



The Pelagic In situ Observation System (PELAGIOS) to reveal biodiversity, behavior, and ecology of elusive oceanic fauna

Henk-Jan Hoving¹, Svenja Christiansen², Eduard Fabrizio¹, Helena Hauss¹, Rainer Kiko¹, Peter Linke¹, Philipp Neitzel¹, Uwe Piatkowski¹, and Arne Körtzinger^{1,3}

¹GEOMAR, Helmholtz Centre for Ocean Research Kiel, Düsternbrooker Weg 20, 24105 Kiel, Germany

²^{TS1}University of Oslo, Blindernveien 31, 0371 Oslo, Norway

³^{TS2}Christian Albrecht University Kiel, Christian-Albrechts-Platz 4, 24118 Kiel, Germany

Correspondence: Henk-Jan Hoving (hhoving@geomar.de)

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Abstract. ^{CE1 CE2 TS3} There is a need for cost-efficient tools to explore deep-ocean ecosystems to collect baseline biological observations on pelagic fauna (zooplankton and nekton) and establish the vertical ecological zonation in the deep sea. The Pelagic In situ Observation System (PELAGIOS) is a 3000 m rated slowly (0.5 m s^{-1} ^{TS4}) towed camera system with LED illumination, an integrated oceanographic sensor set (CTD ^{CE3}-O₂) and telemetry allowing for online data acquisition and video inspection (low definition). The high-definition video is stored on the camera and later annotated using annotation software and related to concomitantly recorded environmental data. The PELAGIOS is particularly suitable for open-ocean observations of gelatinous fauna, which is notoriously under-sampled by nets and/or destroyed by fixatives. In addition to counts, diversity, and distribution data as a function of depth and environmental conditions (T, S, O₂), in situ observations of behavior, orientation, and species interactions are collected. Here, we present an overview of the technical setup of the PELAGIOS as well as example observations and analyses from the eastern tropical North Atlantic. Comparisons to MOCNESS ^{CE4} net sampling and data from the Underwater Vision Profiler are provided and discussed.

oceanographic expeditions, oceanic communities of macrozooplankton and micronekton have been sampled using nets (Wiebe and Benfield, 2003). Such sampling has revealed a community typically consisting of crustaceans, cephalopods, fishes, and some sturdy and commonly found gelatinous fauna (Benfield et al., 1996). Underwater observations in the open ocean via scuba diving (Hamner et al., 1975) and later via submersibles (Robison, 1983; Robison and Wishner, 1990) and in situ camera systems (Biard et al., 2016; Picheral et al., 2010) revealed that a variety of organisms are much more abundant in the open ocean than previously estimated from net sampling (Robison, 2004). This was particularly true for fragile gelatinous zooplankton, a diverse taxonomic group of different phyla, including the ctenophores and medusae (Remsen et al., 2004; Haddock, 2004) as well as polychaetes (Christiansen et al., 2018), Rhizaria (Biard et al., 2016), and pelagic tunicates (Remsen et al., 2004; Neitzel, 2017), which often are too delicate to be quantified using nets as they are damaged beyond identification, or they are easily destroyed by the use of common fixatives.

Underwater (in situ) observations in the pelagic ocean not only revealed a previously unknown community, they also allowed the collection of fine-scale distribution patterns of plankton in relation to biotic and abiotic factors (e.g., Haslob et al., 2009; Möller et al., 2013; Hauss et al., 2016) as well as information on posture, interactions, and behavior (Hamner and Robison, 1992; Robison, 2004; Robison, 1999; Hoving et al., 2017). Submersibles have proven to be valuable instruments to study deep-sea pelagic biology (e.g., Robison, 1987 ^{TSS}; Bush et al., 2007; Hoving et al., 2013; 2016). Us-

1 Introduction

The open-ocean pelagic zones include the largest, yet least explored habitats on the planet (Robison, 2004; Webb et al., 2010; Ramirez-Llodra et al., 2010). Since the first

ing video transecting methodology, pelagic ROV ^{CE5} surveys have been applied to study inter- and intra-annual variation in mesopelagic zooplankton communities (Robison et al., 1998; Hull et al., 2011) and to explore deep pelagic communities in 5 different oceans (Youngbluth et al., 2008; Hosia et al., 2017; Robison et al., 2010). However, due to high costs as well as technological and logistical challenges, regular submersible operations are still restricted to very few institutes and geographical locations. Hence, there is a need for the develop- 10 ment of additional more cost-effective methodologies to explore and document deep-sea communities via in situ observations.

In the last decades, a variety of optical instruments has been developed to image and quantify plankton in situ (Ben- 15 field et al., 2007). The factors that typically differentiate the available plankton imaging technologies are the size fraction of the observed organisms, illumination type, resolution of collected images or video, depth rating, deployment mode (e.g., autonomous, towed, CTD-mounted), and towing speed. 20 Examples of instruments include the autonomous Underwater Vision Profiler (UVP5; Picheral et al., 2010), the Light-frame On-sight Key species Investigations (LOKI; Schulz et al., 2010) and towed plankton recorders (ISiS ^{CE6}; Cowen and Guigand, 2008; for a review, see Benfield et al., 2007). 25 These instruments can be deployed from ships of opportunity and collect detailed information on fine-scale distribution and diversity patterns of particles and plankton. The data reveal biological patterns on a global scale (Kiko et al., 2017) and of previously underappreciated plankton species 30 (Biard et al., 2016). More recently, optical (and acoustic) instruments have been combined with autonomous gliders and rapidly increasing spatial resolution (Ohman et al., 2019).

Various towed camera platforms have been developed that can obtain video transect observations above the deep sea 35 floor. Examples are the TowCam (WHOI), the DTIS (Deep Towed Imaging system, NIWA), the WASP vehicle (Wide Angle Seafloor Photography), OFOS (Ocean Floor Observation System, GEOMAR), and the more recent version OFOBS (Ocean Floor Observation and Bathymetry System; 40 Purser et al., 2018). All these instruments are used for video or photo transects of the seafloor, with a downward looking camera and typically a set of lasers for size reference. However, published descriptions of optical systems, other than ROVs and submersibles that visualize macrozooplankton and 45 micronekton (> 1 cm) in the water column undisturbed by a filtering device or cuvette are, to the best of our knowledge, restricted to one (Madin et al., 2006). The Large Area Plankton Imaging System (LAPIS) is the only towed system that was developed for the documentation of larger organisms 50 in the water column (Madin et al., 2006). LAPIS visualizes organisms between 1 and 100 cm, it combines a high-resolution color digital CCD ^{CE7} camera using progressive scanning interline-transfer technology with flashing strobes, and it is towed at 1 knot via a fibre optic wire. LAPIS col- 55 lects still images, illumination is sideways, and organisms

have to enter an illuminated volume to be visualized. Deploy- 60 ments in the Southern Ocean enabled the reconstruction of depth distributions of the pelagic fauna (salps, medusae) but also allowed some behavior observations, e.g., the molting of krill (Madin et al., 2006). More publications of data collected with LAPIS are unavailable to our knowledge. In addition 65 to LAPIS ^{CE8}, we wanted to develop a towed pelagic observation system that collects video during horizontal transects (with forward projected light), in a similar way to pelagic ROV video transects, in order to document behavior in addition to diversity, species-specific distribution, and abundance 70 data of pelagic fauna.

The functional requirements for the instrument were (1) the ability to visualize organisms > 1 cm in waters down to 1000 m with high-definition video, (2) the possibility of de- 75 ploying the instrument from ships of opportunity in an autonomous or transmitting mode, (3) for it to be lightweight and practical so it can be deployed easily and safely with two deck persons and a winch operator, (4) to enable the correlation of observations with environmental parameters (S, T, O₂) and other sensor data, and (5) to make observations com- 80 parable to ROV video transects in other reference areas. We present a description of the Pelagic In situ Observation System (PELAGIOS), examples of the kind of biological information it may gather, as well as biological discoveries that 85 have resulted from deployments on research cruises in the eastern tropical North Atlantic.

2 Pelagic In Situ Observation System

2.1 Technical specifications

The PELAGIOS consists of an aluminum frame (length: 2 m) 85 that carries the oceanographic equipment (Fig. 1). White light LED arrays (^{CE9}four LEDs produced at GEOMAR, two LED arrays (type LightSphere of Deep-Sea Power and Light ©), which illuminate the water in front of the system, are mounted on an aluminum ring (diameter: 1.2 m). Power 90 is provided by two lithium batteries (24 V; 32 Ah) in deep-sea housing. High-definition video is collected continuously by a forward-viewing deep-sea camera (type 1Cam Alpha, SubC Imaging ©), which is mounted in the center of the ring. We used the maximum frame rate of 50 frames s⁻¹ 95 but a lower frame rate is possible. A CTD (SBE 19 Sea-CAT, Sea-Bird Scientific ©) with an oxygen sensor (SBE 43, Sea-Bird Scientific ©) records environmental data. A deep-sea telemetry (DST-6, Sea and Sun Technology ©; Linke et al., 2015) transmits video and CTD data to a deck unit on 100 board allowing a low-resolution preview (600 × 480 lines) of the high-definition video that is stored locally on the SD card (256 GB) of the camera. The power from the batteries is distributed to the LEDs via the camera. The 1Cam Alpha camera is programmable in such a way that there is a delay 105 between providing power to the camera (by connecting to

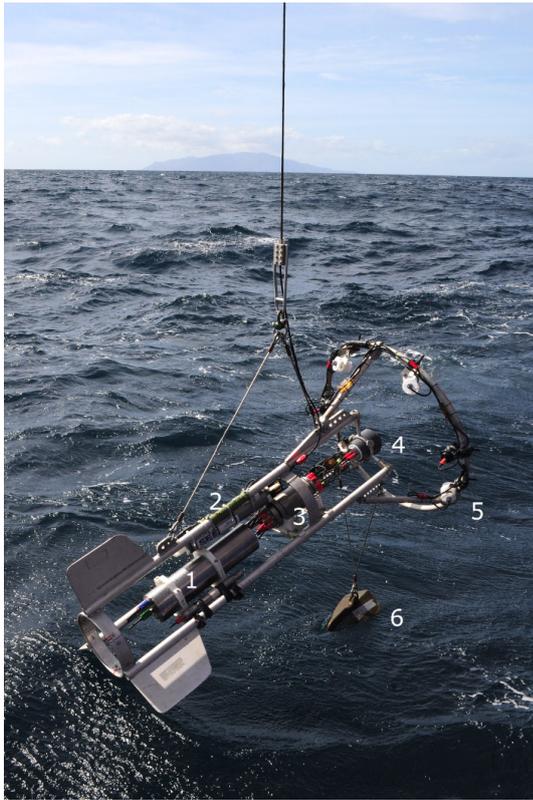


Figure 1. (a) The Pelagic In Situ Observations System (PELAGIOS) with battery (1), CTD (2), telemetry (3), camera (4), LEDs (5), and depressor (6) during deployment from R/V *Poseidon* in February 2018 (photo: Karen Hissmann).

the battery) and the start of recording and switching on the LEDs. This enables the illumination to be turned on only underwater, and prevents overheating of the LED arrays while out of the water. During a cruise with the German research vessel *Maria S. Merian* (MSM 49) we mounted a steel scale bar in front of the camera at a distance of 1 m. The distance between the centers of the white marks on the bar measured 5 cm.

2.2 Video transects

The PELAGIOS is towed horizontally at specified depths of 20–1000 m. The standard towing speed over ground is 1 knot (0.51 m s^{-1}), and the speed is monitored via the ship's navigational system. A video transect at a particular depth can take as long as desired and is terminated by lowering the PELAGIOS to the next desired depth. Maximum deployment time with full batteries is approximately 6 h. The typical transect duration is 10–30 min. The depth of the PELAGIOS can be monitored via online CTD data. Figure 2 shows the trajectories of the PELAGIOS at different depths in the water column during a video transect down to 700 m. The deployment from deck into the water and the reverse is fast and typically

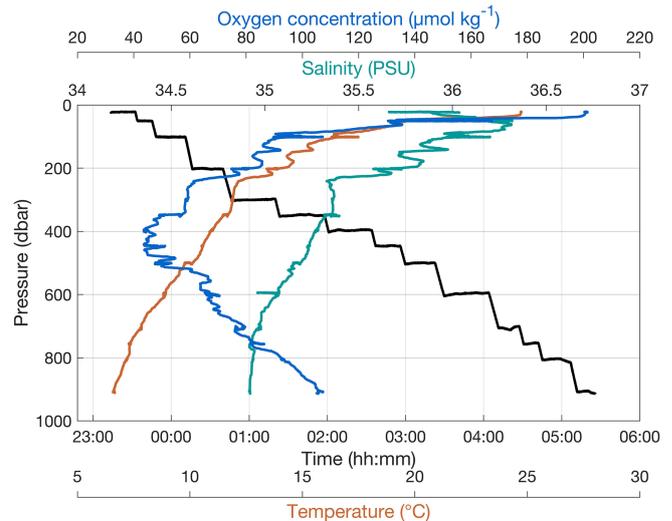


Figure 2. Stairwise trajectory of PELAGIOS through the water column, to the desired depths with concomitantly measured environmental data.

takes only about 5 min (see video clip [CE10](#) in the ESM [CE11](#)). It is possible to deploy PELAGIOS in “blind mode”, where only the depth is monitored using an online depth sensor (e.g., Hydrobios ©) and the video (without transmitted preview) is recorded locally on the camera. The system can be operated completely blind (i.e., with no communication between deck and underwater unit) where the target depth is estimated from the length and angle of the wire put out, and the actual depth is recorded on the system by CTD or an off-line pressure sensor, e.g., SBE Microcat ©.

2.3 Video analysis and curation

After a deployment, the video (consisting of individual clips of 1 h) is downloaded from the camera. Synchronization between video and CTD data is done by setting all instruments to UTC prior to deployment, which allows the data and video to be linked during analysis. The video is annotated using the Video Annotation and Reference System (VARS) developed at the Monterey Bay Aquarium Research Institute (Schlinding and Jacobsen, 2006). This annotation program allows for frame grabs from the video, including time code. A knowledge base allows for the insertion of taxonomic names and hierarchy, and a query allows searching the created database. While many kinds of annotation software are available (for a review, see Gomes-Pereira et al., 2016), we consider VARS to be the most suitable for our purposes since it combines the features of high-resolution video playback with a user-friendly annotation interface and the automatic creation of an annotation database which can easily be accessed through the various search functions and tools of the query. The taxonomic hierarchy and phylogenetic trees in the database are directly applicable to our video transects. Since this soft-

ware was developed by MBARI (CBI2), which also maintains the most extensive databases of deep pelagic observations, it makes communication about and comparison of observations and data practical. Videos are transported on hard drives after an expedition and are transferred for long-term storage on servers maintained by the central data and computing center at GEOMAR, providing instant access to videos and images with metadata description via the media server ProxSys.

2.4 Sample volume

To estimate the sample volume of the PELAGIOS we compared video counts from the PELAGIOS with concomitantly obtained abundance data from an Underwater Vision Profiler (UVP5; Picheral et al., 2010). Four deployments from the R/V *Maria S. Merian* cruise MSM 49 (28 November–21 December 2015, Las Palmas de Gran Canaria, Spain – Mindelo, Cabo Verde) were used for the comparison where a UVP5 was mounted underneath the PELAGIOS. The UVP5 takes between 6 and 11 images per second of a defined volume (1.03 L) and thus enables a quantitative assessment of particle and zooplankton abundances. Objects with an equivalent spherical diameter (ESD) > 0.5 mm are saved as images, which can be classified into different zooplankton, phytoplankton, and particle categories. For the comparison between PELAGIOS and the UVP5, we used the pelagic polychaete *Poebobius* sp., as (1) this organism could be observed well on both instruments, (2) *Poebobius* sp. is not an active swimmer and lacks an escape response and (3) it was locally very abundant, thus providing a good basis for the direct instrument comparison.

The UVP5 images were classified as described in Christiansen et al. (2018). *Poebobius* sp. abundance (ind. (individuals) m⁻³) was calculated for 20 s time bins and all bins of one distinct depth step (with durations of 10–11 min at depths <= 50 m, 19–22 min at depths < 350 m, and 9–11 min at depths >= 350 m) averaged. These mean abundances were compared to the PELAGIOS counts (ind. s⁻¹) of the same depth step. A linear model between the PELAGIOS counts as a function of UVP5 abundance provided a highly significant relationship (linear regression: $p < 0.001$; adjusted $r^2 = 0.69$; Fig. 3). The linear regression slope b (0.116 m³ s⁻¹; standard error 0.01 m³ s⁻¹) between the PELAGIOS-based count (C_{PELAGIOS} , ind. s⁻¹) and mean UVP-based abundance (A_{UVP} , ind. m⁻³),

$$C_{\text{PELAGIOS}} = b \cdot A_{\text{UVP}} + a, \quad (1)$$

was used to estimate the volume recorded per time in cubic meters per second (b) and the field of view in square meters ($b/\text{towing speed}$) recorded by PELAGIOS.

From this calculation it can be derived that PELAGIOS recorded an average volume of 0.116 m³ s⁻¹ at a towing speed of 1 knot (= 0.51 m s⁻¹). A cross-sectional view field of approximately 0.23 m² of PELAGIOS can be expected, compared to a theoretical field of view (FOV)

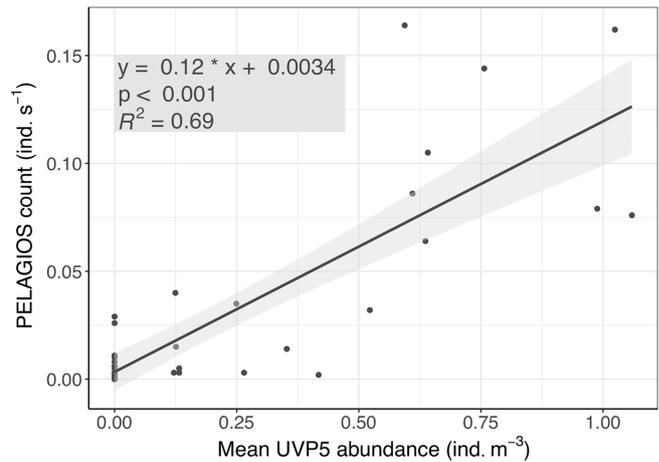


Figure 3. PELAGIOS video counts of *Poebobius* sp. as a function of UVP5-derived abundance on the same transects at two stations on cruise MSM 49 on R/V *Maria S. Merian*.

of 0.45 m² based upon the maximum image dimensions (0.80 m × 0.56 m) at 1 m distance from the lens. We can now calculate the individuals observed by PELAGIOS per time to individuals per volume. To do so we use the number of individuals in one transect and divide this number by the duration of the transect to obtain individuals per minute, and we divide this by 60 to get the individuals per second. From the UVP–PELAGIOS comparison we derived a conversion factor of 6 to calculate the number of individuals per second to the number of individuals per cubic meter. This value is then multiplied by the conversion factor 6 and again multiplied by 1000 to go from cubic meter to 1000 m³.

2.5 Abundance, size, and diversity at an example station, “Senghor NW”

To provide an example of the type of data that can be obtained with the PELAGIOS, we report here on day and night video transects down to 950 m in the eastern tropical North Atlantic, on the northwestern slope of Senghor Seamount (17° 14.2' N, 22° 00.7' W; bottom depth of approximately 1000 m). The results from the video annotations show that faunal abundances depend on the depth of deployment, and time of the day. During two transects of 11 min at 400 m, 226 individuals (1066 ind. 1000 m⁻³) (TS7) were encountered during the day (the three dominant organism groups were fish, euphausiids, and appendicularians) compared to 196 individuals (591 ind. 1000 m⁻³) during the night (the four dominant organism groups are fish, chaetognaths, medusae, and ctenophores). Overall abundance of chaetognaths, decapods, and mysids, and fishes was higher during the night. The peak of euphausiids' abundance at 400 m shifts to the surface at night (Fig. 4). The higher abundance of decapods, mysids, and chaetognaths at night may indicate lateral migration or daytime avoidance. The vertical migration that was

observed for fishes and crustaceans was much less clear for the gelatinous zooplankton groups including medusae and appendicularians (Fig. 4). Ctenophores and siphonophores were abundant in the surface at night (but we did not perform transects at 20 and 50 m during the day), and the thaliaceans migrated vertically and were most abundant in shallow waters at night. The total number of annotated organisms for the daytime transects (total transect time 187 min; max depth 950 m) was 835 compared to 1865 organisms for the longer nighttime transects (total transect time 292 min; max depth 900). The enormous abundance of gelatinous zooplankton (128) annotated organisms (899 ind. 1000 m^{-3}) belonging to the three dominant groups of Ctenophora (53), Siphonophorae (21), and Thaliacea (44) in the topmost layer (20 m) at night is remarkable. Below this layer, the depth profile shows a minimum in numbers of annotated individuals at 100, 200, and 300 m water depth with a smaller peak of 57 gelatinous organisms (299 ind. 1000 m^{-3}) in 450 m. Compared to this, the depth distribution during the daytime shows a more regular, almost Gaussian shape with a maximum of 31 (254 ind. 1000 m^{-3}) and 54 (254 ind. 1000 m^{-3}) gelatinous organisms at 200 and 400 m water depth, respectively.

The faunal observations at station Senghor NW include a wide variety of taxa (Table 1; Figs. 5 and 6), spanning in size from radiolarians to large siphonophores (such as *Praya dubia* and *Apolemia*). Chaetognaths were the dominant faunal group. Typical examples of fragile organisms that were not present or identifiable in the MOCNESS samples from the same cruise (Christiansen et al., 2016; Lüsrow et al., 2019^{TS9}) but which can be efficiently observed by PELAGIOS include large larvaceans (probably *Bathochordaeus* and *Mesochordaeus*), pelagic polychaetes (*Poebobius*, *Tomopteris*) (Fig. 5), and smaller siphonophores (such as *Bargmannia* and *Lilyopsis*^{CE14}; the latter can be easily distinguished by their fluorescent body parts) (Fig. 5). Observed medusae belonged to the genera *Periphylla*, *Halitrephes*, *Haliscera*, *Crossota*, *Colobonaema*^{CE15}, *Solmissus*, and *Solmundella* (Fig. 5). Venus girdles (*Cestum* spp.), *Beroe*, cypripids, and lobate ctenophores (such as *Thalassocalyce inconstans*, *Leucothea*, *Bathyceroe*^{CE16}; see Harbison et al., 1978, for differences in robustness among ctenophores) were encountered at Senghor NW (Fig. 5). Cephalopod observations were rare but small individual cranchiid squids were observed in the upper 50 m at night. Mastigoteuthid squids were observed with their mantle in a vertical orientation and with extended tentacles in waters below 500 m. One large squid, *Taningia danae* was observed during a transit between transecting depths. Other pelagic molluscs include the nudibranch *Phylliroe* and different pteropod species. Observed fishes are snipe eels, hatchet fishes, lantern fishes, and *Cyclothone*. Fishes are among the dominant organisms encountered during PELAGIOS transects, but it is often impossible to identify fishes to species level from the video.

2.6 Individual behavior

In situ observations by PELAGIOS video may reveal direct observations on individual behavior. Decapod shrimps were observed to release a blue or green bioluminescent cloud after performing their tail flip as part of the escape response (Fig. 6d). Potential reproductive behavior was observed for two specimens of krill which were seen in what could be a mating position, and salps were observed to reproduce asexually by the release of salp oozoids (Fig. 6c). Feeding behaviors were observed for large prayid siphonophores and calycophoran siphonophores which had their tentacles extended. *Poebobius* worms were observed with their mucus web deployed to capture particulate matter (Christiansen et al., 2018) (Fig. 6a). Narcomedusae of the genus *Solmissus* were observed with their tentacles stretched up and down, which is a feeding posture (Fig. 5). In situ observations by the PELAGIOS also showed the natural body position of pelagic organisms. Snipe eels were observed in a vertical position with their heads up, while dragonfish and some myctophids were observed in an oblique body position with their head down (Fig. 6b).

3 Discussion

PELAGIOS is a pelagic ocean exploration tool that fills a gap in the array of observation instruments that exist in biological oceanography, as transparent and fragile organisms (> 1 cm) are up to now under-sampled by both net-based and optical systems. The PELAGIOS video transects are comparable to ROV video transects and can be obtained in a cost-effective way. The resulting data can provide information on diversity, distribution and abundance of large (> 1 cm), fragile zooplankton and some nekton and also of rare species. Due to the collection of HD color video, the behavior, color, and position of larger gelatinous planktonic organisms in the water column are documented, which may provide additional ecological information that cannot be obtained by nets or existing plankton recorders. The PELAGIOS system complements gear that is suitable for stratified observations and collections of robust mesozooplankton and micronekton (MOCNESS, Hydrobios Multinet, and others) and optical systems that are suitable for high-resolution sampling of small and abundant organisms (e.g., VPR, UVP5) (e.g., Benfield et al., 2007; Picheral et al., 2010; Biard et al., 2015^{TS11}). The instrument can be deployed with a small team and from vessels of opportunity, in transmission or blind mode. Due to the relatively simple design we experienced limited technical failures, which makes the PELAGIOS a reliable tool for oceanic expeditions. While thus far the system has only been deployed in the open ocean, it can be used in any pelagic environment with water that has reasonable clearance and visibility. The data obtained after annotation of the video can be uploaded into databases (e.g., the large database PAN-

Table 1. Taxonomic groups which were encountered during pelagic video transects in the eastern tropical Atlantic. [TSS](#)

Phylum	Class	Order	Family	Genus
Cercozoa	Thecofilosea			
Radiozoa				
Cnidaria	Hydrozoa	Narcomedusae	Solmundaeginidae Aeginidae	<i>Solmundella</i> <i>Aegina</i> <i>Aeginura</i>
		Trachymedusae	Cuninidae Haliceatidae	<i>Solmissus</i> <i>Haliceas</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 Hippopodiidae	<i>Forskalia</i> <i>Hippopodius</i> <i>Vogtia</i>
			Physophoridae Prayidae	<i>Physophora</i> <i>Craseoa</i> <i>Lilyopsis</i> <i>Praya</i> <i>Rosacea</i>
			Pyrostephidae Resomiidae	<i>Bargmannia</i> <i>Resomia</i>
	Scyphozoa	Coronatae	Atollidae Nausithoidae Peryphyllidae	<i>Atolla</i> <i>Nausithoe</i> <i>Periphylla</i>
Ctenophora	Nuda	Beroida	Beroidae	<i>Beroe</i>
	Tentaculata	Cestida	Cestidae	<i>Cestum</i> <i>Velamen</i>
		Cydippida	Aulacoctenidae Pleurobrachiidae	<i>Aulacoctena</i> <i>Hormiphora</i>
		Lobata	Bathocyroidae Eurhamphaeidae Leucotheidae Ocryopsidae	<i>Bathocyroe</i> <i>Kiyohimea</i> <i>Leucothea</i> <i>Ocryopsis</i>
		Thalassocalycida	Thalassocalycidae	<i>Thalassocalyce</i>
Chaetognatha	Sagittoidea			
Annelida	Polychaeta	Phyllodocida Canalipalpata	Tomopteridae Flabelligeridae	<i>Tomopteris</i> <i>Poeobius</i>
Arthropoda	Malacostraca	Amphipoda Decapoda Euphausiacea		

Table 1. Continued.

Phylum	Class	Order	Family	Genus
		Isopoda	Munnopsidae	<i>Munnopsis</i>
Mollusca	Cephalopoda	Octopoda	Amphitretidae Octopodidae	
		Teuthida	Cranchiidae	<i>Helicocranchia</i>
			Mastigoteuthidae	<i>Mastigoteuthis</i>
			Octopoteuthidae	<i>Octopoteuthis</i> <i>Taningia</i> <i>Sthenoteuthis</i>
			Ommastrephidae	
	Gastropoda	Nudibranchia Pteropoda	Phylliroidae	<i>Phylliroe</i>
Chordata	Appendicularia	Copelata	Oikopleuridae	<i>Bathochordaeus</i> <i>Mesochor- daeus</i>
	Thaliacea	Doliolida Pyrosomatida ^{CE13} Salpida	Pyrosomatidae Salpidae	<i>Pyrostemma</i> <i>Cyclosalpa</i>
	Actinopteri	Anguilliformes Myctophiformes Stomiiformes	Nemichthyidae Myctophidae Gonostomatidae Sternoptychidae	<i>Cyclothone</i>

GAEA) after publication of the results allowing for efficient data sharing and curation.

The clear distribution patterns that we observed in some animal groups (fish, crustaceans, and some gelatinous fauna) after annotating the video transects confirm that established biological processes such as diurnal vertical migration (e.g., Barham, 1963) can be detected in PELAGIOS data and that the distribution data that we observe for the organisms encountered are representative of the natural situation. It has to be noted, though, that while the observed distribution patterns should be representative, care must be taken with regard to abundance estimates of especially actively and fast-swimming organisms. Some fish and crustaceans react to the presence of underwater instrumentation (e.g., Stoner et al., 2008 ^{TS12}). Gear avoidance (e.g., Kaartvedt et al., 2012 ^{TS13}) can lead to an underestimation of abundance, whereas attraction to the camera lights (e.g., Utne-Palm et al., 2018 ^{TS14}; Wiebe et al., 2004 ^{TS15}) would result in an overestimation. The large bioluminescent squid *Taningia danae* seemed to be attracted to the lights of the PELAGIOS, and attraction behavior of this species has been described in other publications (Kubodera et al., 2007). Compared to day transects, the high abundance of gelatinous organisms close to the surface during night is likely to be partly an effect of the higher contrast in the videos of the night transects and better visibility of the gelatinous fauna than during day transects. Therefore, we did not perform transects shallower than 50 m dur-

ing the day. Many of the observed gelatinous fauna might be present as well at shallow depths during daylight but are not detectable in “blue-water conditions”. The difference between the taxa encountered during the day and night transect may also be due to the trapping of organisms at the slopes of Senghor Seamount during the day (Isaacs and Schwartzlose, 1965; Genin, 2004) or by other causes for patchiness (Haury et al., 2000). However, from a methodological point of view it should be noted that while the ship’s towing speed is typically 1 knot, the current speeds at the survey depths may differ, also between day and night. Currents may result in a more or less sampled volume of water and hence a variation in plankton being visualized. Since abundance estimation relies on an accurate determination of the image volume, it needs to be pointed out that it is our aim to better technically constrain the image area in future developments (now derived from UVP quantitative observations) and to include flowmeter measurements.

After annotation, the PELAGIOS video transects may be used to reconstruct species-specific distribution patterns, which can be related to environmental conditions (Neitzel, 2017; Hoving et al., 2019 ^{TS16 TS17}). Such data are also valuable for overlap comparison in distribution patterns of consumers and food items (see, e.g., Haslob et al., 2009; Möller et al., 2012). The data can also be used in biological studies that aim to predict the consequences of a changing ocean with altering oceanographic features and conditions

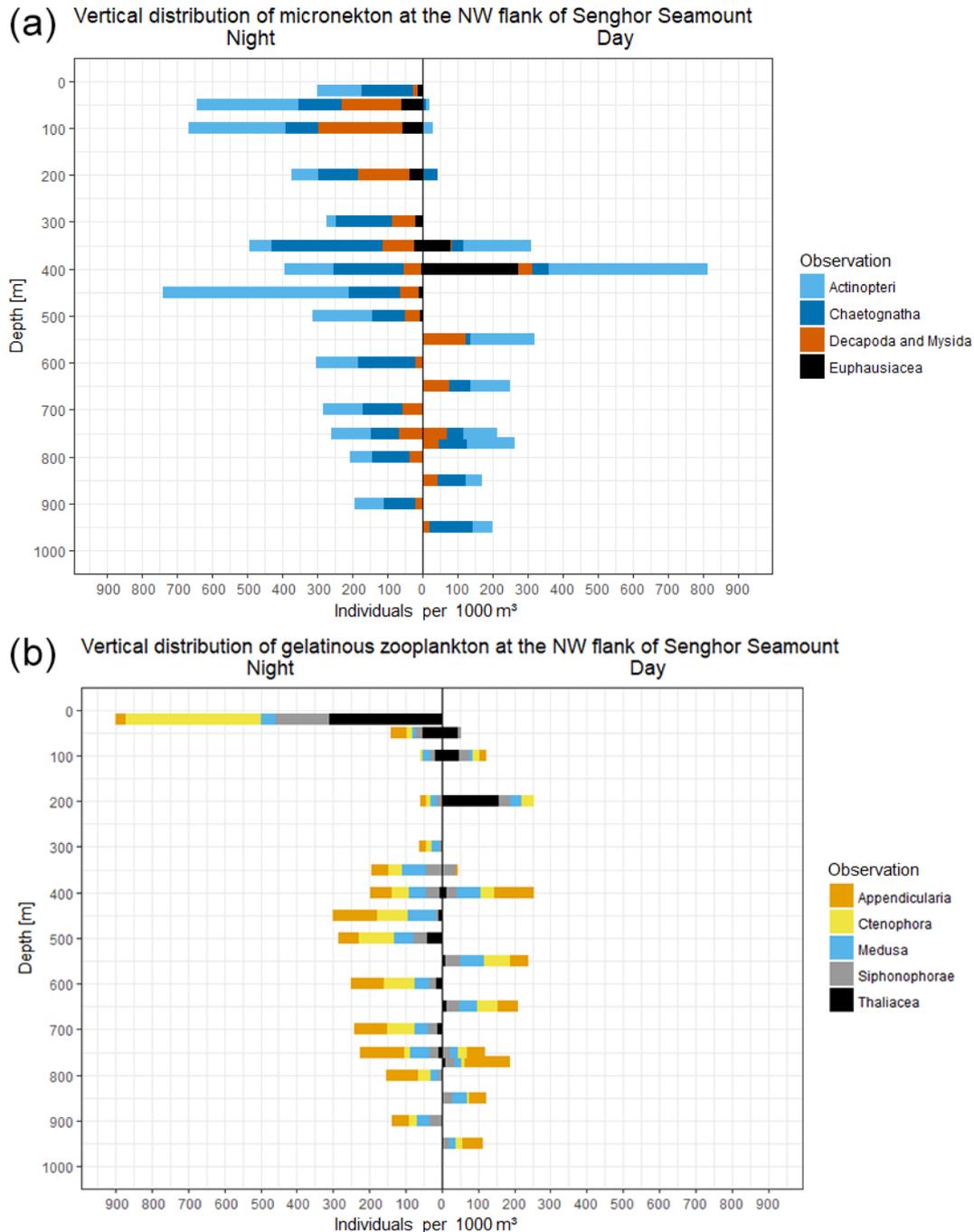


Figure 4. Day and night comparison of faunal observations obtained by PELAGIOS at the northwest flank of Senghor Seamount: (a) [TS10](#) fishes, krill, chaetognaths, and decapods; (b) gelatinous zooplankton groups.

for species' distributions, as has been done for net sampling of mesozooplankton (Wishner et al., 2013). One example of changing oceanographic conditions is the global trend of oxygen loss in the world oceans (Oschlies et al., 2018). Oxygen minimum zones (OMZs) occur naturally in the mesopelagic zone (Robinson et al., 2010), and in different oceans they have been found to expand horizontally

and vertically as a result of climate change (Stramma et al., 2008; [TS18](#); Oschlies et al., 2018). The expansion of OMZs may result in a habitat reduction of the pelagic fauna (e.g., Stramma et al., 2012) or increase the habitat for species with hypoxia tolerance (Gilly et al., 2013). To predict the potential consequences of OMZ expansion for pelagic invertebrates, we investigated the abundance and distribution

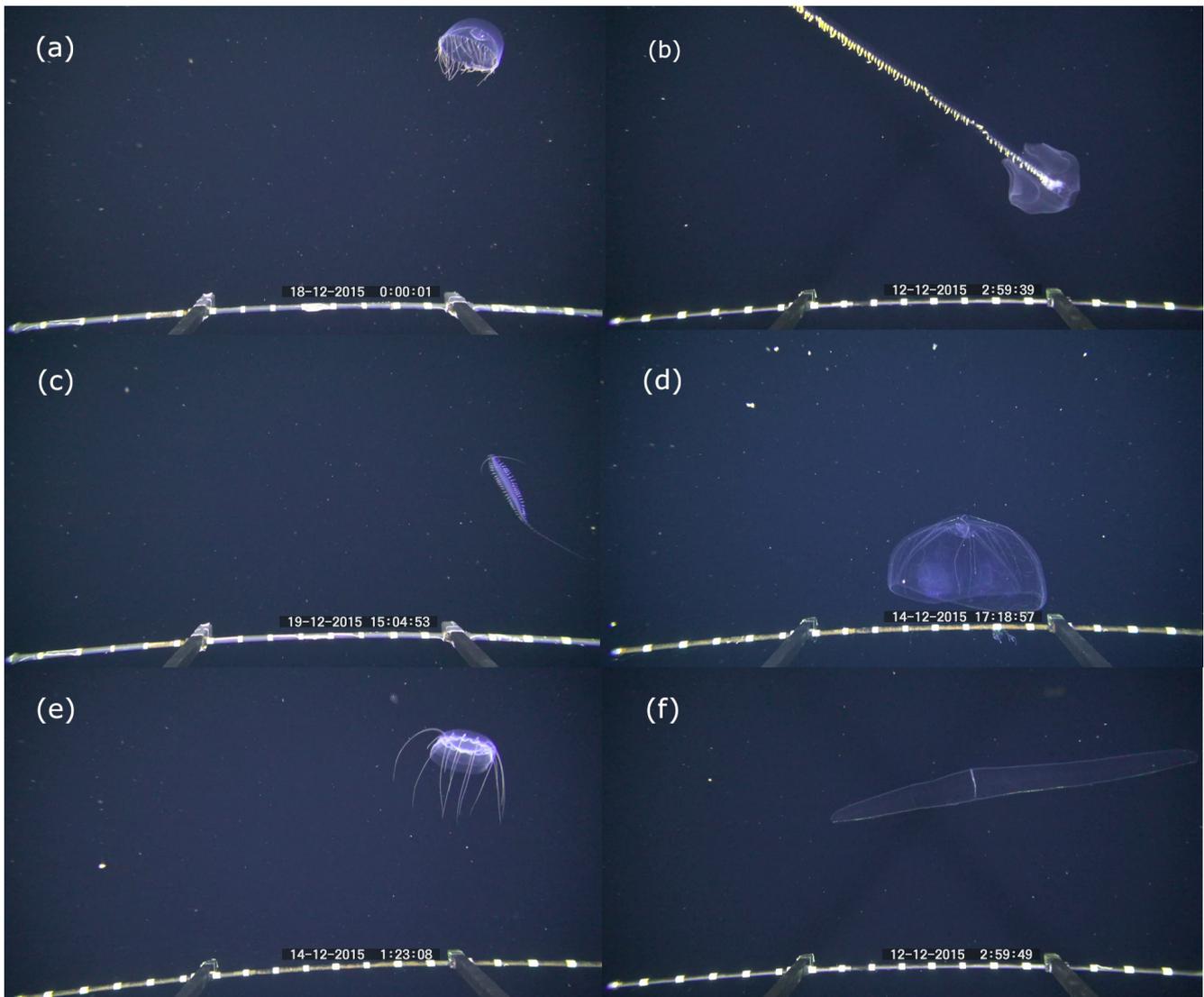


Figure 5. Examples of organisms encountered during pelagic video transects with PELAGIOS during cruise MSM49 in the eastern tropical Atlantic. **(a)** A medusa *Halitrephes* sp. **(b)** A siphonophore *Praya dubia*. **(c)** A tomopterid worm. **(d)** The ctenophore *Thalassocalyce inconstans*. **(e)** The medusa *Solmissus*. **(f)** The ctenophore *Cestum*. The distance between the white bands on the horizontal bar on the bottom of the images is 5 cm.

of distinct large gelatinous zooplankton species, including medusae, ctenophores, siphonophores, and appendicularians, in the eastern tropical North Atlantic using PELAGIOS video transects and correlated the biological patterns to the oxygen gradients (Neitzel, 2017; Hoving et al., 2019).

During various cruises, the UVP5 was mounted underneath the PELAGIOS providing concomitant data on macrozooplankton and nekton (PELAGIOS) as well as particles and mesozooplankton (UVP5). The combination of the two instruments provides a great opportunity to assess both the mesopelagic fauna and particles during one sampling event. The joint deployment of the PELAGIOS and UVP5 also allowed an estimation of the sampled water volume of the

PELAGIOS as described above. The linear relationship between counts of the nonmoving *Poebobius* sp. with UVP5 and the PELAGIOS indicates the comparability of the two different methods for animals in this size class and provides a correction factor to estimate organism abundance (ind. m^{-3}) from PELAGIOS count (ind. s^{-1}) data.

The field of view (FOV) derived from the UVP5 comparison for the PELAGIOS was estimated to be 0.23 m^2 in comparison to 0.45 m^2 based on measurement of the scale bar at 1 m from the camera. The angle of view of the PELAGIOS is 80° , and therefore the field of view (FOV) is much smaller than the FOV of video transects with a wide-angle lens, e.g., by ROV Tiburon (Robison et al., 2010). When comparing

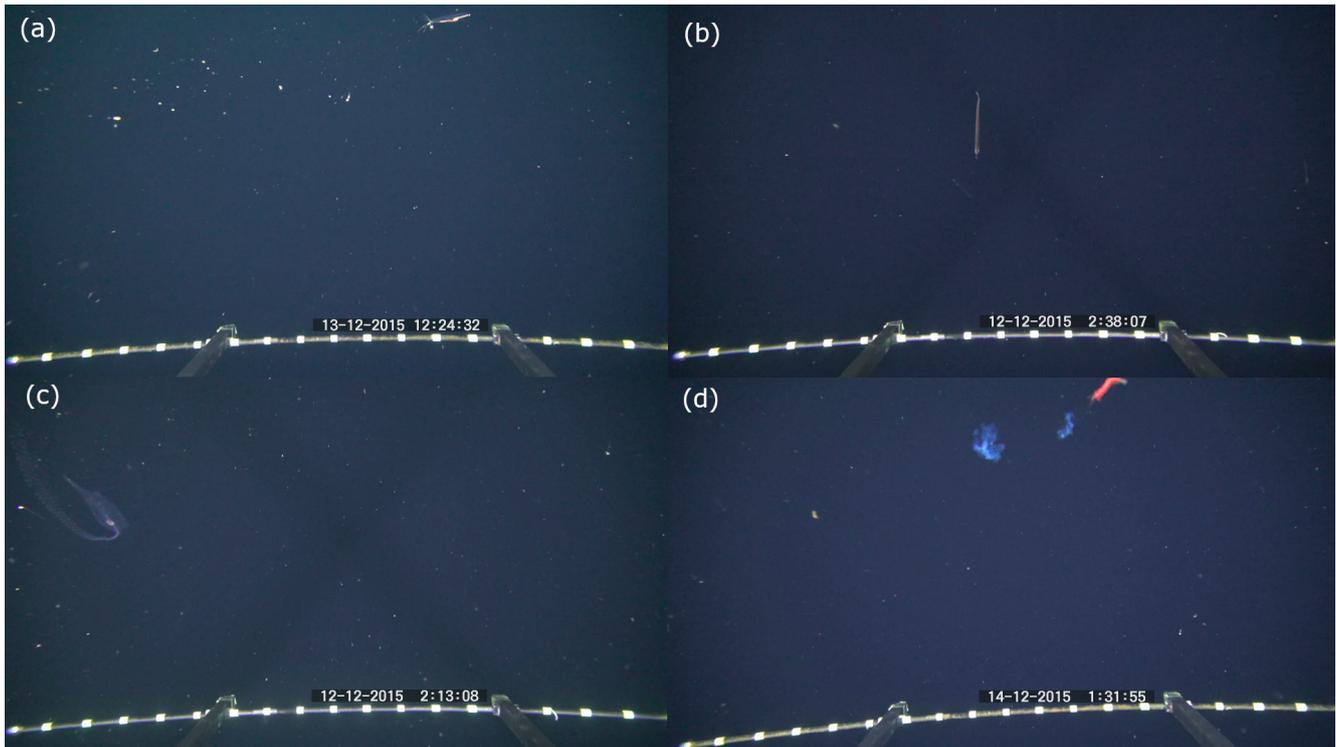


Figure 6. Examples of behaviors observed during pelagic video transects with the PELAGIOS. (a) *Poeobius* sp. in a feeding position with a mucus web (left side of the animal); (b) a dragonfish of the family Stomiidae in a vertical position; (c) a salp releasing a blastozoid chain; (d) a crustacean releasing two bioluminescent clouds while performing an escape response. The distance between the white bands on the horizontal bar on the bottom of the images is 5 cm.

the FOV, it is important to take into account the object that is observed. We provided an estimate of the FOV using *Poeobius* sp., which is a small organism that can be detected only when it is close to the camera. Therefore, the area of the FOV for the quantification of *Poeobius* sp. is smaller than when quantifying larger organisms, and the initial identification distance differs between species (Reisenbichler et al., 2017^{TS19}).

We compared PELAGIOS video transects with MOCNESS net (opening 1 m²) abundance data by integrating the PELAGIOS counts over the respective depth strata of the MOCNESS that happened at the same cruise (Lüskow et al., 2019). The diversity of the gelatinous zooplankton in the total MOCNESS catch is much lower (eight different taxa) (Lüskow et al., 2019) than in the pooled video transects (53 different annotated taxa) at the same station. The ctenophore *Beroe* is an example of a gelatinous organism captured in MOCNESS hauls and also observed on PELAGIOS transects. Normalization and subsequent standardization of the encountered *Beroe* in MOCNESS and PELAGIOS transects show that on the same station and the same depths, PELAGIOS observes 3–5 times more *Beroe* at the three depths where they were encountered by both instruments. Additionally, the PELAGIOS also repeatedly observed *Beroe* at depths where they were not captured by MOCNESS at all

(although there were also depths where PELAGIOS did not observe any *Beroe*). Preliminary comparisons of the data obtained with PELAGIOS and with MOCNESS indicate substantial differences in the documented fauna, a phenomenon also observed in previous comparisons between optical and net data (Remsen et al., 2004). Many more gelatinous taxa were observed during PELAGIOS video transects than were captured in MOCNESS catches at the same station (data presented here, Lüskow et al., 2019) due to the delicate nature of many ctenophores, medusae, and siphonophores, preventing their intact capture by nets. A notable exception are the small and robust calycothoran colonies of the families Diphyidae and Abylidae, which were also captured by MOCNESS. In contrast, avoidance behavior of strongly and fast-swimming jellyfish (e.g., *Atolla*, *Periphylla*), which may escape from the relatively slowly towed PELAGIOS, may explain their increased occurrence in nets compared to video recordings. While PELAGIOS is certainly suitable for visualizing delicate gelatinous fauna, it cannot replace net or ROV sampling since complementary specimen collections are needed to validate the identity of organisms that were observed during PELAGIOS video observations. Therefore, it is desired that net tows with open and closing nets such as Multinet Maxi or MOCNESS are performed in the same areas or that collections during submersible dives are made. An advan-

tage of ROVs over PELAGIOS is the ROV's ability to stop on organisms for detailed close-up recording and potentially the collection of the observed organisms. This is not possible with PELAGIOS as the ship tows the instrument.

5 While the imaging processing pipeline is not as streamlined as in other optical systems that use still images such as the VPR or the UVP5, the potential of the PELAGIOS as an exploration tool is illustrated by the discovery of previously undocumented animals. An example is the ctenophore
10 *Kiyohimea usagi* (Matsumoto and Robison, 1992), which was observed seven times by the PELAGIOS and once by the manned submersible JAGO during cruises in the eastern tropical North Atlantic. This large (> 40 cm wide) lobate ctenophore was previously unknown from the Atlantic Ocean
15 and demonstrates how in situ observations in epipelagic waters can result in the discovery of relatively large fauna (Hoving et al., 2018). Since gelatinous organisms are increasingly recognized as vital players in the oceanic food web (Choy et al., 2017) and in the biological carbon pump (Robison et al., 2005^{TS20}), in situ observations with tools like the PELAGIOS can provide new important insights into the oceanic ecosystem and the carbon cycle. But small gelatinous organisms may also have a large biogeochemical impact on their environment. This was illustrated by the discovery of
25 the pelagic polychaete *Poebius* sp. during the PELAGIOS video transects in the eastern tropical North Atlantic (Christiansen et al., 2018). The observations of the PELAGIOS provided the first evidence for the occurrence of *Poebius* sp. in the Atlantic Ocean. During the R/V *Meteor* cruise M119,
30 *Poebius* was found to be extremely abundant in a mesoscale eddy. Following this discovery, it was possible to reconstruct the horizontal and vertical distribution of Atlantic *Poebius* in great detail using an extensive database of the UVP5 (956 vertical CTD or UVP5 profiles) in the eastern tropical North
35 Atlantic, and to establish that the high local abundance of *Poebius* was directly related to the presence of mesoscale eddies in which they substantially intercepted the particle export flux to the deep sea (Christiansen et al., 2018; Hauss et al., 2016).

40 Future effort should be focused on improving the assessment of the sample volume by integrating technology that can quantify it (e.g., current meters, a stereo-camera setup or a laser-based system). A stereo-camera set up would also allow for size measurements of the observed organisms, which
45 could be beneficial to estimate the biomass of the observed organisms from published size-to-weight relationships. It might also be possible to obtain similar information based on structure-from-motion approaches that proved successful in benthic video imaging (Burns et al., 2015). The PELAGIOS system can also be a platform for other sensors. For example, the PELAGIOS was used to mount and test the TULUMIS multispectral camera (Liu et al., 2018). Future developments include the preparation of the system for deployments down to 6000 m water depth. The integration of acoustic sensors would be valuable to measure the target strength
55

of camera-observed organisms, to estimate gear avoidance or attraction, and to estimate biomass and the abundance of organisms outside the field of view of the camera. We strongly encourage the use of complementary instruments to tackle the relative importance of a wide range of organisms in the oceanic pelagic ecosystem. 60

Data availability. The datasets generated and/or analyzed during the current study are available in the PANGAEA repository: <https://doi.org/10.1594/PANGAEA.902241>^{TS21 TS22}.

Author contributions. This instrument was designed, tested and applied by HJH and EF. RK and HH developed the idea of combining the PELAGIOS with the UVP5. PN and SC analyzed the data in this paper in consultation with HJH, RK, and HH. AK, UP, and PL added valuable input to the further development of the instrument and its application and/or the data interpretation. All authors contributed to writing the paper. All authors approved the final submitted paper.^{CE17} 70

Competing interests. The authors declare that they have no conflict of interest.

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References

- Barham, E. G.: Siphonophores and the deep scattering layer, *Science*, 140, 826–828, 1963.
- Benfield, M. C., Davis, C. S., Wiebe, P. H., Gallagher, S. M., Lough, R. G., and Copley, N. J.: Video Plankton Recorder estimates of copepod, pteropod and larvacean distributions from a stratified region of Georges Bank with comparative measurements from a MOCNESS sampler, *Deep-Sea Res. Pt. II*, 43, 1925–1945, 1996.
- Benfield, M. C., Grosjean, P., Culverhouse, P. F., Irigoien, X., Sieracki, M. E., Lopez-Urrutia, A., Dam, H. G., Hu, Q., Davis, C. S., Hansen, A., Pilskaln, C. H., Riseman, E. M., Schultz, H., Utgoff, P. E., and Gorsky, G.: RAPID: Research on Automated Plankton Identification, *Oceanography*, 20, 172–187, 2007.
- Biard, T., Picheral, M., Mayot, N., Vandromme, P., Hauss, H., Gorsky, G., Guid, L., Kiko, R., and Not, F.: In situ imaging reveals the biomass of giant protists in the global ocean, *Nature*, 532, 504–507, 2016.
- Burns, J. H. R., Delparte, D., Gates, R. D., and Takabayashi, M.: Integrating structure-from-motion photogrammetry with geospatial software as a novel technique for quantifying 3D ecological characteristics of coral reefs, *PeerJ*, 3, e1077, [TS24](#), 2015.
- Bush, S. L., Caldwell, R. L., and Robison, B. H.: Ink utilization by mesopelagic squid, *Mar. Biol.*, 152, 485–494, 2007.
- Choy, C. A., Haddock, S. H. D., and Robison, B. H.: Deep pelagic food web structure as revealed by *in situ* feeding observations, *Proc. R. Soc. B. Biol. Sci.*, 284, 1869, <https://doi.org/10.1098/rspb.2017.2116>, 2017.
- Christiansen, B., Buchholz, C., [TS25](#) Buchholz, F., Chi, X., Christiansen, S., Denda, A., Fabrizio, E., Hauss, H., Hoving, H.-J. T., and Janßen, S.: SEAMOX: The Influence of Seamounts and Oxygen Minimum Zones on Pelagic Fauna in the Eastern Tropical Atlantic, Technical Cruise Report Cruise No. MSM49 28 November–21 December 2015 Las Palmas de Gran Canaria (Spain)-Mindelo (Republic of Cape Verde), 2016.
- Christiansen, S., Hoving, H. J. T., Schütte, F., Hauss, H., Karstensen, J., Körtzinger, A., Schröder, M., Stemmann, L., Christiansen, B., Picheral, M., Brandt, P., Robison, B. H., Koch, R., and Kiko, R.: Particulate matter flux interception in oceanic mesoscale eddies by the polychaete *Poebius* sp., *Limnol. Oceanogr.*, 63, 2093–2109, 2018.
- Cowen, R. K. and Guigand, C. M.: In situ ichthyoplankton imaging system (ISIIS): system design and preliminary results, *Limnol. Oceanogr.-Meth.*, 6, 126–132, 2008.
- Genin, A.: Bio-physical coupling in the formation of zooplankton and fish aggregations over abrupt topographies, *J. Marine Syst.*, 50, 3–20, 2004.
- Gilly, W. F., Beman, J. M., Litvin, S. Y., and Robison, B. H.: Oceanographic and biological effects of shoaling of the oxygen minimum zone, *Annual Review in Marine Science*, 5, 393–420, 2013.
- Gomes-Pereira, J. N., Auger, V., Beisiegel, K., Benjamin, R., Bergmann, M., Bowden, D., Buhl-Mortensen, P., De Leo, F. C., Dionísio, G., Durden, J. M., Edwards, L., Friedman, A., Greinert, J., Jacobsen-Stout, N., Lerner, S., Leslie, M., Nattkemper, T. W., Sameoto, J. A., Schoening, T., Schouten, R., Seager, J., Singh, H., Soubigou, O., Tojeira, I., van den Beld, I., Dias, F., Tempera, F., and Santos, R. S.: Current and future trends in marine image annotation software, *Prog. Oceanogr.*, 149, 106–120, 2016.
- Haddock, S. H. D.: A golden age of gelata: past and future research on planktonic ctenophores and cnidarians, *Hydrobiologia*, 530, 549–556, 2004.
- Haddock, S. H. D.: Comparative feeding behavior of planktonic ctenophores, *Integ. and Comp. Biol.*, 47, 847–853, 2007, [TS26](#)
- Hamner, W. M., Madin, L. P., Alldredge, A. L., Gilmer, R. M., and Hamner, P. P.: Underwater observations of gelatinous zooplankton: sampling problems, feeding biology and behavior, *Limnol. Oceanogr.*, 20, 907–917, 1975.
- Hamner, W. M. and Robison, B. H.: In situ observations of giant appendicularians in Monterey Bay, *Deep-Sea Res.*, 39, 1299–1313, 1992.
- Harbison, G., Madin, L., and Swanberg, N.: On the natural history and distribution of oceanic ctenophores, *Deep-Sea Res.*, 25, 233–256, 1978.
- Harbison, G. R.: Toward a study of the biogeography of the pelagic ctenophores, in: *Pelagic Biogeography*, edited by: Pierrot-Bults, A. C., van der Spoel, S., Zahuranec, B. J., and Johnson, R. K., UNESCO Technical Papers in Marine Science, 49, 112–117, 1986, [TS27](#)
- Haslob, H., Rohlf, N., and Schnack, D.: Small scale distribution patterns and vertical migration of North Sea herring larvae (*Clupea harengus*, Teleostei: Clupeidae) in relation to abiotic and biotic factors, *Sci. Mar.*, 73, 13–22, 2009.
- Haury, L., Fey, C., Newland, C., and Genin, A.: Zooplankton distribution around four eastern North Pacific seamounts, *Prog. Oceanogr.*, 45, 69–105, 2000.
- Hauss, H., Christiansen, S., Schütte, F., Kiko, R., Edvam Lima, M., Rodrigues, E., Karstensen, J., Löscher, C. R., Körtzinger, A., and Fiedler, B.: Dead zone or oasis in the open ocean? Zooplankton distribution and migration in low-oxygen modewater eddies, *Biogeosciences*, 13, 1977–1989, <https://doi.org/10.5194/bg-13-1977-2016>, 2016.
- Hosia, A., Falkenhaus, T., Baxter, E. J., and Pagès, F.: Abundance, distribution and diversity of gelatinous predators along the Mid Atlantic Ridge: A comparison of different sampling methodologies, *PLoS One*, 12, e0187491, [TS28](#), 2017.
- Hoving, H. J. T. and Robison, B. H.: Deep-sea in situ observations of gonatid squid and their prey reveal high occurrence of cannibalism, *Deep-Sea Res. Pt. I*, 116, 94–98, 2016.
- Hoving, H. J. T., Zeidberg, L., Benfield, M., Bush, S., Robison, B. H., and Vecchione, M.: First in situ observations of the deep-sea squid *Grimalditeuthis bonplandi* reveals unique use of tentacles, *Proc. R. Soc. B*, 280, 1769, [TS29](#), 2013.
- Hoving, H. J. T., Bush, S. L., Haddock, S. H. D., and Robison, B. H.: Bathyal feasting: post-spawning squid as a source of carbon for deep-sea benthic communities, *Proc. R. Soc. B*, 284, 20172096, [TS30](#), 2017.
- Hoving, H.-J., Neitzel, P., and Robison, B.: In situ observations lead to the discovery of the large ctenophore *Kiyohimea usagi* (Lobata: Eurhampheidae) in the eastern tropical Atlantic, *Zootaxa*, 4526, 232–238, 2018.
- Isaacs, J. D. and Schwartzlose, R. A.: Migrant sound scatterers: interaction with the sea floor, *Science*, 150, 1810–1813, 1965.

- Hull, P. M., Osborn, K. J., Norris, R. D., and Robison, B. H.: Seasonality and depth distribution of a mesopelagic foraminifer, *Hastigerinella digitata*, in Monterey Bay, California, *Limnol. Oceanogr.*, 56, 562–576, 2011.
- 5 Kiko, R., Biastoch, A., Brandt, P., Cravatte, S., Hauss, H., Hummels, R., Kriest, I., Marin, F., McDonnell, A. M. P., Oschlies, A., Picheral, M., Schwarzkopf, F. U., Thurnherr, A. M., and Stemmann, L.: Biological and physical influences on marine snowfall at the equator, *Nat. Geosci.*, 42, 852–858, 2017.
- 10 Kubodera, T., Koyama, Y., and Mori, K.: Observations of wild hunting behaviour and bioluminescence of a large deep-sea, eight-armed squid, *Taningia danae*, *P. Roy. Soc. B*, 274, 1029–1034, 2007.
- Linke, P., Schmidt, M., Rohleder, M., Al-Barakati, A., and Al-Farawati, R.: Novel online digital video and high-speed data broadcasting via standard coaxial cable onboard marine operating vessels, *Mar. Tech. Soc. J.*, 49, 7–18, 2015.
- 15 Liu, H., Sticklus, J., Köser, K., Hoving, H. J. T., Ying, C., Hong, S., Greinert, J., and Schoening, T.: TuLUMIS – A tunable LED-based underwater multispectral imaging system, *Opt. Express*, 26, 7811–7828, 2018.
- Madin, L., Horgan, E., Gallager, S., Eaton, J., and Girard, A.: LAPIS: A new imaging tool for macrozooplankton, *IEEE J. Oceanic Eng.*, [TS31](#) 1-4244-0115-1/06, 2006.
- 25 Matsumoto, G. I. and Robison, B. H.: *Kiyohimea usagi*, a new species of lobate ctenophore from the Monterey Submarine Canyon, *B. Mar. Sci.*, 51, 19–29, 1992.
- Möller, K. O., John, M. S., Temming, A., Floeter, J., Sell, A. F., Herrmann, J.-P., and Möllmann, C.: Marine snow, zooplankton and thin layers: indications of a trophic link from small-scale sampling with the Video Plankton Recorder, *Mar. Ecol. Prog. Ser.*, 468, 57–69, 2012.
- 30 Neitzel, P.: The impact of the oxygen minimum zone on the vertical distribution and abundance of gelatinous macrozooplankton in the Eastern Tropical Atlantic. MSc Thesis, Christian-Albrechts-Universität Kiel, Germany, 75 pp., 2017.
- Ohman, M. D., Davis, R. E., Sherman, J. T., Grindley, K. R., Whitmore, B. M., Nickels, C. F., and Ellen, J. S.: Zooglider: An autonomous vehicle for optical and acoustic sensing of zooplankton, *Limnol. Oceanogr.-Meth.*, 17, 69–86, 2019.
- 40 Oschlies, A., Brandt, P., Stramma, L., and Schmidtko, S.: Drivers and mechanisms of ocean deoxygenation, *Nat. Geosci.*, 11, 467–473, 2018.
- Picheral, M., Guidi, L., Stemmann, L., Karl, D. M., Iddaoud, G., and Gorsky, G.: The Underwater Vision Profiler 5: An advanced instrument for high spatial resolution studies of particle size spectra and zooplankton, *Limnol. Oceanogr.-Meth.*, 8, 462–473, 2010.
- 45 Purser, A., Marcon, Y., Dreutter, S., Hoge, U., Sablotny, B., Hehemann, L., Lemburg, J., Dorschel, B., Biebow, H., and Boetius, A.: Ocean floor observation and bathymetry system (OFOBS): A new towed camera/sonar system for deep-sea habitat surveys, *IEEE J. Oceanic Eng.*, 44, 87–99, 2018.
- Ramirez-Llodra, E., Brandt, A., Danovaro, R., De Mol, B., Escobar, E., German, C. R., Levin, L. A., Martinez Arbizu, P., Menot, L., Buhl-Mortensen, P., Narayanaswamy, B. E., Smith, C. R., Tittensor, D. P., Tyler, P. A., Vanreusel, A., and Vecchione, M.: Deep, diverse and definitely different: unique attributes of the world's largest ecosystem, *Biogeosciences*, 7, 2851–2899, <https://doi.org/10.5194/bg-7-2851-2010>, 2010.
- 55 Remsen, A., Hopkins, T. L., and Samson, S.: What you see is not what you catch: a comparison of concurrently collected net, Optical Plankton Counter, and Shadowed Image Particle Profiling Evaluation Recorder data from the northeast Gulf of Mexico, *Deep-Sea Res. Pt. I*, 51, 129–151, 2004.
- Robinson, C., Steinberg, D. K., Anderson, T. R., Arístegui, J., Carlson, C. A., Frost, J. R., Ghiglione, J.-F., Hernández-León, S., Jackson, G. A., Koppelman, R., Quéguiner, B., Ragueneau, O., Rassoulzadegan, F., Robison, B. H., Tamburini, C., Tanaka, T., Wishner, K. F., and Zhang, J.: Mesopelagic zone ecology and biogeochemistry – a synthesis, *Deep-Sea Res. Pt. II*, 57, 1504–1518, 2010.
- 70 Robison, B. H.: Midwater biological research with the WASP ADS, *Mar. Tech. Soc. J.*, 17, 21–27, 1983.
- Robison, B. H.: The coevolution of undersea vehicles and deep-sea research, *Mar. Tech. Soc. J.*, 33, 65–73, 1999.
- 75 Robison, B. H.: Deep pelagic biology, *J. Exp. Mar. Boil. Ecol.*, 300, 253–272, 2004.
- Robison, B. H.: Conservation of Deep Pelagic Biodiversity, *Cons. Biol.*, 23, 847–858, 2009. [TS32](#)
- Robison, B. H. and Wishner, K.: Biological research needs for submersible access to the greatest ocean depths, *Mar. Tech. Soc. J.*, 24, 34–37, 1990.
- 80 Robison, B. H., Reisenbichler, K. R., Sherlock, R. E., Silguero, J. M. B., and Chavez, F. P.: Seasonal abundance of the siphonophore, *Nanomia bijuga*, in Monterey Bay, *Deep-Sea Res. Pt. II*, 45, 1741–1752, 1998.
- Robison, B. H., Sherlock, R. E., and Reisenbichler, K.: The bathypelagic community of Monterey Bay, *Deep-Sea Res. Pt. II*, 57, 1551–1556, 2010.
- Schlining, B. and Jacobsen Stout, N.: MBARI's Video Annotation and Reference System. In: Proceedings of the Marine Technology Society/Institute of Electrical and Electronics Engineers Oceans Conference, Boston, Massachusetts, 1–5, 2006.
- 90 Schulz, J., Barz, K., Mengedoht, D., Hanken, T., Lilienthal, H., Rieper, N., Hoops, J., Vogel, K., and Hirche, H.-J.: A Light-Frame Onsite Key Species Investigation, *IEEE J. Oceanic Eng.*, [TS33](#), 2009. [TS34](#)
- Schulz, J., Barz, K., Ayon, P., Lüdtke, A., Zielinski, O., Mengedoht, D., and Hirche, H. J.: Imaging of plankton specimens with the Lightframe On-sight Keyspecies Investigation (LOKI) system, *J. Eur. Opt. Soc.-Rapid*, 5S, 1–9 [TS35](#), 2010.
- 100 Stramma, L., Prince, E. D., Schmidtko, S., Luo, J., Hoolihan, J. P., Visbeck, M., Wallace, D. W. R., Brandt, P., and Körtzinger, A.: Expansion of oxygen minimum zones may reduce available habitat for tropical pelagic fishes, *Nat. Clim. Change*, 2, 33–37, 2012.
- 105 Webb, T. J., Vanden Berghe, E., and O'Dor, R.: Biodiversity's Big Wet Secret: The Global Distribution of Marine Biological Records Reveals Chronic Under-Exploration of the Deep Pelagic Ocean, *PLoS ONE*, 5, e10223, [TS36](#), 2010.
- Wiebe, P. H. and Benfield, M. C.: From the Hensen net toward four-dimensional biological oceanography, *Prog. Oceanogr.*, 56, 7–136, 2003.
- 110 Wishner, K. F., Outram, D. M., Seibel, B. A., Daly, K. L., and Williams, R. L.: Zooplankton in the eastern tropical north Pacific: Boundary effects of oxygen minimum zone expansion, *Deep-Sea Res. Pt. I*, 79, 122–140, 2013.
- Youngbluth, M., Sørnes, T., Hosiya, A., and Stemmann, L.: Vertical distribution and relative abundance of gelatinous zooplankton, in

situ observations near the Mid-Atlantic Ridge, Deep-Sea Res. Pt. II, 55, 119–125, 2008.

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