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Sunlight modulates the relative importance of heterotrophic bacteria and picophytoplankton in DMSP-sulphur uptake

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There is a large body of evidence supporting a major role of heterotrophic bacteria in dimethylsulphoniopropionate (DMSP) utilisation as a source of reduced sulphur. However, a role for phototrophic microorganisms has been only recently described and little is known about their contribution to DMSP consumption and the potential modulating effects of sunlight. In an attempt to ascertain the relative quantitative roles of heterotrophic bacteria and picophytoplankton in the osmoheterotrophic uptake of DMSP-sulphur upon exposure to natural sunlight conditions, we incubated northwestern Mediterranean waters under various optical filters and used an array of bulk and single-cell activity methods to trace the fate of added ^{35}S-DMSP. Flow cytometry cell sorting confirmed dark ^{35}S uptake by *Prochlorococcus*, *Synechococcus* and heterotrophic bacteria, the latter being the most efficient in terms of uptake on a cell volume basis. Under exposure to full sunlight, however, the relative contribution of *Synechococcus* was significantly enhanced, mainly because of the inhibition of heterotrophic bacteria. Microautoradiography showed a strong increase in the proportion of *Synechococcus* cells actively taking up ^{35}S-DMSP, which, after full sunlight exposure, made up to 15% of total active *Bacteria*. Parallel incubations with ^{3}H-leucine generally showed no clear responses to light. Finally, size-fractionated assimilation experiments showed greater relative cyanobacterial assimilation during the day than at night compared with that of heterotrophic bacteria. Our results show for the first time a major influence of sunlight in regulating the competition among autotrophic and heterotrophic picoplankton for DMSP uptake at both the daily and seasonal time scales.

Introduction

Dimethylsulphide (DMS) is a biogenic volatile compound that is universally present in sea water (Lovelock *et al.*, 1972; Kettle *et al.*, 1999) and represents the major natural source of sulphur to the global troposphere (Bates *et al.*, 1992; Andreae and Crutzen, 1997). The biogeochemical significance of DMS was first suggested when its emissions were found to be a key step in the global sulphur cycle (Lovelock *et al.*, 1972), and research was further encouraged when marine plankton was proposed to play a significant role in climate regulation through the effects of DMS emissions on cloud formation (Charlson *et al.*, 1987).

The biochemical precursor of DMS is dimethylsulphoniopropionate (DMSP), an osmolyte produced by many phytoplankton taxa and released into the dissolved organic matter pool through grazing, viral lysis, algal autolysis or exudation (Stefels, 2000; Simó, 2001). Once in sea water, DMSP may become available as a significant source of carbon and sulphur for other planktonic organisms. Released dissolved DMSP also acts as a direct or indirect (by transformation into DMS) chemical signal for plankton microbes (Seymour *et al.*, 2010), marine invertebrates, fish, birds and mammals (Van Alstyne *et al.*, 2001; Cunningham *et al.*, 2008; DeBose *et al.*, 2008; Nevitt, 2008).

Among the marine organisms directly utilising DMSP, heterotrophic bacteria have been the most extensively studied (Kiene *et al.*, 1999; Simó *et al.*, 2002; Howard *et al.*, 2006; Vila-Costa *et al.*, 2007)

and their capacity to assimilate sulphur from DMSP appears to be widespread among different taxonomic groups (see, for example, González et al., 1999; Malmstrom et al., 2004a; Vila-Costa et al., 2008b; Ruiz-González et al., 2011). Actually, bacterial uptake of dissolved DMSP and partial assimilation of its sulphur is thought to be the dominant mechanism for DMSP degradation in the pelagic ocean (Kiene et al., 2000; Zubkov et al., 2002). Other transformation processes include cleavage into DMS and carbon products mediated by the DMSP producers themselves and bacteria (Stefels, 2000; Sunda et al., 2002), accumulation or assimilation by zooplankton grazing on phytoplankton (Dacey and Wakeham, 1986; Archer et al., 2003; Tang and Simó, 2003; Saló et al., 2009) and direct uptake and assimilation by eukaryotic phytoplankton (Vila-Costa et al., 2006; Ruiz-González et al., submitted) and cyanobacteria (Malmstrom et al., 2005; Vila-Costa et al., 2006).

Among the aforementioned transformation processes, little is known about the quantitative role of non-DMSP-producing photosynthetic organisms. Because of their autotrophic lifestyle, the contribution of phytoplankton to the turnover of DMSP was expected to be minimal, but Synechococcus were reported to be major consumers of DMSP and methanethiol with contributions comparable to other bacterial groups (Malmstrom et al., 2005). Similarly, Prochlorococcus, diatoms and photosynthetic picoeukaryotes are also able to take up and assimilate a remarkable fraction of DMSP (Vila-Costa et al., 2006), suggesting that in the appropriate circumstances, they could compete with heterotrophic bacteria for this substrate.

An important implication of DMSP utilisation by phytoplankton is the possible effect of light on DMSP consumption processes. As algae are affected by variations in energy supply because of changes in the quantity and quality of light available, processes such as DMSP-sulphur assimilation may also be coupled to this periodicity. Only in two studies has light been shown to affect the assimilation of DMSP-sulphur by phototrophs, providing variable degrees of light-driven stimulation of the uptake (Malmstrom et al., 2005; Vila-Costa et al., 2006). Therefore, light may be playing an important role in regulating the relative contributions of heterotrophic and phototrophic uptake to total DMSP consumption. So far, however, experiments had been conducted in the absence of ultraviolet radiation (UVR, 280–400 nm), and hence it is likely that the contribution of phytoplankton relative to that of heterotrophic bacterioplankton (generally less UVR protected) had been underestimated. UVR has recently been found to be a significant factor modifying the fate of DMSP through either inhibition of microbial consumption (Slezak et al., 2001, 2007) or stimulation of its production, release (Sunda et al., 2002; Slezak and Herndl, 2003; Archer et al., 2010) or uptake by autotrophs (Malmstrom

et al., 2005; Vila-Costa et al., 2006; Ruiz-González et al., submitted). Incubations under natural sunlight conditions are necessary to properly assess the shares of the different components of the microbial food web in the use of this widespread substrate.

Our aim in this study was to assess for the first time the role of sunlight, including UVR, in DMSP uptake by picoplankton through experiments conducted with plankton communities sampled at the Blanes Bay Microbial Observatory and adjacent offshore waters of the northwestern Mediterranean. We used an array of multiple bulk and specific activity methods including flow cytometry cell sorting, size-fractionated assimilation and microautoradiography combined with RNA probing after samples were exposed to different light spectrum conditions. As sunlight has the potential to trigger the autotrophic activity and simultaneously inhibit heterotrophic bacterial activity, our hypothesis was that exposure to enhanced natural solar radiation would favour picophytoplankton in their competition for DMSP uptake against heterotrophic bacteria.

Materials and methods

Study area and sample collection
Water samples were collected either from a shallow (20 m depth) coastal station (the Blanes Bay Microbial Observatory (BBMO)) located 800 m offshore or during a cruise aboard the RV 'García del Cid' between 18 and 26 September 2007 at two stations, one located on the continental shelf in the vicinity of the BBMO (station C) and another one offshore over a 2000 m deep-water column between the BBMO and Mallorca (northwestern Mediterranean, Table 1). Surface samples (0.5 m) from the BBMO were collected with a Niskin Go-flow bottle (5 l), prefiltered through a 200 μm-mesh size net, and transported under dim light to the lab for the three experiments carried out on 5 August 2003 (experiment (exp.) 2), 9 July 2008 and 30 September 2008 (exps. 3 and 4, respectively). Water samples during the cruise (4 or 48 m depth) were taken with a 12-Niskin-bottle rosette attached to the CTD (conductivity, temperature, depth recorder; exps. 1, 5 and 6).

Experimental design
Different types of incubations were carried out with added trace ^{35}S-DMSP, kindly donated by RP Kiene (University of South Alabama, Dauphin Island Sea Lab, Dauphin Island, AL, USA). At several occasions, parallel incubations with ^{3}H-leucine (Amersham, Piscataway, NJ, USA; 161 Ci mmol^{-1}) were also done for comparative purposes as it is widely used as a measurement of bacterial heterotrophic production (Kirchman et al., 1985). Only in exps. 3 and 4, the photosynthetic active radiation (PAR) and UVR doses were monitored during incubations with a PUV-2500 radiometer

Table 1 Sampling locations, methodology and date of each experiment

Station	Exp.	Techniques	Date (day/month /year)	Longitude	Latitude	Depth (m)	HB (10^5 per ml)	Syn (10^4 per ml)	Prochl (10^4 per ml)	UVB ($kJ\,m^{-2}$)
D	1	Sorting (dark)	24/09/07	2° 51.06′E	40° 39.60′N	48	6.9	2.1	17.8	—
BBMO	2	Sorting (light)	05/08/03	2° 48.03′E	41° 39.90′N	0.5	8.5	7.2	0.2	21.0
BBMO	3	MAR-CARD-FISH	09/07/08	2° 48.03′E	41° 39.90′N	0.5	8.5	2.3	—	22.0
BBMO	4	MAR-CARD-FISH	30/09/08	2° 48.03′E	41° 39.90′N	0.5	7.2	4.7	3	11.3
C	5	Diel cycle (Assim)	18–19/09/07	2° 47.58′E	41° 40.08′N	4	4.8	0.5	0.4	—
D	6	Diel cycle (Assim)	23–24/09/07	2° 51.06′E	40° 39.60′N	4	6.0	1.5	1.3	—

Abbreviations: Assim, size-fractionated ^{35}S assimilation from added ^{35}S-DMSP; BBMO, Blanes Bay Microbial Observatory; MAR-CARD-FISH, microautoradiography combined with catalysed reporter deposition-fluorescence *in situ* hybridisation. The abundances of heterotrophic bacteria (HB), *Synechococcus* (Syn) and *Prochlorococcus* (Prochl) are given for the initial water sample. Integrated ultraviolet B (UVB) radiation during experiments were determined with a PUV radiometer (exps. 3 and 4) or estimated from the irradiance values collected at the Malgrat de Mar meteorological station (exp. 1, see Materials and methods).

(Biospherical Instruments, San Diego, CA, USA). In exp. 1, UVB levels were estimated from the irradiance values obtained from a meteorological station located 5 km southwest from the BBMO sampling station (Malgrat de Mar, Catalan Meteorological Service, SCM). The UVB doses are shown in Table 1.

Exp. 1: dark incubation for cell sorting. Water was collected around midnight from 48 m depth at the offshore station D. This was the depth where the cell abundance of *Prochlorococcus* (1.8×10^5 cells per ml; Table 1) was large enough to allow sorting. A single 50 ml sample was amended with ^{35}S-DMSP ($815\,Ci\,mmol^{-1}$, 0.03 nM final concentration (conc.)) and incubated in the dark for 6 h at *in situ* temperature (ca. 17 °C). After exposure, 5 ml subsamples were fixed with 1% paraformaldehyde (PFA) + 0.05% glutaraldehyde (final conc.), flash-frozen in liquid nitrogen and stored at −80 °C. Killed controls were prepared by addition of the fixative 30 min before the addition of the radioisotope, and were simultaneously incubated with the live samples.

Exps. 2–4: light manipulation experiments. Spherical quartz glass bottles of 50 or 100 ml were used to incubate surface water samples (0.5 m depth) collected in the Blanes Bay for 4 h under different light conditions. In exps. 3 and 4, samples were amended with trace ^{35}S-DMSP (453 and $23\,Ci\,mmol^{-1}$, 0.08 and 1 nM final conc., respectively) or ^{3}H-leucine ($161\,Ci\,mmol^{-1}$, 0.5 nM final conc.) immediately before exposure, whereas in exp. 2, they were added after the light incubations. In this experiment (no. 2), the conditions were: (1) full sunlight spectrum; (2) full spectrum minus UVB; (3) full spectrum minus the whole of UVR, that is, PAR only; and (4) darkness (wrapped with aluminium foil inside a black plastic bag). In exps. 3 and 4, the conditions were the same except for that treatment (2) was excluded. For the removal of UVB radiation (that is, PAR + UVA treatment), one layer of the plastic foil Mylar-D (150 μm thickness, 50% transmission at 325 nm) was used. For PAR-only treatments, bottles from exp. 2 were wrapped with one

layer of a vinyl chloride foil (50% transmittance at 405 nm; CI Kasei Co., Tokyo, Japan), and bottles from exps. 3 and 4 were covered with two layers of Ultraphan URUV (0.1 mm thickness, 50% transmittance at 380 nm; Digefra, Munich, Germany). All bottles were incubated 4 cm below the water surface inside a black tank with circulating sea water to maintain *in situ* temperature. After sunlight exposure, 50–100 ml subsamples from exp. 2 were incubated for 4 h with trace additions of ^{35}S-DMSP (0.1 nM, specific activity $130–350\,Ci\,mmol^{-1}$) or ^{3}H-leucine ($161\,Ci\,mmol^{-1}$, 0.5 nM final conc.) in the dark in acid-cleaned glass serum vials. Samples were fixed with 1% PFA + 0.05% glutaraldehyde (final conc.), flash-frozen in liquid nitrogen and stored at −80 °C. Live samples from exps. 3 and 4 were fixed overnight with PFA (1% final conc.) at 4 °C in the dark. Aliquots of 10–15 ml were filtered through 0.22 μm polycarbonate filters (GTTP, Millipore Iberica, Madrid, Spain), rinsed with milli-Q water, air dried and stored at −20 °C until processing.

Exps. 5 and 6: day/night cycles. During the September 2007 cruise, two diel cycle studies of the assimilation of ^{35}S-DMSP by different size fractions were conducted. For that purpose, 50 ml surface water samples (4 m depth) were collected every 4 h during two 24 h periods and trace concentrations of ^{35}S-DMSP ($815\,Ci\,mmol^{-1}$, 0.8 pM final conc.) were added. Samples were then incubated in 50 ml quartz flasks with the radioisotope for 6 h at *in situ* light and temperature conditions inside a black tank with circulating sea water. Killed controls were prepared in 30 ml Teflon flasks by adding PFA (1% final conc.) before the addition of the radioisotope. After exposure, the incorporation of substrate was stopped by fixing samples overnight with PFA (1% final conc.) at 4 °C in the dark.

Flow cytometry cell sorting
In exps. 1 and 2, different populations were identified and sorted using a FACSCalibur flow cytometer-cell sorter (Becton Dickinson, Franklin Lakes, NJ, USA). Sorted cells were collected onto

0.2 μm nylon filters and assayed by liquid scintillation counting. We used the 'single cell sort' mode of the instrument and sorted between 100 000 and 400 000 heterotrophic bacteria, between 30 000 and 130 000 *Synechococcus* and between 30 000 and 90 000 *Prochlorococcus* (the latter in exp. 1 only). Assimilation of ^{35}S or ^3H-leucine in killed samples was 2–3% of the value in live samples.

Microautoradiography combined with catalysed reporter deposition-fluorescence in situ *hybridisation (MAR-CARD-FISH)*
Filters with retained cells from exps. 3 and 4 were hybridised following the CARD-FISH protocol (Pernthaler *et al.*, 2002). Two horseradish peroxidase probes were used to specifically identify most *Eubacteria* (Eub338-II-III, Amann *et al.*, 1990; Daims *et al.*, 1999) and the cyanobacterial genus *Synechococcus* (Syn405, West *et al.*, 2001). Hybridisations were done on sections of the filters at 35 °C overnight. Smaller pieces from each hybridised section were cut and stained with 4,6-diamidino-2-phenylindole (DAPI, 1 μg ml^{-1}) to estimate the relative abundance of each group before applying the microautoradiography, for which 500–1900 DAPI-stained cells were manually counted within 10 to 25 fields. For microautoradiography, we followed the protocol described in Vila-Costa *et al.* (2007) and filters were exposed inside black boxes at 4 °C until development (3 days for ^3H-leucine and ^{35}S-DMSP in exp. 3, and 5 days for ^3H-leucine and 7 days for ^{35}S-DMSP in exp. 4). Slides were developed as previously reported (Vila-Costa *et al.*, 2007), stained with DAPI (1 μg ml^{-1}) and 500–700 hybridised cells were counted manually under an epifluorescence microscope.

Isotope assimilation during incubations
Triplicate aliquots of samples from exps. 5 and 6 (previously prefiltered through 3 μm pore-sized filters to exclude larger organisms; SSWP, Millipore) were sequentially filtered through 0.65 and 0.2 μm pore-sized filters (DAWP and GNWP, respectively; Millipore) and rinsed with 0.2 μm filtered sea water. Macromolecules were precipitated by treating filters with 5 ml of cold trichloroacetic acid 5% for 5 min. The filters were then rinsed with milli-Q water, placed into 5 ml of scintillation cocktail (Optiphase HiSafe 2, Perkin Elmer, Madrid, Spain) and counted with a Beckman scintillation counter (Beckman Coulter Inc., Barcelona, Spain). Incorporation of ^{35}S-DMSP and ^3H-leucine in PFA-killed controls was always <1.5% of that in live samples.

Results

Flow cytometry cell sorting of samples exposed to fractional sunlight (exps. 1 and 2)
Flow cytometry cell sorting of samples amended with ^{35}S-DMSP was used to investigate the relative role of heterotrophic bacteria and cyanobacteria in the uptake of DMSP-sulphur. Exps. 1 and 2 were conducted with water from the offshore station D and the coastal BBMO site, respectively. Among cyanobacteria, *Synechococcus* occurred in high numbers at both stations, but *Prochlorococcus* only occurred in sufficient numbers at a depth of ca. 50 m in station D, exp. 1 (Table 1). Cell sorting in this experiment showed that heterotrophic bacteria, *Prochlorococcus* and *Synechococcus* had the capability of assimilating ^{35}S from DMSP (Figure 1a). On a per-cell basis, the most important DMSP-sulphur assimilators were *Synechococcus* (Figure 1b), which showed 8 times more disintegrations per minute (d.p.m.) per cell than heterotrophic bacteria. However, when these values were normalised to total cell volume (assuming a volume ratio HB/Prochl/Syn = 1 : 5 : 10), heterotrophic bacteria were the most relevant consumers of ^{35}S-DMSP followed by *Synechococcus* and *Prochlorococcus* (Figure 1c).

Incubation of samples from the Blanes Bay under increasing short-wave radiation doses (exp. 2, Figure 2) resulted in a significant reduction of the contribution of heterotrophic bacteria to total ^{35}S-DMSP uptake (Tukey's test, $P<0.05$), from 84% in the dark to 17% after full sunlight exposure. Conversely, *Synechococcus* did not seem to be negatively affected by full sunlight exposure, and even a slight photostimulation of their uptake was apparent, yet not significant at $P<0.05$ (Figure 2a).
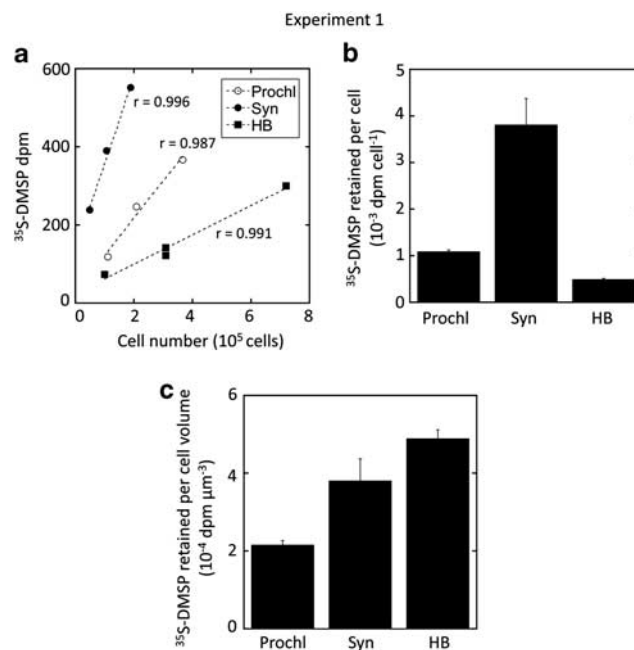

Figure 1 Experiment 1. (**a**) ^{35}S-DMSP retained in cells (d.p.m.) as a function of the number of sorted cells: *Prochlorococcus* (Prochl), *Synechococcus* (Syn) and heterotrophic bacteria (HB). (**b**) Average ^{35}S-DMSP retained per cell (d.p.m. per cell). (**c**) Same as (**b**) but scaled to cell volume (d.p.m. per μm³), assuming that *Prochlorococcus* and *Synechococcus* are 5 and 10 times larger, respectively, than heterotrophic bacteria. Shown are means ± standard errors.
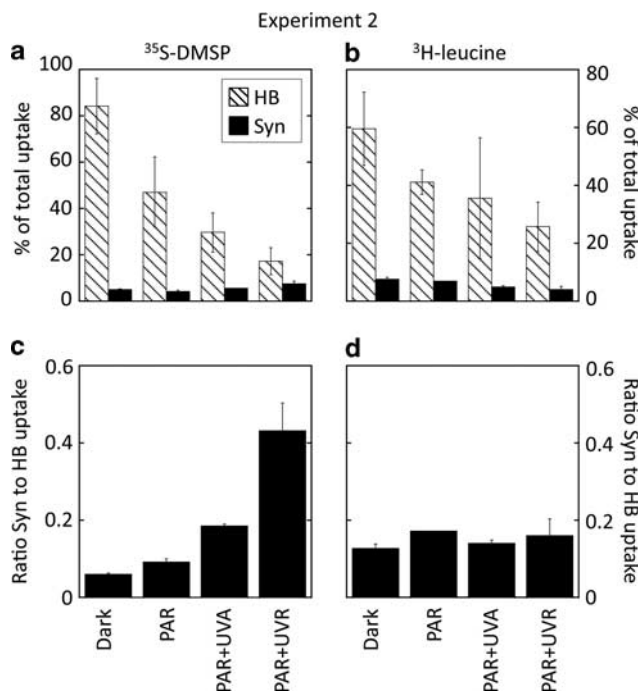
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Figure 2 Experiment 2. (Top panels) Contribution of sorted heterotrophic bacteria and *Synechococcus* cells to total uptake of (**a**) ³⁵S-DMSP or (**b**) ³H-leucine under different light conditions. Shown are means ± standard errors. (Bottom panels) Ratio between the contribution of *Synechococcus* (Syn) and heterotrophic bacteria (HB) to total uptake of (**c**) ³⁵S-DMSP or (**d**) ³H-leucine.

As a result, the contribution of *Synechococcus* relative to that of heterotrophic bacteria increased after full sunlight (PAR + UVR) exposure up to sevenfold (Figure 2c), accounting for 40% of the heterotrophic bacterial ³⁵S-DMSP uptake. The fraction of assimilated ³⁵S-DMSP that was not retained by *Synechococcus* or by heterotrophic bacteria showed an increased contribution to total uptake toward the full spectrum (reaching up to 75% under PAR + UVR; Figure 2a); yet, we did not specifically analysed which organisms were responsible for this uptake.

When we further assessed the role of light on the efficiency of ³⁵S-DMSP uptake per cell volume, the differences were even greater (Figure 3). Exposure to full sunlight caused a drastic decrease in the d.p.m. per μm³ in heterotrophic bacteria to the extent that they equalled those of *Synechococcus* (Figure 3a). Therefore, under full sunlight, *Synechococcus* cells seemed to be, on average, as efficient in taking up ³⁵S-DMSP as the average heterotrophic bacterium, as shown by the high ratio between the uptake per unit of cell volume of *Synechococcus* and heterotrophic bacteria (Figure 3b).

With ³H-leucine as the added substrate, a decreasing trend in the uptake by both *Synechococcus* and heterotrophic bacteria in all light treatments was observed, although for the latter it was not significant at $P < 0.05$ (Figure 2b). The resulting pattern of the relative contributions was essentially invariable with the sunlight spectrum (Figure 2d).
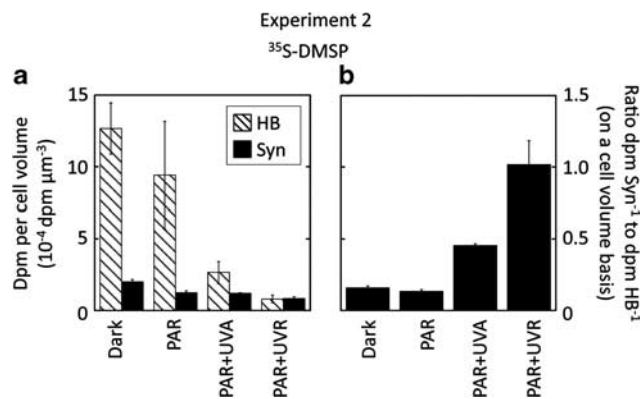

Figure 3 Experiment 2. (**a**) Comparison of ³⁵S-DMSP retained per cell volume (d.p.m. per μm³) between heterotrophic bacteria and *Synechococcus* under different light conditions. Values are means ± standard errors. (**b**) Ratio between the ³⁵S-DMSP uptakes per cell volume of *Synechococcus* (Syn) and heterotrophic bacteria (HB) under different light conditions.

Single-cell assessment of ³⁵S-DMSP and ³H-leucine uptake by MAR-CARD-FISH

The specific differences in the sensitivity to light of heterotrophic bacteria and *Synechococcus* were further assessed by applying the MAR-CARD-FISH technique to samples from the BBMO (exps. 3 and 4). Hybridisation with specific probes showed that *Bacteria* accounted for 88% and 67% of total DAPI counts in exps. 3 (summer) and 4 (autumn), respectively, whereas only 1% and 4% were *Synechococcus* (Ruiz-González, unpublished). When samples of the exp. 3 amended with ³⁵S-DMSP were exposed to the different light conditions (Figure 4a), the number of *Bacteria* active at ³⁵S uptake significantly decreased upon full sunlight exposure compared with both dark and PAR treatments (Tukey's test, $P < 0.05$), whereas the number of active *Synechococcus* strongly increased from ca. 10% to up to 80%, reaching 15% of total active *Bacteria* after full sunlight exposure (Figure 5a) when multiplied by their abundances. In exp. 4, conversely, both *Bacteria* and *Synechococcus* were stimulated in their ³⁵S uptake upon exposure to light (Figure 4b); however, the increase of *Synechococcus* (by eightfold) was much greater than that of bacteria, which resulted in a light-driven increased contribution of the former to the total numbers of cells active in ³⁵S uptake (Figure 5b).

The number of *Bacteria* active in the uptake of ³H-leucine in both experiments remained unaffected by light conditions, showing high percentages (70–80%) all through incubations (Figures 4c and d). Similarly, *Synechococcus* from the autumn experiment (exp. 4) were not affected by solar radiation levels (Figure 4d), whereas in the summer (exp. 3) they were stimulated by PAR exposure compared with darkness. Inclusion of UVR caused a subsequent 30% decrease in the number of active cells (all differences being significant according to the Tukey's test, $P < 0.05$; Figure 4c).
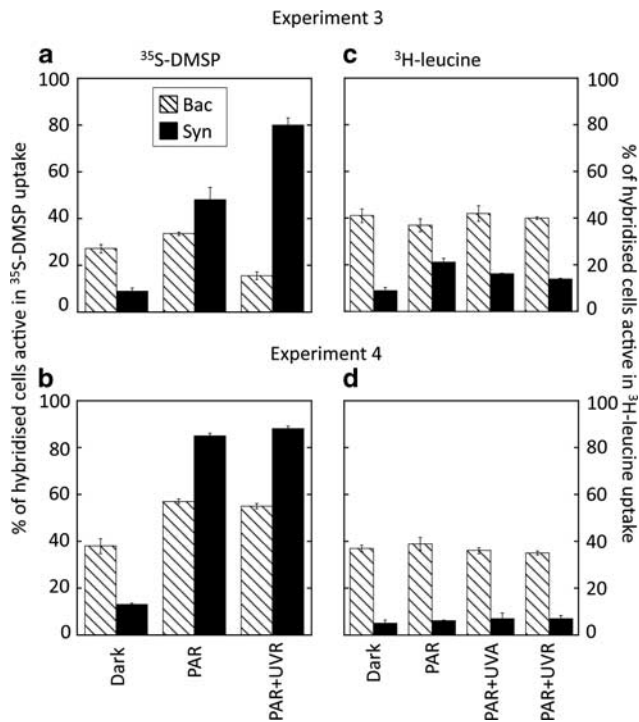
Figure 4 Experiments 3 (top panels) and 4 (bottom panels). Percentage of *Bacteria* (Bac) and *Synechococcus* (Syn) cells taking up ^{35}S-DMSP (**a**, **b**) or ^3H-leucine (**c**, **d**) as quantified by MAR-CARD-FISH after exposure to different light conditions. ^{35}S-DMSP incubations lacked the PAR + UVA treatment.
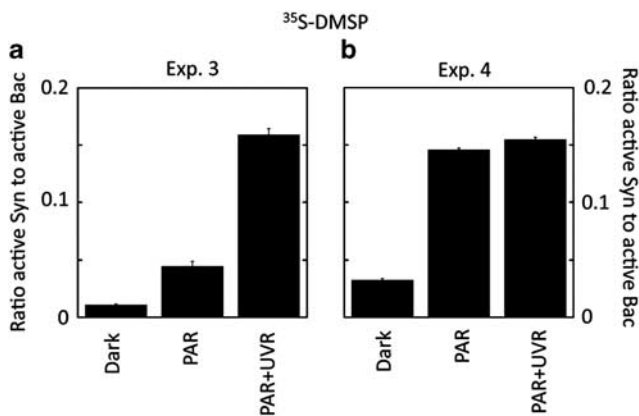

Figure 5 Experiments 3 (**a**) and 4 (**b**). Ratio between the total number of active *Synechococcus* (Syn) and the total number of active *Bacteria* (Bac) in ^{35}S-DMSP uptake upon different light conditions. Values were calculated by multiplying the percentage of active cells within each group by their total abundances.

Diel variation in the bulk assimilation of ^{35}S-DMSP by size-fractionated plankton

A further assessment of the relative uptake and assimilation of ^{35}S-DMSP was conducted by size-fractionating samples through 0.65 and 0.22 µm filters, both after prefiltration through 3 µm. Results were subtracted to calculate the contribution of organisms sized either 0.65–3 µm (mostly cyanobacteria and picoeukaryotes) or 0.22–0.65 µm (mostly heterotrophic bacteria). Two experiments at sea were
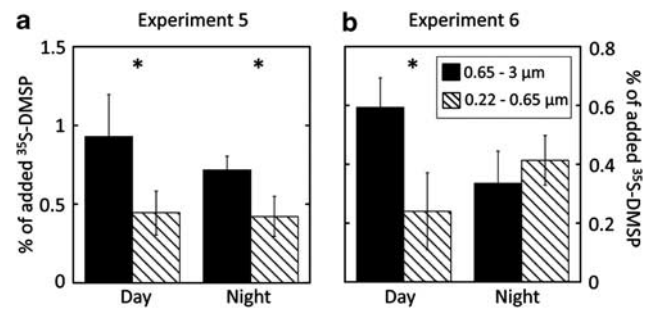

Figure 6 Experiments 5 (**a**) and 6 (**b**). Day- and night-averaged percentages of assimilated ^{35}S-DMSP by organisms sized 0.65–3 µm (black bars) and 0.22–0.65 µm (dashed bars) as measured every 4 h during a 24 h cycle. Values are mean ± standard errors of 3–4 data points. *Significant differences between both size fractions (analysis of variance (ANOVA), $P < 0.05$).

carried out during 24 h cycles, one in shelf waters (exp. 5) and the other in open-sea waters (exp. 6). Incubations were done under natural full sunlight conditions. No clear patterns were observed throughout the cycles when plotting single data points measured every 4 h (data not shown); however, when day and night samples were separately averaged for each fraction, we found that during the day, the larger fraction (containing cyanobacteria) assimilated significantly more substrate than the smaller bacterial fraction (analysis of variance, $P < 0.05$), whereas at night these differences were either lower (cycle 1, Figure 6a) or null (cycle 2, Figure 6b).

Discussion

It is now recognised that the marine picophytoplankton communities composed of *Synechococcus, Prochlorococcus* and small eukaryotic phytoplankters dominate the photoautotrophic plankton over vast tracks of the world's oceans. However, whereas their contribution to global primary production is well documented (Waterbury *et al.*, 1986; Partensky *et al.*, 1999), their role in the consumption of dissolved organic compounds, although recognised, has been much less intensely studied (see, for example, Zubkov and Tarran, 2005; Michelou *et al.*, 2007; Mary *et al.*, 2008).

Results from our dark incubation (exp. 1) further confirmed that similarly to heterotrophic bacteria, both *Prochlorococcus* and *Synechococcus* may benefit from using a reduced sulphur source such as DMSP, probably because it saves the energy required to reduce the abundant sulphate (Kiene *et al.*, 1999). This is the second study reporting ^{35}S-DMSP uptake by a natural *Prochlorococcus* population, after Vila-Costa *et al.* (2006). Studies with cultured and natural assemblages of heterotrophic bacteria showed that DMSP and glycine betaine share the same membrane transporter (Kempf and Bremer, 1998; Kiene *et al.*, 1998).

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Putative glycine betaine transporter genes have been found in the genomes of culture representatives of both *Prochlorococcus* and *Synechococcus* (Palenik *et al.*, 2003; Rocap *et al.*, 2003), thus supporting the observed capacity of these widespread photosynthetic taxa to take up and assimilate DMSP. We found that the amount of radioisotope incorporated per cyanobacterial cell was larger than that per heterotrophic bacterium, consistent with their larger size. On a per biovolume (proxy to biomass) basis, and in agreement with the observations of Vila-Costa *et al.* (2006), heterotrophic bacteria were the most efficient at incorporating DMSP, although they were closely followed by *Synechococcus* and, to a lesser extent, by *Prochlorococcus.* As the sample was collected in the night and it was incubated in the absence of light, it is likely that the observed uptake efficiencies among the studied groups may change in the presence of light because the cyanobacterial heterotrophic uptake of DMSP can be stimulated upon illumination, as suggested by the present and previous works (Malmstrom *et al.*, 2005; Vila-Costa *et al.*, 2006).

Although often considered ecologically unimportant, recent studies indicate that cyanobacterial photoheterotrophy might significantly influence the flux of dissolved organic matter in the euphotic zone of marine ecosystems. As an example, Church *et al.* (2004, 2006) attributed their observed light enhancement of bacterial production to *Prochlorococcus* photoheterotrophy in the North Pacific gyre, and these same organisms were also responsible for ~30% of methionine turnover in the Arabian Sea (Zubkov *et al.*, 2003). Likewise, some *Synechococcus* have been shown to assimilate amino acids (Willey and Waterbury, 1989; Paerl, 1991), yet their contribution to methionine uptake was lower than that of *Prochlorococcus* in the Arabian Sea (Zubkov *et al.*, 2003).

By using flow cytometry cell sorting (exp. 2), we found that exposure to diverse sunlight treatments caused differential effects on *Synechococcus* and heterotrophic bacteria. These two groups had been previously studied in Blanes Bay with regard to their UVR sensitivities. Heterotrophic bacterial activity is generally negatively affected by UVR (Herndl *et al.*, 1993; Sommaruga *et al.*, 1997), although this varies among bacterial taxa (Alonso-Sáez *et al.*, 2006), whereas *Synechococcus* exhibit high resistance (Sommaruga *et al.*, 2005). In agreement with the previous observations, we found that upon PAR and particularly full sunlight exposure, heterotrophic bacteria were inhibited in their ^{35}S-DMSP uptake whereas *Synechococcus* were not. Consequently, the relative role of *Synechococcus* as a DMSP sink became more important under full solar radiation than under dark conditions (Figure 2). Furthermore, on a per-cell volume basis, *Synechococcus* cells equalled the amount of radioisotope incorporated by heterotrophic bacteria (Figure 3), suggesting an important role of sunlight in

regulating the fate of dissolved DMSP and the physiological use of DMSP-sulphur.

Interestingly, we observed that the fraction of assimilated ^{35}S-DMSP not associated with *Synechococcus* or heterotrophic bacteria showed a greater contribution to total uptake toward the full sunlight spectrum (up to 75%, Figure 2a). We have no direct hint of which organisms may be responsible for this large proportion of ^{35}S-DMSP assimilation; Vila-Costa *et al.* (2006) and Ruiz-González *et al.* (submitted) have reported notable uptake activity by large eukaryotic phytoplankton, particularly diatoms. Actually, the amount of ^{35}S-DMSP assimilated by the unknown organisms increased with PAR and PAR + UVA and decreased with respect to those treatments under full sunlight (data not shown), a response similar to that observed in polar diatoms by Ruiz-González *et al.* (submitted). This suggests an important role of eukaryotic organisms in the DMSP fluxes, which still deserves further research.

Assimilation of ^{3}H-leucine was simultaneously measured for comparative purposes as leucine is considered a universal substrate for heterotrophic bacteria (Kirchman *et al.*, 1985) that can also be incorporated by cyanobacteria (Kamjunke and Jähnichen, 2000; Mary *et al.*, 2008). Unlike with ^{35}S-DMSP-amended samples, solar radiation did not seem to alter the relative contributions of both heterotrophic bacteria and cyanobacteria to ^{3}H-leucine uptake (Figure 2d). There are studies showing that light can stimulate the uptake of amino acids by phototrophic organisms (Zubkov and Tarran, 2005; Michelou *et al.*, 2007; Mary *et al.*, 2008), but none studied the process in the presence of UVR.

MAR-CARD-FISH allowed for a visual analysis of the single-cell uptake activity of both *Bacteria* and *Synechococcus* from Blanes Bay in exps. 3 and 4. As a result of light exposure, the relative contribution of *Synechococcus* to the number of ^{35}S-DMSP-assimilating cells strongly increased, whereas that of heterotrophic bacteria either decreased upon UVR exposure (exp. 3, early July) or increased, but less than *Synechococcus* (exp. 4, late September). Moreover, in spite of their low abundances, active *Synechococcus* accounted for 15% (exps. 3 and 4) of active *Bacteria* under full sunlight, a proportion comparable to those observed for other major bacterial groups in diverse ecosystems (Vila *et al.*, 2004; Malmstrom *et al.*, 2004b; Vila-Costa *et al.*, 2007), and for *Synechococcus* in the Northwest Atlantic and the Gulf of Mexico (Malmstrom *et al.*, 2005). Hence, according to this significant assimilation capacity and their widespread distribution (Waterbury *et al.*, 1979), *Synechococcus* are likely to be an important sink for marine DMSP. Interestingly, a higher contribution of *Synechococcus* was observed in exp. 3 than in exp. 4, in accordance with higher UVB doses measured in the former (Table 1). Yet, any hypothesis about *Synechococcus* heterotrophy dependence on irradiance levels remains to be

tested. Once again, this pattern was not reflected in ³H-leucine uptake. Only in exp. 3, *Synechococcus* seemed to respond to light changes, showing an increase in the proportion of active cells upon PAR exposure and a further decrease when UVR was included. This light stimulation of ³H-leucine uptake by *Synechococcus* has been recently observed at the BBMO across different seasons except autumn, and hence has the corresponding UVR-induced inhibition, which was stronger in the spring and summer periods coinciding with higher UVR doses (Ruiz-González, unpublished). Despite the higher percentages of ³H-leucine-assimilating *Synechococcus* cells found in our light incubations, their contribution to total active *Bacteria* was never higher than 3%. Altogether, these results indicate that heterotrophy in *Synechococcus* is strongly dependent not only on sunlight spectrum and maybe intensity, but also on the type of substrate considered.

Shipboard exps. 5 and 6 served to further explore the potential osmoheterotrophic competition between cyanobacteria and heterotrophic bacteria at the sub-daily scale. Over the two 24 h periods, a trend toward higher ³⁵S-DMSP assimilation by the cyanobacteria-containing, larger picoplankton fraction (that is, 0.65–3 µm) was found during the light hours, whereas at night, it was more evenly distributed between the two size fractions. Interestingly, in most cases, the average ³⁵S-DMSP assimilation in the 0.65–3 µm fraction was significantly higher than that in the 0.22–0.65 µm fraction, pointing to a more important contribution of picophytoplankton (and maybe attached bacteria) than that of free-living heterotrophic bacteria to total DMSP-sulphur assimilation. However, size fractionation is an inaccurate method to assess the distribution of assimilation owing to imperfect size segregation, inclusion of detritus and overlook of bacterial aggregates. Furthermore, the larger fraction included, beside cyanobacteria, picoeukaryotic cells that could have also assimilated ³⁵S-DMSP (Vila-Costa *et al.*, 2006) or could have ingested labelled bacteria. Whether these complementary players also respond positively to light is unknown. In any case, these results support the aforementioned observations at the single-cell level that light shifts DMSP-sulphur assimilation away from the clear dominance of heterotrophic bacteria usually found in dark incubations.

As a photosynthetic cell, *Synechococcus* is subjected to diel variations in energy supply over the light/dark cycle, and many physiological processes, such as specific enzyme transcription, DNA synthesis or cell division, are coupled to this periodicity (Wyman, 1999; Jacquet *et al.*, 2001). Hence, diel variations in *Synechococcus* and other picophytoplankters' activities are expected to result in shifts in the relative dominance of phototrophs versus heterotrophs in the uptake of DMSP-sulphur throughout the light/dark cycle. Additionally, similar shifts may also occur at the seasonal scale.

Considering that in Blanes Bay, both the highest concentrations of dissolved DMSP (Vila-Costa *et al.*, 2008a) and the maximum abundances of *Synechococcus* (Agawin *et al.*, 1998; Schauer *et al.*, 2003) occur in the highly irradiated waters of late spring and summer, one might expect competition between cyanobacteria and heterotrophic bacteria (and possibly larger phototrophs; Vila-Costa *et al.*, 2006) for DMSP-sulphur to be maximal in summer and minimal in autumn and winter.

Overall, our results confirm that marine, free-living, unicellular cyanobacteria (that is, *Prochlorococcus* and *Synechococcus*) from the Mediterranean Sea are able to take up DMSP and assimilate its sulphur, and all our different experimental approaches agreed with an increased contribution of *Synechococcus* and probably some picoeukaryotes to ³⁵S-DMSP uptake relative to heterotrophic bacteria under light exposure compared with dark conditions. These results suggest that the DMSP dynamics in oceanic surface waters are severely influenced by solar UVR through differential inhibition or stimulation of the microbial consortia responsible for most of the DMSP consumption. Our findings stress the generally overlooked role of phytoplankton as DMSP consumers under realistic light conditions and the need for further research. Interestingly, the pronounced effects of the light-driven activation of *Synechococcus* did not show up in the uptake of ³H-leucine. Determining the reasons for this differential regulation of substrate uptake by light may help better understand and predict the microbial use of labile dissolved organic matter in the surface ocean.

Acknowledgements

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