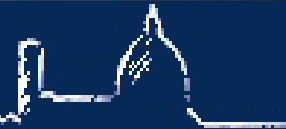


**Exotopic protrusions and
Ellobiopsids' infection on freshwater
copepods:
another invasion from marine
sites?**

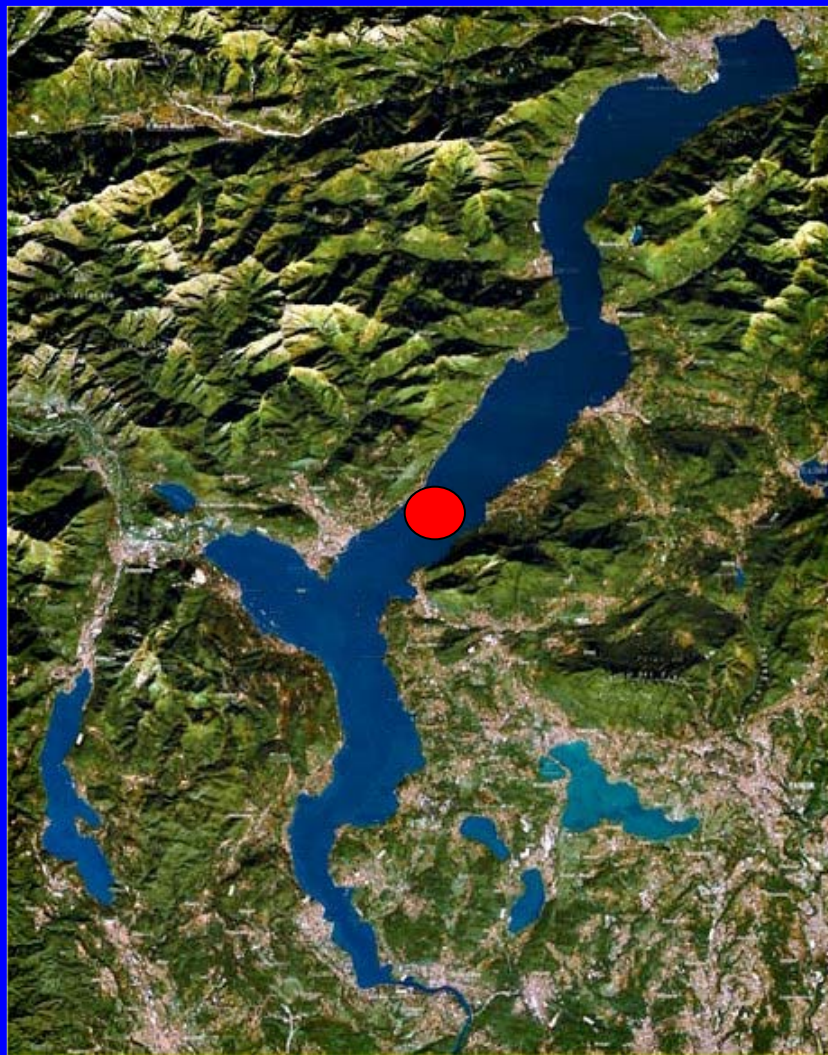


Università degli Studi di Firenze



Biological Invasions in Inland Waters

International Workshop - Florence, May 5-7, 2005



Commissione internazionale per la protezione delle acque italo-svizzere
Ricerche sull'evoluzione del Lago Maggiore

**Long-term zooplankton records: 1978-
data from 1909**

100 μm

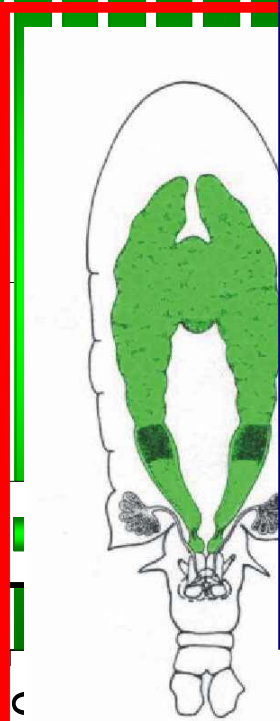
100%

50%

0%

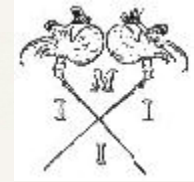
'48
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'91
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Memorie
dell'Istituto Italiano di Idrobiologia

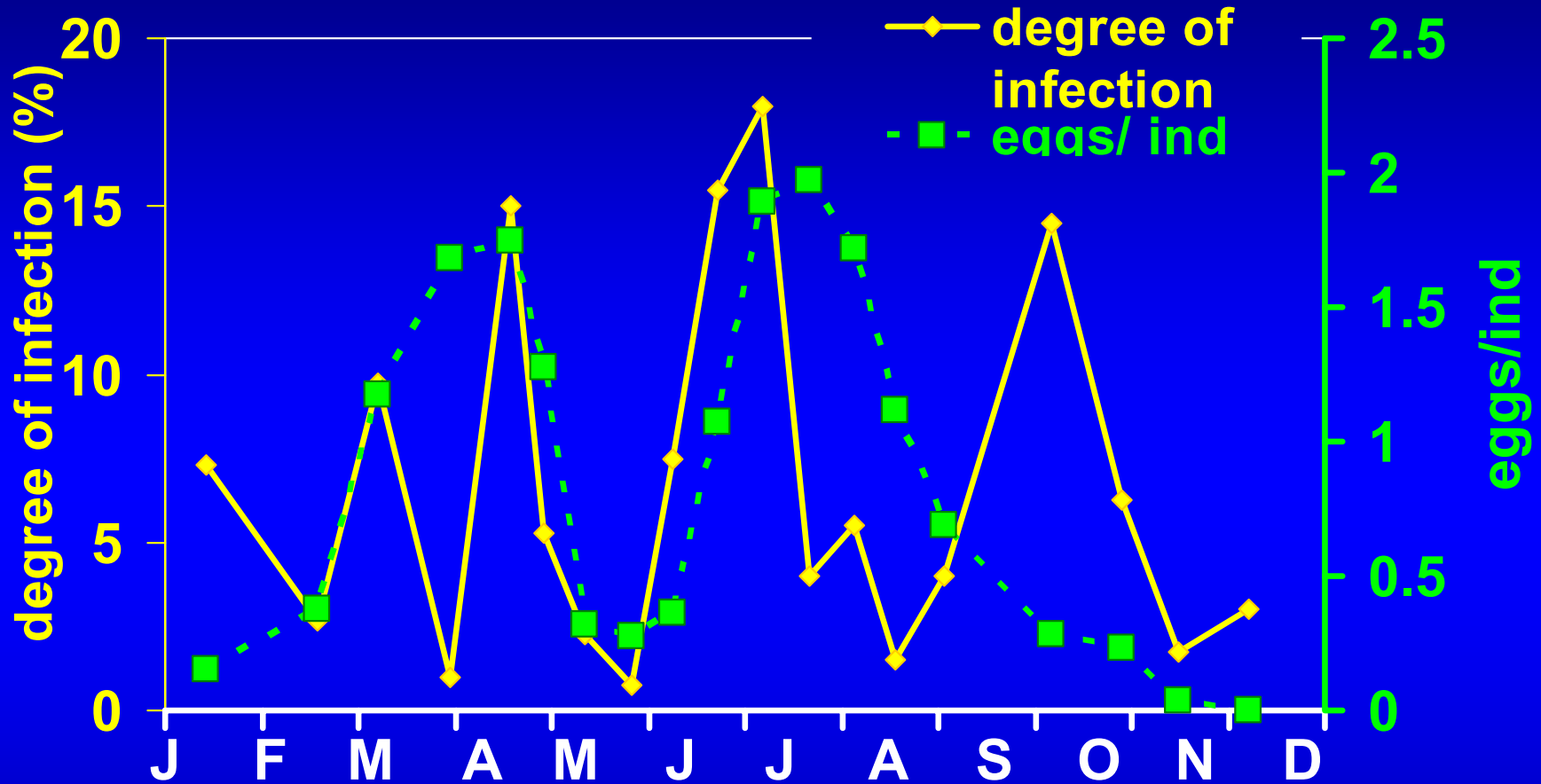
International Journal of Limnology



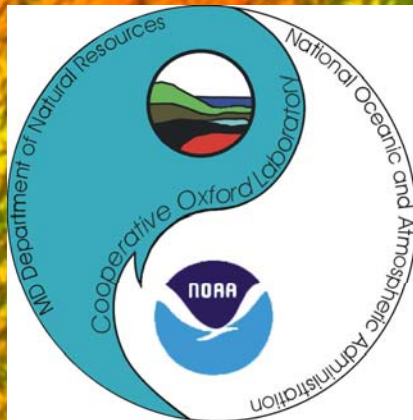
Volume
54

- **level of infection of the population, i.e. the percentage of infected/total number of animals**
- **prevalence, i.e. the proportion of infections on different species and/or developmental stages**

Specificity of the host: adults or pre-adults of *Eudiaptomus padanus*



Infection dynamics related to egg production and adults' share



Cooperative Oxford
Laboratory

Gretchen Messick

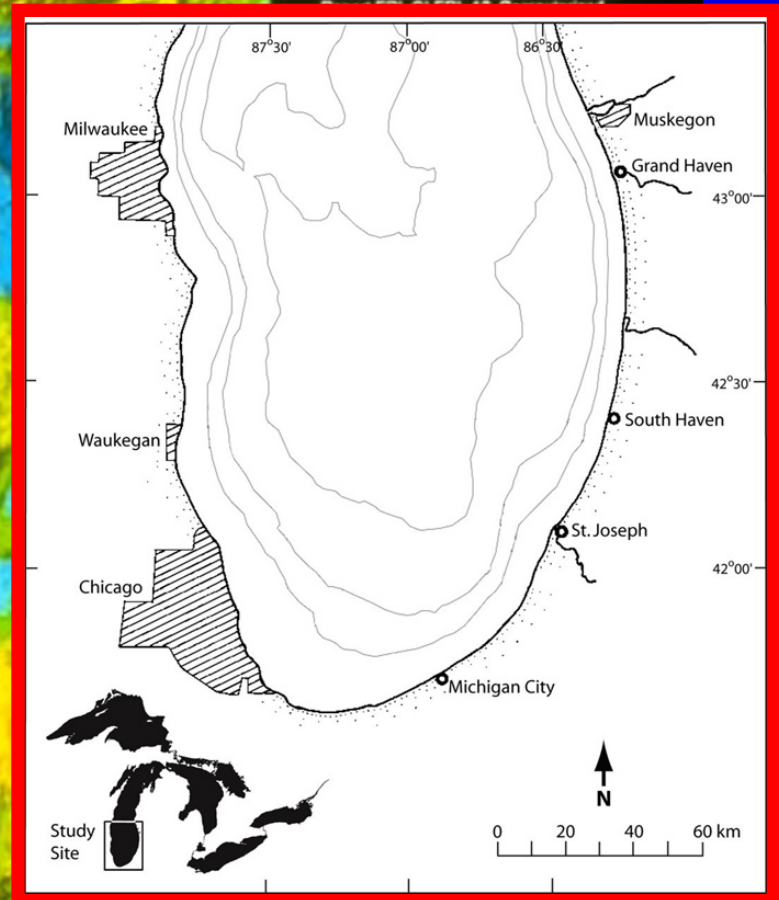
whereas the distance in the south and west. The distance in the north and west of the scene.



Great Lakes Environmental
Research Laboratory

Data sources:

- USGS 30 sec. topography from NOAA Geophysics of North America CD ROM.
- Lake bathymetric data from NOAA Data



LES
PÉRIDIINIENS PARA
MORPHOLOGIE, REPRODUCT
ETHOLOGIE

PAR
ÉDOUARD CHATTON

Chef de laboratoire à l'Institut Pasteur, Paris.
Maître de Conférences à la Faculté des Sciences, Strasbourg

Arni Litt, Zoology
Department,
University of
Seattle, Washington

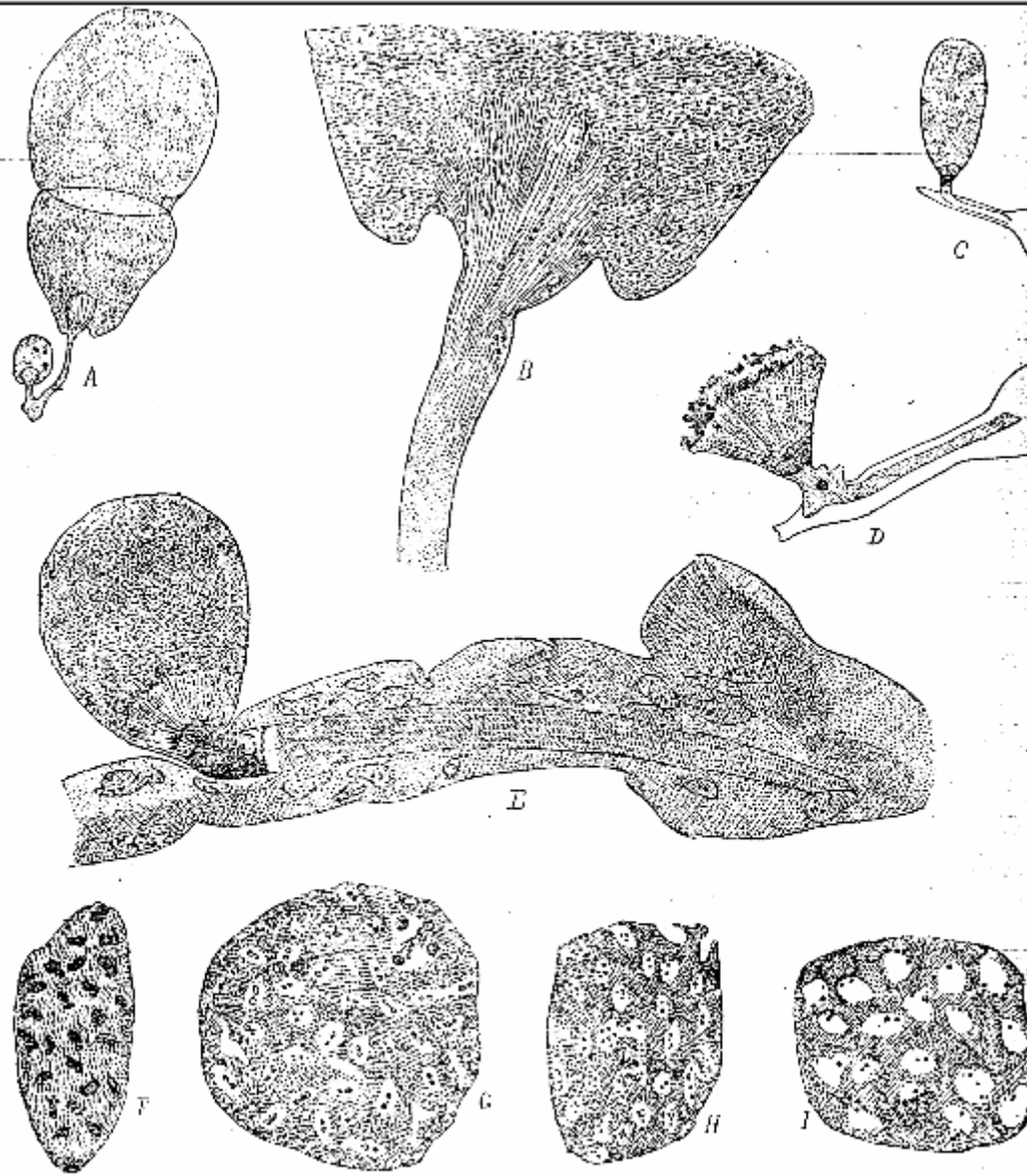


FIG. CXXVI. M. CAULLERY (1910). *Peridinium Chattoni*: a, coupe longitudinale du stade le plus avancé, montrant la séparation des deux parties proximale et distale et la fixation par un pédoncule à un appendice de l'hôte coupé transversalement ($\times 45$); b, coupe de la portion basilaire du parasite précédent ($\times 280$); c, stade très-jeune ($\times 280$); d, portion basilaire du même ($\times 750$); e, stade moyen montrant la tige de fixation engagée dans l'autosome de l'hôte ($\times 280$); f, fragments de coupes d'*Peridinium* à divers stades au point de l'aspect des noyaux chromosomiques ($\times 1125$).

«Coutiere (1911) a créé la famille des Ellobiopsidae par en ensemble de Protistes parasites des crustacés pelagique, dont le type est l' *Ellobiopsis chattoni* CAULLERY (1910)....»

***E. chattoni*, « Type de l'espèce parasite externe de *Calanus helgolandicus* à Banyuls-sur-Mer.... *Calanus helgolandicus* est un Copépode rare en Banyuls-sur-Mer »**

« La Méditerranée ne semble pas être l'habitat de prédilection du *Calanus helgolandicus*, ni le milieu le plus propre à l'expansion de son parasite

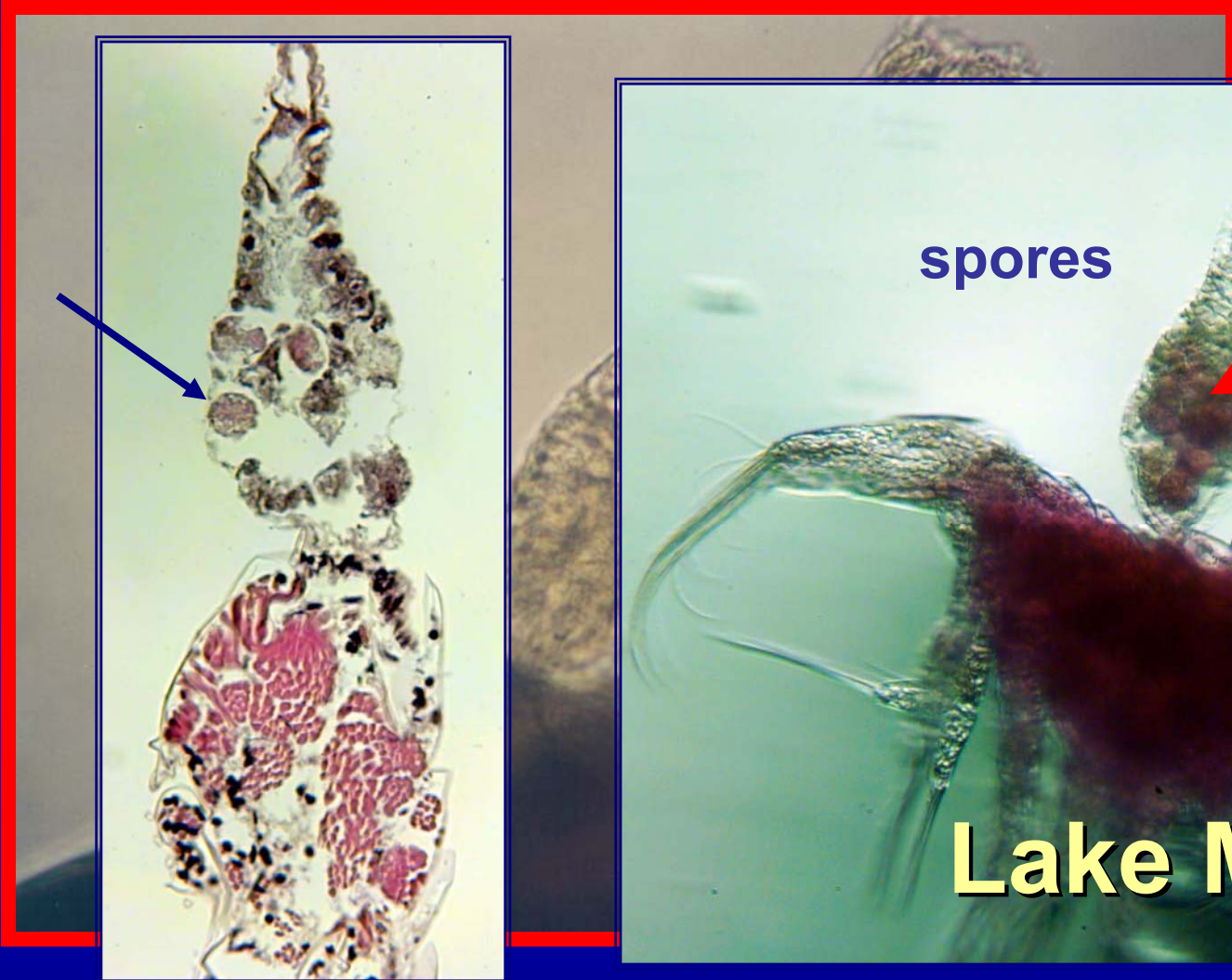
L'un et l'autre paraissent être surtout des formes septentrionales :

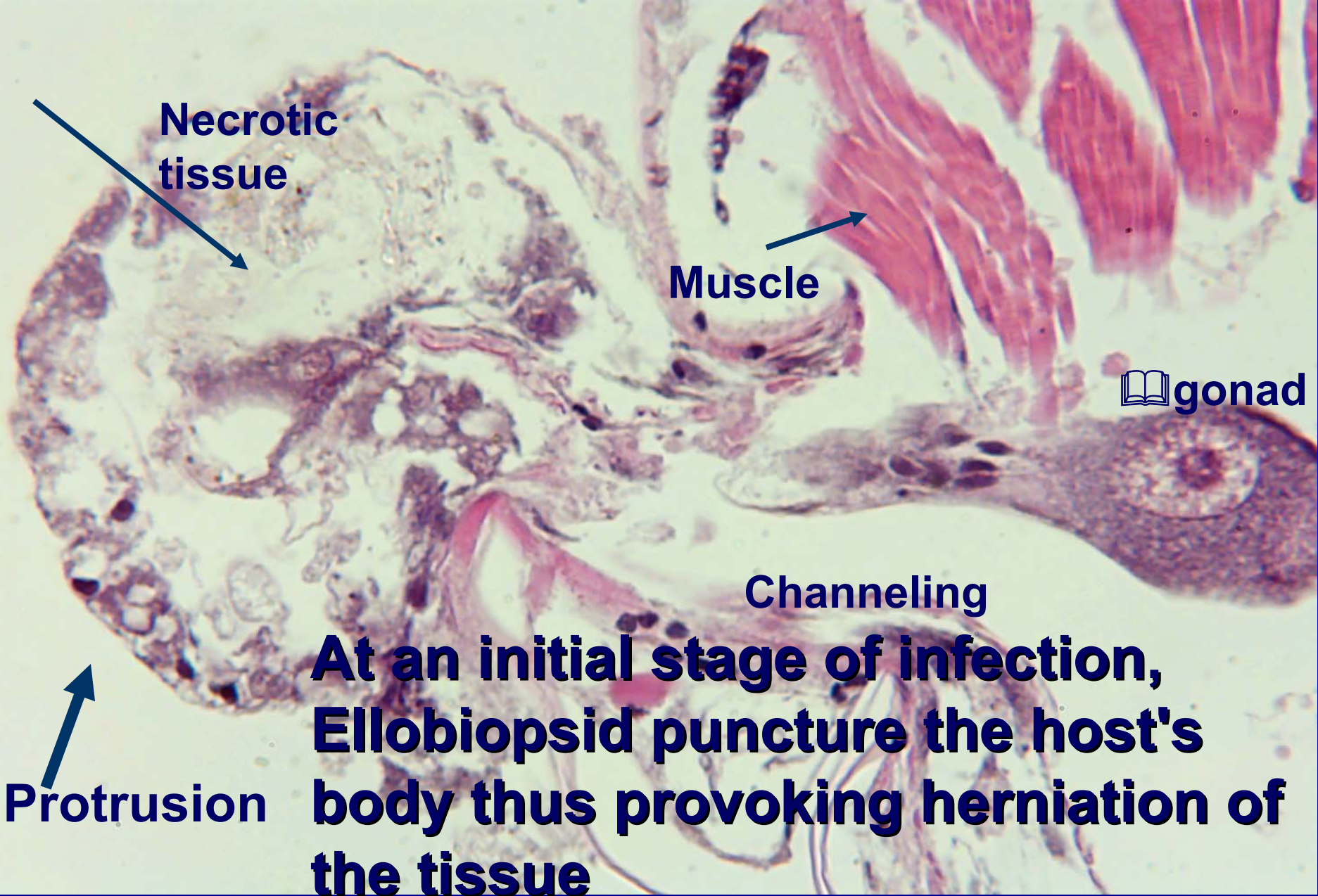
Loch Fyne, Scotland (Scott, 1896)

North Sea and the Baltic on *Calanus finmarchicus*, *Pseudocalanus* (?), *Acartia clausi* (Apstein, 1911)

- They were well known from marine copepods
- No record from freshwater sites

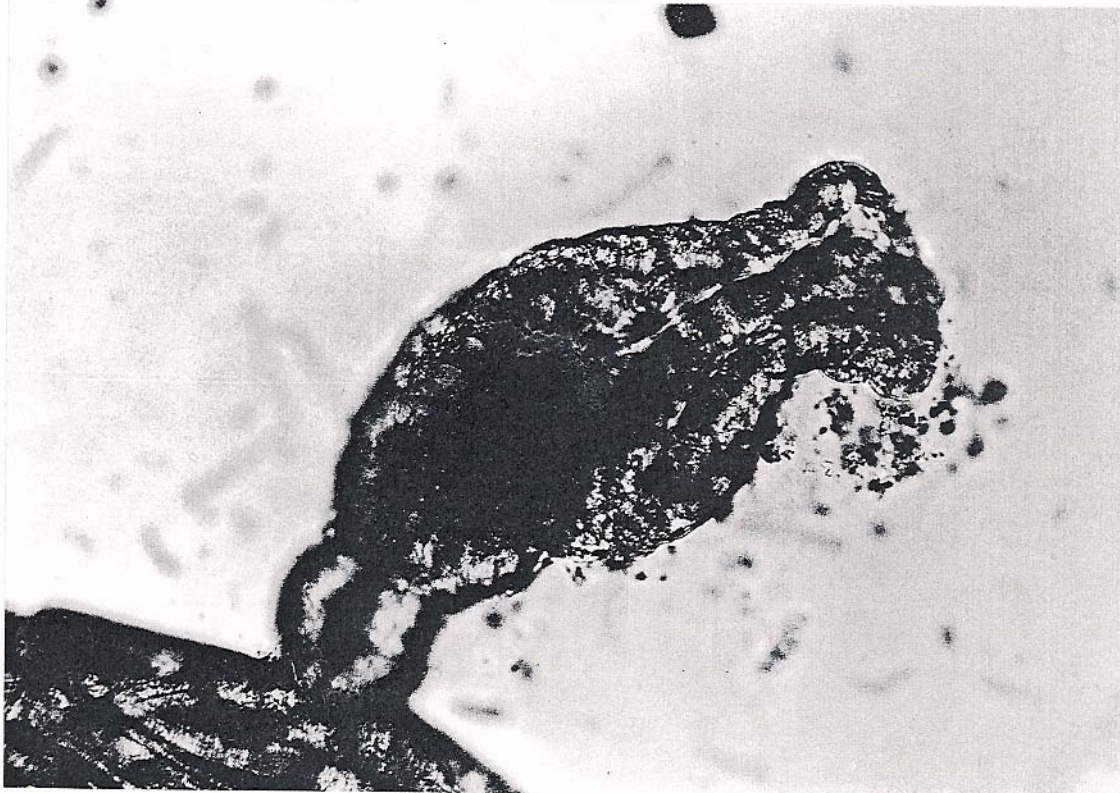
Lake Maggiore





At an initial stage of infection, Ellobiopsid puncture the host's body thus provoking herniation of the tissue

Channeling and necrotic tissue



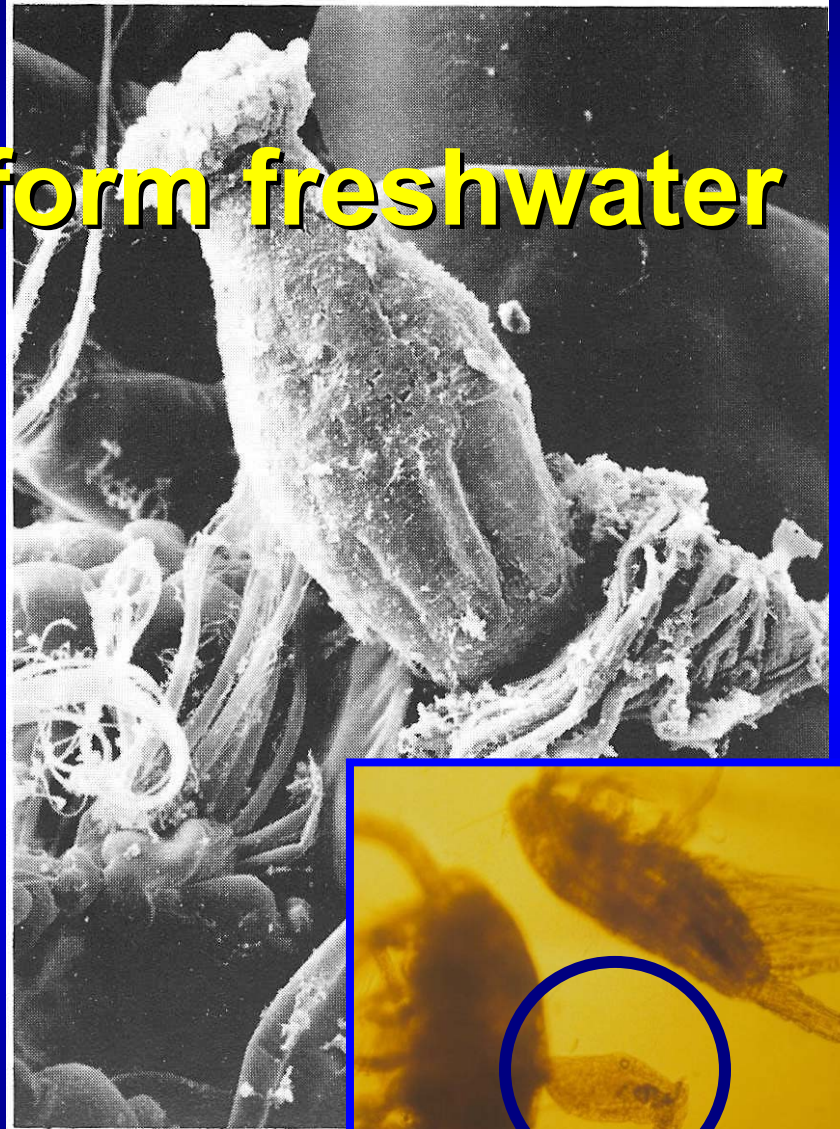
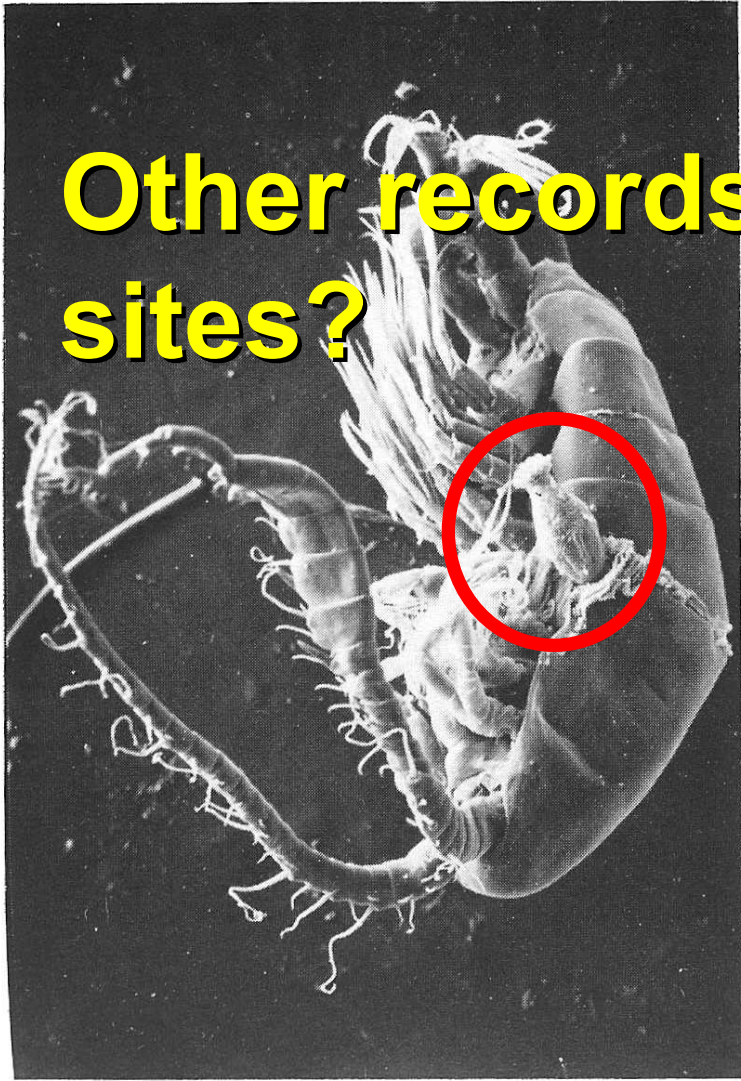
**Any
records
from
marine
sites in
Italy?**

YES

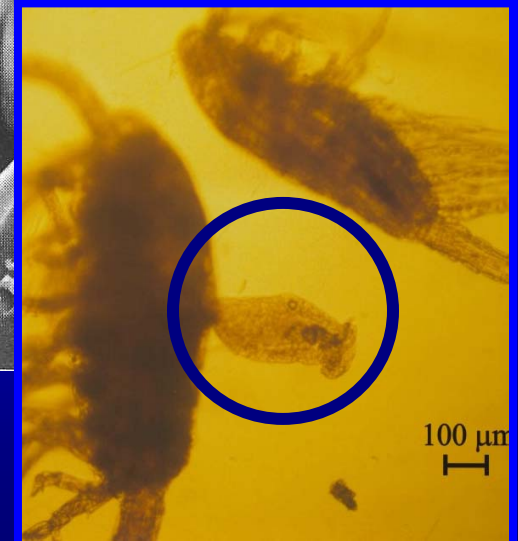
**Ellobiopsis on *Acartia latisetosa* from Mar
Piccolo di Taranto; photo by Giovanni
Fanelli, CNR ITT, now CNR ISMAR, Taranto**

Lake Midmar, South Africa

Other records from freshwater sites?



Lake Maggiore, Italy



First record of a freshwater calanoid *Tropodiptomus spectabilis* (Kiefer, 1929) (Crustacea, Copepoda) as host of an ellobiopsid parasite

Nancy A. Rayner and Eleanor M. King

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Abstract. Ellobiopsids have been recorded as parasites of marine pelagic Crustacea. An unidentified ellobiopsid has been discovered for the first time in freshwater, its host being a calanoid, *Tropodiptomus spectabilis* (Kiefer), in South Africa.

basing their decision on the morphology of the zoospores of *Phaeocystis* spp. Kane, which have the unique flagellation of the dinoflagellates. No records can be found in the literature of ellobiopsids having been found on freshwater Crustacea. Host species are usually euphausiids, mysids and amphipods (Vader, 1973). Jepps (1937) recorded *Ellobiopsis chattoni* Caullery as a parasite of *Calanus finmarchicus* Gunnerus, a marine calanoid of the Clyde Sea area, and noted that it was first recorded by Scott in the Firth of Clyde in 1896.

An ellobiopsid parasite has been found to parasitise *Tropodiptomus spectabilis*, the dominant calanoid (Rayner, 1981) in Lake Midmar (29°30'S; 30°12'E), a warm-temperate oligotrophic impoundment situated at an altitude of 1044 m in the Natal midlands, Republic of South Africa. At full supply the surface area of the lake is 15.59 km² and the mean depth 11.4 m (Breen, 1983).

Tropodiptomus spectabilis (Kiefer, 1929) is similar to *Ellobiopsis chat*



Pergamon

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0959-8030(94)E0001-G

THE PARASITIC DINOFLAGELLATES OF MARINE CRUSTACEANS

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Gloucester Point, VA, 23062, USA

Abstract. Parasitic dinoflagellates have recently emerged as significant disease agents of commercially important crustaceans. For example, epizootics of *Hematodinium* have seriously affected certain crab and lobster fisheries. The parasitic dinoflagellates of crustaceans are, however, relatively unknown. Marine crustaceans are parasitized by two orders of dinoflagellates: the Blastodinida and the Syndinida. Crustaceans are also parasitized by the Paradinida and the Ellobiopsidae, taxa that have close historical ties and possible taxonomic affinities with the dinoflagellates. The taxonomy and life history patterns of the different parasitic species are largely dictated by their host-parasite relationships. For example, sporulation in the blastodinids occurs internally but is completed externally with the expulsion of spores via the anus of the host. The egg-parasitic chytridiodinids sporulate externally after destroying their host egg. The tissue-dwelling syndinids have plasmodia that sporulate internally and generally kill their hosts upon the expulsion of the dinospores. Unfortunately, complete life cycles have not been elucidated for any of the parasitic forms, hence characteristics of the life cycles must be applied cautiously to the systematics of the taxa. For example, gamogony and the presence of resting cysts are only known from a few species; they probably occur in most species. Further work on the life cycles of the parasitic dinoflagellates of crustaceans should concentrate on establishing the life cycles of representative species from each order or family. Parasitic dinoflagellates infect copepods, amphipods, mysids, euphausiids, and decapods. Their pathogenicity varies with their invasiveness in the host. The gut-dwelling blastodinids are relatively benign, while the chytridiodinids kill their host egg. Members of the pervasive Syndinida and Paradinida are overtly pathogenic and insidiously ramify throughout the hemal sinuses and organs of their hosts. Members of the Ellobiopsidae vary from the commensal *Ellobiocystis* to the overtly parasitic *Thalassomyces*. Host castration and feminization are common pathologic results of infection by these parasites. The severity of the castration is dependent upon the invasiveness of the parasitic species and the duration of the infection, while the degree of feminization is related to the stage at which the host acquires the infection. Most of the parasitic dinoflagellates occur in epizootics in their host populations. Recent epizootics of *Hematodinium* spp. have had severe effects on crustacean fisheries in Alaska, Virginia, and Scotland, and may potentially result in changes to the benthic communities of the hosts. The epizootics are often associated with host-parasite systems that occur in regions with unique hydrological features, such as fjords or poorly draining estuaries with shallow sills. These regions are ideal for the application of a "landscape" ecology approach that could lead to a better understanding of the epizootiology of parasitic dinoflagellates and other marine pathogens.

Keywords. Epizootiology, Life cycle, Taxonomy, Pathology, *Actinodinium*, *Atelodinium*, *Blastodinium*, *Chytridiodinium*, *Dissodinium*, *Ellobiocystis*, *Ellobiopsis*, *Hematodinium*, *Paradinium*, *Parallobiopsis*, *Rhizellobiopsis*, *Schizochytridinium*, *Syltodinium*, *Syndinium*, *Thalassomyces*, *Trypanodinium*

literature on
marine
environments
and or that
dealing with fish
diseases

“included in the review are two protistan taxa of unknown affinities, the Paradina and the Ellobiopsidae. Their inclusion is warranted for comparative purposes and to illustrate the unresolved questions on their affinities to the dinoflagellates”.

Differences among the genera are based on the **structure** of attachment (invasive or non-invasive, and single or multiple) as well as the **number of trophomeres** as well as **gonomeres** and the way they sporulate

• *Ellobiocystis*.....a single trophomere attached to the host via a noninvasive mucoid stalk, and 1-2 gonomeres arising from the trophomere

• *Ellobiopsis*possessing a trophomere with a single, invasive peduncle for host attachment ; multiple gonomeres arising from the trophomere

• *Parallobiopsis coutieri* ... A trophomere with a sucker-like holdfast and a simple, invasive peduncle; a single gonomere developing from the trophomere

“the genus *Ellobiocystis* does not penetrate into the body of its host, and the location around the buccal appendages suggest a commensal relation with their hosts” ...

Little is known on the pathology of *E. chattoni* infections. The penetration of the invasive stalk into the host causes localized damage to the surrounding musculature

- The life cycle is also briefly outlined

“A flagellated spore presumably settles close to...the host, developing into a trophomere, which in *Ellobiopsis* penetrates through the host cuticle, apparently functioning in absorption”...in *E. fagei*, sporulation starts as the gonomere becomes tightly constricted from the trophomere” and the spore are released, whose phate is unknown”

While being known since a very long time for marine copepods, it seems to be relatively new for freshwater copepods

possible explanations for this:

- a general lack of interest on structures by fw zooplanktologists**
- a basic ignorance of the marine literature by the fw zooplanktologists**
- while being an old phenomenon for marine specimens, is it a relatively new one for freshwater copepods?**



PO RIVER CATCHMENT BASIN

- | | | | |
|-----|------------|-----|-----------------|
| 1 | MONTORFANO | 19 | COMABBIO |
| 5 | CANDIA | 21 | S.M. VALVESTINA |
| 5* | SIRIO | 22 | ENDINE |
| 6 | GARDA | 22* | PUSIANO |
| 7 | GARLATE | 23 | ANNONE |
| 8 | PIANO | 23* | IDRO |
| 10 | MAGGIORE | 23^ | SEGRINO |
| 11 | MONATE | 25 | MORO |
| 13 | MERGOZZO | 25* | VIVERONE |
| 14 | AVIGLIANA | 27 | GHIRLA |
| 14* | LUGANO | 27* | VARESE |
| 15 | COMO | 28 | ALSERIO |
| 15* | ORTA | 29 | SARTIRANA |
| | | 30 | MANTOVA |



EUROPE

Barents Sea

Norwegian Sea

400 mi

400 km

Atlantic Ocean

North Sea

Volga

Ural Mountains

Shannon

Baltic Sea

Thames

Rhine

Elbe

Oder

Vistula

Don

Volga

Seine

Danube

Dnieper

Bay of Biscay

Loire

Po

Danube

Black Sea

Caspian Sea

Tagus

Rhone



Marine sites

Mediterranean Sea

ASIA

AFRICA





Marine sites

Ellobiopsidae

<i>Ellobiocystis caridarum</i>	<i>P. pacifica</i>	<10–84.0	Auke Bay, Alaska	54
<i>Ellobiopsis chattoni</i>	<i>Calanus finmarchius</i>	0.3	Loch Striven, Scotland	45
	<i>Metridia longa</i>	5.0–22.4	Kachemak Bay, Alaska	57
	<i>Undinula vulgaris</i>	8.3	Bay of Bengal, India	50
	<i>U. vulgaris</i>	26.0	Zanzibar Channel off Africa	90



Freshwater sites

- Lake Midmar, South Africa**
- Lake Maggiore, Italy**
- Lake Michigan**
- Small inland lakes in Michigan**

Invasive animals established in the **Great Lakes** drainage since the mid-**1980s** originate mainly from the Ponto-Caspian area

Common name	Year of Discovery	Endemic region	Mode of transfer	Probable donor region
Ruffe	1986	Ponto-Caspian	Ballast water	Danube River
Zebra mussel	1988	Ponto-Caspian	Ballast water	Baltic Sea
Quagga mussel	1989	Ponto-Caspian	Ballast water	Black Sea
Rudd	1989	Eurasia	Bait release	--
Round goby	1990	Ponto-Caspian	Ballast water	Black Sea
Tubenose goby	1990	Ponto-Caspian	Ballast water	Black Sea
New Zealand mudsnail	1991	New Zealand	Ballast water	Baltic Sea
Blueback herring	1995	Atlantic, N.A.	Canal	Atlantic N.A.
<i>Echinogammarus</i> amphipod	1994	Ponto-Caspian	Ballast water	Baltic Sea
<i>Acineta noticrae</i> ciliate	1997	Eurasia	Ballast water	Black Sea
<i>Cercopagis</i> waterflea	1998	Ponto-Caspian	Ballast water	Baltic Sea
<i>Daphnia lumholtzi</i>	1999	Africa, Asia, Aust.	Fish ?	Ohio Reservoirs
Schizopera borutzkyi	1999	Ponto-Caspian	Ballast water	Danube River
Heteropsyllus nr. nunni	1996	Atlantic N.A.	?	Atlantic N.A.
Sphaeromyxa sevastopoli protis	1994	Black Sea	Ballast water	Black Sea
Scolex pleuronectis cestode	1994	Black Sea	Ballast water	Black Sea
Ichthyocotylurus pileatus trematode	1994	Black Sea	Ballast water	Black Sea
Nitocra incerta copepod	1999	Ponto-Caspian	Ballast water	Black Sea
3 testate rhizopod spp.	2003	Ponto-Caspian	Ballast water	Eurasia

is it possible that the more recent records from freshwater sites are the result of an invasion from the marine sites? Like reported for other invasions?

Further investigations on a broader scale are needed, to verify this hypothesis

Mechanisms of international transport for these parasites:

- Infected copepods moving in ballast water?
- Spores carried on air currents?
- Other dispersal mechanisms?



Thank you for your attention