Demosponges from the Northern Mid-Atlantic Ridge shed more light on the diversity and biogeography of North Atlantic deep-sea sponges

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In July–August 2004, the Mid-Atlantic Ridge Ecosystems (MAR-Eco) expedition collected Demospongiae (Porifera) from the Northern Mid-Atlantic Ridge (MAR) north of the Azores, between 41°N and 61°N. Demosponges were found at 10 stations, at depths ranging from 753 to 3046 m. Twenty-two species were identified: 17 Tetractinellida, one Polymastiida, one Suberitida, two Poecilocosclerida and one Dendroceratida. The study of this material is an opportunity to revise the taxonomy and the North Atlantic distribution of each of these deep-sea species. Some species are particularly rare and poorly known (Tetilla longipilis, Tetilla sandalina, Craniochilus azorica, Polymastia corticata) and two are new to science: Forcepia (Forcepia) toxifer sp. nov. and Iotroata paravaridens sp. nov. This study suggests that the MAR is not a longitudinal barrier for the dispersal of deep-sea demosponges while on the contrary, the Charlie-Gibbs Fracture Zone (CGFZ) may be a latitudinal border for the dispersal of deep-sea demosponges, due to great depths and currents.

**Keywords:** Porifera, Demospongiae, taxonomy, Mid-Atlantic Ridge, Charlie-Gibbs Fracture Zone, amphi-Atlantic, MAR-Eco, Tetractinellida, Geodia, new species

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**INTRODUCTION**

The Mid-Atlantic Ridge (MAR) between Iceland (63°N) and the Azores (36°–39°N) spans across 3000 km. About mid-way between Iceland and the Azores (around 52°N), the Charlie-Gibbs Fracture Zone (CGFZ) offsets the MAR about 300 km, with depths reaching 4500 m (Felley et al., 2008), thereby opening the deepest connection between North-East and North-West Atlantic waters (Figure 1). Furthermore, a northern branch of the North Atlantic Current (NAC) crosses the Mid-Atlantic Ridge from west to east just over the CGFZ while being under the influence of colder subpolar waters on its northern border. The CGFZ area is thus considered a major latitudinal transition zone in the environment and biodiversity of the MAR (Vecchione et al., 2010) and is now part of the world’s first network of marine protected areas (MPAs) by both OSPAR (Oslo-Paris Convention) and the NEAFC (North-East Atlantic Fisheries Commission) (O’Leary et al., 2012). Therefore, the poorly explored MAR and CGFZ in particular are of considerable interest for deep-sea biodiversity and biogeography studies. The Mid-Atlantic Ridge Ecosystems (MAR-Eco) project (2001–2010, www.mar-eco.no), part of the ’Census of Marine Life’ (www.coml.org), was aimed at studying the patterns and processes of the ecosystems of the northern Mid-Atlantic. In July–August 2004, the Norwegian RV ‘G.O. Sars’ collected benthic samples on the MAR between the Azores and the southern region of the Reykjanes Ridge. Most of the material has already been identified and published, notably in special issues of ’Deep-Sea Research Part II’ and ‘Marine Biology Research’ in 2008 (http://www.tandfonline.com/toc/smar20/4/1-2, http://www.sciencedirect.com/science/journal/09670645/55/1, both accessed 15 April 2014). Taxa examined and studied in these special issues include fish, fish parasites, Cetacea, corals, copepods, planktonic cnidarians and ctenophores, Holothuroidea, poutrelsid sea urchins, Isopoda (Aegidae), Ophiuroidea, Anthozoa, Asteroidea, Echiura, Sipuncula, Brachiopoda and Hexactinellida. Additional taxon checklists of benthic organisms, including a preliminary demosponge species list of our identifications, were added and discussed by Gebruk et al. (2010). In the present paper, we update and refine this checklist while thoroughly describing and discussing the taxonomy of each species. Indeed, the study of this material is an opportunity to revise some poorly known Atlantic deep-sea species. We also compiled distribution maps for most of these species. They will form the basis of
a discussion on the biogeography of deep-sea demosponges in the North Atlantic.

MATERIALS AND METHODS

Sponge sampling

Sponge specimens were collected by bottom trawl on board the RV ‘G.O. Sars’ during Leg 2 (4 July – 5 August) of the 2004 MAR-Eco expedition (cf. cruise report at http://www.mar-eco.no/sci/cruises/expedition_report_rv_g.o.sars_2004, accessed 8 March 2014). The trawl was a Campelen 1800 shrimp trawl with a 22 mm mesh size cod-end liner and a 12–17 m by 4.5 m opening at 50 m doorspread. Environmental data associated with the trawls were near-bottom measurements obtained with a CTD sensor (Søiland et al., 2008). For a detailed station list with coordinates, depths and environmental data, and for a full account of the approach and methods for sampling of the benthic fauna during this expedition, see Bergstad & Gebruk (2008). Demosponges were found at 10 stations, at depths ranging from 753 to 3046 m (Table 1, Figure 1): two stations were on the north-west of the CGFZ (stations 72 and 70), two shallower stations (<1000 m) were on MAR seamounts (stations 65 and 53), and finally six stations were north of the Azores (stations 40, 42, 44, 46, 50 and 52). Specimens were preserved in 4% buffered formaldehyde on board and later transferred to 70% ethanol. Specimens are stored at room temperature in the Bergen Museum (ZMBN).

Morphology studies

To collect the spicules, sponge tissue was digested in nitric acid on a microscope glass slide. Spicules were then washed with water and with ethanol 96% and mounted in Euparal; 25 spicules per spicule type were measured, unless otherwise stated. Measurements of all spicules were made with a light microscope. Width of triaene rhabdomes was measured right under the cladomes. Width of the oxeas was measured in the middle. For Thenea species, we only measured microscleres because megascleres have been shown to be similar in North Atlantic Thenea (Cárdenas & Rapp, 2012). Some of these spicules were placed on a stub, coated with a gold/palladium mix and observed with the ZEISS Supra 55VP and JEOL JSM-840A scanning electron microscopes (SEM) at the Laboratory of Electron Microscopy, University of Bergen. Thick sections (100–800 μm) were made with a diamond wafering blade and a low speed saw using an Agar Low Viscosity Resin kit (Agar Scientific) in accordance with the manufacturer’s mixing instructions to make a hard embedding medium. Digital pictures of these sections were taken with a Nikon.
camera fixed to a stereomicroscope (Leica M216 A). Thick sections, spicule preparations and SEM stubs are stored in the Bergen Museum. With respect to taxonomy, our study will follow the demosponge classification proposed by Morrow & Ca´rdenas (2015).

Distribution maps of species

All the records obtained from our identifications and the literature were compiled and mapped with GeoMapApp v. 3.3.9 (http://www.geomapapp.org), using the North Polar base map projection and the default Global Multi-Resolution Topography Synthesis (Ryan et al., 2009). When the latitude/longitude information was missing but the locality was given, we reconstructed the geographic coordinates using Google Earth. Distribution maps of boreo-arctic Geodia species have been updated from Cárdenas et al. (2013), by adding the MAR-Eco records and records from the Reykjanes Ridge (Copley et al., 1996). Additional records were also added for (i) Geodia atlantica from the Kerry Head Reefs cruise CV13012 (August 2013, chief scientist: Louise Allcock, material sorted by C. Morrow, identifications by PC) and for (ii) Geodia macandrewii from the Porcupine Bank (CE13008 campaign, June 2013, RV ‘Celtic Explorer’ using the ROV ‘Holland I’, chief scientist: Louise Allcock, material sorted by C. Morrow, identifications by PC). Maps of boreo-arctic Thenea species and Poecilastera compressa have also been updated from Steenstrup & Tendal (1982).

Table 1. List of the 22 Demospongiae collected by the 2004 MAR-Eco expedition. Temperature, salinity and oxygen data obtained with near-bottom measurements with a CTD sensor (Søiland et al., 2008).

<table>
<thead>
<tr>
<th>Stations</th>
<th>Area</th>
<th>North of Azores</th>
<th>F</th>
<th>H</th>
<th>CGFZ (north-west)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean depth (m)</td>
<td>2961</td>
<td>2078</td>
<td>1742</td>
<td>3046</td>
<td>2600</td>
</tr>
<tr>
<td>Temperature (°C)</td>
<td>3.16</td>
<td>3.93</td>
<td>5.01</td>
<td>3.48</td>
<td>3.79</td>
</tr>
<tr>
<td>Salinity (psu)</td>
<td>34.92</td>
<td>34.97</td>
<td>35.06</td>
<td>34.93</td>
<td>34.95</td>
</tr>
<tr>
<td>Oxygen (mL L⁻¹)</td>
<td>5.94</td>
<td>5.78</td>
<td>5.38</td>
<td>5.89</td>
<td>5.82</td>
</tr>
</tbody>
</table>

ASTROPHORINA
Geodiidae
- Geodia atlantica* 5
- Geodia barretti* 4
- Geodia hentscheli* 1
- Geodia cf. hentscheli 7
- Geodia macandrewii* 3
- Geodia nodastrella* 1
- Geodia phlegreae* 1
- Stelletta tuberosa* 2
- Stelletta rhaphidiophora* 2
- Stryphinus fortis* 1

Ancorinidae
- Stelletta rhaphidiophora* 1

SIPHIOPHORINA
Tetillidae
- Craniella azorica 2
- Tetilla longipilis 1
- Tetilla sandalina 1

POLYMASTIIDA
Polymastiidae
- Polymastia corticata* 1

SUBERITIDA
Stylocordylidae
- Stylocordyla borealis* 1

POECICLOSCLERIDA
Coelosphaeridae
- Forcepia (Forcepia) toxsfera sp. nov. 1

DENDROCERATIDA
Dictyodendrillidae
- Spongionella aff. pulchella 1

CGFZ, Charlie Gibbs Fracture Zone; F, Faraday Seamount; H, Hecate Seamount.

*Amphi-Atlantic species.
and Cárdenas & Rapp (2012), with new records from the CE13008 campaign, the NEREIDA campaigns (see acknowledgements for more details, material sorted by M. Best, identifications by PC), as well as groundfish bottom trawl surveys carried out by the Spanish Institute of Oceanography (IEO) and the European Union (EU) on board the Spanish RV ‘Vizconde de Eza’. We also added records of specimens examined in the collections of the Zoological Museum in Copenhagen (ZMUC): this includes records of *P. compressa* (Danish Ingolf Expedition 1895–1896), *Stryphnum fortes* (Ingolf), *Tetilla longipilis* (Ingolf, BIOICE 1991–2004), *Stelleta tuberosa* (Ingolf), *Stelleta rhaphidiophora* (Ingolf, Lynch 1972–1973, BIOICE, Paamui 1994) and *Stylocordyla boralis* (BIOICE, BIOFAR 1987–1993, Michael Sars 1902). We also re-examined *Tethsea* spp. specimens from museums in Monaco (MOM) and Stockholm (SMNH). More records came from underwater picture databases; that of the Marine area database for Norwegian waters (MAREANO, http://www.mareano.no) and that of habitat mapping studies in the West Shetland Channel (pictures courtesy of K. L. Howell) (Howell et al., 2010). We also used the Global Biodiversity Information Facility (GBIF) website (http://www.gbif.org, accessed 22 April 2014) to retrieve records of *Stylocordyla boralis* from the eastern US coast, from specimens stored at the Yale Peabody Museum (2009; specimen data records available through distributed digital resources). The compiled records used for the species distribution maps – including geographic positions (decimal degrees coordinates), museum collection or reference, temperature and salinity when available – can be found in the Supplementary material.

**Abbreviations**

The following abbreviations are used for the institutions from which we have examined material, or where material will be deposited. BIO: Bedford Institute of Oceanography, Halifax, NS, Canada; GNM: Göteborg Natural History Museum, Sweden; Mc: National Museums, Northern Ireland, Holywood; MNHN: Museum of Natural History, Paris, France; MOM: Oceanographic Museum of Monaco, Monaco; NHM: The Natural History Museum, London, UK; PC: Private Collection, Department of Biology, UoB, Norway (HTR, PC); RMNH: The State Museum of Natural History, Leiden, the Netherlands; SMF: Senckenberg Museum, Frankfurt, Germany; SMNH: Swedish Museum of Natural History, Stockholm; UPSZMC: The Zoological Museum of Uppsala, Sweden; ZMAPOR: The Zoological Museum, University of Amsterdam, the Netherlands; ZMUC: The Zoological Museum, Copenhagen, Denmark; ZMBN: Bergen Museum, Norway.

**RESULTS**

Table 1 lists the 22 demosponge species identified per station: 14 species belonged to the Astrophorida and three to the Spirophorida so that a total of 17 species belonged to the order Tetractinellida. The other five species belonged to the orders Polymastiida (one), Suberitida (one), Poecilosclerida (two) and Dendroceratida (one). An advantage we had when studying this collection is that most of the Astrophorina identified in this collection had been revised using morphological (Cárdenas & Rapp, 2012; Cárdenas et al., 2013) and molecular data (Cárdenas et al., 2011) so we will often refer to these revisions for additional information. Specimens cited in the aforementioned papers have been used as comparative material.

**SYSTEMATICS**

Class DEMOSPONGIAE Sollas, 1885

Subclass HETEROSCLEROMORPHA Cárdenas et al., 2012

Order TETRACTINELLIDA Marshall, 1876

Suborder ASTROPHORINA Sollas, 1887

Family GEODIIDAE Gray, 1867

Genus *Geodia* Lamarck, 1815

*Geodia atlantica* (Stephens, 1915) (Figures 2A & 3)

**MATERIAL**

Five specimens from the same trawl, Station 50–373, 2610 m: ZMBN 105633, 105637, 105639, 105641–42.

**OUTER MORPHOLOGY AND SKELETON**

**Organization** (Figures 3A–C)

Massive subspherical specimens which can be flattened (105639, 105642). 105633 is about 3 cm in diameter. Specimens 105639, 105641 and 105642 stick together along with one *Stryphnum fortes* and one *Stelletta tuberosa* (Figure 3A). Uniporal oscules and ciri: pioral pores. Colour in ethanol is whitish to light brown. Surface is smooth and slightly bumpy (‘deflated’ appearance). Cortex is 0.5 mm thick. Thick sections were made for 105639 and 105642. The choanosomes of 105639 and 105642 are filled with numerous foraminifera. In these specimens also, anatriaenes are rare and oyster-like are very numerous (as in the holotype) (Figure 3C). Skeleton organization is similar to that described in Cárdenas et al. (2013).

**SPICULES** ZMBN 105639 (Figure 3D–H)

*Megascleres*:

(a) oxeas, straight or bent, length: 3125–3325–3675 μm; width: 32–35–40 μm. (b) orthotriaenes, rare dichotriaenes, straight or slightly bent rhodome, rhodome length: 1525–2237.5–2600 μm (N = 6); width: 51–60.5–70 μm (N = 27); orthotriaene clad length: 290–376.1–480 μm (N = 18); protoclad length: 150–210 μm (N = 2); deuteroclad length: 75–230 μm (N = 2). (c) anatriaenes, straight or slightly bent rhodome, width: 18.19.3–20 μm (N = 3); clad length: 205–230–245 μm (N = 3).

*Microscleres*:

(d) sterrasters, spherical, length: 84–99.7–120 μm (N = 21), smooth rosettes. (e) spheroidactines, small actines, 7.5–10–12.5 μm in diameter. (f) oyster-like, small actines, diameter: 29–36.4–83 μm. (g) oyster-like II, small actines, usually with a larger centrum than oyster-like I, diameter: 12.5–21.8–35 μm.

**BATHYMETRIC RANGE**

65–2610 m (Cárdenas et al., 2013; this study).

**DISCUSSION**

Spicule measurements and morphologies fit with the description of this species (Cárdenas et al., 2013). But the asters are clearly more strongly spinous than in the NEA specimens (including the type). Also, the sterrasters are spherical, like in NWA specimens, whereas they were more

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elongated in NEA specimens. Although dichotriaenes are usually rare in this species, they are particularly common in specimens 105633 and 105637. The external morphology is a bit different from more northern specimens as well: darker colour and ‘deflated’ surface appearance. To conclude, as suggested before (Cárdenas et al., 2013) there could be a southern morphotype of *G. atlantica*, to which the MAR-Eco specimens belong.
Geodia barretti Bowerbank, 1858
(Figures 2B & 4)

MATERIAL
Four specimens, all from Station 70–385, 1650 m: ZMBN 105662, 105665, 105667, 105672.

OUTER MORPHOLOGY AND SKELETON ORGANIZATION (FIGURE 4A, B)

Massive subspherical white specimens. 105662 is the largest specimen, about 7 cm in diameter; 105672 is the smallest specimen, about 1 cm in diameter. All specimens have a single preosculate and cribriporal pores. 105667 is growing on Stelletta rhaphidiophora. Cortex 0.4–0.5 mm (105662). Thick sections were made for 105662, the skeleton organization is in accordance with previous observations (Cárdenas et al., 2013). Radial crystalline structures (‘spherulites’) observed in the sections, these structures are about 12–25 μm in diameter with fan-shaped fibrous structures, with a rather smooth clear surface appearance. They are very abundant just below the cortex and around canals. They sometimes group in very confused masses.

SPICULES, ZMBN 105662 (FIGURE 4C–I)
Megascleres: (a) oxeas I, straight or bent, length: 1075–2530.3–4275 μm; width: 12.5–39.2–62 μm. (b) oxeas II (=microxeas), straight or slightly bent, sometimes slightly centrotyloate, length: 190–255.3–390 μm; width: 6–8.8–11 μm. (c) dichotriaenes, rhabdome length: 2175–3375 μm (N = 2); width: 80–109.2–130 μm; protoclad length: 150–211.6–310 μm; deuteroclad length: 110–216.3–320 μm. (d)
meso/protriaenes, rare, rhabdome length: up to 2300 μm; width: 7.5–15 μm (N = 2); clad length: 60 μm; central clad length: 98 μm. Microscleres: (e) sterrasters, elongated, rosettes with warts, length: 96–104–114 μm, width: 91–96.3–106 μm. (f) strongylasters, rough actines, diameter: 4–5.5–6.2 μm in diameter. (g) oxyasters I, rough actines, diameter: 36.6–56.8–78 μm. (i) oxyasters II, rough actines, diameter: 8–15.4–23.5 μm.

BATHYMETRIC RANGE
30–2000 m (Cárdenas et al., 2013).

DISCUSSION
Spicule measurements and external morphologies are perfectly in accordance with the description of the species (Cárdenas et al., 2013). A large category of oxyaster is present, as in other G. barretti which have been collected at depths deeper than 1000 m (Cárdenas & Rapp, 2013; Cárdenas et al., 2013). The radial crystalline structures, thought to be calcareous, are common in many Tetillidae (e.g. Cárdenas et al., 2009) but to our knowledge this is the first time they are reported in Geodiidae.

Geodia hentscheli Cárdenas et al., 2010
(Figures 2C, 5 & 6)

MATERIAL
One specimen from Station 72–386, 2534 m: ZMBN 105680.

OUTER MORPHOLOGY AND SKELETAL ORGANIZATION (FIGURE 5)
Massive subspherical whitish specimen, about 5 cm in diameter. Single preosculum and cribriform pores. Cortex is
0.5–0.6 mm thick, with a rather thick ectocortex (200 μm). Skeleton organization is similar to that observed in other specimens (Cárdenas et al., 2013). Crystalline structures were observed, these structures are about 50–87 μm in diameter with radial fibrous organization (very ‘bushy’ and confused appearance), and unclear borders. They are significantly larger and clearly different from the ones observed in G. barretti. They are present below the cortex and in the choanosome. Light brown granulated cells (or vacuoles?) are moderately present throughout the cortex as well as just below it; they are 10–20 μm in diameter.

**MATERIAL**

Seven specimens from Station 50–373. 2598 m: ZMBN 105632, 105634–36, 105638, 105645 (two specimens).

**COMPARATIVE MATERIAL EXAMINED**

ZMBN 25668, two wet specimens, Michael Sars 1910 Expedition, Station 53, 34.98333, 2615–2865 m, 08/06/1910, identified by E. Arnesen as ‘Isops sp.?;’ RMNH Por1458, Azores, 38.1, 24.81667, 2400–3100 m, CANCAP-V Expedition.

**OUTER MORPHOLOGY AND SKELETON ORGANIZATION (FIGURE 7)**

Specimens are massive spherical sponges with a smooth regular surface and a brown colour (in ethanol). 105638 has an atypical oblongated shape. 105652 is attached to G. atlantica (105633). As in G. hentscheli, uniporal oscules are concentrated in the preoscule; every oscule is surrounded by ridges which can be very developed. The cribitoporal pores are scattered over the rest of the body. The sterraster layer in 105645 is 0.8–1 mm thick with a poorly developed ectocortex (45 μm thick). Light brown granulated cells (or vacuoles) are moderately present throughout the cortex as well as below it; they are 10–20 μm in diameter.

**BATHYMETRIC RANGE**

130–2534 m (Cárdenas et al., 2013; this study).

**DISCUSSION**

We identified the MAR-Eco specimen as *G. hentscheli* and not *G. barretti* because it has no oxeas I, sterrasters with ‘cauliflower’ surface (Figure 6D, G), and many promesotriaenes. But this specimen is also slightly different from more northern specimens. The observed brown cells were not found in comparative material. The MAR-Eco specimen has elongated sterrasters which is unusual compared with more northern specimens. There are no irregular sterrasters, commonly found in Arctic specimens. The sterrasters are also fairly large for this species, but similar sizes were found in Davis Strait specimens. The microxeas are thinner: average of 5.5 μm averages of 8–16 μm in comparative material from the NEA and NWA (Cárdenas et al., 2013; Table 4). Finally, the hook-like spines on the strongylasters had never been observed before (spines in the paratype ZMB Por 7551 strongylasters are simple), but SEM observations of more specimens is required to see how significant this character is. We reexamined thick sections made on comparative material (paratype ZMB Por 7551, Iceland specimen from Ingolf Expdt. Station 78, ZMBN 77925 from the Greenland Sea) to look for crystalline structures. Only ZMBN 77925 had such structures, but with a different morphology: smaller (32–50 μm), less confused, with fewer fibres and often with two opposite fans (unlike the MAR-Eco specimen, fixed in formalin, ZMBN 77925 had been directly fixed in ethanol 96%).

**Fig. 5.** *Geodia hentscheli* Cárdenas et al., 2010, ZMBN 105680.
found but these are usually found in the fur of *G. hentscheli* and here no fur was observed, maybe due to the trawling collecting method. The sterrasters are spherical, never irregular or with a ‘cauliflower’ aspect (a common feature in *G. hentscheli*). The dark brown colour seems to be the rule for this southern population whereas *G. hentscheli* is more often whitish or greyish; brown cells were observed only in the sections of *G. hentscheli* (105680), never in the *G.

Fig. 6. Spicules of *Geodia hentscheli* Cárdenas et al., 2010, ZMBN 105680 and *Geodia cf. hentscheli,* specimen ZMBN 105645: (A) Dichotriaene (*G. hentscheli*); (B) Protriaene (*G. hentscheli*); (C) Dichotriaenes, small orthotriaene and sterrasters (*G. cf. hentscheli*); (D, E) Close-up of sterrasters showing the rosettes (*G. hentscheli*); (F) Close-up of sterrasters showing the rosettes (*G. cf. hentscheli*); (G) sterrasters (*G. hentscheli*); (H) sterraster (*G. cf. hentscheli*); (I) Strongylasters (*G. hentscheli*); (J) Oxyster (*G. hentscheli*); (K) Strongylasters (*G. cf. hentscheli*); (L) Oxysters (*G. cf. hentscheli*).
hentscheli specimens revised by Cárdenas et al. (2013). No budding was observed (vs occasional budding in G. hentscheli) but again, we may have seen too few specimens. The oxyasters are also fairly large but these sizes are also found in G. hentscheli (Cárdenas et al., 2013). Likewise for the thicker cortex (1 mm vs 0.5 mm usually), it has been found in some G. hentscheli from Davis Strait (Cárdenas et al., 2013). SEM observations of the strongylasters show that they also have hook-like spines, such as the ones observed in 105680. All in all, this southern morphotype of G. hentscheli may represent a separate southern species, which diverged from its Arctic counterpart. But they seem closer morphologically to the MAR-Eco G. hentcheli 105680. Instead of creating a new species, we prefer to wait for genetic data to take taxonomic action. We have found two other specimens belonging to this southern morphotype in the Bergen Museum (ZMBN 25668) and in Naturalis (RMNH 1458): both were collected in the Azores area at 2400 m depth.

Geodia macandrewii Bowerbank, 1858
(Figures 2D, 8 & 9)

MATERIAL
Three specimens from Station 70–385, 1650 m: ZMBN 105661, 105666, 105669.

OUTER MORPHOLOGY AND SKELETON ORGANIZATION (FIGURE 8A, B)
Massive subspherical, whitish specimens with smooth surfaces. Specimens 105661, 105666 and 105669 are respectively around 12/4/1.5 cm in diameter. Cortex of 105661 is 1 mm thick. Thick sections of 105661 were made, skeleton organization is similar to that observed in other specimens (Cárdenas et al., 2013) except that anatriaenes are more abundant below the cortex, associated with the orthotriaene bundles. Some sub-circular crystalline structures were observed, about 130–240 μm in diameter with radial fibrous organization (very ‘bushy’ and confused appearance), and unclear borders. They are similar but larger than the ones observed in G. barretti, and are present below the cortex and in the choanosome.

SPICULES, ZMBN 105661 (FIGURES 8G, G & 9)
Megascleres: (a) oxeas I, straight or bent, length: 4000–5280–7000 μm (N = 10); width: 25–48.7–60 μm (N = 10). (b) oxeas II (=microxeas), straight or slightly bent, not centrotylote, length: 250–335.5–485 μm; width: 3.5–8.4–12.5 μm. (c) orthotriaenes, rhabdome length: 1100–4762.5–6550 μm (N = 10); width: 20–82.8–105 μm; clad length: 300–739–1042 μm. (d) anatriaenes, very common, with sometimes a swelling or spine on top of the cladome (22–175 μm long), rhabdome length >13500 μm; width: 3–19.7–40 μm; clad length: 15–137.5–280 μm. (e) promesotriaene, rhabdome length >5700 μm; width: 10–32–45 μm (N = 10); clad length: 47–237.4–500 μm (N = 9); central clad length: 55–152.2–240 (N = 4). Microscleres: (f) sterrasters, spherical, warty rosettes, 125–155.2–180 μm in diameter. (g) spheroxyasters, rough essentially at the tip of actines, 5–10–15 μm in diameter. (h) oxeasters, rough essentially at the tip of actines, diameter: 10–22.4–47 μm.

BATHYMETRIC RANGE
157–1900 m (Cárdenas et al., 2013).
Fig. 8. *Geodia macandrewii* Bowerbank, 1858: (A) ZMBN 105661; (B) Thick section (ZMBN 105661), notice the bundles of orthotriaenes and anatriaenes under the cortex; (C–G) SEM pictures of spicules (ZMBN 105666): (C) Oxea I; (D) Oxea II (*¼ microxea*); (E) Anatriaene; (F) Prototriaene; (G) Orthotriaene. (C–F: same scale).

Fig. 9. Microscleres of *Geodia macandrewii* Bowerbank, 1858, ZMBN 105666: (A) Sterraster with fully developed warty rosettes; (B) Sterraster with developing rosettes; (C) Oxyasters; (D) Close-up of the sterraster in A showing the fully developed warty rosettes; (E) Close-up of the sterraster in B showing the hilum and the developing rosettes; (F) Spheroxyasters.
The large size of the oxyasters is typical of individuals living deeper than 1000 m (Cárdenas et al., 2013). The sterrasters measured in 105661 are the smallest sized ones ever found for this species and specimens 105666 and 105669 also had sterrasters in similar size ranges: *G. macandrewii* sterrasters are usually more than 200 mm in diameter (Cárdenas et al., 2013). We reexamined thick sections of ZMBN 77924 (*G. macandrewii* from Korsfjord, Norway, fixed in ethanol 70%) but could not find any crystalline structures.

**DISCUSSION**

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*Geodia megastrella* Carter, 1876

(fig. 10)

**MATERIAL**

One specimen from Station 50–373, 2600 m: ZMBN 105646.

**OUTER MORPHOLOGY AND SKELETON ORGANIZATION (FIGURE 10A, B)**

Subspherical specimen, 2 cm in diameter. Colour in ethanol is brown. A single small prediscule on the top surface, cribriporal pores all over the rest of the surface. The cortex is 1.2–1.7 mm thick, very tough, and supported by triaenes and oxeas which form a 3.5 mm thick layer; below, oxeas occur in confusion. Oxyasters I can be found in high abundance in the choanosome. Microxeas can be found in the ectocortex and the choanosome. Large granulated cells about 25 μm in size can be found in the upper part of the endocortex, they contain many brown vacuoles inside. Some crystalline structures were observed in the choanosome, about 30–87 μm in diameter with clear radial fibrous organization and ‘hairy’ border. They are of similar size and shape to the ones observed in *G. hentscheli* (105680) but the fibres seem to be thinner and less confused.

**SPICULES, ZMBN 105646 (FIGURE 10C–G)**

**Megascleres**: (a) oxeas I, bent, some slightly centrotylote, length: 840–1671.9–2750 μm (N = 13); width: 15–26.3–35 μm (N = 13). (b) oxeas II (=microxeas), straight, a few are centrotylote, length: 195–290.2–350 μm; width: 4–7.6–11 μm. (c) orthotriaenes, and very few dichotriaenes, rhabdome length: 1750–2193.7–3000 μm (N = 4); width: 40–55.3–70 μm (N = 14); protoclad length: 120 μm (N = 1); deuteroclad length: 220 μm (N = 1). (d) promesotriaene, only one found, rhabdome width: 20 μm; clad: 110 μm; central clad: 300 μm. **Microscleres**: (e) sterrasters, spherical, 150–163.5–190 μm in diameter, hilum ~10 μm in diameter,
warty rosettes. (f) strongylasters, rough actines, 5.5–8.5–12 μm in diameter. (g) oxyasters I, 3–8 rough actines, diameter: 47–93.3–150 μm. (h) oxyasters II, 7–9 thin rough actines, diameter: 19–28–42 μm.

**BATHYMETRIC RANGE**

200–2600 m (Topsent, 1911; this study).

**DISCUSSION**

The external morphology of the MAR-Eco specimen from Station 50 (north of the Azores) is very similar to the specimen described and illustrated by Topsent (1928) from Madeira at 2380 m, and the specimen identified as 'Sidonops sp.' by Arnesen (1920) from the Ibero-Moroccan Gulf at 1215 m (ZMBN 25652, re-examined for this study). A still photograph from a video by the manned-submersible MIR 1 above 1700 m depth in the CGFZ (Felley et al., 2008, Supplementary material) shows globular sponges with a single preoscule that look very much like *G. megastrella*, so this species may also be present in the CGFZ (dive coordinates 52°58′N 35°01′W). No distribution map was made for this species found between Ireland and the Azores in the NEA and between the New England seamounts and Florida in the NWA (P. Cárdenas, unpublished results) since we suspect it to be a species complex (Cárdenas et al., 2011) which needs to be properly revised before anything can be said about its biogeography. This is the first time that microxeas are observed to be occasionally centrotylote in *G. megastrella*. Occasional centrotylote microxeas may be a synapomorphy of the species belonging to the Depressiogeodia clade (*G. barretti, G. hentscheli* and *G. megastrella* complex) even though it has also been observed in *G. macandrewii*, albeit more rarely (Cárdenas et al., 2013).

*Geodia nodastrella* Carter, 1876

(Figures 2E & 11, Table 2)

**MATERIAL**

One specimen from Station 50–373, 2600 m: ZMBN 105644.

**COMPARATIVE MATERIAL EXAMINED**

*Geodia nodastrella*, MOM-INV-22575 (04-0118) (wet specimen) and MNHN-DT846 (slide), Station 198 (1888), 800 m, 38.44028, −30.98611, original det. as *G. barretti* var. *nodastrella* (Topsent, 1892, p. 48); MNHN-DT845, two slides for two different specimens from Station 882 (one of the slides has written ‘spécimen monstrueux’ on it because the spicules...
are atypical, therefore the identification of this specimen remains doubtful), 38.06111, –28.57917, 98 m, original det. as *G. eosaster* (Topsent, 1904, p. 67); MNHN-DT84 (slide), Station 899, 37.95, –29.42583, 200 m, original det. as *G. eosaster* (Topsent, 1904); MNHN-DT90, off Morocco, Station 1116, 31.725, –10.77917, 2165 m, det. by Topsent (1928); MNHN-DT1259, West of Flores, Azores, Station 2210, 39.41667, –31.375, 1229 m, det. by Topsent (1928); MNHN Balgim campaign (Boury-Esnault et al., 1994), CP63–E2 (three specimens), 35.50167, –7.70928, 1510 m, CP98–E41, 34.48681, –7.68536, 1747 m, ZMBN 25660 (two specimens), Station 24, 35.56667, –7.58333, 1615 m, original det. as 'Sidonops sp.' (Arnesen, 1920), PC67–679, AVILES 0511, DRO305011, 43.91862, –5.7658, 799 m. BANGAL 0710, DRO610810, 42.66298, –11.94938, 920 m; DRO3090810, 42.7271, –11.83727, 797 m; DRO1080810, 42.6811, –11.61076, 999 m; DRO7120810, 42.81035, –11.72252, 999 m. ZMAMOR 20372, Mingulay Reef, field#D63/20, 56.80663, –7.43173, 151–159 m, det. by P. Cárdenas. UPSZMC-BIO, NEREIDA 2009–2016; UPSZMC 78295, DR7, 48.25610, –44.03084, 1339 m; DR18, 47.26186, –43.53101, 1079 m; DR19, 47.16431, –43.476328, 1137 m; DR20, 47.07224, –43.44914, 1122 m; DR21, 46.84606, –43.71764, 870 m; DR22, 46.84001, –43.64039, 956 m; DR23, 46.77485, –43.86510, 1127 m; DR24, 46.69422, –43.96859, 1104 m; UPSZMC 78296, DR64, 46.42378, –44.85475, 1091 m; UPSZMC 78294, DR70, 45.88667, –45.56916, 916 m. Geodia divisoria, holotype, MOM–INV 22553 (04–1333) (wet specimen) and MNHN DT–1299 (slide), Madeira, Station 3113, 32.57917, –17.09167, 1700 m. **OUTER MORPHOLOGY AND SKELETON**

**ORGANIZATION (FIGURE 11A, B)**

Massive spherical specimen (6 cm in diameter), white in ethanol, fairly hispid on one side, not compressible. Regular surface with no obvious large openings but small cribriporal pores and oscules are present. Cortex is 0.5–0.6 mm thick and fairly easy to cut. The positions of the different euasters are clear on the thick sections. Ectocortex is 200–250 μm thick, with sub-radial microxeas, numerous strongylasters (especially packed in the eustome) and few spherasters. The fibrous endocortex is 500–600 μm and packed with sterrasters; it is supported by large dichotriastes. Protriaenes are crossing the cortex with their cladomes beyond the surface. In the choanosome, there are sparse oxyasters. Some irregular crystalline structures were observed in the choanosome, about 87–175 μm in length with radial fibrous organization. They are similar to the ones observed in *G. macandrewii* but more irregular in shape and less ‘dirty’.

**SPICULES, ZMBN 105644 (FIGURE 11C–I)**

**Megascleres** (a) oxeas I, straight or fairly bent, sometimes somewhat veriform, length: 2050–3203–4100 μm (N = 20); width: 25–32.7–38 μm (N = 20). (b) oxeas II (microxeas), straight or slightly bent, not centrotolyte, length: 260–370.7–500 μm; width: 4–6.2–8 μm. (c) dichotriaste, straight rhombast, deuteroclads are often curved, rhombast length: 2500–3802.5–4750 μm; width: 45–73.4–82 μm; protoclad length: 100–121.3–170 μm; deuteroclad length: 100–258.8–400 μm. (d) anastriaenes, rhombast length >8500 μm; width: 3–160–35 μm; clad length: 15–137.5–280 μm. (e) protriaenes and promesotriaenes, rhombast length >6500 μm; width:
14–32.7–45 μm (N = 23); clad length: 150–296.3–430 μm (N = 23); central clad length: 160–232.8–310 (N = 7). Microscleres: (i) spherasters, spherical, rosettes with warts, 68–77.1–83 μm in diameter. (g) strongylasters, spiny, 6–9.4–15 μm in diameter. (h) spherasters, blunt to truncated actines with spiny tips, 20–25.7–40 μm in diameter. (i) oxyasters, usually with many thin actines, spiny, 10–18.3–25 μm in diameter.

**BATHYMETRIC RANGE**
98–2600 m (Topsent, 1904; this study).

**DISCUSSION**
This study is an opportunity to revise the morphology of this poorly known, albeit fairly common, Lusitanian species. Many specimens photographed just after collection during the NEREIDA campaign (courtesy of F. J. Murillo) and the BANGAL cruise (courtesy of P. Rios) show that the natural external colour of this species is whitish to light brown. This species always has dichotriaenae, which often have ‘wavy’ deuteroclads. As previously observed by Topsent (1904) and Stephens (1915), the easasters have variable morphologies. The ectocortical strongylasters often have an inflated centrum (spherostrongylaster) and are then difficult to separate from the smallest spherasters; in other specimens they also sometimes have long actines with inflated tips as in tylasters (MNHN-DT1090, NEREIDA DR07-043b). The spherasters are most often quite spherical but, sometimes, they are less inflated; their actine tips are also variable with sometimes blunt truncated tips, and sometimes sharp conical tips. As for the choanosomal oxyasters they sometimes have thin long actines, and sometimes wider conical actines. For instance, in the specimen from the Azores (Station 198) studied by Topsent (1892, 1904) and in the Balgim specimens, we found both morphologies of oxyasters.

Stephens (1915) states that the spherasters are placed just below the cortex but our sections of the MAR-Eco specimen and MOM 04-0118 clearly show that spherasters are present in the ectocortex as well (i.e. above the sterrater layer). Actually, in the MAR-Eco specimen, they are only present in the ectocortex. Topsent (1904) notices that spherasters can become rare. We also observed this, and even found specimens where the characteristic spherasters were missing (e.g. BANGAL PC581, PC579, ZMBN 25660). In these specimens without spherasters, we did find many oxyspherasters but it is unclear if they are deflated spherasters or inflated oxyasters (since spherasters and oxyasters essentially have similar sizes).

The size of the oxyasters is also very variable, they can vary from 20–24 μm (Stephens, 1915) to 30–40 μm (MNHN-DT844) as in the MAR-Eco specimen and even up to 72 μm (BANGAL PC579, ZMBN 25660). The continuum of sizes makes it impossible to delimitate two size categories. The important variation in size of the choanosomal oxyasters has already been observed in all Atlantic boreo-arctic *Geodia*, and may be related to the depth and/or the silica concentration (Cárdenas & Rapp, 2013; Cárdenas et al., 2013). We should stress here that specimens without spherasters and/or large oxyasters were confirmed to be *G. nodastrella* with external morphology, other spicules, as well as with molecular data (P. Cárdenas, unpublished results). We also noticed that the MAR-Eco specimen has spherical spherasters, whereas they can be ellipsoid in the comparative material (e.g. BANGAL PC581, MNHN-DT846). The size of the spherasters varied between 68 and 115 μm which is in the same range as most boreo-arctic *Geodia*, except for the very large spherasters of *G. macandrewii* (Cárdenas et al., 2013). Finally, another variation concerned the microxeas found in the ectocortex: the MAR-Eco specimen has fairly longer microxeas (260–500 μm) than in previous measurements (167–350 μm) (Table 2).

With all these variations in mind, we re-examined the holotype of *Geodia barretti divaricans* from Madeira (MOM-INV-002282, wet specimen and MNHN DT-1299, spicule preparation). The wet specimen is a small hispid fragment attached to coral, it is the only specimen known of this species. We made thick sections from the type and measured easasters (Table 2). It occurred to us that the spicule repertoire was very close to that of *G. nodastrella*. The oxyasters can be found in various sizes, the smaller ones (15–22.5 μm) can be found just below the cortex and the larger ones (27–70 μm) are very numerous throughout the choanosome. Topsent (1928) surprisingly overlooked the small oxyaster sizes, even though we found some on Topsent’s slide (MNHN-DT1299). The small oxyasters with an inflated centrum and spiny actines look similar to the ones we observed in specimens of *G. nodastrella* from Galicia (BANGAL). The length of the microxeas in *G. divaricans* (210–525 μm) are actually closer to the ones from our MAR-Eco specimen (260–500 μm). *Geodia divaricans* was also characterized by inflated rhombomes of the dichotriaenae and rare flattened anatriaeas (Topsent, 1928). However, slight rhambome swellings were also observed in the dichotriaenae of *G. nodastrella* from Galicia and more or less flattened anatriaeas (although not as flattened as in *G. divaricans*) were also observed in the Irish specimens (Stephens, 1915). Finally, the absence of spherasters as discussed previously is possible in *G. nodastrella*. So, in our opinion, no specific cortical or spicule characters really remain to keep the valid status of *G. divaricans*. Therefore, we formally propose that *G. divaricans* Topsent, 1928 is a junior synonym of *G. nodastrella* Carter, 1876. The *G. nodastrella* recorded by Burton (1934) in Greenland has been re-identified as *G. hentscheli* (Cárdenas et al., 2013). We also re-examined the two specimens (ZMBN 25660) identified as ‘Sidonops sp.’ by Arnesen (1920); these were collected quite near the Balgim CJP53 station where *G. nodastrella* was reported. Their external morphology (large spherical sponges with cibarial pores/ osules) clearly matches that of *G. nodastrella* and their spicules match those of *G. divaricans* (no spherasters, very large oxyasters). The other sponge identified as ‘Sidonops sp.’ (ZMBN 25652) is in fact *G. megastrella* (see above).

*Geodia phlegreae* (Sollas, 1880)
(Figures 2F & 12)

**MATERIAL**
Five specimens from Station 72–386, 2534 m: ZMBN 105675-78, 105681.

**OUTER MORPHOLOGY AND SKELETON**
**ORGANIZATION** (FIGURE 12A–E)
Specimens are spherical to subspherical sponges between 1.5 and 6 cm in diameter, with remains of hispidity on the sides, which are purple (possibly coloured by the encrusting sponge *Hexadella dedritifera* Topsent, 1913). 105678 has an Hexactinellida growing on it (Figure 12B). Oscules and pores are uniporal. Thick sections of 105675 were made. Cortex is...
moderately thick: 1.1–1.2 mm. Skeleton organization is in accordance with previous descriptions (Cárdenas et al., 2013). Oxyasters are very abundant in the choanosome. Sub-circular crystalline structures were observed in the choanosome, about 45–112 μm in diameter with radial clear fibrous organization.

**Spicules, ZMBN 105675 (FIGURE 12F–H)**

Megascleres: (a) oxeas, length: 3375–4154.5–4750 μm, width: 60–71.5–85 μm. (b) orthotriænes, rhabdome length: up to 3750 μm; width: 70–82.2–90 μm; orthotriæne clad length: 430–771.4–1032 μm. Microscleres: (c) sterrasters, subspherical, smooth rosettes, length: 90–101.9–116 μm. (d) sphera-sters, spiny, 13–17.7–27 μm in diameter. (e) oxyasters, smooth, 20–42.6–70 μm in diameter.

**Bathymetric Range**

40–3000 m (Cárdenas et al., 2013).

**Discussion**

We identified these specimens as *G. phlegraei* and not *G. parva* — its sister species from the Arctic (Cárdenas et al., 2013) — based on the external morphology (thick cortex, regular smooth surface) and fairly large sterrasters. But we note however that these sterrasters are spherical as in some NWA specimens, and not elongated as in NEA specimens (Cárdenas et al., 2013). We re-examined sections of *G. phlegraei* ZMBN 77929 (Korsfjord, Norway) for crystalline structures, and we found many: small (27–37 μm) dirty sub-circular ones were very abundant below the cortex, larger ones (similar to the ones observed in the MAR-Eco specimen) were observed deeper in the choanosome.

*Stelletta tuberosa* (Topsent, 1892) (FIGURES 13A & 14)

**Note**

We place this *Stelletta* in the Geodiidae based on molecular phylogeny results from Cárdenas et al. (2011). It seems that, like *Stelletta tuberosa*, many species of *Stelletta* are actually *Geodia* species that have lost their sterrasters, they group in a *Geostellettap* clade (named according to the rules of the PhyloCode). Before reallocating these *Stelletta* species in a new Geodiinae genus, we are waiting for more sequences of *Stelletta* species to have a better morphological understanding of this *Geostellettap* clade.
Fig. 13. Distribution maps. T, type locality; CGFZ, Charlie-Gibbs Fracture Zone. Maps generated with GeoMapApp 3.3 (http://geomapapp.org).
MATERIAL

COMPARATIVE MATERIAL EXAMINED
MNHN-DT835 (slide), '1888-122’, det. by Topsent. MNHN-DT1896 (slide), Station 183-1896, det. by Topsent. MNHN-DT3012 (slide), off Galicia, Spain, Station 3012, 42.44167, 29.48333, 1680 m, recorded in Topsent (1928). MNHN-DCL4066, Bay of Biscay, 4400 m, BIOGAS V expedition (Centob), det. by P. Cárdenas. ZMAPOR 21665, South Azores Seamounts, 26°16.098’N 38°09.931’W, 2110 m, EMEPC/G3/08.10, det. by P. Cárdenas. RMNH Por1454, CANCAP-V Expedition, SNMH #474, Swedish Arctic Exp. 1871, Station 37, 53.56667, −52.01667, 1792 m, det. by P. Cárdenas. UPSZMC 78301, Flemish Cap, NEREIDA 2009, field#DR06-38, 48.291, −44.06, 1348 m, det. by P. Cárdenas. UPSZMC 78302, Flemish Cap, NEREIDA 2009, field#DR07-33, 48.256, −44.03, 1339 m, det. by P. Cárdenas. UPSZMC 78269, Flemish Cap, NEREIDA 0609, Station BC89, 48.974, −45.272, 1574 m, det. by P. Cárdenas; ZMUC, Ingolf Expedition, Station 78, Reykjanes Ridge, 60.61667, −27.86667, 1505 m, Station 90, Denmark Strait, 64.75, −29.1, 1070 m, det. by P. Cárdenas.

OUTER MORPHOLOGY AND SKELETON ORGANIZATION (FIGURE 14A–E)
Massive subspherical usually with a single large oscule but specimen 105610 has four. Large conspicuous sphincters can be found around the oscules. Colour in ethanol is brownish. Thick sections were made for 105649 and 105650. The cladoomes of the triaenes and oxyasters (large and small sizes) are embedded in an ectosome about 200–250 μm thick. Below the ectosome there is a lighter area ≏ 500 μm thick with large subcortical canals and where oxyasters are more scarce. Finally, the choanosome is rich in oxyasters. Large oxeas tend to join the rhabdomes of the triaenes thus making thick radial bundles in the choanosome. Small crystalline structures (27–37 μm long), radial star shaped to elongated shape were commonly found in the sections of 105649, only in the sub-ectosomal area and especially in its canals. Many Globigerina (planktonic Foraminifera) can be found in the choanosome.

SPICULES, ZMBN 105649 (FIGURE 14F, G)
Megascleres: (a) large oxeas, length: 4000–4600–5100 μm (N = 5), width: 40–53–65 μm; (b) dichotriaenes (plagiotriaenes are rare), massive, rhabdome was usually straight but sometimes bent, it often had a distinct swelling below the cladome, rhabdome length: 1525–2552–3800 μm (N = 8), width: 70–115.5–145 μm, protoclad: 80–137.9–
250 μm, deuteroclad: 100–289.5–390 μm. Microscleres: (c) oxyasters, spiny, diameter: 16–34.1–57 μm.

BATHYMETRIC RANGE
454–4400 m (Topsent, 1892; this study).

DISCUSSION
This is the Stelletta sp. recorded by Murillo et al. (2012) off Newfoundland (F. J. Murillo, personal communication). We have also identified this species in the NEREIDA 2009 material collected in the same area (DR06, DR07, BC89). Larger oxyasters were measured in the other specimens, 105611 had oxyasters sizes up to 66 μm while 105679 had oxyasters sizes up to 68 μm. In the specimens described by Topsent (1892, 1904, 1928) from the Azores, and some we examined from Newfoundland, the actines of oxyasters usually have a tiny inflated tip. The MAR-Eco specimens, the Ingolf Expedition specimens, some Newfoundland specimens (UPSZMC 78269) and the specimen from Bay of Biscay (MNHN-DCL4066) and the one of the South Azores Seamounts (ZMAPOR 21665) had larger oxyasters (up to 96 μm) with thinner and pointier actines. The Rhabdomes of Newfoundland specimens are swollen (UPSZMC 78302) (although not as much as in the southern MAR-Eco specimens) or not (#474, UPSZMC 78301, 78269). So at this point, it is important to emphasize that the inflated tip of actines and the swelling of rhabdomes are not diagnostic characters of this species, they may be absent. Anatriaenes (in 105670) and pro/mesotriaenes (in 105679) were observed. The Bay of Biscay specimen (MNHN-DCL4066) and the one of the South Azores Seamounts (ZMAPOR 21665) had larger oxyasters (up to 96 μm) with thinner and pointier actines.

Family ANCORINIDAE Schmidt, 1870
Genus Stelletta Schmidt, 1862
Stelletta rhaphidiophora Hentschel, 1929
(Figures 13B & 15)

MATERIAL
Two specimens from Station 70–385, 1650 m: ZMBN 105664, 105668. One specimen from Station 72–386, 2548 m: ZMBN 105682.

COMPARATIVE MATERIAL EXAMINED
Stelletta rhaphidiophora, ZMBN 85222, Iceland, 67.71972, −19.9, BIOICE 3661, 604 m; ZMBN 85223, H2Deep 2008

**OUTER MORPHOLOGY AND SKELETON**

**ARRANGEMENT** *(Figure 15A, B)*

Massive subspherical, very hispid. Diameter of the specimens range between 6 cm (105664), 2 cm (105668) and 4 cm (105682). A small *Geodia barretti* (106565) is growing on 105664 *(Figure 15A)*. Surface colour in ethanol is whitish (sometimes dirty brown due to the sediments trapped in the hispid layer), choanosome colour is light brown. Oscules and pores not visible. Thick sections of 105682 were made. Cortex is 2–2.5 mm thick. Vitreous cortex, light greyish:

- **(a)** large oxeas, most were broken, up to 6500 μm in length, 80 μm in width; **(b)** massive triaenes with more or less forward oriented clads, large triaenes look like protiaenes, clads are usually divided in two but not always, rhabdome was straight or usually bent, rhabdome length: 1750–4010–6000 μm (N = 15), width: 45–83.9–125 μm, protoclad: 60–149.2–250 μm, deuteroclad: 100–305.2–480 μm, clad (not divided): 780–1500 μm; **(c)** small plagiotriaenes, usually found in the lower portion of bundles, they may be young triaenes or a separate category, rhabdome length: 800–900–1100 μm (N = 4), width: 20–26.2–30 μm (N = 4), clad: 140–182.5–270 μm (N = 4). **Microscleres:**

**BATHYMETRIC RANGE**

241–2548 m *(Klitgaard & Tendal, 2004; this study).*

**DISCUSSION**

In the original description of *S. rhaphidiophora*, there are two categories of anatriaenes, characteristic flattened ones and more usual ones *(Hentschel, 1929).* Although we did not find the flattened anatriaenes in any of our specimens, we are sure of our identification since the rest of the spicule morphologies and measurements perfectly match the original description and our comparative material. We did not find the common anatriaenes in 105682 but we found two in 105664: rhabdome length >4100 μm, width: 30 μm, clad: 130–160 μm. This is the first time SEM observations are made for this species. It shows that the strongylasters have spined actines *(Figure 15F)*, which are not visible with the optical microscope, and thus not reported in the original description. As observed in the boreo-arctic *Geodia* species *(Cárdenas et al., 2013)*, there is some variation in the maximum size choanosomal oxyasters can reach. Type material has oxyasters that reach 40 μm *(Hentschel, 1929)*, but in the MAR-Eco specimens the oxyasters only reach 27.5 μm. In ZMBN 85222 from Iceland *(Hentschel, 1929)* and UPSZMC 78297–78298 from Davis Strait *(~850 m)*, oxyasters respectively reach 65 and 56 μm and have an inflated centre. On the other hand, specimens from the Greenland Sea, near the Schultz Massive seamount and the Arctic Mid-Atlantic ridge *(ZMBN 85223, 1600–1760 m)* have smaller oxyasters *(up to 25 μm)* like in the MAR-Eco specimens; but they are in very low numbers, unlike the MAR-Eco specimens. ZMBN 85223 also has very few trichodragmas, rare and small triaenes. It is interesting to note that *Geodia hentschelii* *(ZMBN 77925)* collected from the same locality at similar depth has a similar phenotype: low number of oxyasters, rare and smaller triaenes *(Cárdenas et al., 2013, Figure 12D, Table 3)* so in our opinion environmental parameters around the Schultz Massive seamount clearly influence these spicule morphologies, abundances and sizes.

The MAR-Eco specimens considerably extend the range of this Arctic species southwards. Furthermore, our comparative material from the Davis Strait extends its range to the West. *Stelletta rhaphidiophora* is an amphi-Atlantic Arctic species while its sister-species, *Stelletta normani* Sollas, 1880 is typically boreal *(P. Cárdenas, unpublished data). In the field, these species can usually be distinguished by looking at their colour and a cross-section in their cortex with the naked eye. *Stelletta normani* has a clear double-layered cortex while *S. rhaphidiophora* is hairy white. *Stelletta normani* has a clear double-layered cortex with *(i)* a bright white layer *(packed with trichodragmas)* and *(ii)* a vitreous grey layer *(fibrous layer). Stelletta rhaphidiophora* has more of a vitreous grey single-layered cortex since the upper white layer is less obvious and bright *(because it never has as much trichodragmas)*.

**Genus Stryphnus Sollas, 1886**

*Stryphnus foris* *(Vosmaer, 1885)* *(Figures 13C & 16)*

**MATERIAL**

One specimen from Station 50–373, 2600 m: ZMBN 105640.

**COMPARATIVE MATERIAL EXAMINED**

*Stryphnus foris*, ZMAPOR 02189, holotype, wet specimen, Willem Barents Sea Expedition 1880–1881, Barents Sea, 71.91667, 20.51667, 327 m *(schizoholotype, RMNH 59, not seen)*; ZMUC, Ingolf Expedition, Station 78, Reykjanes Ridge, 60.61667, −27.86667, 1505 m; ZMBN 82977, Korsfjord, Norway, 60.161667, 5.174722, 200–400 m; ZMBN 85224, Langenuen, Norway, 59.88133, 5.51667; ZMBN 89724 and 89727, Spitsbergen, 80.534167, 15.351667, 215 m; UPSZMC 78303, NEREIDA 2009, field#DR03-12, Flemish Cap, 48.3873883, −44.200891, 1324 m; UPSZMC 78304, NEREIDA 2009, field#DR15-31, Flemish Cap, 47.628715,
243.53382667, 1358 m; UPSZMC 153297, Ramso¨, Kosterfjorden, Sweden, ~200 m, original det. by H. Alander in 1935 as S. ponderosus var. rudis; field#BDV1132, Celtic Explorer 13008, off Western Ireland, Porcupine Bank, ROV 15, 54.056, −12.5548, 1401 m; PC794, field#BDV1514, Celtic Explorer 13008, off Western Ireland, Globan Spur, ROV 45, 49.0101, −10.9449, 654 m. Stryphnus ponderosus, MC3395, Rathlin Island, 55.30797, −6.26878, 29 m.

OUTER MORPHOLOGY AND SKELETON ORGANIZATION (FIGURE 16A, B)
The single irregular specimen about 7 cm wide ‘glues’ together three G. atlantica (105639, 105641, 105642) and one Stelletta tuberosa (105648) (Figures 3A & 16A). Colour is dark brown in ethanol. Surface is rough. Radial organization of the skeleton at the surface, more confused in the choanosome. The cladomes of short-shafted dichotriaenes and plagiotriaenes reach a thin fibrous ectosome and sometimes slightly cross it. We could not find sanidasters in the ectosome. Below this ectosome and between the bundles of triaene rhabdomes there are many subectosomal canals. The ectosome and the subectosomal canals make a light layer 1 mm thick. A few anatriaenes and mesoanatriaenes were also occasionally found in this layer but usually crossed the ectosome (on sections #1 and #2). In the choanosome, there is a dense accumulation of oxyasters and oxeas in a confused arrangement. Large granular cells are present in the ectosome but are not obvious (they are not coloured).

SPICULES, ZMBN 105640 (FIGURE 16C–G)
Megascleres: (a) oxeas, slightly bent, length: 1575–2118–2750 µm, width: 22–37.5–55 µm. (b) dichotriaenes, rhabdome is straight and relatively short, clads are sometimes slightly forward oriented, clads can be irregular with trifurcations, rhabdome length: 390–897–1250 µm, width: 25–59.1–70 µm, protoclad: 90–148.8–270 µm, deuteroclad: 110–227.8–300 µm. (c) plagiotriaenes, their rhabdomes are similar to those of the dichotriaenes, clad: 300–425–580 µm (N = 10). (d) anatriaenes, rare, with widely open clads, rhabdome width: 30–39.1–45 µm (N = 6), clads: 200–246.7–300 µm (N = 8). (e) mesoanatriaenes, rare, with widely open clads and an additional clad on top of the rhabdome, rhabdome width: 15–21.3–27 µm (N = 3), clads: 115–183.3–235 µm (N = 3), central clad length: 105–160 µm (N = 2).

Microscleres: (e) spiny oxyasters, with sharp actines, none are reduced to two actines, smaller sizes have more actines and a large centrum, diameter: 17–41–67 µm; (f) spiny sanidasters to amphisanidasters, length: 10–11.8–15 µm.

Fig. 16. Stryphnus fortis (Vosmaer, 1885), ZMBN 105640: (A) Specimen overgrowing Geodia atlantica, close up of Figure 3A; (B) Thick section; (C) Oxea; (D) Plagiotriaenes and oxyaster (ox); (E) Dichotriaene; (F) Oxyasters and amphisanidasters; (G) Amphisanidasters.
BATHYMETRIC RANGE
157–2600 m (Klitgaard & Tendal, 2004; this study).

DISCUSSION
Historically, the microscleres in *Stryphnos fortis* were called ‘amphistyles’ (Sollas, 1888) because they are somewhat symmetrical but we prefer to use ‘sanidaster’ and ‘amphisanidaster’ (the latter term was coined by Kelly & Smith (2012)) to emphasize that they are homologous to the ones found in *Asteropus* and *Ancorina*.

The morphology of the MAR-Eco specimen broadly agrees with the description of *S. fortis* but a few differences with the comparative material (including the type) were noted. The sanidasters in the type are slightly longer (12–12.8–15, N = 10) and the proportion of amphisanidasters (vs sanidasters) is higher; indeed, the sanidasters in the MAR-Eco specimen often have additional actines on the shaft. We confirm that the type has only plagiotriaenes, as noted by Vosmaer (1885). Although most of the *S. fortis* comparative material we examined (from Norway, Sweden and Flemish Cap) had both plagiotriaenes and dichotriaenes, it was always with a majority of plagiotriaenes. Our specimen has on the contrary a higher proportion of dichotriaenes, which seems to be a characteristic of the southern population of *S. fortis* (Topsent, 1904; Boury-Esnault et al., 1994). Vosmaer (1885) states in his original description that oxyasters come in two sizes. However, this appears to be incorrect as we found only one size when we examined the holotype (ZMAPOR 02189). The oxyasters in the type are fairly large (42–60.9–75 μm, N = 10) compared with the ones observed in the MAR-Eco specimen (~17–30 μm), with many actines and a large centrum. More unexpected is the presence of ana-triaenes, a spicule never observed before in this species, and usually never found in the genus *Stryphnos*; anastaenae have only been found in one atypical New Zealand species rightly called *Stryphnos atypicus* Kelly & Smith (2012). Finally, the cortical arrangement in the type and the Norwegian specimen ZMBN 82977 is much more confused than in the MAR-Eco specimen. In the type and ZMBN 82977 some triaenes are somewhat radially disposed but they usually cross the ecosome; many other triaenes can be found in all other directions, including in the choanosome. To conclude, the MAR-Eco specimen is slightly different from typical boreal *S. fortis* but the examination of more southern specimens is required to see if these differences are consistent with a separate population or even species.

According to our observations and comparison with specimen Mc3395 of *S. ponderosus*, we confirm that *S. fortis* is a valid species, and not a synonym of *S. ponderosus* as suggested by some (Burton, 1926; Koltun, 1966). *Stryphnos ponderosus* has smaller oxyasters (usually never larger than 25 μm in diameter), it is a shallower species (0–200 m) that lives in temperate waters of the North-East Atlantic (British Isles, Ireland, France, Spain) and Mediterranean Sea, it is often covered by the sponge *Desmacella annexa* Schmidt, 1870 (Topsent, 1928, Lévi, 1950), which is the case of Mc3395. On the other hand, *S. fortis* is a deep-sea amphi-Atlantic species (200–2598 m) found from the boreo-arctic region to the Azores, it is often covered by the deep-sea yellow sponge *Hexadella detrifera* Topsent, 1913. Since *S. ponderosus* and *S. fortis* mainly differ by the size of the oxyasters and mega-scleres, we wonder whether they could be conspecific, their spicule differences being a direct consequence of depth and silica concentration in the environment, as shown in other Astrophorina (Cárdenas & Rapp, 2013). However, molecular data shows that the two species are genetically different (Cárdenas et al., 2011), there is a 2 bp. difference in the Folmer COI fragment, and 1 bp. in the 28S (C1–D3) fragment. So, in our opinion, *S. fortis* and *S. ponderosus* are sister-species that diverged recently from a common ancestor that colonized shallow waters from the deep-sea (or the reverse). This new environment may have influenced the size of the oxyasters. Today the shallow and deep-sea populations may not be genetically connected anymore (this should be tested with a wider sampling) and have retained their respective morphologies.

After examining material collected during the deep-sea NEREIDA 2009 campaign off Newfoundland, we concluded that the *S. ponderosus* recorded off Newfoundland is in fact *S. fortis* (Murillo et al., 2012; Kenchington et al., 2013; Kutt et al., 2013). For example, UPSZMC 78303 from the Flemish Cap has oxyasters which are 30–44.5–75 μm (N = 10) in diameter. Likewise, *S. ponderosus* recorded from the Atlantic boreo-arctic region, as a major component of boreal sponge grounds, is in fact *S. fortis* (Hougaard et al., 1991; Klitgaard, 1995; Klitgaard & Tendal, 2004).

Vosmaer (1885) gave no measurements for the spicules of *S. fortis*, so it was originally difficult to identify. *Stryphnos rudis* Sollas, 1888 was described as a new species based on the fact that it had dichotriaenes and plagiotriaenes (vs only plagiotriaenes in the type of *S. fortis*). It was collected in the Korsfjord in Norway, where we have collected extensively. The *Stryphnos* specimens that we found there were similar to the paratype of *S. fortis* so we confirm, as suggested by Topsent (1904), that *S. rudis* is a junior synonym of *S. fortis*. Likewise, it was initially thought that *S. ponderosus* only had dichotriaenes so when specimens were collected with dichotriaenes and plagiotriaenes, they were referred to as *S. ponderosus* var. *rudis* Sollas, 1888 (Topsent, 1894, Lévi, 1950; Uriz, 1981). We now know that the proportion of dicho/plagiotriaenes can vary in *S. ponderosus*, so *S. ponderosus* var. *rudis* becomes a synonym of *S. ponderosus* (except for the records of Alander (1942) in Sweden which belong to *S. fortis*).

Family VULCANELLIDAE Cárdenas et al., 2011
Genus Poecillastra Sollas, 1888
*Poecillastra compressa* (Bowerbank, 1866) (Figures 13D & 17)

MATERIAL
One specimen from Station 65–382, 753 m: ZMBN 105660.

OUTER MORPHOLOGY AND SKELETON
ORGANIZATION (FIGURE 17A)
Irregular lamellar shape, 5 cm long, about 1 cm thick. White specimen in ethanol. Skeleton organization is in complete accordance with the description of Norwegian specimens in Cárdenas & Rapp (2012).

SPIECULES, ZMBN 105660 (FIGURE 17B–F)
*Mesicles* (a) oxeas I, stout, smooth, usually slightly curved, but rarely double-curved, a few modified to styles, or even to strongyles, length: 300–1234.7–2600 μm; width: 3–
18.1–27 μm. (b) oxeas II, thin and pointy, length: 1950–2033–2150 μm (N = 3); width: 6.5–7–7 μm (N = 4). (c) short-shafted orthotriaenes, sometimes pseudocalthrops (rhabdome = clads), often irregular clads, rhabdome/clad length: 91–199.7–357 μm; rhabdome/clad width: 18 μm (N = 1). Microscleres: (d) microxeas, in high numbers, microspiny, straight or bent, centrotyle or not, length: 95–165.8–201 μm; width: 5.7–7.5 μm. (e) spiraster to metaster (frequent), microspiny and fairly fat actines, length: 13–18.2–25 μm; width: 10–14.8–21 μm. (f) plesiaster, 3–6 actines, spiny actines, diameter: 24–37.3–57.5 μm.

**BATHYMETRIC RANGE**

0–1740 m (Topsent, 1928, Sarà, 1964).

**DISCUSSION**

The presence of strongyles in our specimen is intriguing; they have never been recorded before in this species (Cárdenas & Rapp, 2012), the rest of the spicules agree well with those of *P. compressa*. More specimens and molecular data are needed to eventually test the status of these Mid-Atlantic ridge populations. Despite numerous records, this species has never been recorded to this day beyond the MAR, in the NWA.

**MATERIAL**

Six specimens from Station 53–375, Faraday seamount, 990 m: ZMBN 105653–57, 105659.

**COMPARATIVE MATERIAL EXAMINED**

*Thenea levis*, UPSZMC 155199, Flemish Cap, NEREIDA 2009, field#DR06-046B, 48.291, −44.06, 1348 m, det. by P. Cárdenas. UPSZMC 155198, Flemish Cap, NEREIDA 2009, field#DR12-062, 47.77, −43.57, 1462 m, det. by P. Cárdenas. UPSZMC 155197, Flemish Cap, NEREIDA 2009, field#DR18-039, 47.26, −43.53, 1079 m, det. by P. Cárdenas. UPSZMC 78289, PA2010-009, set. 155, West Greenland, 66.85, −58.45, 847 m. Additional specimens from the Norwegian coast (Cárdenas & Rapp, 2012).

**OUTER MORPHOLOGY (FIGURE 18A)**

105653–105654 are the only large specimens with the typical elongated shape, they are respectively 5 and 3 cm long. The
other specimens are small, subspherical and possible fragments.

**SPICULES, ZMBN 105653 (FIGURE 18B–F)**

*Megascleres*: (a) oxeas. (b) dichotriaenes, deutoclads are sometimes bent towards each other. (c) common anatriaenes. *Microscleres*: (d) plesiasters, 2–6 minutely spined actines, common, length of actine: 24–63.2–116 μm; actine thickness: 8–20 μm. (e) spined spirasters to metasters, common, length: 16–20.5–24 μm.

**BATHYMETRIC RANGE**

190–1480 m (Steenstrup & Tendal, 1982; Cárdenas & Rapp, 2012).

**DISCUSSION**

No protriaenes were observed. In 105654 plesiasters are sometimes reduced to two actines, a feature not observed in Norwegian specimens but fairly common in our NWA specimens (UPSZMC 78289, 155197, 155199). Plesiasters of the MAR-Eco specimens are smaller than the ones measured in a shallower Norwegian specimen (actine length: 30–83.2–145 μm, ZMBN 85230, off Korsfjord, 300 m). We are currently missing spicule characters to properly discriminate *T. levis* from the other common North Atlantic *Thenea*. The SEM observations of the MAR-Eco specimens confirmed that large plesiasters were more minutely spined in *T. levis* (Cárdenas & Rapp, 2012) but this character is impossible to see without a SEM. We made two new observations: (1) the plesiaster actines in *T. levis* are ‘fat’ or bullet-shaped (in the MAR-Eco and comparative material), which is not the case of the plesiasters in *T. muricata/schmidtii/valdiviae*; (2) anatriaenes in *T. levis* are on average more ‘open’ and with shorter clads than in *T. muricata/schmidtii/valdiviae*. If confirmed these two spicule characters may be used to discriminate *T. levis* from the other North Atlantic *Thenea*. We extend the western distribution of *T. levis* by reporting its first presence off Newfoundland (Flemish Cap) at depths of 1079 m to 1462 m.

*Thenea cf. schmidtii* Sollas, 1886 (Figures 13F & 19)

**MATERIAL**

Two specimens from 42–368, 2078 m: ZMBN 105618-19. Six specimens from Station 44–369, 1742 m: ZMBN 105620-25. One specimen from Station 70–385, 1650 m: ZMBN 105671.

**OUTER MORPHOLOGY AND SKELETON ORGANIZATION (FIGURE 19A–C)**

Subspherical, fairly hispid sponges, 1.5–3 cm in diameter. Cribriporal oscules in the equatorial area and single naked oscule on the top surface. Thick sections of 105621 and 105671 were made: the skeleton organization is highly similar in both specimens. Large plesiasters are very abundant and uniformly distributed.

**SPICULES, ZMBN 105621 AND 105671 (FIGURE 19D–K)**

*Megascleres*: (a) oxeas. (b) dichotriaenes with pointy bent rhabdome. (c) few anatriaenes. (d) many protriaenes found in 105624. *Microscleres*: (e) plesiasters, minutely to fairly spined, very common, 2–7 actines (105621), 4–7 actines (105671), length of actine: 28–89–188 μm (105621), 28–89.8–170 μm (105671) and up to 204 μm (105622); actine thickness: 4–16 μm. (f) metasters to spirasters, spined, common, length: 24–29.9–36 μm (105621), 24–32.2–41 μm (105671).

**BATHYMETRIC RANGE**

349–4020 m (Topsent, 1904).

**Fig. 18. Thenea levis** von Lendenfeld, 1907; ZMBN 105653: (A) Specimen; (B) Dichotriaene; (C) Anatriaene; (D) Large plesiasters; (E) Smaller plesiasters; (F) Spirasters.
DISCUSSION

42–368/44–369 specimens and 70–385 (105671) come from distant stations and the morphology of the spicules are slightly different and we decided to present both in Figure 19D–K. Plesiasters reduced to two actines were not recorded in this species before (Sollas, 1888; Cárdenas & Rapp, 2012) but they are common in the 42–368/44–369 specimens, except for 105671. This is a tentative identification since without genetic data, it is still difficult to decide whether to call this species *T. muricata* or *T. schmidti* (its southern sister-species). Cárdenas & Rapp (2012) have shown that there are genetic differences between the two species (using the C1–D2 fragment of 28S), a result which still needs to be confirmed with additional specimens and independent molecular markers. They have also shown that clear morphological characters are still missing to distinguish both species. We decided to call our specimens *T. cf. schmidti* because the morphology of the MAR-Eco specimens agrees more with the original description of this species (Sollas, 1888). All specimens had abundant large plesiasters. Plesiasters have a much wider range than in *T. valdiviae*. It is difficult to separate the largest metasters/spirasters and the smallest plesiasters, there is a continuum (whereas in *T. levis*, the spiraster category is quite distinct).

**Figure 19. Thenea cf. schmidti** Sollas, 1886, (A–G) ZMBN 105641; (H–K) ZMBN 105671: (A) Side view showing the cribriporal pores in the equatorial area; (B) Top view showing the single naked oscule on the top surface; (C) Thick sections, large plesiasters are very abundant and uniformly distributed; (D) Anatriaene; (E) Plesiasters, one reduced to two actines; (F) Metasters; (G) Spirasters; (H) Anatriaene; (I) Dichotriaene; (J) Plesiasters; (K) Spiraster.

**Thenea cf. valdiviae** von Lendenfeld, 1907

(Figures 20A & 21)

**MATERIAL**

Seven specimens from Station 40–367, 2961 m: ZMBN 105613–14. 7 specimens from Station 46–372, 3046 m: ZMBN 105627–31. 1 specimen from Station 52–374, 2977 m: ZMBN 105652.

**COMPARATIVE MATERIAL EXAMINED**


**OUTER MORPHOLOGY AND SKELETON ORGANIZATION**

*Thenea valdiviae*, Sub-globular, slightly flattened, sometimes with a triangular shape. Fairly hispid surface. One to several oscules with large meshed sieves (Figure 20B). Equatorial poral area (also sieved). Specimens all have a dirty colour in ethanol. Thick sections of 105631 were made. Large dichotriaenes are found at the surface, with their cladomes supporting the ektosome in a very regular fashion (Figure 20D). Bundles of oxeas...
surround the triaenes and cross the ectosome. Spirasters and plesiasters are very abundant throughout the choanosome. Foraminifera (Globigerina especially) are abundant in the choanosome. Sediments are also abundant.

**SPICULES, ZMBN 105631, UNLESS OTHERWISE STATED** (FIGURE 20E–G)

*Megascleres:* (a) oxeas. (b) dichotriaenes with pointy bent rhabdome. (c) proatriaenes, rare (only found in 105614, 105627 and 105628). *Microscleres:* (d) plesiasters, common, usually with 5–7 actines, minutely spined, length of actine: 44–55.3–64 μm or 52–63.3–77 (in 105613); actine thickness: 3–5 μm. (e) amphisters (rare spirasters observed, no metasters observed), common, spiny, length: 20–26.6–32 μm but they can be up to 47 μm long (105613).

**BATHYMETRIC RANGE**

100–3046 m (Ca´rdenas & Rapp, 2012; this study).

**DISCUSSION**

Steenstrup & Tendal (1982) consider the species to be dimorphic with a spherical arctic form and a more flattened hemispherical boreal form. Ca´rdenas & Rapp (2012) later showed that there are two COI (Folmer fragment) haplotypes for this species: the Arctic COI haplotype is identical to the COI of *T. muricata* while the boreal haplotype has 1 bp. difference with the COI of *T. muricata*. With respect to their external morphologies, the MAR-Eco specimens are similar to the boreal form of *T. valdiviae*, also collected in western Norway (Ca´rdenas & Rapp, 2012). Proatriaenes were rarely observed and anatriaenes were not found; however, both of these megascleres do seem to be rare in this species (Ca´rdenas & Rapp, 2012). We further noted significant spicule differences with boreo-arctic comparative material. Plesiasters are much more common than in boreo-arctic specimens (from the Barents Sea, the Greenland Sea and the NWA) (Ca´rdenas & Rapp, 2012). These plesiasters have on average more actines (which are
thinner) than boreo-arctic specimens. Above all, the most noticeable difference is the presence of amphiasters (instead of spirasters), which have never been observed in this species before (including in the type from the Faroe Islands). Metasters were absent in the MAR-Eco specimens, although they are present, albeit in small numbers, in boreo-arctic specimens. For these reasons (plesiaster abundance and morphology, amphiasters, metaster absence), we decided to cautiously identify these specimens under the name *T. cf. valdiviae*.

The MAR-Eco specimens would be the deepest *T. valdiviae* ever collected (3046 m). These would also be the southernmost record for this species, usually considered to be boreo-arctic. While reviewing the distribution of this species we examined pictures of specimens from Stations 960 (394 m) and 1052 (440 m) collected in the Norwegian Sea and Barents Sea and identified by Topsent (1913) as *T. muricata* (pictures courtesy of M. Bruni, MOM). All these specimens are clearly *T. valdiviae* with sieved oscules. Specimens from Station 922 could not be retrieved (M. Bruni, personal communication) but since Topsent treated them also as *T. muricata* we assume they were also *T. valdiviae*. We also report here the first records of this species off Newfoundland (in the Flemish Cap). Actually, the specimen from the Flemish Cap identified as *T. muricata* by Murillo et al. (2012, Figure 6I) is also *T. valdiviae*: the oscule clearly has a sieve.

Suborder SPYROPHORINA Bergquist & Hogg, 1969
Family TETILLIDAE Sollas, 1886
Genus *Craniella* Schmidt, 1870

**Craniella azorica** (Topsent, 1913)

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**Material**

Two specimens from Station 70–385, 1650 m: ZMBN 105673-74.

**Comparative Material Examined**

*Craniella azorica*, MOM-INV-22534 (04 0889), lectotype here designated, large specimen represented by Topsent (1913, pl. III), paralectotype (smaller specimen), both wet specimens, Azores, near Terceira Island, Station 866, 38.88056, −27.21806, 599 m; MNHN-DT2877, two type slides (one from each type).
OUTER MORPHOLOGY AND SKELETON

ARRANGEMENT (FIGURE 22A, B)
Both specimens are small spherical hispid sponges, 1 cm and 1.5 cm in diameter. Surface colour is light greyish in ethanol. Cortex colour is light grey, choanosome colour is brown. Cortex is 0.7–1 mm thick. 105673 has a triangular fringe of spicules on its surface which could be a closed oscule (such as the ones we observed in a boreal species: Craniella zelandica (Carter, 1872)). Thick sections of 105674 were made. Oxeas I (and few pro/anatriaenes) form large radial tight bundles that cross the ectosome. Many protiaenes, and fewer anatriaenes appear in the extension of these bundles, at the surface of the sponge, thus forming the hispidity. We also observe a double-layered cortex: (i) an upper-layer with a thin ectosome (∼50 μm) made of large cells, and many sub-ectosomal canals, no spicules, (ii) a fibrous layer filled with a crisscross pattern of paratangential oxeas II. These oxeas II are absent in the choanosome. No embryos were observed in the thick sections.

SPICULES, ZMBN 105674, UNLESS OTHERWISE STATED (FIGURE 22C–E)
Megascleres: (a) oxeas I, large, a bit thicker on one end (=anisoactinate), length: 2325–3587.8–4320 μm (N = 16), width: 28–38.9–60 μm (N = 16); (b) oxeas II, slightly bent or sometimes double bent, slightly anisoactinate, few are very slightly centrotylote, length: 470–690.8–840 μm, width: 20–28.5–36 μm; length: 550–713.2–850 μm, width: 25–29.3–32 μm (105673); (c) protiaenes (sometimes modified to prodiaenes), equal clads, clads are sometimes subdivided at their tips, whip-like end of rhabdome, rhabdome length: 4720–5805–6800 μm (N = 8), width: 28–32.1–40 μm, clads: 140–209.3–260 μm; (d) anatriaenes, rhabdome length more than 9800 μm, width: 8–10.4–12 μm, clads: 70–85.7–100 μm.

BATHYMETRIC RANGE
599–1650 m (Topsent, 1913; this study).

DISCUSSION
Our specimens agree well with the type material. The cortical oxeas II of the MAR-Eco specimens are slightly shorter and wider than in the lectotype (520–809.6–1010 × 18–23.8–32 μm). The only noticeable difference is that oxeas II in our specimens are slightly bent (or double bent in 105674) whereas they are very straight in both types. Topsent (1913) had noted that protiaenes could have irregular clads, we observed that these clads can even subdivide, as in the Balgim specimens (Boury-Esnault et al., 1994). As for the anatriaenes, their cladomes have an umbrella shape, similar to the ones found in the paralectotype (and not as open as in the
We observed spermatic cysts in the choanosome (sections of 105674, collected on 26 July 2004), but we did not observe embryos (in either specimen).

The MAR-Eco specimens greatly extend the northern distribution range for this species, until now only recorded from the Azores (Topsent, 1913, 1928) and the Ibero-Moroccan gulf (Balgim campaign) (Boury-Esnault et al., 1994). The MAR-Eco specimens were found in the same depth range as the Balgim specimens (1510 m).

Genus *Tetilla* Schmidt, 1868
*Tetilla longipilis* Topsent, 1904
(Figures 20C & 23)

**MATERIAL**
One specimen from Station 40–367, 2961 m: ZMBN 105612.
Two specimens from Station 42–368, 2078 m: 105616–17.
Seven specimens from Station 50–373, 2600: ZMBN 105643.

**COMPARATIVE MATERIAL EXAMINED**
*Tetilla longipilis*, MOM-INV-22536 (04 0515), lectotype here designated, specimen represented by Topsent (1904, plate II), MOM-INV-22537 (04 0515), 12 paralectotypes, Azores, SE of Flores, Station 698, 39.18333, 230.74444, 1846 m; MNHN-DT880, slide of type material (not clear to which syntype it belongs); PC484, EMEPC2007, field#D37A-Ma10, Plato Seamount, 33.1509, 28.7905, 1716 m. *Tetilla polyura*, PC522, Ecosystem Barents Sea 2007 campaign, Station 2663, Barents Sea.

**OUTER MORPHOLOGY AND SKELETON ARRANGEMENT (FIGURE 23A–E)**
Massive subspherical, up to 12 cm in diameter, highly hispid. Specimens (105643) have a long greyish and brownish ‘fur’ (Figure 23A, B). 105616 (Figure 23C) is less hairy and has a large base made of spicules; it also has a small apical oscule. Pores not visible. Choanosome is light brownish in ethanol. Thick sections of 105616 (Figure 23D, E) and 105617 were done, revealing identical skeleton organizations. Oxeas I (and few pro/anatriaenes) form large radial tight bundles that cross the ectosome to form the typical ‘fur’. Protariaenes and anatriaenes are very abundant in the ‘fur’, which is full of sediments. Just below the surface where sigmaspires tend to accumulate, there is a fibrous layer 120–200 μm thick, where some anatriaenes can also be observed. Below, the choanosome is full of oxeas II sometimes paratangential to the surface (they do not form bundles), and sometimes without a particular orientation. Sigmaspires are also very common there.

![Fig. 23. *Tetilla longipilis* Topsent, 1904: (A–B) ZMBN 105643 cut in half; (C) ZMBN 105617, notice the large base made of spicules only; the small single oscule is on the top (not visible on this picture); (D) Thick section from ZMBN 105617 showing the confused disposition of the oxeas II in the choanosome; (E) Close up of the ectosome in D. The fibrous layer lower boundary is shown with a drawn black line; (F) Thick section of lectotype MOM-INV-22536 showing the unusual criss-cross layer of oxeas II below the ectosome; (G) Protariaene (ZMBN 105617); (H) Anatriaene (ZMBN 105617); (I) Sigmaspere (ZMBN 105612).](https://www.cambridge.org/core)
SPICULES, ZMBN 105616 (FIGURE 23G–I)

*Megascleres:* (a) oxeas I, large, a bit thicker on one end (= anisoactinate), length: 7650–10807.1–13250 μm (N = 14), width: 60–84.6–100 μm (N = 14); (b) oxeas II, usually straight and rarely bent, both ends are equal (=isoactinate), some modified to styles, length: 410–831.5–1475 μm, width: 8.5–16.9–30 μm; (c) protoriaenes, rhombode length: 660–3820–8650 μm (N = 18), width: 5–132.6–240 μm, clads: 50–132.6–240 μm; (d) anatriaenes, rhombode length: 11000–11750–13250 μm (N = 3), width: 5–12.7–30 μm, clads: 18–71.5–140 μm. *Microscleres:* (e) spiny sigmaspires, ‘s’ or ‘c’ shape, sometimes with a slight central swelling, length: 14–16.5–22.5 μm.

**BATHYMETRIC RANGE**

999–2960 m (this study).

**DISCUSSION**

This is only the second record of this species, if we exclude a record of *Tetilla aff. longipilis* (van Soest et al., 2007) which has been re-identified as *Craniella* sp. (P. Cárdenas, unpublished results). Along with this second record we bring additional boreal records from specimens we identified from the Danish-Ingolf Expedition 1895–1896 and BIOICE (both are ZMUC collections). Furthermore, we identified this species on pictures of specimens trawled in Hatton Bank (NW of Rockall Bank), and referred to as *Craniella* sp. (Durán Muñoz et al., 2009). Koltun (1966) was wrong in synonymizing *T. longipilis* with *T. polyura* Schmidt, 1870; they have clearly different external morphologies and spicule differences, so they are both valid species. Furthermore, *T. longipilis* is Lusitanian and boreal, *T. polyura* is an arctic species. We considerably extend the northern range of *T. longipilis* since we report it for the first time in boreal waters, thanks to specimens we identified in the Ingolf Expedition and BIOICE.

The external morphology and spicule measurements of the MAR-Eco specimens agree well with the type series described from the Azores. Our thick sections of 105616 and 105617 (Figure 23D, E) were compared with the ones we made from the lectotype (Figure 23F), collected in the Azores at 1846 m depth. The skeleton arrangements are similar but below the fibrous layer (300–400 μm thick), the oxeas II in the lectotype are more regularly arranged and concentrated in a paratangential manner making a loose layer about 300 μm thick. This layer is made of shorter oxeas II (around 600 μm long) than the other oxeas II in the choanosome (around 1000 μm long). Further observations of cross-sections in many specimens suggest that this layer of oxeas II, which makes a thin whitish layer visible to the naked eye, is more or less present depending on where the section is made. This layer is not characteristic of a *Tetilla* and reminds the microxea layer of *Craniella* species (Figure 23F). Another character that reminds of *Craniella* is the presence of embryos in the type series; however, they are bigger and less abundant than in *Craniella* species (Topsent, 1904). We did not find them in the MAR-Eco specimens, nor in the Ingolf and BIOICE material that we examined. Based on the presence of cortical oxeas, Lehnert & Stone (2011) suggest the transfer of this species to *Craniella* but the thick sections made on the lectotype and the MAR-Eco specimens clearly show that even though *T. longipilis* has a denser arrangement of oxeas II below its fibrous layer, it has no true double-layered cortex, as in other *Craniella* species sensu Sollas (1888) (cf. description of *Craniella azorica*). So in our opinion, although this species might represent an evolutionary intermediate step between *Tetilla* and *Craniella*, we propose to keep it in *Tetilla* until additional data are obtained, and a revision of *Tetilla* and *Craniella* is made.

*Tetilla sandalina* Sollas, 1886

(FIGURES 20D & 24)

**MATERIAL**

One specimen from 44–369, 1742 m: ZMBN 105626.

**COMPARATIVE MATERIAL EXAMINED**

*Tetilla polyura*, PC522, Ecosystem Barents Sea 2007 campaign, Station 2663, Barents Sea; ZMUC, Godthaab Expedition 1928, off Labrador, 55, −56.56667, 314 m, original det. by Brøndsted (1933) as *Tetilla sandalina*.

**OUTER MORPHOLOGY (FIGURE 24A)**

This is a small fragment of a larger sponge. No cortex is visible. A large spicule tuft is present on the surface which could represent an oscule.

![Fig. 24. *Tetilla sandalina* Sollas, 1886, ZMBN 105626: (A) Specimen; (B) Oxeas I; (C) Protoriaene; (D) Sigmaspires and raphid-like oxeas III.](https://www.cambridge.org/core/terms .https://doi.org/10.1017/S0025315415000983)
spicules, ZMBN 105626 (Figure 24B, D)
Megascleres: (a) oxeas I, large, a bit thicker on one end (=anisoactinate), length: 1210–2124.3–2900 μm, width: 14.5–22.2–30 μm; (b) oxeas II, straight or slightly bent, both ends are equal (=isooactinate), length: 580–658.6–785 μm, width: 15–17.1–20 μm; (c) oxeas III, raphide-like, flexuous, 242–250 μm, width: 2–3 μm; (d) protariaenes, rhabdome with a whip-like end, rhabdome length: 1300–1967.6–3200 μm (N = 17), width: 2.5–5.2–7.5 μm, unequal cladome with two small clads and a much longer one, small clads: 5–24.3–40 μm, longer clad: 52–160.4–225 μm. Microscleres: (e) sigmaspires, ‘s’ or ‘c’ shape, length: 15–19.7–25 μm, width: 1 μm.

bathymetric range
1742–1818 m (Sollas, 1888; this study).

discussion
We do not have a complete specimen but it is still enough to confirm the validity of this species. Indeed, both Topsent (1923) and Koltun (1966) suggest that T. sandalina is a synonym of T. polyura but after examining both species, we conclude that, although they are obviously morphologically close, they are clearly distinct. They both share the typical ‘2 short clads + 1 long clad’ protariaene cladome and the raphide-like oxeas (called ‘trichodal oxeas’ by Sollas (1888)) but in T. sandalina, the sigmaspires are not centrotylote as in T. polyura. Furthermore, as in the original description, we could not find any anatriaenes (whereas they are very commonly found in T. polyura). One difference with the original description is that Sollas (1888) reports that one or both of the shortest clad can disappear; we have not observed this in our specimen. This is only the second record of this species, originally collected at a similar depth (1818 m) in the Azores during the Challenger expedition. We examined the ZMUC specimen of T. sandalina collected off Labrador (314 m depth) from Brøndsted (1933) and re-identified it as T. polyura (it has centrotylote sigmaspires, smaller than in T. sandalina: 12–15 μm); this is not surprising since T. polyura has been recorded in the Flemish Cap at similar depths (F. J. Murillo, personal communication).

Order POLYMASTIID order Morrow & Cárdenas, 2015
Family POLYMASTIIDAE Gray, 1867
Genus POLYMASTIA Bowerbank, 1864
Polymastia corticata Ridley & Dendy, 1886
(Figures 25 & 26)

Material
One specimen from 70–385, 1650 m: ZMBN 105663.

Comparative material examined

outer morphology and skeleton
Organization (Figure 26A, B)
Massive cushion-shaped specimen. Surface is nearly smooth, with exhalant and inhalant papillae of different sizes, irregularly distributed. Colour of the surface, papillae and choanosome is pale pink in alcohol. Cortex dense, leathery, easy detachable. Choanosome friable. Exhalant papillae conical, slightly flattened, 2.8 mm in basal diameter and 0.4 mm in apical diameter. Inhalant papillae are nearly cylindrical. Main choanosomal skeleton is a reticulation of bundles of principal styles. Additional choanosomal skeleton made by free-scattered intermediary styles. The cortex is 2 mm thick, made of two overlapping layers: the external palisade of small styles and the internal confused and dense mass of intermediary styles mixed with small styles.

spicules, ZMBN 105663 (Figure 26C–E)

bathymetric range
1113–2390 m (this study).

discussion
The type locality of P. corticata is in the South-West Atlantic, between Pernambuco and Bahia (Figure 25), at 2194 m depth, and yet our specimen very closely matches this species. Similar connections between the northern MAR and the South-West Atlantic have been reported for deep-sea Hexactinellida.
The specimens described by Topsent (1892, 1904) in the Azores are similar although he reports only one type of papillae. The specimen SMF 9633 (Lesser Antilles, Kahouanne Basin, 1127 m) was identified by D. Janussen (Meyer & Kuever, 2008); we did not examine this specimen but its 28S (D1–D2) sequence (GenBank#EU005552) groups with the sequences of other identified *P. corticata* (A. Plotkin, personal communication).

**Order SUBERITIDA Chombard & Boury-Esnault, 1999**

**Family STYLOCORDYLIDAE Topsent, 1892**

**Genus Stylocordyla Thomson, 1873**

**Stylocordyla borealis** (Loveén, 1868)  
(Figures 27 & 28)

**MATERIAL**

One specimen from 50–373, 2600 m: ZMBN 105647.

**COMPARATIVE MATERIAL EXAMINED**


**OUTER MORPHOLOGY (FIGURE 28A)**

Typical pedunculate shape with long stalk. One osculum (3 mm in diameter), surrounded by a spicule fringe, is present on the upper side of a conical body. The head is somewhat flattened on top (length of head: 2 cm). Length of stalk: 9 cm. Brown in ethanol.

**SPICULES, ZMBN 105647 (FIGURE 28B–E)**

*Megascleres*: (a) oxeas I, straight, usually slightly centrotyloic, length: 776–1011–1367 μm (N = 23), width: 10.4–14.8–20.8 μm (N = 23). (b) oxeas II, more or less centrotyloic, straight to slightly bent, some are asymmetrical with a sharp bent, length: 316–503–684 μm, width: 3.9–5.7–10.4 μm.  
(c) microxeas, some are faintly centrotyloic, usually straight, length: 80–93–116 μm, width: 2.3–3.2–4.6 μm. Spicules were sampled from the body and stalk.

**Fig. 26.** *Polymastia corticata* Ridley & Dendy, 1886, ZMBN 105663: (A) Specimen; (B) Thick section; (C) Principal styles; (D) Intermediary styles; (E) small cortical tylostyles.

**Fig. 27.** Distribution map of *Stylocordyla borealis* (Loveén, 1868). T, type locality; CGFZ, Charlie-Gibbs Fracture Zone. Map generated with GeoMapApp 3.3 (http://geomapapp.org).
BATHYMETRIC RANGE
22 – 2600 m (this study).

DISCUSSION
This specimen’s morphology agrees well with the redescription of the species and the type material (Uriz et al., 2011). The only difference we note is that the osculum is not present on the top surface but is slightly off to the side, as in *Stylocordyla longissima* Sars, 1872, a species later synonymized with *S. borealis* (Hansen, 1885; Vosmaer, 1885; Arndt, 1913). This species is widespread across the deep North Atlantic boreo-arctic region, on either side of the Atlantic. However, this is its first record in the Azores region. Many GBIF (www.gbif.org) records (from the Yale Peabody Museum, material not seen) suggest that this species is very common on the eastern US coast shelf down to the Bahamas (Figure 27). Schmidt (1880) also recorded this species from Grenada (290 m). We found a specimen from Olfafsjörður (Iceland, BIOICE campaign, Station 2193) collected at only 18–22 m depth, which is the shallowest record for this species. The dubious identity of the small *S.

Fig. 28. *Stylocordyla borealis* (Lovén, 1868), ZMBN 105647. (A) Specimen; (B) Oxeas I; (C) Oxeas II; (D) Microxeas.

Fig. 29. *Forceptia (Forcepia) toxfera* sp. nov., holotype, ZMBN 105651: (A) Holotype; (B) Forceps; (C) Chelae; (D) Sigma; (E) Styles, sigmas and oxeote toxas (arrows); (F) Style; (G) Tylote.
**Expedition to Greenland 1883**, 228 m.

Holotype, east coast of Greenland, number 579, Sofia SNHM-Type-8629 (wet specimen), SNHM-113069 (slide), *thielei* (Lundbeck, 1905), ZMUC-DEM-171 (slide), syntype, Ingolf Expedition Station 73, South of Iceland, 62.96667, 23.46667, 915 m.

**Outer morphology (Figure 29a)**

Massive subspherical, 3 cm in diameter, fairly compressible, slightly velvety surface, mixed with sand and bivalve debris, whitish in ethanol.

**Spicules (Figure 29b–g)**

*Megascleres*: (a) styles, slightly bent, very often with a stronger bent below the blunt end, completely smooth or with very few spines (1–3), length: 622–684–735 μm, width: 10–25–29 μm. (b) oxeote toxas, less common than styles, smooth, rarely modified to oxeas, length: 770–857.6–920 (N = 17), width: 20–22–25 μm (N = 17). (c) tylotes, length: 413–486–520 μm, width: 7.8–9.9–10.4 μm, tyle: 9.1–10.3–13.0 μm.

*Microscleres*: (d) arcuate isochelae, length: 22–37.9–55 μm, width: 5–7 μm, shaft (side view): 2.5–4.5 μm. (e) sigmas, very common, plane, ’C’ shaped, more rarely ’S’ shaped, hook-shaped ends, rarely one of the ends is divided in two (seen 3), length: 105–125.2–175 μm, width: 5–7 μm. (f) forceps, not common, length of long branch: 43–52.8–65 μm (N = 11), length of short branch: 25.7–30.4–35.5 μm (N = 20).

**Etymology**

This species is named after the large oxeote toxas: ‘tosafera’ means ‘that carries toxas’.

**Bathymetric range**

2252–2977 m (Topsent, 1904; this study).

**Discussion**

The spicule assemblage of the MAR-Eco specimen is quite similar to the one found in *Forcepia (Forcepia) groenlandica* and *Forcepia (Forcepia) thielei*, respectively described from the east coast and from the west coast of Greenland. Both type materials were re-examined for this study and we also made new spicule preparations from the type of *F. (F.) groenlandica*. Actually, our specimen is even closer to the specimen borealis specimen (BMNH.1882.7.28.73) collected off Bahia (Brazil) at only 12–36 m depth by the Challenger ( Ridley & Dendy, 1887) would need to be revised.

**Order POECILOSCLERIDA** Topsent, 1928

**Family COELOSPHAERIDAE** Dendy, 1922

**Genus Forcepia** Carter, 1874

*Forcepia (Forcepia) toxafera* sp. nov. (Figure 29, Table 3)

<table>
<thead>
<tr>
<th>Material</th>
<th>Depth (m)</th>
<th>Styles (length/width)</th>
<th>Oxeote toxas (length/width)</th>
<th>Tylotes (length/width)</th>
<th>Isochelae (length/thickness)</th>
<th>Sigmas (length/thickness)</th>
<th>Forceps (short/long branch)</th>
</tr>
</thead>
</table>

Means are in bold; other values are ranges. N = 30 unless stated otherwise between parentheses, or unless measurements come from other studies. *Measurements from this study.

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described by Topsent (1904) under the name F. (F.) groenlandica, and collected at great depths (2252 m) in the Azores. Topsent’s specimen and ours are clearly conspecific (Table 2) and differ from F. (F.) groenlandica: (i) the styles are larger, (ii) the sigmas of our specimens are plane (vs more or less contorted), (iii) the styles are smooth with occasional spines (vs fairly spiny) and (iv) there are large oxeote toxas, absent in F. (F.) groenlandica. The nearly smooth styles also make F. (F.) toxafaera sp. nov. close to the Arctic species F. (F.) thielei but there are differences: (i) the styles can have a few spines (vs no spines at all in F. (F.) thielei), (ii) the forceps of our specimen are less densely spined, (iii) the short branches of the forceps are straight (vs bent in F. (F.) thielei), and again (iii) there are oxeote toxas. All these differences support the creation of a new species, named after the large oxeote toxas that were overlooked by Topsent (1904) but we found them to be common (albeit in lower numbers than the styles) when we re-examined his specimen (MOM INV-22577).

After F. (F.) groenlandica and F. (F.) thielei, F. (F.) toxafaera sp. nov. is the third species of Forcepia with a single category of forceps, which are asymmetrical. There are three other North Atlantic Forcepia species with asymmetrical forceps but they also have another category of larger symmetrical forceps: Forcepia (Leptolabis) brunnea (Topsent, 1904), F. (L.) assimilis (Lundbeck, 1910) and F. (L.) forcipula (Topsent, 1904). Forcepia (L.) brunnea and F. (L.) assimilis may actually be the same species according to Topsent (1928).

**SPICULES (FIGURE 30B-E)**

Megascleres: (a) styles, smooth, straight to very slightly curved, length: 5.46–612–673 µm, width: 6.5–16.6–23.4 µm. (b) tylothes (the swellings of the ends can be very faint but are always present), straight, length: 367–401–454 µm, width: 5.2–7.0–9.1 µm, tyle: 6.5–7.9–9.1 µm. Microscleres: (c) unguiferous chelae I with 5–6 pointy teeth, length: 49–58–68 µm, alae span: 7.8–10.1–13.0 µm, shaft: 3.9–4.5–5.2 µm. (d) polydentate chelae II, with shaft more or less strongly bent, 7–9 teeth with sub-rounded ends, length: 14–20–25 µm, shaft (side view): 1.3 µm, alae span: 5.2–8.0–10.4 µm.

**ETYMOLOGY**

This species is named ‘paravaridens’ to underline its presumed phylogenetic affinities with the Arctic species *Iotroata varidens*.

**DISCUSSION**

According to their definition, Iotrochotidae species should have birotula microscleres (van Soest, 2002). However, our new species does not have birotulas but two sizes of anchorate chela. We nonetheless decide to allocate this species to the genus *Iotroata* for the following reasons: (1) the spicule repertoire of this species closely matches that of other *Iotroata* species (smooth styles, tylothes and unguiferous chelae) and (2) birotulas are probably derived from anchorate chelae (van Soest, 2002). So we consider here that the smaller size of anchorate chelae in our new species is homologous to the small category of birotulas in other *Iotroata* species. Furthermore, this new species is very similar with respect to spicule morphology and sizes to *Iotroata varidens* (Lundbeck, 1910), a deep-sea species described from the Denmark Strait at 567 m depth. The main difference between the two species is the presence of birotulas in *I. varidens* (size of birotulas: 15–21 µm) vs polydentate anchorate chela in *I. paravaridens* (size of chela: 14–25 µm).

**TYPE MATERIAL**

**Holotype**: ZMBN 105615, Station 40–367 (42.91667, –30.33333), 2961 m.

**OUTER MORPHOLOGY (FIGURE 30A)**

Small subspherical, and presumably complete specimen (1.5 cm in length); brownish, full of debris and sand.

**Fig. 30. Iotroata paravaridens** sp. nov., holotype, ZMBN 105615: (A) Holotype; (B) Chelae II; (C) Chelae I; (D) Styles; (E) Tylote.
One specimen from 53–375, Faraday Seamount, 990 m: ZMBN 105658.

Comparative material examined
Spongionella pulchella, GNM-Spongiaria-801, 803, 804, Svenska Artprojektet, Swedish West coast; Spongionella sp., Ifremer Caracole 2001, specimen K14h, Rockall Bank, 867 m; Spongionella sp., GNM-Spongiaria-802, Svenska Artprojektet, Swedish West coast; Spongionella gracilis (Vosmaer, 1883), 209S, personal collection of J. Vacelet (Endoume Marine Station, Marseille), Cap Kavensur, Tunisia, 37 m.

Outer morphology and skeleton (figure 31a)
Small whitish cup-shaped specimen with a conulose surface. Specimen is 2 cm in diameter with a single oscule (~3 mm in diameter). Very regular network of primary and thinner perpendicular secondary fibres, pale yellow in colour. Distance between primary fibres is 550–1150 μm long. A few tertiary fibres interconnect secondary fibres. Fibres are stratified (no distinct pith observed). Eurypylous chambers are large, usually more than 100 μm in length and up to 190 μm. Twelve embryos (~200 μm in diameter each) were observed in the choanosome of the sections.

Fibres (figure 31b)
(a) Primary fibres, stratified, 27–45 μm wide, length of extension beyond surface: 250–400 μm, (b) secondary fibres, stratified, 12–20 μm wide, (c) tertiary fibres, more rare, around 10 μm.

Discussion
Four species of Spongionella are recorded from the North-East Atlantic/Mediterranean area: S. pulchella (Sowerby, 1806), S. gracilis (Vosmaer, 1883), S. ramodigitata (Topsent, 1901) and S. depressa Topsent, 1929. Based on our measurements of the primary and secondary fibres and on the external morphology, our specimen is closer to S. pulchella. But the distance between the primary fibres (550–1150 μm) is much bigger than usual (200–350 μm), which is a character found in S. depressa. Moreover, S. pulchella is usually found between 40 and 380 m, the deepest record being off Monaco (Vacelet, 1969). So we tentatively identify this specimen as S. aff. pulchella, while waiting for additional specimens from this area and a revision of this group in the North Atlantic/Mediterranean Sea which should test if North Atlantic and Mediterranean populations are conspecific and if deep-sea and shallow S. pulchella are conspecific. While investigating Spongionella species, we came across the description of Spongionella brandtii (Miklucho-Maclay, 1870) from the Sea of Okhotsk (Arctic Pacific), currently a synonym of S. pulchella according to previous authors (von Lendenfeld, 1888; Koltun, 1959). But S. brandtii has a characteristic smooth foliaceous shape (termed ‘discoidal’ by Koltun) with concentric trenches (Koltun, 1959, plate 41), whereas S. pulchella is often a rather thick lamella with a minute conulose surface (not smooth). We suggest that S. brandtii type material should be re-examined carefully because we suspect this species to be valid.

Discussion
We have identified in the MAR-Eco collection 22 species of Demospongiae, two of which are new for science (Table 1): Forcepia (Forcepia) toxafera sp. nov. and Iotroata paravaridens sp. nov. Seventeen out of the 22 belong to the Tetractinellida order, a group thought to have originated in the deep sea (Cárdenas et al., 2011). In comparison, 16 species of Hexactinellida sponges were identified in the MAR-Eco material with 13 species new for science (Menshenina et al., 2007; Tabachnick & Menshenina, 2007; Tabachnick & Collins, 2008). Only one specimen of Calcarea sponge was found in the MAR-Eco collection: a new calcaronean species (to be described). Station 50 (north of the Azores) was the richest in terms of demosponge.
species and biomass: eight species (including four Geodia species) and 24 specimens indicating a sponge ground in this area. Station 70 (north-west of the CGFZ) was also fairly diverse with seven species and 13 specimens. Station 70 actually had the highest diversity of Hexactinellida with nine species identified (Gebruk et al., 2010). Overall, Station 70 (at the southern tip of Reykjanes Ridge) had not only the highest sponge diversity, but also the highest benthic species richness (Gebruk et al., 2010).

The MAR is not a longitudinal barrier for deep-sea demosponges

According to our distribution maps (Figures 2, 13, 20, 25 & 27), 68% of the MAR-Eco species (15 species out of the 22) are amphi-Atlantic. We further note that both boreo-arctic species (with distributions north of the CGFZ) and Lusitanian species (with distributions south of the CGFZ) can be amphi-Atlantic. Although Tetilla longipilis has been essentially collected on the MAR, apart from a record in Hatton Bank (NEA), we have identified specimens collected in the Denmark Strait and on the western side of the MAR so it is likely present further west, and we suspect it to be a true amphi-Atlantic species. Since this is a poorly known species it could have easily been overlooked in the NWA. On the other hand, Poccellastra compressa seems genuinely restricted to the NEA and the MAR, since it has numerous records and is a relatively easy species to identify. The rest of the NEA-restricted species are either new species (I. paravaridens sp. nov., F. (F.) toxafera sp. nov.) or poorly known deep-sea species (i.e. with very few records) with distribution ranges south of the CGFZ (C. azorica, T. sandalina, S. aff. pulchella). Therefore these species might very well be amphi-Atlantic but for now they are considered restricted to the NEA and/or the MAR. 68% of amphi-Atlantic sponge species is well in agreement with a faunistical review showing that 61% of the 80 species collected in the Reykjanes Ridge (Madreporaria, Cirripedia, Echinoidea, Asteroidea and Brachiopoda), north of the CGFZ, were amphi-Atlantic (Mironov & Gebruk, 2006). The conspecificity of boreo-arctic amphi-Atlantic Geodia species has been further confirmed with genetic data (Cárdenas et al., 2013). Seemingly, the status of the amphi-Atlantic MAR-Eco species found south of the CGFZ would need to be tested with genetic data. To conclude, the high proportion of amphi-Atlantic species suggest that the MAR is not a barrier to the dispersal of deep-sea Demospongiae. On the contrary, deep-sea demosponges may use the MAR and neighbouring continental margins to disperse at bathyal depths, to avoid the much deeper abyssal depths (i.e. large abyssal basins) where they are absent (Figures 2, 13, 20, 25 & 27).

The CGFZ, a potential latitudinal border for deep-sea demosponges

Some MAR-Eco species are restricted to the boreal and/or Arctic waters and thus stay north of the CGFZ: Stelletta rhaphidiophora, Geodia macandrewii, Geodia phlegreaei, Thenea levis. Other MAR-Eco species are restricted to Lusitanian waters and thus stay south of the CGFZ or at the most on the northern part of the CGFZ: Geodia nodastrella, Thenea schmidtii, Craniali azorica, Tetilla sandalina. Finally, eight species can be found north and south of the CGFZ: Geodia atlantica, Geodia hentscheli, Stelletta tuberosa, Stryphnum forsy, Poccellastra compressa, Thenea valdiviae, Tetilla longipilis and Stylocordyla borealis. This is not surprising since most (65%) of the species identified during a study in the Reykjanes Ridge had distribution ranges further south than 52°N (Mironov & Gebruk, 2006). However, in the case of five out of these eight species, our study showed that the MAR southern populations (i.e. south of the CGFZ) were somewhat morphologically different from the northern populations. Southern populations of G. atlantica, G. hentscheli, T. valdiviae, S. forsy and P. compressa had clear external morphology and/or spicule differences. More specimens from this area as well as genetic data are now required to confirm these observations and test if these southern populations represent sister-species. Actually, T. schmidtii is probably another example of this since it may represent a southern sister-species of T. muricata, hence our difficulty to discriminate them morphologically. Genetically, the mitochondrial cytochrome oxidase I (COI) Folmer fragment is not enough to differentiate these two populations but the 28S (C1–D2) marker is more promising (Cárdenas & Rapp, 2012). Likewise, F. (F.) toxafera sp. nov. and I. paravaridens sp. nov. are probably sister species of more northern Greenland species, respectively F. (F.) groenlandica and I. varidens. Overall, these observations suggest that the CGFZ area represents a major biogeographic barrier for deep-sea demosponges, which limits gene flow between northern and southern populations. This barrier is not limited to the two deep fractures per se (about 4500 m deep) since some of the southern species (C. azorica, S. tuberosa) were actually sampled slightly north of the CGFZ (Station 70 and 72) but it is more likely in combination with environmental changes in the CGFZ area. It is also interesting to notice that two (G. hentscheli, S. rhaphidiophora) out of the five species are strict Arctic species, which manage to extend their southern distribution range to the CGFZ, probably by following the southward flow of cold deep waters (Iceland-Scotland Overflow Water: ISOW) along the Reykjanes Ridge.

The CGFZ area has already been considered a major biogeographic transition zone for many planktonic, pelagic or benthic organisms, (Mironov & Gebruk, 2006; Gebruk et al., 2010; Vecchione et al., 2010; Alt et al., 2013). However, one should keep in mind that the CGFZ may not be a boundary for all marine organisms: Kongsrud et al. (2013) do not find a difference in the species composition of benthi polychaetes north and south of the CGFZ and wish for more samples to settle the matter. Watling et al. (2013) delineated global lower benthal biogeographic provinces (801–3500 m) but were unsure of the boundaries for many of them. In our opinion, the CGFZ could be considered as the MAR biogeographic boundary between the two lower benthal provinces BY2 (North Atlantic Boreal) and BY4 (North Atlantic) (Watling et al., 2013).

The distribution of deep-sea sponges in the North Atlantic

Numerous studies from shallow water demosponges suggest that sponge larvae are generally short-lived, thus suggesting that short-distance dispersal may be the rule in shallow waters (Uriz & Turon, 2012). There are currently no behavioural or ecological studies on deep-sea demosponge larvae.
One study on the Hexactinellida *Oospacas minuta* (considered to be a Mediterranean deep-sea species) showed that most larvae settle and metamorphose into the juvenile sponge within 12–24 h after release from the parent (Leyes et al., 2007). So deep-sea sponge larvae may have similar behaviour and short-life expectancies as shallow species and thus have short-dispersal potential. Keeping this in mind, knowing that the CGFZ represents a border between two deep-sea sponge faunas partially hints at the environmental parameters that might influence and limit the distribution of deep-sea sponges. Temperature, salinity and oxygen are very similar at the north and the south of the CGFZ (Table 1) but other environmental parameters related to depth may influence the distributions of deep-sea sponges.

Even though we acknowledge that the distribution maps presented in this study closely mirror overall sampling efforts, one should remember that the North Atlantic is probably the best studied and explored deep-sea area in the world. So we can consider most of these distribution maps (for all the Astrophorina: *Thenea* spp., *Geodia* spp., *P. compressa*, *S. fortis*) as good approximations of the current distribution of these species. Although these maps suggest widespread distributions for these deep-sea sponges, depth clearly seems to be a limiting factor. *Geodia barretti* is the most widely distributed species of our collection, probably because it is the most common large *Geodia* of the North Atlantic, with one of the widest bathymetric ranges (30–2000 m) (Cárdenas et al., 2013). *Geodia philogrea* was commonly observed and collected at 3000 m on Orphan Knoll (north of Flemish Cap) (Cárdenas et al., 2013) and we examined a specimen of *Geodia megastrella* collected at 4152 m depth on the Atlantic continental margin off France (MNHN-DCL2857); both of these species were present in the MAR-Eco material. *Thenea valdiviae* also has a large track record since it is easily collected in soft sediments, with an even wider bathymetric range (100–3046 m). So most deep-sea demosponge species have bathymetric records that rarely go beyond 3000 m depth, which thus probably reflects a true lower limit. Of course, there are a few typical deep-sea North Atlantic demosponges that live in the deeper abyssal plains (Barthel & Tendal, 1993), but these species were not found in the MAR so our discussion will focus on bathyal demosponges. The 3000 m depth limit of bathyal demosponges was confirmed by direct observations when exploring the 4500 m deep CGFZ with manned-submersibles, *Geodia* (called ‘round sponges’ in Felley et al. (2008)) were not found deeper than 3000 m and most were found shallower than 2500 m depth (Felley et al., 2008; Gebruk & Krylova, 2013). Hexactinellida are also fairly common and diverse around 2500 m depth but they can also occur deeper than 3000 m (Felley et al., 2008; Gebruk & Krylova, 2013). Interestingly, most corals in the CGFZ (Felley et al., 2008) also share this lower limit of 3000 m. So depth is certainly a limiting factor for the dispersal and distribution of most Demospongiae and Hexactinellida. If these species are restricted to the lower bathyal depth layer (800–3500 m) it means they can disperse without much depth restrictions in the boreo-arctic region, especially in the Shetland-Faroe-Iceland-south Greenland arc where depth does not exceed 3500 m (Figure 1). This would explain why we have so many amphi-Atlantic boreo-arctic sponge species. On the other hand, between 50°N and 30°N in the North Atlantic, deep basins with plains at abyssal depths (3500–6500 m) (Figure 1) would greatly limit or even prevent the dispersal of these animals, which usually have short-lived larvae. For instance, at 48°N, the MAR is about 1000 km from the closest continental margins (Flemish Cap to the west, Porcupine Bank to the east) (Figure 1). And yet, our results showed that some Lusitanian species were amphi-Atlantic so they manage to disperse across the Atlantic. These deep-sea demosponges would then maybe use the numerous North Atlantic seamounts (Figure 1) and the MAR as ‘stepping stones’ above the wider ocean basins, although this scenario has never been extensively tested in the North Atlantic and the few genetic studies made tend to give mixed results (Cho & Shank, 2010; Rowden et al., 2010). It is however a fact that *Geodia* species are commonly found on the slopes and tops of seamounts (e.g. seamounts south of the Azores or off Portugal), or oceanic islands (Bermudas, Azores).

The CGFZ is characterized by two main deep rift valleys 35 km apart; they can be up to 4500 m deep, but are only 10 km wide (Figure 1) (Priede et al., 2011). CGFZ sponges significantly show clumped distributions, which could only be partially accounted for by their preference for hard substrates (Felley et al., 2008). So it is possible that either larvae are attracted to other conspecific individuals or that larvae settle close to the parents, a common feature in shallow sponge larvae (Uriz & Turon, 2012). Furthermore, the fact that MAR sponges are mainly restricted to depths <3000 m and that dispersal may be over short distances, the 10 km wide/4500 m deep rifts may represent a true barrier for larval dispersal. But since Lusitanian species do make it to the northern edge of the CGFZ, larvae must find a way to cross in 1–2 days if we consider them short-lived. Other parameters may be limiting their dispersal further north.

In addition to depth, the distribution of Demospongiae may also be limited by suitable habitats, which may be rocky outcrops (e.g. for *Geodia* species, *P. compressa*) or soft bottoms (for *Thenea* or *Stylocordyla*). Soft bottoms are the most widely distributed habitats in the MAR so they cannot be a limiting factor. As for rocky outcrops, although they only represent 5% of the MAR, essentially in steep slopes (>30°) flanking the MAR (Niedzielski et al., 2013), they are present throughout the MAR and may therefore not significantly limit the dispersal of sponges which prefer these habitats.

Deep-sea currents may also influence deep-sea sponge dispersal and distribution. In the North Atlantic, the Gulf Stream gives the eastward North Atlantic Current (NAC) which crosses the MAR around the CGFZ (Søiland et al., 2008). The northern edge of the NAC (temperate waters) runs alongside the subpolar front (colder waters) which shifts between 48°N and 53°N (Søiland et al., 2008). This subpolar front is correlated with a clear faunal discontinuity (Vecchione et al., 2016). Meanwhile, there is an opposite deep current of overflowing high salinity bottom cold water coming from the Arctic: the Iceland-Scotland Overflow Water (ISOW). The ISOW runs southward along the flank of the Reykjanes Ridge and explains why deep-sea Arctic sponges (*S. rhaphidiophora* and *G. hentscheli*) reach the CGFZ. In the CGFZ area, the ISOW current flows mainly from east to west (in the opposite direction of the NAC) (Priede et al., 2011) and may therefore limit the dispersal to the north of species sensitive to cold waters. Furthermore, it has been argued that the ISOW does not act as a dispersal route across the ridge since substantial differences in megafaunal invertebrate...
community composition (sponges not included) within the northern part of the CGFZ were observed (Alt et al., 2013), but this needs to be confirmed with a much wider sampling. To conclude, even though the ISOW is the main current along the northern MAR up to the CGFZ, its influence on deep-sea sponge dispersal and distribution is poorly understood. One of the main reasons may be that the ISOW itself is still not fully understood (Kanzow & Zenk, 2014).

SUPPLEMENTARY MATERIAL
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