

# Effect of light on the trematode Himasthla elongata: from cercarial behaviour to infection success

Simao Correia, Rosa Freitas, Xavier de Montaudouin, Luisa Magalhaes

## ▶ To cite this version:

Simao Correia, Rosa Freitas, Xavier de Montaudouin, Luisa Magalhaes. Effect of light on the trematode Himasthla elongata: from cercarial behaviour to infection success. Diseases of Aquatic Organisms, 2021, 146, pp.23-28. 10.3354/dao03616 . hal-04187299

# HAL Id: hal-04187299 https://hal.science/hal-04187299

Submitted on 28 Aug 2023  $\,$ 

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

- 1 Light effect on the trematode *Himasthla elongata* From cercarial behaviour to
- 2 infection success
- 3 Simão Correia<sup>1</sup>, Rosa Freitas<sup>1</sup>, Xavier de Montaudouin<sup>2</sup>, Luísa Magalhães<sup>1\*</sup>
- <sup>4</sup> <sup>1</sup> CESAM, Departamento de Biologia, Universidade de Aveiro, Campus Universitário
- 5 de Santiago, 3810-193, Aveiro, Portugal
- <sup>6</sup> <sup>2</sup> Université de Bordeaux, CNRS, EPOC, EPHE, UMR 5805, Station Marine, 2 rue
- 7 du Pr Jolyet, F-33120 Arcachon, France
- 8 Running page head:
- 9 \*Corresponding author:
- 10 Tel.: +351 234 370 350; fax. +351 234 372 587
- 11 E-mail address: <u>luisa.magalhaes@ua.pt</u>
- 12 Address: Universidade de Aveiro, Departamento de Biologia, Campus Universitário
- de Santiago, 3810-193 Aveiro, Portugal.
- 14

#### 1 ABSTRACT

Cerastoderma edule (cockle), a socio-economically important bivalve of the 2 3 northeast Atlantic, is host to several trematodes, including Himasthla elongata. In the 4 complex life cycle of this trematode, cercariae (free-living stages) emerge from the first intermediate host, a snail, to infect cockles as second intermediate host. During 5 their short lifespan (i.e.less than one day), cercariae have to ensure a successful 6 host-to-host transmission using the surrounding water as transference medium and 7 8 therefore, they are exposed and impacted by different environmental conditions, including abiotic factors. Being light:dark cycle one of the major drivers of aquatic life 9 behaviour, this study aimed to determine light influence on cercaria and their second 10 host behaviour based on two hypotheses. By having a benthic second intermediate 11 host, cercariae will display a photonegative orientation and, on the other hand, the 12 13 host behaviour will not be influenced by light conditions. Results showed cercariae display a photopositive orientation (first hypothesis rejected), displaying movements 14 towards light. The second host behaviour (evaluated by oxygen consumption) was 15 similar among conditions, i.e. dark vs. light (hypothesis accepted), but acquired more 16 parasites when experimentally infested in the dark treatment. This host light-17 18 dependent infection can be explained by a change on cercarial behaviour when exposed to a light stimulus that increased their infection success. This study 19 highlights trematode responses to external conditions may be focused on the life 20 cycle successful completion rather than altered by the host habitat. It was 21 emphasized the light influence on cercarial behaviour resulting in an increased 22 infection success that may be decisive on trematodes population dynamics and 23 distributional range. 24

### 1 KEYWORDS: Cerastoderma edule; Parasitism; photosensitivity; dark: light

2 cycle; oxygen consumption;

#### 1 1. INTRODUCTION

2 Trematodes are prevalent macroparasites in coastal waters (Lauckner 1983) 3 that display a heteroxenous life cycle (infecting up to three hosts) with alternation 4 between parasitic and free-living stages (Bartoli & Gibson 2007). Cercariae, one of these free-living stages, develop by asexual multiplication inside the first 5 intermediate host, usually a mollusc, from which they are emitted and display a short 6 lifespan (< 48 hours) to infect the following host (de Montaudouin et al. 2016). These 7 larvae form an essential component of the ecosystem, representing a high fraction of 8 marine biomass (Lambden & Johnson 2013) and production (Thieltges et al. 2008). 9 Cercariae can exert a strong influence on the structure, dynamics and function of 10 food webs, altering their topology or lineage density (Dunne et al. 2013). Cercarial 11 activity is strongly influenced by environmental abiotic conditions, among which light 12 13 can be determinant. While studying cercariae from different trematode species, authors have been demonstrating their photosensitivity, either moving towards or 14 away from light. Cercariae that are photosensitive, have developed photoreceptors 15 (or proteins with analogous functions) to facilitate light perception (Haas 1992). The 16 development of such responses might be related to the trematode life cycle and the 17 18 habit of the hosts involved. Indeed, when the trematode downstream host is a benthic organism, cercariae usually remain near the sediment (positive geotaxis) and 19 usually respond negatively to a light stimulus, moving away from it (Combes et al. 20 1994). Additionally, cercarial behaviour can display a synchronized activity with the 21 daily active period of its host (Combes et al. 1994, Esch et al. 2001, Combes et al. 22 23 2002). Nonetheless, there are several behavioural patterns that can be combined to influence the cercarial performance (Morley 2020). For example, there are other 24

external factors, such as the force of gravity, that can act together with light on
 defining cercarial swimming preferences (Kennedy 1979, Combes et al. 1994).

3 Cercariae of trematode species presenting three-host life cycles transform into metacercariae after penetrating the second intermediate host. The pathological 4 effects of this trematode parasitic stage on the host depend on the number of 5 accumulated cysts (Desclaux et al. 2004) as well on environmental abiotic factors 6 such as pollutants (de Montaudouin et al. 2010, Paul-Pont et al. 2010). This complex 7 8 interaction between host-metacercariae dynamics and environmental conditions have already been reported to cause host mortality (Desclaux et al. 2004, Thieltges 9 2006). 10

The knowledge about behavioural responses of trematode cercariae towards 11 abiotic conditions (such as light) is still scarce, especially in what concerns the 12 13 consequences of such responses on infection success and subsequent modulation of host population dynamics. In the present study, the trematode species Himasthla 14 elongata and its second intermediate host, the bivalve Cerastoderma edule (edible 15 cockle) were used as host-parasite model. This trematode uses the common 16 periwinkle, Littorina littorea, an intertidal gastropod of the medio- and supralittoral 17 18 zone (Eschweiler et al. 2009), as first intermediate host, and infects the second intermediate host, the infaunal intertidal bivalve Cerastoderma edule, by inhalation 19 (Wegeberg et al. 1999). The present study aimed to experimentally assess the 20 influence of light (dark vs. light exposure) on the cercarial behaviour, cercariae 21 infection success and cockle susceptibility to infection. The hypotheses were: (1) due 22 23 to the second intermediate host endobenthic habit, cercariae of H. elongata will display a photonegative orientation response; (2) cockles filtering activity will not be 24

- 1 influenced by light and (3) cercarial performance and infection success will be
- 2 influenced by light.

#### 1 2. MATERIAL AND METHODS

2

#### 2.1. Organisms collection and maintenance

3 Littorina littorea snails were previously collected in Texel, The Netherlands, and exposed to a temperature boost to incite cercariae emergence. Afterwards, the 4 trematode metacercariae was identified to species level through morphological 5 analysis by infecting cockles and following de Montaudouin et al. (2009). Infected 6 snails were kept in laboratory in an aquarium with artificial seawater (salinity 35), at a 7 temperature of  $14 \pm 1$  °C and with a natural photoperiod (12:12 hours light/dark). 8 During the maintenance period, snails were fed with Ulva sp. ad libitum. To obtain 9 cercariae, infected snails were transferred to individual containers with artificial 10 seawater (salinity 35) and exposed to a temperature boost (approximately 24 °C) 11 and constant illumination (approximately 6 hours). Cercariae were then counted 12 under a stereomicroscope and collected with a micropipette. 13

Individuals of *C. edule*, ranging from 13 to 17 mm shell length, were collected in the Mira channel of the Ria de Aveiro coastal lagoon, Portugal. Cockles were transported to the laboratory and acclimated for one week at controlled conditions of salinity (30), temperature (18 °C) and photoperiod (12:12 hours light/dark) and were fed with Algamac Protein Plus ® by Aquafauna, a heterotrophic and phototrophic species mixture of macroalgae, at a concentration of 730 cells  $\mu$ L<sup>-1</sup> day<sup>-1</sup> (Pronker et al. 2015).

21

22

#### 2.2. Cercarial behaviour experiment

To study cercarial photosensitivity, a pre-designed oval shape aquarium (50 x 30 x 5 cm, 10 cm channel width) was used to create a light gradient (Appendix 1). The aquarium was built with half of transparent acrylic and half of black opaque acrylic. The aquarium was filled with artificial seawater at salinity 35, maintained at 18 °C and kept in a dark room exposed to a horizontal light source while covered by a black opaque lid so the only light input was horizontally. Light was supplied by a 7volt, 1.5 watts LED bulb during the whole experiment. Maximum light intensity obtained was 50  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup>, similar to light intensities observed in clear shallow waters (dos Santos et al. 2015). Light intensity was measured with an Apogee handheld MQ-200 Quantum meter with a separate sensor.

In each experiment (same experiment was performed in duplicate), a total of 8 9 1000 cercariae (maximum: 6 hours old) were released in two different releasing areas (areas D and E, corresponding to mid position and intermediate light intensity). 10 After 4 hours, cercariae were assigned to 8 different areas of similar volume and 11 counted through image analysis. Three digital pictures per area were obtained using 12 a high resolution photographic camera and cercariae were counted manually and 13 using OpenCFU software. The percentage of cercariae present in each area was 14 then calculated. Area A was exposed to maximum light intensity (50  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup>), 15 while area H was exposed to minimum light intensity (dark – 0  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup>). The 16 remaining six areas were exposed to equal light intensity in a two-by-two pattern. 17 Areas B and C were exposed to a light intensity of 35  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup>, areas D and E to 18 23  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup> of light intensity and areas F and G to a total light intensity of 6  $\mu$ mol 19  $m^{-2} s^{-1}$ . 20

A previous study performed with a trematode parasite of the same family revealed that cercariae mobility tend to decrease severely after 12 hours (de Montaudouin et al. 2016). Therefore, in the present study, a total 10 hours of lifespan (6 hours after emission + 4-hour exposure) was selected to ensure cercariae mobility until the end of the experiment.

2

#### 2.3. Cockles oxygen consumption

3 To test the light effect on cockles filtering activity, oxygen consumption used as a proxy of this activity (Wegeberg et al. 1999), was measured through simple 4 static respirometry. After acclimation period, twelve cockles of similar shell length 5 (between 15 and 17 mm) were individually placed in 1L respirometric chambers. All 6 chambers were filled to their maximum capacity with artificial seawater at salinity 30, 7 to avoid formation of bubbles. In order to allow cockles acclimation to the 8 respirometric chamber conditions, individuals were left for 1 hour inside the 9 respirometric chamber. Six of these chambers were covered in aluminium foil, 10 preventing the entrance of light, and the other six were exposed to natural light 11 condition (approximately 25  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup>). In addition, four other respirometric 12 chambers (two covered in aluminium foil and two uncovered) were used as control, 13 i.e. no organisms to account for background oxygen consumption. Each chamber 14 was equipped with an oxygen sensor. Oxygen concentration was measured twice, in 15 16 equal time intervals (1 hour), by contacting the oxygen sensor with a multi-channel optic fiber oxygen meter (Multi channel oxygen meter, PreSens, GmbH, 17 Regensburg, Germany) and reading the concentration value after stabilization in the 18 PreSens Measurement Studio software. 19

Oxygen consumption was calculated as the difference between 2 reading points (separated by a period of two hours) and expressed in ppmO<sub>2</sub> gram of cockle fresh weight<sup>-1</sup> h<sup>-1</sup>. Cockles fresh weight was calculated using the shell length: weight relationship previously determined for the area where cockles were sampled (S. Correia unpubl. data) and described as:

25  $\log W = 3.1886 \log L + 3.7577, R^2 = 0.92$ 

1 Where, *W* is cockles fresh weight, *L* is the shell length (in mm) and  $R^2$  is the 2 coefficient of determination.

3

4

2.4. Cercariae infection success

After cockles acclimation period, an experiment using twenty-four replicates 5 was performed in duplicate. Previously, a subsample of 15 cockles were dissected 6 and observed under a stereomicroscope to obtain natural infection intensities at the 7 8 sample site. To minimize uncontrolled effects, similar sized cockles (ranging from 13 to 17 mm), were placed separately into 50 mL glass flasks at two different conditions 9 (light vs. dark) in a dark room. Twelve cockles were exposed to a vertical light source 10 supplied by a 7-volt, 1.5 watts LED bulb with a light intensity of 50  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup>. The 11 flasks of the other 12 cockles were covered in aluminium foil and exposed to dark. 12 During this experiment, water medium was maintained at 18 °C and salinity 30. 13

A total of 600 cercariae, 25 per flask, were collected and used to individually infect each cockle. Experiments lasted for 48 hours, the time required for cercariae to encyst (de Montaudouin et al. 2016). After, metacercariae of *H. elongata* were counted under a stereomicroscope by squeezing cockles' flesh between two glass slides.

19

#### 20 2.5 Data analysis

21 Concerning cercarial behaviour experiment, two chi-squared tests were 22 performed, one to check the homogeneity among experiment duplicates and the 23 second to test the data adjustment between the expected and the observed 24 frequencies of cercariae per area.

For cockles oxygen consumption, a Mann-Whitney U test was used to assess the differences in terms of oxygen consumption among conditions (light vs. dark), whereas for cercariae infection success experiment, a Kruskal-Wallis H test followed by Dunn-Bonferroni post hoc test were performed to compare cockles infection between light conditions and natural infection (light vs. dark vs. natural infection).

All non-parametric statistical analyses were performed using the IBM SPSS
Statistics software v.25 after testing the parametric statistical assumptions (Levene's
and Shapiro-Wilk tests for homoscedasticity and normality, respectively) and failing
samples normality.

10

#### 11 3. RESULTS AND DISCUSSION

12 3.1. Cercarial behaviour experiment

After 4 hours exposure, 83.2 % and 81.8 % of total released cercariae were 13 retrieved from each duplicate of the cercarial behaviour experiment. Cercariae of 14 Himasthla elongata displayed a photopositive orientation behaviour. From the total 15 cercariae found, 80.3 % ± 2.1 standard deviation (SD) were detected in area A, 16 corresponding to the highest light intensity (50  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup>). Area H (0  $\mu$ mol m<sup>-2</sup> s<sup>-1</sup>), 17 was identified as the area with the lowest percentage of cercariae abundance (3.0 % 18  $\pm$  0.9 SD of total cercariae retrieved). The remaining cercariae were found in the 19 areas located between maximum light intensity and dark. In detail, areas B and C 20 presented 9.3 % ± 0.8 SD of cercariae final count, while 3.8 % ± 0.4 SD were found 21 in the areas D and E. In the areas F and G, areas positioned closer to the dark 22 condition, the number of cercariae found corresponded to 3.8  $\% \pm 0.8$  SD of the final 23 count. The chi-squared test showed that the cercariae proportion retrieved by area 24 did not follow a uniform distribution ( $\chi^2$  (7) = 8412.72; p < 0.001), i.e. there were 25

significantly more cercariae in some areas compared to others. Cercarial behaviour 1 showed to be the same among experiment duplicates ( $\chi^2$  (7) = 8.444; p = 0.295). 2 Since H. elongata first host, Littorina littorea, is in an upstream (supralittoral) and 3 therefore more lighted area compared to cockles, the second host, it was 4 hypothesized a photonegative attraction by cercariae to optimize cockle infection. 5 6 However, the behavioural pattern found, i.e. the cercariae movement towards light, allows to reject the postulated hypothesis. These results were the opposite found by 7 8 other authors for H. elongata (Prokofiev 2006) and H. rhigedana (Fingerut et al. 2003), possibly due to a stronger geotactic response that supress cercarial 9 photosensitivity or, in the case of H. rhigedana, due to its second host 10 11 (Pachygrapsus crassipes) higher activity at night (Morgan et al. 2006). In fact, H. elongata cercarial orientation towards light may still be related to the host habit, in 12 this case, its intertidal habitat. While being attracted by light, the cercariae of H. 13 elongata can settle in a water column with higher light incidence and lower water 14 15 column height, characteristics of an intertidal area. Coupled with the positive 16 geotactic behaviour that this trematode species already demonstrated (Nikolaev et al. 2017), the orientation positively influenced by light can promote the chances to 17 infect cockles. This behaviour has already been described for other cercariae 18 species infecting intertidal hosts (e.g. Kennedy 1979). Accordingly, higher trematode 19 infection levels of cockles from intertidal areas compared to subtidal ones have 20 already been described in the literature (Correia et al. 2020). At the same time, 21 Himasthla spp. are also natural parasites of mussels (Galaktionov et al. 2015), which 22 inhabit in a higher position of the littoral zone, and therefore a photopositive 23 attraction can benefit the infection of these hosts, i.e. Himasthla spp. could be more 24 25 adapted to infect upstream positioned hosts.

2

#### 3.2. Cockles oxygen consumption and cercariae infection success

Cockles oxygen consumption was not significantly different between light conditions (Z(16) = -1.435, p = 0.151). In fact, it is already described in the literature that if they are submersed, cockles can filter actively regardless light conditions (Newell & Bayne 1980).

Results of the present study showed that when cockles were not exposed to 7 light, they displayed higher infection levels (mean infection = 14.29 metacercariae 8  $cockle^{-1} \pm 4.14$  SD) compared to cockles exposed to a vertical light source (mean 9 infection = 10.16 metacercariae cockle<sup>-1</sup>  $\pm$  4.34 SD) and the sampling site natural 10 infection (mean infection = 2.07 metacercariae cockle<sup>-1</sup>  $\pm$  2.15 SD) (*H*(2) = 33.751, *p* 11 > 0.001). Differences between groups were confirmed by post hoc tests (Light (L) vs. 12 Natural conditions (NC): Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX; p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. NC: Z = XXXX, p < 0.001; Dark (D) vs. (D) vs. NC: Z = XXXXX, p < 0.001; Dark (D) vs. 13 0.001; L vs. D: Z = -2.598, p = 0.009). Cockles were experimentally infested in low 14 volume containers to promote host-parasite contact. However, given that no 15 differences were found in the oxygen consumption of cockles exposed to light 16 compared to those maintained in dark conditions, and that cercariae of Himasthlidae 17 18 family (family of the studied trematode species) are passive invaders, i.e. cercariae infect the second intermediate host by being ingested through their feeding activity 19 (Wegeberg et al. 1999, Galaktionov & Dobrovolskij 2003), the results lead to infer 20 that recorded infection differences are rather dependent on cercarial behaviour 21 towards distinct light conditions than on cockles individual filtering activity. Indeed, 22 when exposed to a light source, cercarial movements can be enhanced (Haas 1992) 23 which, due to the absence of water current in the experimental containers used, 24 could have led to a rise in the water column. Additionally, the low water column 25

height could have neglected the geotactic effect. Consequently, these vertical 1 2 movement decreased the chance of contact, departing cercariae from the feeding 3 area of the second intermediate host. On the other hand, in the absence of a light stimulus, cercariae swim without orientation, slowly sinking in the water column 4 (Feiler & Haas 1988, Haas et al. 1990). In this way, the cercariae tended to approach 5 the feeding area of its host, increasing the chance of contact and consequent 6 infection. When in close contact with the host, the previously described passive 7 8 behaviour of cercariae shifts to an inciting active infection (Galaktionov & Dobrovolskij 2003), increasing the probability of infection, thus explaining the results 9 obtained. Nonetheless, further studies should be taken into account to fully 10 understand the effect of depth. 11

The cercarial behaviour reported in this study allows *H. elongata* to use light as a guidance mechanism to disperse throughout the aquatic system and seek for a suitable habitat for its life cycle completion, promoting the infection of cockles more susceptible to be predated by the final host (shorebirds). Meanwhile, during night, with lower light incidence, cercariae would save their energies, sinking to its second host active area, waiting to be inhaled by cockles.

18

19

#### 4. CONCLUSION

In conclusion, the present study showed that by displaying a photopositive sensitivity, cercariae of *Himasthla elongata* can increase their dispersion ability, invading the most suitable habitats where the second intermediate host lives. It is important to highlight that this study clearly demonstrated a higher infection success in cockles exposed to dark compared to cockles maintained in the light. This higher infection can only be ascribed to cercarial behaviour since host filtering activity is not

affected by light. These results support the theory that trematode responses besides 1 2 being dependent on the second intermediate host type of habitat (benthic vs. pelagic 3 habitat) are also subordinated by the will to complete the life cycle. In this particular case, as the second intermediate host is widely distributed along the whole tidal 4 gradient (from lighter to darker areas) cercariae can use external stimuli (such as 5 light) to relocate themselves towards areas where the second intermediate host is 6 more likely to be predated by its final host (shorebirds in case of the species under 7 8 study) and, therefore be able to proceed the life cycle.

- 9
- 10

#### 11 ACKNOWLEDGMENTS

Simão Correia and Luísa Magalhães benefited from research fellowships 12 13 funded by INTERREG-ATLANTIC program through the research project COCKLES 14 (EAPA\_458/2016 COCKLES Co-Operation for Restoring CocKle SheLlfisheries & its 15 Ecosystem-Services in the Atlantic Area). Rosa Freitas was funded by national funds (OE), through FCT - Fundação para a Ciência e a Tecnologia, I.P., in the scope of 16 the framework contract foreseen in the numbers 4, 5 and 6 of the article 23, of the 17 18 Decree-Law 57/2016, of August 29, changed by Law 57/2017, of July 19. This work was supported by the research project COCKLES (EAPA\_458/2016 COCKLES Co-19 Operation for Restoring CocKle SheLlfisheries & its Ecosystem-Services in the 20 Atlantic Area). Thanks are due to FCT/MCTES for the financial support to CESAM 21 (UIDB/50017/2020+UIDP/50017/2020), through national funds. 22

### 1 REFERENCES

2	Bartoli P, Gibson DI (2007) Synopsis of the life cycles of Digenea (Platyhelminthes)
3	from lagoons of the northern coast of the western Mediterranean. Journal of
4	Natural History 41:1553-1570
5	Combes C, Bartoli P, Théron A (2002) Trematode transmission strategies. In: Lewis
6	EE, Campbell F, Sukhdeo MVK (eds) The Behavioural Ecology of Parasites.
7	CAB International, Wallingford, UK.
8	Combes C, Fournier A, Mone H, Theron A (1994) Behaviors in trematode cercariae
9	that enhance parasite transmission - patterns and processes. Parasitology
10	109:S3-S13
11	Correia S., Picado A., de Montaudouin X., Freitas R., Rocha RJM., Dias JM,
12	Magalhães L (2020). Parasite assemblages in a bivalve host associated with
13	changes in hydrodynamics. Estuarine and Coasts 848
14	de Montaudouin X, Blanchet H, Desclaux-Marchand C, Lavesque N, Bachelet G
15	(2016) Cockle infection by Himasthla quissetensis - I. From cercariae
16	emergence to metacercariae infection. Journal of Sea Research 113:99-107
17	de Montaudouin X, Paul-Pont I, Lambert C, Gonzalez P, Raymond N, Jude F,
18	Legeay A, Baudrimont M, Dang C, Le Grand F, Le Goic N, Bourasseau L,
19	Paillard C (2010) Bivalve population health: Multistress to identify hot spots.
20	Marine Pollution Bulletin 60:1307-1318
21	Desclaux C, de Montaudouin X, Bachelet G (2004) Cockle Cerastoderma edule
22	population mortality: role of the digenean parasite Himasthla quissetensis.
23	Marine Ecology Progress Series 279:141-150

1	dos Santos CDB, Gaudig C, Fritsche M, dos Reis MA, Kirchner F (2015) An
2	evaluation of artificial fiducial markers in underwater environments. OCEANS
3	2015 - Genova, Genova, Italy 1-6
4	Dunne JA, Lafferty KD, Dobson AP, Hechinger RF, Kuris AM, Martinez ND,
5	McLaughlin JP, Mouritsen KN, Poulin R, Reise K, Stouffer DB, Thieltges DW,
6	Williams RJ, Zander CD (2013) Parasites Affect Food Web Structure Primarily
7	through Increased Diversity and Complexity. Plos Biology 11
8	Esch GW, Curtis LA, Barger MA (2001) A perspective on the ecology of trematode
9	communities in snails. Parasitology 123:S57-S75
10	Eschweiler N, Molis M, Buschbaum C (2009) Habitat-specific size structure
11	variations in periwinkle populations (Littorina littorea) caused by biotic factors.
12	Helgoland Marine Research 63:119-127
13	Feiler W, Haas W (1988) Host-finding in Trichobilharzia ocellata cercariae: swimming
14	and attachment to the host. Parasitology 96:493-505
15	Fingerut JT, Zimmer CA, Zimmer RK (2003) Larval swimming overpowers turbulent
16	mixing and facilitates transmission of marine parasite. Ecology 84:2502–2515
17	Galaktionov KV, Dobrovolskij AA (2003) The Biology and Evolution of Trematodes:
18	an essay on the Biology, Morphology, Life Cycles, Transmissions, and
19	Evolution of Digenetic Trematodes, Vol. Kluwer Academic Publishers
20	Galaktionov K., Bustnes JO, Bårdsen BJ, Wilson JG, Nikolaev KE, Sukhotin AA,
21	Skírnisson K, Saville DH, Ivanov MV, Regel KV (2015) Factors influencing the
22	distribution of trematode larvae in blue mussels Mytilus edulis in the North
23	Atlantic and Arctic Oceans. Marine Biology 162:193-206
24	Haas W (1992) Physiological analysis of cercarial behavior. Journal of Parasitology
25	78:243-255

Haas W, Granzer M, Brockelman CR (1990) *Opisthorchis viverrini* - Finding and
 recognition of the fish host by the cercariae. Experimental Parasitology
 71:422-431

Kennedy MJ (1979) The responses of miracidia and cercariae of *Bunodera mediovitellata* (Trematoda: Allocreadiidae) to light and to gravity. Canadian
 Journal of Zoology 57:603-609

- Lambden J, Johnson PTJ (2013) Quantifying the biomass of parasites to understand
   their role in aquatic communities. Ecology and Evolution 3:2310-2321
- 9 Lauckner G (1983) Diseases of Mollusca: Bivalvia. In: Kinne O (ed) Diseases of
   10 Marine Animals, Book 2. Hamburg, Biologische Helgoland, Germany
- Morgan SG, Spilseth SA, Page HM, Brooks AJ, Grosholz ED (2006) Spatial and
   temporal movement of the lined shore crab *Pachygrapsus crassipes* in salt
   marshes and its utility as an indicator of habitat condition. Marine Ecology
   Progress Series 314:271-281
- Morley NJ (2020) Cercarial swimming performance and its potential role as a key
   variable of trematode transmission. Parasitology 144:1369–1374

Newell RIE, Bayne BL (1980) Seasonal changes in the Physiology, Reproductive
 condition and Carbohydrate content of the cockle *Cardium* (= *Cerastoderma*)
 *edule* (Bivalvia: Cardiidae). Marine Biology 56:11-19

Nikolaev KE, Prokofiev VV, Levakin IA, Galaktionov KV (2017) How the position of
 mussels at the intertidal lagoon affects their infection with the larvae of
 parasitic flatworms (Trematoda: Digenea): A combined laboratory and field
 experimental study. Journal of Sea Research 128: 32-40.

Paul-Pont I, Gonzalez P, Baudrimont M, Jude F, Raymond N, Bourrasseau L, Le
 Goic N, Haynes F, Legeay A, Paillard C, de Montaudouin X (2010) Interactive

1	effects of metal contamination and pathogenic organisms on the marine
2	bivalve Cerastoderma edule. Marine Pollution Bulletin 60:515-525
3	Prokofiev VV (2006) Strategies Of The Animal Host Infection With Trematode
4	Cercariae: An Attempt Of Analysis In Marine Coastal And Lake Ecosystems
5	Of Northwestern Russia (Dissertation). Zoological Institute of the Russian
6	Academy of Sciences, St.Petersburg (in Russian).
7	Pronker AE, Peene F, Donner S, Wijnhoven S, Geijsen P, Bossier P, Nevejan NM
8	(2015) Hatchery cultivation of the common cockle (Cerastoderma edule L.):
9	from conditioning to grow-out. Aquaculture Research 46:302-312
10	Thieltges DW (2006) Parasite induced summer mortality in the cockle Cerastoderma
11	edule by the trematode Gymnophallus choledochus. Hydrobiologia 559:455-
12	461
13	Thieltges DW, de Montaudouin X, Fredensborg B, Jensen KT, Koprivnikar J, Poulin
14	R (2008) Production of marine trematode cercariae: a potentially overlooked
15	path of energy flow in benthic systems. Marine Ecology Progress Series
16	372:147-155
17	Wegeberg AM, de Montaudouin X, Jensen KT (1999) Effect of intermediate host size
18	(Cerastoderma edule) on infectivity of cercariae of three Himasthla species
19	(Echinostomatidae, Trematoda). Journal of Experimental Marine Biology and

20 Ecology 238:259-269

1 Figure 1. Schematic representation of cercariae light behaviour aquarium subdivided 2 in the 8 areas of light gradient. A: Side View; B: Top View; C: Front View. Cercariae were released in areas 4 and 5. 3 4 5 Figure 2. Experimental design of the cercariae infection success assay. Each 6 condition contained 12 replicates with 25 cercariae released per replicate. 7 8 Figure 3. Percentage of cercariae found in each experimental area of the behaviour aquarium. Area 1 presented the maximum luminosity, while area 8 was exposed to dark. 9 The remaining areas had an equal luminosity two-by-two. 10









Figure 2





