An ecophysiological explanation for manganese enrichment in rock varnish


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Desert varnish is a dark rock coating that forms in arid environments worldwide. It is highly and selectively enriched in manganese, the mechanism for which has been a long-standing geological mystery. We collected varnish samples from diverse sites across the western United States, examined them in petrographic thin section using microscope imaging techniques, and investigated the associated microbial communities using 16S amplicon and shotgun metagenomic DNA sequencing. Our analyses described a material governed by sunlight, water, and manganese redox cycling that hosts an unusually aerobic microbial ecosystem characterized by a remarkable abundance of photosynthetic Cyanobacteria in the genus Chroococcidiopsis as the major autotrophic constituent. We then showed that diverse Cyanobacteria, including the relevant Chroococcidiopsis taxon, accumulate extraordinary amounts of intracellular manganese—over two orders of magnitude higher manganese content than other cells. The speciation of this manganese determined by advanced paramagnetic resonance techniques suggested that the Cyanobacteria use it as a catalytic antioxidant—a valuable adaptation for coping with the substantial oxidative stress present in this environment. Taken together, these results indicated that the manganese enrichment in varnish is related to its specific uptake and use by likely founding members of varnish microbial communities.

Rock varnish (also called desert varnish) is a thin, dark coating found on exposed surfaces of rocks in arid environments that is composed primarily of clay minerals and manganese and iron oxides (1–3). It has long been recognized as a geochemical puzzle (4–6) and has received considerable scientific and popular interest due to its widespread occurrence (7), association with archaeo logical petroglyphs (8–10), use in age dating (11–17), potential as a paleoclimate proxy (16, 18–20), and comparisons to rock coatings on Mars (21–24). However, many interpretations and applications of varnish hinge on understanding the mechanism of its formation, which remains unknown.

The most perplexing aspect of varnish is its extremely high enrichment in manganese. Varnish characteristically contains 10–30 wt% MnO—two to three orders of magnitude higher manganese content than typical underlying rocks or the surrounding dust from which much of the mass comprising varnish originates (2, 3, 25). Other major elements including iron, silicon, aluminum, magnesium, sodium, and titanium, although abundant, are not enriched.

Diverse microorganisms are known to be associated with varnish, but whether or not they play a role in its origin has been fiercely debated (26). Numerous processes, both abiotic (e.g., dust deposition, water leaching, photochemical manganese oxidation) and biological (e.g., microbial mediation of binding and cementation, microbial manganese oxidation), have been proposed to contribute (summarized in SI Appendix, Text), but since varnish grows very slowly—at most tens of microns over a thousand years (27)—empirical demonstration has not been attainable. While many of these processes may be relevant to varnish formation, none of them satisfactorily explains the highly and selectively enriched manganese content in varnish.

In this paper, we stepped back from the various paradigms that have been previously proposed and considered varnish formation from a holistic perspective. Synthesizing results from physical, chemical, and biological analyses, we reevaluated the relationship between varnish microflora and their environment and developed a hypothesis linking specific manganese accumulation to environmental adaptations of major members of the varnish microbial community.

**Results**

**Varnish Is Governed by Sunlight, Water, and Manganese Redox Cycling.** We collected varnish on a range of rock types from seven field areas across the western United States (SI Appendix, Fig. S1). Varnish occurs on diverse lithologies in different settings, yet there are some common developmental patterns. A close relationship between varnish and sunlight has been established—varnish develops preferentially on lit rather than shaded surfaces (28–30). Although found in arid environments, therein varnish develops preferentially with the availability of water, such as in shallow depressions on rock surfaces where dew accumulates (31) and along runoff streaks down cliff faces (32). These observations provided circumstantial evidence that light and water play important roles in varnish formation.

**Significance**

**Rock varnish** is a prominent feature of desert landscapes and the canvas for many prehistoric petroglyphs. How it forms—and, in particular, the basis for its extremely high manganese content—has been an enduring mystery. The work presented here establishes a biological mechanism for this manganese enrichment, underpinned by an apparent antioxidant strategy that enables microbes to survive in the harsh environments where varnish forms. The understanding that varnish is the residue of life using manganese to thrive in the desert illustrates that, even in extremely stark environments, the imprint of life is omnipresent on the landscape.


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The authors declare no competing interest.

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To understand better the physical processes controlling varnish development, we examined depositional textures in petrographic thin sections using backscatter scanning electron microscopy (SEM). Varnish cross-sections revealed micrometer-scale, sub-horizontal laminations that reflect its accretionary mode of growth (Fig. 1B and SI Appendix, Fig. S2) (3). In all varnish samples examined, we observed laminae with crinkly to columnar or domal textures that mark an emergent topography similar to that of stromatolites—macroscopic sedimentary structures commonly understood as mineralized residue of ancient microbial mats (33). In certain stromatolites, these textures have been interpreted in terms of light-dependent growth, models of which come from studies of coral growth (34). On a topographically irregular surface, relative highs receive more light while relative lows get shaded; thus, the highs grow higher forming columnar features. In varnish, these microtextures supported the relationship with sunlight that has been documented with macroscale field observations and strengthened the evidence for a role for light in varnish genesis.

Textural data also supported the previously documented relationship with water by suggesting that varnish formation involves manganese redox cycling through a soluble phase. Desert dust samples that we collected at varnish sites contained manganese as both trace Mn^{2+} in igneous minerals and manganese oxide particles (SI Appendix, Fig. S2E). We observed detrital particles embedded in varnish, but high-resolution chemical imaging by energy dispersive spectroscopy (EDS) and nanoscale secondary ion mass spectrometry (NanoSIMS) showed that these particles are largely silicate minerals, while the manganese-rich oxide material that characterizes varnish is in the laminated cement itself (SI Appendix, Figs. S3 and S4). Since manganese is water soluble in its divalent form but not as higher-valent oxides, redox cycling is required to mobilize and reprecipitate the manganese oxide in dust to form the accreting oxide cements.

The manganese oxide mineral phase in varnish has been described as poorly crystalline birnessite (2)—a phase composed of manganese octahedra organized in layers (35); it is formally Mn^{4+}O_2 but can incorporate a substantial fraction of Mn^{3+} instead of Mn^{4+} when accompanied by charge balance with heteroatoms (Na^+, K^+, Ba^{2+}, etc.) between the layers (36). Using synchrotron X-ray absorption near-edge structure (XANES) spectroscopy and multiple-energy “redox” mapping at the manganese K-edge, we found that varnish birnessite is not homogenous in its redox properties. Although predominantly Mn^{4+}, it contains common discrete microscale domains with variable and considerable mixtures of Mn^{3+} (Fig. 1C–E and SI Appendix, Fig. S2). This heterogeneity is consistent with the view that manganese redox cycling occurs within varnish, and the distribution of these domains indicates that such cycling is not restricted to surficial processes contributing to the ongoing accretion of varnish, but rather also characterizes the ecosystem that exists within well-developed varnish.

Numerous manganese redox cycling processes occur in Earth surface environments. Manganese oxidation with atmospheric O_2

![Fig. 1. Stromatolitic microtextures and manganese redox states in varnish developed on a basalt flow at Babbitt Ranch, AZ. (A) Photograph of varnished surface with a freshly broken face revealing underlying rock. (B) SEM image of a cross-section through the varnish–rock interface showing accretionary laminations that establish stromatolitic columns and domes. (C) Synchrotron X-ray microprobe map showing manganese distribution. Varnish is massively enriched in manganese relative to underlying rock. (D) Manganese redox map. Manganese in basalt is entirely Mn^{2+}, while varnish is predominantly Mn^{4+} with spatially varying domains richer in Mn^{3+}. (E) Point spectra taken across a manganese redox gradient. Inset table shows least-squares fits quantifying the components of each spectrum.](https://doi.org/10.1073/pnas.2025188118)
is thermodynamically favorable and can be catalyzed by metal oxide surfaces or bacterial and fungal enzymes (37–40). Manganese reduction can also be catalyzed by microbial processes, notably anaerobic respiration (41), and with an appropriate electron donor present (e.g., organic carbon) photochemical reduction of manganese oxides occurs readily even in aerobic settings (42–44). While any of these processes might contribute to varnish development and the manganese redox heterogeneity we observed, the relationship between varnish and sunlight raised the hypothesis that photochemistry—and perhaps photobiology—might play particularly important roles.

To visualize organic matter in varnish, we used NanoSIMS imaging of sulfur. $^{32}$S appeared throughout varnish, notably concentrated in micrometer-scale particles that sit along varnish laminations and are distinct from detrital grains observed in the $^{28}$Si and $^{57}$Fe mass channels (SI Appendix, Fig. S4A). To confirm that this sulfur reflects organics rather than just sulfur-bearing mineral phases, we used XANES spectroscopy to assess electronic structure at the sulfur K-edge. The sulfur in varnish displayed complex speciation; sulfur is present in organic forms observed in biological material (variable mixtures of thiols with disulfides, sulfoxides, and sulfonates) as well as sulfate salts (SI Appendix, Fig. S4B). The fact that varnish is rich in organics and exposed to light—conditions that promote manganese photoreduction—and yet the manganese is maintained largely as Mn$^{4+}$ oxides supports the view that dynamic redox cycling occurs within varnish.

The Varnish Microbial Community Is Characterized by Cyanobacteria.

To investigate the microbial diversity in varnish, we extracted DNA from varnish samples along with samples of surrounding surface soils for comparison. 16S rRNA gene amplicon sequencing revealed a varnish-specific microbial community that is distinct from surrounding soils, but common among varnishes from different rock types and locations (SI Appendix, Figs. S5–S7). The taxa we recovered are consistent with previous studies of varnish microbiology (45–52) (Fig. 2). 16S sequences assigned to the Xenococcaceae (either $\textit{Chroococcidiopsis}$ or unassigned below the family level) were recovered from 48 out of 49 varnish samples and accounted for 25.9% of all sequence reads and 98.7% of cyanobacterial reads recovered from varnish. In contrast, the Xenococcaceae represented only 0.06% of all reads and 1.4% of cyanobacterial reads from soil. The single most abundant unique $\textit{Chroococcidiopsis}$ sequence variant accounted for 8% of all reads from varnish and was completely absent from soil samples. Top BLAST hits for this sequence in the National Center for Biotechnology Information (NCBI) nr database included a remarkable representation of hits from previous varnish studies, and 99.6% identity with isolate Ryu 1–3 from the University of the Ryukyus in Okinawa, Japan (53), which we obtained for further study. Additionally, epifluorescence microscopy on varnish flakes revealed sarcinoid clusters of cells that we obtained for further study. Additionally, epifluorescence microscopy on varnish flakes revealed sarcinoid clusters of cells that might play an important role in the formation of varnish itself.

By abundance and function, we concluded that $\textit{Chroococcidiopsis}$ are extremely important taxa in varnish and are likely founding autotrophic members of the community. When considered in this context, the physical evidence that sunlight and water play important roles in varnish development can be interpreted to suggest that these Cyanobacteria, which grow with light and water, might be involved in the formation of varnish itself.

Metagenomic data also revealed that the varnish microbial communities mark a highly aerobic ecosystem, far more so than typical sediments or soils. No obligate anaerobic metabolisms were represented by the gene content of our MAGs—consistent with the known physiologies of the major taxa identified in our 16S analysis. Indeed, nearly all MAGs contained high-potential bioenergetic systems utilized in aerobic respiration as well as reactive oxygen species detoxification systems. Of the 38 MAGs
we recovered from varnish samples, we observed heme-copper 
O_2 reductase and/or bd O_2 reductase complexes in 37 of them, a 
superoxide dismutase in 34 of them, and a catalase in 24 of them.
These aerobic adaptations are unsurprising considering the prox-
imity of this community to atmospheric O_2 and exposure to solar 
irradiation; they are biochemical attestations to the high degrees 
of oxidative stress encountered in this harsh environment.

**Cyanobacteria Accumulate Manganese Likely as a Nonenzymatic 
Antioxidant System.** The significance of cyanobacterial taxa as 
abundant keystone members of the varnish ecosystem suggested 
to us a previously unexplored connection between the microbial 
community and the manganese content of varnish. The model 
freshwater cyanobacterium *Synechocystis* sp. PCC 6803 has been 
shown to accumulate a massive pool of intracellular Mn^{2+}, up to 
10^7 atoms per cell, which when averaged over cell volume is the 
equivalent concentration of 100 mM manganese—four orders of 
magnitude higher than their growth medium (54). We hypothe-
sized that if this hyperaccumulation of manganese occurs broadly 
in the Cyanobacteria—particularly those taxa that are dominant 
members of the varnish community—then this physiological pecu-
liarity might underpin varnish development.

We examined two strains of *Chroococcidiopsis* (PCC 7433 and 
Ryu 1–3, the closest cultured relative to the varnish sequences), 
*Synechocystis* sp. PCC 6803, and *Gloeobacter violaceus* PCC 7421, 
a very deep branching member of the photosynthetic Cyanobacteria 
(SI Appendix, Fig. S9), along with noncyanobacterial model or-
ganisms *Escherichia coli* K12 and *Shewanella oneidensis* MR-1 for 
comparison. Using inductively coupled plasma mass spectrometry 
(ICP-MS) to measure total cellular manganese, we observed 
dramatic accumulation in all cyanobacterial strains; the highest values were seen in Ryu 1–3, with over two orders of 
magnitude greater manganese content than that seen in *E. coli* or 
*S. oneidensis* (Fig. 3A).

To probe the speciation of this copious intracellular manganese, 
we combined electron paramagnetic resonance (EPR), electron 
nuclear double resonance (ENDOR), and electron spin echo en-
velope modulation (ESEEM) spectroscopies. The EPR spectra of 
all Cyanobacteria examined showed that the majority of the Mn^{2+} 
pool exists in soluble high-symmetry complexes with low–molecular-
weight ligands (denoted H–Mn^{2+}) and not in low-symmetry com-
plexes with strongly chelating ligands or bound to proteins (denoted 
L–Mn^{2+}) (Fig. 3B and SI Appendix, Fig. S10 A and B). 31P, 1H 
ENDOR, and 14N ESEEM measurements along with 31P ENDOR 
on labeled cells further revealed that this manganese binds unde-
tectably low amounts of phosphate and nitrogenous ligands, and is 
instead predominantly bound to carboxylato ligands (Fig. 3C 
and SI Appendix, Fig. S10 C and D and Text). Notably, previously 
studied organisms with abundant cellular H–Mn^{2+} exhibited sig-
ificant contributions from phosphate and nitrogenous ligands 
(55, 56); this difference makes the Mn^{2+} pool in Cyanobacteria 
distinctive.

This speciation data implied the functional role for Mn^{2+} in 
cytophysiology as a small-molecule antioxidant system. Certain H–Mn^{2+} complexes are known to act as effective 
catalytic antioxidants of superoxide and peroxide (57, 58), and 
accumulation of H–Mn^{2+} has been shown to predict oxidative 
stress resistance (55, 59). This physiology is well documented in 
radioactive-resistant taxa such as *Deinococcus radiodurans* (60, 61), 
but was not previously known in Cyanobacteria. The H–Mn^{2+} 
accumulation shown here helps explain how the Cyanobacteria 
in varnish cope with the extremely aerobic, arid, and irradiated 
environments where varnish is found. Taken together with the 
high abundance of cyanobacterial taxa associated with varnish, 
the establishment of this cyanobacterial manganese-based anti-
oxidant system provides an adaptive biological mechanism be-
hind the enrichment of manganese in rock varnish.

**Discussion**

Varnish develops in environments that are extremely harsh, 
where protective strategies against irradiation and oxidative 
stress are essential for life to survive. The presence of the varnish 
microbial community has been noted in such terms; for example, 
varnish provides a habitat for microbial life shielded by oxide 
minerals that absorb UV radiation (62). However, up until now, 
how and why the varnish—with its high and specific enrichment 
in manganese—got there in the first place was a mystery. Based 
on the data presented here, we propose that varnish itself is a 
hallmark of life prevailing in these extreme environments.

We showed that Cyanobacteria of the genus *Chroococcidiopsis* 
are intimately and abundantly associated with varnish. Moreover, 
being the main primary producers of the ecosystem, these or-
ganisms must establish prior to heterotrophic taxa that depend 
on them for organic substrates, implicating them as likely founding 
members of the microbial community. Evidence tying sunlight and 
water to varnish development provided further, albeit circum-
stantial, evidence that photosynthetic Cyanobacteria play a fun-
damental role. We then demonstrated that diverse Cyanobacteria, 
including *Chroococcidiopsis*, accumulate substantial quantities of 
H–Mn^{2+}. This phenomenon is well established as an antioxidant 
strategy that enables tremendous oxidative stress resistance (58, 
59, 63); indeed, there is no other known physiological purpose for 
such elevated manganese concentrations. The cyanobacterial H–Mn^{2+} pool that we observed is comparable in magnitude to 
the most radiation-resistant organisms known (60).
Many previous studies have focused on a mechanism of manganese oxidation as the key to varnish formation. However, in such aerobic environments, manganese oxidation may proceed through numerous pathways with both biological (SI Appendix, Text and Fig. S8) and abiotic (including photochemical and autocatalytic) mechanisms. Rather than oxidation, we argue that it is the selective enrichment of manganese that represents a process of singular importance to the development of varnish. The manganese hyperaccumulation that we observed in *Chroococcidiopsis* provides a simple and effective, ecologically relevant, physiological explanation for this manganese enrichment.

Thus, we propose a hypothesis for varnish formation (Fig. 4). These Cyanobacteria grow on sunlit rock surfaces with intermittent access to water, sequestering high concentrations of manganese in their cells and exploiting the unique redox chemistry of manganese complexes as a catalytic antioxidant system that enables their survival in such a harsh environment. When they die, the residue from their biomass provides an enriched manganese source that is ultimately oxidized to form the oxide mineral cement that comprises varnish. Since varnish forms over timescales of millennia, a well-developed varnish sample represents the time-integrated manganese accumulation of many, many generations of cells, which are sparsely distributed at any given time. This was demonstrated in our analyses of varnish samples in petrographic thin section—the Mn$^{2+}$/Mn$^{4+}$ content of any living cells is negligible next to thousands of years of accumulated Mn$^{2+}$/Mn$^{4+}$ mineral. In addition to solving the mystery of manganese enrichment, cyanobacterial exudates and necromass also supply fresh organic carbon to the varnish ecosystem. This provides a growth substrate for the heterotrophic microbes that inhabit varnish, as well as an effective electron donor for photochemical manganese reduction.

Taken together, our results place the activity of extremophilic Cyanobacteria as a key driver of both the physical and biological development of varnish ecosystems.

Fig. 4. Natural history of varnish. (A) Manganese is delivered largely as oxide particles in windborne dust, reduced by either photochemical or biological processes, and taken up by *Chroococcidiopsis* cells for use as a catalytic antioxidant. (B) *Chroococcidiopsis* grows with light and water, fixing carbon and trapping accumulated manganese. (C) Dust material not adhered to the rock surface is removed by wind or precipitation. (D) When *Chroococcidiopsis* cells die, the manganese-rich residue left behind by their biomass is oxidized to generate the manganese oxides that comprise varnish. This oxidation could be biologically catalyzed and/or abiotic. (E) Products from cyanobacterial photosynthesis serve as substrates for heterotrophic community members. (F and G) Manganese redox cycling continues in developed varnish, with abundant O$_2$ as an electron acceptor (F) and organic matter as an electron donor (G).

**Methods**

**Study Locations and Sampling.** Samples were taken from seven field locations across the southwestern United States, ranging from arid to semiarid climates, corresponding to rock varnish Type I and Type II designated by Macholdt et al. (16). Geology and lithology (Table S1) included mafic, felsic, and sedimentary rocks. Varnished rocks were collected with ethanol-sterilized gloves into Whirl-Pak sample bags, and surface soil samples were collected using sterile spatulas into Falcon tubes. Ultrathin sections cut orthogonal to the varnished surface were prepared by David Mann at High Mesa Petrographics (Los Alamos, NM). The thin sections revealed varnishes ranging in thickness from ∼5 to ∼100 μm, likely representing hundreds to thousands of years of development (27). For DNA sampling, varnished rocks were returned to the laboratory, gently rinsed with sterile nanopure water using a 50-mL syringe and 16-gauge needle, and allowed to dry. For DNA extraction, varnish was removed from rock surfaces by scraping with flame-sterilized steel brushes, spatulas, and dental picks, and collected in a weigh boat. Each varnish sample was obtained from a separate rock. Dust analysis was conducted on a fine grained (clay-silt sized) fraction of surface soil from the Barstow field location.

**SEM/EDS/NanoSIMS.** SEM and EDS analyses were conducted in the Caltech Geological and Planetary Sciences Division Analytical Facility on a FEI Quanta 3D Field Emission SEM, with a Robinson-type backscatter electron detector and an Oxford X-Max SDD X-ray EDS system. Varnish was imaged at 10–20 kV with working distances of 7–10 mm and magnifications of 500–4,500×. NanoSIMS analyses were conducted in the Caltech Microanalysis Center, on a Cameca NanoSIMS 50-30 using a 50-nm raster primary ion beam with 50-nm ion current of 1 pA, and dwell time of 3.5 ms/pixel. The masses of $^{14}$C, $^{24}$Mg, $^{34}$S, $^{30}$P, $^{18}$O, $^{54}$Mn$^{16}$O, and $^{57}$Fe$^{16}$O were collected. Petrographic thin sections were coated with 20-nm graphite using a Turbo carbon evaporator for SEM/EDS, and with 40-nm gold using a Cressington HR metal sputter coater for NanoSIMS. EDS and NanoSIMS images were examined using ImageJ.

**Synchrotron X-Ray Spectroscopy.** Synchrotron analyses were conducted at the Stanford Synchrotron Radiation Lightsource, on X-ray microprobes at beamlines 2-3 for the manganese K-edge and 14-3 for the sulfur K-edge. The 2-3 beam was energy calibrated using the pre-edge feature of KMnO$_4$ at 6,543.34 eV, and 14-3 was calibrated using the pre-edge feature of Na$_2$SO$_4$ at 2472.02 eV. Multiple energy maps for producing images of manganese redox state were collected at 6,553, 6,557, 6,559, 6,562, and 6,570 eV, with 3-μm resolution. For both spectra and maps, least-squares fitting was done using a pseudo-Voigt function of Gaussian and Lorentzian components. The resulting fractions of Mn$^{2+}$, Mn$^{3+}$, Mn$^{4+}$ and Fe$^{3+}$O$^2$O$^-$ were calculated using the fit results and contributions of each component to the overall spectrum. The maps were reduced and fit using the SIXPACK (67) and SMAK (68) software packages, respectively (https://www.sams-xrays.com/). Colormaps were converted to viridis using fixthejet (https://fixthejet.eclife.org/).

**DNA Extraction and Sequencing.** Genomic DNA was extracted from 10 to 50 mg varnish scrapings or soil using the FastDNA SPIN kit for soil (MP Biomedicals). DNA yields were quantified using a Qubit 2.0 fluorometer with the HS dsDNA assay kit (Thermo Fisher Scientific).

To generate 16S rRNA gene libraries, the V3–V4 hypervariable region of the 16S gene was amplified using degenerate primers (341–806 pair) from ∼12.5 ng of genomic DNA with KAPA HiFi HotStart ReadyMix (denaturation at 95 °C for 3 min, 20 cycles of 95 °C for 30 s, 55 °C for 30 s, and 72 °C for 30 s, and a final extension of 72 °C for 5 min before holding at 4 °C). A second round of PCR added Nextera XT v2 indexes (Illumina) (denaturation at 95 °C for 3 min, 8 cycles of 95 °C for 30 s, 55 °C for 30 s, and 72 °C for 30 s and a final extension of 72 °C for 5 min before holding at 4 °C). A second round of PCR added Nextera XT v2 indexes (Illumina) (denaturation at 95 °C for 3 min, 8 cycles of 95 °C for 30 s, 55 °C for 30 s, and 72 °C for 30 s and a final extension of 72 °C for 5 min before holding at 4 °C). The amplicons were cleaned up using AMPure XP beads (Beckman Coulter). A no-template control was processed and did not show a band in the amplicon region. The amplicons were pooled and sequenced on the Illumina MiSeq platform generating paired end 301-bp reads using the MiSeq reagent kit v3 (600 cycles) (Illumina).

Two representative varnish DNA samples (sample 24, sandstone from Babbitt Ranch, AZ, and sample 41, basalt from Mesa Prieta, NM) were selected for metagenomic sequencing. These samples were selected based on 16S amplicon data as likely candidates to recover high-quality genomes for the major taxa in the varnish community. No aspects of the communities in these two samples were outliers, and together they represent samples from two different locations and very different rock types.

Illumina shotgun libraries were prepared using the Next Ultra DNA II library preparation kit (New England Biolabs). DNA was fragmented using a Covaris E220; the ends were made blunt and adapters and indexes added.
onto the fragments to generate Illumina libraries, which were eluted in DNA elution buffer (Zymo). Libraries were quantified using the KAPA Illumina/ Universal Quantification kit, normalized based on qPCR tables containing the frequencies of each unique sequence variant per sample. Quality filtering, denoising, merging of paired end reads, and chimera removal were done using DADA2 (70). The QIME2 q2-feature-classifier plugin was used to align the sequences against the Greengenes 13.8 database (71) and assign taxonomy. One sample that returned <2,000 total reads was omitted from downstream analyses. NMDS and ANOSIM analyses were done by calculating a Bray dissimilarity matrix using the vegan ecology package in R (72). LEfSe analysis was done using the Microbiome Analyst tool (73) with default settings.

For the phylogenetic tree presented in this paper, sequences were aligned with the SINA aligner (74) and converted from fasta format to phylip format with SEAVIEW (75). The phylogenetic tree was constructed using FigTree.

**Metagenomic Data Processing.** Taxonomic assignments of raw metagenomic reads were done using the MG-RAST analysis platform (77). Metagenomic sequence read quality control, de novo assembly, and binning of MAGs was performed largely on the KBase platform (78). Combinations of read preprocessing, assembly, and binning methods were tested and evaluated based on the quality and quantity of final MAGs. Raw preprocessing included no processing (raw reads), Bloom Filter Read Error Correction (79) v. r181 (drop_unique_kmer-reads < 1, kmer_size = 33), Trimmomatic (80) v. 0.39, LEADING 3 TRIMMING 3, SPLITnę WINDOW14 15 MINLEN 125, and BBMap (81) v7.3793 (bbduk.sh -Xmx1g, k = 15, mink = 10, trimr = r, tbo). Processed reads were assembled with MEGAHIT (82) v2.4.2 (metalarge), MetaSPAdes (83) v1.2.4 (K-mer sizes 33, 55, 77, 99, and 127), and IDBA-UD (84) v1.0.4. Assemblies were performed with minimum contig size of 2,000 bp. MAGs were constructed with two automated binning tools, MaxBin2 (85) v2.2.4 and MetaBAT (86) v1.7 with default settings except for minimum contig lengths of 2,500 bp. All MAGs from combinations of the above setting were evaluated for quality using the Checkm software v1.0.18 on KBase with default settings. Sufficiently high-quality bins were defined using the Parks et al. (87) cutoff of completeness > 50% contamination > 0%. Taxonomy was assigned using GTDB-Tk v.0.3.2 using gtdbtk release 49 (88). The number of bins passing this cutoff and their phylogenetic affiliations were used to compare between the different parameters described above. The greatest number of quality bins was achieved with Trimmomatic, MetaSPAdes, and MetaBAT2 for sample 21, and Trimmomatic + BBmap + MetaSPAdes, and MetaBAT2 for sample 24. Other combinations of parameters produced a subset of less complete versions of the final bins, not completely different MAGs. Genome annotation was conducted using the RASTkr algorithm (89) v.0.1.11 in KBase, as well as a local implementation of the KFOamSc (90) software for the KEGG database. The canonical metabolic pathways encoded by these genomes were parsed using KEGG-decoder (91), and additional analyses of genes of interest, including implementation of the KOfamScan (90) software for the KEGG database.

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**Fluorescence Microscopy.** Fluorescence microscopy was performed at the Center of Integrated Nanotechnologies at Los Alamos National Laboratory. Cavity well microscope slides (Globe Scientific) were used to mount varnish flattened to 50 mL with nanopure water. ICP-MS analysis was conducted in the Stanford Synchrotron Radiation Lightsource (SSRL, SLAC National Accelerator Laboratory) with a 50-mL Nano Spray source (Syage Lab, U.S.A.) using a 10-μm tip and 100% acetonitrile elution buffer (Zymo). Libraries were quantified using the KAPA Illumina/Universal Library Quantification kit, normalized based on qPCR

**EPR/ENDOR/ESR Spectroscopy.** Concentrated cell suspensions in 40% glycerol were loaded into custom-made quartz EPR tubes, and then flash-frozen in liquid N2. Paramagnetic resonance spectroscopy was conducted in the Stanford Synchrotron Radiation Lightsource (SSRL, SLAC National Accelerator Laboratory). The 35-GHz continuous-wave (CW) EPR spectra were recorded using a laboratory-built EPR spectrometer (94). Absorption-display EPR spectra of frozen cells and manganese standards were collected in the “rapid passage” mode at 2 K as previously described (56, 59) (microwave [MW] frequency, 34.9 GHz; MW power, 1 mW; temperature, 2 K; modulation amplitude, 1 G; time constant, 64 ms; scan rate, 1 mG/min).

Pulsed ENDOR/ESR spectra were recorded using a laboratory-built 35-GHz pulsed EPR spectrometer (95). All spectra were recorded at 2 K using an immersion helium cryostat. 31P, 1H Davies ENDOR spectra were recorded using the pulse sequence  t = τ − t2 − τ − t − τ − τ − t − τ − τ − τ − t − τ − τ − t − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − t − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − τ − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t − t -