

Anoxic carbon flux in photosynthetic microbial mats as revealed by metatranscriptomics

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Photosynthetic microbial mats possess extraordinary phylogenetic and functional diversity that makes linking specific pathways with individual microbial populations a daunting task. Close metabolic and spatial relationships between Cyanobacteria and Chloroflexi have previously been observed in diverse microbial mats. Here, we report that an expressed metabolic pathway for the anoxic catabolism of photosynthate involving Cyanobacteria and Chloroflexi in microbial mats can be reconstructed through metatranscriptomic sequencing of mats collected at Elkhorn Slough, Monterey Bay, CA, USA. In this reconstruction, *Microcoleus* spp., the most abundant cyanobacterial group in the mats, ferment photosynthate to organic acids, CO₂ and H₂ through multiple pathways, and an uncultivated lineage of the Chloroflexi take up these organic acids to store carbon as polyhydroxyalkanoates. The metabolic reconstruction is consistent with metabolite measurements and single cell microbial imaging with fluorescence in situ hybridization and NanoSIMS.

Subject Category: integrated genomics and post-genomics approaches in microbial ecology

Keywords: metatranscriptomics; NanoSIMS; anoxic carbon flux; fermentation; glycogen

Introduction

Linking biogeochemical processes observed in natural microbial communities with their associated metabolic pathways and assigning these pathways to specific groups remains a daunting task, particularly as community complexity increases (Kunin et al., 2008a). Photosynthetic microbial mats found in marine and hypersaline environments are stratified microbial communities that harbor extraordinary phylogenetic and functional diversity, harboring up to 42 phyla, including many candidate divisions (Nicholson et al., 1987; Ley et al., 2006; Kunin et al., 2008b; Dillon et al., 2009). Extensive measurements have been performed on these communities, documenting biogeochemical cycling of carbon,

nitrogen and sulfur (Jorgensen and Des Marais, 1986; Canfield and Marais, 1993; Hoehler et al., 2001; Hoehler et al., 2002; Decker et al., 2005). The microbial populations involved in these biogeochemical cycles have been primarily inferred based on the physiology of isolates and expression of individual functional genes (Omeregíe et al., 2004; Green et al., 2008; Orphan et al., 2008). However, a comprehensive understanding of element cycling in the mats requires reconstruction of complete metabolic pathways expressed by specific microbial groups.

Photosynthetic microbial mats perform light driven carbon fixation during daylight hours (Bebout et al., 2002). Photosynthate is accumulated, often as glycogen, through both oxygenic and anoxygenic photosynthesis (Nicholson et al., 1987). This photosynthate is fermented to organic acids and H₂ under anoxic conditions at night (Hoehler et al., 2002; Burow et al., 2012). The flux through these fermentation products varies up to four orders of magnitude between day and night (Hoehler et al., 2001). Recent studies of mats collected at Elkhorn Slough, Monterey Bay, CA, USA, has provided persuasive evidence that

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Microcoleus spp., the most abundant cyanobacterial group in the mats, are the dominant fermenters (Burow *et al.*, 2012). However, the pathways through which *Microcoleus* spp. ferment photosynthate are unknown, as are microbial groups and pathways that may further metabolize the organic acids derived from fermentation. Relative measures of activity comparing rRNA transcripts to genes in samples from the Elkhorn Slough mats have indicated that *Chloroflexi* are among the most active microbial group under dark, anoxic conditions (Burow *et al.*, 2012). Studies in thermophilic mat systems collected in Yellowstone National Park, which are much less complex than marine and hypersaline mats, have suggested a link between anoxic fermentation of accumulated photosynthate by *Cyanobacteria* and carbon uptake by *Chloroflexi* (van der Meer *et al.*, 2003, 2005, 2007). The intimate physical association of *Cyanobacteria* and *Chloroflexi* was demonstrated in hypersaline mats collected from Guerrero Negro, Baja California, where fluorescence *in situ* hybridization was used to visualize *Chloroflexi* filaments within the exopolysaccharide sheaths of *Microcoleus* spp. (Ley *et al.*, 2006). The close links between *Cyanobacteria* and *Chloroflexi* observed in other microbial mat environments suggest that this association may be a general phenomenon observed in mats and that this association may play an important role in anoxic carbon cycling.

Metatranscriptomic analysis has been used to characterize active microbial communities in aquatic and terrestrial environments (Poretsky *et al.*, 2005; Frias-Lopez *et al.*, 2008; Gilbert *et al.*, 2008; Urich *et al.*, 2008; Turnbaugh *et al.*, 2010). Gene expression from microbial assemblages has been correlated with biogeochemical data and the phylogeny of expressed genes affiliated with specific populations (Poretsky *et al.*, 2009, 2010; McCarren *et al.*, 2010; Hollibaugh *et al.*, 2011; Liu *et al.*, 2011; Mou *et al.*, 2011; Shi *et al.*, 2011). In engineered microbial systems where complexity is reduced, the expression of specific pathways has been defined and correlated with environmental conditions (He *et al.*, 2010). Metatranscriptomics of microbial mats offer the possibility of defining key metabolic pathways in the biogeochemical cycling of carbon and of establishing a link in the carbon cycle between *Cyanobacteria* and *Chloroflexi*.

Here, we demonstrate that metatranscriptomic data obtained from Elkhorn Slough microbial mat samples collected under dark anoxic conditions were dominated by transcripts from *Cyanobacteria* and *Chloroflexi*. This density of sequenced transcripts allowed us to reconstruct pathways for anoxic carbon catabolism that link fermentation in the *Cyanobacteria* with organic acid uptake and carbon storage in the *Chloroflexi*. These metabolic pathway inferences were supported by ¹³C-acetate isotopic labeling studies of the Elkhorn Slough mat coupled with single cell microbial imaging.

Materials and methods

Sample site and collection

The sampling site was located in the Elkhorn Slough estuary in Central California, USA, at 36N 48' 46.61" (decimal degree Lat 36.81295) and -121S 47' 4.89" (decimal degree Long -121.7847). Microbial mats were sampled on 10 January 2009 and transported to a greenhouse facility at NASA Ames Research Center within 1–2 h. In the greenhouse, mat pieces were placed in acrylic aquaria transparent to ultraviolet radiation and covered with water for ca. 20 h before the beginning of a diel cycle study (Bebout *et al.*, 2002; Burow *et al.*, 2012). Diel cycle studies were carried out under natural solar irradiance, and the water temperature was kept constant at ca. 18 °C (*in situ* average). The data presented result from diel cycle studies carried out on the 12th/13th January and 13th/14th January 2009.

Biogeochemical assays

Replicate vials were prepared identically as follows: small subcores (11-mm diameter, 15-mm depth or vertically sectioned for depth profile analyses) were cut from whole sections of intact microbial mat and placed in serum vials with 4 ml of field site water. Serum vials were capped with butyl rubber stoppers. The 10.5-ml headspace of the serum vials was left as air for light/daytime incubations and was thoroughly flushed with N₂ (gas and liquid phase degassed) for dark/nighttime incubations. H₂ was allowed to accumulate in three replicate vials and was repeatedly sampled at specific time points over the course of the diel experiment. Organic acids were allowed to accumulate in replicate vials (separate to vials in which H₂ was measured), with three vials sacrificed at each time point. Sections of microbial mat were also incubated individually in triplicate vials as described above after sectioning to identify the location of H₂ production. Sectioning using a sterile scalpel blade yielding the following mat layers; 0–2 mm, 2–4 mm and 4–15 mm. To measure H₂, 25 µl of headspace gas was withdrawn by volumetric syringe and injected directly onto a gas chromatograph with quantification by a mercuric oxide detector. To analyze organic acids, the entire liquid phase (4 ml) was sampled (with the associated incubation sacrificed). Liquid was filtered through 0.2 µm syringe-driven filters for storage in glass vials at -20 °C. Organic acids (C1–5) were quantified via high-pressure liquid chromatography (Albert and Martens, 1997).

Nucleic acid extraction from Elkhorn Slough mat cores

Ten mat cores of 1-cm diameter were flash frozen in liquid nitrogen at multiple time points during the diel cycle and stored at -80 °C until further processing. RNA of two samples (BN; 2100 hours, 12 January 2009; 4 h after sunset, and EN; 0700

hours, 13 January 2009, 20 min before dawn) was extracted from the uppermost 2 mm of 5 mat cores by combining phenol–chloroform extraction with parts of the RNeasyMini kit (Qiagen, Valencia, CA, USA). Per core, biomass was transferred in a tube containing 0.5 ml RLT buffer and homogenized using a rotor-stator homogenizer (Omni International, Kennesaw, GA, USA). The suspension was then bead-beated with zirconium beads (200 μ m, OPS Diagnostics, Lebanon, NJ, USA) and the cell debris and beads pelleted. The supernatant was extracted with phenol–chloroform–isoamyl alcohol (125:24:1, pH 4.5, Ambion, Austin, TX, USA). The aqueous phase was run through the gDNA eliminator spin column to remove genomic DNA and further purified following the RNeasyMini kit protocol. Extracted RNA was treated with DNase using the TURBO DNA-free kit according to the protocol (Ambion). Amplification, sequencing and sequence analysis of 16S rRNA genes and transcripts is described in Supplementary Information.

Selective removal of rRNA and cDNA synthesis

The MICROBEnrich and MICROBExpress Kits (Ambion) were used to remove the ribosomal RNAs (rRNA) by a subtractive hybridization approach, thereby enriching the messenger RNA (mRNA) in the total RNA pool. In the MICROBEnrich protocol, seven additional eukaryotic capture probes, and in the MICROBExpress Kit, five additional cyanobacterial capture probes were used. The sequences can be found in Supplementary Information. Approximately 400 ng of mRNA were linearly amplified with the MessageAmp II-Bacteria Kit (Ambion) according to the manufacturer's instructions. In all, 4 μ g of the amplified, antisense RNA (aRNA) were converted to double-stranded cDNA with random hexamers in multiple replicates using the SuperScript Double-Stranded cDNA Synthesis Kit (Invitrogen, Carlsbad, CA, USA) and purified with DNA Purification Spin Columns.

The quality and quantity of total RNA, mRNA, aRNA and cDNA were verified by measurements on the NanoDrop-1000 Spectrophotometer (NanoDrop Technologies, Wilmington, DE, USA), with the Qubit fluorometer (Life Technologies, Grand Island, NY, USA) and the Bioanalyzer (Agilent Technologies, Santa Clara, CA, USA).

Metatranscriptomic sequencing and analysis

Double-stranded cDNA of the two samples (BN, 12 January 2009, 2100 hours and EN, 13 January 2009, 0700 hours) were sequenced at the Department of Energy Joint Genome Institute (JGI, Walnut Creek, CA, USA) on a 454 Genome Sequencer FLX Instrument (454 Life Sciences, Branford, CT, USA). After sequencing, ribosomal RNA reads were detected by BLAST search and removed from the data set. Non-rRNA sequences were analyzed for replicate sequences using the open-source program

CD-HIT provided as a web-based tool at <http://microbiomes.msu.edu/replicates> according to a previously described protocol (Li and Godzik, 2006; Gomez-Alvarez *et al.*, 2009). Replicates were defined as sequences sharing >99% nucleotide identity, with an allowable length difference of 2 bp, and a requirement that the first 3 bp of the replicate sequences be identical.

The BLASTX program in BLAST version 2.2.21 was used with dereplicated non-rRNA reads as query sequences against all amino-acid sequences in the microbial RefSeq release 45 (January 2011) database (Altschul *et al.*, 1990; Pruitt *et al.*, 2009). RefSeq BLAST hits with bitscores below 40 were removed, and the results were processed using MEGAN version 4.32 with default settings (Huson *et al.*, 2007). Taxonomic assignments based on NCBI taxonomy and functional assignments based on SEED and KEGG were analyzed using graphical tools within MEGAN (Ogata *et al.*, 1999; Overbeek *et al.*, 2005).

Completeness of sampling estimates were made using Chao and ACE estimators in MOTHUR version 1.21.1 (Schloss *et al.*, 2009). Read clusters used as input for MOTHUR were generated in two ways. First, reads with a top BLASTX hit to the same RefSeq protein sequence were considered part of the same mRNA transcript cluster and used to estimate absolute coverage. Second, reads unambiguously assigned by MEGAN to a single SEED functional category or species (as defined by NCBI taxonomy) were considered part of the same functional or species cluster and used to calculate functional and species sampling completeness, respectively.

FISH-SIMS

Small subcores (11-mm diameter, 15-mm depth) of the microbial mats were cut from whole sections of intact microbial mat and placed in serum vials with 4 ml water obtained from the field site. Serum vials were capped with butyl rubber stoppers and the headspace of the vials (10.5 ml) was thoroughly flushed with N₂ (gas and liquid phase degassed) for dark/nighttime incubations. A stable isotope tracer, [2-¹³C]-acetate (99% ¹³C, Sigma, St Louis, MO, USA), was added to the liquid phase of vials containing live cores at a final concentration of 0.2 mM. As a control, cores pretreated with 4% paraformaldehyde (final concentration) were also incubated with 0.2 mM [2-¹³C]-acetate. The cores were incubated under dark, anoxic conditions for 10 h. The incubation was stopped by adding 4% paraformaldehyde to the live cores. Preliminary NanoSIMS analysis revealed that single cells in the live cores were labeled but not in the killed cores pretreated with paraformaldehyde. The method used to label *Chloroflexi* with fluorescent oligonucleotides was previously described (Woebken *et al.*, 2012). *Chloroflexi* were targeted using equimolar amounts of the FISH (fluorescence *in situ* hybridization) probes CFX1223 and GNSB-941 (Björnsson *et al.*, 2002). High-resolution secondary ion mass spectrometry

(SIMS) was performed at LLNL with a Cameca NanoSIMS 50. The $^{13}\text{C}/^{12}\text{C}$ ratio was measured using $^{12}\text{C}_2$ and $^{13}\text{C}^{12}\text{C}$ corrected for the dimer abundances (Finzi-Hart *et al.*, 2009).

Data deposit

The dereplicated metatranscriptomic sequences with rRNA sequences removed can be downloaded from IMG/M (<http://img.jgi.doe.gov/cgi-bin/m/main.cgi>) under the names Elkhorn Slough cyanobacterial mat night (2100 hours Metatranscriptome CGUI) (BN library) and Elkhorn Slough cyanobacterial mat day (0700 hours transcriptome CGUN) (EN library) at the tab 'FASTA nucleic acid file for all scaffolds'. 16S rRNA sequences obtained in this study are deposited under GenBank accession numbers JX002103–JX002655.

Results

Microbial mat characterization

H_2 production was $>10\times$ higher at night in the upper phototrophic layer (0–2 mm) in the Elkhorn Slough mat compared with H_2 produced during the day or in the lower mat layers (Supplementary Figure 1a). Acetate, formate and propionate were concomitantly produced with H_2 , whereas lactate, butyrate and valerate were not detected (200 nM detection limit; Supplementary Figure 1b). H_2 and organic acid measurements observed for the microbial mat sampled in January 2009 were similar to those previously reported from this field site in November 2009 (Burow *et al.*, 2012).

The microbial community composition in the upper layer was assessed using amplicon pyrosequencing of the V8 region of SSU rRNA genes from a time point taken before sunrise (EN; 0700 hours, 13 January 2009); sequences grouping with 16 distinct phyla were observed (Supplementary Table 1 and Figure 1a). The active microbial community was investigated by amplicon pyrosequencing of the V8 region of SSU rRNA from cDNA derived from RNA extracted at the same time point (EN) and at the beginning of the dark period (BN; 2100 hours; 12 January 2009). The active communities were subsets of the total number of phyla (11 and 13 phyla, respectively) and dominated by *Cyanobacteria*. *Chloroflexi* sequences were present at low abundance in both the DNA and cDNA pyrotag libraries.

Nearly full-length 16S rRNA sequences were consistent with amplicon pyrosequencing of the cDNA (Figure 1a). Cyanobacterial sequences clustered into a few groups (Figure 2), of which the majority were related to *Microcoleus chthonoplastes* PCC 7420 and a cyanobacterium recently isolated from the Elkhorn Slough mats (ESFC-1) (Woebken *et al.*, 2012). The *Chloroflexi*-related sequences were only distantly related to cultured representatives, but were closely related to sequences from uncultivated *Chloroflexi* recovered from hypersaline environment at Guerrero Negro, Baja California (Figure 3) (Ley *et al.*, 2006).

Metatranscriptome sequence analysis

A total of 590 783 and 492 302 metatranscriptome sequence reads were generated for the BN and EN

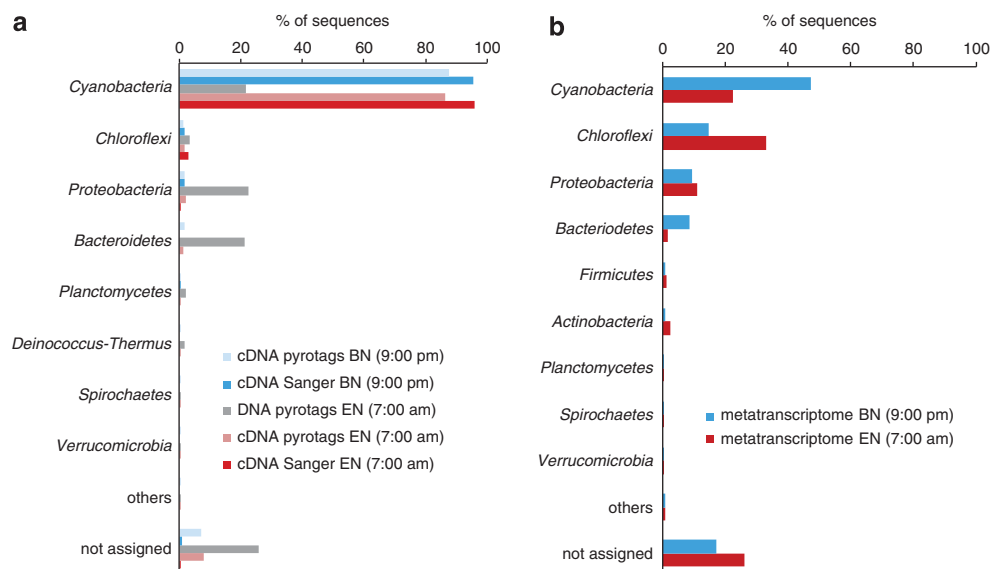


Figure 1 (a) Comparison of nearly full-length Sanger and pyrotag amplicon 16S rRNA sequences of the phototrophic layer from Elkhorn Slough mat analyzed using the RDP classifier (Wang *et al.*, 2007) with a confidence threshold of $\geq 80\%$. Unassigned reads are sequences that could not be classified with a confidence threshold of $\geq 80\%$ on the phylum level. (b) Taxonomic classification of metatranscriptome reads analyzed by MEGAN using the RefSeq database (NCBI). Unassigned reads have an unknown taxonomic affiliation.

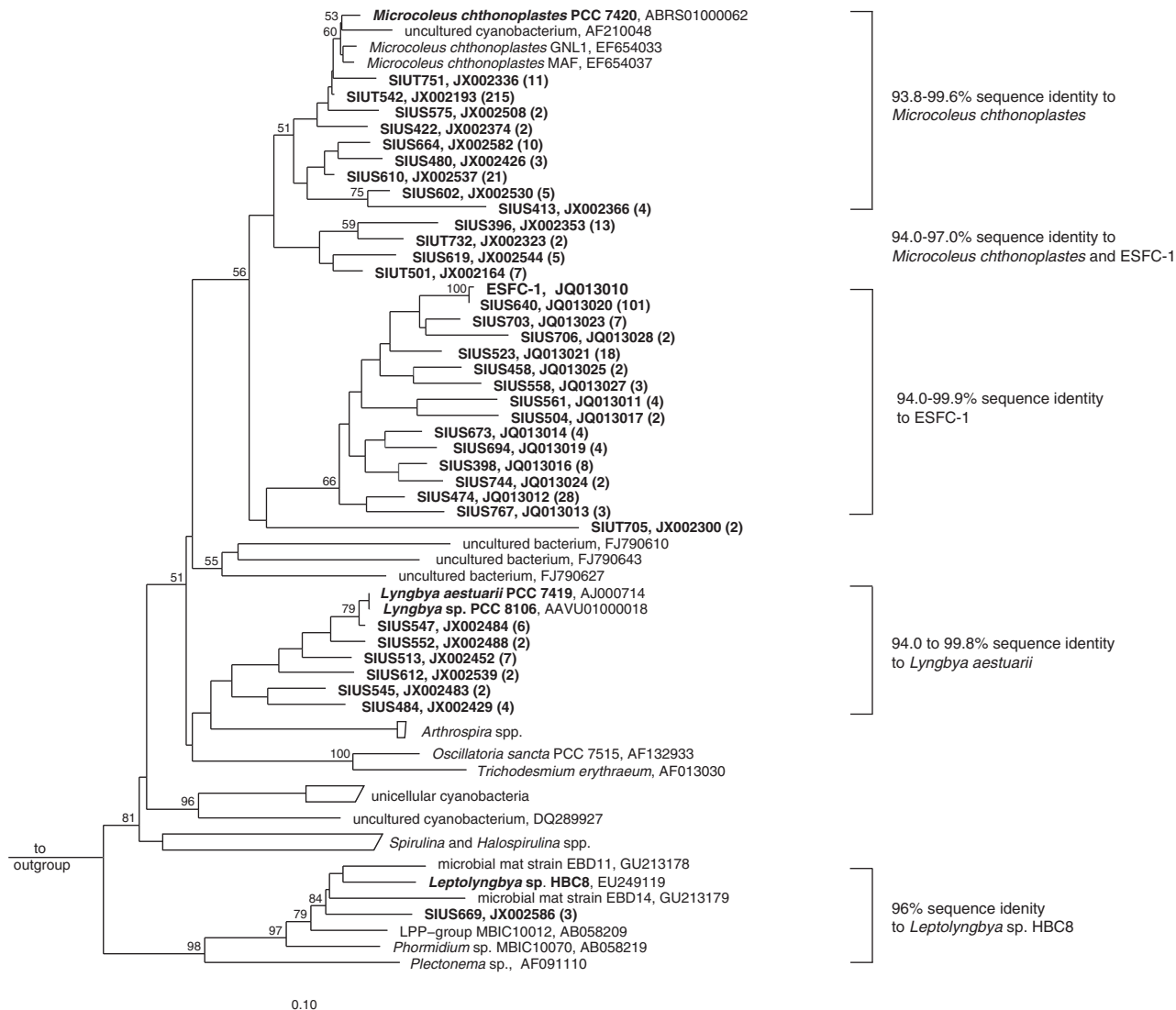


Figure 2 Neighbor joining tree of *Cyanobacteria*-related nearly full-length 16S rRNA sequences from cDNA samples recovered from BN (SIUS) and EN (SIUT) samples. OTU₉₈ representatives of OTUs that contain more than one sequence (numbers of sequences per OTU in parenthesis) are depicted in the tree. Bootstrap values calculated with the PhyML algorithm that were $\geq 50\%$ are displayed in the tree. Scale bar represents 10% estimated sequence divergence.

time points of which 43% (BN) and 32% (EN) were non-ribosomal sequences (Supplementary Table 2). After removal of these rRNA sequences and dereplication, 244 004 (BN) and 144 017 (EN) sequences remained for further analysis.

Coverage of the total pool of unique mRNA transcripts in the samples was analyzed in three ways: absolute, functional and species-level taxonomic. Absolute coverage was estimated by BLAST against the microbial RefSeq database, where reads yielding top BLAST hits from the same RefSeq amino-acid sequence were considered part of the same mRNA transcript. This RefSeq BLAST is likely to underestimate transcript diversity, as only 64.6% (BN) and 68.7% (EN) of reads for each sample showed significant similarity (bitscore > 40) to any RefSeq sequence (Supplementary Table 3).

Collector's curves showed that sampling of the total number of unique mRNA transcripts is far from complete (Supplementary Figure 2a).

Estimations of functional and species-level coverage were obtained using the unambiguous assignment by MEGAN of reads to SEED functional categories or species. Although only 37.9–40.4% of reads were given a functional assignment and 15.6–22.9% given an unambiguous species-level assignment, both time points were sampled relatively completely with respect to the functional and species-level diversity presently described in RefSeq (Supplementary Table 3; Supplementary Figure 2b). These results are likely a reflection of the limited scope of the SEED and RefSeq databases compared with the full natural range of functions and species, rather than that of the true

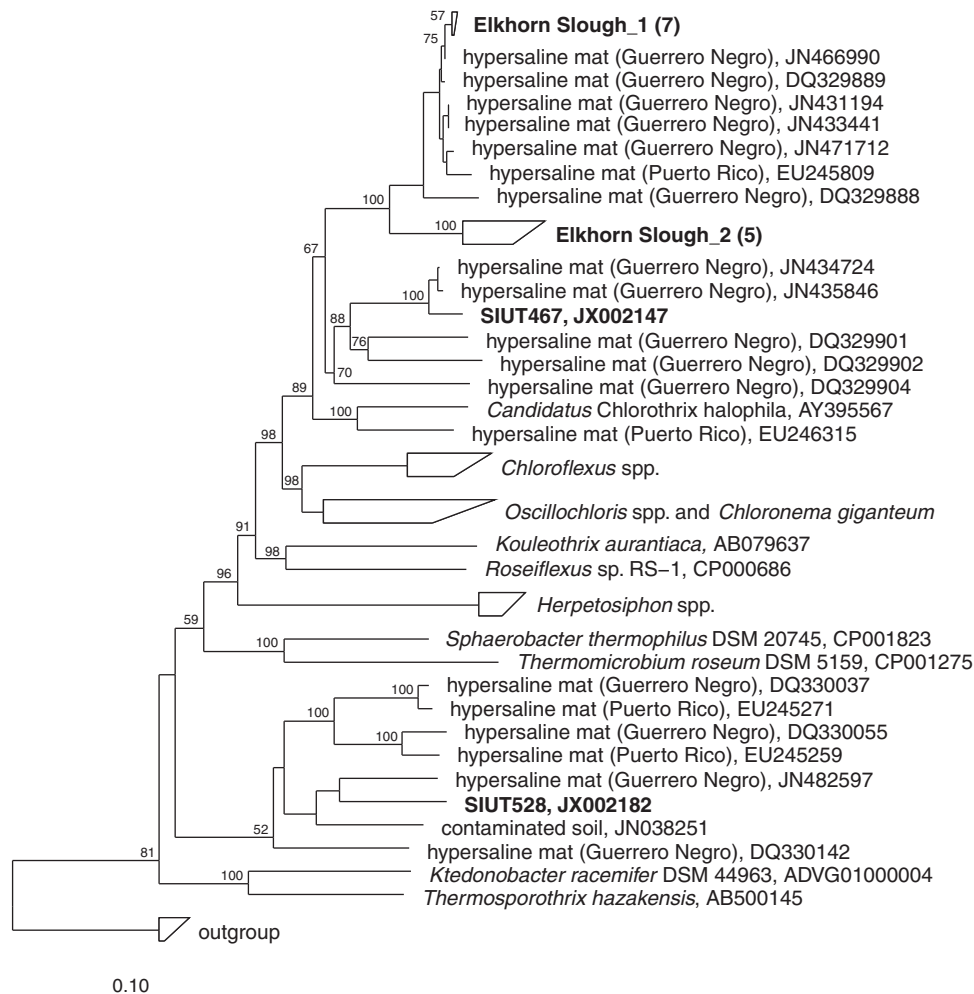


Figure 3 Neighbor joining tree of *Chloroflexi*-related nearly full-length 16S rRNA sequences from cDNA samples BN (SIUS) and EN (SIUT). Elkhorn Slough mat sequences group with sequences retrieved from other microbial mat sites and are only distantly related with cultured *Chloroflexi*. Bootstrap values calculated with the PhyML algorithm that were $\geq 50\%$ are displayed in the tree. Scale bar represents 10% estimated sequence divergence.

completeness of sampling. Moreover, although functions and species were well sampled independently, by no means were all the functions for every species sampled as well. However, these results do mean that these metatranscriptomic samples are broadly representative of those functions and species present in both the sample and SEED/RefSeq databases.

Abundant taxa in metatranscriptomes

The taxonomic composition of the microbial mat was assessed using the MEGAN software package for all protein-coding transcripts. Both metatranscriptomes were dominated by reads assigned to *Cyanobacteria* and *Chloroflexi*, the former constituting 22–47% of the reads, the latter 15–33% (Figure 1b). *Proteobacteria* and *Bacteroidetes*-affiliated reads were the next abundant groups. A large proportion of the reads were recruited to the *M. chthonoplastes* PCC 7240 genome. Of the 8294 predicted protein-coding genes in the *M. chthonoplastes* genome, 4030

(49%) were detected in the BN library and 1821 (24%) were detected in the EN library. Estimates of richness using Chao and ACE indices predicted the expression of 4978 and 4948 *M. chthonoplastes* protein-coding genes in the BN transcriptome, suggesting 81% (Chao) or 80% (ACE) coverage. These indices predicted 62% (Chao) and 48% (ACE) coverage in the EN transcriptome. Transcripts associated with *Microcoleus* spp. represented 24% ($n = 37\,524$) of all BN transcripts with matches to the RefSeq database ($n = 157\,525$) and 8% ($n = 7581$) of all EN transcripts with matches to the RefSeq database ($n = 98\,894$). These reads are not necessarily all derived from a single *Microcoleus* relative, and may reflect a diverse pool of *Microcoleus* spp. that are more closely related to *M. chthonoplastes* PCC 7240 than to any other sequenced cyanobacterial genome. Unlike the *Microcoleus* spp., the *Chloroflexi* populations present in the mat samples did not have a closely related genome for fragment recruitment. Within the metatranscriptome reads

affiliated with the phylum *Chloroflexi*, 96% were assigned to the class *Chloroflexi* at both time points, and ca. 85% of those reads further to the order *Chloroflexales* (BN; $n=18\,769$ and EN; $n=26\,981$, respectively; Supplementary Figure 3).

Anoxic carbon catabolism by *Cyanobacteria* and *Chloroflexi*

Transcript reads from *Microcoleus* spp. associated with carbohydrate metabolism were overrepresented when compared with carbohydrate reads observed for the whole community (BN; $n=18\%$ and EN; $n=22\%$; Supplementary Figure 4). To elucidate the carbon catabolic pathways in the *Microcoleus* populations in the mat samples,

fermentation pathways for conversion of glycogen were reconstructed from the *M. chthonoplastes* genome (Figure 4a; Supplementary Tables 4 and 5). Genes for glycogen phosphorylase and the EMP (Embden-Meyerhof-Parnas) pathway coded for proteins for the conversion of stored photosynthate, glycogen, to pyruvate. Genes for pyruvate transformation via three pathways were identified: to lactate (lactate dehydrogenase), to formate and acetyl-CoA (pyruvate-formate lyase) and to acetyl-CoA and CO_2 (pyruvate-ferredoxin oxidoreductase). Genes to transform acetyl-CoA to ethanol (alcohol/aldehyde dehydrogenase) and acetate (acetate kinase) were identified. No annotated phosphotransacetylase was detected and no reads that cluster with other cyanobacterial phosphotransacetylases were

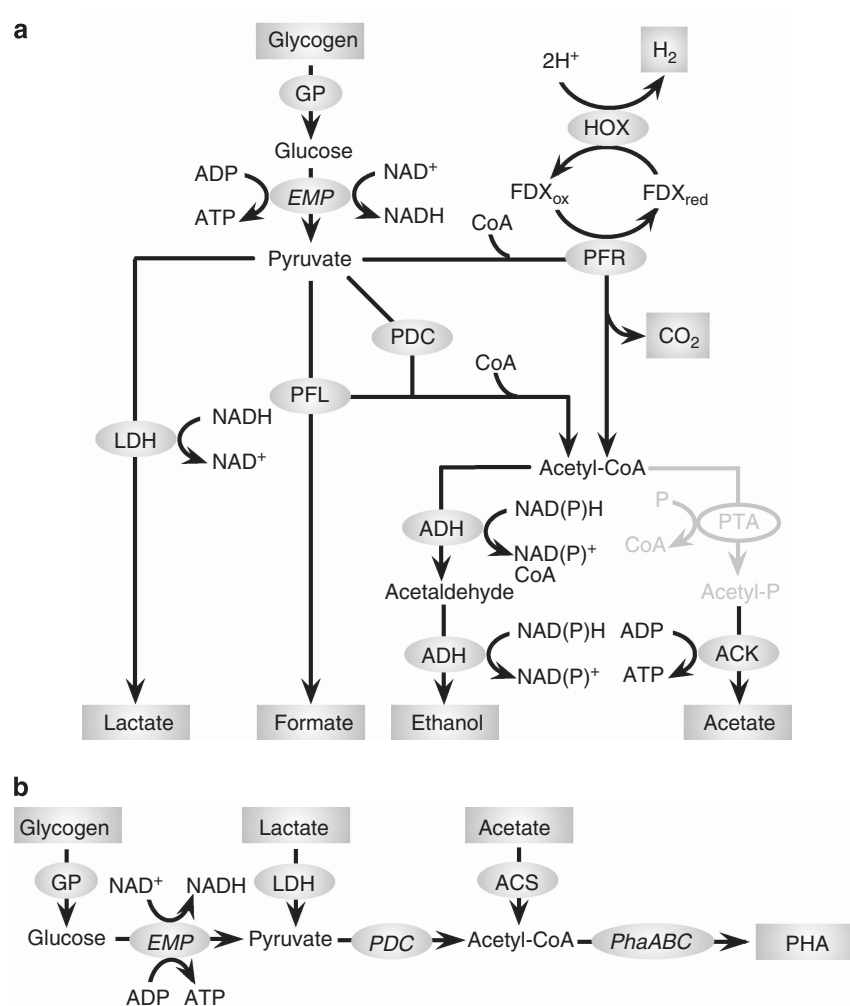


Figure 4 (a) Reconstruction of anoxic low-molecular-weight metabolite producing pathways based on the *M. chthonoplastes* genome. Abbreviations: ACK, acetate kinase; ADH, alcohol/aldehyde dehydrogenase; GP, glycogen phosphorylase; HOX, bidirectional [NiFe] hydrogenase; LDH, lactate dehydrogenase; PFL, pyruvate-formate lyase; PFR, pyruvate:ferredoxin oxidoreductase. Phosphotransacetylase (PTA) was not identified. (b). Reconstruction of glycogen to PHA conversion pathway in sequenced *Chloroflexales* genomes. Abbreviations: PDC, pyruvate dehydrogenase complex; PhaA, acetoacetyl-CoA transferase; PhaB, acetoacetyl-CoA reductase; PhaC, polyhydroxyalkanoate synthase. Detailed assignments for these genes are presented in the Supplementary Information (Supplementary Tables 4 and 5). Quantitative measurements of recruited reads from BN and EN data sets are presented in Figure 5 and Supplementary Table 6.

identified in the transcript reads. The catalytic [NiFe] Hox hydrogenase was identified as part of a predicted seven gene operon with NAD(P)H dehydrogenase subunits (*hoxEFUYH*), a predicted aldehyde/alcohol dehydrogenase (*adhE*) and a hypothetical protein. Transcripts for all the genes described above were observed in the BN and EN libraries (Figure 5; Supplementary Table 6). Higher numbers of reads for the fermentative pathway genes were observed in the BN sample, consistent with the higher level of *Microcoleus* expression at the beginning of the dark period. The *Microcoleus* genes were the majority of all the cyanobacterial-associated reads assigned to genes in the fermentation pathway.

A number of *Chloroflexales*-associated reads were assigned as acetate- and lactate-permeases, consistent with the hypothesis that these microbes take up the organic acids excreted during fermentation by *Microcoleus* spp. (Figure 4b; Supplementary Tables 4–6). Organic acid uptake is often associated with production of polyhydroxyalkanoates (PHAs) that act as carbon storage compounds and uptake of acetate has been shown to be involved in the conversion of glycogen to PHAs in bacteria responsible for EBPR (enhanced biological phosphorus removal) (Mino *et al.*, 1998). Pathways for the biosynthesis of PHAs were reconstructed from multiple *Chloroflexales* genomes and transcripts were identified for *phaA* (acetoacetyl-CoA transferase), *phaB* (acetoacetyl-CoA reductase) and *phaC* (PHA synthase) in both the BN and EN libraries

(Figure 5). The identification of *Chloroflexales*-associated transcripts for glycogen phosphorylase in both libraries implies that reducing equivalents required for PHA production in these bacteria derive from the breakdown of glycogen. Transcripts for acetyl-CoA synthetase (*acs*), which converts acetate to acetyl-CoA, were also detected in both libraries. Accumulation of glycogen by the *Chloroflexi* is proposed to occur during anoxygenic photosynthesis to store carbon and reducing equivalents for dark, anoxic metabolism (Sirevag and Castenholz, 1979; Taffs *et al.*, 2009). These reducing equivalents are generated by glucose metabolism through the EMP pathway to produce pyruvate, which is oxidized to acetyl-CoA by pyruvate dehydrogenase, which is detected in both libraries.

¹³C-Acetate uptake by *Chloroflexi* observed by FISH-SIMS

The reconstructed pathway described above suggested that the *Chloroflexi* in the Elkhorn Slough mat assimilated organic acids under dark, anoxic conditions. Single cell analysis using NanoSIMS was used to test the specific hypothesis that acetate, the most abundant organic acid observed in the mats, was assimilated by the *Chloroflexi*. Mat samples were incubated with 0.2 mM ¹³C-acetate during the dark portion of the diel cycle and *Chloroflexi* cells in these samples were stained by the CARD-FISH procedure with *Chloroflexi*-specific oligonucleotide probes (Figure 6A) and identified by

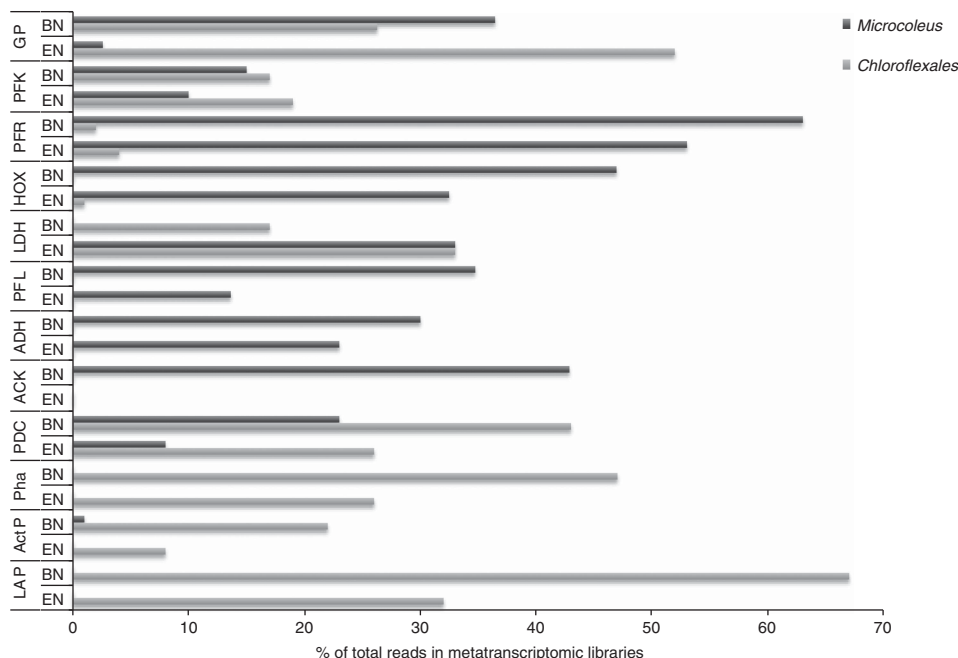


Figure 5 Comparison of reads involved in carbon catabolic pathways in BN and EN libraries. Reads representing transcripts for the *Microcoleus* spp. fermentation pathway and the *Chloroflexales* glycogen to PHA conversion. The reads are presented as a relative proportion of the total reads assigned to this SEED functional category. Pha (*phaCAB* and *HOX* (*hoxEFUYH*) represent the sum of multiple transcripts. Actual numbers of reads recruited to the genes in each category are presented in the Supplementary Information (Supplementary Table 6).

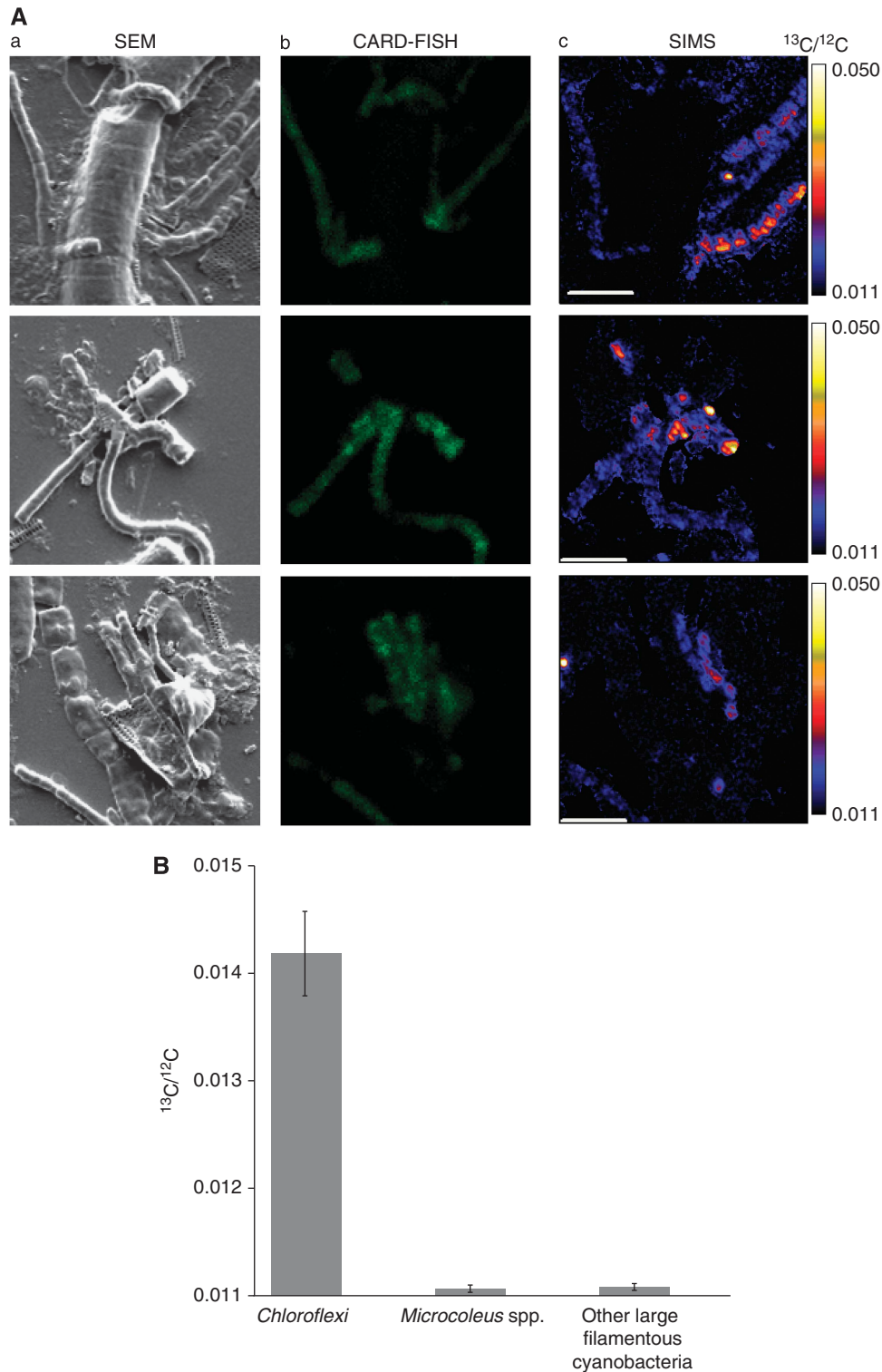


Figure 6 (A) ^{13}C -acetate uptake under dark, anoxic conditions within the microbial community of the Elkhorn Slough mat. (a) Scanning-electron micrographs (SEM). (b) CARD-FISH micrographs using CF mix probes (CFX1223 and GNSB-941) targeting *Chloroflexi* in green. (c) Ion micrographs of $^{13}\text{C}/^{12}\text{C}$ ratios of the same community members imaged by SEM and CARD-FISH. Scale bar = 5 μm . (B) ^{13}C -acetate uptake under dark, anoxic conditions by different microbial groups in the Elkhorn Slough mat. *Chloroflexi* ($n=40$) were identified using CF mix CARD-FISH probes. *Microcoleus* spp. ($n=12$) and other *Cyanobacteria* ($n=12$) were identified by their morphological characteristics. Error bars represent the standard error of the mean. Note that the probes do not distinguish *Chloroflexi* from *Chloroflexales* so these results are discussed at the phylum level.

confocal laser scanning microscopy. This analysis revealed a morphological diversity consistent with the phylogenetic diversity detected in nearly full-length 16S rRNA sequences from Elkhorn Slough mats (Figure 3), in which the sequence identity of the *Chloroflexi* phylotypes ranged from 76.6% to 89.6%. The identified *Chloroflexi* were targets for NanoSIMS analysis measuring $^{13}\text{C}/^{12}\text{C}$ in individual bacterial cells. These FISH-SIMS experiments demonstrated specific uptake of ^{13}C -labeled acetate into *Chloroflexi* cells ($\delta^{13}\text{C} = 260 \pm 35\text{‰}$; $n = 40$), with the observed ^{13}C -label in *Microcoleus* ($-15.8 \pm 3.0\text{‰}$; $n = 12$) and other cyanobacterial cells ($-14.0 \pm 2.8\text{‰}$; $n = 12$) close to natural abundance (Figure 6B). Additional single cells were enriched in ^{13}C above natural abundance, but did not stain with the *Chloroflexi*-specific probes.

Discussion

In this study, we have taken a highly diverse microbial system and used metatranscriptomic sequencing to identify genes in the two most abundant taxa, *Cyanobacteria* and *Chloroflexi*, which are responsible for anoxic catabolism of photosynthate. The success of this reconstruction is remarkable because of the relatively modest amount of sequencing (~ 400 MB) used. It is highly unlikely that a comparable amount of metagenomic sequencing would have provided a resolved metabolic pathway because the diversity of the community would prevent assembly of large contiguous sequences (Kunin *et al.*, 2008b). Metatranscriptomic sequencing focuses on the most highly expressed genes, which can resolve important metabolic processes. Central to the success of this strategy was the availability of isolate genome sequences for *Microcoleus* and the *Chloroflexales* that facilitated metabolic pathway reconstruction and fragment recruitment of transcripts. A notable discrepancy in the metatranscriptomic data obtained from Elkhorn Slough samples is the difference in relative abundance of the rRNA transcripts, which identify *Cyanobacteria* as the dominant active population, compared with the mRNA transcripts, which fluctuate between *Cyanobacteria* and *Chloroflexi* as the dominant populations. This phenomenon has been observed, though less dramatically, in metatranscriptomic studies of a microbial mat from Mushroom Spring in Yellowstone (Liu *et al.*, 2011). The dominance of cyanobacterial rRNA may reflect the synthesis of large numbers of ribosomes by cyanobacterial cells, which have larger cell sizes in comparison to other bacterial cells. Cell size and ribosome content have previously been directly correlated in prokaryote cells (Ecker and Kokaisl, 1969). Alternatively, it could reflect an amplification bias in the PCR reaction with cDNA from the mats. However, since two different forward primers were

used for pyrotag sequencing and Sanger clone libraries, this explanation is not very likely.

Combining the metatranscriptomic data and metabolic pathway reconstruction with metabolite and single cell measurements enabled us to propose an anoxic catabolic pathway for carbon fixed during the light period by oxygenic and anoxygenic photosynthesis. In this pathway, photosynthate accumulated during the day is stored as glycogen by both *Microcoleus* spp., derived from oxygenic photosynthesis and the *Chloroflexi*, derived from anoxygenic photosynthesis. Under dark anoxic conditions, *Cyanobacteria* depolymerize and ferment the glycogen, excreting H_2 and organic acids and generating ATP for cellular maintenance. *Chloroflexi* depolymerize glycogen, generating reducing equivalents to transform organic acids cross-fed from the *Cyanobacteria* to PHAs, generating ATP for cellular maintenance. Our analysis does not exclude other pathways for anoxic metabolism of organic acids, such as sulfate reduction by δ -proteobacteria, but these pathways could not be completely reconstructed from the metatranscriptomic data sets (Risatti *et al.*, 1994).

Reconstruction of fermentation pathways from the *M. chthonoplastes* genome identified multiple pathways for pyruvate fermentation, all of which were expressed in the mats samples analyzed by metatranscriptomics. Therefore, the predominant organic acid end products of pyruvate fermentation were predicted to be acetate, formate and lactate. Acetate and formate were observed, consistent with the predicted activity of pyruvate ferredoxin oxidoreductase (PFR), pyruvate formate-lyase and pyruvate dehydrogenase complex. However, no lactate was observed, consistent with both the low number of observed lactate dehydrogenase transcripts and the detection of reads for lactate permeases from the *Chloroflexi*. Propionate production was also observed, which suggests that *Microcoleus* may have a pathway to ferment pyruvate to propionate, similar to pathways present in propionibacteria (Himmi *et al.*, 2000). Interestingly, no genes in the *Microcoleus* genome or cyanobacterial transcripts in the two data sets could be assigned as a phosphotransacetylase despite the identification of multiple pathways in *Microcoleus* to produce acetyl-CoA and the production of high levels of acetate in the mat samples. This observation suggests that an unassigned protein may substitute for phosphotransacetylase in *Microcoleus* fermentation. The recruitment of metatranscriptomic reads provides persuasive evidence that H_2 is generated by a NAD(P)H-dependent Hox hydrogenase, consistent with previous studies which utilized PCR and microarray-based methods (Burow *et al.*, 2012, Marshall *et al.*, 2012). The expression of PFR may imply that reducing equivalents for H_2 are generated by oxidation of pyruvate to acetyl-CoA, as is observed for [FeFe] hydrogenases (Schut and Adams, 2009). However, the specific interactions between PFR

and the Hox hydrolase are unknown, despite the co-occurrence of PFR with Hox in all available cyanobacterial genomes (Carrieri *et al.*, 2011).

An important hypothesis arising from analysis of the metatranscriptomic data was that the *Chloroflexi* should take up organic acids under dark, anoxic conditions. The linkage of organic acid uptake to anoxic conversion of glycogen to PHA has previously been observed in bacteria performing EBPR during wastewater treatment (Mino *et al.*, 1998). The postulated link between fermentation products excreted by *Microcoleus* and carbon compounds taken up by *Chloroflexi* is consistent with observations of cross-feeding between these populations in other microbial mats. Glycogen fermentation and organic acid excretion by *Cyanobacteria* has been demonstrated in thermophilic *Synechococcus*-dominated mats found in alkaline siliceous hot springs (Octopus Spring and Mushroom Spring) in Yellowstone National Park using both ^{14}C -labeling and transcript analysis, which demonstrated upregulation of *Synechococcus* fermentation genes (Nold and Ward, 1996, Steunou *et al.*, 2006). In these mats, thermophilic *Chloroflexales* related to *Roseiflexus castenholzii* are also abundant and pulse labeling experiments with $^{13}\text{CO}_2$ and $^{14}\text{CO}_2$ provided evidence for the transfer of cyanobacterial-derived photosynthate from *Synechococcus* to the *Roseiflexus* populations (van der Meer *et al.*, 2005). The intermediacy of acetate was demonstrated by lipid analysis of *Synechococcus* mat samples incubated with ^{13}C -acetate. However, this acetate incorporation was observed during the morning, in contrast to observations with the *Microcoleus*-dominated mats described here, and may be directly linked to photoheterotrophic growth. The cross-feeding of *Synechococcus* and *Roseiflexus* populations in the Yellowstone mats has served as the basis for developing *in silico* models of mat carbon metabolism that are broadly consistent with the anoxic carbon metabolism observed at Elkhorn Slough (Taffs *et al.*, 2009). Though accumulation of PHAs by mat dwelling *Chloroflexi* has never been directly demonstrated, studies of microbial mats found in the Great Sippewissett Salt Marsh in Massachusetts, USA which contain *Chloroflexi* populations, have shown that PHAs accumulate at night and decrease during the day (Rothermich *et al.*, 2000). The accumulation of PHA in the dark was putatively linked to bacterial breakdown of glycogen, which supports the model for the Elkhorn Slough mats derived from metatranscriptomic data. Additionally, studies of carbon metabolism in *Chloroflexus aurantiacus* have shown that this model organism produces both glycogen and PHA (Sirevag and Castenholz, 1979).

Acetate uptake and incorporation into PHA was demonstrated in single cells of uncultured *Accumulibacter* spp., a microbial population important in EBPR, by MAR-FISH using ^{14}C -labeled acetate (Burow *et al.*, 2008). Evidence for acetate uptake by

single cells affiliated with the *Chloroflexi* was obtained by application of FISH-SIMS to the mat samples using ^{13}C -labeled acetate and *Chloroflexi*-specific oligonucleotide probes. Other single cells that were not affiliated with the *Chloroflexi* were observed with enriched $^{13}\text{C}/^{12}\text{C}$ ratios, indicating that additional bacterial groups assimilated acetate under anoxic conditions. Future work will focus on cultivating *Chloroflexi* from the Elkhorn Slough mats to test the proposed linkage of acetate uptake to glycogen conversion to PHA and identifying other group involved in the anoxic metabolism of organic acids derived from fermentation.

In conclusion, metatranscriptomic sequencing of samples obtained from microbial mats has demonstrated that *Cyanobacteria* and *Chloroflexi* are the most active groups under dark, anoxic conditions. Genomic reconstruction and fragment recruitment of transcripts was used to reconstruct a pathway for the catabolism of photosynthate that linked fermentation by *Microcoleus* with PHA production by the *Chloroflexi*. This work highlights the utility of metatranscriptomics to define pathways for element cycling in complex microbial communities.

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