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# **Annelid polychaetes experience metabolic acceleration as other Lophotrochozoans: inferences on the life cycle of *Arenicola marina* with a Dynamic Energy Budget model**

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1 Annelid polychaetes experience metabolic acceleration as other  
2 Lophotrochozoans: inferences on the life cycle of *Arenicola*  
3 *marina* with a Dynamic Energy Budget model

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9 **Abstract**

10 *Arenicola marina* is a polychaete (Lophotrochozoan) displaying a complex benthopelagic life  
11 cycle with two larval dispersal phases, only partially described up to now. A Dynamic Energy  
12 Budget (DEB) model was applied to the species in order to reconstruct its life cycle and growth  
13 under *in situ* environmental conditions. Two types of DEB models are usually applied to other  
14 Lophotrochozoans displaying similar life cycles: the standard (std-) model, applied to polychaetes  
15 (5 entries among the 1524 of the Add-my-Pet database on the 18/10/2018), and the abj-model,  
16 which includes an acceleration of metabolism between birth and metamorphosis, and which has  
17 been applied to most molluscs (77 abj- entries out of the 80 mollusc entries) enabling better fit  
18 predictions for the early life stages. The parameter estimation was performed with both models to  
19 assess the suitability of an abj-model for *A. marina*. The zero-variate dataset consisted of length  
20 and age data at different life cycle stages, the lifespan, the maximum observed length, and the wet  
21 weight of an egg. The uni-variate dataset consisted of two growth experiments from the literature  
22 at two food levels and several temperatures, laboratory data of oxygen consumption at several  
23 temperatures, and fecundity for different lengths. The predictions of the abj-model fitted better to  
24 the data (SMSE = 0.29). The acceleration coefficient was *ca* 11, which is similar to mollusc values.  
25 The field growth curves and the scaled functional responses (as a proxy of food levels) were suitably  
26 reconstructed with the new parameter set. The reconstruction of the early life-stages chronology  
27 according to *in situ* environmental conditions of a temperate marine ecosystem indicated a first  
28 dispersal phase of 5 days followed by a 7 months temporary settlement before a second dispersal  
29 phase in spring, at the end of metamorphosis. We emphasize the need for using abj-models for  
30 polychaetes in future studies.

31 **Key words**

32 Bioenergetics, lugworm, growth, oxygen consumption, life-history traits, dispersal

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### 33 Introduction

34 *Arenicola marina* (Linnaeus, 1758) is a marine polychaete (Lophotrochozoan, Annelida) inhab-  
35 iting most intertidal soft sediments from the Arctic to the Mediterranean. The species is intensively  
36 dug for bait by recreational fishermen (Blake, 1979; De Cubber et al., 2018; Watson et al., 2017)  
37 and the comparison between harvest efforts and observed populations abundance has evidenced  
38 the need for some regulation of this activity in some places (De Cubber et al., 2018). In aquacul-  
39 ture, *A. marina* is also reared for bait (Olive et al., 2006), and more recently, for its particular  
40 haemoglobin that might represent a valuable blood substitute for humans in the future (Rousselot  
41 et al., 2006) and which is already used for organs conservation before transplantation. However,  
42 its complex benthic-pelagic life cycle with two dispersal phases before recruitment has made the  
43 description of the early life stages and their chronology complicated, and still little is known about  
44 the development of *A. marina* between the trochophore larva stage and the benthic recruitment  
45 (Farke and Berghuis, 1979a, b; Newell, 1948; Reise, 1985). Moreover, the literature regarding *A.*  
46 *marina*'s life cycle and growth is quite ancient (mostly from 1979) and since 1990, it has been found  
47 that two cryptic species actually exist and might live in sympatry: *A. marina* and *A. defodiens*.  
48 Therefore ancient life-history description has to be used with caution (Cadman and Nelson-Smith,  
49 1990).

50 The 'Dynamic Energy Budget' (DEB) theory quantifies the energy allocation to growth and  
51 reproduction of an individual during its life cycle according to environmental conditions such as  
52 temperature and food availability (Kooijman, 2010) even in species with complex and numerous  
53 life-stages (Llandres et al., 2015). Twelve primary parameters are sufficient for the implementation  
54 of a standard (std-) DEB model. However, among the assumptions implied in std-DEB models,  
55 some, like isomorphism during growth, the fact that growth always follows a typical Von Bertalanffy  
56 growth curve, or the presence of three life stages (embryo, juvenile and adult) are not found in every  
57 species. Therefore, extensions of the std-model (implying the use of more parameters) were created  
58 (Kooijman, 2014) accounting for deviations from typical development implied by the std-model,  
59 like foetal development, acceleration of metabolism, or extra life stages.

60 As of October 2018, Add-my-Pet (AmP) database estimated DEB parameters for 1524 animal  
61 species (Marques et al., 2018). Among these entries, only 11 were annelid species, 4 of them being  
62 polychaetes species for which std-models were applied. The closest phylum with a large amount  
63 of data is the molluscs' phylum (over 80 entries), also presenting a larval stage. Indeed, annelids  
64 and molluscs belong to the Lophotrochozoan clade and both, after the embryogenesis, lead to a  
65 trochophore larval stage. Mostly abj-DEB models have been applied only for the mollusc phylum,  
66 which are an extension of std-models considering an acceleration of metabolism between birth (first  
67 feeding) and metamorphosis (end of the change of shape) and are applied to most species with a  
68 larval phase (Kooijman, 2014). Although polychaete species often present a larval phase during  
69 their life cycle, until now, abj-models were not applied to this taxa.

70 A std- entry for *A. marina* is present in the AmP database and enables predictions of the  
71 growth and reproduction of the species. However, more than half of the dataset used for the  
72 parameter estimation consists of unpublished data (time since birth at puberty and maximum  
73 reproduction rate taken from Marlin: <https://www.marlin.ac.uk/> and lifespan and ultimate total  
74 length taken from Wikipedia: <https://www.wikipedia.org/>), guessed data (wet weight at birth and  
75 puberty, ultimate wet weight), or data related to other species (age at birth from *A. cristata* and  
76 *A. brasiliensis*) (AmP entry: Bas Kooijman. 2015. AmP *Arenicola marina*, version 12/07/2015).  
77 We therefore completed the data set with literature, experimental and field data, and implemented  
78 a new parameter estimation for the species using both a std- and an abj-DEB models.

79 The objectives were:

- 80 (1) to calibrate a DEB model for *A. marina* based on a reliable and complete dataset and  
81 adapted to its life cycle features (and therefore to compare the relevance of the use a std- or  
82 an abj-DEB model for this species)
- 83 (2) to make predictions about the chronology of the early life stages of *A. marina* and the growth  
84 potential according to the environmental conditions
- 85 (3) to compare the parameters of the DEB models implemented for *A. marina* with the other  
86 Lophotrochozoan species' parameters and discuss the advantages of the use of an abj-model  
87 for this species.

## 88 1. Material and Methods

### 89 1.1. The DEB theory and its implementation for *Arenicola marina*

#### 90 1.1.1. The model

91 The DEB theory describes the energy flows within an organism between three compartments  
92 (state variables) : the reserve ( $E$ ), the structure ( $V$ ), and the maturity ( $E_H$ ) or the reproduction  
93 buffer (offsprings) ( $E_R$ ) according to its life stage in order to describe its energy allocation to  
94 growth and reproduction for a given food level and at a reference temperature  $T_{ref}$  (Fig. 1). The  
95 three differential equations linked to the state variables are obtained from the expression of the  
96 different fluxes (Table 1)(Kooijman, 2010; Van der Meer, 2006).

97 Temperature corrections are made to the rates considered by the model in the equation of  
98 fluxes (e.g. the surface-area specific maximum assimilation rate,  $\{\dot{p}_{Am}\}$  ( $\text{J}\cdot\text{cm}^{-2}\cdot\text{d}^{-1}$ ), the energy  
99 conductance,  $\dot{v}$  ( $\text{cm}\cdot\text{d}^{-1}$ ), the specific volume-linked somatic maintenance rate,  $[\dot{p}_M]$  ( $\text{J}\cdot\text{cm}^{-3}\cdot\text{d}^{-1}$ ),  
100 and the maturity maintenance rate coefficient,  $\dot{k}_J$  ( $\text{d}^{-1}$ ), see Tables 1 and 4). Indeed, when the  
101 temperature  $T$  (K) is different from the reference temperature  $T_{ref}$  (taken to be 293.15 K) these  
102 rates are multiplied by the correction given in Equation (1), where  $T_A$  is the Arrhenius temperature  
103 (K),  $\dot{k}_1$  the rate of interest at  $T_{ref}$  and  $\dot{k}$  the rate of interest at  $T$ .

$$\dot{k}(T) = \dot{k}_1 \cdot \exp\left(\frac{T_A}{T_{ref}} - \frac{T_A}{T}\right) \quad (1)$$

104 The links between observable metrics (physical length and wet weight) and the DEB model  
 105 quantities are made with the shape coefficient  $\delta$  (varying between  $\delta = \delta_{Me}$  for embryos and  $\delta = \delta_M$   
 106 after metamorphosis), the density of wet structure  $d_V$  ( $g.cm^{-3}$ ), of wet reserve  $d_E$  ( $g.cm^{-3}$ ) and of  
 107 dry reserve  $d_{Ed}$  ( $g.cm^{-3}$ ), the specific chemical potential of reserve  $\mu_{Ed}$  ( $J.Cmol^{-1}$  of reserve), and  
 108 the molar weight of reserve  $w_{Ed}$  ( $g.Cmol^{-1}$ ) (Table 1). Here, we assumed that  $d_V = d_E = 1 g.cm^{-3}$ ,  
 109  $d_{Ed} = 0.16 g.cm^{-3}$ ,  $\mu_{Ed} = 550000 J.Cmol^{-1}$  and that  $w_{Ed} = 23.9 g.Cmol^{-1}$ .

### 110 1.1.2. Adaptation to *Arenicola marina*'s life cycle

111 The spawning event of *Arenicola marina* happens in late summer or early autumn (De Cubber  
 112 et al., 2018; Watson et al., 2000). After the external fertilization, the embryo develops in the female  
 113 gallery up to the post-embryonic stage, the trochophore larva, which is able to move vertically in  
 114 the water column (Fig. 2, Farke and Berghuis, 1979a, b). At this time, the shape is changing  
 115 from ovoidal (oocytes, with a shape coefficient  $\delta = \delta_{Me}$ ) to cylindrical when the trochophore larva  
 116 gradually acquires new setiger becoming a metatrochophore larva (of shape coefficient  $\delta < \delta_{Me}$ ).  
 117 The metatrochophore larva is released in the water column when it reaches 3 setigers and is  
 118 transported by currents during several days. In lugworms, embryos and larvae are lecithotrophic,  
 119 living on maternal reserve and therefore supposed not to be able to feed (the maturity threshold  
 120  $E_H$  did not yet reach its value for birth:  $E_H < E_H^b$ , where birth is the time when individuals start  
 121 to feed). Therefore, there is no feeding or assimilation flux during the embryo and larval stage and  
 122  $dE/dt = -\dot{p}_C$  (Table 1). Moreover, these young stages do not have enough complexity yet to be  
 123 able to produce gametes and the reproduction flux goes to maturity (Table 1), which represents in  
 124 this case the acquisition of complexity of the individual (Fig. 1).

125 The metatrochophore larva settles and begins eating as post-larva, when the gut appears func-  
 126 tional ( $E_H = E_H^b$ ), either on mussel beds, macroalgae or sheltered soft sediment bottoms (Fig.  
 127 2). At this point, it lives inside a mucus tube stuck to the bottom and feeds on the particles  
 128 deposited on the tube and around it, as well as on suspended particles (Farke and Berghuis, 1979a,  
 129 b; Newell, 1949; Reise, 1985; Reise et al., 2001). During this temporary settlement period, the  
 130 post-larva continues to gradually acquire new setigers up to the 19 final setigers found in adults  
 131 (the shape coefficient  $\delta$  keeps on decreasing until it reaches the shape coefficient value of the adults  
 132  $\delta_M$ ), developing a proboscis in the way of the adults (Farke and Berghuis, 1979a, b; Newell, 1949).  
 133 These morphological changes are assimilated to metamorphosis (up to when the maturity threshold  
 134  $E_H$  reaches its value at metamorphosis:  $E_H = E_H^j$ ). During this period, a metabolic acceleration  
 135 (Kooijman, 2014) was considered, which is supposed to happen in most species that have a lar-  
 136 val phase, frequently coinciding with morphological metamorphosis (Marques et al., 2018), and  
 137 resulting in an exponential growth of the organism between the first feeding and the end of meta-  
 138 morphosis. From birth ( $E_H = E_H^b$ ), feeding and assimilation are not null anymore, but individuals  
 139 are not able yet to produce gametes ( $\dot{p}_R = 0$ ).

140 When metamorphosis ends, a second dispersal phase of unknown period occurs in the water

141 column and the newly juvenile lugworm settles on intertidal areas colonized by adults' lugworms,  
142 where it changes its mode of nutrition, becoming psammivorous like the adults (Beukema and De  
143 Vlas, 1979)(Fig. 2). The shape coefficient value stops changing, the growth starts to be isomorphic  
144 and follows the Von Bertalanffy growth curve for a constant scaled functional response (Kooijman,  
145 2010), but it is not yet able to reproduce like the adults (since the maturity threshold  $E_H$  did not  
146 reach its value at puberty yet:  $E_H < E_H^p$ ).

147 Finally, the adults acquire the ability to reproduce (which is when  $E_H > E_H^p$ ) and the energy  
148 flow formerly allocated to maturity is transferred to a reproduction buffer (offsprings) that empties,  
149 in the case of *Arenicola marina*, once a year in early autumn, during the spawning event.

## 150 1.2. Compilation of data for *Arenicola marina* and parameter estimation

### 151 1.2.1. Zero-variate and uni-variate data from the literature

152 *Zero-variate data from the literature.* An important part of the zero-variate dataset found in the  
153 literature was composed of data taken from a larval culture performed by Farke and Berghuis  
154 (1979) before 1990, when the two species *Arenicola marina* and *A. defodiens* were not yet delimited  
155 (Cadman and Nelson-Smith, 1993): the lengths at trochophore larva, at birth (first feeding) and at  
156 metamorphosis with their associated ages (Table 5). Although the lengths data seem quite accurate  
157 (plates and pictures), the chronology description made by the authors remains vague. The precise  
158 time line had thus to be estimated from sometimes quite confused date references and we gave a  
159 weight of 0.5 to this data in the parameter estimation procedure. In the larval culture performed  
160 by Farke and Berghuis (1979), the temperature varied from 8 to 16 °C, so a mean temperature of  
161 12 °C was used for the data taken from this experiment.

162 The second part of the zero-variate dataset from the literature was collected after 1990. First,  
163 the age for the occurrence of the trochophore larva at 10 °C was communicated by S. Gaudron  
164 from unpublished *in vitro* fertilization experiments. The maximum observed trunk length (good  
165 biometric estimate, see De Cubber et al., 2018) was observed by S. Gaudron on a specimen kept  
166 in the Animal Biology Collection of the Sorbonne University (France). Finally, the age and length  
167 at puberty, the oocyte diameter and the lifespan were previously acquired by the authors at the  
168 same study site (De Cubber et al., 2018). The temperature used for this data was the mean  
169 temperature of the seawater over the year 2017 (13 °C, SOMLIT data: [http://somlit-db.epoc.u-](http://somlit-db.epoc.u-bordeaux1.fr/)  
170 [bordeaux1.fr/](http://somlit-db.epoc.u-bordeaux1.fr/), bottom coastal sampling point at Wimereux). The age and the trunk length at  
171 puberty corresponded to a first mature adult of 2.5 cm and 1.5 years old. All the age data estimated  
172 from length analysis were given a weight of 0.5 in the parameter estimation procedure considering  
173 their potentially low accuracy. For all zero-variate data the  $f$  value was set to 1, considering that  
174 only the "best individuals" were used.

175 *Uni-variate data from the literature.* The uni-variate dataset retrieved from the literature consisted  
176 in the datasets of two growth experiments:

- 177 - One growth experiment in which trunk length was measured at four different temperatures  
178 (5, 10, 15 and 20 °C) under two different food conditions (fed and unfed) taken from De  
179 Wilde and Berghuis (1979) (8 treatments). The corresponding  $f$  values were set at  $f_{fed} = 0.8$   
180 and  $f_{unfed} = 0.1$  in view of growth comparisons made by the authors in the same study.
- 181 - One growth experiment in which wet weight was measured at one temperature varying be-  
182 tween 16 and 20°C under two different conditions (fed and unfed) taken from Olive et al.  
183 (2006) (2 treatments). Temperature was set at 19.5°C and the  $f$  values were left free for both  
184 conditions.

185 For these two growth experiments, the temperature and feeding conditions met before the start  
186 of the experiment were not known so we had to assume the levels of reserve and structure at the  
187 beginning of the experiment. Therefore, predictions of growth could only be made considering a  
188 physical trunk length  $TL_w(0)$  at the beginning of the experiment and a physical wet weight  $W_w(0)$   
189 at the beginning of the experiment equalling to the one of the experiment.

#### 190 1.2.2. Laboratory experiments and field data

191 Additional reproductive data (reproduction rate as a function of trunk length and wet weight  
192 of an egg), growth data (trunk length over time) and oxygen consumption data (oxygen consump-  
193 tion as a function of wet weight) were acquired by the authors in the laboratory and from field  
194 observations between 2016 and 2018 in order to complete the dataset collected from the literature.

195 *Study area and sampling strategy.* Lugworms were collected at Wimereux (N 50°46'14" and E  
196 01°36'38"), Le Touquet (N 50°31'07" and E 01°35'42") and Fort Mahon (N 50°20'31" and E  
197 01°34'11"), located in the Eastern English Channel (Hauts-de-France, France)(Table 2). More  
198 details on the sites are given in De Cubber et al. (2018). For the oxygen consumption experiment  
199 (Exp. A), the lugworms were collected at Wimereux from the high mediolittoral to the high in-  
200 fralittoral part of the foreshore (Fig. 2), in order to collect all the different age groups and sizes  
201 (De Cubber et al., 2018), on the sandy beach part, using a shovel. Collection happened three times  
202 between May and July 2018 in order to follow the summer increase of the seawater temperature  
203 of the English Channel (Table 2). For the reproductive data (Exp. B), ripe females of *A. marina*  
204 were collected at Wimereux, Le Touquet and Fort Mahon using a shovel or a bait pump (Decathlon  
205 ltd.) during the spawning period of each year (Table 2). For the growth experiment (Exp. C),  
206 young individuals of *A. marina* were collected at Wimereux on the high mediolittoral part of the  
207 foreshore with a shovel (De Cubber et al., 2018) at the end of May 2018 (Table 2, see more details  
208 in De Cubber et al., 2018).

209 *Laboratory measurements.* After each sampling, all lugworms were put in separate containers filled  
210 with seawater. Individuals of *Arenicola marina* were maintained in the laboratory during 24 h at  
211 the temperature of the English Channel at Wimereux at the time of their collection (12, 15 and

212 20.5 °C) for the oxygen consumption experiment (Exp. A), and at 15 °C otherwise (Exp. B and  
213 C), in a cold room, to allow gut to be devoided of their content prior to observations (Watson et  
214 al., 2000). Biometric measurements consisted in total length, trunk length (more reliable, see De  
215 Cubber et al., 2018 and De Wilde and Berghuis, 1979), and in wet weight measurements.

216 *Experiment A: Oxygen consumption.* The oxygen consumption rates of lugworms were recorded  
217 as a proxy of metabolic activity (Galasso et al., 2018). Metabolic rates can vary between two  
218 fundamental physiological rates, one minimal maintenance metabolic rate (the standard metabolic  
219 rate) and one maximum aerobic metabolic rate (the active metabolic rate) (Galasso et al., 2018;  
220 Norin and Malte, 2011). In order to recreate these two situations of activity in the laboratory, and  
221 avoid any over- or underestimation of the metabolic rate, the oxygen consumption of lugworms was  
222 measured under two different conditions in which their metabolic activity was supposed close to  
223 the standard metabolic rate on one hand, and close to the active metabolic rate on the other hand.  
224 In the condition in which lugworms were supposed to experience a standard metabolic rate, around  
225 30 of the collected individuals were transferred into Eppendorfs or Falcon centrifuge tubes (5 ml  
226 or 50 ml according to the size of the worms) half-filled with sand from Wimereux burnt at 550°C  
227 during 5 h, and with twice-filtered seawater (TFSW, 0.45 µm and 0.22 µm), enabling the lugworms  
228 to burry. The sediment was well mixed before the transfer in order to avoid air bubbles inclusions  
229 between sediment grains. In the condition in which lugworms were supposed to experience an active  
230 metabolic rate, around 30 of the collected individuals were transferred into centrifuge tubes filled  
231 with TFSW only, where they were constantly trying to burry (no sand). Blanks were also made  
232 for both conditions (centrifuge tubes without lugworms). Lugworms were acclimatized 24 hours  
233 at the experimental temperature in order to allow them to burrow when possible and relax. For  
234 each condition, centrifuge tubes were oxygenated using an air pump, and refilled with oxygenated  
235 TFSW when needed. At this point, lugworms in the "active" condition experienced regularly extra  
236 stress due to water movements. The oxygen content was then measured using a microelectrode  
237 Unisense® OX500 coupled to a picoammeter (Unisense PA 2000, Denmark). The data acquisition  
238 was performed using the software InstaCal® and the tubes were then rapidly hermetically closed  
239 with Parafilm® M. For the 50 ml centrifuge tubes, measurement was renewed three times every 10  
240 to 15 minutes after opening the Parafilm® M lid for a few seconds and homogenizing the water.  
241 For the 5 ml tubes, only two measurements were made at the beginning of the experiment and  
242 after 1 h given the low oxygen consumption observed. Before every measurement series, the whole  
243 system was calibrated (measurements of 100% and 0% oxygenated TFSW) and the salinity of the  
244 TFSW used for the experiment was measured using a refractometer. The temperature of the cold  
245 room was followed throughout the duration of the experiment. After the experiment, lugworms  
246 from the sand condition were sieved out of their tubes and maintained 24 h to allow gut contents  
247 to be devoided prior to biometric measurements. All lugworms were then measured (trunk length  
248 and total length) and weighed (wet weight).



249 *Experiment B: Reproductive data.* All oocytes were collected, from females that had been previ-  
250 ously weighted and measured, in a 60  $\mu\text{m}$  sieve, rinsed with TFSW and placed in a 5 ml Eppendorf  
251 tube filled with TFSW (Table 2). A triplicate of 20  $\mu\text{L}$  of the homogenized solution were then put  
252 on a microscope slide and the oocytes were counted under the microscope. When fecundity was  
253 estimated for each female, the supernatant was removed and the Eppendorf tubes were weighted  
254 with and without oocytes.

255 *Experiment C: Growth experiment.* The growth experiment lasted for two months in a controlled  
256 room (temperature, photoperiod) at the Wimereux Marine Station (University of Lille, France)  
257 under a recirculating custom seawater system (Fig. 3). In the custom system, one aquarium  
258 tray was dedicated to water filtering and two aquaria held the lugworm growing experiment. The  
259 seawater, directly pumped from the sea, was kept several days in the filtering aquarium containing  
260 fine and coarse filter foam, crushed pozzolana and oyster shells and kept in the dark (Fig. 3).  
261 10% of the seawater contained in the two growing aquaria was renewed every day or every second  
262 day with the water of the filtering aquarium. Two external filters (Eheim professional 4+ 250)  
263 and pumps allowed the circulation and additional filtration of the seawater system (Fig. 3a). A  
264 lightening system consisting in two light ramps (Alpheus Radiometrix 13C1001C) mimicking the  
265 external light intensity and photoperiod was added to the system (Fig. 3a), air pumps (Air pump  
266 8000 and Eheim 400 from Europrix ltd., not represented on Fig. 3) linked to home-made finely  
267 punctured pipes allowed the oxygenation of the system. The temperature was kept around 15°C ( $\pm$   
268 1°C). Each of the two growing aquarium trays were holding each twice 3 boxes filled with sediment  
269 burnt at 60°C during 24 h and lugworms (Fig. 3b). The first 3 boxes closer to the seawater arrival  
270 were dedicated to the unfed condition, the next 3 to the fed condition. A small waterfall between  
271 them prevented the seawater (and food) to circulate in the opposite direction of the main current,  
272 thus no food could reach the unfed condition. The design of the boxes and of the separations  
273 prohibited the worms to leave their box and to circulate from one condition to another condition  
274 (Figs. 3a, b). All lugworms were measured and only individuals ranging from 0.4 cm to 1.6 cm of  
275 trunk length were selected. Twelve batches of 30 individuals were made with the same size (trunk  
276 length) range. Each batch was placed in a separated box within the experimental set up (Fig. 3b).  
277 Feeding occurred twice at  $t = 0$  and  $t = 35$  days with yeast wastes (obtained from Brasserie du  
278 pays Flamand ltd., a local brewery) inserted within the sediment with 20 ml syringes (between 1.8  
279 and  $3.6 \cdot 10^{10}$  cells added per box) (Olive et al., 2006). One batch of lugworms of each condition in  
280 both aquaria was withdrawn at the beginning of the experiment, after 35 days and after 62 days,  
281 kept 24 h in the cold room and weighted and measured.

282 *Data analyses.* All data analyses were performed on Matlab R2015b. For the oxygen consumption  
283 experiment (Exp. A), for each measurement (blanks included), the associated percentage of oxygen

284 within the tube was calculated according to the Equation (2).

$$O_{2 \text{ measured}}(\%) = \frac{O_{2 \text{ measured}}(V) - O_{2 \text{ min}}(V)}{O_{2 \text{ max}}(V) - O_{2 \text{ min}}(V)} \cdot 100 \quad (2)$$

285 With  $O_{2 \text{ measured}}(V)$  the oxygen measured,  $O_{2 \text{ min}}(V)$  the oxygen measured for 0% of oxygen, and  
286  $O_{2 \text{ max}}(V)$  the oxygen measured for 100% of oxygen. The oxygen content ( $\mu\text{mol.L}^{-1}$ ) was then  
287 calculated according to the temperature (T, in  $^{\circ}\text{C}$ ), salinity (S, in  $\text{‰}$ ) and the water content of  
288 each tube according to Aminot and K erouel (2004, see on pages 110-118). The blank effect was  
289 deleted, and the individual oxygen consumption ( $\mu\text{mol.h}^{-1}$ ) was then calculated as the inverse of  
290 the slope of the linear regression of the evolution of the oxygen content over time. Both conditions  
291 were analyzed together to consider an average level of activity.

292 For the reproduction data acquisition, the fecundity (F) was calculated for each female according  
293 to Equation (3) (with n the mean of the three counts).

$$F = \frac{n}{4 \cdot 10^{-3}} \quad (3)$$

294 Since spawning happens only once a year for *A. marina*, the reproduction rate for each female  
295 was calculated as the fecundity divided by the number of days in one year and plotted against the  
296 female trunk length (uni-variate data). The wet weight of an egg was calculated as the total weight  
297 of oocytes divided by fecundity (zero-variate data).

### 298 1.2.3. Parameters estimation

299 The parameters estimation of the DEB models was done using the covariation method described  
300 by Lika et al. (2011), using the dataset shown in Table 3. The estimation was completed using the  
301 package DEBtool (as described in Marques et al., 2018) on the software Matlab R2015b using both  
302 a std-DEB model and an abj-DEB model, in order to select the best fit model and to compare the  
303 parameter obtained with both models.

304 The parameter estimation procedures were evaluated by computing the Mean Relative Errors  
305 (MRE), varying from 0, when predictions match data exactly, to infinity when they do not, and  
306 the Symmetric Mean Square Errors (SMSE), varying from 0, when predictions match data exactly,  
307 to 1 when they do not (<http://www.debtheory.org>).

### 308 1.3. Inferring environmental conditions from biological data and vice versa

#### 309 1.3.1. Functional scaled response associated to growth data

310 The parameters of abj-model for *Arenicola marina* (best fit model), as well as two different  
311 growth datasets, were used to validate the model and infer the environmental conditions (in terms  
312 of food levels) of these datasets. The first growth dataset was taken from Beukema and De Vlas  
313 (1979). It represents seasonal changes in mean individual dry weight (9-year averages) in small  
314 lugworms from two populations of the Wadden Sea. The second dataset consists of the observations  
315 of wet weight and trunk length of the experiment C. Since the results of the latest experiment  
316 seemed to indicate that food was lacking from  $t = 35$  days to  $t = 62$  days and since no significant

317 difference between the two feeding conditions were observed, the abj-model applied in this study  
318 was used to reconstruct the scaled functional response ( $f$ ) as a proxy of food levels during the  
319 whole experiment for the two conditions. Predictions on these different growth experiments were  
320 made at one temperature but for feeding conditions varying from  $f = 0.02$  to  $f = 1$ . The best  
321 fit predictions were chosen as the ones presenting the smallest sum of squares of the differences  
322 between observations and predictions.

### 323 1.3.2. Life cycle chronology under *in situ* environmental conditions

324 The abj-DEB model for *Arenicola marina* was used to reconstruct the chronology of the early  
325 life stages of the species under the *in situ* environmental conditions of Wimereux (Eastern English  
326 Channel, Hauts-de-France), as well as its growth in wet weight and trunk length, and compared  
327 them with optimal food and temperature conditions ( $f = 1$  and  $T = 20$  °C).

328 *Local environmental conditions.* The *in situ* temperature of the year 2017 were taken from SOM-  
329 LIT. As a first approximation, the scaled functional response  $f$  was guessed from monitoring of the  
330 phytoplankton within the Eastern English Channel (Lefebvre et al., 2011) showing higher abun-  
331 dances in spring and autumn, as generally observed in the North Atlantic temperate ocean (Miller  
332 and Wheeler, 2012, Fig. 11.7).

333 *Chronology of the early life-stages and associated lengths.* The parameters of the abj-model pre-  
334 viously estimated were used to predict age and length at trochophore larva stage, birth, meta-  
335 morphosis and puberty under the non-optimal environmental conditions of Wimereux previously  
336 defined.

337 *Growth predictions.* The evolution of the compartments of reserve, structure and reproduction  
338 buffer from the fertilization to the lifespan  $a_m$  and further was calculated according to the equations  
339 of Table 1. For each environmental condition, the ages for all the life stages were predicted as  
340 previously and a temperature correction was applied when the temperature was different from 20  
341 °C. The values of  $E$ ,  $V$  and  $E_R$  over time were then converted into wet weight and/or physical  
342 trunk length with the equations found in Table 1.

### 343 1.4. Comparison of the DEB parameters of *Arenicola marina* with other Lophotrochozoan species

344 The parameters found with the abj-DEB model for *Arenicola marina* were compared with the  
345 ones found with the std-model, as well as with the parameters of other molluscs and annelid species.  
346 The parameters collected were taken from the Add-my-Pet collection (AmP) (Marques et al., 2018)  
347 using the function prtStat of the AmPtool package used on Matlab R2015b. All values were given  
348 for a reference temperature  $T_{ref}$  of 20 °C. The most complete data set for molluscs is for the  
349 gastropod *Lymnaea stagnalis* (completeness = 5). The maximum completeness value for annelids  
350 in AmP is 2.8, and is found in two species of polychaetes and four species of clitellates. The least  
351 complete data set for molluscs is for the symbiotic bivalve *Thyasira cf. gouldi* (completeness = 1.5)

352 and the least complete data set for annelids is for the polychaete *Capitella teleta* (completeness =  
353 1.5).

354 Some of the primary parameters of the two models for *A. marina* were not compared given the  
355 lack of data for these parameters (e.g. the searching rate  $\{\dot{F}_m\}$ , the digestion and the reproduction  
356 efficiencies  $\kappa_X$  and  $\kappa_R$ ), as in Kooijman and Lika (2014). The acceleration factor  $s_M$  of *A. marina*  
357 was calculated as  $s_M = L_j/L_b$ , with  $L_b$  the structural length at birth and  $L_j$  the structural length at  
358 the end of the metamorphosis, and compared with the one of other species. For the species showing  
359 a metabolic acceleration ( $s_M > 1$ ), the infinite length  $L_\infty$  was calculated as  $L_\infty = L_m \cdot s_M$ , with  $L_m$   
360 the maximum structural length ( $L_m = \kappa \cdot \frac{\{p_{Am}\}}{[p_M]}$ ). The energy conductance after metamorphosis  
361  $v_j$  and the maximum assimilation rate after metamorphosis  $\{p_{Am}\}_j$  were calculated as  $v_j = v_b \cdot s_M$   
362 and  $\{p_{Am}\}_j = \{p_{Am}\}_b \cdot s_M$ , with  $v_b$  the energy conductance at birth and  $\{p_{Am}\}_b$  the maximum  
363 assimilation rate at birth (Kooijman, 2014; Kooijman and Lika, 2014). All ten parameters, as well  
364 as the expectations based on the general animal (Kooijman, 2010, Table 8.1), were represented as  
365 functions of  $L_\infty$  for all the considered species.

## 366 2. Results

### 367 2.1. Parameter estimation

#### 368 2.1.1. Parameters of the model

369 The completeness of the models was set at 4.2 following Lika et al. (2011) according to the  
370 dataset used in the parameters estimation (Table 3). The implementation of the parameter es-  
371 timation of the std-DEB model provided a Mean Relative Error (MRE) of 0.30 and Symmetric  
372 Mean Square Error (SMSE) of 0.38 (Marques et al, 2018). The implementation of the parameter  
373 estimation of the abj-DEB model provided a MRE of 0.23 and SMSE of 0.29. In addition to the  
374 fact that the abj-model provided a better fit to the data set, it appears that the std-model largely  
375 underestimates the age and length at birth,  $a_b$  and  $L_b$  (the relative errors, RE, are respectively  
376 0.91 and 0.83), as well as the age when then trochophore larva appears,  $a_{tr}$ , (RE = 0.60). For both  
377 models, the values of the fraction of the metabolized energy allocated to soma,  $\kappa$ , appeared equal  
378 (Table 4). The specific somatic maintenance rate,  $[p_M]$ , and the maximum assimilation rate at  
379 birth,  $\{p_{Am}\}_b$ , and at metamorphosis,  $\{p_{Am}\}_j$ , were respectively five, twenty and two times higher  
380 with the std-model than with the abj-model. However, the maturation thresholds for the occurring  
381 of the trochophore larva,  $E_H^{tr}$ , for birth,  $E_H^b$ , and for puberty,  $E_H^p$ , and the energy conductance at  
382 metamorphosis,  $\dot{v}_j$ , appeared higher with the abj-model, that considered a metabolic acceleration  
383 rate between birth and metamorphosis,  $s_M$ , around 11 (Table 4).

#### 384 2.1.2. Observations vs predictions

385 *Zero-variate data.* For 9 of the 12 zero-variate observations of the estimation procedure with the  
386 abj-model, the predicted values were close to the observed ones (RE  $\leq$  0.27) (Table 5). The last  
387 three predictions for the age at birth  $a_b$ , the age at puberty  $a_p$  and the total length at birth  $L_b$

388 showed higher relative errors (RE  $\sim$  0.65). The predictions obtained with the std-model estimation  
389 procedure were overall less well adjusted to the zero-variate observations with 50% of the predictions  
390 associated RE higher than 0.45 (Table 5). For instance, the age at birth  $a_b$ , the age when the  
391 trochophore larva is first observed  $a_{tr}$  and the age at puberty  $a_p$  were highly underestimated with  
392 the std-model estimation procedure (RE respectively of 0.91, 0.6 and 0.74), as well as the total  
393 length at birth  $L_b$  and the length when the trochophore larva is first observed  $L_{tr}$  (RE respectively  
394 of 0.83 and 0.45).

395 *Uni-variate data.* The RE of the uni-variate data set ranged from 0.06 to 0.41 with the abj-  
396 DEB model, and from 0.08 to 0.42 with the std-DEB model, with, in both cases, the highest  
397 values corresponding to the fit to the length-weight data collected on individuals of highly variable  
398 reserve and reproduction buffer levels, and to the oxygen consumption data set (most scattered  
399 values) (Figs. 4, 5, 6, 7). In both cases, the oxygen consumption increased with the increase of  
400 temperature (Fig. 4). The values of the shape coefficients  $\delta_M$  varied for *a priori* the same measure  
401 of the trunk length between 0.14 and 0.20 with the abj-DEB model and between 0.09 and 0.13 with  
402 the std-DEB model according to the authors (Fig. 7), which is due to the lack of rigid measurable  
403 parts in *Arenicola marina* that could be used as a proxy for length.

## 404 2.2. Reconstruction of environmental conditions with the abj-model for *Arenicola marina* from bi- 405 ological data and vice versa

### 406 2.2.1. Scaled functional response

407 *From a field growth dataset.* The abj-model provided a good fit for the field growth data taken  
408 from Beukema and De Vlas (1979) for the two studied sites (Fig. 8a). The values of the scaled  
409 functional response  $f$  were shown to evolve on both sites during the year, with the highest values  
410 during spring and late summer periods compared to winter period (Fig. 8a).

411 *From laboratory growth data.* Overall, the abj-model provided a good fit for the growth data  
412 obtained in the laboratory (Exp. C), although growth was slightly underestimated between  $t = 0$   
413 and  $t = 35$  d, and slightly overestimated between  $t = 35$  and  $t = 62$  d (Fig. 8b). The reconstruction  
414 of the scaled functional response  $f$  provided indications on the fact that the food levels within the  
415 sediment between  $t = 35$  d and  $t = 62$  d might have been really low and did not allow an optimal  
416 growth.

### 417 2.2.2. Chronology and growth during the life cycle according to the environmental conditions

418 *In situ environmental conditions.* The seawater temperatures ranged from 5.5 to 20 °C at Wimereux,  
419 with the highest temperature between July and September and the lowest temperature between  
420 January and February (Fig.9a). The scaled functional response was supposed to range from 0.3 to  
421 0.95 with higher values in spring and autumn and lower values in summer and winter (Fig.9b).

422 *Chronology of the first life stages.* The abj-model predicted an age at trochophore larva stage  $a_{tr}$   
423 of 10.3 days and an age at birth  $a_b$  (used as an approximation of the age at the first settlement)  
424 of 15.5 days at Wimereux, considering the environmental conditions presented in Fig. 9 (Table  
425 6), suggesting a first dispersal phase in between these two events of around 5 days. The age at  
426 the end of metamorphosis  $a_j$  was predicted to be 208 days (a little less than 7 month) in local  
427 environmental conditions, which means around mid April for a spawning period in mid September.  
428 The age and trunk length at puberty of the lugworms of Wimereux,  $a_p$  and  $TL_p$ , were predicted  
429 to be respectively 373.2 days and 3.5 cm.

430 *Wet weight and trunk length growth predictions according to the environmental conditions.* The  
431 total wet weight of *Arenicola marina* (considering the structure, reserve and reproduction buffer  
432 compartments) predicted by the model at the maximum age  $a_m$  was around 20 times superior in  
433 optimal conditions ( $f = 1$  and  $T = 20^\circ\text{C}$ , around 400 g) compared to *in situ* conditions recorded  
434 at Wimereux ( $f = 0.4$  and  $T = 13^\circ\text{C}$ , around 20 g) (Fig. 10). The total trunk length of *A. marina*  
435 predicted by the model was more than twice superior in optimal conditions ( $f = 1$  and  $T = 20^\circ\text{C}$ ,  
436 around 33 cm) than in the environmental conditions recorded at Wimereux ( $f = 0.4$  and  $T = 13^\circ\text{C}$ ,  
437 around 14 cm) (Fig.11).

### 438 2.3. Comparison of the abj-DEB parameters of *Arenicola marina* with other Lophotrochozoan species

439 In annelids and molluscs, the maximum assimilation rate,  $\{\dot{p}_{Am}\}$ , increased with the maxi-  
440 mum structural length as expected, and more markedly after metamorphosis and the associated  
441 metabolic acceleration phase (Fig. 12). The values for *Arenicola marina* with the abj-model ap-  
442 peared lower than those of most of the other polychaetes and clitellates species before and after  
443 metamorphosis, except for the values of *Urechis caupo* (echiurian species), the only other annelid  
444 species for which an abj-model was applied.

445 The value of the maximum assimilation rate after metamorphosis,  $\{\dot{p}_{Am}\}_j$ , of *A. marina* with  
446 both models was close to the one expected for the generalized animal, and followed the tendency  
447 found in most of the others mollusc species, which was not the case for the other polychaetes species,  
448 mostly showing higher values. The allocation fraction to soma,  $\kappa$ , was higher for *A. marina* ( $\sim$   
449 0.92) than the one expected for the generalized animal (0.8), and did not appear inconsistent with  
450 the values of  $\kappa$  calculated in molluscs species (Fig. 12). The energy conductance value,  $\dot{v}$ , for *A.*  
451 *marina* appeared lower in the abj- than in the std-model before metamorphosis, but the opposite  
452 happened after metamorphosis, where the abj-model's value was higher than the generalized animal  
453 but closer to molluscs' values. The specific somatic maintenance costs values,  $[\dot{p}_M]$ , of *A. marina*  
454 were much lower than those predicted for most of the other species of annelids (except for *Urechis*  
455 *caupo*) but were close to the one of the generalized animal and are consistent with the values for  
456 the molluscs species (Fig. 12). The value of the costs of structure,  $[E_G]$ , of *A. marina* appeared  
457 equal to those of the other annelids' and most of the molluscs' species (Fig. 12). The value of the

458 maturity maintenance rate coefficient,  $k_j$ , of *A. marina* was equal to those of the other annelids'  
459 and of most of the molluscs' species (Fig. 12). The values of the maturity thresholds for birth,  
460 metamorphosis and puberty,  $E_H^b$ ,  $E_H^j$ , and  $E_H^p$ , of the abj-model for *A. marina* were lower than  
461 those of the generalized animal but similar to most of the mollusc species' values (Fig. 12).

### 462 3. Discussion

463 In the present study, we successfully estimated the parameters of both a std- and an abj-DEB  
464 model for the lugworm *Arenicola marina*, combining the use of literature, experimental and field  
465 data. We found that the abj-model was more appropriate for modelling *A. marina*'s energy budget  
466 and life cycle and implemented it under field conditions to reconstruct feeding levels as well as *A.*  
467 *marina*'s growth and life cycle chronology.

#### 468 3.1. Physiological implications of the std- and the abj- parameter estimation results

469 Major differences in the organisms physiology were implied by the parameter results obtained  
470 with a faster metabolism for *Arenicola marina* with a std-DEB model compared to an abj-DEB  
471 model. Indeed,  $\{\dot{p}_{Am}\}_b$ ,  $\{\dot{p}_{Am}\}_j$ ,  $[\dot{p}_M]$  and  $\dot{v}_b$  appeared higher with the std- parameter estimation,  
472 and  $\dot{v}_j$  higher with the abj- parameter estimation. First, a higher value of the maximum assim-  
473 ilation rate  $\{\dot{p}_{Am}\}$  implies a higher value of the assimilation flux from the same amount of food,  
474 and a higher value of the energy conductance  $\dot{v}$  implies a larger mobilization flux (Agüera et al.,  
475 2015). The reserve capacity  $[E_m]$ , defined by the ratio  $[E_m] = \{\dot{p}_{Am}\} / \dot{v}$  (Montalto et al., 2014) was  
476 found to be 16766 J.cm<sup>-3</sup> with the std- parameter estimation compared to 1177 J.cm<sup>-3</sup> with the  
477 abj- parameter estimation (considering a temperature of 20°C). In comparison,  $[E_m]$  values for ac-  
478 celerating molluscs species were estimated around 4500 J.cm<sup>-3</sup> and  $[E_m]$  values for non-accelerating  
479 molluscs species were estimated around 11600 J.cm<sup>-3</sup> (Add-my-Pet collection consulted in Novem-  
480 ber 2018). Second, a higher value of the volume-specific maintenance costs,  $[\dot{p}_M]$ , implies a higher  
481 level of energy needed for the same amount of structure acquired. The comparison of the parame-  
482 ter estimation of the abj- and std- models therefore resulted on the one hand, with the parameter  
483 estimation of the std-model, in one organism able to store more energy in the reserve compart-  
484 ment, but also using more energy for the maintenance of its structure, and on the other hand,  
485 with the parameter estimation of the abj-model, in one organism able to store less energy in the  
486 reserve compartment, but using less energy for the maintenance of its structure. Indeed, although  
487 the predictions of the std- and abj- versions of the model were quite similar (except for the early  
488 life-stages predictions), they implied really different bioenergetics in two kinds of organisms storing  
489 and using energy differently.

#### 490 3.2. Implications of using an abj-model for *Arenicola marina* in relation with its biology and ecology

491 For *Arenicola marina*, the abj-model gave better fit results than the std-model (smaller MRE  
492 and SMSE), even when only few observations within the data set accounted for the acceleration

493 period (only the zero-variate observations  $a_j$  and  $L_j$  were added, but no uni-variate observations  
494 made between birth and metamorphosis). The presence of a metabolic acceleration between birth  
495 and metamorphosis in *A. marina* might be related to its benthopelagic life cycle. Indeed, acceler-  
496 ating species have longer incubation time (before birth) than non-accelerating species (Kooijman,  
497 2014; Kooijman et al., 2011), which might be linked to the presence of a larval dispersal phase, since  
498 a lower metabolism (in comparison with a non-accelerating species, or with juvenile or adult from  
499 the same species) allows for more dispersal time, especially when dispersal rate mainly depends  
500 on passive water transport (Kooijman, 2014). This seems in accordance with the presence of a  
501 dispersal phase happening before birth for *A. marina* and with the fact that predictions of  $a_{tr}$  and  
502  $a_b$  of the std- model presented in this study appeared much smaller than observations, compared to  
503 predictions made by the abj-model. In lugworms, the gradual change of feeding behaviour between  
504 the first feeding at birth, the temporary settlement between birth and metamorphosis, and the  
505 semi-permanent settlement after metamorphosis on the foreshores inhabited by adults (Farke and  
506 Berghuis, 1979a, b) might be a mechanism of the increase of the metabolic acceleration  $s_M$ , also  
507 increasing the resulting specific assimilation rate (since  $\{\dot{p}_{Am}\}_j = s_M \cdot \{\dot{p}_{Am}\}_b$ ) of the individual.  
508 The increase of the organic matter concentration within the water column during spring (spring  
509 blooms), before the second dispersal phase when metamorphosis is almost completed, might also  
510 play a role in the increase of the specific assimilation rate, increasing the amount of food available  
511 for the same feeding effort.

### 512 3.3. Phylogenetic implications of the use of abj-DEB models for polychaetes

513 The metabolic acceleration rate value for *A. marina* ( $\sim 11$ ) falls in the range of what can be  
514 found in mollusc species (for more than 95% of the mollusc species,  $1 \leq s_M \leq 27$  in the AmP  
515 database), which seems consistent with the fact that polychaetes and molluscs both belong to the  
516 Lophotrochozoan clade, having both a common trochophore larval stage after the embryogenesis.  
517 However, although all annelids are part of the Lophotrochozoan clade, they do not all share the  
518 presence of at least one larval dispersal phase during their life cycle, and therefore, might not  
519 all experience a metabolic acceleration during their life cycle. As an example, clitellates have a  
520 direct development with no larval phase (related to their terrestrial habitat) and std-models might  
521 show better fit for these species that may not experience a metabolic acceleration during their life  
522 cycle. From an evolutionary point of view, metabolic acceleration might first have been common  
523 to all Lophotrochozoans and secondarily lost in clitellate species (as suggested by Marques et al.,  
524 2018, for other taxa). Nevertheless, since some species with no larval phase might also experience  
525 a metabolic acceleration (Kooijman, 2014), and since metabolic acceleration seem common to a  
526 large part of the species belonging to the Lophotrochozoan clade (Kooijman, 2014; Marques et  
527 al., 2018), a comparison of the use of both abj- and std- models for clitellate species should be  
528 considered.



529 3.4. Energy budget and *in situ* life cycle predictions

530 The predictions on the chronology of *Arenicola marina*'s life cycle stages under the *in situ*  
531 environmental conditions met at Wimereux (metamorphosis completed at around 7 months, in  
532 mid April) seemed in accordance with observations made by De Cubber et al. (2018), who spotted  
533 the first recruits of the species (e.g. juveniles after metamorphosis) in May at the same site.  
534 This would suggest a second dispersal period of less than one month if lugworms migrate after  
535 metamorphosis. Moreover, the age and length at puberty of the lugworms at the Wimereux site  
536 were predicted with the abj-model to be respectively 373.2 days and 3.5 cm, which is close to the  
537 observations made by De Cubber et al. (2018) with a length at first spawning (after the acquisition  
538 of maturity) of 3.8 cm and an age of 1.5 to 2.5 years. Newell (1949, 1948) reported the presence  
539 of *A. marina* metatrochophore larvae close to birth (and thus close to the first settlement stage)  
540 with 3 to 4 setigers and around 0.034 cm of length around 2 to 3 weeks after the occurrence of the  
541 spawning event at Whistable (UK) (limit between the English Channel and the North Sea). His  
542 observations also seem in accordance with the abj-model predictions. Indeed, the age and length at  
543 birth predictions at Wimereux in October were of 15.5 days and 0.034 cm. Since temperatures are  
544 lower in November in the English Channel and even lower in the North Sea, birth might have been  
545 slightly delayed in their study and their observations seem to be in accordance with the abj-model  
546 implemented in our study. Observations of post-larvae in mucus tubes were commonly made on  
547 fucus and pebbles areas until the end of February (Benham, 1893; Newell, 1949, 1948) and up to  
548 April in some cases (Newell, 1949). First settlements of juveniles on adult grounds were reported  
549 by Newell (1949, 1948) at the end of April or beginning of May, which is in accordance with our  
550 model predictions (after the age at metamorphosis, which is around 5 months-old) and correspond  
551 to a dispersal period after metamorphosis of a maximum of one month.

552 The biggest individuals of *A. marina* collected at the studied sites might give indications on the  
553 *in situ* environmental conditions met by the lugworms on these sites. Indeed, at Wimereux, the  
554 heaviest individual collected by the authors between 2015 and 2018 weighted 10 g and the longest  
555 one measured 15.2 cm of trunk length (data not shown), which is in accordance with the length and  
556 weight predicted by the abj-model for *Arenicola marina* at an age of 5 to 6 years old (age of the last  
557 cohort calculated by De Cubber et al. (2018)) for  $f = 0.4$  and  $T = 13^{\circ}\text{C}$ . At Le Touquet (Eastern  
558 English Channel, De Cubber et al., 2018), the heaviest individual collected weighted 53.1 g and  
559 the longest one measured 20.2 cm of trunk length (data no shown), and at Fort Mahon (Eastern  
560 English Channel, De Cubber et al., 2018), the heaviest individual collected weighted 26 g and the  
561 longest one measured 18.4 cm of trunk length (data not shown). Since no major difference between  
562 the seawater temperature at the three different sites exist, the main difference was possibly the food  
563 availability. The comparison of these biometric values with the ones predicted by the abj-model  
564 (around 600 g of maximum wet weight and 35 cm of maximum trunk length for  $f = 1$  and  $T =$   
565  $13^{\circ}\text{C}$ ) seems to indicate that  $f$  was higher at Le Touquet and Fort Mahon compared to Wimereux.

566 In the different sites of the Eastern English Channel cited previously, De Cubber et al. (2018)  
567 showed that the lugworms' recreational harvest in 2017 removed more than 500 000 lugworms and  
568 represented a total retail value of around 232 447 euros. The need for implementing management  
569 measures was also evidenced for at least one beach by these authors. Knowing the food levels of the  
570 different sites might then enable predictions with the abj-model on the *in situ* ages and lengths at  
571 puberty, which could help managers to implement relevant regulations if needed such as a relevant  
572 harvest minimum size limit on the different sites showing highly variable food levels and maximum  
573 lengths and weights.

### 574 3.5. Possible future extensions of the model

575 In order to provide the best model possible for *Arenicola marina* further adjustments could be  
576 implemented linked to the species life cycle and habitats. First, defining the temperature tolerance  
577 range of the species could improve the abj-model by applying better temperature corrections.  
578 Growth experiments from Farke and Berghuis (1979) seem to point out a higher boundary of  
579 the temperature tolerance range  $T_H$  around 25 °C. Other studies suggest a lower boundary of  
580 the temperature tolerance range  $T_L$  under 5°C (Sommer et al., 1997; Wittmann et al., 2008),  
581 but no Arrhenius temperatures beyond the temperature tolerance ( $T_{AH}$  and  $T_{AL}$ ) range could  
582 be calculated yet. Further experiments on growth or respiration under temperatures beyond the  
583 temperature tolerance range could be performed to define  $T_{AH}$  and  $T_{AL}$  and thus improve the  
584 temperature correction.

585 During their life cycle, the different stages of *A. marina* inhabit different marine habitats with  
586 different ranges of temperature variation. From the metatrochophore to the post-larval stage the  
587 lugworms inhabit the subtidal area where seawater temperature does not fluctuate that much daily,  
588 compared to the intertidal areas inhabited by the juveniles and adults, where temperature can  
589 change dramatically during one day. As an example, a variation of 15°C was recorded within  
590 the sediment at the Wimereux site in November 2017 (Fig. 13). In this study, the Arrhenius  
591 temperature was calculated from the oxygen consumption rate of juveniles living on the upper  
592 shore. We hypothesize that a different Arrhenius temperature may exist for the larval and post-  
593 larval stages living in habitats with a more stable temperature, as suggested by Kooijman (2010).  
594 Further experiments could be implemented on larvae in order to record physiological rates and  
595 estimate their Arrhenius temperature.

596 Monaco and McQuaid (2018) highlighted the interest of adding to the temperature correction  
597 an aerial exposure term  $M_d$  (linked to tidal height and the position of organisms on the shore) in  
598 foreshore habitats showing wide fluctuations in temperature and desiccation. Given the intense  
599 variations experienced by juvenile and adult lugworms (Fig. 13), it might be interesting to add an  
600 aerial exposure term for the species. Indeed, the underestimation of growth by the model compared  
601 to our observation of growth of juveniles in the laboratory (Exp. C) might be linked to the fact  
602 that no tide was simulated and lugworms stayed immersed during all the experiment time, without

603 the stress brought by high temperature variations and aerial exposure. However, it was found that  
604 lugworms gradually migrate down the shore while growing (De Cubber et al., 2018), so the aerial  
605 exposure correction, if implemented, should gradually decrease during the life cycle of the organism  
606 as well.

607 DEB models as implemented here enable to reconstruct the growth and the reproduction of  
608 a species at the individual level. However, in order to be used in a population context, DEB  
609 theory can be associated to individual-based models (IBM) in order to explore properties of both  
610 individual life-history traits and population dynamics (Bacher and Gangnery, 2006; Martin et al.,  
611 2012). The association of the abj-model developed here, and providing predictions on the duration  
612 of the larval dispersal phase, with biophysical larval dispersal models (Nicolle et al., 2017) could  
613 also allow the understanding of the populations' connectivity in the area and thus give valuable  
614 information for the conservation of the species.

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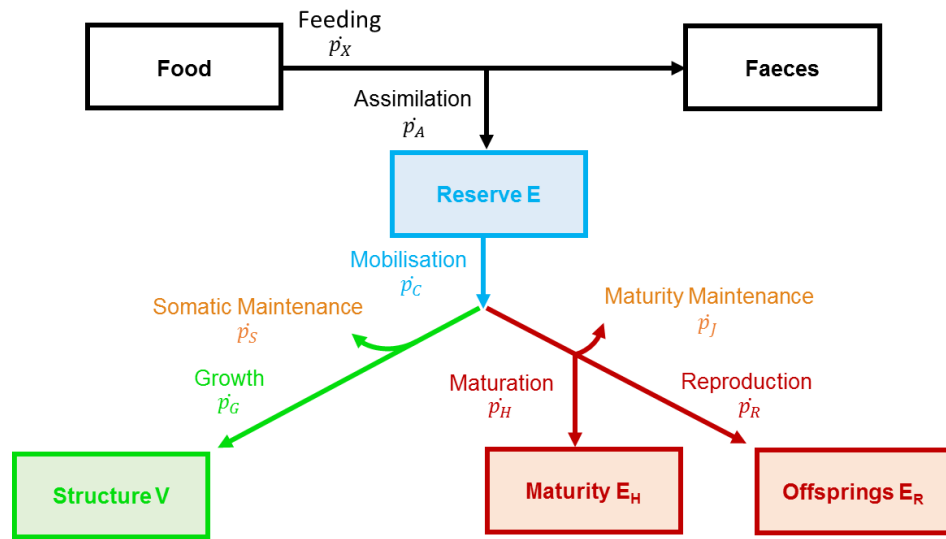


Figure 1: Schematic representation of DEB model and associated state variables and fluxes, adapted from Kooijman (2010). Boxes are the state variables: 1) reserve  $E$  (J); 2) the structural volume,  $V$  ( $\text{cm}^3$ ); 3) the cumulated energy invested in Maturity,  $E_H$  (J) or in reproduction  $E_R$  (J). Arrows are energy flows in  $\text{J}\cdot\text{d}^{-1}$ . Details of  $\dot{p}_X$ ,  $\dot{p}_A$ ,  $\dot{p}_C$ ,  $\dot{p}_S$ ,  $\dot{p}_G$ ,  $\dot{p}_J$ ,  $\dot{p}_H$  and  $\dot{p}_R$  are given in Table 1.



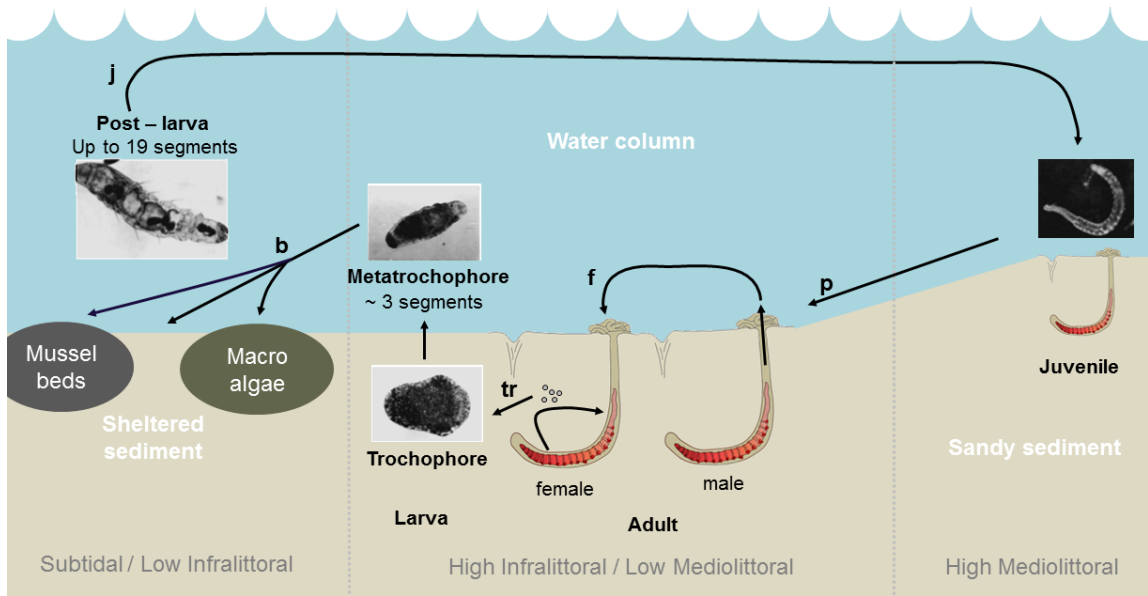


Figure 2: Life cycle of *Arenicola marina* and associated habitats. f stands for fertilization; tr for when the trochophore larva appears; b for birth (e.g. first feeding, as described in the DEB theory); j for the end of metamorphosis; and p for puberty. Adapted from Farke and Berghuis (1979a, 1979b), Reise (1985) and Reise et al. (2001). Pictures of the different life stages of *A. marina* are taken from Farke and Berghuis (1979a).

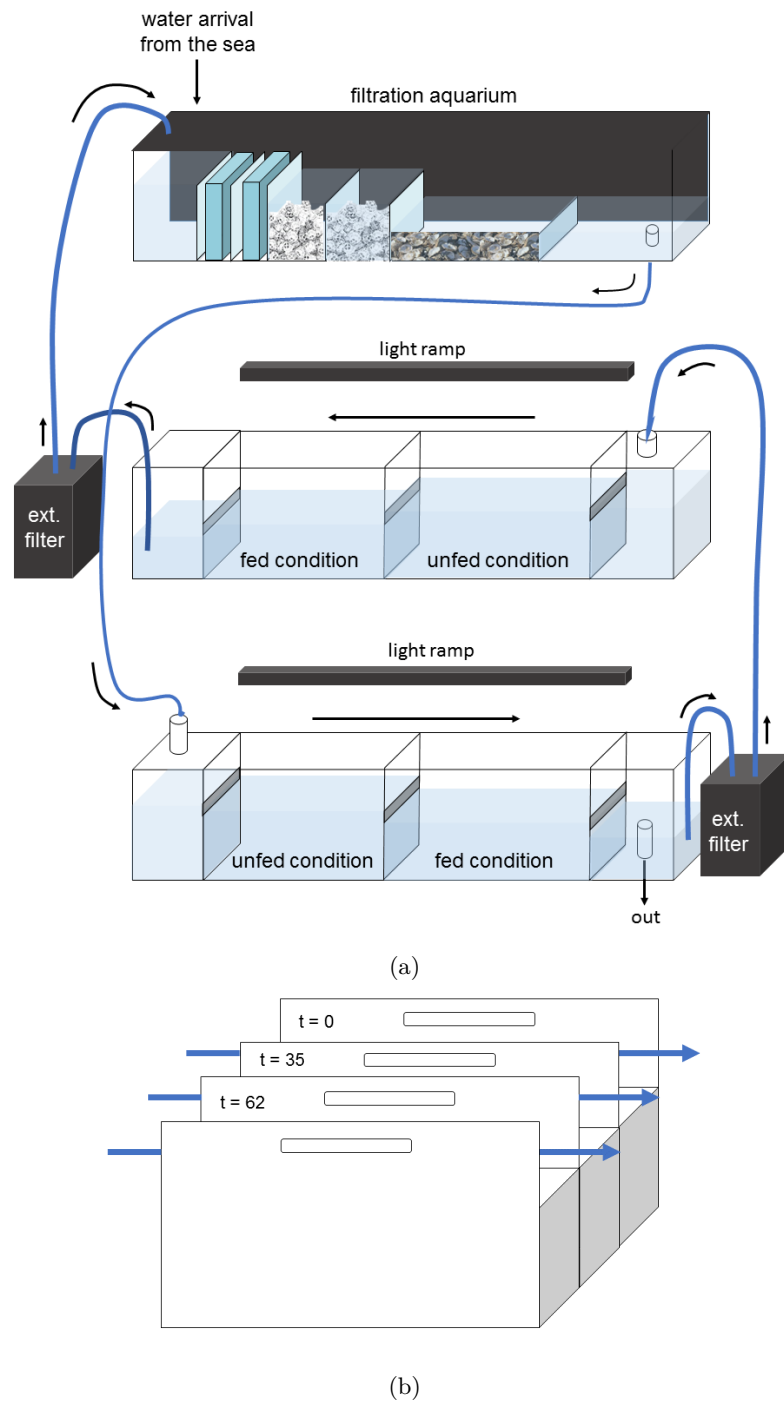


Figure 3: Circulation, filtration of seawater and lightening (a) and boxes (b) of the custom *Arenicola marina* growing system with recirculating seawater (Experiment C). The filtration aquarium tray (a, top aquarium) contained (from left to right): fine and coarse filter foam, crushed pozzolana and oyster shells. The boxes (b) were placed in each of the fed and unfed condition in the remaining aquarium trays and removed at  $t = 0$  (first feeding of the fed condition),  $t = 35$  and  $t = 62$  days. The light ramps consisted in Alpheus Radiometrix 13C1001C, and the external (ext.) filters in Eheim professional 4+ 250. The oxygenation ramps placed in the last two aquarium trays are not represented.

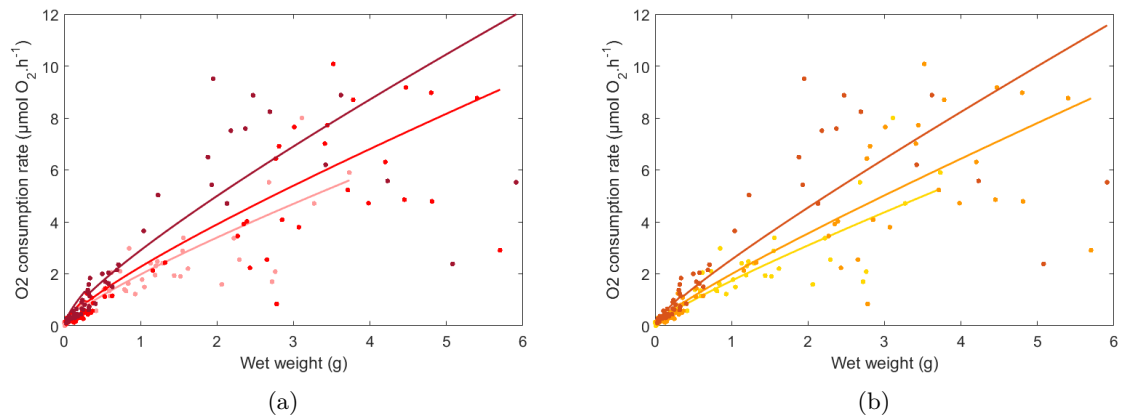


Figure 4: Data (dots) and predictions (lines) of the oxygen consumption of the abj-DEB model (a) and the std-DEB model (b) of *Arenicola marina* measured by the authors (Exp. A) as a function of wet weight at three different temperatures (from light to dark red: 12, 15 and 20.5°C). The respective relative errors from 12 to 20.5°C were 0.29, 0.38 and 0.40 with the abj-model (b) and 0.28, 0.33, and 0.38 with the std-model (b).

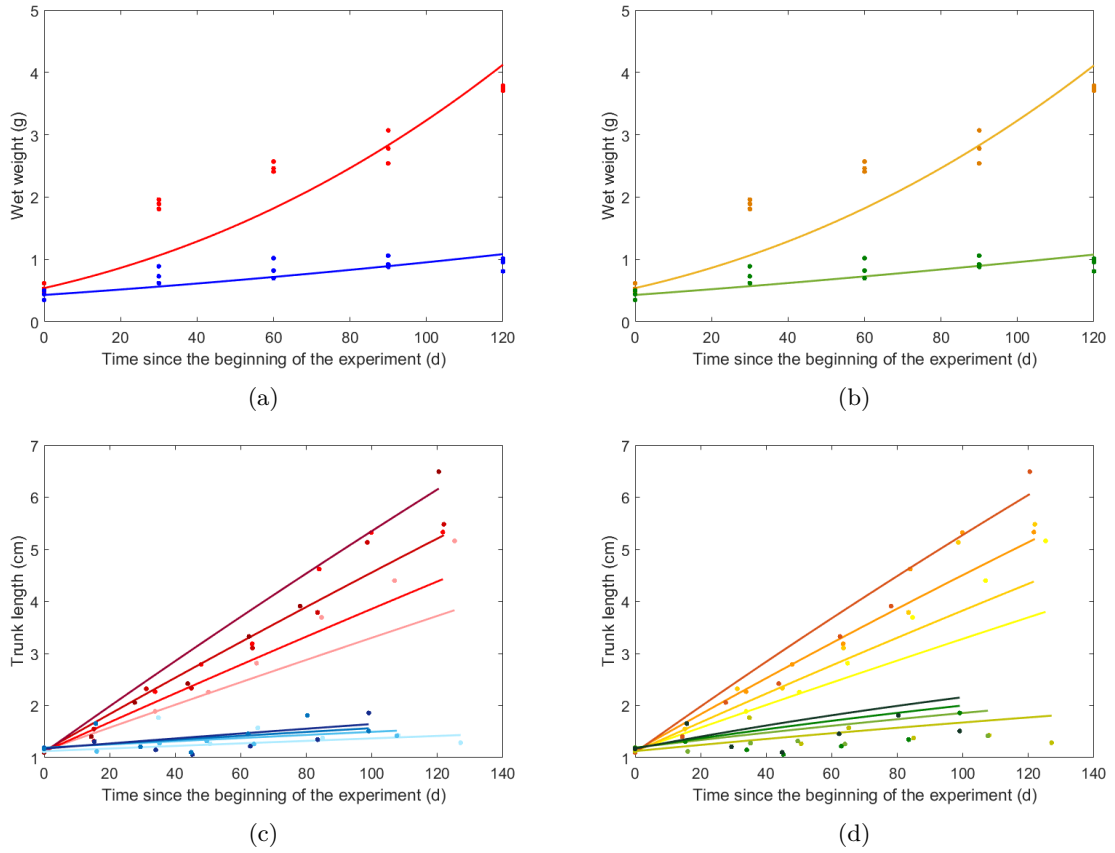
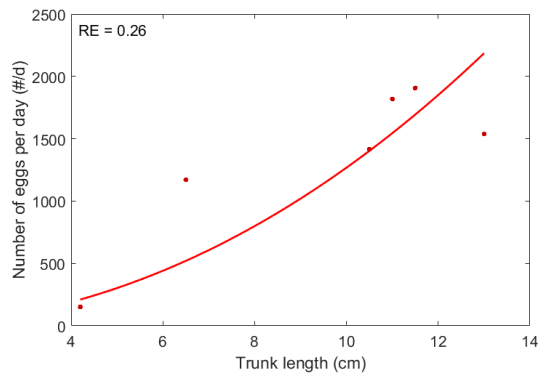
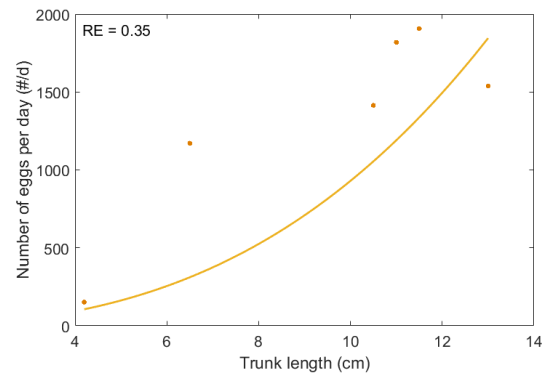


Figure 5: Data (dots) and predictions (lines) of the growth of *Arenicola marina* juveniles in wet weight (a,b) and in trunk length (c,d) using both an abj-DEB model (a,c) and a std-DEB model (b,d). Data from (a,b) was taken from Olive et al.(2006). *A. marina* was reared in fed (red and orange) and unfed (blue and green) conditions between 12 and 20°C. The respective relative errors (RE) for the growth curves in fed and unfed conditions were 0.19 and 0.15 (a) 0.18 and 0.15 (b). Data from (c,d) were taken from De Wilde and Berghuis (1979). *A. marina* was reared in fed (red and orange) and unfed (blue and green) conditions at four different temperatures (from light to dark: 5, 10, 15 and 20°C). The respective (RE) for the growth curves in fed conditions at 5, 10, 15 and 20°C were 0.15, 0.17, 0.07 and 0.13 with the abj-model (c) and 0.16, 0.18, 0.08 and 0.12 with the std-model (d). The respective RE for the growth curves in unfed conditions at 5, 10, 15 and 20°C were 0.11, 0.06, 0.13 and 0.12 with the abj-model (c) and 0.17, 0.19, 0.23 and 0.24 with the std-model (d).



(a)



(b)

Figure 6: Data (dot) and prediction (line) of the reproduction rate of *Arenicola marina* collected by the authors (Exp. B) (Eastern English Channel, France, see Table 2) as a function of trunk length using both an abj-DEB model (a) and a std-DEB model (b). RE stands for relative error.

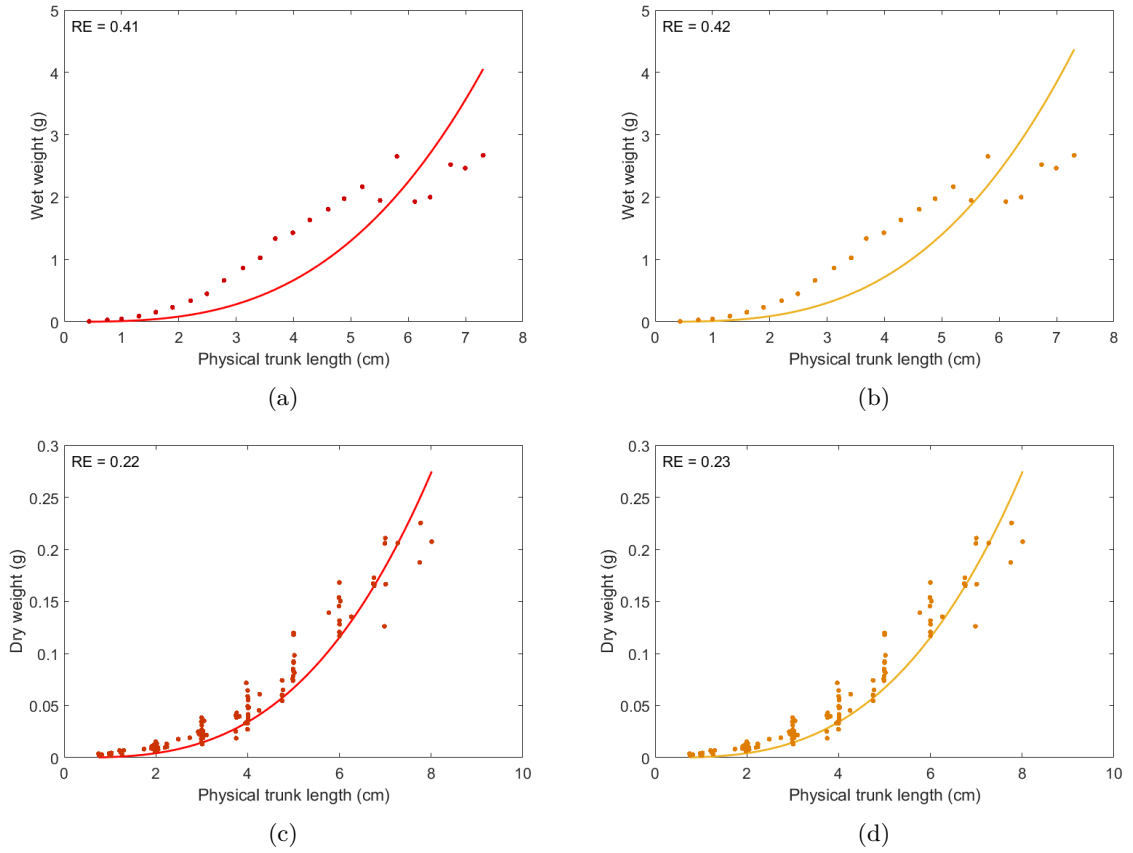
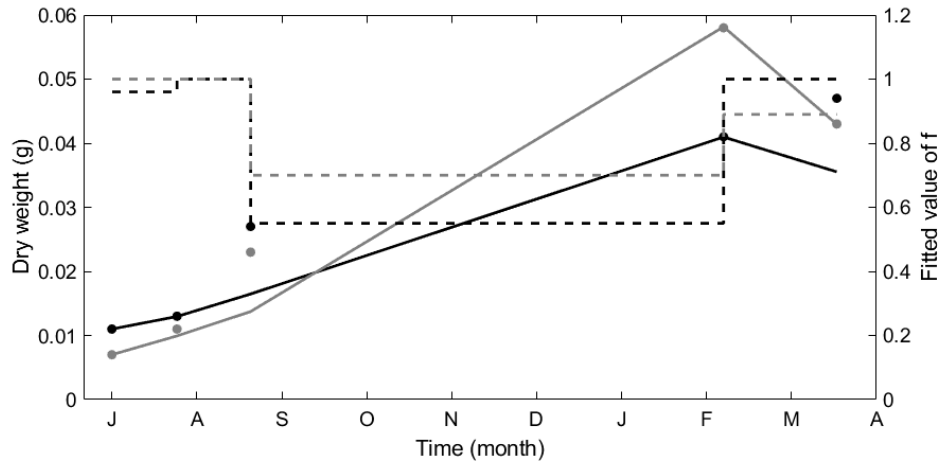
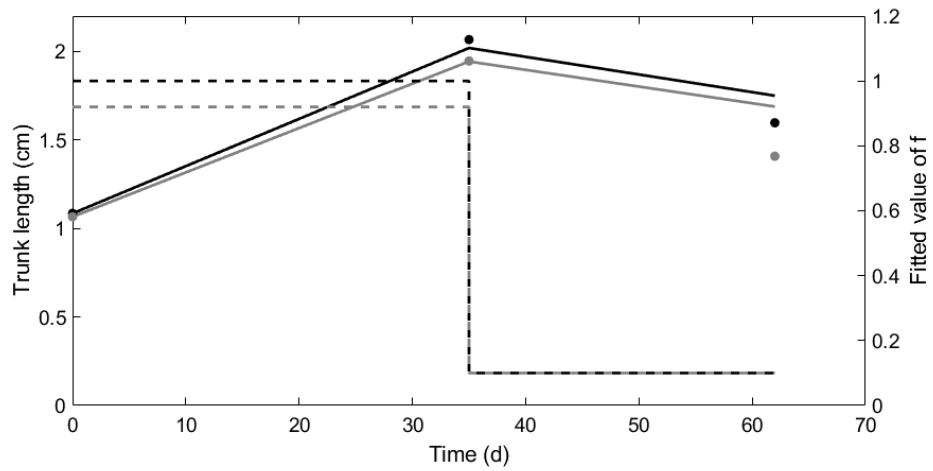


Figure 7: Data (dots) and predictions (lines) of the wet weight as a function of trunk (a,b) and total (b,c) length for *Arenicola marina* individuals collected at Wimereux (this study) and of the dry weight as a function of trunk length (e,f) for *A. marina* (data from De Wilde and Berghuis (1979)) using both an abj-DEB model (a,c,e) and a std-DEB model (b,d,f). The corresponding values of the shape coefficient are: (a)  $\delta_M = 0.20$  (b)  $\delta_M = 0.13$  (c)  $\delta_M = 0.14$  (d)  $\delta_M = 0.09$ . RE stands for relative error.

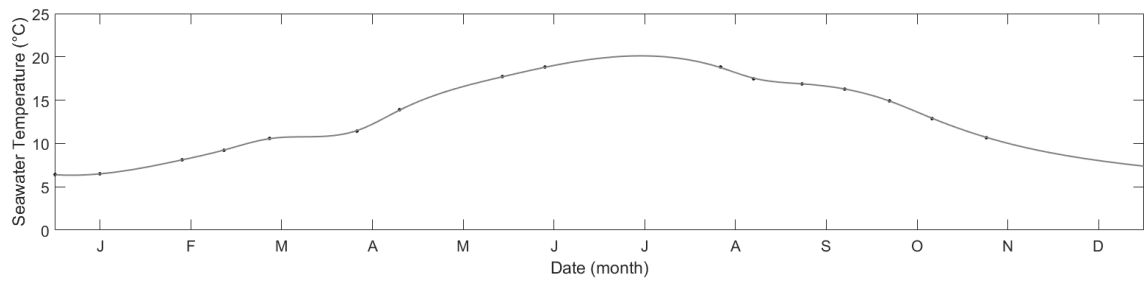


(a)

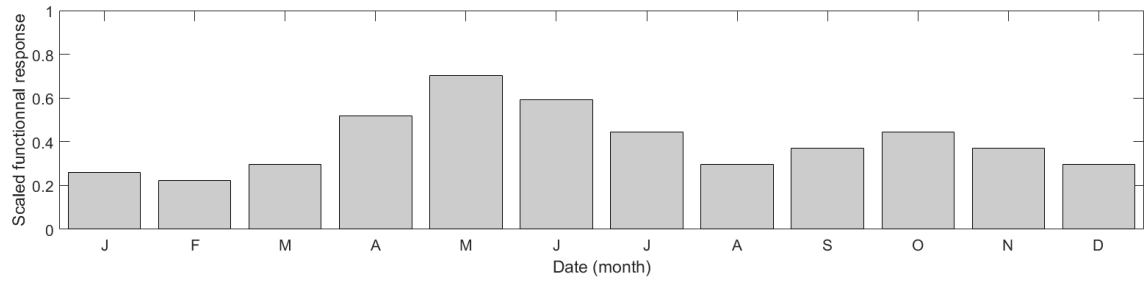


(b)

Figure 8: Reconstruction of the scaled functional response value ( $f$ , dashed lines) from field (a) and experimental (b) data (dots) with the abj-model for *Arenicola marina* from this study (lines are the model predictions). The observations of a field growth survey of *A. marina* (dots) are taken from Beukema and De Vlas (1979) in two beaches of the Wadden Sea (black and grey). The associated predictions of the DEB model (lines) for the best fitted values of  $f$  (dashed lines) are represented. Sea surface temperature were taken from Van Aken (2008). The laboratory observations (dots) on growth in trunk length of *Arenicola marina* in different feeding conditions (fed in black and unfed in grey) are associated to the growth predictions of the DEB model (lines) with the fitted values of the scaled functional response  $f$  (dashed lines) for  $T = 16.5^{\circ}\text{C}$ .



(a)



(b)

Figure 9: *In situ* temperature of the seawater at Wimereux (Hauts-de-France, Eastern English Channel) during the year 2017 (a), and estimated scaled functional response  $f$  at this site (b), used for the predictions of the chronology of the first life stages of the life cycle of *Arenicola marina* and of the wet weight and trunk length growth of the species at this site.



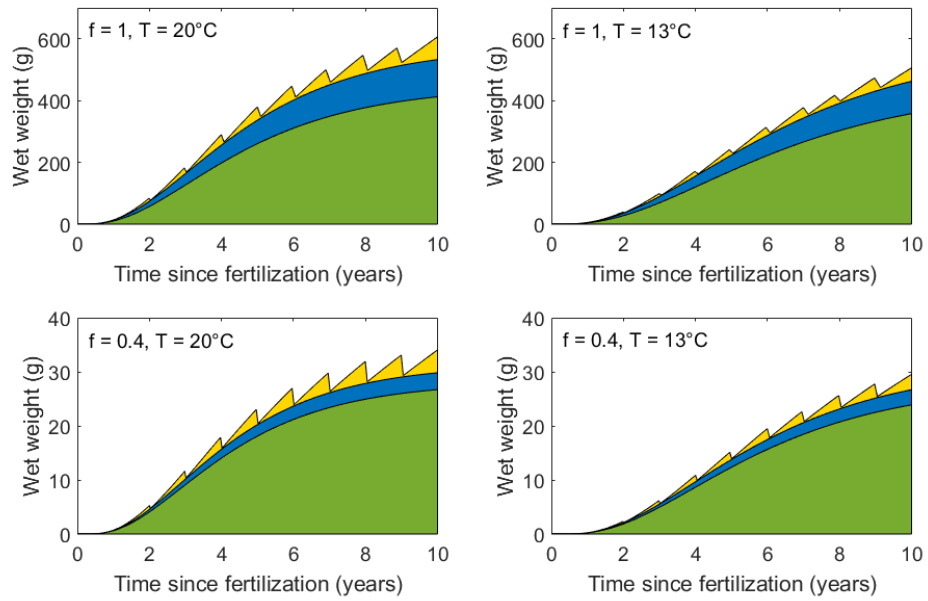


Figure 10: Predictions of the abj-DEB model of the evolution of the wet weight of the structure  $S$  (green), the reserve  $E$  (blue) and the reproduction buffer  $E_R$  (yellow) compartments of *Arenicola marina* under different environmental conditions from fertilization time:  $f = 1$  (food available *ad libitum*) &  $T = 20^\circ\text{C}$ ;  $f = 1$  &  $T = 13^\circ\text{C}$ ;  $f = 0.4$  &  $T = 20^\circ\text{C}$ ;  $f = 0.4$  &  $T = 13^\circ\text{C}$  (mean environmental conditions found at Wimereux).

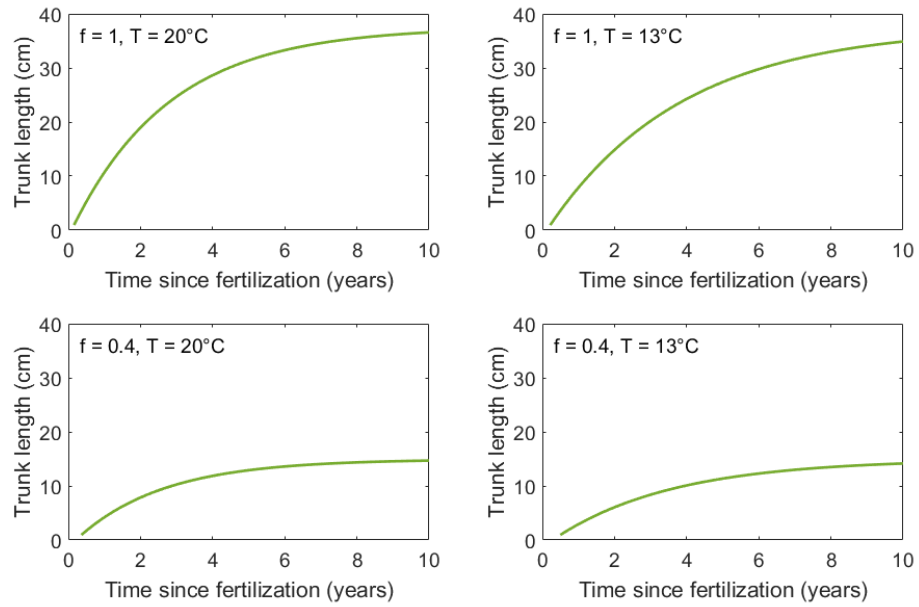


Figure 11: Predictions of the abj-DEB model of the evolution of the trunk length of *Arenicola marina* under different environmental conditions ( $f = 1$  &  $T = 20^\circ\text{C}$ ;  $f = 1$  &  $T = 13^\circ\text{C}$ ;  $f = 0.4$  &  $T = 20^\circ\text{C}$ ;  $f = 0.4$  &  $T = 13^\circ\text{C}$ ) from the age at puberty  $a_p$ .

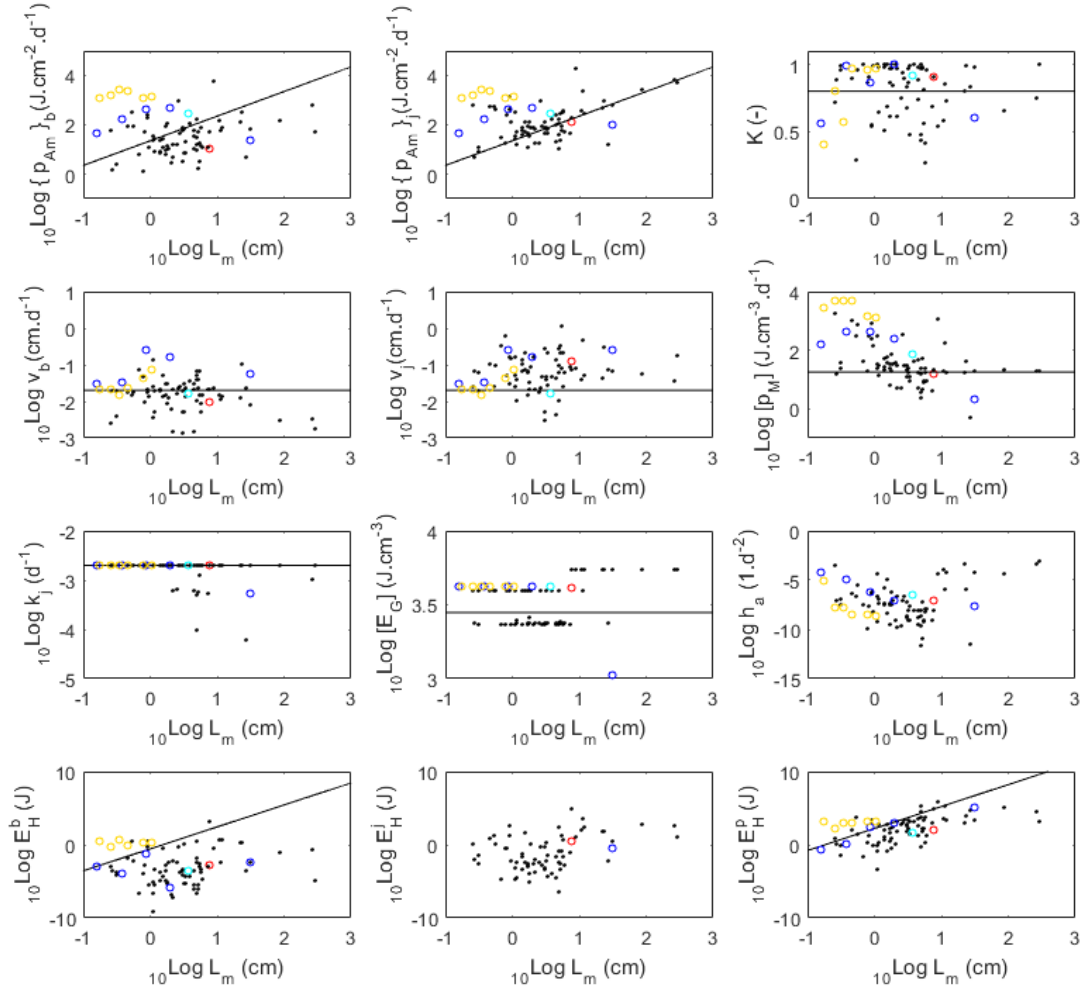


Figure 12: Comparison of log-log plots of  $\{p_{Am}\}_b$  at birth and  $\{p_{Am}\}_j$  after the metamorphosis,  $\kappa$ ,  $v_b$  at birth and  $v_j$  after the metamorphosis,  $[p_M]$ ,  $k_j$ ,  $[E_G]$ ,  $h_a$ ,  $E_H^b$ ,  $E_H^j$ , and  $E_H^p$  in Mollusca (black), Annelida Clitellata (yellow), Annelida Polychaeta (blue), standard model values for *Arenicola marina* (cyan) and abj model values from this study (red).  $\{p_{Am}\}$  is the maximum assimilation rate,  $\kappa$  the fraction of mobilised reserve allocated to soma,  $v$  the energy conductance,  $[p_M]$  the specific somatic maintenance costs,  $k_j$  the maturity maintenance rate coefficient,  $[E_G]$  the costs of structure,  $h_a$  the Weibull ageing acceleration, and  $E_H^b$ ,  $E_H^j$ , and  $E_H^p$  the maturity thresholds for birth, metamorphosis and puberty. The lines correspond to expectations on the basis of the generalized animal (Kooijman, 2010, Table 8.1).

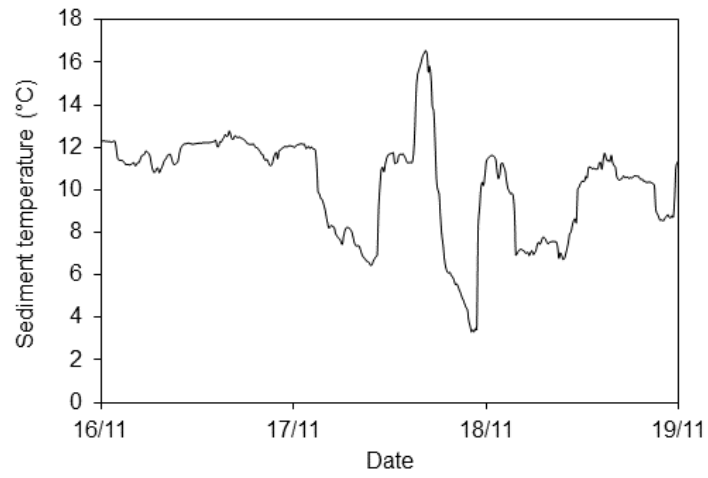


Figure 13: Sediment temperature (upper shore, 10 cm deep) recorded every 10 min at Wimereux (Hauts-de-France, Eastern English Channel) between the 16 of November and the 19 of November 2017 with a HOBO probe.

Table 1: State variables, fluxes, metric relationships, acceleration and shape coefficient of the abj-DEB model and associated mathematical expressions (Kooijman, 2014; Kooijman and Lika, 2014; Kooijman, 2010; Van der Meer, 2006).  $L$  is the structural length (cm) with  $L = V^{1/3}$ , and  $L_b$  and  $L_j$  are the structural lengths at birth and metamorphosis respectively.  $d_V$  is the density of wet structure,  $d_E$  the density of wet reserve,  $d_{Ed}$  the density of dry reserve,  $\mu_{Ed}$  the specific chemical potential of reserve and  $w_{Ed}$  the molar weight of dry reserve.  $L_w(t)$  is the physical total length at time  $t$  of the organism and  $TL_w(t)$  its physical trunk length.  $W_w(t)$  is the wet weight at time  $t$  of the organism. If  $E_H^j = E_H^b$  the abj- model reduces to the std- model.

State variables	Reserve	$\frac{dE}{dt} = \dot{p}_A - \dot{p}_C$
	Structure	$\frac{dV}{dt} = \frac{\dot{p}_G}{[E_G]}$
	Maturity	if $E_H < E_H^p$ $\frac{dE_H}{dt} = \dot{p}_H$ ; else $\frac{dE_H}{dt} = 0$
	Allocation to reproduction	if $E_H \geq E_H^p$ , $\frac{dE_R}{dt} = \kappa_R \cdot \dot{p}_R$ ; else $\frac{dE_R}{dt} = 0$
Fluxes	Ingestion	$\dot{p}_X = \frac{\dot{p}_A}{\kappa_X}$
	Assimilation	$\dot{p}_A = \{\dot{p}_{Am}\} \cdot s_M \cdot f \cdot V^{2/3}$
	Mobilisation	$\dot{p}_C = E \cdot \frac{\dot{v} \cdot s_M \cdot V^{2/3} \cdot [E_G] + \dot{p}_S}{\kappa \cdot E + V \cdot [E_G]}$
	Somatic maintenance costs	$\dot{p}_S = [\dot{p}_M] \cdot V$
	Maturity maintenance costs	$\dot{p}_J = \dot{k}_J \cdot E_H$
	Growth	$\dot{p}_G = \kappa \cdot \dot{p}_C - \dot{p}_S$
	Reproduction	$\dot{p}_R = (1 - \kappa) \cdot \dot{p}_C - \dot{p}_J$
	Maturity	$\dot{p}_H = (1 - \kappa) \cdot \dot{p}_C - \dot{p}_J$
Metric relationships	Physical length (cm)	$L_w(t) = \frac{V(t)^{1/3}}{\delta}$
	Wet weight (g)	$W_w(t) = d_V \cdot V(t) + (E(t) + E_R(t)) \cdot \frac{w_{Ed} \cdot d_E}{\mu_{Ed} \cdot d_{Ed}}$
Acceleration coefficient	if $E_H < E_H^b$ $s_M = 1$ ; if $E_H^b \leq E_H < E_H^j$ $s_M = L/L_b$ ; else $s_M = L_j/L_b$ if $E_H \geq E_H^j$	
Shape coefficient	if $E_H < E_H^b$ $\delta = \delta_{Me}$ ; if $E_H^b \leq E_H < E_H^j$ $\delta = \delta_{Me} + (\delta_M - \delta_{Me}) \cdot (\frac{L - L_b}{L_j - L_b})$ ; else $\delta = \delta_M$ if $E_H \geq E_H^j$	

Table 2: Abiotic and biometric data related to the samples of *Arenicola marina* collected at Wimereux, Le Touquet and Fort Mahon and used later on for the parameter estimation of a DEB-model for *A. marina*

Type of data	Number of samples	Collection date	Temperature (°C)	Wet weight (g)		Trunk length (cm)	
				range	mean	range	mean
Oxygen consumption	39	16/05/2018	12	0.00 - 3.73	$1.10 \pm 1.00$	0.36 - 5.60	$2.85 \pm 1.52$
	63	13/06/2018	15	0.02 - 5.70	$1.39 \pm 1.69$	0.80 - 7.30	$2.95 \pm 1.82$
	55	25/07/2018	20.5	0.03 - 5.91	$0.92 \pm 1.33$	0.90 - 6.80	$2.64 \pm 1.50$
Growth	290	26/05/2018	13	0.00 - 0.11	$0.05 \pm 0.02$	0.40 - 1.60	$1.10 \pm 0.20$
Reproduction	9	Sept. to Nov. 2016 to 2018	13	2.30 - 17.60	$6.10 \pm 5.60$	4.20 - 13.00	$7.40 \pm 3.70$

Table 3: Data used in the abj- and std- model parameters estimations for *Arenicola marina* among the available dataset. The age and length at metamorphosis were only used for the abj-model parameter estimation.

Type of data	Data	References
Zero-variate	age at trochophore larva	Pers. comm. from S. Gaudron
	age at birth	Farke and Berghuis (1979)
	age at metamorphosis	Farke and Berghuis (1979)
	age at puberty	De Cubber et al. (2018)
	lifespan	Beukema and De Vlas (1979), De Cubber et al. (2018)
	egg diameter	Watson et al (1998), De Cubber et al. (2018)
	total length of the trochophore larva	Farke and Berghuis (1979)
	total length at birth	Farke and Berghuis (1979)
	total length at metamorphosis	Farke and Berghuis (1979)
	trunk length at puberty	De Cubber et al. (2018)
	total maximum length	Pers. comm. from S. Gaudron (Sorbonne Univ.)
	wet weight of an egg	This study
Uni-variate	TL-Ww	This study
	TL-Wd	De Wilde and Berghuis (1979)
	t-TL (4 temperatures, 2 feeding conditions)	De Wilde and Berghuis (1979)
	t-Ww (2 feeding conditions)	Olive et al.(2006)
	Ww-O2 (3 temperatures, experimental conditions)	This study
	TL-R	This study

Table 4: Summary of the primary and some auxiliary parameters provided by the parameter estimation of the std- and the abj-DEB models for *Arenicola marina*

Parameter	Symbol	Value		Unit
		std-model	abj-model	
Reference temperature <sup>1</sup>	$T_{ref}$	293.15	293.15	K
Fraction of food energy fixed in reserve <sup>1</sup>	$\kappa_X$	0.80	0.80	-
Arrhenius temperature	$T_A$	3800	3800	K
Energy conductance <sup>2</sup>	$\dot{v}$ ( $\dot{v}_j$ )	1.67 e <sup>-02</sup> (-)	9.79 e <sup>-03</sup> (0.12)	cm.d <sup>-1</sup>
Allocation fraction to soma	$\kappa$	0.92	0.92	-
Reproduction fraction fixed in eggs <sup>1</sup>	$\kappa_R$	0.95	0.95	-
Volume specific costs of structure	$[E_G]$	4173	4127	J.cm <sup>-3</sup>
Maturation threshold for the trochophore larva	$E_H^{tr}$	2.73 e <sup>-04</sup>	8.44 e <sup>-04</sup>	J
Maturation threshold for birth	$E_H^b$	2.73 e <sup>-04</sup>	1.27 e <sup>-03</sup>	J
Maturation threshold for metamorphosis	$E_H^j$	-	1.94	J
Maturation threshold for puberty	$E_H^p$	38.62	104.50	J
Weibull ageing acceleration	$\ddot{h}_a$	3.08 e <sup>-07</sup>	6.69 e <sup>-08</sup>	d <sup>-2</sup>
Gompertz stress coefficient <sup>1</sup>	$s_G$	1.00 e <sup>-04</sup>	1.00 e <sup>-04</sup>	-
Acceleration rate <sup>3</sup>	$s_M$	-	11.46	-
Maximum assimilation rate <sup>2</sup>	$\{\dot{p}_{Am}\}$ ( $\{\dot{p}_{Am}\}_j$ )	280.08 (-)	10.62 (130.63)	J.cm <sup>-2</sup> .d <sup>-1</sup>
Specific somatic maintenance rate	$[\dot{p}_M]$	69.89	15.82	J.cm <sup>-3</sup> .d <sup>-1</sup>
Maturity maintenance rate <sup>1</sup>	$\dot{k}_J$	2.00 e <sup>-03</sup>	2.00 e <sup>-03</sup>	d <sup>-1</sup>
Specific density of wet structure <sup>1</sup>	$d_V$	1	1	g.cm <sup>-3</sup>
Specific density of wet reserve <sup>1</sup>	$d_E$	1	1	g.cm <sup>-3</sup>
Specific density of dry reserve <sup>1</sup>	$d_{Ed}$	0.16	0.16	g.cm <sup>-3</sup>
Specific chemical potential of dry reserve <sup>1</sup>	$\mu_{Ed}$	550000	550000	J.Cmol <sup>-1</sup>
Molar weight of dry reserve <sup>1</sup>	$w_{Ed}$	23.9	23.9	g.Cmol <sup>-1</sup>

<sup>1</sup> Fixed parameters. The values were taken from the generalized animal (Kooijman, 2010).

<sup>2</sup> The values inside brackets are the ones after metamorphosis when using the abj-model:  $\dot{v}_j = s_M \cdot \dot{v}$  and  $\{\dot{p}_{Am}\}_j = s_M \cdot \{\dot{p}_{Am}\}$

<sup>3</sup>  $s_M$  is given for a scaled functional response of 1 after metamorphosis



Table 5: Summary of the zero-variate observations values and associated predictions and relative errors (RE) obtained with both the abj- and the std-DEB models for *Arenicola marina*

Data	Symbol	Value	Predictions (RE)		Unit
			std-model	abj-model	
age at trochophore larva	$a_{tr}$	7	2.769 (0.60)	7.7 (0.1)	d
age at birth	$a_b$	30	2.774 (0.91)	10.52 (0.65)	d
age at metamorphosis	$a_j$	78	-	89.68 (0.15)	d
age at puberty	$a_p$	548	142.9 (0.74)	174.5 (0.68)	d
lifespan	$a_m$	2190	2194 (0.02)	2462 (0.12)	d
egg diameter	$L_0$	0.02	0.020 (0.02)	0.021 (0.07)	cm
total length of the trochophore larva	$L_{tr}$	0.025	0.014 (0.45)	0.019 (0.20)	cm
total length at birth	$L_b$	0.08	0.014 (0.83)	0.023 (0.71)	cm
total length at metamorphosis	$L_j$	0.89	-	0.85 (0.05)	cm
trunk length at puberty	$TL_p$	2.5	3.2 (0.28)	3.17 (0.27)	cm
maximum trunk length	$TL_i$	34	27.58 (0.19)	37.4 (0.10)	cm
wet weight of an egg	$Ww_0$	$4.78 \text{ e}^{-6}$	$4.45 \text{ e}^{-6}$ (0.07)	$5.15 \text{ e}^{-6}$ (0.08)	g

Table 6: Predictions on the chronology and lengths of different life cycle stages of *Arenicola marina* according to the *in situ* environmental conditions at Wimereux (Hauts-de-France, Eastern English Channel) made by the abj-DEB model.

Event	Length (cm)	Age (d)
Trochophore larva	0.021	10.29
Birth (first feeding)	0.034	15.51
End of the metamorphosis	1.12	208.26
Puberty	3.50	373.19