Implications of hyperparasitism for studies into the myxosporean infections of oligochaetes





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Background:

While examining myxosporean infected oligochaetes, Janiszewska (1957) noted that when co-infections with other parasites occurred the myxosporean infection was notably 'slight'. Here I report on a novel hyperparasitic microsporidian affecting an Aurantiactinomyxon-type myxosporean and discuss how such infections could affect experimental results.



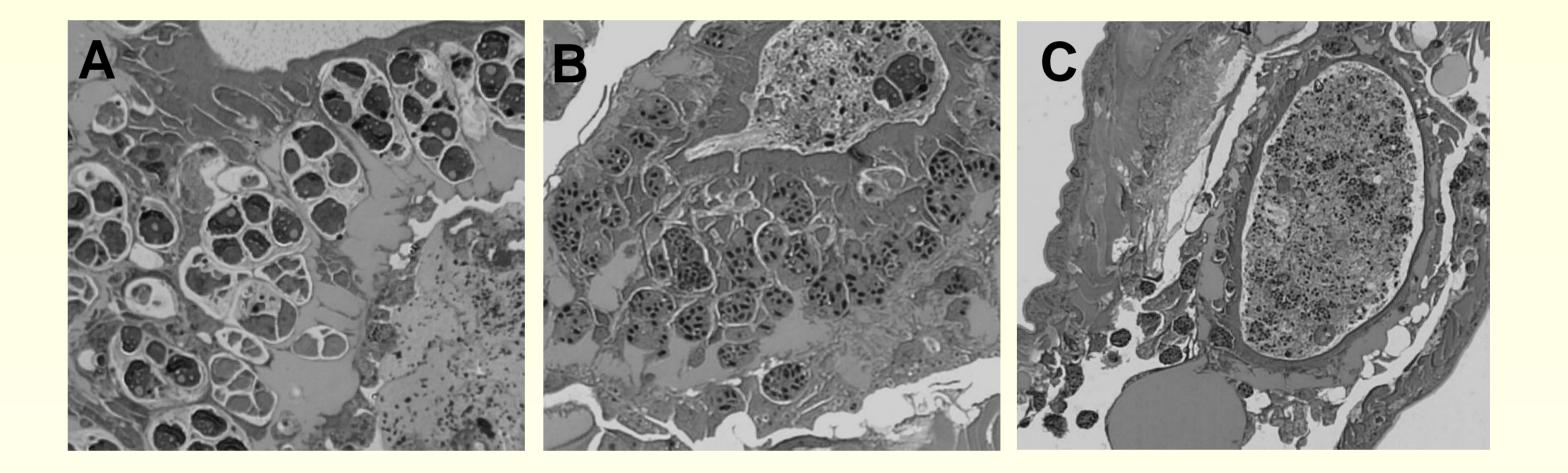
Materials and Methods:

Sediment was collected from a trout pond in Scotland. It was passed through graded sieves to remove silt and the remainder placed into a bucket prior to sorting. The material was placed into shallow dishes and oligochaetes removed to a small beaker. They were then placed individually into cell wells and examined for actinospore production using an inverted microscope.

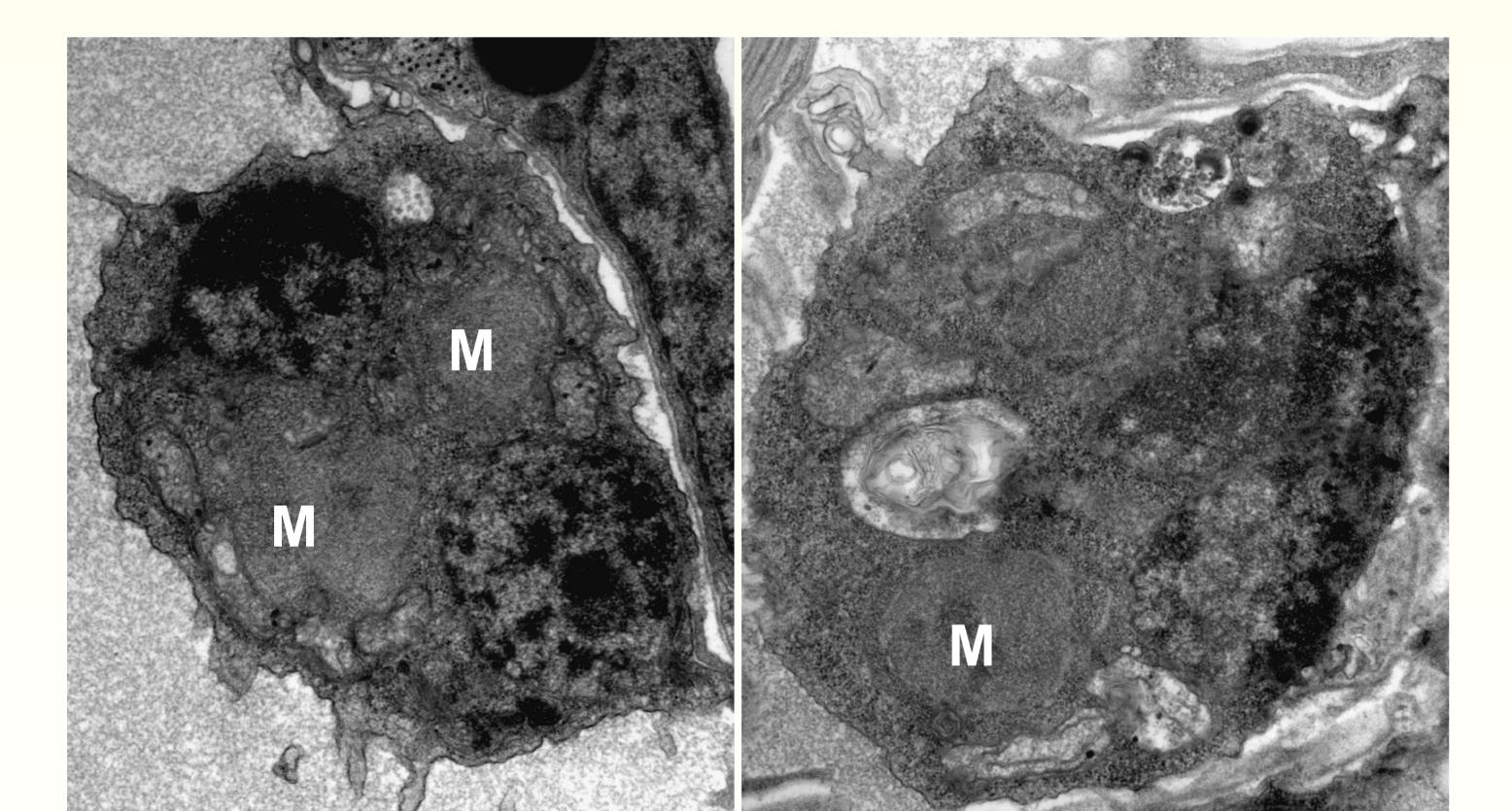
Infected worms were fixed 7,14 and 28 days post-sorting and examined ultrastructurally.

Results:

An Aurantiactinomyxon-type myxosporean was found infecting 6 of 672 oligochaetes, as determined by actinospore release. All of these worms were co-infected with a Infected binucleate stage containing mature spores migrating through tissue to lumen of intestine.



hyperparasitic microsporidian. This parasite infected the binucleate stages* of the myxosporean. At 7 days postsorting all worms were releasing large quantities of actinospores, but the microsporidian infection was progressive, eventually stopping all actinospore release in the worms by 28 days post-sorting.



Progression of infection post-sorting
A) 7 days - many developing actinospores present
B) 14 days - most myxosporeans infected with microsporidian
C) 28 days - no actinospore production

Conclusions:

At least one microsporidian exists that hyperparasitises myxosporeans within oligochaetes. This infection reduces and stops actinospore development.

Current myxosporean study techniques, which pool oligochaetes and/or monitor actinospore release, could be compromised. Especially if confounding infections have a <u>direct</u> life cycle.

Literature reports on instances of



*see poster 83

Funded by

Sudden actinospore cessation. Microsporidian-like 'schizogonic' stages present in samples. Difficulty replicating transmission studies Myxosporean 'susceptible' *Tubifex* becoming 'resistant' after exposure to 'resistant' lineages. Large spatial/temporal shifts in actinospore production

All of these results can be explained by confounding infections such as the microsporidian reported here.

BRS

Acknowledgement

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Reference:. Janisewska, J. (1957) *Zoologica Poloniae*. 8. 3-34.