

Seasonal Dynamics of Epiphytic Microbial Communities on Marine Macrophyte Surfaces

Marino Korlević^{1*}, Marsej Markovski¹, Zihao Zhao², Gerhard J. Herndl^{2,3}, Mirjana Najdek¹

1. Center for Marine Research, Ruđer Bošković Institute, Croatia

2. Department of Functional and Evolutionary Ecology, University of Vienna, Austria

3. NIOZ, Department of Marine Microbiology and Biogeochemistry, Royal Netherlands Institute for Sea Research, Utrecht University, The Netherlands

*To whom correspondence should be addressed:

Marino Korlević

G. Paliaga 5, 52210 Rovinj, Croatia

Tel.: +385 52 804 768

Fax: +385 52 804 780

e-mail: marino.korlevic@irb.hr

Running title: Seasonal dynamics of epiphytic communities

1 Summary

2 Surfaces of marine macrophytes (seagrasses and macroalgae) are inhabited by diverse microbial
3 communities. Most studies focusing on epiphytic communities of macrophytes did not take into
4 account temporal changes or applied low sampling frequency approaches. Illumina sequencing
5 of the V4 16S rRNA region was performed to determine the seasonal dynamics of epiphytic
6 communities of the seagrass *Cymodocea nodosa* and invasive macroalga *Caulerpa cylindracea*.
7 Leaves and thalli were sampled in a meadow of *Cymodocea nodosa* invaded by the invasive *Caulerpa*
8 *cylindracea* and in a monospecific settlement of *Caulerpa cylindracea* located in the proximity of
9 the meadow at monthly intervals. For comparison the ambient prokaryotic plankton community was
10 also characterized. At the OTU level, the microbial community composition differed between the
11 ambient water and the epiphytic communities exhibiting host-specificity. Also, successional changes
12 were observed connected to the macrophyte growth cycle. Taxonomic analysis, however, showed
13 similar high rank groups in the ambient water and the epiphytic communities, with the exception
14 of *Desulfobacterota*, which were only found on *Caulerpa cylindracea*. *Cyanobacteria* showed
15 seasonal changes while other high rank taxa were present throughout the year. Phylogenetic groups
16 present throughout the year constituted most of the sequences, while less abundant taxa showed
17 seasonal patterns connected to the macrophyte growth cycle. Taken together, epiphytic microbial
18 communities of the seagrass *Cymodocea nodosa* and the macroalgae *Caulerpa cylindracea* appear
19 to be host-specific and contain taxa that undergo successional changes.

20 **Introduction**

21 Marine macrophytes (seagrasses and macroalgae) are important ecosystem engineers forming
22 close associations with microorganism belonging to all three domains of life (Egan *et al.*, 2013;
23 Tarquinio *et al.*, 2019). Microbes can live within macrophyte tissue as endophytes or can form
24 epiphytic communities on surfaces of leaves and thalli (Egan *et al.*, 2013; Hollants *et al.*, 2013;
25 Tarquinio *et al.*, 2019). Epiphytic and endophytic microbial communities exhibit a close functional
26 relationship with the macrophyte host. It was proposed that this close relationship constitutes a
27 holobiont, an integrated community where the macrophyte organism and its symbiotic partners
28 support each other (Margulis, 1991; Egan *et al.*, 2013; Tarquinio *et al.*, 2019).

29 Biofilms of microbial epiphytes can contain diverse taxonomic groups and harbor cell
30 abundances from 10^2 to 10^7 cells cm^{-2} (Armstrong *et al.*, 2000; Bengtsson *et al.*, 2010; Burke
31 and Thomas *et al.*, 2011). In such an environment a number of positive and negative interactions
32 between the macrophyte and the colonizing microorganisms have been described (Egan *et*
33 *al.*, 2013; Hollants *et al.*, 2013; Tarquinio *et al.*, 2019). Macrophytes can promote growth of
34 associated microbes by nutrient exudation, while in return microorganisms may support macrophyte
35 performance through improved nutrient availability, phytohormone production and protection from
36 toxic compounds, oxidative stress, biofouling organisms and pathogens (Egan *et al.*, 2013; Hollants
37 *et al.*, 2013; Tarquinio *et al.*, 2019). Besides these positive interactions, macrophytes can negatively
38 impact the associated microbes such as pathogenic bacteria by producing reactive oxygen species
39 and secondary metabolites (Egan *et al.*, 2013; Hollants *et al.*, 2013; Tarquinio *et al.*, 2019).

40 All these ecological roles are carried out by a taxonomically diverse community of
41 microorganisms. At higher taxonomic ranks a core set of macrophyte epiphytic taxa was described
42 consisting mainly of *Alphaproteobacteria*, *Gammaproteobacteria*, *Desulfobacterota*, *Bacteroidota*,
43 *Cyanobacteria*, *Actinobacteriota*, *Firmicutes*, *Planctomycetota*, *Chloroflexi* and *Verrucomicrobiota*
44 (Crump and Koch, 2008; Tujula *et al.*, 2010; Lachnit *et al.*, 2011). In contrast, at lower taxonomic

45 ranks host specific microbial communities were found (Lachnit *et al.*, 2011; Roth-Schulze *et al.*,
46 2016). Recently, it was shown that even different morphological niches within the same alga had a
47 higher influence on the composition of the bacterial community than biogeography or environmental
48 factors (Morrissey *et al.*, 2019). While the microbial community composition varies between host
49 species, metagenomic analyses revealed that the majority of the microbial functions are conserved
50 (Burke and Peter Steinberg *et al.*, 2011; Roth-Schulze *et al.*, 2016). This discrepancy between the
51 microbial taxonomic and functional composition might be explained by the lottery hypothesis. It
52 postulates that an initial random colonization step takes places from a set of functionally equivalent
53 taxonomic groups resulting in taxonomically different epiphytic communities sharing a core set of
54 functional genes (Burke and Peter Steinberg *et al.*, 2011; Roth-Schulze *et al.*, 2016). In addition,
55 some of the variation in the reported data could be attributed to different techniques used in these
56 studies, such as different protocols for epiphytic cell detachment and/or DNA isolation, as no
57 standard protocol has been yet established to study epiphytic communities (Ugarelli *et al.*, 2019;
58 Korlević *et al.*, submitted).

59 The majority of studies describing macrophyte epiphytic microbial communities did not include
60 possible seasonal changes (Crump and Koch, 2008; Lachnit *et al.*, 2009; Burke and Thomas *et*
61 *al.*, 2011; Roth-Schulze *et al.*, 2016; Ugarelli *et al.*, 2019). If seasonal changes were taken into
62 account, low temporal frequency, applied methodologies and/or limited number of analyzed host
63 species did not allow a high taxonomic resolution (Tujula *et al.*, 2010; Lachnit *et al.*, 2011; Miranda
64 *et al.*, 2013; Michelou *et al.*, 2013; Mancuso *et al.*, 2016). In the present study we describe the
65 seasonal dynamics of bacterial and archaeal communities on the surfaces of the seagrass *Cymodocea*
66 *nodosa* and siphonous macroalgae *Caulerpa cylindracea* determined on a mostly monthly scale.
67 Bacterial and archaeal epiphytes were sampled in a meadow of *C. nodosa* invaded by the invasive
68 *C. cylindracea* and in a locality of only *C. cylindracea* located in the proximity of the seagrass
69 meadow. For comparison, the microbial community of the ambient seawater was also characterized.

70 **Results**

71 Sequencing of the 16S rRNA V4 region yielded a total of 1.8 million sequences after quality
72 curation and exclusion of sequences without known relatives (no relative sequences) and eukaryotic,
73 chloroplast and mitochondrial sequences (Table S1). A total of 35 samples originating from
74 epiphytic archaeal and bacterial communities associated with surfaces of the seagrass *C. nodosa*
75 and the macroalga *C. cylindracea* were analyzed. In addition, 18 samples (one of the samples was
76 sequenced twice) originating from the ambient seawater were also processed for comparison. The
77 number of reads per sample ranged between 8,408 and 77,463 sequences (Table S1). Even when
78 the highest sequencing effort was applied the rarefaction curves did not level off as commonly
79 observed in high-throughput 16S rRNA amplicon sequencing approaches (Fig. S1). Following
80 quality curation and exclusion of sequences as mentioned above reads were clustered into 28,750
81 different OTUs at a similarity level of 97 %. Read numbers were normalized to the minimum
82 number of sequences (8,408, Table S1) through rarefaction resulting in 17,157 different OTUs with
83 371 to 2,071 OTUs per sample (Fig. S2). To determine seasonal changes in richness and diversity
84 the observed number of OTUs, Chao1, ACE, Exponential Shannon (Jost, 2006) and Inverse Simpson
85 were calculated after normalization through rarefaction. Generally, richness estimators and diversity
86 indices showed similar trends. On average, higher values were found for *C. cylindracea* (mixed
87 [Number of OTUs, $1,685 \pm 127.6$ OTUs] and monospecific [Number of OTUs, $1,735.2 \pm 166.6$
88 OTUs]) than for *C. nodosa* (Number of OTUs, $1,055.2 \pm 212.0$ OTUs) and lowest values were
89 obtained for the microbial community of the ambient seawater (Number of OTUs, 533.4 ± 142.1
90 OTUs) (Fig. S2). Seasonal changes did not reveal such large dissimilarities. *C. nodosa* communities
91 showed a slow increase towards the end of the study, while *C. cylindracea* (mixed and monospecific)
92 communities were characterized by slightly higher values in spring and summer than in autumn and
93 winter (Fig. S2).

94 To determine the proportion of shared archaeal and bacterial OTUs and communities sampled
95 in different environments the Jaccard's Similarity Coefficient on presence-absence data and

96 Bray-Curtis Similarity Coefficient, respectively, were calculated. Coefficients were determined after
97 normalization through rarefaction and binning of samples from the particular environment. The
98 highest proportion of shared OTUs and community was found between mixed and monospecific *C.*
99 *cylindracea* (Jaccard, 0.35; Bray-Curtis, 0.77), while lower shared values were calculated between
100 ambient seawater and epiphytic communities (Fig. 1). Shared proportion between *C. nodosa* and *C.*
101 *cylindracea* were approximately in-between the values of mixed and monospecific *C. cylindracea*.
102 To assess seasonal changes in the proportion of shared OTUs and communities the Jaccard's and
103 Bray-Curtis Similarity Coefficients were calculated between consecutive sampling points (Fig. 2).
104 Both coefficients showed similar trends. Temporal proportional changes were more pronounced for
105 ambient seawater than for *C. nodosa* and especially *C. cylindracea* associated communities (Fig. 2).
106 In addition, only 0.4 – 1.0 % of OTUs from each surface associated community were present at
107 all seasons. These persistent OTUs constituted a high proportion of total sequences (40.7 – 51.7
108 %). To further disentangle the environmental and seasonal community dissimilarity a Principal
109 Coordinates Analysis (PCoA) based on Bray-Curtis distances and OTU abundances was applied. A
110 clear separation between ambient seawater and surface associated communities was found (Fig. 3).
111 In addition, a separation of epiphytic bacterial and archaeal communities based on host species
112 was detected. This separation was further supported by ANOSIM ($R = 0.96, p < 0.001$). Seasonal
113 changes of *C. nodosa* associated communities indicated a separation between spring, summer and
114 autumn/winter samples (ANOSIM, $R = 0.54, p < 0.01$). For *C. cylindracea* associated communities
115 a separation between summer and autumn/winter/spring samples was observed that was, however,
116 not as strong as for *C. nodosa* associated communities (ANOSIM, $R = 0.29, p < 0.01$) (Fig. 3).

117 The taxonomic composition of both, macrophyte associated and ambient seawater communities
118 was dominated by bacterial ($99.1 \pm 2.1 \%$) over archaeal sequences ($0.9 \pm 2.1 \%$) (Fig. 4). Higher
119 relative abundances of chloroplast related sequences were only observed in surface associated
120 communities, with higher values in autumn/winter ($37.2 \pm 11.2 \%$) than in spring/summer
121 ($20.9 \pm 9.7 \%$) (Fig. S3). Generally, at higher taxonomic ranks (phylum-class), epiphytic and
122 ambient seawater microbial communities were composed of similar bacterial taxa. Ambient

123 seawater communities were mainly comprised of *Actinobacteriota*, *Bacteroidota*, *Cyanobacteria*,
124 *Alphaproteobacteria*, *Gammaproteobacteria* and *Verrucomicrobiota*. Communities associated with
125 *C. nodosa* consisted additionally of *Planctomycetota* contributing more in summer 2018 than in
126 other seasons. In addition, communities from mixed and monospecific *C. cylindracea* were similar
127 and characterized by the same groups as ambient seawater and *C. nodosa* communities with the
128 addition of *Desulfobacterota* (Fig. 4). Larger differences between environments and host species
129 were observed at lower taxonomic ranks (Figs. 5 – 9).

130 *Cyanobacteria* related sequences comprised, on average, $5.5 \pm 4.4\%$ of total sequences (Fig. 5).
131 Higher proportions were found for *C. nodosa* ($16.4 \pm 5.3\%$) and *C. cylindracea* mixed (7.7 ± 3.9
132 %) and monospecific ($7.8 \pm 2.4\%$) associated communities in autumn and for ambient seawater
133 communities in winter ($8.8 \pm 7.5\%$). Large taxonomic differences between surface associated
134 and ambient seawater cyanobacterial communities were observed. Ambient seawater communities
135 were mainly comprised of *Cyanobium* and *Synechococcus*, while surface associated communities
136 were comprised of *Pleurocapsa* and sequences within the class *Cyanobacteriia* that could not be
137 further classified (no relative *Cyanobacteriia*) (Fig. 5). In addition, seasonal changes in surface
138 associated communities were observed in *Pleurocapsa* and no relative *Cyanobacteriia* comprising
139 larger proportions in autumn and winter and *Acrophormium*, *Phormidesmis* and sequences without
140 known relatives within the *Nodosilineaceae* (no relative *Nodosilineaceae*) in spring and summer
141 (Fig. 5).

142 Sequences classified as *Bacteroidota* comprised, on average, $19.2 \pm 5.5\%$ of all sequences
143 (Fig. 6). Similarly to *Cyanobacteria*, large differences in the taxonomic composition between
144 ambient seawater and surface associated communities were found (Fig. 6). The ambient seawater
145 community was characterized by the NS4 and NS5 marine groups, uncultured *Cryomorphaceae*,
146 uncultured *Flavobacteriaceae*, NS11-12 marine group, *Balneola*, uncultured *Balneolaceae* and
147 *Formosa*. In contrast, in surface associated communities *Lewinella*, *Portibacter*, *Rubidimonas*,
148 sequences without known relatives within the *Saprospiraceae* (no relative *Saprospiraceae*),

149 uncultured *Saprospiraceae*, (sequences without known relatives within the *Flavobacteriaceae*
150 (no relative *Flavobacteriaceae*)and uncultured *Rhodothermaceae* were found. Some groups
151 showed minor seasonal changes such as no relative *Flavobacteriaceae* whose sequences were
152 more abundant from November 2017 until June 2018. In contrast, uncultured *Rhodothermaceae*
153 showed higher proportions from June 2018 until the end of the study period. Surface associated
154 *Bacteroidota* communities were very diverse as observed in the high proportion of taxa clustering
155 as other *Bacteroidota* (Fig. 6).

156 On average, *Alphaproteobacteria* were in comparison to the other high rank taxa the largest
157 taxonomic group, comprising 29.2 ± 12.0 % of all sequences (Fig. 7). In accordance to the above
158 described taxa, large differences between ambient seawater and surface associated communities
159 were observed. Ambient seawater communities were composed mainly of the SAR11 clade,
160 AEGEAN-169 marine group, SAR116 clade, sequences without known relatives within the
161 *Rhodobacteraceae* (no relative *Rhodobacteraceae*), HIMB11 and the OCS116 clade, while
162 surface associated communities were composed mainly of no relative *Rhodobacteraceae* and to
163 a lesser degree of *Pseudoahrensia*, *Amylibacter* and sequences without known relatives within
164 the *Alphaproteobacteria* (no relative *Alphaproteobacteria*) and *Hyphomonadaceae* (no relative
165 *Hyphomonadaceae*). Representatives of no relative *Rhodobacteraceae* comprised on average 40.6
166 ± 23.2 % of all alphaproteobacterial sequences in the epiphytic community (Fig. 7). In addition,
167 *Amylibacter* was detected mainly in *C. nodosa* from November 2017 until March 2018.

168 Sequences related to *Gammaproteobacteria* comprised on average 18.6 ± 3.9 % of all
169 sequences (Fig. 8). Similar to above mentioned taxa, large taxonomic differences between ambient
170 seawater and surface associated communities were found. Ambient seawater communities were
171 mainly comprised of the OM60 (NOR5) clade, *Litoricola*, *Acinetobacter* and the SAR86 clade,
172 while epiphytic communities were mainly composed of sequences without known relatives within
173 the *Gammaproteobacteria* (no relative *Gammaproteobacteria*) and *Granulosicoccus*. Beside
174 these two groups specific to all three epiphytic communities, *C. nodosa* was characterized by

175 *Arenicella*, *Methylotenera* and sequences without known relatives within the *Burkholderiales* (no
176 relative *Burkholderiales*), while *Thioploca*, *Reinekea* and sequences without known relatives within
177 *Cellvibrionaceae* (no relative *Cellvibrionaceae*) were more specific to both mixed and monospecific
178 *C. cylindracea*. In addition, *Arenicella* was more pronounced in November and December 2017,
179 while no relative *Burkholderiales* and *Methylotenera* were characteristic for the period from March
180 until May 2018. For the *C. cylindracea* specific taxa no relative *Cellvibrionaceae* and *Reinekea*
181 showed seasonality and were characteristic for samples originating from June to October 2018.
182 In addition, similar to *Bacteroidota*, a large proportion of the surface associated community was
183 grouped as other *Gammaproteobacteria* indicating high diversity within this group (Fig. 8).

184 *Desulfobacterota* were specific for *C. cylindracea*. In the mixed and monospecific *C.*
185 *cylindracea* communities the proportion of *Desulfobacterota* was $25.7 \pm 11.2\%$ and $24.0 \pm 4.3\%$,
186 respectively (Fig. 9). In contrast, in ambient seawater and *C. nodosa* communities the contribution
187 of *Desulfobacterota* was only $0.1 \pm 0.08\%$ and $1.0 \pm 0.7\%$, respectively. In *C. cylindracea* the
188 community consisted mainly of *Desulfatitalea*, *Desulfobulbus*, *Desulfopila*, *Desulforhopalus*,
189 *Desulfotalea*, SEEP-SRB4, uncultured *Desulfocapsaceae* and sequences without known relatives
190 within the *Desulfobacteraceae* (no relative *Desulfobacteraceae*), *Desulfobulbaceae* (no relative
191 *Desulfobulbaceae*) and *Desulfocapsaceae* (no relative *Desulfocapsaceae*) (Fig. 9).

192 **Discussion**

193 Surfaces of marine macrophytes harbor biofilms consisting of diverse microbial taxa (Egan
194 *et al.*, 2013; Tarquinio *et al.*, 2019). No standard protocol has been developed to study these
195 macrophyte-associated microbes (Ugarelli *et al.*, 2019). Different procedures for removal of
196 microbial cells from host surfaces are described, such as host tissue shaking (Nõges *et al.*, 2010),
197 scraping (Uku *et al.*, 2007), swabbing (Mancuso *et al.*, 2016) and ultrasonication (Cai *et al.*,
198 2014). All these methods result in different removal efficiencies but none was enabling a complete
199 removal of attached microbial cells based on our experience. In the present study, we applied a
200 removal protocol (Korlević *et al.*, submitted) based on direct cellular lysis (Burke *et al.*, 2009).
201 The application of a direct lysis procedure coupled with a high sampling frequency and Illumina
202 amplicon sequencing has enabled us to described in detail the bacterial and archaeal communities
203 associated with the surfaces of two marine macrophytes, *C. nodosa* and *C. cylindracea*.

204 In the present study, highest richness was observed for *C. cylindracea* (mixed and monospecific)
205 followed by *C. nodosa* and lowest richness was found in ambient seawater microbial communities.
206 Higher richness of microbial communities associated with seagrasses than in ambient seawater
207 was described earlier and could be attributed to a larger set of inhabitable microniches existing
208 on macrophyte surfaces than in the ambient seawater (Ugarelli *et al.*, 2019). The highest richness
209 observed for *C. cylindracea* might be partly due to its contact with the sediment. The stolon of *C.*
210 *cylindracea* is attached to the sediment surface with rhizoids and thus, the stolon and rhizoids are in
211 a direct contact with the sediment. In addition, seasonal differences in richness observed for surface
212 attached communities indicated a slightly higher richness in spring and summer. This pattern could
213 be explained by a higher macrophyte growth in these two seasons than in autumn and winter (M.
214 Najdek, personal communication; Zavodnik *et al.*, 1998; Ruitton *et al.*, 2005). During their main
215 growth season in spring and summer macrophytes exhibit a more dynamic chemical interaction
216 with the surface community probably causing an increase in the number of inhabitable microniches
217 (Borges and Champenois, 2015; Rickert *et al.*, 2016).

218 We observed a strong differentiation between the surface attached and ambient seawater
219 communities at the level of OTUs, in agreement with most published studies (Burke and Thomas *et*
220 *al.*, 2011; Michelou *et al.*, 2013; Roth-Schulze *et al.*, 2016; Mancuso *et al.*, 2016; Crump *et al.*,
221 2018; Ugarelli *et al.*, 2019). This indicates that marine macrophytes are a selecting factor from
222 the pool of microbial taxa present in the ambient seawater, modifying the microbial community
223 once the macrophyte associated microbial biofilm develops (Salaün *et al.*, 2012; Michelou *et*
224 *al.*, 2013). In contrast, Fahimipour *et al.* (2017) report in a global study of *Zostera marina*,
225 similarities between the microbial community developed on leaves and in the ambient seawater.
226 The discrepancy between our data and the study of Fahimipour *et al.* (2017) could be explained
227 by different seagrass species, methodological variations or biogeographic trends as Fahimipour
228 *et al.* (2017) analyzed samples from different locations throughout the northern hemisphere. It is
229 possible that the microbial communities in ambient seawater and on leaves from the same location
230 are differing but are still more similar to each other when compared to other sampling locations.
231 Indeed, it was found that prokaryotic communities vary substantially between different sampling
232 sites (Bengtsson *et al.*, 2017). When the taxonomic composition at high ranks was analyzed no
233 such strong differentiation was noticed. Phyla and classes such as *Actinobacteriota*, *Bacteroidota*,
234 *Cyanobacteria*, *Alphaproteobacteria*, *Gammaproteobacteria* and *Verrucomicrobiota* were found
235 in both ambient seawater as well as macrophyte associated, in agreement with previous studies
236 (Burke and Thomas *et al.*, 2011; Egan *et al.*, 2013; Michelou *et al.*, 2013). In contrast, when low
237 taxonomic ranks were analyzed (i.e., family and genus) a strong differentiation was observed (Figs.
238 5 – 9). A similar differentiation at lower microbial taxonomic ranks between ambient seawater and
239 macrophytes was described for other macrophyte species as well (Egan *et al.*, 2013; Michelou *et al.*,
240 2013; Ugarelli *et al.*, 2019).

241 Beside differences between ambient seawater and surface associated microbial communities,
242 it is unclear whether the prokaryotic epiphytic community is host-specific or whether there are
243 generalist taxa characteristic to all or many macrophytes (Egan *et al.*, 2013). Similar to previously
244 described differences between microbial communities in the ambient seawater and on macrophytes,

245 at high taxonomic ranks no major difference between the microbial communities associated with
246 different hosts was observed. The only high rank phylum that was differing between *C. nodosa*
247 and *C. cylindracea* was *Desulfobacterota*, with more abundant sequences in the *C. cylindracea*
248 associated community. As already mentioned, the rhizoids and part of the stolon are in contact
249 with the sediment. Thus *Desulfobacterota* are probably a part of the epiphytic community that
250 was in contact with the sediment. Similar high rank taxa found in this study were described to be
251 specific for other species of macrophytes (Burke and Thomas *et al.*, 2011; Lachnit *et al.*, 2011;
252 Mancuso *et al.*, 2016; Bengtsson *et al.*, 2017). In contrast to high taxonomic ranks, a substantial
253 differentiation between host specific communities was found supporting the notion that macrophyte
254 associated microbial communities might be host-specific. Host-specificity was also observed for
255 other species of macroalgae and seagrasses (Lachnit *et al.*, 2011; Roth-Schulze *et al.*, 2016; Ugarelli
256 *et al.*, 2019; Morrissey *et al.*, 2019). Taken together, at high taxonomic ranks a core set of taxa
257 could be described that is characteristic for all or many macrophytes, while at low taxonomic ranks
258 a community specific to host species was identified (Figs. 3 and 4) (Egan *et al.*, 2013).

259 Seasonal changes in richness in the epiphytic community were substantial as indicated by the
260 proportion of OTUs ($\leq 1.0\%$) present at every sampling date. These persistent OTUs, however,
261 were accounting for a high proportion of sequences ($\leq 51.7\%$) (Fig. 2). A very similar proportion
262 of persistent OTUs was reported in high-frequency sampling studies describing seasonal changes in
263 picoplankton (Gilbert *et al.*, 2009, 2012). In comparison to the seawater community, a lower degree
264 of seasonal shifts was observed for the macrophyte surface associated communities. It appears that
265 microniches at the surfaces of macrophytes are providing more stable conditions than the ambient
266 seawater. At the level of OTUs seasonal changes of *C. nodosa* and *C. cylindracea* associated
267 communities were identified that could be linked to the growth cycle of the seagrass and macroalgae
268 (M. Najdek, personal communication). *C. nodosa* was characterized by a spring community
269 during maximum seagrass proliferation, a summer community during the highest standing stock of
270 *C. nodosa* and an autumn/winter community during the decay of seagrass biomass. In contrast, *C.*
271 *cylindracea* started to proliferate in late spring and was characterized only by a summer community

272 during high growth rates and by an autumn/winter/spring community when the biomass was at
273 the peak and decaying thereafter. Similar seasonal changes in the epiphytic community were also
274 described for other macroalgae (Tujula *et al.*, 2010; Lachnit *et al.*, 2011). Higher seasonal stability
275 of *C. cylindracea* surface communities than in *C. nodosa* was also observed in the higher proportion
276 of shared communities between two consecutive sampling dates in *C. cylindracea*.

277 Chloroplast sequence abundances were higher in autumn/winter than in spring/summer. This
278 pattern is not surprising as seagrasses harbor more algal epiphytes during autumn/winter than in
279 spring/summer (Reyes and Sansón, 2001). Furthermore, we used an adapted DNA isolation protocol
280 that is known to partially co-extract DNA from planktonic eukaryotes (Korlević *et al.*, 2015).
281 Strong seasonal fluctuations of high rank epiphytic taxa were not observed, with the exception of
282 *Cyanobacteria*. Cyanobacterial sequences were more pronounced in November and December than
283 in spring and summer. In the months of high cyanobacterial sequence abundances the majority of
284 sequences from this group were classified as *Pleurocapsa*, a group known to colonized different
285 living and non-living surfaces (Burns *et al.*, 2004; Longford *et al.*, 2007; Mobberley *et al.*, 2012;
286 Reisser *et al.*, 2014). It is possible that during periods of low metabolic activity there is a reduced
287 interaction and selection of the epiphytic community by the seagrass, causing leaves to become
288 a suitable surface for nonspecific colonizers (Zavodnik *et al.*, 1998). *Pleurocapsa* was replaced
289 in spring and summer by *Acrophormium*, *Phormidesmis* and sequences without known relatives
290 within the *Nodosilineaceae*. A study of coastal microbial mats found also a higher proportion
291 of *Nodosilineaceae* sequences in summer, while *Phormidesmis* sequences were at their peak in
292 autumn (Cardoso *et al.*, 2019). Other high rank taxa did not exhibit strong successional patterns. In
293 every analyzed group, with the exception of *Desulfobacterota*, taxa present throughout the year in
294 similar proportions and season specific taxa could be identified (Figs. 6 and 9). Within *Bacteroidota*
295 different groups within the family *Saprospiraceae* (e.g. *Lewinella*, *Portibacter* and *Rubidimonas*)
296 were detected across all seasons. Members of this family are often found in association with
297 macrophytes and it is suggested that they are involved in the hydrolysis and utilization of complex
298 carbon sources (Burke and Thomas *et al.*, 2011; McIlroy and Nielsen, 2014; Crump *et al.*, 2018).

299 In contrast, the families *Flavobacteriaceae* and *Rhodothermaceae* showed seasonal patterns, with
300 *Flavobacteriaceae* being more pronounced from November to June and *Rhodothermaceae* from
301 June to October (Fig. 6). Within *Alphaproteobacteria* the family *Rhodobacteraceae* comprised the
302 majority of sequences throughout the year (Fig. 7). This metabolically versatile family is often
303 associated with macrophyte surfaces and usually is one of the most abundant groups (Burke and
304 Thomas *et al.*, 2011; Michelou *et al.*, 2013; Pujalte *et al.*, 2014; Mancuso *et al.*, 2016). In addition,
305 *Hyphomonadaceae* were found in all samples. Interestingly, some of the species within this group
306 contain stalks on their cells, which can be used to attach to the macrophyte surface (Weidner *et al.*,
307 2000; Abraham and Rohde, 2014). Within the *Gammaproteobacteria*, sequences without known
308 representatives were the most pronounced group present throughout the year (Fig. 8). In addition,
309 *Granulosicoccus* was also found in almost all samples. *Gammaproteobacteria* are often a major
310 constituent of macrophyte epiphytic communities (Burke and Thomas *et al.*, 2011; Michelou *et al.*,
311 2013; Crump *et al.*, 2018). Beside these two groups, other less abundant, taxa showed seasonal and
312 host-specific patterns. For example, *C. cylindracea* harbored *Thioploca*, a known sulfur sediment
313 bacteria and *Cellvibrionaceae*, a family with cultured members known as polysaccharide degraders
314 (Jørgensen and Gallardo, 1999; Xie *et al.*, 2017). *Desulfobacterota* were found only associated with
315 *C. cylindracea* and no group within this phylum showed seasonal patterns (Fig. 9). The presence of
316 this phylum only on *C. cylindracea* is to be expected as part of the epiphytic community is in direct
317 contact with the sediment. The *Desulfobacterota* community was dominated by *Desulfatitalea* and
318 no relative *Desulfocapsaceae*, known sulfate sediment groups (Kuever, 2014; Higashioka *et al.*,
319 2015).

320 In temperate zones, marine macrophytes are exhibiting growth cycles, so it is not surprising that
321 the associated epiphytic microbial community is undergoing partial seasonal changes. In the present
322 study, we could identify in every analyzed high rank taxa phylogenetic groups present throughout
323 the year, comprising most of the sequences and a lower proportion of taxa showing seasonal
324 patterns connected to the macrophyte growth cycle (Figs. 4 and 9). Studies focusing on functional
325 comparisons between communities associated with different hosts showed that the majority of

326 functions could be found in every community, indicating functional redundancy (Roth-Schulze *et*
327 *al.*, 2016). This difference between phylogenetic variability and functional stability was explained by
328 the lottery hypothesis assuming an initial random colonization step performed by a set of functionally
329 equivalent taxonomic groups (Burke and Peter Steinberg *et al.*, 2011; Roth-Schulze *et al.*, 2016).
330 It is possible that functional redundancy is a characteristic of high abundance taxa detected to be
331 present throughout the year, while seasonal and/or host-specific functions are an attribute of taxa
332 displaying successional patterns. Further studies connecting taxonomy with functional properties
333 will be required to elucidate the degree of functional redundancy or specificity in epiphytic microbial
334 communities.

335 **Experimental procedures**

336 **Sampling**

337 Sampling was performed in the Bay of Funtana, northern Adriatic Sea (45°10'39" N,
338 13°35'42" E). Leaves and thalli were sampled in a meadow of *Cymodocea nodosa* invaded by the
339 invasive *Caulerpa cylindracea* (mixed settlement) and in a monospecific settlement of *Caulerpa*
340 *cylindracea* located in the proximity of the meadow at approximately monthly intervals from
341 December 2017 to October 2018 (Table S1). Leaves and thalli were collected by diving and
342 transported to the laboratory in containers placed on ice and filled with seawater collected at
343 the sampling site. Upon arrival to the laboratory, *C. nodosa* leaves were cut into sections of 1 –
344 2 cm, while *C. cylindracea* thalli were cut into 5 – 8 cm long sections. Leaves and thalli were
345 washed three times with sterile artificial seawater (ASW) to remove loosely attached microbial cells.
346 Ambient seawater was collected in 10 l containers by diving and transported to the laboratory where
347 the whole container volume was filtered through a 20 µm net. The filtrate was further sequentially
348 filtered through 3 µm and 0.2 µm polycarbonate membrane filters (Whatman, United Kingdom)
349 using a peristaltic pump. Filters were briefly dried at room temperature and stored at –80 °C.
350 Seawater samples were also collected approximately monthly from July 2017 to October 2018.

351 **DNA isolation**

352 DNA from surfaces of *C. nodosa* and *C. cylindracea* was isolated using a previously modified
353 and adapted protocol that allows for a selective epiphytic DNA isolation (Massana *et al.*, 1997;
354 Korlević *et al.*, submitted). Briefly, leaves and thalli are incubated in a lysis buffer and treated
355 with lysozyme and proteinase K. Following the incubations, the mixture containing lysed epiphytic
356 cells was separated from the leaves and thalli and extracted using phenol-chloroform. Finally, the
357 extracted DNA was precipitated using isopropanol. DNA from seawater picoplankton was extracted

358 from 0.2 µm polycarbonate filters according to Massana *et al.* (1997) with a slight modification.
359 Following the phenol-chloroform extraction, 1/10 of chilled 3 M sodium acetate (pH 5.2) was added.
360 DNA was precipitated by adding 1 volume of chilled isopropanol, incubating the mixtures overnight
361 at –20 °C and centrifuging at 20,000 × g and 4 °C for 21 min. The pellet was washed twice with
362 500 µl of chilled 70 % ethanol and centrifuged after each washing step at 20,000 × g and 4 °C for 5
363 min. Dried pellets were re-suspended in 50 – 100 µl of deionized water.

364 **Illumina 16S rRNA sequencing**

365 Illumina MiSeq sequencing of the V4 region of the 16S rRNA gene was performed as described
366 previously (Korlević *et al.*, submitted). The V4 region of the 16S rRNA gene was amplified using
367 a two-step PCR procedure. In the first PCR, the 515F (5'-GTGYCAGCMGCCGCGTAA-3')
368 and 806R (5'-GGACTACNVGGGTWTCTAAT-3') primers from the Earth Microbiome Project
369 (<http://press.igsb.anl.gov/earthmicrobiome/protocols-and-standards/16s/>) were used (Caporaso *et*
370 *al.*, 2012; Apprill *et al.*, 2015; Parada *et al.*, 2016). These primers contained on their 5' end a tagged
371 sequence. Purified PCR products were sent for Illumina MiSeq sequencing at IMGM Laboratories,
372 Martinsried, Germany. Prior to sequencing at IMGM, the second PCR amplification of the two-step
373 PCR procedure was performed using primers targeting the tagged region incorporated in the first
374 PCR. In addition, these primers contained adapter and sample-specific index sequences. Beside
375 samples, a positive and negative control for each sequencing batch was sequenced. The negative
376 control comprised PCR reactions without DNA template, while for a positive control a mock
377 community composed of evenly mixed DNA material originating from 20 bacterial strains (ATCC
378 MSA-1002, ATCC, USA) was used. Sequences obtained in this study have been deposited in the
379 European Nucleotide Archive (ENA) at EMBL-EBI under accession number PRJEB37267.

380 **Sequence analysis**

381 Obtained sequences were analyzed on the computer cluster Isabella (University Computing
382 Center, University of Zagreb) using mothur (version 1.43.0) (Schloss *et al.*, 2009) according
383 to the MiSeq Standard Operating Procedure (MiSeq SOP; https://mothur.org/wiki/MiSeq_SOP)
384 (Kozich *et al.*, 2013) and recommendations provided by the Riffomonas project to enhance data
385 reproducibility (<http://www.riffomonas.org/>). For alignment and classification of sequences the
386 SILVA SSU Ref NR 99 database (release 138; <http://www.arb-silva.de>) was used (Quast *et al.*,
387 2013; Yilmaz *et al.*, 2014). Pipeline data processing and visualization was done using R (version
388 3.6.0) (R Core Team, 2019) combined with packages vegan (version 2.5-6) (Oksanen *et al.*, 2019),
389 tidyverse (version 1.3.0) (Wickham *et al.*, 2019) and multiple other packages (Xie, 2014, 2015,
390 2020; Neuwirth, 2014; Xie *et al.*, 2018; Y. Xie, 2019b, 2019a; Allaire *et al.*, 2019; Zhu, 2019). The
391 detailed analysis procedure including the R Markdown file are available in the GitHub repository
392 (https://github.com/mkorlevic/Korlevic_EpiphyticDynamics_EnvironMicrobiol_2020). Based on
393 the ATCC MSA-1002 mock community included in the analysis an average sequencing error rate
394 of 0.01 % was determined, which is in line with previously reported values for next-generation
395 sequencing data (Kozich *et al.*, 2013; Schloss *et al.*, 2016). In addition, the negative controls
396 processed together with the samples yielded on average only 2 sequences after sequence quality
397 curation.

398 **Acknowledgments**

399 This work was founded by the Croatian Science Foundation through the MICRO-SEAGRASS
400 project (IP-2016-06-7118). We would like to thank Margareta Buterer for technical support, Paolo
401 Paliaga for help during sampling and the University Computing Center of the University of Zagreb
402 for access to the computer cluster Isabella.

403 **References**

- 404 Abraham, W.R. and Rohde, M. (2014) The family *Hyphomonadaceae*. In *The Prokaryotes: Alphaproteobacteria and Betaproteobacteria*. Rosenberg, E., DeLong, E.F., Lory, S., Stackebrandt, E., and Thompson, F. (eds). Berlin, Heidelberg: Springer-Verlag, pp. 283–299.
- 405
- 406
- 407 Allaire, J.J., Xie, Y., McPherson, J., Luraschi, J., Ushey, K., Atkins, A., et al. (2019) Rmarkdown: Dynamic documents for R.
- 408
- 409 Apprill, A., McNally, S., Parsons, R., and Weber, L. (2015) Minor revision to V4 region SSU rRNA 806R gene primer greatly increases detection of SAR11 bacterioplankton. *Aquat Microb Ecol* **75**: 129–137.
- 410
- 411
- 412 Armstrong, E., Rogerson, A., and Leftley, J. (2000) The abundance of heterotrophic protists associated with intertidal seaweeds. *Estuar Coast Shelf Sci* **50**: 415–424.
- 413
- 414 Bengtsson, M.M., Bühler, A., Brauer, A., Dahlke, S., Schubert, H., and Blindow, I. (2017) Eelgrass leaf surface microbiomes are locally variable and highly correlated with epibiotic eukaryotes. *Front Microbiol* **8**: 1312.
- 415
- 416
- 417 Bengtsson, M., Sjøtun, K., and Øvreås, L. (2010) Seasonal dynamics of bacterial biofilms on the kelp *Laminaria hyperborea*. *Aquat Microb Ecol* **60**: 71–83.
- 418
- 419 Borges, A.V. and Champenois, W. (2015) Seasonal and spatial variability of dimethylsulfoniopropionate (DMSP) in the Mediterranean seagrass *Posidonia oceanica*. *Aquat Bot* **125**: 72–79.
- 420
- 421
- 422 Burke, C., Kjelleberg, S., and Thomas, T. (2009) Selective extraction of bacterial DNA from the surfaces of macroalgae. *Appl Environ Microbiol* **75**: 252–256.
- 423

424 Burke, C., Steinberg, P., Rusch, D., Kjelleberg, S., and Thomas, T. (2011) Bacterial
425 community assembly based on functional genes rather than species. *Proc Natl Acad Sci U S A* **108**:
426 14288–14293.

427 Burke, C., Thomas, T., Lewis, M., Steinberg, P., and Kjelleberg, S. (2011) Composition,
428 uniqueness and variability of the epiphytic bacterial community of the green alga *Ulva australis*.
429 *ISME J* **5**: 590–600.

430 Burns, B.P., Goh, F., Allen, M., and Neilan, B.A. (2004) Microbial diversity of extant
431 stromatolites in the hypersaline marine environment of Shark Bay, Australia. *Environ Microbiol* **6**:
432 1096–1101.

433 Cai, X., Gao, G., Yang, J., Tang, X., Dai, J., Chen, D., and Song, Y. (2014) An ultrasonic
434 method for separation of epiphytic microbes from freshwater submerged macrophytes. *J Basic
435 Microbiol* **54**: 758–761.

436 Caporaso, J.G., Lauber, C.L., Walters, W.A., Berg-Lyons, D., Huntley, J., Fierer, N., et al.
437 (2012) Ultra-high-throughput microbial community analysis on the Illumina HiSeq and MiSeq
438 platforms. *ISME J* **6**: 1621–1624.

439 Cardoso, D.C., Cretoiu, M.S., Stal, L.J., and Bolhuis, H. (2019) Seasonal development of a
440 coastal microbial mat. *Sci Rep* **9**: 9035.

441 Crump, B.C. and Koch, E.W. (2008) Attached bacterial populations shared by four species of
442 aquatic angiosperms. *Appl Environ Microbiol* **74**: 5948–5957.

443 Crump, B.C., Wojahn, J.M., Tomas, F., and Mueller, R.S. (2018) Metatranscriptomics and
444 amplicon sequencing reveal mutualisms in seagrass microbiomes. *Front Microbiol* **9**: 388.

- 445 Egan, S., Harder, T., Burke, C., Steinberg, P., Kjelleberg, S., and Thomas, T. (2013) The
446 seaweed holobiont: Understanding seaweed-bacteria interactions. *FEMS Microbiol Rev* **37**:
447 462–476.
- 448 Fahimipour, A.K., Kardish, M.R., Lang, J.M., Green, J.L., Eisen, J.A., and Stachowicz, J.J.
449 (2017) Global-scale structure of the eelgrass microbiome. *Appl Environ Microbiol* **83**: e03391–16.
- 450 Gilbert, J.A., Field, D., Swift, P., Newbold, L., Oliver, A., Smyth, T., et al. (2009) The
451 seasonal structure of microbial communities in the Western English Channel. *Environ Microbiol* **11**:
452 3132–3139.
- 453 Gilbert, J.A., Steele, J.A., Caporaso, J.G., Steinbrück, L., Reeder, J., Temperton, B., et al.
454 (2012) Defining seasonal marine microbial community dynamics. *ISME J* **6**: 298–308.
- 455 Higashioka, Y., Kojima, H., Watanabe, T., and Fukui, M. (2015) Draft genome sequence of
456 *Desulfatitalea tepidiphila* S28bF^T. *Genome Announc* **3**: e01326–15.
- 457 Hollants, J., Leliaert, F., De Clerck, O., and Willems, A. (2013) What we can learn from sushi:
458 A review on seaweed-bacterial associations. *FEMS Microbiol Ecol* **83**:
- 459 Jost, L. (2006) Entropy and diversity. *Oikos* **113**: 363–375.
- 460 Jørgensen, B.B. and Gallardo, V.A. (1999) *Thioploca* spp.: filamentous sulfur bacteria with
461 nitrate vacuoles. *FEMS Microbiol Ecol* **28**: 301–313.
- 462 Korlević, M., Markovski, M., Zhao, Z., Herndl, G.J., and Najdek, M. Selective DNA and
463 protein isolation from marine macrophyte surfaces.
- 464 Korlević, M., Pop Ristova, P., Garić, R., Amann, R., and Orlić, S. (2015) Bacterial diversity in
465 the South Adriatic Sea during a strong, deep winter convection year. *Appl Environ Microbiol* **81**:
466 1715–1726.

- 467 Kozich, J.J., Westcott, S.L., Baxter, N.T., Highlander, S.K., and Schloss, P.D. (2013)
- 468 Development of a dual-index sequencing strategy and curation pipeline for analyzing amplicon
- 469 sequence data on the MiSeq Illumina sequencing platform. *Appl Environ Microbiol* **79**: 5112–5120.
- 470 Kuever, J. (2014) The family *Desulfobulbaceae*. In *The Prokaryotes: Deltaproteobacteria and*
- 471 *Epsilonproteobacteria*. Rosenberg, E., DeLong, E.F., Lory, S., Stackebrandt, E., and Thompson, F.
- 472 (eds). Berlin, Heidelberg: Springer-Verlag, pp. 75–86.
- 473 Lachnit, T., Blümel, M., Imhoff, J.F., and Wahl, M. (2009) Specific epibacterial communities
- 474 on macroalgae: Phylogeny matters more than habitat. *Aquat Biol* **5**: 181–186.
- 475 Lachnit, T., Meske, D., Wahl, M., Harder, T., and Schmitz, R. (2011) Epibacterial community
- 476 patterns on marine macroalgae are host-specific but temporally variable. *Environ Microbiol* **13**:
- 477 655–665.
- 478 Longford, S., Tujula, N., Crocetti, G., Holmes, A., Holmström, C., Kjelleberg, S., et al. (2007)
- 479 Comparisons of diversity of bacterial communities associated with three sessile marine eukaryotes.
- 480 *Aquat Microb Ecol* **48**: 217–229.
- 481 Mancuso, F.P., D'Hondt, S., Willems, A., Airoldi, L., and De Clerck, O. (2016) Diversity
- 482 and temporal dynamics of the epiphytic bacterial communities associated with the canopy-forming
- 483 seaweed *Cystoseira compressa* (Esper) Gerloff and Nizamuddin. *Front Microbiol* **7**: 476.
- 484 Margulis, L. (1991) Symbiogenesis and symbioticism. In *Symbiosis as a Source of*
- 485 *Evolutionary Innovation: Speciation and Morphogenesis*. Margulis, L. and Fester, R. (eds).
- 486 Cambridge, Massachusetts: The MIT Press, pp. 1–14.
- 487 Massana, R., Murray, A.E., Preston, C.M., and DeLong, E.F. (1997) Vertical distribution and
- 488 phylogenetic characterization of marine planktonic *Archaea* in the Santa Barbara Channel. *Appl*
- 489 *Environ Microbiol* **63**: 50–56.

- 490 McIlroy, S.J. and Nielsen, P.H. (2014) The family *Saprospiraceae*. In *The Prokaryotes: Other*
491 *Major Lineages of Bacteria and the Archaea*. Rosenberg, E., DeLong, E.F., Lory, S., Stackebrandt,
492 E., and Thompson, F. (eds). Berlin, Heidelberg: Springer-Verlag, pp. 863–889.
- 493 Michelou, V.K., Caporaso, J.G., Knight, R., and Palumbi, S.R. (2013) The ecology of microbial
494 communities associated with *Macrocystis pyrifera*. *PloS One* **8**: e67480.
- 495 Miranda, L.N., Hutchison, K., Grossman, A.R., and Brawley, S.H. (2013) Diversity and
496 abundance of the bacterial community of the red macroalga *Porphyra umbilicalis*: Did bacterial
497 farmers produce macroalgae? *PloS One* **8**: e58269.
- 498 Mobberley, J.M., Ortega, M.C., and Foster, J.S. (2012) Comparative microbial diversity
499 analyses of modern marine thrombolitic mats by barcoded pyrosequencing. *Environ Microbiol* **14**:
500 82–100.
- 501 Morrissey, K.L., Çavas, L., Willems, A., and De Clerck, O. (2019) Disentangling the influence
502 of environment, host specificity and thallus differentiation on bacterial communities in siphonous
503 green seaweeds. *Front Microbiol* **10**: 717.
- 504 Neuwirth, E. (2014) RColorBrewer: ColorBrewer palettes.
- 505 Nõges, T., Luup, H., and Feldmann, T. (2010) Primary production of aquatic macrophytes and
506 their epiphytes in two shallow lakes (Peipsi and Võrtsjärv) in Estonia. *Aquat Ecol* **44**: 83–92.
- 507 Oksanen, J., Blanchet, F.G., Friendly, M., Kindt, R., Legendre, P., McGlinn, D., et al. (2019)
508 Vegan: Community ecology package.
- 509 Parada, A.E., Needham, D.M., and Fuhrman, J.A. (2016) Every base matters: Assessing small
510 subunit rRNA primers for marine microbiomes with mock communities, time series and global field
511 samples. *Environ Microbiol* **18**: 1403–1414.

- 512 Pujalte, M.J., Lucena, T., Ruvira, M.A., Arahal, D.R., and Macián, M.C. (2014) The family
513 *Rhodobacteraceae*. In *The Prokaryotes: Alphaproteobacteria and Betaproteobacteria*. Rosenberg,
514 E., DeLong, E.F., Lory, S., Stackebrandt, E., and Thompson, F. (eds). Berlin, Heidelberg:
515 Springer-Verlag, pp. 439–512.
- 516 Quast, C., Pruesse, E., Yilmaz, P., Gerken, J., Schweer, T., Yarza, P., et al. (2013) The SILVA
517 ribosomal RNA gene database project: Improved data processing and web-based tools. *Nucleic
518 Acids Res* **41**: D590–D596.
- 519 R Core Team (2019) A language and environment for statistical computing, Vienna, Austria:
520 R Foundation for Statistical Computing.
- 521 Reisser, J., Shaw, J., Hallegraeff, G., Proietti, M., Barnes, D.K.A., Thums, M., et al. (2014)
522 Millimeter-sized marine plastics: A new pelagic habitat for microorganisms and invertebrates. *PLoS
523 One* **9**: e100289.
- 524 Reyes, J. and Sansón, M. (2001) Biomass and production of the epiphytes on the leaves of
525 *Cymodocea nodosa* in the Canary Islands. *Bot Mar* **44**: 307–313.
- 526 Rickert, E., Wahl, M., Link, H., Richter, H., and Pohnert, G. (2016) Seasonal variations in
527 surface metabolite composition of *Fucus vesiculosus* and *Fucus serratus* from the Baltic Sea. *PLoS
528 One* **11**: e0168196.
- 529 Roth-Schulze, A.J., Zozaya-Valdés, E., Steinberg, P.D., and Thomas, T. (2016) Partitioning of
530 functional and taxonomic diversity in surface-associated microbial communities. *Environ Microbiol
531* **18**: 4391–4402.
- 532 Ruitton, S., Verlaque, M., and Boudouresque, C.F. (2005) Seasonal changes of the introduced
533 *Caulerpa racemosa* var. *cylindracea* (Caulerpales, Chlorophyta) at the northwest limit of its
534 Mediterranean range. *Aquat Bot* **82**: 55–70.

- 535 Salaün, S., La Barre, S., Santos-Goncalvez, M.D., Potin, P., Haras, D., and Bazire, A. (2012)
- 536 Influence of exudates of the kelp *Laminaria digitata* on biofilm formation of associated and
- 537 exogenous bacterial epiphytes. *Microb Ecol* **64**: 359–369.
- 538 Schloss, P.D., Jenior, M.L., Koumpouras, C.C., Westcott, S.L., and Highlander, S.K. (2016)
- 539 Sequencing 16S rRNA gene fragments using the PacBio SMRT DNA sequencing system. *PeerJ* **4**:
- 540 e1869.
- 541 Schloss, P.D., Westcott, S.L., Ryabin, T., Hall, J.R., Hartmann, M., Hollister, E.B., et al.
- 542 (2009) Introducing mothur: Open-source, platform-independent, community-supported software for
- 543 describing and comparing microbial communities. *Appl Environ Microbiol* **75**: 7537–7541.
- 544 Tarquinio, F., Hyndes, G.A., Laverock, B., Koenders, A., and Säwström, C. (2019) The seagrass
- 545 holobiont: Understanding seagrass-bacteria interactions and their role in seagrass ecosystem
- 546 functioning. *FEMS Microbiol Lett* **366**: fnz057.
- 547 Tujula, N.A., Crocetti, G.R., Burke, C., Thomas, T., Holmström, C., and Kjelleberg, S. (2010)
- 548 Variability and abundance of the epiphytic bacterial community associated with a green marine
- 549 *Ulvacean* alga. *ISME J* **4**: 301–311.
- 550 Ugarelli, K., Laas, P., and Stingl, U. (2019) The microbial communities of leaves and roots
- 551 associated with turtle grass (*Thalassia testudinum*) and manatee grass (*Syringodium filiforme*) are
- 552 distinct from seawater and sediment communities, but are similar between species and sampling
- 553 sites. *Microorganisms* **7**: 4.
- 554 Uku, J., Björk, M., Bergman, B., and Díez, B. (2007) Characterization and comparison of
- 555 prokaryotic epiphytes associated with three East African seagrasses. *J Phycol* **43**: 768–779.
- 556 Weidner, S., Arnold, W., Stackebrandt, E., and Pühler, A. (2000) Phylogenetic analysis
- 557 of bacterial communities associated with leaves of the seagrass *Halophila stipulacea* by a
- 558 culture-independent small-subunit rRNA gene approach. *Microb Ecol* **39**: 22–31.

- 559 Wickham, H., Averick, M., Bryan, J., Chang, W., McGowan, L.D., François, R., et al. (2019)
- 560 Welcome to the tidyverse. *J Open Source Softw* **4**: 1686.
- 561 Xie, Y. (2015) Dynamic Documents with R and knitr, 2nd ed. Boca Raton, Florida: Chapman
- 562 and Hall/CRC.
- 563 Xie, Y. (2014) Knitr: A comprehensive tool for reproducible research in R. In *Implementing*
- 564 *Reproducible Computational Research*. Stodden, V., Leisch, F., and Peng, R.D. (eds). New York:
- 565 Chapman and Hall/CRC, pp. 3–32.
- 566 Xie, Y. (2019a) Knitr: A general-purpose package for dynamic report generation in R.
- 567 Xie, Y. (2019b) TinyTeX: A lightweight, cross-platform, and easy-to-maintain LaTeX
- 568 distribution based on TeX Live. *TUGboat* **40**: 30–32.
- 569 Xie, Y. (2020) TinyTeX: Helper functions to install and maintain 'TeX Live', and compile
- 570 'TeX' documents.
- 571 Xie, Y., Allaire, J.J., and Grolemund, G. (2018) R Markdown: The Definitive Guide, 1st ed.
- 572 Boca Raton, Florida: Chapman and Hall/CRC.
- 573 Xie, Z., Lin, W., and Luo, J. (2017) Comparative phenotype and genome analysis of *Cellvibrio*
- 574 sp. PR1, a xylanolytic and agarolytic bacterium from the Pearl River. *BioMed Res Int* **2017**:
- 575 6304248.
- 576 Yilmaz, P., Parfrey, L.W., Yarza, P., Gerken, J., Pruesse, E., Quast, C., et al. (2014) The
- 577 SILVA and "All-Species Living Tree Project (LTP)" taxonomic frameworks. *Nucleic Acids Res* **42**:
- 578 D643–D648.
- 579 Zavodnik, N., Travizi, A., and de Rosa, S. (1998) Seasonal variations in the rate of
- 580 photosynthetic activity and chemical composition of the seagrass *Cymodocea nodosa* (Ucr.) Asch.
- 581 *Sci Mar* **62**: 301–309.

583 **Figure legends**

584 **Fig. 1.** Proportion of shared bacterial and archaeal OTUs (Jaccard's similarity coefficient) and
585 shared bacterial and archaeal communities (Bray-Curtis similarity coefficient) between assemblages
586 associated with the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C. cylindracea*
587 (mixed and monospecific settlement) and communities in the ambient seawater.

588 **Fig. 2.** Proportion of shared bacterial and archaeal communities (Bray-Curtis similarity coefficient)
589 and shared bacterial and archaeal OTUs (Jaccard's similarity coefficient) between consecutive
590 sampling dates and from the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C.*
591 *cylindracea* (mixed and monospecific settlement) and in the ambient seawater.

592 **Fig. 3.** Principal Coordinates Analysis (PCoA) of Bray-Curtis distances based on OTU abundances
593 of bacterial and archaeal communities from the surfaces of the macrophytes *C. nodosa* (mixed
594 settlement) and *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater.
595 Samples from the same environment or same season are labeled in different colors. The proportion
596 of explained variation by each axis is shown on the corresponding axis in parentheses.

597 **Fig. 4.** Taxonomic classification and relative contribution of the most abundant (> 1 %) bacterial
598 and archaeal sequences on the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C.*
599 *cylindracea* (mixed and monospecific settlement) and in the ambient seawater. NR – No Relative
600 (sequences without known relatives within the corresponding group)

601 **Fig. 5.** Taxonomic classification and relative contribution of the most abundant (> 1 %)
602 cyanobacterial sequences on the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C.*
603 *cylindracea* (mixed and monospecific settlement) and in the ambient seawater. The proportion
604 of cyanobacterial sequences in the total bacterial and archaeal community is given above the
605 corresponding bar. NR – No Relative (sequences without known relatives within the corresponding
606 group)

607 **Fig. 6.** Taxonomic classification and relative contribution of the most abundant (> 2 %) sequences
608 within the *Bacteroidota* on the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C.*
609 *cylindracea* (mixed and monospecific settlement) and in the ambient seawater. The proportion of
610 sequences classified as *Bacteroidota* in the total bacterial and archaeal community is given above the
611 corresponding bar. NR – No Relative (sequences without known relatives within the corresponding
612 group)

613 **Fig. 7.** Taxonomic classification and relative contribution of the most abundant (> 2 %)
614 alphaproteobacterial sequences on the surfaces of the macrophytes *C. nodosa* (mixed settlement)
615 and *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater. The
616 proportion of alphaproteobacterial sequences in the total bacterial and archaeal community is given
617 above the corresponding bar. NR – No Relative (sequences without known relatives within the
618 corresponding group)

619 **Fig. 8.** Taxonomic classification and relative contribution of the most abundant (> 2 %)
620 gammaproteobacterial sequences on the surfaces of the macrophytes *C. nodosa* (mixed settlement)
621 and *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater. The
622 proportion of gammaproteobacterial sequences in the total bacterial and archaeal community is given
623 above the corresponding bar. NR – No Relative (sequences without known relatives within
624 the corresponding group)

625 **Fig. 9.** Taxonomic classification and relative contribution of the most abundant (> 1 %) sequences
626 within the *Desulfobacterota* on the surfaces of the macrophytes *C. nodosa* (mixed settlement) and
627 *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater. The proportion
628 of sequences classified as *Desulfobacterota* in the total bacterial and archaeal community is given
629 above the corresponding bar. NR – No Relative (sequences without known relatives within the
630 corresponding group)

Jaccard's Similarity Coefficient

<i>Caulerpa cylindracea</i> (Mixed)	0.28		
<i>Caulerpa cylindracea</i> (Monospecific)	0.27	0.35	
Seawater	0.12	0.10	0.10

Bray-Curtis Similarity Coefficient

<i>Caulerpa cylindracea</i> (Mixed)	0.40		
<i>Caulerpa cylindracea</i> (Monospecific)	0.38	0.77	
Seawater	0.06	0.05	0.06

Fig. 1. Proportion of shared bacterial and archaeal OTUs (Jaccard's similarity coefficient) and shared bacterial and archaeal communities (Bray-Curtis similarity coefficient) between assemblages associated with the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C. cylindracea* (mixed and monospecific settlement) and communities in the ambient seawater.

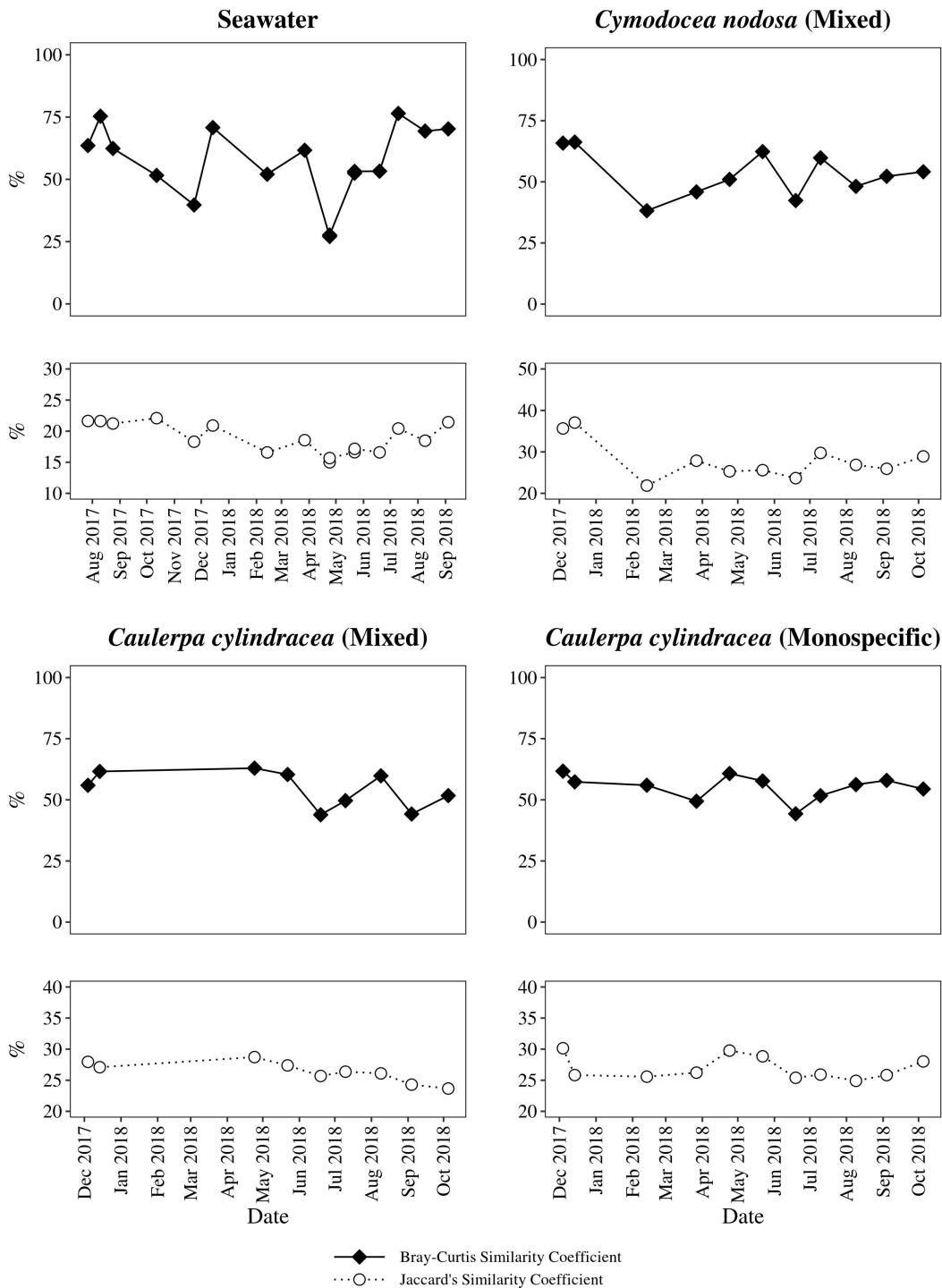


Fig. 2. Proportion of shared bacterial and archaeal communities (Bray-Curtis similarity coefficient) and shared bacterial and archaeal OTUs (Jaccard's similarity coefficient) between consecutive sampling dates and from the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater.

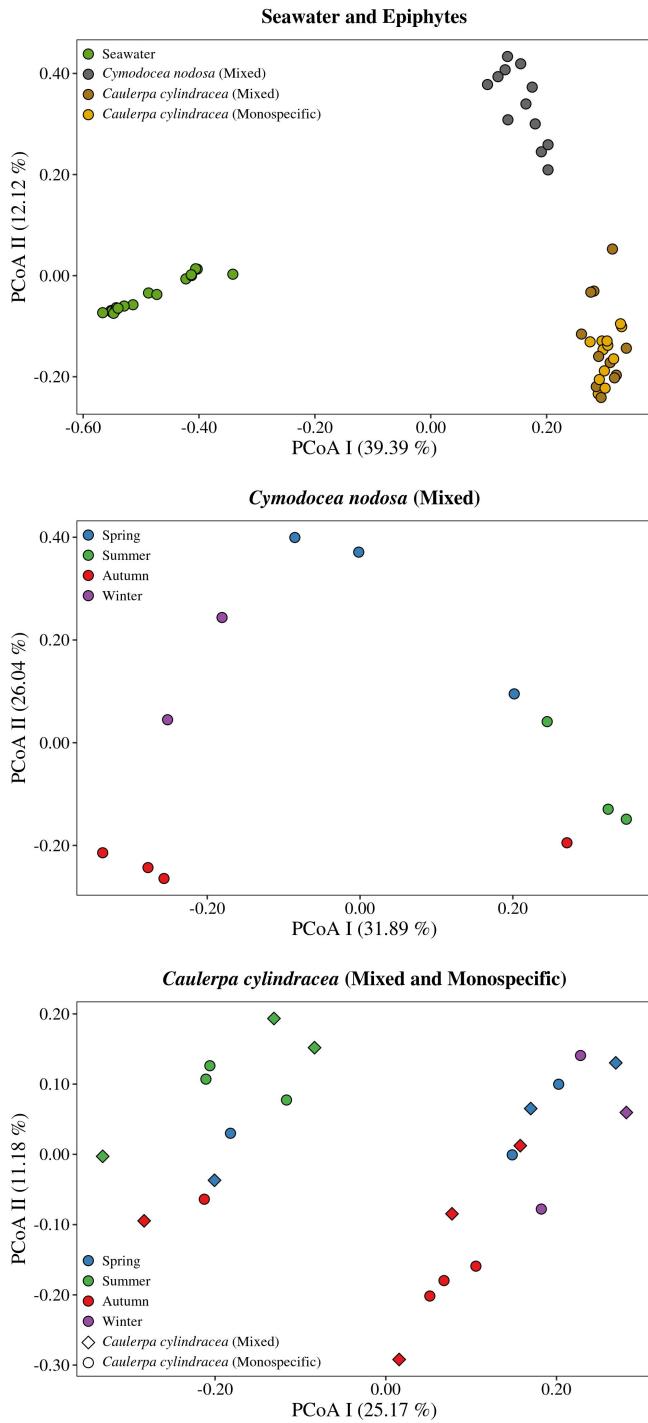


Fig. 3. Principal Coordinates Analysis (PCoA) of Bray-Curtis distances based on OTU abundances of bacterial and archaeal communities from the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater. Samples from the same environment or same season are labeled in different colors. The proportion of explained variation by each axis is shown on the corresponding axis in parentheses.

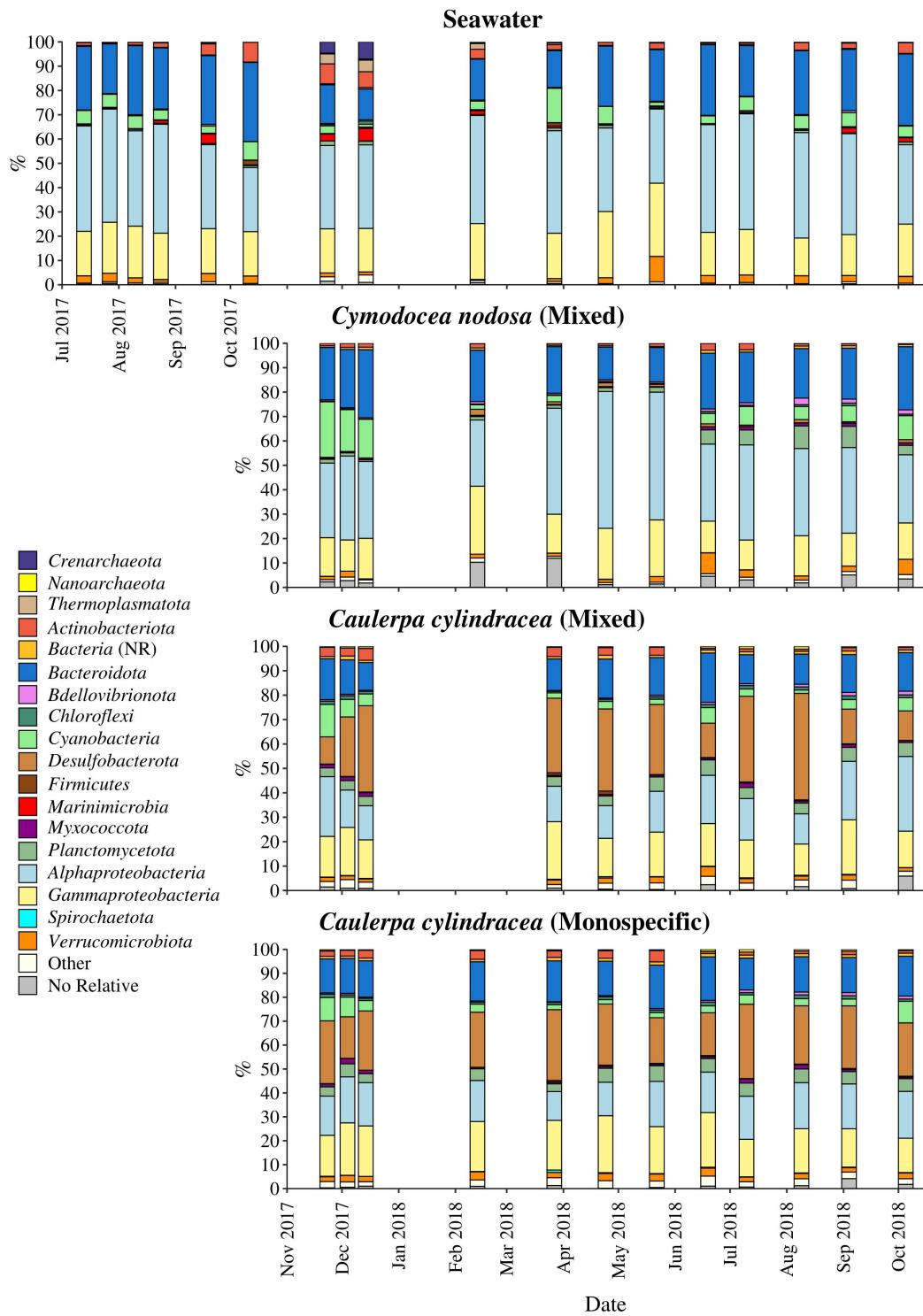


Fig. 4. Taxonomic classification and relative contribution of the most abundant (> 1 %) bacterial and archaeal sequences on the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater. NR – No Relative (sequences without known relatives within the corresponding group)

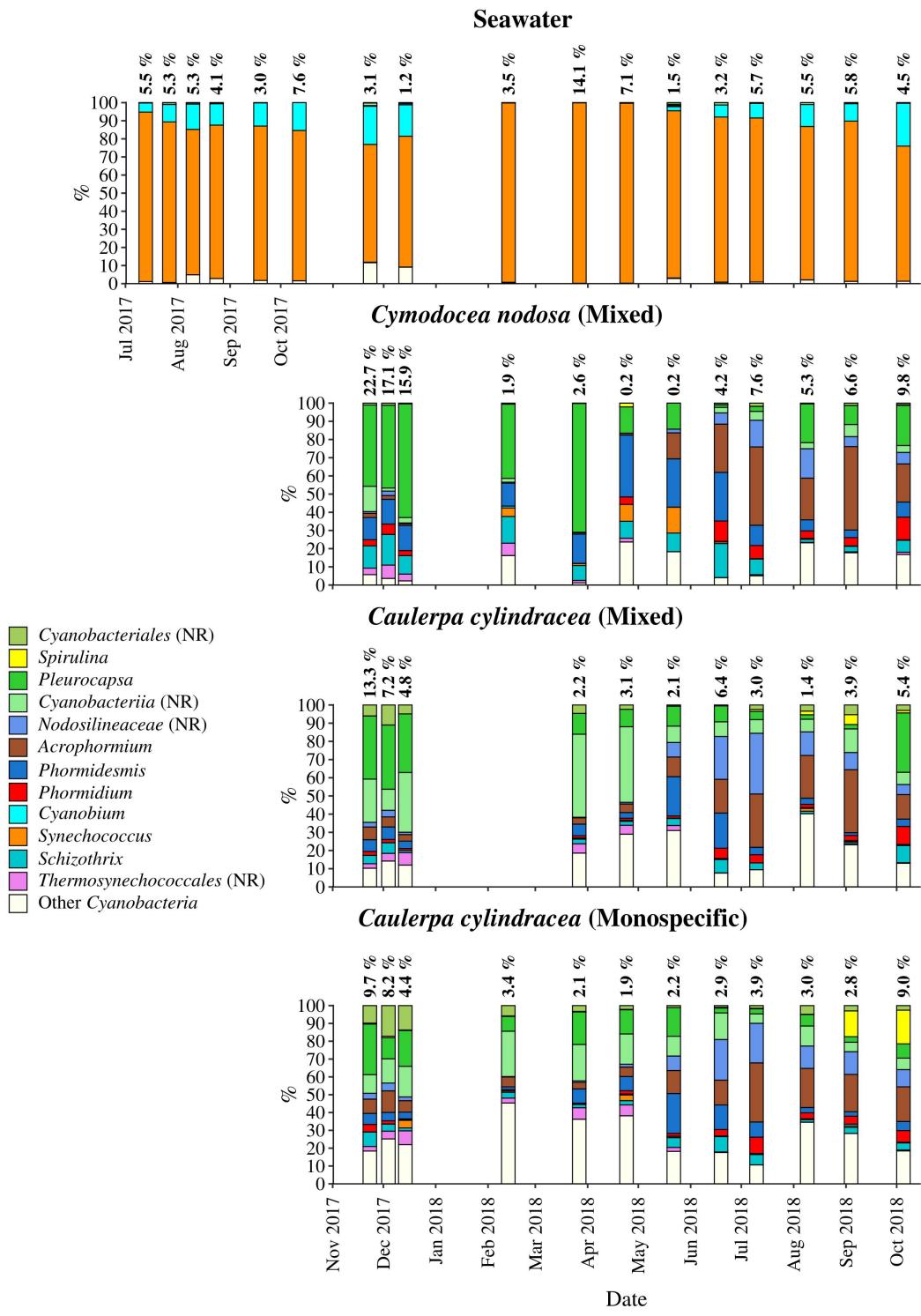


Fig. 5. Taxonomic classification and relative contribution of the most abundant (> 1 %) cyanobacterial sequences on the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater. The proportion of cyanobacterial sequences in the total bacterial and archaeal community is given above the corresponding bar. NR – No Relative (sequences without known relatives within the corresponding group)

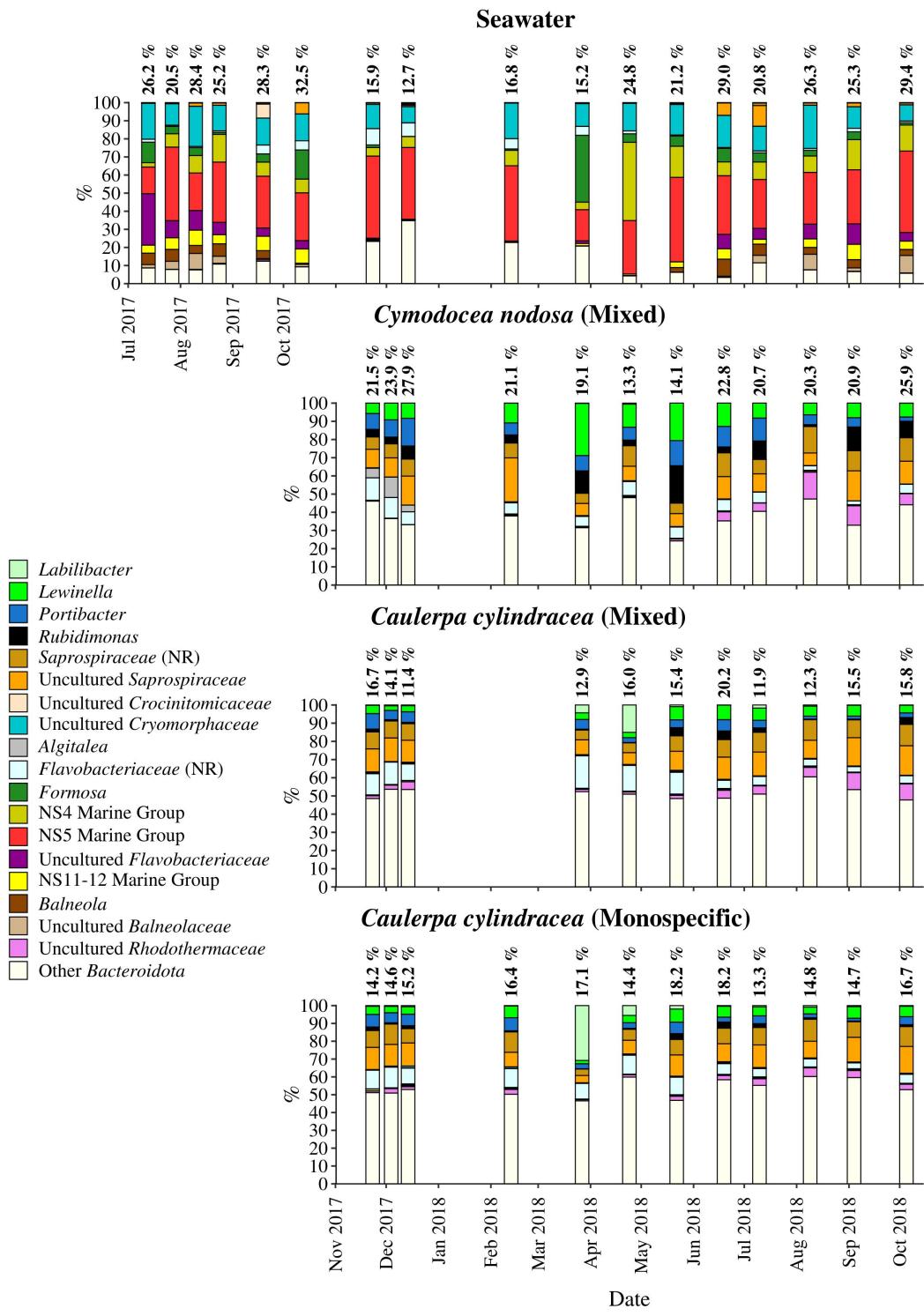


Fig. 6. Taxonomic classification and relative contribution of the most abundant (> 2 %) sequences within the *Bacteroidota* on the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater. The proportion of sequences classified as *Bacteroidota* in the total bacterial and archaeal community is given above the corresponding bar. NR – No Relative (sequences without known relatives within the corresponding group)

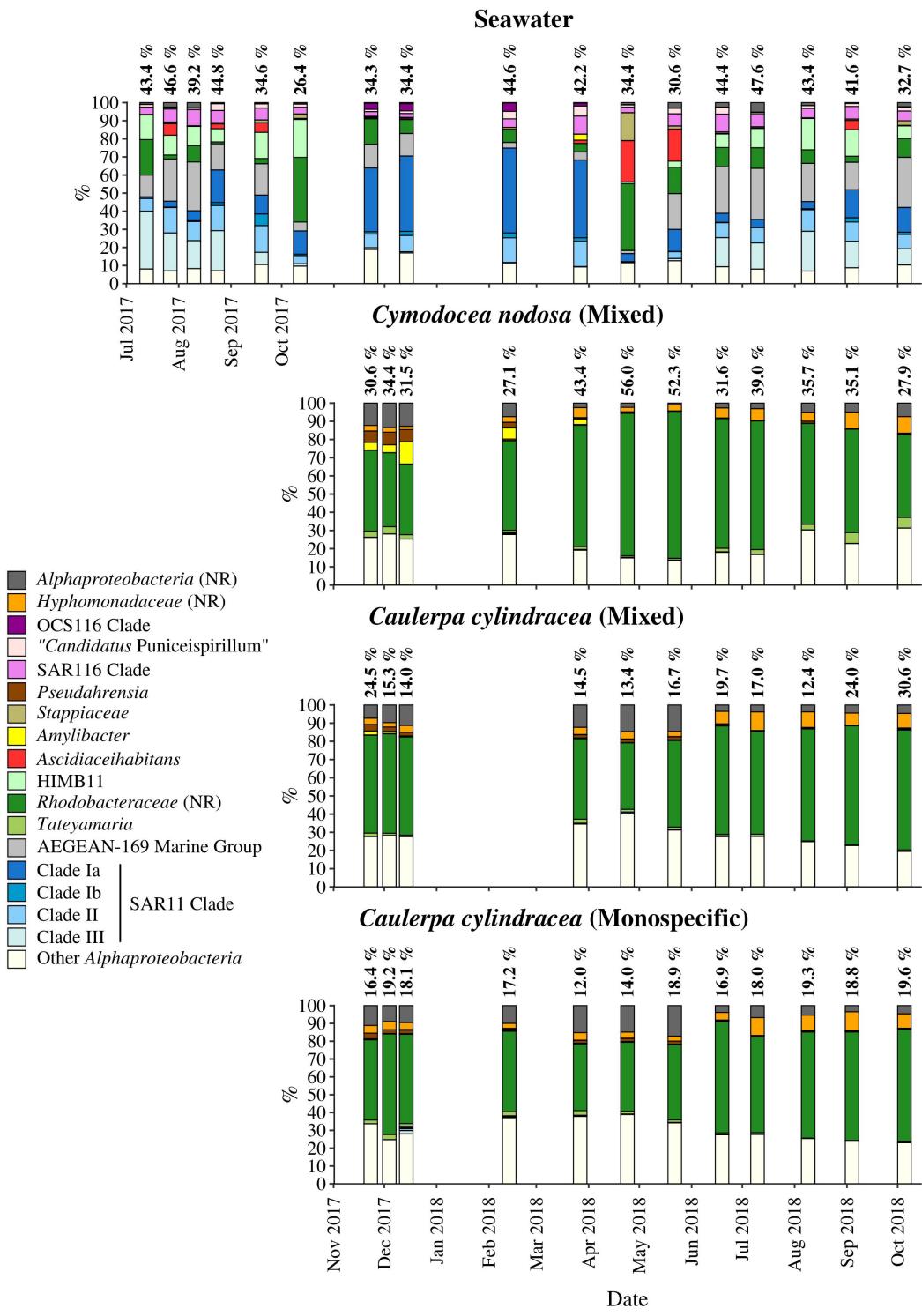


Fig. 7. Taxonomic classification and relative contribution of the most abundant (> 2 %) alphaproteobacterial sequences on the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater. The proportion of alphaproteobacterial sequences in the total bacterial and archaeal community is given above the corresponding bar. NR – No Relative (sequences without known relatives within the corresponding group)

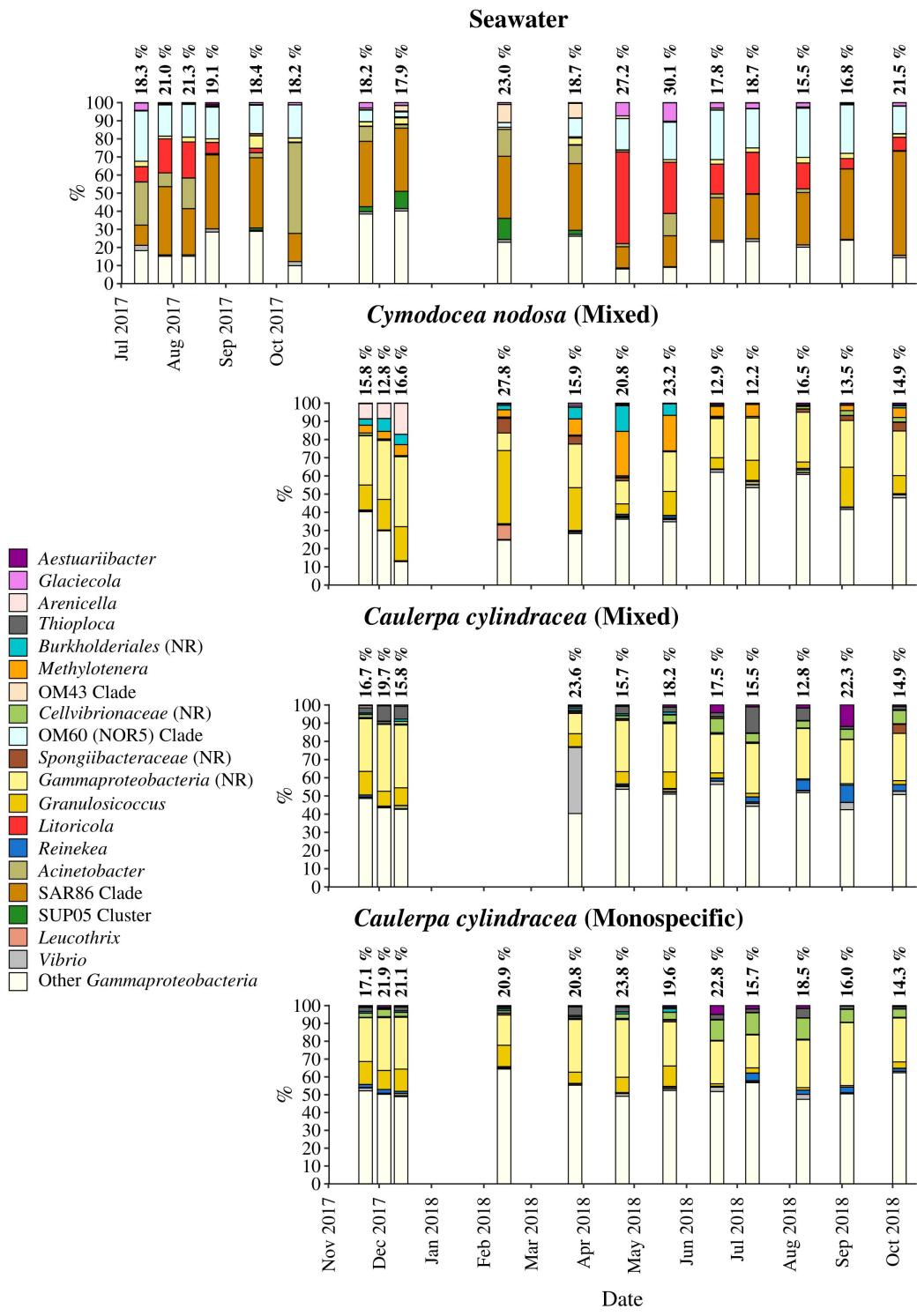


Fig. 8. Taxonomic classification and relative contribution of the most abundant (> 2 %) gammaproteobacterial sequences on the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater. The proportion of gammaproteobacterial sequences in the total bacterial and archaeal community is given above the corresponding bar. NR – No Relative (sequences without known relatives within the corresponding group)

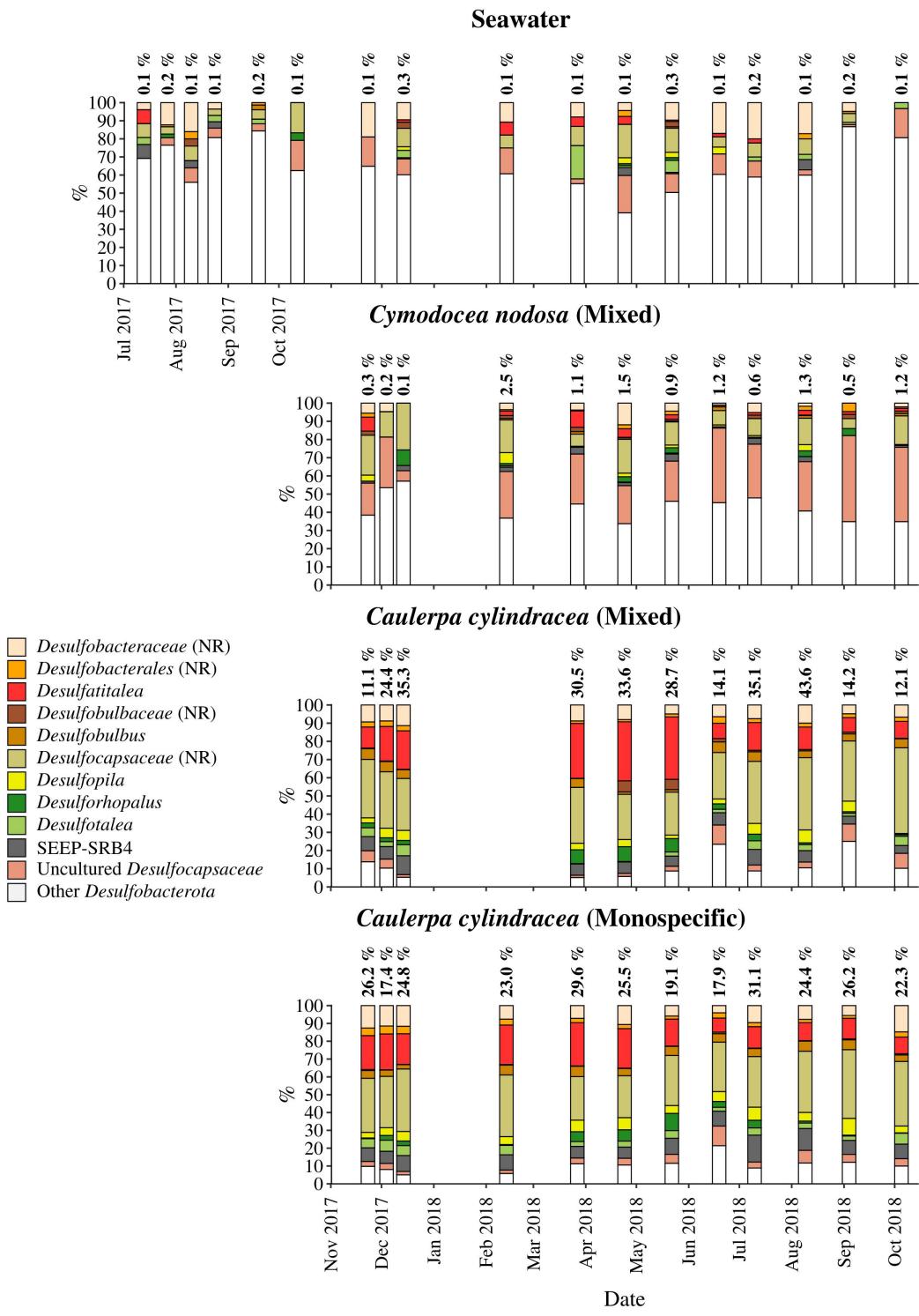


Fig. 9. Taxonomic classification and relative contribution of the most abundant (> 1 %) sequences within the *Desulfobacterota* on the surfaces of the macrophytes *C. nodosa* (mixed settlement) and *C. cylindracea* (mixed and monospecific settlement) and in the ambient seawater. The proportion of sequences classified as *Desulfobacterota* in the total bacterial and archaeal community is given above the corresponding bar. NR – No Relative (sequences without known relatives within the corresponding group)