

Single Cell Analysis Reveals a New Microsporidia-Host Association in a Freshwater Lake

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1	Single cell analysis reveals a new Microsporidia-host
2	association in a freshwater lake
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23 Abstract

24

Microsporidia are a large group of obligate intracellular eukaryotic parasites. Recent studies 25 26 suggest that their diversity can be huge in freshwater lake ecosystems especially in the <150 µm 27 size fraction. However, little is known about their hosts and therefore their impact on the trophic food web functioning. In this study, single cell analysis and fluorescence microscopy were used 28 29 to detect new host-parasite association within rotifer communities in lake Aydat (France). Our 30 analysis showed the existence of a potential new species belonging to the Crispospora genus able of infecting the rotifer Kellicottia with a high prevalence (42.5%) suggesting that 31 Microsporidia could have a great impact on the rotifer populations regulation in lakes. 32 33 **Keywords** 34

35 Microsporidia, host-parasite association, single cell, TSA-FISH, lake, microbial ecology

Microsporidia are obligate intracellular eukaryotic parasites related to Fungi. They are mostly 37 38 known to parasitize a broad range of hosts among the animal kingdom in different environments 39 [1]. Recent metabarcoding studies revealed an unexpected diversity in lake ecosystems especially in the <150 µm size fraction suggesting that Microsporidia could have an impact on 40 the planktonic communities at the base of the food web [2, 3]. However, they have rarely been 41 42 described as parasites of protists and microzooplankton. In marine environments, they are 43 mainly known to parasite gregarines (e.g. [4]), but they were also shown in association with Paramyxea [5]. In freshwaters, only a few observations showed Microsporidia parasiting 44 45 ciliates with only two microsporidian species characterized by morphological and molecular 46 approaches (Euplotespora binucleata and Globosporidium paramecii parasitizing the ciliates Euplotes woodruffi and Paramecium aurelia, respectively [6, 7]). Among microzooplankton, 47 Microsporidia have been observed in the rotifers Brachionus calicyflorus as well as in 48 49 Asplanchna priodonta and A. brightwellii in both river and freshwater lakes [8, 9] however these observations (no molecular data) remain sporadic and for most at low prevalence. 50 51 Moreover, with co-occurrence network and microscopic observations, a more recent study 52 suggested that Microsporidia could infect a broader range of rotifers [3]. We therefore used, in 53 this study, single cell analysis as well as Tyramide Signal Amplification - Fluorescent In Situ 54 Hybridization (TSA-FISH) to detect new host-parasite association within microzooplankton communities in a lake ecosystem (lake Aydat, Massif Central, France) and especially (i) 55 confirm the hypothesis raised about the broader range of rotifers parasitized by Microsporidia 56 57 and (ii) characterize by molecular information and microscopic observation both partners.

Zooplankton was sampled with vertical tows of 20 μ m-mesh plankton net at the central point of the lake (depth = 14 m) on the 18/05/2021 (see Supplementary Material). 17 rotifers, identified morphologically as probably belonging to the genera *Kellicottia* (n = 13) and *Keratella* (n = 4), were isolated by micromanipulation and individually placed in a 96 wells

plate before Whole Genome Amplification (WGA) (see Supplementary Material). The Single 62 Amplified Genomes (SAGs) were then screened for the presence of Microsporidia by the 63 amplification of the V1-V3 region of the microsporidian gene coding for the SSU rRNA 64 according to Chauvet et al. [3]. The sequences obtained after cloning and sequencing were 65 compared to those available in public databases and included in a phylogenetic tree (see 66 67 Supplementary Material). This analysis showed that on the 17 isolated rotifers, 6 Kellicottia 68 had a positive PCR amplification signal with Microsporidia primers however no Microsporidia amplification was recorded in Keratella. All Microsporidia sequences were affiliated to the 69 70 genus Crispospora (Fig. 1) which is affiliated to the clade IV2; Arlequin group (defined in 71 Dubuffet et al. [2]), and well represented (~ 50% of the total OTUs) in metabarcoding studies 72 conducted in lake Aydat [2, 3]. With 97% identity with Crispospora chironomi (the only known species of the genus so far) these sequences could represent a new species of the Crispospora 73 74 genus (Fig. S2a). Indeed, although thresholds for species delineation may vary, an identity close 75 to 100% is suggested for taxa separation at the species level for Microsporidia [2, 11]. 76 Crispospora chironomi is known to parasite chironomids eggs [12] that can be found in aquatic 77 environments especially when they are eutrophic and anthropized [13].

On the Microsporidia positive SAGs, rotifers were characterized by amplification of the COI using the 30F and 885R primers set according to Zhang *et al.* [14] with some modifications (see Supplementary Material). The rotifer affiliation analysis revealed that this potential new species of *Crispospora* is able to parasite the genus *Kellicottia* (Fig. S1). Even though *Kellicottia* sequences grouped into two clades in the phylogenetic analysis, species determination is challenging due to high intraspecific variation of identity percent which can be higher than 20% for rotifers (Fig. S2b) [15].

To visualize and quantify Microsporidia-rotifer associations, TSA-FISH was performed on the rest of the plankton net sample, using a Microsporidia specific probe (USP01) according to

Chauvet et al. [3] (see Supplementary Material). Positive signals were obtained in Kellicottia 87 88 rotifers with a surprisingly high prevalence (42.5%, total rotifers observed = 73) (Fig. 2). Microsporidia were so far recorded in a small number of rotifers [8, 9]. Microsporidium sp. and 89 Microsporidium asperospora have been observed in Brachionus calicyflorus, and 90 Microsporidium polygona in Asplanchna priodonta and A. brightwellii in freshwater 91 92 ecosystems most of the time at low prevalence (e.g. Brachionus calvciflorus by Microsporidium 93 sp. on 1.2% of the host population, [9]). However, these descriptions were only morphological and neither the site of infection nor the microsporidian spore or intracellular form were 94 95 described. Here, we identified an infection which could be in the rotifer's stomach or 96 pseudocoelom. The intracellular forms of the Microsporidia were on average 1.53 µm long, most being oval, some spherical. These findings were consistent with the described shape of 97 the only known Crispospora species (i.e. Crispospora chironomi), with two shapes identified: 98 99 short oval and spherical, with a size of $1.5 \times 2.5 \mu m$ and $1.5 \times 2 \mu m$, respectively [12].

This finding is highly relevant since Microsporidia, along with hyphomycetes and oomycetes 100 101 [17, 18] could be an important regulator of these populations in lakes. Rotifers are indeed an 102 important component of plankton in aquatic ecosystems playing fundamental role in both 103 microbial loop and classical food web and as a link in energy flow between primary producers 104 (algae, cyanobacteria) and higher trophic levels (zooplankton, shellfish, fish) [16]. Kellicottia 105 can reach 600 ind./l in lake Aydat and is the most abundant rotifers along with Keratella [10]. Microsporidia sequences retrieved during this study had an identity >99% with an highly 106 107 abundant OTU identified in the same lake in a previous study where the temporal dynamic of 108 the Microsporidia diversity was analyzed over one year [3] (Fig. S2a). This Crispospora OTU 109 and the only OTU affiliated to Kellicottia presented the same pattern of temporal variations 110 (data not shown). This co-variation and the high prevalence of this association suggest an impact on the dynamic of *Kellicottia* in the lake however this needs to be confirmed for example 111

- by the culture of both partners which will also allow the characterization of the developmental
- 113 cycle of this potential new species of Microsporidia.

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186 Competing Interests

187 The authors declare they have no competing interests.

188

189 Availability of data and material

190 All data and material are available in this publication and additional information online

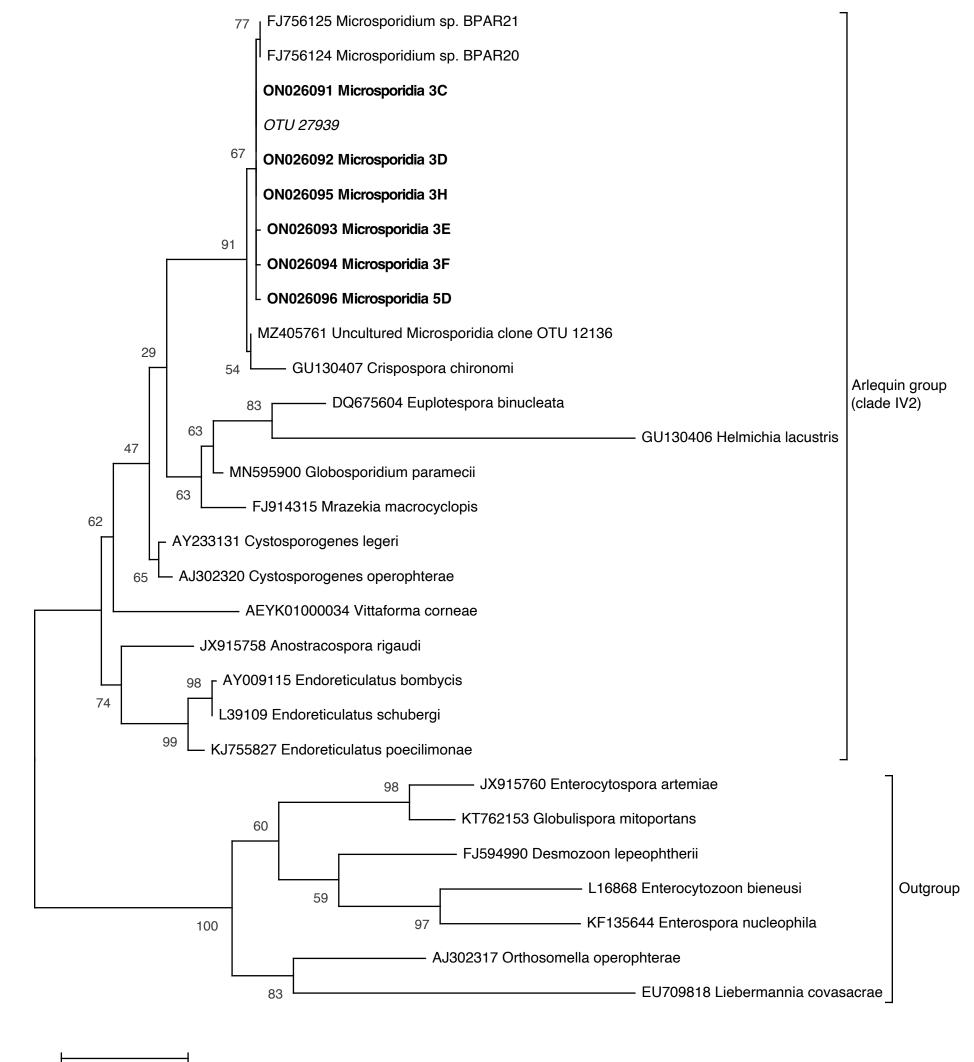
- 191 (supplementary material and supplementary figures). Sequences are available on GenBank,
- accession numbers: from ON026091 to ON026096 for the microsporidian sequences and from
- 193 ON033621 to ON033624 for the rotifers.

194 Figure Legends

195

196	Fig. 1 Maximum Likelihood phylogeny of microsporidian sequences.
197	We included in this phylogeny several sequences from undescribed microsporidian species
198	(found through BLAST search) and a sequence found extensively in a previous study (in italic,
199	[3]), in addition to described species, all belonging to the Arlequin group (clade IV2).
200	Sequences from this study are given in bold. Outgroup species are representative species of
201	other groups belonging to clade IV2. Numbers next to branches indicate the ML bootstrap
202	support (100 replicates).
203	
204	Fig. 2 Microsporidia (targeted by TSA-FISH with the USP01 probe, [3]) infecting a rotifer
205	(genus Kellicottia).
206	Under UV showing nucleus after DAPI staining (a), under blue light showing Microsporidia in
207	green fluorescence (b), overlay (c). Magnification on microsporidian cells inside a rotifer with
208	a confocal microscope (UV (d), blue light (e), overlay (f)).

209 Scale bars, 20 µm



0.10

