Giant protistan parasites on the gills of cephalopods (Mollusca)

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ABSTRACT: Large Protista of unknown taxonomic affinities are described from 3 species of coleoid squids, and are reported from many other species of cephalopods. The white to yellow-orange, ovoid cyst-like parasites are partially embedded within small pockets on the surface of the gills, often in large numbers. Except for a holdfast region on one side of the large end, the surface of the parasite is elaborated into low triangular plates separated by grooves. The parasites are uninucleate; their cytoplasm bears lipid droplets and presumed paraglycogen granules. Trichocysts, present in a layer beneath the cytoplasmic surface, were found by transmission electron microscopy to be of the dinoflagellate type. Further studies are needed to clarify the taxonomic position of these protists.

INTRODUCTION

Cephalopods harbor a diversity of metazoan and protozoan parasites (Hochberg 1983). In this study we used light and electron microscopy to characterize a group of unusual parasites that are found on the gills of numerous species of pelagic cephalopods, but which apparently have never been described. We identify them as large Protista of unknown taxonomic affinities bearing trichocysts of the type that is characteristic of a wide variety of dinoflagellates.

MATERIALS AND METHODS

Two coleoid squids Moroteuthis robusta were collected in 1982 from the vicinity of the Friday Harbor Laboratories, San Juan Island, Washington, USA. The first specimen (dorsal mantle length, DML = 100.5 cm) was netted in June by a local fisherman. The second specimen (DML = 127 cm) was found in July in an advanced state of decomposition on a beach. The parasites were fixed in 3% glutaraldehyde (Ladd) in 0.1 M phosphate buffer (pH 7.3) with 0.35 M sucrose (room temp. 1 h), rinsed briefly in buffer with sucrose, and postfixed on ice in 1% osmium tetroxide in 0.1 M phosphate buffer (pH 7.3) with 0.35 M sucrose (1 h). These specimens were embedded immediately in epoxy resin (see below). One specimen each of the coleoid squids Abralia trigonura and Histioteuthis dolfini were trawled near Oahu, Hawaii, in March, 1980. Gill parasites from the former were fixed in formalin; those from the latter were fixed in osmium tetroxide. Parasites from both were stored in 70% ethanol until 1982, then osmicated as described above and prepared for light microscopy (LM), transmission electron microscopy (TEM) or scanning electron microscopy (SEM).

Specimens for sectioning were dehydrated in an ethanol series, transferred through 3 changes of propylene oxide, and embedded in epoxy resin. Parasites from the first specimen of Moroteuthis robusta were originally embedded in Spurr's medium. Because difficulties were encountered with the resin during sectioning, the embedding medium was removed with methyl ethyl ketone (Fullam). The specimens were dehydrated in a methanol series, then transferred through ethanols to propylene oxide, and were re-embedded in Medcast (Pelco). All other parasites were embedded directly in Medcast. For LM, 1 µm sections were stained in 1% methylene blue in 1% borax. For TEM, silver sections were stained in saturated aqueous uranyl acetate for 12 min, and in 0.1% lead citrate for 3 min; they were examined with a Philips 300 transmission electron microscope.

For SEM, doubly fixed specimens were dehydrated in an ethanol series, transferred through 2 changes of
RESULTS

Protists of the type described here have been found (by F. G. H.) on the gills of many species of cephalopods in addition to those examined in the present study: the squids Heteroteuthis hawaiiensis, Pyroteuthis addolux, Pterygioteuthis microlampas, Chiroteuthis picteti, Ctenopteryx sicaula, Octopoteuthis nielseni and the octopus Eledonea pygmaea from the central North Pacific Ocean off Hawaii; the squid Abraliopsis felis, Histiotethis heteropseis, Chiroteuthis calyx, Gonatopsis borealis, Gonatus sp., Berryteuthis magister, Galiteuthis sp., the octopus Japatella dia phana and the vampire squid Vampyroteuthis infer nalis from the eastern North Pacific Ocean; and the squid Loliguncula brevis from the Gulf of Mexico. The parasites are obviously common in pelagic cephalopods.

When the mantle cavity is exposed, the cystlike parasites are easily visible to the naked eye. Their color varies from white to yellow-orange depending on the host species. They are partially embedded within small pockets on the surface of the gills. Although fixed to the gills at this stage of the life cycle, the parasites are easily removed. They occur along the full length of the gill on both the outer and inner demibranchs, between the gill lamellae and on the exposed surfaces. At present there is no evidence that they cause necrosis, although the holdfast penetrates the gill tissue.

External morphology

The parasites from all 3 species of squids are similar in morphology. They are ovoid in shape, and most of the surface is elaborated into low triangular plates. The latter are arranged in groups around numerous foci (Fig. 1 to 3). The number of plates per focus varies from 4 to 7. The holdfast, or area that attaches to the host, is located on one side of the large end; it lacks plates and bears 2 small papillae, each with an opening (arrowhead) at the apex (Fig. 2). The tapered end of the parasite bears an elongate pore of unknown function (Fig. 4).

Parasites from the different species of squids differ from one another in details of the plates and in size. Shallow indentations on all sides of some plates of the Histiotethis dolfini parasite (Fig. 1) are distinctive. The plates of the Moroteuthis robusta parasite are relatively smooth and raised into short ridges at each corner (Fig. 3). The grooves between plates are broad and shallow compared with some of the grooves of the H. dolfini parasite (Fig. 1). Parasites from Abrallia trigonura have flat, smooth plates, although fracturing and exfoliation due presumably to formalin fixation raise the possibility that this is an artifact. Parasites from M. robusta (l = 1.62 mm ± 0.23 mm standard deviation, w = 1.05 mm ± 0.19 mm standard deviation, n = 40) were somewhat larger than those from A. trigonura (l = 1.10 mm, w = 0.80 mm, n = 1), and much larger than those from H. dolfini (l = 0.56 mm, w = 0.36 mm, n = 1).

Internal morphology

Sections reveal that the parasites are unicellular. A single large nucleus is located centrally in cross sections (Fig. 5). It has a relatively electron-lucent flocculent matrix and numerous basophilic bodies. The cytoplasm is liberally supplied with lipid droplets and with electron-dense granules that appear to be paraglycogen. The cytoplasm is covered by a wall that gives the plates their integrity; lobes of cytoplasm extend into the convolutions of the wall (Fig. 5). Fixation of the cytoplasm and nucleus was poor in all cases, which suggests that the wall is resistant to penetration by the reagents utilized. The cytoplasmic surface is immediately underlain by a layer of trichocysts which are perpendicular to the extracellular wall, and which in some specimens were remarkably well preserved. No substantive differences were found between the trichocysts of the squid gill parasites and those of the freeliving dinoflagellate Gonyaulax catenella (see below).

The trichocysts are rod-shaped (approximately 1.7 μm long and 0.3 μm wide) and bound by a dense...
Fig 6 to 13 Trichocysts Scale bars = 200 nm. Fig 6 to 8 Note electron-dense core with conical head and group of filaments. Fig 6 Protist from Moroteuthis robusta Arrowhead dense plaque Fig 7 Protist from Abralia trigonura Fig 8. The dinoflagellate Gonyaulax catenella Fig 9 Protist from M robusta Arrowheads hoops in trichocyst membrane Fig 10. The dinoflagellate C catenella Arrowheads hoops in trichocyst membrane Fig 11 to 13 Trichocyst core showing longitudinal periodicity. Fig 11 Protist from Histioteuthis doliaut Fig 12. The dinoflagellate G catenella Fig 13 Protist from M robusta Fig 14 Ejected trichocyst of protist from M robusta At right, longitudinal section Arrowheads indicate transverse periodicity. Arrow: cross section Scale bar = 50 nm
membrane. The latter encloses an electron-dense core that has a conical head, and a group of filaments that extend from the head to several dense plaques near the limiting membrane of the parasite (Fig. 6 & 7). The core and filaments of trichocysts from parasites of *Moroteuthis robusta* (Fig. 6) and *Abralia trigonura* (Fig. 7) have notably similar counterparts in trichocysts of the dinoflagellate *Gonyaulax catenella* (Fig. 8). The membrane limiting each trichocyst bears hoops (Fig. 9 & 10), which measure 30 nm between centers in the *M. robusta* parasite (Fig. 9), and 23 nm in the dinoflagellate (Fig. 10). A longitudinal periodicity of the core was occasionally well preserved (Fig. 11 to 13), measuring, between centers of the longitudinal members, 9.3 nm in the *Histiotethis doleleini* protozoan (Fig. 11), 10.5 nm in the dinoflagellate (Fig. 12), and 8.8 nm in the protozoan from *M. robusta* (Fig. 13). Ejected trichocysts were found in parasites from the decayed *M. robusta* that was acquired in July 1982. The threads are square in cross section (about 60 nm per side); the longitudinal axis displays a transverse periodicity of 55 nm (Fig. 14).

**DISCUSSION**

The external morphology of the parasites provides few clues to their taxonomic position. The lack of external cilia or flagella does not exclude the possibility that they are ciliates or flagellates, as encysted stages of these groups typically lack motile organelles. Given that the hosts are highly mobile, it is probable that the life cycle includes an as yet undiscovered motile stage. The presence of an apical pore also suggests that the parasites may be dinoflagellates, and furthermore, that they are a cyst stage (Dodge 1985). The nucleus of dinoflagellates typically contains visible organized chromosomes, which are not evident in our specimens, but this is not a routine feature (Hollande & Corbel 1982).

Amoebae typically lack trichocysts, but an estuarine amoeba described by Page (1979) contains a trichocyst-like body. This differs from the trichocysts reported here in that longitudinal striae are not visible in the mature stage, the core head is not conical, and the organelle membrane does not bear hoops. Thus it is unlikely that the gill parasites are sarcodines.

Present evidence argues against the possibility that the squid gill parasites are the cysts of ciliates, although many squids harbor extremely large endosymbiotic ciliates in the digestive tract (Hochberg unpubl.). Cysts of ciliates typically exhibit nuclear dualism (Grimes 1973, Walker & Maugel 1980, Walker et al. 1980), whereas the gill parasites appear to have a single large nucleus. Unlike those described here, mature trichocysts of normal ciliates lack a conical end on the core, longitudinal periodicity, and filaments, and furthermore the trichocyst membrane lacks hoops (Yusa 1965, Bradbury 1966, Bannister 1972, Esteve 1974, Anderer & Hausmann 1977, Peck 1977, Njine & Didier 1980). The core of pre-trichocysts of *Paramecium caudatum* has very fine longitudinal striations that are lost as the trichocysts develop (Yusa 1963), and trichocysts of mutant *P. aurelia* have very fine longitudinal striations (Pollack 1974). The large mature trichocysts of normal *Paramecium* spp., however, are quite unlike those described in the present study. A tiny bundle of filaments is present in the trichocyst of *Drepanomonas dentata* (Hausmann & Mignot 1975), but the accompanying 4 dense rod-like structures appear to be unique. A bounding membrane in *Neobursaridium*
gigas bears large hoops (Dragesco 1968), but the core has a transverse periodicity and the tip is dense as in *Paramecium* spp.

This paper brings attention to a group of giant protistan parasites on the gills of cephalopods, and provides ultrastructural evidence that their trichocysts are similar to those of dinoflagellates. Differences in size and plate morphology of parasites obtained from different species of cephalopods suggest that many species of these protists exist. Further information about the cytology and life stages is needed and will require study of more fresh hosts of various ages, and, perhaps, culturing of the parasites.

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LITERATURE CITED


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