New type of xiphidiocercariae (Digenea: Microphalloidea) from South Vietnam

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Abstract: We found unusual digenean intramolluscan stages, sporocysts and cercariae, in gastropods Sulcospira dautzenbergiana (Morelet) (Caenogastropoda: Pachychilidae) from Southern Vietnam and named them Cercaria cattieni 1. These cercariae have a stylet and thus belong to the Xiphidiata. However, such combination of characters as extremely large body size and I-shaped excretory bladder has not been found before in any other xiphidiocercariae. We obtained COI, ITS1, 5.8S + ITS2, and 28S rDNA sequences for C. cattieni 1. The latter allowed us to specify the phylogenetic position of the discovered cercariae: C. cattieni 1 falls within the superfamily Microphalloidea and is most closely grouped to Pachyspondilus irroratus (Rudolphi, 1819) (Pachyspondilidae), the sea turtle parasite. Information on the family Pachyspondilidae is limited. Judging from the molecular phylogeny, C. cattieni 1 might be the larva of the Pachyspondilidae, documented for the first time.

Keywords: Digenea, xiphidiocercariae, sporocysts, Xiphidiata, Microphalloidea, Pachyspondilidae, 28S rDNA, ITS1, ITS2, COI

The contemporary diversity of the Digenea comprises around 18,000 species (Bray 2008), but most of these are known only from sexual adults (maritae). Some data on the life cycle are available for 5–10% of the species (Cribb et al. 2003). Stages in the intermediate hosts are greatly understudied, and for many taxa (even at the family level) the first intermediate host remains unknown, as well as structure of cercariae and parthenogenetic generations. Since early 1990’s molecular data have been a powerful tool to recognize species and link life cycle stages of digeneans, and together with morphological descriptions these methods are very helpful to elucidate digenean diversity (Blasco-Costa and Poulin 2017).

During examination of the fauna of digenean larvae in freshwater gastropods in the Cát Tiên National Park, Southern Vietnam (11.4305N, 107.4294E). Snails were dissected to detect digenean infection. Sporocysts and cercariae found were rinsed in freshwater, fixed in 70% ethanol and later transferred to 96% ethanol for further morphological and molecular analyses. Video of living cercaria was made with a dissecting microscope and a camera Canon A720.

For morphological description sporocysts and cercariae were stained either with Ehrlich’s haematoxylin or Heidenhain’s iron haematoxylin. Picric acid was used to destain worms after Heidenhain’s haematoxylin, and 70% ethanol with 0.1 M HCl after Ehrlich’s haematoxylin. Toluidine blue was used to identify mucoid substances that give metachromatic staining. After staining, samples were dehydrated in graded alcohols and mounted in Bio-Mount medium (Bio Optic, Milan, Italy).

Photographs of the whole-mounted cercariae and sporocysts were made using a compound microscope Leica DM 2500 (Leica Microsystems, Wetzlar, Germany) and a camera Nikon DS F1i in bright field and with differential interference contrast microscopy (DIC). Measurements were made in Fiji software (Schindelin et al. 2012). All measurements are in micrometres the range of values is followed by mean in parentheses.
For molecular analysis two sporocysts were removed from 96% ethanol and dried; DNA was extracted from each of them separately by incubation in 200 μl 5% Chelex® 100 resin (BioRad, Hercules, California, USA) solution with 0.2 mg/ml proteinase K at 56 °C overnight; 8 min at 90 °C; and centrifugation at 16,000 g for 10 min. Supernatant containing DNA was then transferred to a new tube and stored at -20 °C. Amplifications were performed in 25 μl reaction mixtures containing 17 μl Super-Q® water, 5 μl ScreenMix-HS reaction mix (Evrogen, Moscow, Russia), 0.5 μl of each forward and reverse primer (10 pmol/μl), and 2 μl of the DNA template. PCR products were stained with SYBR® Green (Invitrogen, Carlsbad, California, USA), size-separated by electrophoresis in a 1% agarose gel, and visualised using ChemiDoc MP (BioRad, USA). Primers that we used to amplify and sequence the 28S rDNA, ITS1, 5.8S rDNA+ITS2 and COI are listed in Table 1. Sequencing was performed on ABI PRISM 3500xl (Applied Biosystems, Foster City, California, USA).

Data were analysed using Geneious® 11.1.5 (https://www.geneious.com). BLAST was used for preliminary assessment of new sequences. Relevant 28S rDNA sequences from GenBank were used in alignments and phylogenetic reconstructions. The best substitution model was determined as GTR+I+G with BIC in jModelTest 2.1.10 (Darriba et al. 2012). MrBayes v.3.2.6 (Ronquist et al. 2012) was used to build a tree via the Bayesian inference method. *Brachycladium goliath* (van Beneden, 1858) (family Brachycladiidae) (KR703279) served as an outgroup. To estimate genetic distances, we used a Maximum Composite Likelihood model in MEGA 7 (Kumar et al. 2016); standard errors were obtained by a bootstrap procedure (1,000 replications).

**RESULTS**

Of 41 *S. dautzenbergiana* dissected, one snail was infected with daughter sporocysts containing unidentified xiphidiocercariae – *Cercaria cattieni* 1.

**Superfamily Microphalloidea Ward, 1901**

*Cercaria cattieni* 1

Fig. 1. *Cercaria cattieni* 1. A – sporocyst; B–F – cercariae; B – body structure; C – general view; D, E – stylet, ventral (D) and lateral (E) view; F – scheme of cercarial embryo with mucoid glands (grey).
Sporocysts white, elongate, 799–1,628 × 291–369 (1,124 × 336), contain cercariae at different stages of development.

Cercariae (measurements based on 21 fixed specimens; Figs. 1B–E, 2B–I):
- Body 481–758 (594) long, 285–460 (369) wide, pear-shaped, narrow at fore end. Hindbody wide, with folded margins. Tegument with simple spines, most densely arranged in forebody; largest spines in midbody. Tail simple, 603–841 (697) long, 69–101 (93) wide at base, with smooth tegument. Oral sucker subterminal, spherical, 69–103 × 66–102 (86 × 85). Its rim armed with simple spines thicker than body spines, their size reduces posteriorly (Fig. 2D). Stylet 43–52 (48) long, 5–8 (7) wide at base, single-pointed, with anterior thickening (Figs. 1D,E, 2G). Ventral sucker spherical, 142–196 × 140–204 (160 × 165); 4–5 irregular rows of spines (thicker than body spines) lie at sucker rim, around opening, surrounded by smooth zone with 6 large papillae (probably, sensory receptors) (Fig. 2E).

Prepharynx short; pharynx 28–47 × 20–41 (37 × 28); oesophagus very short. Caecal primordia composed of single row of cells (Fig. 2F), 3–6 (4) wide, extending nearly to level of excretory vesicle. Paired cephalic ganglia conspicuous, posterior to oral sucker, interconnected by dorsal commissure between pharynx and oral sucker. Dorsal and ventral pairs of longitudinal nerve chords visible along whole body. Penetration glands numerous, in forebody, anterolateral to ventral sucker (Fig. 2F). Ducts of penetration glands proceed along dorsal surface of oral sucker and open as two groups, lateral to stylet pouch (Fig. 2G,H).

Excretory vesicle I-shaped, 164–232 × 34–67 (189 × 47). Main collecting ducts starting from excretory vesicle anterolaterally (Fig. 2I). Flame-cell formula not determined. Primordia of testes symmetrical, 12–15 (13) in diameter, anterolateral to excretory vesicle. Cirrus sac primordium anterior to ventral sucker. Primordia of uterus, Laurer’s channel and ovary also visible (Fig. 2I). Cercariae swim with their body contracted, thus taking saucer-shape; elongated tail beats vigorously (Supplementary material).

Mucoid glands found in cercarial embryos as four pairs of cells, anterior, lateral and posterior to ventral sucker (Fig. 1F, 3A). Later these cells form numerous dendrites remaining close to ventral surface (Fig. 3B, C). At a certain stage, part of mucoid transferred into region of stylet pouch (Fig. 3D). Next, metachromatic staining disappears from stylet pouch, being found on cercarial surface (Fig. 3E). Mucoid from rest of glands transferred into body tegument, mostly in middle third of body (Fig. 3F–H).

We obtained sequences of three fragments of the rDNA and the mitochondrial COI gene for C. cattieni 1 (Table 1). None of these sequences had any close BLAST hits. Preliminary 28S rDNA-based phylogenetic analysis (covering
a broad range of taxa within the Xiphidiata) suggested focusing the dataset on the Microphalloidea and particularly the Renicolidae. The final alignment included 40 sequences and was 1,226 bp long, including 39 gaps. The resulting phylogenetic tree (Fig. 4) featured two major clades corresponding to the Microphalloidea and Plagiocrioceridae. One of the lineages within the Microphalloidea included the families Renicolidae, Eucotylidae, the unknown family with unidentified cercariae from Nigeria (KX022508 and KX022509, Awharitoma and Enabulele 2018), and also *C. cattieni* forming a well-supported clade with *Pachypsolus irroratus* (Rudolphi, 1819) (family Pachypsolidae) (AY222274).

**DISCUSSION**

Tropical fauna of digeneans is apparently much understudied and no wonder that some findings may be strikingly unlike anything previously described. Among digeneans found in freshwater gastropods of South-East Asia (Sewell 1922, Ito et al. 1962, Ditrich et al. 1992, Dechruksa et al. 2007, Jayawardena et al. 2011, Besprozvannyk et al. 2013, Chontananarth and Wongsawad 2013, and others) nothing similar to *Cercaria cattieni* has been reported. Survey of the literature considering digenean larvae in other geographical regions and other types of environment (marine, terrestrial) produced the same result. According to the classification of Lühe (1909), *C. cattieni* has to be placed into the armate type of xiphidiocercariae. However, such combination of characters as I-shaped excretory bladder, extremely large body size and advanced primordium of the reproductive system is very unusual and has not been described previously for any xiphidiocercariae.

The process of mucoid formation in *C. cattieni* has similarities with other xiphidiocercariae. Numerous dendrites of mucoid glands were previously described in *Cercaria longistyla* McCoy, 1929 by Kruidenier (1953). Four pairs of mucoid glands were found in various xiphidiocercariae (Galaktionov and Malkova 1994, Shchenkov 2012, Shchenkov et al. 2019).

According to the results of the molecular genetic analysis, *C. cattieni* certainly belongs to the Microphalloidea, and within this superfamily to the clade comprising the Pachypsolidae, Eucotylidae and Renicolidae with high nodal support. Close relationship between *C. cattieni* and *Pachypsolus irroratus* gives us a clue to the systematic position of these cercariae. The family Pachypsolidae was established for the only genus *Pachypsolus* Looss, 1901 by Yamaguti (1958) with sexual adults inhabiting sea turtles and freshwater crocodilians (Blair 2008). No life cycles of the Pachypsolidae have been elucidated. In the digenean phylogeny inferred by Olson et al. (2003) this family is a sister group to the Renicolidae + Eucotylidae, and our data confirm this general topology.

The member of the family Pachypsolidae appears as the closest relative of *C. cattieni* so far. Some of the morphological characteristics support similarity of *C. cattieni* cercariae and adults of *Pachypsolus* to a certain degree. The shared features are the tegument with spines, position of suckers, short oesophagus, position of testes, ovary and cirrus sac, and I-shaped excretory vesicle (Blair 2008; present data). However, they differ in the body shape and digestive system: species of *Pachypsolus* lack a prepharynx and their caeca are inflated and have anterior diverticula. These comparisons should be treated critically because allometric growth during the development from cercaria to sexual adult may substantially affect morphology.

![Fig. 3. Toluidine blue staining of *Cercaria cattieni*. A – cercarial embryo, lateral view, early stage of mucoid glands (mg) formation; B – cercarial embryos at different stage, metachromatic staining shows mucoid glands; C – later cercarial embryo (tail lost during fixation), metachromatic staining of mucoid glands with numerous dendrites; D – mucoid staining of stylet pouch; E – mucoid on the surface of the fore region; F, G – mucoid in the midbody, beneath musculature and tegument; ventral nerve chords (vnc) are visible among mucoid; H – midbody, mucoid transferred to the tegument. os – oral sucker; vs – ventral sucker; t – tail. A, D, E, G, H – differential interference contrast.](image-url)
We suppose that the new type of xiphidiocercariae *C. cattieni* is related or even belongs to the *Pachypo-*

solidae. Another option is that it forms a separate branch within the clade comprising the Pachypo-

solidae, Eucotyli-

dae and Renicolidae. When sequences from more taxa are added to the tree in future, its topology, including the posi-

tion of *C. cattieni*, may change. Recent discovery of new lineages within the Microphalloidea that cannot be placed in any existing family (Awharitoma and Enabulele 2018, Shchenkov et al. 2020) along with our data suggest that current understanding of the diversity in this superfamily is very limited, even at the high taxonomic level. To better resolve the position of *C. cattieni*, more molecular data on the Pachypsolidae are required.

**Supplementary material.** The video of living cercariae is available at DOI: 10.13140/RG.2.2.10264.62726

**Acknowledgements.** The authors are grateful to Frank Koehler (Australian Museum, Sydney) for his help with host identification. A.G. was supported by the research programme of the Zoological Institute of the Russian Academy of Sciences (project number AAAA-A19-119020690109-2). We thank the research resource centre “Molecular and Cell Technologies” of Saint Petersburg State University for providing sequencing facilities.

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Cite this article as: Krupenko D., Gonchar A., Kremnev G., Efeykin B., Krapivin V. 2020: New type of xiphidiocercariae (Digenea: Microphalloidea) from South Vietnam. Folia Parasitol. 67: 033.