

Detection of the Free-Living Forms of Sulfide-Oxidizing Gill Endosymbionts in the Lucinid Habitat (*Thalassia testudinum* Environment)

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Target DNA from the uncultivable *Codakia orbicularis* endosymbiont was PCR amplified from sea-grass sediment. To confirm that such amplifications originated from intact bacterial cells rather than free DNA, whole-cell hybridization (fluorescence in situ hybridization technique) with the specific probe Symco2 was performed along with experimental infection of aposymbiotic juveniles placed in contact with the same sediment. Taken together, the data demonstrate that the sulfide-oxidizing gill endosymbiont of *Codakia orbicularis* is present in the environment as a free-living uncultivable form.

All of the thioautotrophic endosymbioses occurring in marine invertebrates that have been examined so far are unambiguously confined to the gamma subdivision of the *Proteobacteria*, where they form a single coherent cluster (13, 14). These endosymbionts have to infect each new host generation successfully either by a vertical transmission from parents to offspring (6) or by an environmental transmission that involves the infection of the next host generation from an environmental stock of a free-living symbiont form (3).

The endosymbiont transmission mode has been elucidated for only few species due to the difficulty in cultivating symbionts and raising the invertebrate hosts from the egg to the adult stage in the laboratory. The environmental transmission mode was strongly suggested to occur in two oligochaetes (19) and in a few vestimentiferans based on molecular data obtained from mature gonads (8) and on ultrastructural observations of embryonic and larval stages (4, 35, 42). In bivalves, the symbiont transmission mode appears to be family specific, as suggested previously (23): vertical in Solemyidae (7, 33) and Vesicomidae (8, 9) and environmental in Lucinidae (21, 23, 25, 26).

The shallow-water tropical lucinid *Codakia orbicularis* is the only marine invertebrate with chemoautotrophic bacterial endosymbionts for which the environmental transmission mode has unequivocally been demonstrated, by using experimental infections of aposymbiotic juveniles (21, 24) with unsterilized sediment collected from sea-grass beds as an inoculum. These experiments have suggested the presence of a free-living but uncultivable form of the gill endosymbiont in the sediment.

Recently, fluorescence in situ hybridization (FISH) with spe-

cific labeled oligonucleotide probes that target intracellular rRNA were described for the direct identification of individual bacterial cells within their natural environment (1, 2). This approach offers the advantage of circumventing the requirement for cultivation to identify and distinguish bacterial cells in environmental samples either directly (in situ) or after preliminary extraction (2). FISH probes have been successfully applied to the hydrothermal environment for the identification of yet-unculturable filamentous bacteria (41), ϵ -proteobacteria invertebrate epibionts (39), and thermophilic bacteria from deep-sea hydrothermal chimneys (28) but have never been used to locate chemoautotrophic endosymbionts prior to their colonizing invertebrate hosts.

MATERIALS AND METHODS

Sediment was collected from low-sulfide-containing *Thalassia testudinum* sea-grass beds and subdivided into three parts. The first part was used for experimental infections, the second was used for DNA extraction, and the third was

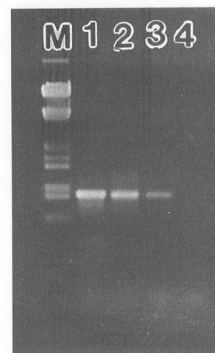


FIG. 1. PCR detection of *C. orbicularis* free-living symbiont form in sediments. Lanes: M, DNA marker; 1, gill; 2, *T. testudinum* sea-grass sediment; 3, mangrove sediment; 4, negative control. PCR products are located between 831 and 983 bp according to the sizes of standard bands.

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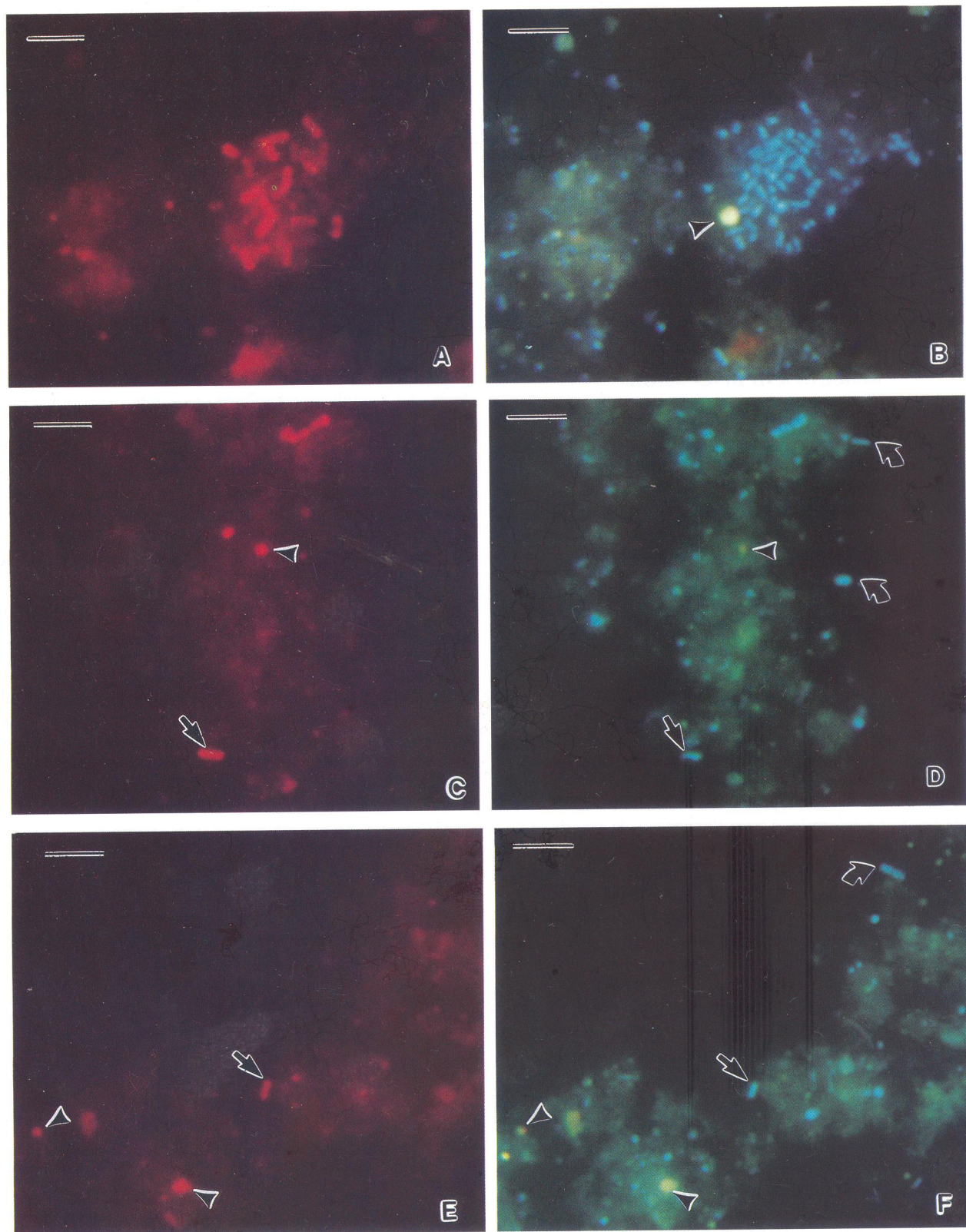


FIG. 2. Whole-cell hybridization of cells extracted from *T. testudinum* sediment. (A) Hybridization with eubacterial probe EUB338; (B) identical microscopic field for DAPI staining; (C and E) hybridization with the *C. orbicularis* symbiont-specific probe Symco2; (D and F) identical microscopic fields for DAPI staining. Arrowheads, nonbacterial sediment particles; straight arrows, free-living form of *C. orbicularis* gill endosymbiont; curved arrows, environmental bacteria. Bars, 5 μm .

fixed in 4% paraformaldehyde in 4× phosphate-buffered saline for FISH experiments.

Bacterial cells were extracted from sea-grass bed sediment as described elsewhere (28) before hybridization as described by Heddi et al. (30). Two oligonucleotides probes were used: EUB338 (5'-GCTGCTCCCGTAGGAGT-3'), targeting most members of the eubacteria (1, 11), and the *C. orbicularis* symbiont-specific probe Symco2 (5'-TACAGAGGGTCGCCAACCCGTG-3'; *Escherichia coli* positions 1247 to 1268) (21).

Total nucleic acids were extracted from low and highly reduced sediments according to the protocol for DNA extraction from soil described by Zhou et al. (43) before PCR amplifications with the specific *C. orbicularis* symbiont primer set (21, 23). To protect against unspecific hybridization that could occur when amplifying total DNA extracted from the environment, each of the symbiont ribosomal DNA (rDNA) targets amplified from sediment was purified and sequenced independently. PCR products from three independent amplifications were pooled, sequenced, and manually aligned with the previously published complete 16S rDNA sequence of the *C. orbicularis* gill endosymbiont (EMBL accession number X84979) (15). A total of 730 nucleotide positions were utilized in this analysis.

RESULTS AND DISCUSSION

PCR amplifications performed with the specific *C. orbicularis* symbiont primer set (Symco1-1492r) on sea-grass bed sediments and on mangrove swamp samples produced DNA fragments around the expected size of 872 bp (Fig. 1). No amplification products were obtained from the negative control reactions.

The oligonucleotide sequence of the specific primer Symco1 had been designed from 16S rDNA sequences deposited in database libraries. Such analyses could not be sufficient enough to guarantee a strong specificity of the primer sequences for studying the microbial diversity in the environment, as the sequences available represent only a small percentage of the environmental bacteria. Direct sequence analysis indicated that the PCR products amplified from sediment contained a single detectable sequence. A total of 730 nucleotides were sequenced for each sample analyzed, corresponding to positions 638 to 1002 and 1126 to 1492 of the *E. coli* nomenclature (5). These 16S rDNA sequences were identical at all 730 nucleotide positions determined from the previously examined *C. orbicularis* symbiont, indicating that the PCR fragments correspond to the gill endosymbiotic DNA.

However, PCR amplifications may also result from free DNA or dead symbionts rather than from free-living intact cells. Such a DNA could be released in the environment from a recently deceased clam close to the sediment collection spot or due to the feeding activity of clam predators that could release symbionts in the sediment. Therefore, FISH experiments were used to confirm the PCR results. Fluorescent probes have successfully been used in various studies of environmental bacteria from soils, lakes, or marine environments (10, 12, 20, 28, 38, 40) and also in studies of various animal symbioses (30, 31, 34).

Here, we used the symbiont-specific probe Symco2, which has good accessibility to its target site along the 16S rRNA sequence (18), i.e., 60%, compared to only 5% for Symco1 (*E. coli* positions 638 to 656).

Most of the bacteria extracted hybridized with the universal probe EUB338, which was used as positive control (Fig. 2A), when compared to the cells stained with the DNA fluorescent dye 4',6'-diamidino-2-phenylindole (DAPI) (Fig. 2B). Only a few bacteria were hybridized with the specific probe Symco 2

(Fig. 2C and E) compared to DAPI staining (Fig. 2D and F). Such cells likely represent the free-living form of the *C. orbicularis* gill endosymbiont.

The gill endosymbionts of *C. orbicularis* are generally rod shaped, large (up to 5 μm), and characterized by sulfur granules located in the periplasmic space (17). The free-living form of *C. orbicularis* gill endosymbiont extracted from *T. testudinum* sediment appears as small rods (1 to 2 μm) (Fig. 2C and E). Thus, the observation of the symbiont structure appears to reflect modifications that occur from extracellular to intracellular life styles. The most striking difference is in the bacterial size, which can increase two- to fivefold inside the bacteriocytes. This phenomenon seems to occur in most animal bacteriocyte-inducing symbioses and may result from bacterial growth deregulation under intracellular conditions (37).

Experimental infections of aposymbiotic juveniles of *C. orbicularis*, obtained as described previously (22), were also performed with crude sediment collected in the same sea-grass beds. All juveniles were infected by symbiosis-competent bacteria from the sediment, while juveniles from the negative control remained aposymbiotic, indicating that no contamination had occurred in the laboratory during the experiments. This demonstrates that the sediment used for in situ hybridization contained viable bacteria that were able to initiate symbiosis with aposymbiotic host juveniles.

The data presented in this study provide the first evidence for the free-living form of this symbiont species in the environment and confirm environmental symbiont acquisition for the other lucinids colonized by the *C. orbicularis* symbiont (15, 16, 26, 27). Thus, lucinid endosymbionts may exhibit evolutionary features different from those of other invertebrate (solemyid and vesicomid bivalves) and particularly insect symbioses, where endosymbionts are transmitted strictly vertically (7, 8, 33), which results in bacterial genome A+T bias and severe genome size reduction (29, 32, 36).

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