

Fe(III) mineral formation and cell encrustation by the nitrate-dependent Fe(II)-oxidizer strain BoFeN1

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ABSTRACT

Understanding the mechanisms of anaerobic microbial iron cycling is necessary for a full appreciation of present-day biogeochemical cycling of iron and carbon and for drawing conclusions about these cycles on the ancient Earth. Towards that end, we isolated and characterized an anaerobic nitrate-dependent Fe(II)-oxidizing bacterium from a freshwater sediment. The 16S rRNA gene sequence of the isolated bacterium (strain BoFeN1) places it within the β -Proteobacteria, with *Acidovorax* sp. strain G8B1 as the closest known relative. During mixotrophic growth with acetate plus Fe(II) and nitrate as electron acceptor, strain BoFeN1 forms Fe(III) mineral crusts around the cells. The amount of the organic cosubstrate acetate present seems to control the rate and extent of Fe(II) oxidation and the viability of the cells. The crystallinity of the mineral products is influenced by nucleation by Fe minerals that are already present in the inoculum.

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INTRODUCTION

Microbial reduction of ferric iron [Fe(III)] plays a major role in mineralization of organic carbon in freshwater and marine environments as well as in soils (Lovley, 1991; Thamdrup, 2000). The product of this metabolism, i.e. ferrous iron [Fe(II)], is reoxidized to Fe(III) at neutral or alkaline pH by molecular oxygen (O₂) in a chemical reaction, whereas under acidic conditions, Fe(II) remains stable (Stumm & Morgan, 1996). Micro-organisms can catalyse the aerobic oxidation of Fe(II) at acidic (Wood, 1988; Blake *et al.*, 1993; Blake & Johnson, 2000) and neutral pH (Hanert, 1981; Wood, 1988; Emerson & Revsbech, 1994; Emerson, 2000) and successfully compete with chemical oxidation. The discovery of phototrophic (Widdel *et al.*, 1993) and nitrate-dependent Fe(II) oxidizers (Straub *et al.*, 1996; Benz *et al.*, 1998) showed that micro-organisms can close the iron cycle not only under oxic but also under anoxic conditions, thus increasing the importance of Fe(III) as electron acceptor for carbon mineralization under anoxic conditions.

Fe(II) oxidation under O₂-free conditions not only plays a role in modern biogeochemical cycling of iron and carbon. Also for a long period in early Earth history virtually no O₂ was

available for chemical or microbial oxidation of Fe(II). Therefore, it is likely that anaerobic, O₂-independent oxidation of Fe(II) also contributed to Fe cycling in the past, as evidenced by the deposition of Precambrian iron mineral deposits (Banded Iron Formations) during a period in Earth history preceding the presumed advent of oxygenic photosynthesis (Hartman, 1984). To date, the mechanism of anaerobic Fe(II) oxidation that has been discussed most in this context has been Fe(II) oxidation by anoxygenic phototrophs (Widdel *et al.*, 1993; Croal *et al.*, 2004; Kappler *et al.*, 2005). However, anaerobic oxidation of Fe(II) by nitrate could also have contributed to Fe cycling in the past, although most likely at a stage in Earth history following the rise of O₂, given that nitrate is a highly oxidized compound that formed in substantial amounts only after molecular oxygen was present (Beaumont & Robert, 1999).

The biochemistry of Fe(II) oxidation has been studied in detail with aerobic acidophilic Fe(II) oxidizers (Blake & Johnson, 2000); however, much less is known about the mechanisms of neutrophilic Fe(II) oxidation. The neutrophilic aerobic Fe(II) oxidizers *Gallionella* and *Leptothrix* produce extracellular organic polymers that nucleate Fe(III) precipitates. Examples include the stalks typical of *Gallionella ferruginea* and

sheaths typical of *Leptothrix ochracea* (Emerson & Revsbech, 1994). It has been suggested that these microbes use these strategies to avoid encrustation of the metabolizing cells that would lead to impaired substrate uptake and metabolite release, and might even cause cell death (Hanert, 1981; Hallberg & Ferris, 2004). Nothing is known so far about the mechanism of anaerobic Fe(II) oxidation. In particular, it is not clear where in the cell or at the cell membrane Fe(II) is oxidized, how the bacteria deal with the poor solubility of the product of their metabolism, i.e. the Fe(III), and how they avoid or overcome encrustation. A microenvironment with lowered pH values in close cell vicinity was observed around colonies of phototrophic Fe(II) oxidizers fixed in semisolid agarose (Kappler & Newman, 2004) in a similar manner as observed for microaerophilic Fe(II) oxidizers (Rodén *et al.*, 2004; D. Emerson, personal communication). Such acidification could help explain why these microbes do not form a Fe(III) mineral crust around their cells.

While it was demonstrated that phototrophic and microaerophilic Fe(II) oxidizers are truly lithoautotrophic microorganisms (Ehrenreich & Widdel, 1994; Emerson & Moyer, 1997; Sobolev & Rodén, 2004), most nitrate-dependent Fe(II) oxidizers isolated so far need an organic cosubstrate for growth (Straub *et al.*, 1996, 2004; Benz *et al.*, 1998; Lack *et al.*, 2002). Only some psychrophilic, neutrophilic Fe-oxidizing bacteria isolated from the deep sea (Edwards *et al.*, 2003) and some nitrate-dependent Fe(II)-oxidizing enrichment cultures (Benz *et al.*, 1998; Straub & Buchholz-Cleven, 1998) oxidize Fe(II) without the addition of organic substrates. However, it has not been demonstrated that these cultures can be cultured successively for several generations with Fe(II) as the sole electron donor, and currently, no pure culture of lithotrophic nitrate-dependent Fe(II) oxidizers is available. The question still remains whether Fe(II) oxidation is beneficial for mixotrophic Fe(II) oxidizers at all, or whether it is just a side activity that could even be harmful for the cells because of cell encrustation.

In this study, we tried to determine how the presence of an organic cosubstrate affects the rate and extent of Fe(II) oxidation, cell encrustation and identity of the Fe(III) minerals precipitated by a newly isolated mixotrophic nitrate-dependent Fe(II)-oxidizing bacterial strain.

MATERIALS AND METHODS

Isolation of organism and 16S rDNA sequence analysis

The nitrate-dependent Fe(II)-oxidizing strain BoFeN1 was isolated from Lake Constance littoral sediments. Sediment samples were taken at 1–2 m water depth at the southern shore of the north-western arm of Lake Constance (Überlinger See), close to the island Mainau where the sediment is slightly sandy and covered with limestone precipitates. Plastic core tubes (8 cm diameter) were immersed into the sediment with a special

handle. Inoculum material for direct dilution in agar shake tubes was taken at 1 cm sediment depth, at the transition zone between brownish surface sediment and grey deeper sediment. Oxygen penetrated into this sediment down to about 4 mm depth (Gerhardt *et al.*, 2005). Agar shake dilutions (Widdel & Bak, 1992) were run in bicarbonate-buffered oxygen-free nonreduced mineral medium (Ehrenreich & Widdel, 1994), with 2 mM acetate, 5 mM FeSO₄ and 5 mM NaNO₃ as substrates. Controls were performed with acetate plus nitrate or with ferrous sulfate plus nitrate as substrates. Colonies appearing after 4–5 weeks of incubation at 28 °C were diluted again in agar shake cultures and transferred into liquid medium after checking for purity by microscopic observation.

16S rDNA from 3 µL cell suspension was amplified by a standard polymerase chain reaction (PCR) method using primers 8F (5′–3′: AGAGTTTGATCCTGGCTCAG) and 1492R (5′–3′: GGTTACCTTGTTACGACTT). The PCR program was as follows: 95 °C for 5 min and then 33 cycles at 95 °C for 30 s, 50 °C for 40 s and 72 °C for 2 min, followed by a 10-min extension time at 72 °C. Amplification was confirmed by agarose gel electrophoresis (1% agarose). For sequencing of the 16S rDNA gene of strain BoFeN1, the primers 8F, 1492R, and the primers 50F (5′–3′: AACACATGCAAGTCGAACG), 356F (5′–3′: ACTCCTACGGGAGGCAGCA), 515F (5′–3′: GTGCCAGCMGCCGCGGTAA) and 805F (5′–3′: ATTAGATACCCTGGTAGTC) were used. Sequencing was performed at the DNA Sequencing Core Facility at the Beckman Institute at Caltech. The resultant sequences were edited and aligned using the ASSEMBLYLIGN™ software (IBI, Kodak Co., New Haven, CT, USA). Neighbour-joining, parsimony and maximum likelihood phylogenetic trees were constructed using the ARB freeware package (O. Strunk and W. Ludwig, Technical University of Munich; <http://www.arb-home.de/>) running within the Linux environment, and the trees were compared to determine the relative robustness of the resulting phylogenetic tree topologies. The tree shown in Fig. 1 was constructed after the maximum-likelihood method using TreePuzzle within the ARB software package with 1388 positions considered (TreePuzzle is a quartet maximum-likelihood method for reconstructing tree topologies). Phylogenetic tree layout-editing was performed using TREEVIEW 1.6.6 for Windows (Page, 1996).

Media and growth conditions

Strain BoFeN1 was cultivated in freshwater mineral medium prepared after Ehrenreich & Widdel (1994). The medium was buffered at pH 6.8 with bicarbonate. Vitamin and trace element solutions (1 mL each) and a vitamin B12 solution (1 mL) were added (for details and concentrations see Ehrenreich & Widdel, 1994). This medium was used for growth with nitrate and acetate in the presence or absence of Fe(II). Bottles were closed with butyl rubber stoppers, crimped, and the headspace was flushed with N₂/CO₂ (v/v, 80/20).

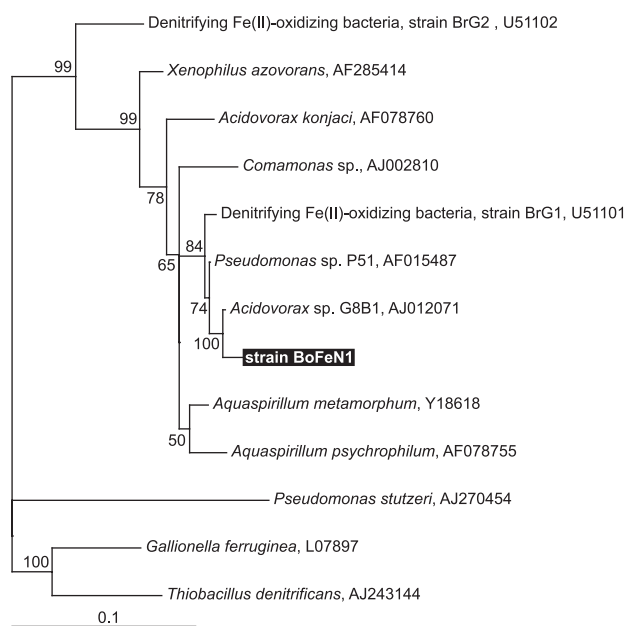


Fig. 1 16S rRNA sequence comparison of strain BoFeN1 with other strains on the basis of databank information. The tree was constructed after the maximum-likelihood method using TreePuzzle within the ARB software package with 1388 positions considered. For analyses of aligned protein sequences, 10 000 puzzling steps were employed. Accession numbers are listed after the bacterium. The scale bar represents evolutionary distance as 0.1 change per nucleotide position. The numbers given are bootstrap values.

For experiments with dissolved Fe(II), the medium was transferred into an anoxic glove box, and 5–10 mL of an anoxic 1 M FeCl₂ stock solution was added followed by precipitation of a whitish-grey precipitate, probably consisting of vivianite and siderite formed from phosphate and bicarbonate present in the medium. After 24 h the medium was filtered (0.2 µm, cellulose nitrate, Millipore) leaving a clear solution with ~2–6 mM dissolved Fe(II). In uninoculated controls, no further precipitation was observed for the duration of the incubations. This method allowed analysis of the Fe(III) precipitates formed by the Fe(II)-oxidizing bacteria in the absence of Fe(II) precipitates in the background. The medium (25 mL) was transferred into 58-mL serum bottles that were closed with butyl rubber stoppers, crimped and flushed with N₂-CO₂ (v:v, 80:20). Cultures were incubated at 30 °C.

Growth of strain BoFeN1 with nitrite as electron acceptor was tested with 2 mM acetate as electron donor and 5 mM NO₂⁻ as electron acceptor; the turbid culture was transferred subsequently several times.

Analytical methods

For analysis of dissolved iron, 200 µL culture suspension was withdrawn with a syringe and filtered over nylon (0.22 µm) filter tubes (Costar, Corning, NY, USA) in an anoxic glove box. The filtrate was analysed for total dissolved Fe and for

dissolved Fe(II) with the ferrozine assay after incubation in the presence or absence of the reducing agent hydroxylamine hydrochloride (Stookey, 1970). For analysis of total iron, 100 µL suspension was added to 900 µL 6 M HCl, and incubated for 1 h. After filtration over 0.22 µm nylon filter tubes, total Fe and total Fe(II) were determined as above. All iron measurements with the ferrozine assay were carried out in triplicates. The standard deviation (SD) for these triplicate measurements was usually smaller than the SD calculated for different independent experiments. For analysis with ion chromatography (IC) and high-pressure liquid chromatography (HPLC), 1 mL samples from culture suspension were exposed to air and vortexed to rapidly oxidize remaining Fe(II) to avoid further reaction with nitrite. After complete oxidation of Fe(II), samples were filtered in nylon filter tubes (0.5 mL volume, 0.22 µm) to remove all precipitates and cells. Acetate was analysed by HPLC with an Aminex HPX-87H column (Bio-Rad, Hercules, CA, USA) at 25 °C. Phosphoric acid (30 mM) was used as eluent in combination with a UV detector (absorbance followed at 210 nm wavelength). IC analysis was performed on an IonPac column (ASII, 4 × 250 mm, Dionex, Sunnyvale, CA, USA) at 1 mL min⁻¹ flow rate at 35 °C with a conductivity detector.

N₂O and NO were quantified by gas chromatography coupled to mass spectrometry (GC-MS) directly from the culture headspace.

Mineral analyses and microscopy

BET surface area

The surface area of biogenic Fe(III) mineral precipitates was determined using the multipoint BET nitrogen adsorption technique (Micromeritics (Norcross, GA, USA) Gemini 2360 surface area analyser). After centrifugation of 300 mL culture and treatment of the precipitates with bleach (~5% Na-hypochlorite, 8 h, room temperature) to oxidize and remove associated organic matter, the precipitates were washed with water and air dried, and the BET surface area was analysed. No change in mineral identity by this procedure was observed by X-ray diffraction.

Powder X-ray diffraction

Samples of culture suspensions were withdrawn with a syringe in an anoxic glove-box and were centrifuged to recover the precipitates. To remove remaining dissolved Fe(II) (to avoid oxidation and formation of nonbiogenic Fe(III) precipitates) the microbially produced precipitates were washed with water that had been degassed under vacuum, flushed with N₂ and equilibrated under an anoxic atmosphere for several days. After spreading on a glass disk, the precipitates were dried inside the glove-box. XRD spectra were obtained on a Scintag Pad V X-ray Powder Diffractometer (Scintag Inc., Cupertino, CA, USA) using Cu-Kα radiation operating at 35 kV and 30 mA with a θ-2θ goniometer equipped with a germanium solid-state detector. Each scan used a 0.04° step size from 10° to 80° with

a counting time of 2 s per step. Structures were identified by comparison to spectra in the PCPDFWIN program, © JCPDS-International Centre for Diffraction Data, 1997 (Newton Square, PA, USA) as well as to spectra of synthetic ferrihydrite and goethite.

Scanning electron microscopy (SEM)

Per sample, a 100 µL culture suspension was withdrawn with a syringe, centrifuged and washed with 300 µL ultrapure anoxic water to remove medium salts. One drop of this suspension was spread on a carbon-coated copper grid (200 mesh, 3 mm, SPI, West Chester, PA, USA) and dried under an anoxic atmosphere in the glove-box. Samples were examined by a LEO 1550VP Field Emission SEM with an Oxford INCA energy dispersive X-ray spectroscopy unit. The system was operated at 1–15 kV for high resolution secondary electron imaging and elemental analysis. Because of reduced beam penetration, the low voltage SEM (<5 kV) was particularly useful in surface imaging of the uncoated fine particles.

Environmental scanning electron microscopy (ESEM)

Samples for environmental scanning electron microscopy (ESEM) were withdrawn from cultures and placed directly on an ESEM sample holder. After the precipitates/cells settled, the overlying liquid was carefully removed by wicking with a paper tissue. The residue was washed with nanopure anoxic water to remove salts present in the medium to avoid precipitation during the analysis. Samples were examined by a FEI ESEM (FEI Company, Hillsboro, OR, USA) at 20 kV under wet mode conditions using the gaseous secondary electron (GSE) detector. The stage temperature was 4 °C with a chamber pressure of around 530 Pa to maintain a relative humidity of 85% at the sample. The elemental composition of the remaining mineral precipitates was determined with an EDAX detector for 60 s lifetime analysis.

Transmission electron microscopy (TEM)

Per sample, 5 mL of culture supernatants was centrifuged, fixed in 2.5–3% (v/v) glutaraldehyde for 12 h at 4 °C and washed three times in water. Half of the samples were stained with 2% osmium tetroxide and 2% uranyl acetate, while the other half were left unstained. Stained and unstained specimens were dehydrated in ethanol and acetone before embedding in EPON resin. After 24 h at 60 °C, samples were cut on a MT-X Ultramicrotome with a 55° Diatome diamond knife to 60 nm thickness; ultrathin sections were placed on 200 mesh copper grids. The prestained samples were subsequently poststained with 2% uranyl acetate for 3 min before final imaging. An Akashi EM-002B microscope operating at 100 kV was used for microscopy and EDS. The area sampled by the Oxford spectrum analyser was approximately 8.8 nm² at 83 kV. Acquisition rates were maintained at 10–20% dead time with 60 s lifetime. The electron beam was defocused at the condenser lens to maintain counting rates

below 1 kHz and lifetime efficiency >95%. The EDS patterns were recorded on an INCA 3.04 Microanalysis Suite and were digitized for analysis.

Confocal laser scanning microscopy

Samples were stained with the Live-Dead dye (L-7012, Molecular Probes, Eugene, OR, USA) for imaging on an inverted Zeiss LSM 510 confocal microscope in the Biological Imaging Center at Caltech, Pasadena, CA, USA.

RESULTS

Strain isolation and identification

Strain BoFeN1 was isolated from rusty-brown colonies that developed in the highest positive tubes in agar shake dilution series after 4–5 weeks of incubation. The number of colonies corresponded to a total of 10⁵–10⁶ colony-forming units per ml of acetate-plus-ferrous-iron-oxidizing nitrate reducers in the inoculum material. Numbers of lithotrophic, acetate-independent nitrate-reducing iron oxidizers were about two orders of magnitude lower.

16S rRNA gene sequence analysis of strain BoFeN1 (Fig. 1) showed that this organism is a β-Proteobacterium, and the closest relative to this strain by 16S rRNA comparison is *Acidovorax* sp. strain G8B1 (98.8% identity, 1388 nucleotides considered). A nitrate-dependent Fe(II)-oxidizing freshwater strain (strain BrG1) also affiliated with the genus *Acidovorax*, but isolated from a different environment, has been described earlier (Straub *et al.*, 2004).

Oxidation of Fe(II) by strain BoFeN1

Strain BoFeN1 grew heterotrophically by oxidizing acetate with nitrate as electron acceptor. In the presence of several mM both of acetate and of dissolved Fe(II), both the acetate and the ferrous iron were oxidized and Fe(III) minerals were produced (Fig. 2A,B). No dissolved Fe(III) could be detected in the cultures. In the absence of nitrate or in uninoculated control experiments, no oxidation of Fe(II) was observed (Fig. 2A). In the presence of both Fe(II) and acetate, acetate oxidation started and a significant lag phase preceded Fe(II) oxidation (Fig. 2B). Nitrite appeared at 1.0–1.5 mM concentrations but only after Fe(II) oxidation started and not as long as only acetate was oxidized; nitrite was not formed at all in the absence of Fe(II) when strain BoFeN1 oxidized acetate with nitrate as electron acceptor (data not shown). In cultures amended with dissolved Fe(II), acetate and nitrate, traces of N₂O but no NO could be detected in the headspace. Strain BoFeN1 was able to grow also with acetate as electron donor and NO₂⁻ as electron acceptor (data not shown).

Strain BoFeN1 needed an organic cosubstrate for fast and successive Fe(II) oxidation. At least 1 mM acetate was necessary to oxidize 4 mM dissolved Fe(II) completely to Fe(III)

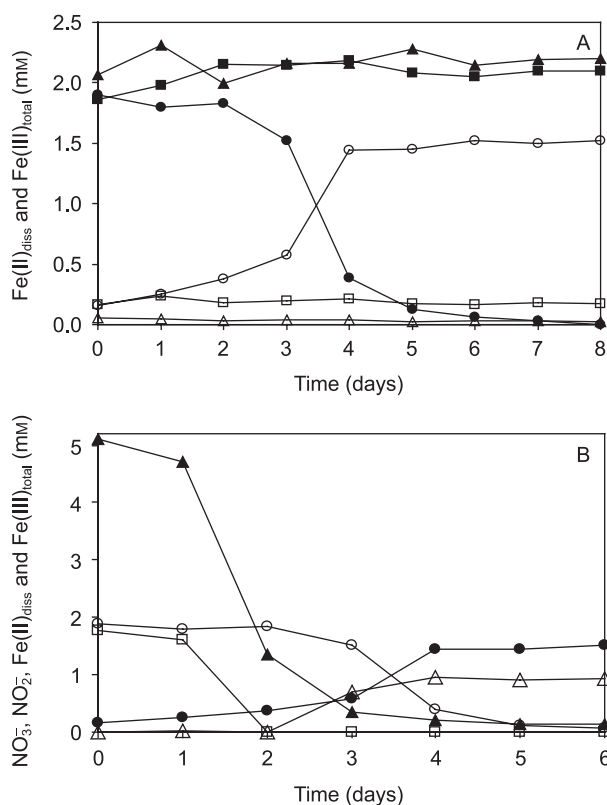


Fig. 2 Oxidation of dissolved Fe(II) by growing cultures of strain BoFeN1 (A) Representative graphs showing Fe(II) oxidation: dissolved Fe(II) (closed symbols) and Fe(III)_{total} (open symbols) in the presence of Fe(II), acetate, nitrate (●,○), in the absence of nitrate (□,■) and in uninoculated controls (▲,△). (B) Oxidation of acetate and dissolved Fe(II) and formation of nitrite by strain BoFeN1: acetate (□), Fe(II)_{diss} (○), Fe(III)_{total} (●) nitrate, NO₃⁻ (▲), nitrite, NO₂⁻ (△). All cultures were inoculated from an exponentially growing culture of strain BoFeN1 (2 mM Fe(II), 2 mM acetate and 5 mM NO₃⁻). Because of slightly different lag phases, the results from four independent experiments showing the same principle trend are not averaged. Instead, one representative data set is shown.

(Fig. 3). When we provided more acetate, the rate of Fe(II) oxidation increased slightly; omitting the acetate slowed down the oxidation rate and caused incomplete Fe(II) oxidation (Fig. 3). This effect was even more pronounced after a further transfer to medium containing Fe(II) but no acetate. After a second transfer, only negligible Fe(II) oxidation was observed at all.

Cell encrustation and cell viability

Confocal laser microscope images showed that bacteria grown with 4 mM dissolved Fe(II) plus 1 or 2 mM acetate have a thick crust of minerals around their cell membranes (Fig. 4). In 4 to 6-day-old cultures growing in the presence of Fe(II), a significant percentage of the cells were stained as 'dead cells' by the dead-live stain (Fig. 4). In contrast, 6-day-old cultures that were growing with acetate plus nitrate (in the absence of Fe(II)) showed almost 100% living cells (data not shown).

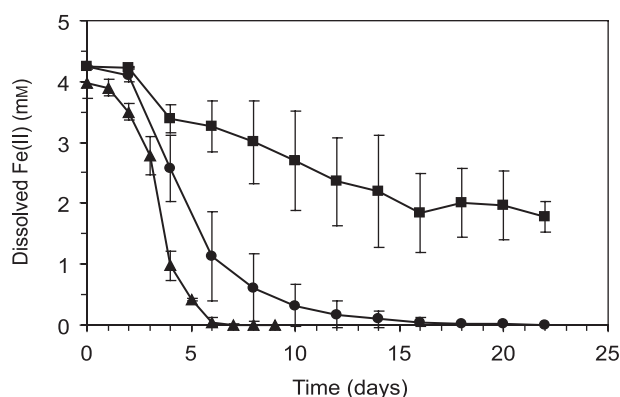


Fig. 3 Dependence of Fe(II) oxidation by strain BoFeN1 on acetate: oxidation of dissolved Fe(II) in the presence of 5 mM NO₃⁻ and either 2 mM (▲), 1 mM acetate (●) or in the absence of acetate (■). The cultures were inoculated from an exponentially growing BoFeN1 culture (4 mM dissolved Fe(II), 2 mM acetate and 5 mM NO₃⁻). The graphs shown each represent the mean of two independent cultures (± SD).

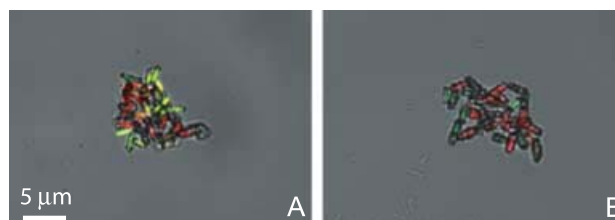


Fig. 4 Confocal microscopy pictures of 4-day-old dead-live stained cells of strain BoFeN1 (green = alive; red = dead) grown with dissolved Fe(II) (4 mM), nitrate (5 mM) and (A) 2 mM acetate and (B) 1 mM acetate showing the viability of BoFeN1 cells and different levels of cell-mineral encrustation.

Environmental scanning electron micrographs and light microscopic observations indicated that shortly after inoculation of Fe(II)/acetate/nitrate-containing medium (before Fe(II) oxidation started) many free-floating, non-encrusted cells were present (data not shown). Mineral crusts around the cells were formed as soon as Fe(II) oxidation started (Fig. 5A), and cells were completely encrusted after all Fe(II) present in the cultures was oxidized (Fig. 5B,C).

Fe(III) mineral formation and biomineralization

After examining the influence of acetate on iron(II) oxidation, we were interested in determining the effect of the presence of pre-existing Fe(III) minerals on the mineralogy of nascent crystals. This was carried out by following the mineralogy of Fe(III) minerals precipitated by subcultures incubated with acetate/nitrate plus Fe(II) and inoculated either from an acetate/nitrate or from an acetate/nitrate plus Fe(II) starter culture. X-ray diffraction analysis of the precipitates showed the presence of the crystalline Fe(III) mineral goethite in cultures inoculated from Fe-/acetate-grown cultures. The broad background peaks present in this spectrum suggest

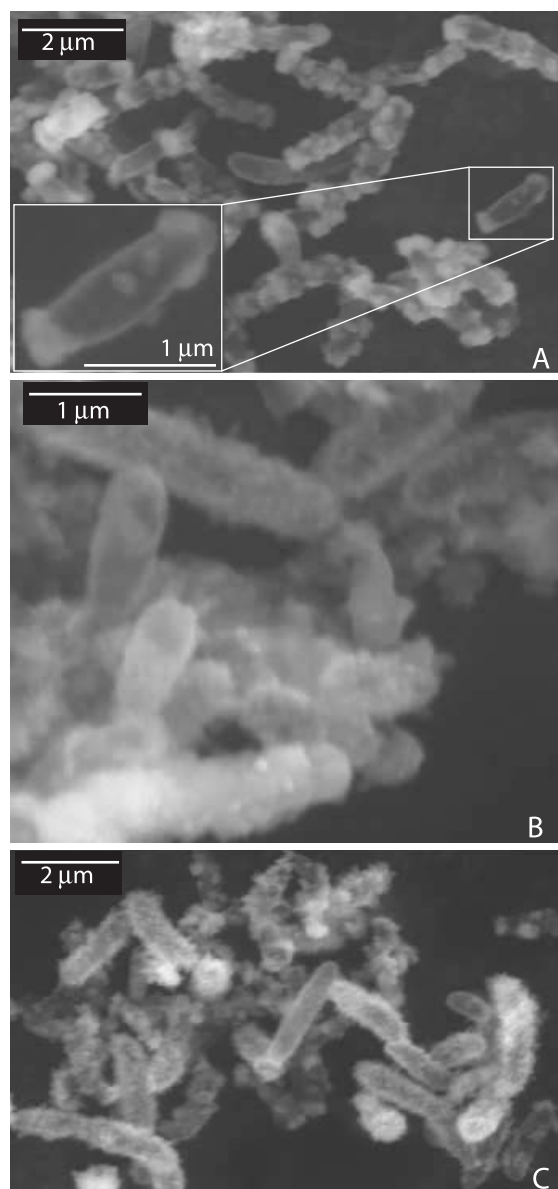


Fig. 5 Environmental scanning electron microscope photographs of cells of strain BoFeN1 grown with dissolved Fe(II) (4 mM), nitrate (5 mM) plus acetate (2 mM) and inoculated from a culture grown on the same substrates before. Samples were taken after 3.5 days (A), after 11 days (B), and after 30 days (C). The inset (A) shows a close-up of a cell with almost no mineral precipitates on the cell surface facilitating the comparison to the encrusted cells in Figs 5B,C.

that probably also poorly crystalline iron mineral phases are present. In contrast, inoculation with cells grown with acetate in the absence of Fe(II) led only to poorly crystalline ferric iron (hydr)oxides (Fig. 6). Within 65 days, we did not observe further transformation of the minerals to other Fe(III) minerals or a further increase in crystallinity. Energy-dispersive X-ray spectroscopy (coupled to scanning electron microscopy) of Fe minerals present in cultures inoculated with either acetate- or Fe/acetate-pregrown cells gave strong signals

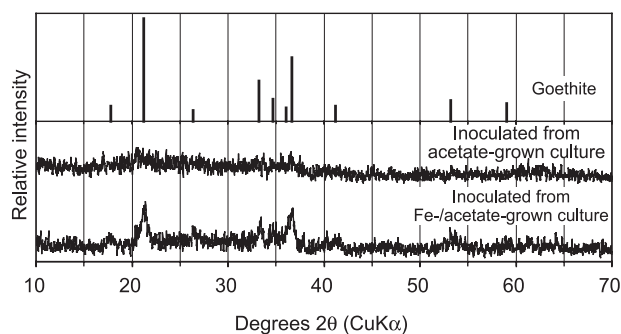


Fig. 6 X-ray diffractograms of precipitates collected from BoFeN1 cultures oxidizing dissolved Fe(II) when inoculated from a culture grown either with Fe(II), nitrate plus acetate ('Fe-/acetate-grown') or grown with nitrate plus acetate ('acetate-grown'). Reference data for goethite from the International Centre for Diffraction Data (ICDD) are included for comparison.

for iron and oxygen (data not shown). No significant signals for other elements such as phosphorus could be detected, confirming the presence of ferric iron (hydr)oxides as the main oxidation products. BET surface area analysis of poorly crystalline ferric iron (hydr)oxides obtained from a culture inoculated with an acetate-grown culture yielded a surface area of $158 \text{ m}^2 \text{ g}^{-1}$.

Scanning and transmission electron micrographs of cells from cultures inoculated with cells grown either with acetate or with Fe/acetate confirmed that the cell encrustation and the difference in crystallinity of the Fe(III) mineral product depended on the history of the inoculum. When cells grown without Fe(II) were transferred to medium containing Fe(II) plus acetate, the precipitates covering the cells were small, consisting of irregularly shaped plates and needles (Fig. 7A). In contrast, when cells grown in the presence of Fe(II) were transferred to fresh medium in which they again were oxidizing Fe(II), the precipitates around the cells were larger and more crystalline as suggested already by the XRD results (Fig. 7B,C and Fig. 8).

DISCUSSION

Enzymatically catalysed vs. nitrite-dependent, chemical Fe(II) oxidation

In cultures of the nitrate-reducing bacterial strain BoFeN1 amended with acetate plus nitrate, we observed oxidation of acetate with concomitant consumption of nitrate; nitrite was never observed in such cultures. However, if cultures were amended with Fe(II) in addition to acetate and nitrate, we observed not only oxidation of both Fe(II) and acetate but also concomitant accumulation of up to 1.5 mM nitrite. Because Fe(II) can be oxidized also chemically by nitrite (Moraghan & Buresh, 1977), one of the initial questions was whether the observed Fe(II) oxidation was catalysed enzymatically or whether Fe(II) was oxidized chemically by nitrite produced during acetate oxidation. Because nitrite was

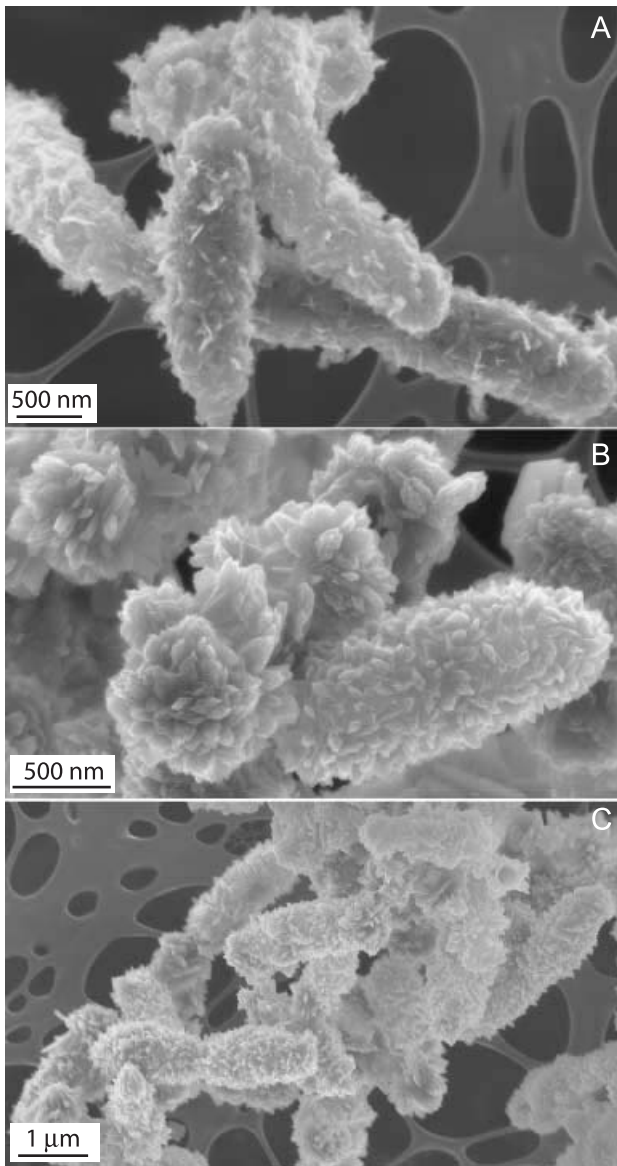


Fig. 7 Scanning electron micrographs of Fe(III)-precipitates produced by strain BoFeN1 when inoculated from a culture (A) grown with acetate (2 mM) plus nitrate (5 mM) and (B–C) grown with acetate (2 mM) plus nitrate (5 mM) plus dissolved Fe(II) (4 mM).

never formed in the absence of Fe(II), this suggests that nitrite formation is a consequence of, but not the reason for, Fe(II) oxidation. The observed formation of nitrite during nitrate-dependent Fe(II) oxidation but not during nitrate-dependent acetate oxidation might be a consequence of the entry of electrons from the redox couples $\text{Fe}(\text{OH})_3/\text{Fe}^{2+}$ and $\text{CO}_2/\text{acetate}$ at different standard redox potentials (~ 0 mV vs. -290 mV; Thauer *et al.*, 1977; Thamdrup, 2000). Entry of electrons at different redox potential levels could change the relative rates of the enzymatically catalysed reductions of the different nitrogen species, leading to the accumulation of intermediates such as the nitrite.

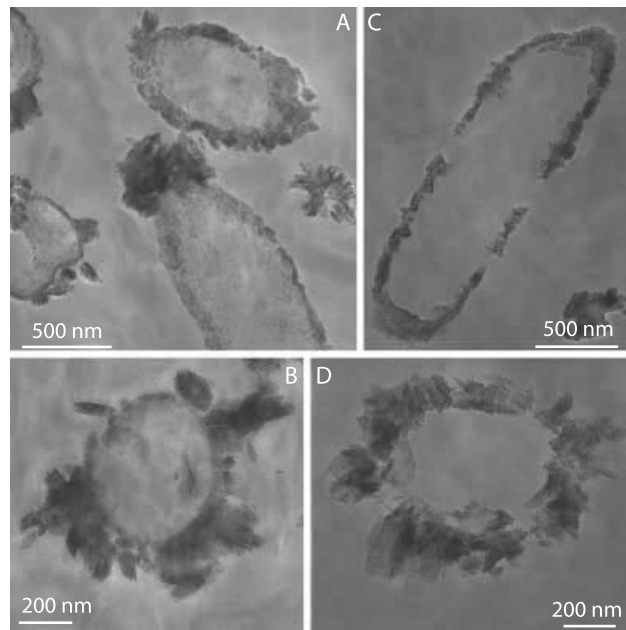


Fig. 8 Transmission electron micrographs of stained (A,B) and nonstained (C,D) thin sections of cells of strain BoFeN1 grown with dissolved Fe(II) (4 mM), nitrate (5 mM) and acetate (2 mM), and inoculated with a culture pregrown on the same substrates. Samples were collected after complete oxidation of Fe(II).

Comparing the rates of Fe(II) oxidation observed in BoFeN1 cultures with rates of chemical Fe(II) oxidation with nitrite also supports our view that Fe(II) is oxidized in an enzymatic reaction: in sterile water amended with 0.5 mM nitrite and 15 mM Fe(II) at pH 6.0, about ~ 0.07 mM nitrite and at pH 8 all nitrite disappeared within 24 h (Moraghan & Buresh, 1977). The growth medium we used for cultivation of nitrate-dependent Fe(II)-oxidizing bacteria contains several mM concentrations of different mineral salts such as sodium bicarbonate and calcium chloride as well as low concentrations of vitamins and trace elements that probably influence the oxidation extent and the oxidation rates. Such growth medium showed no significant Fe(II) oxidation with nitrite concentrations of ~ 1 mM; only with nitrite concentrations of 5 mM, Fe(II) is oxidized chemically at a rate of $\sim 8 \mu\text{M h}^{-1}$ (Benz *et al.*, 1998), and with 2 mM nitrite at $\sim 7 \mu\text{M h}^{-1}$, including a fast initial oxidation of $\sim 40 \mu\text{M h}^{-1}$ during the first 24 h (Straub *et al.*, 1996), which still cannot explain the Fe(II) oxidation rates we observed in our experiments ($50\text{--}75 \mu\text{M h}^{-1}$). We therefore conclude that the oxidation of Fe(II) in our experiments is catalysed enzymatically, and is not due to an abiotic reaction of nitrite with Fe(II).

Beneficial and harmful effects of Fe(III) mineral precipitation

Strain BoFeN1 oxidizes Fe(II) mixotrophically in the presence of acetate and nitrate, forming crusts of Fe(III) minerals around the cells. This is different from the nitrate-dependent

Fe(II)-oxidizing cells described by Benz *et al.* (1998) who saw cells associated with the Fe(III) particles but not encrusted with minerals. However, these authors did not provide electron micrographs, and light microscopy could be misleading because cells that were totally encrusted and thus hidden in the minerals could have been overlooked. Moreover, these experiments were performed with brackish water medium with about 1% salt content, and the higher concentration of counter ions might have influenced the ferric oxide precipitation. Neutrophilic chemolithoautotrophic Fe(II)-oxidizers that use molecular oxygen instead of nitrate as electron acceptor were isolated from the deep sea and from a pH-neutral groundwater and were also shown to be associated but not encrusted with Fe minerals after Fe(II) oxidation (Emerson & Moyer, 1997; Edwards *et al.*, 2003).

It is unclear whether precipitation of Fe(III) minerals in close vicinity to the cells, or even at the cell surface, is harmful to the cells by virtue of limiting substrate diffusion and uptake as is commonly assumed (Hanert, 1981; Hallberg & Ferris, 2004). At first glance, it would appear to make sense for the cell to avoid cell encrustation by locating the Fe(III) mineral precipitates distant to its surface because encrustation might limit substrate uptake. Evidence in support of this comes from a study with *Acinetobacter* spp., where a reduced aerobic respiration rate was observed for the cells when they were encrusted with iron (MacRae & Celo, 1975). Production of extracellular organic material that functions to adsorb Fe(III) and nucleate precipitation (as is known for the aerobic Fe(II) oxidizers *Gallionella ferruginea* or *Leptothrix ochracea*, Hanert, 1981; Hallberg & Ferris, 2004) could be one strategy to avoid cell encrustation. An alternative strategy could be the release of dissolved Fe(III), either complexed by organic ligands or in a colloidal state, that diffuses away and precipitates at a distance to the cell. This was suggested for microaerobic Fe(II) oxidizers that produced several mM dissolved or colloidal Fe(III) (Sobolev & Roden, 2001 and 2002). A low-pH microenvironment as observed for phototrophic and microaerophilic Fe(II) oxidizers (Kappler & Newman, 2004; Roden *et al.*, 2004; D. Emerson, personal communication) could facilitate the release of such dissolved iron species.

Neither electron nor light microscopy of cultures of strain BoFeN1 gave any indication for the involvement of extracellular organic material. We were also never able to detect dissolved Fe(III) in Fe(II)-oxidizing cultures of strain BoFeN1. That strain BoFeN1 obviously can overcome encrustation, and grow when fresh medium with nitrate, acetate and Fe(II) is provided compels us to reconsider whether encrustation is as harmful as generally assumed. From a thermodynamic perspective, the formation of Fe(OH)₃ in close proximity to the cells, compared to the release and loss of Fe(III) as dissolved species, e.g. Fe³⁺, may even be favourable. Calculating the Gibbs free energy for nitrate-dependent Fe(II) oxidation with either Fe³⁺ or Fe(OH)₃ as the product (pH 7, 4 mM Fe²⁺, 10 mM NO₃⁻ and 10 μM Fe³⁺ in the case of Fe³⁺ as the product)

yields a significantly lower value when Fe(III) is released as Fe³⁺ (-54.6 kJ mol⁻¹ NO₃⁻) compared to nitrate-dependent Fe(II) oxidation with precipitation of Fe(OH)₃ (-381.3 kJ mol⁻¹ NO₃⁻). Future experiments determining growth yield and the encrustation of cells grown in chemostat experiments at limiting supply of Fe(II) and acetate could help answer the question whether Fe(II) oxidation in general and encrustation in particular are harmful or beneficial for the cells.

Our results suggest that acetate is oxidized (at least partially) before Fe(II) oxidation starts. This could help the cells to overcome cell encrustation and could lead initially to non-encrusted cells in the cultures that get encrusted concomitant with the start of Fe(II) oxidation. A preference for acetate over Fe(II) was observed also (i) with other nitrate-dependent Fe(II) oxidizers by Benz *et al.* (1998) who saw fast depletion of acetate in the beginning (and significant increase of protein content) followed by slow Fe(II) oxidation (and slow increase in protein content) and (ii) for *Azospira oryzae* (formerly known as *Dechlorosoma suillum*), which can oxidize Fe(II) mixotrophically with nitrate as electron acceptor (Chaudhuri *et al.*, 2001; Tan & Reinhold-Hurek, 2003). The cessation of Fe(II) oxidation in the absence of acetate, the ability to overcome cell encrustation in the presence of acetate and the decreasing cell viability during mineral crust formation (observed by confocal laser microscopy of dead-live stained cells) show that continuous and extensive Fe(II) oxidation by mixotrophic Fe(II) oxidizers such as strain BoFeN1 depends on an organic cosubstrate.

Mechanism of Fe(III) mineral formation

Looking at chemical Fe(II) oxidation experiments (e.g. Cornell *et al.*, 1989) might help to better understand the mechanism of Fe(III) mineral precipitation following microbially catalysed Fe(II) oxidation. In abiotic Fe(II) oxidation, initially, mono- and dinuclear species [FeOH]²⁺ and [Fe₂(OH)₂]⁴⁺ are formed that first transform to colloidal polymers and poorly ordered ferric hydroxides (ferrihydrite, aggregates of spherical particles of ~2–5 nm diameter), and then either transform by internal rearrangement to haematite (Fe₂O₃) (solid-state conversion) or by decomposition of the polymers into low molecular species that transform further to crystalline forms such as goethite ('dissolution–reprecipitation mechanism'). A low-pH microenvironment as observed for phototrophic and microaerophilic Fe(II) oxidizers (Kappler & Newman, 2004; Roden *et al.*, 2004) could not only facilitate the release of dissolved iron species but also could catalyse the fast transformation of ferrihydrite to goethite that is possible only via a dissolution–reprecipitation mechanism. Additionally, the fast formation of goethite concomitant with the formation of a thick crust around the cells may be controlled also by an autocatalytic process: mineral-surface-OH groups are electron-donor ligands that increase the electron density at the metal centre, thus stabilizing Fe(III) vs. Fe(II) and leading to

faster oxidation of Fe(II) (Wehrli *et al.*, 1989). This effect is much stronger with goethite than with ferrihydrite (Elsner *et al.*, 2003). Thus, the formation of goethite at the cell surface and binding of Fe(II) to the goethite surface would increase the electron density at the bound Fe(II), thus catalysing and increasing the rate of Fe(II) oxidation that may work via electron transfer through the mineral crust to the cell surface where the electrons could be accepted by the cell. Electron transfer from adsorbed Fe(II) to the bulk iron(III) oxide inducing growth of an Fe(III) layer on the oxide surface would abolish the need for the Fe(II)-oxidizing microbe to be in direct contact with the dissolved Fe(II). This idea is supported by a study that demonstrated electron transfer between Fe(II) adsorbed to Fe(III) oxides and the underlying iron(III) oxide (Williams & Scherer, 2004).

The observed formation of goethite by strain BoFeN1 differs from previous studies (Straub *et al.*, 1996, 1998, 2004; Benz *et al.*, 1998) in which the formation of poorly crystalline ferrihydrite by nitrate-dependent Fe(II) oxidizers that were transferred continuously on medium containing Fe(II) was described. While it is possible that those strains used different mechanisms for Fe(II) oxidation and Fe(III) release, the slightly different medium composition and different concentration ratios of acetate:Fe(II):nitrate may also have influenced the mineral formation. The fact that goethite formed in our experiments only in cultures inoculated with Fe-grown cultures suggests that not only a potential low-pH microenvironment but also the presence of minerals in the inoculum that might act as nucleation sites is involved in the formation of the more crystalline Fe(III) mineral goethite. A recent study on the effect of oxidation rate on microbial nitrate-dependent Fe(III) mineral formation by Senko *et al.* (2005) showed that low initial Fe(II) oxidation rates can also lead to the formation of more crystalline Fe(III) minerals.

Chemical oxidation of Fe(II) by molecular oxygen (O₂) showed that the mineralogy of the produced Fe(III) oxides is affected by the presence of dissolved compounds such as bicarbonate, phosphate and silica (Cornell & Schwertmann, 2003), whereas the presence of nonmetabolizing cells had no effect on the mineralogy but influenced the size, the spatial organization and the morphology of the particles (Chatellier *et al.*, 2001, 2004). Other authors found that in the presence of bacterial cells (*Bacillus subtilis*, *Bacillus licheniformis*), the formation of crystalline Fe(III) minerals was hindered and ferrihydrite formed instead (Mavrocordatos & Fortin, 2002). These chemical oxidation experiments in the presence of nonmetabolizing cells as well as experiments of fast chemical Fe(II) oxidation (Cornell & Schwertmann, 2003) favour the production of poorly crystalline Fe(III) oxides. The formation of goethite by strain BoFeN1 and the observed variations in mineral formation observed for different nitrate-dependent Fe(II) oxidizers show that not only medium composition, concentrations of cosubstrates and incubation conditions but also Fe(II) oxidation rates and presence of nucleation

sites influence and may even control the mineralogy of the Fe(III) phases produced during microbially catalysed Fe(II) oxidation.

Ecological and environmental implications of Fe(III) mineral formation by nitrate-dependent Fe(II) oxidation

The formation of poorly crystalline Fe(III) minerals such as ferrihydrite or higher crystalline Fe(III) minerals such as goethite by Fe(II)-oxidizing bacteria has important ecological and environmental consequences. The less crystalline Fe(III) minerals that are produced by some nitrate-dependent Fe(II)-oxidizing bacteria were shown to represent an excellent substrate for Fe(III)-reducing bacteria (Straub *et al.*, 1998), and Fe cycling by coupling microaerobic or nitrate-dependent Fe(II) oxidizers to Fe(III)-reducing bacteria was demonstrated (Sobolev & Roden, 2002; Straub *et al.*, 2004). Because more crystalline Fe(III) minerals are difficult to reduce by Fe(III)-reducing bacteria (Lovley *et al.*, 2004), the production of crystalline Fe(III) minerals might prevent microbial Fe(III) reduction and might be the reason for the presence of crystalline Fe(III) minerals in anoxic environments (Cornell & Schwertmann, 2003).

Understanding the factors that influence or even control the iron cycle is important because redox reactions involving Fe play a major role for the fate of organic and inorganic pollutants in our environment. First, Fe(III)-reducing microbes can couple the reduction of Fe(III) to the oxidation of organic pollutants such as aromatic hydrocarbons (see review by Lovley & Anderson, 2000), and the presence of more crystalline and less reducible Fe(III) minerals may restrict or even hinder such degradation. Second, Fe(III) mineral precipitation can lead to coprecipitation of toxic metal ions, e.g. arsenic (Fuller *et al.*, 1993). Coprecipitation and thus immobilization of arsenic with less or nonreducible Fe(III) minerals would be particularly desirable to avoid re-reduction of the Fe(III) that would be coupled to a release of the toxic metal ions. Third, the surfaces of Fe(III) (hydr)oxides adsorb nutrients such as phosphate and metal ions such as arsenate/arsenite (Jenne, 1968; Hering, 1995). Ferric iron minerals of higher crystallinity have a significantly lower surface area (up to 50-fold lower than the poorly crystalline ferrihydrite, Cornell & Schwertmann, 2003) and thus a lower capacity of binding e.g. phosphate and arsenic. Reduction of nutrient- or metal-loaded ferrihydrite coupled to reoxidation producing goethite would consequently release a significant amount of either nutrients or toxic metal ions.

Knowing the mechanisms and controlling factors of anaerobic Fe(II) oxidation is important not only for understanding present-day biogeochemical cycling of iron. Precambrian oceans contained considerable amounts of dissolved ferrous iron, and at least in the beginning of that period almost no molecular oxygen (O₂) was present as a terminal electron acceptor for microbial Fe(II) oxidation. This raises the question

how the Fe(II) was oxidized and precipitated in ancient Banded Iron Formations. Besides phototrophic Fe(II) oxidation, nitrate-dependent oxidation of Fe(II) so far represents the only known microbially catalysed mechanism to oxidize Fe(II) in the absence of O₂. Moreover, it presents a mechanism for Fe(II) oxidation even in dark environments. Modelling studies aiming at understanding the relative contributions of diverse processes to Fe(II) oxidation in the past will benefit from including nitrate-dependent Fe(II) oxidation, in addition to anoxygenic phototrophs (Widdel *et al.*, 1993; Kappler *et al.*, 2005) and neutrophilic, O₂-dependent Fe(II) oxidizers (Emerson & Moyer, 1997; Edwards *et al.*, 2003).

In summary, mixotrophic nitrate-dependent Fe(II)-oxidizing bacteria oxidize Fe(II) with different rates and encrust themselves with different Fe(III) minerals, depending on the conditions and in particular depending on the presence of an organic cosubstrate. The identity of the produced Fe(III) minerals controls iron cycling and the fate of essential nutrients as well as the fate of harmful pollutants. Therefore, a thorough understanding of the factors controlling the mineralogy in this microbially catalysed process is the key for future studies.

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