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The HMA-LMA Dichotomy Revisited: an Electron Microscopical Survey of 56 Sponge Species

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Abstract. The dichotomy between high microbial abundance (HMA) and low microbial abundance (LMA) sponges has been long recognized. In the present study, 56 sponge species from three geographic regions (greater Caribbean, Mediterranean, Red Sea) were investigated by transmission electron microscopy for the presence of microorganisms in the mesohyl matrix. Additionally, bacterial enumeration by DAPI-counting was performed on a subset of samples. Of the 56 species investigated, 28 were identified as belonging to the HMA and 28 to the LMA category. The sponge orders Agelasida and Verongida consisted exclusively of HMA species, and the Poecilosclerida were composed only of LMA sponges. Other taxa contained both types of microbial associations (*e.g.*, marine Haplosclerida, Homoscleromorpha, Dictyoceratida), and a clear phylogenetic pattern could not be identified. For a few sponge species, an intermediate microbial load was determined, and the microscopy data did not suffice to reliably determine HMA or LMA status. To experimentally determine the HMA or LMA status of a sponge species, we therefore

recommend a combination of transmission electron microscopy and 16S rRNA gene sequence data. This study significantly expands previous reports on microbial abundances in sponge tissues and contributes to a better understanding of the HMA-LMA dichotomy in sponge-microbe symbioses.

Introduction

Sponges (Porifera) represent an evolutionarily ancient phylum with a fossil record dating back to Precambrian times (Li *et al.*, 1998). Today, sponges are important components of the marine benthos and play an important role in the coupling of benthic and pelagic environments owing to their immense filter-feeding capacities (Bell, 2008; de Goeij *et al.*, 2013). Within their mesohyl tissues, many sponges harbor a great diversity of symbiotic microorganisms from the three domains of life: Archaea, Bacteria, and Eukaryota. To date, representatives from more than 28 bacterial phyla (including candidate phyla such as Poribacteria and Tectomicrobia) and two archaeal lineages were identified from marine sponges (Hentschel *et al.*, 2012; Schmitt *et al.*, 2012; Simister *et al.*, 2012). The vast majority of sponge-associated microbes remain uncultivated and are thus functionally largely uncharacterized (Taylor *et al.*, 2007).

The presence of microorganisms in marine sponge tissues has been known for almost a century. Dosse (1939) and

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Abbreviations: HMA, high microbial abundance; LMA, low microbial abundance.

Levi and Porte (1962) were among the first to describe microorganisms in the sponge mesohyl matrix using transmission electron microscopy. It was soon discovered that while some sponge species harbored dense microbial consortia within their mesohyl tissues, the mesohyl of other species from the same habitat were notably devoid of microorganisms (Reiswig, 1974; Vacelet and Donadey, 1977, Wilkinson, 1978). Accordingly, two general categories were identified that were termed “bacterial sponges” and “non-symbiont harboring, normal sponges” (Reiswig, 1981). Later on, the terms “low microbial abundance” (LMA) and “high microbial abundance” (HMA) sponges were coined to acknowledge the additional presence of archaea in sponge tissues (Hentschel *et al.*, 2003). A typical HMA sponge contains 10^8 to 10^{10} microorganisms/g sponge tissue, which can make up to 20%–35% of the sponge biomass (Reiswig, 1981; Webster *et al.*, 2001; Hentschel *et al.*, 2012); in contrast, only 10^5 to 10^6 bacteria/g sponge tissue are found in LMA sponges, which is roughly equivalent to the microbial abundances in seawater (Hentschel *et al.*, 2006). This pattern extends to reproductive propagules in that the larvae of HMA sponges contain dense bacterial assemblages at the larval center, while the interior of LMA sponge larvae is largely free of microbes (Ereskovsky and Tokina, 2004; Maldonado, 2007; Schmitt *et al.*, 2007; Gloeckner *et al.*, 2013a, b). Vertical microbial transmission from the parent to the larva, a hallmark of symbiotic host-microbe associations (Bright and Bulgheresi, 2010), is now considered an important and presumably evolutionarily ancient component of HMA sponge symbioses. The microbial community is likely complemented by horizontal acquisition of microbes from seawater, although this process has never been demonstrated (Schmitt *et al.*, 2008a; Webster *et al.*, 2010).

Besides bacterial abundances, there are also noticeable differences in microbial diversity between HMA and LMA sponges. Several studies, employing a variety of 16S rRNA gene-based methods, consistently demonstrated a lower microbial diversity in LMA than in HMA sponges (Weisz *et al.*, 2007; Kamke *et al.*, 2010; Erwin *et al.*, 2011; Schmitt *et al.*, 2012; Giles *et al.*, 2013; Gloeckner *et al.*, 2013a; Poppell *et al.*, 2013; Moitinho-Silva *et al.*, 2014). Each LMA species was dominated by a large clade of Proteobacteria (Alpha-, Beta-, or Gamma-) or Cyanobacteria (genus *Synechococcus*), and there was little overlap between the LMA sponge microbiomes under investigation. The HMA sponge communities, however, showed more phylum-level diversity, with Proteobacteria, Chloroflexi, Acidobacteria, Actinobacteria, candidate phylum Poribacteria, and other phyla as dominant community members. Differences extend also to the microbial physiology of the respective microbial consortia, particularly with respect to nitrogen metabolism (Bayer *et al.*, 2008; Schlappy *et al.*, 2010; Ribes *et al.*, 2012). Distinct differences were further noted regarding the

distribution of certain polyketide synthase genes (*sup-APKS*) that were found in all HMA but not in the LMA sponges under investigation (Hochmuth *et al.*, 2010). Clearly, much remains to be learned about the metabolism, physiology, and function of sponge-associated microbial consortia, particularly in the context of the HMA or LMA dichotomy.

With respect to animal morphology, Vacelet and Donadey (1977) observed early on that the HMA sponges generally display a denser mesohyl, narrower aquiferous canals, and smaller choanocyte chambers than their LMA counterparts. In other words, the HMA sponges appear to be less well “irrigated.” The narrower canals and smaller choanocyte chambers may result in a reduced water flow when compared to LMA sponges (Weisz *et al.*, 2008; Schlappy *et al.*, 2010). In the present study, we revisited the original observations on microbial abundances in sponges; we used electron microscopy complemented by DAPI cell-counting. Altogether, 56 demosponge species from four different geographic locations were investigated. This survey significantly expands previously published datasets on HMA and LMA sponges (Hentschel *et al.*, 2003; Schmitt, 2007; Schmitt *et al.*, 2008a; Weisz *et al.*, 2008), and the findings are interpreted in a taxonomic framework of the animal hosts.

Materials and Methods

Sponge collection

Scuba divers from the research vessels R/V *Seward Johnson* and *F. G. Walton Smith*, Harbor Branch Oceanographic Institution, Florida, collected sponges from various locations around the Bahamas (termed “BAH”) over the years 2003–2013 (Table 1). Sponges were further obtained by scuba diving in 2004 in Key Largo, Florida, USA (25°01'N, 80°23'W) (“FL”); offshore Rovinj, Croatia (45°08'N, 13°64'E) (“MED”) in 2012; at Souda Bay, Crete, Greece (35°31'N, 24°09'E) (“MED”) in 2013; and at other Mediterranean locations. Sponge samples were further collected at Fsar reef, Thuwal, Red Sea, Saudi Arabia (22°23'N, 39°03'E) (“RS”) in 2010 (Table 1). Three to five sponge individuals were collected per species and brought to the surface in separate resealable plastic bags. The samples were processed as described below within a few hours after sampling.

Electron microscopy

Freshly collected sponge material of a few cubic millimeters in size was fixed in 2.5% glutaraldehyde/phosphate-buffered saline for 12 h, rinsed three times for 20 min each in PBS, and postfixed in 2% osmium tetroxide/PBS for 12 h. Several pieces per individual (hereafter termed “technical replicates”) were dehydrated in an ethanol series (30%, 50%, 70%, 90%, 3×100%), incubated three times for 20

Table 1

Sponge collection sites

| Location | Collection Site | Sponge Species |
|----------------------|--|---|
| BAH, Bahamas Islands | Little San Salvador 24°34.39'N; 75°58.00'W | <i>Agelas citrina</i> , <i>Agelas dilatata</i> , <i>Aiolochoiria crassa</i> , <i>Siphonodictyon coralliphagum</i> , <i>Iotrochota birotulata</i> , <i>Plakortis</i> sp. |
| | San Salvador 24°01.14'N; 74°32.68'W | <i>Agelas dispar</i> , <i>Aplysina cauliformis</i> var. thick, <i>Aplysina insularis</i> , <i>Verongula gigantea</i> , <i>Ptilocaulis</i> sp., <i>Cliona varians</i> |
| | Sweetings Cay 26°36.0'N; 77°52.60'W | <i>Cribrochalina vasculum</i> , <i>Dysidea etheria</i> |
| | Chub Cay 25°23.365'N; 77°52.127'W | <i>Aplysina cauliformis</i> var. thin, <i>Monanchora arbuscula</i> |
| | Grand Bahama Island 25°1.282'N; 77°34.56'W | <i>Aplysina archeri</i> , <i>Aplysina fistularis</i> , <i>Chalinula moliitba</i> |
| | Exuma Cay 24°25.642'N; 76°40.464'W | <i>Myrmekioderma gyroderma</i> , <i>Sphaciospongia vesparium</i> , <i>Erylus formosus</i> |
| MED, Mediterranean | Bimini 25°45.316'N; 79°18.061'W | <i>Batzella rubra</i> , <i>Cinachyrella alloclada</i> |
| | Great Inagua 21°05.945'N; 73°40.216'W | <i>Plakortis lita</i> , <i>Svenzea zeai</i> |
| | Cat Cay 25°31.480'N; 79°17.938'W | <i>Callyspongia plicifera</i> |
| | Rovinj, Croatia 45°05'N, 13°38'E | <i>Aplysina aerophoba</i> , <i>Chondrosia reniformis</i> , <i>Axinella polypoides</i> , <i>Tethya aurantium</i> , <i>Suberites domuncula</i> , <i>Dysidea avara</i> |
| | Marseille, France 43°11'48.92'N; 5°21'48.62'E | <i>Aplysina cavernicola</i> , <i>Oscarella lobularis</i> |
| FL, Florida | Banyuls-sur-Mer, France 42°29'N; 03°08'E | <i>Crambe crambe</i> |
| | Souda, Crete, Greece 36°76.759'N; 24°51.422'E | <i>Petrosia</i> sp., <i>Acanthella acuta</i> , <i>Axinella cannabina</i> |
| FL, Florida | Key Largo, USA 24°56.863'N; 80°27.230'W | <i>Agelas wiedenmayeri</i> , <i>Xestospongia muta</i> , <i>Ircinia felix</i> , <i>Ircinia strobilina</i> , <i>Smenospongia aurea</i> , <i>Ecytoplasia ferox</i> , <i>Scopalina ruetzleri</i> , <i>Tedania ignis</i> , <i>Mycale laxissima</i> , <i>Niphates erecta</i> , <i>Niphates digitalis</i> , <i>Amphimedon compressa</i> , <i>Callyspongia vaginalis</i> , <i>Aplysina lacunosa</i> |
| Red Sea | Fsar Reef, Jeddah, Saudi Arabia 22°23.096'N; 39°02.856'E | <i>Amphimedon ochracea</i> , <i>Crella cyathophora</i> , <i>Stylissa carteri</i> , <i>Xestospongia testudinaria</i> |

min each in propylene oxide and polymerized in Epon 812 (Serva, Germany) for 4 days at 60 °C. The embedded sponge pieces were sectioned with an ultramicrotome (OM U3, C. Reichert, Austria). For contrasting, 70–80-nm thick sections were post-stained with 0.5% uranyl acetate in methanol for 10 min and Reynolds lead citrate for 5 min. The resulting sections were investigated by electron microscopy (Zeiss EM 10, Zeiss, Germany). Several different images of three biological specimens were inspected for each species.

Bacterial quantification protocol

A piece of sponge tissue of about 1 g was removed with an ethanol-sterilized scalpel blade and rinsed three times with 0.2- μ m filter-sterilized seawater. The tissue cube was cut so that one side always represented the surface tissue. A 10 \times dilution was obtained by adding 1 ml of sponge tissue to 9 ml of 0.2- μ m filter-sterilized seawater. The tissue was homogenized with a mortar and pestle and poured through Nitex (100- μ m pore size) to remove unground tissue pieces. The suspension was fixed in paraformaldehyde to a final concentration of 3.7% and stored at 4 °C until use. The tissue remainders were rinsed off the Nitex sheet with 0.5-ml filter-sterilized seawater and homogenized, using a

mortar and pestle, in 4.5 ml of filter-sterilized seawater. The suspension was poured through Nitex, fixed in 3.7% paraformaldehyde final concentration, and stored at 4 °C until use. Dilutions ranging from 10⁻¹ to 10⁻³ were prepared from each homogenate. One milliliter of each dilution was stained with 0.7 μ g/ml DAPI (4,6-Diamidino-2-phenylindole) final concentration in the dark for 30 min. The DAPI stock solution (100 μ g ml⁻¹) was prepared weekly and stored at 4 °C. A volume of 1 ml of stained homogenate was added to 9 ml of filter-sterilized seawater and filtered onto a black, 0.2- μ m polycarbonate membrane 25 mm in diameter (Millipore, Germany) that was supported by an 0.45- μ m cellulose nitrate filter (Schleicher und Schuell, Germany). Vacuum (<10 cm Hg) was applied carefully with a hand pump. The filters were washed once with filter-sterilized seawater and subsequently rinsed with 3 ml of 70% ethanol. The 0.2- μ m polycarbonate filter was then mounted with Citifluor (Citifluor Ltd., UK) onto a microscope slide. Bacterial numbers were determined after epifluorescence microscopy using a 100 \times magnification oil lense (AxioLab microscope, Zeiss, Germany). Three independent specimens were processed for each species. Each sample represents an average bacterial number from 10 different counting fields. For each sample, the numbers of bacteria and

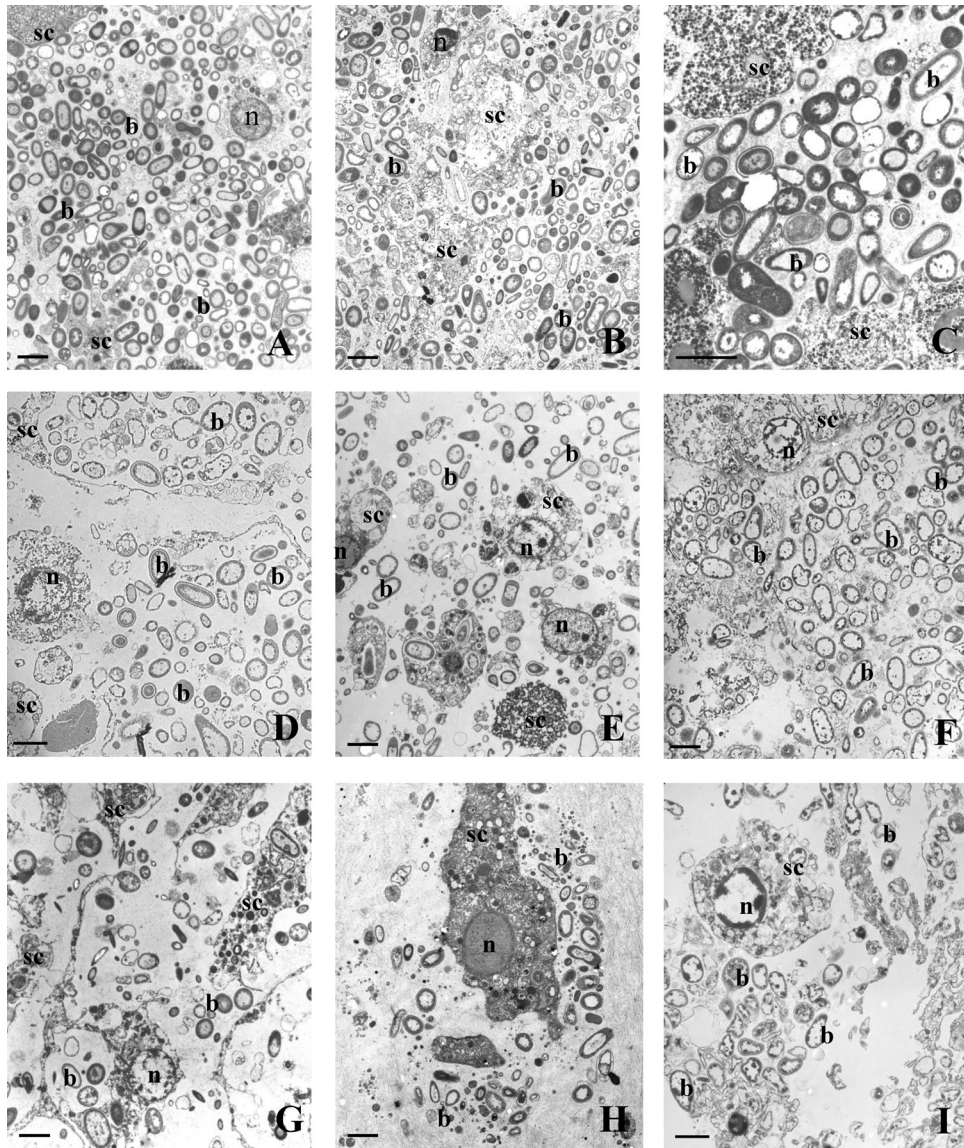


Figure 1. Transmission electron microscopy of selected high microbial abundance sponge species. A = *Aplysina aerophoba*, B = *Spheciospongia vesparium*, C = *Aiolochroia crassa*, D = *Petrosia* sp., E = *Xestospongia testudinaria*, F = *Plakortis lita*, G = *Agelas dispar*, H = *Ircinia felix*, I = *Svenzea zeai*. Scale bar, 2 μ m; b, bacteria; n, nucleus; sc, sponge cell.

nuclei were counted, and the results from the first and second homogenate were summed. The number of cyanobacteria was counted using the red and green fluorescent filter set.

18S rRNA sponge phylogeny

Nearly full-length 18S rRNA gene sequences published in NCBI GenBank (<http://www.ncbi.nlm.nih.gov>) were analyzed. In a few cases, the 18S rRNA gene sequences were unavailable and the sequence of a closely related congeneric (and in the case of *Batzella rubra*, of a confamilial sequence) were therefore substituted, taking the current

changes in demosponge classification into consideration (Redmond *et al.*, 2013). Altogether 45 sponge species were included in the phylogenetic tree, and the GenBank accession numbers are provided in Figure 3. The sequences were aligned using the Sponge Genetree Server with 18S rRNA gene secondary structure information included in the analysis (Erpenbeck *et al.*, 2008). Positions that could not be aligned were excluded from further analyses. Maximum-likelihood reconstruction was inferred with RAxML 7.2.5 (Stamatakis, 2006) using the GTRGAMMI model of nucleotide substitution as suggested by jModeltest ver. 0.1.1. (Darriba *et al.*, 2012) under the Akaike Information

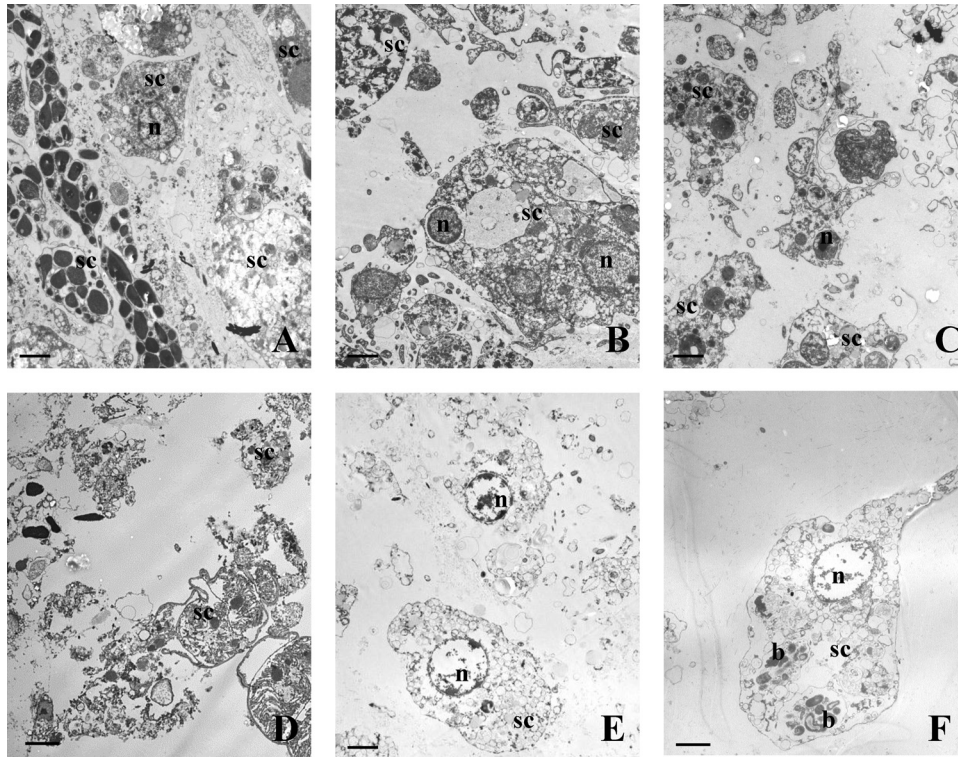


Figure 2. Transmission electron microscopy of selected low microbial abundance sponges. A = *Erylus formosus*, B = *Monanchora arbuscula*, C = *Scopalina ruetzleri*, D = *Amphimedon ochracea*, E = *Axinella cannabina*, F = *Acanthella acuta*. Scale bar, 2 μm ; b, bacteria; n, nucleus; sc, sponge cell.

Criterion (Akaike, 1974) and 100 fast bootstrap replicates. Analyses were performed on the 64-node Linux cluster of the Molecular Geo- and Palaeobiology Labs, Ludwig Maximilian University, Munich.

Results and Discussion

In the present study, 56 sponge species were inspected by transmission electron microscopy (TEM) for the presence of microorganisms in the mesohyl matrix. Of the 41 sponge species examined from the greater Caribbean (Bahamas and Florida locations, Table 1), 24 were identified as high microbial abundance (HMA) and 17 as low microbial abundance (LMA) (Tables 2, 3). Of the 12 species sampled from the various Mediterranean locations, four were identified as HMA and eight as LMA. From the Red Sea collection site, one sponge was identified as HMA and three as LMA. For previous TEM-based surveys of HMA and LMA sponge patterns, the reader is referred to the literature: Hentschel *et al.* (2003) and references cited therein; Schmitt *et al.* (2008b); Weisz *et al.* (2008); Popell *et al.* (2013). The question arises whether the sponges fall into strict HMA or LMA categories or whether they rather represent a continuum. Judging from our data, the HMA-LMA dichotomy may be best described as a continuum with a highly bimodal distribution, in the sense that most investigated

species are found at the extreme ends (either HMA or LMA) of the continuum.

The HMA sponge tissues contained dense and morphologically diverse microorganisms that are located largely extracellularly in the mesohyl matrix (Fig. 1A–I). In extreme cases, the microbial cells are much more abundant than sponge cells, as shown here for *Aplysina aerophoba* (Fig. 1A) and *Sphaciospongia vesparium* (Fig. 1B) as well as elsewhere, for example, for the order Verongida (Vacelet, 1975; Friedrich *et al.*, 1999). Our data support the results of Weisz *et al.* (2008) but disagree with those of Popell *et al.* (2013) in that *S. vesparium* is classified here as an HMA sponge. While the potential for phenotypic plasticity in microbial amount cannot be ruled out, we have found microbial abundance to be a highly conserved trait on the level of sponge species.

In general, there is a remarkably stable presence of certain bacterial morphotypes, of which some appear to contain intracellular compartments and unusual membrane structures (Fig. 1C). Some of these originally described morphotypes (*i.e.*, types A and C according to Vacelet, 1975, and Friedrich *et al.*, 1999) can readily be identified in the present pictures of HMA sponge tissue. The amount of bacteria in the HMA sponge tissues can, however, be variable, ranging from densely packed mesohyl tissues for the order

Table 2

A compilation of high microbial abundance (HMA) sponge species

| Species | Collection Site* | Transmission Electron Microscopy Reference† | Higher Taxon§ |
|--|------------------|---|------------------------|
| <i>Agelas citrina</i> | BAH | Wehrl 2006 | Agelasida |
| <i>Agelas dilatata</i> | BAH | Wehrl 2006 | Agelasida |
| <i>Agelas dispar</i> | BAH | This study | Agelasida |
| <i>Agelas wiedenmayeri</i> | FL | Wehrl 2006; Schmitt <i>et al.</i> 2008b | Agelasida |
| <i>Aiolochoxia crassa</i> | BAH | This study | Verongida |
| <i>Aplysina aerophoba</i> | MED | Hentschel <i>et al.</i> 2003; Siegl <i>et al.</i> 2008 | Verongida |
| <i>Aplysina archeri</i> | BAH | Wehrl 2006 | Verongida |
| <i>Aplysina cauliformis</i> , thick morphotype | BAH | Wehrl 2006 | Verongida |
| <i>Aplysina cauliformis</i> , thin morphotype | BAH | Wehrl 2006 | Verongida |
| <i>Aplysina cavernicola</i> | MED | Wehrl 2006; Friedrich <i>et al.</i> 1999, 2001 | Verongida |
| <i>Aplysina fistularis</i> | BAH | Wehrl 2006; Gloeckner 2013 | Verongida |
| <i>Aplysina insularis</i> | BAH | Wehrl 2006 | Verongida |
| <i>Aplysina lacunosa</i> | FL | Wehrl 2006 | Verongida |
| <i>Chondrosia reniformis</i> | MED | Wehrl 2006 | Chondrosida |
| <i>Cribrochalina vasculum</i> | BAH | Schiller 2006 | (marine) Haplosclerida |
| <i>Ectyoplasia ferox</i> | FL | Schmitt <i>et al.</i> 2008a,b; Gloeckner <i>et al.</i> 2013 | Raspailiidae |
| <i>Ircinia felix</i> | FL | Schmitt <i>et al.</i> 2007 | Dictyoceratida |
| <i>Ircinia strobilina</i> | FL | Schmitt 2007 | Dictyoceratida |
| <i>Myrmekioderma gyroderma</i> | BAH | Gloeckner 2013 | “Halichondrida” |
| <i>Petrosia</i> sp. | MED | This study | (marine) Haplosclerida |
| <i>Plakortis lita</i> | BAH | This study | Homoscleromorpha |
| <i>Plakortis</i> sp. | BAH | Laroche <i>et al.</i> 2007 | Homoscleromorpha |
| <i>Siphonodictyon coralliphagum</i> | BAH | Schiller 2006; Schmitt <i>et al.</i> 2008b | (marine) Haplosclerida |
| <i>Smenospongia aurea</i> | FL | Schmitt <i>et al.</i> , 2008b; Gloeckner 2013 | Dictyoceratida |
| <i>Sphaciospongia vesparium</i> | BAH | This study | “Hadromerida” |
| <i>Svenzea zeai</i> | BAH | This study | “Halichondrida” |
| <i>Verongula gigantea</i> | BAH | Wehrl 2006 | Verongida |
| <i>Xestospongia muta</i> | FL | Wehrl 2006; Hentschel <i>et al.</i> 2006 | (marine) Haplosclerida |
| <i>Xestospongia testudinaria</i> | RS | This study | (marine) Haplosclerida |

* BAH, Bahama Islands; FL, Florida; MED, Mediterranean; RS, Red Sea.

† Some transmission electron microscopy data were reported in Master’s (Schiller, 2006) and Ph.D. theses (Wehrl, 2006; Schmitt, 2007; Gloeckner, 2013).

§ Higher taxon names in quotation marks indicate orders recognized as non-monophyletic which may be subject to emendings in the future.

Verongida (Fig. 1A) to moderately dense microbial consortia such as those of the taxa *Ircinia* or *Agelas* (Fig. 1G, H). However, additional literature reports based on 16S rRNA gene sequencing and inspections of microbial abundances in larval tissues clearly identified these species as HMA sponges (Schmitt *et al.*, 2007, 2008a,b; Schmitt, 2007). Experimental artefacts may have arisen for sponges that were difficult to cut with a diving knife (*i.e.*, *Ircinia*), or for sponges having tissues that are less cohesive (*i.e.*, *Xestospongia*). However, the consistency between technical replicates (representing samples from the same individual) and between biological replicates (representing samples from different individuals) is remarkably high in our experience.

Relative to HMA sponge species, the mesohyl of LMA sponges was noticeably devoid of microorganisms (Fig. 2). A few intracellular bacteria in various stages of digestion were present that likely represent food bacteria (Fig. 2F). Occasionally, bacterial morphotypes were observed in TEM pictures of LMA sponges (Gloeckner *et al.*, 2013a); how-

ever, the morphotype diversity appeared unlike that of HMA sponges in that the compartmentalized cells and those with unusual membranes were missing (Vacelet, 1975; Friedrich *et al.*, 1999). Therefore, on the basis of electron microscopical observations, the combination of bacterial abundance and morphotype diversity determines whether a sponge species belongs to the HMA or LMA category. However, in some sponges with intermediate bacterial abundances, TEM may not be sufficient to determine whether a given species is HMA or LMA, and in these exceptional cases, additional methods are needed.

When the status of a sponge species as either HMA or LMA was equivocal on the basis of microscopical observations, an independent line of evidence was sought in microbial enumeration by staining with DAPI (Table 4). Sponge tissue homogenates were used for bacterial and cyanobacterial quantification, and numbers were either expressed in total or as ratios relative to the number of sponge nuclei. Further, sponge homogenates were screened for the

Table 3

A compilation of low microbial abundance (LMA) sponge species

| Species | Collection Site* | Transmission Electron Microscopy Reference† | Higher Taxon§ |
|-------------------------------|------------------|---|------------------------|
| <i>Acanthella acuta</i> | MED | This study | “Halichondrida” |
| <i>Amphimedon compressa</i> | FL | Angermeier <i>et al.</i> 2012 | (marine) Haplosclerida |
| <i>Amphimedon ochracea</i> | RS | This study | (marine) Haplosclerida |
| <i>Axinella cannabina</i> | MED | This study | “Halichondrida” |
| <i>Axinella polypoides</i> | MED | Wehr 2006 | “Halichondrida” |
| <i>Batzella rubra</i> | BAH | Gloeckner 2013 | Poecilosclerida (s.s.) |
| <i>Callyspongia plicifera</i> | BAH | Gloeckner 2013 | (marine) Haplosclerida |
| <i>Callyspongia vaginalis</i> | FL | Schiller 2006; Wehr 2006 | (marine) Haplosclerida |
| <i>Chalinula molitba</i> | BAH | Schiller 2006; Wehr 2006 | (marine) Haplosclerida |
| <i>Cinachyrella alloclada</i> | BAH | Gloeckner 2013 | Spirophorida |
| <i>Cliona varians</i> | BAH | Schiller 2006 | “Hadromerida” |
| <i>Crambe crambe</i> | MED | Wehr 2006 | Poecilosclerida (s.s.) |
| <i>Crella cyathophora</i> | RS | Giles <i>et al.</i> 2012 | Poecilosclerida (s.s.) |
| <i>Dysidea avara</i> | MED | Wehr 2006 | Dictyoceratida |
| <i>Dysidea etheria</i> | BAH | Schiller 2006 | Dictyoceratida |
| <i>Erylus formosus</i> | BAH | This study | Astrophorida |
| <i>Iotrochota birotulata</i> | BAH | Wehr 2006 | Poecilosclerida (s.s.) |
| <i>Monanchora arbuscula</i> | BAH | This study | Poecilosclerida (s.s.) |
| <i>Mycale laxissima</i> | FL | Wehr 2006 | Poecilosclerida (s.s.) |
| <i>Niphates digitalis</i> | FL | Schiller 2006; Wehr 2006 | (marine) Haplosclerida |
| <i>Niphates erecta</i> | FL | Wehr 2006 | (marine) Haplosclerida |
| <i>Oscarella lobularis</i> | MED | Gloeckner <i>et al.</i> 2013 | Homoscleromorpha |
| <i>Ptilocaulis</i> sp. | BAH | Wehr 2006 | “Halichondrida” |
| <i>Scopalina ruetzleri</i> | FL | Wehr 2006; Gloeckner 2013 | “Halichondrida” |
| <i>Stylissa carteri</i> | RS | Giles <i>et al.</i> 2013 | “Halichondrida” |
| <i>Suberites domuncula</i> | MED | Wehr 2006 | “Hadromerida” |
| <i>Tedania ignis</i> | FL | Schiller 2006; Wehr 2006 | Poecilosclerida (s.s.) |
| <i>Tethya aurantium</i> | MED | Wehr 2006 | “Hadromerida” |

* BAH, Bahama Islands; FL, Florida; MED, Mediterranean; RS, Red Sea.

† Some transmission electron micrography data were reported in Master’s (Schiller 2006) and PhD theses (Wehr 2006; Schmitt 2007; Gloeckner 2013).

§ Higher taxon names in quotation marks indicate orders recognized as non-monophyletic and may be subject to emendings in the future.

presence or absence of bacteria, using defined HMA and LMA sponge homogenates as controls. The sponge species subjected to this analysis formed two distinct groups of HMA and LMA sponges, which mirrored the TEM observations. *Siphonodictyon coralliphagum* was the only exception, in which the low bacterial numbers obtained by DAPI staining indicated it was an LMA sponge, but the TEM observations identified it as an HMA sponge (Schmitt, 2007). The DAPI method can be prone to errors; for example, if the bacterial cells are disrupted upon tissue homogenization, particularly as a result of the presence of cytotoxic secondary metabolites. However, this problem happened only once in 56 species investigated and appeared to be a rare scenario. The ratio of bacteria per sponge cell nuclei was variable, ranging from 17 to 95 bacterial cells per sponge cell nucleus for *Xestospongia muta* and *Agelas insularis*, respectively, and from 0 to 5 bacterial cells per sponge cell nucleus for the LMA sponges (Table 4). These data support the TEM observations that HMA sponges contain variable amounts of microorganisms in the mesohyl

tissues. Major disadvantages of the DAPI-staining method were the large amount of background staining and the limited applicability to sponges with high intrinsic autofluorescence. Also, some sponge tissues were difficult to macerate and yielded clumpy homogenates, making it difficult to determine precise numbers. In these cases, confocal microscopy on sponge tissue sections, as recently employed by Ribes and coworkers (Ribes *et al.*, 2012), is a suitable alternative for microbial visualization and quantification in sponge tissues.

Several recent publications addressed the HMA-LMA dichotomy by 16S rRNA gene sequencing. Whether obtained by clone libraries (Kamke *et al.*, 2010; Giles *et al.*, 2013), DGGE (Weisz *et al.*, 2007; Gloeckner *et al.*, 2013a; Poppell *et al.*, 2013), terminal restriction fragment length polymorphism (T-RFLP) (Erwin *et al.*, 2011), or amplicon sequencing (Schmitt *et al.*, 2012; Moitinho-Silva *et al.*, 2014), the data consistently revealed a different bacterial composition in LMA sponges than in HMA sponges. Similarly, in comparative DGGEs, the LMA sponges

Table 4

Quantification of bacteria, cyanobacteria, and sponge nuclei by DAPI staining (per ml sponge homogenate)

| Species | Bacteria Mean \pm SE | Cyanobacteria Mean \pm SE | Bacteria Total Mean \pm SE | Nuclei Mean \pm SE | Ratio* |
|---------------------------------------|---------------------------------------|---------------------------------------|---------------------------------------|---------------------------------------|--------|
| High microbial abundance sponges† | | | | | |
| <i>Agelas citrina</i> | $7.0 \times 10^8 \pm 2.1 \times 10^8$ | $2.6 \times 10^6 \pm 2.0 \times 10^6$ | $7.0 \times 10^8 \pm 2.1 \times 10^8$ | $3.2 \times 10^7 \pm 5.7 \times 10^6$ | 22.16 |
| <i>A. dilatata</i> | $1.9 \times 10^9 \pm 8.7 \times 10^8$ | $1.5 \times 10^6 \pm 9.6 \times 10^5$ | $1.9 \times 10^9 \pm 8.7 \times 10^8$ | $5.7 \times 10^7 \pm 1.7 \times 10^7$ | 33.40 |
| <i>Aplysina archeri</i> | $1.4 \times 10^9 \pm 1.9 \times 10^7$ | $1.1 \times 10^7 \pm 5.8 \times 10^6$ | $1.4 \times 10^9 \pm 2.4 \times 10^7$ | $3.5 \times 10^7 \pm 1.8 \times 10^7$ | 39.16 |
| <i>A. insularis</i> | $6.2 \times 10^9 \pm 4.2 \times 10^8$ | $2.9 \times 10^8 \pm 7.3 \times 10^7$ | $6.5 \times 10^9 \pm 4.9 \times 10^8$ | $6.8 \times 10^7 \pm 2.2 \times 10^6$ | 95.71 |
| <i>Cribochalina vasculum</i> | $5.5 \times 10^9 \pm 2.3 \times 10^9$ | $1.1 \times 10^9 \pm 1.7 \times 10^7$ | $6.6 \times 10^9 \pm 2.4 \times 10^7$ | $2.1 \times 10^8 \pm 3.3 \times 10^7$ | 31.71 |
| <i>Xestospongia muta</i> | $7.0 \times 10^9 \pm 5.5 \times 10^8$ | $1.2 \times 10^9 \pm 2.8 \times 10^8$ | $8.2 \times 10^9 \pm 7.7 \times 10^8$ | $4.8 \times 10^8 \pm 4.1 \times 10^7$ | 16.96 |
| <i>Ircinia felix</i> | $1.0 \times 10^9 \pm 7.3 \times 10^7$ | $5.7 \times 10^8 \pm 7.1 \times 10^6$ | $1.6 \times 10^9 \pm 7.4 \times 10^7$ | $6.1 \times 10^7 \pm 1.2 \times 10^7$ | 25.67 |
| <i>Plakortis</i> sp. | $4.3 \times 10^9 \pm 8.0 \times 10^8$ | $3.6 \times 10^7 \pm 1.6 \times 10^7$ | $4.3 \times 10^9 \pm 7.9 \times 10^8$ | $4.8 \times 10^7 \pm 1.6 \times 10^7$ | 89.23 |
| <i>Ectyoplasia ferox</i> | $8.7 \times 10^9 \pm 1.7 \times 10^8$ | n.d. | $8.7 \times 10^9 \pm 1.7 \times 10^8$ | $3.8 \times 10^8 \pm 2.6 \times 10^7$ | 22.94 |
| Low microbial abundance sponges‡ | | | | | |
| <i>Iotrochota birotulata</i> | $2.4 \times 10^9 \pm 2.8 \times 10^8$ | n.d. | $2.4 \times 10^9 \pm 2.8 \times 10^8$ | $5.2 \times 10^8 \pm 4.3 \times 10^7$ | 4.67 |
| <i>Siphonodictyon coralliphagum</i> § | $2.0 \times 10^8 \pm 1.5 \times 10^7$ | n.d. | $2.0 \times 10^8 \pm 1.5 \times 10^7$ | $2.2 \times 10^8 \pm 6.2 \times 10^7$ | 0.93 |
| <i>Dictyonella funicularis</i> | n.d. | n.d. | n.d. | $3.2 \times 10^8 \pm 1.4 \times 10^7$ | 0.00 |
| <i>Tedania ignis</i> | n.d. | n.d. | n.d. | $8.9 \times 10^8 \pm 5.2 \times 10^7$ | 0.00 |
| <i>Chalinula molitba</i> | $3.5 \times 10^6 \pm 3.5 \times 10^6$ | n.d. | $3.5 \times 10^6 \pm 3.5 \times 10^6$ | $6.1 \times 10^8 \pm 7.4 \times 10^7$ | 0.01 |
| <i>Niphates digitalis</i> | n.d. | n.d. | n.d. | $4.6 \times 10^8 \pm 1.3 \times 10^8$ | 0.00 |
| <i>Amphimedon compressa</i> | n.d. | n.d. | n.d. | $1.4 \times 10^8 \pm 2.3 \times 10^7$ | 0.00 |
| <i>Callyspongia vaginalis</i> | $2.2 \times 10^6 \pm 1.6 \times 10^6$ | $1.8 \times 10^6 \pm 1.8 \times 10^6$ | $4.0 \times 10^6 \pm 2.0 \times 10^6$ | $5.5 \times 10^8 \pm 1.1 \times 10^7$ | 0.01 |
| <i>Ptilocaulis</i> sp. | n.d. | n.d. | n.d. | $3.7 \times 10^8 \pm 3.2 \times 10^7$ | 0.00 |
| <i>Cliona varians</i> | n.d. | n.d. | n.d. | $2.0 \times 10^8 \pm 2.1 \times 10^7$ | 0.00 |
| <i>Dysidea etheria</i> | $3.1 \times 10^6 \pm 1.2 \times 10^6$ | $1.3 \times 10^6 \pm 7.6 \times 10^5$ | $4.4 \times 10^6 \pm 1.9 \times 10^6$ | $1.9 \times 10^8 \pm 8.1 \times 10^6$ | 0.02 |
| Caribbean seawater | | | $3.5 \times 10^5 \pm 5.7 \times 10^4$ | | |

* Ratio of total bacteria/nuclei.

† The following additional sponges were identified as HMA by qualitative DAPI screening: *Ectyoplasia ferox*, *Myrmekioderma gyroderma*, *Agelas dispar*, *Ircinia felix*, *Ircinia strobilina*, *Smenospongia aurea*, *Aplysina cauliformis*, *Aplysina fistularis*, *Verongula gigantea*, *Aiolochroia crassa*, *Geodia neptuni*, *Chondrosia collectrix*, *Calyx podatypa*.‡ The following additional sponges were identified as LMA by qualitative DAPI screening: *Erylus formosus*, *Mycale (Arenochalina) laxissima*, *Batzella rubra*, *Monanchora arbuscula*, *Scopalina ruetzleri*, *Callyspongia plicifera*, *Callyspongia vaginalis*, *Cinachyrella alloclada*.§ For *Siphonodictyon coralliphagum*, the low bacterial numbers determined by DAPI staining contradict its status as an HMA sponge, as determined by transmission electron microscopy (Schmitt *et al.*, 2008b).

n.d. = not detected.

consistently displayed less complex banding patterns than their HMA counterparts (Weisz *et al.*, 2007; Gerce *et al.*, 2011; Poppell *et al.*, 2013). Even small 16S rRNA gene clone libraries were sufficient to detect the major phylogenetic lineages (Moitinho-Silva and Hentschel, unpubl. data). 16S rRNA gene data have therefore proven to be very useful to infer the HMA or LMA status of the host sponge.

The host phylogeny based on nearly full-length 18S rRNA gene sequences corroborates the current molecular phylogenetic hypotheses of demosponges (*e.g.*, Morrow *et al.*, 2012; see also Redmond *et al.* (2013) for the most comprehensive 18S rRNA gene phylogeny). However, as the new nomenclature is yet to be finalized, we refer to taxon names as currently used in the World Porifera Database (van Soest *et al.*, 2014). Several clades were recovered that in the present sample set consisted exclusively of HMA taxa (*Verongida*, *Agelasida (Agelas)*), while the *Poecilo-*

sclerida (sensu stricto) clade with representatives from six families consisted exclusively of LMA species (Fig. 3). For the remaining clades, of which many orders and families are currently under redefinition (see, *e.g.*, Redmond *et al.*, 2013), distinct HMA/LMA distribution patterns were not recognized. Additional and distinct patterns might be revealed on lower taxonomic levels given a more representative taxonomic sampling. It is noteworthy, however, that the HMA/LMA characteristics are conserved in closely related species over time and space when collected from different geographic regions (Wilkinson, 1978; Montalvo and Hill, 2011).

The question remains open as to what causes the HMA-LMA dichotomy. A survey of the literature revealed no apparent correlation with host defense status (Chanas and Pawlik, 1995; Pawlik *et al.*, 1995), with reproductive mode (oviparous *vs.* viviparous), or with ecological parameters, as

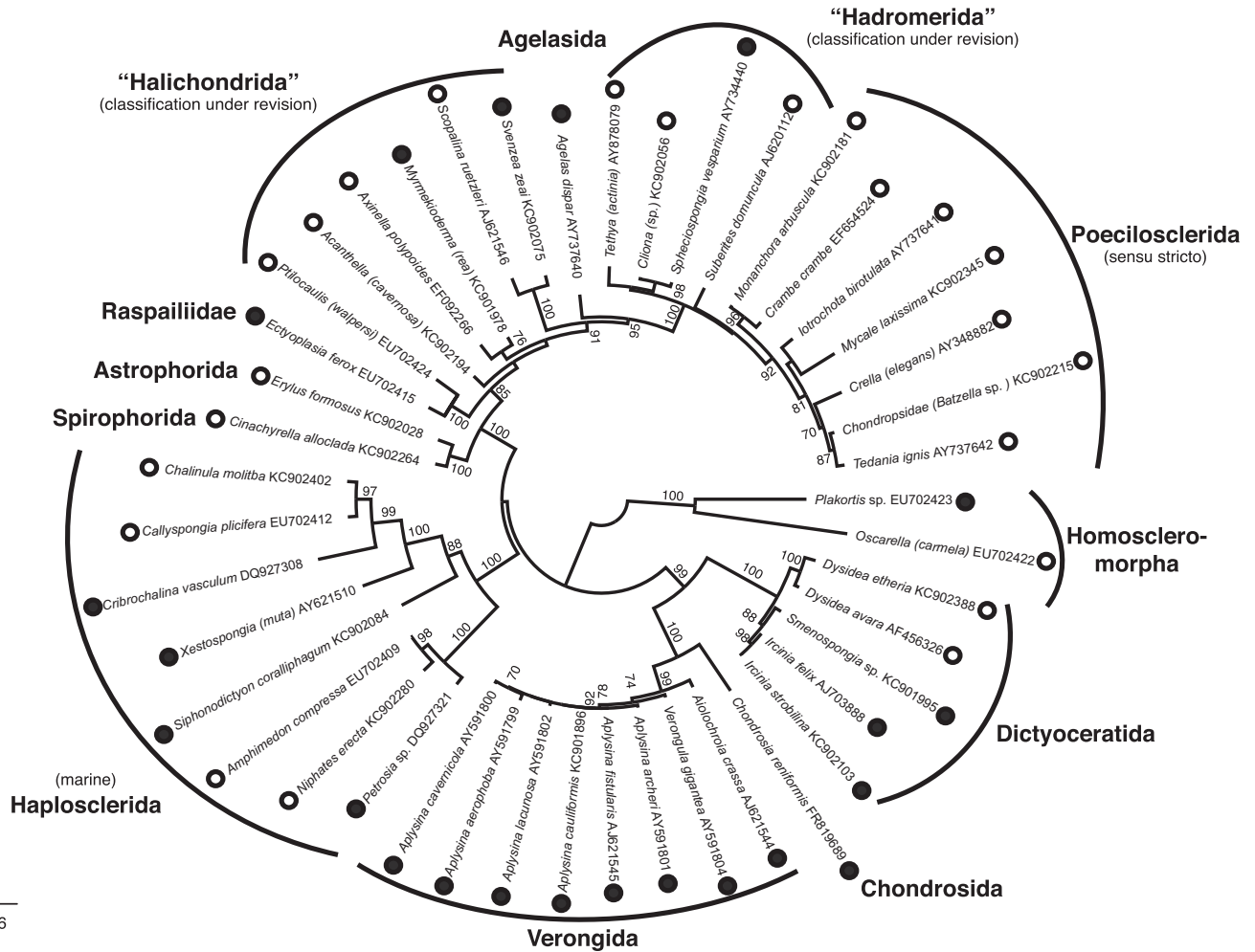


Figure 3. Maximum-likelihood reconstruction of the sponge hosts using almost complete 18S rRNA gene sequences as a phylogenetic marker. Taxon names in parentheses describe the closest related taxon that was used when the 18S rRNA gene sequence of a given host species was unavailable. Numbers at the branches are bootstrap probabilities > 70%. The taxon names are followed by the NCBI GenBank accession numbers. Black filled dots depict high microbial abundance sponges; unfilled dots, low microbial abundance sponges. Scale bar, substitutions/site.

both types of sponges coexist in the same habitat (Schiller, 2006). A phylogenetic signature is present only to a moderate extent. At this point it appears most likely that sponge morphology is an important determining factor. The HMA sponges are frequently large and massive and generally have a firm touch and fleshy consistency, while LMA sponges are generally smaller and feel fragile, soft, brittle, or tough (U.H., pers. obs.). Similarly, the architecture of the sponge interior plays a determining role (Vacelet and Donadey, 1977; Weisz *et al.*, 2008). A higher choanocyte chamber density in LMA sponges was reported by Poppell (2013) and Schlappy (2010). A current hypothesis states that LMA sponges invest more energy into feeding structures, whereas the nutrition of HMA sponges is supplemented by their microbial symbionts (Poppell *et al.*, 2013). With respect to the postulated role of symbionts in the “sponge loop” (de

Goeij *et al.*, 2013), an improved understanding of the different feeding strategies of the sponge “holobiont” is clearly a worthwhile undertaking. It is further safe to speculate that HMA sponges are morphologically adapted to house microbial consortia within their tissues. In evolutionary terms, the question arises whether the sponges are preconditioned to host microbes or whether the animal tissue morphology is a consequence of it containing the microbes. If preconditioned, it is conceivable that the extracellular matrix (ECM) of HMA sponges may be structurally altered to accommodate the presence of sponge symbionts; and *vice versa*, that the sponge symbionts may have mechanisms to survive within ECM, modifying it with their activities. Indeed, a recent study provides compelling evidence obtained by single-cell genomics that poribacterial sponge symbionts can degrade ECM for nutritional purposes (Kamke *et al.*, 2013).

Although the present study is the most comprehensive survey for HMA and LMA patterns to date, more investigations are needed with greater taxonomic depth, including analysis of specimens from other sponge classes besides Demospongiae; and including more locations, such as the Great Barrier Reef, the deep-sea, and the polar seas. With regard to methodologies, we propose a combination of transmission electron microscopy and 16S rRNA gene sequence data to reliably determine the HMA or LMA status of the host sponge. The latter is particularly recommended when TEM data yield ambiguous results. The present sponge survey will help pave the way for a functional understanding of the HMA-LMA dichotomy in sponges.

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