

# *Geothrix fermentans* Secretes Two Different Redox-Active Compounds To Utilize Electron Acceptors across a Wide Range of Redox Potentials

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The current understanding of dissimilatory metal reduction is based primarily on isolates from the proteobacterial genera *Geobacter* and *Shewanella*. However, environments undergoing active Fe(III) reduction often harbor less-well-studied phyla that are equally abundant. In this work, electrochemical techniques were used to analyze respiratory electron transfer by the only known Fe(III)-reducing representative of the *Acidobacteria*, *Geothrix fermentans*. In contrast to previously characterized metal-reducing bacteria, which typically reach maximal rates of respiration at electron acceptor potentials of 0 V versus standard hydrogen electrode (SHE), *G. fermentans* required potentials as high as 0.55 V to respire at its maximum rate. In addition, *G. fermentans* secreted two different soluble redox-active electron shuttles with separate redox potentials (−0.2 V and 0.3 V). The compound with the lower midpoint potential, responsible for 20 to 30% of electron transfer activity, was riboflavin. The behavior of the higher-potential compound was consistent with hydrophilic UV-fluorescent molecules previously found in *G. fermentans* supernatants. Both electron shuttles were also produced when cultures were grown with Fe(III), but not when fumarate was the electron acceptor. This study reveals that *Geothrix* is able to take advantage of higher-redox-potential environments, demonstrates that secretion of flavin-based shuttles is not confined to *Shewanella*, and points to the existence of high-potential-redox-active compounds involved in extracellular electron transfer. Based on differences between the respiratory strategies of *Geothrix* and *Geobacter*, these two groups of bacteria could exist in distinctive environmental niches defined by redox potential.

While most of our understanding regarding electron transfer to metals is derived from a small number of *Shewanella* and *Geobacter* species, a diversity of *Bacteria* and *Archaea* are known to use insoluble Fe(III) oxides as terminal electron acceptors (19, 29, 43, 50). In particular, sequences similar to those for an Fe(III)-reducing isolate from the *Acidobacteria* phylum (sometimes referred to as group 8 *Acidobacteria*) are consistently detected in 16S rRNA surveys of aquifers undergoing Fe(III) reduction, as well as in Fe(III)-rich sediments undergoing uranium bioreduction (2, 8, 59), petroleum oxidation (1), technetium reduction (40), and arsenic reduction (21, 22). In many of these environments, such acidobacterial sequences are more abundant than *Geobacter* sequences, yet the basis for their competitiveness with *Geobacter* is not known.

In general, members of the *Acidobacteria* are ubiquitous in both 16S and metagenomic surveys of soils, often representing over 50% of detected bacterial sequences (15, 23, 25, 45, 49) which are classified into 26 acidobacterial subdivisions (2, 7, 14, 17, 18, 34, 58). Cultivated *Acidobacteria* are slow-growing, nutritionally versatile oligotrophs able to use a variety of carbon sources and that typically prefer microaerophilic conditions. The exception is the Fe(III)-reducing anaerobic isolate *Geothrix fermentans*, originally obtained from the Fe(III)-reducing zone of a petroleum-contaminated aquifer (10).

To transfer electrons to extracellular compounds, bacteria must make contact with electron acceptors and/or secrete redox-active mediators. Both strategies have their advantages: flavin mononucleotide (FMN) actively secreted by *Shewanella* accelerates metal reduction (12, 37, 47, 54), and phenazines secreted by *Pseudomonas* facilitate movement of electrons to aerobic edges of a biofilm (13, 20, 56, 57). Direct mechanisms, such as those used by *Geobacter* (29, 41, 52), benefit only cells in contact with the

acceptor and may also electrically link daughter cells to the metal, often allowing electron transfer over longer distances than soluble shuttles (52). Each strategy has potential tradeoffs; a contact-based strategy may represent a metabolic burden in terms of multiple structural and extracellular redox proteins to interface with unpredictable metal surfaces in the environment, while secreted molecules could be expensive to produce and be limited by diffusion.

*G. fermentans* reduces Fe(III) trapped within porous beads (42), and supernatants from *G. fermentans* cultures accelerate Fe(III) reduction by other bacteria (4), providing evidence that this bacterium may secrete a soluble compound as part of its metal reduction strategy. Both quinone-like compounds and metal chelators have been reported to occur in *G. fermentans* supernatants. However, the identity of any soluble agents, their redox potentials, and their relative importance to electron transfer under conditions relevant to subsurface growth remain unknown (42).

Within the last 5 years, a new set of electrochemical tools has emerged that allows monitoring of electron transfer from living cells. Compared to growth with insoluble metals, electrode potentials can be precisely controlled, allowing rates of electron transfer from cells to be measured without accumulation of reduced end products confounding results (36, 38). Such analyses show that

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*Geobacter* strains transfer electrons beyond the cell membrane at potentials as low as  $-0.2$  V versus the standard hydrogen electrode (SHE) and reach their maximal rate of respiration at  $-0.1$  V. These low values help explain the competitiveness of *Geobacter* in permanently anoxic environments. In *Shewanella* strains, direct electron transfer to electrodes can be measured, but the ability to easily remove the medium surrounding attached cells reveals that an indirect redox shuttling mechanism based on secreted FMN is responsible for nearly  $\sim 80\%$  of respiratory activity by biofilms under these conditions (37).

In this work, the electron transfer strategy of *G. fermentans* was studied for the first time using poised electrodes and was found to be different from that of any previously described organism. Specifically, *G. fermentans* electron transfer reached less than 30% of maximum rates between the commonly studied redox potential window of  $-0.2$  and  $0$  V and instead required potentials above  $0.55$  V to support maximal rates. Removal of supernatants from electrode-grown biofilms reduced the rate of electron transfer, and cell-free supernatants from both Fe(III)-grown and electrode-grown *G. fermentans* cultures showed evidence of two independent soluble electron shuttles. Neither of these activities appeared to be due to a metal chelator. One of the two redox shuttles, active in the low-potential window of  $-0.2$  V, was identified as riboflavin, while the higher-potential compound remains unidentified. This dual redox potential strategy reveals that flavin-based electron shuttles are used by Fe(III)-reducing bacteria outside the *Proteobacteria* and suggests that anaerobic *Acidobacteria* may be more competitive in higher-potential environments, such as near oxic-anoxic interfaces.

## MATERIALS AND METHODS

**Bacterial strains and culture media.** *Geothrix fermentans* (ATCC 700665) was grown anaerobically at  $20^\circ\text{C}$  unless otherwise indicated in minimal medium containing the following (per liter):  $0.38$  g KCl,  $0.2$  g  $\text{NH}_4\text{Cl}$ ,  $0.069$  g  $\text{NaH}_2\text{PO}_4 \cdot \text{H}_2\text{O}$ ,  $0.04$  g  $\text{CaCl}_2 \cdot 2\text{H}_2\text{O}$ , and  $0.2$  g  $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$ . Ten milliliters of mineral mix containing  $0.1$  g  $\text{MnCl}_2 \cdot 4\text{H}_2\text{O}$ ,  $0.3$  g  $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ ,  $0.17$  g  $\text{CoCl}_2 \cdot 6\text{H}_2\text{O}$ ,  $0.1$  g  $\text{ZnCl}_2$ ,  $0.04$  g  $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ ,  $0.005$  g  $\text{AlK}(\text{SO}_4)_2 \cdot 12\text{H}_2\text{O}$ ,  $0.005$  g  $\text{H}_3\text{BO}_3$ ,  $0.09$  g  $\text{Na}_2\text{MoO}_4$ ,  $0.12$  g  $\text{NiCl}_2$ ,  $0.02$  g  $\text{NaWO}_4 \cdot 2\text{H}_2\text{O}$ , and  $0.10$  g  $\text{Na}_2\text{SeO}_4$  per liter was added before autoclaving. Lactate at  $20$  mM was supplied as the electron donor and  $40$  mM fumarate as the soluble electron acceptor for routine growth, and  $100$  mM ferrihydrite ( $\text{FeOOH}$ ) was added for metal-reducing conditions. Lactate at  $20$  mM as an electron donor and a poised electrode as an electron acceptor were used for all electrochemical experiments. All media were adjusted to pH 6.8, buffered with  $2$  g/liter  $\text{NaHCO}_3$ , made anaerobic by being flushed with oxygen-free  $\text{N}_2/\text{CO}_2$  (80:20, vol/vol), and sealed with butyl rubber stoppers prior to autoclaving. One milliliter of a filter-sterilized anaerobic vitamin mixture was added to all cultures before inoculation. The vitamin mixture contained  $0.002$  g biotin,  $0.002$  g folic acid,  $0.01$  g pyridoxine HCl,  $0.005$  g thiamine,  $0.005$  g nicotinic acid,  $0.005$  g pantothenic acid,  $0.0001$  g vitamin  $\text{B}_{12}$ ,  $0.005$  g *p*-aminobenzoic acid, and  $0.005$  g thioctic acid per 1 liter.

**Electrode preparation and assembly.** Carbon cloth (B-1/B WO W000179 E-TER; PEMEAS Fuel Cell Technologies/BASF, Somerset, NJ) was cut into 2- by 2-cm electrodes (geometric surface area,  $8\text{ cm}^2$ ). Carbon cloth electrodes were soaked sequentially in deionized water,  $0.1$  N NaOH, deionized water,  $0.1$  N HCl, and ethanol to remove contaminants. This treatment was repeated, and electrodes were stored in deionized water. Electrodes were attached to  $0.1$ -mm platinum wire. A bioreactor, containing a carbon cloth working electrode, a platinum counter-electrode, and a saturated calomel reference electrode, was prepared as described previously (36). Sterile anaerobic minimal medium (5 ml) lacking electron donors or acceptors was added to autoclaved bioreactors, and

anaerobic conditions were maintained by a constant flow of humidified oxygen-free  $\text{N}_2/\text{CO}_2$  (80:20, vol/vol) at  $20^\circ\text{C}$  and connected to a potentiostat (VMP; Bio-Logic, Knoxville, TN) (36). After equilibration, 5 ml of a culture entering stationary phase (optical density at  $600\text{ nm}$  [ $\text{OD}_{600}$ ], 0.5) was added to the electrochemical cell (a 50% inoculum).

**Electrochemical analysis.** The parameters for cyclic voltammetry of attached biofilms were as follows: equilibrium time =  $5$  s, scan rate =  $1$  mV/s,  $E_{\text{initial}}$  ( $E_i$ ) =  $-0.645$  V versus SHE, and  $E_{\text{final}}$  ( $E_f$ ) =  $0.745$  V versus SHE; the current was averaged over the last 80% of each poised voltage as described previously by Marsili et al. (36). Two sweeps were performed, with data from the second sweep reported. By changing the potential slowly ( $1$  mV/s), data were collected across a range of applied redox potentials, reflecting steady-state electron transfer from cells to electrodes.

Analytical electrochemistry was also performed on supernatants in duplicate between an  $E_i$  of  $-0.645$  V versus SHE and an  $E_f$  of  $0.745$  V versus SHE at a scan rate of  $100$  mV/s. This higher scan rate was chosen to rapidly oxidize or reduce redox active compounds near the electrode and enhance the sensitivity of detection. Data collected from the second sweeps were analyzed and processed using SOAS (16).

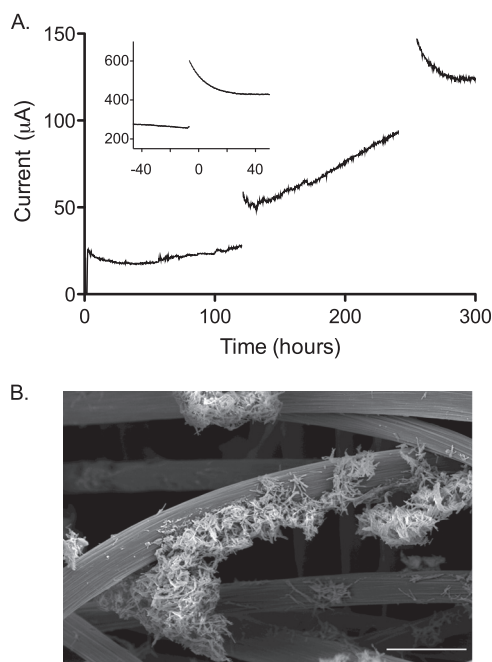
**Biomass measurements.** Biomass was determined using the bicinchoninic acid protein assay (Thermo Scientific, Rockford, IL). After electrochemical analysis, electrodes were harvested and gently washed in minimal medium to remove loosely attached planktonic cells. The electrodes were disassembled and incubated in  $1$  ml of  $0.2$  M NaOH solution at  $96^\circ\text{C}$  for  $20$  min to remove attached biomass. Electrodes were also harvested, fixed in 3% glutaraldehyde, and dehydrated in increasing concentrations of ethanol (25%, 50%, 75%, 95%, and 100%) for  $10$  min. The electrodes were critical-point dried, sputter coated, and imaged using a S3500N scanning electron microscope (Hitachi, Japan).

**Collection of culture supernatants.** Electrode-grown supernatants were harvested by carefully removing the surrounding medium into a sterile anaerobic tube with an 80%  $\text{N}_2$ –20%  $\text{CO}_2$  headspace, centrifuged anaerobically to remove planktonic cells, and stored at  $-20^\circ\text{C}$  after the headspace was flushed with 80%  $\text{N}_2$ –20%  $\text{CO}_2$ . Supernatants collected from FeOOH-grown cultures were centrifuged anaerobically at  $1,500$  rpm to remove insoluble FeOOH. The supernatant was anaerobically mixed with  $0.5$  g of Chelex 100 resin (Bio-Rad, Hercules, CA) per  $10$  ml of supernatant in a sterile anaerobic tube overnight at  $4^\circ\text{C}$  on a shaker. The supernatant was centrifuged to remove resin and stored in a sterile anaerobic tube or passed through an octadecyl-functionalized silica gel ( $\text{C}_{18}$ ) column anaerobically in a glove box. The effluent was stored at  $-20^\circ\text{C}$  in a sterile anaerobic tube. Hydrophobic compounds bound to the  $\text{C}_{18}$  column were collected by adding 100% methanol and stored at  $-20^\circ\text{C}$  for further analysis.

**HPLC.** Cell-free electrode- and FeOOH-grown supernatants were analyzed by high-performance liquid chromatography (HPLC) using a protocol previously described (37). An Eclipse XDB-C18 column ( $4.6$  mm by  $525$  mm and  $5$ - $\mu\text{m}$  particle size) from Agilent Technologies, Santa Clara, CA, was used. A fluorescence detector with an excitation wavelength of  $440$  nm and an emission wavelength of  $525$  nm was used to detect fluorescent compounds.

## RESULTS

**Conditions to minimize cell lysis.** *G. fermentans* culture supernatants have previously been reported to contain soluble factors that accelerate reduction of ferrihydrite (42). While some bacteria actively excrete redox shuttles (12, 37, 54, 57), an alternative explanation is always the possibility of cell lysis releasing intracellular compounds. When *G. fermentans* cultures were grown at  $30^\circ\text{C}$ , over 50% of cells lysed within 2 days of reaching stationary phase. Reducing the growth temperature to  $25^\circ\text{C}$  slowed this process, and when cells were grown at  $20^\circ\text{C}$ , the  $\text{OD}_{600}$  remained stable after more than  $200$  h of incubation in stationary phase (see Fig. S1 in the supplemental material). While cell turnover during pro-



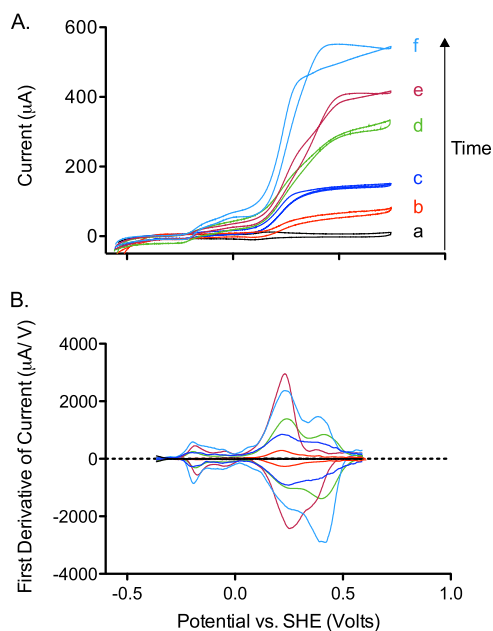
**FIG 1** Typical increase in electron transfer rate by electrode-attached *G. fermentans*. (A) Carbon fiber electrodes (8-cm<sup>2</sup> geometric surface area) were poised at 0.55 V versus SHE and paused every 120 h for voltammetry analysis. The inset shows the increase in capacitance behavior over time after pausing current collection of a 500-h biofilm. (B) SEM image of a *G. fermentans* biofilm on a carbon fiber electrode, 1 month after inoculation, showing sparse coverage and microcolony formation. The scale bar represents 25 µm.

longed incubations is a complex process, growth at 20°C was chosen for all further electrochemical studies to minimize the possibility of compounds being rapidly released that could create significant artifacts (27).

**Colonization of electrodes by *G. fermentans* requires higher potentials.** Experiments were initially conducted using carbon fiber electrodes poised at 0.2 V versus SHE, as these redox potentials are typically used to study metal-reducing *Geobacter* and *Shewanella* (3, 36, 37, 38). After 2 weeks of incubation, a small increase in electron transfer was detected, but it rarely exceeded ~5 µA/cm<sup>2</sup> under these conditions. Voltammetry of these partially colonized electrodes revealed that the rate of electron transfer increased significantly when the electrode was raised to higher potentials. Based on these data, new growth experiments were initiated with electrodes poised at 0.55 V versus SHE.

When the electrode was poised at a higher potential, a 50% inoculum of cells demonstrated a rapid initial rise in electron transfer rate (Fig. 1A), followed by a slow increase over the next 300 to 400 h. Inoculated bioreactors required >2 weeks to reach stable rates of electron transfer to electrodes. For this study, 15 independent biofilms were cultivated at higher potential (0.55 V), analyzed electrochemically, and used to produce supernatants for further analysis.

During colonization, data collection was paused at regular intervals for electrochemical analysis. For example, sweeping slowly between low and high potentials at 1 mV/s allowed measurement of electron transfer rates from bacteria to the electrode across a wide range of electron acceptor redox potentials. After electrodes were returned to the standard potential for growth (0.55 V), tem-



**FIG 2** (A) Cyclic voltammometry (1 mV/s) of *G. fermentans* biofilms collected over time at 100 (b), 240 (c), 360 (d), 480 (e), and 640 (f) h of growth on carbon fiber electrodes. A voltammogram of the uninoculated carbon fiber electrode is shown in black (a). (B) First derivatives of cyclic voltammograms from panel A, facilitating visualization of midpoint potentials in catalytic waves.

porary surges of electron transfer from the biofilm were typically observed. In younger cultures, this surge decayed within 5 h, but up to 30 h was needed to fully discharge older cultures. Representative data from early and late stages of growth, illustrating this “capacitance” phenomenon where excess electrons were stored during pauses in current collection, are shown in Fig. 1A.

Routine *G. fermentans* biofilms allowed to grow over 640 h at 0.55 V achieved a density of 50 to 100 µA/cm<sup>2</sup>, a value 10-fold lower than what is observed for *Geobacter* strains, even when electrode roughness is considered. When attached biomass was measured, rates of electron transfer averaged 0.6 mA/mg protein (range = 0.3 to 1.1 mA/mg; higher values from older cultures), a range also an order of magnitude lower than that of *Geobacter sulfurreducens* but similar to rates achieved by *Shewanella oneidensis* (3, 38). Scanning electron microscopy (SEM) of fully colonized electrodes revealed clusters of microcolonies distributed between fibers, with exposed electrode still visible even after 1 month of growth (Fig. 1B).

Glassy carbon, polished graphite, gold, indium tin oxide (ITO), and graphitic paper electrodes were also screened in triplicate reactors for attachment and colonization of *G. fermentans* using the higher-potential conditions. None of these materials significantly improved colonization or exceeded current densities of 5 µA/cm<sup>2</sup>. None of these materials altered the potential required for growth or the two different potential windows observed during voltammetry (such as those described in Fig. 2). Standard high-temperature cleaning treatments also did not significantly alter colonization or growth rates on carbon fiber electrodes (55).

**Cyclic voltammetry reveals two different potentials that stimulate electron transfer.** Low-scan-rate cyclic voltammetry performed on biofilms of electrode-reducing bacteria such as *Geobacter* (36, 38, 48), *Rhodospseudomonas palustris* DX-1 (60), and

*Thermincola ferriacetica* (35) always produces a sigmoidal response that increases steeply above voltages of  $-0.2$  V versus SHE and reaches a maximum by  $0$  V versus SHE. The response of *G. fermentans* differed from those of previously studied isolates and microbial fuel cell enrichments (9, 44, 51, 53). Most notable was how electron transfer from *G. fermentans* responded at two separate redox potentials. In addition, the majority of the increase in electron transfer rate occurred well above potentials of  $0.2$  V versus SHE. A representative time series is shown in Fig. 2A.

At all time points, these two characteristic windows were observed; one centered at approximately  $-0.2$  V, and the second centered at  $0.3$  V. First-derivative analysis of data (Fig. 2B) often revealed a third inflection point at  $0.4$  V, which increased in intensity in older biofilms. Together, this series of *G. fermentans* electrode-grown biofilms showed that while *G. fermentans* could initiate electron transfer at potentials as low as  $-0.2$  V, cells could not reach their maximum rates of sustained electron transfer unless provided with an electron acceptor near potentials of  $0.55$  V versus SHE, over half a volt higher than with any previously characterized organism. These data also revealed an unusual pattern of multiple independent features within this wide potential window.

**Soluble mediators are involved at both low and high potentials.** Earlier work with indirect assays based on acceleration of Fe(III) reduction implied that *G. fermentans* releases at least one soluble compound (4, 42). To test for the presence of soluble electron shuttles, the medium surrounding electrode-attached biofilms was removed and replaced with sterile anaerobic medium containing only the electron donor. In all cases, an immediate drop in current production was detected. Electron transfer could always be recovered by readdition of the original medium, even after anaerobic centrifugation or filtration to remove planktonic cells.

The absolute magnitude of the current drop after fresh medium was added increased with the age of the culture, varying from as low as 50% to as high as 75%. Data from a 3-week-old biofilm grown at  $0.55$  V are shown in Fig. 3A. These experiments verified that soluble compounds facilitate electron transfer between the cell surface and the electron acceptor, accounting for the majority of electron transfer. Surprisingly, these voltammetry experiments revealed changes in both redox potential windows in all experiments. This indicated that soluble compounds were involved in both low- and high-potential electron transfer mechanisms (Fig. 3B and C).

**Evidence for two independent redox-active compounds.** As medium replacement always decreased electron transfer by *G. fermentans* in both potential windows, this raised two possibilities. Either a single soluble compound was produced, with two distinct redox potentials, or two independent compounds were accumulating. As the relative amount of electron transfer in each window appeared to vary independently with culture age, growth data supported the hypothesis of two independent compounds.

An electrochemical assay was used to detect electrochemically active compounds in filtered supernatants. Voltammetry of sterile supernatants obtained from biofilms of *G. fermentans* always revealed two distinct redox peaks, at the two potential windows observed for electron transfer by whole cells (Fig. 4A). These assays verified the presence of redox mediators in culture supernatants.

Differential chromatographic separation was able to selectively remove one of these compounds, proving that two different me-

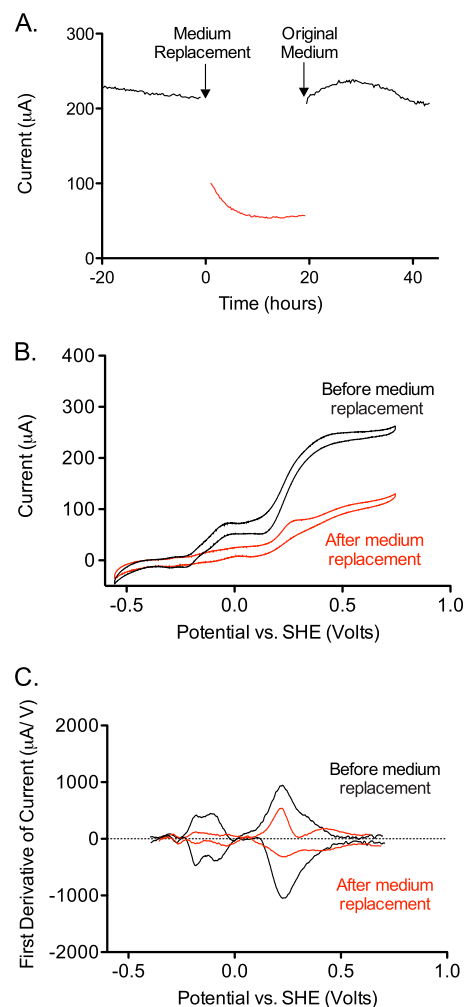
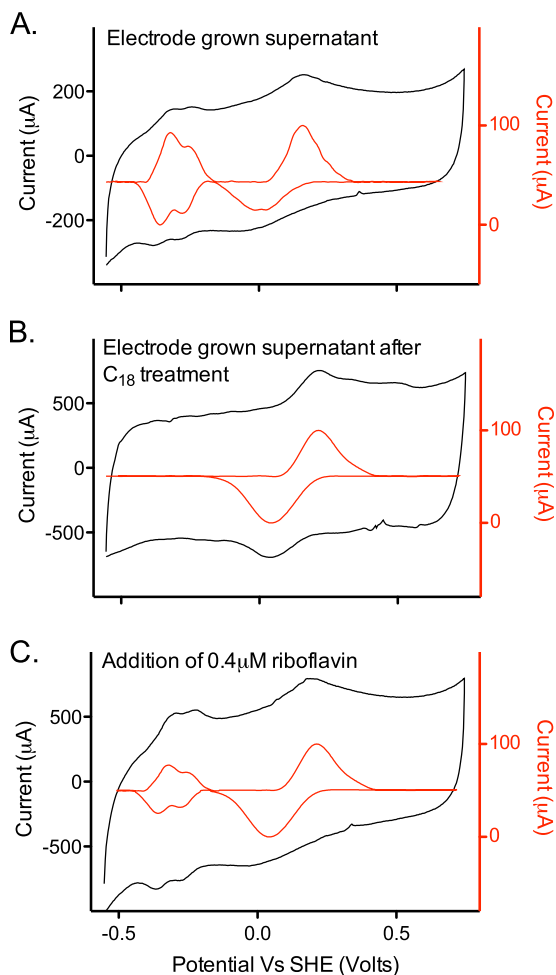


FIG 3 Evidence for two soluble mediators. (A) Rate of electron transfer by an established *G. fermentans* biofilm (black trace), subsequent drop in electron transfer upon medium replacement (red trace), and recovery of electron transfer to the original level after readdition of original medium following centrifugation (black trace). (B) Cyclic voltammograms performed before addition of fresh medium (black) and after addition of fresh medium (red) show changes in the catalytic waves in both low- and high-potential regions. (C) First-derivative analysis of voltammograms from panel B used to identify potential windows affected by medium replacement.

diators were present. Hydrophobic affinity columns, such as octadecyl-functionalized silica gel ( $C_{18}$ ), removed the compound associated with the lower potential window ( $-0.2$  V versus SHE) without altering the higher potential peak (Fig. 4B). When analyzed by HPLC, a single compound was detected in all supernatants that eluted at exactly the same time as authentic riboflavin standards, and this compound was not present in  $C_{18}$ -treated samples.

Unlike for *Shewanella*, which first secretes flavin adenine dinucleotide (FAD) and enzymatically processes it to FMN, no FMN or FAD could be detected in cell-free supernatants of *G. fermentans* cultures. Elution of  $C_{18}$  columns with methanol detected only a compound that eluted in HPLC as riboflavin, and the fluorescence emission profile of this eluted compound also was identical to that of riboflavin. When pure riboflavin was added back to  $C_{18}$ -treated

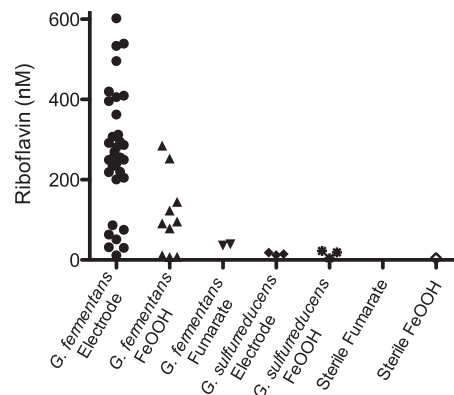


**FIG 4** Analytical electrochemistry evidence for two independent mediators. Supernatants collected from electrode-grown biofilms were centrifuged anaerobically to remove planktonic cells and analyzed using graphite electrodes. (A) Supernatant containing both the low- and high-potential-redox-active compounds. (B) Same supernatant after elution from a  $C_{18}$  column, showing the presence of only the higher-potential-redox-active compound. (C) Supernatant from panel B, after addition of  $0.4 \mu\text{M}$  riboflavin. Both compounds were detected in electrode-grown and FeOOH-grown cultures but not in fumarate-grown cultures. All scan rates were  $100 \text{ mV/s}$ . Insets (red traces) are baseline subtracted.

supernatants, the low-potential redox peak returned in electrochemical assays (Fig. 4C).

Throughout these studies, the higher-redox-potential compound showed no affinity for binding to graphite electrodes, hydrophobic matrices (e.g., compare high potential peaks in Fig. 4A and 4B), or filters used to sterilize supernatants. Treatment with high concentrations of commercial proteases had no effect on the size or location of this peak, suggesting that this mediator was not protein based. Treatment with metal-binding resins also had no effect on the size or location of this residual redox activity, suggesting that this redox peak was not due to a chelator or metal.

When riboflavin was depleted from these samples by  $C_{18}$  treatment, the supernatant containing the remaining high potential compound demonstrated a characteristic UV fluorescence with an excitation maximum of  $229 \text{ nm}$ . Emission by this fluorescent compound shifted to lower wavelengths after chemical reduction



**FIG 5** Ranges of riboflavin concentrations in cultures as detected by HPLC. Electrode-grown *G. fermentans* cultures always accumulated riboflavin, with the levels generally increasing with length of incubation. Riboflavin was always detected in FeOOH-grown cultures, but recovery was typically lower. Fumarate-grown *G. fermentans* cultures always contained less than  $20 \text{ nM}$  riboflavin, similar to *Geobacter* supernatants collected from electrode- and FeOOH-grown cultures.

of samples. This redox activity and fluorescence were not detected in supernatants from fumarate-grown cultures. These observations were consistent with a hydrophilic fluorescent compound reported previously for *G. fermentans* supernatants (42).

**Both redox shuttles are also produced in Fe(III)-grown *G. fermentans* cultures.** All electrode-grown and FeOOH-grown supernatants were reanalyzed using HPLC for the presence of riboflavin. All electrode-grown cultures contained riboflavin, although the concentration varied widely, from  $\sim 200 \text{ nM}$  (after 1 to 2 weeks of incubation) to  $\sim 600 \text{ nM}$  (after 3 to 4 weeks of incubation) (Fig. 5). Supernatants from FeOOH-grown cultures also contained a range of riboflavin concentrations, with levels between  $75 \text{ nM}$  and  $300 \text{ nM}$  detected. Since riboflavin is known to bind FeOOH and to spontaneously degrade upon exposure to light (37), these values represent a lower boundary of riboflavin concentrations. In contrast, *G. fermentans* cultures grown using soluble electron acceptors always had less than  $20 \text{ nM}$  riboflavin, a level at the limit of detection and similar to trace levels found in *G. sulfurreducens* supernatants from fumarate-, FeOOH-, and electrode-grown cultures. FeOOH-grown supernatants collected from four recently isolated *Geothrix* spp. (all within 95% 16S rRNA similar to *G. fermentans*; kindly provided by Z. Shelobolina) also contained riboflavin in the range of  $100$  to  $150 \text{ nM}$ , providing preliminary evidence that secretion of riboflavin may be conserved in related strains.

Sterile supernatants from FeOOH-grown *G. fermentans* cultures were also analyzed to determine if the high-redox-potential compound was also secreted under these conditions. As previous studies detected significant levels of soluble Fe(III) in *G. fermentans* cultures reducing FeOOH (42), it was possible that chelators maintaining trace minerals in a soluble state could create shuttling-like activity in our assays. To limit this possibility, all supernatants analyzed in this study were treated with metal-binding resins to remove soluble metals. Electrochemistry of sterile supernatants from FeOOH-grown cultures revealed that FeOOH-grown cultures always produced both riboflavin and the fluorescent high-potential compound. When these sterile, metal-depleted supernatants from FeOOH-grown cultures were added

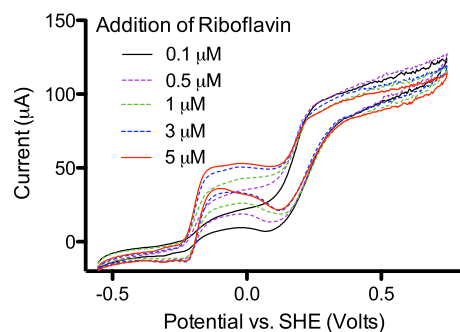


FIG 6 Dose response of an electrode-grown biofilm of *G. fermentans*. Addition of riboflavin, up to concentrations of 5  $\mu\text{M}$ , affected electron transfer only at the potential of  $-0.2$  V versus SHE, without stimulating overall current production.

back to washed biofilms of *G. fermentans*, electron transfer to electrodes was stimulated in the same two potential windows as electrode-grown supernatants (data not shown). This confirmed that redox-active compounds produced during growth with electrodes were also produced during more environmentally relevant conditions, such as growth with FeOOH.

**Stimulation of one pathway is at the expense of the other.** The presence of two redox-active compounds raised the question of whether the two mediators competed for electrons from the same supply or were part of independent additive pathways. The experiments where riboflavin was depleted from supernatants and added back to growing biofilms required medium exchanges, elution of medium through columns, and extensive reactor manipulation to execute. At the very least, switching respiration on and off introduced the artifact of the biofilm capacitance described in Fig. 1A, and the requirement for discharging the biofilm before additional electrochemical measurements could be obtained. These manipulations limited the sensitivity of the assay.

To alter redox shuttle concentrations under more controlled conditions, riboflavin was added stepwise to biofilms, allowing the concentration of the high-potential mediator and attached biomass to remain constant. Addition of riboflavin in this manner only increased the rate of electron transfer at the redox potential of riboflavin, without altering the total rate of electron transfer by the biofilm (Fig. 6). Thus, when riboflavin was present, electrons could be transferred at a lower potential, but at the expense of flux available to flow through the higher-potential pathway.

## DISCUSSION

Acidobacterial sequences related to *Geothrix* are regularly detected in Fe(III)-reducing environments, particularly in zones recently stimulated to cause bioprecipitation of U(VI) (9, 59). For example, *Geothrix*-like *Acidobacteria* were present in unamended sediments from the Oak Ridge Field Research Center and increased after lactate was injected into wells (5). Microcosm experiments from the same site also found *Geothrix*-like *Acidobacteria* comprising between 22 and 90% of 16S rRNA sequences after stimulation, depending on the electron donor (32). While such experiments also consistently detect *Geobacter*-like organisms, there is little information to address how these Fe(III)-reducing bacteria might coexist in the environment. These electrochemical data show that these organisms have fundamentally different ap-

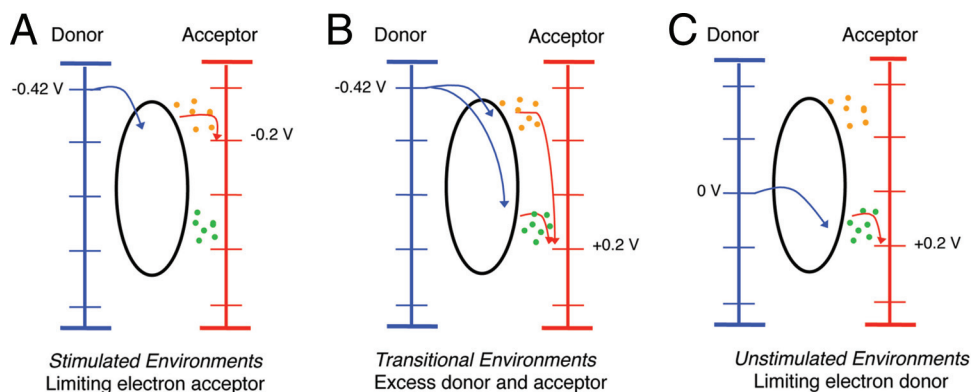
proaches, in terms of respiration rates, need for direct contact, and the redox potentials at which they can operate.

**A two-mediator, two-redox-potential window strategy.** To characterize the Fe(III) reduction strategy of *G. fermentans*, electrochemistry was used to control the redox potential of its electron acceptor during growth. These experiments revealed a dependence on higher redox potentials not previously observed in Fe(III)-reducing bacteria (Fig. 2 and 3). Sterile filtered supernatants from both electrode-grown and FeOOH-grown cultures confirmed that two separate redox-active compounds were secreted that stimulated electron transfer of *G. fermentans* (Fig. 4). The low-potential mediator demonstrated a high affinity for hydrophobic resins, facilitating separation and identification of the mediator as riboflavin. The other electron transfer mediator secreted by *Geothrix* had a much higher potential ( $\sim 0.3$  V versus SHE) than natural phenazine or isoalloxazine ring-based flavin compounds and was much more hydrophilic. Unlike flavins that bind tightly to graphite electrodes (37), the high-potential compound was easily washed from electrodes. Nevin and Lovley (42) reported that *G. fermentans* supernatants contained a water-soluble compound which fluoresced under UV light and that migrated in thin-layer chromatography similarly to model quinones. We also observed UV fluorescence in fractions from both electrode biofilms and Fe(III)-grown cultures. Even after riboflavin was removed, a hydrophilic high-potential redox-active compound remained. This behavior is consistent with substituted hydrophilic quinones, but as over 1,500 naturally produced quinones exist, the exact identity of this compound remains speculative. Purification of this compound will aid in the study of its reactivity with Fe(III) or other possible terminal acceptors under environmental conditions.

Because the low- and high-potential compounds could be easily separated, it was also possible to alter the concentration of each mediator in supernatants and correlate their presence with electron transfer at their characteristic redox potential. These experiments indicated that the effects of the two mediators were not additive. When additional riboflavin was added to biofilms, electron transfer rates at lower potentials increased but at the expense of electron transfer at higher potentials, resulting in the same total flux.

**Comparisons with other redox shuttles.** Some mediators appear to be secreted by bacteria to increase electron transfer to oxygen. Examples include phenazines (midpoint potential [ $E^{\circ}$ ] =  $-0.040$  to  $-0.170$  versus SHE) secreted by *Pseudomonas* spp., which improve survival of biofilm cells in anaerobic zones (56). Strains of *Bifidobacterium* grow to higher yields in the presence of 2-amino-3-carboxy-1,4-naphthoquinone ( $E^{\circ} = -0.071$  V) (a substituted hydrophilic quinone secreted by *Propionibacteria*), by shuttling electrons to oxygen and increasing NADH oxidation (61).

In contrast, flavins have been shown to participate in reactions related to reduction of metals. FMN secreted by *Shewanella* spp. ( $E^{\circ} = -0.21$  V) enhances reduction of many external metals and azo dyes and accounts for approximately 80% of electron transfer to extracellular acceptors (11, 37, 54). Gradients of oxidized flavins allow motile *Shewanella* strains to locate insoluble acceptors in space via energy taxis (26). Flavins secreted by plant roots are hypothesized to increase microbial rhizosphere availability of Fe(III) (46), and flavins in both yeast extract and intestinal contents promote electron transfer to oxygen (24) and electrodes (39). The finding that members of this phylogenetically distant acidobacterial group secrete riboflavin at concentrations similar



**FIG 7** Examples of environmental niches that could exist as a function of varying electron donor and acceptor concentrations if an organism can take advantage of multiple redox potentials. (A) Environments where donor is available, but acceptor concentrations (or metal redox state) limit the acceptor redox potential to  $-0.2$  V. (B) Environments with excess electron donors and acceptors, such as during laboratory growth, shortly after stimulation of an oxidized habitat with electron donor, or with electrodes poised at high potentials. (C) Environments where donor concentrations are limiting but the environment is oxidized, such as in low-carbon sediments or near oxic-anoxic interfaces.

to those secreted by *Shewanella*, specifically under Fe(III)-reducing conditions, shows that use of this versatile electron shuttle may be more widely distributed throughout the bacterial domain than previously recognized.

**What is the benefit of using higher-potential electron acceptors?** A final question is the ecological implication of electron transfer in the higher-potential domain. One benefit to higher-potential electron acceptors, in general, is the opportunity to capture more ATP. Based on the lowest potential at which *Geobacter*'s respiratory chain operates ( $-0.22$  V), and the redox potential of acetate ( $-0.28$  V), the poor  $\Delta G^\circ$  of this redox couple ( $-46$  kJ/mol acetate) explains why *Geobacter* has a growth yield consistent with only capturing  $<1$  ATP per acetate (33, 38). However, this low yield also means that in low-redox-potential environments, such as permanently anoxic zones, *Geobacter* can still respire and compete even when environmental conditions lower the  $\Delta G$  of acetate oxidation to values in this range (Fig. 7A and B).

If *Geothrix* were to capture additional ATP via electron flow to higher-potential electron acceptors (such as above an  $E^\circ$  of  $0.2$  V), a nearly 600% higher yield would be expected. In practical terms, *Geothrix* should reach OD<sub>600s</sub> of at least 0.3 (instead of the  $\sim 0.05$  observed) when respiring Fe(III)-citrate. On electrodes, instead of producing  $\sim 100$   $\mu\text{g}$  protein/ $\text{cm}^2$  from oxidation of 10 mM acetate (standard for *Geobacter*), one would expect over 600  $\mu\text{g}/\text{cm}^2$  from the equivalent amount of lactate. Yet, in all of our experiments, and all reported incubations with *Geothrix* and Fe(III) (10, 42), there is no evidence that *Geothrix* achieves significantly higher ATP yields from respiration.

In the absence of any evidence for increased ATP production, and based also on the fact that this slow-growing organism persists in unstimulated habitats where electron donor concentrations are low, another option is a strategy to enable survival under conditions where a low  $\Delta G$  exists. Oligotrophic sediments lacking a constant input of electron donor are higher in potential and interspersed with oxygenated zones. An organism able to shuttle electrons to freshly precipitated Fe(III) or Mn(IV) oxides, or low levels of oxygen, could grow at very low donor concentrations, but only if that organism conserved a small amount of energy from the process. By keeping only a small amount of energy for itself, an organism like *Geothrix* could position its metabolic strategy to respire at very low electron donor concentrations, as shown in Fig. 7C.

It was coupling of hydrogen partial pressure measurements with thermodynamic calculations that led Lovley and Goodwin (31) to explain zones where respiration was dominated by the presence of a single electron acceptor. In the typical version of this model, electron donors such as  $\text{H}_2$  are released from fermentation of organic matter. Strong electron acceptors such as  $\text{NO}_3$  provide a sufficiently negative  $\Delta G^\circ$  to support growth at  $\text{H}_2$  levels below 1 nM, a concentration unfavorable for reduction of Fe(III) and  $\text{SO}_4$ . This concept of “threshold” donor concentrations was confirmed by measurements of pure cultures, typically under conditions of excess electron acceptor (6, 28, 30).

The observation that *G. fermentans* can utilize higher-potential acceptors, and actively releases a compound which would be beneficial only at higher redox potentials, provides an example of an organism which may gain a competitive advantage under different conditions than typical permanently anoxic Fe(III)-reducing zones. Whether this high-potential window evolved for oxidized metals, shuttling to oxygen, or other acceptors remains to be discovered. Future enrichments targeting bacteria that have evolved to survive in such thermodynamic windows may need to control both redox potential and donor concentrations, to avoid organisms like *Geobacter* shifting conditions to where they can become dominant. In addition, work characterizing communities at oxic-anoxic interfaces, or in subsurface environments where this niche could exist, should consider the possibility that not all metal-reducing bacteria function best at low environmental redox potentials.

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