Sulfur bacteria promote dissolution of authigenic carbonates at marine methane seeps

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Abstract
Carbonate rocks at marine methane seeps are commonly colonized by sulfur-oxidizing bacteria that co-occur with etch pits that suggest active dissolution. We show that sulfur-oxidizing bacteria are abundant on the surface of an exemplar seep carbonate collected from Del Mar East Methane Seep Field, USA. We then used bioreactors containing aragonite mineral coupons that simulate certain seep conditions to investigate plausible in situ rates of carbonate dissolution associated with sulfur-oxidizing bacteria. Bioreactors inoculated with a sulfur-oxidizing bacterial strain, Celeribacter baekdonensis LH4, growing on aragonite coupons induced dissolution rates in sulfidic, heterotrophic, and abiotic conditions of 1773.97 (±324.35), 152.81 (±123.27), and 272.99 (±249.96) μmol CaCO3 • cm⁻² • yr⁻¹, respectively. Steep gradients in pH were also measured within carbonate-attached biofilms using pH-sensitive fluorophores. Together, these results show that the production of acidic microenvironments in biofilms of sulfur-oxidizing bacteria are capable of dissolving carbonate rocks, even under well-buffered marine conditions. Our results support the hypothesis that authigenic carbonate rock dissolution driven by lithotrophic sulfur-oxidation constitutes a previously unknown carbon flux from the rock reservoir to the ocean and atmosphere.

Introduction
The atmospheric flux of oceanic methane is mitigated through the sulfate-dependent anaerobic oxidation of methane (AOM), which serves as the dominant sink for methane in the oceans [1]. Approximately 10–20% of methane oxidized by AOM is incorporated into authigenic mineral deposits [2–5], which are estimated to account for 11–15% of the 11–15 Tmol•year⁻¹ carbonate accumulation estimated for continental shelf sediments [6]. Seafloor authigenic carbonate deposits sequester carbon in what is thought to be a geologically stable phase. However, modern and ancient methane seep carbonates frequently exhibit signs of dissolution on the seafloor such as pits, holes, and micro-etchings of exposed surfaces, which have been suggested to be the result of aerobic methane oxidation, sulfur oxidation, or the combined effect of both [7–11]. These altered surfaces commonly co-occur with bacterial mats at the sediment/water interface that contain sulfur-oxidizing bacteria. Sulfur-oxidizing bacteria predominantly oxidize hydrogen sulfide (H₂S) or thiosulfate (S₂O₃²⁻) completely to sulfate (SO₄²⁻), or to intermediates that can be further oxidized to sulfate, although other possibilities exist [11–13]. Complete sulfide oxidation to sulfate with O₂ is an acid-producing reaction:

$$\text{H}_2\text{S} + 2\text{O}_2 → \text{SO}_4^{2-} + 2\text{H}^+$$  (I)

Likewise, thiosulfate oxidation to sulfate with O₂ is an acid producing reaction that consumes, and produces, equal
mole\ of\ O_2,\ and\ H^+,\ respectively,\ as\ reaction\ I:
\[
S_2O_3^{2-} + H_2O + 2O_2 \rightarrow 2SO_4^{2-} + 2H^+ \tag{II}
\]

Acid production by sulfide oxidation has been shown to mediate carbonate dissolution in freshwater systems including caves and aquifers [14–16], and in the built environment [17–19]. However, the potential for sulfur oxidation to induce dissolution of carbonates in marine systems that are well-buffered against bulk acidification has not been investigated.

Through rRNA gene sequencing of mat samples collected from the Del Mar East Methane Seep Field, we show that abundant and diverse sulfur-oxidizing bacteria colonize carbonate surfaces. We then use biofilm experiments to compare the dissolution rates of carbonate coupons under abiotic conditions with those that occur in the presence of biofilms of sulfur-oxidizing bacteria in order to estimate plausible dissolution rates in nature. pH gradients within mineral-attached biofilms were then measured using laser scanning confocal microscopy to assess a potential mechanism of dissolution in which acid generated by sulfur oxidation overcomes buffering within the biofilm microenvironment. Finally, we use our experimentally-derived dissolution rates to estimate the potential impact of seafloor weathering of carbonates by sulfur-oxidizing bacteria on global carbonate rock reservoirs.

Materials and methods

Del Mar methane seep carbonate community analyses

The carbonate rock sample used for microbial community analyses was collected from Del Mar East Seep, Dive SO 177, (32°54.25456764 N, 117°46.9408327 W) at a depth of 1032.06 m. The site is in the northern portion of the San Diego Trough, about 50 km west of San Diego, California. Visible features of the seep include carbonate boulders and pavements colonized by orange and white bacterial mats, clam beds, and curtains of methane bubbles [20]. For a more in-depth discussion of the Del Mar Methane Seep, please see Grupe et al., (2015). A small chunk of the rock was sealed in a Mylar bag and shipped on dry ice to the University of Minnesota (Twin Cities). Upon arrival, the carbonate sample was temporarily stored at −80 °C until sampling for DNA extractions. All tools used for sampling biomass were autoclave-sterilized prior to use, and all work was performed in a LabConco A2 biological safety cabinet. Sterile aluminum foil was placed over ice packs to provide a cold and clean working environment. Three tubes were designated for the top of the rock, and three tubes were designated for the bottom of the rock. Biomass was scraped from the respective positions and placed in tubes. Top sample tubes had a biomass weight between 76.5–97.6 mg, and bottom sample tubes had a biomass weight between 30.7–59.3 mg. DNA was extracted from the seep samples using the ZymoBIOMICS DNA miniprep kit (Zymo, Irvine, CA). Nuclease-free water from the kit was processed alongside the samples as a negative control for iTag sequence analyses. The amplification of DNA and the generation iTag libraries of the V4 hypervariable region was performed by the University of Minnesota Genomics Center as previously described [21]. The samples and negative control were sequenced on ¼ of a lane of MiSeq for paired end 2x300 bp reads. Primers and adapters were removed with Cutadapt v. 2.10 [22]. The paired end reads were processed and assembled using DADA2 v1.16.0 [23]. The maximum expected error rate was 2 and reads detected as phiX were removed prior to error detection, merging of pairs and chimera detection. Taxonomic assignment was performed using the Silva database v. 138 [24]. Bioinformatic and statistical analyses was performed using tools in PhyloSeq 1.32.0 [25]. R package Decontam 1.8.0 [26] identified contaminating amplicon sequence variants (ASVs). Thus, ASVs of the genera *Escherichia/Shigella*, *Haemophilus*, *Streptococcus*, and the clade Chloroflexi S085 were bioinformatically removed from analyses. ASVs with statistically different abundances between the top and bottom surfaces were detected with DESeq2 v. 1.28.0.

Scanning electron microscopy of seep carbonates

The carbonate rock sample used for SEM imaging was collected from the Lasuen Knoll Seep (Dive SO 170, 33° 23.57489996 N, 118°0.39814252 W) at a depth of 279.705 meters. A small chunk of carbonate rock with filtered bottom water and 25% vol/vol glutaraldehyde was stored at 4 °C, shipped on ice to the University of Minnesota (Twin Cities), and then stored at 4 °C. Several small pieces of the carbonate rock were broken off and placed in an 8-well plate. The carbonate pieces in the 8-well plate were rinsed, and subjected to an ethanol dehydration series as follows: 50% for 2 h, 70% overnight, 80% for 15 min, 95% (x2) for 15 min, and 100% (x2) for 15 min. Carbonate samples were then subjected to critical point drying on a Tousimis Model 780 A Critical-Point-Dryer following standard procedures, followed by sputter coating 1–2 nm Iridium in a Leica ACE600 Sputter Coater. Lastly, carbonate samples were visualized at 1.0 kV on a Hitachi SU8230 Field Emission Gun Scanning Electron Microscope.

Continuous flow bioreactor experiments

To investigate the potential for sulfur-oxidizing bacteria to dissolve carbonate minerals, we used flow-through biofilm
reactors (herein referred to as bioreactors: CDC; CBR 90–3 CDC Biofilm Reactor). Bioreactors contained eight polypropylene coupon holder rods that each accommodate three 12.7 mm diameter coupons. The lid and coupon holder rods were mounted in a 1 L glass vessel with side-arm discharge port at ~400 mL. A liquid medium (described below and Supplementary Table 1) was circulated through the bioreactor, while mixing was generated by a magnetic stir bar at 80 RPM. Sampling of the coupons was conducted aseptically by removing individual coupon holders and harvesting the coupons, while replacing the removed coupon holder with a sterile rubber plug. Experiments were 21 days in duration. On the final day of bioreactor experiments, one coupon was transferred to a treatment imaging flow cell (herein referred to as flow cell: BioSurface Technologies; Model FC 310 Treatment Imaging Flow Cell) in order to visualize and measure the in vivo pH of the biofilm. The flow cell is an autoclavable polycarbonate plastic cell with a recessed inner circle for installing coupons with biofilms grown in CDC Biofilm Reactors, and is equipped with barbed influent and effluent connectors, and topped with a coverslip for in vivo imaging.

The bioreactor medium consisted of two solutions, salt solution 1 & 2, and six additional components: vitamin solution, trace element solution, yeast extract, sodium bicarbonate, sodium thiosulfate, and sodium metasilicate. One 10 L carboy was prepared by autoclaving 5 L of salt solution 1 and one 10 L carboy was prepared by autoclaving 4.1 L (with an additional 839 mL ddH2O) of salt solution 2 at 121 °C for 80 min. The salt solutions were then aseptically pumped together, followed by the addition of sterile stocks of 10 mL 1000X vitamin solution, 10 mL 1000X trace element solution, 10 mL (1000 g/L) yeast extract, 11 mL 1 M NaHCO3, 10 mL 1 M Na2S2O3, and 20 mL 100 mM Na2SiO3 through a 0.2 μm filter.

The final medium had a pH of ~7.85 and contained approximately 1.1 mM HCO3−, 1 mM S2O32−, and 9.14 mM Ca2+, with a saturation state with respect to aragonite (Ωaragonite) of ~0.5, similar to that measured near the sediment/water interface in certain seep environments where carbonate dissolution was observed to occur [7, 27]. The saturation state of aragonite was calculated using CO2SYS_v2.1 [28]. Ingredients for both salt solutions and the additional components added to the medium are presented in Supplementary Table 1. Measured saturation indices throughout bioreactor experiments are presented in Supplementary Table 2.

Mineral preparation for bioreactor experiments

Bulk mineral specimens of aragonite were obtained from D. J. Minerals (Butte, Montana). X-ray powder diffraction (XRD) of the aragonite sample, and peak matching indicates the mineral is nearly 100% aragonite. Samples used for the bioreactor were cored with a diamond coring bit on a drill press, cut to ~1–3 mm thickness and smoothed with a diamond saw. Coupons were then scored with a diamond tipped scribe pen to assign a reference code. After scoring, coupons were placed in a muffle furnace (NEY 2-525 Series II) at 500 °C for 4 h to remove residual organic matter, followed by weighing to 0.01 mg (CAHN 29 Automatic Electrobalance) and stored in a desiccator until being mounted in coupon holder rods.

Assembling the bioreactor

Weighed coupons were mounted in coupon holder rods, and fastened in place with plastic screws to avoid mineral chipping, as occurred with the stock metal screws. The bioreactor lid was equipped with a bubble trap on the media inflow port, a 0.45 μm airport, and a lure-lock covered in foil for inoculating via syringe. The entire bioreactor was autoclaved at 121 °C for 45 min.

Microbial inoculant

*Celeribacter baekdonensis* strain LH4 was used as the pure-culture inoculant for the bioreactor experiments. *C. baekdonensis* strain LH4 is a colorless, chemolithoheterotrophic, sulfur-oxidizing bacterium, belonging to the Rhodobacterales within the Alphaproteobacteria. It was isolated from marine sediments collected from a methane seep/brine pool at Green Canyon Block 233 [29] (Lat/Long: 27° 43.4392’ N, 91° 16.7638’ W, Depth: 648 m) in the Gulf of Mexico. Strain LH4 produces acid from thiosulfate oxidation, likely via the sox pathway (soxA/B/C/D/XY/Z) [30]. Strain LH4 also contains other genes related to sulfur oxidation, including four copies of sulfide quinone oxidoreductase ORFs and three copies that encode flavocytochrome c oxidases. Strain LH4 was grown up in the final medium, centrifuged at 10,000 G for 10 min, and washed 2X. OD590 for experiments was on average ~0.07 (~2.3×108 cells/mL based on cell counts and growth curves yielding the equation y = 3×109(x) + 2×107 where y equals cells/mL, and x equals OD590) with an inoculum volume of 20 mL.

Bioreactor experiments

Experiments were run to measure the dissolution of aragonite in a medium with a saturation state of 0.5 with respect to aragonite, similar to that measured near the sediment/water interface in seep environments [27]. Unlike in the sediments where AOM-generated alkalinity is high and carbonate actively precipitates, at the sediment/
water interface previous studies have measured a notable drop in alkalinity [11, 31–36] and carbonate dissolution was observed to occur [7–9, 11, 27]. In total, six continuous-flow bioreactor experiments were run: three biotic experiments under identical conditions at 10 °C, pH 7.85, 1.1 mM HCO3−, 1 mM S2O3−2, and yeast extract; two uninoculated controls run under the same conditions as above to obtain an abiotic dissolution rate in undersaturated conditions; and one biotic experiment was run under identical conditions, minus the addition of S2O3−2 to elucidate the impact of heterotrophy on mineral stability.

10 L carboys containing the final medium and sterile bioreactors were assembled aseptically in a class A2 biological safety bench. Approximately 325 mL of the media was pumped into the bioreactor, followed by inoculating with 20 mL of pure culture and ~600 μL of fresh microbial culture. The luer-lock was removed and a 0.2 μM filter was added in its place. Bioreactors were then placed in a 10 °C refrigerator on a stir plate at 80 RPM.

Inoculated bioreactors were run in batch phase for 4 h to promote ample attachment of cells to aragonite coupon surfaces. Following the batch phase, the bioreactor was moved back to the clean hood where one coupon holder rod was removed and replaced with a sterile rubber stopper. Two coupons were placed in the incinerator at 500 °C for 4 h, followed by weighing for mass loss. The additional coupon was fixed in 4% PFA for 2 h, washed, and stained with DAPI [4,6-diamidino-2-phenylindole] for 30 min. Stained coupons were washed and mounted on coverslips (0.17 mm thickness) with DPX mountant for cell counting. After batch phase harvesting, the pump was turned on to the max flow rate (31 mL/min) and the first 100 mL out of the reactor was collected to measure pH, alkalinity, [Ca2+], and OD590. pH and alkalinity were measured using a Hanna Instruments total alkalinity mini titrator (HI-84431) and pH meter (HI 1131B) following the manufacturers protocol (Hanna Instruments, Woonsocket, RI, USA). [Ca2+] was measured using a Hach hardness test kit (product #2063900) following to the manufacturer’s protocol (Hach, Loveland, CO, USA). OD590 was measured on a Thermo Spectronic 20D + (Thermo Fisher Scientific, Waltham, MA, USA), where 1 mL of sterile medium was added to a cuvette to blank the spectrophotometer, and then 1 mL of the collected outflow was placed in a cuvette and measured. The pump was kept at the max flow rate for approximately 3–4 h after batch phase to remove planktonic cells and the additional HCO3− added during batch phase. Directly after flushing the bioreactor, water chemistry measurements were taken as they were above, and then the flow rate was dropped to 10 mL/min. Henceforth, the flow rate was only adjusted to keep the bulk fluid at approximately the pH and alkalinity of the starting conditions (max flow rate of 21 mL/min during week 3). To ensure consistency of the bulk fluid chemistry, pH and alkalinity measurements were taken twice daily (8–12 h intervals), just before changing carboys. Carboys were changed about every 8–12 h, depending on the current flow rate. To change carboys, the following steps were taken: (1) The pump was briefly turned off, and sterile aluminum foil was wrapped around the outflow tubing. (2) The bioreactor and currently connected carboy were moved to the biological safety bench on a roll cart. (3) Tubing that connected the carboy to the bioreactor was disconnected, and the male-port of the tubing still connected to the bioreactor was submerged in sterile 70% EtOH. (4) A new 10 L carboy containing the final medium was removed from the 10 °C refrigerator. (5) Aluminum foil covering the female-port of the tubing on the carboy was removed, and sprayed with sterile 70% EtOH. (6) Tubing from the bioreactor and carboy were then connected and placed on the roll cart, and moved back to the 10 °C refrigerator. (7) The pump was turned back on and flow resumed. One coupon holder rod was harvested every four days for 21 days, where two coupons were incinerated and weighed for mass loss and one coupon was fixed and stained for cell counting as described above. Surface area of aragonite coupons were estimated using the diameter of the coupon, the density of aragonite (2.93 g/cm3), and the mass of the coupon before and after bioreactor experiments. Density and mass were used to estimate the volume of the coupon, followed by deriving an estimated surface area. Using the mass of coupons before and after experiments, we calculated the total number of moles lost per unit area. Averaging these data out over time we were able to determine a dissolution rate in units of μmol CaCO3 • cm−2 • hr−1, which we then converted to μmol CaCO3 • cm−2 • yr−1 (8,760 h/year).

**Confocal microscopy**

We utilized the flow cell in conjunction with the ratiometric dye C-SNAFL-1 [5-[6]-carboxyseminaphthofluorescein] and Hoechst 33342 for measuring the in vivo pH of 21-day-old biofilms, and visualizing the biofilm, respectively. Coupons transferred to the flow cell were connected to a peristaltic pump at 3 mL/min until imaging occurred.

**Spectrofluorometric assays**

To calibrate C-SNAFL-1 photon excitations and to evaluate the dye’s potential use in measuring pH in biofilms, 1 mL aliquots of the final media with 10 mM HEPES buffer were adjusted to pH 5.0, 5.2, 5.6, 6.0, 6.4, 6.8, 7.2, 7.6, and 8.0. The final probe concentration was ~1.09 μM of
C-SNAFL-1. Additional calibrations were performed with the addition of 4% (vol/vol) *C. baekdonensis* strain LH4.

\[ \text{Ratio} = \frac{(\text{Ex}_{488} - \text{Ex}_{488\,\text{bkgd}})}{(\text{Ex}_{561} - \text{Ex}_{561\,\text{bkgd}})} \quad (\text{IV}) \]

Confocal scanning laser microscopy

All biofilm images, pH and ratio calibrations in the flow cell were collected using a Nikon TiE inverted microscope equipped with an A1RSi confocal scan head (Nikon). Hoechst 33342 fluorescence was excited with a 405 nm laser and collected between 425 and 475 nm using a regular PMT to visualize the biofilm. SNAFL emission was collected between 570 and 620 nm with a GaAsP PMT alternating excitation with the 488 and 561 nm lasers to obtain a ratiometric image. This image was used to calculate the pH across the field (below). 10X/0.45 PlanApo and 20X/0.75 PlanApo VC objectives were used to collect 512- by 512-bit resolution z-stacks. Depths in this paper are the distance from the substratum (base of biofilm) to the focal plane (distance furthest from the biofilm).

To visualize the biofilms, flow was suspended and coupons were stained with Hoechst 33342 (2 drops/mL medium) for 15 min and then flushed. C-SNAFL-1 was added to the medium at a final concentration of 20 μM. Flow resumed with C-SNAFL-1 in the medium and then was suspended for no more than 15 min to minimize the accumulation of acidic metabolites, which could artificially modify the local pH. Image planes were set up to visualize the coupon-attached biofilm and the bulk media adjacent to the coupon in the same image. Biofilms were excited at 405 nm, and emission was detected between 425 and 475 nm. pH microenvironment images were captured using dual excitation at 488 nm and 561 nm, and single emission at 600 nm. To determine the z-stack image parameters, we first excited the sample at 405 nm to visualize the base of the biofilm, and then switched to the dual excitation channels to determine the end of the focal plane (e.g. the coverslip; point furthest away from the biofilm in the z-dimension where the signal diminishes). After determining the z-distance to image, we imaged the biofilm at 405 nm, and then the pH of the microenvironment at 488/561 nm. The pH of the microenvironment was determined by calculating the ratio of excitation intensities (i.e., pixel values) between the two channels (Ex\textsubscript{488 nm}/Ex\textsubscript{561 nm}). All intensities were determined using NIS Elements software.

Calibration standards for CSLM were performed in the flow cell with a sterile aragonite coupon and media with 20 μM C-SNAFL-1 and 10 mM HEPES buffer at pH 8.0, 7.8, 7.7, 7.55, 7.2, 7.0, and 6.8. Calibrations were performed with identical microscope settings as in the experiment above. Titration curves of pH versus ratios were calculated using NIS Elements software according to Eq. IV and were used to convert C-SNAFL-1 excitation ratios to pH values.

Image analysis

pH gradients within the biofilm and surrounding microenvironment were evaluated with NIS Elements software by using kymographs on the ratio images (Ex\textsubscript{488 nm}/Ex\textsubscript{561 nm}). Each kymograph was -500 μm in length (x) and ~100 μm in depth (z). The resulting kymograph image had 4–5 regions of interest selected, including 2–3 spanning the biofilm into the bulk fluid directly above, and 2-3 in the bulk fluid adjacent to the biofilm. Regions of interest varied in z-depth depending on the thickness of the biofilm, though all were 13 μm in width (x).

Estimation of CaCO\textsubscript{3} dissolved from various seep locations

Seep locations

PDC1 – Dive, SO 162; Site, Point Dume; Feature, Chimney complex 1; Latitude, 33 56.45753693; Longitude, 118 50.70879299; Depth (m), 729.522

PDC2 – Dive, SO 164; Site, Point Dume; Feature, Chimney complex 2; Latitude, 33 56.50074983; Longitude, 118 50.64165898; Depth (m), 725.723

PDC3 – Dive, SO 164; Site, Point Dume; Feature, Chimney complex 3; Latitude, 33 56.45174234; Longitude, 118 50.70728425; Depth (m), 728.966

Determining total surface area

Each image taken by submersible at Point Dume had two scale bars generated at the time of imaging, one in the background and one in the foreground, with each representing 10 cm. These two scale bars, in conjunction with a designed macro in Fiji/ImageJ were used to interpolate the total surface area within each image. The macro is based on the equation $y - y_1 = m(x - x_1)$. The slope, $m$, was determined using the equation $((y_2 - y_1)/(x_2 - x_1))$, where $y_2 = 10 \text{ cm/(length in pixels of the foreground scale bar)}$, $y_1 = 10 \text{ cm/(length in pixels of the distant scale bar)}$, $x_2 = (y\text{-coordinate of the foreground scale bar})$, $x_1 = (y\text{-coordinate of the distant scale bar})$. Each image was imported into Fiji/ImageJ, and scaled to 1333 × 750 pixels. All scales were removed to put all measurements in pixel values. Centroid and integrated density were added to set measurements. Each image was duplicated and the duplicated images were changed to 32-bit images. The macro was then run on each respective image. A polygon was then drawn around the visible area within each image, and added to the ROI manager. The ROIs were added to their respective duplicate and measured. These data gave us the total visible surface area present in each image.
Determining surface area of exposed carbonates and carbonated-attached bacterial mats

Polygons were drawn around all exposed carbonates within the visible region. These ROIs were then added to their respective duplicated image, and the total area of exposed carbonates was measured. Carbonate ROIs that contained small regions in which bacterial mats were not visible had “negative” ROIs drawn around them. When applicable, these negative ROIs were subtracted from the total exposed carbonate measurement, resulting in the total mat-on-rock surface area measurement. It is likely that bacteria are also colonizing the surfaces on which no obvious mat can be observed, but removal of these areas from consideration was done to provide a conservative lower estimate of bacterial coverage.

Moles of carbonate rock dissolved was calculated as follows

Experimentally-derived carbonate dissolution rate = 1773.97 \mu \text{mol CaCO}_3 \cdot \text{cm}^{-2} \cdot \text{yr}^{-1}.

Apply dissolution rate to seep surface area covered by rock (e.g. Hydrate Ridge) to obtain the annual dissolution; 17.7397 \text{ mol CaCO}_3 \cdot \text{m}^{-2} \cdot \text{yr}^{-1} \cdot 3.00E+05 \text{ m}^2 = 5.32E+06 \text{ mol CaCO}_3 \cdot \text{yr}^{-1}.

Apply 92.77% average carbonate-attached bacterial mat percent coverage (Supplementary Table 3) to annual dissolution; (5.32E+06 \text{ mol CaCO}_3 \cdot \text{yr}^{-1}) \cdot (0.9277) = 4.94E+06 \text{ mol CaCO}_3 \cdot \text{yr}^{-1}.

We then applied these calculations to the remaining five seep sites in Table 1 to obtain the annual amount of carbonate dissolved per seep.

Results and discussion

Carbonate-attached mats contain sulfur-oxidizing bacteria

Mats of sulfur-oxidizing bacteria have been recorded at many localities worldwide including on the surface of authigenic carbonates at methane seeps [37–39]. Large sulfur-oxidizing bacteria of the family Beggiatoaceae may form conspicuous mats; while the presence of other sulfur oxidizers can be inferred from chemical profiles indicating the presence of sulfide oxidation at the sediment-water interface [40, 41]. In addition to the presence of macroscopic mats that can be observed by submersible, some previous molecular studies have also identified sulfur-oxidizing bacteria associated with submerged carbonate nodules [42–44] and carbonates exposed at the sediment-water interface [45, 46]. These studies demonstrate that sulfur-oxidizing bacteria are enriched in exposed carbonates.
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S. Marthioploca spp. [49] of the Beggiatoales, Cocleimonas spp. [50] and other Thiotrichales, Woeseia spp. of the Steroidobacteriales [51], Thiohalophilus spp. [52] as well as other Ectothiorhodospirales. But other sulfur-oxidizing ectotypes were present including ASVs within the Campylobacteriales (esp. Sulfovorum and Sulfurimonas spp.) [53] and within the Rhodobacterales, (Roseobacteria clade NAC1 [54] and unclassified Rhodobacteraceae). Interestingly, our model strain, C. baekdonensis strain LH4 is a member of the Roseobacteria group, possesses only the complete Sox pathway, but some members of this clade also possess the reverse Dsr pathway making this clade metabolically versatile in their capacity to oxidize sulfide and thiosulfate [55]. The prevalence of specific ectotypes of sulfur-oxidizing bacteria likely depends on the concentration of sulfide and other nutrients such as available carbon substrates, but filamentous communities dominated by Gammaproteobacteria, esp. Ca. Marithrix, are proposed to be well-established mat communities vs. early successional communities dominated by the Campylobacterota (formally known as Epsilonproteobacteria) [45, 56]. By scanning electron microscopy, we also show that some putative filamentous sulfur bacteria are associated with pitting features on the rock surface (Fig. 2). Combined with previous descriptions from the literature [38, 42, 43, 57, 58], our results support the hypothesis that carbonate surfaces at methane seeps host diverse sulfur-oxidizing bacteria.

Biofilm reactor experiments

To simulate plausible dissolution rates that could be occurring in biofilms associated with natural marine seep carbonates, we conducted a series of reactor experiments to investigate the influence of a bacterial biofilm on carbonate dissolution rates. For these experiments, we inoculated bioreactors containing carbonate rock coupons with a pure culture of C. baekdonensis strain LH4 [30], a chemolithoautotrophic alphaproteobacterium that we isolated from a marine methane seep. C. baekdonensis LH4 is capable of lithotrophic growth via complete oxidation of \( \text{S}_2 \text{O}_3^{2-} \) to \( \text{SO}_4^{2-} + 2\text{H}^+ \). It can also grow heterotrophically on yeast extract, thus we were able to separate the presence of the microbes from their metabolism and show that their metabolism is the mechanism of dissolution rather than the physical attachment. In total, six biofilm reactors were run under continuous flow for three weeks. Three independent bioreactors (designated “thiosulfate” in Fig. 3) were run with the addition of \( \text{S}_2 \text{O}_3^{2-} \) to
induce a sulfur-oxidizing metabolism that is known to generate acidity. Two independent uninoculated control bioreactors were run with the addition of S$_2$O$_3^{2-}$ to determine the abiotic dissolution rate of aragonite. Additionally, one bioreactor was inoculated, but without S$_2$O$_3^{2-}$ (designated “heterotrophic” in Fig. 3) to measure the influence of a heterotrophic metabolism (specifically the production of CO$_2$ from organic carbon oxidation) on aragonite dissolution rates. Aragonite dissolution rates for each test condition – thiosulfate (sulfidic), heterotrophic, and abiotic control – were 1773.97 (±324.35) μmol CaCO$_3$ cm$^{-2}$ yr$^{-1}$, 152.81 (±123.27) μmol CaCO$_3$ cm$^{-2}$ yr$^{-1}$, and 272.99 (±249.96) μmol CaCO$_3$ cm$^{-2}$ yr$^{-1}$, respectively. The bulk solutions pH (~7.75-90) and alkalinity (~1.1-3 mEq/L) of the biofilm reactors remained nearly constant for three weeks in all six reactors. However, these results show that the production of sulfuric acid by S$_2$O$_3^{2-}$ oxidation increases the aragonite dissolution rate by nearly a factor of 12, and 7, when compared to CO$_2$ production from organic carbon oxidation, and abiotic dissolution, respectively. We interpret this dissolution to primarily result from protons produced from lithotrophic sulfur-oxidation creating an acidic microenvironment, a hypothesis that we further tested by measuring the pH within the biofilm.

**Imaging biofilm acidification**

Previous work in non-marine settings has demonstrated that metabolic acid production can create acidic microenvironments within a biofilm that can lead to mineral dissolution [14]. In order to test the hypothesis that lithotrophic sulfide oxidation can also create acidic microenvironments under carbonate-buffered marine conditions, we used laser scanning confocal microscopy in conjunction with a pH sensitive fluorophore, 5-[6]-carboxyseminaphthofluorescein (C-SNAFL-1), to visualize and measure the pH of the aragonite-attached biofilms.
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Fig. 3 Mass loss of aragonite coupons in inoculated sulfidic and heterotrophic bioreactors, as well as uninoculated control bioreactors. Each data point represents one aragonite coupon from a bioreactor (Thiosulfate, \( n = 41 \); Heterotrophic, \( n = 17 \); Abiotic Control, \( n = 34 \)). The mass loss (initial mass – final mass) was converted from mg to \( \mu \)mol CaCO\(_3\) • cm\(^{-2}\) and plotted against the total time the coupon was in the bioreactor. Linear regressions were performed for each test condition to extrapolate an annual dissolution rate. The line of best fit for each condition is plotted (dashed colored lines), and error bars are s.e. for the coupons from that time point.

In biofilms grown in the presence of \( S_2O_3^{2-} \) we observed a pH gradient starting at the mineral surface, progressing through the biofilm and into the medium above (Fig. 4). At the mineral surface, the site of microbial attachment, the pH ranged from 7.68–7.54 (Fig. 4C–E). Within the biofilm the pH decreased to the lowest observed value of 7.33 (Fig. 4D). Just outside of the biofilm and in the bulk solution, the pH increased to 7.44–57 (Fig. 4C–E). Adjacent to the biofilm and mineral, in the bulk solution of the flow cell, the pH increased in the xy dimensions (Fig. 4), and to an even greater extent in the xyz dimensions, reaching a maximum value of 7.76 (Fig. 4D), nearly the pH value of inflowing media (~7.80).

We interpret this pH gradient to represent the influence of the acid produced by thiosulfate oxidation produced by the following reaction:

\[
S_2O_3^{2-} + H_2O + 2O_2 \rightarrow 2SO_4^{2-} + 2H^+ \quad (II)
\]

These protons are then neutralized by the dissolution of the aragonite mineral by the following reaction:

\[
2H^+ + 2CaCO_3 \rightarrow 2HCO_3^- + 2Ca^{2+} \quad (III)
\]

Resulting in the following net reaction:

\[
2CaCO_3 + S_2O_3^{2-} + 2O_2 + H_2O \rightarrow 2SO_4^{2-} + 2HCO_3^- + 2Ca^{2+} \quad (IV)
\]

We believe that dissolution and subsequent proton neutralization is occurring at the biofilm-mineral interface, as this explains the pH values observed at the mineral surface. The lowest pH value recorded, 7.33, occurs in the middle of the biofilm—in the region that is most isolated from the buffering capacity of the aragonite mineral and overlying medium. The pH increase from the biofilm to the overlying medium is attributed to the large buffering capacity of the flow medium, which is meant to simulate the practically unlimited acid-buffering capacity of the bulk ocean. We were unable to obtain pH values further away (z-dimension) from the biofilm and minerals surface due to physical limitations in the microscopy setup. However, the bulk pH values of the flow cell and bioreactor are near inflow conditions (~7.80), indicating that the acidity produced by sulfur-oxidizing bacteria is confined to the microenvironment. Under heterotrophic conditions (growth without \( S_2O_3^{2-} \)), no pH gradient was observed (Fig. 5). Starting at the minerals surface, going through the biofilm and into the bulk medium above the biofilm, the largest pH change was 0.01 units. Furthermore, in the bulk medium adjacent to the mineral and biofilm, the largest pH change was 0.02 units. In both sulfidic and heterotrophic conditions, aragonite-attached biofilms were nearly identical in thickness (Supplementary Fig. 4).

These results suggest the following: (1) oxidation of \( S_2O_3^{2-} \), and perhaps other reduced sulfur species, leads to the production of sulfuric acid that can dissolve aragonite even under buffered marine conditions; and (2) acidity generated by oxidizing \( S_2O_3^{2-} \) is greatest near the mineral surface, and is largely confined to that microenvironment.

Implications for the global carbon cycle

Our bioreactor experiments and observations of seep samples reveal the potential for sulfur-oxidizing bacteria to contribute to carbon flux from the carbonate rock reservoir to the ocean and atmospheric carbon reservoirs.

Using existing data on authigenic carbonate rock surface coverage at cold seeps, as well as ground-truth images from the Point Dume cold seep and sulfur-oxidation-induced carbonate dissolution rates generated from our experiments, we calculated an initial estimate of the amount of carbon that can be dissolved in various well-characterized cold seeps. Using multibeam, sidescan sonar, and ground-truth images of the seafloor, six previously published site descriptions [57–62] were used to estimate the percent coverage of rocks relative to the size of the active seep. Three ground-truth images from Point Dume—designated PDC1, PDC2, and PDC3 (Supplementary Figs. 5 and 6, and Fig. 6, respectively)—were used to estimate the area and percent coverage of microbial mats on exposed carbonates at these sites. These three sites comprise ~43 m\(^2\) in total area, with ~25 m\(^2\) of exposed carbonates, or ~57% rock: total area-coverage (Supplementary Table 3). Bacterial mats
attached to exposed carbonates total ~23 m², or ~93% mat:rock coverage (Supplementary Table 3). While the exact percent coverage of bacterial mats on exposed carbonates at the six sites used to make these estimations are unknown, all sites have reported extensive bacterial mat coverage throughout the active seep [45, 57–63]. In this context, we ascribed the bacterial mat percent coverage calculated in the PDC images to each of the six sites, along with the sulfur-oxidation-induced carbonate dissolution rate from this study to estimate the amount of carbon dissolved from seep carbonates at these locations (Table 1).

These initial estimates come with considerable uncertainty, such as the true extent of biofilm coverage on seep carbonates and their respective metabolisms, including in situ rates of sulfide oxidation, the total surface area of exposed carbonates, percent composition of carbonate-mineral phases, and the total number of seeps globally. While the amount of methane-derived carbonate precipitated globally is not well constrained, it has recently been estimated that 1.7 Tmol•yr⁻¹ (range of 0.6-3.6 Tmol•yr⁻¹) of methane-derived authigenic carbonate precipitates in diffusive settings [6], which is similar to previous estimates of 1 Tmol•yr⁻¹ and 1.5 Tmol•yr⁻¹ [5, 64]. Although the number of cold seeps globally is poorly constrained, thousands of cold seeps have recently been detected, with tens of thousands more predicted [5, 65, 66]. Using the estimated average microbially-mediated dissolution per cold seep from the six locations in Table 1 (based on our experimentally-derived rates) and extrapolating that to 10,000 seeps, we estimate the amount of carbonate dissolved via sulfide-oxidation to be 6.06×10¹⁰±1×10¹⁰ mol C•yr⁻¹, or 7.28×10⁸ (±1.20×10⁸) kg C•yr⁻¹. For comparison, estimated anthropogenic changes resulting in sediment-water interface calcite dissolution is 4×10¹¹ (±1×10¹¹) kg C•yr⁻¹ [67]. And it is important to note, that anthropogenic contribution to calcite dissolution mostly occurs in distinct “hot spots” where anthropogenic CO₂ is introduced to bottom waters, i.e. northwestern Atlantic Ocean, and around islands in the Pacific and Indian Oceans. Marine saturation states of aragonite, which is more soluble than calcite, are already responding strongly to anthropogenic inputs of CO₂ [68] and
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are projected to continue to rapidly decline, but not to the extent that most seep carbonates would be affected within this century [69].

While these estimates come with considerable uncertainty, even with lower amounts of mat coverage in some areas, these fluxes constitute an annual geological lability of 3.56% (range: 1.38–11.92%) of the methane-derived carbon that is thought to be sequestered in stable form as authigenic carbonates. Additionally, some mats are long-lived and grow up to several centimeters thick and are known to span hundreds of meters in diameter, which may allow for more acid to accumulate in the biofilm and subsequently dissolve more carbon than estimated here [39, 45, 70, 71]. It should be noted that this estimation is contingent upon carbonates being exposed at the surface where sulfur-oxidizing bacteria can colonize carbonates, and have O₂ available as an electron acceptor. Community composition and available organic substrates likely influence dissolution rates as well. For example, thiosulfate-oxidizing chemoorganotrophs produce tetrathionate that generates hydroxide and this

**Fig. 5** Confocal laser scanning microscopy pH imaging of biofilm attached to aragonite coupon under heterotrophic conditions. C-SNAFL-1 stained *C. baekdondensis* LH4 biofilm on aragonite coupon under heterotrophic (control) conditions. A 3D view of the aragonite coupon and bulk solution inside the flow cell. The bulk solution is colored and the coupon is the black space directly under it. Orange lines (A, B) are kymograph transects that correspond to their respective images in C, D and E. B 2D view of the aragonite coupon showing where kymograph lines are drawn. C–E. Kymograph profiles show no pH gradient and remain stable at the mineral surface, through the biofilm, above the biofilm, and within the bulk medium.

**Fig. 6** Estimating coverage of carbonates and biofilms from Point Dume Complex 3 (PDC3). Biofilms of sulfur-oxidizing bacteria colonize seep carbonates. Scale bars in a are both 10 cm. The scale bars were used to generate a macro (c) that corrects for perspective and computes the area in a region of interest (ROI). ROIs are outlined in yellow (b, c). The total visible area was calculated first by creating an ROI around the visible area of the photo. ROIs were drawn around all exposed carbonates to generate the total rock coverage. Smaller ROIs were drawn within larger ROIs where biofilms are not present and then subtracted from the total rock coverage to generate total biofilm coverage. Area measurements were converted from cm² to m², and then converted to percent coverage (mat:rock, mat:total area, rock: total area).
could act to counter acid production [72], although tetra-
thionate is short lived in sediments [12]. Also, intracellular sulfur globule producing sulfur-oxidizers such as Ca. Marithrix sequester reduced sulfur. The environmental triggers and the rates of sulfur globule oxidation to sulfuric acid are currently unknown in natural seep settings. Likewise, autotrophy, heterotrophy and fermentation are competing processes that could affect the saturation state of the carbonates through the uptake and release of inorganic carbon. The influence of these metabolisms should be considered in future to work to understand the dynamics of biofilm communities on the dissolution of carbonates. Additionally, our bioreactor experiments examine the effects of lithotrophic dissolution on aragonite, which is a major mineral phase in many seep carbonates. However, other carbonate phases present in carbonate assemblages at methane seeps are less soluble and future work will be required to determine the influence that sulfur-oxidation may have on minerals such as calcite and dolomite. These results demonstrate a novel microbially-mediated aspect of carbon cycling at methane seeps in the form of seafloor carbonate weathering mediated by lithotrophic acid production. Furthermore, they underscore the need for improved quantification of the number of methane seeps globally, the true extent of exposed carbonates in methane seeps, and the extent and composition of microbial communities that colonize these carbonates.

Data availability

Unaltered FASTQ files are deposited in the NCBI Sequence Read Archive, Bioproject PRJNA693332, and the assembled reads are available through the Data Repository of the University of Minnesota (https://conservancy.umn.edu/handle/11299/218173).

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Author contributions

J.J.M. and P.R.G. collected field samples. D.J.L., P.R.S., E.R., and J.V.B. performed experiments and/or microscopy. B.E.F. performed all bioinformatic and genomic analyses. B.E.F., and J.V.B. conceived of the study. D.J.L., B.E.F., and J.V.B. wrote the manuscript.

Compliance with ethical standards

Conflict of interest

The authors declare that they have no conflict of interest.

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