

1 **Length-weight relationships of mesopelagic fishes from the**  
2 **equatorial and tropical Atlantic waters: influence of environment**  
3 **and body shape**

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21 **Abstract**

22 Length-weight relationships (LWR) were estimated for 36 mesopelagic fish species collected  
23 from the equatorial and tropical Atlantic encompassing several oceanographic regions:  
24 oligotrophic, equatorial, Cape Blanc, Cape Verde and the Canary Islands. The sample was  
25 composed of myctophids (25 specimens), gonostomatids (5), sternoptychids (3), stomiids (2)  
26 and phosichthyids (1). The species were clustered according to body shape: ‘short and deep’  
27 (sternoptychids), ‘elongate’ (gonostomids, stomiids and some phosichthyids) and ‘fusiform’  
28 (myctophids and some phosichthyids). Three types of weight and LWRs were considered: wet  
29 weight ( $WW$ ), eviscerated wet weight ( $eWW$ ) and eviscerated dry weight ( $eDW$ ). The study  
30 demonstrated that most species present a positive allometric growth, independent of the weigh  
31 used. However, the allometric value varied in 40-50% of species depending on the type of  
32 weight considered. Significant variations linked to fish morphology were found in the  
33 relationship between the slope and intercept of the LWR equation. Significant differences were  
34 also noted in the water content linked to fish body shape. Based on the distributions of several  
35 species we compare their fitness between oceanographic regions using the relative condition  
36 factor ( $K_{rel}$ ). Except for *Diaphus brachycephalus* (oligotrophic vs equatorial waters) and  
37 *Lampanyctus alatus* (equatorial, Cape Blanc, Cape Verde and the Canary Islands), no regional  
38 significant differences were observed in the species analysed.

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42 **KEYWORDS**

43 mesopelagic fishes, oceanographic gradient, morphology, size-weight growth, Atlantic Ocean

## 44 1. INTRODUCTION

45 Mesopelagic fishes have a worldwide distribution from the Arctic to the Antarctic (Krefft, 1974;  
46 Hulley, 1981), although species richness and annual production are commonly greater in  
47 subtropical and tropical regions (Gjøsaeter & Kawaguchi, 1980). They are generally small to  
48 medium-size fishes, including a high diversity of species, numerically dominated by  
49 bristlemouths (family Gonostomatidae) and other Stomiiformes, and by lanternfishes (Order  
50 Myctophiformes), the majority of which live in the ocean's twilight zone (by definition between  
51 200 and 1000 m). Some, particularly myctophids, undertake diel vertical migrations following  
52 their prey into the epipelagic zone to feed at night (Sutton *et al.*, 2008, 2013; Davison *et al.*,  
53 2013; Bernal *et al.*, 2015; Choy *et al.*, 2015; Wang *et al.*, 2019). During daytime, and mainly  
54 to avoid predators, they descend into deeper waters mainly to avoid predators where they  
55 undertake digestion and excretion. This results in a substantial contribution to total vertical  
56 carbon flux from the surface to the deep ocean (Ikeda *et al.*, 2008; Davison *et al.*, 2013; Hudson  
57 *et al.*, 2014; Draze and Sutton, 2017). Thus, these fishes play a key role in food web interactions  
58 linking primary consumers (e.g., copepods and macro-zooplankton) with both pelagic and  
59 deep-sea fish species, as well as other marine species such as marine mammals and birds  
60 (Springer *et al.*, 1999; Pereira *et al.*, 2011; Smith *et al.*, 2011; Trueman *et al.*, 2014). The high  
61 species richness of stomiiform and myctophiform fishes has contributed to the lack of more  
62 detailed information on the basic aspects of fish biology, including length-weight relationships  
63 (LWR) and condition factors at the species level (Fock & Ehrich, 2010; Battaglia *et al.*, 2010,  
64 2015; Jiang *et al.*, 2017; Fock & Czudaj, 2018; Wang *et al.*, 2018).

65 LWR's have been used extensively to gather information on growth, ontogenetic changes,  
66 population dynamics and in trophic ecology studies to reconstruct and estimate the biomass of  
67 partially digested prey (Pauly, 1993; Verdiell-Cubedo *et al.*, 2006; Battaglia *et al.*, 2010, 2016).

68 In general, the parameters for LWRs differ among species due to multiple factors, including the  
69 fish body shape (Schneider *et al.*, 2000; Kulbicki *et al.*, 2005; Froese, 2006; Jellyman *et al.*,  
70 2013), whilst intra-specific differences may be due to gonad maturity, gender, diet, stomach  
71 fullness, health, preservation techniques, season and habitat (Beyer, 1987; Sakuma *et al.*, 1999).  
72 Even, the sampling protocols (e.g., fresh, frozen or preserved in solution) can lead to variations  
73 in LWR estimations (Eduardo *et al.*, 2019). Some applications of LWRs include the ecosystem-  
74 modelling approach to obtain confident production-over-biomass estimates (Christensen &  
75 Pauly, 1992; Pauly *et al.*, 2000; Christensen & Walters, 2004; Torres *et al.*, 2012). In this sense,  
76 LWR's from similar species and different regions are commonly used for the biomass  
77 estimation of mesopelagic fish species from acoustics (Fock & Ehrich, 2010; Fock & Czudaj,  
78 2018), despite possible differences in allometric relationships (Atkinson, 1989). Similarly, the  
79 condition factor is considered as a proxy of fitness in determining the population health and  
80 variability of individuals and populations (Millar & Hickling, 1990; Lloret *et al.*, 2002, 2014;  
81 Pazianoto *et al.*, 2016; Brosset *et al.*, 2018). But there is a lack of information for mesopelagic  
82 fishes (Watanabe & Kawaguchi, 2003).

83 The subtropical western Atlantic Ocean is one of the most oligotrophic regions of the global  
84 ocean (Morel *et al.*, 2010), whereas the eastern central Atlantic is one of the four most  
85 productive regions of the world, due to upwelling (Mittelstaedt, 1983). The ocean between these  
86 two regions is characterized by the convergence of water masses originating in both the  
87 southern and northern hemispheres, and results in a complex oceanic current system (Stramma  
88 & Schott, 1999). Moreover, south of the Cape Verde archipelago, the mesopelagic layers are  
89 characterized by the presence of an oxygen minimum zone (OMZ), with dissolved oxygen  
90 concentrations corresponding to intermediate hypoxia values (Ekau *et al.*, 2010; Moffitt *et al.*,  
91 2014; Olivar *et al.*, 2017). Previous investigations in the area have shown that some  
92 mesopelagic fishes are associated with particular waters masses, while others occur across the

93 entire region (Olivar *et al.*, 2017, 2018). These differences in the oceanographic features may  
94 also have an influence on the trophic transfer efficiencies from plankton to fishes, which can be  
95 indirectly evaluated from body condition (Le Cren, 1951; Simpkins & Hubert, 2000; Stevenson  
96 & Woods 2006; Wilson & Nussey, 2010).

97 This study attempts: (i) to provide information on LWR's of 36 mesopelagic fish species  
98 collected in the tropical and equatorial Atlantic waters by using different weight variables, so  
99 as to facilitate effective use by scientists working on mesopelagic biomass estimation; (ii) to  
100 assess the changes in the fish health between oceanographic regions using the Relative  
101 Condition Factor (Le Cren, 1951); and (iii) to evaluate how the body shape influences the *a* and  
102 *b* parameters of LWR equation, as well as the water content of species. These types of  
103 information can provide useful evidence to improve the understanding of how species and  
104 individuals are structured in open waters.

105

## 106 2. MATERIAL AND METHODS

### 107 2.1. Sample collection

108 Specimens were collected during a cruise carried out in April 2015 across the equatorial and  
109 tropical Atlantic, from near the Brazilian coast (13°S 38.3°W) to south Canary Islands (28°N  
110 15.6°E), on board R/V Hespérides. A total of 12 stations were occupied both day and night, and  
111 fish samples were obtained in different strata of the water column, from 800 m to the surface  
112 (Olivar *et al.*, 2017). Ship speed was 2 knots (1 m/s) and haul duration at each strata ranged  
113 from 10 to 30 min. The sampling gear deployed was a pelagic midwater trawl, the  
114 “Mesopelagos” net (35-m<sup>2</sup> mouth opening; total length 58 m, with graded-mesh netting from  
115 30 mm at mouth to 4 mm towards the cod end) (more details in Olivar *et al.*, 2017).

116 The fresh samples were analysed on board. Fishes were identified to the lowest possible  
117 taxon and kept frozen at -20°C until transference to the laboratory, where their identifications  
118 were checked, and individual measurements of length and weight were taken. Standard length  
119 (*SL*) was recorded to the nearest 1 mm, using digital calipers, and whole body wet weight (*WW*)  
120 to the nearest 0.0001 g, using a digital balance. Individuals were then dissected, the digestive  
121 tract, stomach and gonads were removed, and the eviscerated wet weight (*eWW*) of each  
122 specimen was taken. To eliminate water content, eviscerated specimens were freeze-dried in a  
123 Telstar LyoAlfa 6 freeze dryer for 24 h, and the eviscerated dry weight of each specimen was  
124 then recorded (*eDW*). Water content was calculated as the difference between *eWW* and *eDW*,  
125 expressed as % *eWW*, to highlight their effect on the estimation of LWRs.

126

## 127 **2.2. Length-weight relationships**

128 Size and weight data were fitted using a power function:  $y = ax^b$ , where  $x$  is *SL* and  $y$  the weight  
129 parameter (*WW*, *eWW* or *eDW*). The slope,  $b$ , is the allometric growth factor and the intercept,  
130  $a$ , is the expected value of  $y$  at  $x=1$  (Gould, 1966). Data from the logarithmically transformed  
131 equations were then adjusted by the method of least squares. The values of  $b$ , their 95%  
132 confidence intervals (95% CI), and the coefficient of determination ( $r^2$ ) were calculated  
133 according to the methods of Sokal & Rohlf (1979). In addition, a Student's- $t$ -test was used to  
134 evaluate the isometric growth ( $b=3$ ), and whether  $b$  value is significantly higher or lower  
135 indicating a positive or negative allometry, respectively (Gould, 1966; Margalef, 1974; Pauly,  
136 1984).

137 To assess the influence of body morphology on the intercept ( $a$ ) and slope ( $b$ ) parameters of  
138 the LWR (using *eDW*), a linear regression was estimated for three groups of fishes defined on  
139 the basis of their general body forms: 'short and deep' (Sternorhynchidae), 'elongate'

140 (Gonostomatidae and Stomiidae) and ‘fusiform’ (Phosichthyidae and Myctophiidae). Statistical  
141 differences between the slopes were tested using an ANCOVA analysis. In addition, the average  
142 water content of each fish group was compared using a Kruskal-Wallis test (non-parametric  
143 test).

144

### 145 **2.3. Relative condition factor**

146 The fish condition was estimated using the Relative Condition Factor ( $K_{rel} = (W_o/aSL^b)$ ; [Le Cren, 1951](#)), which relates the observed body weight ( $W_o = eDW$ ) of a given individual with the  
147 predicted by the length–weight relationship ( $aSL^b$ ). The parameters  $a$  and  $b$  were obtained from  
148 the regional LWR derived by pooling data for all regions for each species separately ([Efitre et al., 2009](#)).  
149 Calculation of Fulton’s Condition Index ( $Kn = 100W/L^3$ ; [Fulton, 1911](#)) was omitted  
150 because the growth of mesopelagic fishes was not isometric (see results). The Relative Weight  
151 index ([Wege & Anderson, 1978](#); [Froese, 2006](#)) was not used because there are no studies based  
152 on this index supporting a population differentiation of mesopelagic fishes in open ocean.  
153

154 In order to perform the regional comparisons in the  $K_{rel}$ , regions of the study area were  
155 defined according to oceanographic parameters described in [Olivar et al. \(2017\)](#): oligotrophic  
156 (stations #1-3), equatorial (stations #4-6), Cape Verde islands (stations #7-10), Cape Blanc  
157 upwelling (station#11), Canary Islands (station#12) (Figure 1). For these comparisons, only 9  
158 species (*Chauliodus danae*, *Ceratoscopelus warmingii*, *Diaphus brachycephalus*, *Diaphus mollis*,  
159 *Diaphus rafinesquii*, *Lampanyctus alatus*, *Lepidophanes guentheri*, *Myctophum nitidulum* and  
160 *Vinciguerria nimbaria*) were considered to have a sufficient number of  
161 individuals ( $n > 10$ ) by region and LWRs with high correlations ( $r^2 > 0.90$ ). The normality and  
162 homogeneity of variances in the  $K_{rel}$  data were checked for each species by region using the  
163 Shapiro-Wilk’s test and Bartlett’s test, respectively. Depending on the Gaussian distribution of

164 data, variations of  $K_{rel}$  between two regions were compared by the Student's  $t$ -test (parametric  
165 test), or the Mann–Whitney's  $U$  Test (non-parametric test). To analyse more than two regions,  
166 a Kruskal-Wallis test followed by *a posteriori* Dunn test was performed with the package  
167 *dunn.test* v.1.3.5. in R (Dino, 2017).

168 All statistical analyses and graphical representations were conducted with the software R (R  
169 Core Team, 2016).

170

### 171 3. RESULTS

172 A total 1277 individuals belonging to 36 species from 5 families were analysed. The family  
173 Myctophidae was represented by 11 genera and 25 species, followed by Gonostomatidae, with  
174 2 genera and 5 species; Sternoptychidae, with 3 genera and 3 species; Stomiidae, with 2 genera  
175 and 2 species, and Phosichthyidae, with 1 genus and 1 species (Table 1).

176 The determination coefficients ( $r^2$ ) of the fitted equations were generally high (mean value  
177 of  $0.991 \pm 0.001$ ), ranging from 0.613 for *Cyclothone pseudopallida* (Gonostomatidae) to 0.997  
178 for *Bolinichthys photothorax* (Myctophidae). Excluding the extreme case presented by *C.*  
179 *pseudopallida*, 77% of the relationship have  $r^2 > 0.950$  (Table 2 for *eDW*; Table S1 for *WW* and  
180 Table S2 for *eWW*, Supplementary material).

181 In terms of growth, the LWRs for *WW* revealed that 57.2% of the species showed isometric  
182 growth, the 30.5% positive allometry ( $b > 3$ ) and the 11.1% negative allometry ( $b < 3$ ). A similar  
183 tendency was also detected when fitting *eWW* (52.7%, 33.3% and 13.8%, respectively). The  
184 main differences in growth patterns between the LWRs for *WW* or *eWW* were observed for  
185 *Vinciguerria nimbaria* (Phosichthyidae), which presented a higher  $b$  when fitting total weight,  
186 and for *Lepidophanes guentheri* (Myctophidae) and *Sternoptyx diaphana* (Sternoptychidae)



187 with lower  $b$  using eviscerated data. By contrast, LWRs for  $eDW$  showed a slight increment in  
188 the proportion of species with positive allometry (47.2% isometric, and 38.8% and 13.8%,  
189 positive and negative allometry, respectively). Irrespective of the fitted data, the lowest  
190 allometric coefficients were always found for *Cyclothone* spp., together with the myctophid  
191 *Diaphus vanhoeffeni*. The highest allometric growth coefficient was always in *Polyipnus polli*,  
192 and in myctophids such as *Benthosema glaciale*, *Ceratoscopelus warmingii*, *Diaphus*  
193 *brachycephalus*, *D. holti*, *D. metopoclampus*, *D. rafinesquii*, *Lampanyctus alatus*, *L. pusillus*,  
194 *Lepidophanes guentheri*, *Lobianchia dofleini*, *Myctophum affine*, *M. nitidulum*, and *M.*  
195 *punctatum* (Table 2 for  $eDW$ ; Table S1 for  $WW$  and Table S2 for  $eWW$ , Supplementary  
196 material). The 30.5% species showed an allometric coefficient higher for LWRs fitting with  
197  $eDW$  in comparison with  $WW$ , and 27.8% in relation to  $eWW$ . Nevertheless, 52.8% and 59.3%  
198 of cases provided similar  $b$  values ( $\pm 0.1$ ). In general, the greatest differences between the  
199 growth patterns were observed for 6 myctophids: *Lampanyctus pusillus*, *L. alatus*, *Diaphus*  
200 *metopoclampus*, *Benthosema glaciale*, *Ceratoscopelus warmingii* and *Diaphus holti*; and for  
201 the gonostomatid *Cyclothone pseudopallida*. In these cases, the higher growth rates ( $b$ ) of the  
202 RLWs were estimated for  $eDW$ . In contrast, a negative difference between  $b$  values was noted  
203 for: *Sigmops elongatus* (Gonostomatidae), *Notolychnus valdiviae* (Myctophidae),  
204 *Argyropelecus sladeni* (Sternoptychidae), and *Chauliodus danae* (Stomiidae). Strong  
205 correlations were found between the values of  $b$  and  $\log a$ , varying significantly between fish  
206 body shapes ( $F_{2,32} = 3.486$ ,  $P = 0.044$ , Figure 2). The slope was larger in elongated shapes ( $b =$   
207  $-1.878$ ) and smaller in short and compressed fishes as sternoptychids ( $b = -1.056$ ).

208 Intra-specific differences between growth patterns within species were higher when  
209 comparing  $eWW$  and  $eDW$  than between  $WW$  and  $eWW$ , which revealed the important  
210 contribution of water content. The species with higher water content ( $> 80\%$ ) were *Cyclothone*  
211 *pallida*, *C. acclinidens*, *Lampanyctus nobilis*, *Chauliodus danae*, *Sigmops elongatus* and

212 *Stomias boa boa* (Table 3). By contrast, *Notolychnus valdiviae*, *Polyipnus polli*, *Lampanyctus*  
213 *pusillus*, *Benthoosema glaciale* and *Diaphus fragilis* had values lower than 75% (Table 3). The  
214 elongated shaped fishes were characterized by higher water content (mean and standard  
215 deviation,  $81.50 \pm 1.85$ ) than fusiform fishes ( $76.41 \pm 4.11$ ), and the short and deep shaped  
216 species ( $75.99 \pm 4.06$ ). A high variability was found between groups (Kruskal-Wallis test,  $\chi^2 =$   
217  $12.81$ ,  $df = 3$ ,  $P = 0.002$ ).

218 The mesopelagic fishes showed similar average values of  $K_{rel}$  ranging between 0.792  
219 (*Diaphus brachycephalus* from the oligotrophic region) and 1.120 (*D. rafinesquii* from the  
220 Cape Blanc upwelling region). Regional associations in the average values of  $K_{rel}$  were found  
221 for *D. brachycephalus* ( $t$ -test,  $t = 8.817$ ,  $df = 84$ ,  $P < 0.0001$ ) with lower values ( $0.792 \pm 0.185$ )  
222 in oligotrophic waters than in equatorial region ( $0.902 \pm 0.076$ ) (Table 4), and in *Lampanyctus*  
223 *alatus* (Kruskal-Wallis test,  $\chi^2 = 17.309$ ,  $df = 3$ ,  $P = 0$ ), showing an increase of average  $K_{rel}$  for  
224 regions closer to African coast. The equatorial region differed from the Cape Blanc upwelling  
225 region and Canary Islands, but it was not dissimilar to Cape Verde Islands region. The  $K_{rel}$  value  
226 of Cape Verde Islands only presented differences with the average of the Canary Islands; and  
227 the Cape Blanc upwelling region and Canary Islands also reached similar average of condition  
228 (Table 4).

229

#### 230 **4. DISCUSSION**

231 The present study contributes to knowledge of mesopelagic fishes by reporting on LWR  
232 equations for 36 species, thereby establishing the effect of different weights in the allometry of  
233 LWR's, and by comparing the environmental effect in the relative condition factor. In general,  
234 our findings seem to reinforce the theory that the growth pattern is a feature identifying each

235 species (Mayrat, 1970) since 24 of the 36 of mesopelagic fishes analysed demonstrated similar  
236 allometric relationships, independent of the weight measure used. Some species reached a  
237 higher  $b$  value when fitting  $WW$  instead of  $eWW$ , reflecting the influence of full guts and, to a  
238 lesser extent, gonadal mass (e.g., *Vinciguerria nimbaria* *Chauliodus danae* and *Cyclothone*  
239 *livida*), but in other species showed the opposite effect (e.g., *Lobianchia dofleini*, *Lepidophanes*  
240 *guentheri* and *Diaphus vanhoeffeni*). Nevertheless, the use of eviscerated dry weight is always  
241 recommended for LWR's, since it more accurately reflects better the muscular growth,  
242 irrespective of the trophic behaviour (full or empty guts) or the gonadal weight (important at  
243 maturation) (Pauly 1984; Froese, 2006). In other instances, such as for reconstruction and  
244 estimation of biomass of partially digested prey in studies of trophic ecology, information on  
245  $WW$  would also be relevant. Allometric coefficients ( $b$  for  $eDW$ ) in the present study were  
246 within the expected range (2.5-3.5) for fishes (Froese, 2006), mostly ranging between 2.952  
247 (25% percentile) and 3.384 (75% percentile). The most atypical allometric coefficient was  
248 observed for the myctophid *Lampanyctus pusillus*, 4.247, which could be due to the small size  
249 range analysed.

250 The species and sizes ranges of myctophids and hatchetfishes studied here showed a positive  
251 allometric growth, which implies a faster growth in body mass than in body length, i.e., more  
252 robust body and with a greater amount of muscle mass. This feature may help in the daily  
253 feeding vertical migrations, to the surface in myctophids and to the shallower mesopelagic  
254 depths in hatchetfishes (Olivar *et al.*, 2012, 2016, 2017). The opposite pattern with negative  
255 allometric growths, i.e., faster growth in length than in weight, was mostly observed in  
256 stomiiform species such as *Chauliodus danae*, *Cyclothone acclinidens*, *C. livida*, *C. pallida* and  
257 *Sigmops elongatus*, which live in deeper waters, and which have an elongate shape and higher  
258 water content. This finding is in accordance with previous studies showing an association  
259 between the proportion of water content and distribution depth of these fishes (Childres &

260 Nygaard, 1973; Neighbors & Nafpaktitis, 1982; Bailey & Robison, 1986; Stickney & Torres,  
261 1989; Pakhomov *et al.*, 1996), and is also accompanied by a decrease in skeletal ash content  
262 (see Childres & Nygaard, 1973). The main reason of this somatic growth is probably due to  
263 absence of extensive vertical migrations in most of these species (Badcock and Merret, 1976;  
264 Olivar *et al.*, 2017). In general, the allometric pattern found in our species did not differ from  
265 those in other studies, although slight variations can be due to exogenous and endogenous  
266 factors (e.g., Kimmerer *et al.*, 2005; Jobling, 2008; Mazumder *et al.*, 2016), as well as the type  
267 of length measurement (standard or total), and size-range analysed. For instance, Fock & Ehrich  
268 (2010) gave a wide list of LWR's for mesopelagic fishes from the North Atlantic. Their  
269 estimation of allometric coefficients were smaller than in our study (for *eDW*) for those species  
270 differing in the size range, for example in *B. glaciale* (3.647 vs 3.020), *D. dumerilli* (3.076 vs  
271 3.018), *D. metopoclampus* (3.684 vs 3.074), *L. dofleini* (3.448 vs 2.609) and *N. bolini* (3.212 vs  
272 2.331). However, they were similar when the size ranges were similar as for example in *D. holti*  
273 (3.356 vs 3.350), *M. punctatum* (3.363 vs 3.448) and *S. boa boa* (3.081 vs 3.184). For fishes  
274 with similar fish size-ranges, our allometric patterns were larger than in other geographical  
275 areas, for example in the Mediterranean Sea for *Diaphus holti* (3.356 vs 3.102) and  
276 *Lampanyctus pusillus* (4.247 vs 2.296) (Battaglia *et al.*, 2010), and in the North-western Pacific  
277 Ocean for *Ceratoscopelus warmingii* (3.537 vs 3.153) (Wang *et al.*, 2018).

278 Food availability and physical factors have a strong influence on the growth and condition  
279 factors (Le Cren, 1951). Olivar *et al.* (2017) found higher environmental gradients in terms of  
280 temperature (22°C of difference) and salinity (3 PSU) between the sea surface and deep waters  
281 in the subtropical western Atlantic Ocean than eastern zone (10°C and 1 PSU, respectively).  
282 Although this environmental variation may affect the energetic balance and growth of  
283 specimens, any difference was detected in most species, except for *Lampanyctus alatus*. The  
284 most probable reason for the increase of the  $K_{rel}$  along the Atlantic transect may be related to

285 the visual system, a more generalist pattern being characterized by a major visual field  
286 favouring the detection of preys in all directions (de Buserrolles *et al.*, 2014, de Buserrolles &  
287 Marshall, 2107), and which is more numerous in the western sector. Finally, low values of  $K_{rel}$   
288 ( $< 1$ ) for some species (e.g., *Chauliodus danae*, *Diaphus brachycephalus*, *D. mollis* and  
289 *Vingiguerria. nimbaria*) were noted in all regions, which suggests two hypotheses: a) the  
290 energetic cost of diel vertical migration (DVM) may be higher in these species; or b) the  
291 energetic requirement may be less. Unfortunately, information on the food conversion  
292 efficiency on fish growth is not available for many mesopelagic fish species. However, the  
293 intraspecific and regional variability found in our study reinforces the importance of  
294 investigations into ecological and energetic demands of deep-sea organisms (Siebel & Drazen,  
295 2007; Irigoien *et al.*, 2014; Belcher *et al.*, 2019).

296

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302

## 303 **Contributions**

304 MPO and PAH were responsible for the species identifications; CLP, conceived the initial idea  
305 and wrote the main paper in collaboration with the other co-authors; CLP, MPO and VTA  
306 conducted the statistical analysis. All authors discussed results and implications, providing  
307 significant inputs to the manuscript at all stages.

308

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## 515 **Legends**

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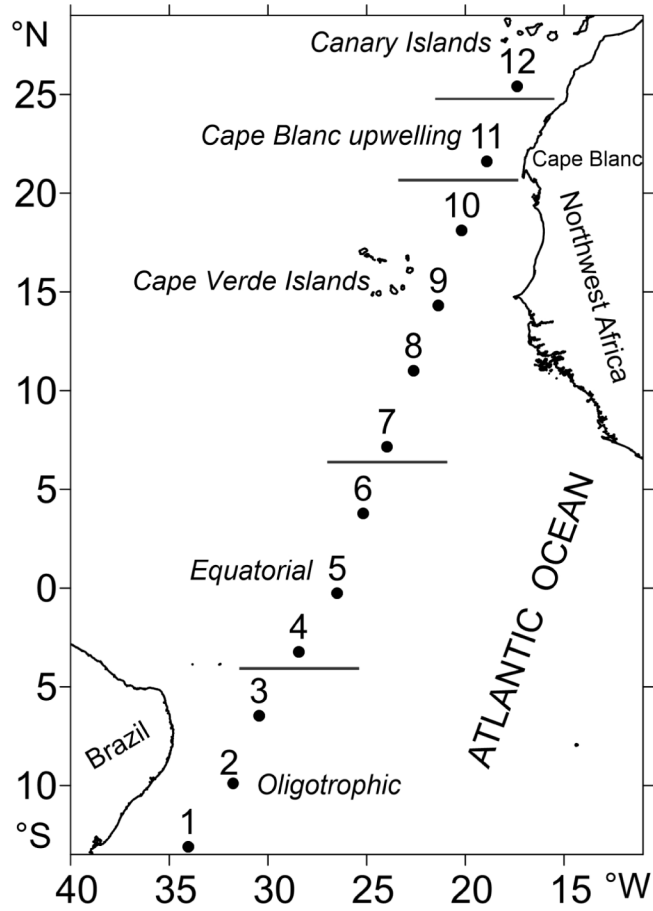
517 **FIGURE 1** Location of the stations sampled during the survey across the equatorial and tropical  
518 Atlantic during April 2015, and oceanographic regions according to [Olivar et al. \(2017\)](#).

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520 **FIGURE 2** Relationships between the regression coefficients  $\log a$  and  $b$  of LWR equation for  
521 different body shapes (short-deep, elongate and fusiform) at the species level.

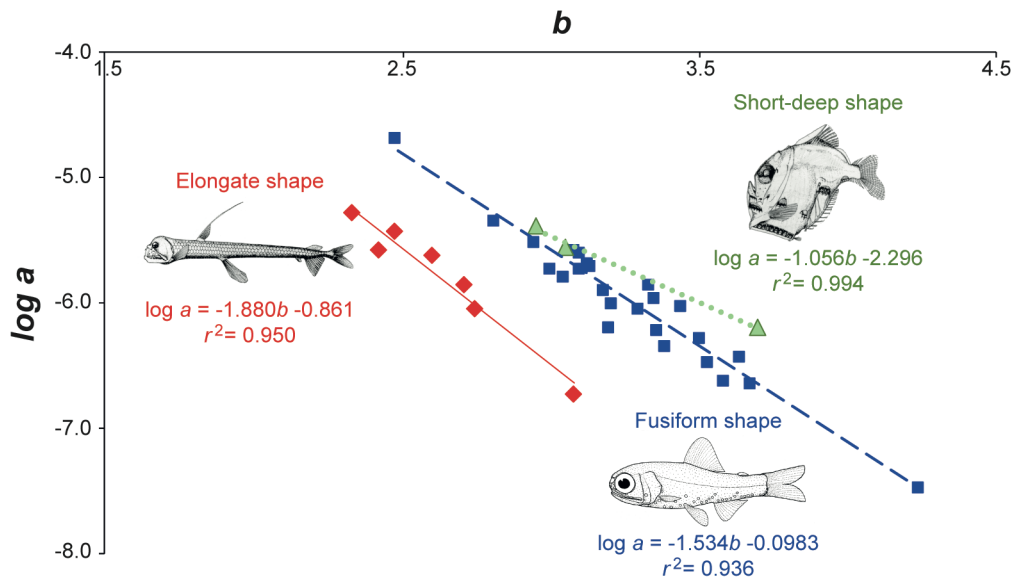
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**TABLE 1** List of mesoplegic fishes from equatorial and tropical Atlantic waters analyzed in the present study. Region indicates the sampling origin according to Olivar *et al.* (2017): CB, Cape Blanc; CI, Canary Islands; CV, Cape Verde; O, oligotrophic; E, equatorial. Taxa were ordered according to Nelson *et al.* (2016)

Order	Family	Genera	Species	Author	Region	
Stomiiformes	Gonostomatidae	<i>Cyclothone</i>	<i>Cyclothone acclinidens</i>	Garman, 1899	CV, CB	
			<i>Cyclothone livida</i>	Brauer, 1902	CV, CB	
			<i>Cyclothone pallida</i>	Brauer, 1902	CV, CB	
			<i>Cyclothone pseudopallida</i>	Mukhacheva, 1964	CV	
	Phosichthyidae	<i>Gonostoma</i>	<i>Sigmops elongatus</i>	(Günther, 1878)	E, CV, CB, CI	
			<i>Vinciguerria nimbaria</i>	(Jordan & Williams, 1895)	CV, CB	
	Sternoptychidae	<i>Argyropelecus</i>	<i>Argyropelecus sladeni</i>	Regan, 1908	E, CV, CB	
			<i>Polypnus polli</i>	Schultz, 1961	CV, CB	
			<i>Sternoptyx diaphana</i>	Hermann, 1781	CV, CB	
	Stomiidae	<i>Chauliodus</i>	<i>Chauliodus danae</i>	Regan & Trewavas, 1929	E, CV, CB	
			<i>Stomias boa boa</i>	(Risso, 1810)	CV	
	Myctophiformes	Myctophidae	<i>Benthoosema</i>	<i>Benthoosema glaciale</i>	(Reinhardt, 1837)	CB
				<i>Benthoosema suborbitale</i>	(Gilbert, 1913)	CV, CI
			<i>Bolinichthys photothorax</i>	(Parr, 1928)	O, E, CV	
<i>Ceratoscopelus warmingii</i>			(Lütken, 1892)	O, E, CV		
<i>Diaphus</i>			<i>Diaphus brachycephalus</i>	Tåning, 1928	O, E, CV, CB	
			<i>Diaphus dumerilii</i>	(Bleeker, 1856)	E	
			<i>Diaphus fragilis</i>	Tåning, 1928	E	
			<i>Diaphus holti</i>	Tåning, 1918	CV, CB	
			<i>Diaphus metopoclampus</i>	(Cocco, 1829)	E, CI	
			<i>Diaphus mollis</i>	Tåning, 1928	O, E, CV	
			<i>Diaphus problematicus</i>	Parr, 1928	E	
			<i>Diaphus rafinesquii</i>	(Cocco, 1838)	CB, CI	
			<i>Diaphus vanhoeffeni</i>	(Brauer, 1906)	CV	
			<i>Hygophum macrochir</i>	(Günther, 1864)	CB (30)	
			<i>Lampanyctus</i>	<i>Lampanyctus alatus</i>	Goode & Bean, 1896	E, CV, CB, CI
				<i>Lampanyctus nobilis</i>	Tåning, 1928	O, E
				<i>Lampanyctus pusillus</i>	(Johnson, 1890)	E, CB, CI
<i>Lepidophanes guentheri</i>			(Goode & Bean, 1896)	O, E, CV		
<i>Lobianchia dofleini</i>			(Zugmayer, 1911)	CV, CB, CI		
<i>Myctophum</i>			<i>Myctophum affine</i>	(Lütken, 1892)	CV	
			<i>Myctophum nitidulum</i>	Garman, 1899	CV, CI	
			<i>Myctophum punctatum</i>	Rafinesque, 1810	CB	
<i>Notolychnus valdiviae</i>			(Brauer, 1904)	E, CV		
<i>Notoscopelus</i>			<i>Notoscopelus bolini</i>	Nafpaktitis, 1975	CB	
			<i>Notoscopelus resplendens</i>	(Richardson, 1845)	E, CV, CB, CI	

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**TABLE 2** Length-weight parametres for mesopelagic species (ordered alphabetically) from the equatorial and tropical Atlantic waters. *a*, intercept; *a*<sup>+</sup>, positive allometry; *a*<sup>-</sup>, negative allometry; *b*, allometry coefficient (slope); *eDW*, eviscerate body dry weight; *i*, isometry; *n*, sample size; *SL*, standard length; *r*<sup>2</sup>, coefficient of determination; 95% CL of *b*, confidence interval

Species	n	Range <i>SL</i> (mm)	Range <i>eDW</i> (g)	<i>a</i>	<i>b</i>	Inferior 95% CL of <i>b</i>	Superior 95% CL of <i>b</i>	<i>r</i> <sup>2</sup>	Growth model
<i>Argyroleucus sladeni</i>	27	15 – 43	0.012 – 0.186	0.00000414	2.950	2.472	3.428	0.866	i
<i>Benthoosema glaciale</i>	15	15 – 35	0.005 – 0.145	0.00000035	3.647	3.486	3.809	0.990	a+
<i>Benthoosema suborbitale</i>	29	16 – 33	0.012 – 0.103	0.00000198	3.135	2.868	3.401	0.955	i
<i>Bolinichthys photothorax</i>	6	21 – 51	0.021 – 0.335	0.00000121	3.183	2.944	3.421	0.997	i
<i>Ceratoscopelus warmingii</i>	47	16 – 64	0.006 – 0.796	0.00000032	3.537	3.443	3.631	0.992	a+
<i>Chauliodus danae</i>	43	25 – 212	0.023 – 3.519	0.00000139	2.710	2.554	2.867	0.968	a-
<i>Cyclothone acclinidens</i>	22	31 – 46	0.005 – 0.051	0.00000524	2.337	1.849	2.825	0.768	a-
<i>Cyclothone livida</i>	32	23 – 37	0.007 – 0.030	0.00000372	2.477	2.011	2.941	0.798	a-
<i>Cyclothone pallida</i>	24	25 – 54	0.009 – 0.073	0.00000235	2.600	2.270	2.929	0.924	a-
<i>Cyclothone pseudopallida</i>	19	23 – 35	0.004 – 0.017	0.00000258	2.420	1.596	3.244	0.693	i
<i>Diaphus brachycephalus</i>	100	12 – 47	0.002 – 0.749	0.00000131	3.339	3.194	3.482	0.956	a+
<i>Diaphus dumerilii</i>	30	31 – 57	0.089 – 0.628	0.00000254	3.076	2.865	3.285	0.970	i
<i>Diaphus fragilis</i>	10	46 – 79	0.183 – 1.022	0.00000155	3.056	2.740	3.371	0.984	i
<i>Diaphus holti</i>	91	11 – 50	0.002 – 0.491	0.00000106	3.356	3.250	3.460	0.978	a+
<i>Diaphus metopoclampus</i>	10	19 – 40	0.009 – 0.182	0.00000022	3.684	3.219	4.147	0.977	a+
<i>Diaphus mollis</i>	30	29 – 55	0.073 – 0.571	0.00000247	3.093	2.912	3.272	0.979	i
<i>Diaphus problematicus</i>	9	45 – 72	0.223 – 0.787	0.00000437	2.820	2.441	3.199	0.978	i
<i>Diaphus rafinesquii</i>	125	11 – 70	0.004 – 1.425	0.00000242	3.107	3.042	3.170	0.987	a+
<i>Diaphus vanhoeffeni</i>	17	25 – 36	0.054 – 0.154	0.00001968	2.488	1.856	3.119	0.825	i
<i>Hygophum macrochir</i>	30	20 – 42	0.018 – 0.223	0.00000190	3.138	2.973	3.302	0.982	i
<i>Lampanyctus alatus</i>	142	16 – 57	0.006 – 0.362	0.00000023	3.594	3.492	3.696	0.972	a+
<i>Lampanyctus nobilis</i>	19	28 – 74	0.022 – 0.619	0.00000061	3.205	2.906	3.503	0.968	i
<i>Lampanyctus pusillus</i>	22	24 – 36	0.019 – 0.132	0.00000003	4.247	3.638	4.855	0.914	a+
<i>Lepidophanes guentheri</i>	35	19 – 61	0.008 – 0.521	0.00000043	3.391	3.181	3.601	0.970	a+
<i>Lobianchia dofleini</i>	30	13 – 30	0.006 – 0.118	0.00000090	3.448	3.081	3.813	0.930	a+
<i>Myctophum affine</i>	34	14 – 47	0.005 – 0.328	0.00000049	3.509	3.384	3.633	0.990	a+
<i>Myctophum nitidulum</i>	25	15 – 74	0.006 – 1.233	0.00000084	3.308	3.219	3.396	0.996	a+
<i>Myctophum punctatum</i>	28	16 – 69	0.007 – 1.068	0.00000057	3.363	3.252	3.472	0.993	a+
<i>Notolychnus valdiviae</i>	27	15 – 22	0.019 – 0.123	0.00000289	2.956	2.956	3.577	0.816	i
<i>Notoscopelus bolini</i>	11	21 – 28	0.015 – 0.040	0.00000092	3.212	2.664	3.760	0.951	i
<i>Notoscopelus resplendens</i>	20	20 – 70	0.016 – 1.085	0.00000179	3.111	2.941	3.280	0.988	i
<i>Polyipnus polli</i>	34	16 – 43	0.019 – 0.627	0.00000064	3.697	3.431	3.962	0.962	a+
<i>Sternoptyx diaphana</i>	29	9 – 41	0.002 – 0.334	0.00000277	3.054	2.879	3.228	0.978	i
<i>Stomias boa boa</i>	20	53 – 153	0.041 – 1.073	0.00000019	3.081	2.855	3.305	0.979	i
<i>Sigmops elongatus</i>	35	45 – 151	0.029 – 0.962	0.00000089	2.746	2.608	2.883	0.980	a-
<i>Vinciguerrria nimbaria</i>	32	14 – 50	0.005 – 0.226	0.00000177	3.005	2.932	3.077	0.996	i

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**TABLE 3** Mean water content (and standard deviation, sd) expressed as % of eviscerated wet weight in mesopelagic fishes from the equatorial and tropical Atlantic waters

<b>Species</b>	<b>mean <math>\pm</math> sd</b>
<i>Notolychnus valdiviae</i>	58.99 $\pm$ 19.97
<i>Polyipnus polli</i>	71.31 $\pm$ 2.95
<i>Lampanyctus pusillus</i>	73.50 $\pm$ 3.73
<i>Benthoosema glaciale</i>	73.63 $\pm$ 3.59
<i>Diaphus fragilis</i>	74.67 $\pm$ 0.80
<i>Diaphus vanhoeffeni</i>	75.14 $\pm$ 1.29
<i>Myctophum nitidulum</i>	75.19 $\pm$ 2.04
<i>Diaphus problematicus</i>	75.34 $\pm$ 1.20
<i>Lobianchia dofleini</i>	75.70 $\pm$ 2.28
<i>Myctophum affine</i>	75.76 $\pm$ 2.00
<i>Diaphus dumerilii</i>	76.20 $\pm$ 1.67
<i>Vinciguerrria nimbaria</i>	76.22 $\pm$ 1.47
<i>Notoscopelus resplendens</i>	76.49 $\pm$ 1.44
<i>Lepidophanes guentheri</i>	76.50 $\pm$ 2.79
<i>Diaphus holti</i>	76.58 $\pm$ 3.44
<i>Diaphus brachycephalus</i>	76.89 $\pm$ 1.71
<i>Diaphus mollis</i>	77.49 $\pm$ 1.34
<i>Benthoosema suborbitale</i>	77.60 $\pm$ 1.47
<i>Diaphus rafinesquii</i>	77.69 $\pm$ 2.01
<i>Argyropelecus sladeni</i>	78.15 $\pm$ 2.01
<i>Lampanyctus alatus</i>	78.24 $\pm$ 2.97
<i>Sternoptyx diaphana</i>	78.51 $\pm$ 2.90
<i>Diaphus metopoclampus</i>	79.05 $\pm$ 2.86
<i>Cyclothone pseudopallida</i>	79.07 $\pm$ 2.93
<i>Bolinichthys photothorax</i>	79.37 $\pm$ 1.30
<i>Myctophum punctatum</i>	79.53 $\pm$ 2.18
<i>Cyclothone livida</i>	79.69 $\pm$ 2.02
<i>Notoscopelus bolini</i>	79.77 $\pm$ 1.13
<i>Ceratoscopelus warmingii</i>	79.84 $\pm$ 2.49
<i>Hygophum macrochir</i>	79.93 $\pm$ 1.51
<i>Cyclothone pallida</i>	80.66 $\pm$ 2.19
<i>Cyclothone acclinidens</i>	81.21 $\pm$ 2.37
<i>Lampanyctus nobilis</i>	81.54 $\pm$ 1.51
<i>Chauliodus danae</i>	82.65 $\pm$ 5.82
<i>Sigmops elongatus</i>	83.35 $\pm$ 2.42
<i>Stomias boa boa</i>	83.90 $\pm$ 1.64

**TABLE 4** Statistical comparison of Relative Condition Factor ( $K_{rel}$ ) between oceanographic regions of the study area (Olivar *et al.*, 2017) for mesopelagic fish species from the equatorial and tropical Atlantic waters. CB, Cape Blanc; CI, Canary Islands; CV, Cape Verde; E, equatorial;  $eWD$ , eviscerated dry weight (mg) ;  $LWR$ , length-weight relationship;  $n$ , number of specimens; ns, non-significant;  $KW$ , Kruskal-Wallis test; O, oligotrophic;  $sd$ , standard deviation;  $SL$ , standard length (mm); U, Mann-Whitney U test

Species	Region	n	SL range	eDW range	LWR equation	$r^2$	$K_{rel}$ mean (sd)	Statistical analysis		
								test	df	P
<i>Chauliodus dane</i>	E	12	40-176	0.024-1.758	$\log eWD = 2.867 \log SL - 6.182$	0.993	0.936 (0.090)	$t$ -test = -0.294	33	ns
	CV	23	49-176	0.047-3.520						
<i>Ceratoscopelus warmingii</i>	E	23	16-64	0.071-0.796	$\log eWD = 3.610 \log SL - 6.581$	0.992	1.072 (0.185)	$U = 129, z = -1.552$		ns
	CV	16	36-64	0.105-0.768						
<i>Diaphus brachycephalus</i>	O	34	30-53	0.101-0.606	$\log eWD = 3.264 \log SL - 5.767$	0.972	0.792 (0.067)	$t$ -test = -6.817	84	< 0.0001
	E	53	16-53	0.015-0.750						
<i>Diaphus mollis</i>	E	12	34-52	0.134-0.572	$\log eWD = 3.182 \log SL - 5.748$	0.985	0.914 (0.076)	$t$ -test = 1.169	20	ns
	CV	10	29-54	0.074-0.566						
<i>Diaphus rafinesquii</i>	CB	90	30-70	0.101-1.425	$\log eWD = 3.079 \log SL - 5.573$	0.984	1.120 (0.094)	$t$ -test = 0.063*	47	ns
	CI	34	30-58	0.118-0.773						
<i>Lepidophanes guentheri</i>	O	16	19-44	0.008-0.137	$\log eWD = 3.479 \log SL - 6.513$	0.984	0.998 (0.187)	$t$ -test = -1.312	28	ns
	E	14	34-61	0.077-0.522						
<i>Myctophum nitidulum</i>	CV	10	15-74	0.101-0.238	$\log eWD = 3.308 \log SL - 6.074$	0.984	1.070 (0.102)	$U = 72, z = -0.166$		ns
	CI	15	18-57	0.118-0.231						
<i>Vinciguerria nimbaria</i>	CV	17	37-50	0.101-0.238	$\log eWD = 3.005 \log SL - 5.752$	0.996	0.891 (0.083)	$t$ -test = 0.088	30	ns
	CB	15	14-20	0.118-0.231						
<i>Lampanyctus alatus</i>	E	22	27-48	0.027-0.217	$\log eWD = 3.594 \log SL - 6.648$	0.972	1.067 (0.151)	$t$ -test = 0.088	30	ns
	CV	69	16-57	0.006-0.358			1.103 (0.152)			
	CB	25	34-51	0.072-0.363			1.195 (0.149)			
	CI	26	39-53	0.150-0.347			1.234 (0.171)			

\* $t$ -test for unequal variance

**TABLE S1** Length-weight parameters for mesopelagic species (ordered alphabetically) from the equatorial and tropical Atlantic waters. *a*, intercept; *a*+, positive allometry; *a*-, negative allometry; *b*, allometry coefficient (slope); *WW*, whole body wet weight; *i*, isometry; *n*, sample size; *SL*, standard length; *r*<sup>2</sup>, correlation of determination; 95% CL of *b*, confidence interval

Species	n	Range <i>SL</i> (mm)	Range <i>WW</i> (g)	<i>a</i>	<i>b</i>	Inferior 95% CL of <i>b</i>	Superior 95% CL of <i>b</i>	<i>r</i> <sup>2</sup>	Growth model
<i>Argyroleucus sladeni</i>	27	15 – 43	0.061 – 1.054	0.00000925	3.223	2.692	3.753	0.862	i
<i>Benthoosema glaciale</i>	15	15 – 35	0.034 – 0.597	0.00000539	3.251	3.100	3.400	0.989	a+
<i>Benthoosema suborbitale</i>	29	16 – 33	0.069 – 0.527	0.00001615	2.982	2.592	3.371	0.943	i
<i>Bolinichthys photothorax</i>	6	21 – 51	0.126 – 1.625	0.00001123	3.022	2.607	3.435	0.990	i
<i>Ceratoscopelus warmingii</i>	47	16 – 64	0.038 – 3.865	0.00000442	3.284	3.195	3.371	0.992	a+
<i>Chauliodus danae</i>	43	25 – 212	0.062 – 29.083	0.00000267	2.976	2.868	3.083	0.987	i
<i>Cyclothone acclinidens</i>	22	31 – 46	0.039 – 0.340	0.00004436	2.243	1.779	2.705	0.772	a-
<i>Cyclothone livida</i>	32	23 – 37	0.040 – 0.195	0.00000599	2.846	2.340	3.351	0.815	i
<i>Cyclothone pallida</i>	24	25 – 54	0.051 – 0.434	0.00001276	2.616	2.210	3.020	0.891	i
<i>Cyclothone pseudopallida</i>	19	23 – 35	0.030 – 0.087	0.00005320	2.016	1.356	2.675	0.710	a-
<i>Diaphus brachycephalus</i>	100	12 – 47	0.014 – 3.557	0.00000906	3.243	3.123	3.362	0.967	a+
<i>Diaphus dumerilii</i>	30	31 – 57	0.423 – 2.681	0.00001689	2.981	2.763	3.199	0.966	i
<i>Diaphus fragilis</i>	10	46 – 79	0.871 – 4.415	0.00000971	2.976	2.798	3.152	0.995	i
<i>Diaphus holti</i>	91	11 – 50	0.020 – 2.367	0.00001677	3.006	2.887	3.124	0.966	i
<i>Diaphus metopoclampus</i>	10	19 – 40	0.066 – 0.943	0.00000398	3.353	2.914	3.791	0.975	i
<i>Diaphus mollis</i>	30	29 – 55	0.378 – 2.883	0.00001912	2.972	2.814	3.129	0.982	i
<i>Diaphus problematicus</i>	9	45 – 72	0.971 – 3.568	0.00001890	2.834	2.498	3.168	0.983	i
<i>Diaphus rafinesquii</i>	125	11 – 70	0.027 – 6.080	0.00003096	2.850	2.788	2.913	0.985	a-
<i>Diaphus vanhoeffeni</i>	17	25 – 36	0.265 – 0.691	0.00007869	2.530	2.088	2.972	0.908	a-
<i>Hygophum macrochir</i>	30	20 – 42	0.115 – 1.192	0.00002150	2.926	2.772	3.079	0.982	i
<i>Lampanyctus alatus</i>	142	16 – 57	0.047 – 1.795	0.00000488	3.199	3.106	3.290	0.971	a+
<i>Lampanyctus nobilis</i>	19	28 – 74	0.150 – 3.349	0.00000431	3.167	2.945	3.388	0.982	i
<i>Lampanyctus pusillus</i>	22	24 – 36	0.085 – 0.517	0.00000136	3.573	3.063	4.081	0.915	a+
<i>Lepidophanes guentheri</i>	35	19 – 61	0.056 – 2.564	0.00000642	3.108	2.932	3.282	0.975	i
<i>Lobianchia dofleini</i>	30	13 – 30	0.036 – 0.508	0.00001128	3.130	2.811	3.449	0.935	i
<i>Myctophum affine</i>	34	14 – 47	0.025 – 1.910	0.00000371	3.375	3.109	3.640	0.955	a+
<i>Myctophum nitidulum</i>	25	15 – 74	0.025 – 5.943	0.00000350	3.334	3.209	3.458	0.993	a+
<i>Myctophum punctatum</i>	28	16 – 69	0.042 – 4.582	0.00000545	3.221	3.131	3.311	0.995	a+
<i>Notolychnus valdiviae</i>	27	15 – 22	0.019 – 0.123	0.00000181	3.484	2.342	3.859	0.410	i
<i>Notoscopelus bolini</i>	11	21 – 28	0.085 – 0.221	0.00000719	3.101	3.484	1.720	0.905	i
<i>Notoscopelus resplendens</i>	20	20 – 70	0.065 – 4.832	0.00000837	3.108	2.894	3.320	0.981	i
<i>Polyipnus polli</i>	34	16 – 43	0.072 – 2.313	0.00000227	3.723	3.463	3.982	0.964	a+
<i>Sternoptyx diaphana</i>	29	9 – 41	0.014 – 1.861	0.00001472	3.104	2.900	3.308	0.973	i
<i>Stomias boa boa</i>	20	53 – 153	0.330 – 8.212	0.00000158	3.042	2.781	3.302	0.971	i
<i>Sigmops elongatus</i>	35	45 – 151	0.165 – 7.879	0.00000075	3.205	3.021	3.387	0.975	a+
<i>Vinciguerria nimbaria</i>	32	14 – 50	0.027 – 1.339	0.00000416	3.245	3.156	3.333	0.995	a+

**TABLE S2** Length-weight parameters for mesopelagic species (ordered alphabetically) from the equatorial and tropical Atlantic waters. *a*, intercept; *a*<sup>+</sup>, positive allometry; *a*<sup>-</sup>, negative allometry; *b*, allometry coefficient (slope); *WW*, eviscerate body wet weight; *i*, isometry; *n*, sample size; *SL*, standard length; *r*<sup>2</sup>, correlation of determination; 95% CL of *b*, confidence interval

Species	n	Range <i>SL</i> (mm)	Range <i>eWW</i> (g)	<i>a</i>	<i>b</i>	Inferior 95% CL of <i>b</i>	Superior 95% CL of <i>b</i>	<i>r</i> <sup>2</sup>	Growth model
<i>Argyropelecus sladeni</i>	27	15 – 43	0.053 – 0.910	0.00000844	3.208	2.681	3.735	0.863	i
<i>Benthoosema glaciale</i>	15	15 – 35	0.027 – 0.528	0.00000367	3.323	3.179	3.467	0.991	a+
<i>Benthoosema suborbitale</i>	29	16 – 33	0.060 – 0.478	0.00001402	2.993	2.584	3.401	0.943	i
<i>Bolinichthys photothorax</i>	6	21 – 51	0.116 – 1.520	0.00000994	3.032	2.611	3.452	0.990	i
<i>Ceratoscopelus warmingii</i>	47	16 – 64	0.033 – 3.368	0.00000364	3.303	3.214	3.391	0.992	a+
<i>Chauliodus danae</i>	43	25 – 212	0.059 – 22.652	0.00000266	2.957	2.855	3.059	0.988	i
<i>Cyclothone acclinidens</i>	22	31 – 46	0.033 – 0.274	0.00004776	2.183	1.738	2.635	0.774	a-
<i>Cyclothone livida</i>	32	23 – 37	0.039 – 0.146	0.00001491	2.538	2.032	3.045	0.777	i
<i>Cyclothone pallida</i>	24	25 – 54	0.047 – 0.380	0.00001526	2.540	2.135	2.944	0.885	a-
<i>Cyclothone pseudopallida</i>	19	23 – 35	0.027 – 0.083	0.00005324	1.986	1.178	2.794	0.613	a-
<i>Diaphus brachycephalus</i>	100	12 – 47	0.011 – 3.175	0.00000692	3.285	3.164	3.406	0.967	a+
<i>Diaphus dumerilii</i>	30	31 – 57	0.387 – 2.399	0.00001871	2.928	2.716	3.140	0.966	i
<i>Diaphus fragilis</i>	10	46 – 79	0.748 – 3.857	0.00000711	3.019	2.799	3.239	0.992	i
<i>Diaphus holti</i>	91	11 – 50	0.012 – 2.102	0.00000965	3.134	3.002	3.265	0.962	a+
<i>Diaphus metopoclampus</i>	10	19 – 40	0.064 – 0.802	0.00000330	3.349	2.979	3.718	0.982	i
<i>Diaphus mollis</i>	30	29 – 55	0.358 – 2.678	0.00001476	3.016	2.857	3.175	0.983	i
<i>Diaphus problematicus</i>	9	45 – 72	0.869 – 3.234	0.00001349	2.888	2.473	3.304	0.975	i
<i>Diaphus rafinesquii</i>	125	11 – 70	0.026 – 5.829	0.00002696	2.864	2.802	2.925	0.986	a-
<i>Diaphus vanhoeffeni</i>	17	25 – 36	0.248 – 0.584	0.00018969	2.238	1.740	2.735	0.860	a-
<i>Hygophum macrochir</i>	30	20 – 42	0.097 – 1.047	0.00001658	2.964	2.806	3.122	0.981	i
<i>Lampanyctus alatus</i>	142	16 – 57	0.040 – 1.720	0.00000418	3.214	3.121	3.306	0.971	a+
<i>Lampanyctus nobilis</i>	19	28 – 74	0.128 – 3.097	0.00000311	3.224	2.965	3.483	0.976	i
<i>Lampanyctus pusillus</i>	22	24 – 36	0.079 – 0.448	0.00000230	3.389	2.916	3.861	0.918	i
<i>Lepidophanes guentheri</i>	35	19 – 61	0.025 – 2.331	0.00000156	3.443	3.212	3.674	0.965	a+
<i>Lobianchia dofleini</i>	30	13 – 30	0.030 – 0.464	0.00000849	3.184	2.855	3.512	0.934	i
<i>Myctophum affine</i>	34	14 – 47	0.021 – 1.491	0.00000254	3.440	3.291	3.589	0.986	a+
<i>Myctophum nitidulum</i>	25	15 – 74	0.021 – 5.137	0.00000243	3.400	3.281	3.518	0.994	a+
<i>Myctophum punctatum</i>	28	16 – 69	0.036 – 3.863	0.00000438	3.232	3.137	3.326	0.995	a+
<i>Notolychnus valdiviae</i>	27	15 – 22	0.019 – 0.123	0.00000278	3.303	2.498	3.759	0.424	i
<i>Notoscopelus bolini</i>	11	21 – 28	0.077 – 0.195	0.00000594	3.129	3.302	1.587	0.933	i
<i>Notoscopelus resplendens</i>	20	20 – 70	0.060 – 4.427	0.00000756	3.114	2.908	3.320	0.983	i
<i>Polyipnus polli</i>	34	16 – 43	0.060 – 2.070	0.00000207	3.719	3.453	3.985	0.962	a+
<i>Sternoptyx diaphana</i>	29	9 – 41	0.009 – 1.604	0.00000729	3.236	3.038	3.434	0.977	a+
<i>Stomias boa boa</i>	20	53 – 153	0.312 – 7.306	0.00000159	3.013	2.739	3.292	0.967	i
<i>Sigmops elongatus</i>	35	45 – 151	0.147 – 7.149	0.00000064	3.220	3.064	3.376	0.982	a+
<i>Vinciguerrria nimbaria</i>	32	14 – 50	0.023 – 0.964	0.00000594	3.073	2.989	3.158	0.995	i