A Novel Enrichment Culture Highlights Core Features of Microbial Networks Contributing to Autotrophic Fe(II) Oxidation Coupled to Nitrate Reduction

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Keywords
Fe(II) oxidation · Denitrification · Gallionellaceae · Noviherbaspirillum · Meta’omics

Abstract
Fe(II) oxidation coupled to nitrate reduction (NRFO) has been described for many environments. Yet very few autotrophic microorganisms catalysing NRFO have been cultivated and their diversity, as well as their mechanisms for NRFO in situ remain unclear. A novel autotrophic NRFO enrichment culture, named culture BP, was obtained from freshwater sediment. After more than 20 transfers, culture BP oxidized 8.22 mM of Fe(II) and reduced 2.42 mM of nitrate within 6.5 days under autotrophic conditions. We applied metagenomic, metatranscriptomic, and metaproteomic analyses to culture BP to identify the microorganisms involved in autotrophic NRFO and to unravel their metabolism. Overall, twelve metagenome-assembled genomes (MAGs) were constructed, including a dominant Gallionellaceae sp. MAG (>71% relative abundance). Genes and transcripts associated with potential Fe(II) oxidizers in culture BP, identified as a Gallionellaceae sp., Noviherbaspirillum sp., and Thiobacillus sp., were likely involved in metal oxidation (e.g., cyc2, mtoA), denitrification (e.g., nirK/S, norBC), carbon fixation (e.g., rbcL), and oxidative phosphorylation. The putative Fe(II)-oxidizing protein Cyc2 was detected for the Gallionellaceae sp. Overall, a complex network of microbial interactions among several Fe(II) oxidizers and denitrifiers was deciphered in culture BP that might resemble NRFO mechanisms in situ. Furthermore, 16S rRNA gene amplicon sequencing from environmental samples revealed 36 distinct Gallionellaceae taxa, including the key player of NRFO from culture BP (approx. 0.13% relative abundance in situ). Since several of these in situ-detected Gallionellaceae taxa were closely related to the key player in culture BP, this suggests that the diversity of organisms contributing to NRFO might be higher than currently known.

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Published by S. Karger AG, Basel

Introduction
Oxidation of ferrous iron [Fe(II)] and reduction of ferric iron [Fe(III)] contribute to biogeochemical carbon, nitrogen, oxygen and sulphur cycles and can be mediated by specific microorganisms. There are three physiological types of Fe(II)-oxidizing bacteria (FeOB), namely, micro-
aerophiles, photoferrotrophs, and nitrate-reducing Fe(II) oxidizers. Autotrophic nitrate-reducing Fe(II) oxidizers have the ability to fix carbon dioxide (CO₂) and to oxidize Fe(II) coupled with complete or incomplete denitrification and can, thus, influence the fate of the groundwater pollutant nitrate and the greenhouse gases nitrous oxide (N₂O) and carbon dioxide (CO₂). However, detailed knowledge about the enzymatic mechanisms of autotrophic Fe(II) oxidation coupled to nitrate reduction (NRFO) in the environment is lacking.

Based on several laboratory studies on autotrophic and mixotrophic nitrate-reducing Fe(II)-oxidizing bacteria [ZoBell and Upham, 1944; Rabus and Widdel, 1995; Straub et al., 1996; Buchholz-Cleven et al., 1997; Edwards et al., 2003; Beller et al., 2006; Kumaraswamy et al., 2006; Weber et al., 2006; Su et al., 2015; Zhang et al., 2015; Zhou et al., 2016], it was proposed that true autotrophic nitrate-reducing Fe(II)-oxidizing cultures: (i) have no need of an additional organic carbon source, (ii) maintain Fe(II) oxidation over several (>3) transfers without organic carbon addition, (iii) confirm the growth of cells with only Fe(II), nitrate, and CO₂ provided, and (iv) show CO₂ uptake by the incorporation of labelled CO₂ into biomass during Fe(II) oxidation [Bryce et al., 2018]. To date, there has only been one example of a stable autotrophic nitrate-reducing Fe(II)-oxidizing bacterial culture that fits all of the above criteria: enrichment culture KS [Straub et al., 1996; He et al., 2016; Tominski et al., 2018a, b]. Culture KS was initially described by Straub et al. [1996], originates from Bremen, Germany, and is composed of FeOB, i.e., an unclassified Gallionellaceae sp., and denitrifiers, i.e., Rhodanobacter sp. and Bradyrhizobium sp. [He et al., 2016]. These FeOB and denitrifiers in culture KS showed that species interdependencies are used as a survival strategy under the carbon-limited but Fe(II)- and nitrate-rich culture conditions [Huang et al., 2021]. Yet, it remains unknown whether additional autotrophic nitrate-reducing Fe(II)-oxidizing cultures, aside from culture KS, can be cultivated to expand the diversity of known key players of autotrophic NRFO.

Nitrate-reducing Fe(II)-oxidizing bacteria were found in various types of environments [Straub and Buchholz-Cleven, 1998; Laufer et al., 2016] and were studied as treatments for contaminated groundwater [Su et al., 2016; Zhang et al., 2016; Kiskira et al., 2017] and constructed wetlands (i.e., wastewater treatment systems) [Song et al., 2016]. Furthermore, microorganisms of the family Gallionellaceae were widely found in a variety of natural environments [Wang et al., 2009; McBeth et al., 2013; Emerson et al., 2016; Fabisch et al., 2016; Jewell et al., 2016; Li et al., 2017]. In addition, several observations pointed towards an involvement of Gallionellaceae spp. in NRFO processes at different habitats [Emerson et al., 2016; Jewell et al., 2016; Bethencourt et al., 2020]. However, it remained largely unknown whether these FeOB are dependent on other microbially enriched culture members, similar to the interdependencies described for culture KS [He et al., 2016; Huang et al., 2021].

Here, we explored a novel autotrophic NRFO enrichment culture, named culture BP (Bremen Pond), that originated from freshwater sediment in Bremen, Germany, and was obtained in 2015 from a pond close to the sampling site of culture KS. We studied the physiology of culture BP by analysing its geochemistry, cell numbers, and microbial community composition (via 16S rRNA gene amplicon sequencing). In addition, we used metagenomic, metatranscriptomic, and metaproteomic (meta’omics) analysis to: (i) identify the key players of NRFO in culture BP, (ii) study their potential metabolism, particularly for Fe(II) oxidation, denitrification, and carbon fixation, (iii) explore possible microbial interspecies interactions in culture BP, and (iv) compare the more recent culture BP that might resemble environmental NRFO processes more closely to the long-term established culture KS. We furthermore performed DNA- and RNA-based 16S rRNA (gene) amplicon sequencing of six sites at the original habitat of culture BP to (v) assess the in situ diversity and relative abundance of potential key players involved in NRFO at the field site.

**Results**

**Physiology and Microbial Community Composition of the Novel Enrichment Culture BP**

To explore the diversity of cultivable autotrophic FeOB contributing to NRFO in the environment, we sampled a freshwater habitat, a pond in Bremen (Germany), which is in the vicinity of the origin of the existing NRFO culture KS (a connected ditch nearby). We established the novel culture BP, maintained under the same autotrophic growth conditions as culture KS [Tominski et al., 2018a]. Culture BP was grown under autotrophic conditions (i.e., with 10 mM Fe(II) as an electron donor and 4 mM nitrate as an electron acceptor in bicarbonate-buffered medium) and under heterotrophic conditions (i.e., with 5 mM acetate as an electron donor and 4 mM nitrate as an electron acceptor) as alternative growth condition for three successive transfers in triplicates, respectively. Samples for meta’omics analyses were collected on
the 2nd day under autotrophic conditions, and at 18 h under heterotrophic conditions; optimum sampling points were determined based on geochemical and short-read 16S rRNA gene amplicon sequencing analyses conducted in parallel.

Under autotrophic conditions, 8.22 mM of Fe(II) was oxidized in 6.5 days with an average Fe(II) oxidation rate of 1.66 mM/day during the active growth phase (i.e., between 23 and 90 h) (Fig. 1a). This co-occurred with a reduction of 2.42 mM of nitrate and an average nitrate reduction rate of 0.53 mM/day, without detectable nitrite production (Fig. 1a). The average Fe(II)oxidized/nitrate-reduced stoichiometric ratio was 3.4, and cell numbers increased from 1.04 × 10^6 cells/mL to 4.31 × 10^6 cells/mL within 3.75 days (Fig. 1c). Under heterotrophic conditions, 3.4 mM of acetate was consumed with an average...
acetate oxidation rate of 2.7 mM/day during the exponential growth phase (i.e., between 11 and 37 h). This co-occurred with a reduction of 4.0 mM nitrate and an average nitrate reduction rate of 3.5 mM/day. Nitrite increased from 0.02 to 0.8 mM from 11 to 18 h, but decreased again afterwards (Fig. 1b). The average ratio for acetate_{oxidized}/nitrate_{reduced} was 0.85, which is higher than the theoretical stoichiometry value of 0.625, suggesting that acetate was oxidized not only for energy generation, but also used for biomass production and nitrate was eventually reduced completely to N\textsubscript{2}. Throughout the experiment, cell numbers increased from 3.89 × 10\textsuperscript{4} cells/mL to 3.68 × 10\textsuperscript{7} cells/mL (corresponding to approx. up to 3.5 × 10\textsuperscript{7} cells/mL).

Under autotrophic conditions, an amplicon sequencing variant (ASV) classified as *Gallionellaceae* sp. dominated with 71–78% relative abundance on days 1, 2, 4, and 7 (corresponding to approx. up to 3.1 × 10\textsuperscript{6} cells/mL; Fig. 1c). In addition to short-reads, long-read 16S rRNA gene amplicon sequencing was performed to obtain improved taxonomic identification of the microbial populations in culture BP (long-read and short-read 16S rRNA gene sequences are presented in the online suppl. data S1 and S2, respectively; for all online suppl. material, see www.karger.com/doi/10.1159/000517083). Based on long-reads, the novel unclassified *Gallionella*ceae sp. in culture BP shared 97.88% sequence identity with the unclassified *Gallionella*ceae sp. in culture KS and 96.17% with *Ferruginium kumadai* (online suppl. Table S1). Other abundant ASVs were classified as *Noviherbaspirillum* (9–15%; corresponding to approx. up to 5.3 × 10\textsuperscript{5} cells/mL), *Geothrix* (2–7%; approx. up to 1.9 × 10\textsuperscript{6} cells/mL), *Rhodoferax* (ASV-1 with 2–3%; approx. up to 7.5 × 10\textsuperscript{5} cells/mL), *Thiobacillus* (two ASVs with 0.4–1.6%; approx. up to 4.1 × 10\textsuperscript{4} cells/mL), and *Rhodoblastus* (1–5%; approx. up to 1.9 × 10\textsuperscript{5} cells/mL; Fig. 1c). Under heterotrophic growth conditions, a *Rhodoferax* sp. (ASV-1) dominated with 84–94% relative abundance (corresponding to approx. up to 3.5 × 10\textsuperscript{7} cells/mL) throughout the 3rd transfer under heterotrophic conditions, while lower relative abundances were detected for ASVs classified as *Noviherbaspirillum* (2–11%; approx. up to 8.6 × 10\textsuperscript{5} cells/mL), *Rhodoferax* (ASV-2, only 1 nucleotide difference to ASV-1; 2–3%; approx. up to 1.3 × 10\textsuperscript{6} cells/mL), *Gallionellaceae* (up to 0.1%; approx. up to 3.2 × 10\textsuperscript{4} cells/mL), and *Geothrix* (up to 0.03%; approx. up to 4.0 × 10\textsuperscript{3} cells/mL; Fig. 1d). The long-read 16S rRNA gene amplicon sequence of *Rhodoferax* sp. was closely related to the *Curvibacter delicatus* strain NBRC 14919 (98.56% sequence identity; online suppl. Table S1).

We identified highly similar 16S rRNA gene sequences of the *Rhodoblastus* sp. using short- and long-read 16S rRNA amplicon sequencing compared to the metagenome-assembled genome (MAG; see below). However, for the *Gallionellaceae* sp., *Noviherbaspirillum* sp., *Rhodoferax* sp., *Thiobacillus* sp., *Geothrix* sp. and *Betaproteobacteria* bacterium, only short- and long-read sequencing offered 16S rRNA gene sequences of the respective taxa. The phylogenetic relationships of the 16S rRNA gene sequences of the amplicon sequencing approach and the MAG is shown in the online supplementary Figure S1.

**General Characteristics of the MAGs in Culture BP**

In order to study the genomic potential of key players involved in NRFO and to use this information as reference for our met’omics approach, twelve MAGs were retrieved from the metagenome of culture BP (online suppl. data S3). Among these twelve MAGs, six were high quality with completeness >90% (*Gallionellaceae* sp., *Noviherbaspirillum* sp., *Rhodoferax* sp., *Rhodoblastus* sp., *Thiobacillus* sp. and *Betaproteobacteria* bacterium; online suppl. Table S1), two had medium quality with 88% and 64.8% completeness (*Ramlibacter* sp. and *Geothrix* sp., respectively; online suppl. Table S1), and four were low quality with <50% completeness (*Geothrix* sp., *Mesorhizobium* sp. and two unclassified MAGs), according to the definition of Bowers et al. [2017]. The four low-quality MAGs and one medium-quality MAG ( *Geothrix* sp.) did not contain Fe(II) oxidation genes and showed only partial genes for denitrification. Thus, these five MAGs were not explored further. Additionally, the classification of the *Betaproteobacteria* bacterium MAG was vague, and it did not express any denitrification genes in culture BP. Therefore, we mainly focused on the remaining six MAGs, i.e., the *Gallionellaceae* sp., *Noviherbaspirillum* sp., *Rhodoferax* sp., *Rhodoblastus* sp., *Thiobacillus* sp. and *Ramlibacter* sp., representing up to 98% of the microbial community during the Fe(II) oxidation phase. These MAGs harboured genes associated with key biogeochemical cycles, including carbon, nitrogen, oxygen, and iron, in particular Fe(II) oxidation (Fig. 2). In order to add more evidence to gene-level information, we conducted metatranscriptomic and metaproteomic analyses to gain additional evidence at RNA and protein levels. The detection of a protein confirmed its presence and therefore suggested that the catalysed reaction of this particular protein was likely happening. Statistical meta’omics data are presented in the online supplementary Table S2 and full detected transcript and protein data are shown in the online supplementary data S4.
**Fig. 2.** The relative and estimated absolute abundance (calculated using total cell numbers and short-read 16S rRNA gene amplicon sequencing relative abundance data) of related ASVs under autotrophic conditions accompanied by fold changes (log2) of key transcripts and proteins involved in Fe(II) oxidation, CO₂ fixation, denitrification, and potential oxygen respiration genes under autotrophic conditions compared to heterotrophic conditions. Genes with several copy numbers are listed multiple times. MT, metatranscriptomic analysis; MP, metaproteomic analysis. Stars indicate significant changes (adjusted \( p \leq 0.05 \)). A white “A” indicates the protein was only detected under autotrophic conditions. A white “H” indicates the protein was only detected under heterotrophic conditions. “#a” indicates two short-read *Rhodoferax* ASVs were combined, which had 100 and 99.6% sequence identity, respectively, to the long-read *Rhodoferax* sp. (only one nucleotide difference). “#b” indicates two short-read *Thiobacillus* ASVs were combined, which had a difference of three nucleotides to each other.

<table>
<thead>
<tr>
<th>Abundance of ASV</th>
<th>Gallionellaaceae</th>
<th>Novihelvotaespirillum</th>
<th><em>Rhodoferax</em> #a</th>
<th><em>Thiobacillus</em> #b</th>
<th><em>Rhodobacter</em></th>
<th><em>Ramlibacter</em></th>
</tr>
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<tbody>
<tr>
<td>Relative</td>
<td>78.0%</td>
<td>14.1%</td>
<td>2.2%</td>
<td>1.4%</td>
<td>1.1%</td>
<td>0.3%</td>
</tr>
<tr>
<td>Absolute (cells/ml)</td>
<td>2.2×10⁶</td>
<td>4.1×10⁶</td>
<td>6.2×10⁴</td>
<td>4.1×10⁴</td>
<td>3.1×10⁴</td>
<td>7.7×10³</td>
</tr>
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</table>
Fe(II) Oxidation

To explore the pathways of Fe(II) oxidation in culture BP, we searched for homologues of known and putative Fe(II) oxidation genes in the six MAGs, such as cyc2, mtoA, and mofA [Corstjens et al., 1997; Castelle et al., 2008; El Gherianny et al., 2009; Liu et al., 2012; He et al., 2017]. Specifically, the cyc2 gene was identified in the MAGs of the Gallionellaceae sp., Rhodoblastus sp., Thiobacillus sp., and Ramlibacter sp. At the transcript level, cyc2 was detected for the Gallionellaceae sp. and Thiobacillus sp. with significantly higher abundance (i.e., amount of detected transcript) under autotrophic conditions compared to heterotrophic conditions. In addition, the Cyc2 protein from the Gallionellaceae sp. was detected under autotrophic conditions (Fig. 2). To uncover the evolutionary relationship of Cyc2 between culture BP and other studied organisms, we performed phylogenetic analysis of Cyc2 amino acid sequences, including those of the Gallionellaceae sp., Rhodoblastus sp., and Ramlibacter sp. (online suppl. Fig. S2). The Cyc2 of the Gallionellaceae sp. in culture BP and of Sideroxydans lithotrophicus ES-1 were closely related to each other. Overall, there appeared to be distinct clusters within the Cyc2 tree, which consisted of Gallionellaceae spp. and Zetaproteobacteria spp. (related to Mariprofundus spp.), respectively (online suppl. Fig. S2). In addition, mtoAB genes were identified for the Noviherbaspirillum sp. MAG and the transcripts were at a significantly higher abundance (Fig. 2). Phylogenetic analysis of MtoAB from Noviherbaspirillum sp. demonstrated that they are closely related to the MtoAB from other proposed microaerophilic FeOB (online suppl. Fig. S3) [Liu et al., 2012]. Particularly for MtoB, there appeared to be a distinct cluster for Noviherbaspirillum (online suppl. Fig. S3). Furthermore, mofA genes were identified in the Thiobacillus sp. MAG and the transcripts were also of significantly higher abundance under autotrophic compared to heterotrophic conditions (Fig. 2).

Denitrification (nirK/S, norBC)

To investigate the mechanisms of denitrification in culture BP, we identified genes encoding the nitrate reductase (narGHI), nitrite reductase (nirK or nirS), nitric oxide reductase (norBC), and nitrous oxide reductase (nosZ) in the MAGs of Noviherbaspirillum sp., Thiobacillus sp., Rhodoblastus sp., and Ramlibacter sp. Among these organisms, the transcripts of detected denitrification genes had significantly higher abundance under autotrophic conditions compared to heterotrophic conditions (Fig. 2). Although the denitrification pathway was incomplete in the Gallionellaceae sp., it appeared to contribute to nitrite reduction and nitric oxide reduction. Both nitrite reductase (nirK and nirS) and nitric oxide reductase (norB and norC) transcripts from the Gallionellaceae sp. were detected at significantly higher abundance under autotrophic conditions compared to heterotrophic conditions, and the proteins of NirK/S were detected under autotrophic conditions (Fig. 2). Since the transcripts and proteins of these nirK/S genes in the MAGs of the dominant Gallionellaceae sp. were detected under autotrophic conditions (Fig. 2), this suggests that the reduction of nitrite coupled Fe(II) oxidation in culture BP might have occurred enzymatically. However, an abiotic contribution to the reduction of nitrite and nitric oxide by Fe(II) cannot be ruled out. Alternatively, it is possible that the Gallionellaceae sp. was expressing nitrite reductase genes to detoxify the products derived from nitrate reduction performed by other community members.

Carbon Fixation

Carbon fixation is essential for the growth of autotrophic microorganisms; thus, we explored carbon fixation pathways in culture BP under autotrophic conditions. All six MAGs contained genes involved in carbon metabolism pathways, i.e., the reductive pentose phosphate (CBB) cycle, glycolysis, tricarboxylic acid (TCA) cycle, and pentose phosphate pathway (PPP; online suppl. Table S3). More specifically, genes responsible for carbon fixation (rbcL/S), encoding ribulose-1,5-bisphosphate carboxylase-oxygenase (RuBisCO), were identified in the MAGs of the Gallionellaceae sp., Noviherbaspirillum sp., Rhodoblastus sp., Thiobacillus sp., and Ramlibacter sp. (Fig. 2; online suppl. Fig. S4). The transcripts of rbcL were detected at a significantly higher level under autotrophic conditions compared to heterotrophic conditions for under autotrophic conditions. Grey font and lines indicate incomplete pathway of transcripts detected under autotrophic conditions. Intense blue colour shows all proteins of the complete pathway detected under autotrophic conditions. Lighter blue colour shows incomplete pathway of proteins detected under autotrophic conditions.

Fig. 3. Overview of the meta’omics results for the extracellular electron transfer system, denitrification, carbon metabolism, and oxidative phosphorylation in the three proposed key players for Fe(II) oxidation: the Gallionellaceae-BP sp. (a), Noviherbaspirillum sp. (b), and Thiobacillus sp. (c) under autotrophic conditions. Black font and lines indicate complete pathway of transcripts detected (For figure see next page.)
Microbial Network in a Novel NRFO Enrichment Culture

**Gallionellaceae-BP sp.**
- Partial denitrification
- Oxidative phosphorylation
- Fe(II) oxidation

**Noviherbaspirillum sp.**
- Denitrification
- Oxidative phosphorylation
- Fe(II) oxidation

**Thiobacillus sp.**
- Denitrification
- Oxidative phosphorylation

Transcripts of complete pathway detected under autotrophic conditions

Proteins of complete pathway detected under autotrophic conditions

Transcripts of incomplete pathway detected under autotrophic conditions

Proteins of incomplete pathway detected under autotrophic conditions
Gallionellaceae sp., Noviherbaspirillum sp., Thiobacillus sp., and Rhodoblastus sp., as well as rbcS for the Noviherbaspirillum sp. and Thiobacillus sp. (Fig. 2). However, none of the RuBisCO-related proteins were detected (Fig. 2). Phylogenetic analysis of the large-chain RuBisCO (RbcL; online suppl. Fig. S4A) and small-chain RuBisCO (RbcS; online suppl. Fig. S4B) revealed that the RbcL of the Gallionellaceae sp. from culture BP and culture KS were closely related to each other as well as to other FeOB (online suppl. Fig. S4A). This indicates that the RbcL of the Gallionellaceae sp. in culture BP might have the same function as the RbcL of the Gallionellaceae sp. in culture KS (online suppl. Fig. S4A).

Amino Acid Biosynthesis Pathways

Essential amino acids are vital for the survival of individual isolated organisms and, therefore, the biosynthesis of the essential amino acids of the bacterial community members in culture BP was analysed. Except for the Geothrix sp., all community members had most, if not all, essential amino acid biosynthesis pathways at least partially encoded. However, the gene encoding histidine isomerase (hisA) was detected at medium confidence for the MAG of the Gallionellaceae sp. and Rhodoferax sp. with the GapMind tool [Price et al., 2020]. The genes encoding ribose-phosphate diphosphokinase (prs) for histidine biosynthesis and homoserine kinase (thrB) for threonine biosynthesis were missing in the MAG of the Thiobacillus sp. Furthermore, the gene encoding glutamine synthetase (glnA) for glutamine biosynthesis was missing in the MAG of the Rhodoblastus sp. In addition, the genes encoding homoserine dehydrogenase (hom) and vitamin B12-dependent methionine synthase (metH) for methionine biosynthesis were missing in the MAG of the Ramlibacter sp. Almost all of the essential amino acid biosynthesis pathways in the MAGs of both Gallionellaceae sp. and Noviherbaspirillum sp. were expressed (i.e., the transcripts were detected).

Energy (ATP) Generation and Electron Transport Phosphorylation

In order to identify energy generation pathways in culture BP, we explored the different levels of gene expression and proteins for terminal electron acceptors. The genes involved in oxidative phosphorylation complexes I–V, were found in the MAGs of the Gallionellaceae sp., Noviherbaspirillum sp., Rhodoferax sp., Thiobacillus sp., Rhodoblastus sp., and Ramlibacter sp. Under autotrophic conditions, the Gallionellaceae sp., Noviherbaspirillum sp., and Rhodoblastus sp. had significantly higher expression of these respiration genes in comparison to heterotrophic conditions (Fig. 2). On the protein level, partial oxidative phosphorylation proteins were detected under autotrophic conditions for the Gallionellaceae sp. and Noviherbaspirillum sp. (Fig. 2, 3). In addition, cbb3-type and aa3-type cytochrome c oxidase-related transcripts and proteins of complex IV, which typically have high and low affinities to oxygen, were detected under both conditions (Fig. 2), even though culture BP was grown under anoxic conditions.

In situ Microbial Community Analysis

In order to identify the in situ distribution and relative abundance of the enriched populations in culture BP, field studies were conducted in 2017. The microbial community was investigated at the sampling site of the origin of culture BP (and culture KS). To this end, sediment cores of a pond and a ditch were obtained, and DNA- and RNA-based short-read 16S rRNA (gene) sequencing was performed from a total of 50 samples. At the sampling sites, 37 different ASVs were classified as Gallionellaceae (20 ASVs were detected at the pond and 23 ASVs at the ditch; online suppl. Fig. S6); of these, eight were identified as Gallionella spp., eleven as Sideroxydans spp., three as Candidatus Nitrotoga spp. and fifteen as unclassified spp. (online suppl. Fig. S5). In the environmental samples, among these ASVs assigned to the family Gallionellaceae (accounting for up to 3.13% relative abundance for the whole family; online suppl. Fig. S6), nine DNA-based and two RNA-based samples, corresponding to three sediment cores, revealed the same ASV (100% sequence identity) as the dominant Gallionellaceae sp. ASV in culture BP. It accounted for up to 0.13% relative abundance and was found both at the pond and at the ditch (online suppl. Fig. S6). Therefore, the Gallionellaceae sp. in culture BP (i.e., the investigated sequence) was detectable, relatively widespread, and likely active at the sampling sites. The 37 different Gallionellaceae ASVs were furthermore quite diverse and affiliated, based on phylogenetic analysis, for instance with Sideroxydans lithotrophicus ES-1, Gallionella capsiferriformans ES-2, and Ferrigenium kumadai (online suppl. Fig. S5). The other members in culture BP, i.e., the Rhodoferax sp., Thiobacillus sp., and Rhodoblastus sp., were also found at the pond and the ditch with up to 0.13% relative abundance for identical ASVs as in culture BP, respectively, and up to 2.0% relative abundance for the whole genus, i.e., Rhodoblastus (online suppl. Fig. S6). Neither the Noviherbaspirillum sp. and Ramlibacter sp. in culture BP, nor the Gallionellaceae sp. in culture KS, were found within these sediment samples.
However, our sampling procedure might just have missed them, in terms of the methodological detection limit, sampling location, or season.

**Discussion**

**A Novel Autotrophic Nitrate-Reducing Fe(II)-Oxidizing Enrichment Culture**

Microbial NRFO is considered to be an important pathway for nitrate removal in organic-poor, Fe-abundant environments, whereby microorganisms gain electrons from Fe(II) for carbon fixation and for energy generation (by nitrate reduction) which is utilized for growth, ultimately transforming CO₂ into biomass [Weber et al., 2006]. The enrichment of two autotrophic NRFO cultures (i.e., the culture KS and in this study culture BP) from two close-by organic-rich environments suggests that this process might be more widespread than expected and might therefore be occurring in diverse habitats.

The physiology in growing cultures of culture BP was slightly different from the results for culture KS described in previous studies. The ratio of Fe(II)\text{oxidized}/nitrate\text{reduced} was reported in the range of 4.3–4.8 in culture KS [Straub et al., 1996; Tominski, 2016; Tominski et al., 2018a], which was higher than the Fe(II)\text{oxidized}/nitrate\text{reduced} ratio of 3.4 in this study. Theoretically, the stoichiometry suggests that the Fe(II)\text{oxidized}/nitrate\text{reduced} ratio should be around 5 or even higher than 5, when the electrons required from Fe(II) for carbon fixation are also considered, potentially indicating incomplete denitrification or an additional electron donor (i.e., traces of dissolved organic carbon, DOC, in the growth medium) [Huang et al., 2021]. However, cell numbers increased from \(1.04 \times 10^6\) cells/mL to \(4.31 \times 10^6\) cells/mL within 4 days, and therefore the electrons must not only have derived from the DOC but also from Fe(II) oxidation.

In this study, we show that culture BP fulfils the following three criteria of autotrophic NRFO [Bryce et al., 2018]: (i) there is no need for an additional organic carbon substrate for continuous growth, (ii) the culture is able to maintain Fe(II) oxidation over several transfers without organic carbon addition, (iii) there is confirmed growth of cells with only Fe(II), nitrate, and CO₂ provided. Additionally, our meta’omics data demonstrated that culture BP not only has the genetic potential but also expresses carbon fixation genes. Especially the \(rbcL\) gene of the *Gallionellaceae* sp. in culture BP (hereafter *Gallionellaceae*-BP sp.) was closely related to the \(rbcL\) gene of the *Gallionellaceae* sp. in culture KS (hereafter *Gallionellaceae*-ae-KS sp.), which was proven to fix \(^{13}\)C-labelled inorganic carbon [Tominski et al., 2018b]. Culture BP is therefore a novel autotrophic nitrate-reducing Fe(II)-oxidizing enrichment culture that provides the opportunity to study NRFO under more environmentally relevant conditions, i.e., in a more complex microbial network with probably less laboratory evolutionary influence, compared to the long-term established culture KS.

**Role of Gallionellaceae for Fe(II) Oxidation in Culture BP and beyond**

In this study, the dominant *Gallionellaceae*-BP sp. in culture BP (51–87% relative abundance) was identified as Fe(II) oxidizer and was shown to be closely related to known FeOB, such as the unclassified *Gallionellaceae*-KS sp. from culture KS, a nitrate-reducing Fe(II)-oxidizing culture enriched from the same freshwater habitat as culture BP [Straub et al., 1996]. Comparing meta’omics analysis of culture KS from previous studies [He et al., 2016; Huang et al., 2021] with those of culture BP in this study, some interesting differences were observed for the *Gallionellaceae* spp. Although both the *Gallionellaceae*-KS sp. and *Gallionellaceae*-BP sp., likely perform Fe(II) oxidation, partial denitrification, and carbon fixation, the genes encoding these functions differed. For example, the proposed Fe(II) oxidation genes, \(cyc2\), \(mtoAB\), and \(mofAB\) were identified in the *Gallionellaceae*-KS sp. [He et al., 2016; Huang et al., 2021]. In addition, the transcripts of \(cyc2\), \(mtoAB\) and \(mofAB\) were all detected, and the MofA protein of the *Gallionellaceae*-KS sp. was shown to be present [Huang et al., 2021]. For the *Gallionellaceae*-BP sp. \(cyc2\) was detected at DNA, RNA, and protein levels in this study. In a previous study [Castelle et al., 2008], the outer membrane-bound cytochrome c Cyc2 of acidophilic FeOB, *Acidithiobacillus ferrooxidans*, was purified and shown to be responsible for performing Fe(II) oxidation. Therefore, combined with the fact that the *Gallionellaceae*-BP sp. dominates culture BP, these results indicate that the Cyc2 of the *Gallionellaceae*-BP sp. likely plays the main role for Fe(II) oxidation in culture BP under autotrophic conditions. As for denitrification, the *Gallionellaceae*-BP sp. harboured the genes encoding nitrite reductase (\(nirK/S\)) and nitric oxide reductase (\(norBC\)), whereas the *Gallionellaceae*-KS sp. harboured the genes encoding nitrate reductase (\(narGHJ\)) and nitrite reductase (\(nirK/S\)). In contrast to the *Gallionellaceae*-KS sp., the *Gallionellaceae*-BP sp. possesses an NO reductase and can reduce NO to N₂O. However, both *Gallionellaceae* spp. in culture BP and culture KS require other flanking community members to complete denitrification (i.e., they are miss-
ing the nosZ gene). To investigate whether nitrite or NO can serve as electron acceptors and generate energy in the Gallionellaceae-BP sp., further experiments are required. Moreover, we detected coxABC encoding for aa3-type cytochrome c oxidases and belonging to the Gallionellaceae-BP sp. and additional community members in culture BP, both at the transcript level and some at the protein level. Interestingly, coxABC are normally observed under oxic conditions [Krab and Slater, 1979]. In contrast, in our cultivation system under autotrophic and heterotrophic conditions, we used N2/CO2 in the headspace and amended with FeCl2 prior to inoculation, which would have consumed any atmosphere O2 contamination during the preparation procedure. However, one possible scenario might be that the coxABC genes in the organisms in culture BP do not require oxygen to be expressed. Alternatively, there might be a so far undetected internal oxygen production pathway in at least one of the organisms in culture BP, similar to the proposed aerobic methane oxidation under anoxic conditions [Ettwig et al., 2010]. The latter hypothesis was already discussed for culture KS by He et al. [2016]. Nevertheless, these possible scenarios remain highly speculative at this point and require further investigation.

Based on 16S rRNA gene sequences, the Gallionellaceae-BP sp. was closely related to Ferrigenerium kumadai, and Sideroxydans lithotrophicus strain ES-1. While Ferrigenerium kumadai was isolated from rice paddy soils [Khalifa et al., 2018; Nakagawa et al., 2020], Sideroxydans lithotrophicus strain ES-1 was isolated from a groundwater-fed iron seep [Emerson and Moyer, 1997]. It is apparent, when combining our results with other studies, that Gallionellaceae spp. play important roles as FeOB within freshwater ecosystems. Furthermore, our in situ survey indicated a high diversity of Gallionellaceae spp. at the field site, suggesting that, in addition to the microbial populations in culture KS and culture BP, there might be a higher diversity of organisms contributing to NRFO in freshwater environments and other habitats than currently known from cultivation approaches.

Contribution of Noviherbaspirillum sp. and Thiobacillus sp. to NRFO in Culture BP

The 2nd most abundant organism in culture BP under autotrophic conditions was the Noviherbaspirillum sp. (9–15%). Based on 16S rRNA gene sequences, the top three closest related isolates were Noviherbaspirillum autotrophicum strain TSA66 [Ishii et al., 2017], Noviherbaspirillum agri K-1-15 [Chaudhary and Kim, 2017], and Noviherbaspirillum denitrificans TSA40 [Ishii et al., 2017] (online suppl. Table S1). Among these isolates, only Noviherbaspirillum autotrophicum strain TSA66 was reported to be able to grow autotrophically by using H2 as an energy source to perform denitrification [Ishii et al., 2017]. However, a gene search on the Joint Genome Institute’s Integrated Microbial Genome (JGI/IMG) database showed that the genome of Noviherbaspirillum autotrophicum strain TSA66 has genes encoding RuBiSCO but does not have Fe(II) oxidation gene homologues, either mtoA or cyc2, which were detected in the Noviherbaspirillum sp. from culture BP. Therefore, the Noviherbaspirillum sp. from culture BP might have unique capabilities for NRFO within this genus.

Another potential FeOB in culture BP was the Thiobacillus sp., which accounted for only up to 1.6% relative abundance in culture BP and possessed the potential Fe(II) oxidation gene homologues cyc2 and mofA. The top three closest related isolates based on 16S rRNA gene sequences with an available genome sequence were Thiobacillus thioparus strain THI 111, Thiobacillus thioparus DSM 505, and Thiobacillus thioparus strain Starkey [Boden et al., 2012] (online suppl. Table S1). Additionally, Thiobacillus denitrificans ATCC 25259 was previously proposed as an autotrophic nitrate-reducing Fe(II)-oxidizing organism [Straub et al., 1996; Beller et al., 2006]. Interestingly, only Thiobacillus thioparus DSM 505 has the Fe(II) oxidation gene cyc2, and neither Thiobacillus denitrificans ATCC 25259 and Thiobacillus denitrificans RG nor Thiobacillus thiophilus DSM 19892 revealed potential Fe(II) oxidation genes. The copper resistance genes, copAB, i.e., the homologues of the putative Fe(II)-oxidizing genes, pcoAB [He et al., 2017], were detected in Thiobacillus denitrificans ATCC 25259, Thiobacillus denitrificans RG, Thiobacillus denitrificans DSM 12475, and Thiobacillus thiophilus DSM 19892 [Beller et al., 2006, 2013]. We therefore suggest that the Thiobacillus sp. in culture BP was also actively contributing to NRFO.

Interestingly, in culture KS, there seems to be no community member with a similar role to that of the Noviherbaspirillum sp. and Thiobacillus sp. in culture BP. For example, the 2nd most abundant organism in culture KS under autotrophic conditions, a Rhodanobacter sp., was the main organism cooperating with Gallionellaceae-KS, which is likely able to perform NRFO under autotrophic conditions [He et al., 2016; Huang et al., 2021]. The Rhodanobacter sp. likely performs Fe(II) oxidation and complete denitrification but does not have the ability to conduct carbon fixation [Huang et al., 2021]. Therefore, the isolation and the study of Noviherbaspirillum sp. and Thiobacillus sp. in culture BP, which likely contribute to
Fe(II) oxidation and CO₂ fixation, would be beneficial in order to broaden our knowledge of autotrophic nitrate-reducing Fe(II)-oxidizing bacteria.

Expanding the Microbial Network for NRFO in Culture BP

Besides the Gallionellaceae-BP sp., Noviherbaspirillum sp., and Thiobacillus sp., other flanking members potentially also play a relevant role in culture BP: Rhodoblastus sp. (up to 5% relative abundance), Rhodoferax sp. (up to 3%), and Ramlibacter sp. (up to 1%). We identified Fe(II) oxidation gene homologues in the MAGs of the Rhodoblastus sp. (cyc2 and mtoA) and the Ramlibacter sp. (cyc2 and mtoB), indicating that they can potentially play a role in Fe(II) oxidation, despite the fact that cyc2 transcripts were not detected for the Rhodoblastus sp., and neither cyc2 nor mtoB transcripts for the Ramlibacter sp. were detected at significant levels. Furthermore, we observed that the genes necessary to perform full denitrification were present in these three flanking members, i.e., the Rhodoblastus sp., Rhodoferax sp., and Ramlibacter sp., and some of them were detected at significantly higher transcript levels. Regarding carbon fixation, there was a complete CBB reductive pentose phosphate cycle detected in both the Rhodoblastus sp. and Ramlibacter sp., while only a partial cycle was found for the Rhodoferax sp. (online suppl. Table S3). As a result, although the relative abundance of these three flanking community members accounted for less than 5% (respectively) throughout the autotrophic growth conditions, our findings suggest that they play partial roles in NRFO and carbon fixation (Fig. 4).

To summarize, there were five potential Fe(II)-oxidizing microbial key players in culture BP: the Gallionellaceae-BP sp., Noviherbaspirillum sp., Thiobacillus sp., Rhodoblastus sp., and Rhodoferax sp. (Fig. 4). Among these, the Gallionellaceae-BP sp., Noviherbaspirillum sp., and Thiobacillus sp. may not only perform Fe(II) oxidation but also carbon fixation (Fig. 3, 4). The resulting fixed organic carbon was likely provided to other heterotrophic community members and used for their growth (Fig. 4). During denitrification, the nitrate may be reduced by flanking community members, while the Gallionellaceae-BP sp. may contribute to partial denitrification, i.e., ni-

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**Fig. 4.** Overview of the proposed microbial interactions in culture BP, mainly essential for the organisms’ survival under autotrophic conditions. The depicted putative reactions are based on the meta’omics data for Fe(II) oxidation (red arrows and font), denitrification (blue arrows and font), and CO₂ fixation (black arrows and font), for the Gallionellaceae-BP sp., Noviherbaspirillum sp., Thiobacillus sp., Rhodoblastus sp., and Rhodoferax sp. The cells in blue indicate FeOB with carbon fixation ability, in grey indicate heterotrophs, and in green indicate that the organism can conduct carbon fixation and potentially Fe(II) oxidation. The detected transcripts are in italic font, and a combination of both detected transcripts and proteins are with underlined font. The transcripts with significantly higher expression under autotrophic conditions compared to heterotrophic conditions are in bold font.
trite and nitric oxide reduction, and the remaining flanking community members complete the denitrification (Fig. 4). Therefore, in order to survive and indeed thrive, the dominant *Gallionellaceae*-BP sp. has to cooperate with other community members under autotrophic conditions (Fig. 4). In return, the heterotrophic community members may have to rely for their survival on Fe detoxification and carbon fixation likely performed by the *Gallionellaceae*-BP sp., *Noviherbaspirillum* sp., and *Thiobacillus* sp. (Fig. 4). In addition, our findings highlighted potential interdependencies between the microbial community members regarding amino acid biosynthesis that might be crucial for their survival as well.

Comparing the microbial network of culture BP with that of culture KS, the most active species in culture KS were the *Gallionellaceae*-KS sp. and the *Rhodanobacter* sp. under autotrophic conditions [Huang et al., 2021]. However, in culture BP, there were several active populations involved in NRFO under autotrophic conditions, including the *Gallionellaceae*-BP sp., *Noviherbaspirillum* sp., *Thiobacillus* sp., *Rhodoblastus* sp., *Rhodoferax* sp., and *Ramilibacter* sp. (Fig. 2, 3). Therefore, in contrast to culture BP, culture KS is likely characterized by a reduced microbial network, which may be due to long-term cultivation and selection of specialized organisms. The occurrence of the lower relative abundance, yet active, organisms (i.e., the *Rhodoblastus* sp., *Rhodoferax* sp., and *Ramilibacter* sp.) in culture BP expands the microbial network contributing to NRFO processes, which might be more similar to the complex microbial interactions in the environment.

**NRFO Core Features of Complex Microbial Networks including *Gallionellaceae* spp.**

There seems to be several core features that might be similar for autotrophic NRFO processes carried out by complex microbial networks including *Gallionellaceae* spp. (Fig. 4): (i) NRFO requires Fe(II) oxidation features, with several possible electron transfer pathways (i.e., different putative Fe(II)-oxidizing proteins might be used); (ii) during denitrification, the *Gallionellaceae* spp. may contribute to partial denitrification (e.g., reducing nitrate to nitric oxide or nitrite to nitrous oxide), while the flanking microbial community members complete the denitrification; (iii) fixed organic carbon from the autotrophic lifestyle of *Gallionellaceae* spp., is provided to other heterotrophic community members and used for their growth; therefore, the key FeOB (i.e., the *Gallionellaceae* sp.) has to cooperate with other community members under autotrophic conditions; however (iv) the identity of these flanking community members seems interchangeable. We suggest that these core features are relevant for NRFO processes in the environment as well, that NRFO is widespread, including organic-rich habitats, and that several microbial key players of NRFO *in situ* thrive in complex microbial networks and are yet-to-be discovered.

**Experimental Procedures**

_Cultivation, Analytical Methods, and Cell Counts_

Culture BP originated from a freshwater pond at the Max Planck Institute for Marine Microbiology, located in Bremen, Germany, sampled in 2015 (described in the online suppl. material). Since then, culture BP was transferred with 10% (v/v) inoculum, including more than 20 transfers per year from 2018, growing with 25 mL anoxic headspace (the N₂/CO₂ ratio was 90/10), unfiltered, bicarbonate-buffered medium, containing 10 mM FeCl₂, 4 mM NaNO₃, vitamins, and trace elements with a final pH of 6.9–7.2 in 58 mL serum bottles [Hegler et al., 2008; Blöthe and Roden, 2009]. The incubation temperature was 28°C in the dark. Under heterotrophic conditions, 5 mM acetate was used instead of 10 mM FeCl₂ as the electron donor. Analytical methods of Fe(II), Fe(total), nitrate, nitrite and acetate concentrations were described by Tominiski et al. [2018a]. Prior to cell counts, a mineral dissolution step was conducted, using the protocols obtained from personal communication with Stefanie Becker, as described in the online supplementary material. Cells were stained with BacLight Green stain (Thermo Fisher Scientific, 1 mM stain/1 mL sample) and cell numbers were determined with an Attune NxT Flow Cytometer (Thermo Fisher Scientific) with the instrument setup adapted from the procedure of Schmidt et al. [2020], as described in the online supplementary material. Estimated absolute cell numbers were calculated by multiplying flow cytometer-based total cell numbers with short-read 16S rRNA gene amplicon sequencing relative abundance data (described below).

**Experimental Setup, Biomass Sampling, DNA/RNA Co-Extraction and Protein Extraction of Culture BP**

Both autotrophic and heterotrophic conditions were used with triplicates, respectively, in the meta’omics experiment (described in the online suppl. material). Samples were taken on the 2nd day from cultures grown under autotrophic conditions. For the control group, it was deemed essential to incubate the cultures for some time prior to the main experiment to avoid protein carryover from remaining proteins of Fe(II) oxidation. Therefore, under heterotrophic conditions, pre-cultures were first grown for two transfers sequentially with 10 and 1% (v/v) inoculum for at least 24 h each. In a second step, the 3rd transfer with 1% (v/v) inoculum under heterotrophic conditions was used for the experimental setup and samples were taken after approximately 18 h. DNA, RNA, and protein samples were collected from culture BP grown under autotrophic and heterotrophic conditions. DNA/RNA co-extraction was performed according to an upscaled version of Lueders et al. [2004] (described in the online suppl. material) and used for meta’omics and short-read 16S rRNA (gene) amplicon sequencing. DNA for long-read 16S rRNA gene amplicon sequencing was
extracted using the DNEasy UltraClean Microbial Kit (QIAGEN, Hilden, Germany), according to the user manual. Protein extraction was performed according to Spät et al. [2015], as described in the online supplementary material.

**Sampling and DNA/RNA Co-Extraction for in situ Analysis**

For environmental sampling, six cores were obtained, three from a pond and three from a ditch near the Max Planck Institute for Marine Microbiology in Bremen. The sediment cores were separated into six different depths: 0–0.5, 0.5–1, 1–2, 2–4, 4–7, and 7–10 cm. DNA and RNA were co-extracted using the method published by Griffiths et al. [2000] with modifications described in the online supplementary material.

**Short-Read 16S rRNA (Gene) Amplicon Sequencing**

DNA from culture BP as well as DNA and cDNA (cDNA synthesis is described in the SI) from sediments were used for amplification with universal primers, i.e., 515F: GTGYYACCMGGG-GCGGTGAA [Parada et al., 2016] and 806R: GGCATACNVGGGT-WTCTAAT [Apprill et al., 2015] fused to Illumina adapters. The PCR, library preparation, and sequencing are detailed in the online supplementary material. Quality control, reconstruction of 16S rRNA gene sequences, and taxonomic annotation was performed with nf-core/ampliseq v1.1.0 [Ewels et al., 2020; Straub et al., 2020] as outlined in the online supplementary material.

**Long-Read 16S rRNA Gene Amplicon Sequencing**

DNA from culture BP was used for amplification with primers 27F and 1492R. PacBio Sequel SMRT long-read amplicon sequencing was performed at the Helmholtz Zentrum München, Germany. Circular consensus sequences were analysed with DADA2 v1.10.0 [Callahan et al., 2016] Callahan et al., 2019 in R v3.5.1 [R Development Core Team, 2018]. Further details are described in the online supplementary material.

**Metagenome Analysis and Draft Genome Recovery**

Library preparation and shotgun Illumina sequencing of culture BP grown under autotrophic and heterotrophic conditions were performed by GeGaT,Tuebingen, Germany. Raw read quality control, assembly, metagenome assembled genome binning and taxonomic annotation was performed with nf-core/mag v1.0.0 (https://nf-co.re/mag, DOI: 10.5281/zenodo.3589528) [Ewels et al., 2020]. Further details are described in the online supplementary material. Characteristics of the assembled metagenome can be found in the online supplementary Table S2. Twelve MAGs were obtained from the metagenome, of which eleven had determined completeness and four had between 11 and 45% completeness. Selected MAGs are presented in the online supplementary Table S1.

The assembled metagenome and MAGs were uploaded to the Joint Genome Institute’s Integrated Microbial Genome and Microbiome Expert Review (IMG/MER) pipeline (IMGAP v5.0.8) for annotation (available online at: https://img.jgi.doe.gov/cgi-bin/mer/main.cgi) [Chen et al., 2019]. FeGenie [Garber et al., 2020] and the IMG database [Chen et al., 2019] were used to search for potential Fe(II) oxidation genes, i.e., cyc2, mtoAB, pcoAB, mofA [He et al., 2017], for the closely related species of the community in culture BP.

**Metatranscriptome Analysis**

For RNA analysis, DNase treatment, library preparation including bacterial ribodepletion (with Illumina Stranded Total RNA Prep with Ribo-Zero Plus kit), and sequencing with 2 × 75 bp and 6–53 MiO clusters per sample were performed by Microsynth AG (Balghach, Switzerland) using triplicate samples of culture BP grown under autotrophic and heterotrophic conditions, respectively. For data analysis, nf-core/rnaseq v1.4.2 (https://nf-co.re/rnaseq) [Ewels et al., 2020] performed quality control, mapping on the metagenome, and differential abundance testing, as detailed in the online supplementary material. The definition of a significant difference was postulated for transcripts with Benjamini and Hochberg-adjusted \( p \leq 0.05 \). A summary of features of the metatranscriptome is described in online supplementary Table S2.

**Phylogenetic Analysis, Figure Illustration, and Data Availability**

Phylogenetic analysis of Cyc2, MtoA/B, RbcL/S protein sequences of culture BP and 16S rRNA genes of Gallionellaceae spp., identified in the environmental samples were conducted with MEGA X [Kumar et al., 2018] as described in the online supplementary material. The amino acid biosynthesis pathways were determined by using the tool: GapMind: Automated Annotation of Amino Acid Biosynthesis (https://papers.genomics.lbl.gov/cgi-bin/gapView.cgi) [Price et al., 2020]. The physiological growth plots, cell count plots, heat map, and bubble plots (Fig. 1, 2; online suppl. Fig. S6) were constructed via R v3.6.1 and its working interface RStudio (https://www.R-project.org/ and http://www.rstudio.com/) [RStudio Team, 2019; R Development Core Team, 2020]. Figure illustrations of cell shapes (Fig. 3, 4) were adapted from “Icons > Cell Membranes > Simplified Bilayer Membranes” and “Icons > Cell Structures > Organelles,” respectively, using BioRender.com (2021; https://app.biorender.com/biorender-templates). The datasets presented in this study can be found in online repositories. The names of the repositories and accession numbers are described below and in the online supplementary Table S4. Raw sequencing data and metagenome assembly were deposited at the Sequence Reads Archive (SRA; https://www.ncbi.nlm.nih.gov/bioproject/PRINA693457). The mass spectrometry proteomics data have been deposited to the ProteomeXchange Consortium via the PRIDE [Perez-Riverol et al., 2019] partner repository with the dataset identifier PXD023710. The IMG metagenome ID for BP culture is 3300036710 and the corresponding accession numbers of MAGs are: Gallionellaceae sp. 2831290873; Novitheraspirillum sp. 2831285802; Rhodofex sp. 2840079448; Thiobacillus sp.
2840071692; *Rhodoblastus* sp. 2840074988; *Ramlibacter* sp. 2840082686; *Geothrix* sp. 2840087968, and *Betaproteobacteria* bacterium 2840090052 (online suppl. Table S1 and S2) [Chen et al., 2019]. The other four MAGs did not reach the quality criteria for submission to IMG.

**Acknowledgements**

We thank Shun Li for obtaining the sediment sample for cultivation, Jens Harder, Ingrid Kunze, Timm Bayer, Lea Sauter, and Casey Bryce for assistance during sampling, Tillmann Lueders, Zhe Wang, and Natalia Jakus for PacBio library preparation and sequencing, Irina Droste-Borel for metaproteomic measurement, Verena Nikeleit, Ellen Roehm, and Franziska Schaedler for acetate, nitrate and nitrite measurement, as well as Stefanie Becker for developing the cell count methods for flow cytometry.

**Statement of Ethics**

As no human or animal research materials were involved in this study, the paper is exempt from ethical committee approval.

**Conflict of Interest Statement**

The authors have no conflicts of interest to declare.

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**Funding Sources**

This work was supported by the German Research Foundation (Deutsche Forschungsgemeinschaft, DFG)-funded research train- ing group RTG 1708 “Molecular Principles of Bacterial Survival Strategies.” Daniel Straub was funded by the Institutional Strategy of the University ofTuebingen (DFG, ZUK63). Andreas Kappler is funded by the DFG under Germany’s Excellence Strategy, cluster of Excellence EXC2124 (project ID 390838134). Nia Blackwell was funded by the Collaborative Research Center 1253 CAMPOS (Project 5: Fractured Aquifers) from the DFG (grant No. SFB 1253/1 2017). Sara Kleindienst is funded by an Emmy-Noether fel- lowship from the DFG (grant No. 326028733).

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Y.-M.H.: experimental design and conduct, metagenomic, metatranscriptomic and 16S rRNA amplicon sequencing data interpretation, manuscript writing and revising, final approval of the version to be published. D.S.: experimental design, metagenomic, metatranscriptomic and 16S rRNA amplicon sequencing bioinformatics analysis, manuscript writing and revising, final approval of the version to be published. A.K.: experimental design, supervision, laboratory resources, manuscript revising, final approval of the version to be published. N.S.: metagenomic and 16S rRNA amplicon sequencing data interpretation, manuscript writing and revising, final approval of the version to be published. N.B.: experimental design, supervision, manuscript revising, final approval of the version to be published. S.K.: experimental design, supervision, funding acquisition, laboratory resources, manuscript revising, final approval of the version to be published.

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