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Low frequency Raman Spectroscopy for micron-scale and *in vivo* characterization of elemental sulfur in microbial samples

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Elemental sulfur (S(0)) is an important intermediate of the sulfur cycle and is generated by chemical and biological sulfide oxidation. Raman spectromicroscopy can be applied to environmental samples for the detection of S(0), as a practical non-destructive micron-scale method for use on wet material and living cells. Technical advances in filter materials enable the acquisition of ultra-low frequency (ULF) Raman measurements in the 10–100 cm⁻¹ range using a single-stage spectrometer. Here we demonstrate the potency of ULF Raman spectromicroscopy to harness the external vibrational modes of previously unrecognized S(0) structures present in environmental samples. We investigate the chemical and structural nature of intracellular S(0) granules stored within environmental mats of sulfur-oxidizing γ -Proteobacteria (*Thiothrix*). *In vivo* intracellular ULF scans indicate the presence of amorphous cyclooctasulfur (S₈), clarifying enduring uncertainties regarding the content of microbial sulfur storage globules. Raman scattering of extracellular sulfur clusters in *Thiothrix* mats furthermore reveals an unexpected abundance of metastable β -S₈ and γ -S₈, in addition to the stable α -S₈ allotrope. We propose ULF Raman spectroscopy as a powerful method for the micron-scale determination of S(0) structure in natural and laboratory systems, with a promising potential to shine new light on environmental microbial and chemical sulfur cycling mechanisms.

Elemental sulfur S(0) is prevalent across a diverse number of sulfide-rich environments from geothermal¹ and cold sulfidic springs², deep-sea hydrothermal vents³, to salt marshes⁴, marine sediments^{5,6}, pyroclastic ash emissions⁷, mangrove swamps⁸, deep-sea hydrocarbon seeps⁹, and waste-water treatment plants¹⁰. Its presence in the environment is often associated with microbial oxidation of reduced sulfur compounds¹¹. Elemental sulfur can also be formed through abiogenic processes when sulfide is oxidized by molecular oxygen, possibly catalyzed by oxidized metals¹². S(0) plays a central role in the sulfur biogeochemical cycle, and is used as an energy source by a variety of sulfur-oxidizing, sulfate-reducing, and sulfur-disproportionating bacteria¹³.

Under Earth's surface conditions, S(0) exists in many forms, from cyclic octamers (S₈ rings) that crystallize in different structures or allotropes, to sulfur chains with varying numbers of S-S bonds, and polysulfides (S_n²⁻)^{5,14,15}. Characterization of S(0) in natural samples can be undercut by instabilities in its frangible chemical and structural properties. The allotropic enantiotropy of S₈, its ability to catenate¹⁶, and many possible reduction or oxidation reactions make sulfur a difficult material for analysis. These difficulties have so far prevented a consensus on the speciation and structure of biological S(0) stored as intracellular globules in bacteria, which has been described as “S₈ rings”¹⁷, “solid S₈”¹⁸, “microcrystalline S₈”¹⁹, “liquid sulfur”²⁰, “a mixture of polysulfides and cyclooctasulfur”²¹, and sulfur chains associated with “unidentified organic residues”²². Considerations on the changing nature of microbial sulfur storage, where “the speciation of stored sulfur varies under different

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ecophysiological conditions²¹, and the variance in sulfur speciation—cyclooctasulfur, sulfur chains, and polythionates—contingent on the sulfur-oxidizing metabolism²³, suggest the necessity of *in vivo* methods to finally elucidate this question.

A series of publications has debated the application of synchrotron-based X-ray absorption near edge structure spectroscopy (XANES) at the sulfur K-edge to investigate the nature of intracellular sulfur globules^{17,18,22–24}. Shortcomings of this method include possible distortions of the XANES spectra due to experimental artifacts¹⁷, the inability to discriminate different crystal structures of S₈, and the infeasibility of *in vivo* measurements. X-ray diffraction (XRD) allows differentiation between S₈ structures, but as a bulk method, it requires an abundance of material and does not provide spatial information on S(0) distribution within a sample. Micro- and nano-scale analyses by scanning electron microscopy (SEM), and transmission electron microscopy (TEM) also present significant drawbacks, since the vacuum conditions of the SEM and TEM chambers can cause the sublimation of S(0), while the electron beam can rapidly transform or burn sulfur material¹⁹.

Raman spectromicroscopy provides a non-destructive analytical technique in studies probing the chemical composition and speciation of biological sulfur. Raman scattering circumvents many of the problems associated with other characterization methods, as measurements can be collected on solid, liquid, and live samples at room temperature and atmospheric pressure. Additionally, Raman mapping produces high spatial (~1 μm) and spectral resolution analyses for detailed insights on the distribution and speciation of sulfur in the sample material. Characteristic internal vibrational (molecular) spectra make S(0) particularly easy to detect and characterize with Raman scattering^{9,19,21,25–27}.

Innovation in Raman filter technology over the past decade has enabled newfound access to the ultra low-frequency (ULF) modes of the Raman spectrum²⁸. The introduction of volume Bragg gratings has sanctioned measurements vicinal to the frequency of the laser line²⁹, reaching approximately 10 cm⁻¹. However, these new tools and advancements in the spectroscopy field have hitherto not been integrated within the earth and environmental science disciplines.

Most, if not all, biochemical, geomicrobiological, and kinetics studies using Raman analysis^{8,19,21,27,30,31} only focus on the properties of the S₈ internal frequency regions of the Raman spectrum (>80 cm⁻¹). In the absence of ULF spectral region, past research characterizing S(0) inclusions of sulfur bacteria through Raman spectroscopy relied on shifts in peak position, full width at half maximum (FWHM), and relative peak intensity^{19,21}, which produced ambiguous conclusions on the chemical speciation of the biomineralized S(0), and no information on crystal structure. We argue that an examination of Raman spectra of sulfur beyond the intramolecular vibrational modes is required to discriminate between different possible sulfur forms in biological S(0).

Our work uses low-frequency (<100 cm⁻¹) ULF Raman scattering as a tool to collect micro-scale spectral information on the external molecular structure and crystal phonons of S₈. Combined with analyses on the internal mode peak assignments, this study explores novel spectral identification of metastable allotropic forms of S₈, while investigating the *in vivo* speciation and structure of intracellular S(0) in a natural community of sulfur bacteria. Comprehensive inspection of Raman low frequency phases improves our ability to pinpoint and characterize the amorphous or crystalline forms of S(0) within environmental samples, while probing the potential sources of these sulfur particles.

Understanding the nature and distribution of sulfur could better constrain the conditions required for S(0) formation and preservation in biogeochemical processes and the mechanisms of microbial and chemical sulfur cycling.

Results

Reference spectra – External and internal S₈ frequencies, molten sulfur, and polysulfides. *Solid, S₈ allotropes.* Raman spectroscopy detects the distinct vibrational modes of the S₈ molecular structure. Figure 1 is a plot of the Raman spectra of three reference solid S₈ allotropes. Low frequency solid S₈ peak assignments measured in this study can be found in Supplementary Table 1.

All three solid S₈ allotropes have similar internal, high frequency vibrations with a few minor spectral variations. Overall, scans of the reference S₈ samples indicate characteristic Raman active peaks, with internal vibrations indicated by features at 153 cm⁻¹, 220 cm⁻¹, and 473 cm⁻¹. The vibrational modes at ~150 cm⁻¹ and 220 cm⁻¹ represent asymmetric S-S bending and symmetric S-S bending respectively. The peak at 473 cm⁻¹, with a shoulder feature at ~467 cm⁻¹, represents S-S bond stretching of the S₈ ring structure³². A broad feature at ~440 cm⁻¹ is present in all solid allotropes which also corresponds to S-S stretching modes; the α-S₈ spectrum presents a more defined peak at this position. The internal vibrations of metastable β-S₈ mirror the other solid allotropes, however, the β-S₈ asymmetric bending and the S-S stretching peaks present a minor broadening of FWHM in comparison to the other spectra. Additionally, in all normalized solid reference spectra, the relationship between the two bending features remains consistent, where the asymmetric bending peak (~150 cm⁻¹) is invariably less intense than the symmetric bending mode (~220 cm⁻¹). The three reference allotropes also present distinct asymmetric bending (~150) peaks, including variation in FWHM, relative intensities, and minor peak components, however, these peaks differences are inconsistent across several measurements.

The intra-molecular vibrations of γ-S₈ are consistent with the other S₈ reference allotropes, with a minor but well-defined peak present at 248 cm⁻¹ in both the γ-S₈, and the α-S₈, scans. γ-S₈ presents a shoulder at ~216 cm⁻¹ where α-S₈ and β-S₈ indicate a distinct peak in same position. The γ-S₈ S-S stretching peak at 473 cm⁻¹ is broadened and shows a more pronounced shoulder feature at 468 cm⁻¹ than β- and α-S₈.

The low frequency wavenumber region reveals major spectral differences between the solid S₈ references. The external vibrations of α-S₈ in the low frequency range are expressed by prominent peaks positioned at: 28 cm⁻¹, 44 cm⁻¹, 51 cm⁻¹, 63 cm⁻¹, and a doublet at 82 and 88 cm⁻¹. Orthorhombic α-S₈ is easily distinguished from the other S₈ allotropes in the identification of a prominent librational feature at 28–30 cm⁻¹ (depending on the scan)³³. Several bands display doublet structures that can be ascribed to the crystal-field effects of rhombic S₈¹⁵.

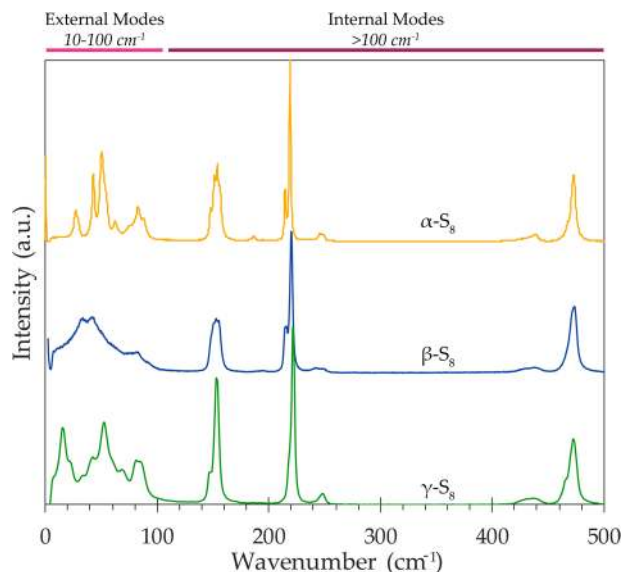


Figure 1. Raman spectra of reference standards of solid cyclooctasulfur allotropes: α -S₈ (yellow), β -S₈ (blue), and γ -S₈ (green), all normalized to the 473 cm⁻¹ peak. Wavenumber ranges for external and internal vibrational modes are indicated.

The main low frequency vibrations of β -S₈ yield broad, widened doublet feature with peaks at: 33 cm⁻¹, 42 cm⁻¹, 82 cm⁻¹, and a shoulder at 60 cm⁻¹ (Fig. 1). The low frequency spectrum measured in this work shares similar features with the β -S₈ spectra published in previous Raman scattering studies^{34,35}. Here we present the first published γ -S₈ Raman spectrum which includes both external and internal modes (Fig. 1). The inter-molecular vibrations of γ -S₈ in the low frequency region include features at: 15 cm⁻¹, 22 cm⁻¹, 33 cm⁻¹, 41 cm⁻¹, 52 cm⁻¹, 68 cm⁻¹, and a doublet peak at 83 and 85 cm⁻¹. The external diagnostic peak of γ -S₈ at 15 cm⁻¹ is the lowest wavenumber feature apparent in any of the solid S₈ reference allotropes.

Molten S₈ – Liquid sulfur. High temperature *in-situ* Raman experiments probing the composition and structure of molten sulfur are presented in Fig. 2. Measurements of powder α -S₈ standard initiate the experiment, with Raman spectra acquired as temperature increases incrementally from 20 to 160 °C. The phase transition from α -S₈ to β -S₈ to liquid sulfur (Fig. 2a) produces extensive variation in the ULF region of the Raman spectrum, while it generates only subtle spectral changes in the internal frequencies. The loss of crystallinity and long-range order is apparent in the ULF spectra, as the definitive peaks of the α - and β -S₈ lattice vibrations meld into a smooth, wing feature near the Rayleigh line in the molten sulfur scans. The wing feature is a spectral signal representative of amorphous, disordered materials, referred to as the Boson peak³⁶, and it is a characteristic of liquid sulfur in the external modes. Supplementary Fig. 1 compares the spectrum of molten sulfur with the spectrum of the aluminum foil on which the sample was analyzed to verify sulfur as the source of the Boson peak signal. Additionally, the torsional vibration band at the crystal lattice boundary, which is a strong doublet peak in the α -S₈ spectra at 82 and 88 cm⁻¹, and a distinct peak feature respectively in the β -S₈ spectra, morphs into a soft shoulder on the Boson peak in the molten sulfur spectra (Fig. 2a), indicating the loss of torsional vibration.

Subtle spectral changes occur in the internal vibrations as sulfur transforms from its solid to liquid state. Overall, a decrease in signal-to-noise ratio between the spectrum of crystalline solid (Fig. 2b) and the molten sulfur spectrum (Fig. 2c) can be attributed to the phase change and subsequent differences in the Raman scattering cross-section of the sulfur. The FWHM of the molten sulfur peaks increases marginally over the course of heating. The multiple Raman components of the 220 cm⁻¹ S-S bending vibration disappear into a singular peak feature around 100 °C, the melting point of β -S₈ (Fig. 2a). This peak remains relatively more intense than the S-S stretching peak at 473 cm⁻¹ across all spectra in the *in-situ* experiment.

Several chemical physics, soft matter, and kinetics studies explore the unique liquid-liquid “ λ ” transition of sulfur at T _{λ} = 159 °C, where liquid sulfur polymerizes into long chain structures (S _{μ})^{37–41}. This transition is illustrated by changes in the 473 cm⁻¹ peak region at elevated temperatures, with the development of an expansive shoulder around 461 cm⁻¹ in the 140–160 °C molten spectra (Fig. 2a,c). This feature has been associated with the polymerization of cyclooctasulfur rings, a process whereby the scission of S-S bonds and subsequent concatenation yield long polymeric diradical S _{μ} chains^{37–40,42,43}. However, the higher frequency, 467 cm⁻¹ shoulder of the α -S₈ standard in Fig. 2b, is assigned to an S-S stretching vibration of the α -S₈ crystal in Raman polarization studies^{33,44}. An increase in the intensity of the 430 cm⁻¹ peak at high temperatures (>150 °C) (Fig. 2a,c) also signals the increasing polymer content and the progression of polymerization⁴⁵.

Some polarization studies investigate the extent of sulfur polymerization by comparing the depolarization ratio of Raman lines in the high frequency range to identify differences in chain length and molecular weight species³⁸. This study does not explore the depolarization ratio, but it notes the presence of symmetric stretch vibrations (461 cm⁻¹) of the polymeric S _{μ} component present in the >140 °C molten spectra in Fig. 2c.

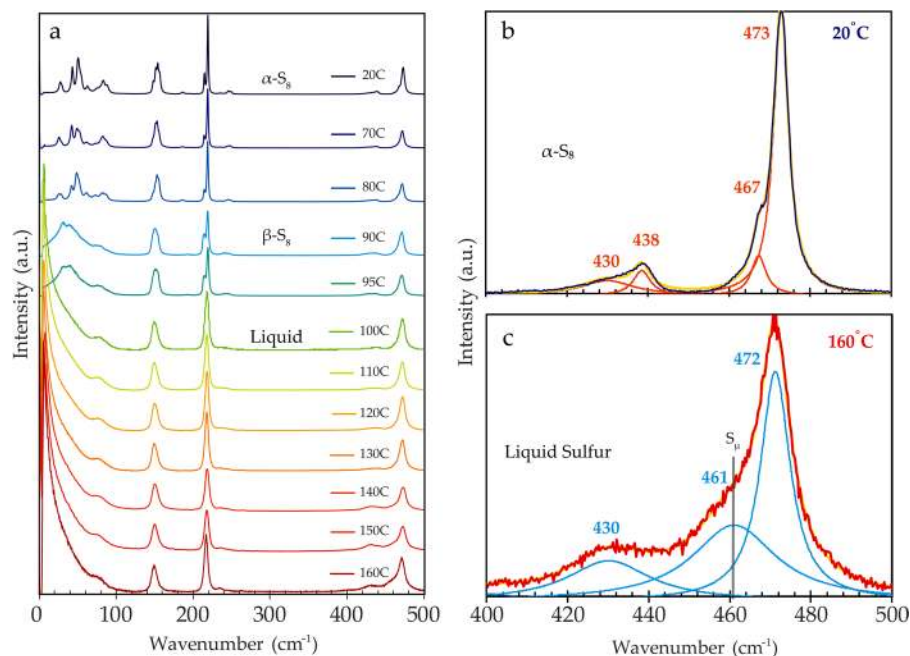


Figure 2. (a) Raman spectra from an *in-situ* experiment at increasing temperatures. The experiment begins with solid α -S₈, with an observable transition to β -S₈, and then a transition to liquid S₈. Spectra are normalized to the 150 cm⁻¹ peak; (b) A plot of the 20 °C α -S₈ spectrum and (c) the 160 °C liquid sulfur spectrum normalized to the characteristic 473 cm⁻¹ S₈ feature, showing Lorentzian functions (orange and blue respectively) and the cumulative fit in yellow. Plot (c) shows the presence of a shoulder feature (461 cm⁻¹), which denotes the accretion of disordered polymeric sulfur chains (S _{μ}) in the high temperature liquid sulfur.

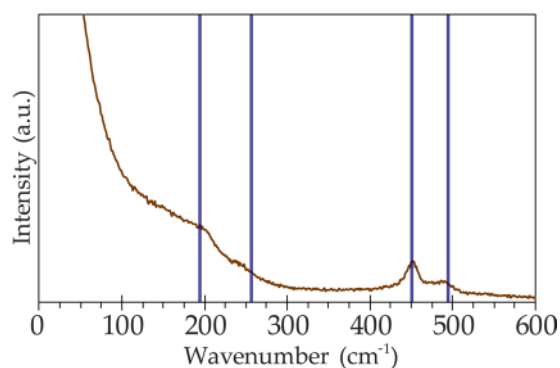


Figure 3. Spectrum of an aqueous polysulfide solution. Blue markers delineate positions of previously published polysulfide species bands^{46,47}.

Polysulfides standard. The Raman spectrum of a polysulfide solution is shown in Fig. 3. The plotted blue lines in Fig. 3 indicate published peak positions for Na₂S₄^{46,47} and similar values to the observed bands of S-S bond stretching in H₂S₄ polysulfanes¹⁵. The aqueous polysulfide spectrum presents generally weak and broad peaks, with the exception of the more pronounced 450 cm⁻¹ feature, which is characteristic for aqueous Na₂S₄⁴⁶. Our reference aqueous polysulfide scan lacks the diagnostic oxoanion shoulder at 337 cm⁻¹, which suggests a pool of mostly reduced S_n²⁻ species⁴⁷. The undefined spectral wing in the low frequency range is also consistent with a liquid solution.

S(0) storage globules in *Thiothrix* sp. *Living Thiothrix* cells. *In vivo* Raman spectroscopy of *Thiothrix* sp. was performed on samples examined immediately after field collection. Movement of the *Thiothrix* filaments under the microscope ensured the cells were alive throughout the *in vivo* Raman analysis. Complementary SEM analyses show that the sulfur globules, approximately 1 μ m in diameter, stack within the enclosed sheath structure of the *Thiothrix* cells (Fig. 4). These sulfur inclusions rest within invaginations of the cytoplasmic membrane, as described in previous studies⁴⁸.

In vivo Thiothrix Raman spectra present the characteristic S₈ structure previously described for the crystalline S₈ standards and molten sulfur in the internal modes (>100 cm⁻¹). The spectra present a definitive Rayleigh wing

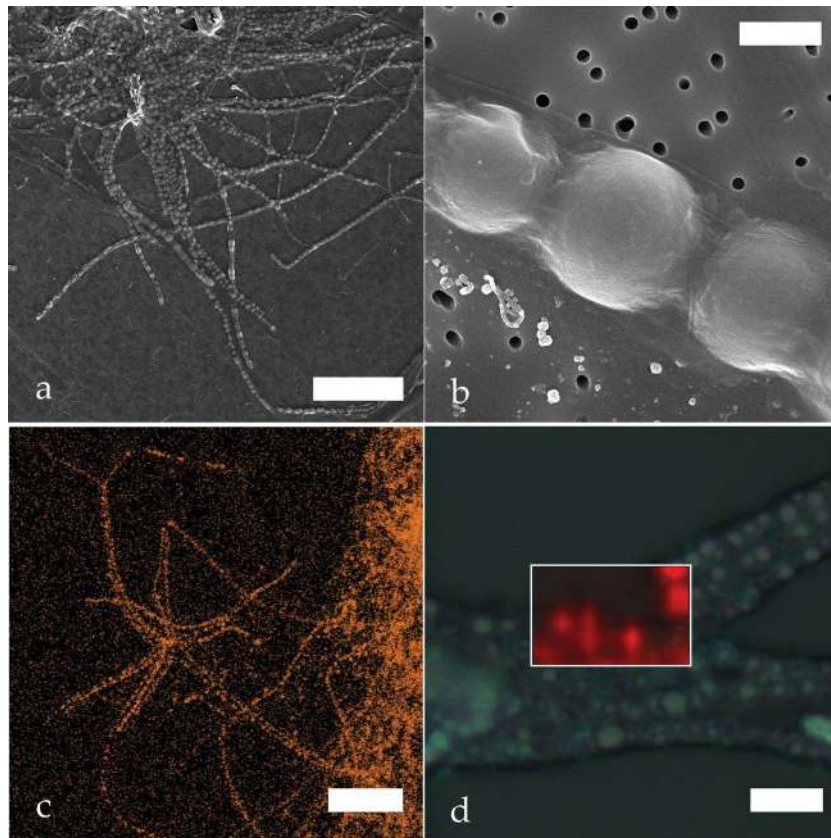


Figure 4. Intracellular sulfur globules in *Thiothrix* sp. (a) Scanning electron micrographs of *Thiothrix* filaments; (b) Close-up on *Thiothrix* intracellular sulfur globules; (c) SEM-EDS elemental map with *Thiothrix* sulfur globules highlighted in orange; (d) Raman map (white rectangle) of a bundle of *Thiothrix* filaments, where the areas mapped in red could be fitted with the biosulfur spectrum shown in Fig. 5. Scale bars: (a) 25 μm (b) 1 μm (c) 25 μm (d) 5 μm .

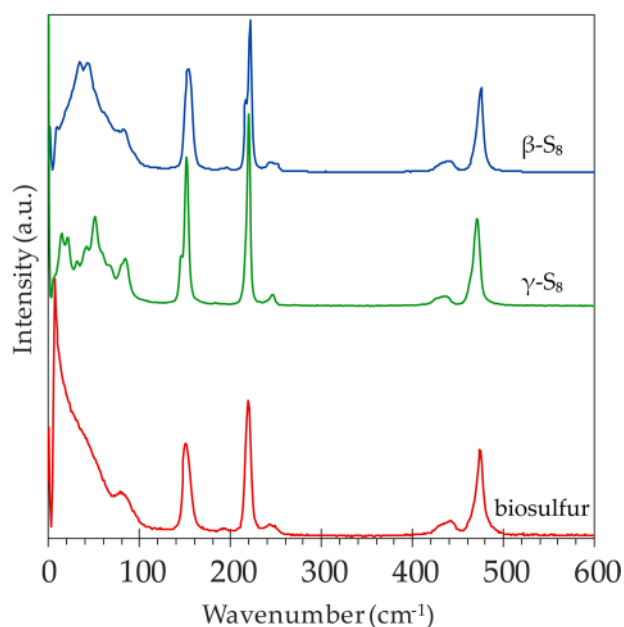


Figure 5. A plot of the sulfur allotrope spectra selected from mapped regions of several *Thiothrix* samples in varying conditions. The colors of the spectra correspond with the colors of the areas mapped in Figs 4, 6 and 7.

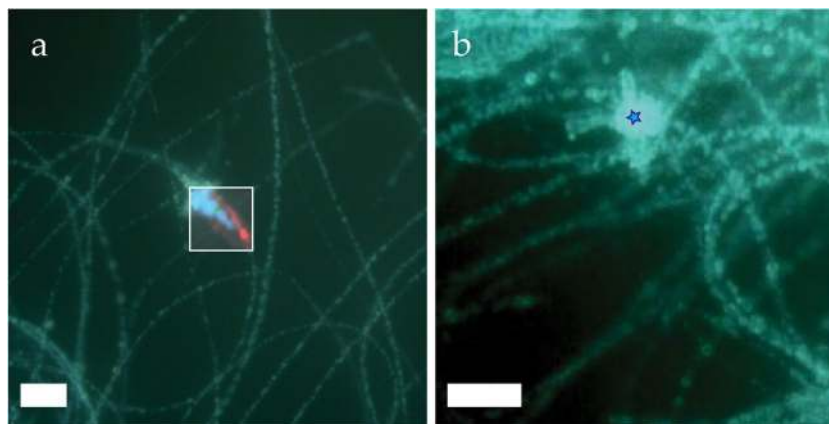


Figure 6. Aged *Thiiothrix* in stored at 5 °C (a) Raman map (white rectangle) overlay on optical microscope image, showing extracellularly precipitated β -S₈ (blue) adjacent to *Thiiothrix* filaments containing globules of amorphous biosulfur (red); (b) Cluster of extracellular beta sulfur marked with blue star. Scale bars: (a) 10 μ m (b) 5 μ m.

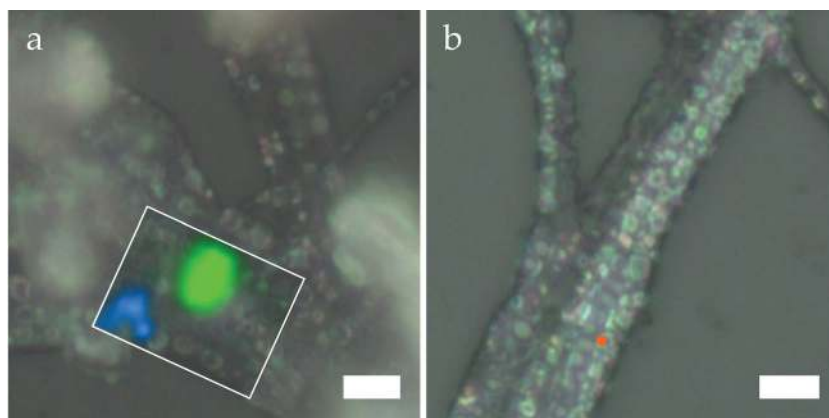


Figure 7. (a) Raman map (white rectangle) overlay on optical microscope image of externally precipitated β -S₈ (blue) and γ -S₈ (green) resting atop dried *Thiiothrix* filaments (b) Two-week dehydrated *Thiiothrix*, lodged with bright spherules mostly void of S(0). An intact globule (orange marker) with residual biosulfur could be identified. Scale bars: 4 μ m.

feature throughout the ULF region indicating amorphous or glassy disordered sulfur molecules (Fig. 5); the torsional peak around 80 cm^{-1} is positioned approximately five wavenumbers lower than the crystalline S₈ allotropes (76 cm^{-1} instead of 82 cm^{-1}) and it is relatively less pronounced in comparison to our reference S₈ scans. The internal peaks have similar FWHM to the reference solid allotropic standards (Fig. 5). However, the individual S₈ Raman components present in the main characteristic bending-stretching modes of solid allotropes, seen as shoulders on the 150 cm^{-1} and 220 cm^{-1} peaks, are absent in the biosulfur globules and could indicate differences in the intra-molecular bonds of the biosulfur structure.

Aged *Thiiothrix*. To identify potential changes in the structure or speciation of the intracellular *Thiiothrix* biosulfur with time after field collection, we measured samples stored in Falcon tubes in a refrigerator (at 5 °C), in original well water, for 17 days. Analyses of aged *Thiiothrix* samples present spectral shifts compared with live samples, probably due to chemical changes occurring during storage of the samples. *Thiiothrix* cells aged in native sulfidic well water at 5 °C likely experience oxidizing conditions through diffusion of oxygen from the headspace and the imperfect seal of the Falcon tube in which they are stored. For storage periods of hours to days, the sulfur inclusions are still visible within *Thiiothrix* cells, and the spectral S₈ bands are nearly indistinguishable from the spectra of fresh, living *Thiiothrix*. However, after one or several months of prolonged storage, intracellular sulfur globules disappear, leaving behind empty sheaths. This could be due to the oxidation of S(0) by *Thiiothrix* following the depletion of sulfide from the storage medium^{49,50}.

Extracellular β -S₈ crystals formed in a *Thiiothrix* sample aged for 17 days in the original well water (Fig. 6). These sulfur precipitates, ranging in diameter from one to five micrometers, often develop proximal to the filaments still containing intracellular amorphous S(0) globules. The observations of S(0) oxidation and precipitation of extracellular S(0) with different structures argue for the necessity to limit time between sample collection and

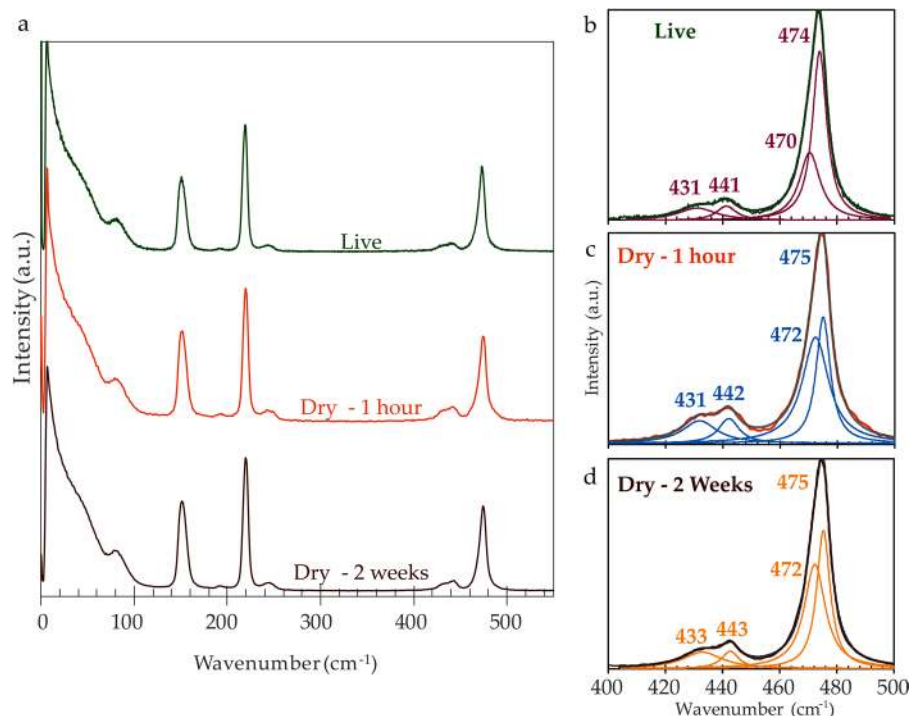


Figure 8. (a) The spectra of intact *Thiothrix* sulfur globules measured through a range of hydration states: live (top), dried on slide for one-hour (middle), and dried on slide for two weeks (bottom); (b–d) Peak fits in the 400–500 cm^{-1} range. Experimental data corresponds to color of the spectrum plotted in 8a; each fit profile is shown in gray; and Lorentzian peaks are shown in purple, blue, and orange. Spectra in (a) and (b–d) were normalized to the characteristic S_8 473 cm^{-1} peak.

Raman analyses when characterizing sulfur in environmental microbial samples, as already discussed in another study¹⁹.

Dried *Thiothrix*. Raman analyses on freshly collected *Thiothrix* cells dried for two to three hours on a microscope slide indicate no detectable change in sulfur composition. The S(0) inclusions remain amorphous, stabilized within the cell membrane.

Two-week dried *Thiothrix* samples, stored on microscope slides housed in foil-wrapped petri dishes, show extensive alteration in chemical and structural composition. Notably, Raman scattering on the bright intracellular features (Fig. 7b) produces featureless spectra from 0–1200 cm^{-1} with a strong fluorescence background signal, which could indicate some organic remainder (data not shown). These spherical inclusions are possibly empty vesicles of former sulfur storage globules⁵¹. Some filaments in the same samples however featured intact amorphous S(0) globules, albeit sparsely distributed, despite the weeks-long dehydration (Fig. 7b). This residual amorphous biosulfur in the dried samples co-occurs with $>2 \mu\text{m}$ crystals of extracellular metastable S_8 allotropes proximal to the cells. Figure 7a shows a dense cluster of γ - S_8 adjacent to smaller β - S_8 crystals resting atop S(0)-depleted *Thiothrix* filaments.

Raman analysis on the living, aged, and dried *Thiothrix* samples shows the absence of sulfur in any other oxidation state besides S(0). Sulfur oxoanion species, including sulfite, thiosulfate, and sulfate, have Raman band assignments in the 500 to 1100 cm^{-1} range, which were not observed in our scans performed into higher Raman frequency ranges. In particular, the symmetric S=O stretch vibration of sulfate at 960–1000 cm^{-1} ⁵² is undetectable in dozens of scans performed in this study, even in aged or dried samples.

A more detailed spectral exploration of intact sulfur globules remaining in the dried *Thiothrix* samples uncovers slight shifts in shoulder features within the low frequency modes and changes in the higher wavenumber internal vibrations (Fig. 8). The softened 80 cm^{-1} torsional peak of the living *Thiothrix* in Fig. 8a becomes slightly more prominent in the dried samples' spectra. The relative intensities of the main S_8 peaks are mostly conserved during drying, except for a slight increase in the intensity of the asymmetric S-S bending peak (150 cm^{-1}) in the dried samples.

Figure 8b–d present peak fits in the 400–500 cm^{-1} wavenumber range of the spectra shown in Fig. 8a, and illustrate some changes in the S_8 stretching modes of living versus dried (1 hour and 2 weeks) *Thiothrix* sulfur globules. The positions of the Lorentzian functions used to fit the spectra of the dried *Thiothrix* at 1 hour and 2 weeks indicate vibrations of the S-S stretching peak, with components at 472 and 475 cm^{-1} . These band positions are commonly attributed to the stretching of the S_8 ring species⁵³. These components are shifted into lower frequencies in the *in vivo* *Thiothrix* spectrum (470 and 474 cm^{-1}) and this could indicate an increase in bond length. The spectral feature at 470 cm^{-1} shown in the *in vivo* sample spectrum can be attributed to either short

sulfur chains, designated as S_{ub}^{39} , or to the S_8 ring species⁴⁰. The *in vivo* and dried *Thiothrix* spectra lack the diagnostic shoulder feature of long, polymeric sulfur chains (S_{ub}) at 460 cm^{-1} , prominent in the molten sulfur shown in Fig. 2b.

Peak fits of the *in vivo* and dried *Thiothrix* samples show two features at approximately 430 and 440 cm^{-1} in Fig. 8b–d. The positions of these stretching vibrations are consistent between the *in vivo* and 1 hour dried *Thiothrix*, but the peak positions shift to a slightly higher range (1 to 2 cm^{-1}) in the 2 weeks dried sample. This trend could coincide with the drying process.

Discussion

The present study highlights the advantages of using low-frequency Raman in the detection and characterization of elemental sulfur in microbial and environmental samples. This work supplements the limited literature on sulfur Raman external modes^{33–35,44}, and presents the first low-frequency Raman spectrum of γ - S_8 published so far. Several (geo)microbiological investigations have used Raman spectromicroscopy to confirm the presence of S(0) in environmental samples or laboratory cultures^{8,9,19,21,26,27}. However, these previous studies did not include low-frequency measurements, limiting analyses of the phonon vibrations of the crystal lattice and hence structural determinations of the sulfur. Indeed, while only minor differences can be observed in the high frequency ($>100\text{ cm}^{-1}$) Raman signature of different S_8 allotropes, Raman measurements below 100 cm^{-1} enable clear determination of S_8 crystalline structures, as well as non-crystalline forms of sulfur (Figs 1 and 2). While structural characterization of crystalline S(0) can also be achieved by bulk XRD analysis, this method is unsuitable for the detection of amorphous or liquid S(0), nano-crystalline S_8 allotropes, and allotropes present in small quantities. For instance, our XRD analysis of aged *Thiothrix* samples failed to detect crystalline S_8 phases, while our ULF Raman measurements revealed the presence of micron-sized β - S_8 crystallites (Fig. 6). Furthermore, Raman spectromicroscopy, as a micron-scale method, enables spatial localization (mapping) of the analyzed particles and determination of their morphology, and can potentially be correlated with sub-micron scale imaging techniques such as SEM for high resolution analyses. Spatial correlation of particle morphology and S(0) crystal structure can also be achieved by electron diffraction methods, but only ULF Raman spectromicroscopy enables such investigations to be performed on wet or live samples under room temperature and atmospheric pressure conditions. The importance of *in vivo* and wet analyses when characterizing the structure of microbial or environmental S(0) was demonstrated here by the precipitation of extracellular monoclinic β - S_8 and γ - S_8 crystals in aged and dried *Thiothrix* samples (Figs 6 and 7).

Application of ULF Raman spectromicroscopy to S(0) detection and characterization in natural samples paves the way for future investigations of metastable monoclinic S_8 allotropes in the environment. β - and γ - S_8 have so far only been described in a few low-temperature natural settings^{2,54,55}. It was recently demonstrated that β - and γ - S_8 can be formed by sulfide oxidation in the presence of organics, through a process called S(0) organomineralization⁵⁶. The stabilizing effect of organics on monoclinic S_8 allotropes could explain the precipitation of β - and γ - S_8 in the proximity of the microbial cells in our aged and dried *Thiothrix* samples (Figs 6 and 7). In the future, a more widespread application of low-frequency Raman analyses on natural samples might reveal that this process is more frequent than previously thought.

Further ULF Raman measurements can also be used to explore complex interactions or conditions promoting the precipitation of metastable S_8 allotropes. Hydration sequences appear to impact the formation of γ - S_8 in the environment. Localities experiencing cyclical hydration-dehydration phases host conditions favorable to γ - S_8 crystallization: in evaporitic systems where gelled EPS matrix embeds crystals in cyanobacteria biofilms perched atop the water table⁵⁵, in supraglacial perennial subsurface springs where EPS deposits encase sulfur rhombs⁵⁴, in sewage sludge experiments where micro-crystals encrust dehydrated sewage granules⁵⁷, in our experimental dried *Thiothrix* sample where γ - S_8 particles form adjacent to cellular sheaths, and in other dried microbial microscope samples that yield prismatic, spindled sulfur crystals¹⁹. Low frequency investigations could help determine potential links between drying processes and the precipitation of metastable gamma sulfur in environmental samples.

Low-frequency Raman spectromicroscopy can also be employed to solve uncertainties in the ongoing debate on the composition of microbial biosulfur. This question has been the object of several studies using synchrotron-based X-ray absorption spectroscopy (XANES) on a variety of sulfur bacteria, without resulting in a consensus. It has, for instance, been proposed that microbial sulfur was composed: of sulfur chains capped by unidentified organic residues²², simple solid S_8 ¹⁸, globules of varied speciation – cyclooctasulfur, sulfur chains, and polythionates – contingent on the sulfur-oxidizing metabolism²³, and a core of S_8 crowns enclosed in a hydrophilic envelope¹⁷. Analytical difficulties of XANES techniques leading to spectral distortions and artifacts further complicate this debate¹⁷. Our *in vivo* Raman analyses of intracellular S(0) globules in *Thiothrix* sp. showed diagnostic features of the internal vibrational modes of solid S_8 , in addition to a Boson peak in the external modes, indicative of disordered material. This suggests that globular biosulfur lacks long range order, countering previous Raman findings of microcrystalline biosulfur deposits^{19,58}. The low frequency spectra of the *Thiothrix* biosulfur resemble the spectral features of liquid sulfur measured in our *in-situ* high temperature experiment (Fig. 2). However, the contribution of long polymeric sulfur, visible by the pronounced shoulder at 461 cm^{-1} in liquid sulfur – is absent from the biosulfur spectra. The 470 cm^{-1} peak observed in live *Thiothrix* cells could signal shorter polymeric chain species^{38,39}, but this spectral component has also been assigned to S_8 ring species in other work⁴⁰. Considering the difficulty in resolving or assigning the 470 cm^{-1} peak, our analyses suggest that intracellular S(0) globules in *Thiothrix* sp. are mostly composed of disordered S_8 rings, with the possible presence of short polymeric diradical species.

Another Raman study identified S_8 as the main S(0) form in sulfur globules produced in *Beggiatoa* filaments, but only in subpopulations located at the sulfide-oxygen interface of gradient tubes or in early growth stage cultures²¹. *Beggiatoa* mats forming in a deeper sulfide-rich, anoxic zone, and freshwater gradient cultures presented

S_8 Raman active peaks with an additional feature at 272 cm^{-1} , and a range of S-S stretching vibrations ($\sim 470\text{ cm}^{-1}$) respectively, suggesting a composite of S_8 rings and linear polysulfides (S_n^{2-}) within the globules²¹. The characteristic polysulfide or polysulfane peaks were not detected in our Raman spectra of live *Thiothrix*, and these differences in polysulfide content could clue different S(0) metabolic pathways in *Beggiatoa* versus *Thiothrix*. Unlike *Thiothrix*, *Beggiatoa* can reduce stored S(0) into sulfide under short-term anoxia^{59,60}, which could explain the presence and detection of polysulfide intermediates in the *Beggiatoa* measurements. That said, the transience and reactivity of polysulfide anions make them challenging to analyze, and inter-ion interactions in sulfide solutions can perturb the vibrational frequencies⁴⁷ – arguing for a multidimensional approach in measuring S_n^{2-} species.

XRD studies have suggested the presence of “liquid-like” sulfur in extracted microbial sulfur globules²⁰, though these results face skepticism over the stability of liquid sulfur at room temperature and pressure¹⁸. Recent XANES data suggest that biosulfur particles have a density of $2.0\text{ g}\cdot\text{cm}^{-3}$, overturning earlier assumptions of low, $1.2\text{ g}\cdot\text{cm}^{-3}$ density globules and assigning a density value akin to crystalline α - S_8 ¹⁷. The more recent assessment of biosulfur density – though contentious – supports our results suggesting globules dominated by closely stowed S_8 crowns. Overall, our results are in agreement with other studies and Raman analysis indicative of intracellular microbial sulfur in the form of compact S_8 rings^{26,50,61}. Raman external mode vibration measurements in this work augment the current understanding of intracellular S(0) globules by verifying the disordered, amorphous inter-molecular structure of encapsulated S_8 rings.

This proof-of-concept study demonstrates the capacity of low frequency Raman spectromicroscopy to reveal micro-scale S(0) speciation and structure in environmental and biological samples. Our work opens up the opportunity for an examination of the conditions and mechanism of precipitation and stabilization of metastable sulfur allotropes in the environment, and for future studies on the composition, formation and use of S(0) inclusions in different sulfur bacteria.

Materials and Methods

Sample collection and preparation. *Microbial mats.* Samples of microbial mats dominated by *Thiothrix* sp. – a sulfur-oxidizing, Gammaproteobacterial genus – were collected from an abandoned oil and gas well in Centre County, Pennsylvania. Fluorescence *in situ* hybridization (FISH) and cell morphology were used to identify the primary mat populations. FISH was carried out as described in Macalady *et al.* (2006)⁶² with commercially synthesized oligonucleotides (Sigma-Genosys). Slides were counterstained with 4',6'-diamidino-2-phenylindole (DAPI). Microbial mat samples were dominated by rosettes of thin filaments, 1–1.5 μm in diameter, with intracellular S(0) inclusions that hybridized with the EUBMIX and GAM42a probes. FISH images are shown in Supplementary Fig. 2.

Several sampling excursions were performed from winter (December 2017) through spring (late May 2018). An estimated 60 mL of the microbial mat proximal to the sulfidic outflow was pipetted into sterile Falcon tubes during each site visit. Samples collected and analyzed on the same day were centrifuged at low speeds ($<5000\text{ rpm}$) and rinsed three times with ultrapure 18.2 M Ω water. Samples assessed in subsequent days were stored at 5°C with oxygenated headspace in original well water. These samples were rinsed with ultrapure 18.2 M Ω water on the day of analysis. All samples were manipulated in a biosafety cabinet under laminar flow conditions to minimize contamination.

Preparation of living cells for *in vivo* Raman spectroscopy analyses mirrored the procedure used by Berg *et al.* (2014)²¹: a 20 μL droplet of rinsed sample was positioned on a glass microscope slide and secured by a taped coverslip to preserve the bacterial micro-habitat during data collection. “Dried” samples used un-sealed coverslips and were dry prior to analysis. We tested the repeatability of the Raman measurements on several collections of *Thiothrix* samples and on all sulfur standards.

Raman spectroscopy. Raman spectroscopy of all samples and standards was conducted using a Horiba LabRam HR Evolution Raman Vis-NIR spectrometer coupled with a HeNe 633 nm laser source, an ideal excitation energy for applications aiming to minimize sample damage and requiring increased wavelength stability. The LabRam system utilizes an Olympus BXM-ILHS optical microscope and IDS uEye 3 MPix video camera/software. BraggGrate notch filters allowed the collection of low frequency ($5\text{--}10\text{ cm}^{-1}$) signal. Most spectra in the study were collected using a groove density of 600 g/mm; 1800 g/mm grating was employed on certain samples and on all standards to better resolve both ULF vibrations and internal mode spectral features. The confocal pinhole aperture was set at 50 μm , providing a spectral resolution of $\sim 0.8\text{ cm}^{-1}$. The acquisition range of the spectrometer was centered at 456 cm^{-1} to incorporate anti-Stokes and Stokes measurements (-200 cm^{-1} to 1000 cm^{-1}) and verify features in the low frequency range. Reference spectra plotted in both regions can be found in Supplementary Fig. 3. The spectrometer was calibrated with a silicon standard prior to each Raman session.

The point measurements and Raman maps were carried out using either an apochromatic 50x or a 100x objective, while the reference standard spectra were collected using a 50x long working distance (LWD) objective. This analysis achieved a spatial resolution of approximately 1 μm . Point spectra were collected with two accumulations and 10 to 15-second exposure time; Raman mapping was conducted with one, 10-second accumulation, scanned in defined 0.75–0.80 μm x-y steps on an automated stage. Laser power was approximately 1–1.3 mW at the sample for both single point scans and mapping to avoid sample damage. Power levels documented in different sulfur-Raman studies provided a framework for the basic parameters in this research^{19,21}. Additionally, to avoid potential photothermal effects and to address potential overlap with the absorption edge of long elemental sulfur chains, several power percentages were tested to optimize laser threshold power.

The LabSpec6 platform was used to process Raman spectra. Baseline correction was performed on raw spectra using polynomial fitting in LabSpec6 to remove fluorescence and background signal. Raman mapping incorporated the Classical Least Squares (CLS) fitting function in LabSpec6 to highlight the distribution of sulfur allotropes within a mapped region of interest. A representative spectrum was selected as the reference per each

mapped sulfur component. Overlays of Raman maps on optical images provided visual correlation between the microscopic features and the molecular spectra. Optical microscope images were exported from LabSpec6 and processed using the ImageJ package.

The OriginPro software package enabled the nonlinear curve fitting of certain spectral features. Spectra were fit to Lorentzian peaks using a method based on the Levenberg-Marquardt iterative algorithm in order to detect overlapping features in the high frequency regions⁶³.

Reference sulfur compounds. *S₈ Allotropes.* All crystalline sulfur solids were recorded with Raman point measurements. The same samples were concurrently analyzed with X-ray powder diffraction to confirm their crystal structure (Supplementary Fig. 4).

Sulfur powder (Alfa Aesar 99% pure sulfur) provided the standard for orthorhombic, α -S₈ sulfur. Orthorhombic sulfur is the thermodynamically stable allotrope at room temperature and pressure (Supplementary Fig. S5). For Raman spectroscopy, α -S₈ powder was measured on a glass slide.

β -S₈ is a metastable S₈ allotrope that forms at temperatures above 96 °C (see Supplementary Fig. S5). β -S₈ can be obtained from the quenched melt of molten α -S₈³⁹ and this monoclinic sulfur commonly forms elongated, pointed crystals. Approximately 10 grams of pure sulfur powder (Alfa Aesar 99%) were emplaced in a test tube and melted with a torch. The liquified sulfur was quenched in ice water and formed a dark brown, tacky solid. Approximately one cm² of this quenched sulfur was analyzed with Raman on a glass slide and by XRD within 30 minutes of solidifying to avoid re-crystallization of α -S₈.

γ -S₈ is a rare monoclinic allotrope of S₈ that forms yellowish-white elongated, spiny crystals⁶⁴. Akin to β -S₈, gamma sulfur is stable at high temperatures and pressures (Supplementary Fig. S5). The slow oxygenation of a sulfide solution (500 μ M Na₂S, Spectrum Chemical) containing yeast extract (5 g.L⁻¹, Fisher Scientific) for three weeks yielded stable particulate γ -S₈ in a process described as S(0) organomineralization⁵⁶. This precipitate was collected by centrifugation and rinsed with 18.2 M Ω water. A 20 μ L volume of this concentrated slurry was pipetted onto a glass slide and allowed to dry before Raman investigation. XRD analyses confirmed the presence of γ -S₈ as the only form of crystalline S(0) in two separate samples prepared through the same organomineralization protocol.

Molten sulfur. *In-situ* molten sulfur experiments were conducted on the same Horiba LabRam HR Evolution setup paired with a Linkam temperature-controlled stage (HFS600E) and the LNP95 cooling pump. Nitrogen gas was used to purge the stage system headspace. A 50x LWD objective was used with the Linkam stage. The stage was calibrated using a thermocouple to align actual temperature measurements with the recorded output of the Linkam control system. Synchronous temperature adjustments were recorded, corresponding with each incremental measurement of the heating experiment.

Sulfur powder (Alfa Aesar 99% sulfur) was loaded on a section of aluminum foil, enclosed, purged with N₂, and then heated on the silver block stage body. The experiment was initiated at 70 °C, followed by 10 °C increases, to reach a final temperature of 160 °C. Additionally, the pivotal temperature point of allotropic conversion from α -S₈ to β -S₈ was measured at 95 °C⁶⁵. Sample drying could not be mitigated with this stage configuration. Sample loss and the evaporation of molten sulfur above 170 °C prevented higher temperature measurements.

Polysulfides. A polysulfide reference solution was formulated following the recipe used in Berg *et al.*, (2014), where a solution of 5.8 g of elemental sulfur per 100 mL 18.2 M Ω water was mixed with 5.06 g Na₂S · 9H₂O at a final pH of 9.5 in an anaerobic chamber and passed through a 0.2 μ m filtration system²¹. Press-to-seal silicone isolators (Electron Microscopy Services) were used for Raman measurements. DuoScan averaging mode captured aqueous polysulfides spectra acquisitions over a 30 μ m² swath of sample, at approximately 1.3 mW, using a 200 μ m confocal hole, 50x LWD objective, 633 nm laser, and 600 gr/mm.

Complementary techniques. Scanning electron microscopy supplemented Raman data with high resolution imaging of the microbial samples. Secondary electron images were generated at an accelerating voltage of 7 kV, with a working distance (WD) of approximately 3 mm, using an FEI Nova NanoSEM630 instrument. Energy dispersive spectroscopy (EDS) analyses were performed at 15 kV and a 5 mm WD using an Oxford Instruments UltimMax detector and the data were analyzed using the Oxford AZtec platform. EDS maps and spectra confirmed microbial sulfur globules content and distribution.

For SEM/EDS analysis, samples were rinsed with 18.2 M Ω water, and deposited onto 0.22 μ m Whatman Nucleopore filters, using a syringe filter holder dispensing system (25 mm, Pall Laboratory). Filters were placed on double-sided carbon tape atop aluminum stubs. All samples were sputter coated with iridium (12 nm) to prevent charging.

X-ray Diffraction data was collected on rinsed samples deposited on a single crystal miscut Si holder. Samples were analyzed using a PANalytical Empyrean diffractometer paired with a PIXcel^{3D} detector, and CuK α ($\lambda = 1.5406$ Å) incident X-ray radiation. Scans were conducted over a 2θ range of 5–70°, and analyses used a step size of 0.025°, a time of 96.4 seconds per step, and a current density of 40 mA. X-ray diffraction data were analyzed with Jade software. Recorded spectra were compared with reference spectra from the International Centre for Diffraction database.

Data Availability

The datasets generated and/or analyzed during the current study are available from the corresponding author on reasonable request.

References

- Macur, R. E. *et al.* Microbial community structure and sulfur biogeochemistry in mildly-acidic sulfidic geothermal springs in Yellowstone National Park. *Geobiology* **11**, 86–99 (2013).
- Lau, G. E. *et al.* Low-temperature formation and stabilization of rare allotropes of cyclooctasulfur (β -S₈ and γ -S₈) in the presence of organic carbon at a sulfur-rich glacial site in the Canadian High Arctic. *Geochim. Cosmochim. Acta* **200**, 218–231 (2017).
- Taylor, C. D., Wirsén, C. O. & Gaillardet, F. Rapid Microbial Production of Filamentous Sulfur Mats at Hydrothermal Vents Rapid Microbial Production of Filamentous Sulfur Mats at Hydrothermal Vents. *Appl. Environ. Microbiol.* **65**, 2253–2255 (1999).
- Taylor, C. D. & Wirsén, C. O. Microbiology and ecology of filamentous sulfur formation. *Science* (80-). **277**, 1483–1485 (1997).
- Kamyshny, A. & Ferdelman, T. G. Dynamics of zero-valent sulfur species including polysulfides at seep sites on intertidal sand flats (Wadden Sea, North Sea). *Mar. Chem.* **121**, 17–26 (2010).
- Zopf, J., Ferdelman, T. G. & Fossing, H. Distribution and fate of sulfur intermediates - sulfite, tetrathionate, thiosulfate, and elemental sulfur in marine sediments. *Geol. Soc. Am. Spec. Pap.* **379**, 97–116 (2004).
- Mather, T. A., Pyle, D. M. & Oppenheimer, C. Tropospheric volcanic aerosol. *Geophys. Monogr. Ser.* **139**, 189–212 (2003).
- Himmel, D., Maurin, L. C., Gros, O. & Mansot, J.-L. Raman microspectrometry sulfur detection and characterization in the marine ectosymbiotic nematode *Eubostriechus dianae* (Desmodoridae, Stilbonematidae). *Biol. Cell* **101**, 43–54 (2009).
- Eichinger, I., Schmitz-Esser, S., Schmid, M., Fisher, C. R. & Bright, M. Symbiont-driven sulfur crystal formation in a thiotrophic symbiosis from deep-sea hydrocarbon seeps. *Environ. Microbiol. Rep.* **6**, 364–372 (2014).
- Howarth, R. *et al.* Phylogenetic relationships of filamentous sulfur bacteria (Thiothrix spp. and Eikelboom type 021 N bacteria) isolated from wastewater- treatment plants and description of *Thiothrix eikelboomii* sp. nov., *Thiothrix unzii* sp. nov., *Thiothrix fructosivorans* s. *Int. J. Syst. Bacteriol.* **49**, 1817–1827 (1999).
- Kleinjan, W. E., Keizer, A. & Janssen, A. J. H. *Biologically Produced Sulfur*. 167–188, <https://doi.org/10.1007/b12114> (2003).
- Luther, G. W. *et al.* Thermodynamics and kinetics of sulfide oxidation by oxygen: A look at inorganically controlled reactions and biologically mediated processes in the environment. *Front. Microbiol.* **2**, 1–9 (2011).
- Dahl, C. & Friedrich, C. G. eds *Microbial Sulfur Metabolism*. (Springer Berlin Heidelberg), <https://doi.org/10.1007/978-3-540-72682-1> (2008).
- Meyer, B. *et al.* Elemental Sulfur. *Chem. Rev.* **76**, 367–388 (1976).
- Trofimov, B. A., Sinegovskaya, L. M. & Gusarova, N. K. Vibrations of the S-S bond in elemental sulfur and organic polysulfides: A structural guide. *J. Sulfur Chem.* **30**, 518–554 (2009).
- Stuedel, R. Homocyclic sulfur molecules. *Inorg. Ring Syst.* **3594**, 149–176 (1982).
- George, G. N., Gnida, M., Bazylinski, D. A., Prince, R. C. & Pickering, I. J. X-ray absorption spectroscopy as a probe of microbial sulfur biochemistry: The nature of bacterial sulfur globules revisited. *J. Bacteriol.* **190**, 6376–6383 (2008).
- Pickering, I. J. *et al.* Analysis of sulfur biochemistry of sulfur bacteria using X-ray absorption spectroscopy. *Biochemistry* **40**, 8138–8145 (2001).
- Pasteris, J. D., Freeman, J. J., Goffredi, S. K. & Buck, K. R. Raman spectroscopic and laser scanning confocal microscopic analysis of sulfur in living sulfur-precipitating marine bacteria. *Chem. Educ.* **180**, 3–18 (2001).
- Hageage, G. J., Eanes, E. D. & Gherna, R. L. X-ray diffraction studies of the sulfur globules accumulated by *Chromatium* species. *J. Bacteriol.* **101**, 464–469 (1970).
- Berg, J. S., Schwedt, A., Kreutzmann, A. C., Kuypers, M. M. M. & Milucka, J. Polysulfides as intermediates in the oxidation of sulfide to sulfate by *Beggiatoa* spp. *Appl. Environ. Microbiol.* **80**, 629–636 (2014).
- Prange, A. *et al.* *In situ* analysis of sulfur in the sulfur globules of phototrophic sulfur bacteria by X-ray absorption near edge spectroscopy. *Biochim. Biophys. Acta - Gen. Subj.* **1428**, 446–454 (1999).
- Prange, A. *et al.* Quantitative speciation of sulfur in bacterial sulfur globules: X-ray absorption spectroscopy reveals at least three different species of sulfur. *Microbiology* **148**, 267–276 (2002).
- Lee, Y. J. *et al.* *In situ* analysis of sulfur species in sulfur globules produced from thiosulfate by *Thermoanaerobacter sulfurifigens* and *Thermoanaerobacterium thermosulfurigenes*. *J. Bacteriol.* **189**, 7525–7529 (2007).
- Oren, A., Mana, L. & Jehlička, J. Probing single cells of purple sulfur bacteria with Raman spectroscopy: Carotenoids and elemental sulfur. *FEMS Microbiol. Lett.* **362**, 1–6 (2015).
- Eder, S. H. K., Gigger, A. M., Hanzlik, M. & Winklhofer, M. Sub-micrometer-scale mapping of magnetite crystals and sulfur globules in magnetotactic bacteria using confocal Raman micro-spectrometry. *PLoS One* **9**, 1–12 (2014).
- Maurin, L. C., Himmel, D., Mansot, J. L. & Gros, O. Raman microspectrometry as a powerful tool for a quick screening of thiotrophy: An application on mangrove swamp meiofauna of Guadeloupe (F.W.I.). *Mar. Environ. Res.* **69**, 382–389 (2010).
- Smith, G. P. S., Huff, G. S. & Gordon, K. C. Investigating Crystallinity Using Low Frequency Raman Spectroscopy: Applications in Pharmaceutical Analysis. *Spectroscopy* 1–8 (2016).
- Parrott, E. P. J. & Zeitler, J. A. Terahertz time-domain and low-frequency Raman spectroscopy of organic materials. *Appl. Spectrosc.* **69**, 1–25 (2015).
- Garcia, A. A. & Druschel, G. K. Elemental sulfur coarsening kinetics. *Geochem. Trans.* **15**, 1–11 (2014).
- Barletta, R. E., Gros, B. N. & Herring, M. P. Analysis of marine biogenic sulfur compounds using Raman spectroscopy: Dimethyl sulfide and methane sulfonic acid. *J. Raman Spectrosc.* **40**, 972–981 (2009).
- Harvey, P. D. & Butler, I. S. Raman spectra of orthorhombic sulfur at 40 K. *J. Raman Spectrosc.* **17**, 329–334 (1986).
- Becucci, M., Bini, R., Castellucci, E., Eckert, B. & Jodl, H. J. Mode Assignment of Sulfur α -S₈ by Polarized Raman and FTIR Studies at Low Temperatures. *J. Phys. Chem. B* **101**, 2132–2137 (1997).
- Gautier, G. & Debeau, M. Spectres de diffusion Raman du soufre β monoclinique. *Spectrochim. Acta Part A Mol. Spectrosc.* **32**, 1007–1010 (1976).
- Gorelik, V. S. & Tochilin, S. D. Raman Scattering in an Opal – Sulfur Composite **43**, 846–847 (2007).
- Kalampounias, A. G., Andrikopoulos, K. S. & Yannopoulos, S. N. Probing the sulfur polymerization transition *in situ* Raman spectroscopy. *J. Chem. Phys.* **118**, 8460–8467 (2003).
- Kalampounias, A. G., Andrikopoulos, K. S. & Yannopoulos, S. N. Rounding of the sulfur living polymerization transition under spatial confinement. *J. Chem. Phys.* **119**, 7543–7553 (2003).
- Andrikopoulos, K. S., Kalampounias, A. G., Falagara, O. & Yannopoulos, S. N. The glassy and supercooled state of elemental sulfur: Vibrational modes, structure metastability, and polymer content. *J. Chem. Phys.* **139** (2013).
- Zhang, L., Ren, Y., Liu, X., Han, F. & Evans-Lutterodt, K. Chain Breakage in the Supercooled Liquid – Liquid Transition and Re-entry of the λ -transition in Sulfur. *Sci. Rep.* 1–10, <https://doi.org/10.1038/s41598-018-22775-y> (2018).
- Andrikopoulos, K. S., Kalampounias, A. G. & Yannopoulos, S. N. Confinement effects on liquid-liquid transitions: Pore size dependence of sulfur's living polymerization. *Soft Matter* **7**, 3404–3411 (2011).
- Stuedel, R. Liquid Sulfur Liquid Sulfur. 81–116, <https://doi.org/10.1007/b12111> (2003).
- Stillinger, F. H. & Weber, T. A. Chemical Reactions in Liquids: Molecular Dynamics Simulation for Sulphur. *J. Chem. Phys.* **85**, 6460 (1986).
- Biermann, C., Winter, R., Benmore, C. & Egelstaff, P. Structural and dynamic properties of liquid sulfur around the λ -transition. *J. Non. Cryst. Solids* **232–234**, 309–313 (1998).
- Stuedel, R. & Eckert, B. *Molecular Spectra of Sulfur Molecules and Solid Sulfur Allotropes. Topics in Current Chemistry* **231**, (Springer, 2003).

45. Kozhevnikov, V. F., Payne, W. B., Olson, J. K., McDonald, C. L. & Inglefield, C. E. Physical properties of sulfur near the polymerization transition. *Etc* **7379** (2015).
46. Janz, G. J., Roduner, E., Coutts, J. W. & Downey, J. R. Raman Studies of Sulfur-Containing Anions in Inorganic Polysulfides. Barium Trisulfide. *Inorg. Chem.* **15**, 1751–1754 (1976).
47. Khan, S. A., Hughes, R. W. & Reynolds, P. A. Raman spectroscopic determination of oxoanions in aqueous polysulfide electrolyte solutions. *Vib. Spectrosc.* **56**, 241–244 (2011).
48. Larkin, J. M. & Strohl, W. R. Beggiatoa, *Thiothrix*, and Thioploca. *Annu. Rev. Microbiol.* **37**, 341–367 (1983).
49. Okabe, S., Ito, T., Sugita, K. & Satoh, H. Succession of Internal Sulfur Cycles and Sulfur-Oxidizing Bacterial Communities in Microaerophilic Wastewater. *Biofilms.* **71**, 2520–2529 (2005).
50. Maki, J. S. Bacterial intracellular sulfur globules: Structure and function. *J. Mol. Microbiol. Biotechnol.* **23**, 270–280 (2013).
51. Dahl, C. & Prange, A. Bacterial Sulfur Globules: Occurrence, Structure and Metabolisms. In *Inclusions in Prokaryotes*, <https://doi.org/10.1007/7171> (2006).
52. Ben Mabrouk, K., Kauffmann, T. H., Aroui, H. & Fontana, M. D. Raman study of cation effect on sulfate vibration modes in solid state and in aqueous solutions. *J. Raman Spectrosc.* **44**, 1603–1608 (2013).
53. Becucci, M., Castellucci, E., Foggi, P., Califano, S. & Dows, D. A. Temperature dependence of vibrational relaxation processes in sulfur crystals: Effect of isotopic impurities. *J. Chem. Phys.* **96**, 98–109 (1992).
54. Gleeson, D. F. *et al.* Biosignature Detection at an Arctic Analog to Europa. *Astrobiology* **12**, 135–150 (2012).
55. Douglas, S. & Yang, H. Mineral biosignatures in evaporites: Presence of rosickyite in an endoevaporitic microbial community from Death Valley, California. *Geology* **30**, 1075–1078 (2002).
56. Cosmidis, J., Nims, C., Diercks, D. & Templeton, A. S. Formation and stabilization of elemental sulfur through organomineralization. *Geochim. Cosmochim. Acta* **247**, 59–82 (2019).
57. Poffet, M. S., Käser, K. & Jenny, T. A. Thermal Runaway of Dried Sewage Sludge Granules in Storage Tanks. *Chim. Int. J. Chem.* **62**, 29–34 (2008).
58. Ghosh, W. & Dam, B. Biochemistry and molecular biology of lithotrophic sulfur oxidation by taxonomically and ecologically diverse bacteria and archaea. *FEMS Microbiol. Rev.* **33**, 999–1043 (2009).
59. Kreuzmann, A.-C. & Schulz-Vogt, N. H. N. Oxidation of Molecular Hydrogen by a Chemolithoautotrophic ABSTRACT. *Appl. Environ. Microbiol.* **82**, 2527–2536 (2016).
60. Nelson, D. C. & Castenholz, R. W. Use of reduced sulfur compounds by *Beggiatoa* sp. *J. Bacteriol.* **147**, 140–154 (1981).
61. Marnocha, C. L., Levy, A. T., Powell, D. H., Hanson, T. E. & Chan, C. S. Mechanisms of extracellular S₀ globule production and degradation in *Chlorobaculum tepidum* via dynamic cell-globule interactions. *Microbiol. (United Kingdom)* **162**, 1125–1134 (2016).
62. Macalady, J. L. *et al.* Dominant Microbial Populations in Limestone-Corroding Stream Biofilms, Frasassi Cave System, Italy. *Applied and Environmental Microbiology* **72**(8), 5596–5609 (2006).
63. Data Analysis: Curve Fitting. *OriginPro User Guide*. OriginLab Corporation, Northampton, MA (2019). Available at www.originlab.com/doc/User-Guide/.
64. Schaefer, H. F. & Palmer, G. D. Plastic and Allotropic Forms of Sulfur. *J. Chem. Educ.* **17**, 473 (1940).
65. Stolz, M. *et al.* The structural properties of liquid and quenched sulphur II. *J. Phys. Condens. Matter* **6**, 3619–3628 (1994).

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Author Contributions

J.C. and C.N. designed the study. C.N., B.C., J.M. and J.C. conducted field sample collections. C.N. performed laboratory and Raman analyses with guidance from M.W. M.W. performed *in-situ* molten sulfur Raman analysis. B.C. conducted FISH analysis. C.N., J.C., J.M. and M.W. contributed to the scientific interpretation of the data. C.N. and J.C. wrote the manuscript text and prepared the figures. All authors reviewed, contributed to, and approved the final manuscript.

Additional Information

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