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# Fish shrinking, energy balance and climate change

**Running title:** Energy balance and fish shrinking

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**Keywords:** size decline, energy expenditure, feeding behaviour, temperature, small pelagic  
fish, respirometry, experiments

## 19 **Abstract**

20 A decline in size is increasingly recognised as a major response by ectothermic species to  
21 global warming. Mechanisms underlying this phenomenon are poorly understood but could  
22 include changes in energy balance of consumers, driven by declines in prey size coupled with  
23 increased energy demands due to warming. The sardine *Sardina pilchardus* is a prime  
24 example of animal shrinking, European populations of this planktivorous fish are undergoing  
25 profound decreases in body condition and adult size. This is apparently a bottom-up effect  
26 coincident with a shift towards increased reliance on smaller planktonic prey. We investigated  
27 the hypothesis that foraging on smaller prey would lead to increased rates of energy  
28 expenditure by sardines, and that such expenditures would be exacerbated by warming  
29 temperature. Using group respirometry we measured rates of energy expenditure indirectly,  
30 as oxygen uptake, by captive adult sardines offered food of two different sizes (0.2 or 1.2 mm  
31 items) when acclimated to two temperatures (16°C or 21°C). Energy expenditure during  
32 feeding on small items was tripled at 16°C and doubled at 21°C compared to large items,  
33 linked to a change in foraging mode between filter feeding on small or direct capture of large.  
34 This caused daily energy expenditure to increase by ~10% at 16°C and ~40% at 21°C on small  
35 items, compared to large items at 16°C. These results support that declines in prey size  
36 coupled with warming could influence energy allocation towards life-history traits in wild  
37 populations. This bottom-up effect could partially explain the shrinking and declining  
38 condition of many small pelagic fish populations and may be contributing to the shrinking of  
39 other fish species throughout the marine food web. Understanding how declines in prey size  
40 can couple with warming to affect consumers is a crucial element of projecting the  
41 consequences for marine fauna of ongoing anthropogenic global change.

## 42 **1. Introduction**

43 Ongoing global warming constitutes a major threat for biodiversity, especially in marine  
44 ecosystems, with some scenarios of future temperature increases reaching +5°C in 2100  
45 (IPCC, 2013, 2014; Orr et al., 2005). For ectotherms such as fishes, warming results in large  
46 increases in their physiological rates (Clarke & Fraser, 2004; Seebacher et al., 2015). While  
47 ongoing global warming might therefore be expected to boost growth rates in ectotherms  
48 (Morrongiello et al., 2019; Seebacher et al., 2015), it has in fact been correlated with a  
49 progressive decline in adult body size of many fish species in the wild (e.g. in Baudron et al.,  
50 2014; Gardner et al., 2011; Sheridan & Bickford, 2011, but see Audzijonyte et al., 2020). The  
51 factors that contribute to this shrinking of fishes are poorly understood; it is coherent with  
52 how warming might affect macroecological phenomena such as Bergmann's rule and James'  
53 rule, and is associated with a significantly higher proportion of younger age classes and a  
54 generalised decline in individual size-at-age in populations (Daufresne et al., 2009). Shrinking  
55 of fishes may also be linked to the Temperature-Size rule (TSR), the phenomenon whereby  
56 warm temperatures cause more rapid early growth of ectotherms but a decline in their final  
57 adult size, when compared to conspecifics reared in a cooler regime (Atkinson, 1994). The  
58 mechanisms involved in the TSR remain to be elucidated, it is observed in wild populations  
59 but can also be reproduced under controlled conditions in the laboratory (Forster et al., 2012;  
60 Horne et al., 2015).

61 Although there has been recent theoretical focus on whether the TSR relates to respiratory  
62 physiology (Verberk et al., 2021), early work focussed upon whether changes in energy  
63 budget and allocation may be a major driver of fish shrinking with warming (Gardner et al.,  
64 2011; Pauly et al., 2010). The availability and quality of food resources can affect individual

65 growth rates and adult body size through energy trade-offs among growth, survival and  
66 reproduction (Stearns, 1989, 1992), and such effects may be exacerbated if energy  
67 requirements are increased by warming. That is, it is unlikely that temperature per se is the  
68 only variable involved in the size decline in wild populations, since few exceptions to this rule  
69 are spreading across years, in particular studies that investigated food resources as a driver  
70 explaining the TSR (e.g. in Diamond & Kingsolver, 2010; Lee et al., 2015; Ljungström et al.,  
71 2020; Millien et al., 2006).

72 Temperature and food resources are both environmental variables whose variations can  
73 challenge an individual's energy balance and that can drive fish life-history traits through  
74 physiological processes. Thus, while the higher physiological rates due to warming cause an  
75 increase in energy demands, energy availability for marine fishes is predicted to decline due  
76 to climatic stressors that affect primary production and marine animal biomass (Ariza et al.,  
77 2022; Bopp et al., 2005; Daufresne et al., 2009; Lotze et al., 2019). Ocean warming can amplify  
78 vertical stratification and limit nutrient mixing (Roemmich & McGowan, 1995) which causes  
79 declines in plankton abundance at the base of the food web and leads to communities  
80 dominated by smaller-sized species and individuals (Bopp et al., 2005, 2013; Daufresne et al.,  
81 2009; Richardson & Schoeman, 2004; Ward et al., 2012).

82 The first impacts of such changes at low trophic levels could be observed on planktivorous  
83 species, such as small pelagic fishes (e.g. Brosset et al., 2017; van Beveren et al., 2014). These  
84 species represent about 25 % of worldwide fishery landings by weight (FAO, 2018), supporting  
85 the economy of several countries (Alheit et al., 2009; Fréon et al., 2005). Fluctuations of their  
86 populations can have critical economic and social consequences, as observed following the  
87 collapse of the Peruvian anchovy in the early 1970s (Alheit et al., 2009; Allison et al., 2009;

88 Schwartzlose et al., 1999). Population fluctuations of small pelagics are being exacerbated by  
89 ongoing global change (Brochier et al., 2013; Shannon et al., 2009), so these species represent  
90 key models to evaluate energetic mechanisms underlying shrinking of adult fish size.

91 In fact, small pelagic planktivorous fishes in the Mediterranean Sea are a major example of  
92 shrinking (Albo-Puigserver et al., 2021; Brosset et al., 2017). There is an ongoing and profound  
93 decrease in individual body size and condition of sardine (*Sardina pilchardus*) and anchovy  
94 (*Engraulis encrasicolus*), which appears to be a consequence of bottom-up control mediated  
95 by changes in plankton composition and abundance (Brosset et al., 2016; Saraux et al., 2019).  
96 This was associated with a major regime change in the mid-2000s, with shifts of nutrient  
97 inputs, water mixing and plankton production (Feuilloley et al., 2020). Since 2008, these  
98 species' diet has shifted progressively from large prey (> 1 mm, especially cladocerans) to  
99 increased reliance on smaller prey (< 1 mm, especially copepods), which indicates changes in  
100 the plankton community towards smaller species (Brosset et al., 2016). Smaller zooplankton  
101 can be less nutritious (Zarubin et al., 2014), so a decline in zooplankton size could entrain a  
102 decrease in rates of energy acquisition by their predators. Identifying a clear mechanistic link  
103 between a decrease in plankton size and fish growth, and ultimately population dynamics, is  
104 crucial since fish shrinking is spreading to new ecosystems and species (see Bensebaini et al.,  
105 2022; Véron et al., 2020).

106 Challenges to energy balance when prey become smaller could be further exacerbated in  
107 fishes if prey size also influences foraging behaviour. Here, the sardine is also an interesting  
108 model species. Sardines spontaneously modify their feeding behaviour according to the size  
109 of their prey, using diffuse filter-feeding when prey is small but direct capture when prey is  
110 large (Garrido et al., 2007, 2008). A recent long-term experiment on captive sardines showed

111 that, for the same food ration, a reduction in food size could significantly impair growth and  
112 body condition (Queiros et al., 2019). Sardines filter feeding on small particles had to consume  
113 twice as much as those capturing large particles to achieve the same growth and body  
114 condition (Queiros et al., 2019). We suspected that the two foraging modes had different  
115 energetic costs for the same degree of resource acquisition, with costs being higher for  
116 sustained aerobic swimming during filter-feeding compared to brief bursts of swimming to  
117 capture prey (Costalago & Palomera, 2014; Queiros et al., 2019). At the same time, food  
118 availability could be highly significant in the wild, filtration could be effective in very rich areas  
119 such as upwellings whereas particulate feeding might be more advantageous in areas with  
120 lower prey density (Costalago et al., 2015).

121 The current study focused on this complex predator-prey interaction in a captive population  
122 of adult sardines. We investigated the hypothesis that foraging on smaller prey would lead to  
123 increased rates of energy expenditure by sardines, and that these energy requirements would  
124 be exacerbated with warming temperature. To assess the energetic consequences of feeding  
125 sardines on prey of different sizes and at different abundances, we used group respirometry  
126 to measure rates of oxygen uptake and provided prey as commercial pellets of two different  
127 sizes at a range of ration levels. We compared animals acclimated to two temperatures within  
128 the species' thermal range, either a cool 16°C or warm 21°C. Thus, the effects on oxygen  
129 consumption of particle size and temperature were investigated according to 5 scenarios: (1)  
130 change from large to small particles at cool temperature; (2) change from large to small  
131 particles at warm temperature; (3) rise in temperature with fish fed on large particles; (4) rise  
132 in temperature with fish fed on small particles, and (5) change from large to small particles  
133 while also increasing temperature. To that end, we focussed on overall daily energetic costs

134 but also a careful comparison of energetic costs incurred during and after feeding for each  
135 scenario.

136

## 137 **2. Material and methods**

### 138 **2.1. Animal capture and husbandry**

139 Sardines were captured by commercial purse-seiner and transferred to the IFREMER Palavas-  
140 les-Flots research station, with the same fishing and husbandry procedures as described tailed  
141 in Queiros et al. (2019). Over the first week, sardines were acclimated to tanks and weaned  
142 onto commercial aquaculture pellets. They were fed with a mixture of *Artemia* nauplii and  
143 commercial aquaculture pellets (mix of 0.2 and 1.2 mm diameter), with increasing  
144 proportions of pellets and decreasing proportions of *Artemia* throughout the week,  
145 concluding exclusively with pellets. After 2-3 weeks, sardines were transferred into indoor  
146 1m<sup>3</sup> holding tanks, until experimentation. Water temperature was not set during this period  
147 but followed natural fluctuations from 15 to 20°C (SST at the time of capture was 14°C).

### 148 **2.2. Experimental design**

149 Eighty sardines were distributed among 8 experimental tanks in groups of 10 animals (volume  
150 50 L), to ensure similar distributions of body mass and condition among tanks (Fig. S1), and  
151 fish densities comparable to those of Queiros et al. (2019). Fish were acclimated to the new  
152 tanks while temperature was gradually changed from 19°C to either 16°C or 21°C over one  
153 week. Before the experiments began, fish were fed with commercial pellets twice a day, a mix  
154 of 0.2 and 1.2 mm to avoid preference bias for pellet size. These eighty sardines were used



155 for both experiments 1 and 2, described below, and these two experiments were performed  
156 sequentially in the same setup.

157 The tanks were modified to function as open automated respirometers (McKenzie et al., 2007,  
158 2012; Queiros et al., 2021) using the principles of cyclical intermittent stopped flow  
159 (Steffensen, 1989), as described below. Four tanks were held at each of the two  
160 temperatures, each set of four was supplied by water from a single reservoir where water  
161 temperature was regulated by an Ice 3000 (Aquavie) at 16°C or by a Red Line heater (Zodiac)  
162 at 21°C. Water in the reservoir was vigorously aerated, to maintain oxygen saturation and  
163 ensure thorough mixing. Water was delivered to tank respirometers by submersible pumps  
164 (Eheim 3400); within each respirometer the water was also gently but thoroughly mixed by a  
165 submersible pump (Newa Maxi 500) to avoid any thermal or oxygen gradients (see  
166 Supplementary Material and Fig. S2).

167 All respirometers were exposed to a 12L:12D photoperiod (L: light, D: darkness) with a natural  
168 sunlight spectrum and 30 minute progressive dawns and sunsets. Individual total length and  
169 body mass was measured every two weeks under anaesthesia (140 mg L<sup>-1</sup> benzocaine). To  
170 estimate total tank biomass each day, body mass gain (or loss) was assumed to be linear  
171 between successive bi-weekly measures. Total biomass was then used to adjust rations and  
172 to calculate oxygen consumption. No mortality was observed during the experiments.

## 173 **2.3. Protocols**

### 174 ***2.3.1. Experiment 1: Effects of prey size, prey abundance and temperature on daily*** 175 ***energy expenditure.***

176 To investigate effects of prey size, we offered sardines one of two commercial pellets that  
177 had similar composition in terms of lipids and proteins but differed in size, being either 0.2

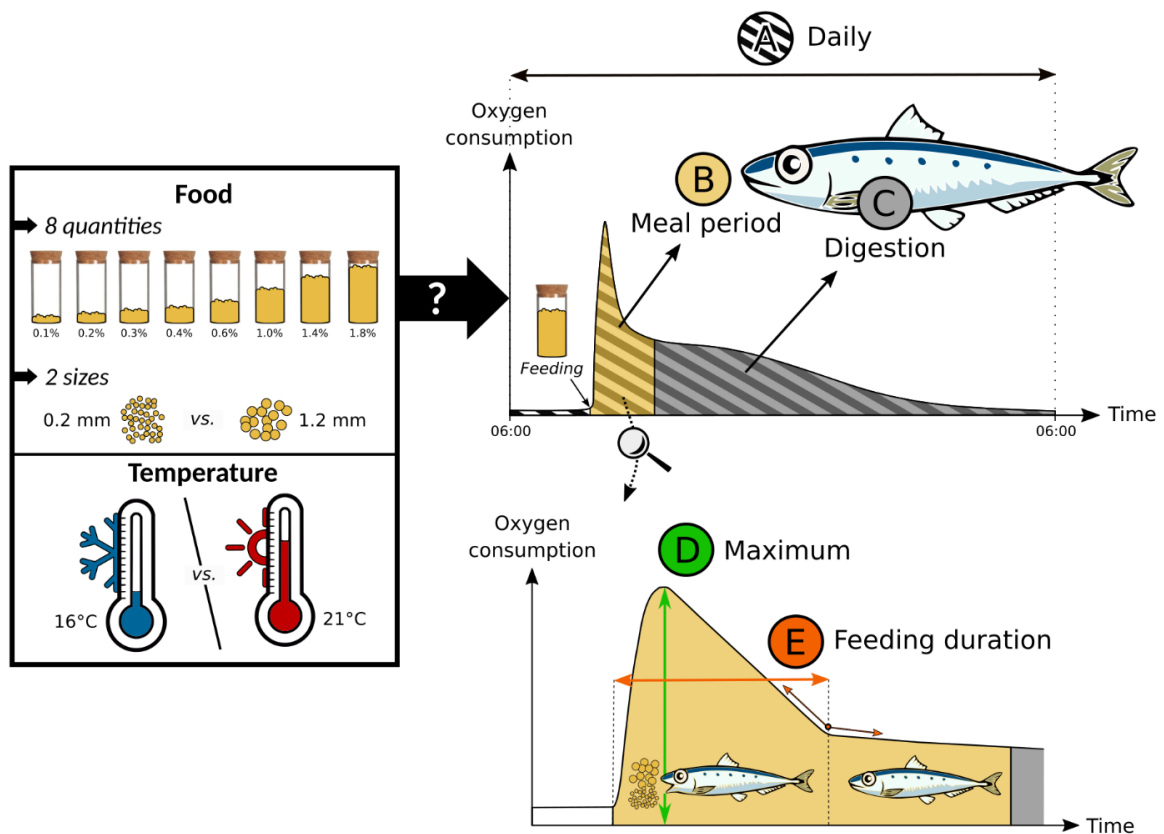
178 mm or 1.2 mm in diameter, for a period of six weeks. These sizes fall within the natural range  
179 of sardine prey (Nikolioudakis et al., 2012), but elicit two markedly different foraging modes,  
180 being either filtering on 0.2 mm pellets or particulate capture of 1.2 mm pellets (Queiros et  
181 al., 2019). Eight prey abundances were studied, as pellet rations ranging from 0.1 to 1.8 % of  
182 the total fish mass per tank: 0.1%, 0.2%, 0.3%, 0.4%, 0.6%, 1.0%, 1.4%, 1.8%. The combination  
183 of two sizes and eight rations resulted in 16 feeding treatments for each of the 2 temperatures  
184 (Figure 1). Sardines were fed once a day at 09:00 in the morning. Daily feeding treatment for  
185 a tank was randomly assigned but comprised 2 replicates of each feeding treatment per tank  
186 over the entire experiment (i.e. 8 replicates per feeding treatment x temperature).

187 The cyclical measures of oxygen uptake rate ( $MO_2$  in  $mg\ kg^{-1}\ h^{-1}$ ) provided an indirect estimate  
188 of metabolic rate and, therefore, energy use, while the sardines fed, digested and exhibited  
189 diurnal patterns of spontaneous activity. Methodological details are provided below. A  
190 continual cycle of 15 min stopped flow to measure  $MO_2$  alternated with 15 min flush with  
191 aerated water was used, except at feeding when flow was stopped for 30 min (at rations of  
192 0.1%, 0.2%, 0.3% and 0.4%) or 60 min (at rations of 0.6%, 1.0%, 1.4% and 1.8%), to ensure the  
193 entire ration was consumed before flushing. Food was distributed 5 minutes after flow was  
194 stopped, when water level had stabilized in all respirometers.

195 Bias due to behavioural responses to the act of feeding (e.g. anticipation caused by human  
196 presence near tanks at the typical feeding time) was controlled for by sham-feeding events,  
197 where the typical feeding gestures were performed but no food was provided. These shams  
198 were performed twice a day (9:00 am and 2:00 pm) for 2 days in all tanks, in the middle and  
199 at the end of experiment 1.

200           **2.3.2. Experiment 2: Effects of prey size, prey abundance and temperature on**  
201           **features of foraging behaviour.**

202   In this experiment we studied features of the foraging modes, filtration or particulate capture,  
203   in more detail, considering duration and maximum intensity (Figure 1). To this end, sardines  
204   were fed twice a day (9:00 am and 2:00 pm) for 3 weeks with 8 treatments: one of two food  
205   sizes (0.2 and 1.2 mm) at four rations (0.1, 0.2, 0.3 and 0.4% of tank biomass). The  
206   combination of two sizes and four rations resulted in 8 feeding treatments for each of the 2  
207   temperatures (16°C and 21°C). Based on the results of Experiment 1, rations were chosen not  
208   to cause satiety. Similar to Experiment 1, MO<sub>2</sub> was measured throughout and for 30 min  
209   during feeding (food distributed after 5 min). Combining these data with those of Experiment  
210   1 (Fig. S3), we obtained a total of 24 replicates per feeding treatment x temperature for food  
211   rations between 0.1% and 0.4%. Any bias due to behavioural responses to the act of feeding  
212   were assessed by two sham events, as described above.



213

214 *Figure 1: Conceptual framework of the two experiments on cocktail effects of food size (0.2*  
 215 *and 1.2 mm), food ratios (between 0.1% and 1.8% of the total biomass in tank) and*  
 216 *temperature (16°C and 21°C) on energy expenditure of sardines (daily [A], during the meal*  
 217 *period [B], during digestion [C], on the maximal intensity during feeding [D]) and on the*  
 218 *duration of the feeding activity [E].*

#### 219 **2.4. Respirometry**

220 Water oxygen levels were recorded every 5 seconds in the tank respirometers, with an O<sub>2</sub>  
 221 optode (Oxy-10 mini; PreSens Precision Sensing GmbH, /www.presens.de) and associated  
 222 software (Pre-Sens Oxy 4v2). Water O<sub>2</sub> saturation never fell below 70% during the 15 min of  
 223 stopped flow and never below 60% after feeding. Saturation was rapidly restored when the  
 224 tanks were flushed with a flow of aerated water from the reservoir.

225 Oxygen uptake by the sardines caused a linear decline in water O<sub>2</sub> concentration over time  
226 during each stopped flow phase ('closed phase'). The MO<sub>2</sub> was calculated in mg O<sub>2</sub> kg<sup>-1</sup> h<sup>-1</sup>,  
227 using least-square regression of the slope, considering oxygen solubility at the appropriate  
228 temperature (measured continuously) and salinity (measured daily); tank volume (50 L), and  
229 fish biomass (McKenzie et al., 2007). Only slopes with R<sup>2</sup> ≥ 0.95 were kept for further analyses  
230 (<5% of slopes were removed from analyses). Gas exchange across the water surface being  
231 negligible, no correction was applied when estimating sardine oxygen consumption  
232 (McKenzie et al., 2007; Queiros et al., 2021).

## 233 **2.5. Respirometry data analyses**

### 234 ***2.5.1. Basal and daily oxygen consumption***

235 Basal O<sub>2</sub> uptake rate of day<sub>i</sub> was expressed as the lowest 15%-quantile (Chabot et al., 2016a)  
236 of the daily O<sub>2</sub> consumption of the previous day (from 06:00 a.m. day<sub>i-1</sub> to 06:00 a.m. day<sub>i</sub>).  
237 This rate of oxygen uptake was then used as a baseline for calculating daily oxygen  
238 consumption on day<sub>i</sub>, expressed in mg O<sub>2</sub> kg<sup>-1</sup> d<sup>-1</sup>, as an increase from this basal rate. This  
239 normalisation avoided bias linked to a change in fish biomass during experiments, short-term  
240 effects of a previous meal, or a small change in temperature, salinity, minor human  
241 disturbance, etc. Daily MO<sub>2</sub> was calculated as the area under the curve (AUC) of MO<sub>2</sub> over  
242 time, from 06:00 a.m. and for 24 hours (Figure 1, point A) using the '*DescTools*' package in R  
243 (Andri Signorell 2021). The AUC was calculated over two periods: (i) raw data from 06:00 a.m.  
244 until noon, to catch the peak of oxygen consumption observed during the meal period and (ii)  
245 smoothed values of the oxygen consumption after 12:00 a.m. (oxygen consumption  
246 smoothed using *lowess* function) to avoid outliers due to, for example, minor disturbance in  
247 the room, that might distort daily estimations (Fig. S3).

248 **2.5.2. Oxygen consumption during feeding**

249 When focusing on effects of a meal on MO<sub>2</sub>, these were calculated relative to a control  
250 baseline that was estimated as the mean of the preceding 2.5 hours. This was done to avoid  
251 bias when either lights were turned on 1.5 hour before the 1<sup>st</sup> daily meal, or there were  
252 remnant effects of digestion of that 1<sup>st</sup> meal for the 2<sup>nd</sup> meal period. Since it took up to 2  
253 minutes to feed all tanks (i.e. between 5 and 7 minutes after the beginning of the closed  
254 phase), we first needed to establish the start of the feeding event for each tank. To do so, we  
255 identified a break in the rate of oxygen decline in the water during the initial minutes of the  
256 'closed phase', using the 'segmented' package (Muggeo, 2008). Once this was identified,  
257 oxygen consumption was calculated, in mg O<sub>2</sub> kg<sup>-1</sup> h<sup>-1</sup>, as the linear decline of oxygen  
258 concentration from there until 2 min before the end of the 'closed phase' (Figure 1, point B).

259 **2.5.3. Oxygen consumption during digestion**

260 The start of the digestion period was considered to begin 90 min after the start of the meal  
261 period, this being the maximal duration of the feeding and then flush periods across the  
262 different rations. Thus, with feeding at 09:00, the oxygen consumption during digestion was  
263 calculated, in mg O<sub>2</sub> kg<sup>-1</sup> d<sup>-1</sup>, as the AUC of the oxygen consumption over time between 10:30  
264 a.m. of day<sub>i</sub> and 06:00 a.m. of day<sub>i+1</sub> (Figure 1, point C). This oxygen consumption was  
265 expressed as an increase from the basal O<sub>2</sub> uptake of day<sub>i</sub> as estimated above.

266 **2.5.4. Maximal oxygen consumption during feeding**

267 To reveal dynamics of metabolic rate after feeding (Fig. S5), MO<sub>2</sub> was estimated as a moving  
268 average at 30 second intervals during the closed feeding period, using linear regressions over  
269 1 minute on smoothed data for 12 measures of tank oxygen concentration. This revealed the

270 maximum oxygen consumption, in  $\text{mg O}_2 \text{ kg}^{-1} \text{ h}^{-1}$ , achieved during each meal period (Figure 1,  
271 point D).

#### 272 ***2.5.5. Duration of the feeding period***

273 To estimate feeding duration, in minutes, we identified the end of the meal as the breakpoint  
274 when oxygen concentration stopped decreasing severely after feeding, taken to indicate the  
275 end of feeding-related activity (Figure 1, point E). That is, a broken-line regression was  
276 performed on oxygen consumption values calculated every 30 seconds (also every 30 seconds  
277 over 1 minute), starting at the peak of oxygen consumption as estimated above.

#### 278 **2.6. Statistical analyses**

279 Effects of food rations, prey (particle) size and temperature, on oxygen consumption and  
280 feeding duration, were assessed using linear mixed-effects models. We built a series of  
281 models including three fixed effects (food size, food ration and temperature), as well as their  
282 interactions. Because of variability among tanks within each food ration x food size x  
283 temperature treatment, we also introduced a random tank intercept effect. The best-fitting  
284 model was selected based on the lowest  $\text{AIC}_c$  values (Burnham and Anderson, 2002) following  
285 Zuur et al. (2009). When the difference between these models in  $\text{AIC}_c$  ( $\Delta\text{AIC}_c$ ) was lower than  
286 two, the most parsimonious model was selected (Burnham and Anderson, 2002). Food ration  
287 was log-transformed for models of  $\text{MO}_2$  during feeding and maximal  $\text{MO}_2$  during feeding.  
288 Then, food ration was second order polynomial transformed to model feeding activity.

289 Finally, the effects of prey (particle) size and temperature on oxygen consumption were  
290 investigated according to 5 scenarios: (1) a change from large to small particles at cool  
291 temperature; (2) a change from large to small particles at warm temperature; (3) a rise in  
292 temperature with fish fed on large particles; (4) a rise in temperature with fish fed on small

293 particles, and (5) a change from large to small particles while also increasing temperature (see  
294 arrows in Figure 2). As such effects also depend on the food ration when the interaction with  
295 food ration was significant, we performed pairwise comparison to test significance of  
296 scenarios using selected best-fitting models as previously described. Results of scenarios over  
297 food ration are expressed as absolute and relative increases. Results are indicated as mean  
298 [95% CI]. Upper and lower 95% CI values of relative differences over food ration were  
299 calculated following Kohavi et al. (2009).

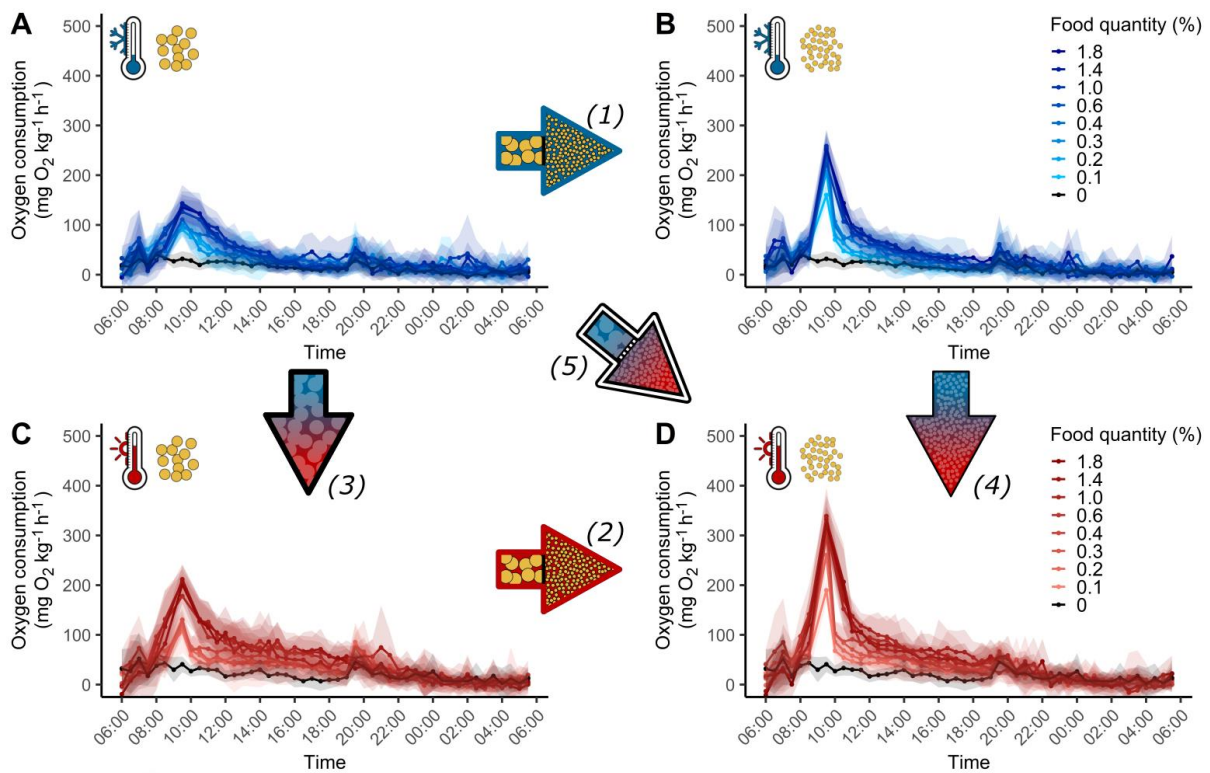
300 All data analyses were performed under R (R Core Team, 2020) and linear mixed-effects  
301 models were built using the 'lme4' package (Bates et al. 2015). All statistical tests were  
302 considered significant at p-values < 0.05.

303

### 304 **3. Results**

305 When fasted, MO<sub>2</sub> was low and statistically similar throughout the day (black curve in Figure  
306 2). When sardines were fed, MO<sub>2</sub> peaked during the feeding period after 09:00, then  
307 decreased for the rest of the day for all feeding treatments and both temperatures. Oxygen  
308 consumption increased with food ration and rearing temperature but, during feeding at both  
309 temperatures, MO<sub>2</sub> was higher when feeding on the small particles (Figure 2).





310

311 *Figure 2: Median oxygen consumption over time according to food ration for the 4*  
 312 *experimental treatments: cool temperature and large particles (A), cool temperature and*  
 313 *small particles (B), warm temperature and large particles (C) warm temperature and small*  
 314 *particles (D). Black lines are for days of fasting, blue and red represent cool (16°C) and warm*  
 315 *(21°C) temperatures, respectively. Darker lines represent higher rations of food. Arrows*  
 316 *represent the 5 scenarios for which oxygen consumption were compared: (1) change from*  
 317 *large particles to small particles at cool temperature; (2) change from large particles to small*  
 318 *particles at warm temperature; (3) rise in temperature for fish fed on large particles; (4) rise*  
 319 *in temperature for fish fed on small particles, and (5) change from large particles to small*  
 320 *particles while also increasing temperature.*

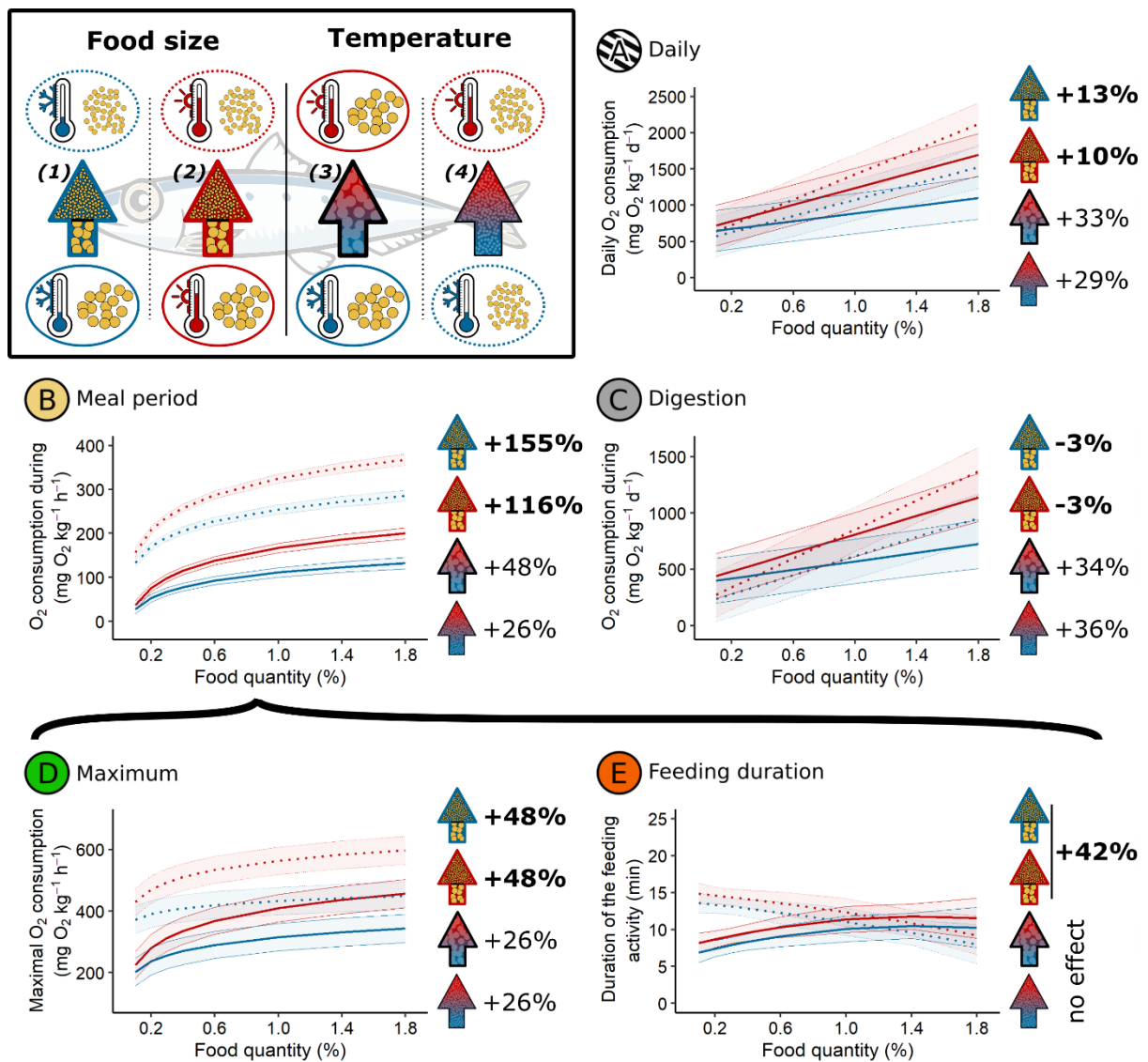
321

### 322 3.1. Daily oxygen consumption

323 During fasting days, median daily MO<sub>2</sub> (i.e. the AUC relative to basal daily oxygen  
324 consumption) was the lowest, demonstrating the clear effects that feeding and/or digestion  
325 exerted on daily energy expenditure (Figure 2, Fig. S6).

326 The best linear mixed-effect model included double interactions between food ration and  
327 food size and between food ration and temperature (Tables S1 and S2, Fig. S6, S7). An increase  
328 in ration consistently caused a significant increase in daily oxygen consumption when  
329 considering either food size or temperature. When considering only food size effects, slopes  
330 were significantly different ( $p < 0.001$ ) and the increase was smaller for large particles (slope  
331 [95% CI]; 420 [324;515] mg of O<sub>2</sub> kg<sup>-1</sup> d<sup>-1</sup>) than for small particles (714 [618;810] mg of  
332 O<sub>2</sub>/kg/d, graph *Daily* in Figure 3, Table S3).

333 When comparing large to small particles over all food rations, scenarios (1) and (2) were not  
334 significant since food size x temperature interaction was not retained during model selection.  
335 Daily MO<sub>2</sub> exhibited a mean [95% CI] relative increase of 13 [1;37]% for small particles at  
336 16°C, while this increase was 10 [3;22]% at 21°C (graph *Daily* in Figure 3, Table 1).



337

338 *Figure 3: Smooth functions of [A] daily, [B] while feeding, [C] while digesting, [D] maximal*

339 *oxygen consumptions (relative to basal oxygen consumption, see details in Material and*

340 *Methods) and [E] feeding duration according to the food ration for the 4 experimental*

341 *treatments: cool temperature and large particles (solid blue lines), cool temperature and small*

342 *particles (dotted blue lines), warm temperature and large particles (solid red lines) and warm*

343 *temperature and small particles (dotted red lines). Arrows represent the mean relative*

344 *increase of the oxygen consumption/feeding duration according to 4 scenarios summarized in*























345 *top-left panel: scenario 1 = meal modification from large particles to small particles at cool*

346 *temperature, (2) meal modification from large particles to small particles at warm*

347 *temperature, (3) increasing temperature when fish fed on large particles and (4) increasing*  
348 *temperature when fish fed on small particles. Only food size x food ration had a significant*  
349 *effect on feeding duration.*

350

351 *Table 1: Absolute and relative differences (estimates and 95% confidence intervals) for the 5 scenarios*  
 352 *for which oxygen consumption were compared: (1) change from large particles to small particles at*  
 353 *cool temperature; (2) change from large particles to small particles at warm temperature; (3) rise in*  
 354 *temperature for fish fed on large particles; (4) rise in temperature for fish fed on small particles, and*  
 355 *(5) change from large particles to small particles while also increasing temperature. Scenarios were*  
 356 *tested using pairwise comparisons. Both differences were calculated by pairwise comparisons and 95%*  
 357 *CI of relative differences were estimated following Kohavi et al. 2009. Absolute differences are given*  
 358 *in mg of O<sub>2</sub> kg<sup>-1</sup> d<sup>-1</sup> for daily and during digestion, in mg of O<sub>2</sub> kg<sup>-1</sup> h<sup>-1</sup> for maximal and during feeding,*  
 359 *and in minutes for feeding duration.*

Period	Scenario	Relative change (%)	Absolute change
Daily	 <b>Scenario (1):</b> 16°C   Large ⇒ Small	13 [1;37]	108 [6;211]
	 <b>Scenario (2):</b> 21°C   Large ⇒ Small	10 [3;22]	108 [6;211]
	 <b>Scenario (3):</b> Large   16°C ⇒ 21°C	33 [6;87]	265 [-290;821]
	 <b>Scenario (4):</b> Small   16°C ⇒ 21°C	29 [8;68]	265 [-265;821]
	 <b>Scenario (5):</b> 16°C + Large ⇒ 21°C + Small	46 [10;121]	373 [-179;926]
Meal period	 <b>Scenario (1):</b> 16°C   Large ⇒ Small	155 [46;397]	132 [120;143]
	 <b>Scenario (2):</b> 21°C   Large ⇒ Small	116 [59;209]	146 [135;157]
	 <b>Scenario (3):</b> Large   16°C ⇒ 21°C	48 [11;128]	41 [24;58]
	 <b>Scenario (4):</b> Small   16°C ⇒ 21°C	26 [16;37]	56 [38;73]
	 <b>Scenario (5):</b> 16°C + Large ⇒ 21°C + Small	220 [68;561]	187 [170;205]
Digestion	 <b>Scenario (1):</b> 16°C   Large ⇒ Small	-3 [-13;-1]	-18 [-112;76]
	 <b>Scenario (2):</b> 21°C   Large ⇒ Small	-3 [-6;-1]	-18 [-112;76]
	 <b>Scenario (3):</b> Large   16°C ⇒ 21°C	34 [-4;183]	178 [-212;569]
	 <b>Scenario (4):</b> Small   16°C ⇒ 21°C	36 [-5;214]	178 [-212;569]
	 <b>Scenario (5):</b> 16°C + Large ⇒ 21°C + Small	31 [-3;164]	160 [-228;548]
Maximal	 <b>Scenario (1):</b> 16°C   Large ⇒ Small	48 [24;84]	134 [111;157]
	 <b>Scenario (2):</b> 21°C   Large ⇒ Small	48 [29;75]	170 [148;193]
	 <b>Scenario (3):</b> Large   16°C ⇒ 21°C	26 [12;46]	72 [-15;159]
	 <b>Scenario (4):</b> Small   16°C ⇒ 21°C	26 [16;39]	108 [21;195]
	 <b>Scenario (5):</b> 16°C + Large ⇒ 21°C + Small	87 [47;148]	242 [155;329]
Duration	  Large ⇒ Small	42 [-11;339]	4 [2;6]

360

### 361        **3.2. Oxygen consumption during feeding**

362        During fasting days, median MO<sub>2</sub> at the time of sham feeding was centered on zero (Fig. S8),  
363        indicating that the increase in MO<sub>2</sub> observed during all true feeding events resulted from  
364        actual energy expenditure to feed and not from behavioural responses by the sardines to  
365        feeding gestures.

366        The MO<sub>2</sub> during feeding was significantly related to the three double interactions (food ration  
367        x food size, food ration x temperature and food size x temperature, Tables S4 and S5, Fig. S8,  
368        S9). When considering only food size effects, slopes were significantly different ( $p < 0.001$ )  
369        and the increase was smaller for large particles (slope [95% CI]; 95 [85;105] mg of O<sub>2</sub> kg<sup>-1</sup> h<sup>-1</sup>)  
370        than for small particles (129 [119;139] mg of O<sub>2</sub> kg<sup>-1</sup> h<sup>-1</sup>, Table S3).

371        Food size had a strong and significant effect on MO<sub>2</sub> during feeding. When sardines fed on  
372        small particles at 16°C, their mean [95% CI] MO<sub>2</sub> was almost multiplied by 2.5 by comparison  
373        to large particles ( $p$ -value  $< 0.001$ ), rising by 155 [46;397]%, while it doubled in sardines  
374        feeding on small particles, rising by 116 [59;209]% compared to large particles at 21°C ( $p$ -  
375        value  $< 0.001$ , graph *Meal period* in Figures 3, Table 1).

### 376        **3.3. Oxygen consumption during digestion**

377        Similar to the daily MO<sub>2</sub>, best linear mixed-effect model for MO<sub>2</sub> during digestion included  
378        double interactions between food ration and food size and between food ration and  
379        temperature (Tables S6 and S7, Fig. S10, S11). Slopes differed significantly ( $p < 0.001$ ); the  
380        lowest slope was estimated for sardines on large particles (slope [95% CI]; 300 [213;388] mg  
381        of O<sub>2</sub> kg<sup>-1</sup> d<sup>-1</sup>) while the highest slope was obtained for sardines on small particles (533  
382        [445;621] mg of O<sub>2</sub> kg<sup>-1</sup> d<sup>-1</sup>, graph *Digestion* in Figure 3).

383 When comparing large to small particles averaged over all food rations, scenarios (1) and (2)  
384 were not significant since interaction between food size and temperature was not retained in  
385 the selected model. Indeed, mean [95% CI] MO<sub>2</sub> during digestion decreased by 3 [-13,-1]% at  
386 16°C, while this decrease was 3 [-6;-1]% at 21°C (graph *Digestion* in Figure 3, Table 1).

### 387 **3.4. Maximal consumption during feeding**

388 Similar to the MO<sub>2</sub> during feeding, maximal MO<sub>2</sub> during feeding was significantly correlated  
389 with the three double interactions (Tables S8 and S9, Fig. S12, S13). Slopes differed  
390 significantly ( $p = 0.001$ ); the lowest slope was estimated for sardines on small particles (slope  
391 [95% CI]; 87 [67;106] mg of O<sub>2</sub> kg<sup>-1</sup> h<sup>-1</sup>) while the highest slope was obtained for sardines on  
392 large particles (132 [113;152] mg of O<sub>2</sub> kg<sup>-1</sup> h<sup>-1</sup>, graph *Maximum* in Figure 3, Table S3).

393 Food size had a strong and significant effect on maximal MO<sub>2</sub> during feeding since food size  
394 x temperature interaction was retained during model processing. When sardines fed on small  
395 particles at 16°C, their mean [95% CI] maximal MO<sub>2</sub> rose by 48 [24;84]% by comparison to  
396 large particles, while such increase was 48 [29;75]% at 21°C (graph *Maximum* in Figure 3,  
397 Table 1).

### 398 **3.5. Feeding duration**

399 Contrary to the previous MO<sub>2</sub> features, the selected model for the feeding duration included  
400 interaction between food ration and food size but not with temperature since food ration x  
401 temperature and food size x temperature interaction were not retained (Tables S10 and S11,  
402 Fig. S14, S15). Slopes differed significantly between the two food sizes ( $p = 0.004$ ). Indeed, the  
403 feeding duration decreased with increasing food ration when sardines fed on small particles  
404 (slope [95% CI]; -3 [-6;0] min) while it increased when sardines fed on large particles (4 [1;7]  
405 min, Table S3).

406 When comparing large to small particles averaged over all food rations, scenarios (1) and (2)  
407 were not significant since interaction between food size and temperature was not retained in  
408 the selected model. Mean [95% CI] feeding duration increased by 42 [-11;339]% (graph  
409 *Feeding duration* in Figure 3, Table 1).

### 410 **3.6. Temperature effects**

411 Temperature had significant effects on all oxygen consumptions and on the feeding duration  
412 (Tables S1 to S11). Indeed, food ration x temperature interaction was included in all selected  
413 models on MO<sub>2</sub> and food size x temperature interaction was included in models on MO<sub>2</sub>  
414 during feeding and maximal MO<sub>2</sub> during feeding. Moreover, in the model on feeding  
415 duration, temperature was retained without its interactions (Tables S10).

416 When studying interaction of food ration and temperature, slopes were significant different  
417 between the two temperatures in all MO<sub>2</sub> models and they were always smaller at cool than  
418 at warm temperature. Thus, slopes were smaller at 16°C than at 21°C for daily MO<sub>2</sub>, MO<sub>2</sub>  
419 during digestion, MO<sub>2</sub> during feeding, and maximal MO<sub>2</sub> during feeding (Table S3).  
420 Surprisingly, slopes were very similar when considering either large particles or cool  
421 temperature effects (e.g. slopes [95% CI] for daily MO<sub>2</sub>, 420 [324;515] and 413 [317;509],  
422 respectively) and either small particles or warm temperature effects (for daily MO<sub>2</sub>, 714  
423 [618;810] and 721 [626;817], respectively, suggesting similar effects of prey shrinking and  
424 temperature warming over food ration on MO<sub>2</sub> (see Table S3).

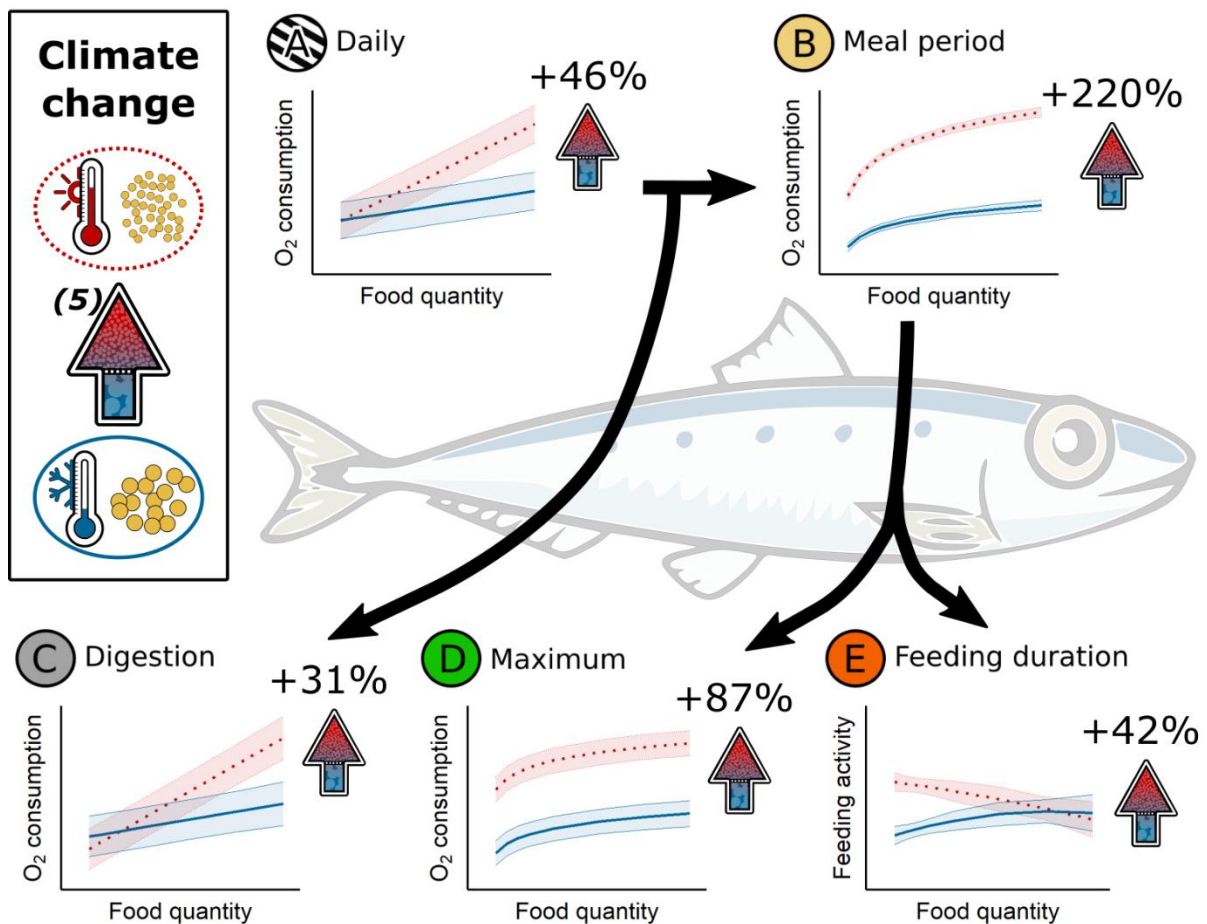
425 When comparing cool to warm conditions, in scenarios (3) and (4) there was no significant  
426 effect of temperature on daily MO<sub>2</sub>, MO<sub>2</sub> during digestion or on feeding duration, because  
427 food size x temperature interaction was not retained within selected models. On the other  
428 hand, warming effects were significant on MO<sub>2</sub> during feeding (p-values < 0.001) but only



429 scenario (4) was significant on maximal MO<sub>2</sub> during feeding (p-value = 0.02). Thus, the  
430 temperature change from 16°C to 21°C caused mean MO<sub>2</sub> during feeding to increase by 26  
431 [16;37]% in fish fed on small particles, and by 49 [11;129]% in fish fed on large particles. This  
432 temperature change caused mean maximal MO<sub>2</sub> during feeding to increase by 26 [16;39]%  
433 in sardines fed with small particles (Figure 3, Table 1).

### 434 **3.7. Cocktail effects of the global warming**

435 Smaller particle size and higher temperature resulted in a mean [95% CI] daily MO<sub>2</sub> increase  
436 of 46 [10;121]%, representing an increase of 373 [-179;926] mg of O<sub>2</sub> kg<sup>-1</sup> d<sup>-1</sup>. This increase  
437 was caused by the significant multiplication by 3 of the MO<sub>2</sub> during the meal period (220  
438 [68;561]%, representing 187 [170;205] mg of O<sub>2</sub> kg<sup>-1</sup> h<sup>-1</sup>) and higher MO<sub>2</sub> during the digestion  
439 (31 [-3;163]%, representing 160 [-228;548] mg of O<sub>2</sub> kg<sup>-1</sup> d<sup>-1</sup>). Moreover, such change caused  
440 an increase of the maximal MO<sub>2</sub> during feeding by 87 [47;148]%, representing an increase of  
441 242 [155;329] mg of O<sub>2</sub> kg<sup>-1</sup> h<sup>-1</sup>, and a longer feeding period (42 [-11;339]%, representing 4  
442 [2;6] min without temperature effect, Figure 4, Table 1).



443

444 *Figure 4: Smooth functions of [A] daily, [B] while feeding, [C] while digesting, [D] maximal*  
 445 *oxygen consumptions and [E] feeding duration according to the food ration for the 2*  
 446 *experimental treatments representing past and future environmental conditions, i.e. cool*  
 447 *temperature and large particles (solid blue lines) and warm temperature and small particles*  
 448 *(dotted red lines), respectively. Arrows represent the mean relative increase of the oxygen*  
 449 *consumption/feeding duration according to the global warming (scenario 5), i.e. meal*  
 450 *modification from large particles to small particles with increasing temperature, and*  
 451 *summarized in top-left panel (feeding duration is not significantly affected by temperature*  
 452 *increase, see Results).*

453

#### 454 **4. Discussion**

455 This study investigated how a modification of food resources under climate warming might  
456 jeopardize energy balance of small pelagic species, using sardines in the Mediterranean Sea  
457 as a case study. To do so, we used in-vivo group respirometry to investigate the effects of  
458 prey (food) size and availability (ration) on sardine energy expenditure, and how this was  
459 influenced by temperature. Our results demonstrate that both food size and temperature had  
460 significant effects on multiple measures of energy expenditures, over daily and hourly  
461 timescales. While temperature significantly increased expenditures overall, food size had a  
462 major impact on energy expenditure for activity during feeding itself. That is, the results  
463 indicate that food resources and temperature are major environmental drivers that can  
464 dramatically increase energy expenditures of fishes and disturb their energy balance in a  
465 scenario of future climate change, in warmer waters with smaller prey. As such, the results  
466 also provide experimental evidence that such challenges to energy balance may contribute to  
467 the ongoing shrinking of fish populations.

468 Daily oxygen consumption measured in this study was corrected against a baseline of  
469 standard metabolism (Chabot et al., 2016a), so represents daily energy expenditure on  
470 activity. The results suggested greater expenditure for days where sardines fed on small  
471 particles, due to higher oxygen consumption during either feeding and/or digestion. The very  
472 marked increase in oxygen consumption during actual feeding on small items must reflect  
473 different costs of foraging mode, with filtering being more expensive than particulate feeding.  
474 While this confirms our hypothesis and helps explain the decreased growth and body  
475 condition of sardines fed for an extended period on small items (Queiros et al., 2019), the  
476 magnitude of the effect is quite remarkable. Both MO<sub>2</sub> and duration of the meals provide a

477 more detailed understanding of the widely different energy costs of the two foraging modes.  
478 First, the higher maximal MO<sub>2</sub> when fish fed on small particles indicates greater energy  
479 requirements for the continuous aerobic swimming in filter-feeding compared to rapid bursts  
480 to capture large particles (Costalago & Palomera, 2014). Queiros et al. (2019) had already  
481 noted that the duration of feeding activity was longer when sardines fed on small particles, it  
482 presumably represents the time needed to filter the entire tank volume and, therefore, might  
483 not be expected to change much with ration. It is interesting therefore that feeding duration  
484 on small particles was in fact lower at low or high rations than at intermediate ones. Low  
485 duration at low ration might suggest rapid loss of interest if food acquisition was very poor,  
486 while at high ration it might indicate satiation. For particulate feeding, more particles to catch  
487 should translate into longer duration, which was observed until a ration threshold where a  
488 plateau would indicate satiation. Overall, we expected feeding duration to be longer on small  
489 particles at low ration but longer on large particles, but this was only true for rations below  
490 0.6 %, after which duration was similar for both particle sizes. Finally, all these results indicate  
491 that higher energy expenditure by sardines filter feeding on small particles can explain why  
492 they would have to eat twice as much as when feeding on large pellets to achieve similar  
493 growth or body condition (Queiros et al. 2019).

494 Our finding that oxygen consumption during the digestion increased with the food ration, for  
495 both particle sizes, presumably reflects the so-called specific dynamic action of feeding (SDA)  
496 response (McCue, 2006). This reflects the energy needed for the digestion, absorption and  
497 assimilation of a meal (Chabot et al., 2016b), hence the energetic 'costs of growth'. Therefore,  
498 larger meals require greater energy investment but then provide a great return in terms of  
499 tissue accretion and growth (Fu et al., 2005a, 2005b; Jordan & Steffensen, 2007; Norin & Clark,

2017). The fact that a doubling of ration from 0.4% to 0.8% only caused a 17% or 55% increase in apparent SDA (based on estimated slopes), in fish fed large items at 16 °C or small items at 21°C, respectively, might seem limited. This increase with doubling of ration is low compared to other fish species (see Secor, 2009). Furthermore, the high surface area to volume ratio of small particles should speed up digestion by promoting enzymatic processes and, thereby, reduce a part of digestion costs (discussed in Legler et al., 2010). On the other hand, a large SDA response can indicate that lots of nutrients were assimilated, notably amino acids for protein synthesis, with high costs of turning these into tissues but that reflect robust growth (Fraser & Rogers, 2007; McCue, 2006; Secor, 2009). That is, a large SDA would imply good growth, which is coherent with the fact that sardines fed on large particles exhibited higher growth and greater condition in previous studies (see Queiros et al. 2019).

There is another mechanism that might increase energy expenditure during 'digestion' of large particles, being the costs of recovery from rapid bursts of anaerobic swimming used for prey capture. The metabolic cost of such recovery, so-called 'excess post-exercise oxygen consumption' (EPOC) can be divided into 3 phases in fishes: rapid, plateau, slow (Zhang et al., 2018). While the rapid phase is very short (< 1 hour), both plateau and slow phases can require several hours to return to standard metabolism. This can be more than 10 hours for salmon although the duration is certainly species dependent (C. G. Lee et al., 2003; Li et al., 2020; Plambech et al., 2013; Svendsen et al., 2012; Zhang et al., 2018). Considering that we estimated energy expenditure of digestion starting at 1.5 hours after providing the meal, this would omit the rapid phase of EPOC, although the phenomenon may have contributed to the final phases of metabolic costs of feeding. A potential role for EPOC in costs of feeding on large prey remains to be proven, since studies on individual sardines are technically extremely

523 challenging. Overall, oxygen consumption due to digestion was lower for small compared to  
524 large particles, but the magnitude of the difference was much less than for the activity costs  
525 of feeding. Therefore, daily energy expenditure was mostly affected by what happened during  
526 the meal. The benefit of digesting small prey (in terms of energy expenditure for a same food  
527 ration) remained too weak to counterbalance the increased energy to capture shrinking  
528 plankton in the wild. Furthermore, the warm temperature also significantly increased  
529 metabolic rates, leading to overall higher energy expenditure during both meals and digestion  
530 (Clarke & Fraser, 2004; Seebacher et al., 2015), whatever the food size or ration. Higher  
531 energetic cost for digestion at warm temperatures has been reported for tunas, another  
532 species that swims continuously (Klinger et al., 2016) although relationships between costs of  
533 digestion and temperature are not necessarily linear (McKenzie et al., 2013; Tirsgaard et al.,  
534 2015).

535 Our study applied relatively short thermal acclimation times, which might tend to  
536 overestimate temperature effects. When natural populations are allowed to acclimatise over  
537 generations, baseline metabolism may show a much less marked effect of temperature (e.g.  
538 Wootton et al. 2022). Such intergenerational experiments are not feasible for the  
539 Mediterranean sardine because their life cycle cannot be completed in captivity. Our  
540 experimental temperatures (16°C and 21°C) were well within the range that sardines have  
541 experienced in the Gulf of Lions over the last 40 years (12 - 24°C ; Feuilleley et al. 2020) and  
542 our rate of temperature change was slow (< 0.5°C/day) allowing acclimation at an ecologically  
543 realistic pace. Furthermore, by expressing oxygen consumption as a relative increase from  
544 MO<sub>2</sub> baseline (the baseline was estimated daily as the lowest 15%-quantile rate for daily MO<sub>2</sub>  
545 and MO<sub>2</sub> during digestion, and as the mean of the preceding 2.5 hours for MO<sub>2</sub> while

546 feeding), effects of temperature on baseline metabolism were been taken into account in our  
547 study. Finally, although the effects of temperature were significant, potential acclimation  
548 across a few generations leading to similar baseline metabolism between generations would  
549 reinforce our results on the effects of food size.

550 Although food size had only quite minor effects on daily energy use, increasing it by 10 [3;22]%  
551 in fish fed on small particles at 21°C, long-term effects may be significant. Furthermore,  
552 sardines may feed continuously in the wild, not only once or twice a day, which would  
553 increase consequences of differences in energy expenditure during feeding. In the wild,  
554 sardines face predation and pathogens that require energy expenditure. Therefore, higher  
555 daily energy expenditure for feeding may well impair energy balance in the wild, resulting in  
556 less energy allocated towards survival and growth. For instance, lower swimming  
557 performance due to low energy reserves (e.g. swimming endurance (Martínez et al., 2003,  
558 2004)) could isolate leaner individuals from schools, leading to a vicious circle, with lower  
559 food foraging and thus reinforcing lower energy reserves. Nonetheless, calorie-restricted  
560 sardines display better phenotypic plasticity to face fasting, which improves their ability to  
561 reduce their metabolic energy expenditures during long-term fasting (Queiros et al., 2021).  
562 Further, mitochondria from sardines fed with small particles exhibited lower basal oxidative  
563 activity but higher efficiency of ATP production than those fed with large particles, a  
564 mechanism that should help them spare energy (Thoral et al., 2021). Nevertheless, although  
565 sardines may display plasticity or adaptation that ameliorates the energetic consequences of  
566 smaller prey and warmer temperatures, the situation of sardine populations in the Gulf of  
567 Lions remains very concerning.

568

569 **Conclusion**

570 This study supports the hypothesis of bottom-up control to explain the profound shrinking of  
571 small pelagic fish communities in the Gulf of Lions and is a hypothesis worth exploring to  
572 explain the spread of this phenomenon throughout the Mediterranean (Albo-Puigserver et  
573 al., 2021; Brosset et al., 2017) to new ecosystems, and to species higher in the food web  
574 (Bensebaini et al., 2022; Véron et al., 2020). Altogether, the results indicate that energy  
575 balance can be a major mechanism explaining shrinking of fish populations globally. Declines  
576 in prey size could impact the energy balance of individuals when their energy expenditures  
577 are increased by warmer temperatures, with future projections of prey resources predicting  
578 a decline of prey biomass and quality.



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583 improve the manuscript.

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586

587 **Data availability**

588 Data and scripts are available here: <https://zenodo.org/record/8413664>.

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881

## Supplementary material

882

883

### 884 **Material and methods**

885 Each respirometer was shaped by a vertical PVC pipe and included three perforated semi-cylinders of  
886 PVC pipe in its interior section (Fig. S2). An external pump was connected to this system to collect  
887 water from one semi-cylinder and deliver it to the two other semi-cylinders, so water was constantly  
888 mixed in the respirometer (yellow arrows in Fig. S2). Outside of measurement period, respirometer  
889 water was renewed with water pumped from continuously aerated buffer tanks (300 L) into the same  
890 hemispheres that the external pump to maintain dissolved O<sub>2</sub> concentrations close to saturation (i.e.  
891 100 %)(white arrows in Fig. S2). Excess of water was collected through a PVC pipe and returned to its  
892 given buffer tanks. These buffer tanks were supplied with the same seawater and the temperatures  
893 were maintained equal to either 16°C (cool) or 21°C (warm), maintaining constant temperature in  
894 respirometers. Pumps in buffer tanks were controlled by an electrical timer and were turned off during  
895 measurements and water level settled at the overflow to provide a constant volume (50 L). Large buffer  
896 tanks were continuously supplied with water pumped from the sea, filtered through sand filter (30–40  
897 µm) and sterilized with UV light. The photoperiod was adjusted to 12-12h cycle.

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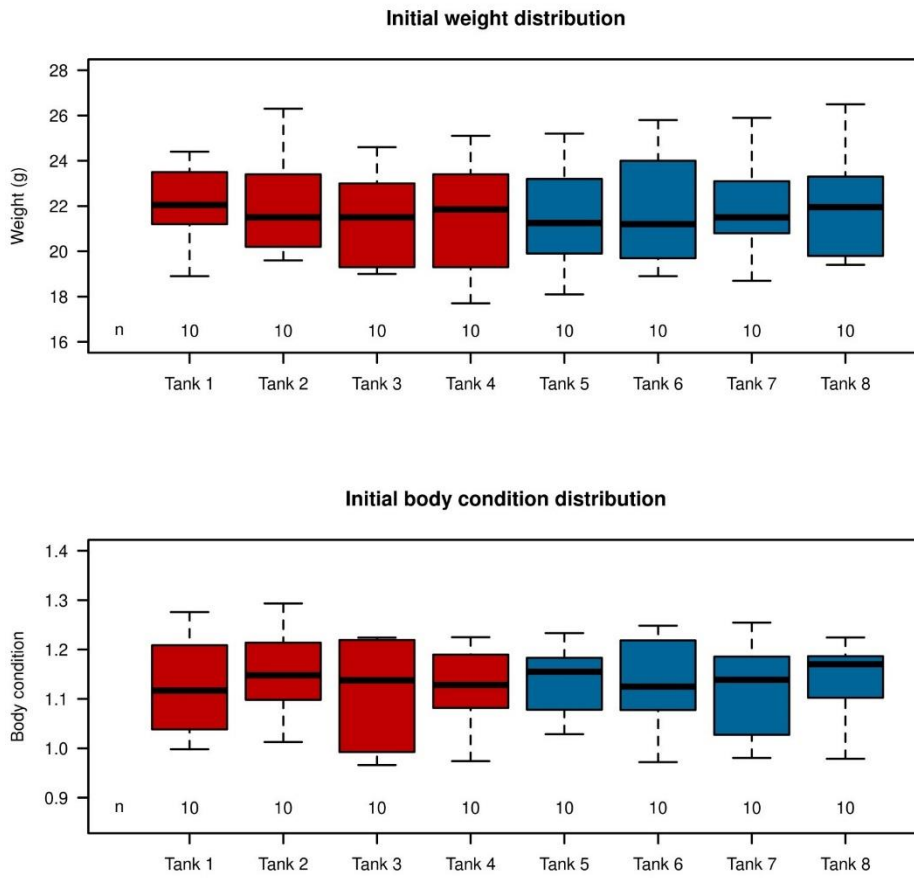
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905 **Figures**



906

907

908 Fig. S1: Distribution of body mass (A) and condition (B) of sardines for each tank at the beginning of the experiment.

909 Colors refer to the water temperature in each tank i.e. either 21°C (in red) or 16°C (in blue). The number of

910 individuals is given by n at the bottom of the panels.

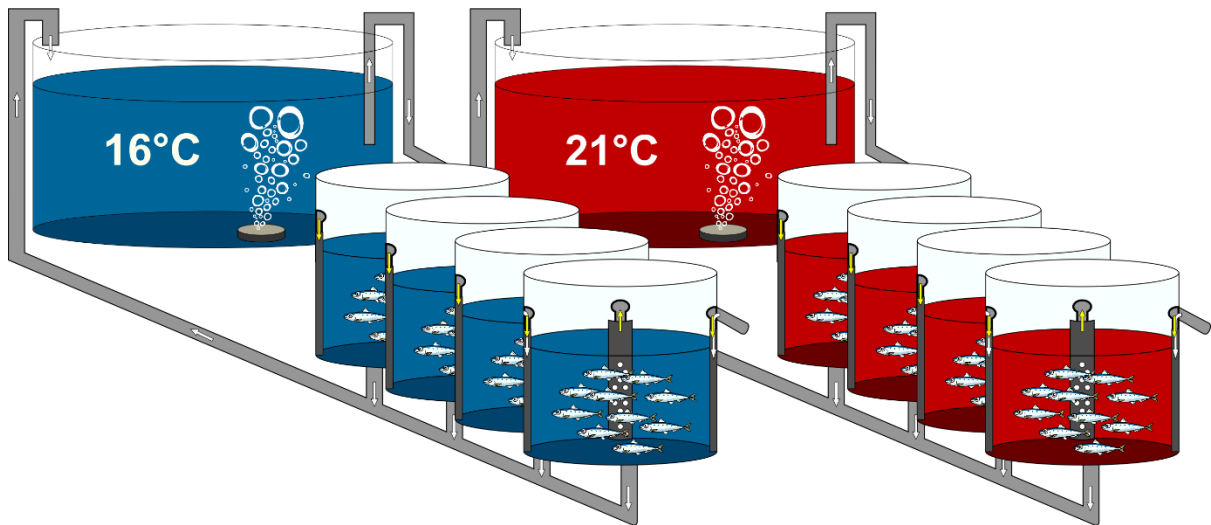
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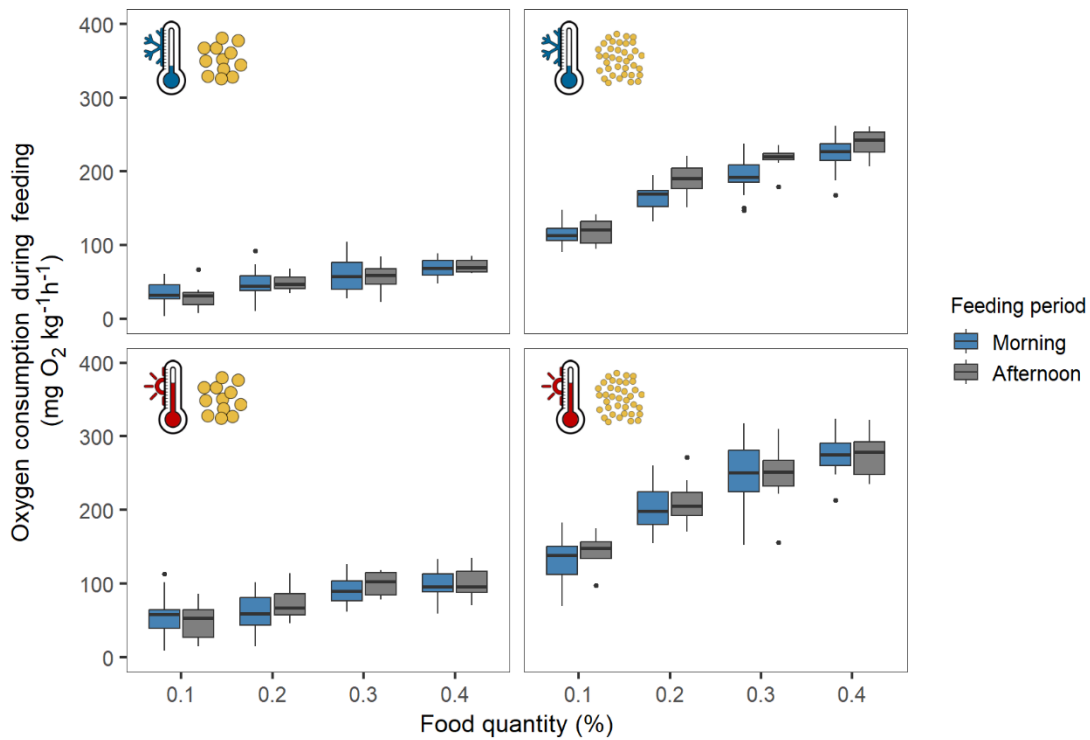
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918 *Fig. S2: The entire setup was composed by (i) 2 large aerated buffer tanks that homogenized water quality between*  
919 *tanks and maintained constant water temperature over time and (ii) 8 small experimental tanks. Each 50L*  
920 *experimental tank, in which 10 individuals were reared, was modified as automated respirometer with a continuous*  
921 *water flow that ensured the homogeneity of the water parameters (indicated by yellow arrows). The renewal and*  
922 *excess water flows are indicated by white arrows. The two colors refer to the two water temperatures: 21°C in red*  
923 *and 16°C in blue.*

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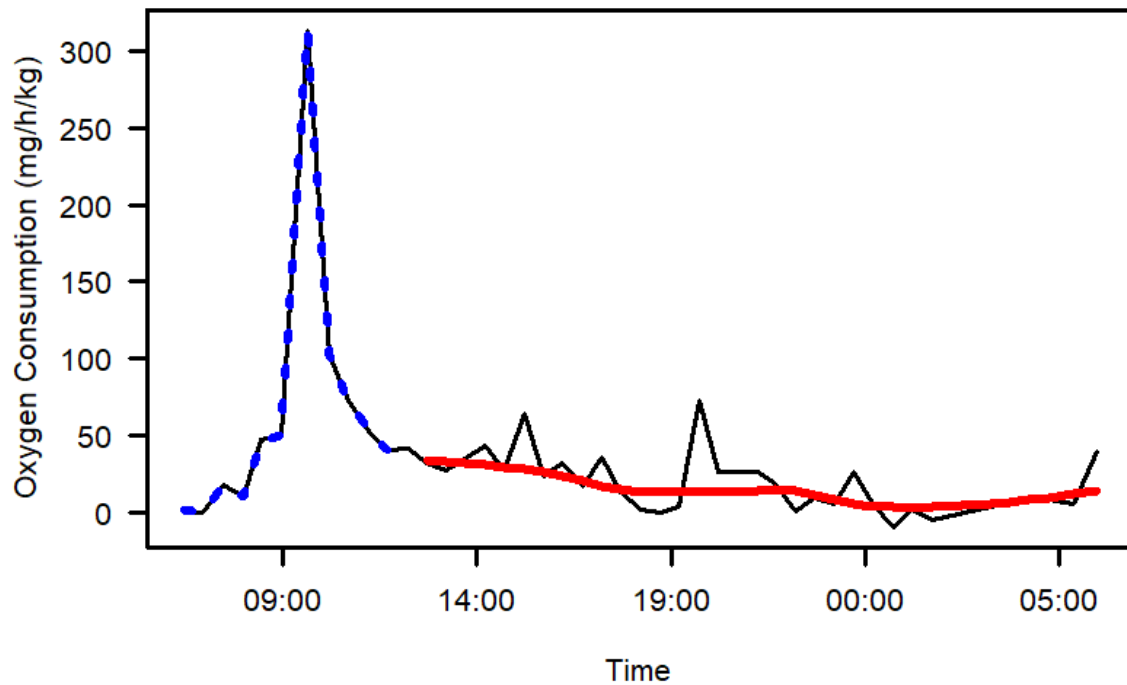
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929 *Fig. S3: Boxplots of oxygen consumption during the 1<sup>st</sup> meal in the morning (in blue) and the 2<sup>nd</sup> meal in the*  
 930 *afternoon (in grey) for the four feeding treatment x temperature (cool temperature and large particles, cool*  
 931 *temperature and small particles, warm temperature and large particles and warm temperature and small particles).*  
 932 *Oxygen consumptions have been calculated relative to a control baseline that was estimated as the mean of the*  
 933 *preceding 2.5 hours. This calculation avoided bias when lights were turned on 1.5 hour before the 1<sup>st</sup> daily meal,*  
 934 *and also potential remnant effects of digestion of that 1<sup>st</sup> meal for the 2<sup>nd</sup> meal period. No significant effect of the*  
 935 *period (morning vs. afternoon) on the oxygen consumption over feeding treatment x temperature was found*  
 936 *(ANOVA,  $p = 0.63$ ).*

937

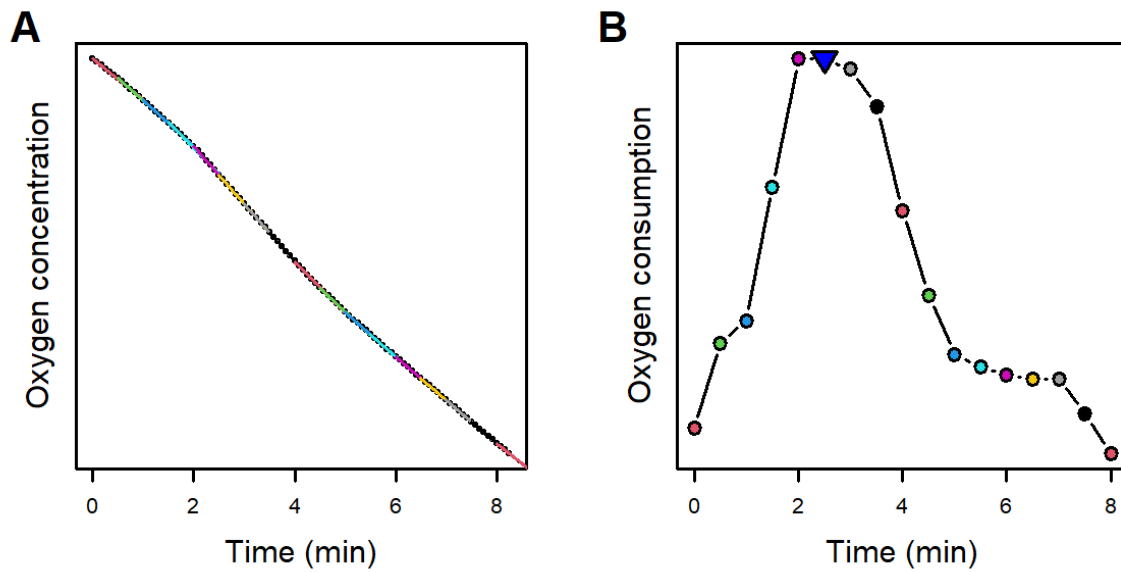
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940 *Fig. S4: Oxygen consumption over time from 06:00 a.m. and for 24 hours. The black line represents the original*  
941 *data of the oxygen consumption. The estimation of the daily oxygen consumption was calculated over a*  
942 *combination of two periods: (1) following original data from 06:00 a.m. until noon to catch the peak of oxygen*  
943 *consumption observed during the meal period (dashed blue line) and (2) smoothed values of the oxygen*  
944 *consumption after noon to avoid any outlier (e.g. peak at 19:30) that could distort daily consumption estimation*  
945 *(solid red line).*

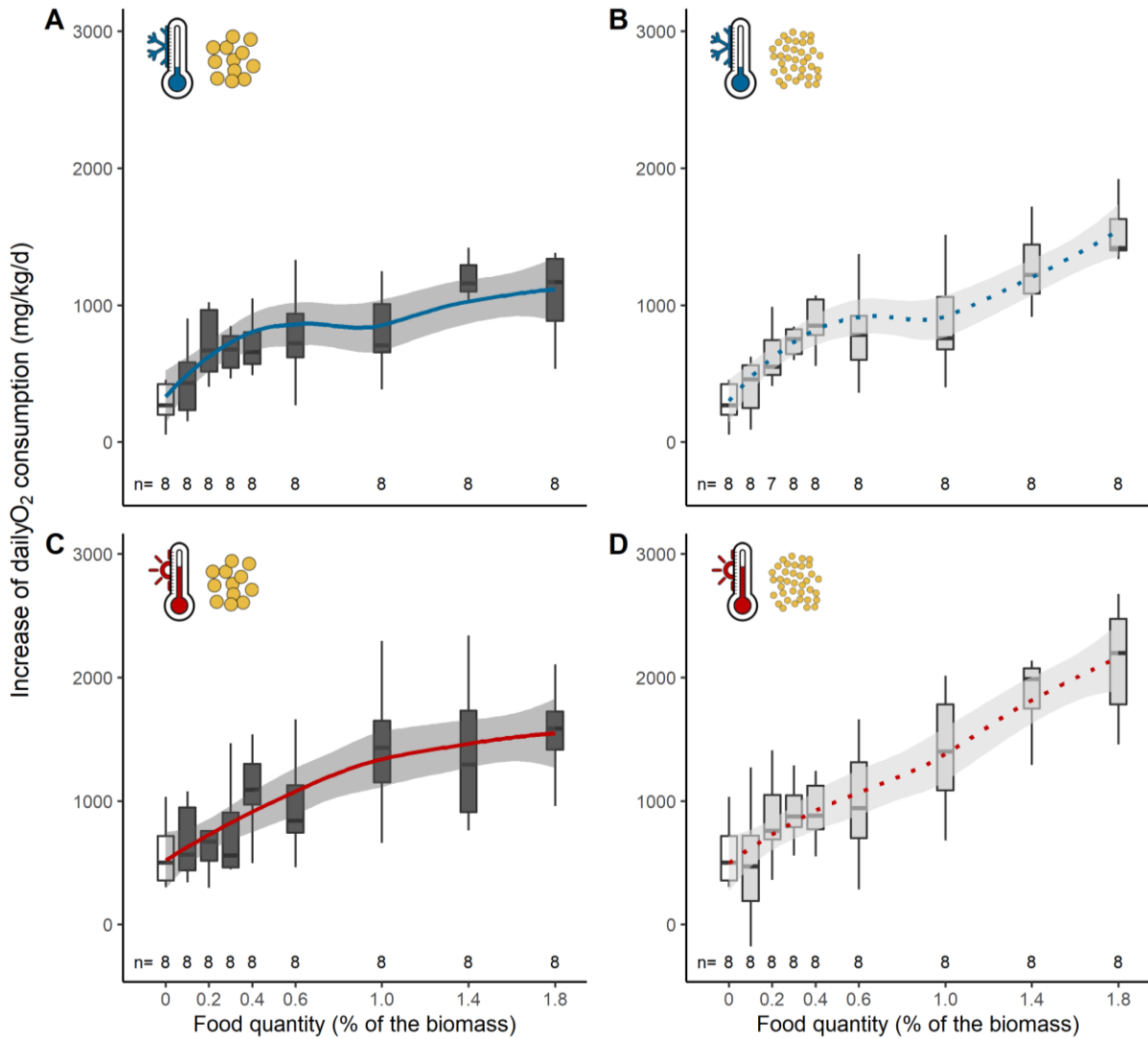
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*Fig. S5: Smoothed oxygen concentration (A) and oxygen consumption (B) after the start of the meal. Oxygen consumptions were estimated every 30 seconds using linear regressions over 1 minute on smoothed oxygen concentrations (lines in A and corresponding points in B). The blue triangle represents the maximal oxygen consumption reached during the meal.*



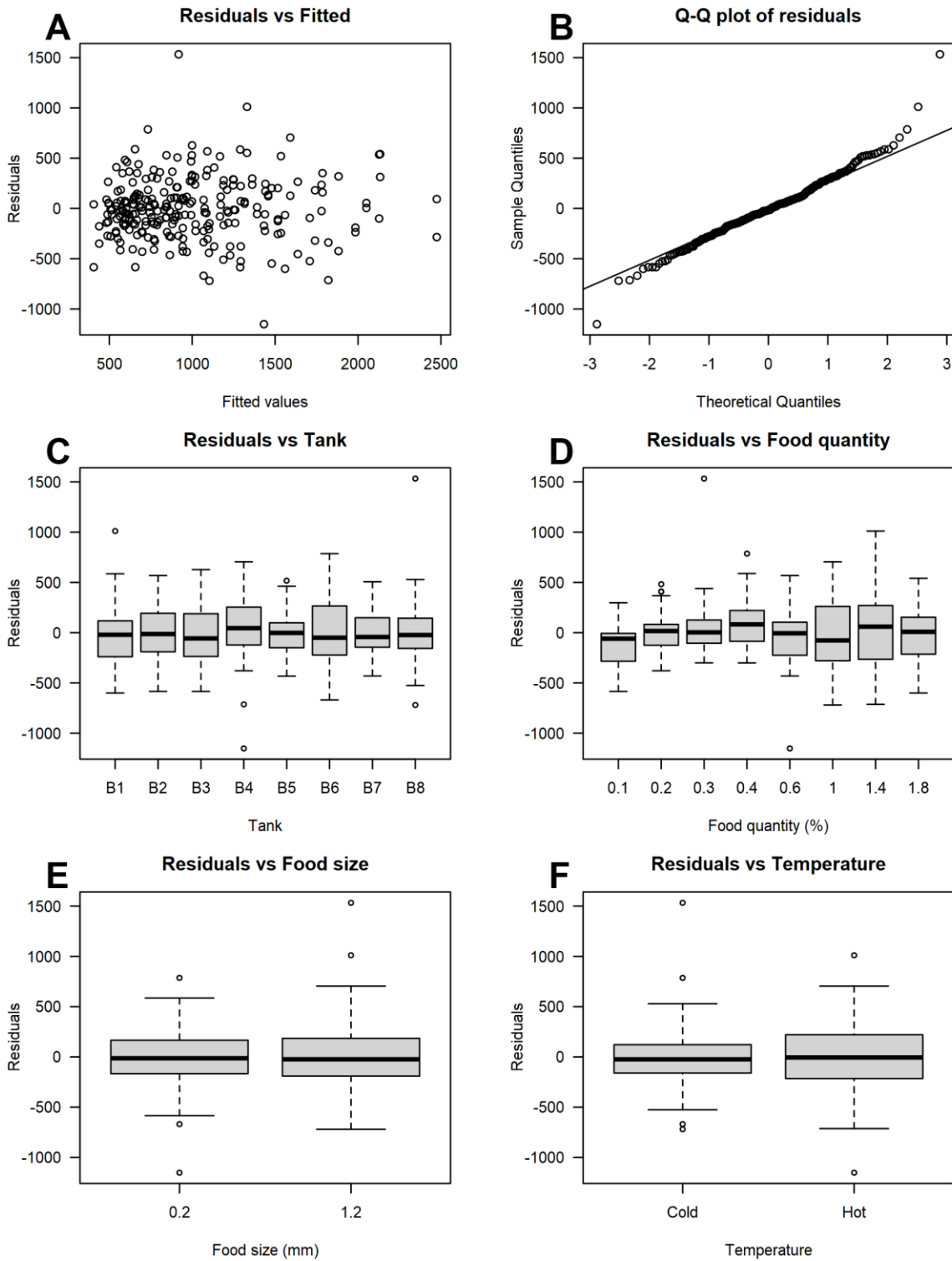


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958 *Fig. S6: Boxplot of the daily oxygen consumption increase (relative to basal oxygen consumption and due to*  
 959 *movement, food foraging and digestion) according to the food ration for the 4 experimental treatments: cool*  
 960 *temperature and large particles (A), cool temperature and small particles (B), warm temperature and large particles*  
 961 *(C) warm temperature and small particles (D). Dotted (for small particles) and solid lines (for large particles)*  
 962 *represent the smooth increase of the median daily oxygen uptake in relation to the food ration. Grey bands*  
 963 *represent 95% confidence intervals of smooth increases. White colour represents data during days of fasting, while*  
 964 *light and dark grey represent small and large particles, respectively. Blue and red colours represent cool (16°C)*  
 965 *and warm (21°C) temperatures, respectively. Only slopes (i.e. oxygen consumption) with R squared  $\geq 0.95$  were*  
 966 *used. Sample size (i.e. number of days) is given as 'n' at the bottom of each panel. Outliers were removed for*  
 967 *clarity purpose.*

968



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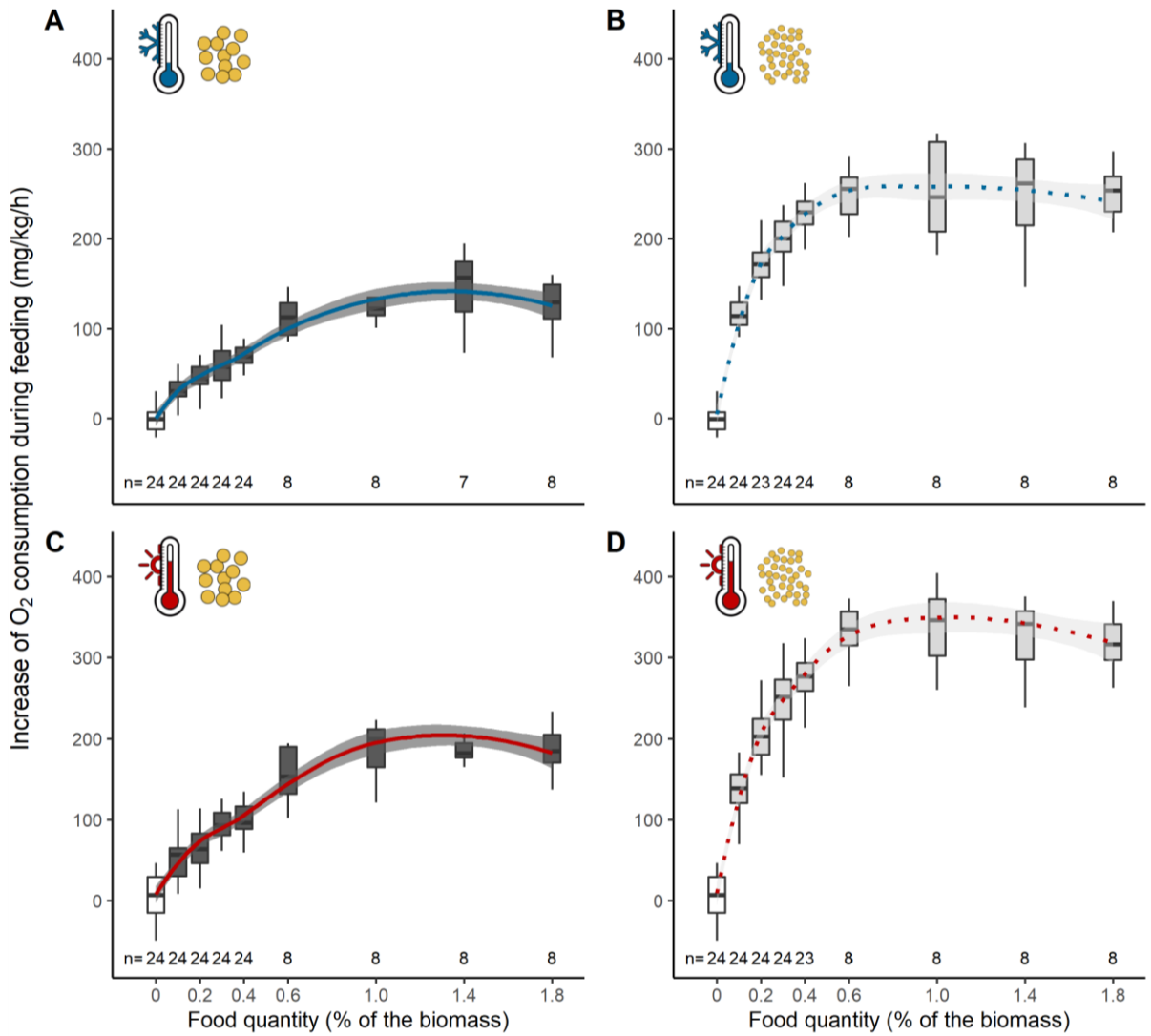
971 *Fig. S7: Diagnostic plots of the selected model for the daily oxygen consumption. A: Fitted values versus residuals.*

972 *B: Q-Q plot of the residuals. C: Residuals versus Tank ID. D: Residuals versus Food ration. E: Residuals versus*

973 *Food size. F: Residuals versus Temperature.*

974

**B** Meal period

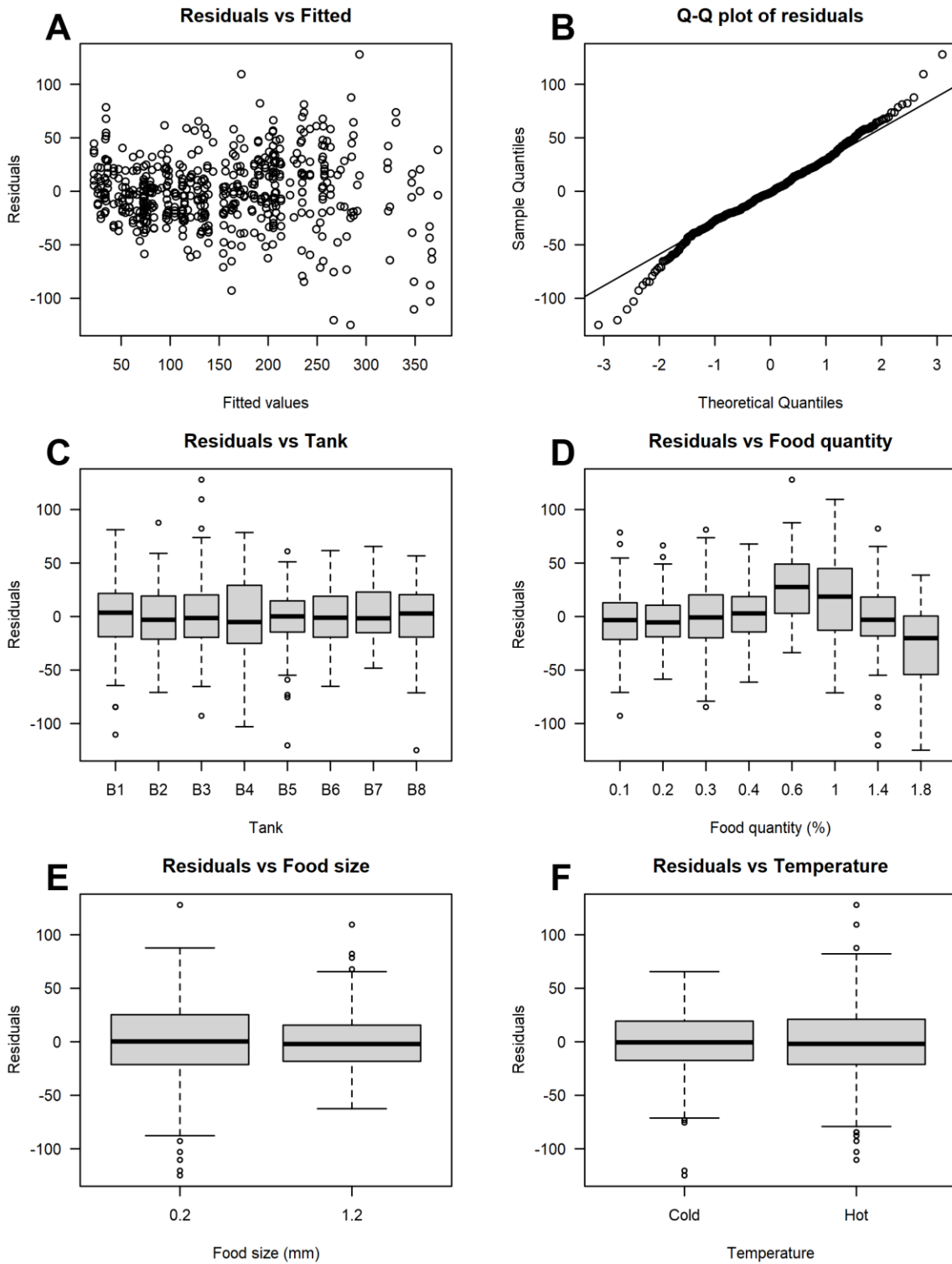


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977 *Fig. S8: Boxplot of the oxygen consumption increase during feeding (relative to the previous 2.5h oxygen*  
 978 *consumption) according to the food ration for the 4 experimental treatments: cool temperature and large particles*  
 979 *(A), cool temperature and small particles (B), warm temperature and large particles (C) warm temperature and*  
 980 *small particles (D). Dotted (for small particles) and solid lines (for large particles) represent the smooth increase of*  
 981 *the median daily oxygen uptake in relation to the food ration. Grey bands represent 95% confidence intervals of*  
 982 *smooth increases. White colour represents data during days of fasting, while light and dark grey represent small*  
 983 *and large particles, respectively. Blue and red colours represent cool (16°C) and warm (21°C) temperatures,*  
 984 *respectively. Only slopes (i.e. oxygen consumption) with R squared  $\geq 0.95$  were used. Sample size (i.e. number of*  
 985 *meals) is given as 'n' at the bottom of each panel. Sample size is greater for food ration below 0.6% because*  
 986 *results from experiments 1 and 2 are gathered here (n = 8 and n = 16 for experiment 1 and experiment 2,*  
 987 *respectively). Outliers were removed for clarity purpose.*

988



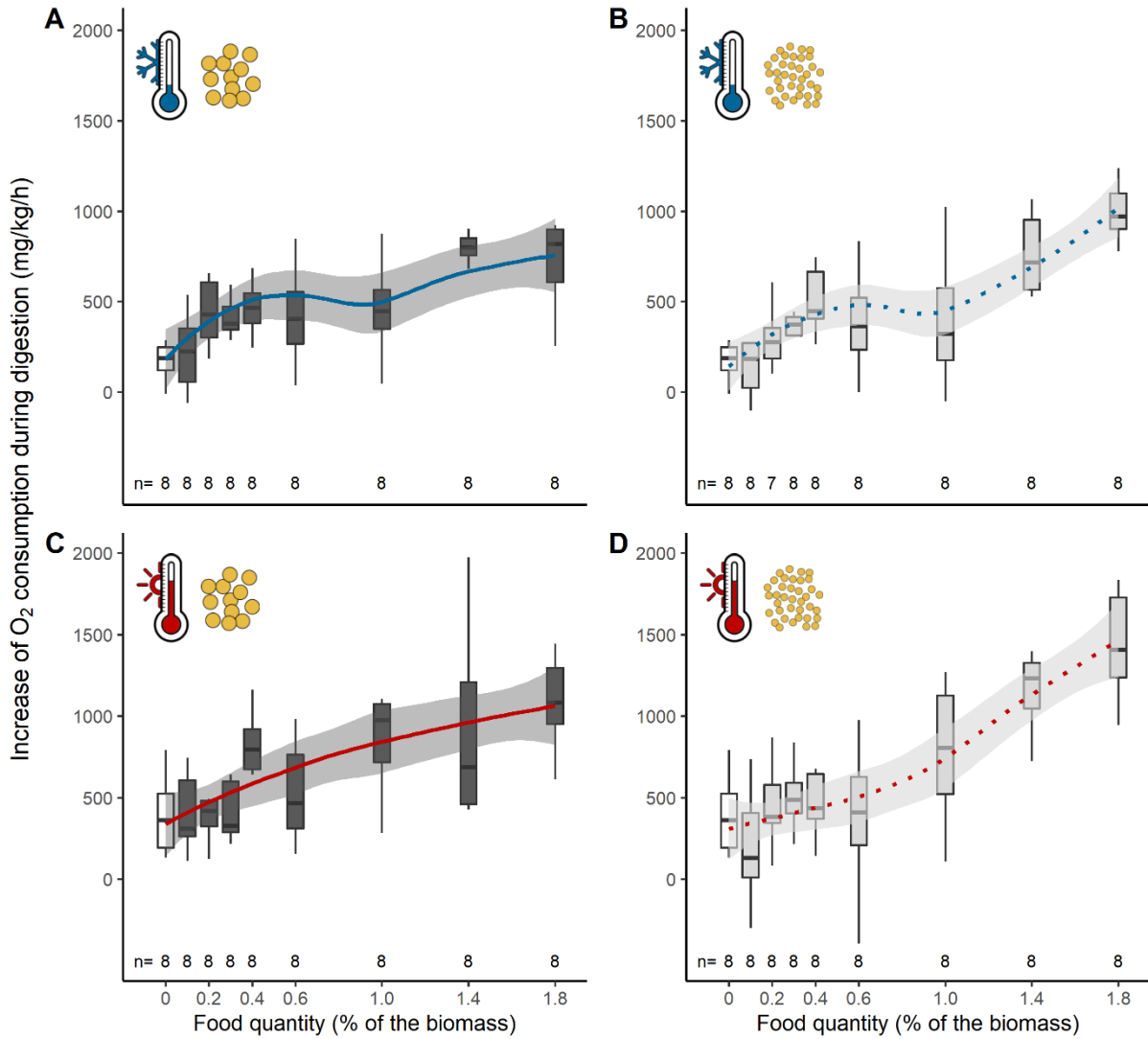
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991 *Fig. S9: Diagnostic plots of the selected model for the oxygen consumption during feeding. A: Fitted values versus*  
 992 *residuals. B: Q-Q plot of the residuals. C: Residuals versus Tank ID. D: Residuals versus Food ration. E: Residuals*  
 993 *versus Food size. F: Residuals versus Temperature.*

994

# C Digestion

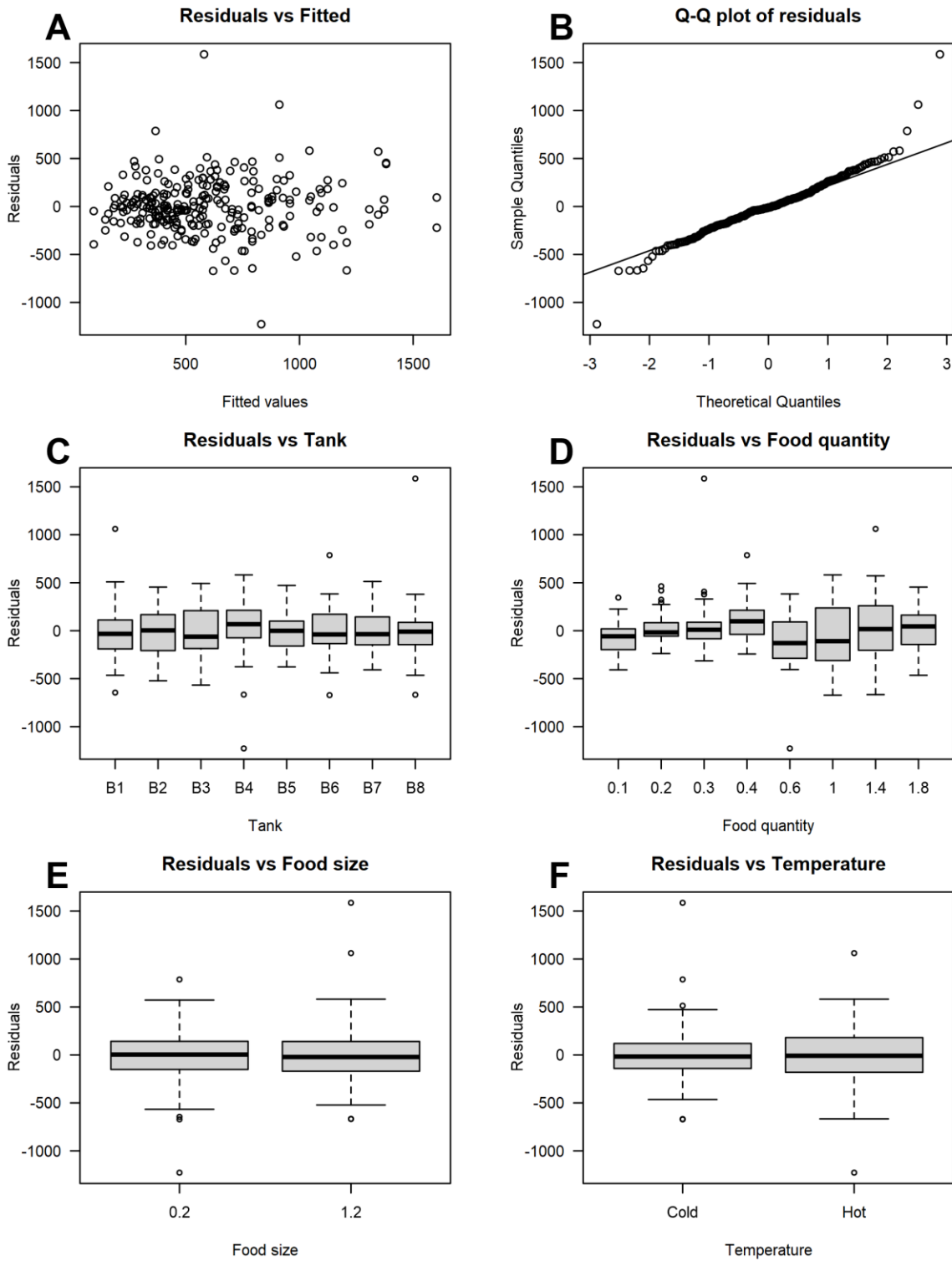


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997 *Fig. S10: Boxplot of the oxygen consumption increase (relative to basal oxygen consumption) during the digestion*  
 998 *(here 1.5 hour after the beginning of the meal period) according to the food ration for the 4 experimental treatments:*  
 999 *cool temperature and small particles (A), cool temperature and large particles (B), warm temperature and small*  
 1000 *particles (C) warm temperature and large particles (D). Dotted (for small particles) and solid lines (for large*  
 1001 *particles) represent the increase of the median oxygen uptake during digestion in relation to the food ration. Grey*  
 1002 *bands represent 95% confidence intervals of smooth increases. White color represents data during days of fasting,*  
 1003 *while light and dark grey represent small and large particles, respectively. Blue and red colors represent cool (16°C)*  
 1004 *and warm (21°C) temperature, respectively. Sample size (i.e. number of meals) is given as 'n' at the bottom of*  
 1005 *each panel.*

1006



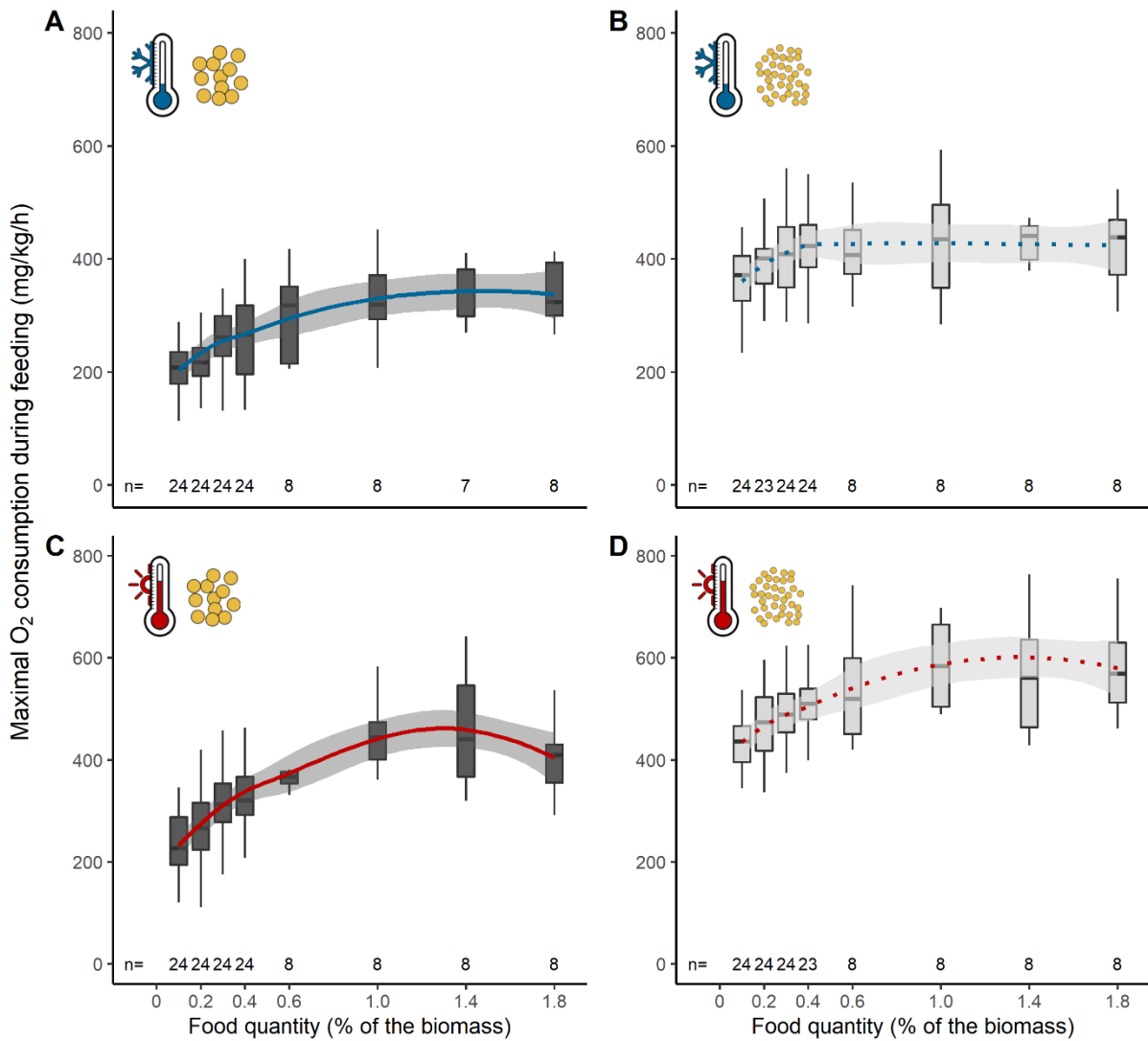
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1009 *Fig. S11: Diagnostic plots of the selected model for the oxygen consumption during digestion. A: Fitted values*  
 1010 *versus residuals. B: Q-Q plot of the residuals. C: Residuals versus Tank ID. D: Residuals versus Food ration. E:*  
 1011 *Residuals versus Food size. F: Residuals versus Temperature.*

1012

# D Maximum

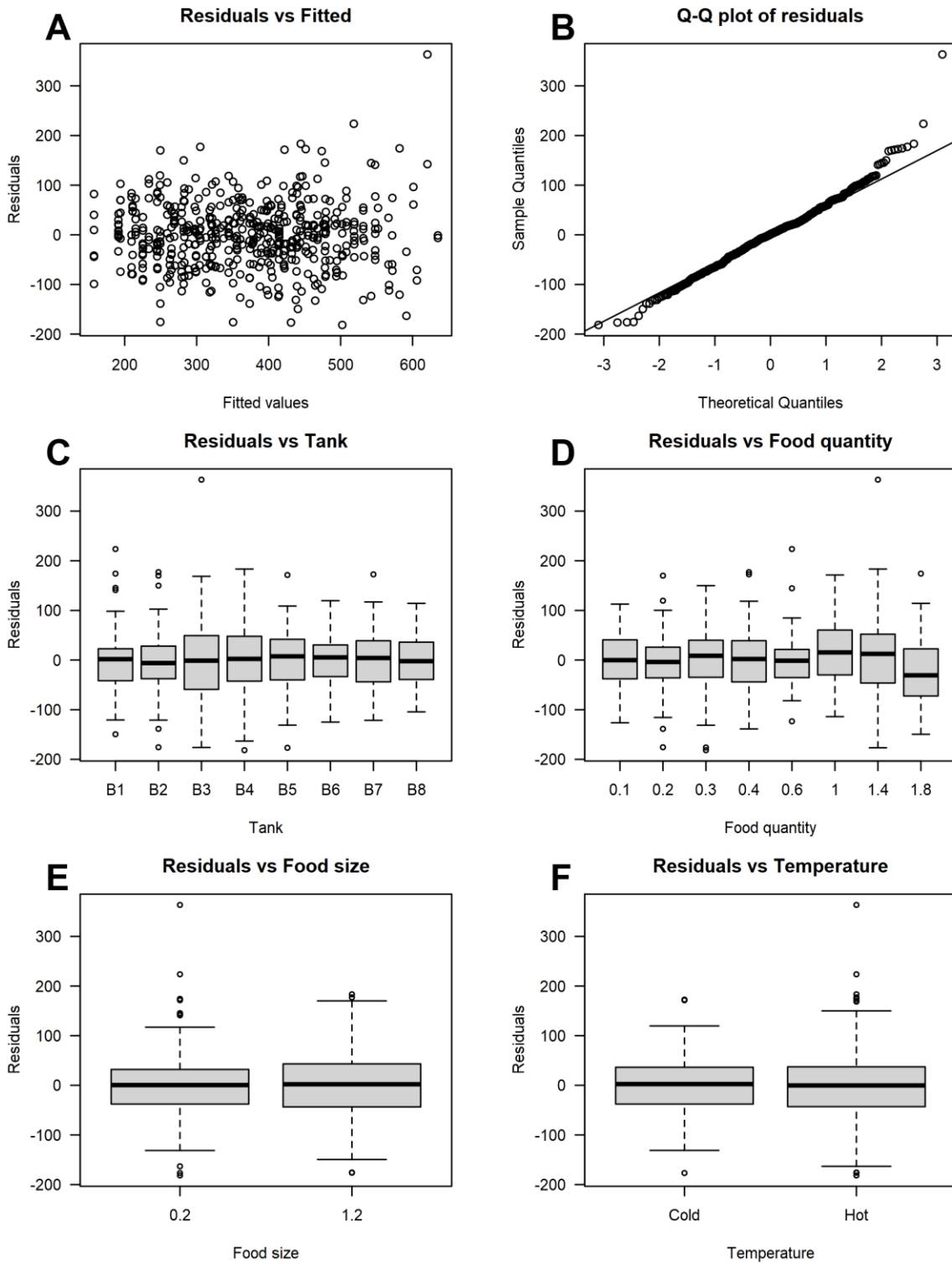


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1015 *Fig. S12: Boxplot of the maximal oxygen consumption increase (relative to basal oxygen consumption) during the*  
 1016 *meal period according to the food ration for the 4 experimental treatments: cool temperature and small particles*  
 1017 *(A), cool temperature and large particles (B), warm temperature and small particles (C) warm temperature and*  
 1018 *large particles (D). Dotted (for small particles) and solid lines (for large particles) represent the increase of the*  
 1019 *median of the maximal oxygen uptake on relation to the food ration. Light and dark grey represent small and large*  
 1020 *particles, respectively. Grey bands represent 95% confidence intervals of smooth increases. Blue and red colors*  
 1021 *represent cool (16°C) and warm (21°C) temperature, respectively. Sample size (i.e. number of meals) is given as*  
 1022 *'n' at the bottom of each panel. Sample size is greater for the 4 smallest quantities (i.e. 0.1, 0.2, 0.3 and 0.4%)*  
 1023 *because oxygen consumption obtained from experiments 1 and 2 were pooled here.*

1024



1025

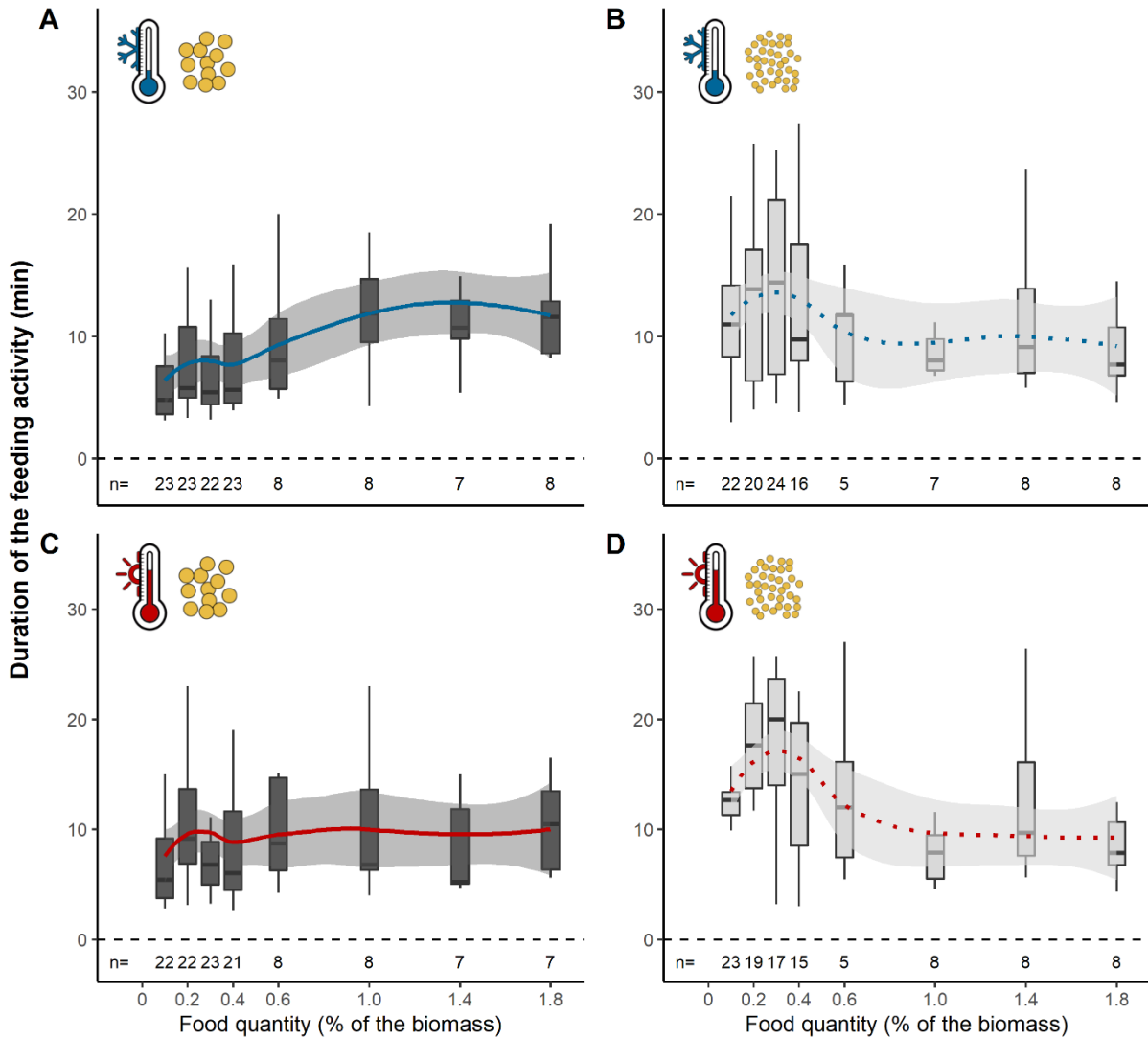
1026

1027 *Fig. S13: Diagnostic plots of the selected model for the maximum oxygen consumption during feeding. A: Fitted*  
 1028 *values versus residuals. B: Q-Q plot of the residuals. C: Residuals versus Tank ID. D: Residuals versus Food*  
 1029 *ration. E: Residuals versus Food size. F: Residuals versus Temperature.*



1030

**E** Feeding duration

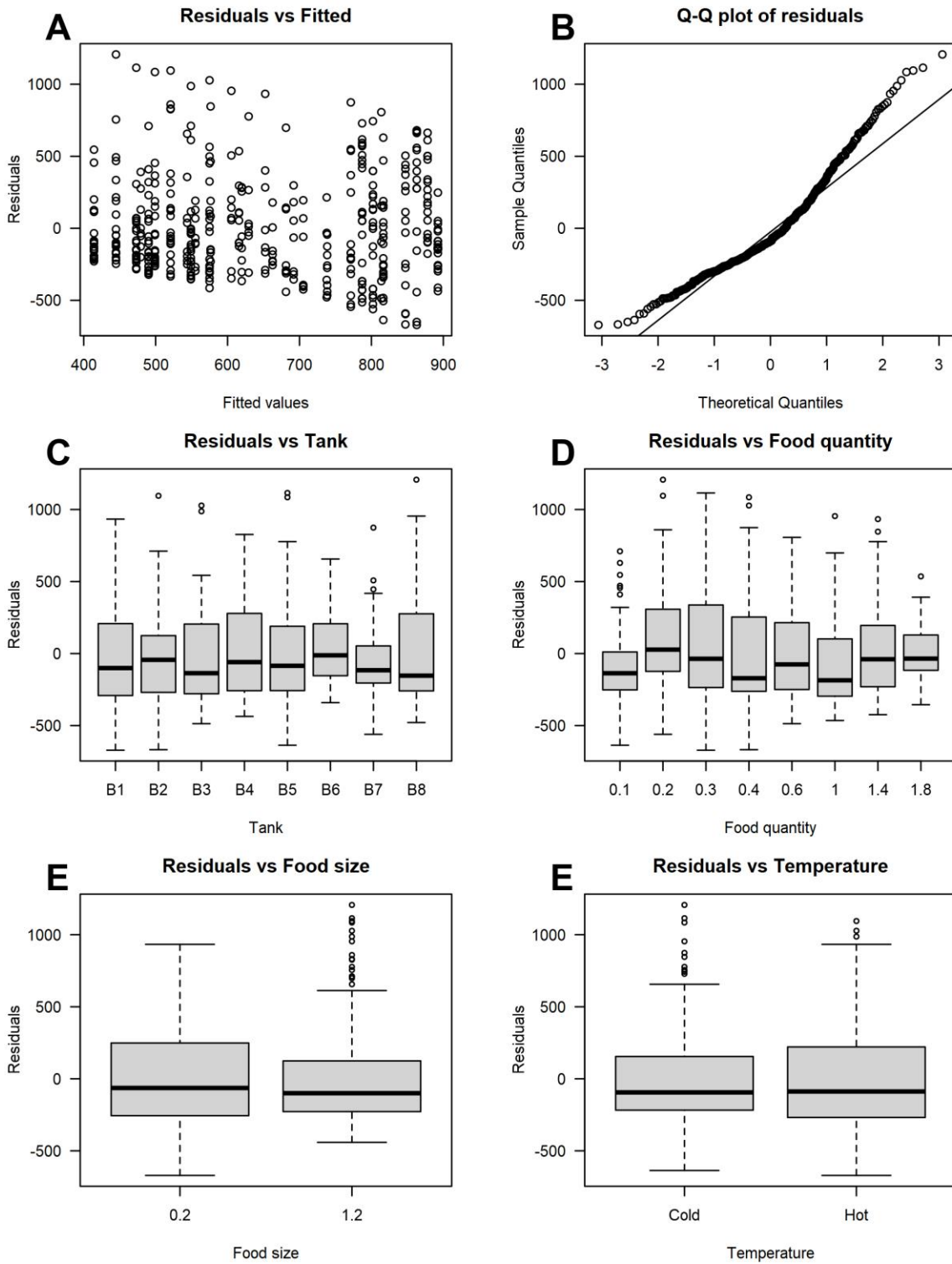


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1033 *Fig. S14: Boxplot of the feeding duration according to the food ration for the 4 experimental treatments: cool*  
 1034 *temperature and small particles (A), cool temperature and large particles (B), warm temperature and small particles*  
 1035 *(C) warm temperature and large particles (D). Dotted (for small particles) and solid lines (for large particles)*  
 1036 *represent the variations of the median feeding duration on relation to the food ration. Grey bands represent 95%*  
 1037 *confidence intervals of smooth increases. Light and dark grey represent small and large particles, respectively.*  
 1038 *Blue and red colors represent cool (16°C) and warm (21°C) temperature, respectively. Sample size (i.e. number of*  
 1039 *meals) is given as 'n' at the bottom of each panel. Sample size is greater for the 4 smallest quantities (i.e. 0.1, 0.2,*  
 1040 *0.3 and 0.4%) because oxygen consumption obtained from experiments 1 and 2 were pooled here. When a feeding*  
 1041 *duration could not be calculated (e.g. no breakpoint), data was removed from this analysis.*

1042



1043

1044 *Fig. S15: Diagnostic plots of the selected model for the feeding duration. A: Fitted values versus residuals. B: Q-Q*  
 1045 *plot of the residuals. C: Residuals versus Tank ID. D: Residuals versus Food ration. E: Residuals versus Food*  
 1046 *size.*

1047 **Tables**

1048

1049 *Table S1: Results of model selection on the daily oxygen consumption (df, logLik, AICc and ΔAICc).*

1050 *The best-fitting model was selected based on the lowest AICc values (Burnham and Anderson, 2002)*

1051 *following Zuur et al. 2003. When difference in AICc (ΔAICc) was lower than two between these models,*

1052 *the more parsimonious model was selected (Burnham and Anderson, 2002). The selected models is*

1053 *represented in bold.*

1054

Models	df	logLik	AICc	ΔAICc
<b>~ Food ration x Temperature + Food ration x Food size</b>	<b>8</b>	<b>-1837.9</b>	<b>3692.3</b>	<b>0.0</b>
~ Food ration x Temperature + Temperature x Food size + Food ration x Food size	9	-1837.7	3694.2	1.8
~ Food ration x Food size x Temperature	10	-1837.2	3695.4	3.1
~ Food ration x Temperature + Food size	7	-1846.9	3708.2	15.9
~ Food ration x Food size + Temperature	7	-1847.7	3709.8	17.5
~ Food ration x Temperature + Temperature x Food size	8	-1846.7	3710.1	17.8
~ Food ration x Food size	6	-1849.2	3710.7	18.4
~ Food ration x Food size + Temperature x Food size	8	-1847.5	3711.6	19.3
~ Food ration x Temperature	6	-1850.3	3713.0	20.7
~ Food ration + Food size + Temperature	6	-1856.1	3724.5	32.2
~ Food ration + Food size	5	-1857.6	3725.4	33.1
~ Food ration + Food size x Temperature	7	-1855.9	3726.4	34.0
~ Food ration + Temperature	5	-1859.3	3728.8	36.4
~ Food ration	4	-1860.8	3729.7	37.4
~ Food size + Temperature	5	-1939.6	3889.4	197.1
~ Food size	4	-1941.0	3890.2	197.9
~ Temperature	4	-1941.3	3890.7	198.4
~ Food size x Temperature	6	-1939.5	3891.4	199.1
~ 1	3	-1942.7	3891.6	199.3

1055

1056

1057 *Table S2: Results of the selected mixed effect model on the daily oxygen consumption (Estimates, 95%*  
 1058 *confidence intervals and p-values). Estimations of the predictors of all other fixed effects were based*  
 1059 *on the estimations of treatment 'warm temperature and small particles'. The selected model is*  
 1060 *presented using REML estimation.*

1061

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**Fixed effects:**

<i>Predictors</i>	<i>Estimates</i>	<i>95% CI</i>	<i>p-value</i>
(Intercept)	555.39	313.23 – 797.54	<0.001
Food ration	868.58	751.42 – 985.74	<0.001
Food size [Large]	106.34	-19.32 – 232.00	0.097
Temperature [Cool]	-41.06	-371.97 – 289.84	0.807
Food size [Large] x Food ration	-294.84	-430.28 – -159.40	<0.001
Temperature [Cool] x Food ration	-308.42	-443.85 – -172.98	<0.001

**Random Effects:**

$\sigma^2$	100091.73		
$\tau_{00 \text{ Tank}}$	48311.87		
ICC	0.33		
$N_{\text{Tank}}$	8		
Observations	255		
Marginal $R^2$ / Conditional $R^2$	0.490 / 0.656		

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1062

1063

1064 *Table S3: Results of linear mixed-effects models on the daily, while feeding, while digesting, maximal*  
 1065 *oxygen consumptions (relative to basal oxygen consumption, see details in Material and Methods) and*  
 1066 *feeding duration according to the food ration for food size and temperature. Slopes were provided after*  
 1067 *being back-transformed for models on feeding and maximal oxygen consumptions. Slopes are given in*  
 1068 *mgO<sub>2</sub>/kg/d for daily MO<sub>2</sub> and during digestion, in mgO<sub>2</sub>/kg/h for oxygen consumption during feeding*  
 1069 *and maximal MO<sub>2</sub>, and in min for feeding duration.*

1070

Period	Treatment	Slope	[95% CI]
Daily	 Large particles	420	[324;515]
	 Small particles	714	[618;810]
	 At 16°C	413	[317;509]
	 At 21°C	721	[626;817]
Meal period	 Large particles	95	[85;105]
	 Small particles	129	[119;139]
	 At 16°C	91	[81;101]
	 At 21°C	133	[123;143]
Digestion	 Large particles	300	[213;388]
	 Small particles	533	[445;621]
	 At 16°C	306	[218;394]
	 At 21°C	527	[439;615]
Maximal	 Large particles	132	[113;152]
	 Small particles	87	[67;106]
	 At 16°C	78	[58;97]
	 At 21°C	141	[122;161]
Duration	 Large particles	4	[1;7]
	 Small particles	-3	[-6;0]

1071

1072 *Table S4: Results of model selection on the oxygen consumption during feeding (df, logLik, AICc and*  
 1073 *ΔAICc). The best-fitting model was selected based on the lowest AICc values (Burnham and Anderson,*  
 1074 *2002) following Zuur et al. 2003. When difference in AICc (ΔAICc) was lower than two between these*  
 1075 *models, the more parsimonious model was selected (Burnham and Anderson, 2002). The selected*  
 1076 *models is represented in bold. Food ration was log-transformed for model selection.*

1077

Models	df	logLik	AICc	ΔAICc
~ log(Food ration) x Food size x Temperature	10	-2505.3	5031.0	0.0
<b>~ log(Food ration) x Temperature + Temperature x Food size + log(Food ration) x Food size</b>	<b>9</b>	<b>-2506.4</b>	<b>5031.1</b>	<b>0.1</b>
~ log(Food ration) x Temperature + log(Food ration) x Food size	8	-2509.3	5035.0	4.0
~ log(Food ration) x Temperature + Temperature x Food size	8	-2517.4	5051.1	20.1
~ log(Food ration) x Temperature + Food size	7	-2520.3	5054.8	23.7
~ log(Food ration) x Food size + Temperature x Food size	8	-2523.3	5062.9	31.9
~ log(Food ration) x Food size + Temperature	7	-2526.1	5066.5	35.5
~ log(Food ration) + Food size x Temperature	7	-2533.5	5081.2	50.2
~ log(Food ration) + Food size + Temperature	6	-2536.2	5084.6	53.6
~ log(Food ration) x Food size	6	-2536.3	5084.7	53.7
~ log(Food ration) + Food size	5	-2546.4	5102.9	71.8
~ Food size + Temperature	5	-2794.5	5599	568.0
~ Food size x Temperature	6	-2793.6	5599.4	568.3
~ Food size	4	-2804.3	5616.8	585.7
~ log(Food ration) x Temperature	6	-2917.4	5847.0	815.9
~ log(Food ration) + Temperature	5	-2920.6	5851.2	820.2
~ log(Food ration)	4	-2929.2	5866.5	835.5
~ Temperature	4	-3004.7	6017.6	986.5
~ 1	3	-3012.6	6031.3	1000.3

1078

1079

1080

1081 *Table S5: Results of the selected mixed effect model on the oxygen consumption during feeding*  
 1082 *(Estimates, 95% confidence intervals and p-values). Estimations of the predictors of all other fixed*  
 1083 *effects were based on the estimations of treatment ‘warm temperature and small particles’. The selected*  
 1084 *model is presented using REML estimation. Food ration was log-transformed before regression.*

1085

<b>Fixed effects:</b>			
<i>Predictors</i>	<i>Estimates</i>	<i>95% CI</i>	<i>p-value</i>
(Intercept)	324.08	314.62 – 334.98	<0.001
log(Food ration)	72.58	67.01 – 78.72	<0.001
Food size [Large]	-156.87	-168.88 – -146.69	<0.001
Temperature [Cool]	-70.01	-83.52 – -56.74	<0.001
log(Food ration) x Food size [Large]	-15.81	-23.06 – -9.52	<0.001
log(Food ration) x Temperature [Cool]	-20.29	-27.06 – -13.52	<0.001
Food size [Large] x Temperature [Cool]	14.05	2.76 – 25.94	0.015
<b>Random Effects:</b>			
$\sigma^2$	1106.87		
$\tau_{00 \text{ Tank}}$	29.07		
ICC	0.03		
$N_{\text{Tank}}$	8		
Observations	509		
Marginal R <sup>2</sup> / Conditional R <sup>2</sup>	0.863 / 0.867		

1086

1087

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1089

1090 *Table S6: Results of model selection on the oxygen consumption during digestion (df, logLik, AICc and*  
 1091 *ΔAICc). The best-fitting model was selected based on the lowest AICc values (Burnham and Anderson,*  
 1092 *2002) following Zuur et al. 2003. When difference in AICc (ΔAICc) was lower than two between these*  
 1093 *models, the more parsimonious model was selected (Burnham and Anderson, 2002). The selected*  
 1094 *models is represented in bold.*

1095

Models	df	logLik	AICc	ΔAICc
<b>~ Food ration x Temperature + Food ration x Food size</b>	<b>8</b>	<b>-1813.3</b>	<b>3643.1</b>	<b>0.0</b>
~ Food ration x Temperature + Temperature x Food size + Food ration x Food size	9	-1813.3	3645.3	2.1
~ Food ration x Food size x Temperature	10	-1812.7	3646.3	3.2
~ Food ration x Temperature	6	-1820.1	3652.6	9.5
~ Food ration x Food size + Temperature	7	-1819.4	3653.2	10.1
~ Food ration x Food size	6	-1820.8	3653.9	10.8
~ Food ration x Temperature + Food size	7	-1820.0	3654.5	11.4
~ Food ration x Food size + Temperature x Food size	8	-1819.4	3655.3	12.2
~ Food ration x Temperature + Temperature x Food size	8	-1820.0	3656.6	13.5
~ Food ration + Temperature	5	-1826.0	3662.2	19.1
~ Food ration	4	-1827.3	3662.9	19.7
~ Food ration + Food size + Temperature	6	-1825.8	3664.0	20.9
~ Food ration + Food size	5	-1827.2	3664.7	21.6
~ Food ration + Food size x Temperature	7	-1825.8	3666.1	23.0
~ Temperature	4	-1887.8	3783.8	140.7
~ 1	3	-1889.2	3784.4	141.3
~ Food size + Temperature	5	-1887.8	3785.8	142.7
~ Food size	4	-1889.1	3786.4	143.3
~ Food size x Temperature	6	-1887.8	3787.9	144.8

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1098 *Table S7: Results of the selected mixed effect model on the oxygen consumption during digestion*  
 1099 *(Estimates, 95% confidence intervals and p-values). Estimations of the predictors of all other fixed*  
 1100 *effects were based on the estimations of treatment ‘warm temperature and small particles’. The selected*  
 1101 *model is presented using REML estimation.*

1102

<b>Fixed effects:</b>			
<i>Predictors</i>	<i>Estimates</i>	<i>95% CI</i>	<i>p-value</i>
(Intercept)	212.24	33.29 – 391.20	<b>0.020</b>
Food ration	643.27	536.00 – 750.54	<b>&lt;0.001</b>
Food size [Large]	187.13	72.08 – 302.18	<b>0.002</b>
Temperature [Cool]	-17.63	-257.48 – 222.22	0.885
Food ration x Food size [Large]	-232.56	-356.57 – -108.56	<b>&lt;0.001</b>
Food ration x Temperature [Cool]	-221.02	-345.02 – -97.01	<b>0.001</b>
<b>Random Effects:</b>			
$\sigma^2$	83904.80		
$\tau_{00 \text{ Tank}}$	22833.11		
ICC	0.21		
$N_{\text{Tank}}$	8		
Observations	255		
Marginal R <sup>2</sup> / Conditional R <sup>2</sup>	0.410 / 0.536		

1103

1104 *Table S8: Results of model selection on the maximal oxygen consumption during feeding (df, logLik,*  
 1105 *AICc and  $\Delta$ AICc). The best-fitting model was selected based on the lowest AICc values (Burnham and*  
 1106 *Anderson, 2002) following Zuur et al. 2003. When difference in AICc ( $\Delta$ AICc) was lower than two*  
 1107 *between these models, the more parsimonious model was selected (Burnham and Anderson, 2002).*  
 1108 *The selected models is represented in bold. Food ration was log-transformed for model selection.*

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Models	df	logLik	AICc	$\Delta$ AICc
<b>~ log(Food ration) x Temperature + Temperature x Food size + log(Food ration) x Food size</b>	<b>9</b>	<b>-2865.6</b>	<b>5749.5</b>	<b>0.0</b>
~ log(Food ration) x Food size x Temperature	10	-2865.4	5751.3	1.7
~ log(Food ration) x Temperature + log(Food ration) x Food size	8	-2870.3	5756.9	7.4
~ log(Food ration) x Temperature + Temperature x Food size	8	-2870.7	5757.8	8.3
~ log(Food ration) x Temperature + Food size	7	-2875.4	5764.9	15.4
~ log(Food ration) x Food size + Temperature x Food size	8	-2875.7	5767.7	18.2
~ log(Food ration) x Food size + Temperature	7	-2880.3	5774.8	25.3
~ log(Food ration) + Food size x Temperature	7	-2880.8	5775.7	26.2
~ log(Food ration) x Food size	6	-2884.1	5780.4	30.8
~ log(Food ration) + Food size + Temperature	6	-2885.2	5782.6	33.1
~ log(Food ration) + Food size	5	-2889.0	5788.2	38.6
~ Food size x Temperature	6	-2975.1	5962.5	212.9
~ Food size + Temperature	5	-2978.1	5966.3	216.7
~ Food size	4	-2981.9	5971.9	222.3
~ log(Food ration) x Temperature	6	-3099.5	6211.2	461.7
~ log(Food ration) + Temperature	5	-3103.4	6216.9	467.4
~ log(Food ration)	4	-3107.1	6222.2	472.7
~ Temperature	4	-3147.1	6302.2	552.7
~ 1	3	-3150.8	6307.6	558.1

1110

1111 *Table S9: Results of the selected mixed effect model on the maximal oxygen consumption during*  
 1112 *feeding (Estimates, 95% confidence intervals and p-values). Estimations of the predictors of all other*  
 1113 *fixed effects were based on the estimations of treatment 'warm temperature and small particles'. The*  
 1114 *selected model is presented using REML estimation. Food ration was log-transformed before*  
 1115 *regression.*

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**Fixed effects:**

<i>Predictors</i>	<i>Estimates</i>	<i>95% CI</i>	<i>p-value</i>
(Intercept)	563.38	525.07 – 601.69	<b>&lt;0.001</b>
log(Food ration)	57.86	46.18 – 69.53	<b>&lt;0.001</b>
Food size [Large]	-154.24	-176.38 – -132.11	<b>&lt;0.001</b>
Temperature [Cool]	-130.42	-183.56 – -77.29	<b>&lt;0.001</b>
log(Food ration) x Food size [Large]	22.10	8.59 – 35.61	<b>0.001</b>
log(Food ration) x Temperature [Cool]	-31.11	-44.62 – -17.60	<b>&lt;0.001</b>
Food size [Large] x Temperature [Cool]	36.14	13.02 – 59.26	<b>0.002</b>

**Random Effects:**

$\sigma^2$	4404.37		
$\tau_{00 \text{ Tank}}$	1208.92		
ICC	0.22		
$N_{\text{Tank}}$	8		
Observations	509		
Marginal R <sup>2</sup> / Conditional R <sup>2</sup>	0.650 / 0.726		

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1119 *Table S10: Results of model selection on the feeding duration (df, logLik, AICc and ΔAICc). The best-*  
 1120 *fitting model was selected based on the lowest AICc values (Burnham and Anderson, 2002) following*  
 1121 *Zuur et al. 2003. When difference in AICc (ΔAICc) was lower than two between these models, the more*  
 1122 *parsimonious model was selected (Burnham and Anderson, 2002). The selected models is represented*  
 1123 *in bold. Food ration was second order polynomial transformed for model selection.*

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Models	df	logLik	AICc	ΔAICc
~ poly(Food ration) x Temperature + poly(Food ration) x Food size	11	-3283.3	6589.2	0.0
~ poly(Food ration) x Temperature + Temperature x Food size + poly(Food ration) x Food size	12	-3282.5	6589.6	0.4
<b>~ poly(Food ration) x Food size + Temperature</b>	<b>9</b>	<b>-3286.0</b>	<b>6590.4</b>	<b>1.2</b>
~ poly(Food ration) x Food size + Temperature x Food size	10	-3285.2	6591.0	1.7
~ poly(Food ration) x Food size x Temperature	14	-3282.3	6593.5	4.3
~ poly(Food ration) x Food size	8	-3288.7	6593.7	4.5
~ Food size + Temperature	5	-3299.6	6609.3	20.1
~ poly(Food ration) x Temperature + Food size	9	-3295.8	6609.9	20.7
~ Food size x Temperature	6	-3298.9	6610.0	20.8
~ poly(Food ration) x Temperature + Temperature x Food size	10	-3295.0	6610.5	21.3
~ poly(Food ration) + Food size + Temperature	7	-3298.5	6611.3	22.1
~ poly(Food ration) + Food size x Temperature	8	-3297.8	6612.0	22.7
~ Food size	4	-3302.0	6612.2	22.9
~ poly(Food ration) + Food size	6	-3301.0	6614.2	25.0
~ Temperature	4	-3326.4	6661.0	71.8
~ 1	3	-3328.5	6663.1	73.9
~ poly(Food ration) x Temperature	8	-3323.7	6663.7	74.5
~ poly(Food ration) + Temperature	6	-3325.9	6664.1	74.8
~ poly(Food ration)	5	-3328.0	6666.2	77.0

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1132 *Table S11: Results of the selected mixed effect model on the feeding duration (Estimates, 95%*  
 1133 *confidence intervals and p-values). Estimations of the predictors of all other fixed effects were based*  
 1134 *on the estimations of treatment 'warm temperature and small particles'. The selected model is*  
 1135 *presented using REML estimation. Food ration was second order polynomial transformed before*  
 1136 *regression.*

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**Fixed effects:**

<i>Predictors</i>	<i>Estimates</i>	<i>95% CI</i>	<i>p-value</i>
(Intercept)	906.09	795.90 – 1016.28	<b>&lt;0.001</b>
poly(Food ration, deg = 2) [coef 1]	-133.39	-526.26 – 259.49	0.505
poly(Food ration, deg = 2) [coef 2]	-34.19	-249.79 – 181.40	0.755
Food size [Large]	-448.82	-595.85 – -301.80	<b>&lt;0.001</b>
Temperature [Cool]	-75.92	-139.61 – -12.23	<b>0.020</b>
poly(Food ration, deg = 2) [coef 1] x Food size [Large]	474.35	-69.23 – 1017.92	0.087
poly(Food ration, deg = 2) [coef 2] x Food size [Large]	-82.83	-381.48 – 215.82	0.586

**Random Effects:**

$\sigma^2$	118832.80
$\tau_{00 \text{ Tank}}$	0.00
$N_{\text{Tank}}$	8

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Observations	453
Marginal R <sup>2</sup> / Conditional R <sup>2</sup>	0.169 / NA

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