

Microbially Promoted Solubilization of Steel Corrosion Products and Fate of Associated Actinides

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Research Objective

The U.S. Department of Energy (DOE) statements of need call for “biological and physical chemical parameters for effective decontamination of metal surfaces using environmentally benign aqueous-based biopolymer solutions and microbial processes with potential for decontaminating corroding metal surfaces.” Improved understanding of the fundamental processes of microbial reductive dissolution of iron oxide scale on corroding carbon steel will support assessment and potential application of an environmentally benign and cost-effective strategy for in situ decontamination of structural metal surfaces and piping.

This research is designed to develop a safe and effective biological approach for decontaminating mild and stainless steels that were used in the production, transport, and storage of radioactive materials.

Research Progress and Implications

This report summarizes research progress made during the 3-year tenure for this project. An extension with minimal carryover funds was requested and granted to complete and submit manuscripts for publication. During this research, we have

- Demonstrated that Fe(III)-reducing bacteria reduce Pu(IV) (insoluble) to Pu(III) (soluble).
- Confirmed that the bacteria sorb and accumulate trivalent cations, such as Pu(III).
- Demonstrated that bacteria attached to oxide surfaces are very difficult to remove. Concluded that recovery of bacteria with sorbed Pu(III) would be impractical.
- Demonstrated that Fe(II) and reduced quinone-like compounds, which are both products of anaerobic respiration, can reduce chemically reduce solid Pu(IV) to dissolved Pu(III).
- *Conceptualized a bead-based system that effectively removes Pu from iron oxides and accumulates Pu(III) in beads of sodium alginate that can be easily separated from the bulk aqueous phase.*

Dissimilatory iron-reducing bacteria enzymatically reduce and dissolve iron oxides, which are common components of corrosion films, and release soluble species of plutonium, Pu(III). Consistent with our previous hypothesis, cell surfaces sorb Pu(III) and remove it from the bulk aqueous phase. However, we incorrectly hypothesized that bacteria with sorbed actinides could be easily detached and recovered from the surfaces that they had colonized and enzymatically altered. In fact, we have demonstrated that although cells do naturally detach from oxide

surfaces during their growth cycle, they leave behind negatively charged reactive portions of their outer surface that are strong sorbants for cations. Without a means for recovering both intact bacteria, their subcellular products and associated contaminants, the use of iron-reducing bacteria for decontaminating corroded steel surfaces would not be feasible. Hence, we have targeted an approach that avoids direct contact and attachment of cells to the corrosion films but allows for reduction, dissolution, and sorption of corrosion products and associated actinides.

Description of Bead-Based Treatment

Iron reducing bacteria are encapsulated in small beads of sodium alginate. Encapsulation prevents direct contact between the bacteria and the contaminated oxide surface. Anthraquinone disulfonate (AQDS) is used as a dissolved electron shuttle to carry electrons from the bacteria to Fe(III) and Pu(IV) on the corrosion film. AQDS reduces Fe(III) to Fe(II) and Pu(IV) to Pu(III). The reduced forms of these metals are very soluble and partition to the aqueous phase. The bacterial surface and the sodium alginate sorb and accumulate Fe(II) and Pu(III). The beads, which now contain most of the Pu(III), can be easily separated from the bulk aqueous phase. The benign process requires no hazardous chemicals or extreme pH conditions.

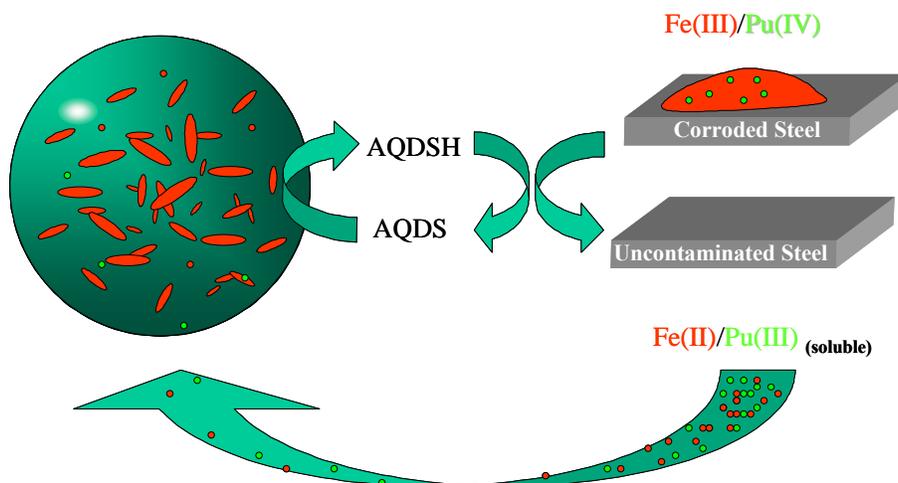


Figure 1. This illustrates a conceptual model of a bead-based system for decontaminating corroded steels. Metal-reducing bacteria are enrobed in porous alginate beads. Oxidized anthraquinone disulfonate, AQDS, which will serve as a dissolved electron shuttle between immobilized cells and elements in the corrosion film, diffuses into the beads and is enzymatically reduced by the bacteria. The reduced AQDSH diffuses out of the bead and chemically reduces and dissolves Fe(III) and Pu(IV) in the corrosion film. Soluble Fe(II) and Pu(III) sorb to cationic exchange sites within the alginate beads. The beads and accumulated actinides can then be easily separated from the bulk aqueous phase and the uncontaminated steel.

Carryover funds were insufficient to evaluate the bead-based system for removing Pu from contaminated corrosion. However, the system was tested for its ability to remove dissolved radionuclides (uranium and technetium) from aqueous media. Figure 2 illustrates that iron-reducing bacteria embedded in a porous alginate matrix reduce dissolved Tc(VII) (as pertechnetate ion) to poorly soluble TcO₂. Vials 1 and 2 contain alginate beads that are blackened by TcO₂ precipitates. Vial 3 served as a control and contains beads with no cells. All of the Tc(VII) remained in the aqueous phase. Vials containing dissolved U(VI) in lieu of Tc(VII) yielded similar results. These results clearly demonstrate the potential for removing dissolved radionuclides from solution using an environmentally benign bead-based approach.

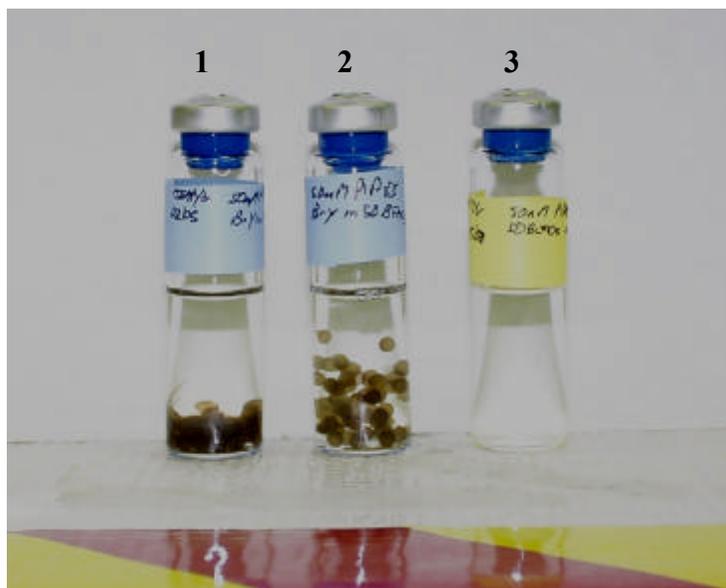


Figure 2. Vials demonstrating the ability of iron-reducing bacteria in alginate beads to remove dissolved Tc(VII) from solution. 100 μ M of Tc(VII) were reduced and precipitated within the beads in vials 1 and 2. Vial 3 served as a negative control and lacked bacterial cells. Beads without bacteria are difficult to see because they contain none of the dark Tc(IV) precipitate.

Information Access

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