Warming the phycosphere: differential effect of temperature on the use of diatom-derived carbon by two copiotrophic bacterial taxa

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Significance Statement

Phytoplankton-derived exudates represent the main carbon source for heterotrophic bacteria in coastal waters. During microphytoplankton blooms, some fast-growing bacterial groups colonize the phycosphere of microphytoplankton cells, which provides them with a faster access to recently fixed DOC. Ocean warming may alter the bacterial cell-specific uptake of algal-derived C, with important biogeochemical implications, but quantifying C fluxes of microphytoplankton-associated bacteria is challenging as it requires resolving microbial cell activities below the microscale spatial level. Using nanoSIMS analysis, we quantified the impact of experimental warming on algal-derived C uptake by two model copiotrophic taxa (Rhodobacteraceae and Flavobacteria), which were abundant components of the algal/particle-associated fraction year-round at a coastal station of the Bay of Biscay. We found that Flavobacteria were the main contributor to total C uptake under high diatom abundance, and experimental warming enhanced their uptake activity up to 6-fold.

Summary

Heterotrophic bacteria associated with microphytoplankton, particularly those colonizing the phycosphere, are major players in the remineralization of algal-derived carbon. Ocean warming might impact DOC uptake by microphytoplankton-associated bacteria with unknown biogeochemical implications. Here, by incubating natural seawater samples at 3 different temperatures we analyzed the effect of experimental warming on the abundance and C and N uptake activity of Rhodobacteraceae and Flavobacteria, two bacterial groups typically associated with microphytoplankton. Using NanoSIMS single-cell analysis we quantified the temperature-sensitivity of these two taxonomic groups to the uptake of algal-derived DOC in the microphytoplankton-associated fraction with ^{13}C -bicarbonate and ^{15}N -leucine as tracers. We found that cell-specific ^{13}C uptake was similar for both groups ($\sim 0.42 \text{ fg C h}^{-1} \mu\text{m}^{-3}$), but Rhodobacteraceae were more active in ^{15}N -leucine uptake. Due to the higher abundance of Flavobacteria associated with microphytoplankton, this group incorporated 4-fold more carbon than Rhodobacteraceae. Cell-specific ^{13}C uptake was influenced by temperature, but no significant differences were found for ^{15}N -leucine uptake. Our results show that the contribution of Flavobacteria and Rhodobacteraceae to C assimilation increased up to 6-fold and 2-fold, respectively, with an increase of 3°C above ambient temperature, suggesting that warming may differently affect the contribution of distinct copiotrophic bacterial taxa to carbon cycling.

Introduction

Marine phytoplankton contribute 46% of the biosphere's net primary production (Field, 1998; Behrenfeld *et al.*, 2001) and represent one of the most important sources of dissolved organic carbon (DOC)

(Norrman *et al.*, 1995; Moran, 2015). In particular, coastal microphytoplankton blooms dominated by fast growing diatoms are responsible for a major share of global primary production (Joiris *et al.*, 1982; Caffrey *et al.*, 1998). DOC compounds released during these blooms are actively taken up by heterotrophic bacteria (Chrost and Faust, 1983; Meon and Kirchman, 2001), which can i) transfer this organic matter to higher trophic levels through the microbial loop, ii) convert it into more refractory carbon or iii) release it to the atmosphere as CO₂ through bacterial respiration (Azam, 1998; Jiao *et al.*, 2010). Trophic interactions between bacteria and microphytoplankton are influenced by the composition of bacterial communities (Pinhassi *et al.*, 2004; Grossart *et al.*, 2005; Klindworth *et al.*, 2014), as distinct taxonomic groups have different roles in the uptake of algal-derived DOC (Teeling *et al.*, 2012). Some environmental parameters such as temperature may also have a key impact on these interactions, by promoting the C fluxes between both bacterial and phytoplankton communities (Arandia-Gorostidi *et al.*, 2017b).

Remarkably, most of the studies addressing the bacterial-phytoplankton coupling have focused exclusively on free-living communities (Duarte *et al.*, 2005; Hoppe *et al.*, 2008; von Scheibner *et al.*, 2014), overlooking the heterotrophic bacteria found in the vicinity of microphytoplankton cells, i.e., the phycosphere. The phycosphere is defined as the region immediately surrounding the algal cells, which is highly enriched with organic molecules exuded by the microphytoplankton cell (Seymour *et al.*, 2017). It has been proposed that bacteria associated with microphytoplankton may have stronger physical and metabolic interactions with microphytoplankton cells than their free-living counterparts (Rooney-Varga *et al.*, 2005; Grossart *et al.*, 2007), since the uptake of an important fraction of the microphytoplankton exudates takes place in this limited area where bacteria have fast access to recently fixed C (Smriga *et al.*, 2016). A previous study has indicated that increasing temperatures may have a higher impact on the C

uptake by bacteria associated to phytoplankton than by their free-living counterparts (Arandia-Gorostidi *et al.*, 2017b). However, the temperature-response of different bacterial taxa to the uptake of DOC is largely unknown, mostly due to the difficulties in coupling taxonomic identification and cell-specific substrate uptakes in small-scale environments such as the phycosphere.

In this work we combined taxonomic identification by CARD-FISH with stable isotope probing and nanoSIMS analysis to determine the effect of experimental warming on the cell-specific DOM assimilation-rates of two ecologically relevant bacterial groups typically found in association with microphytoplankton: Flavobacteria and Rhodobacteraceae (Buchan *et al.*, 2005; Gómez-Pereira *et al.*, 2010; Fernández-Gómez *et al.*, 2013; Buchan *et al.*, 2014). As explained in Buchan *et al.* (2014), the Rhodobacteraceae family includes the *Roseobacter* clade, which contains a large number of genera (>50), and marine Flavobacteria contains representatives of the Flavobacteriales order, including two families (Flavobacteriaceae and Cryomorphaceae). Both, Rhodobacteraceae and Flavobacteria groups have been described as major players in the use and remineralization of algal-derived DOC (Buchan *et al.*, 2014), but their relative contribution to the consumption of photosynthates is unknown. Coinciding with high diatom abundance at *in situ* conditions, we quantified the single-cell uptake activity of both leucine and algal-derived DOC uptake by both bacterial taxonomic groups in order to assess differences in their potential contribution to carbon and nitrogen fluxes under increased temperature conditions.

Results

*Environmental analysis of particle-associated bacterial taxa under different chlorophyll *a* conditions*

In order to identify bacterial taxa that were preferentially associated to particles (including microphytoplankton cells), we analysed the bacterial community composition in the particle-associated (fraction $>3\ \mu\text{m}$), and the free-living fraction (fraction between $3\ \mu\text{m}$ and $0.22\ \mu\text{m}$). In a total of 11 samples collected during 3-years (2009-2012), encompassing from relatively low ($0.42\ \mu\text{g L}^{-1}$) to high ($1.81\ \mu\text{g L}^{-1}$) Chl *a* concentrations, we found that in the particle-associated fraction the Flavobacteriales order (within Flavobacteria) was generally the most abundant group (Figure 1B). Its average contribution to total abundance (22%) was significantly higher than Rhodobacterales (11%, Wilcoxon test, $p\text{-value}=0.02$), while the latter was more abundant than other groups such as Alteromonadales (4.6%, Wilcoxon test, $p\text{-value}=0.008$). Occasionally, other groups such as Sphingomonadales (Flavobacteria) were also abundant (up to 38.6% in May 2010), although their average contribution (6.1%) was clearly below Rhodobacterales and Flavobacteriales. While different taxa dominated the particle-associated and free-living communities (Supplementary Figure S1), the difference between the relative abundance of Flavobacteriales and Rhodobacterales was not significant.

Experimentally induced growth of microphytoplankton and bacterial communities in temperature-controlled experiments

To determine the temperature sensitivity of growth of microphytoplankton and microphytoplankton-associated bacteria, we carried out 3 temperature-controlled incubation experiments in January, February and March 2014 using natural seawater. *In situ* analysis of environmental variables showed similar temperatures in the three months (12.5 to 13.0°C), while the initial chlorophyll *a* concentration varied

substantially, from ca. $1 \mu\text{g L}^{-1}$ in January and February to $5.4 \mu\text{g L}^{-1}$ in March (Supplementary figure S2A), which was the highest value measured during 3 years sampling at the study site. Accordingly, the *in situ* abundance of microphytoplankton in March was one order of magnitude higher as compared to January and February.

We followed the growth of phytoplankton in the three experiments, until the abundance of phytoplankton cells stabilized in the incubations, and before they started declining (4, 5, 1 days in January, February and March, respectively). We observed a substantial increase in the growth of microphytoplankton in the ambient temperature treatment of January and February (from ca. 20 ± 7 cells mL^{-1} to 3150 and 2378 cells mL^{-1}). Although *in situ* microphytoplankton abundance in March was highest (144 ± 9 cells mL^{-1}) only a slight increase was observed after 12 h of incubation (to 165 ± 15 cells mL^{-1} , Figure 2), followed by a decrease in phytoplankton abundance, indicating that the microphytoplankton community sampled in March could have been already decaying.

The community composition of the microphytoplankton present in the three incubations was also different. *Thalassiosira* spp. dominated in January and February (90% and 70% of total microphytoplankton, respectively, Supplementary figure S3) and *Lauderia annulata* dominated in March (mean 59% of microphytoplankton cells larger than $5 \mu\text{m}$). In January and February, other groups such as *Asterionellopsis* spp. (4% and 10%, respectively) or *Cylindrotheca* spp. (2% and 9%, respectively) were also present, while *Guinardia* spp. or *Chaetoceros* spp. were present at relatively low abundance in March (9% and 1%). Dinoflagellates were only present during the experiment of March (22%, Supplementary figure S3).

The abundance of microphytoplankton-associated bacteria was also analyzed by epifluorescence microscopy. Only cells found in the vicinity of the phycosphere were quantified, including bacteria attached to the phytoplanktonic cells or to the exopolymers surrounding them (Supplementary figure S4). Bacterial abundance was analyzed at the initial time and when phytoplankton abundance was maximum during the incubations (132 h, 108 h and 12 h in January, February and March, respectively). We found that at the initial time of the experiments, phytoplankton-associated bacteria were only present in March while in January and February, due to the very low phytoplankton abundance at *in situ* conditions, no cells with associated bacteria were found using the epifluorescence microscope. The abundance of Flavobacteria and Rhodobacteraceae in March (as cells targeted by CFB315 and Ros537 probes) was 490 and 80 cells mL⁻¹, respectively (Figure 2), together representing the 72% of all particle-associated bacterial counts in that month and confirming their high abundance in this fraction at the single-cell level. The abundance of microphytoplankton-associated bacteria at the ambient temperature treatment increased during the three experiments (Figure 2). This increase was especially noticeable for Flavobacteria, which consistently exceeded the abundance of Rhodobacteraceae at peak microphytoplankton values for the three experiments (1.8-fold higher abundance of Flavobacteria in January and 3.7-fold higher in February and March, Figure 2). Total bacterial abundance also increased in the free-living fraction, particularly in January and February. The abundance of Flavobacteria between the initial and final times in the free-living fraction was 45 and 15 times higher in January and February, while the abundance in March increased 2-fold. In contrast, in a microphytoplankton-free control where microphytoplanktonic cells were removed by size fractionation, the bacterial abundance was one order of magnitude lower at the end of all the incubations in comparison to the cells incubated with microphytoplankton (Supplementary Figure S5). Such lower abundance in the absence of microphytoplankton was more noticeable for

Flavobacteria than for Rhodobacteraceae, where abundances were up to 28 and 10 times higher, respectively, in the treatments with phytoplankton in comparison to the micro-phytoplankton-free treatment.

The abundance of total microphytoplankton cells and the bacteria associated to them increased along the experimental gradient from low to warm treatments in the three experiments (ANOVA, p -value <0.001 , Figure 3). Bacterial abundances were 2-fold higher in the warm treatment in comparison to the cold one in January and March and 6-fold higher in February. Such increases in abundance were particularly noticeable for Flavobacteria in March, when after only 12 hours of incubation 2.2 times more cells were found in the warm treatment as compared with the cold one, while Rhodobacteraceae abundance was 1.6 times higher. On the contrary, we did not observe any warming effect in the free-living fraction for any of the two taxonomic groups during the 2 to 6-day incubations.

Effect of temperature on ^{13}C and ^{15}N incorporation by microphytoplankton and microphytoplankton-associated bacteria

We carried out stable isotope probing (SIP) incubations in the experiment of March, when the sampling was coincident with high *in situ* *Chl a* concentration (supplementary figure S2A) and the abundance of diatoms and bacteria associated to them was the maximum of the three experiments. We used ^{13}C -bicarbonate and ^{15}N -leucine incubations (after 12 hours of the initial time) to measure bulk uptake by elemental-analysis mass spectrometry (EA-MS), and cell-specific uptake by microphytoplankton and bacteria by nanoSIMS. We found a significant ^{13}C enrichment in bulk samples, with an average uptake of 0.86 ± 0.34 atom percent excess (APE), while no effect in ^{13}C APE was found for different temperature treatments. In the cell-specific analysis (Figure 4, Supplementary figure S6), we found that the major

fraction of ^{13}C -bicarbonate was preferentially assimilated by microphytoplankton cells, which showed uptake rates significantly different from 0 (Wilcoxon test, p -value <0.001 , with all 36 microphytoplankton cells analyzed actively incorporating ^{13}C -bicarbonate). On the contrary, ^{15}N -leucine was preferentially assimilated by microphytoplankton-associated heterotrophic bacteria, showing significant incorporation rates (Wilcoxon test, p -value $=0.015$, with 89.7% of 155 bacteria analyzed actively incorporating ^{15}N -leucine). Yet, we also found that ^{13}C incorporation by heterotrophic bacteria during the incubation, presumably transferred from the primary C-fixing organisms (i.e. microphytoplankton), was significant (Wilcoxon test, p -value <0.001). For the cell-specific ^{13}C and ^{15}N uptake measurements using nanoSIMS, bacteria were mostly analyzed in the surroundings of *Lauderia annulata* and *Chaetoceros spp.* diatoms, since the phycosphere of these species harbored the highest bacterial abundance while in other abundant groups such as dinoflagellates we virtually did not find any associated microbes. Yet, there was not any difference in ^{13}C uptake between bacteria associated with these two different diatom species (Supplementary figure S7). ^{15}N -leucine incorporation in Rhodobacteraceae was one order of magnitude higher than in Flavobacteria (Wilcoxon test, p -value $=0.008$, Figure 5B). By contrast, both Flavobacteria and Rhodobacteraceae showed almost identical cell-specific ^{13}C assimilation rates (0.42 ± 0.30 and $0.43 \pm 0.15 \text{ fg } \mu\text{m}^{-3} \text{ h}^{-1}$, respectively, Figure 5A).

We did not observe any clear effect of warming on microphytoplankton cell-specific ^{13}C incorporation (Supplementary Figure S8). However, we found that ^{13}C assimilation was significantly higher in the warm treatment for both bacterial groups (Wilcoxon test, p -value <0.001 , Figure 5A), while no differences in ^{15}N incorporation were found between the distinct temperature treatments for any group (Figure 5B). Due to the low abundance of Rhodobacteraceae in the cold treatment (below $200 \text{ cells mL}^{-1}$) and the

associated difficulty to detect individual cells for the nanoSIMS analysis, for this treatment, we only show assimilation rates for Flavobacteria in Figure 5.

Finally, we also calculated the contribution of each bacterial taxon to total C and N assimilation (Figure 5C, D). We found that ^{13}C total assimilation by Flavobacteria was 4-fold higher than Rhodobacteraceae at ambient temperature, while warming increased this difference up to 6-fold. Assimilation of ^{15}N -leucine by Rhodobacteraceae was 5 times higher than Flavobacteria at ambient temperature ($0.64 \pm 0.16 \text{ fg mL}^{-1} \text{ h}^{-1}$ and $0.11 \pm 0.004 \text{ fg mL}^{-1} \text{ h}^{-1}$ respectively), but due to the higher cell abundance of Flavobacteria in the warm treatment, no significant differences in total ^{15}N -leucine assimilation were found between the two groups.

Discussion

Marine Flavobacteria and Rhodobacteraceae are two of the most abundant and ecologically relevant bacterial groups in coastal waters (Kirchman, 2002; Buchan *et al.*, 2005), with a well-known opportunistic behavior during phytoplankton blooms (Riemann *et al.*, 2000; Tada *et al.*, 2011; Teeling *et al.*, 2012; Buchan *et al.*, 2014). Previous works have also shown that both bacterial taxa are major components of phytoplankton-associated communities (Grossart *et al.*, 2005; Rooney-Varga *et al.*, 2005) in agreement with our findings at the site of study, where they consistently exceeded the abundance of other typical model copiotrophs such as Alteromonadales. The higher, or at least faster response of Flavobacteria to microphytoplankton growth compared with Rhodobacteraceae also agrees with previous works showing differential responses of various taxonomic groups to increasing microphytoplankton biomass (Tada *et al.*, 2011), and suggests that Flavobacteria may have some ecological advantage over Rhodobacteraceae when competing for newly released DOC.

Moreover, although the initial abundances of both groups in the free-living fraction were similar, when analysing bacteria colonizing microphytoplankton senescent cells (Supplementary figure S9), we observed that Flavobacteria represented on average 80% of the microphytoplankton-associated bacteria, while Rhodobacteraceae contributed only 13%. Members of Flavobacteria have shown a high specialization in the degradation of polymeric compounds (Bauer *et al.*, 2006; Gomez-Pereira *et al.*, 2012) and a well described gliding motility, which may be key when colonizing microphytoplankton cells (Kiorboe *et al.*, 2002). The fact that OTUs affiliated to both groups were simultaneously found in the free-living and particle-associated fractions (Supplementary Figure S10) highlights this colonization ability, indicating that some free-living cells were able to switch between both lifestyles, a process already described to be important in the bacterial dispersion process in the deep sea (Mestre *et al.*, 2018).

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The main aim of our study was to analyse the impact of temperature on the uptake activity of these two copiotrophic taxa thriving in the phycosphere. According to the predictions of the latest IPCC report, climate change may result in surface ocean temperatures 1-3°C warmer by the end of this century (IPCC, Pörtner et al., in press). Thus, to test the temperature sensitivity of Flavobacteria and Rhodobacteraceae over a realistic range in this scenario, we used 3°C above ambient temperature as the warm treatment. Additionally, we also included a colder treatment (3°C below ambient temperature) in order to have a range as broad as possible without putting the communities into thermal stress (Huete-Stauffer *et al.*, 2016; Arandia-Gorostidi *et al.*, 2017a). Since short incubation experiments do not consider potential long-term effects such as bacterial acclimation in response to slow environmental change, caution should be taken when interpreting results obtained from this type of experiments. We deem our results as a way to test the temperature sensitivity of current microbial communities (Kirchman *et al.*, 2005; Huete-Stauffer *et al.*, 2015) and generate plausible hypotheses about the directions of change, which in some cases are confirmed by time-series records (e.g. see Morán *et al.* 2015 about the shrinkage of bacterial cell size with warming).

In the short-term incubations carried out in this study, we found that experimental warming noticeably increased the abundance of microphytoplankton-associated bacteria, while the effect on the free-living community was less pronounced. Such larger effect of associated bacteria may be explained by at least two mechanisms: (i) increased chemotaxis ability to newly fixed DOC by the microphytoplankton-associated bacteria (Smriga *et al.*, 2016), which would also promote a faster response to warming as the temperature-sensitivity of bacteria is strongly related to the availability of resources (López-Urrutia and Morán, 2007; Berggren *et al.*, 2010; Moran *et al.*, 2018), and/or (ii) a higher exudation of organic compounds by diatoms at higher temperatures, including transparent exopolymers particle (TEP) (Claquin

et al., 2008; Wohlers *et al.*, 2009; Seebah *et al.*, 2014), which would facilitate bacterial aggregation (Mari and Kiørboe, 1996; Gardes *et al.*, 2011). In any case, we found that the impact of experimental warming was distinct for the entire associated bacterial community in comparison to the two taxa analysed and that it also exerted a noticeable larger effect on Flavobacteria than on Rhodobacteraceae, indicating a differential response of both taxonomic groups in terms of abundance, as previously observed in free-living communities (von Scheibner *et al.*, 2014; Arandia-Gorostidi *et al.*, 2017a). Adaptations to detect and exploit DOC hotspots have been well described in both Rhodobacteraceae and Flavobacteria (Miller *et al.*, 2004; Buchan *et al.*, 2005; Fernández-Gómez *et al.*, 2013). Yet, the observed higher response of Flavobacteria to warming suggests that this group may be better adapted to use the newly released DOC within the phycosphere. Interestingly, in a previous study at the same site, we observed that the response to increasing temperature of free-living Rhodobacteraceae was stronger than that of Flavobacteria (Arandia-Gorostidi *et al.*, 2017a), suggesting that the response to warming may differ in each fraction. However, since in the latter study the effect of warming was exclusively tested on the growth rates (not on abundances) of the free-living community, the results of both studies are not fully comparable.

In terms of cell-specific uptake activity, the higher ^{13}C incorporation of microphytoplankton-associated bacteria with increasing temperatures agrees with a previous study (Arandia-Gorostidi *et al.*, 2017b), where only total bacterial communities were analyzed, disregarding potential differences based on taxonomic affiliation. Here, we found a similar response in ^{13}C assimilation rates for the two taxa targeted, Flavobacteria and Rhodobacteraceae, indicating a comparable ability to incorporate algal-derived DOC per cell volume, in contrast to the differences in absolute cell abundance found for both groups. These results indicate an imbalance between growth rates and uptake activity for each phylogenetic group. This difference could be explained by (i) a higher growth efficiency of some specific

clades within Flavobacteria, which with a similar DOC incorporation were able to generate more bacterial biomass, or (ii) a better adaptation of Flavobacteria at detecting new DOC hotspots. Regardless of the underlying mechanism, here we show that due to their higher abundance in the microphytoplankton-associated fraction, Flavobacteria were quantitatively the major consumers of algal-derived DOC and that this role was significantly enhanced by warming.

We cannot rule out that part of the bacterial incorporation of ^{13}C could be due to anaplerotic assimilation of CO_2 (Roslev *et al.*, 2004; Hesselsoe *et al.*, 2005; Alonso-Sáez *et al.*, 2010). However, such incorporation generally represents a minor fraction of total biomass production (between 1 and 6.5%, Roslev *et al.*, 2004; Hesselsoe *et al.*, 2005). Examples of exceptionally high anaplerotic C assimilation have been demonstrated for some gammaproteobacterial taxa under starvation conditions (Alonso-Sáez *et al.*, 2010), but it is unlikely that microphytoplankton-associated bacteria experience carbon starvation. Furthermore, the C uptake rates observed in this work are of the same order of magnitude of those reported by earlier studies analyzing algal-derived DOC incorporation by heterotrophic bacteria (Adam *et al.*, 2016; Arandia-Gorostidi *et al.*, 2017b). The fact that bacterial abundance was consistently higher in the presence of microphytoplankton (as compared to microphytoplankton-free controls), also supports the view that most of the ^{13}C assimilation originated from algal-derived DOC.

In the case of microphytoplankton, the lack of effect of temperature on bulk (measured by EA-MS) and cell-specific ^{13}C -bicarbonate incorporation was reflected in the low impact of warming on microphytoplankton abundance. We cannot rule out that factors not tested in this work such as nutrient availability or differences in the physiological state of the cells may impact this result. However, our results are consistent with a previous study where picophytoplankton dominated the algal communities at

the same coastal site (Arandia-Gorostidi *et al.*, 2017b) and other studies showing a lack of increase in particulate primary production with warming (Huete-Stauffer *et al.*, 2017). On the contrary, warming has been related with a relative increase in the release of newly fixed C as dissolved compounds (Morán *et al.*, 2006; Engel *et al.*, 2011; Yvon-Durocher *et al.*, 2012; Huete-Stauffer *et al.*, 2017), which may explain the clear effects of temperature on the C uptake activity of microphytoplankton-associated bacteria. Altogether, warming may result in important imbalances of the carbon budget in coastal waters, ultimately transferring more carbon to heterotrophic bacteria in relation to autotrophic cells.

In terms of ^{15}N -Leucine assimilation, the high contribution of Rhodobacteraceae agrees with the high amino-acid uptake activity observed for this group in the free-living fraction as compared to Flavobacteriaceae in marine samples (Alonso and Pernthaler, 2006; Alonso-Sáez and Gasol, 2007; Mayali *et al.*, 2014). Contrary to ^{13}C incorporation, cell-specific ^{15}N -leucine assimilation rates were not enhanced by warming. However, ^{15}N assimilation rates could indeed have been underestimated because other N pools such as ammonium, which represents an important N source for heterotrophic bacteria (Fouilland *et al.*, 2007), were not targeted. Besides, the low activity in leucine uptake by Flavobacteria may have hinder the impact of warming on the amino acid uptake by an important fraction of the microphytoplankton-associated bacterial community. Despite these limitations, the larger temperature effect observed in C relative to N uptake may indicate a differential impact of warming to the incorporation of distinct DOM sources, which would have important implications for element cycling in the ocean under warmer conditions.

In a previous work at the same site, a significant effect of warming on the exchange of C and N between picophytoplanktonic eukaryotes and attached heterotrophic bacteria was found (Arandia-Gorostidi *et al.*,

2017b). While picophytoplankton-heterotrophic bacteria interactions can be relevant particularly in the oligotrophic ocean (Agawin *et al.*, 2000; Jardillier *et al.*, 2010), diatom community have been described as the major annual source of organic C for coastal bacterial communities (Smith *et al.*, 1995; Klindworth *et al.*, 2014). In support of this idea, the bacterial C uptake rates reported here under high diatom abundance conditions were 2 to 3 times higher than in the earlier study, when phytoplankton was dominated by small eukaryotes (Arandia-Gorostidi *et al.*, 2017b), even if the ambient temperature was virtually the same (ca. 12°C). Although the difference may be related to the different methods used to calculate the assimilation rates, such results were consistent when we re-calculated the assimilation rates using the equations proposed by Popa *et al.* (2007), since in Arandia-Gorostidi *et al.* (2017b) we did not consider isotope correction. However, despite both studies were performed at the same site and under similar environmental conditions, algal-derived DOC uptake and its temperature dependence may also depend of factors no tested in any of both studies, such as inorganic nutrient availability. Thus, additional studies are required to fully understand the consequences of changes in autotrophic community composition on the C assimilation by their associated bacteria.

In summary, our results show that warming enhanced the use of algal-derived DOC by diatom-associated bacteria. We found that under experimental conditions, the abundance of two copiotrophic bacterial taxa showed a strong increase in response to temperature when associated with microphytoplankton cells, but not in the free-living fraction, while paradoxically most studies on microphytoplankton-bacteria interactions have traditionally focused on the free-living counterparts. Flavobacteria consistently overtook Rhodobacteraceae in terms of biomass production and algal-derived DOC consumption, indicating a dominant role of the former group as DOC-consumers, favored under warm conditions. However, some experimental limitations need to be considered, such as the difference of microphytoplankton abundance

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and their physiological state between the experiments, which may have an impact in the DOC pool available for bacteria and, hence, in their response to temperature (López-Urrutia and Morán, 2007). The difference in incubation time (due to the differences in the growth of microphytoplankton in the three experiments) may have also resulted in a longer acclimation period in the first two experiments, which may ultimately influence the future bacterial response to warming. Yet, we found that in the three experiments, the response of microbial abundance to temperature was consistent, indicating that regardless of *in situ* conditions, temperature may have an important impact on the abundance of microphytoplankton-associated bacteria. Although many uncertainties about the impact of climate change on the microphytoplankton-bacteria coupling remain, our work represents a step forward towards elucidating the interactions taking place in the complex environment of the phycosphere. Latest technological advancements such as nanoSIMS have allowed the precise determination of C and N incorporation within this niche, which may impact not only the individual cells but also the global cycling of these elements at ecosystem level (Seymour *et al.*, 2017). Here, we demonstrated that experimental changes in temperature impacted the composition of the microphytoplankton-associated bacteria and the DOM cycling within the phycosphere, which highlights the importance of this microenvironment to study the future consequences of global change for ocean carbon cycling.

Experimental procedures

DNA samples collection and processing

During a 3-year *in situ* analysis (December 2009-December 2012) surface seawater was collected monthly at the coastal station E2 (43.675°N, 5.578°W) located 13 km off Gijón/Xixón (Spain) in the southern Bay of Biscay. Five-liter Niskin bottles were filled with seawater and transported to the lab in the dark within the next 6 hours. Seawater was sequentially filtered through 3 µm pore-size polycarbonate filters (GTTP, 25mm, Millipore, Eschborn, Germany), to retain bacteria associated to particles and microphytoplankton cells, and through 0.2 µm pore size filters to collect the free-living fraction (Alonso-Sáez *et al.*, 2015). Filters were stored at -80°C before DNA extraction. Other parameters such as temperature, chlorophyll *a* and nitrate concentration were measured in the lab. Chlorophyll *a* concentration was determined by filtering 200 mL of seawater on 0.22 µm pore-size polycarbonate filters. Filters were frozen at -20°C and processed within 2 weeks as described in Calvo-Díaz and Morán (2006). Samples for nitrate concentration were determined with a Technicon autoanalyzer as indicated in Calvo-Díaz and Morán (2006).

From monthly samples collected along a 3-year period, a total of 11 samples corresponding to months with different concentrations of chlorophyll *a* were selected for DNA sequencing of both the microphytoplankton-associated and free-living fractions. DNA extraction and 16S rDNA genes amplification were carried out following Alonso-Sáez *et al.* (2015) using general bacterial primers 341F and 805R (Herlemann *et al.*, 2011). Triplicate PCR products were purified using a QIAquick purification Kit (Qiagen, Hilden, Germany) and amplicons were sequenced using a 454 FLX+ platform (Lifesequencing). Sequences were analyzed using the MOTHUR platform (Schloss *et al.*, 2009) as

described in Alonso-Sáez *et al.* (2015). Samples were rarefied to 2535 sequences per sample, obtaining a total of 55770 sequences for analysis. Sequences have been deposited in the European Nucleotide Archive project PRJEB6399 (accession numbers ERS1836182- ERS1836193 for the microphytoplankton-associated sequences deposited from this study).

Temperature-controlled experiments setup

Seawater was collected at the E2 station in January 26th, February 19th and March 22nd in 2014. The collected water was pre-filtered by 200 µm pore size filters to remove mesozooplankton and transported to the laboratory in 20 L Nalgene polycarbonate bottles (Nalgene, Rochester, NY, USA) within 6 hours from collection. At the arrival in the laboratory, seawater was distributed into polycarbonate bottles for incubation experiments (2 L per bottle). Additionally, we prepared a control with 2 L of seawater filtered by 0.8 µm pore size cartridge filters (Pall, Ilfracombe, United Kingdom) in order to remove microphytoplankton cells. This treatment is referred as microphytoplankton-free treatment. We incubated the water samples at 3 different temperatures: i) ambient temperature (12.5-13°C), ii) 3°C below (9.5-10 °C, cold treatment) and iii) 3°C above ambient temperature (15.5-16°C, warm treatment). Two duplicate bottles were incubated at each temperature and the additional microphytoplankton-free treatment bottle at ambient temperature. The temperature-controlled incubations were carried out at the natural photoperiod of each month, under saturating PAR irradiance (ca. 150 µmol photons m⁻² s⁻¹) and lasted up to 7 days.

Each day, 25 mL were removed from the bottles and fixed with 2% of Lugol solution for microphytoplankton taxonomy analysis (identifying cells with a diameter larger than 5 µm). Microphytoplankton cells were identified up to genus or species level in most of the cases, using an inverted microscope (Lund *et al.*, 1958) with a final magnification of x20. To determine total

microphytoplankton abundance, a minimum of 300 cells were count when possible, with an error of 11.5% for a 95% CI. When microphytoplankton abundance was very low (as at in situ conditions of January and February) a minimum of 100 cells were count (20% error).

At the initial time of the incubations and when microphytoplankton abundance reached maximum abundance, we filtered additional volumes onto 0.2 μm pore-size filters (5 mL) and 3 μm pore-size filters (100 mL) to quantify the abundance of Rhodobacteraceae and Flavobacteria in the microphytoplankton-associated and free-living fractions by Catalyzed Reporter Deposition Fluorescence *In situ* Hybridization (CARD-FISH). CARD-FISH analysis was performed following Pernthaler *et al.* (2002). Hybridizations were carried out using 9 μL the Horseradish peroxidase (HRP) labeled probes (50 $\text{ng } \mu\text{L}^{-1}$) Ros537 and CF319 (Manz *et al.*, 1996; Eilers *et al.*, 2001) diluted in 900 μL of hybridization buffer with 55% formamide. Fluorescently-labeled tyramide Alexa Fluor® 488 (1 mg mL^{-1} , 46°C, 30 min, ThermoFisher, Germany) was used for the amplification step. The hybridized cells were further stained with 4',6-diamidino-2-phenylindole (DAPI) at 1 $\mu\text{g mL}^{-1}$ and were quantified using an epifluorescence microscope Leica DM5500B. In order to provide absolute cell-abundance counts, cell abundance of Flavobacteria and Rhodobacteraceae was calculated based on the number of probe-specific hybridized cells per mL of filtered seawater, while abundance of the total bacterial community was based on the total DAPI counts. Counts of hybridized bacteria were calculated from 10 randomly chosen microscopic fields. A minimum of 200 DAPI stained cells were counted in the microphytoplankton-associated fraction and 1000 cells in the free-living fraction in order to have a representative estimation of total bacterial abundance.

In order to obtain updated values of coverage of the CF319 and Ros537 probes used in this study, we used the TestProbe 3.0 tool against the SILVA database. According to this analysis, the Ros537 probe almost exclusively targets the Rhodobacterales order (63% of known 16S sequences of this group). At the

family level, Ros537 exclusively targets Rhodobacteraceae family (which is the only marine representative of the Rhodobacterales order identified in our 16S data-set, with 63% coverage), including different genera such as *Roseobacter* (with 95% of coverage), *Sulfitobacter* (81% coverage), *Planktomarina* (98% coverage) or *Donghicola* (90% coverage). Yet, the main OTUs found in our study site were only affiliated to the *Roseobacter* and *Donghicola* genera (Supplementary figure S10). The probe CF319a targets mostly the Bacteroidia class (48% coverage), and two marine Flavobacteria orders: Flavobacteriales (89% coverage) and Sphingobacteriales (88% coverage). Although both orders have been detected at our study site, Flavobacteriales was considerably more abundant (22.3% in average) than Sphingobacteriales (6.1%).

Stable isotope probing and nanoSIMS analysis

For quantifying cell-specific C and N uptake for both microphytoplankton associated and free-living bacterial groups, water samples were collected in March (22nd, 2014) when the highest chlorophyll values were recorded. The freshly collected water samples were incubated for 12 hours at different temperatures (see temperature-controlled experiment above) followed by subsampling of 250 mL from each replicate of the different temperature treatments (+3°C, ambient temperature and -3°C) to duplicate Pyrex glass bottles for the stable isotope probing (SIP) incubations. To this volume, we added 300 $\mu\text{mol L}^{-1}$ of ^{13}C -bicarbonate (99 atom% ^{13}C , Cambridge Isotope Laboratories Inc., Cambridge, MA, USA), equivalent to 15% ^{13}C after dilution with ambient HCO_3^- , and 50 nmol L^{-1} of ^{15}N -leucine (Cambridge Isotopes). The SIP incubations lasted for 12 h at each of the corresponding temperatures.

For the nanoSIMS analysis, 100 mL from each replicate and each temperature treatment was fixed at the end of the SIP incubation with 4% formaldehyde for 3 hours at room temperature and filtered through

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gold-palladium coated polycarbonate filters (type GTTP; pore size, 3 μm ; diameter, 47 mm; Millipore, Eschborn, Germany). Filters were stored at -80°C until analysis. First, filters were hybridized using the HRP-labeled nucleotide probes as described above. The amplification step was done using either fluorine containing fluorescently labeled tyramides Oregon Green® 488 (ThermoFisher, Germany) or the Alexa Fluor® 488 (ThermoFisher, Germany) following the protocol by Musat *et al.* (2008). Hybridized cells were identified and mapped by using a laser microdissection system (LMD, Zeiss, Germany) equipped with an inverted epifluorescence microscope. Several areas containing target cells were mapped in the LMD system using an air-objective with 60X magnification as previously described by Alonso *et al.* (2012).

The LMD marked areas were further analyzed with a nanoSIMS 50L (Cameca, Gennevilliers, France) extracting negative secondary ions upon sample sputtering with primary Cs^+ ions. The 2 pA beam of 16 keV Cs^+ ions was focused into about 70 nm spot at the sample surface during the analysis. Samples were analyzed in a 512×512 px raster over $30 \times 30 \mu\text{m}^2$ areas with 5 msec dwell time per pixel. Before the analysis, the sample surface was treated with 10 nA of low-energy (50 eV) Cs^+ beam over $80 \times 80 \mu\text{m}^2$ area for 10 minutes. The secondary ions species were analyzed for their mass to charge ratio (m/z) and detected using the seven available collectors: $^{12}\text{C}^-$, $^{13}\text{C}^-$, $^{19}\text{F}^-$, $^{12}\text{C}^{14}\text{N}^-$, $^{12}\text{C}^{15}\text{N}^-$ and $^{13}\text{C}^{14}\text{N}^-$, $^{31}\text{P}^-$, $^{32}\text{S}^-$. Two secondary ion species ($^{12}\text{C}^{15}\text{N}^-$ and $^{13}\text{C}^{14}\text{N}^-$) were detected by switching the deflector voltage in combined analysis mode. The mass resolving power (MRP) was checked to be between 7000 and 10000 with the exit slit width of 70 μm , 20 μm entrance slit width, 200 μm aperture slit and with the energy slit cutting about 30% of secondary ions in high-energy tail of their energy distribution. Under these analysis conditions, we observed that cells got sputtered completely within 10 scans. Scans 3-8 were considered for the analysis employing LANS software (Polerecky *et al.*, 2012) allowing for the accumulation of the

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scans with lateral drift correction and quantitative analysis of isotopic ratios ($^{13}\text{C}/^{12}\text{C}$, $^{13}\text{C}^{14}\text{N}/^{12}\text{C}^{14}\text{N}$ and $^{12}\text{C}^{15}\text{N}/^{12}\text{C}^{14}\text{N}$). Identification of bacterial cells in the nanoSIMS images was performed by using $^{12}\text{C}^{14}\text{N}$ and ^{31}P channels, which allowed to differentiate the cells from the background signal, while identification of hybridized cells was done by comparison with the epifluorescence images. To determine cell-specific isotope ratios, regions of interest (ROIs) were drawn inside the cells, avoiding the cell-surrounding areas. To quantify the natural abundances of $^{13}\text{C}^{14}\text{N}/^{12}\text{C}^{14}\text{N}$ and $^{12}\text{C}^{15}\text{N}/^{12}\text{C}^{14}\text{N}$ in our samples, we analyzed the isotopic ratios on non-labeled samples collected just before the SIP incubation of March. Changes in the isotope composition during stable isotope labeling measured by nanoSIMS were generally considered quantitatively to calculate assimilation rates of single cells, following the calculations provided in Stryhanyuk *et al.* (2018). Assimilation rates were calculated as the fraction (K_A) of carbon or nitrogen assimilated into a cell during its incubation in isotope-labeled growth substrate:

$$K_A = \frac{R_f - R_i}{R_i + 1} \times \frac{R_{gs} + 1}{R_{gs} - R_f}$$

Where R_{gs} is the isotope ratio of the growth substrate during incubation, R_i is the initial isotope ratio before incubation, and R_f is the isotope ratio after incubation. With the isotope ratios (R) defined as heavy-to-light ratio for carbon isotopes, $R = ^{13}\text{C}/^{12}\text{C}$, the fractions of heavy and light isotopes (D_{heavy} and D_{light}) were expressed as follows:

$$D_{heavy} \equiv D = \frac{^{13}\text{C}}{^{13}\text{C} + ^{12}\text{C}} = \frac{R}{R + 1} \text{ and } D_{light} = \frac{^{12}\text{C}}{^{13}\text{C} + ^{12}\text{C}} = \frac{1}{R + 1}.$$

For performing the calculations, we considered a value of 64.6% of C dilution due to the CARD-FISH treatment (Musat *et al.*, 2014). In the case of nitrogen, since N loss during the treatment could not be

modeled, we decided to omit any N correction. Besides isotope correction, we also considered the following parameters during the calculations: the fraction of the isotopes in chemicals used for the treatment and the isotope ratio after the CARD-FISH. Cell biovolume (in μm^3) was estimated from the bacterial diameter measured from the nanoSIMS images and assuming a spherical form. The partial density of carbon and nitrogen for bacteria ($111.9 \pm 45.5 \text{ fg C } \mu\text{m}^{-3}$ and $21.5 \pm 5.6 \text{ fg N } \mu\text{m}^{-3}$, Fukuda *et al.*, 1998) and for microphytoplankton ($140.0 \text{ fg C } \mu\text{m}^{-3}$ and $10.0 \text{ fg N } \mu\text{m}^{-3}$, Putt and Stoecker, 1989; Montagnes and Franklin, 2001) were used to obtain the final C and N assimilation rates per unit (μm^3) of cell volume. We also calculated C and N assimilation rates per cell (Supplementary Table S1). In addition, C and N assimilation rates were also determined according to the calculations described by Popa *et al.* (2007), without considering any isotope correction, in order to compare our results with ones obtained in Arandia-Gorostidi *et al.* (2017b) (Supplementary Table S2).

To determine the assimilation of carbon and nitrogen by Flavobacteria and Rhodobacteraceae cells, we calculated total C and N uptake in fg mL^{-1} by multiplying the assimilation rate per volume by the total biovolume of each taxonomic group.

Analysis of bulk C assimilation by an elemental analyser

In addition to cell-specific analysis, at the end of the SIP incubation, 50 mL of each bottle were filtered through pre-combusted GF/F filters and stored at -80°C to determine $^{13}\text{C}/^{12}\text{C}$ isotope ratio by an elemental-analysis mass spectrometry (EA-MS). Several round slices of each GF/F filters (with a diameter of 0.5 cm) were packed in small cups and dried overnight at 55°C . Dried filter packages were loaded onto an autosampler and assimilation of bulk C was analysed by a Euro EA3000 elemental analyzer (HEKAtech, Germany) equipped with a commercially available combustion reactor filled with

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tungsten oxide and silver cobalt oxide (HE46820995, HEKAtech, Germany) and connected to a gas analyser (GAM 445, Balzers AG, Balzers, Liechtenstein) using caffeine as standard for isotope correction. Bulk C assimilation was determined as atom percent enrichment (APE) over the natural abundance, as described in the following equation:

$$APE = (D_f - D_i) * 100$$

Where $D_{f/i}$ is the fraction of ^{13}C at the final and initial sampling time, respectively

$$D_{f/i} = \frac{^{13}\text{C}}{^{12}\text{C} + ^{13}\text{C}}$$

Conflict of interest: The authors declare that they have no conflict of interest.

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Figure legends

Figure 1. (A) Bacterial community composition in the particle-associated fraction ($>3\ \mu\text{m}$ pore-size filters) expressed as percentage of total reads. Monthly *in situ* values of Chl *a* concentration and temperature are showed with a green dashed line and a black line respectively. (B) Bacterial community composition as percentage of total reads for the free-living fraction (fraction collected between $3\ \mu\text{m}$ and $0.22\ \mu\text{m}$ pore-size filters).

Figure 2. Abundance of microphytoplankton cells (lines), total heterotrophic bacteria (quantified by epifluorescence microscope using DAPI staining) and the two taxonomic groups (quantified by CARD-FISH) in the free-living (A, B, C) and microphytoplankton-associated fractions (D, E, F) during the experimental incubations of January, February and March 2014 at ambient temperature. Note that the scale of the y axis differs in the incubations of March (C,F).

Figure 3. Microphytoplankton (left panel) and total heterotrophic bacteria (middle panel) abundances at the end of the experiments of January, February and March. The right panel shows the abundance of Flavobacteria and Rhodobacteraceae in the free-living and microphytoplankton-associated fractions for the three temperature treatments. Incubation times differ for each incubation (132 h for January, 108 h for February and 12 h for March). Amb. T refers to ambient temperature, Cold to 3°C below ambient temperature and Warm to 3°C above ambient temperature. Error bars indicate standard deviation.

Figure 4. Images showing Flavobacteria (upper panels) and Rhodobacteraceae (lower panels) cells under epifluorescence microscopy (A, B) and their respective nanoSIMS image of atomic percent enrichment (APE) of ^{13}C (C, D) and ^{15}N (E, F), indicating cell-specific isotopic uptake over the natural abundance of each isotope. White scale-bars equal $3\ \mu\text{m}$.

Figure 5. Boxplots of carbon (A) and nitrogen (B) assimilation rates in the microphytoplankton-associated fraction of Flavobacteria and Rhodobacteraceae at each temperature treatment and total carbon (C) and nitrogen (D) assimilation. Amb. T refers to ambient temperature, Cold to 3°C below ambient temperature and Warm to 3°C above ambient temperature. Error bars indicate standard deviation.

Figure 1

Figure 2

Figure 3

Figure 4

Figure 5