Antibiotic Resistance Genes from Mat and Planktonic Microbiomes of Antarctic Lake Fryxell and Lake Bonney, and a Mat-forming Culture of *Rhodoferax antarcticus*

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ABSTRACT

The Antarctic Taylor Valley Lakes Fryxell and Bonney harbor oligotrophic microbial communities that are separated geographically from other aquatic systems. Their microbiomes include planktonic as well as lift-off mat communities that float to the underside of the perennial ice cover and eventually emerge at the surface. We investigated the antibiotic resistance genes (ARGs) from metagenomes of lift-off mats emerging from ice, from filtered water samples of Lake Fryxell, and from filtered water samples of Lake Bonney. ARG sequence markers were designed by ShortBRED-Identify using the Comprehensive Antibiotic Resistance Database (CARD). The overall proportion of ARG hits in the metagenomes was found to be similar to that found in temperate-zone rural water bodies with moderate human inputs (0.0002-0.0007%). The specific ARGs found showed distinct distributions for the two lakes, and for mat versus planktonic sources. An enrichment culture of *Rhodoferax antarcticus* from a Lake Fryxell mat sample showed a mat-forming phenotype not previously reported for this species. Its genome showed no ARGs associated with Betaproteobacteria, but had ARGs consistent with a *Pseudomonas* minor component. The Antarctic lake mats and water showed specific ARGs distinctive to the mat and water sources, but overall ARG levels were similar to those of temperate water bodies.

INTRODUCTION

One of Earth's coldest dry deserts, the Antarctic Taylor Valley possesses a string of stratified lakes with unique geochemistry and microbial communities (1–4). The microbial genomic analysis of these lakes remains limited (5–7). There are a number of reports on antibiotic resistance genes (ARGs) in Antarctic soil or marine bacteria (7–11) but few studies of ARGs in Antarctic lake water or their mat communities (12). The prevalence of ARGs in relatively pristine habitats is of compelling importance for humans and for environmental study (13, 14). We report on the taxonomic composition and ARG prevalence in the water of Lake Fryxell and Lake Bonney, and in the phototrophic mat communities of Lake Fryxell.

Along the Taylor Valley, 80 km across the Ross Sea from McMurdo Station, lies Lake Fryxell, 10 km downstream of Lake Bonney (**Fig. 1A**). No permanent stream connects the two lakes; their main hydrology involves glacial meltwater and an underground brine aquifer (15). Their water ecosystems are entirely microscopic, with no fauna larger than nematodes. The lakes and their ecosystems are studied as models for ancient Mars (16) and as indicators of climate change (17).

Lakes Fryxell and Bonney have permanent stratified layers in the water column that give rise to oligotrophic microbial communities (2, 18, 19). The lack of turnover allows for the growth of benthic microbial mats that form flame-shaped towers several centimeters tall (20, 21). The microbial mats have been studied with primary emphasis on the role of cyanobacteria such as *Microcoleus* and *Oscillatoria* (20, 22) although metagenomes reveal a high abundance of proteobacteria (5) and protists (18, 23). Known as "lift-off" mats, the benthic biofilms reach upward, buoyed by oxygen bubbles, until pieces break off and float upward to the undersurface of the ice cover (24, 25). During winter, water freezes beneath the mat fragments, adding to the

ice layer, but above the lake the valley's dry winds ablate the ice, exposing the dessicated mats (**Fig. 1B**). Surviving five years trapped in ice, the mat material remains viable, and psychrophiles may continue growing (26). When mat organisms reach the surface of ablated ice, they can blow off and enter "moats" of melted ice that occur in summer, surrounding the permanent ice in the main part of the lake. The mat material carries communities that include microscopic arthropods and nematodes; thus, the mats can act as vectors for transport of entire ecosystems (24).

Molecular genetic data on Dry Valley microbes remained limited until recently (19, 22, 27). In December, 2014, the austral summer, we obtained water samples from Fryxell and Bonney as well as desiccated mat samples emerging from the ice cover of Fryxell. We applied tools of metagenome analysis to characterize diversity, and we sequenced a novel genome from a mat-forming *Rhodoferax* enrichment culture that differs from the commonly reported cyanobacterial mats.

We surveyed our mat and water metagenomes for the presence of ARGs, referenced to the Comprehensive Antibiotic Resistance Database (CARD) (28). Antibiotic resistance is an ancient, widespread phenomenon naturally and historically occurring in all environments (13, 14). Non-anthropogenic processes select for ARGs in pristine habitats; for example, cyanobacterial blooms drive increases in bacterial ARG prevalence and diversity (29). Inputs of human origin, even at relatively modest levels, can add substantially to the native ARG pool as well as add additional types of ARGs (8, 10, 12).

RESULTS

Taxa abundance of mat and water samples.

We characterized the taxa abundance from three microbial sources: the water columns of Lake Fryxell (FRY-01, 02, 03) and Lake Bonney (BON-01, 02, 03), and the lift-off mat from the ice surface of Lake Fryxell (Mat 01 – 06) (**Fig. 2**). Metagenomic DNA was sequenced by Illumina MiSeq. The 150-kb sequence reads were assigned to taxonomic classes using the Kraken 2 classifier pipeline (30, 31) with taxa percentages calculated via the Bracken abundance estimator (32) (described below under Methods).

The ice-surface mat samples (Mat-01 through Mat-06) showed substantial amounts of cyanobacteria, consistent with previous reports for Fryxell mats obtained from benthic samples (5). For Mat-01, 02, 03, 06 the cyanobacterial assignments were primarily Oscillatoriales and Nostocales. Mat-04 and Mat-05 however showed depletion of Oscillatoriales, with predominant abundance of Betaproteobacteria. Throughout the six mat samples, other taxa with significant abundance included Actinobacteria, Bacteroidetes, and Alphaproteobacteria, with smaller abundance of Planctomycetes, Bacteroidetes and Firmicutes. Thus the ice-trapped, air-exposed community showed composition similar to that of benthic samples (5).

The filtered water from both lakes Fryxell and Bonney had almost no cyanobacteria but a high proportion of eukaryotes, in some cases more than 50% of the reads classified by Kraken2/Bracken. Microbial eukaryotes including phototrophs and mixotrophs play important roles in the lake ecology (18). Abundant bacterial taxa included Actinobacteria, Bacteroides, Betaproteobacteria, and Alphaproteobacteria, similar to previous reports of Fryxell water (19) and other glacier-associated water bodies (26).

The validity of taxa classifier pipelines is highly dependent on their algorithm and taxa database. For comparison with Kraken2/Braken output, our mat and water reads were mapped to core taxonomic markers using the MetaPhlAn2 pipeline (33, 34) (**Fig. 3**). Four of the six mat samples showed mainly cyanobacteria, consistent with longstanding reports of cyanobacterial mats (20–22). Consistent with the Kraken analysis, Mat-04 and Mat-05 samples showed major amounts of Betaproteobacteria and Alphaproteobacteria, with depletion of cyanobacteria. For the filtered water samples, MetaPhlAn2 markers include limited content of Eukarya, and the pipeline did not find the eukaryotic taxa predicted by Kraken2 (**Fig. 2**). The filtered water from both lakes showed mainly Actinobacteria, with some Betaproteobacteria. Overall MetaPhlAn2 appeared less effective than Kraken 2/Bracken at classifying Antarctic taxa.

ARG composition characterized using ShortBRED.

Gene sequences encoding various forms of antibiotic resistance, including genomic loci as well as mobile elements, are collected in the Comprehensive Antibiotic Resistance Database (CARD) (28). We used the ShortBRED pipeline to identify ARGs in our samples, based on a set of marker peptides matching CARD sequences. The percentage of reads from each sample that matched ARGs ranged from 0.0001-0.0006% (**Fig. 4**). The overall ARG abundance was simlar amongst the three sample classes (Fryxell mat, Fryxell water, Bonney water). It was comparable to the total ARG abundance of temperate-zone rural water bodies with moderate human inputs, in a study where samples were analyzed using the same ShortBRED marker set (35).

The specific ARGs identified by ShortBRED were sorted by abundance across samples (**Table 1;** Supplementary file **Table S1**). Eight of the top 20 sorted ARGs showed a difference among the three sample classes (Kruskal-Wallis test, $P \leq 0.05$); of these results, 7 of 8 are

predicted to be significant. Of the top-ranked ARGs, BUT-1, and vancomycin-resistance genes *vanYA*, *vanTG* and *vanI* appeared mainly in the water column. BUT-1 is class-C beta-lactamase reported in *Buttiauxella agrestis*, an environmental Gammaproteobacterium found in environmental water sources (36, 37). The vancomycin resistance genes occur in *Enterococcus* and other Gram-positives (38). By contrast, the beta-lactamase AAC(3)-Ia (39) occurred only in the mat samples. AAC(3)-Ia is encoded on *Pseudomonas aeruginosa* integrons as well as in other Proteobacteria. Other ARGs showed possible differences in prevalence between Lake Bonney and Lake Fryxell water; for example, the Streptomyces-associated beta-lactamase AAC(3)-VIIa (40) was more abundant in Bonney water whereas *vanYA* was more abundant in Fryxell water.

Overall, the three sample classes each had distinctive ARG-abundance signatures. Ten of the top 20 ranked ARGs were associated with Proteobacteria, which roughly corresponds with the proportion of bacterial taxa abundances predicted by Kraken2/Bracken (**Fig. 2**).

Enrichment of Mat-forming *Rhodoferax antarcticus*

The relatively high proportions of Betaproteobacteria and Alphaproteobacteria in MAT-04 differed from that of other samples, and the color of the sample material was more red than green. We cultured the various samples in Harwood PM mineral salts medium with 10 mM succinate (41, 42), anaerobically (in closed Pyrex tubes) at 10°C with illumination (approximately 10% PAR). From MAT-04 red spots of biofilm were obtained. Approximately 0.05g portions were twice subcultured for two weeks. The organism grew entirely as a mat, forming large red blobs with a “red nose” appearance. In addition the culture extended as a red film-like mat growing along the inside of the culture tube (**Fig. 5A**). There was no evidence of

planktonic growth (cloudiness) or cell motility within the culture medium. It was not possible to obtain isolated colonies, but serial culture of the "red noses" showed consistent biofilm formation and Gram-stain morphology (**Fig. 5B**). This finding is consistent with an earlier report that the Fryxell sedimentary biofilm includes a layer of phototrophic purple bacteria beneath the cyanobacterial layer (43). Ours is the first report of such biofilm cultured from material at the ice surface.

The MAT-04 biofilm culture was subjected to genome analysis. DNA was sequenced and short reads were used to BLAST the NCBI database. The predominant hits were to the genome of *Rhodoferax antarcticus* ANT.BRT, a Betaproteobacterium originally isolated from a saltwater pond at Cape Royds, Ross Island, Antarctica (44, 45). Kraken 2/Bracken analysis of sequenced reads agreed with this classification, assigning the samples as 95% *R. antarcticus* with about 1% *Pseudomonas.*

We used the *breseq* pipeline (46) to assemble reads from two samples (7-RN1 and 8-RN2) whose DNA sequence reads were assembled to the *R. antarcticus* ANT.BR reference genome (Supplementary file, **Table S2** and **S3**). These tables present "mutations," that is, all differences from the ANT.BR reference sequence that show 100% read coverage. The two samples, 7-RN1 and 8-RN2, showed 490 and 473 sequence differences respectively. A large part of the observed sequence differences from reference were silent mutations, the same ones in both samples. These observations confirm the close relatedness of our culture to the published strain.

The *R. antarcticus* enrichment culture (designated *R. antarcticus* JLS) was tested for ARG abundance using ShortBRED (**Table 2**). More than thirty ARGs showed hits, but nearly all were associated with *Pseudomonas aeruginosa*. By contrast, no ARGs matched those from

CARD database that were associated with *Rhodoferax.* This finding suggests that a wide range of naturally-occurring ARGs remain to be discovered in polar strains.

Characterization of the mat-forming *R. antarcticus* JLS

Despite the high sequence identity, the *R. antarcticus* JLS enrichment culture differed from the original reported *R. antarcticus* ANT.BR (44) in important respects. Unlike the highly motile *R. antarcticus* ANT.BR, our culture showed no motility and little sign of planktonic single-celled growth. Instead, the subcultured material grew entirely as a biofilm, in globular spots and as a film-like growth along the interior surface of the glass tube.

The *R. antarcticus* JLS biofilm was further tested for growth range of pH and salinity. The biofilm was subcultured anaerobically with illumination, in a malate-succinate *Rhodoferax* medium modified from that of references (44, 47) as described under Methods. The pH was maintained by inclusion of 100 mM of sulfonate buffers with pKa appropriate for each pH value: pH 6, pH 7, and pH 8, all adjusted with KOH (**Fig. 6A, B, C**). After 45 days, at pH 7, the bacteria formed globules as well as a red coating along the glass. At pH 6, spots of biofilm grew slowly, and at pH 8 little growth was seen. Growth was also tested for cultures buffered at pH 7 with NaCl amendment (**Fig. 6D, E, F**). The fullest growth was seen in the absence of NaCl amendment (the core medium contains approximately 10 mM Na^+ ions). Less growth was seen with added NaCl (23 mM or 46 mM).

DISCUSSION

The mat lift-off samples we obtained from the ice surface showed a range of DNA taxa consistent with those reported for samples obtained from benthic mats (5). Given that our collected mat organisms had survived five years within ice, followed by air dessication and prolonged UV radiation exposure, it is impressive how much of the Antarctic microbes retain viability with intact DNA. Even ciliated protists and invertebrate worms were obtained alive from samples cultured after months of storage at -80°C (not shown). It is likely that some of these organisms are psychrophiles that continue to grow within the ice (26).

Previous studies emphasize the cyanobacterial content of lift-off mats, primarily *Microcoleus* and *Nostoc* (20, 22). While most of our lift-off samples showed abundance of cyanobacteria, one sample yielded cultures of 99% *R. antarcticus.* This indicates that portions of the lower layer of *Rhodoferax* mat (43), along with the cyanobacterial upper layer, can break off and form lift-off patches that emerge from the ice. Furthermore, our *R. antarcticus* JLS obtained from Lake Fryxell showed mat-forming morphology very different from the motile single cells of *R. antarcticus* ANT.BR isolated from Cape Royds.

The prevalence of eukaryotic sequences in the lake water metagenomes is consistent with previous reports that protists play important roles in the Taylor Valley lake communities (18, 23). Protists can regulate bacterial ARG composition in terrestrial communities (48), so this factor may be of interest when assessing the lake water ARG pools.

Identification of antibiotic resistance genes from relatively pristine environments is of interest for several reasons. Long before human introduction of high-dose antibiotics, environmental bacteria evolved multidrug pumps to efflux toxic products of their own metabolism, as well as antimicrobial substances produced by their competitors (13, 49). Many

antibiotics possess signalling capabilities and other unknown functions. Phylogeny dates the origin of beta-lactamases to hundreds of millions of years ago. Vancomycin resistance genes are found in 30,000-year-old permafrost (50). But the specific kinds of ARGs found in environmental sources may differ from those prevalent in human microbiomes, those conferring resistance to the drugs we depend on for therapy (51).

In polar regions, previous metagenomic studies reveal a range of naturally occurring ARGs. Surface soils of the Antarctic Mackay Glacier region show ARGs encoding multidrug pumps, beta-lactamases and aminoglycoside inactivators, largely associated with Gram-negative bacteria (52). A study of Tibetan soils showed a number of abundant ARGs, most notably encoding vancomycin resistance (53). Our study of ARGs from Antarctic lakes adds to this picture, showing that both water and microbial mat sources contain familiar ARGs, and likely contain others not yet discovered in Antarctic strains.

METHODS

Sample collection and culture.

Microbial communities were sampled in December, 2014 from two meromictic lakes of the Taylor Valley, Victoria Land, Antarctica. Microbial lift-off mat was collected from Lake Fryxell ice surface, within the GPS area of: 77.60491, 163.16315; 77.60473, 163.16290; 77.60463, 163.16405; 77.60495, 163.16495. Each sample consisted of a separate tuft of dessicated microbial mat, collected with alcohol-sterilized forceps and stored at 20°C (4 weeks) then at -80°C (indefinitely).

Lake water was sampled from the permanent chemoclines of Lake Fryxell (77.605 S, 163.163 E, 9 m depth) and Lake Bonney, east lobe (77.719 S, 162.283 E, 15 m depth). Water

samples were collected in 1-liter cubitainers pre-washed in 10% HCl. Each water sample was concentrated by filtration through 0.45 µm filter.

Enrichment culture was performed using Harwood PM medium supplemented with 10 mM succinate (41, 54). Screwcap pyrex tubes were filled with medium and inoculated with approximately 0.05 g dessicated material from sample Mat-04. Tubes were incubated under approximately 10% PAR at 10°C for five weeks. Portions of biofilm were serially subcultured for two-week periods, then frozen at -80°C.

Testing for growth range of pH and NaCl amendment was performed using a *Rhodoferax* medium modified from references (44, 47). The medium contained: yeast extract (0.5 g/l), EDTA (20 mg/l), malic acid (4 g/l), $(NH_4)_2SO_4$ (1 g/l), $MgSO_4 \cdot 7H_2O$ (200 mg/l), $FeSO_4 \cdot 7\ H_2O$ (12 mg/l), K_2HPO_4 (0.9 g/l), KH_2PO_4 (0.6 g/l), 10 mM sodium succinate, 100 mM MOPS (3-(N-morpholino)propanesulfonic acid) adjusted to pH 7 with KOH. In addition, 1ml/l of a trace elements solution was added [H_3BO_3 (2.8 g/l), $MnSO_4 \cdot H_2O$ (1.6 g/l), $Na_2MoO_4 \cdot 2H_2O$ (0.76 g/l), $ZnSO_4 \cdot 7\ H_2O$ (240 mg/l), $Cu(NO_3)_2 \cdot 3\ H_2O$ (40 mg/l), $CoCl_2 \cdot 6\ H_2O$ (200 mg/l)]. For pH 6, the MOPS buffer was replaced with 100 mM MES (2-(*N*-morpholino)ethanesulfonic acid) and for pH 8, the buffer was 100 mM TAPS ([tris(hydroxymethyl)methylamino]propanesulfonic acid).

DNA isolation and sequencing.

Mat DNA was extracted by the PowerBiofilm kit (MO BIO). From lake water filters, DNA was extracted using an MP FastDNA SPIN DNA kit (MP Biomedicals, CA) following the manufacturer's instructions (23). Shotgun metagenomic library construction and sequencing were carried out at DOE Joint Genome Institute (JGI) using standard protocols on the Illumina

HiSeq 2500 platform. The raw reads were quality filtered using JGI standard protocols and 1.2 Tb total sequences were obtained.

Cultured biofilm DNA was isolated by the PowerBiofilm kit (MO BIO). Sequencing was performed by the Michigan State University Genomics Core. Libraries were prepared using the Illumina TruSeq Nano DNA Library Preparation Kit. Sequencing was performed on an Illumina MiSeq done in a 2x250bp format using an Illumina 500 cycle v2 reagent cartridge. The raw reads were quality filtered and 1 Tb total sequences were obtained.

Identification of ARGs by ShortBRED.

ShortBRED (55) is a pipeline that identifies protein family sequences by generating specific peptide markers from a database of interest, then screening the markers for specificity against the UniProt universal database of protein sequences (56). The shortbred_identify command scans a reference database for protein family-specific peptide sequences. It was used to scan UniRef (the UniProt protein reference database) for sequences of the ARGs in version 3.0.7 of the Comprehensive Antibiotic Resistance Database (CARD) (28). The markers were then used by the shortbred_quantify command to assign ARGs in the 150-kb reads from lake water and mat samples.

Trimmomatic.

The Trimmomatic pipeline (57) removes adapters and low-quality sequence from Illumina reads. It counts the number of reads per FASTQ file, and was used to determine the total reads per sample metagenome.

Taxa classification by Kraken 2/Bracken.

The 150-kb sequence reads from mat and water metagenomes were assigned to taxonomic classes using the Kraken 2 classifier pipeline (30, 31). Kraken associates genomic *k*-mers (short sequence strings) in the reads with lowest common ancestor taxa. The read classifications were then used to calculate taxa percentages via the Bracken abundance estimator (32). The *k*-mer length was set at 150. For the Kraken 2-Bracken pipeline, the reference database used was the 5/17/2021 Standard Collection accessed at: https://benlangmead.github.io/aws-indexes/k2.

Taxa profiling by MetaPhlAn2.

Mat and water metagenomes were mapped to core taxonomic markers using the MetaPhlAn2 pipeline (33, 34). MetaPhlAn2 assigns short reads to taxa using a set of marker genes identified from approximately 17,000 microbial reference genomes.

Genome assembly by *breseq*.

The pipeline *breseq* 0.35.6 (46) was used to assemble reads from two samples of the enrichment culture obtained from Mat-04. The reads were mapped to the *Rhodoferax antarcticus* ANT.BRT reference genome (45).

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Author Contributions Statement

ST designed the bioinformatic analysis of ARGs and drafted the manuscript. GH characterized the *Rhodoferax* enrichment culture and contributed to the manuscript. DB ran the bioinformatic pipelines and performed statistics. JLS conceived the central concept, collected the mat samples and isolated DNA, mentored students, and completed the manuscript.

Conflict of Interest Statement

The authors declare that the submitted work was carried out with no personal, professional or financial relationships that could potentially be construed as a conflict of interest.

Contribution to the Field Statement

Our study is the first to report metagenomic analysis of antibiotic resistance genes from the water column and from ice-surface lift-off mats of the Taylor Valley lakes, a major study site for Antarctic microbiology. We are the first to report a mat-forming enrichment culture of *Rhodoferax* species from an Antarctic lift-off mat.

Availability of data and materials

Sequence read files from lake water and mat samples are deposited at NCBI under SRA accession numbers SRP104818, SRP104821, SRP098041, SRP098040, SRP098042,

SRP104817, SRP098044, SRP104822, SRP098050, SRP104819, SRP104820, SRP104823. Sequence read files from cultured *Rhodoferax antarcticus* are deposited at NCBI under SRA accession number PRJNA736311.

TABLES

Table 1. Mat and water antibiotic resistance genes (ARGs) identified by ShortBRED marker peptides*

ARG Family	MAT-01	MAT-02	MAT-03	MAT-04	MAT-05	MAT-06	FRY-01	FRY-02	FRY-03	BON-01	BON-02	BON-03	TOTAL	P
ARO_3004294|BUT-1 *Buttiauxella agrestis*	0	1	2	0	0	0	554	1720	967	266	262	323	4095	0.027
ARO_3002541|AAC(3)-VIIa *Streptomyces rimosus*	89	235	114	173	152	54	44	31	31	323	313	231	1790	0.038
ARO_3002528|AAC(3)-Ia *Pseudomonas aeruginosa*	302	189	131	492	238	196	0	0	0	0	0	0	1548	0.022
ARO_3003167|CTX-M-159 Proteobacteria	36	9	25	1032	111	18	1	0	0	0	0	3	1235	0.052
ARO_3001461|OXA-156 *Pandoraea pulmonicola*	43	18	75	146	70	36	14	15	19	70	60	53	619	0.099
ARO_3002894|otrC *Streptomyces rimosus*	40	122	16	27	13	23	162	73	81	11	20	10	598	0.079
ARO_3003599|OXA-443 *Ralstonia mannitolilytica*	41	260	13	42	33	16	64	23	12	2	3	4	513	0.061
ARO_3001338|SHV-100 *Klebsiella pneumoniae*	58	1	49	144	56	44	4	1	5	10	6	11	389	0.241
ARO_3002955|vanYA *Enterococcus faecium*	0	3	2	1	1	0	101	59	47	19	29	31	293	0.027
ARO_3001855|ACT-35 *Enterobacter cloacae*	4	7	26	32	37	5	30	22	22	40	22	30	277	0.214
ARO_3000410|sul1 Proteobacteria	72	0	32	19	34	62	1	0	0	0	0	0	220	0.199
ARO_3003022|dfrB3 *Klebsiella oxytoca*	13	4	9	19	23	8	54	21	18	9	12	18	208	0.069
ARO_3002972|vanTG *Enterococcus faecalis*	1	0	1	0	2	0	10	9	13	50	53	55	194	0.027
ARO_3001517|OXA-329 *Acinetobacter calcoaceticus*	2	2	1	2	0	1	76	52	24	1	15	5	181	0.054
ARO_3002701|cmr *Rhodococcus fascians*	6	5	17	22	19	2	39	47	16	3	2	2	180	0.038
ARO_3003723|van *Desulfitobacterium hafniense*	0	0	0	0	0	0	53	50	40	9	16	10	178	0.024
ARO_3003942|abcA *Staphylococcus aureus*	31	75	2	9	17	20	0	0	0	7	4	5	170	0.055
ARO_3003583|basS *Pseudomonas aeruginosa*	12	1	30	8	7	12	1	0	0	47	32	19	169	0.046
ARO_3004652|Erm(O)-lrm *Streptomyces lividans*	28	8	16	37	35	10	8	7	4	5	2	5	165	0.061
ARO_3001299|tlrB_tylosin *Streptomyces fradiae*	18	8	10	30	26	8	2	2	0	18	18	22	162	0.034

*Class differences for ARG abundance were tested for significance ($P \leq 0.5$) using the Kruskal-Wallis test.

Table 2. *Rhodoferax* enrichment culture ARGs identified by ShortBRED*

A. *Rhodoferax* (95%), *Pseudomonas* (1.5%)	Hits	Marker Length	B. *Rhodoferax* (95%), *Pseudomonas* (0.7%)	Hits	Marker Length
ARO_3003679|TriA__Pseudomonas_aeruginosa_PAO1	8	242	ARO_3003680|TriB__Pseudomonas_aeruginosa_PAO1	3	144
ARO_3001796|OXA-50__Pseudomonas_aeruginosa	4	116	ARO_3003031|mexW__Pseudomonas_aeruginosa_PAO1	1	56
ARO_3003681|TriC__Pseudomonas_aeruginosa_PAO1	3	113	ARO_3002320|KPC-10__Acinetobacter_baumannii	1	66
ARO_3003710|mexL__Pseudomonas_aeruginosa_PAO1	3	134	ARO_3003583|basS__Pseudomonas_aeruginosa	1	290
ARO_3003030|mexV__Pseudomonas_aeruginosa_PAO1	3	176	ARO_3004777|CMH-1__Enterobacter_cloacae	1	25
ARO_3002507|PDC-8__Pseudomonas_aeruginosa	3	155	ARO_3003705|mexN__Pseudomonas_aeruginosa	1	63
ARO_3000802|OprJ__Pseudomonas_aeruginosa	2	36	ARO_3000808|mexI__Pseudomonas_aeruginosa_PAO1	1	118
ARO_3003680|TriB__Pseudomonas_aeruginosa_PAO1	2	144	ARO_3004054|CpxR__Pseudomonas_aeruginosa	1	26
ARO_3000379|OprM__Pseudomonas_aeruginosa_PAO1	2	35			
ARO_3000805|OprN__Pseudomonas_aeruginosa_PAO1	2	102			
ARO_3002985|arnA__Pseudomonas_aeruginosa_PAO1	2	102			
ARO_3003692|mexJ__Pseudomonas_aeruginosa_PAO1	2	120			
ARO_3003031|mexW__Pseudomonas_aeruginosa_PAO1	2	56			
ARO_3004038|emrE__Pseudomonas_aeruginosa_PAO1	2	51			
ARO_3003705|mexN__Pseudomonas_aeruginosa	2	63			
ARO_3003698|mexP__Pseudomonas_aeruginosa	2	85			
ARO_3002645|APH(3)-IIb__Pseudomonas_aeruginosa	2	153			
ARO_3000377|MexA__Pseudomonas_aeruginosa_PAO1	2	106			
ARO_3001214|mdtM__Escherichia_coli_str__K-12	1	71			
ARO_3004077|PmpM__Pseudomonas_aeruginosa_PAO1	1	109			
ARO_3000804|MexF__Pseudomonas_aeruginosa_PAO1	1	102			
ARO_3004072|OpmB__Pseudomonas_aeruginosa_PAO1	1	111			
ARO_3004075|MuxC__Pseudomonas_aeruginosa_PAO1	1	177			
ARO_3003693|mexK__Pseudomonas_aeruginosa_PAO1	1	44			
ARO_3000809|opmD__Pseudomonas_aeruginosa_PAO1	1	116			
ARO_3003682|OpmH__Pseudomonas_aeruginosa_PAO1	1	79			
ARO_3002695|cmlA5__uncultured_bacterium	1	99			
ARO_3003583|basS__Pseudomonas_aeruginosa	1	290			
ARO_3000795|mdtE__Escherichia_coli_str__K-12	1	54			
ARO_3004612|ampH__Escherichia_coli_DH1	1	68			

*Enrichment cultures A and B, taxa proportions were estimated by Kraken2/Bracken pipeline

FIGURES

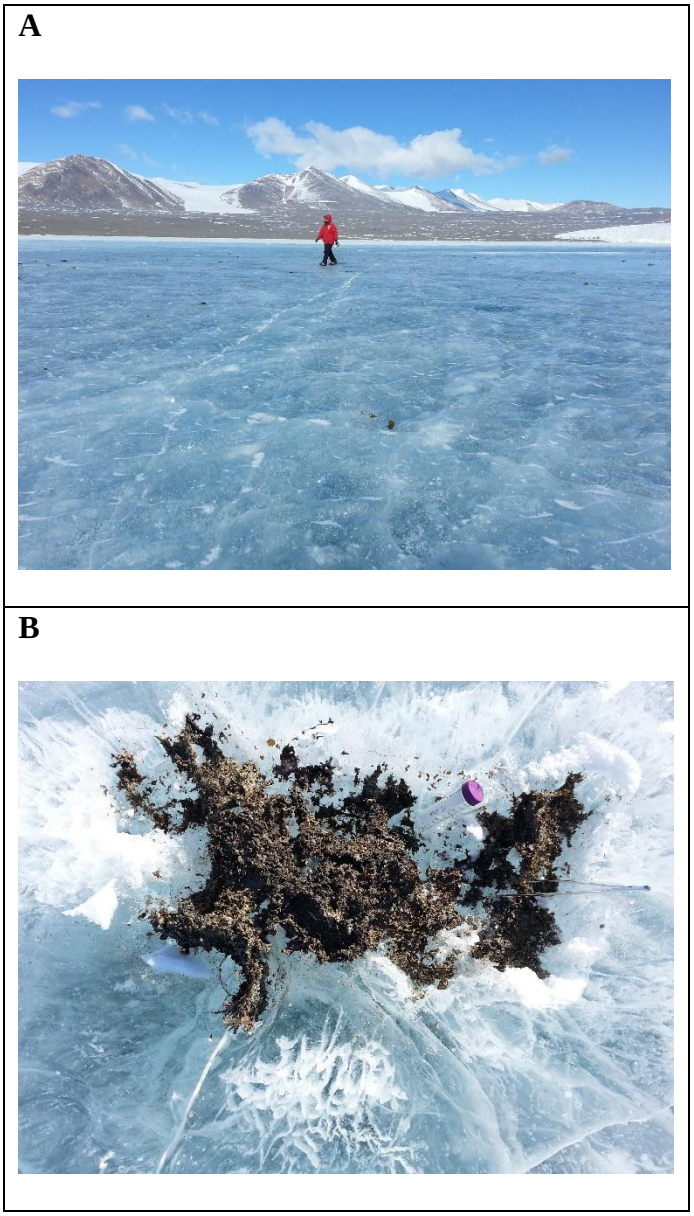

Figure 1. Lake Fryxell uplift mats.

(**A)** Lake Fryxell with permanent ice cover, weathered by katabatic winds; December 10, 2014.

(**B)** Source of MAT-4 DNA. Uplift mat emerges from the ice; approximately 40 cm across.

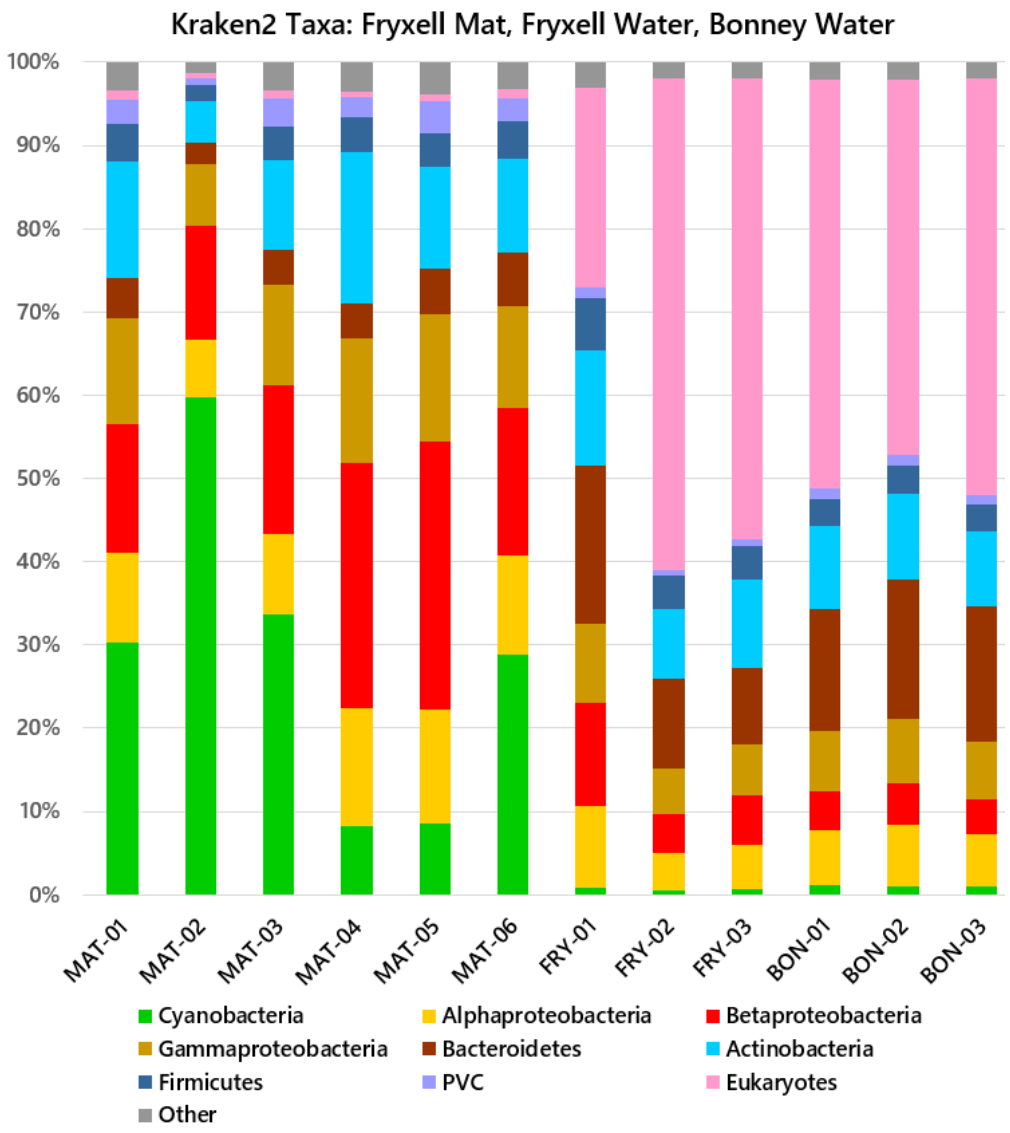

Figure 2. Microbial community composition of mat and water samples. Relative abundance of phyla was determined by read alignment to marker genes using Kraken2/Bracken pipeline. MAT (Lake Fryxell, mat samples); FRY (Lake Fryxell, water samples); BON (Lake Bonney, water samples). The water samples were collected on a 0.45-µm filter. MAT samples were from ice surface, unfiltered. PVC=Planctomycetes-Verrucomicrobia-Chlamydiae

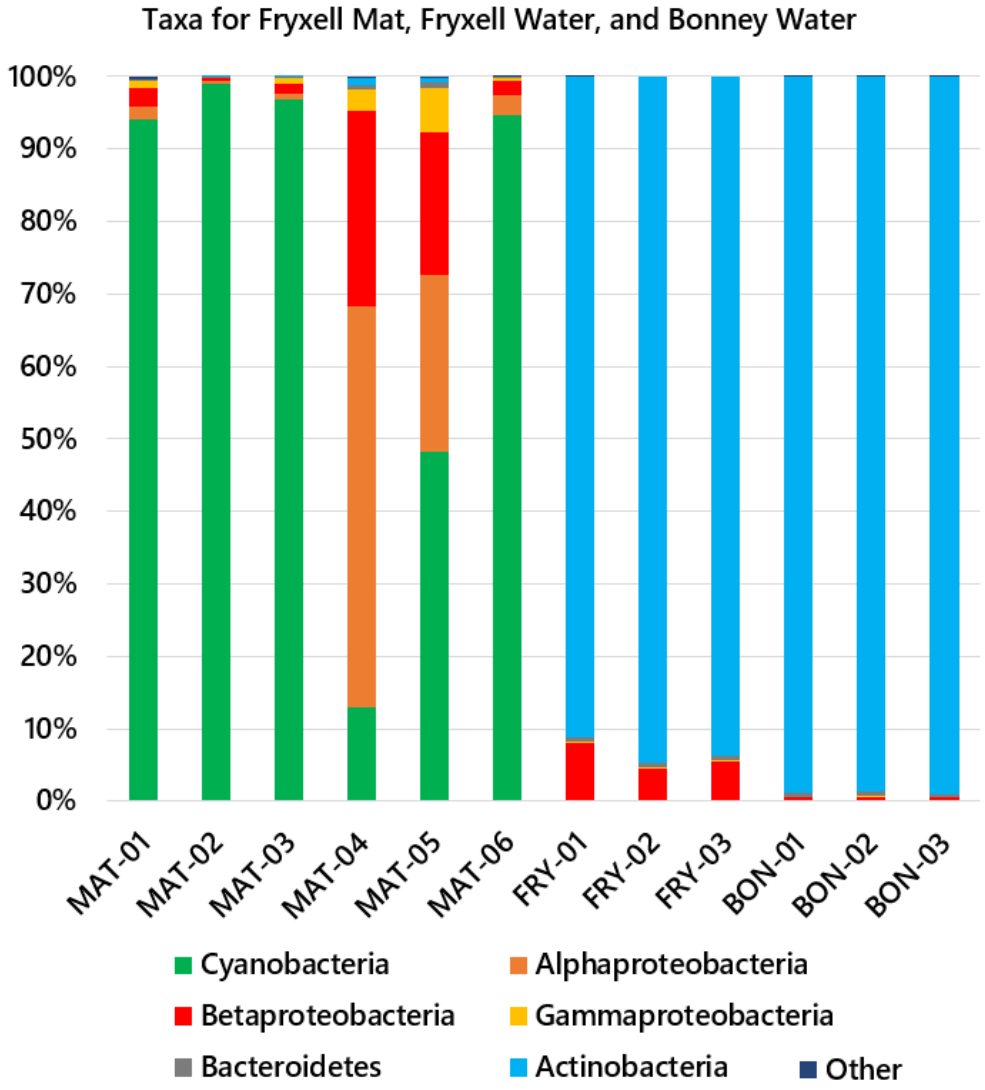

Figure 3. Microbial community composition via MetaPhlAn2. Relative abundance of class-level taxa was determined by read alignment to marker genes using MetaPhlAn2 (33, 34). Samples were as indicated for Figure 2.

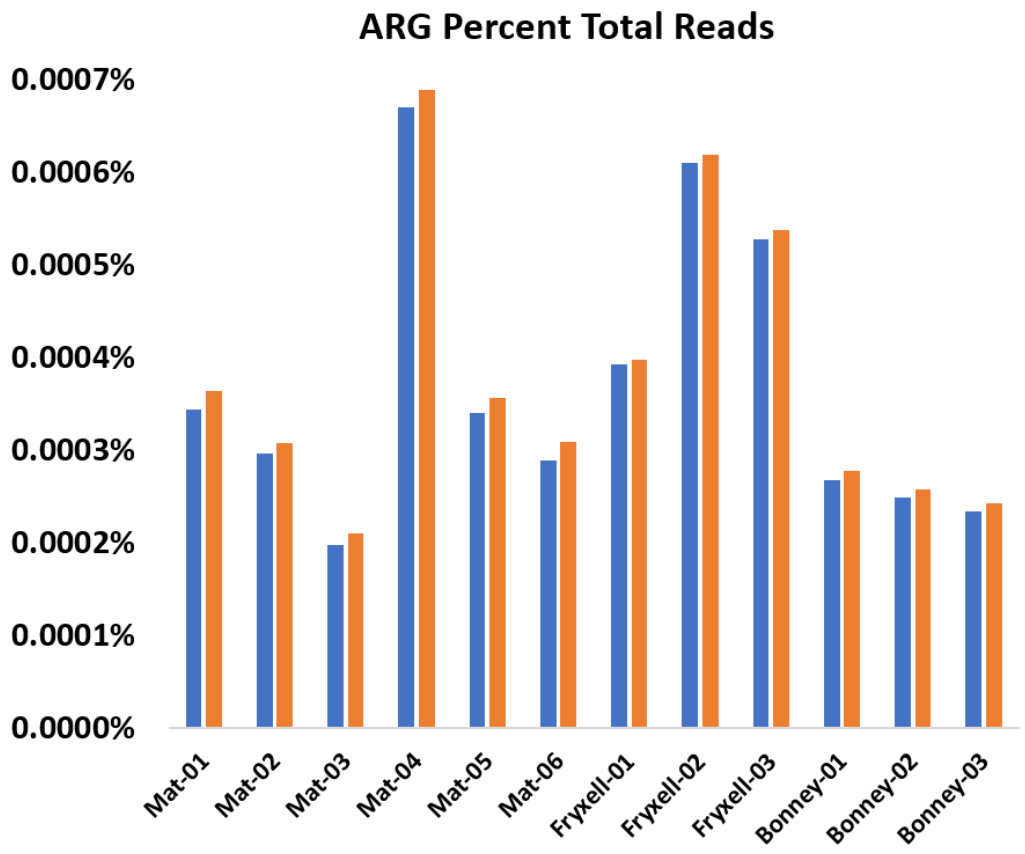

Figure 4. ARG abundance in lake samples. Percentage of reads matched to ShortBRED markers. Total reads were counted using Trimmomatic. Bars indicate total ARG hits per sample (blue) and top 60 ranked ARG hits (orange).

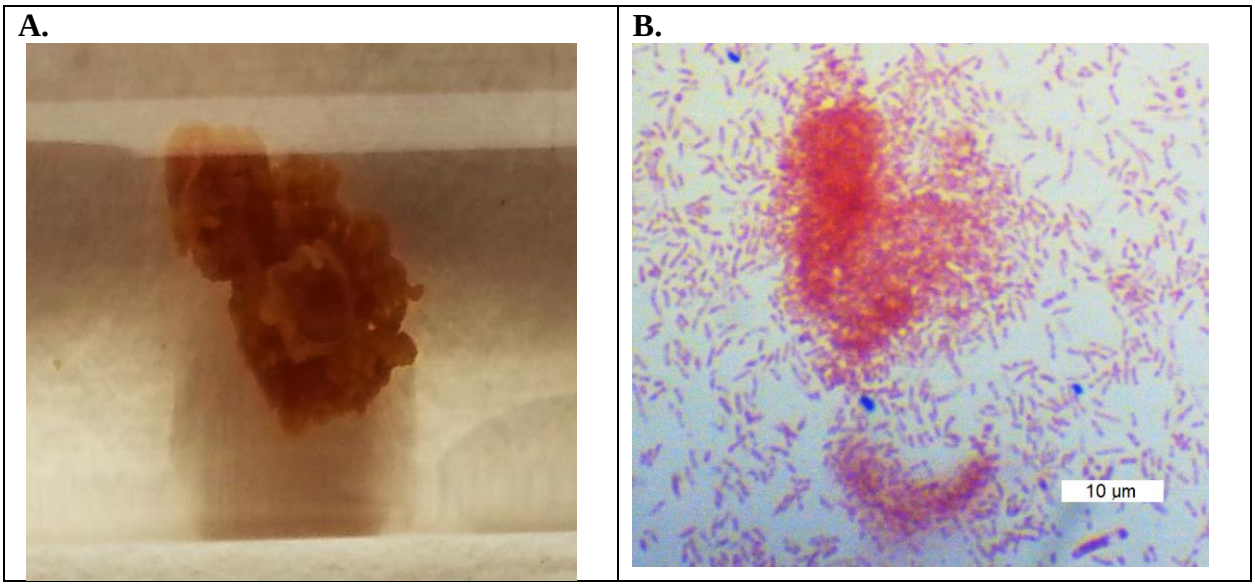

Figure 5. Mat-forming enrichment of Mat-04 *R. antarcticus* JLS.

(A) Biofilm of cultured *R. antarcticus* within closed tube of medium, showing globular form of growth. Initial Mat-04 sample was subcultured serially three times in PM medium, in horizontal closed tubes at 10°C with 30 umoles/m^2s illumination.

(B) Gram stain of culture, 1000X with oil immersion. Courtesy of Emma Stuart-Bates.

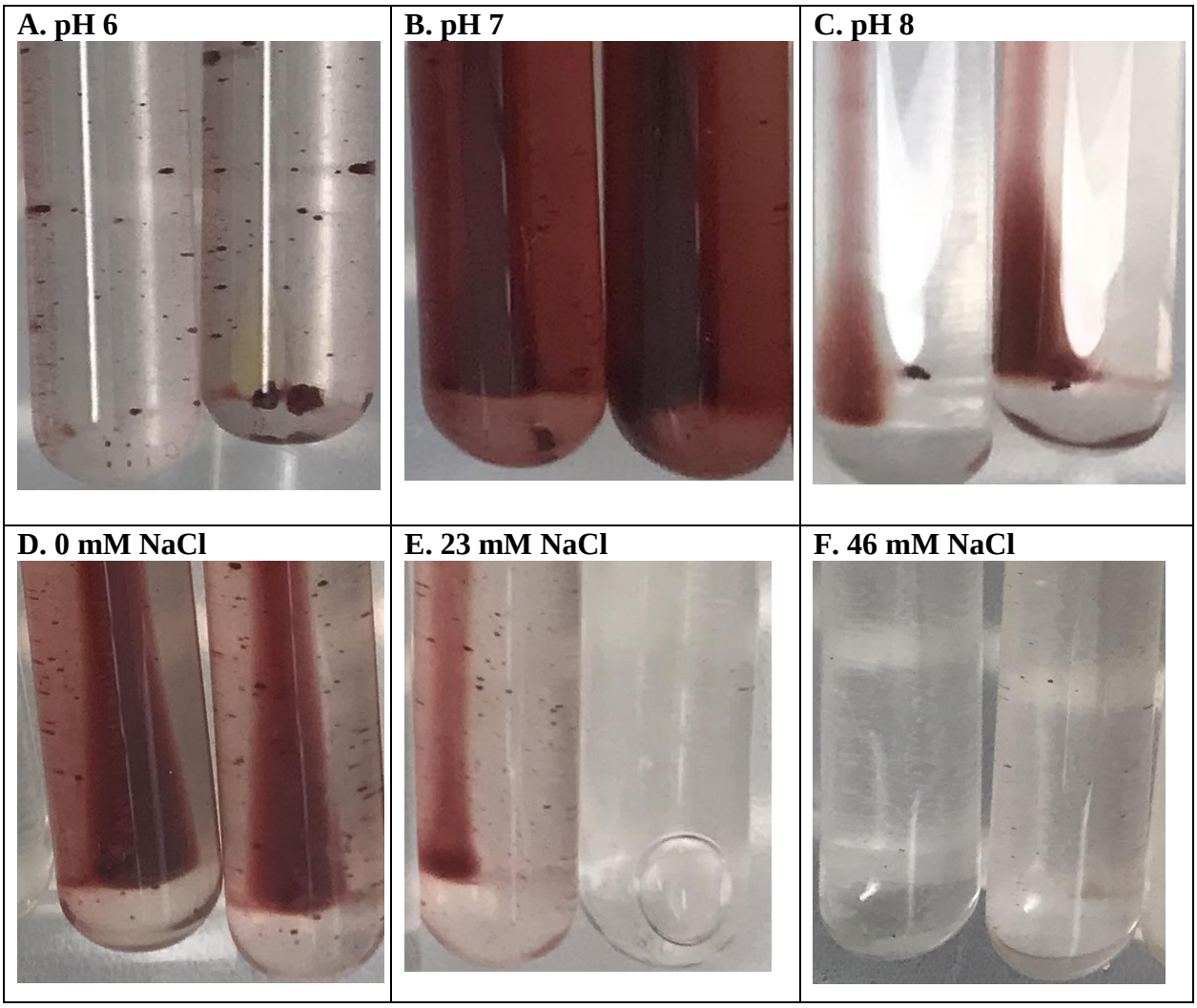

Figure 6. pH and NaCl dependence of *R. antarcticus* JLS.

(A, B, C) Medium PM contained **(A)** 100 mM MES pH 6; **(B)** 100 mM MOPS pH 7; **(C)** 100 mM TAPS pH 8. Cultures were incubated 45 days.

(**D, E, F)** Medium PM was amended with **(D)** no NaCl added; **(E)** 23 mM NaCl; **(F)** 46 mM NaCl. Cultures were incubated 45 days.

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