The biogeochemistry of Pu mobilization and retention

B.D. Honeyman (PI)¹, A.J. Francis², C. J. Dodge², J.B. Gillow² and P.H. Santschi³

¹Environmental Science and Engineering Division, Colorado School of Mines; ²Environmental Sciences Department, Brookhaven National Laboratory; ³Texas A&M University at Galveston









Historically, expectations of minimal aqueous Pu transport of have been confounded by the apparent contradiction:
an understanding of low Pu solubility (based on its inorganic speciation) and
the observed transport of Pu sometimes at substantial distances from its presumed

source.

Of the relatively limited number of papers on Pu environmental speciation, a majority have implicated 'organic matter' as a transport agent^{*}.

This project focuses on the role of bacteria in the production of EPS: as Pu mobilization and immobilization agents.

*(e.g., Nelson *et al.,* 1990; Orlandini *et al.,* 1990; Loyland *et al.,* 2001; Honeyman, 1998; Santschi *et al.,* 2002)

Comparison of U and Pu solubility



Examples of environmental measurements





<u>Pu release from vegetated soils as a function</u> of quality and quantity of DOC



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<u>PAGE (Gel electrophoresis) of ^{239,240}Pu from Soil</u> <u>Resuspension Experiment</u>



Santschi et al. (2002) Environ. Sci. Technol., 36, 3711-3719.

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Pourbaix Diagram (Eh/ pH) diagram for the system Pu-O $_2$ -CO $_2$ -H $_2$ O at 25 °C and 1 bar total pressure for Σ Pu(aq)=10 ⁻⁸ M and total carbonate (C $_T$) = 10 ^{-2.0} M



- A: Aerobic bacteria
- **B:** Denitrifers
- C: Mn-reducers
- D. Fe reducers
- E: Fermenters
- F: Sulfate reducers

Pu mobility / immobility as a transformational process





Key: A. Pu carrier-phase dissolution; B. Trapping of colloidal Pu by EPS;C. Biodegradation of Pu-EPS; D. Enhanced Transport of Pu by EPS.

Characterization of EPS

<u>TEM of <0.45 μm EPS from</u> Shewanella putrefaciens



Bar = 200 nm

TEM image of Clostridium sp.



J.B.Gillow (unpublished)

TEM image of *Clostridium* sp. after 48 hours growth showing polysaccharide surrounding cell (bar = 1000nm)



Objectives

- Determination of amphiphilic character of EPS through evaluation of chemical composition of hydrophilic carbohydrates and uronic acids, and more hydrophobic proteins, through the use of Hydrophobic Interaction Chromatography (HIC).
- Pu(IV,V) partitioning to EPS from Pseudomonas with(w/) and without(w/o) proteins.





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GC-MS Spectra of EPS (C.-C. Hung): Clostridium sp. Exudates (> 6 kDa)





Hydrophobicity and MW determinations:



- Waters HPLC
- Amersham HiTrap, 2 butyl
- 1 ml columns in series



- Waters HPLC
- Tosoh Biosciences
 G4000 PWxl, 7.8 mm
 x 30 cm, particle size
 10 μm, with guard
 column 6 mm x 4 cm

Results HCA, Protein, Charged Groups



Results Molecular Weight (MW) by SEC

Overlain SEC Chromatograms of Pseudomonas florescens





Summary of characterization work:

- The neutral monosaccharides in this EPS consist of rhamnose, fucose, ribose, arabinose, xylose, mannose, galactose and glucose.
- The acidic groups in this EPS are mainly composed of carboxylic acid and minor polyanionic groups, e.g., sulphate and phosphate.
- 45 70 % of total carbohydrates are uronic acids, and total carbohydrates made up 26-31% of organic carbon.
- Besides the neutral and acidic sugars in the EPS, EPS also contained 9 % of proteins (% of carbon), which makes the EPS amphiphatic (amphiphilic).



Pu / EPS complexes

- Evaluation of Pu binding sites ('empirical adequacy'): affinity distribution
- Determination of conditional Pu /EPS complexes: ligand competition method



Comparison of EPS 'ligand' specific concentrations



Bacterial EPS type

Comparison of Pu / EPS binding



Microbially-enhanced Pu mobilization

- Soil analysis
- Batch studies
- 'Static' columns

Radioactive Properties of Key Plutonium Isotopes

Isotope	Half- Life (years)	Specific Activity (Ci/g)	Decay Mode	Alpha (MeV)	Beta (MeV)	Gamma (MeV)
Pu-238	88	17	α	5.5	0.011	0.0018
Pu-239	24,000	0.063	α	5.1	0.0067	<
Pu-240	6,500	0.23	α	5.2	0.011	0.0017
Pu-241	14	100	β	<	0.0052	<
Pu-242	380,000	0.0040	α	4.9	0.0087	0.0014

[US DOE, Plutonium Fact Sheet, ANL, 2001]

^{239,240}Pu: *in situ* ('aged)
²⁴¹Pu: tracer
²⁴²Pu: tracer

µ-XRF of Rocky Flats Soil



[1 mm² map of Fe (red) and Sr (blue) of 'as-rec'd' RF soil; incident beam energy 17.5 KeV] [Microbeam (10 x 20 µm spot) X-ray fluorescence of RF soil elements characteristic of loam/sandy soil]

[Analyses performed at NSLS Envirosuite beamline X27A; note that 'as-rec'd' the soil contained 1.6 ng ^{239,240}Pu g⁻¹ dry wt., 5 mg Fe g⁻¹ dry wt. (0.5 wt. %), 1.97% TOC]

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Inelastic

Compton

Zr

15

Sr



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Biotransformation of Pu in Soil from Rocky Flats

Incubation time = 10 days



Biostimulation of RF soil:

A. The pH dropped to 7 due to metabolism of lactate; and to 5 due to glucose fermentation. 25 days, E_H glucose = -180 mV, lact. = -123 mV.

B. Glucose fermentation released 50 wt. % of the total reducible iron oxide from the soil; lactate metabolism released 3.5 wt. %.



Formation of Pu Colloids in Soil due to Microbial Action



[²⁴²Pu nitrate (1.5 x 10⁻⁶M) added to soil resulted in 1) inorganic colloid formation, 2) sorption to soil and 3) remobilization as a colloid (<0.45 μ m) upon incubation with glucose and lactate. <0.1% of total ²⁴²Pu was in this fraction of unamended samples.

^{239,240}Pu and ²⁴²Pu Fate at 77 Days Incubation



[Indigenous 239,240 Pu and spiked 242 Pu were mobilized (<0.45 μ m) from RF soil with both glucose and lactate; the majority of the ^{239,240}Pu in the 'as-rec'd' soil was associate with the organic and inert fraction.] [mol % of total added] 37

Re-distribution of ²⁴²Pu in Soil at 77 Days Incubation



[²⁴²Pu was recovered with the reducible iron oxide fraction (CBD extraction) in unamended samples, however, it was redistributed to another phase after biostimulation with glucose or lactate]

²⁴²Pu and Total Carbohydrates



[At 77 days, the colloidal ²⁴²Pu was correlated with an increase in suspended carbohydrates (<0.45 μ m); this indicates that microbial exudates may play a role in Pu mobilization in the incubation experiments]

Summary of Soil Biotransformation Studies

- Pu was below detection for microprobe XRF, however discrete Fe phases were observed.
- Biostimulation with glucose released a significant amount of Fe while Fe(II) was readsorbed in lactate amended samples.
- A ²⁴²Pu spike rapidly sorbed to soil, however it was remobilized due to microbial action; under highly reducing, fermentative conditions 0.7% of the total was detected in the <0.45µm, >30kDa fraction.
- The majority of Pu was released with the reducible iron oxide fraction of unamended samples, however this was not the case after biostimulation.
- The indigenous ^{239,240}Pu was remobilized, to a lesser extent than the spike, but this may be due to different mineralogical association (majority resided with the organic and inert fraction).
- There was an increase in soluble carbohydrates in the biostimulated samples implicating microbial exudates in stabilizing Pu in the colloidal fraction.



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'Static column' experiments

Assess Pu mobility under solid / solution ratios appropriate to *in situ* conditions

Static Column Incubation Experiment: 10 vs. 47 Day Incubation



 239,240 Pu = ~70 pCi / g; 0.5 w/v glucose; 0.015% w/v NH₄Cl

Pu transport enhancement by EPS



Sorbed Pu-241 to RF soil for 24hrs then 22 mg/L EPS (~ 10 mg/L OC) injected

EPS Facilitated Pu Mobilization







<u>Summary:</u> <u>Pu mobility as a transformational process</u>



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