

ARTICLE



Sulfoquinovose is a widespread organosulfur substrate for *Roseobacter* clade bacteria in the ocean

Le Liu¹, Xiaofeng Chen¹, Jianing Ye¹, Xiaoyi Ma¹, Yu Han¹, Yajie He¹ and Kai Tang¹✉

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Sulfoquinovose (SQ) is one of the most abundant organosulfur compounds in the biosphere, and its biosynthesis and degradation can represent an important contribution to the sulfur cycle. To date, in marine environments, the microorganisms capable of metabolising SQ have remained unidentified and the sources of SQ are still uncertain. Herein, the marine *Roseobacter* clade bacteria (RCB) *Dinoroseobacter shibae* DFL 12 and *Roseobacter denitrificans* OCh 114 were found to grow using SQ as the sole source of carbon and energy. In the presence of SQ, we identified a set of highly up-regulated proteins encoded by gene clusters in these two organisms, of which four homologues to proteins in the SQ monooxygenase pathway of *Agrobacterium fabrum* C58 may confer the ability to metabolise SQ to these marine bacteria. The sulfite released from SQ desulfonation by FMN-dependent SQ monooxygenase (SmcC) may provide bacteria with reduced sulfur for assimilation, while proteins associated with sulfite production via assimilatory sulfate reduction were significantly down-regulated. Such SQ catabolic genes are restricted to a limited number of phylogenetically diverse bacterial taxa with the predominate genera belonging to the *Roseobacter* clade (*Roseobacteraceae*). Moreover, transcript analysis of Tara Oceans project and coastal Bohai Sea samples provided additional evidence for SQ metabolism by RCB. SQ was found to be widely distributed in marine phytoplankton and cyanobacteria with variable intracellular concentrations ranging from micromolar to millimolar levels, and the amounts of SQ on particulate organic matter in field samples were, on average, lower than that of dimethylsulfoniopropionate (DMSP) by one order of magnitude. Together, the phototroph-derived SQ actively metabolised by RCB represents a previously unidentified link in the marine sulfur cycle.

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INTRODUCTION

SQ is a sulfonated hexose compound that is present in higher plants, algae and photosynthetic bacteria [1]. SQ can be produced via microbial acyl- and glucoside hydrolases acting on sulfoquinovosyl diacylglycerol (SQDG), a sulfolipid that can partially compensate for phosphatidylglycerol deficiency in photosynthetic membranes under phosphate-limiting conditions [2, 3]. Up to 10 billion tons annually, SQ is one of the most abundant organic sulfur compounds in the biosphere [4, 5], and represents a significant fraction of the sulfur cycle.

Within the natural organic sulfur cycle, microorganisms generally decompose sulfur-containing organic compounds in order to grow, yielding sulfur that is further incorporated into living organisms or released into the environment [6, 7]. The bacterial catabolism of SQ has attracted considerable attention because of its significance to the global sulfur flux and the regulation of SQ metabolism in the intestinal track [4, 8, 9]. The existence of microbial SQ degradation was first observed in soil [10]. In addition to the abundant SQ-utilizing bacterial strains from soil environments, cultured isolates have been recovered from plants, sewage sludge, as well as fresh- and hypersaline water, and even the human intestine [8, 9, 11–14]. In parallel to classic glycolysis, bacteria can convert SQ to the three-carbon organosulfonates 2,3-dihydroxypropane-1-sulfonate (DHPS)

and sulfolactate via analogous reactions through three pathways, namely the sulfolactate Entner–Doudoroff (sulfo-ED), the sulfolactate Embden–Meyerhof–Parnas (sulfo-EMP) and its variant (sulfo-EMP2) pathways [8, 11, 14]. The sulfo-EMP pathway is commonly found in Gram-negative intestinal bacteria and the sulfo-EMP2 pathway extends the sulfo-EMP to Gram-positive bacteria [8, 15]. The sulfo-ED pathway was frequently found in freshwater and soil bacteria [11]. Sulfolactate and DHPS can also be produced in aerobic and anaerobic bacteria, respectively, via the transaldolase-dependent sulfolactate pathway (sulfo-transaldolase) pathway [12]. Anaerobic bacteria were also found to carry out transketolase-dependent sulfolactate pathway (sulfo-transketolase) pathway for fermenting SQ, producing isethionate [14]. The sulfonate sulfur derived from these pathways could be further converted into sulfite or sulfide by other bacteria to complete a fully degradative process for SQ [13, 16–19]. Recently, a novel pathway (sulfoquinovose monooxygenase pathway, sulfo-SMO) has been identified as a prevalent SQ metabolic pathway across rhizobial strains, especially in the genera *Agrobacterium* and *Rhizobium* [20]. The sulfo-SMO pathway represents the only known complete degradation pathway for SQ within a single bacterial species, where the desulfonation of SQ into sulfite by a monooxygenase occurs at the first step and the

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subsequent two reduction reactions result in the formation of glucose for bacterial growth [20].

Marine eukaryotic phytoplankton and prokaryotic cyanobacteria have the ability to synthesize SQDG and thus have been proposed as the sources of SQ in the ocean [21]; however, there is no evidence for the presence of free-form SQ in their metabolites. Moreover, the known SQ metabolic pathways have not yet been reported in phytoplankton and cyanobacteria [21]. Accordingly, marine microorganisms can be considered metabolic specialists, having evolved the capacity to catabolize SQ similar to microorganisms in other ecosystems, but to date, the identity of marine SQ-degrading microorganisms remain unclear and the likely relevance of SQ in ecosystem functioning poorly understood. This work identifies such a missing module of the marine organic sulfur cycle by revealing the SQ degradation pathway in marine bacteria as well as the microbial catabolic potential and natural occurrence of SQ in the ocean.

MATERIALS AND METHODS

Chemical synthesis of SQ and DHPS

SQ was synthesized chemically using a previously published protocol [22]. The structure of SQ was verified by ^1H NMR (Fig. S1) and mass spectrometry (Fig. S2). DHPS was synthesized chemically and verified elsewhere [17].

SQ: ^1H NMR (400 MHz, D_2O): δ = 5.24 (d, J = 2.28 Hz, 1H), 4.70 (d, J = 5.28 Hz, 1H), 4.25 (t, J = 6.36 Hz, 1H), 3.82 (t, J = 6.24 Hz, 1H), 3.75 (t, J = 6.32 Hz, 1H), 3.58 (dd, J = 2.44, 6.48 Hz, 1H), 3.52 (dd, J = 6.12, 12.32), 3.43 (d, J = 9.84 Hz, 2H), 3.30 (dd, J = 5.96, 11.56 Hz, 3H), 3.10 (m, 2H) ppm.

Cultivation

The strains *D. shibae* DFL 12 (DSM 16493), *R. denitrificans* OCh 114 (DSM 7001) and *R. pomeroyi* DSS-3 (DSM 15171) were obtained from German Collection of Microorganisms and Cell Cultures (Braunschweig, Germany). *D. shibae* DFL 12 and *R. denitrificans* OCh 114 were grown in modified *Silicibacter* basal medium (pH 7.2) [23] supplemented with SQ (6 mM), glucose (6 mM) or SQ (3 mM) plus glucose (3 mM). *R. pomeroyi* DSS-3 was grown in modified *Silicibacter* basal medium (pH 7.2) [23] supplemented with SQ (6 mM), yeast (0.25 g L^{-1}) or SQ (6 mM) plus yeast (0.25 g L^{-1}). The cultures (100 mL) were incubated in acid-cleaned and pre-sterilized flasks at 28 °C with a shake speed of 160 rpm. At each sampling timepoint, triplicate biological replicates and triplet blank control samples were taken. Samples were taken at intervals to monitor the growth with OD_{600} and the concentrations of SQ, sulfite and sulfate.

Thalassiosira pseudonana 220, *Navicula pelliculosa* 350, *Phaeodactylum tricornutum* 267, *Emiliania huxleyi* CS369, *E. huxleyi* 2090, *E. huxleyi* 1516, *Isochrysis galbana* 079, *I. galbana* 321, *Phaeocystis globosa* 124, *Amphidinium carterae* Hulbert 279, *Alexandrium tamarense* 260, *Chlorella* sp. 309, *Chlorella* sp. 410, *Prochlorococcus* sp. 9301, *Prochlorococcus* sp. 9312, *Synechococcus* sp. XM24, *Synechococcus* sp. 1333, *Synechococcus* sp. YX04-3 were obtained from Center for Collections of Marine Algae (Xiamen University, China). Triplicate axenic phytoplankton cultures were grown in acid-cleaned and sterilized flasks in an incubator. Diatoms, dinoflagellates and haptophytes were grown under the following conditions: light of $100\ \mu\text{E}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$ and temperature of 20 °C. Diatoms were incubated in f/2 media supplemented with Na_2SiO_3 . Dinoflagellates and haptophytes were incubated in f/2 media [24] without Na_2SiO_3 . *Prochlorococcus* and *Synechococcus* were grown on Pro99 media [25] and SN media [26], respectively, with light of $50\ \mu\text{E}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$ and temperature of 20 °C. Samples of phytoplankton cultures at plateau phase were collected to analyze SQ and DHPS concentrations. Cell abundance of diatoms, haptophytes, chlorophyta and *Synechococcus* were measured using an Accuri C6 flow cytometer (BD Biosciences, New Jersey, USA). Cell abundance of *Prochlorococcus* was measured using FACSARIA flow cytometer (BD Biosciences). Cell counts for dinoflagellates were measured using microscopy. Cell volumes of diatoms, haptophytes, chlorophyta and dinoflagellates (Table S1) were based on cell dimensions directly observed under the microscope and calculated using a previously reported method [27]. Cell volumes of cyanobacteria were based on previously reported values of cell dimension and size [28, 29].

Study sites and sample collection

Samples were collected from five sites (A18: 38.42° N, 119.42° E; A31: 38.75° N, 120.42° E; A35: 39.42° N, 119.70° E; A42: 37.75° N, 119.40° N; A45: 38.00° N, 120.42° E) in the Bohai Sea, China, on August 18–30, 2018 (Fig. S3). Seawater samples were collected using 10-L Niskin bottles mounted on a rosette sampler (model 1018, General Oceanics, Florida, USA). Seawater samples were collected from three depths (surface, ~3 m below the surface; middle; and bottom, ~2 m above the bottom), with the depths dependent on the sampling sites. Seawater (0.7–3 L) was pre-filtered through a nylon screen (200 μm), then filtered via a polycarbonate filter (47 mm, 3 μm , Merck Millipore, New Jersey, USA) to collect eukaryotic phytoplankton cells. After collection, the filters were immediately frozen by liquid nitrogen, and stored at $-80\text{ }^\circ\text{C}$ until further extraction. Approximately 0.5 L of seawater was filtered through precombusted (480 °C for 5 h) GF/F glass fiber filter membrane (47 mm, 0.75 μm , Whatman, USA), and the filters were subsequently stored at $-20\text{ }^\circ\text{C}$ until the measurement of particulate organic carbon (POC) with an Elemental Analyzer (Carlo-Erba, Milano, Italy) [30].

Samples were collected from the S51 station (14.00° N, 116.00° E) in the South China Sea, on July 13–14, 2019. Seawater was sampled using 10 L Niskin bottles mounted on rosette sampler (SBE-911 plus, SEA-BIRD, WA, USA). Sampling depth was ~5 m below the surface. Seawater (200 L) was pre-filtered via a polycarbonate filter (144 mm, 3 μm , Merck Millipore), then concentrated to 750 mL using crossflow filters (Sartocon, Göttingen, Germany) equipped with a Sartocon Hydrosart Cassette (0.6 m^2 , 0.2 μm , Sartocon). Concentrated seawater (100 mL) was taken to filter through a 0.2 μm polycarbonate membrane (Merck Millipore). The filters were immediately frozen at $-20\text{ }^\circ\text{C}$ until further extraction of metabolites. The solution of 50% glutaraldehyde was added to the concentrated seawater (1:100 v/v) and then frozen at $-20\text{ }^\circ\text{C}$ until further analysis. The cell abundance of *Prochlorococcus* within seawater was determined by FACSARIA flow cytometer (BD Biosciences) [31].

Seawater samples were collected from a harbor in the Xiamen coastal sea (24.53° N, 118.18° E) using an acid-cleaned plastic bucket. Sampling depth was ~0.5 m below the surface. Seawater (7 L) was filtered via polycarbonate membrane (144 mm, 3 μm , Merck Millipore) followed by a 0.2 μm polycarbonate membrane (Merck Millipore). The 0.2 μm filters were immediately frozen at $-20\text{ }^\circ\text{C}$ until further extraction of intercellular SQ for *Synechococcus* in field sample. The solution of 50% glutaraldehyde was added to the 3 μm -filtered seawater (1:100 v/v) and then frozen at $-20\text{ }^\circ\text{C}$ until further analysis. The cell abundance of *Synechococcus* within seawater was determined by FACSARIA flow cytometer (BD Biosciences) [31]. The 0.2 μm -filtered seawater was used for the extraction of dissolved SQ.

Extraction and sample preparation of metabolites in culture samples

For determination of metabolites in the medium supernatant, 10 μL of supernatant, from 100 μL of medium that had been centrifuged at 12,000 rpm for 15 min, was added to 990 μL of an extraction solvent, composed of acetonitrile, methanol and water (2:2:1 v/v/v) and stored at $-20\text{ }^\circ\text{C}$ for 2 h. The sample was centrifuged at 12,000 rpm for 5 min at 4 °C, and 100 μL of that supernatant was transferred to 1 mL glass vial (CNW, Düsseldorf, Germany).

For determination of intracellular metabolites from phytoplankton and cyanobacteria, cultures were filtered thru a 0.2 μm polycarbonate membrane (Merck Millipore). The filters were cut into pieces with dissecting scissors rinsed with water in 10% formic acid. The filter pieces were transferred to bead beating tubes and 1 mL of cool extraction solvent and approximately 500 μL of silica beads were added. The samples were shaken on a FastPrep-24 Homogenizer (MP Biomedicals, CA, USA) for 40 s and then frozen for 30 min. This step was repeated for a total of two times. The samples were centrifuged at 12,000 rpm for 5 min at 4 °C to remove precipitates. The 100 μL of supernatant was transferred to a 1 mL glass vial.

Extraction and sample preparation of metabolites in field samples

The intracellular metabolites of phytoplankton cells were extracted by acetonitrile:methanol:water (2:2:1 v/v/v, $-20\text{ }^\circ\text{C}$) solution. Filter samples were homogenized using the FastPrep-24 Homogenizer (MP Biomedicals), followed by incubation at $-20\text{ }^\circ\text{C}$ for 2 h. After centrifugation, the supernatant was filtered through a hydrophilic polyvinylidene fluoride membrane filter (13 mm, 0.2 μm , JINTENG, Tianjin, China), and then stored at $-80\text{ }^\circ\text{C}$ until UPLC/MS analysis.

For determination of dissolved SQ in field samples, a 90-mL seawater sample was filtered through a polycarbonate membrane (47 mm, 0.2 µm, Merck Millipore) and then freeze-dried and redissolved in 6 mL acetonitrile. The precipitate was discarded after centrifuging at 5000 g for 10 min. The supernatant was transferred to a 40 mL brown glass vial and dried under a gentle stream of nitrogen at room temperature and redissolved in 100 µL of a mixture of acetonitrile and water (9:1 v/v). The sample was stored at -80 °C until HPLC/MS analysis after centrifuging at 8000 g for 10 min.

HPLC/MS analysis

The concentration of dissolved SQ in sea water was quantified using a Q Exactive high performance liquid chromatography system (Thermo Fisher Scientific, Waltham, USA) fitted with a ZIC-HILIC column (3.5 µm, 200 Å, 150 × 2.1 mm; Merck), which was connected to an orbitrap mass spectrometer (Thermo Fisher Scientific). Electrospray ionization was performed in negative mode ionization with the following parameters: capillary temperature 320 °C, spray voltage 3500 V, sheath gas flow 40 arbitrary units and aux gas flow 10 arbitrary units. The HPLC conditions were as follows: the column was equilibrated with 90% solvent A (100% acetonitrile) and 10% solvent B (10 mM NH₄Ac) for 10 min. The gradient was comprised of an increase from 10% to 35% solvent B over 25 min, then held at 35% for 10 min, 35% to 10% over 0.5 min, and finally held at 10% for 10 min. The flow rate was 0.25 mL/min and the injection volume was 2 µL. Data analyses were performed with the Xcalibur software version 2.0.3 (Thermo Fisher Scientific). Calibration curve for synthetic SQ as standard was used to determine SQ concentrations. The detection limit was 10 nM for SQ with a signal-to-noise ratio of three.

UPLC/MS analysis

The concentrations of SQ, DHPS, DMSP, sulfolactate and isethionate in culture and on particulate organic matter were quantified by ultra-high performance liquid chromatography-mass spectrometry analysis. Metabolite concentrations were generated using an Agilent 6460 triple quadrupole mass spectrometer (Agilent, CA, USA) coupled to an Agilent 1290 Infinity II series UHPLC System (Agilent), with an UHPLC BEH Amide column (1.7 µm, 2.1 mm × 100 mm, Waters, MA, USA). The determination parameters were set up as described in [17]. Agilent MassHunter Work Station Software B.08.00 (Agilent) was used for MRM data acquisition and processing. Calibration curves for individual analytes were generated to determine metabolite concentrations. The detection limits were 0.61 nM for SQ, 0.08 nM for DHPS, 0.02 nM for DMSP, 0.30 nM for sulfolactate and 0.30 nM for isethionate with signal-to-noise ratios of three. Taurine-d4 was used as isotopically labelled internal standard.

The quantification for sulfite

Sulfite of culture supernatant was quantified colorimetrically by the fuchsin sulfite assay [20, 32], which requires solution A (4.0 mg of basic fuchsin was dissolved in 8.25 mL of deionized water at 0 °C, prior to the addition of 1.25 mL of 98% H₂SO₄), solution B (0.32 mL of 36% formaldehyde added into 9.68 mL deionized water at 0 °C) and solution C (1 mL solution A diluted with 7 mL deionized water, prior to addition of 1 mL solution B). Sample (36 µL) was mixed with solution C (174 µL). The mixture was incubated for 10 min at room temperature and the absorbance at 570 nm was measured using a SynergyH1 Multi-Mode Reader (BioTek, Winooski, Vermont). Calibration curve for a Na₂SO₃ standard was used to determine sulfite concentrations. The detection limit was calculated to 1.4 µM for sulfite with the standard deviation method [33].

The quantification for sulfate

Sulfate of culture supernatant was quantified using the ICS 5000 ion chromatography system (Dionex, Thermo Fisher Scientific) with IonPac AS15 column (9 µm, 4 mm × 250 mm, Dionex). The mobile phase was deionized water. Suppressor hydroxide was 36 mM and the current was 90 mA. The flow rate was 1.0 mL/min and the injection volume was 20 µL. Calibration curve for a Na₂SO₄ standard was used to determine sulfate concentrations. The detection limit of sulfate was 7.0 µM with a signal-to-noise ratio of three.

Proteomic analysis

D. shibae DFL 12 was grown in modified *Silicibacter* basal medium with SQ (6 mM) as the sole carbon source. The control group medium was

glucose (3 mM) or acetate (9 mM) as the sole carbon source. Cell pellets were harvested after centrifugation at 12,000 rpm for 15 min. Lysis buffer (8 M urea, 1% protease inhibitor, Cocktail) was used to suspend the cell pellets and the mixture was sonicated three times on ice to lyse cells. Cell debris were removed by centrifugation at 12,000 g at 4 °C for 10 min. Concentration of extracted protein was determined with BCA Protein Assay Kit (Bio-Rad, CA, USA). Trichloroacetic acid was slowly added to the protein solution at a final concentration of 20%. The sample solution was placed at 4 °C for 2 h after vortexing and then centrifuged at 4500 g for 5 min. The supernatant was discarded. The protein precipitates were washed three times with cold acetone. Final concentration of 200 mM triethylamine borane was added to the dry protein precipitates followed by sonication to resuspend the protein pellets. Trypsin (1/50 w/w) was added to the sample solution to digest the protein overnight. The digested protein solution was reduced with dithiothreitol, 5 mM final concentration, for 30 min at 56 °C followed by incubation with 11 mM iodoacetamide at room temperature, in the dark, for 15 min. Digested peptides were desalted using a C18 SPE column (7 mm, volume 3 mL, Sigma, Missouri, USA) and vacuum dried. Peptides were then dissolved in solvent A (0.1% formic acid, 2% acetonitrile). The gradient was comprised of an increase from 6% to 24% solvent B (0.1% formic acid in 100% acetonitrile) over 70 min, 24% B to 35% B over 14 min, then climbing to 80% B in 3 min, and held at 80% B for 3 min, at a constant flow rate of 0.45 µL/min. Peptides were then subjected to capillary ion source followed by mass spectrometry (MS/MS) on a tims TOF Pro (Bruker, Billerica, USA) coupled online to an UPLC system with the spray voltage of 1.75 kV and MS/MS survey scan of 100–1700 *m/z*. Data acquisition was in the PASEF mode. The resulting MS/MS data were searched using MaxQuant (v1.6.15.0) against the genome sequences of *D. shibae* DFL 12 downloaded from the National Center for Biotechnology Information (NCBI) database (<https://www.ncbi.nlm.nih.gov/>). The protease specificity was set to trypsin/P. Two miscleavage events were allowed. The first search and main search of MS tolerance were both set at 20 ppm. MS/MS tolerance was set at 20 ppm. Carbamidomethylation of cysteine was set as a fixed modification. Oxidation of methionine as well as acetylation of protein N-termini were set as variable modifications. A false discovery rate (FDR) of 1.0% was set for protein and peptide-spectrum matching identifications. Differential expression analysis was performed using LFQ intensities, which were transformed to Log₂, and the value of Log₂FC were used to generate heatmap using GraphPad Prism 8.0.2 (GraphPad Software, San Diego, USA). The same protocol was used for the proteomic analysis of *R. denitrificans* OCh 114.

Bioinformatic analysis

The sulfur metabolism-related protein sequences in *D. shibae* DFL 12 (FMN-dependent SQ monooxygenase, SmoC, Dshi_2757; DHPS dehydrogenase, HpsN, Dshi_2034; DMSP demethylase, DmdA, Dshi_2320; L-cysteate sulfolyase, CuyA, Dshi_1439; phosphoadenylylsulfate reductase, CysH, Dshi_1152; sulfite oxidase, SoeB, Dshi_1237; sulfolactate sulfolyase, SuyB, Dshi_2038; sulfoacetaldehyde acetyltransferase, Xsc, Dshi_2045), taurine dioxygenase in *Escherichia coli* K12 (TauD, Eco06-DRAFT_02825) and SQ dehydrogenase in *Pseudomonas putida* SQ1 (SqED, PpSQ1_00090) were BLASTp against the marine prokaryote genome databases MAR [34] with an E-value of 1e⁻⁵. The candidate amino acid sequences of these genes were then submitted to similarity searches against UniRef90 (E-value, 1e⁻⁵) [35] and were manually examined in terms of annotations. MAFFT [36] was used to align retrieved sequences with the query sequences that possessed known three-dimensional structures (Protein Data Bank ID: SmoC, 7OH2; DmdA, 3TFH; CuyA, 4D8T; CysH, 2GOY; TauD, 6EDH) [20, 37–40]. Those sequences containing conserved amino acid residues within the catalytic domains of enzymes were retained (Fig. S4a–e). AlphaFold2 was used to predict the structures of proteins [41, 42]. The structure of the lowest-scoring sequence had highly similar folds to the given protein structure of query sequence with the template modeling score > 0.5 measured by US-align [43, 44] (Fig. S4). IQ-TREE [45] was used to determine phylogenetic relationships between homologous sequences using the maximum likelihood method and 1000 bootstraps (Fig. S4). Thus, the minimum bit score of sequences was selected as a cut-off value of an alignment against the *Tara* Oceans database [46] and Bohai Sea metatranscriptomes [30] (143 for SmoC, 110 for DmdA, 139 for CuyA, 103 for CysH, 104 for TauD, 167 for HpsN, 276 for SoeB, 147 for SuyB, 160 for Xsc, 51 for SqED). The abundance results represented the homologs

fraction in the total gene set for each sample. The abundance of eukaryotes and cyanobacteria were obtained from *Tara* Oceans OTU 18 S V9 version 2 database [47]. The potential key proteins of other SQ degradation pathways (sulfo-ED, sulfo-EMP, sulfo-EMP2, sulfo-transaldolase and sulfo-transketolase) were BLASTp against the Ocean Microbiomics Database [48], which included the metagenomic data (*Tara* Oceans, Malaspina, Biogeotraces, HOT and BATS) [49–51] and publicly available genome data (GORG SAGs, the MAR databases and manually-curated MAGs) [34, 52, 53]. The genomes that contained homologs (sequence identities >40%, simultaneously) of all key proteins for SQ degradation pathways were retained for further analysis. Protein structures were displayed using Pymol 2.4 (Schrodinger, NY, USA). Violin plot was generated by GraphPad Prism 9 (GraphPad Software). Correlation analysis and graph creation were performed by OriginPro (OriginLab, MA, USA). The geographical pattern of *smoC* in the global oceans was visualized with Tableau 2022 (Tableau Software, Seattle, USA). A phylogenetic tree for the distribution of candidate sulfo-SMO genes in publicly available marine bacterial genomes using the concatenated alignment of 39 single-copy orthologous genes shared by bacterial strains was constructed by the Maximum Likelihood method using MEGA 6.0 with LG + F model plus a gamma distribution with eight categories [54]. Bootstrap resampling was performed for 1000 replications. Phylogenetic trees were visualized with Evolview v3 [55].

Heterologous expression and purification of proteins

The genes *smoABC* of *D. shibae* DFL 12 were synthesized by Sangon Biotech (Sangon, Shanghai, China). The synthesized genes were reconstructed into vectors pColdII followed by transformation into *Escherichia coli* BL21 (DE3), which were grown in LB media (50 µg/mL ampicillin) with shaking at 200 rpm and 37 °C until OD₆₀₀ reached 0.8. The culture was cooled to 16 °C and isopropyl β-D-thiogalactopyranoside was added to the culture with a final concentration of 500 µM. The culture was maintained at 200 rpm of shaking at 16 °C for 16 h. Cells were harvested by 12,000 g of centrifugation for 30 min at 4 °C, followed by resuspension with 10 mM Tris-HCl buffer (pH 8.0, 150 mM NaCl). The

suspension was subjected to sonication to disrupt the cells. Cell debris was removed by 12,000 g of centrifugation for 20 min at 4 °C. The supernatant was filtered through a 0.45 µm membrane (Merck Millipore) and then loaded onto AKTA pure (Cytiva, MA, USA) couple with a HisTrap HP column (5 mL, Cytiva). The product fraction was eluted with imidazole-containing (50–500 mM) elution buffer (10 mM Tris-HCl, 150 mM NaCl, pH 8.0) at a flow rate of 1 mL/min. The collected product fraction was further subjected to a desalting column (5 mL, Cytiva) with elution buffer to remove imidazole.

Enzyme assay and product analysis

For SmoA, a 200 µL reaction mixture containing 10 mM Tris-HCl pH 8.0, 150 mM NaCl, 1 mM NADPH, 100 µM FMN and 57 µM SmoA was incubated at 30 °C for 20 min. The absorbance at 340 nm was monitored by SynergyH1 Multi-Mode Reader (BioTek, Winooski, Vermont) in real time. For verifying the enzymatic function of SmoB, a 200 µL reaction mixture containing 10 mM Tris-HCl pH 8.0, 150 mM NaCl, 3 mM NADPH, 20 µM FMN, 11 µM SmoA and 2 µM SmoC was incubated at 30 °C for 3 h. SmoC and SmoA in the reaction mixture were removed by an ultrafiltration tube (1.5 mL, 3000 Dalton, Millipore), followed by the additions of 2 µM SmoB and 0.5 mM NADPH and allowed to continue to incubate for 1 h at 30 °C. After the addition of 990 µL acetonitrile into 10 µL final reaction mixture, the sample was subjected to LC-MS. The LC-MS protocol has been described in the HPLC/MS analysis section. For SmoC, a 200 µL reaction mixture containing 10 mM Tris-HCl pH 8.0, 150 mM NaCl, 5 mM NADPH, 20 µM FMN, 5 mM SQ, 11 µM SmoA and 10 µM SmoC was incubated at 30 °C for 1 h. The 50 µL reaction mixture was taken to detect for the generation of sulfite in the acidic fuchsin assay [56]. The reaction mixture was derivatized with 2, 4-dinitrophenylhydrazine (DNPH) before LC-MS analyses [19]. LC-MS analysis was performed using a UPLC BEH C18 column (1.7 µm, 2.1 mm × 50 mm, Waters, MA, USA) with the following protocol: The column was equilibrated with 95% solvent A (0.1% formic acid) and 5% solvent B (100% acetonitrile) for 10 min. The gradient was comprised of an increase from 5% to 50% solvent B over 40 min, then an increase to 95% solvent B for 5 min, held at 95% for 5 min, then decreased from 95% to 5% over 1 min, and finally held at 5% for 10 min. The flow rate was 0.2 mL/min

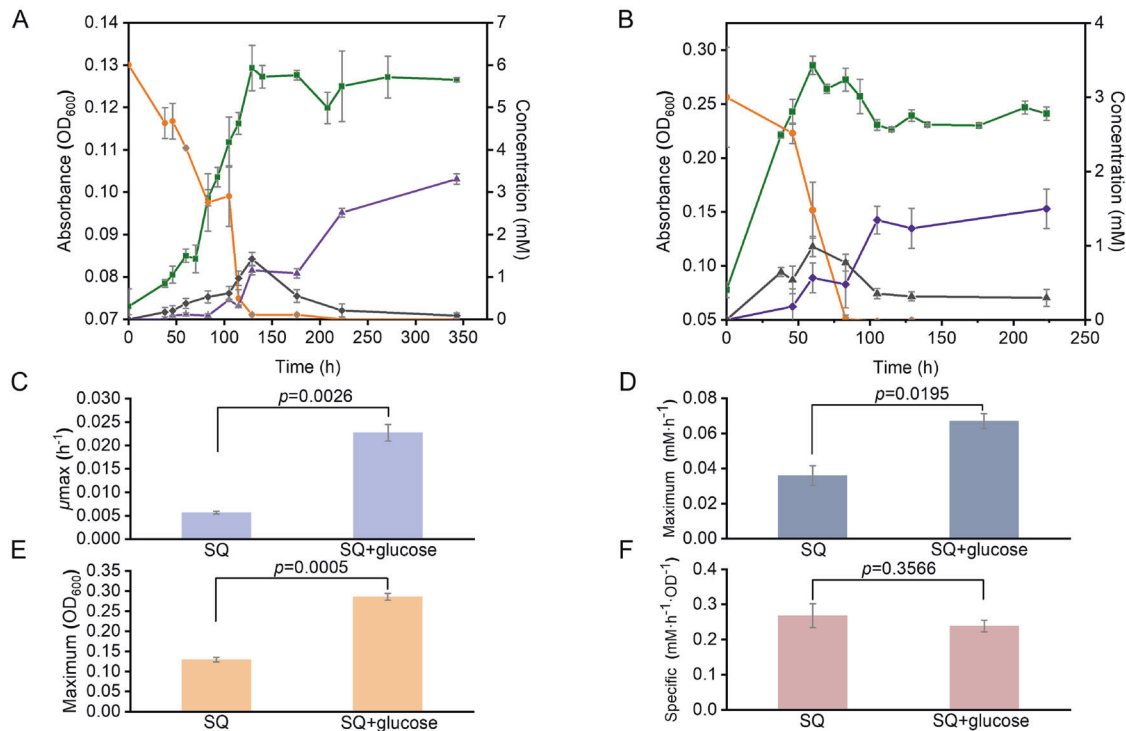


Fig. 1 *D. shibae* DFL 12 utilizes SQ. *D. shibae* DFL 12 utilization of SQ (A) and SQ plus glucose (B) for growth. Optical density (green) of *D. shibae* DFL 12, concentration of SQ in medium (orange), change in concentration of sulfite in medium (purple) and change in concentration of sulfite in medium (black), with respect to time. Maximum growth rate (C) and maximum biomass (E) for *D. shibae* DFL 12 grown on SQ and SQ plus glucose. Maximum (D) and specific (relative to total biomass) (F) SQ consumption rates for *D. shibae* DFL 12 grown on SQ and SQ plus glucose. Error bars represent the standard deviation ($n = 3$). P values were calculated based on t -test to indicate the significance between different groups.

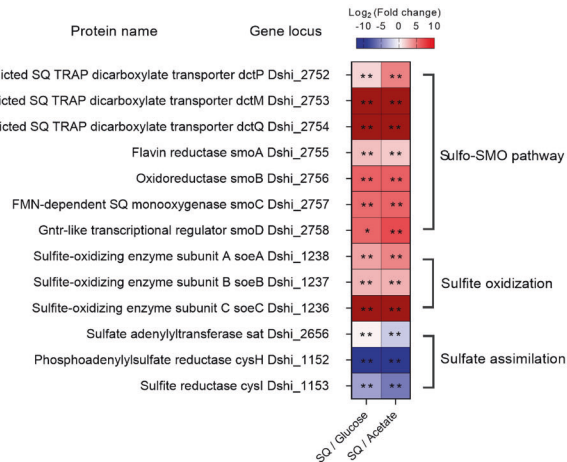
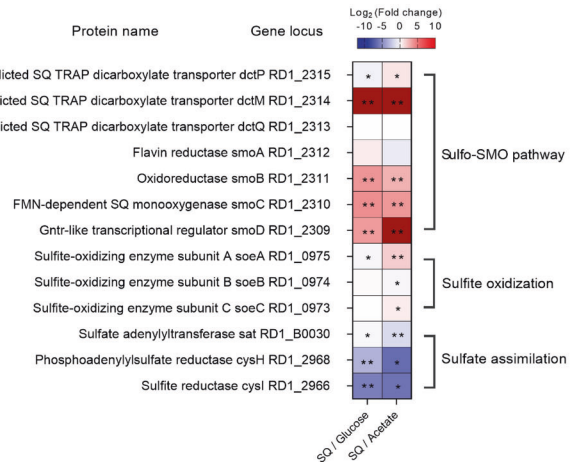
A. *Dinoroseobacter shibae* DFL 12B. *Roseobacter denitrificans* OCh 114

Fig. 2 Comparative proteomic analysis. *D. shibae* DFL 12 (A) and *R. denitrificans* OCh 114 (B) grown on SQ vs glucose and SQ vs acetate. Since each treatment was performed in triplicate, results are shown as average values. Corrected *P*-values (*q*-values) were calculated based on Multiple *t*-test using FDR approach (5%). Significance: **q* < 0.05; ***q* < 0.01.

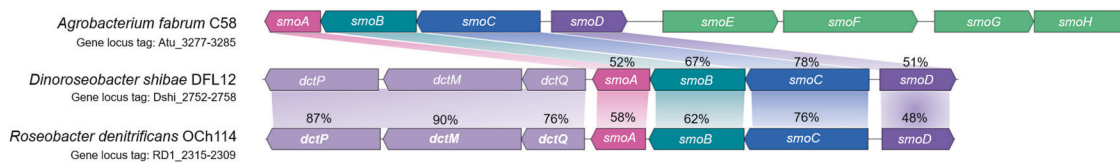


Fig. 3 Gene clusters encoding the proteins involved in the metabolism of SQ, with gene locus tag and annotations, in *A. fabrum* C58, *D. shibae* DFL 12 and *R. denitrificans* OCh114. Gene functions are represented by different colors. ■ *smoA*, gene encoding flavin reductase; ■ *smoB*, gene encoding oxidoreductase; ■ *smoC*, gene encoding FMN-dependent SQ monooxygenase; ■ *smoD*, gene encoding GntR-like transcriptional regulator; *smoEFGH*, genes encoding ABC transporters for SQGro (glycer-1-yl α -sulfoquinovoside) uptake; ■ *dctPMQ*, genes encoding predicted TRAP dicarboxylate transporters for SQ uptake. Homologous genes are connected by colored bars between the genomes. The percentage values on the genes *dctP*, *dctM* and *dctQ* indicate the amino acid similarities between *D. shibae* DFL 12 and *R. denitrificans* OCh 114. The percentage values on the genes *smoA*, *smoB*, *smoC* and *smoD* indicate the amino acid similarities to the *A. fabrum* C58 proteins.

and the injection volume was 2 μ L. Data analyses were performed with the Xcalibur software version 2.0.3 (Thermo Fisher Scientific).

RESULTS AND DISCUSSION

Bacterial culture work

A marine *Roseobacter* clade bacterium, *Dinoroseobacter shibae* DFL 12 [57] was found to have the ability to degrade SQ (Fig. 1). The potential SQ catabolic products DHPS, sulfolactate and isethionate were not detected in culture supernatant. Instead, SQ was utilized completely by *D. shibae* DFL 12 for growth with the concomitant release of sulfite and sulfate into the culture medium (Fig. 1). *D. shibae* DFL 12 exhibited a significantly lower growth rate with SQ as the sole organic substrate compared to that in glucose and acetate ($p < 0.01$, *t*-test) (Fig. S5), suggesting that SQ was a less-preferred substrate for this organism. The bacterium *Ruegeria pomeroyi* DSS-3 displayed a relatively slow-growth rate with either sulfonate DHPS or isethionate as the sole organic substrate as well, but growth was faster when more-labile substrates were co-present with the less-labile sulfonates [58]. Similarly, the maximum growth rate and biomass for *D. shibae* DFL 12 grown on SQ plus glucose were both significantly higher than when grown on SQ alone ($p < 0.01$, *t*-test) (Fig. 1C, E). Meanwhile, the maximum SQ consumption rates for *D. shibae* DFL 12 in SQ plus glucose medium was higher than that in SQ only medium (Fig. 1D); however, the specific SQ degradation rates showed no significant difference (Fig. 1F), suggesting that the enhancement of utilization of SQ in SQ plus glucose relative

to only SQ was most likely due to the increased biomass with growth on glucose (Fig. 1A, B, F, and S4). The simultaneous availability of multiple organic substrates derived from algae enables heterotrophic microorganisms to grow relatively fast, which would presumably be the case within a phycosphere that harbors various labile resources as well as less-labile substrates, such as sulfonates [58].

Comparative proteomic analysis and gene functional verification

Compared to cells grown on glucose and acetate, one cluster of significantly up-regulated proteins in *D. shibae* DFL 12 were clearly discerned during growth with SQ, (locus tag, Dshi_2752-2759; fold-change > 3, *q*-value < 0.0001, Multiple *t*-test) (Fig. 2A; Table S2). The gene cluster was comprised of seven consecutive genes that had previously been annotated in the Integrated Microbial Genomes database [59] as alkanesulfonate monooxygenase, a short-chain oxidoreductase, a flavin reductase, a transcriptional factor, and TRAP dicarboxylate transporter components (Fig. 3). Among which, four encoded proteins were homologous to *SmoABCD* as encoded by a gene cluster in the soil bacterium *A. fabrum* C58 for SQ catabolism (Fig. 3) [20]. Intriguingly, two encoded proteins (Dshi_2756 and 2757; up-regulated 40- to 56-fold) shared substantial amino acid sequence similarity to the well characterized FMN-dependent SQ monooxygenase (*SmoC*) and 6-oxo-glucose oxidoreductase (*SmoB*) (78% and 67%, respectively) that mediate the conversion of SQ to sulfite and glucose in the sulfo-SMO pathway [20].

Another *Roseobacter* clade bacterium *R. denitrificans* OCh 114 [60] possesses the same gene contents as *D. shibae* DFL 12 but arranged in the opposite order (Fig. 3), allowing the use of SQ as a sole organic substrate (Fig. S6). Comparative proteomics of *R. denitrificans* OCh 114 grown on SQ, acetate and glucose showed that the two encoded proteins (locus tag, RD1_2310-2311) homologous to SmoC and SmoB in *A. fabrum* C58 were significantly up-regulated (6- to 16-fold) (Figs. 3, 2B; Table. S2). The transcriptional regulator (SmoD, locus tag, RD1_2309) in *R. denitrificans* OCh 114 was only identified during growth on SQ compared to acetate (Fig. 2B; Table. S2). A clear distinction between the gene clusters of these two RCB and *A. fabrum* C58 was the presence of *dctPMQ* that encodes TRAP transporters in *D. shibae* DFL 12 and *R. denitrificans* OCh 114, instead of *smoEFGH* that encodes a putative ABC-type SQ transporter in *A. fabrum* C58 (Fig. 3). Proteins encoded by *dctM* (locus tag, Dshi_2753) and *dctQ* (locus tag, Dshi_2754) were only identified during *D. shibae* DFL 12 growth on SQ (Fig. 2A; Table S2), and DctP (encoded by RD1_2315, up-regulated 1.4- to 24-fold) and DctM (encoded by RD1_2314, not detected in acetate group) of *R. denitrificans* OCh 114 were significantly up-regulated in the SQ group compared to the glucose and acetate groups (Fig. 2B; Table S2). Moreover, the homologues of TRAP transporter genes *dctPMQ* also exist in other organisms that harbor predicted *smoABCD* gene cluster and are adjacent to these SQ catabolic genes (Fig. S7). This information suggests that *dctPMQ* are probably associated with SQ uptake in *D. shibae* DFL 12 and *R. denitrificans* OCh 114 and further characterization will be required. Thus, these two RCB contained the entire set of genes necessary to specify the catabolism of SQ. Together, the gene clusters Dish_2752-2758 in *D. shibae* DFL 12 and RD1_2309-2315 in *R. denitrificans* OCh 114 were likely important for the growth of these two RCB strains on SQ.

To verify the catalytic capacity of the predicted SQ metabolic enzymes of *D. shibae* DFL 12, an in vitro enzymatic assay was performed. SQ was incubated with SmoC, SmoA, FMN, and NADPH. The release of sulfite was detected by Fuchsin assay (Fig. S8) [14]. DNPH derivatization of reaction mixture generated a mass peak with negative ion m/z 357.0688 on LC-MS, which was consistent with DNPH-6-dehydroglucose (Fig. S9) [14], demonstrating the formation of 6-oxo-glucose. Incubation of SmoB with SQ, SmoA, SmoC, FMN and NADPH led to the appearance of $[M + Na]^+$ ion peak at m/z 203.0524, which was identical to the glucose standard (Fig. S10), proving the formation of glucose. A decrease of absorbance at 340 nm over time after incubation of SmoA with NADPH and FMN (Fig. S11) indicated that FMN was catalyzed by SmoA. These results demonstrate that the enzymes encoded by *smoACB* in *D. shibae* DFL 12 indeed convert SQ to glucose.

The other striking result was comparative proteomic profiling of *D. shibae* DFL 12 and *R. denitrificans* OCh 114, which both showed that growth with SQ strongly down regulated three key proteins of the bacterial sulfate assimilation pathway. Sulfate adenylyltransferases (Sat, encoded by Dshi_2656 in *D. shibae* DFL 12 and RD1_B0030 in *R. denitrificans* OCh 114), which transforms sulfate into an organic intermediate, were down-regulated by 3- and 5-fold in *D. shibae* DFL 12 (1- and 3-fold in *R. denitrificans* OCh 114) compared to growth in glucose and acetate, respectively (Table S2). The severely down-regulated phosphoadenylylsulfate reductase (CysH, encoded by Dshi_1152, not detected in SQ-grown *D. shibae* DFL 12; encoded by RD1_2968, down-regulated by 9- and 91-fold in SQ-grown *R. denitrificans* OCh 114) and sulfite reductase (CysI, encoded by Dshi_1153, down-regulated by 16- and 60-fold in *D. shibae* DFL 12; encoded by RD1_2966, down-regulated by 50- and 71-fold in *R. denitrificans* OCh 114) reduced the conversion of organic intermediate product into sulfide, which is then used for the synthesis of sulfur-containing amino acids, while

the expression of the protein involved in last step of cysteine biosynthesis remained stable (Table S2). This regulation pattern was identical in the presence of the phytoplankton metabolite DHPS [17], which is indicative of the close interaction between sulfonate metabolism and sulfate assimilation. When bacteria utilize sulfonate-derived sulfite for sulfur incorporation, it may save energy, at the equivalent of 2-3 ATP per molecule, compared to sulfate assimilation, showing that organosulfonates serve as a more economical sulfur source than sulfate for bacteria. Accordingly, the sulfate assimilation pathway was down-regulated when bacteria were offered reduced sulfur compounds, similar to the manner of sulfur demand-driven regulation in plant sulfate assimilation [61]. In contrast, as the sulfite anion is a reactive and toxic compound that can cause damage to vital cell components, such as DNA and protein, its level must be carefully controlled [62]. Bacterial sulfite oxidases were up-regulated in the presence of both SQ (Fig. 2) and DHPS catabolism [17], as a means of generating energy via sulfite oxidation but also for detoxifying excess amounts of sulfite.

The distribution of sulfo-SMO pathway in RCB

In contrast to *D. shibae* DFL 12 and *R. denitrificans* OCh 114, *R. pomeroyi* DSS-3 [63] failed to utilize SQ as a sole carbon source (Fig. S12), and no such SQ catabolic genes were found within its genome. Besides *D. shibae* DFL 12 and *R. denitrificans* OCh 114, around 1/3 of currently sequenced marine RCB (recently reclassified as *Roseobacteraceae* of the *Rhodobacterales*) [64] genomes (75 of 248 genomes) contained at least three genes that encode SmoABC enzymes required for complete SQ degradation (Fig. 4). The genetic capability for SQ degradation was also present in a limited number of marine relatives of the alphaproteobacterial *Rhodobacteraceae* of the *Rhodobacterales* (11 species), *Rhizobiales* (10 species), the SAR116 clade (2 species) and three gammaproteobacterial species (Fig. 4).

Marine bacteria are able to catabolize the ubiquitous algal metabolites DMSP and DHPS via DMSP demethylation (*dmdA*-mediated) [65] and three DHPS dehydrogenation reactions (encoded by *hpsN*, *hpsO* and *hpsP*) [17], respectively. These pathways have been found among a wider range of bacterial taxa than SQ degradation, with genes encoding proteins experimentally verified in the *Roseobacter* clade [17] and the SAR11 clade [66, 67]. The majority of candidate SQ-degrading RCB (61 of 75 strains) had predicted *hpsN* genes. More than half of the candidate SQ-degrading RCB contained predicted *dmdA* (51 of 75 strains) and (54 of 75 strains) harbored at least one predicted sulfonate desulfonation pathway that involved the metabolism of sulfolactate, isethionate and cysteate (mediated by *suyAB*, *xsc* and *cuyA*, respectively) (Fig. 4; Table S3). The ability to catabolize multiple naturally occurring organosulfur species might provide an advantage for RCB by allowing them to obtain a steady source of reduced sulfur in various marine environments. However, a total of 11 candidate SQ degraders lacked the capabilities of DMSP demethylation, DHPS dehydrogenation and sulfonates (sulfolactate, isethionate and cysteate) desulfonation (Table S3), suggesting that SQ is a resource that cannot be ignored for members of the *Roseobacter* clade.

Metatranscriptomes

5We further investigated the expression patterns of *smoC*, which encodes an enzyme for SQ desulfonation, across the *Tara* Oceans stations and Bohai Sea. All 186 of the available metatranscriptomic samples from the *Tara* Oceans contained *smoC* transcripts, showing a widespread occurrence of gene expression (Table S4). *Tara* Oceans' *smoC* comprised on average 0.001% of the total prokaryotic metatranscriptomic reads, with levels almost one order of magnitude lower than that of the

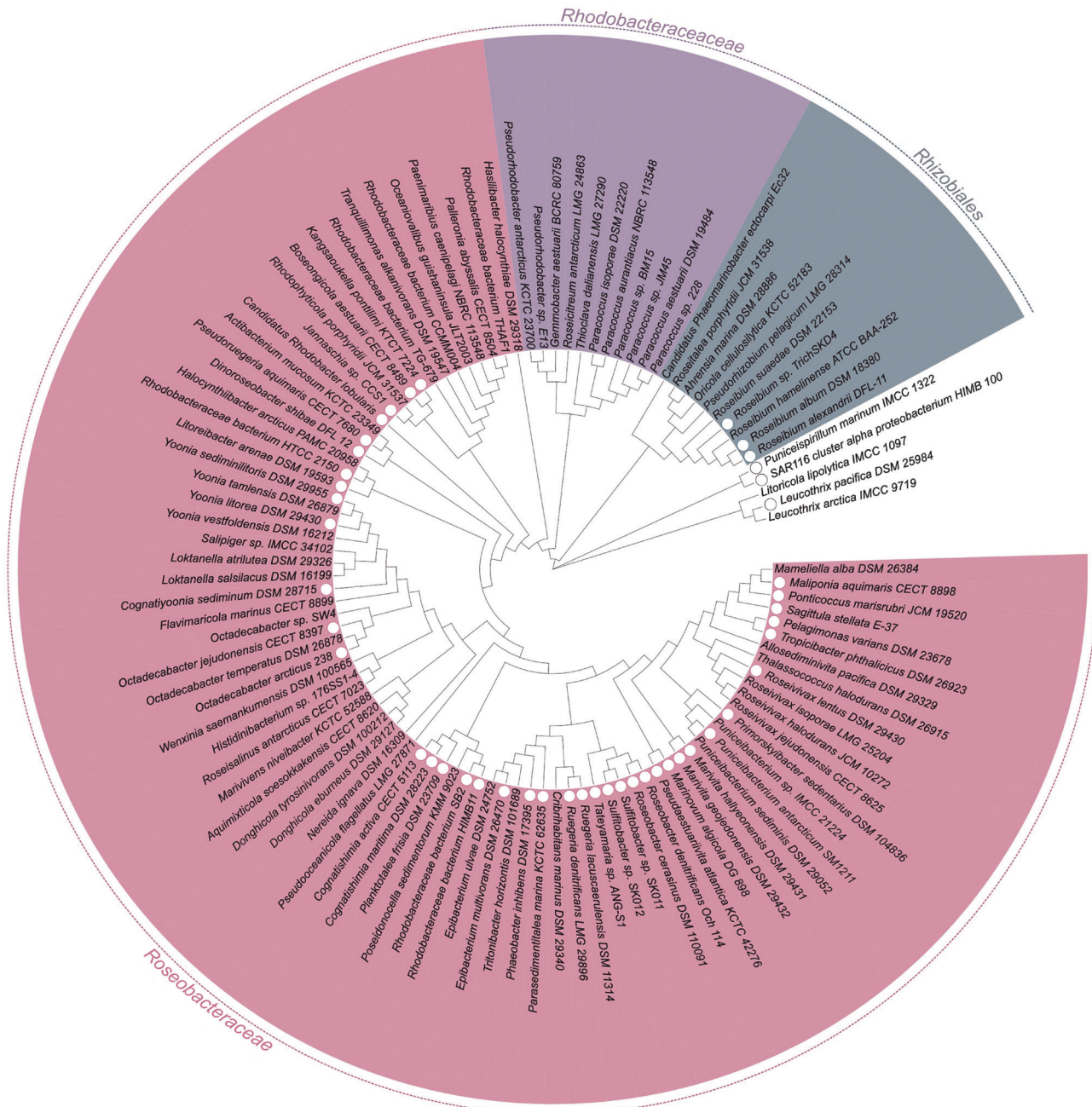


Fig. 4 The distribution of predicted sulfo-SMO SQ metabolic genes in publicly available marine bacterial genomes. The tree was based on the 39 single-copy orthologous genes shared by bacterial strains that possess predicted *smoABC*. The different colors represent different taxa. ■, *Roseobacteraceae*; ■, *Rhodobacteraceae*; ■, *Rhizobiales*. White dots indicate that the species have the genetic potential of metabolising not only SQ, but also harbor the candidate genes for complete DHPs degradation pathways.

dmdA gene (on average 0.023%) and *hpsN* gene (on average 0.013%) which encodes a key enzyme for DHPs catabolism (Table S4) [16]. We also found wide expression of *smoC* from coastal samples in the Bohai Sea (Fig. 5; Table S6). Similar to *Tara* Oceans, *smoC* transcripts was on average 0.001% of the total prokaryotic transcripts in Bohai Sea, and the reads of *hpsN* and *dmdA* were 2- to 41-fold and 5- to 42-fold higher than that of *smoC*, respectively (Table S6). *dmdA* and *hpsN* are not only highly abundant organosulfur compound degrading genes in the marine environment, but are also widespread and expressed among diverse marine bacterial taxa including the *Roseobacter* clade, SAR11 and *Gammaproteobacteria*. Similarly, the

abundance of *smoC* transcripts in each sample was also significantly lower than those of genes involved in the desulfonation of sulfoacetaldehyde (*xsc*) and sulfolactate (*suyB*) that are known to be vital components of a sulfonate-based network between phytoplankton and heterotrophic bacteria ($p < 0.0001$, one-way analysis of variance (ANOVA) (Fig. 5) [67]. Those genes were identified within members of the *Alpha*-, *Gamma*- and *Deltaproteobacteria* [7], representing a wider range of taxonomic origins than *smoC*. In most (144 of 186 samples) *smoC*-containing metatranscriptomic samples of the *Tara* Oceans project, the majority of *smoC* transcripts were from members of the *Roseobacter* clade (Table S4), and the majority of

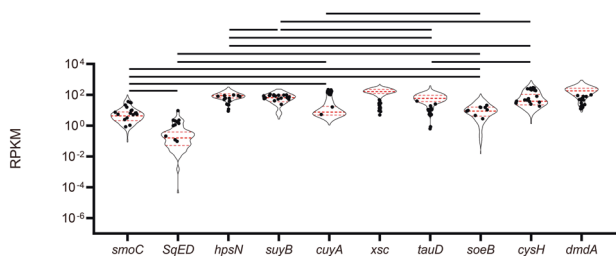


Fig. 5 Expression levels of genes involved in the metabolism of sulfonates, sulfite, DMSP, and sulfate in the Bohai Sea and Tara Oceans data sets. Black dots represent the data from Bohai Sea. Relative abundance of genes transcripts in each sample was estimated with RPKM (reads per kilobase per million mapped reads). *smoC*, FMN-dependent SQ monooxygenase; *SqED*, SQ dehydrogenase; *hpsN*, DHPS dehydrogenase; *suyB*, sulfolactate sulfolylase; *cuyA*, L-cysteate sulfolylase; *xsc*, sulfoacetaldehyde acetyltransferase; *tauD*, taurine dioxygenase; *soeB*, sulfite oxidase; *dmdA*, DMSP demethylase; *cysH*, phosphoadenylylsulfate reductase. Statistical differences in relative abundance by analysis of variance (ANOVA). Black lines connect genes that are not significantly different, otherwise, significantly different ($p < 0.001$).

the total *smoC* transcripts in each sample from Bohai Sea belonged to taxonomic relatives of the *Roseobacter*, suggesting that RCB were the predominate microorganisms carrying and actively expressing *smoC* in the ocean. This taxonomically limited distribution pattern in marine SQ catabolism suggests that SQ might be used for targeted feeding of specific RCB and enhance associations between ubiquitous RCB and phytoplankton especially in the SQ-abundant phycosphere. The homologous genes for other SQ degradation pathways (sulfo-EMP, sulfo-EMP2, sulfo-ED, sulfo-transaldolase and sulfo-transketolase) were not currently found in marine cultivated and uncultivated bacterial genomes, except that the five candidate genes for the sulfo-ED pathway were simultaneously detected in few bacterial genomes from Tara Oceans (Table S7). The transcripts abundance of the predicted *SqED* gene were on average lower than that of *smoC* in Tara Oceans by almost one order of magnitude (Fig. 5; Table S4). These results indicate that the sulfo-SMO pathway and its primary carrier RCB currently appear to be the most prevalent known SQ metabolic pathway and SQ consumers, respectively, in the marine pelagic environment.

The highest levels of *smoC* abundance and expression were detected at the deep chlorophyll maximum layer of the Arctic Ocean (Table S4). Both *dmdA* and *hpsN* transcript abundances displayed significant positive correlation with chlorophyll *a* concentrations, as was *smoC* (Spearman's $r = 0.506$, $p < 0.001$), while *cysH* showed a negative correlation ($r = -0.438$, $p < 0.001$) (Fig. 6), suggesting that marine phytoplankton provided exogenous reduced sulfur for bacterial assimilation. Furthermore, *dmdA* and *hpsN* transcript abundances were both positively correlated with diatoms, haptophytes and chlorophyta, while *cysH* displayed an inverse correlation (Fig. 6). Similarly, the expression level of *smoC* exhibited a strong positive correlation with diatoms ($r = 0.582$, $p < 0.001$) and chlorophyta ($r = 0.595$, $p < 0.001$) and a weak positive correlation with haptophytes ($r = 0.149$, $p < 0.01$) (Fig. 6). Diatoms are particularly prevalent at high latitudes and in upwelling zones [68], while haptophytes, such as the cosmopolitan *Phaeocystis*, are the main primary producers in polar marine environments [69]. Higher levels of *smoC* transcript abundance were observed in the polar waters of the Arctic and Southern Oceans, and the coastal waters of Chile-Peru Current compared to other Tara Oceans stations (Fig. 7; Table S4), indicating a clear

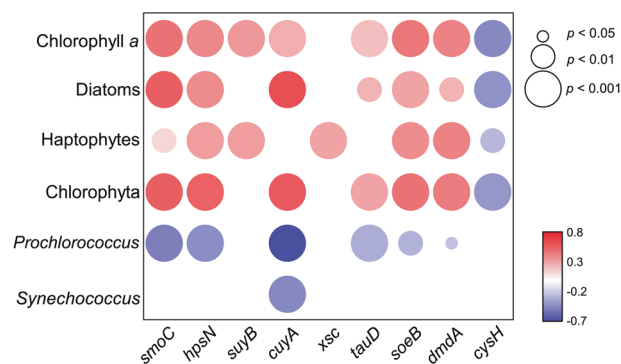


Fig. 6 Correlation analysis among relative abundance of genes transcripts, selected phytoplankton taxa and chlorophyll *a*. *smoC*, FMN-dependent SQ monooxygenase; *hpsN*, DHPS dehydrogenase; *suyB*, sulfolactate sulfolylase; *cuyA*, L-cysteate sulfolylase; *xsc*, sulfoacetaldehyde acetyltransferase; *tauD*, taurine dioxygenase; *soeB*, sulfite oxidase; *dmdA*, DMSP demethylase; *cysH*, phosphoadenylylsulfate reductase. Color represents the Spearman's r . Circle size represents statistical significance ($p \geq 0.05$).

regional pattern for SQ catabolism. Moreover, RCB from high latitudes (60–90 °N and 60–90 °S) contributed a higher proportion of the *smoC* gene transcripts (average 80% of *smoC* transcripts) than that from low latitudes (30 °S–30 °N) (average 60% of *smoC* transcripts) (Table S5), suggesting the relatively higher bacterial catabolism potential for SQ in the polar pelagic ocean.

Cultured plankton and environmental metabolomics

The metabolite SQ was present at micromolar to millimolar cellular concentrations in all pure cultures of phytoplankton and cyanobacteria species investigated in this study, showing that SQ has both marine eukaryotic and bacterial origins (Table 1). The diatom *Navicula pelliculosa* was a high SQ producer with a median intracellular concentration in excess of 1 mM, which was comparable to the concentrations of the two cytosolic structural analogues sulfolactate and DHPS (0.9 ± 0.5 mM and 6.6 ± 1.3 mM, respectively). The other diatoms *Thalassiosira pseudonana* (44.5 ± 13.0 μ M) and *Phaeodactylum tricornutum* (62.8 ± 19.5 μ M) maintained low intracellular concentrations of SQ, which were around 153- and 128-fold lower than cytosolic DHPS concentrations, respectively. Sulfolactate also co-occurred with SQ and DHPS in the cytosol of *T. pseudonana* (113.9 ± 23.6 μ M) and *P. tricornutum* (70.6 ± 15.3 μ M) with SQ concentrations showing similar levels (Table 1). SQ production (97–527 μ M) in the three haptophyte *Emiliania huxleyi* were lower 22- to 87-fold than cytosolic DHPS concentrations, which were more than the co-occurred isethionate concentrations by approximately one order of magnitude (Table 1). The sulfolactate cytosolic concentrations (4–15 μ M) were similar to the levels of isethionate cytosolic concentrations (12–24 μ M) in dinoflagellates (Table 1), which contained substantially lower micromolar concentrations of SQ than the reported hundreds of millimolar concentrations of the organosulfur compound DMSP in these organisms [70, 71]. Sulfolactate and isethionate were also detected in chlorophyta and cyanobacteria, all of which could produce SQ. In conjunction with the active sulfonate conversions in the surface ocean [67], SQ is likely an integral part of the sulfonate network in marine algae. Dissolved SQ was detected in the culture supernatants from all of these photosynthetic plankton with concentrations of up to 31.2 ± 10.9 μ M in diatom culture supernatant and 80.2 ± 65.7 μ M in *Prochlorococcus* culture supernatant. These findings suggested that

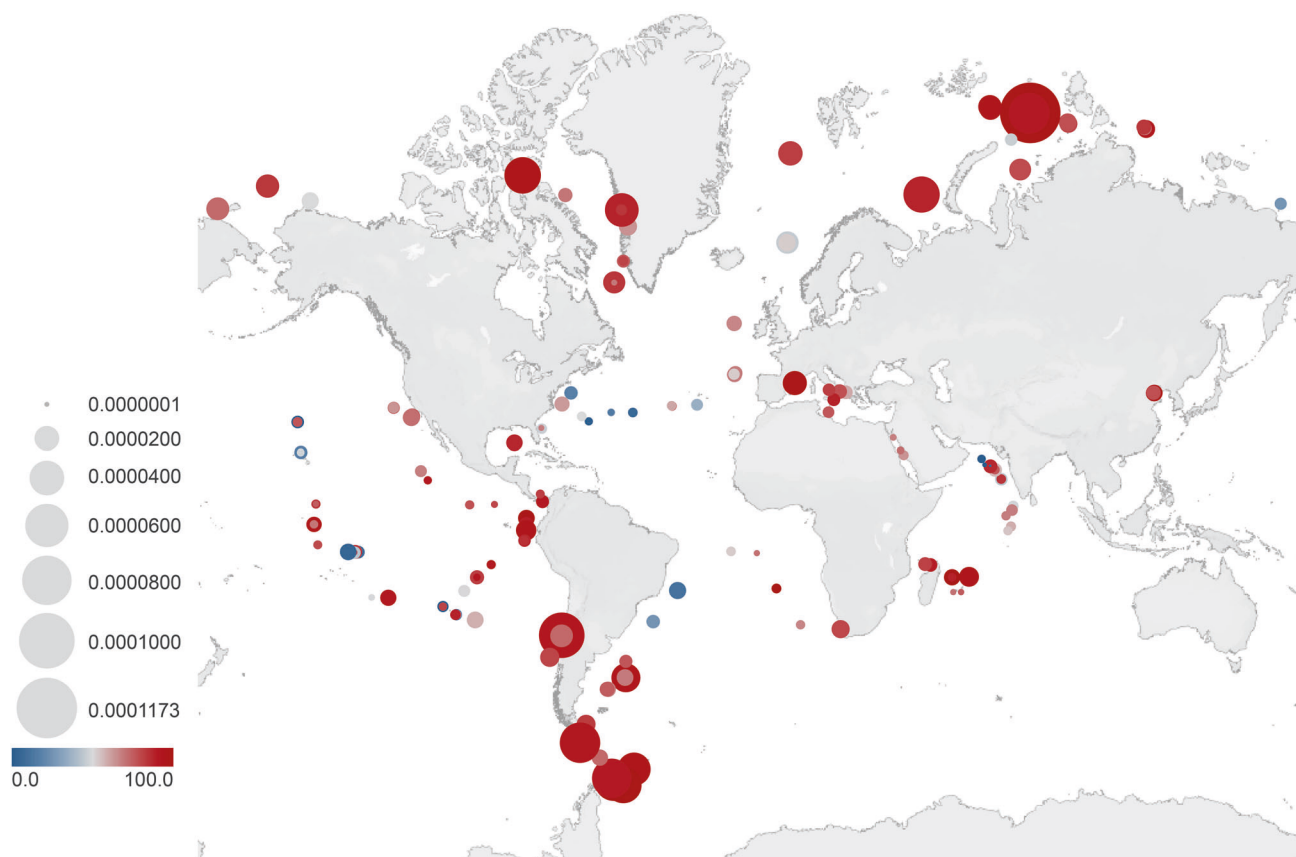


Fig. 7 *smcC* transcripts are geographically widespread throughout the oceans. The size of the circle represents the relative abundance of *smcC* transcripts in each sample. Color bar represents the percent of *smcC* transcripts from RCB.

phytoplankton and cyanobacteria mediated SQ production could be a common and important process in marine surface waters. Consistent with this hypothesis, SQ was detected at multiple coastal sites in the Bohai Sea. Particulate (size $> 3 \mu\text{m}$) SQ concentrations were up to $1.90 \pm 0.76 \mu\text{mol/g}$ POC with an average of $0.88 \pm 0.20 \mu\text{mol/g}$ POC in the water column (Fig. 8), which were ~ 5 -fold lower than DHPS ($8.43 \pm 4.37 \mu\text{mol/g}$ POC) and ~ 10 -fold lower than DMSP ($10.46 \pm 6.95 \mu\text{mol/g}$ POC) on particles that represent the major fraction of total DMSP in most sea waters [72]. Particulate sulfolactate ($0.40 \pm 0.36 \mu\text{mol/g}$ POC) and isethionate ($0.48 \pm 0.17 \mu\text{mol/g}$ POC) were both detected in these coastal sites at averages of 29.6% and 64.0% of the particulate SQ concentrations, respectively (Fig. 8). Dissolved SQ in natural seawater ($0.2 \pm 0.1 \text{ nM}$) was detected (Fig. S13), which was approximately 4-fold lower than the DHPS concentration in the dissolved organic matter pool ($0.7 \pm 0.5 \text{ nM}$) [16], suggesting that unbound SQ was released into seawater by photosynthetic organisms.

SQ concentrations were lower by one order of magnitude compared to DMSP in field samples (Fig. 8). Based on DMSP production that exceeds 10^3 Tg per year in the oceans, the annual particulate SQ production was estimated at teragram amounts. We generated a crude estimate of the annual SQ production in marine cyanobacteria at about 1.3 Tg , based on the total number of cyanobacterial cells (approximately *Prochlorococcus* 3×10^{27} cells and *Synechococcus* 7×10^{26} cells, respectively) [73], in conjunction with the SQ contents of single *Prochlorococcus* and *Synechococcus* cells in the marine environment, which were $1.7 \pm 1.0 \text{ amol cell}^{-1}$ and $0.5 \pm 0.1 \text{ amol cell}^{-1}$, respectively.

The amount of SQDG was higher than SQ by two orders of magnitude in *T. pseudonana*, making up $\sim 9\%$ of the cellular sulfur pool [67], suggesting that phytoplankton contained considerable amounts of SQ in a combined form. Normally, the SQDG degradation processes occurs in the gut and soil environments through glucoside hydrolase from bacteria, and also possibly from fungal, protist and animal origins [4]. Marine SQDG decomposition can occur as particles sink to deeper waters [74]; however, no defined enzyme has yet been linked to the conversion of SQDG to SQ in the ocean. Meanwhile, our results showed that marine algae and cyanobacteria could directly produce SQ and release it into the environment. RCB have no homologs to known SQDG degradation enzymes, but after release of photosynthetic organism-derived SQ into the seawater through viral lysis and active/passive exudation, it can be turned over by RCB through the monooxygenase pathway. We cannot exclude the possibility of the direct uptake and catabolism of SQDG in marine bacterial cells as bacteria can cleave complex carbohydrate compounds through the use of diverse glycosidases [75]. It has been shown that marine cyanobacteria and phytoplankton show a decline in phospholipids with a marked accumulation of SQDG in phosphorus-depleted environments [2, 3]. SQDG was often observed throughout the entire water column in the oligotrophic ocean ranging from 1–4% of organic carbon in sinking particles [74], and in some cases was comparable to the contribution of DMSP [72, 76, 77]. Given this high proportion of SQDG in marine phototrophic organisms and the role that they play in carbon and sulfur export, SQ transformation may be of global significance in the ocean.

Table 1. Analysis of sulfonate metabolites in pure cultures of various plankton.

Species	SQ (μM)	SQ (amol cell^{-1})	DHPS (mM)	DHPS (amol cell^{-1})	Sulfolactate (μM)	Sulfolactate (amol cell^{-1})	Isethionate (μM)	Isethionate (amol cell^{-1})
Diatoms								
<i>Thalassiosira pseudonana</i> 220	44.5 ± 13.0	2.2 ± 0.6	6.9 ± 0.1	344.5 ± 5.0	113.9 ± 23.6	5.7 ± 1.2	—	—
<i>Navicula pelliculosa</i> 350	1055.3 ± 508.1	52.8 ± 25.4	6.6 ± 1.3	328.6 ± 67.4	874.5 ± 549.2	43.7 ± 27.5	—	—
<i>Phaeodactylum tricornutum</i> 267	62.8 ± 19.5	5.8 ± 1.8	8.0 ± 1.3	736.3 ± 120.8	70.6 ± 15.3	6.5 ± 1.4	—	—
Haptophyta								
<i>Emiliania huxleyi</i> CS369	527.4 ± 87.3	21.1 ± 3.5	12.0 ± 2.9	479.4 ± 114.5	1139.3 ± 248.2	45.6 ± 9.9	106.3 ± 23.5	4.3 ± 0.9
<i>E. Huxleyi</i> 2090	97.4 ± 42.6	3.9 ± 1.7	8.5 ± 2.2	339.1 ± 87.5	—	—	17.7 ± 10.6	0.7 ± 0.4
<i>E. Huxleyi</i> 1516	372.6 ± 142.2	14.9 ± 5.7	8.3 ± 1.7	332.6 ± 67.2	41.5 ± 13.4	1.7 ± 0.5	12.9 ± 3.9	0.5 ± 0.2
<i>Isochrysis galbana</i> 079	195.6 ± 25.4	12.7 ± 1.7	—	—	—	—	5.8 ± 2.4	0.4 ± 0.2
<i>I. galbana</i> 321	371.0 ± 75.3	24.1 ± 6.0	—	—	—	—	2.2 ± 1.0	0.1 ± 0.08
<i>Phaeocystis globosa</i> 124	42.7 ± 22.1	4.4 ± 2.3	—	—	—	—	—	—
Dinoflagellates								
<i>Amphidinium carterae</i> Hulbert 279	71.3 ± 14.4	25.4 ± 5.1	—	—	3.5 ± 1.7	1.3 ± 0.6	12.2 ± 1.6	4.4 ± 0.6
<i>Alexandrium tamarense</i> 260	9.7 ± 3.2	114.8 ± 38.7	—	—	14.9 ± 8.0	177.0 ± 95.4	23.8 ± 2.4	283.5 ± 28.7
Chlorophyta								
<i>Chlorella</i> sp. 309	89.7 ± 17.2	8.1 ± 1.5	—	—	601.0 ± 294.7	54.1 ± 26.5	—	—
<i>Chlorella</i> sp. 410	23.3 ± 8.6	2.1 ± 0.8	—	—	—	—	433.4 ± 66.0	39.0 ± 6.0
Cyanobacteria								
<i>Prochlorococcus</i> sp. 9301	4303.7 ± 1631.1	0.8 ± 0.3	—	—	—	—	612.4 ± 72.5	0.11 ± 0.01
<i>Prochlorococcus</i> sp. 9312	27783.5 ± 2796.3	5.4 ± 0.7	—	—	143.7 ± 40.5	0.026 ± 0.007	—	—
<i>Synechrococcus</i> sp. XM24	85.0 ± 4.5	0.060 ± 0.003	—	—	3.4 ± 1.1	0.003 ± 0.001	—	—
<i>Synechrococcus</i> sp. 1333	249.9 ± 50.8	0.2 ± 0.04	—	—	—	—	2.4 ± 0.7	0.002 ± 0.001
<i>Synechrococcus</i> sp. YX04-3	1543.1 ± 512.9	1.1 ± 0.4	—	—	2.2 ± 1.5	0.002 ± 0.001	5.9 ± 1.8	0.004 ± 0.001

The values indicate the estimated cytosolic concentrations of endo-metabolites from biomass. Values are means ± standard errors of the means. —, not detected.

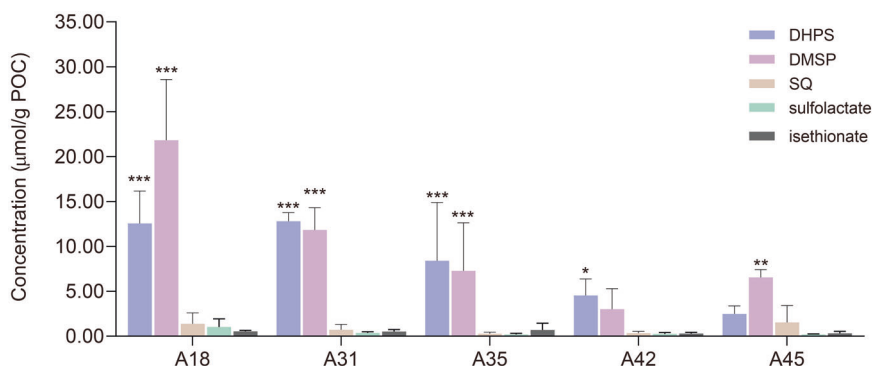


Fig. 8 Particulate concentrations of SQ, DHPS, DMSP, sulfolactate and isethionate (3–200 µm) from selected sites of the Bohai Sea. Sites A18, A31, A35, A42 and A45 were analyzed. The organosulfur concentrations were normalized by the concentrations of particulate organic carbon (POC). Error bars represent the standard deviation ($n = 3$). Asterisks indicate significant difference between SQ and different compounds (Two-way ANOVA; * $p < 0.05$; ** $p < 0.01$; *** $p < 0.001$).

CONCLUSION

Since the identification of the first SQ-degrader thirty years ago, it has been shown that SQ catabolism is widely present in various environments; however, little known is about this process in the ocean. Our study revealed that marine RCB, such as *D. shibae* DFL 12 and *R. denitrificans* OCh 114, not only used the recently identified monooxygenase pathway to catabolize SQ as carbon and energy source, but also the reduced sulfur for assimilation. This pathway was associated primarily with the members of the *Roseobacter* clade. The ubiquitous gene expression for SQ desulfonation throughout the global ocean indicates this molecule likely plays an important and yet overlooked role in the marine sulfur cycle. Furthermore, SQ was found to be widespread in marine phytoplankton and cyanobacteria. The prevalence of SQ in the marine environment and the ability of SQ metabolism in specialized species highlight the importance of RCB as major players in ocean biogeochemical organosulfur transformations.

DATA AVAILABILITY

Proteomic data have been uploaded to the National Omics Data Encyclopedia (NODE, <https://www.biosino.org/node/>) database with the accession number OEP003145.

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AUTHOR CONTRIBUTIONS

KT conceived the study, analyzed data, and wrote the manuscript. LL performed the experiments, analyzed data, wrote and edited the manuscript. XC sampled, performed the experiments, analyzed data, and wrote the manuscript. JY performed the experiments. XM performed the experiments. YuH sampled, and analyzed data. YajieH performed the experiments. All authors have approved the submitted final version.

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COMPETING INTERESTS

The authors declare no competing interests.

ADDITIONAL INFORMATION

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