16S rRNA phylogeny of sponge-associated cyanobacteria

Laura Steindler\textsuperscript{1,2,*}, Dorothée Huchon\textsuperscript{2}, Adi Avni\textsuperscript{1} and Micha Ilan\textsuperscript{2}

\textsuperscript{1} Department of Plant Sciences
\textsuperscript{2} Department of Zoology

George S. Wise Faculty of Life Sciences, Tel Aviv University, Tel Aviv 69978, Israel

Running title: Phylogeny of sponge-associated cyanobacteria

Keywords: Sponge; Cyanobacteria; 16S rRNA; RFLP; phylogeny.

(*) Corresponding author: Tel = 972-3-6409125 / Fax = 972-3-6409380 / e-mail address: lauras@post.tau.ac.il
ABSTRACT

Phylogenetic analyses of 16S rDNA sequences of sponge-associated cyanobacteria showed them to be polyphyletic, implying that they derived from multiple independent symbiotic events. Most of the symbiont sequences were affiliated to a group of *Synechococcus* and *Prochlorococcus* species. However, other symbionts were related to different groups, such as the Oscillorariales.
Although both cyanobacteria and sponges have a very long evolutionary history (1,3), little is known about the identity and phylogeny of cyanobacterial sponge symbionts. For marine sponges symbiosis with cyanobacteria can be obligate or non-obligate according to the sponge species (2). Thus far, all attempts of culturing sponge-associated cyanobacteria have failed (unpublished data, 13). It is thus not known whether sponge-associated cyanobacteria are able to survive outside their host. The most common sponge-associated cyanobacterium was not found in water samples (13, 15). In *Chondrilla australiensis* this cyanobacterium is transmitted vertically (through sponge eggs) (14). Additionally, a study showing co-speciation between *Dysidea* species and their associated cyanobacteria (12) supports the hypothesis that sponges and associated cyanobacteria are coevolving. Few studies sequenced the 16S rRNA gene of sponge-associated cyanobacteria: Diaz (5) identified the first two cyanobacteria from marine sponges, Webb and Maas (16) found that the cyanobacteria inhabiting *Mycale hentscheli* were phylogenetically related to *Cyanobacterium stanieri* and species of *Synechocystis* and *Prochloron*, and Hentschel et al. (7) found that 7 sequences from cyanobacteria inhabiting the sponges *Aplysina aerophoba* and *Theonella swinhoei* could be divided into two clades (*Synechococcus/Prochlorococcus* and *Pleurocapsa*). The aim of the present study was to increase our understanding on the diversity of sponge-associated cyanobacteria, and to determine their phylogenetic position. Sponge samples from 16 species were collected from four locations: The Caribbean (Bahamas, 26°33’N, 77°52’W), Mediterranean (Rapallo, Italy, 44°18’N, 9°12’E), Red Sea (Elat, 31°35’N, 34°54’E), and Western Indian Ocean (Zanzibar, 06°09’S, 39°11’E). Aposymbiotic specimens (specimens that do not contain cyanobacterial symbionts), growing in dark caves or overhangs, were also collected.
for two sponge species (*Petrosia ficiformis* and *Xestospongia muta*). Those samples, collected at short distances from symbiotic specimens, were used as negative controls, to ensure that 16S rDNA sequences were derived from true symbionts and not from surface associated cyanobacteria or digested cyanobacteria. Photosynthetic activity inside the living sponge tissue was tested by pulse amplitude modulated fluorometry (Diving PAM, Walz, Germany). Tissue samples (1 cm³) were rinsed twice in 100% EtOH, and kept in 100% EtOH at 4 ºC. DNA was extracted following the procedure of Bernatzky and Tanksley (4). 16S rDNA was amplified with the primers 361F (5’-GAATTTCGCAATGGGC -3’) and 1459R (5’- GGTAAYGACTTCGGGCRT -3’) (5). 1060-bp fragments were cloned in the PTZ57R/T vector (Fermentas). Twenty clones per individual were amplified using M13 universal primers. The PCR products were digested with restriction enzymes *Apa*I and *Hae*III. One clone was sequenced for each pattern present in more than 10% of the clones. The sequences were deposited in Genbank (accession numbers AY701287-AY701315).

Neither photosynthetic activity, nor amplification of the 16S rRNA gene was obtained for aposymbiotic specimens. On the opposite, photosynthetic activity was recorded for all the other samples. Up to three different cyanobacterial clones were sequenced per individual sponge investigated (with a sequence homology of 90-99.7%). Similarly, Webb and Maas (16) showed four closely related (99.1-99.8%) cyanobacterial clones in the sponge *Mycale hentscheli*. Different cyanobacterial types from the same individual were usually phylogenetically close to each other (<1-2% sequence divergence), but in one case (*Lendenfeldia dendyi*) the symbionts were very divergent: two types were in group 5, while the third type was in group 2, with 10% sequence divergence from the former two (Fig. 1). Closely related cyanobacterial
types should not be the result of Taq-polymerase errors or cloning-bias since only patterns present in more than one clone were sequenced. However, the true cyanobacterial diversity in sponges might be underestimated. Nevertheless our results are in agreement with other studies (13, 16). For example, Usher et al. (13) also identified only one cyanobacterial type in the sponge Petrosia ficiformis.

A maximum likelihood (ML, (6)) analysis was done based on 147 taxa and 1396 nucleotides. The sequences were chosen to include: 1- representatives of the cyanobacteria diversity; 2- all sponge-associated cyanobacteria sequences overlapping with the new 16S rDNA sequences; and 3- representatives of the Synechococcus and Prochlorococcus diversity because sponge symbionts have been suggested to belong to these genera. BLAST searches were also conducted for each sponge sequence in order to include in the analysis the most similar sequences available in the databanks. The GTR model with a gamma distribution (4 categories), and a proportion of invariant site was found to be the most appropriate using Modeltest version 3.06 (9)).

The ML tree was reconstructed in an iterative way using PAUP* version 4.0b10 (11). First a heuristic search was conducted using the ML parameters identified by Modeltest; Modeltest topology as starting tree and NNI branch swapping. The topology found at the end of the search was used to identify new parameters. These new parameters and topology were used to conduct a new search till convergence. The best ML parameters found were then used to compute 500 bootstrap replicates starting with a NJ tree and setting the maximum number of tree to one to reduce computation time.

The phylogenetic tree obtained in this study was in agreement with the results of other studies that divided cyanobacteria into seven or eight major lineages (8, 10), except that group 2 (8) was here paraphyletic (Fig. 1). Sponge-associated
cyanobacteria were found to be polyphyletic. They were divided into thirteen lineages spread among various groups of cyanobacteria (Fig. 1 and 2). The major clade of sponge-associated cyanobacteria (37 strains from 18 different sponge species collected from a wide range of geographic regions: Australia, Caribbean, French and Italian Mediterranean coast, Red Sea, Zanzibar) was strongly affiliated with group 6 (Prochlorococcus, Synechococcus and Microcystis, (8)). Three additional sponge-associated cyanobacteria (from Chondrilla spp. and Haliclona sp.) were also part of group 6, but were closer to free-living Synechococcus than to the other sponge symbionts (Fig. 2). A second group of sponge-associated cyanobacteria (consisting of four sequences from Dysidea herbacea, Dysidea granulosa, Lendenfeldia dendyi and Aplysina gerardogreeni, originating from Guam, Mexican Pacific and Zanzibar) were affiliated to group 2 (morphologically classified in the order Oscillatoriales). A few other sponge-associated cyanobacteria were affiliated with group 5 (Chroococcales, Oscillatoriales, Pleurocapsales and Prochlorales, (8)), but they did not form a monophyletic group (Fig. 1). One additional cyanobacterium (AF295635) from an Australian sponge was found affiliated to the marine group 7 (8). Finally, a sequence of a cyanobacterium associated with the sponge Pseudoaxinella flava appeared unrelated to any major group of cyanobacteria (Fig. 1). The polyphyletic origin of sponge-associated cyanobacteria indicates that they derived from multiple independent symbiotic events, involving several different cyanobacterial types and/or host sponges.

The phylogenetic relationships between the cyanobacterial sequences of group 6 suggest only a partial specificity of cyanobacteria to the host sponges. Two specimens of Petrosia ficiformis, one collected in Italy (this study), and another in France (13), were found to be 99% identical, and closely related (BP = 100%). Three
cyanobacterial types from two Bahamian *Xestospongia muta* specimens were strongly
affiliated to each other and with that from the congeneric species *Xestospongia
proxima* (BP= 100%). Cyanobacteria from two congeneric *Theonella* species
(*swinhoei* and *conica*) were affiliated to each other (BP=77%). Finally, six
cyanobacteria associated with *Chondrilla australiensis* (13, 15) were closely affiliated
(BP = 49%). These results may also support the idea of co-evolution between sponges
and their symbionts. However, cyanobacteria associated to *Theonella swinhoei* from
Japan and the Red Sea clustered in different groups (Fig. 1 and 2). In this large-scale
phylogenetic analysis, sponge-associated cyanobacteria clades deriving from distant
geographic location (as the clades in group 2 and 6) are most probably true-
symbionts, while the symbiotic nature of sponge cyanobacteria that are closely related
to free living species (*e.g.* those in *Pseudoaxinella flava*) remains uncertain.

In the future it will be interesting to examine whether cyanobacterial sponge
symbionts from different lineages perform diverse functions in this symbiosis. In
addition, reconstruction of sponge molecular phylogeny will enable to test for co-
evolution of cyanobacteria and their host sponges.

This work was supported by grant 2000-321 from the United States-Israel
Binational science foundation (BSF) to M. Ilan, S. Beer and J.R. Pawlik. L. Steindler
received a scholarship from the Rieger Foundation for Environmental Studies that
assisted in the travel costs for sampling sponge specimens. S. Beer is thanked for his
editorial comments of the manuscript, and T. Pupko for providing access to his
computers. We wish to thank J.R. Pawlik (University of North Carolina, Wilmington),
who invited L. Steindler, M. Ilan, and S. Beer to participate in research-cruises aboard
the R/V Seward Johnson, which enabled us to collect many samples for this study.
The staff of the Institute of Marine Sciences (Zanzibar) is thanked for the helpful hospitality. We are grateful to C. Cerrano (Università degli studi di Genova – Italy) who helped to collect *Petrosia ficiformis* specimens. S. Zea (Universidad Nacional de Colombia) is acknowledged for assisting in identification of some of the Bahamian sponges. L. Steindler would like to indicate her appreciation to S. Schuster who first introduced her into the “molecular world”.

REFERENCES


FIGURE LEGENDS

FIG. 1: Maximum Likelihood tree of sponge-associated cyanobacteria based on 16S rRNA. Bootstrap values above 50% are indicated. Sponge-associated cyanobacteria (which were named following their host species) are indicated in bold. The group numbers follow Honda et al. (8) and Robertson et al. (10). The triangle represents group 6 detailed in Figure 2. The individual identification number is given at the end of the sequence; different letters represent different clones obtained for the same sponge individual. Sequences from this work are indicated by a star after the accession number. The sampling localities of sponge-associated cyanobacteria are indicated by bars. AU-Australia, BH-Bahamas, GU-Guam, JP-Japan, MX-Mexican Pacific, NZ-New Zealand, ZZ-Zanzibar.

FIG. 2: “Group 6” detailed 16S rRNA-based Maximum Likelihood tree of sponge-associated cyanobacteria. Arrow, to outgroup (cf. Fig. 1). Bootstrap values above 50% are indicated. Sponge-associated cyanobacteria (which were named following their host species) are indicated in bold. The individual identification number is given at the end of the sequence; different letters represent different clones obtained for the same sponge individual. Sequences from this work are indicated by a star after the accession number. The sampling localities of sponge-associated cyanobacteria are indicated by bars. AU-Australia, BH-Bahamas, MS-Mediterranean Sea, RS-Red Sea, ZZ-Zanzibar.