



- 1 The Pelagic In situ Observation System (PELAGIOS) to reveal
- 2 biodiversity, behavior and ecology of elusive oceanic fauna
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12 **1. Abstract**

13 There is a need for cost-efficient tools to explore deep ocean ecosystems to collect baseline
14 biological observations on pelagic fauna (zooplankton and nekton) and establish the vertical
15 ecological zonation in the deep sea. The Pelagic In situ Observation System (PELAGIOS) is a
16 3000 m-rated slowly (0.5 m/s) towed camera system with LED illumination, an integrated
17 oceanographic sensor set (CTD-O₂) and telemetry allowing for online data acquisition and video
18 inspection (Low Definition). The High Definition video is stored on the camera and later annotated
19 using the VARS annotation software and related to concomitantly recorded environmental data.
20 The PELAGIOS is particularly suitable for open ocean observations of gelatinous fauna, which is
21 notoriously undersampled by nets and/or destroyed by fixatives. In addition to counts, diversity
22 and distribution data as a function of depth and environmental conditions (T, S, O₂), in situ
23 observations of behavior, orientation and species interactions are collected. Here we present an
24 overview of the technical setup of the PELAGIOS as well as example observations and analyses
25 from the eastern tropical North Atlantic. Comparisons to MOCNESS net sampling and data from
26 the Underwater Vision Profiler are provided and discussed.

27

28 **2. Introduction**

29 The open ocean pelagic zones include the largest, yet least explored habitats on the planet
30 (Robison, 2004; Webb et al., 2010; Ramirez-Llodra et al., 2010). Since the first oceanographic
31 expeditions, oceanic communities of macrozooplankton and micronekton have been sampled
32 using nets. Such sampling has revealed a community typically consisting of crustaceans,
33 cephalopods, fishes and some sturdy and commonly found gelatinous fauna. Underwater
34 observations in the open ocean via SCUBA diving (Hamner et al., 1975) and later via submersibles



35 (Robison, 1983; Robison and Wishner, 1990) and *in situ* camera systems (Biard et al., 2016,
36 Picheral et al. 2010) revealed that a variety of organisms are much more abundant in the open
37 ocean than previously estimated from net sampling (Robison, 2004, Haddock, 2004; Biard et al.
38 2016, Christiansen et al. 2018). This was particularly true for fragile gelatinous zooplankton, a
39 diverse taxonomic group of different phyla, including the ctenophores, medusae, siphonophorae,
40 thaliaceans, polychaetes, rhizaria and larvaceans, which often are too delicate to be quantified
41 using nets as they are damaged beyond identification, or they are easily destroyed by the use of
42 common fixatives.

43 Underwater (*in situ*) observations in the pelagic ocean not only revealed a previously unknown
44 community, they also allowed the collection of fine scale distribution patterns, information on
45 posture, interactions, and behavior and a better understanding of the ecological context and role of
46 pelagic organisms (Hamner and Robison, 1992; Robison, 2004; Robison, 1999; Biard et al., 2016;
47 Hoving et al., 2017). Submersibles have proven to be valuable instruments to study deep-sea
48 pelagic biology (e.g. Robison, 1987; Bush et al., 2007; Hoving et al., 2013; 2016). Using video
49 transecting methodology, pelagic ROV surveys have been applied to study inter and intra-annual
50 variation in mesopelagic zooplankton communities (Robison et al., 1998) and to explore deep
51 pelagic communities in different oceans (Youngbluth et al., 2008; Hosia et al., 2017; Robison et
52 al., 2010). However, due to high costs as well as technological and logistical challenges, regular
53 submersible operations are still restricted to very few institutes and geographical locations. Hence,
54 there is a need for the development of additional more cost-effective methodologies to explore and
55 document deep-sea communities via *in situ* observations.

56 In the last decades, a variety of optical instruments has been developed to image and quantify
57 plankton *in situ*. The factors that typically differentiate the available plankton imaging



58 technologies are the size fraction of the observed organisms, illumination type, resolution of
59 collected images/video, depth rating, deployment mode (e.g., autonomous, towed, CTD-mounted)
60 and towing speed (Benfield et al., 2007). Examples of instruments include the autonomous
61 Underwater Vision Profiler (UVP) (Picheral et al., 2010), the Lightframe On-sight Key species
62 Investigations (LOKI; Schulz et al 2009) and towed plankton recorders (ISiS; Cowen and Guigand
63 2008; for review see Benfield et al 2007). These instruments can be deployed from ships of
64 opportunity and collect detailed information on fine scale distribution and diversity patterns of
65 particles and plankton. The data reveal biological patterns on a global scale (Kiko et al., 2017) and
66 of previously underappreciated plankton species (Biard et al., 2016).

67 Various towed camera platforms have been developed that can obtain video transect observations
68 above the deep sea floor. Examples are the TowCam (WHOI), the DTIS (Deep Towed Imaging
69 system, NIWA), the WASP vehicle (Wide Angle Seafloor Photography), OFOS (Ocean Floor
70 Observation System, GEOMAR), and the more recent version OFOBS (Ocean Floor Observation
71 and Bathymetry System) (Purser et al., 2018). All these instruments are used for video or photo
72 transects of the seafloor, with a downward looking camera, and typically a set of lasers for size
73 reference. However, published descriptions of optical systems, other than ROVs and submersibles,
74 that visualize macrozooplankton and micronekton (>1 cm) in the water column are, to the best of
75 our knowledge, restricted to one (Madin et al., 2006). The Large Area Plankton Imaging System
76 (LAPIS) is the only towed system that was developed for the documentation of larger organisms
77 in the water column (Madin et al., 2006). LAPIS visualizes organisms between 1 and 100 cm, it
78 combines a low light camera with red illumination, and it is towed at 1 knot via a conducting wire.
79 Deployments in the Southern Ocean enabled the reconstruction of depth distributions of the pelagic



80 fauna (salps, medusae) but also allowed behavior observations, e.g. the moulting of krill (Madin
81 et al., 2006).

82 To establish a baseline in abundance, distribution and diversity of the pelagic fauna in its natural
83 environment, we developed an ocean observation platform for pelagic video transects. The
84 functional requirements for the instrument were the ability to: (1) visualize organisms > 1 cm in
85 waters down to 1000 m, (2) deploy the instrument from ships of opportunity in an autonomous or
86 transmitting mode, (3) make it light and practical so it can be deployed easily and safe with 2 deck
87 persons and a crane operator, (4) enable correlation of observations with environmental parameters
88 (S, T, O₂) and other sensor data, and (5) make observations comparable to ROV video transects in
89 other reference areas. We present a description of the Pelagic In situ Observation System
90 (PELAGIOS), examples of the kind of biological information it may gather, as well as biological
91 discoveries that have resulted from deployments on research cruises in the eastern tropical
92 Atlantic.

93

94 **3. Pelagic In Situ Observation System**

95 **3.1 Technical Specifications**

96 The PELAGIOS consists of an aluminum frame (length = 2 m) that carries the oceanographic
97 equipment (Figure 1). White light LED arrays (4 LEDs produced at GEOMAR, 2 LED arrays type
98 LightSphere of Deep-Sea Power and Light ©) which illuminate the water in front of the system
99 are mounted on an aluminum ring (diameter = 1.2 m). Power is provided by two lithium batteries
100 (24V; 32 Ah) in a deep-sea housing. High-definition video is collected continuously by a forward
101 viewing deep-sea camera (type 1Cam Alpha, SubC Imaging ©) which is mounted in the center of
102 the ring. We used the maximum frame rate of 50 frames s⁻¹ but a lower frame rate is possible. A



103 CTD (SBE 19 SeaCAT, Sea-Bird Scientific) with an oxygen sensor (SBE 43, Sea-Bird Scientific)
104 records environmental data. A deep-sea telemetry (DST-6, Sea and Sun Technology ©; Linke et
105 al., 2015) transmits video and CTD data to a deck unit on board allowing a low resolution preview
106 (600x480 lines) of the high definition video that is stored locally on the SD card (256 GB) of the
107 camera. The power from the batteries is distributed to the LEDs via the camera. The 1Cam Alpha
108 camera is programmable in such a way that there is a delay between providing power to the camera
109 (by connecting to the battery) and the start of recording and switching on the LEDs. This enables
110 the illumination to be turned on only underwater, and prevents overheating of the LED arrays
111 while out of the water. During a cruise with the German research vessel MARIA S. MERIAN
112 (MSM 49) we mounted a steel scale bar in front of the camera at a distance of 1 m. The distance
113 between the centers of the white marks on the bar measured 5 cm.

114 **3.2 Video transects**

115 The PELAGIOS is towed horizontally at specified depths of 20-1000 m. The standard towing
116 speed over ground is 1 knot (0.5 m/s), and the speed is monitored via the ship's navigational
117 system. A video transect at a particular depth can take as long as desired and is terminated by
118 lowering the PELAGIOS to the next desired depth. Maximum deployment time with full batteries
119 is approximately 6 hours. The typical transect duration is 10-30 min. The depth of the PELAGIOS
120 can be monitored via online CTD data. Figure 2 shows the trajectories of the PELAGIOS at
121 different depths in the water column during a video transect down to 700 m. The deployment from
122 deck into the water and the reverse is fast and typically takes only about 5 min (see video clip:
123 [https://www.wissenschaftsjahr.de/2016-17/das-wissenschaftsjahr/die-](https://www.wissenschaftsjahr.de/2016-17/das-wissenschaftsjahr/die-forschungsflotte/forschungsschiff-blogs/unerforschte-meeresgebiete.html)
124 [forschungsschiff-blogs/unerforschte-meeresgebiete.html](https://www.wissenschaftsjahr.de/2016-17/das-wissenschaftsjahr/die-forschungsflotte/forschungsschiff-blogs/unerforschte-meeresgebiete.html)). It is possible to deploy
125 PELAGIOS in 'blind mode', where only the depth is monitored using an online depth sensor (e.g.,



126 Hydrobios ©) and the video (without transmitted preview) is recorded locally on the camera. The
127 system can be operated completely blind (i.e., with no communication between deck and
128 underwater unit) where the target depth is estimated from the length and angle of the wire put out,
129 and the actual depth is recorded on the system by an offline pressure sensor e.g. SBE Microcat ©.
130

131 **3.3 Video analysis and curation**

132 After a deployment, the video (consisting of individual clips of one hour) is downloaded from the
133 camera. Synchronisation between video and CTD data is done by setting all instruments to UTC
134 prior to deployment, which allows the data and video to be linked during analysis. The video is
135 annotated using the Video Annotation and Reference System VARS developed by the Monterey
136 Bay Aquarium Research Institute (Schlining and Jacobsen, 2006). This annotation program allows
137 for frame grabs from the video including time code. A Knowledge Base allows for inserting
138 taxonomic names and hierarchy, and a Query allows for searching the created database. While
139 many kinds of annotation software are available (for review see Gomes-Pereira et al., 2016), we
140 consider VARS the most suitable for our purposes since it combines the features of high resolution
141 video play with a user friendly annotation-interface and the automatic creation of an annotation
142 database which can easily be accessed through the various search-functions and tools of the Query.
143 The taxonomic hierarchy and phylogenetic trees in the database are directly applicable to our video
144 transects. Since this software was developed by MBARI, which also maintains the most extensive
145 databases of deep pelagic observations, it makes communication about and comparison of
146 observations and data practical. Videos are transported on hard drives after an expedition. At
147 GEOMAR, videos are transferred for long term storage on servers maintained by the central data



148 and computing centre at GEOMAR, providing instant access to videos and images with metadata
149 description via the media server ProxSys.

150 .

151 **3.4 Sample volume**

152 To estimate the sample volume of the PELAGIOS we compared video counts from the PELAGIOS
153 with concomitantly obtained abundance data from an Underwater Vision Profiler (UVP5; Picheral
154 et al., 2010). Four deployments from the R/V Maria S. Merian cruise MSM 49 (28.11. -
155 21.12.2015, Las Palmas de Gran Canaria/Spain – Mindelo/Cape Verde) were used for the
156 comparison where a UVP5 was mounted underneath the PELAGIOS (Figure 1). The UVP5 takes
157 between 6-11 images per second of a defined volume (1.03 L) and thus enables a quantitative
158 assessment of particle and zooplankton abundances. Objects with an equivalent spherical diameter
159 (ESD) >0.5 mm are saved as images, which can be classified into different zooplankton,
160 phytoplankton and particle categories. For the comparison between PELAGIOS and the UVP5,
161 we used the pelagic polychaete *Poeobius* sp., as 1) this organism could be observed well on both
162 instruments, 2) *Poeobius* sp. is not an active swimmer and 3) it was locally very abundant, thus
163 providing a good basis for the direct instrument comparison.

164 The UVP5 images were classified as described in Christiansen et al. (2018). *Poeobius* sp.
165 abundance (ind m⁻³) was calculated for 20 s time bins and all bins of one distinct depth step
166 averaged. These mean abundances were compared to the PELAGIOS counts (ind s⁻¹) of the same
167 depth step. A linear model between the PELAGIOS counts as a function of UVP5 abundance
168 provided a highly significant relationship (linear regression: $p < 0.001$, $adjusted\ r^2 = 0.69$; Figure
169 3). The linear regression slope b (0.116 m³ s⁻¹, standard error 0.01 m³ s⁻¹) between the PELAGIOS-
170 based count ($C_{PELAGIOS}$, ind s⁻¹) and mean UVP-based abundance (A_{UVP} , ind m⁻³):



171
$$C_{PELAGIOS} = b * A_{UVP} + a \quad (\text{Equation 1})$$

172 was used to estimate the volume recorded per time in $\text{m}^3 \text{s}^{-1}$ (b) and the field of view in m^2
173 ($b/\text{towing speed}$) recorded by PELAGIOS.

174 From this calculation it can be derived that PELAGIOS recorded an average volume of $0.116 \text{ m}^3 \text{ s}^{-1}$
175 ¹ at a towing speed of 1 knot ($= 0.5144 \text{ m s}^{-1}$). A cross-sectional view field of approximately 0.23
176 m^2 of PELAGIOS can be expected.

177

178 **3.5 Abundance, size and diversity at an example station “Senghor NW”**

179 To provide an example of the type of data that can be obtained with the PELAGIOS, we report
180 here on day and night video transects down to 950 m in the Eastern Tropical North Atlantic, on
181 the northwestern slope of Senghor Seamount ($17^{\circ}14.2'N$, $22^{\circ}00.7'W$; bottom depth of
182 approximately 1000 m). The results from the video annotations show that faunal abundances
183 depend on the depth of deployment, and time of the day. During two transects of 11 minutes at
184 400 m, 232 individuals were encountered during the day (the three dominant organism groups are
185 fish, euphausiids and appendicularians) compared to 208 individuals during the night (the four
186 dominant organism groups are fishes, chaetognaths, medusae and ctenophores). Overall
187 abundance of chaetognaths, decapods and mysids, and somewhat for fishes was higher during the
188 night. The peak of euphausiids' abundance at 400 m shifts to the surface at night (Figure 4). The
189 higher abundance of decapods, mysids and chaetognaths at night may indicate lateral migration or
190 daytime avoidance. The vertical migration that was observed for fishes and crustaceans was much
191 less clear for the gelatinous zooplankton groups including the medusae and appendicularians
192 (Figure 4). Ctenophores and siphonophores were abundant in the surface at night and the
193 thaliaceans migrated vertically and were most abundant in shallow waters at night. The total



194 number of annotated organisms for the daytime transects (total transect time 182 minutes; max.
195 depth 950 m) was 835 compared to 1865 organisms for the nighttime transects. Remarkable is the
196 enormous abundance of gelatinous zooplankton (129 annotated organisms belonging to the three
197 dominant groups of Ctenophora (53), Siphonophorae (32) and Thaliacea (44) in the topmost layer
198 (20 m) at night. Below this layer, the depth profile shows a minimum in numbers of annotated
199 individuals at 100, 200, and 300 m water depth with a smaller peak of 56 gelatinous organisms in
200 450 m. Compared to this, the depth distribution at day time shows a more regular, almost Gaussian
201 shape with a maximum of 47 and 54 gelatinous organisms at 200 and 400 m water depth,
202 respectively.

203 The faunal observations at station Senghor NW include a wide variety of taxa (Table 1). The
204 smallest annotated specimens belonged to the radiolarians. Chaetognaths were the dominant faunal
205 group. Large larvaceans tentatively identified to belong to the genus *Bathochordaeus* and
206 *Mesochordaeus* were also observed. Pelagic polychaetes of the genus *Poeobius* can be easily
207 distinguished and are up to 23 mm long (Christiansen et al., 2018). Other pelagic worms are
208 tomopterid and alciopid worms, the latter can reach 1 m in length. The faunal group with the largest
209 specimens, attaining up to several metres in length, are the siphonophores, including *Praya dubia*
210 and *Apolemia*. Siphonophores of the genus *Bargmannia* and *Lilyopsis* were also observed.
211 *Lilyopsis* can be easily distinguished by their fluorescent body parts. Observed medusae belonged
212 to the genera *Periphylla*, *Halitrephes*, *Haliscera*, *Crossota*, *Colobonaema*, *Solmissus* and
213 *Solmundella*. Lobate ctenophores such as *Thalassocalyce inconstans*, *Leucothea*, *Bathyceroe* are
214 typical examples of organisms that cannot be captured by nets but which can be properly quantified
215 by PELAGIOS. Venus girdles (*Cestum* spp.), *Beroe* and cydippids are other ctenophores that were
216 encountered at Senghor NW. Cephalopod observations are rare but small individual cranchid



217 squids were observed in the upper 50 m at night. Mastigoteuthid squids were observed with their
218 mantle in a vertical orientation and with extended tentacles in waters below 500 m. One *Taningia*
219 *danae* was observed during a transit between transecting depths. Other pelagic molluscs include
220 the nudibranch *Phylliroe* and different pteropod species. Observed fishes are snipe eels, hatchet
221 fishes, lantern fishes and *Cyclothone*. Fishes are among the dominant organisms encountered
222 during PELAGIOS transects but it is often impossible to identify fishes to species level from the
223 video.

224 We compared PELAGIOS video transects with MOCNESS net (opening 1 m²) abundance data by
225 integrating the PELAGIOS counts over the respective depth strata of the MOCNESS. The diversity
226 of the gelatinous zooplankton in the total MOCNESS catch is much lower (8 different taxa) than
227 in the pooled video transects (53 annotated taxa) on the same station. The ctenophore *Beroe* is
228 captured in MOCNESS hauls and also observed on PELAGIOS transects. Normalization and
229 subsequent standardization of the encountered *Beroe* in MOCNESS and PELAGIOS transects
230 shows that on the same station and the same depths, PELAGIOS observes 3.3-4.7 times more
231 *Beroe* at the three depths where they were encountered by both instruments. Additionally, the
232 PELAGIOS also repeatedly observed *Beroe* at depths where they were not captured by MOCNESS
233 at all (although there were also depths where PELAGIOS did not observe any *Beroe*).

234

235 **3.6 Individual behaviour**

236 In situ observations by PELAGIOS video may reveal direct observations on individual behavior.
237 Decapod shrimps were observed to release a blue or green bioluminescent cloud after performing
238 their tail flip as part of the escape response (Figure 5). Potential reproductive behavior was
239 observed for two specimens of krill which were seen in a mating position, and salps were observed



240 to reproduce asexually by the release of salp oozoids (Figure 5). Feeding behaviors were observed
241 for large prayid siphonophores and calycophoran siphonophores which had their tentacles
242 extended. *Poeobius* worms were observed with their mucus web deployed to capture particulate
243 matter (Christiansen et al., 2018). Narcomedusae of the genus *Solmissus* were observed with their
244 tentacles stretched up and down, which is a feeding posture (Figure 5). In situ observations by the
245 PELAGIOS also showed the natural body position of pelagic organisms. Snipe eels were observed
246 in a vertical position with their heads up, while dragonfishes and some myctophids were observed
247 in an oblique body position with their head down.

248

249 **4. Discussion**

250 PELAGIOS is a cost-effective pelagic ocean exploration tool that fills a gap in the array of
251 observation instruments that exist in biological oceanography. The instrument can be deployed
252 with a small team and from vessels of opportunity, in transmission or blind mode. The relatively
253 simple design limits technical failures and makes the PELAGIOS a reliable tool for oceanic
254 expeditions. While thus far the system has only been deployed in the open ocean, it can be used in
255 any pelagic environment with water that has reasonable clearance and visibility. The data obtained
256 after annotation of the video can be uploaded into databases (e.g., Pangaea) after publication of
257 the results allowing for efficient data sharing and curation.

258 The clear signal of the vertical migration in some animal groups (fishes, crustaceans) that we
259 observed during the video transects confirms that established biological processes can be detected
260 in PELAGIOS data, and that the distribution data that we observe for encountered organisms are
261 representative for the natural situation. It has to be noted, though, that while the observed
262 distribution patterns should be representative, care must be taken with regards to abundance



263 estimates of especially actively- and fast-swimming organisms. Some fish and crustaceans react
264 to the presence of underwater instrumentation (e.g. Stoner et al. 2008). Gear avoidance (e.g.
265 Kaartvedt et al. 2012) can lead to an underestimation of abundance, whereas attraction to the
266 camera lights (e.g. Utne-Palm et al. 2018, Wiebe et al. 2004) would result in an overestimation.
267 The large bioluminescent squid *Taningia danae* seemed to be attracted to the lights of the
268 PELAGIOS. Compared to day transects, the high abundance of gelatinous organisms close to the
269 surface during night is likely to be partly an effect of the higher contrast in the videos of the night
270 transects and better visibility of the gelatinous fauna than during day transects. Many of the
271 observed gelatinous fauna might be as well be present at shallow depths during day-light but are
272 not detectable at ‘blue-water-conditions’. The large difference between encountered taxa during
273 the day and night transect may also be explained by the lateral migration of animals towards
274 Senghor seamount at night. However, from a methodological side it should be noted that while the
275 ship’s towing speed is typically 1 knot, the current speeds at the survey depths may differ, also
276 between day and night. Currents may result in more or less sampled volume of water and hence a
277 variation in plankton being visualized. Therefore it is recommended to perform future surveys with
278 a current meter to measure the speed through water.

279 After annotation, the PELAGIOS video transects may be used to reconstruct species-specific
280 distribution patterns, which can be related to environmental gradients. Such data is valuable for
281 studies on overlap in distribution patterns of consumers and food items (e.g. *Poeobius* and
282 particles, ctenophores and krill). The data can also be used in biological studies that aim to predict
283 the consequences of a changing ocean with altering environmental gradients for species’
284 distributions. One example of changing environmental gradients is the global trend of oxygen loss
285 in the world oceans. Oxygen minimum zones (OMZ) are occurring naturally in the mesopelagic



286 zone, and in different oceans they have been found to expand horizontally and vertically as a result
287 of climate change (Stramma et al., 2008). Expansion of OMZs may result in a habitat reduction of
288 the pelagic fauna (e.g., Stramma et al., 2012), or increase the habitat for species with hypoxia
289 tolerance. To predict the potential consequences of OMZ expansion for pelagic invertebrates we
290 investigated the abundance and distribution of distinct large gelatinous zooplankton species,
291 including medusae, ctenophores, siphonophores and appendicularians, in the eastern tropical
292 Atlantic using PELAGIOS video transects and correlated the biological patterns to the oxygen
293 gradients (Neitzel, 2017; Hoving et al., in prep.).

294 Preliminary comparisons of the data obtained with PELAGIOS and with MOCNESS indicate
295 substantial differences in the documented fauna. Many more gelatinous taxa were observed during
296 PELAGIOS video transects than were captured in MOCNESS catches at the same station, with
297 the exception of the small and robust calyphoran colonies of the families Diphyidae and
298 Abylidae. This discrepancy is likely the result of the delicate nature of many ctenophores, medusae
299 and siphonophores, preventing their intact capture by nets. Additionally, avoidance behavior of
300 strong swimming jellyfish (e.g. *Atolla*, *Periphylla*), which escape from the relatively slow moving
301 PELAGIOS, may explain their increased occurrence in nets compared to video recordings. While
302 PELAGIOS is certainly suitable for visualizing delicate gelatinous fauna, it cannot replace net-
303 sampling since complementary specimen collections are needed to validate the identity of
304 organisms that were observed during PELAGIOS video observations. Therefore, it is desired that
305 net tows with open and closing nets such as Multinet Maxi or MOCNESS are performed in the
306 same areas, or that collections during submersible dives are made.

307 The potential of the PELAGIOS as an exploration tool is illustrated by the discovery of previously
308 undocumented animals. An example is the ctenophore *Kiyohimea usagi* (Matsumoto and Robison,



309 1992) which was observed seven times by the PELAGIOS and once by the manned submersible
310 JAGO during cruises in the eastern tropical Atlantic. This large (>40 cm wide) lobate ctenophore
311 was previously unknown from the Atlantic Ocean and demonstrates how in situ observations in
312 epipelagic waters can result in the discovery of relatively large fauna (Hoving et al., submitted).
313 Since gelatinous organisms are increasingly recognized as vital players in the oceanic food web
314 (Choy et al., 2017) and in the biological carbon pump (Robison et al., 2005), in situ observations
315 with tools like the PELAGIOS can provide new important insights into the oceanic ecosystem and
316 the carbon cycle. But small gelatinous organisms may also have a large biogeochemical impact on
317 their environment. This was illustrated by the discovery of the pelagic polychaete *Poeobius* sp.
318 during the PELAGIOS video transects in the eastern Atlantic (Christiansen et al., 2018). The
319 observations of the PELAGIOS provided the first evidence for the occurrence of *Poeobius* sp. in
320 the Atlantic Ocean. During the R/V Meteor cruise M119, *Poeobius* was found to be extremely
321 abundant in a mesoscale eddy. Using an extensive database of the UVP5 (956 vertical CTD/UVP
322 profiles) in the eastern tropical Atlantic it was possible to reconstruct the horizontal and vertical
323 distribution of Atlantic *Poeobius* in great detail and to establish that the high local abundance of
324 *Poeobius* was directly related to the presence of mesoscale eddies in which they possibly
325 intercepted the entire particle flux that was on the way to the deep sea (Christiansen et al., 2018;
326 Hauss et al., 2016).

327 During various cruises, the UVP 5 was mounted underneath the PELAGIOS providing
328 concomitant data on macrozooplankton and nekton (PELAGIOS) as well as particles and
329 mesozooplankton (UVP). The combination of the two instruments provides a great opportunity to
330 assess both the mesopelagic fauna and particles during one sampling event. The joint deployment
331 of the PELAGIOS and UVP also allowed a quantification of the sampled water volume of the



332 PELAGIOS as described above. The linear relationship between counts of the non-moving
333 *Poeobius* sp. with UVP5 and the PELAGIOS indicates comparability of the two different methods
334 and provides a correction factor to estimate organism abundance (ind m^{-3}) from PELAGIOS count
335 (ind s^{-1}) data. The field of view (FOV) for the PELAGIOS was estimated to be 0.23 m^2 . The angle
336 of view of the PELAGIOS is 80° and therefore the field of view (FOV) is much smaller than the
337 FOV of video transects with a wide-angle lens by ROV Tiburon (Robison et al., 2010). When
338 comparing the FOV, it is important to take into account the object that is observed. We provided
339 an estimate of the FOV using *Poeobius* sp., which is a small organism that can be detected only
340 when it is close to the camera. Therefore, the area of the FOV for quantification of *Poeobius* sp. is
341 smaller than when quantifying larger organisms, and the initial identification distance differs
342 between species (Reisenbichler et al., 2017). Future effort should be focused on improving the
343 assessment of the sample volume by integrating technology that can quantify it (e.g. current
344 meters, a stereo-camera setup or a laser-based system). A stereo-camera set up would also allow
345 for size measurements of the observed organisms, which could be beneficial to estimate the
346 biomass of the observed organisms from published size-to-weight relationships. It might also be
347 possible to obtain similar information based on structure-from-motion approaches that proved
348 successful in benthic video imaging (Burns et al., 2015). The PELAGIOS system can also be a
349 platform for other sensors. The PELAGIOS was used to mount and test the TuLUMIS
350 multispectral camera (Liu et al., 2018). Future developments include the preparation of the system
351 for deployments down to 6000 m. The integration of acoustic sensors would be valuable to measure
352 target strength of camera observed organisms, to estimate gear avoidance or attraction and to
353 estimate biomass and abundance of organisms outside the field of view of the camera.

354



355

356 **Author contribution**

357

358 This instrument was designed, tested and applied by Henk-Jan Hoving and Eduard Fabrizious.
359 Rainer Kiko and Helena Hauss developed the idea of combining the PELAGIOS with the UVP5.
360 Philipp Neitzel and Svenja Christiansen analyzed the data in this manuscript in consultation with
361 Henk-Jan Hoving, Rainer Kiko and Helena Hauss. Arne Körtzinger, Uwe Piatkowski and Peter
362 Linke added valuable input to the further development of the instrument and its application
363 and/or the data interpretation. All authors contributed to writing the paper. All authors approved
364 the final submitted manuscript.

365

366 **Data availability**

367 The datasets generated and/or analysed during the current study will be available in the Pangaea
368 repository. A link will be provided when the paper is accepted.

369

370

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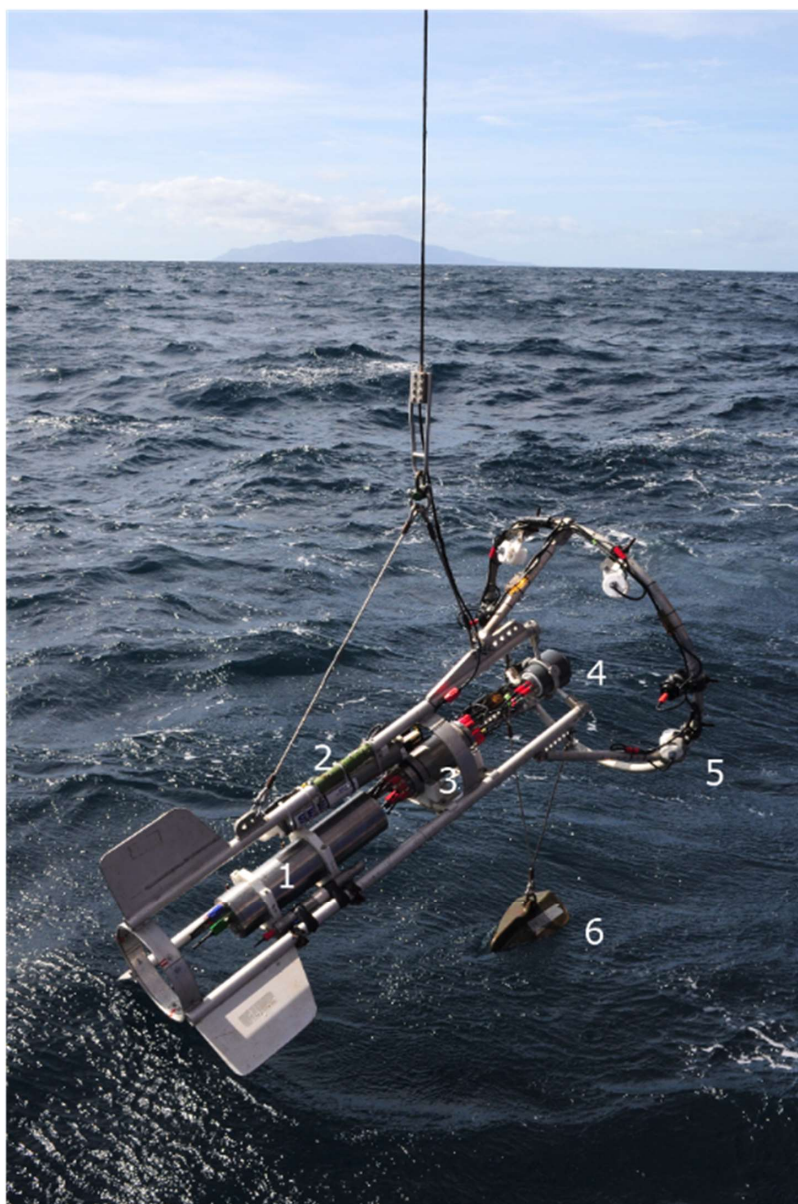
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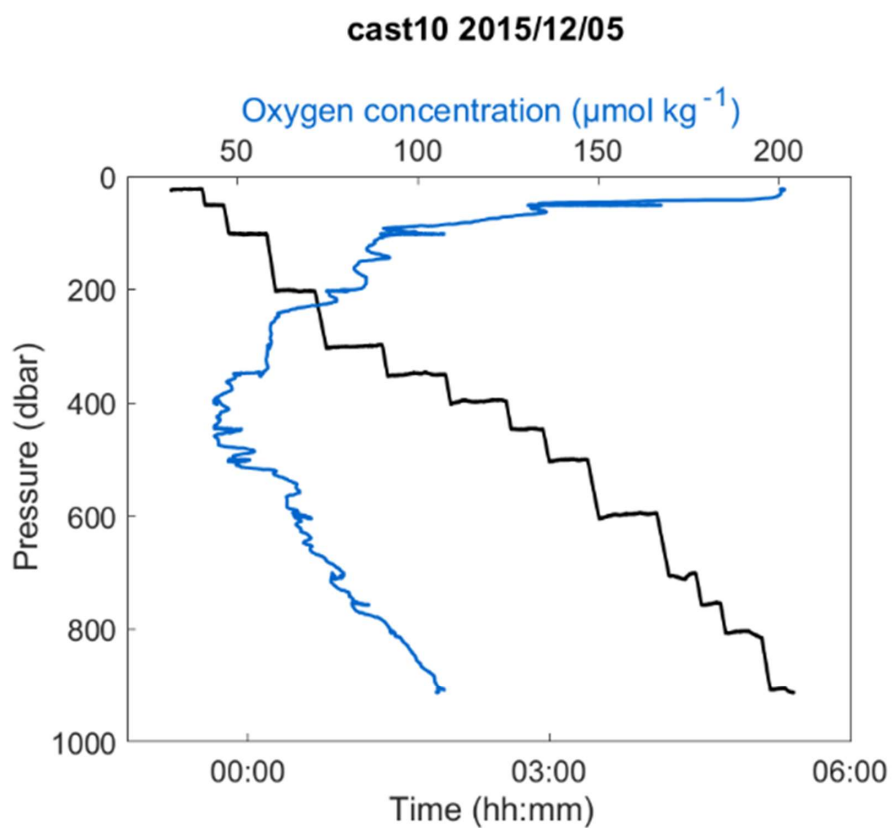
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492 Figure 1: a) The Pelagic in situ observations system with the battery (1), CTD (2), telemetry (3),

493 camera (4), LEDs (5), depressor (6), during deployment from R/V POSEIDON in February

494 2018.

495



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497 Figure 2: Stair wise trajectory of PELAGIOS through the water column, to the desired depth.

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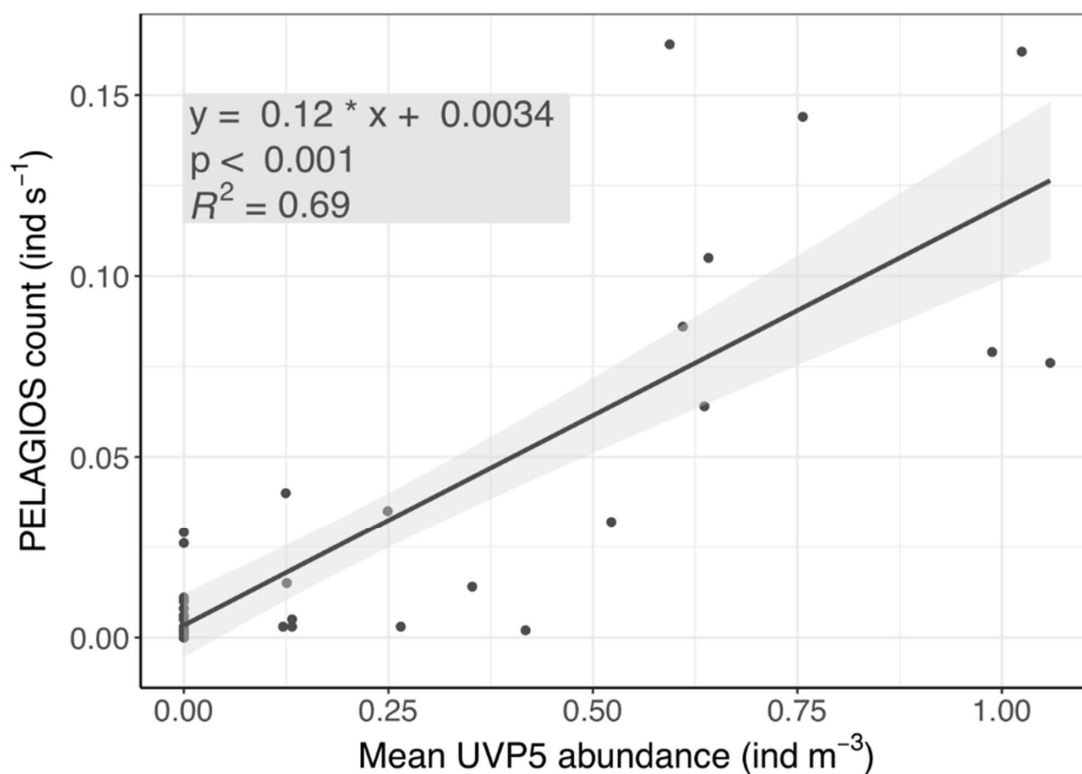
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506 Figure 3: PELAGIOS video counts of *Poeobius* sp. as a function of UVP5-derived abundance on

507 the same transects at two stations on cruise MSM 49 on RV MARIA S. MERIAN.

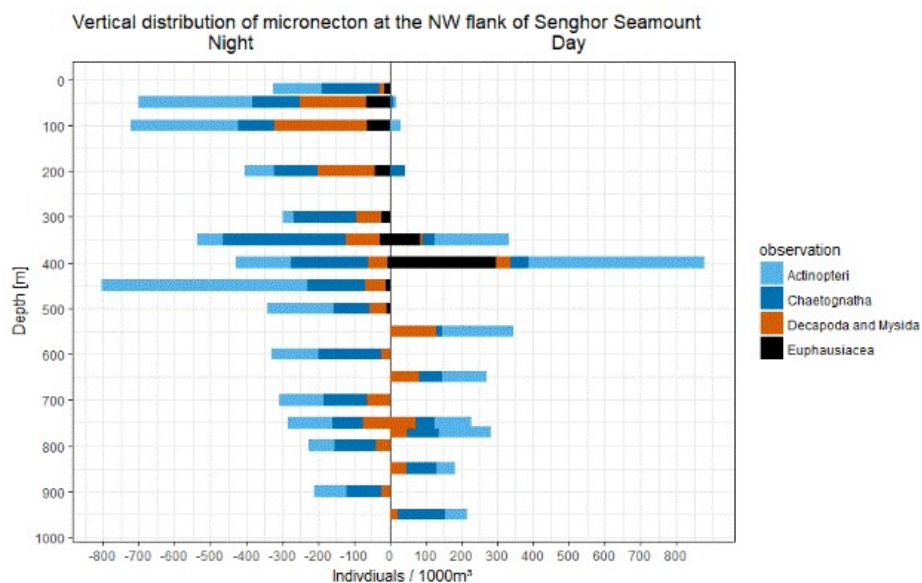
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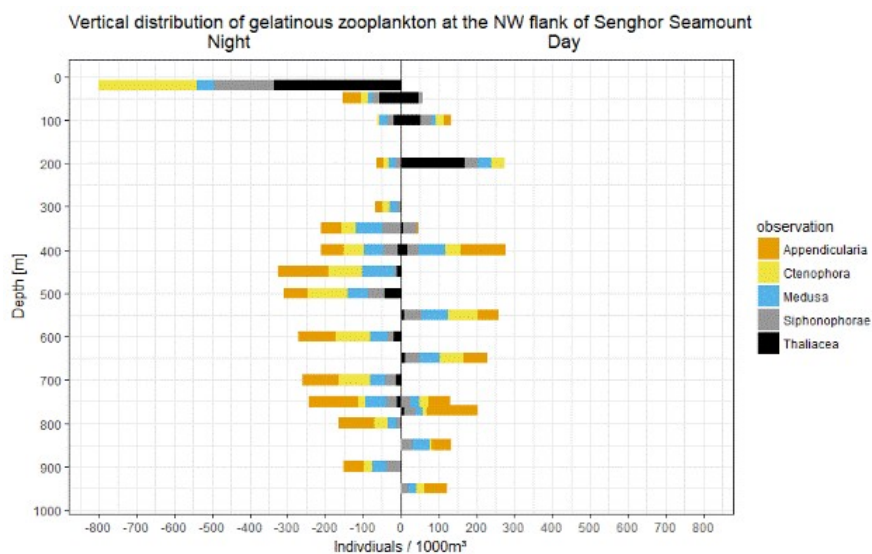
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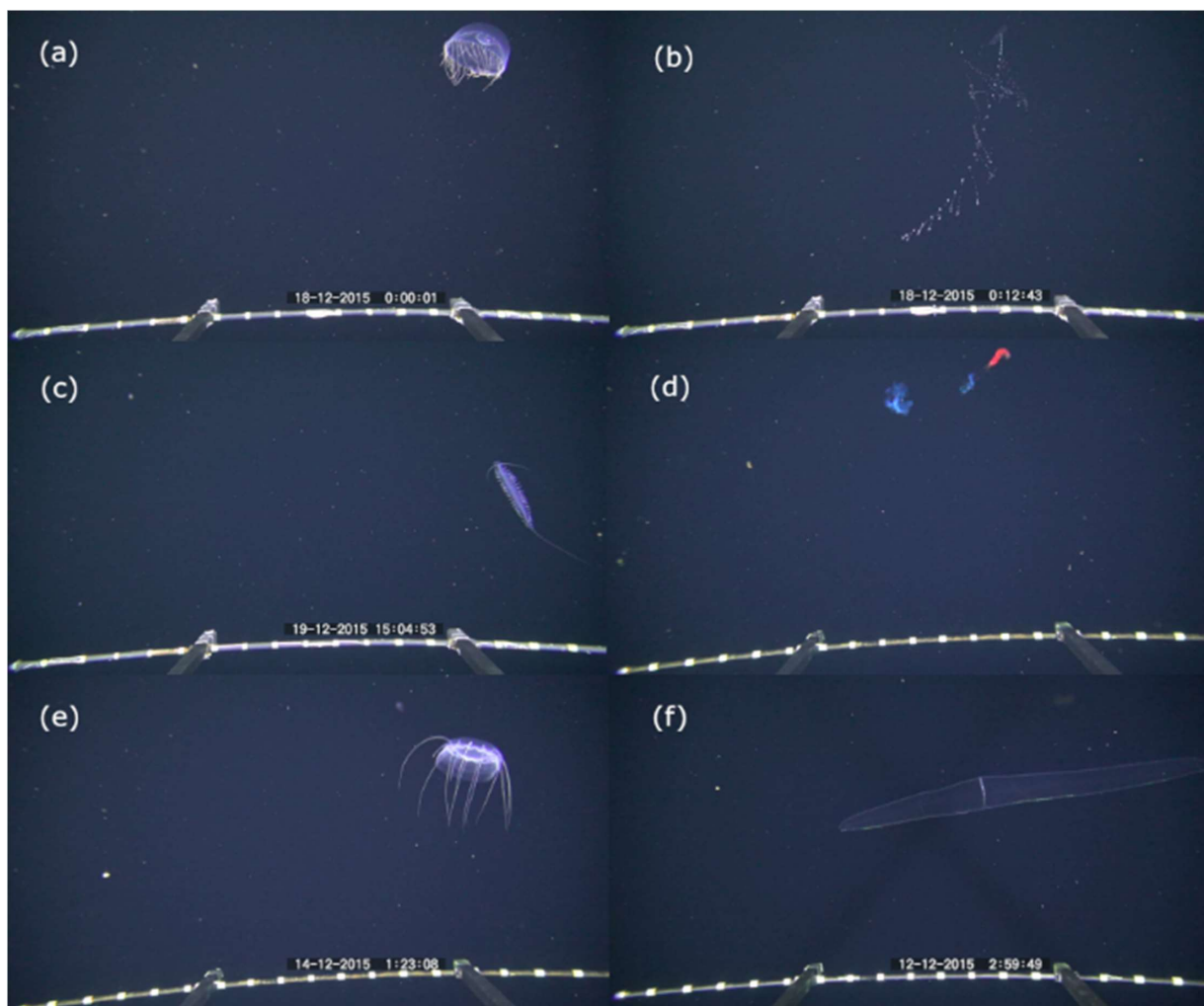
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515 Figure 4: Day and night comparison of faunal observations obtained by PELAGIOS at the North
 516 West flank of Senghor seamount A: fishes, krill, chaetognaths and decapods B: gelatinous
 517 zooplankton groups

518



519

520 Figure 5: example of organisms encountered during pelagic video transects with PELAGIOS
521 during cruise MSM49 in the eastern tropical Atlantic. (a) a medusa *Halitrephes* sp. (b) a
522 calycophoran siphonophore in feeding position (c) a tomopterid worm (d) a crustacean releasing a
523 bioluminescent cloud (e) the medusa *Solmissus* (f) the ctenophore *Cestum*

524



525 Table 1: Taxonomic groups which were encountered during pelagic video transects in the eastern
526 tropical Atlantic.

| Phylum | Class | Order | Family | Genus | | |
|------------|------------------|-------------------|-----------------------|--|---|--|
| Cercozoa | Thecofilosea | | | | | |
| Radiozoa | | | | | | |
| Cnidaria | Hydrozoa | Narcomedusae | Solmundaeginidae | <i>Solmundella</i> | | |
| | | | Aeginidae | <i>Aegina</i> <i>Aeginura</i> | | |
| | | | Cuninidae | <i>Solmissus</i> | | |
| | | Trachymedusae | Halicreatidae | <i>Halicreas</i> <i>Haliscera</i> <i>Halitrephes</i> | | |
| | | | Rhopalonematidae | <i>Colobonema</i> <i>Crossota</i> <i>Rhopalonema</i> | | |
| | | | Geryoniidae | <i>Geryonia</i> <i>Liriope</i> | | |
| | | | Siphonophorae | Agalmatidae | <i>Halistemma</i> <i>Marrus</i> <i>Nanomia</i> <i>Apolemia</i> | |
| | | | | Apolemiidae | | |
| | | | | Diphyidae | | |
| | | Forskaliidae | | <i>Forskalia</i> | | |
| | | Hippopodiidae | | <i>Hippopodius</i> <i>Vogtia</i> | | |
| | | | | | Physophoridae | <i>Physophora</i> |
| | | | | | Prayidae | <i>Craseoa</i> <i>Lilyopsis</i> <i>Praya</i> <i>Rosacea</i> |
| | | | | | Pyrostephidae | <i>Bargmannia</i> |
| | | | | | Resomiidae | <i>Resomia</i> |
| | Scyphozoa | Coronatae | Atollidae | <i>Atolla</i> | | |
| | | | Nausithoidae | <i>Nausithoe</i> | | |
| | | | Peryphyllidae | <i>Periphylla</i> | | |
| Ctenophora | Nuda | Beroida | Beroidae | <i>Beroe</i> | | |
| | Tentaculata | Cestida | Cestidae | <i>Cestum</i> <i>Velamen</i> | | |
| | | Cydippida | Aulacoctenidae | <i>Aulacoctena</i> | | |
| | | | Pleurobrachiidae | <i>Hormiphora</i> | | |
| | | Lobata | Bathocyroidae | <i>Bathocyroe</i> | | |
| | | | Eurhamphaeidae | <i>Kiyohimea</i> | | |
| | | | Leucotheidae | <i>Leucothea</i> | | |
| | Ocryopsidae | <i>Ocryopsis</i> | | | | |
| | Thalassocalycida | Thalassocalycidae | <i>Thalassocalyce</i> | | | |



| | | | | |
|--------------|----------------|--|--|---|
| Chaetognatha | Sagittoidea | | | |
| Annelida | Polychaeta | Phyllodocida Canalipalpata | Tomopteridae Flabelligeridae | <i>Tomopteris</i> <i>Poebius</i> |
| Arthropoda | Malacostraca | Amphipoda Decapoda Euphausiacea Isopoda | Munnopsidae | <i>Munnopsis</i> |
| Mollusca | Cephalopoda | Octopoda Teuthida | Amphitretidae Octopodidae Cranchiidae Mastigoteuthidae Octopoteuthidae Ommastrephidae | <i>Bolitaena</i> <i>Helicocranchia</i> <i>Mastigoteuthis</i> <i>Octopoteuthis</i> <i>Taningia</i> <i>Sthenoteuthis</i> |
| | Gastropoda | Nudibranchia Pteropoda | Phylliroidae | <i>Phylliroe</i> |
| Chordata | Appendicularia | Copelata | Oikopleuridae | <i>Bathochordaeus</i> <i>Mesochordaeus</i> |
| | Thaliacea | Doliolida Pyrosomatida Salpida | Pyrosomatidae Salpidae | <i>Pyrostemma</i> <i>Cyclosalpa</i> |
| | Actinopteri | Anguilliformes Myctophiformes Stomiiformes | Nemichthyidae Myctophidae Gonostomatidae Sternoptychidae | <i>Cyclothone</i> |

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