Introduction

Shewanella species are facultative anaerobic bacteria with a remarkable respiratory versatility, being able to couple the oxidation of organic matter to the reduction of terminal electron acceptors including nitrate (NO$_3^-$), Fe(III), Mn(IV), U(VI) and Tc(VII). To date, reactions involving Fe(III) and Mn(IV) have received the most attention, and here, the metals are thought to be reduced enzymatically by a range of anaerobic and facultatively anaerobic microorganisms, including Shewanella oneidensis MR-1, to poorly soluble species. The redox chemistry of Pu is more complicated, but the dominant oxidation state in most environments is highly insoluble Pu(IV), which can be reduced to Pu(III) which has a potentially increased solubility which could enhance migration of Pu in the environment. Recently it was shown that flavins (riboflavin and flavin mononucleotide (FMN)) secreted by Shewanella oneidensis MR-1 can act as electron shuttles, promoting anoxic growth coupled to the accelerated reduction of poorly-crystalline Fe(III) oxides. Here, we studied the role of riboflavin in mediating the reduction of radionuclides in cultures of Shewanella oneidensis MR-1. Our results demonstrate that the addition of 10µM riboflavin enhances the reduction rate of Tc(vi) to Tc(v), Pu(iv) to Pu(III) and to a lesser extent, Np(v) to Np(IV), but has no significant influence on the reduction rate of U(vi) by Shewanella oneidensis MR-1. Thus riboflavin can act as an extracellular electron shuttle to enhance rates of Tc(vi), Np(v) and Pu(iv) reduction, and may therefore play a role in controlling the oxidation state of key redox active actinides and fission products in natural and engineered environments. These results also suggest that the addition of riboflavin could be used to accelerate the bioremediation of radionuclide-contaminated environments.

Fe(III) substrate; (1) the direct transfer of electrons from the cell surface to the Fe(III) mineral, and (2) the use of low-molecular weight soluble redox mediators or “electron shuttles” to promote extracellular electron transfer. An Fe(III) reduction deficient mutant strain of Shewanella oneidensis MR-1 was found to contain a mutation in a gene (gspD) that encodes the outer membrane porin of a type II secretion system, which is involved in the extracellular translocation of Shewanella oneidensis outer membrane cytochromes MrTC and OmcA. The lack of a properly functioning type II protein secretion system renders the organism incapable of reducing Fe(III) oxides at wild-type rates. However, the Fe(III) reduction activity of the type II protein secretion mutant can be rescued by the addition of exogenous electron shuttling compounds such as anthraquione-2,6-disulfonate (AQDS).

In addition to direct reduction mechanisms via cytochromes, for more than a decade Shewanella strains have also been known to secrete soluble electron shuttles that can play a role in extracellular electron transfer. More recently, the flavin molecules FMN and riboflavin were identified as the electron shuttles secreted by S. oneidensis MR-1 and shown to promote growth via the reduction of poorly soluble Fe(III) oxides. These molecules have also been shown to play a role in mediating the flow of electrons to an anode surface in a microbial...
fuel cell containing *Shewanella oneidensis* MR-1 and *Shewanella* sp. MR-1.\(^{10}\)

*Shewanella oneidensis* MR-1 can also reduce a range of redox active radionuclides. For example U(\(vi\)) is reduced directly under anoxic conditions, resulting in the precipitation of U(\(vi\)) as uraninite (UO\(_2\)).\(^{11-13}\) Recently, it was demonstrated that c-type cytochromes of *Shewanella oneidensis* MR-1 are essential for the reduction of U(\(vi\)) and the subsequent formation of extracellular UO\(_2\) nanoparticles.\(^{12,14}\) In particular, the outer membrane (OM) decaheme cytochrome MtrC, previously implicated in Mn(\(vi\)) and Fe(\(iii\)) reduction, was shown to play a role in transferring electrons to U(\(vi\)). Additionally, deletions of mtrC and/or the gene encoding another outer membrane cytochrome omcA significantly affected the *in vivo* rate of U(\(vi\)) reduction relative to the wild-type MR-1 strain.\(^{12}\) A study on the electron transfer pathway for U(\(vi\)) reduction mediated by FMN, which is secreted by *Shewanella* species, demonstrated that FMN may also act as a mediator during the reduction of U(\(vi\)) to U(\(iv\)), accelerating its reduction.\(^{15}\)

Tc(\(vi\)) in contrast, is generally thought to be reduced via direct interactions with hydrogenase, using hydrogen as an electron donor.\(^{16,17}\) *Shewanella oneidensis* MR-1 is able to catalyse this reaction consistent with the presence of a NiFe hydrogenase in the strain.\(^{15}\) However, this organism can also couple lactate oxidation to a slower rate of enzymatic Tc(\(vi\)) reduction, with c-type cytochromes implicated in electron transfer to the radionuclide with this electron donor.\(^{18}\)

The reductive capability of *Shewanella putrefaciens* (ATCC 8071) (now renamed *S. oneidensis*) has also been used for the reduction of Np(\(vi\)) to lower valence (probably Np(\(v\))). However, the neptunium in these experiments could not be removed from solution in this study via this direct reductive precipitation mechanism. Instead, bioreduction to Np(\(iv\)) by *S. putrefaciens*, together with phosphate liberation by a *Citrobacter* sp. with high phosphatase activity, permitted bioprecipitative removal of Np, presumably as an insoluble Np(\(iv\)) phosphate.\(^{19}\) A later study demonstrated that cell suspensions of *S. oneidensis* were able to enzymatically reduce unchelated Np(\(iv\)) to insoluble Np(\(iv\))(s), and that the addition of citrate enhanced Np(\(iv\)) bioreduction.\(^{20}\)

Finally, the mechanism of plutonium reduction by *S. oneidensis* MR-1 has also been studied, and it was demonstrated that in the absence of complexants, very little Pu(\(iii\)) was produced from the enzymatic reduction of Pu(\(iv\))(OH)\(_4\). By contrast, in the presence of the complexant EDTA, most of the Pu(\(iv\))(OH)\(_4\)(am) was reduced to Pu(\(iii\)).\(^{21,22}\) Clearly, much work remains to be done to fully understand mechanism and environmental consequence of Pu hydrous oxide reduction in this and other organisms.

The aim of the work described in this paper is to determine if flavins can be used to accelerate the reduction of a range of priority radionuclides by *S. oneidensis*. This information is important to help elucidate the mechanisms of radionuclide reduction in flavin-secreting metal-reducing bacteria, and also to help inform biotechnological approaches for the efficient bioremediation of radionuclide contaminated land and/or wastewaters. Wild type cells of *Shewanella oneidensis* MR-1 were used, and in the case of U(\(vi\)) and Tc(\(vi\)), a mutant with an in-frame deletion of gspD (\(\Delta\)gspD) was used to identify the potential role of outer membrane c-type cytochromes.

### Material and methods

#### Bacterial strains and media

*Shewanella oneidensis* MR-1 was obtained from University of Manchester Geomicrobiology group culture collection and the *Shewanella oneidensis* targeted in-frame deletion mutant (\(\Delta\)gspD) was obtained from Thomas DiChristina (Georgia Tech, USA).\(^{6}\) Starter cultures of the bacteria were cultured aerobically in tryptic soy broth (Oxoid CM0876) for 10 h (30 °C) at 125 rpm and then transferred to a *Shewanella* minimal medium.\(^{3}\) In the case of the mutant, 50 µM of the antibiotic rifamycin was added to the medium. The bacteria for experimental cultures were grown to mid-exponential growth in *Shewanella* minimal medium with DL-sodium lactate (100 mM) as the electron donor and sodium fumarate as electron acceptor (20 mM) for 20 h at 30 °C under an atmosphere of N\(_2.\)\(^2\) Cell suspensions were prepared for metal reduction assays by washing the cells twice and resuspending them in 30 mM sodium bicarbonate buffer (pH 7; degassed with N\(_2\) : CO\(_2\) (80 : 20)).

#### U(\(vi\)) reduction assay

U(\(vi\)) reduction assays were conducted in 30 mM sodium bicarbonate buffer (pH 7) containing 500 µM U(\(vi\)) as uranyl acetate. DL-Sodium lactate was used as the electron donor (10 mM) with or without the addition of 10 µM riboflavin as an extracellular electron shuttle. Controls were also prepared which contained no added electron donor or no cells. Sterile solutions of electron donor, riboflavin and *S. oneidensis* MR-1 cell suspension or \(\Delta\)gspD mutant cells were added anaerobically to the autoclaved and degassed buffer solution. Kinetic studies were initiated by the addition of 0.2 ml of a standardised suspension of cells of *S. oneidensis* MR-1 or the \(\Delta\)gspD mutant, resulting in a final assay density of 2 × 10\(^8\) cells per ml. All experiments were conducted in triplicate. The assay tubes were incubated at 30 °C in the dark. At multiple time-points samples were collected and the U(\(vi\)) content of the supernatant (after centrifugation at 1 min, 9000 g) was analysed using a colorimetric assay with 2-(5-bromo-2-pyridylazo)-5-diethylaminophenol.\(^{21}\)

#### Tc(\(vi\)) reduction assay

The kinetics of Tc(\(vi\)) reduction by wild-type *S. oneidensis* MR-1 and a \(\Delta\)gspD mutant were determined using washed cell suspensions. Tc(\(vi\)) reduction assays contained a final concentration of 100 µM pertechnetate (CERCA, France) in pH 7, 30 mM sodium bicarbonate buffer purged with an N\(_2\) : CO\(_2\) (80 : 20) gas mixture. The electron donor used to support radionuclide reduction in this study was DL-sodium lactate (10 mM). In addition, 10 µM riboflavin was added as an electron shuttle. Kinetic studies were initiated by the addition of 0.2 ml of a...
standardised suspension of cells of *S. oneidensis* MR-1 or the Δ*gspD* mutant, resulting in a final assay density of 2 × 10⁶ cells per ml. All experiments were conducted in triplicate and incubated at 30 °C in the dark. At multiple time-points, samples (0.2 ml) were collected and centrifuged for 4 min at 7000g to remove the cells and any precipitated radionuclide. An aliquot of the supernatant (50 µl) was analysed for total Tc concentrations by liquid scintillation counting on a Packard Tri-Carb 2100TR. Another aliquot of the supernatant (100 µl) was also complexed with tetr phenyl arsonium chloride (TPAC) and then the TPAC-TcO₄⁻ complex was extracted from the soluble phase with chloroform, prior to liquid scintillation counting (as above) to quantify the concentration of Tc(Ⅵ) in solution.¹⁸

**Np(Ⅴ) reduction assay**

The kinetics of aqueous Np(Ⅴ) reduction by the wild-type *S. oneidensis* MR-1 were investigated by resting cell assays using washed cell suspensions. All experiments were carried out at pH 7, in 50 mM 3-(N-morpholino)propanesulfonic acid (MOPS) buffer, that had been purged with a N₂:CO₂ (80:20) gas mixture. Three treatments were established (all with 11 mM Np added as NpO₂⁻); (i) abiotic (without cells) containing 10 mM sodium lactate, (ii) biotic (with wild type cells) with 10 mM sodium lactate, and (iii) biotic (with wild type cells) containing 10 mM sodium lactate and 10 mM riboflavin. All medium constituents were sterilised by autoclaving or <0.22 µm filtration (in the case of riboflavin) before use. The biotic treatments were inoculated with washed *S. oneidensis* MR-1 cells, resulting in a final assay density of 4 × 10⁷ cells per ml. All experiments were conducted in triplicate and incubated at 30 °C in the dark. Periodically, samples (~400 µl) were taken aseptically using argon gas purged needles and syringes. Thereafter, samples were centrifuged for 5 min at 7000g, and then diluted into 2% HNO₃ for determination of the total neptunium concentration in solution by inductively coupled plasma mass spectrometry (ICP-MS) with ²³²Th as an internal standard. It was assumed that all Np remaining in solution was present as Np(Ⅴ).

An additional experiment to ascertain the Np oxidation state at the end of the incubation in the biotic (with wild type cells) treatment containing 10 mM sodium lactate and 10 µM riboflavin was also established. Here, the Np concentration of the starting solution was increased to 50 µM whilst all other parameters remained unchanged. After 120 hours of incubation the total amount of neptunium in solution was then measured, and a cell pellet was harvested via centrifugation (5 min at 7000g) under an inert atmosphere for subsequent XAS analysis.

**Neptunium L₃-edge X-ray absorption spectroscopy**

Under an inert atmosphere the Np labelled cell-pellet was mixed with boron nitride to attain a solid sample with ~400 ppm Np. The sample was then carefully packed into an airtight XAS sample tube and stored under an inert atmosphere at −80 °C prior to analysis. Neptunium XAS analysis was conducted at the INE Beamline for actinide research at the ANKA synchrotron light source, Karlsruhe, Germany. Neptunium L₃ edge XANES spectra were collected in fluorescence mode by a 5 element solid-state geranium detector with the sample maintained at cryo-temperatures throughout analysis. Parallel measurements of a Zr foil were also taken in transmission for energy calibration. Spectra summation and analysis was conducted using the ATHENA software.

**Pu(Ⅵ) reduction assay**

Pu(Ⅵ) reduction assays contained a final concentration of 1 mM ²⁴²Pu in 30 mM sodium bicarbonate buffer purged with an N₂:CO₂ (80:20) gas mixture. α-sodium lactate (10 mM) was used as an electron donor, with or without added riboflavin (10 µM). Control solutions contained no cells. All experiments were conducted in triplicate and incubated at 30 °C in the dark. ²⁴²Pu concentrations were measured by liquid scintillation counting.²⁴ Solution concentrations in supernatant samples were measured after centrifugation (14 000g, for 5 min). For ²⁴²Pu experiments, precipitates and cells were collected by centrifugation (14 000g, for 5 min), dissolved in 1 M HCl and UV-visible-near infrared spectra were recorded from the resulting supernatants using the method described by Boukhalfa and coworkers.²¹

**Results**

**The influence of riboflavin on U(Ⅵ) reduction**

*Shewanella oneidensis* MR-1 can reduce U(Ⅵ) under anoxic conditions, which results in the precipitation of uraninite (UO₂).¹¹⁻¹³ Outer membrane-associated c-type cytochromes are thought to be involved in U(Ⅵ) reduction,¹² although the involvement of secreted flavin redox shuttles has not been investigated. To test if riboflavin could enhance the rate of U(Ⅵ) reduction by *S. oneidensis* MR-1, 10 µM of the flavin was added to cell suspensions and the impact on the rate of U(Ⅵ) reduction was quantified. The added riboflavin had no significant influence on the reduction rate of U(Ⅵ) by the anaerobically-grown wild type cells (Fig. 1), suggesting that addition of the endogenous electron shuttle does not play a significant role in enhancing reduction of this radionuclide. The cell pellets collected in these cultures after centrifugation were also stained black, indicating the likely formation of reduced, insoluble uraninite.¹¹ However, the Δ*gspD* mutant that was unable to secrete outer membrane cytochromes required for the reduction of extracellular electron acceptors, reduced U(Ⅵ) at a slower rate than the wild type (Fig. 1). This Δ*gspD* mutant reduced U(Ⅵ) at a slightly faster rate with 10 µM riboflavin added, but the rate of uranium reduction did not reach the reduction rate noted in the cultures of wild type cells.

**The influence of riboflavin on Tc(Ⅶ) reduction**

*Shewanella oneidensis* MR-1 couples the rapid reduction of Tc(Ⅶ) to the oxidation of hydrogen, in common with other well studied Tc(Ⅶ)-reducing cultures, but can also use lactate as an electron donor for this reaction, albeit at a slower
rate. To test if riboflavin accelerated the reduction rate of Tc(VII) by *S. oneidensis* MR-1 with lactate as an electron donor (presumably coupling to electron transfer from c-type cytochromes that are abundant in the cell), 10 µM of riboflavin was added to resting cell suspensions challenged with Tc(VII). The addition of 10 µM of riboflavin enhanced significantly the rate of Tc(VII) reduction catalysed by the wild type *S. oneidensis* MR-1 cells and also the ΔgspD mutant (Fig. 2). In the presence of riboflavin, the MR-1 strain and the ΔgspD mutant were both able to reduce ∼55% of the Tc(VII) within 22 h. In contrast, without riboflavin both cultures reduced only 20% of the Tc(VII) within this time period. Within 48 h ∼80% of the Tc(VII) was reduced by both the MR-1 strain and the ΔgspD mutant, and all the detectable Tc was reduced after 120 hours in the presence of the added flavin. Without added riboflavin, there was a clear difference in the amount of Tc(VII) reduced after 120 hours, with the wild type strain reducing 71% ± 6% compared to only 40% ± 7% of the initial Tc(VII) in cultures of the ΔgspD mutant (Fig. 2). A black cell-associated precipitate was noted in all cultures where Tc(VII) reduction occurred consistent with the formation of insoluble Tc(IV)O₂;

**The influence of riboflavin on Np(V) reduction**

The reduction of Np(V) by *Shewanella* species has been noted previously, although interestingly other Gram-negative Fe(III)-reducing bacteria such as *Geobacter sulfurreducens* cannot reduce this radionuclide. As the mechanism of Np(v) reduction in *Shewanella* species remains poorly characterised,
in this study we investigated the impact of the addition of 10 µM riboflavin on the reduction of Np(V) by *Shewanella oneidensis* MR-1. As shown in Fig. 3, the addition of 10 µM of riboflavin enhanced slightly the rate of reduction of Np(V) by *S. oneidensis* MR-1, which was mainly noticeable at the 24 hour time-point.

The wild type cells of *S. oneidensis* removed >90% of the available Np from solution after incubation for 48 h. As shown in Fig. 4, comparison of XANES spectra collected from experimental cultures, with those from Np(IV) and (V) standards showed spectra from the 50 µM Np(V) experiment end point were consistent with the presence of Np(IV). Specifically, the XANES lacked the multiple scattering feature on the high energy side of the absorption edge which is attributed to scattering along the neptunyl axial oxygens. Indeed, a simple two end-member linear combination model of the sample between the oxic Np(V) and reduced Np(IV) standards, suggested 80% of the Np was speciated as Np(IV). This highlighted the dominance of Np(IV) in these systems with only a modest, if any, contribution from Np(V) sorption, and confirmed the reductive precipitation of Np(V) to Np(IV).

**The influence of riboflavin on Pu(IV) reduction**

Previous studies have shown that washed cell suspensions of *Shewanella oneidensis* MR-1 produce very little Pu(III) enzymatically from Pu(IV)(OH)₄, whereas the presence of the complexant EDTA enhances the reduction of the Pu(IV)(OH)₄(am) to Pu(III), presumably by increasing the solubility and bioavailability of Pu(IV). To determine if the addition of riboflavin enhances the reduction of Pu(IV), 10 µM of riboflavin was added to washed cell suspensions with 1 mM Pu(IV), and the ability of the cells to reduce the radionuclide investigated.
Plutonium was removed rapidly from solution (within 10 min). After 24 h incubation, UV-visible-near infrared spectra of the dissolved cell pellets (Fig. 5) were altered, with increases in the amplitude of diagnostic peaks for Pu(Ⅲ) at 561, 601 and 1025 nm associated with the cell suspensions supplemented with lactate or lactate and riboflavin. The diagnostic peaks for Pu(Ⅲ) at 601 nm and 1025 nm (* ) are characteristic for Pu(Ⅲ) and are present in the samples of wild type cells with and without riboflavin.

Our results also demonstrated that a Shewanella oneidensis ∆gspD mutant reduces U(Ⅵ) and Tc(Ⅶ) at slower rates than the wild-type. The S. oneidensis ∆gspD mutant, that lacks a functioning type II protein secretion systems, is also incapable of reducing iron(oxy)hydr(ox)ides and Mn(Ⅳ) oxides at wild-type rates. These data are to be expected as the outer membrane c-type cytochromes of S. oneidensis MR-1 (e.g. MtrC, also known as OmcB and OmcA) are thought to be essential for lactate-dependent reduction of Fe(Ⅲ), and also U(Ⅵ) and Tc(Ⅶ), presumably mediated at the cell surface. Indeed, these electron transfer proteins are translocated across the outer membrane to this site by the type II protein-secretion systems, is also incapable of reducing iron(oxy)hydr(ox)ides and Mn(Ⅳ) oxides at wild-type rates. These data are to be expected as the outer membrane c-type cytochromes of S. oneidensis MR-1 (e.g. MtrC, also known as OmcB and OmcA) are thought to be essential for lactate-dependent reduction of Fe(Ⅲ), and also U(Ⅵ) and Tc(Ⅶ), presumably mediated at the cell surface. Indeed, these electron transfer proteins are translocated across the outer membrane to this site by the type II protein-secretion systems, is also incapable of reducing iron(oxy)hydr(ox)ides and Mn(Ⅳ) oxides at wild-type rates.

Discussion

The secretion of endogenous flavin electron shuttles by Shewanella species has been shown to promote the dissimilatory reduction of extracellular electron acceptors including poorly soluble Fe(Ⅲ) oxides and anodes in microbial fuel cells. However, the impact of secreted flavin molecules on the redox state of radionuclides has not been studied intensively to date, even though in many cases they may be present at the surface of microbial cells; microbial uptake of actinides into the microbial cell remains a controversial subject, and bioavailable U(Ⅵ) can be sorbed to minerals in contaminated land.

Information provided here is important to help understand the mechanisms of electron transfer to contaminant metals and radionuclides, and to aid in the design and optimisation of bioremediation approaches. Shewanella species have the widest radionuclide substrate range of metal-reducing prokaryotes studied to date, are able to mediate Tc(Ⅶ) reduction via novel non hydrogenase-mediated mechanisms and also have the capacity to enzymatically reduce Np(Ⅵ), unlike the few Geobacter species that have been studied. The aim of our study was to identify novel mechanisms of radionuclide reduction in Shewanella oneidensis MR-1, and in particular to investigate the influence of riboflavin on the reduction of the actinides U(Ⅵ), Np(Ⅵ) and Pu(Ⅳ) as well as the fission product Tc(Ⅶ).

The addition of 10 μM riboflavin increased significantly the reduction rate of Tc(Ⅶ) by wild type cultures of S. oneidensis MR-1, while having a more modest, but significant, impact on the reduction rate of Np(Ⅵ). In addition, more Pu(Ⅳ) was also detected in the presence of riboflavin than without (17% to 3%). Interestingly, there was no significant change in the reduction rate of U(Ⅵ) by S. oneidensis MR-1 when riboflavin was added, suggesting that under the conditions of study, secreted flavins do not play a significant role in mediating U(Ⅵ) reduction by this organism. This is in contrast to a study on a potential flavin mononucleotide (FMN)-mediated electron pathway for microbial U(Ⅵ) reduction by Shewanella putrefaciens (ATCC 8072), which suggested that the addition of FMN promoted U(Ⅵ) reduction. It is noteworthy that there were several key differences between this study and our experiments: another Shewanella strain was used, with FMN (instead of riboflavin), which was added at higher concentrations (50 μM and 500 μM), and in combination with U(Ⅵ) citrate (instead of uranyl acetate in our experiments). Thus direct comparisons are difficult, although these data do suggest that very high concentrations of flavin molecules could enhance U(Ⅵ) reduction in some experimental systems.

Our results also demonstrated that a Shewanella oneidensis ∆gspD mutant reduces U(Ⅵ) and Tc(Ⅶ) at slower rates than the wild-type. The S. oneidensis ∆gspD mutant, that lacks a functioning type II protein secretion systems, is also incapable of reducing iron(oxy)hydr(ox)ides and Mn(Ⅳ) oxides at wild-type rates. These data are to be expected as the outer membrane c-type cytochromes of S. oneidensis MR-1 (e.g. MtrC, also known as OmcB and OmcA) are thought to be essential for lactate-dependent reduction of Fe(Ⅲ), and also U(Ⅵ) and Tc(Ⅶ), presumably mediated at the cell surface. Indeed, these electron transfer proteins are translocated across the outer membrane to this site by the type II protein-secretion pathway. The disruption of the type II secretion system in the S. oneidensis ∆gspD mutant seemingly prevents the delivery of the c-type cytochromes to this key exposed and reactive compartment of the Gram-negative bacterial cell, thus lowering the rate of reduction of U(Ⅵ) and Tc(Ⅶ).

It was demonstrated previously that the addition of the humic analogue and extracellular electron shuttle anthraquinone-2,6-disulphonate (AQDS) reinstates the ability of the ∆gspD mutant to reduce extracellular insoluble Fe(Ⅲ) oxides.

Fig. 5 UV-visible-near infrared absorbance spectra of cell pellets of wild type cells of Shewanella oneidensis MR-1 dissolved in 1 M HCl from cultures containing 1 mM Pu(OH)4 incubated for 24 h at 30 °C and pH 7.0. Lactate was supplied as an electron donor, where indicated, at 10 mM while riboflavin was supplied at 10 μM. Controls had no cells or no electron donor included. Spectral features at 561, 601 and 1025 nm (*) are characteristic for Pu(Ⅲ) and are present in the samples of wild type cells with and without riboflavin.

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Our results showed that the addition of riboflavin to cultures of the ΔgspD mutant also results in an increase in the rate of Tc(vi) reduction, restoring it to the levels noted for the wild-type cells. Thus, the reduction of Tc(vi) may be mediated directly by (reduced) riboflavin, which would have access to c-type cytochromes that were not secreted, for example CymA which is localised to the periplasm of the Gram-negative cell. Similar predictions can be made for the actinides Np(v) and Pu(III), on the basis of enhanced reduction in the wild type cultures, when riboflavin was added. Clearly further work is required on these more radiologically challenging elements, to help define the precise mechanism of bioreduction in Shewanella cultures, and other environmental systems. However, the situation with uranium is distinct, and here the rate of reduction of U(vi) by the ΔgspD mutant did not reach the same rate noted in the wild-type cultures even in the presence of riboflavin. This suggests that the outer membrane c-type cytochromes of S. oneidensis MR-1 most likely play a defining role in the direct reduction of U(vi) at the cell surface which cannot be compensated for by the addition of the electron shuttle. This is consistent with other studies, which have shown that the correct processing of surface exposed outer membrane c-type cytochromes is a key determining factor in extracellular electron transfer in S. oneidensis MR-1. This would not seem to be true of Tc(vi) and potentially Np(v) and Pu(III), and it seems for these radionuclides riboflavin could likely mediate directly between the enzymatic electron transfer machinery of the cell and the radionuclide. This is in contrast to U(vi), which seems unlikely to couple directly to the reduced flavin molecule under the conditions used in our experiments. It is conceivable that the reduced riboflavin-Tc(vi) interactions could be in free solution, or mediated by flavin groups bound to the outer membrane cytochromes as noted recently.

Shewanella species and other microorganisms that actively secrete flavins and use them as electron shuttles could have an adaptive advantage in environments that contain significant concentrations of metals and radionuclides that are not easily to access (e.g. intergran areas or low porosity rocks and mineral assemblages). The concentration of flavins secreted by anaerobically grown cultures of S. oneidensis was measured at the relatively low levels of 0.1 to 0.6 µM, but it is expected that the local concentrations of secreted flavins in microenvironments such as biofilms or micropores in minerals are potentially much higher, approaching the levels used in our experiments. However, the addition of riboflavin (available in large quantities as a relatively inexpensive food additive) could also be used to accelerate the bioremediation of radionuclide contaminated environments, for example in sediments contaminated with Tc(vii) and Np(v). Pure culture experiments such as those described here can help identify the fate of key radionuclides in environmental systems, and the underpinning mechanisms catalysing biocycling reactions. Although it is clear that there is a need for follow up experiments on more complex environmental systems, including those using mixed microbial consortia, realistic groundwater chemistries and appropriate sediment/mineral phases, it should be noted the reductive endpoints for U, Np, and Tc, and potentially Pu noted in these experiments are broadly in line with those recorded in sediment microcosms that have been driven anaerobic by complex, extant microbial communities. With advances in analytical tools for flavin analyses in complex environmental systems, and the use of sensitive meta-omic tools targeting flavin secretion pathways, and other electron transfer systems, it should be possible to assess the precise role of these pathways in controlling radionuclide species in anoxic systems.

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