



Specific epibacterial communities on macroalgae: phylogeny matters more than habitat

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ABSTRACT: Epibacterial communities on thalli of the algal species *Fucus serratus*, *Fucus vesiculosus*, *Laminaria saccharina*, *Ulva compressa*, *Delesseria sanguinea* and *Phycodryis rubens* were analysed using 16S ribosomal RNA gene-based DGGE. Individuals of all species were collected in the Kiel Fjord (Baltic Sea) and in the rocky intertidal of Helgoland (North Sea). DGGE gels as well as cluster and multidimensional scaling analysis based on the DGGE band patterns of the epibacterial community showed significant differences between the epibacterial communities on the investigated algal species both in the Baltic and North Seas. Epibacterial communities differed less between regions than between host species, and were more similar on closely related host species. Results give the first evidence for lineage-specific bacterial associations to algal thalli. Furthermore, the results suggest that these algal species may control their epibiotic bacterial communities.

KEY WORDS: Biofilm · Bacteria · Antifouling · Algae · DGGE

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INTRODUCTION

The brown algae *Fucus serratus*, *F. vesiculosus* and *Laminaria saccharina*, the green alga *Ulva compressa* and the red algae *Delesseria sanguinea* and *Phycodryis rubens* are common representatives of macroalgae in the Baltic and North Seas. Most aquatic organisms, in particular primary producers like macroalgae, interact with their biotic and abiotic environment through their surface (Wahl 2008). Functionally relevant features of this interface are colour, texture, flexibility, transparency, permeability, toxicity, smell and taste. All these characteristics are affected when the alga's surface is colonized by epibionts. Epibiosis, the settlement of organisms on living surfaces, is known to dramatically modify the strength and mode of interaction between a host and its environment (Wahl & Hay 1995, Rohde et al. 2008). After chemical surface conditioning, bacterial colonization is the second of 4 steps of the colonization of a new surface, thus prokaryotes can be regarded as primary colonizers of algal thalli (Wahl 1989). Epibiotic bacteria play a key role in the colonization process of an algal thallus for several reasons:

they are fast colonizers, highly adaptive and capable of quick metabolism of algal exudates. Furthermore, under certain circumstances, epibiotic bacteria may serve as a nutrient source for the algae (Croft et al. 2006) and they may supply growth factors for algae (Matsuo et al. 2005, Tsavkelova et al. 2006). Of course, some epibiotic bacteria on algal thalli also are known as pathogens (Michel et al. 2006), e.g. *Alteromonas* sp. (Vairappan et al. 2001) and *Pseudo alteromonas* sp. (Ivanova et al. 2002).

However, epibiotic bacteria are also supposed to have a profound effect on further colonization processes (Harder et al. 2002, Tait et al. 2005, Dobretsov et al. 2006). Dobretsov & Qian (2002) investigated the effect of several bacterial strains on marine micro- and macrofouling on *Ulva reticulata* and found a *Vibrio* sp. which significantly inhibits settlement and metamorphosis of polychaete larvae. Their results suggest that certain epibacteria may contribute to the host alga's protection against further fouling. On the other hand, a review on this topic suggests that, depending on the composition, biofilms grown on hard substrate promote larval settlement (Wieczorek & Todd 1998). In either

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case, the capacity to control its epibacterial community should be of substantial ecological and selective advantage for the individual alga. Because of the different composition of the thalli of brown, red and green algae, it can be assumed that their thallus surfaces are colonized in differing manners (temporal and spatial distribution on the thallus) by bacteria (Longford et al. 2007) and probably host different epibacterial communities on their thalli. For example, several red algae are known to actively control the bacterial colonization on their thalli. Steinberg et al. (1997) showed that *Delisea pulchra* produces a furanone resembling signal molecules of the acyl homoserine lactone (AHL) regulation system of gram-negative bacteria leading towards a selective enrichment of gram-positive bacteria on thalli of this species. Weinberger et al. (1999) demonstrated another defense mechanism of the red alga *Gracilaria conferta*: this alga is able to secrete hydrogen superoxide when colonized by agarolytic bacteria.

To date there are few studies dealing with the bacterial community composition of different algal species. Culture-based studies on the bacterial epibionts of *Laminaria saccharina* revealed a shift from mesophilic bacterial species in summer to psychrophilic bacteria in winter (Mazurek & Field 1980). This may be due to seasonal variation in either the presence of different bacterial strains in the surrounding water or the ability of bacterial species to attach to the algal surface. Staufenberg et al. (2008) recently analysed the bacterial community attached to *L. saccharina* using DGGE and clone libraries based on PCR-amplified 16S ribosomal RNA (rRNA) gene fragments in the North and Baltic Seas. The authors showed that on young algal tissues there was a close correlation between bacterial communities on different conspecific algae, regardless of the sample origin and season.

Based on the results of these different studies it may be assumed that the epibiotic bacterial community may differ between algal species. Therefore, the present study focused on the analysis of epibacterial communities on different algal individuals belonging to red, green and brown macroalgae of the North and Baltic Seas using 16S rRNA gene-based DGGE.

MATERIALS AND METHODS

Sampling. Three individuals each of 6 marine algal species were collected in August 2004 in the Baltic (Kiel Fjord, 54° 27' 4 N, 10° 12' E) and North Seas (north coast of Helgoland, 54° 11' 5 N, 7° 52' 5 E). The sampling site in the Kiel Fjord was located in the littoral zone in an area protected from strong currents, whereas samples in Helgoland (North Sea) were taken in a littoral zone strongly influenced by tidal changes. Salinity in the

North Sea is around 35‰, compared to 15‰ in the Kiel Fjord. Selected algal species for analysis were *Fucus serratus* Linnaeus, *F. vesiculosus* Linnaeus, *Laminaria saccharina* (Linnaeus) Lamouroux, *Ulva compressa*, *Delleseria sanguinea* (Hudson) Lamouroux and *Phycodryas rubens* (Linnaeus) Batters. These species represent all 3 major macroalgal phyla and were present at both sampling locations during the sampling season. Sampling in the North and Baltic Seas was performed by SCUBA diving. Algal individuals showing a good physiological state were collected individually in sterile Ziploc bags and transported to the laboratory in a cooler (<10°C). Within 2 h after collection, the individual algae were washed 10 times (water exchange after each step) in sterile petri dishes with filtered (0.2 µm pore size) and autoclaved seawater to remove loosely attached bacteria. Rinsed samples were placed in empty, sterile petri dishes. Two cm² of the surface of each algal individual was vigorously swabbed with a sterile cotton-tipped applicator. Only young and clean thallus regions were sampled from *F. vesiculosus*, *F. serratus* and *L. saccharina*, whilst swabs of whole algal surfaces were taken from *U. compressa*, *D. sanguinea* and *P. rubens*. Subsequently, swab tips were transferred to sterile 2 ml vials and frozen at -20°C until further analysis.

DNA extraction. DNA was extracted using the QiaAmp DNA Mini Kit (Qiagen) following the manufacturer's instructions (buccal swab protocol). For elution, DNA-free water (Fluka Biochemica) was used instead of Buffer AE.

PCR amplification. PCR of 16S rRNA genes of bacterial community DNA was performed using PuReTaq Ready-To-Go PCR Beads (GE Healthcare) in a total PCR volume of 25 µl; 10 pmol of each bacterial primer 341F-GC (5'- [CGC CCG CCG CGC GCG GCG GGC GGG GCG GGG GCA CGG GGG GC] CTA CGG GAG GCA GCA G-3') and 534R (5'-ATT ACC GCG GCT GCT GG-3') was used for amplification of suitable fragments for DGGE (Muyzer et al. 1993). A GC-clamp (sequence in square brackets) was attached to the 5' end of the forward primer.

PCR conditions were as follows: initial denaturation at 94°C for 2 min; 15 touchdown cycles starting with an annealing temperature of 65°C for 40 s and an incremental reduction of 1°C per cycle; elongation at 72°C for 40 s; and denaturation at 95°C for 30 s. The touchdown steps were followed by 40 cycles of annealing temperature at 50°C for 40 s, elongation at 72°C for 40 s and denaturation at 94°C for 30 s; a final annealing step was performed at 42°C for 60 s and a final elongation at 72°C for 5 min. The correct size of the amplified DNA fragments was verified by electrophoresis of 10% of the PCR reaction volume in 2% agarose in 1 × TBE buffer.

DGGE was performed using double gradient polyacrylamide gels (Petri & Imhoff 2001). DGGE gels

contained a denaturing gradient from 40 to 80% (100% defined as 7 M urea and 10 M formamide) and an acrylamide (Acrylamide-Bis: 37.5:1) gradient from 6% to 8%. Electrophoresis was run at 60°C for 13.5 h at 80 V in 0.5 × TAE buffer in a CBS Scientific DGGE-2001 system. After electrophoresis, the gel was stained for 45 min in SYBR Gold® (Invitrogen), rinsed for 30 min in 1 × TAE buffer and photographed under UV light.

Statistical analysis. DGGE gels were analysed by the generation of a presence-absence matrix based on the band pattern. All visible bands in every gel lane were taken into account for further calculation using the Primer software v.6.1.9 (Primer-E). To further ensure the comparability of band patterns, only samples from one gel were compared with each other. Bray-Curtis values without transformation were calculated. Sample similarities are shown by cluster analysis and non-metric multidimensional scaling (NMDS). Band positions were assigned to species and the program then compared species composition within and among host alga species using analysis of similarities (ANOSIM). The ANOSIM global test was used for the pairwise comparison of cluster groups. R-values near 1 indicate that similarity within a group is higher than between different groups.

RESULTS

According to the 16S rRNA gene-based DGGE analysis, the epibacterial community composition differed between algal species in the Baltic Sea. This observation was confirmed by the statistical analysis

Table 1. ANOSIM pairwise comparison of epibacterial communities on different algal phyla from the Baltic and North Seas derived from DGGE band pattern analysis. Algal individuals were arranged according to their phylogenetic affiliation into *Phaeophyta*, *Chlorophyta* and *Rhodophyta*. Similarity of DGGE band pattern of the epibacterial community was analysed within and between group identities. R-values near 1 indicate a higher similarity within a group than between different groups

Groups	R	Significance level (%)	Permutations		No. obs.
			Possible	Actual	
Baltic Sea					
Phaeophyta, Chlorophyta	0.981	0.5	220	220	1
Phaeophyta, Rhodophyta	0.989	0.1	5005	999	0
Chlorophyta, Rhodophyta	0.886	1.2	84	84	1
North Sea					
Phaeophyta, Chlorophyta	0.656	0.5	220	220	1
Phaeophyta, Rhodophyta	0.655	0.1	5005	999	0
Chlorophyta, Rhodophyta	0.935	1.2	84	84	1

based on the DGGE band pattern. ANOSIM (global test, $R = 0.877$, $p = 0.001$) showed that epibacterial communities on algal thalli differed significantly at the host species level (Table 1).

With only one exception, *Ulva* individual 1 (U1), the 3 individuals of each species showed a highly similar band pattern (Fig. 1). This result was also reflected in the cluster analysis based on the DGGE band patterns (Fig. 1). In particular, the 3 *Delesseria sanguinea* individuals showed a high similarity of the epibacterial community (>80%). Inter-individual similarities were lower in *Laminaria saccharina* (>50%) and *Phycodrys rubens* (>60%). Interestingly, within the genus *Fucus*, epibacterial communities were very similar and host species did not differ more than conspecific individuals (Fig. 1). At the other phylogenetic extreme, the epibacterial communities significantly differed between the algal phyla *Phaeophyta*, *Rhodophyta* and *Chlorophyta* according to the results

Fig. 1. DGGE gel and cluster analysis based on 16S rDNA amplified epibacterial communities of 6 macroalgal species from the Baltic Sea (each sampled in triplicate). Cluster analysis of DGGE band patterns was performed using the Bray-Curtis similarity index; similarity values are given in %. FS: *Fucus serratus*; FV: *Fucus vesiculosus*; LS: *Laminaria saccharina*; U: *Ulva compressa*; D: *Delesseria sanguinea*; P: *Phycodrys rubens*

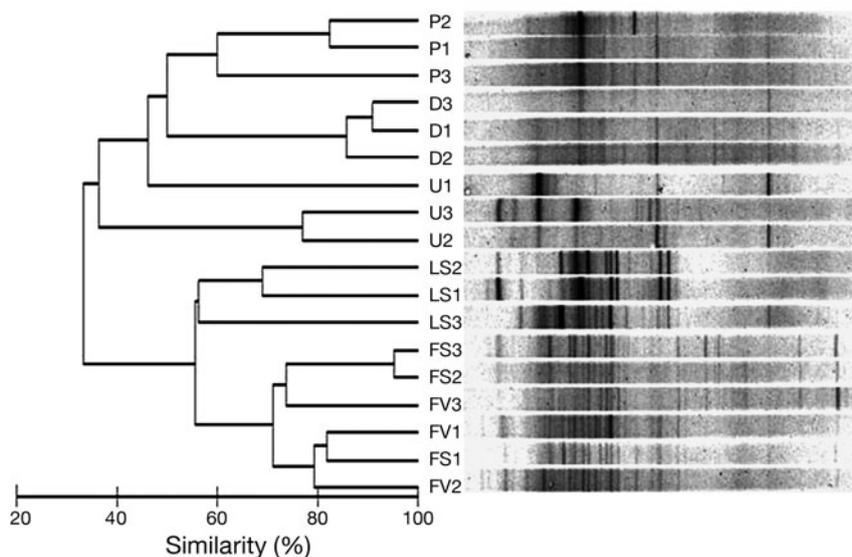


Table 2. ANOSIM pairwise comparison of epibacterial communities on different macroalgal species from the North and Baltic Seas. FV: *Fucus vesiculosus*; LS: *Laminaria saccharina*; U: *Ulva compressa*; D: *Delesseria sanguinea*; P: *Phycodrys rubens*. Each group consists of 6 individual algal samples

Groups	R	Significance level (%)	Permutations		No. obs.
			Possible	Actual	
U, D	0.919	0.2	462	462	1
U, LS	0.92	0.2	462	462	1
U, FV	1.0	0.2	462	462	1
D, LS	0.622	0.2	462	462	1
D, FV	0.936	0.2	462	462	1
LS, FV	0.796	0.2	462	462	1

DISCUSSION

In the present study, a molecular approach using 16S rRNA gene-based DGGE analysis was conducted in order to investigate how the structure of epibacterial communities differed between host species, host phyla and host regional provenance (North and Baltic Seas). In microbial ecology, 16S rRNA gene-based DGGE is widely applied as a method to compare bacterial communities in a variety of habitats including macroalgae (Celussi & Cataletto 2007). Although the information given by a DGGE band pattern is not as detailed as the information obtained, for instance, by a 16S rRNA gene-based clone library, DGGE is considered today to be a valuable method for the comparison of microbial communities in different or changing environments.

The examined marine algal species belong to the phyla *Rhodophyta* (*Delesseria sanguinea* and *Phycodrys rubens*), *Chlorophyta* (*Ulva compressa*) and *Phaeophyta* (*Fucus serratus*, *F. vesiculosus* and *Laminaria saccharina*). Although a certain inter-individual variability was observed even on conspecific algae (e.g. *Ulva* in the Baltic Sea), the general factors contributing to the dissimilarity among epibacterial communities were — with increasing importance — region, host species and host phylum. Host specificity of the associated bacterial community has already been described for red algae (Ashen & Goff 2000), Caribbean corals (Rohwer et al. 2001) and sponges of Australian temperate waters (Taylor et al. 2004). Taylor et al. (2004) analysed bacterial communities of 3 different sponges based on DGGE. They reported a certain variability at all levels, but the highest variation was observed among host species indicating, similarly to the present study, host-specific associations. Surprisingly, in the present study, host specificity was not blurred when algae of very different provenance were pooled. Even though the sampling sites were in the same longitudinal range with similar light regimes and water temperatures, both habitats differ considerably

with regard to salinity, tidal range and, most likely, bacterioplankton composition. Pinhassi et al. (2003) investigated the bacterial community composition of the North and Baltic Seas in the Skagerrak-Kattegat Front and found substantial differences on each side of the front. Staufenberger et al. (2008) investigated the bacterial community of the water surrounding *L. saccharina* individuals in the North and Baltic Seas using 16S rRNA gene-based DGGE analysis and found substantial differences in the bacterial community of the algal thalli from each location. Furthermore, Staufenberger et al. (2008) could detect a specific bacterial community on young thallus parts regardless of the geographic origin or season of sampling. The data from the present study extend the phenomenon of host-specific epibacterial communities to other algal species.

While the composition of epibacterial communities varies within conspecific algae and between regions, this variability is generally lower than the differences between algal species or host phyla. The finding suggests that species-specific properties of the algal surface and/or specific interactions between algae and bacteria may be the driving force for this selectivity.

Although our survey was limited to 6 algal species and only 2 sampling sites in the same temperate region, the results from the present study provide strong support for the hypothesis that the physico-chemical properties of macroalgal thalli may determine the differential settlement and growth of bacteria on their surfaces. There are 3 possible processes which, singly or in combination, may produce algal-specific biofilms: (1) algal propagules may already carry the specific biofilm; (2) specific algal defences may selectively inhibit the growth of biofilms other than that specific to the alga, as was demonstrated for *Delisea pulchra* (Steinberg et al. 1997), or repel already attached agarolytic bacteria (Weinberger et al. 1999); and (3) algal attractants may favour the settlement of certain strains (Pasmore & Costerton 2003). Besides repellent-attractant activity, algae can also affect bacterial growth (stimulate or inhibit). Additionally, physical properties of algae may allow adhesion of some bacterial strains and suppress attachment of others.

The epibacterial communities may in turn have an impact on further interactions between the host alga and its environment (e.g. larval settlement as reviewed by Dobretsov et al. 2006). Micro- and macrofouling is likely to be influenced by the identity and metabolism of epibiotic bacteria, however, any transcutaneous exchanges (light, nutrients, exudates, signalling molecules and other chemicals) may also be affected by the specific surface-attached bacteria (Wahl 2008). Because epibiotic bacteria are known to metabolize algal exudates and produce their own metabolites,

chemical signals of the host algae are very likely modified by their biofilms. Interactions with chemically cueing parasites, pathogens or consumers may be modulated by epibacterial biofilms (Harder et al. 2002, Tait et al. 2005, Dworjanyn & Pirozzi 2008). Thus, an alga's capacity to control composition, and density, of its biofilm is probably everything but trivial. A detailed metagenomic investigation of bacterial communities, as well as a seasonal analysis of the stability of algal associated microorganisms and further investigation of epibacterial communities of macroalgae from different marine habitats (e.g. tropical habitats), should facilitate and improve our understanding of the interaction between macroalgae and bacteria.

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