Nucleophaga in a stained trophozoite of Iodamoeba bütschlii; c, d, Blastocystis hominis (in an unstained smear); e, f, stained Blastocystis hominis; g, an epithelial cell from a faecal smear; h, a polymorphonuclear leucocyte with three ingested erythrocytes. All ×1150 (Kudo).

In a number of parasitic Protozoa, there occur foreign organisms which may be mistaken for food inclusions or chromatin. They are vegetable organisms which were named by Dangeard as Sphaerita and Nucleophaga (Fig. 375, a, b). The former occurs in the cytoplasm and the latter in the nucleus of the host protozoan. These parasites are spherical and about 0.5–1μ in diameter; they are found most frequently in spherical masses composed of varying numbers of individuals. Nucleophaga appears to destroy the host nucleus. Degenerating epithelial cells or leucocytes (Fig. 375, g, h) may simulate parasitic amoebae. Fishes and birds are often infected by Coccidia and when they are consumed as food, the oocysts pass the alimentary canal unchanged and appear in the stools. Sphaerita (Chatton and Brodsky, 1909; Mattes, 1924; Becker, 1926;
The cysts of intestinal Protozoa are, as a rule, distributed throughout the formed faeces and difficult to detect in small portions of the voided specimens. Flecks of mucus in the fluid stool obtained by use of a saline purge may contain more numerous cysts than naturally passed one. Several methods for concentrating cysts for microscopical examination are known. The simplest one is to emulsify thoroughly a small mass of faeces about the size of a lump sugar in a dish by adding a small amount of once-boiled tap water. Add to it about 500 cc. of water and pour the whole emulsion into a glass cylinder, and let it stand for about 15 minutes. Remove the scum floating on the surface and draw off the turbid fluid into another cylinder, leaving the sediment and a little fluid just above it untouched. The majority of cysts are suspended in the drawn-off portion of the emulsion. Centrifuge the fluid, pour off the supernatant fluid and add water. Centrifuge again. Repeat this three times until the supernatant fluid becomes clear. The sediment will be found to contain more numerous cysts than small sample specimens. Bijlmer (1948) finds the following method the most satisfactory. Suspend a fleck of faeces about the size of a pea in a dish with some 33 per cent ZnSO₄. If much debris appear on the surface, filter through a layer of cheese-cloth. The fluid is decanted into a centrifuge tube, and some more ZnSO₄ solution is added to half a centimeter from the top. After centrifuging for 2 minutes, lift a loopful of material from the surface and place on a slide.

B. Permanent preparations

Permanent preparations are employed, as was stated before, to supplement, and not to supplant, fresh preparations. Smear preparations are more frequently studied, while section preparations are indispensable in extensive studies of Protozoa. Various fixatives and stains produce different results, care must be exercised in making and evaluating permanent preparations. Diversity of stained objects (Wenrich, 1941).

a. Smear preparations

Smears are made either on coverglasses or slides. However, coverglass-smears are more properly fixed and require smaller amount of reagents than slide-smears. Greater care must be exercised in handling coverglasses, as they are easily broken. Large free-living Protozoa do not frequently adhere to the glass, since there is not
enough albuminous substance in the culture fluid. If a small drop of fresh egg-white emulsified in sterile distilled water is smeared on the coverglass very thinly with the tip of a clean finger, before mounting material for smear, more specimens will adhere to and remain on the coverglass upon the completion of the preparation. Let the smear lie horizontally for 5–10 minutes or longer.

ParaCTic Protozoa live in media rich in albuminous substances, and therefore, easily adhere to the coverglass in smear. Make uniformly thin smears on coverglasses. If the smears are made from dysenteric or fluid stools, they should be fixed almost immediately. Smears made from diarrhoeic or formed stools by emulsifying in warm salt solution, should be left for a few minutes. In any case, do not let the smear become dry except a narrow marginal zone.

The smears are fixed next. The most commonly used fixative for Protozoa is Schaudinn's fluid. This is made up as follows:

Cold saturated mercuric bichloride (6–7%) 66 cc.
Absolute or 95% alcohol 33 cc.
Glacial acetic acid 1 cc.

The first two can be kept mixed without deterioration, but the acid must be added just before fixation. Fix at room temperature or warmed to 50°C. The fixative is placed in a square Petri dish and the smear is gently dropped on it with the smeared surface facing downward. With a little experience, air bubbles can be avoided and make the smear float on the surface of the fixative. After about one minute, turn it around and let it stay on the bottom of the dish for 5 to 10 more minutes. In case the smear is too thick, a thin coat of vaseline on the upper side of the coverglass will make it to float. About six coverglass-smears may be fixed in the dish simultaneously.

The coverglass-smears are now transferred to a Columbia staining jar for coverglasses, containing 50% alcohol for 10 minutes, followed by two changes for similar length of time. Transfer the smears next to 30% alcohol for 5 minutes, and then to a jar with water, which is now placed under gently running tap water for 15 minutes. Rinse them in distilled water and stain.

Other fixatives frequently used for Protozoa are as follows:

**Bouin’s fluid**

<table>
<thead>
<tr>
<th>Component</th>
<th>Quantity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Picric acid (saturated)</td>
<td>75 cc.</td>
</tr>
<tr>
<td>Formaldehyde</td>
<td>25 cc.</td>
</tr>
<tr>
<td>Glacial acetic acid</td>
<td>5 cc.</td>
</tr>
</tbody>
</table>
Fixation for 5–30 minutes; wash with 70% alcohol until picric acid is completely washed away from the smears.

Sublimate-acetic

Saturated sublimate solution 100 cc.
Glacial acetic acid 2 cc.

This is the original fixative for Feulgen’s nucleal reaction (p. 897). Fixation and after-treatment similar to Schaudinn’s fluid.

Carnoy’s fluid

Absolute alcohol 30 cc.
Glacial acetic acid 10 cc.

Fixation for 5–30 minutes; wash in 95% alcohol.

Osmium tetroxide

The vapor from or the solution itself of 1% Osmium tetroxide may be used. Fixation in 2–5 minutes; wash in running water.

Flemming’s fluid

1% chromic acid 30 cc.
2% osmium tetroxide 8 cc.
Glacial acetic acid 2 cc.

Fixation for 10–50 minutes; wash for one hour or longer in running water.

The most commonly used stain is Heidenhain’s iron haematoxylin, as it is dependable and gives a clear nuclear picture, although it is unsatisfactory for voluminous organisms or smears of uneven thickness. It requires a mordant, ammonio-ferric sulphate (iron alum) and a dye, haematoxylin. Crystals of iron alum become yellow and opaque very easily. Select clear violet crystals and prepare 2% aqueous solution. Haematoxylin solution must be well “ripe.” The most convenient way of preparing it is to make 10% absolute alcohol solution as it does not require ripening. By diluting this stock solution with distilled water, prepare 0.5 or 1% slightly alcoholic solution which will be ready for immediate and repeated use. Smears are left in the mordant in a jar for 1–3 hours or longer. Wash them with running water for 5 minutes and rinse in distilled water. Place the smears now in haematoxylin for 1–3 hours or longer. After brief washing in water, the smears are decolorized in Petri dish in a diluted iron alum, 0.5% HCl in water or 50% alcohol, or saturated aqueous solution of picric acid under the microscope. Upon completion, the smears are
washed thoroughly in running water for about 30 minutes. Rinse them in distilled water. Transfer them through ascending series of alcohol (50 to 95%). If counter-staining with eosin is desired, dip the smears which were taken out from 70% alcohol, in 1% eosin in 95% alcohol for a few seconds, and then in 95% plain alcohol. After two passages through absolute alcohol and through xylol, the smears are mounted one by one on a slide in a small drop of mixture of Canada balsam and xylol. The finished preparations are placed in a drying oven at about 60°C. for a few days.

Other stains that are often used are as follows:

**Delafield’s haematoxylin.** If the stock solution is diluted to 1:5–10, a slow, but progressive staining which requires no decolorization may be made; but if stock solution is used, stain for 1–16 hours, and decolorize in 0.5% HCl water or alcohol. If mounted in a neutral mounting medium, the staining remains true for a long time.

**Mayer’s paracarmine.** In slightly acidified 70% alcohol solution, it is excellent for staining large Protozoa. If over-stained, decolorize with 0.5% HCl alcohol.

**Giemsa’s stain.** Shake the stock solution bottle well. By means of a stopper-pipette dilute the stock with neutral distilled water (5–10 drops to 10 cc.). Smears fixed in Schaudinn’s fluid and washed in neutral distilled water are stained in this solution for 10 minutes to 6 hours to overnight. Rinse them thoroughly in neutral distilled water and transfer them through the following jars in order (about 5 minutes in each): (a) acetone alone; (b) acetone:xylol, 8:2; (c) acetone:xylol, 5:5; (d) acetone:xylol, 2:8; (e) two changes of xylol. The smears are now mounted in cedar wood oil (which is used for immersion objectives) and the preparations should be allowed to dry for a longer time than the balsam-mounted preparations.

**Feulgen’s nucleal reaction.** The following solutions are needed.

(a) HCl solution. This is prepared by mixing 82.5 cc. of HCl (specific gravity 1.19) and 1000 cc. of distilled water.

(b) Fuchsin-sodium bisulphite. Dissolve 1 gm. of powdered fuchsin (basic fuchsin, diamant fuchsin or parafuchsin) in 200 cc. of distilled water which has been brought to boiling point. After frequent shaking for about 5 minutes, filter the solution when cooled down to 50°C. into a bottle and add 20 cc. HCl solution. Cool the solution further down to about 25°C. and add 1 gm. of anhydrous sodium bisulphite. Apply stopper tightly. Decolorization of the solution will be completed in a few hours, but keep the bottle in a dark place for at least 24 hours before using it.

(c) Sulphurous water:
Protozoology

Distilled or tap water 200 cc.
10% anhydrous sodium bisulphite 10 cc.
HCl solution (a) 10 cc.

Feulgen's reaction is used to detect thymonucleic acid, a constituent of chromatin. By a partial hydrolysis, certain purin-bodies in the acid are split into aldehydes which show a sharp Schiff's reaction upon coming in contact with fuchsin-sodium bisulphite. Thus this is a reaction, and not a staining method. Smears fixed in sublimate-acetic or Schaudinn's fluid are brought down to running water, after being placed for about 24 hours in 95% alcohol. Immerse them in cold HCl for one minute, then place them in HCl kept at 60°C. (over a microburner or in an incubator) for 5 minutes, quickly immerse in cold HCl. After rapidly rinsing in distilled water, place the smears in solution (b) for 30-minutes to 3 hours. There is no overstaining. The smears are then washed in three changes (at least 2 minutes in each) of solution (c). Wash them in running water for 30 minutes. If counterstaining is desired, dip in 0.1% light green solution and rinse again in water. The smears are now dehydrated through a series of alcohol in the usual manner and mounted in Canada balsam (Feulgen and Rossenbeck, 1924; Feulgen-Brauns, 1924; Feulgen, 1926; Coleman, 1938; Stowell, 1945).

Silver-impregnation methods. Since Klein (1926) applied silver nitrate in demonstrating the silver-line system of ciliates, various modifications have been proposed.

Dry silver method (Klein, 1926). Air-dried cover glass smears are placed for 6–8 minutes in a 2 per cent solution of silver nitrate and thoroughly washed. The smears are exposed to sunlight for 2–8 hours in distilled water in a white porcelain dish, with occasional control under the microscope. The smears are then washed thoroughly and air-dried; finally mounted in Canada balsam.

Wet silver method (modified after Gelei and Horváth, 1931). The ciliates are fixed in a centrifuge tube for 5–10 minutes in sublimate-formaldehyde solution, composed of saturated corrosive sublimate 95 cc. and formaldehyde 5 cc. The specimens are now washed twice in nonchlorinated water and once in distilled water; they are then treated in 1.5–2 per cent solution of silver nitrate for 5–20 minutes. Without washing, the specimens in the tube are exposed to direct sunlight for 10–60 minutes in distilled water, after which the specimens are washed 4–6 times in distilled water, one minute each. Passing through a gradually ascending alcohol series and xylol, the specimens are mounted in Canada balsam.
Fontana's method. For staining filamentous structures such as the extruded polar filament of microsporidian spores, this method is the most satisfactory one. After air-drying the smears are fixed for 5 minutes in a mixture of formaldehyde, 20 cc.; glacial acetic acid, 1 cc.; and distilled water, 100 cc. After washing in running water, the smears are placed in the following mordant composed of equal parts of 5 per cent tannic acid and 1 per cent carbolic acid, for about 2 minutes at about 60°C. Wash the smears in water and place them for 3–5 minutes in 0.25 per cent solution of silver nitrate warmed to 60°C., to which ammonia has been added drop by drop until a grayish brown cloud appeared. Wash thoroughly and air-dry. After passing through 95 per cent and absolute alcohol, and xylol, the smears are mounted in Canada balsam.

b. Blood film preparations

Thin film. The finger tip or ear lobe is cleaned with 70% alcohol. Prick it with an aseptic blood lancet or a sterilized needle. Wipe off the first drop with gauze and receive the second drop on a clean slide about half an inch from one end (Fig. 376, 1). Use care not to let the slide touch the finger or ear-lobe itself. Quickly bring a second slide, one corner of which had been cut away, to the inner margin of the blood drop (1), and let the blood spread along the edge of the second slide. Next push the second slide over the surface of the first slide at an angle of about 45° toward the other end (2). Thus a thin film of blood is spread over the slide (3). Let the slide lie horizontally and dry, under a cover to prevent dust particles falling on it and to keep away flies or other insects. If properly made, the film is made up of a single layer of blood cells.

Thick films. Often parasites are so few that to find them in a thin film involves a great deal of time. In such cases, a thick film is advocated. For this, 2 to 4 drops of blood are placed in the central half-inch square area, and spread them into an even layer with a needle or with a corner of a slide. Let the film dry. With a little practice, a satisfactory thick smear can be made. It will take two hours or more to dry. Do not dry by heat, but placing it in an incubator at 37°C. will hasten the drying. When thoroughly dry, immerse it in water and dehaemoglobinize it. Air dry again.

Thin and thick film. Often it is time-saving if thin and thick films are made on a single slide. Place a single drop of blood near the center and make a thin film of it toward one end of the slide. Make a small thick smear in the center of the other half of the slide. Dry. When
thoroughly dry, immerse the thick film part in distilled water and de- 
haemoglobinize it. Let the slide dry.

Blood smears must be stained as soon as possible to insure a proper 
staining, as lapse of time or summer heat will often cause poor stain-
ing especially of thick films. Of several blood stains, Giemsa’s and 
Wright’s stains are used here. For staining with Giemsa’s stain, the 
thin film is fixed in absolute methyl alcohol for 5 minutes. Rinse well

Fig. 376. Diagrams showing how a thin blood film is made on a slide.

the slide in neutral distilled water. After shaking the stock bottle 
(obtained from reliable makers) well, dilute it with neutral distilled 
water in a ratio of one drop of stain to 1–2 cc. of water. Mix the solu-
tion and the blood film is placed in it for 0.5–2 hours or longer if 
needed. Rinse the slide thoroughly in neutral distilled water and 
wipe off water with a tissue paper from the underside and edges 
of the slide. Let the slide stand on end to dry. When thoroughly dry, 
place a drop of xylol and a drop of cedar wood oil (used for immersion 
objectives) and cover with a coverglass. The mounting medium
should be absolutely neutral. Do not use Canada balsam for mounting, as acid in it promptly spoils the staining.

For Wright's stain, fixation is not necessary. With a medicine dropper, cover the dried blood film with drops of undiluted Wright's stain, and let the film stand horizontally for 3–5 minutes; then the same number of drops of neutral distilled water is added to the stain and the whole is left for 10–30 minutes. The stain is then poured off and the film is rinsed in neutral distilled water. Dry. Mount in xylol and cedar wood oil.

Use of coverglass on a stained blood film is advocated, since a cedar wood oil mounted slide allows the use of dry objectives which in the hand of an experienced worker would give enough magnification for species determination of Plasmodium, and which will very clearly reveal any trypanosomes present in the film. Furthermore, the film is protected against scratches, and contamination by many objects which may bring about confusion in detecting looked-for organisms.

Films made from splenic punctures for Leishmania or Trypanosoma are similarly treated and prepared.

c. Section preparations

Paraffin sections should be made according to usual histological technique. Fixatives and stains are the same as those mentioned for smear preparations.

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