



# Organic Stabilization of Extracellular Elemental Sulfur in a *Sulfurovum*-Rich Biofilm: A New Role for Extracellular Polymeric Substances?

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### Specialty section:

This article was submitted to  
Microbiological Chemistry and  
Geomicrobiology,  
a section of the journal  
*Frontiers in Microbiology*

Received: 03 June 2021

Accepted: 15 July 2021

Published: 06 August 2021

### Citation:

Cron B, Macalady JL and  
Cosmidis J (2021) Organic  
Stabilization of Extracellular Elemental  
Sulfur in a *Sulfurovum*-Rich Biofilm: A  
New Role for Extracellular  
Polymeric Substances?  
*Front. Microbiol.* 12:720101.  
doi: 10.3389/fmicb.2021.720101

This work shines light on the role of extracellular polymeric substance (EPS) in the formation and preservation of elemental sulfur biominerals produced by sulfur-oxidizing bacteria. We characterized elemental sulfur particles produced within a *Sulfurovum*-rich biofilm in the Frasassi Cave System (Italy). The particles adopt spherical and bipyramidal morphologies, and display both stable ( $\alpha$ -S<sub>8</sub>) and metastable ( $\beta$ -S<sub>8</sub>) crystal structures. Elemental sulfur is embedded within a dense matrix of EPS, and the particles are surrounded by organic envelopes rich in amide and carboxylic groups. Organic encapsulation and the presence of metastable crystal structures are consistent with elemental sulfur organomineralization, i.e., the formation and stabilization of elemental sulfur in the presence of organics, a mechanism that has previously been observed in laboratory studies. This research provides new evidence for the important role of microbial EPS in mineral formation in the environment. We hypothesize that the extracellular organics are used by sulfur-oxidizing bacteria for the stabilization of elemental sulfur minerals outside of the cell wall as a store of chemical energy. The stabilization of energy sources (in the form of a solid electron acceptor) in biofilms is a potential new role for microbial EPS that requires further investigation.

**Keywords:** sulfur, biominerization, organomineralization, extracellular polymeric substances, biofilms, Raman, STXM

## INTRODUCTION

Elemental sulfur [S(0)] is an intermediate of the biogeochemical sulfur cycle found in many natural environments, such as marine sediments, marine and lacustrine water columns, cold or hot springs, hydrothermal environments, salt marshes, and caves (Zerkle et al., 2010; Findlay et al., 2014; Hamilton et al., 2015; Lau et al., 2017; Jørgensen et al., 2019). S(0) is formed by chemical or biological oxidation of more reduced sulfur species, although in low-temperature environments, biological S-oxidation rates are typically more than three orders of magnitude faster than abiotic rates (Luther et al., 2011). A wide diversity of microorganisms can oxidize sulfide, polysulfides, or thiosulfate and precipitate S(0) through both phototrophic and chemotrophic pathways (Kleinjan et al., 2003; Dahl and Prange, 2006). In turn, microbially

formed S(0) can be used as a source of energy for a wide diversity of S-oxidizers, S-reducers, and microorganisms that perform S(0) disproportionation (Warthmann et al., 1992; Dahl, 2020a). Elemental sulfur thus occupies a central and ecologically important role in the biogeochemical sulfur cycle.

Biogenic S(0) is deposited either intracellularly or extracellularly (Kleinjan et al., 2003; Dahl and Prange, 2006; Maki, 2013). Most studies on microbial S(0) biomimetication so far have focused on microorganisms forming intracellular S(0) globules, for instance *Allochromatium vinosum*, *Acidithiobacillus ferrooxidans*, *Thiothrix* spp., or large colorless SOB, such as *Thiomargarita namibiensis*, *Thioploca* spp., or *Beggiatoa* spp. (Brune, 1995; Gray and Head, 1999; Pasteris et al., 2001; Prange et al., 2002; Maki, 2013; Nims et al., 2019; Dahl, 2020b). Many SOB can form S(0) extracellularly, such as green sulfur bacteria (Chlorobiaceae; Gregersen et al., 2011; Marnocha et al., 2016), purple sulfur bacteria of the Ectothiorhodospiraceae family (Then and Trüper, 1983), some purple non-sulfur bacteria (Hansen and van Gemerden, 1972), some lithotrophic sulfur bacteria (Cron et al., 2019), and cyanobacteria (Oren and Shilo, 1979). More work is needed to decipher the formation mechanisms of extracellular S(0) and to characterize its properties.

In sulfide-rich solutions, S(0) is thermodynamically stable only under a very restricted range of natural Eh-pH conditions (e.g., Eh  $-0.05$  to  $0.15$  and pH  $<5$  at  $25^{\circ}\text{C}$  and 1 bar; see Figure 7 in Rickard and Luther, 2007). It is thus not clear how microbial sulfur, particularly when it is extracellular, persists in most circumneutral aqueous environments. Cosmidis et al. (2019) recently showed that interactions with organics are important for the abiotic formation of S(0) minerals, a process termed S(0) organomineralization. Organomineralized sulfur can be found in several metastable crystal structures, including the monoclinic allotropes  $\beta$ -S<sub>8</sub> and  $\gamma$ -S<sub>8</sub>, which are thought to be stabilized by close association with organics. Organic-mineral interactions may also be important in extracellular S(0) formation by bacteria. Indeed, organics produced by the chemolithoautotrophic SOB *Sulfuricurvum* (S.) *kuijense* (Campylobacterota) are needed for extracellular S(0) formation by this organism. The S(0) globules of *S. kuijense* are composed of  $\beta$ -S<sub>8</sub> and  $\gamma$ -S<sub>8</sub> and are coated by organic envelopes that allow them to precipitate under conditions outside of their theoretical thermodynamic stability domain (Cron et al., 2019). Other SOB, such as *Thiobacillus* sp. W5 (Kleinjan et al., 2005) and *Chlorobaculum* (C.) *tepidum* (Hanson et al., 2016; Marnocha et al., 2019), also produce extracellular S(0) globules with organic envelopes, suggesting that organics play a previously overlooked role in microbial S(0) formation and stabilization in nature.

Previous studies describing the importance of organics in S(0) mineralization were based on laboratory experiments, whereas observations from natural environments are still lacking. Elemental sulfur particles with metastable structures and intimate associations with organics were described in a sulfur-rich glacial site in the Arctic, but it could not be determined whether microbial mediation was involved in their formation (Lau et al., 2017). In the present study, we characterized

extracellular S(0) particles formed within microbial biofilms in a subsurface environment dominated by sulfur-cycling bacteria. The Grotto Grande del Vento-Grotta del Fiume (Frasassi) cave system is actively forming in Jurassic limestone in the Apennine Mountains of the Marches Region, Central Italy (D'Angeli et al., 2019). The S(0) minerals described here are found within microbial biofilms in a microaerophilic sulfide-rich stream, Pozzo di Cristalli. Previous full-cycle rRNA and metagenomic approaches identified Campylobacterota most closely affiliated with *Sulfurovum* species as the most abundant organisms in streamer biofilms at this location (Jones et al., 2008; Macalady et al., 2008; Hamilton et al., 2015). *Sulfurovum* oxidize sulfide and thiosulfate to sulfate and extracellular S(0) globules, which can make up more than 60% by weight of their mats. These primary producers serve as the principal source of organic carbon to the subsurface stream ecosystem (Hamilton et al., 2015). The biofilms are attached to rocks in the stream bed where they intersect the water surface. In this environment, sulfidic water turbulently mixes with oxygen in the cave air, providing chemical energy for growth (Macalady et al., 2008).

We characterized the morphology and crystal structure of S(0) particles formed by microbial S-oxidation in these natural *Sulfurovum*-dominated biofilms using scanning electron microscopy (SEM), X-ray diffraction (XRD), and ultra-low-frequency Raman spectromicroscopy. Organics closely associated with the S(0) minerals were also characterized using Fourier-transform infrared (FTIR) spectroscopy and scanning transmission X-ray microscopy (STXM). Our results provide important clues about how organic compounds (e.g., carboxylic acids, amides, aromatics, and aliphatic compounds) excreted by cells are used to encapsulate and stabilize extracellular S(0) within EPS matrices. We discuss the ecological implications of this new role of EPS in sulfur-based microbial ecosystems.

## MATERIALS AND METHODS

### Biofilm Sample Collection

Pozzo di Cristalli is a cave stream that periodically hosts blooms of streamer biofilms primarily composed of Campylobacterota in the genus *Sulfurovum* (Jones et al., 2008; Macalady et al., 2008; Hamilton et al., 2015). The biofilms thrive in turbulent stream riffles near the air-water interface. Sulfide concentration was  $20.5\text{ }\mu\text{M}$  at the time of sampling in September 2016. Streamer sample PC1647 was harvested from customized floating twine supports. In September 2017, a *Sulfurovum*-dominated biofilm (streamer sample PC1718) was collected from a limestone cobble at the same location (Figure 1). Biofilm samples were divided into sterile Falcon tubes and cryotubes. Samples in Falcon tubes were preserved with glutaraldehyde and stored at  $-20^{\circ}\text{C}$ . Cryotube samples were stored at  $-80^{\circ}\text{C}$ .

### X-Ray Diffraction

For XRD, approximatively 1 ml of biofilm sample PC1718 (i.e., the condensed content of a microcentrifuge tube) was rinsed three times with deionized water and deposited on a



**FIGURE 1** | Biofilm collection site in September 2017 with billowing white *Sulfurovum*-rich streamer biofilm on the surface of the stream at center right. The *Sulfurovum*-dominated biofilm is surrounded by a weakly pigmented *Beggiatoa* biofilm growing on the black sediment surface.

single crystal miscut Si holder. Samples were analyzed using a PANalytical Empyrean diffractometer paired with a PIXcel3D detector, and Cu K $\alpha$  ( $\lambda = 1.5406$  Å) incident X-ray radiation. Scans were conducted over a 2 $\theta$  range of 5–70°, and analyses used a step size of 0.025°, a time of 96.4 s per step, and a current density of 40 mA. XRD data were analyzed using MDI JADE software. Mineral identification was performed by comparison with reference XRD patterns from the International Centre for Diffraction Data (ICDD) database.

### Raman Spectromicroscopy

For Raman, 5 ml of frozen unpreserved biofilm (sample PC1718) was rinsed three times with deionized water to remove salts. Samples were either deposited and dried on a microscope slide or kept wet between a microscope slide and cover slip. Raman spectra were collected using a Horiba LabRam HR Evolution Vis–NIR optimized & AIST-NT Scanning Probe, and a Si-based CCD detector (1,024 × 256 pixels). Raman signals were measured in the low-frequency range using BragGrate notch filters (Nims et al., 2019). The spectrometer was calibrated using the 520 cm $^{-1}$  Raman peak of Si prior to analysis. Spectral data were corrected for instrumental artifacts and baseline-subtracted using a polynomial fitting algorithm in LabSpec 6 (Horiba Scientific). The sample spectra were compared with reference Raman spectra for different allotropes of S(0) (Nims et al., 2019).

### Scanning Electron Microscopy

For SEM, biofilm samples were rinsed with deionized water and deposited on polycarbonate filters (GTTP Isopore membrane filters, Merck Millipore, pore size 0.2  $\mu$ m) or on glass slides (for correlative Raman analyses, see below). The samples were allowed to dry at ambient temperature and coated with iridium or gold prior to analysis. For sample PC1718, SEM analyses were conducted on a Field Emission Nova NanoSEM 630 at the Materials Characterization

Laboratory at The Pennsylvania State University. Images were acquired with the microscope operating at 3–7 kV and a working distance (WD) of ~3–5 mm. Energy-dispersive X-ray spectroscopy (EDXS) analyses were performed at 12 kV and WD ~7 mm. For sample PC1647, SEM analyses were conducted on a JSM-7401F field emission scanning electron microscope (FESEM) at the Nanoscale Fabrication Laboratory at the University of Colorado at Boulder. Images were acquired in the secondary electron mode with the microscope operating at 5 kV and a WD of 6 mm, and in the backscattered electron mode at 15 kV and WD 8 mm. EDXS analyses were performed at 20 kV and WD 8 mm.

To correlate Raman data with morphological characterization of the S(0) particles, we designed a correlative Raman-SEM protocol. A frozen unpreserved biofilm sample (PC1718) was rinsed three times with deionized water to remove salts and deposited on a microscope glass slide. The sample was air-dried, and the S(0) particles were first analyzed using Raman. The microscope slide was then coated with iridium and analyzed by SEM on a Field Emission Nova NanoSEM 630 (see details on SEM operations below). Raman-based crystallographic characterization of the S(0) minerals was performed before SEM imaging in order to prevent potential structural alteration in the dry, low-vacuum environment of the SEM chamber.

### Fourier-Transform Infrared Spectroscopy

For FTIR spectroscopy, 1 ml of unpreserved biofilm (sample PC1718) was rinsed three times with deionized water and dried in a vacuum oven at 60°C overnight. Then, 6 mg of dried sample and 100 mg KBr were ground and pelleted. FTIR measurements were conducted on a vertex 70 spectrometer (Bruker Optics) equipped with a deuterated triglycerine sulfate (DTGS) detector and a high-intensity water-cooled Globar source. Spectra were collected at 5 cm $^{-1}$  resolution (2.5 mm aperture) as an average of 100 scans using MVP PRO software (Harrick Scientific). The instrument was purged for 30 min before the first measurement to ensure baseline stability. The FTIR analyses were performed on three separate occasions on distinct aliquots of the biofilm, and the obtained spectra were similar between analyses. The spectra were baseline corrected using the “Rubber Band” algorithm within the OPUS 2.2 software. The experimental spectra were plotted with reference spectra for calcite and quartz (RRUFF database; Lafuente et al., 2015).

### Scanning Transmission X-Ray Microscopy and C K-Edge and S L-Edge X-Ray Absorption Spectroscopy

For STXM, 1 ml of unpreserved biofilm (samples PC1647 and PC1718) was rinsed three times with deionized water. A small drop of the suspension (~3  $\mu$ l) was deposited on a Formvar-coated 200 mesh Cu TEM grid (Ted Pella) and allowed to air-dry at ambient temperature. STXM analyses were performed on beamline 10ID-1 (SM) of the Canadian Light Source (Saskatoon, Canada), and beamline 11.0.2. of the Advanced

Light Source (ALS, Berkeley, CA). The X-ray beam was focused on the samples using a Fresnel zone plate objective and an order-sorting aperture yielding a focused X-ray beam spot of ~30 nm on the samples. After sample insertion in the STXM microscope, the chamber was evacuated to 100 mTorr and back-filled with He at ~1 atm pressure. Energy calibration was achieved using the well-resolved 3p Rydberg peak of gaseous CO<sub>2</sub> at 294.96 eV. Images, maps, and image stacks were acquired in the 260–340 eV (C K-edge) and 155–190 eV (S L-edge) energy ranges.

Scanning transmission X-ray microscopy data were processed using aXis2000 software. Maps of organic C were obtained by subtracting images obtained at 280 eV (i.e., below the C K-edge) and converted into optical density (OD) from OD-converted images at 288.2 eV (1 s→π\* electronic transitions in amide groups). Maps of S were obtained by subtracting OD-converted images obtained at 160 eV (i.e., below the S L-edge) from OD-converted images at 163.5 eV (energy of the S L<sub>3</sub>-edge). X-ray absorption near-edge structure (XANES) spectra were extracted from image stacks as explained in Cosmidis and Benzerara (2014). Different types of objects (e.g., cells, EPS, and sulfur particles) were visually identified in STXM images and maps. Regions of interest were manually selected, and their distinctive XANES spectra were extracted from the stacks and plotted. A linear background correction was applied to the XANES spectra at the C K-edge and the S L-edge, in the 260–280 eV region and 155–160 eV region, respectively, to eliminate the contributions of lower energy absorption edges.

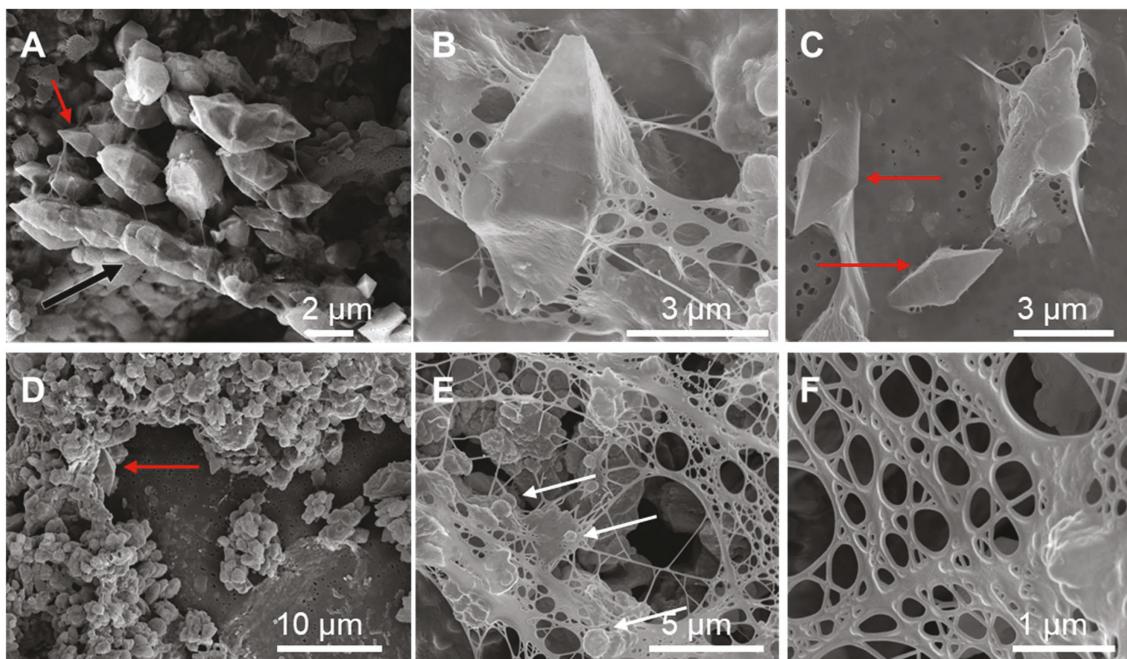
## RESULTS

### Morphology of Elemental Sulfur Particles in *Sulfurovum*-Dominanted Biofilms

Scanning electron microscopy imaging combined with EDXS analyses of the *Sulfurovum*-dominated biofilms revealed abundant sulfur-rich particles within a dense matrix of EPS (Figure 2; Supplementary Figures S1, S2). The particles appear as spheroids and bipyramidal crystals, which are sometimes fused into elongated chains (Figure 2A; Supplementary Figure S1). The particles may also adopt more irregular shapes. The EPS appears either film-like (Figure 2C) or as a web made from a network of thin threads (Figure 2F). Figure 3 shows the size distribution of S(0) spheroids in two biofilm samples [PC1647 ( $n = 1,354$ ) and PC1718 ( $n = 256$ )]. The diameters of the spheroids range from 0.1 to 3.5  $\mu$ m, with an average diameter of 1.31  $\mu$ m for PC1347 and 1.03  $\mu$ m for PC1718. Bipyramids are not abundant enough to plot their size distribution, but range between 1.9 and 4.8  $\mu$ m in length.

### Crystal Structure of Elemental Sulfur X-Ray Diffraction

X-ray diffraction analyses performed on sample PC1718 indicate that the biofilm contains S(0) present as orthorhombic  $\alpha$ -S<sub>8</sub> and monoclinic  $\beta$ -S<sub>8</sub> (Figure 4), as determined by comparison with S(0) XRD patterns from the ICDD database. Calcite and quartz minerals were also detected.



**FIGURE 2 |** Scanning electron microscopy images of sulfur particles associated with EPS in *Sulfurovum*-dominated streamer biofilms. Samples include (A) PC1647 and (B–F) PC1718. Red arrows point to bipyramidal S(0) crystals and white arrows point to S(0) spheroids. (A) shows an example of a chain formed by fused bipyramidal S(0) crystals (black arrow).

## Correlative Scanning Electron Microscopy and Raman Spectromicroscopy

In order to determine the relationship between S(0) crystal structure and particle morphology, Raman spectroscopy correlated with SEM imaging was performed on sample PC1718. Crystal structure identification was performed by comparison with low-frequency Raman spectra for different reference S(0) allotropes (Nims et al., 2019). The  $\alpha$ -S<sub>8</sub> allotrope can be identified using the low-frequency range Raman vibration modes at 28 cm<sup>-1</sup>, 44 cm<sup>-1</sup>, 51 cm<sup>-1</sup>, 63 cm<sup>-1</sup>, and a doublet at 82 cm<sup>-1</sup> and 88 cm<sup>-1</sup> (Figure 5C). The  $\beta$ -S<sub>8</sub> allotrope displays distinctive vibrations in the low-frequency range with a peak positioned at 82 cm<sup>-1</sup>, a doublet at 33 cm<sup>-1</sup> and 42 cm<sup>-1</sup>, and a shoulder at 60 cm<sup>-1</sup>. Spheroids were found to be composed of either  $\alpha$ -S<sub>8</sub> or  $\beta$ -S<sub>8</sub> (Figures 5A,B,D). These two allotropes were sometimes found in close association with each other (Figure 5E). We attempted to determine the crystal structure of S(0) bipyramids, but they could not be found in the sample during Raman analyses due to their relative rarity.

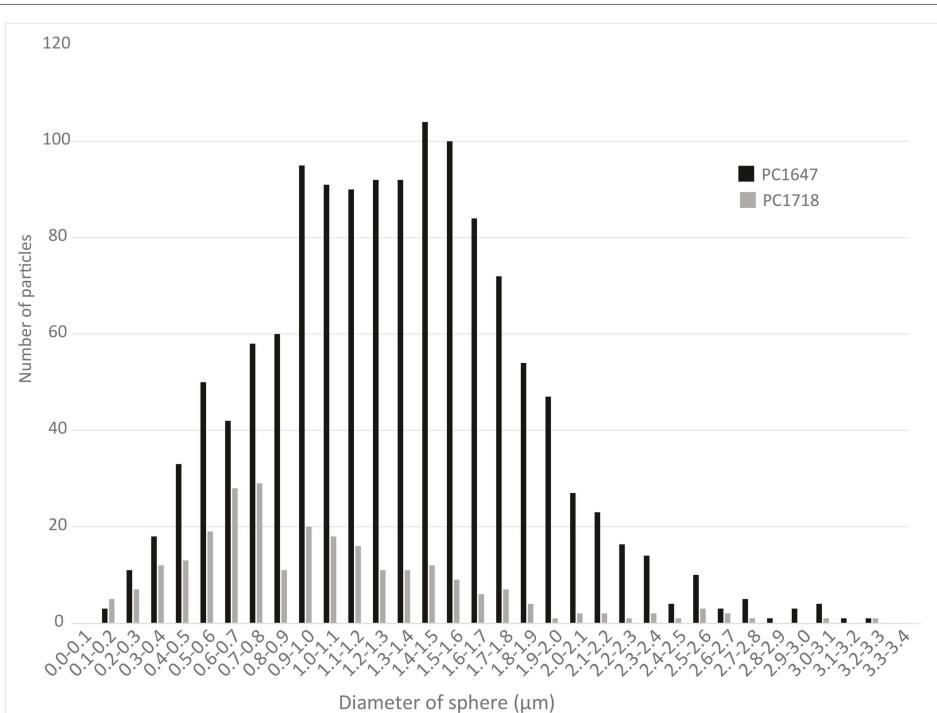
## Association of Elemental Sulfur With EPS and Encapsulating Organics

### Fourier-Transform Infrared Spectroscopy

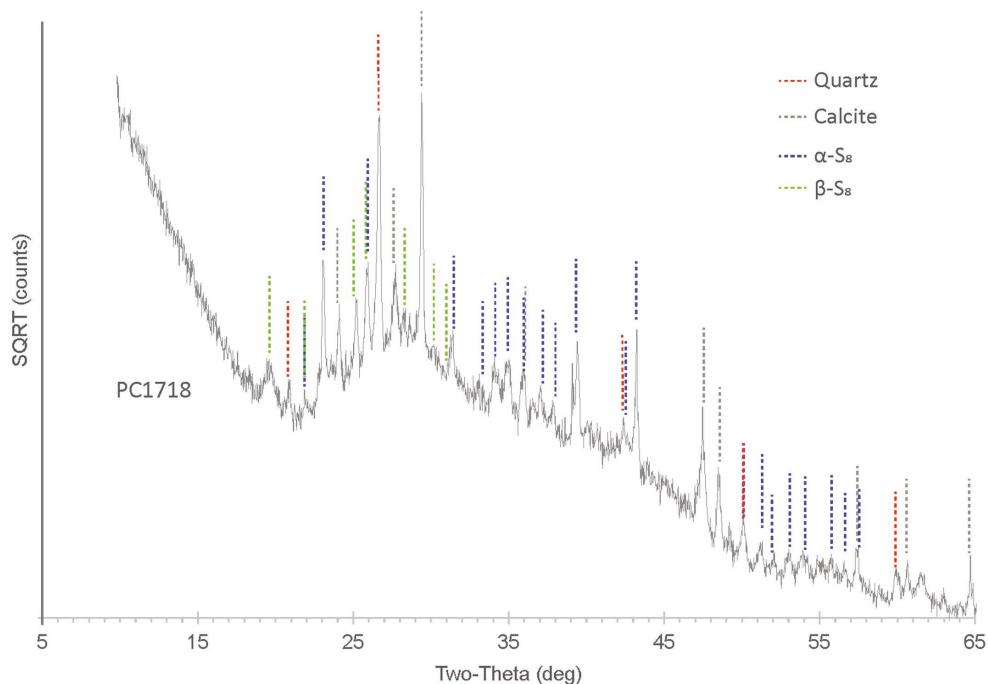
Fourier-transform infrared (FTIR) spectroscopy was performed on the unpreserved PC1718 sample (Figure 6). Reference spectra for calcite and quartz were used to assist in the interpretation of the FTIR data, since these minerals were identified in the sample based on XRD (Figure 4). The sample

spectrum displays a broad band centered around 3,400–3,430 cm<sup>-1</sup> corresponding to O–H stretching frequencies (Supplementary Figure S3). The broad peak around 1,750 cm<sup>-1</sup> corresponds to stretching of COOH or COOR in carboxylic acids and aromatic esters (Artz et al., 2008). Signal in this region is also attributed to C=O stretching in esters and fatty acids (Schmitt and Flemming, 1998). Peak signal around 1,650–1,600 cm<sup>-1</sup> is attributed to stretching of aromatic C=C or asymmetric C–O stretching in COO<sup>-</sup> (carboxylates). The presence of carboxylic acids in our sample is further supported by the presence of a peak at 1,426 cm<sup>-1</sup> originating from symmetric C=O stretching and OH deformation in COOH from carboxylates or carboxylic acid structures (Thomas, 1972). The band at 1,056 cm<sup>-1</sup> is attributed to phosphates in nucleic acids (Orhan Yanikan et al., 2020) and/or to quartz. A sharp peak at 879 cm<sup>-1</sup> is present, commonly interpreted as out of phase ring stretching (ring “breathing”) of aromatics (Artz et al., 2008). Peaks at 470 and 424 cm<sup>-1</sup> are attributed to S–S stretching in S<sub>8</sub> (Meyer, 1976; Steudel and Eckert, 2003). The presence of quartz is confirmed by peaks at 692, 776, and 795 cm<sup>-1</sup> and a shoulder at 1,160 cm<sup>-1</sup>. The peak at 516 cm<sup>-1</sup> is representative of calcite. Quartz peaks may overlap with the molecular vibrations of polysaccharides in the 1,200–900 cm<sup>-1</sup> region, which are typically observed in FTIR spectra of EPS (Naumann et al., 1991).

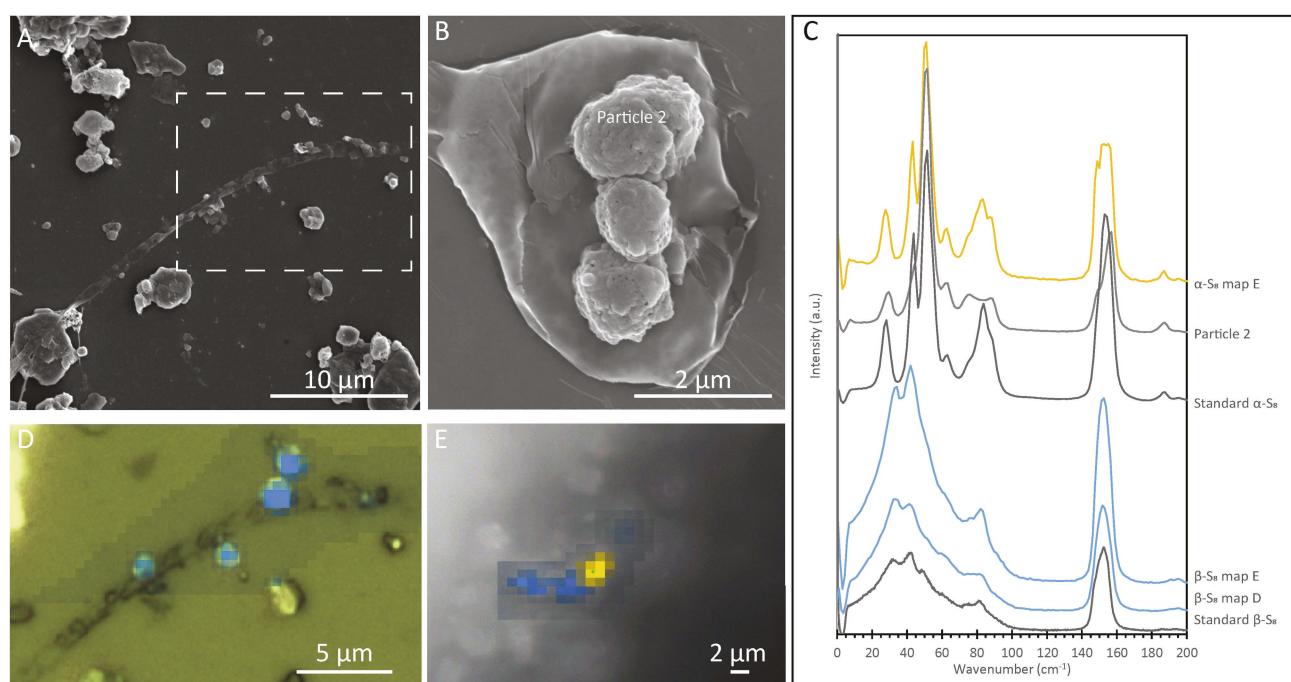
Fourier-transform infrared analysis thus confirmed the presence of quartz, calcite, and S(0) minerals in the biofilm, while signal from the organic material shows a composition dominated by carboxylic acids, carboxylates, and aromatic structures.



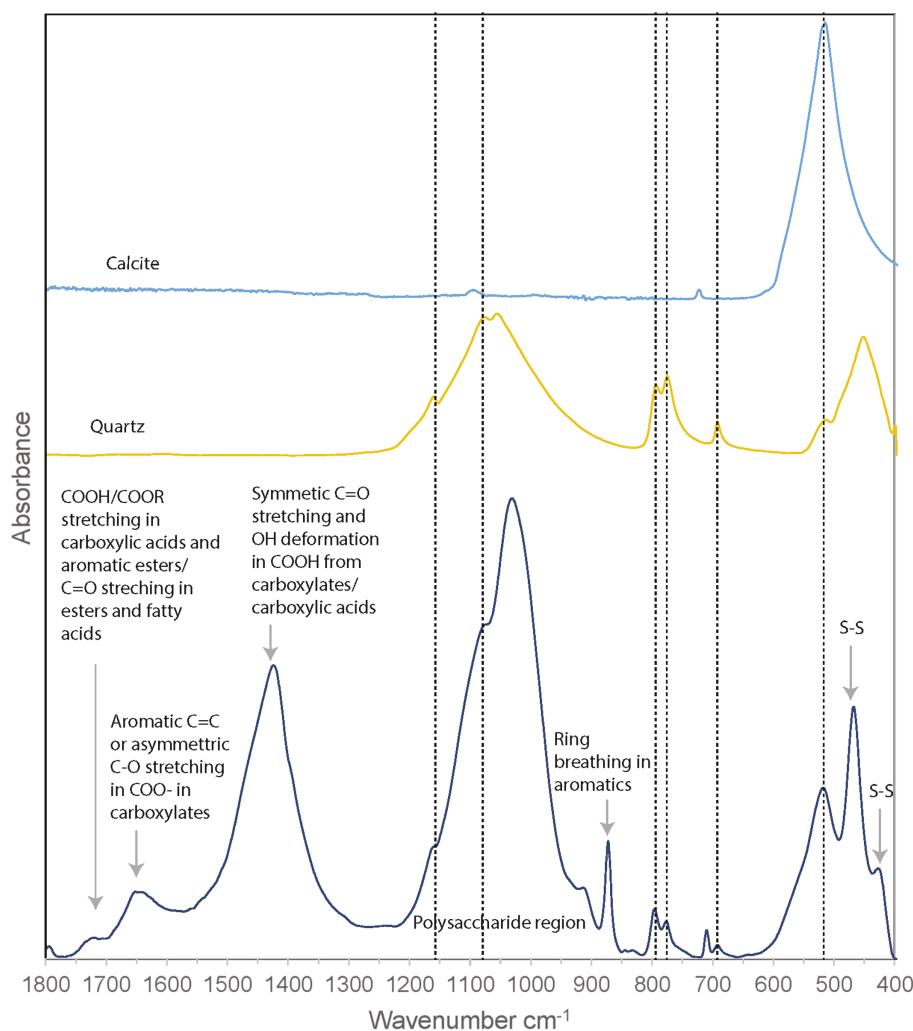
**FIGURE 3** | Histograms depicting the size distributions of spherical S(0) particles in the PC1647 and PC1718 biofilm samples.



**FIGURE 4** | XRD spectrum for biofilm sample PC1718. The peaks in the diffractogram include matches for  $\alpha$ -S<sub>8</sub>,  $\beta$ -S<sub>8</sub>, quartz, and calcite.



**FIGURE 5** | Correlative SEM and Raman analyses of a filamentous cell with extracellular S(0) in the *Sulfurovum*-dominated streamer biofilm (sample PC1718). The sample was air-dried in images (A), (B), and (D), while Raman analyses (E) were performed while the sample was wet. (A) SEM image with a white dashed box indicating the location of the Raman map in (D); (B) SEM image of particles in sample PC1718. " $\alpha$ -S<sub>8</sub> particle 2" indicates where the  $\alpha$ -S<sub>8</sub> Raman spectrum (labeled "Particle 2") in (C) was collected. (C) Ultra-low-frequency Raman spectra. Standard spectra for  $\alpha$ -S<sub>8</sub> and  $\beta$ -S<sub>8</sub> are labeled accordingly. (D) Light microscopy image with an overlay Raman map. The blue areas are rich in  $\beta$ -S<sub>8</sub>. (E) Light microscopy image with a Raman map overlaid. Yellow areas are rich in  $\alpha$ -S<sub>8</sub>, and blue areas are rich in  $\beta$ -S<sub>8</sub>.



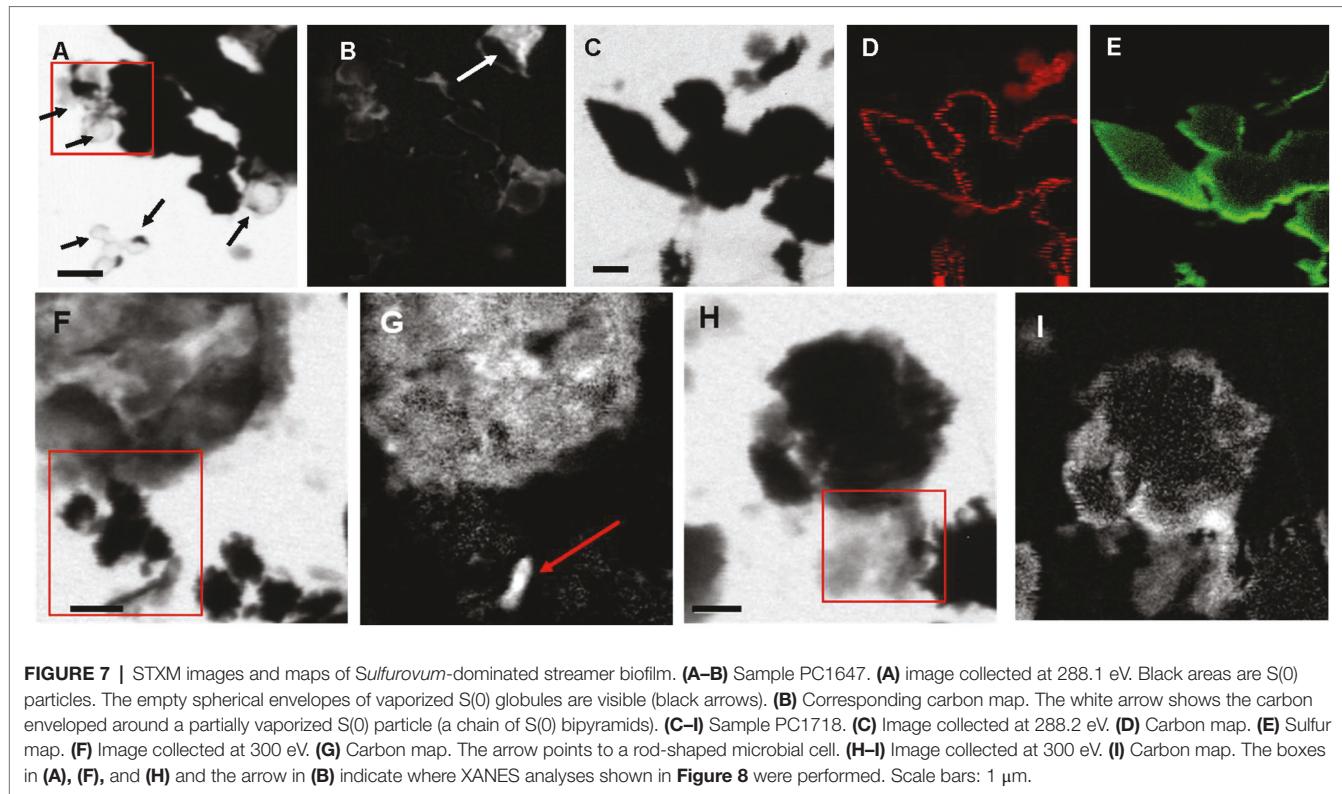
**FIGURE 6 |** FTIR spectrum of *Sulfurovum*-dominated streamer biofilm (sample PC1718), plotted with reference spectra for calcite and quartz. Dashed vertical lines indicate the positions of the peaks corresponding to calcite and quartz.

### Scanning Transmission X-Ray Microscopy at the C K-Edge and S L-Edge

Scanning transmission X-ray microscopy analyses at the S L-edge of the *Sulfurovum*-dominated biofilms (samples PC1647 and PC1718) confirmed that S(0) is the only form of particulate sulfur present in the samples (Supplementary Figure S4). C K-edge analyses showed the presence of bacteria along with two types of extracellular carbon materials: abundant EPS (as already observed in SEM images) and thin envelopes around S(0) particles (Figure 7). The thin organic envelopes are especially visible in STXM images and maps in places where S(0) particles were vaporized due to the low pressure of the STXM chamber and X-ray beam damage (black arrows in Figure 7A).

All C K-edge XANES spectra display peaks at 285–285.2 eV, characteristic of  $1s \rightarrow \pi_{C=C}^*$  transitions in either aromatic or unsaturated carbon; Figure 8). The main peak in all spectra is located at 288.2 eV, corresponding to  $1s \rightarrow \pi^*$  transitions

in amide groups of proteins (Benzerara et al., 2004; Haberstroh et al., 2006; Chan et al., 2011). The presence of amides is in contradiction with FTIR results which did not clearly detect this functional group. This discrepancy may be due to the fact that we focused our STXM analyses on clearly identifiable features, such as microbial cells and organic envelopes around S(0) particles, while the signal from these small features may have been diluted by the signal of the more abundant EPS in bulk FTIR analyses. All spectra furthermore display a shoulder at 287.5 eV, corresponding to  $3s \rightarrow \sigma^*$  transitions in aliphatics (Haberstroh et al., 2006; Lehmann, 2009), and a small peak at 289.4 eV, corresponding to  $1s \rightarrow 3p/\sigma^*$  transitions in hydroxylic groups (Brandes et al., 2004). Some spectra (mostly EPS) have a peak at 288.5 eV, attributed to the  $1s \rightarrow \pi_{C=O}^*$  electronic transitions in carboxylics (Cody et al., 1998; Boyce et al., 2002; Chan et al., 2011). A small peak at 286.6 eV, representative of  $1s \rightarrow \pi^*$  transitions in phenolic groups, ketones, carboxylates,



**FIGURE 7 |** STXM images and maps of *Sulfurovum*-dominated streamer biofilm. **(A–B)** Sample PC1647. **(A)** image collected at 288.1 eV. Black areas are S(0) particles. The empty spherical envelopes of vaporized S(0) globules are visible (black arrows). **(B)** Corresponding carbon map. The white arrow shows the carbon enveloped around a partially vaporized S(0) particle (a chain of S(0) bipyramids). **(C–I)** Sample PC1718. **(C)** Image collected at 288.2 eV. **(D)** Carbon map. **(E)** Sulfur map. **(F)** Image collected at 300 eV. **(G)** Carbon map. The arrow points to a rod-shaped microbial cell. **(H–I)** Image collected at 300 eV. **(I)** Carbon map. The boxes in **(A)**, **(F)**, and **(H)** and the arrow in **(B)** indicate where XANES analyses shown in **Figure 8** were performed. Scale bars: 1  $\mu$ m.

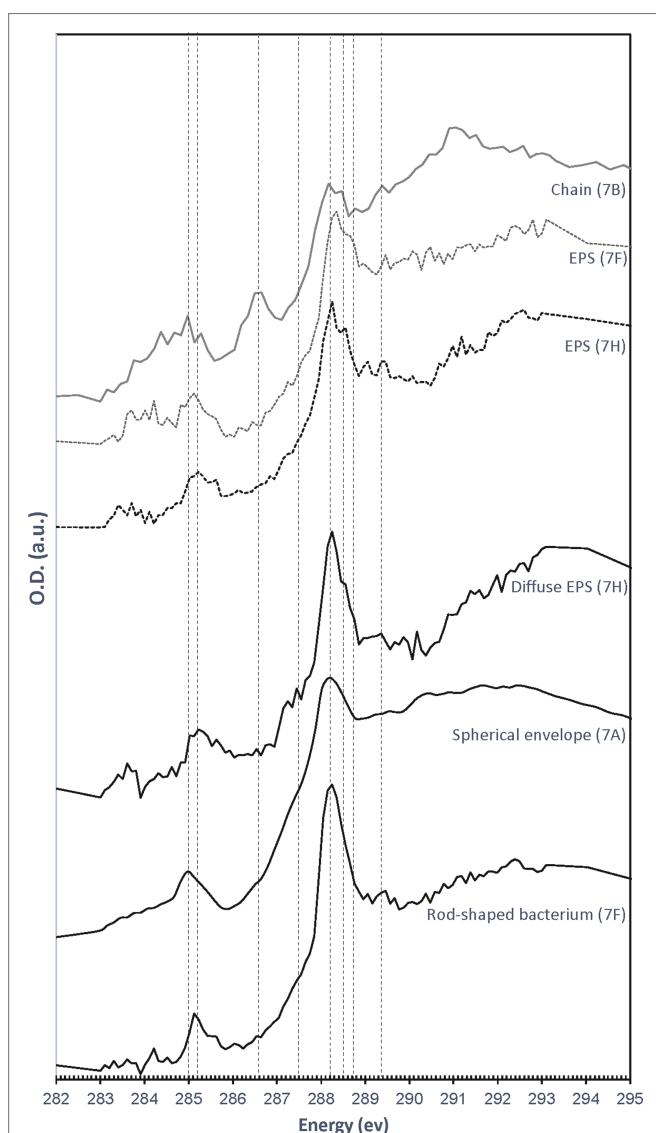
or aldehydes (Myneni, 2002; Lehmann, 2009; Moffet et al., 2011; Cosmidis et al., 2019), is only present in the organic envelope of a S(0) chain (Figures 7B, 8). Spectra of EPS and S(0) envelopes sometimes display a shoulder at 288.7 eV, interpreted as  $\text{C } 1s \rightarrow \pi_{\text{C}=\text{O}}^*$  electronic signature of acidic polysaccharides (Toner et al., 2009; Chan et al., 2011).

## DISCUSSION

### Properties of the Biofilm S(0) Particles

We described S(0) particles forming within biofilms present in microaerophilic, sulfide-rich subsurface stream. The biofilms are dominated by *Campylobacterota* most closely related to *Sulfurovum* species (Jones et al., 2008; Macalady et al., 2008; Hamilton et al., 2015), which are known to oxidize sulfide and thiosulfate to sulfate and/or extracellular S(0) (e.g., Campbell et al., 2006). Microbial sulfide oxidation rates are typically several orders of magnitude higher than chemical oxidation by molecular oxygen (Luther et al., 2011) suggesting that S(0) particles in the Frasassi biofilms are primarily the product of microbial S-oxidation. However, mineral nucleation and growth within biofilms are often influenced by surrounding EPS, which may result in particle shapes, sizes, or crystal structures that differ from those of inorganically precipitated minerals (Tourney and Ngwenya, 2014). Here, we summarize our observations of the S(0) particles in the Frasassi biofilms and suggest how formation within EPS may have influenced S(0) properties. The results are also summarized in Figure 9.

The samples we examined contained both S(0) spheroids and bipyramids, alongside more irregularly shaped particles (Figures 2, 7). Both spherical and bipyramidal morphologies have been observed in microbial biomineralization experiments, microbe-free organomineralization experiments, and inorganically precipitated S(0). Extracellular S(0) spheroids or globules are formed by diverse bacteria (Dahl and Prange, 2006; Marnocha et al., 2016; Cron et al., 2019). S(0) spheroids can also be formed through chemical precipitation of S(0) in the presence (Cosmidis et al., 2019) or in the absence (Marnocha et al., 2019) of organic compounds. Previous studies have suggested that bipyramids are typical of S(0) precipitated in the absence of organics (Steudel, 2003). However, as noted above, bipyramids have also been observed in S(0) organomineralization experiments with humic acids (Cosmidis et al., 2019) and in cultures of *C. tepidum* (Marnocha et al., 2019). The biofilm samples we examined contained S(0) spheroids ranging from 0.2 to 3.3  $\mu$ m (Figure 3) with median particle sizes near 1  $\mu$ m. The PC1647 sample had larger S(0) spheroids. Larger particle size could be controlled by the age of the biofilm, or possibly by the growth of the biofilm on the twine support. Further analysis would be needed to confirm whether biofilm age and the nature of attachment surfaces influence particle size. S(0) particles with sizes ranging from <0.2  $\mu$ m to 10  $\mu$ m have been observed in other natural environments (Findlay et al., 2014; Lau et al., 2017), in microbial cultures (Cron et al., 2019; Marnocha et al., 2019), in inorganically precipitated S(0) (Meyer, 1976; Garcia and Druschel, 2014), and in S(0)



**FIGURE 8 |** C K-edge XANES spectra obtained on the particles and cells shown in **Figure 7**. Dashed lines correspond to absorption energies for different organic functional groups: 285 eV ( $1s \rightarrow \pi_{C=C}^*$  transitions in aromatics or unsaturated carbon), 285.2 eV ( $1s \rightarrow \pi_{C=C}^*$  transitions in aromatics), 286.6 eV ( $1s \rightarrow \pi^*$  transitions in ketones, pyridines, and phenols), 287.5 eV ( $3s \rightarrow \sigma^*$  transitions in aliphatics), 288.2 eV ( $1s \rightarrow \pi^*$  transitions in amides), 288.5 eV ( $1s \rightarrow \pi_{C=O}^*$  transitions in carboxylic acids), 288.7 eV ( $1s \rightarrow \pi_{C=O}^*$  transitions in carboxyls in acidic polysaccharides), and 289.4 eV ( $1s \rightarrow 3p/\sigma^*$  transitions in alcohols, ethers, and hydroxylated aliphatic compounds).

organomineralization experiments (Cosmidis et al., 2019). Particle sizes or morphologies of the S(0) described here are thus not particularly characteristic of their formation within microbial biofilms.

On the other hand, we observed both  $\alpha$ -S<sub>8</sub> and  $\beta$ -S<sub>8</sub> S(0) crystal structures in biofilm sample PC1718 (**Figures 4, 5**). The monoclinic sulfur allotrope  $\beta$ -S<sub>8</sub> is thermodynamically unstable at temperatures lower than 96°C, as opposed to orthorhombic  $\alpha$ -S<sub>8</sub> which is the stable structure at room

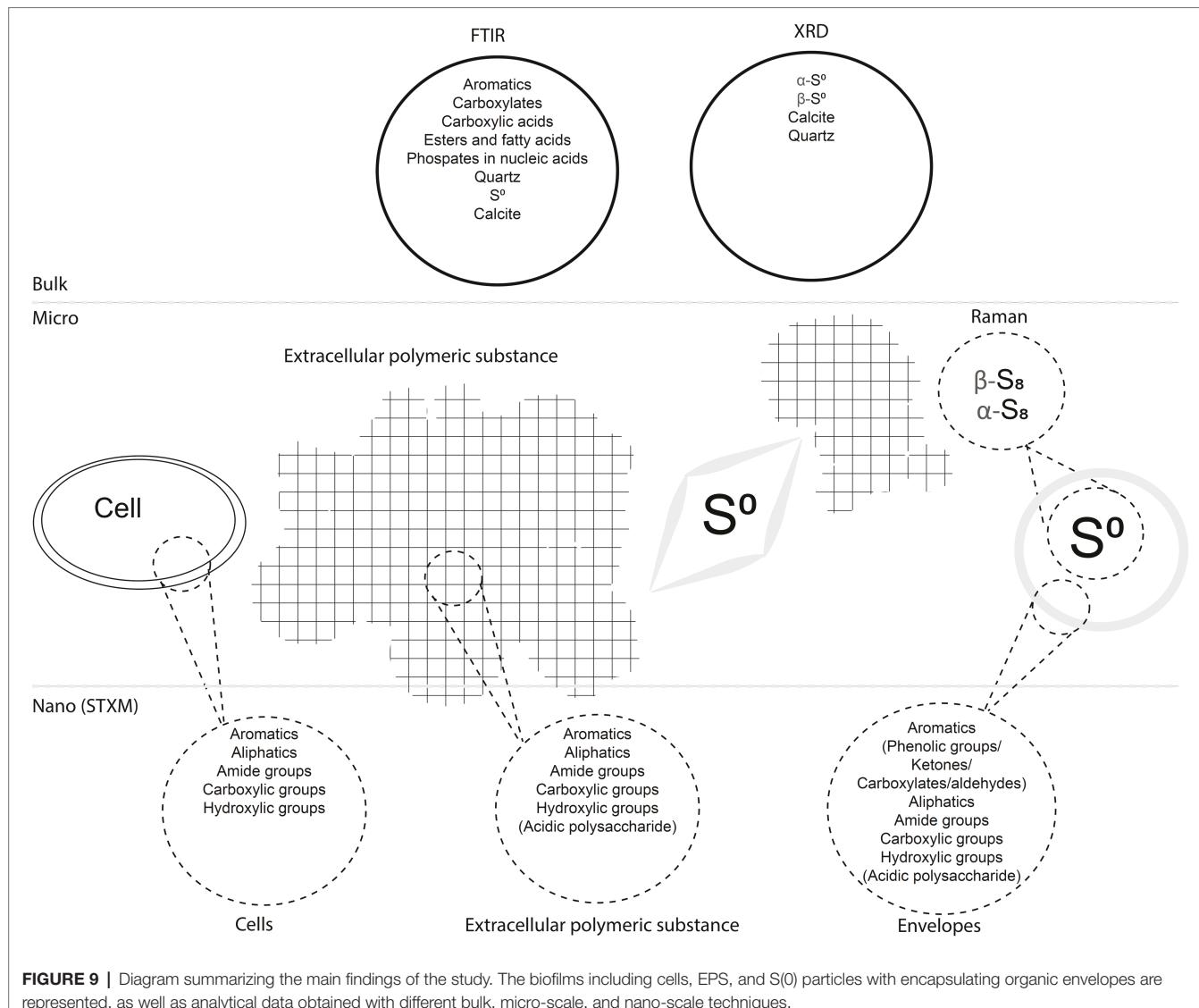
temperature (Steudel, 2003). In laboratory studies, metastable monoclinic S(0) phases were formed abiotically at low temperature in the presence of organics (Guo et al., 2006; Choudhury et al., 2013; Moon et al., 2013; Cosmidis et al., 2019). In microbial incubations, soluble organic compounds produced by *S. kujjiense* were found to play an important role in the formation and stabilization of extracellular  $\beta$ -S<sub>8</sub> particles in cultures and in spent media containing soluble organics (Cron et al., 2019).  $\beta$ -S<sub>8</sub> was also previously described from a low-temperature natural environment, and it was proposed that it was stabilized by its intimate association with organic matter (Lau et al., 2017).

Based on FTIR and XANES, the organic material associated with S(0) in the biofilms has a complex composition dominated by carboxylic acids, amides, aromatics, and aliphatic compounds (**Figures 6, 8**). EPS is typically composed of DNA, polysaccharides, and lipids (Devaraj et al., 2019). These would include polycyclic aromatic compounds and carboxylic substitutes (Flemming and Wingender, 2010). It is important to note that the organic compounds we observed are not merely “associated,” but directly encapsulating S(0) particles (**Figures 7A,B**). Consistent with this observation, previous work describing organics associated with S(0) formed in organomineralization experiments demonstrated the presence of carboxylic (Cosmidis et al., 2019) and amide groups (Cosmidis and Templeton, 2016) in organic envelopes closely encapsulating S(0) minerals. Pure cultures of the *Sulfurovum* relative *S. kujjiense* also produced an amide-rich envelope around extracellular S(0) globules (Cron et al., 2019). Similarly, extracellular S(0) produced by *C. tepidum* was encapsulated in organic envelopes composed of proteins and polysaccharides (Marnocha et al., 2019).

Both organic envelopes around S(0) particles and the presence of the metastable allotrope  $\beta$ -S<sub>8</sub> thus confirm an important role for organic-mineral interactions in extracellular S(0) mineralization. We therefore hypothesize that microbially derived extracellular organics (EPS) are critical for the formation and preservation of S(0) particles in the *Sulfurovum*-dominated biofilms at Frasassi, raising the interesting question below.

## S(0) Organomineralization and Storage in Biofilms: A New Role for EPS?

Extracellular formation of biominerals in close interaction with organic polymeric structures has been documented for different types of systems, for instance precipitation of calcium carbonates in microbial mats (Dupraz et al., 2009) or on diatom EPS (Stanton et al., 2021), or iron-(oxyhydr)oxide mineralization on polymeric bacterial sheaths and stalks (Chan et al., 2009, 2011). While in some cases mineral precipitation on extracellular organics may be “unintended and uncontrolled” (Frankel and Bazylinski, 2003), in other cases specific organic structures are involved in directing extracellular biominerization and this process plays a crucial role in metabolism, growth, and/or survival (Miot et al., 2009; Chan et al., 2016).



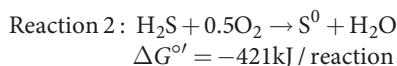
**FIGURE 9** | Diagram summarizing the main findings of the study. The biofilms including cells, EPS, and S(0) particles with encapsulating organic envelopes are represented, as well as analytical data obtained with different bulk, micro-scale, and nano-scale techniques.

In the Frasassi *Sulfurovum*-rich biofilms, EPS appears to influence S(0) formed as a result of microbial S-oxidation, through the stabilization of otherwise unstable mineral phases. Organics have been shown to favor the formation and stabilization of S(0) minerals through S(0) organomineralization (Cosmidis and Templeton, 2016; Cosmidis et al., 2019). S(0) organomineralization may occur with diverse types of organic molecules (e.g., amino-acids, sugars, and humic acids; Cosmidis et al., 2019), as well as with soluble organic compounds produced by different types of bacteria (Cron et al., 2019). It remains to be determined whether specific organics are produced by the Frasassi biofilm microbial community to direct S(0) formation and stabilization. Given that organic matter is energetically costly to produce (Amend et al., 2013; Jayathilake et al., 2017), particularly for autotrophs like *Sulfurovum* (LaRowe and Amend, 2015, 2016), we consider it unlikely that S(0) encapsulation and particle trapping in the biofilm is accidental. Holdfasts in colonizing *Sulfurovum* populations appear

to be made of S(0) rather than organic matter, eliminating a potential role for EPS in biofilm attachment. These observations suggest that binding and encapsulating S(0) in EPS may represent a particular ecological strategy.

We can speculate about the ecological role that EPS and embedded S(0) particles might play in the survival and dispersal of *Sulfurovum* populations in the cave habitat. In this environment, *Sulfurovum*-rich biofilms form only in turbulent stream flow near the air-water interface and are absent when appropriate support structures are unavailable in that position. The entrenched cave streams hosting the biofilms are subject to rapid, weather-related changes in water levels that frequently wash out or drown the biofilms. Thus cells that are producing S(0) under ideal conditions (Reaction 2) may experience rapid changes in O<sub>2</sub> availability. Under O<sub>2</sub> starvation conditions, bioavailable S(0) in the biofilm could enable survival and possibly even growth.

Hamilton et al. (2015) noted that *Sulfurovum* metagenome-assembled genomes (MAGs) obtained from biofilms collected at the same sampling site ubiquitously contained genes that would allow H<sub>2</sub> and formate oxidation with S(0) as an electron acceptor (Reaction 1).



Although Reaction 1 yields much less energy than Reaction 2 under standard conditions (and likely also under *in situ* conditions), the ability to utilize stored S(0) may represent an important ecological advantage to the cave *Sulfurovum* populations, and capturing and storing S(0) in EPS may be an important element in the life strategy of this group. This scenario will be tested using metatranscriptomics in the future work. If so, energy storage through stabilization of electron donors for energy metabolism may be a previously underappreciated role for EPS in biofilms.

Extracellular polymeric substances in the Frasassi biofilms described here appear to be responsible for at least two properties of the S(0) minerals: their crystal structure (with the stabilization of β-S<sub>8</sub>) and organic encapsulation. It is now well established that the properties of S(0) particles play a crucial role in their utilization by bacteria. For instance, particle size, surface area, and S<sup>0</sup> composition and structure affect S(0) oxidation rates by *Thiobacillus albertis* (Laishley et al., 1986). *Allochromatium vinosum* grown on S(0) show a preference for polymeric sulfur over commercial crystalline S<sub>8</sub>, which they are unable to uptake (Franz et al., 2007). Preference for polymeric sulfur utilization over S<sub>8</sub> was also evidenced in natural mats of chemotrophic S-oxidizers (Engel et al., 2007). Incubation experiments of natural freshwater communities with different sulfur sources showed a preference for the utilization of a reactive form of colloidal S(0) – possibly polythionates – over S<sub>8</sub> (Findlay and Kamyshny, 2017). Recently, it was shown that *C. tepidum* can grow from the oxidation of its own biogenic S(0) globules but not from oxidation of commercial sulfur, crystalline S(0), or inorganically precipitated colloidal S(0), which is mineralogically very similar to biogenic S(0) globules but does not have an organic coating (Marnocha et al., 2016, 2019). It is possible that S(0) formation within EPS favors storage in a form more readily utilized by the cells, by favoring metastable S(0) structures and by making S(0) particles hydrophilic, allowing interaction with the cell surface (Marnocha et al., 2019).

## CONCLUSION

We studied a natural *Sulfurovum*-rich S(0)-producing biofilm to investigate processes that influence the precipitation and stabilization of extracellular S(0) in the environment. We found that sulfur particles in the biofilm are encapsulated within

organic envelopes and that some of the particles have an unstable crystal structure (the high-temperature allotrope β-S<sub>8</sub>). These characteristics have also been observed in S(0) produced by organomineralization. Our results suggest that EPS within the biofilm stabilize S(0) particles, preventing their dispersal away from the biofilm and influencing their structural and surface properties. Future studies will be needed in order to determine the ecological importance of this process and its impact on biogeochemical sulfur cycling in the Frasassi cave and in other environments.

## DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/**Supplementary Material**, and further inquiries can be directed to the corresponding author.

## AUTHOR CONTRIBUTIONS

BC, JM, and JC designed the study. BC and JM collected the samples. BC and JC performed the analyses. All authors participated in data interpretation. BC wrote the manuscript with input from JC and JM. All authors contributed to the article and approved the submitted version.

## FUNDING

Support for this study was provided by the Penn State Department of Geosciences through startup funding to JC. BC was supported by the Penn State Department of Geosciences Distinguished Postdoctoral Fellowship. Sample collection and fieldwork were funded by an NSF grant to JM (EAR1252128).

## ACKNOWLEDGMENTS

We thank C. Nims and M. Wetherington (Penn State University) for help with Raman analyses and T. J. Zimudzi (Penn State University) for help with FTIR measurements. The use of the SSRL, SLAC National Accelerator Laboratory, is supported by the US Department of Energy, Office of Science, Office of Basic Energy Sciences under Contract No. DE-AC02-76SF00515. We thank J. Wang for providing support on STXM beamline SM of the Canadian Light Source (CLS, Saskatoon, Canada). CLS is supported by the Canada Foundation for Innovation, Natural Sciences and Engineering Research Council of Canada, the University of Saskatchewan, the Government of Saskatchewan, Western Economic Diversification Canada, the National Research Council Canada, and the Canadian Institutes of Health Research. The authors are grateful to A. Montanari for providing facilities and laboratory space at the Osservatorio Geologico di Coldigioco in Apiro, Italy. Fieldwork was carried out in collaboration with C. Chan, C. Clark, and P. Henri with assistance from

M. Mainiero, S. Mariani, and S. Recanatini. Melissa Peacock (Salish Sea Research Center) is thanked for providing funding to BC for publication fees through the National Science Foundation Tribal Colleges and Universities Program Award No. 1840199.

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## SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fmicb.2021.720101/full#supplementary-material>

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