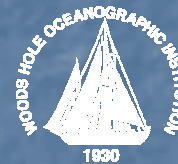


Understanding plutonium variability and mobility in groundwater at the Savannah River Site

K. Buessler¹, M. Dai¹, D. Kaplan² and S. Peterson³

1. *Woods Hole Oceanographic Institution*



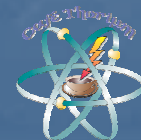
2. *Savannah River National Lab*



3. *Pacific Northwest National Lab*



- with help from S. Pike, J. Kelley, T. Maiti & MANY others



Outline

- **Introduction**

- Plutonium & DOE interests

- **Methods matter**

- Innovative approaches for groundwater Pu studies

- **Savannah River F-area field data**

- Pu concentrations
- Pu colloid abundances
- Pu redox state
- Pu isotope ratios

Variability in Pu isotope ratios, redox state and colloid associations attributable to Pu source effects and groundwater chemistry



Plutonium & DOE Interests

Plutonium is predicted to be insoluble and highly sorbed in subsurface environment, however:

- early studies showed high groundwater colloidal % (Kaplan et al)
- evidence for far field transport (Kersten et al)
- but, new methods show low colloid association (Dai et al)

If Pu variability can be understood/parameterized:

For remediation

- this allows for design strategies to retard Pu, save time and money, develop improved models to predict remediation impacts

For stewardship

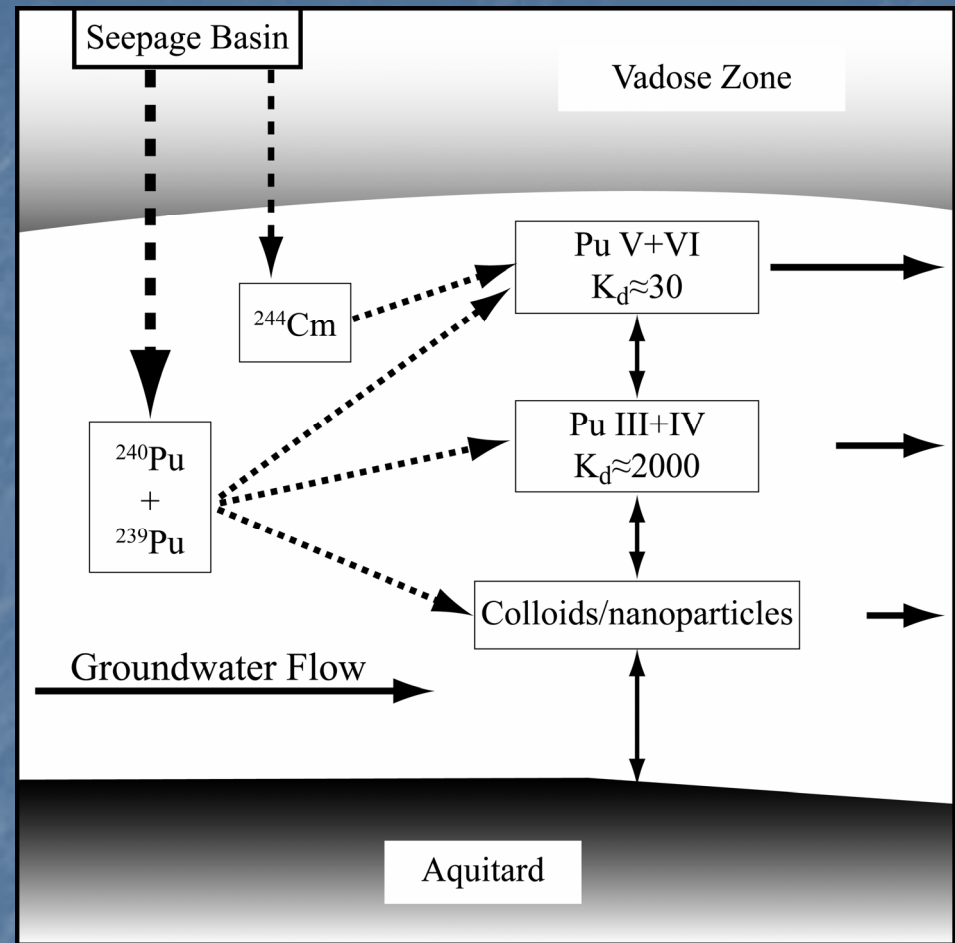
- this allows for better project management, prediction of off site migration & improved monitoring practices



Conceptual diagram of the fate of Pu released into the subsurface environment

Source dependent behavior
- Pu source controls

Source independent behavior
- Pu speciation controls via coupled biogeochemical processes



Fate of Pu is isotope specific and dependent upon temporal changes in groundwater conditions



Methods matter

In groundwater, Pu exists in multiple chemical forms/species
(Pu III, IV, V, VI; solution complexes, colloid associations)
and at concentrations $<10^6$ atoms per liter in different fractions
(= 0.0003 pCi/kg or 0.001 Bq/kg)

Well sampling

- “micro-purge” 150 ml/min & monitor groundwater geochemistry
- Pu concentration increases $>3x$ with standard well sampling (16L/min)



Methods matter

- **Well sampling**
‘micro-purge’ 100’s ml/min & monitor gw geochemistry
- **Redox control**
 - N_2 controls during sampling & processing; immediate in field processing for colloids and redox state
- **Cross-flow filtration**
 - calibration; mass balance checks for loss and contamination

Mass balance (2004)
CFF \pm 2-5%

CFF system
 $>1kD$ to $<0.2 \mu m$ = colloidal

N_2 glove bag

4L sample reservoir
($>1kD$)

Permeate out line
($<1kD$)

1kD Millipore CFF ($1m^2$)

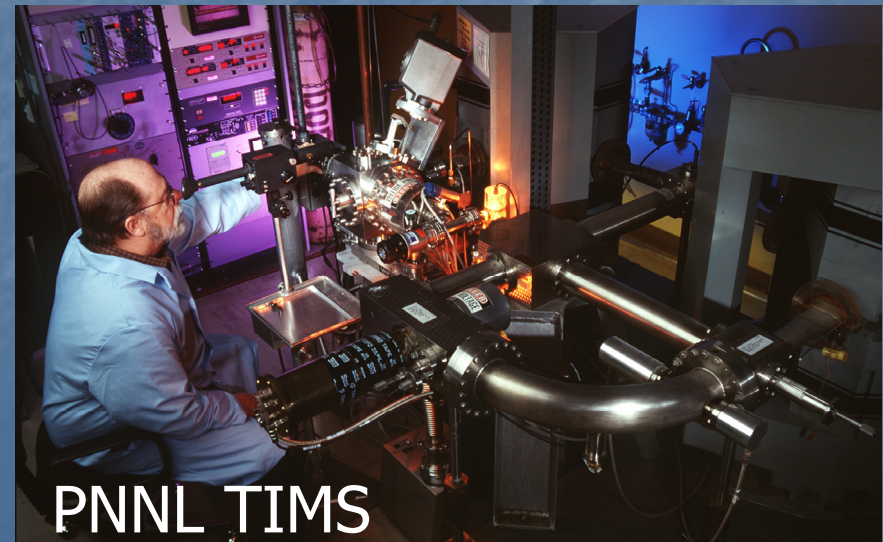
Direct well feed



Methods matter

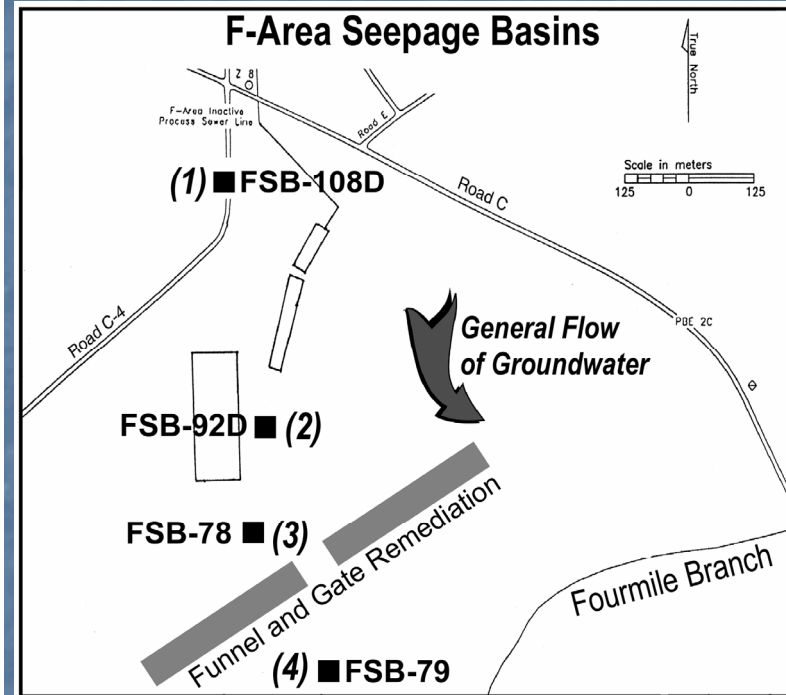
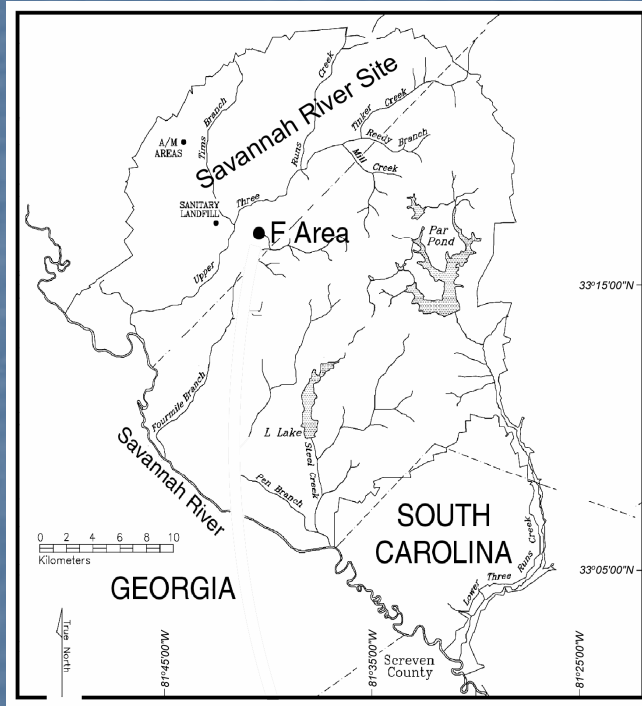
- **Well sampling**
 - 'micro-purge' 100's ml/min & monitor geochemistry
- **Redox control**
 - N₂ controls during sampling & processing; immediate in field processing for colloids and redox state
- **Cross-flow filtration**
 - calibration; mass balance checks for loss and contamination
- **Attention to blanks/TM clean methods**
 - blank levels of 10⁴ atoms/sample
- **Thermal Ionization Mass Spectrometry**
 - identify separate Pu isotopes at environmental levels

Portable clean field lab



SRS F-area Seepage basins

- waste from reactor separation facilities- nitric acid soln.



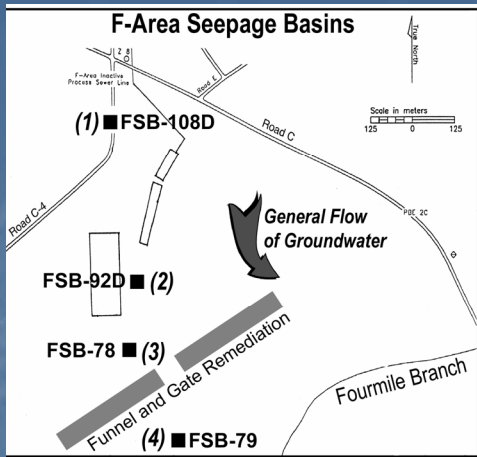
Project sampling in 1998 and 2004

Funnel and gate remediation since 2002/2003

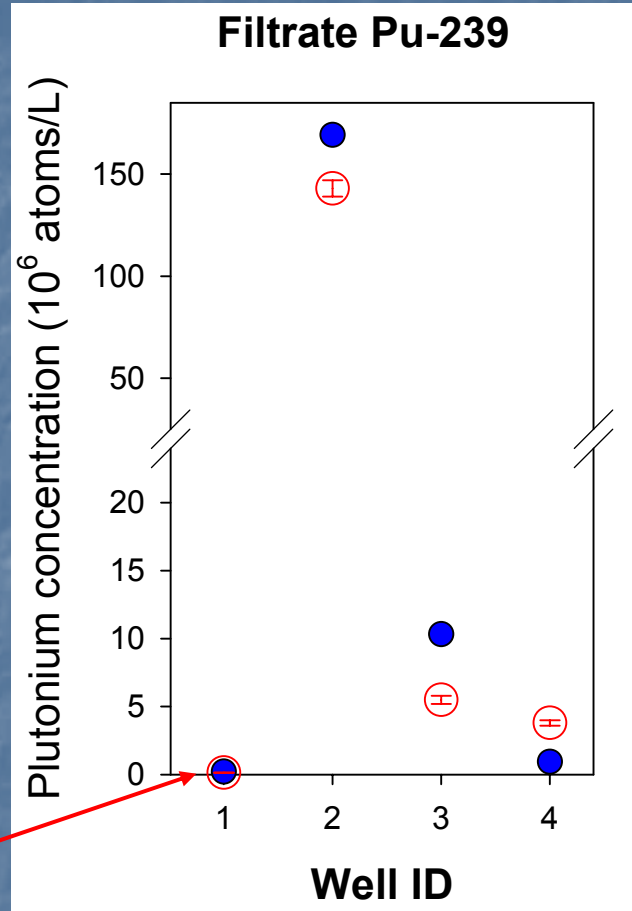
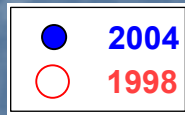
Millions liters of sodium hydroxide-sodium bicarbonate soln.

Increase pH from 4 to 6





Sharp decrease Pu-239 downstream from source

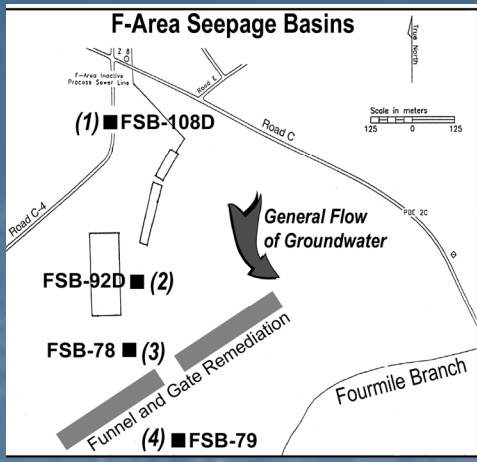


(Filtrate < 0.2 μm)

$^{239}\text{Pu} = 0.14\text{-}0.22 \times 10^6$

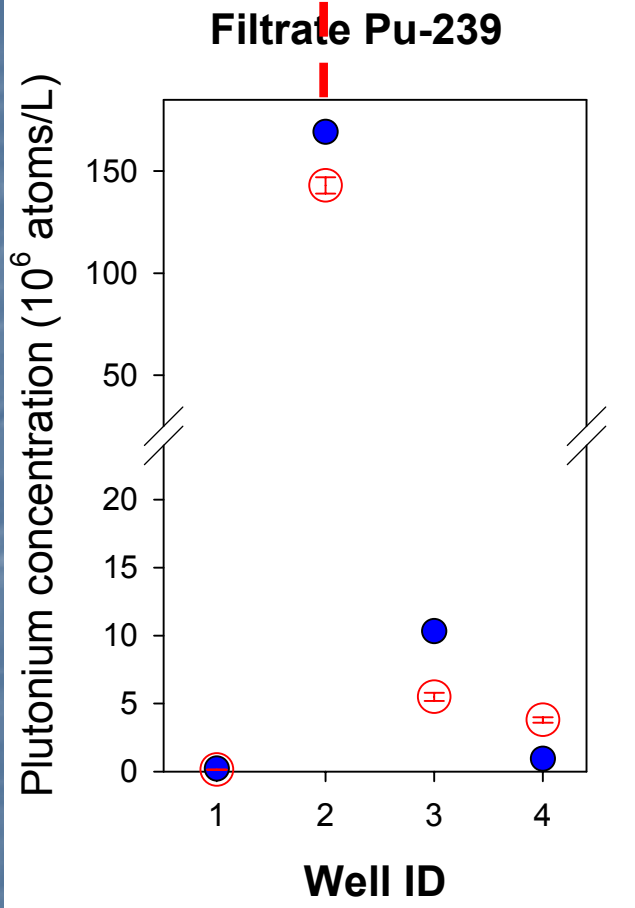
Reproducible data at $10^5 - 10^8$ atom/L levels
 Pu highest in well #2 (near seepage basin)
 Similar in 1998 and 2004 (but not identical)

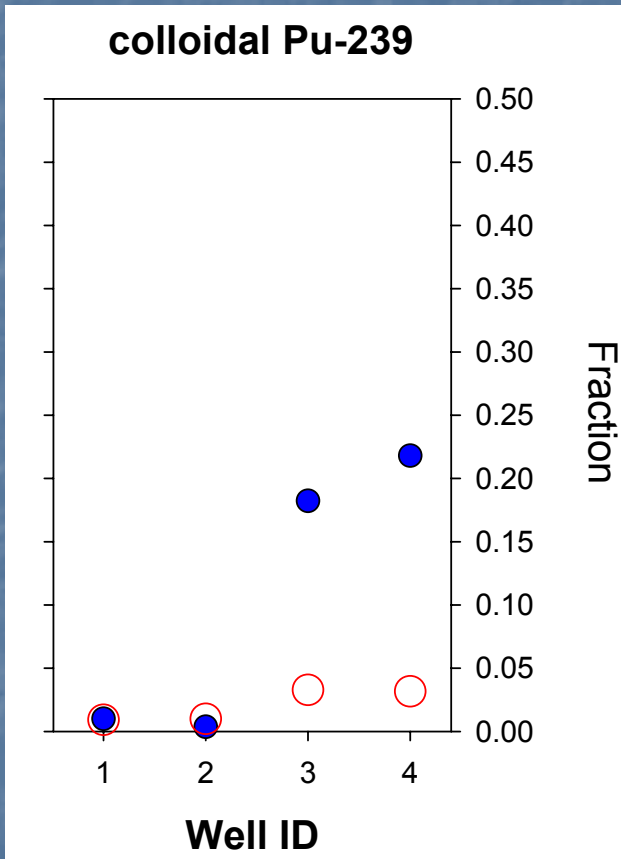
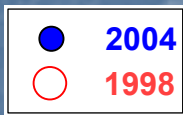
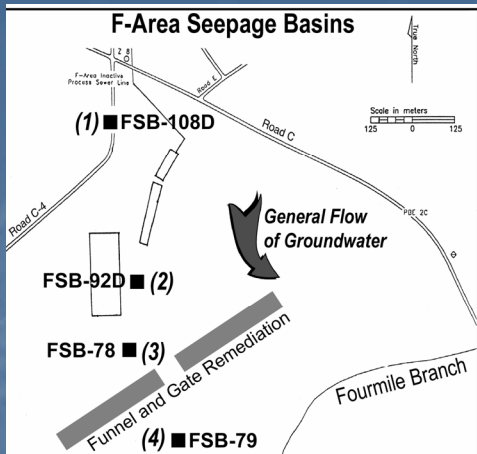




$^{239}\text{Pu} = 500 \times 10^6$ at 16L/min
Well pumping rate important

● 2004
 ○ 1998





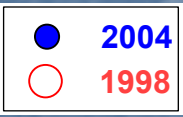
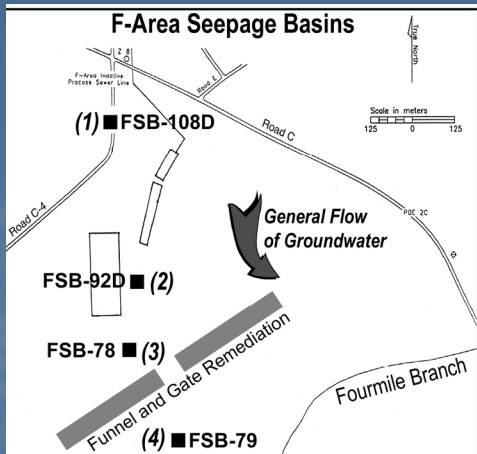
Pu-239 largely <1kD, non-colloidal

% ²³⁹ Pu colloidal	1998	2004
Well 1&2	<1%	<1%
Well 3&4	3%	20%

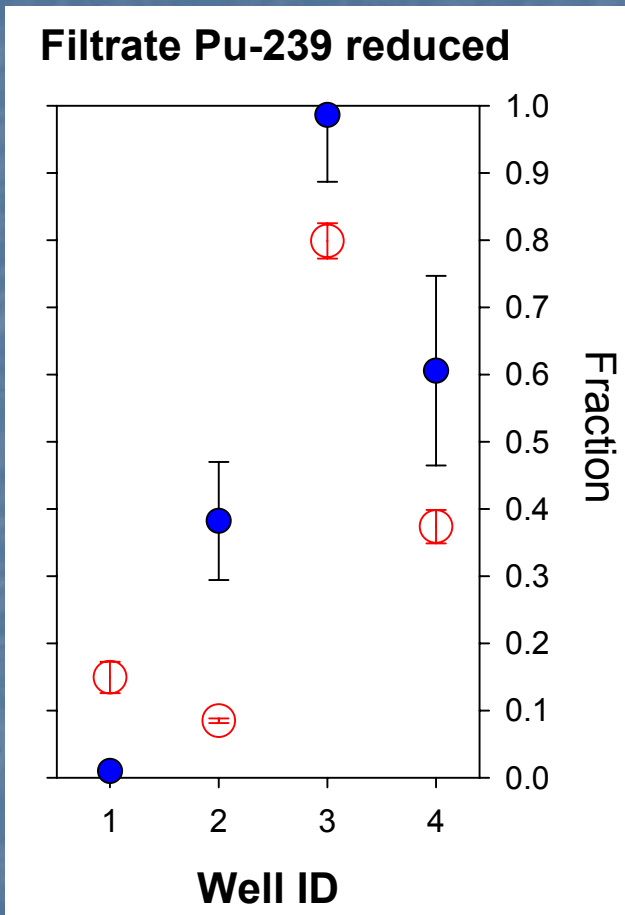
Other sites/same methods
 Hanford K-area 10-30%
 SRS Pond B 40-75%

Significant difference 1998 vs. 2004





Pu-239 fraction reduced higher downstream

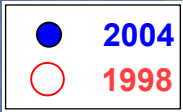
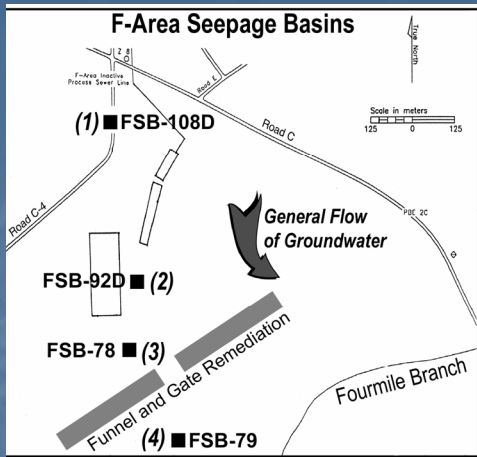


% ²³⁹ Pu reduced	1998	2004
Well 1&2	10-15%	1-40%
Well 3&4	35-80%	60-100%

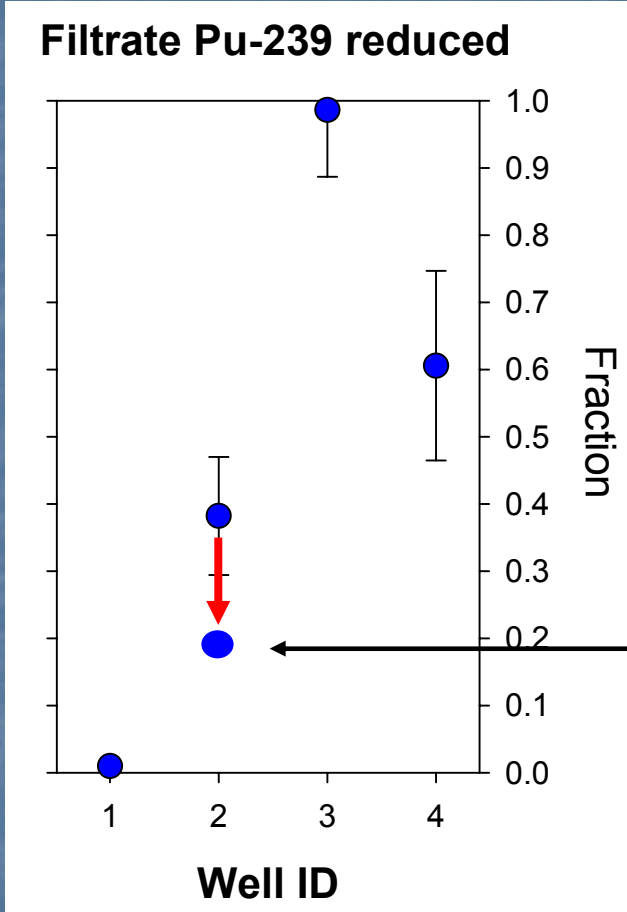
Other sites/same methods
 Hanford K-area 65%
 SRS Pond B 70-100%

Fraction reduced higher in wells with higher colloidal %
 Variable, but some differences 1998 vs. 2004



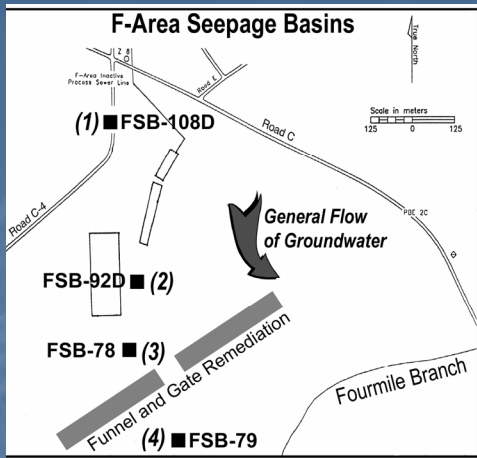


Pu-239 fraction reduced higher downstream



Fraction reduced drops from 38% to 18% after 3 days air exposure

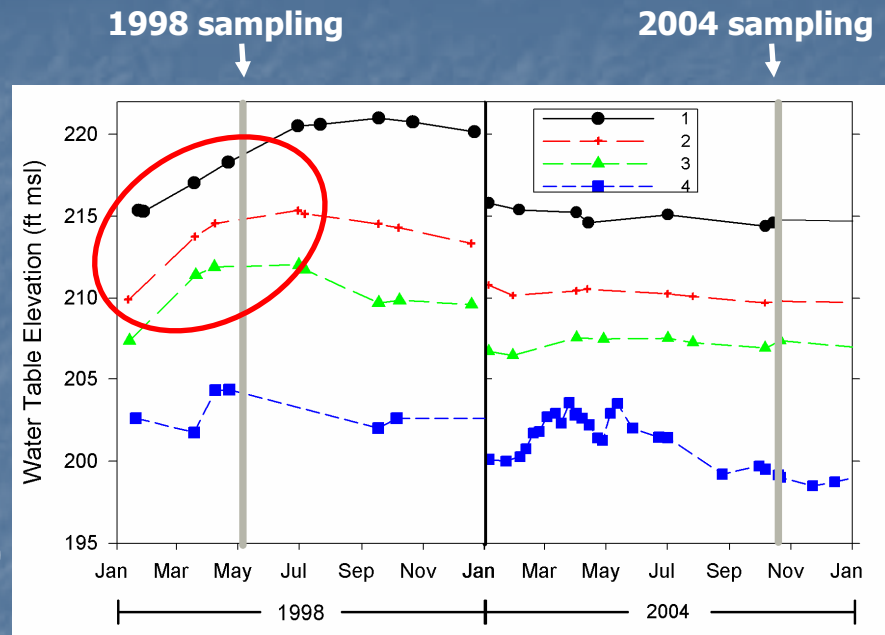




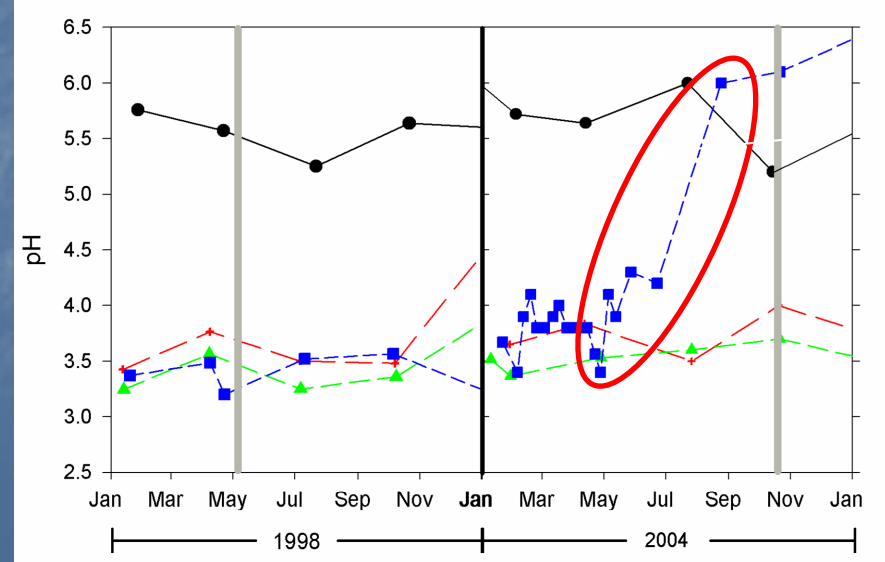
Source independent controls

- did groundwater geochemistry impact ^{239}Pu speciation?

1998
"freshening"
event
(largest in 6 yrs)



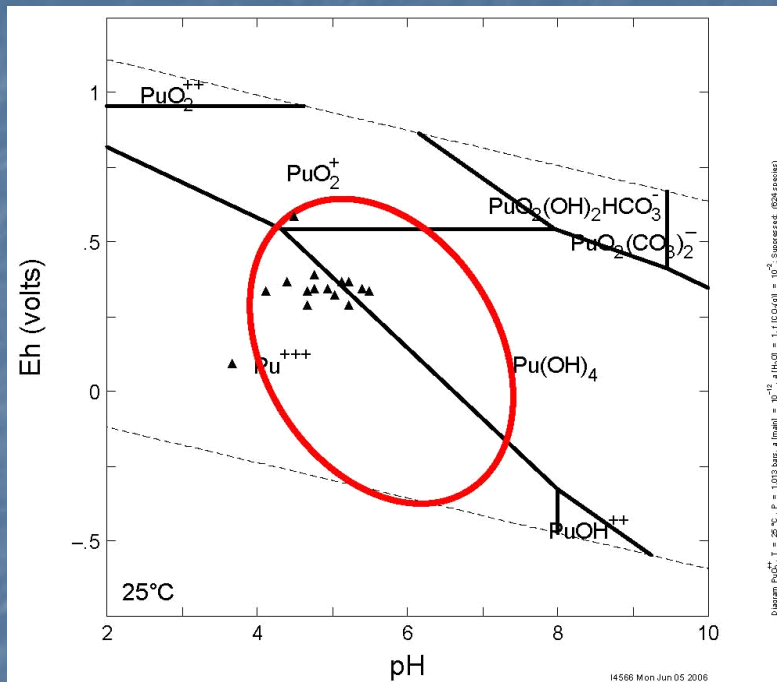
1998
more oxidized,
lower % colloidal



Note remediation
impacts pH well #4
-lower Pu and
esp. ^{244}Cm in 2004

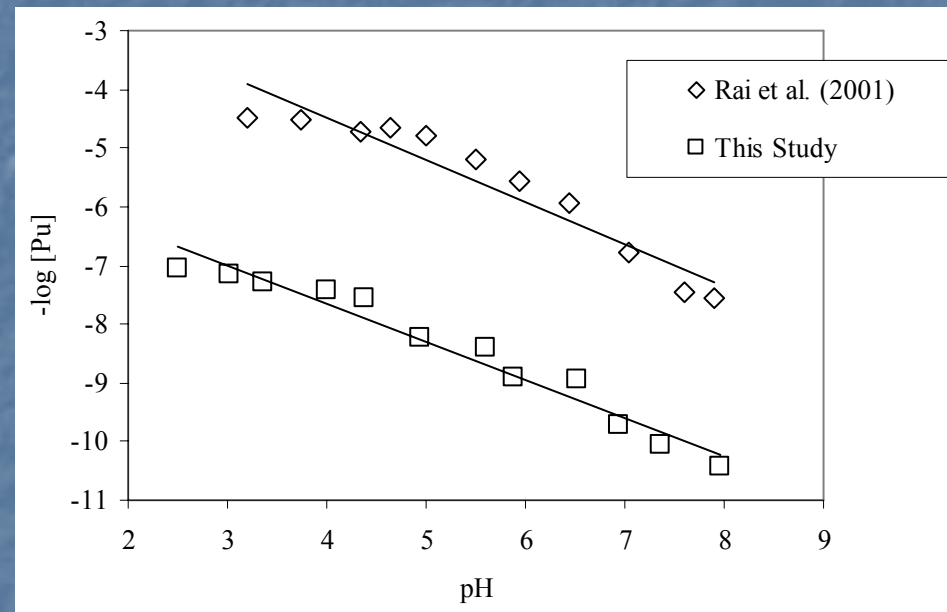


Small changes to groundwater chemistry greatly impact Pu speciation



pH/Eh of F-area plume is Pu(III/IV/V)

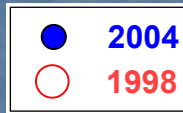
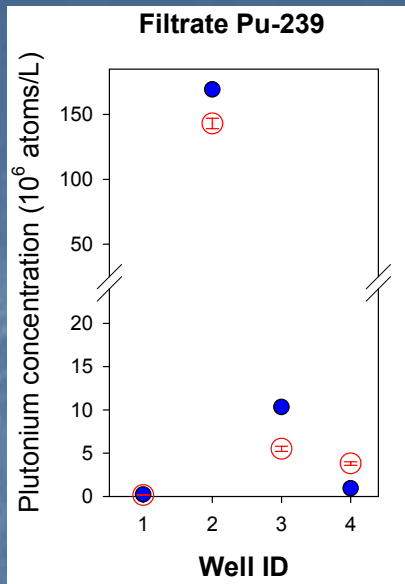
Pu(V) $K_d = 30 \text{ mL/g}$
 Pu(IV) $K_d = 2000 \text{ mL/g}$



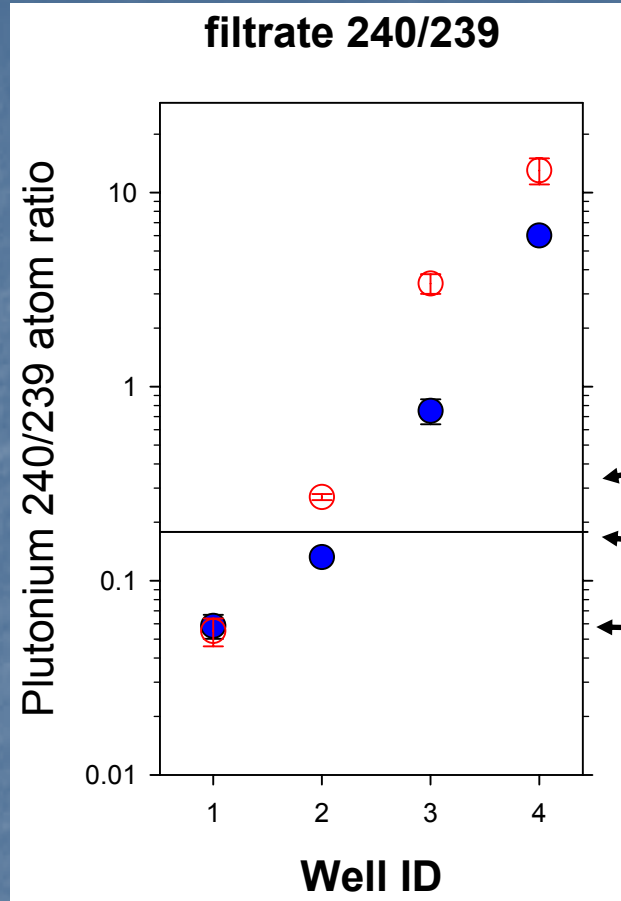
pH decrease from 6 to 5 results in a decrease solubility by 1 order of magnitude



Unusual increases in $^{240}\text{Pu}/^{239}\text{Pu}$ ratio downstream



^{239}Pu $t_{1/2} = 24,100$ yr
 ^{240}Pu $t_{1/2} = 6,560$ yr



$^{240}\text{Pu}/^{239}\text{Pu} > 10$

High yield tests & other reactor products = 0.3-0.4

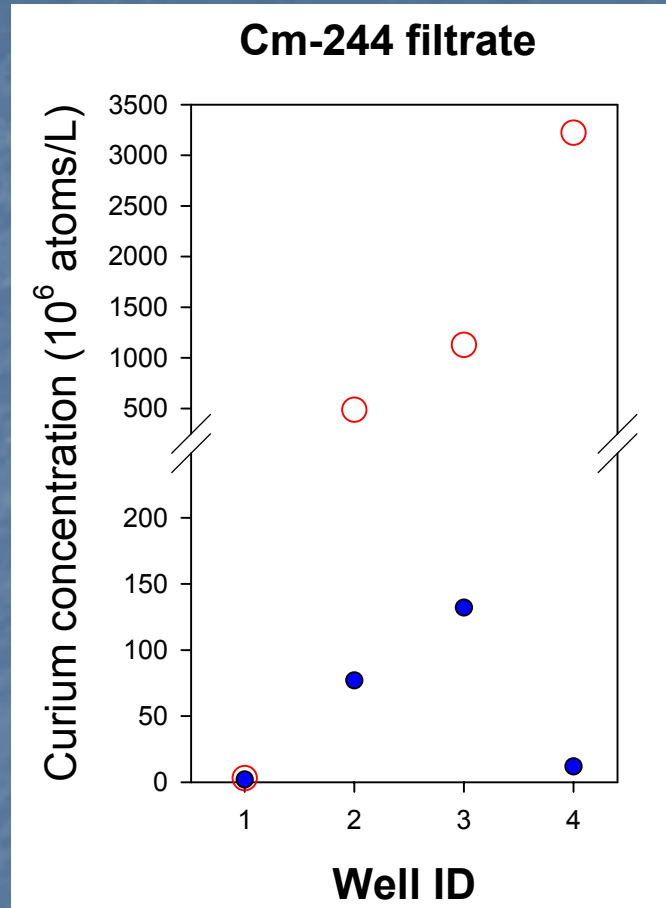
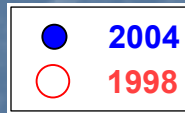
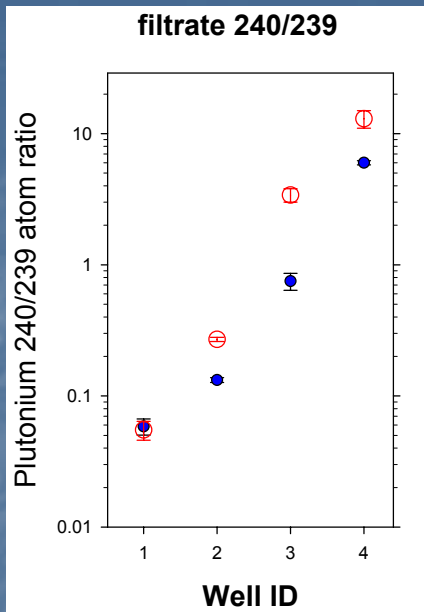
Global fallout = 0.18

Local SRS = 0.06

Local SRS Pu source in background well #1
 What is local source of ^{240}Pu ?
 Preferential transport ^{240}Pu in groundwater?



^{244}Cm - produced at SRS in 1960's

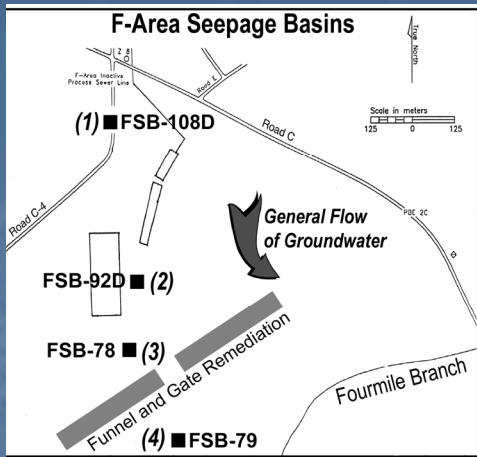


- ^{244}Cm -
- $t_{1/2} = 18.1 \text{ yr}$
- Alpha decay to ^{240}Pu
- Less particle reactive than Pu

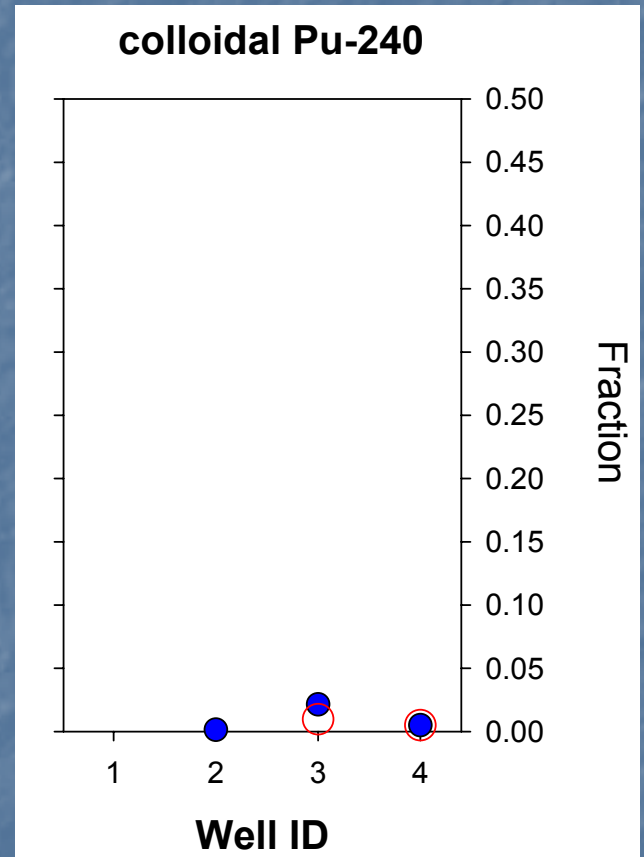
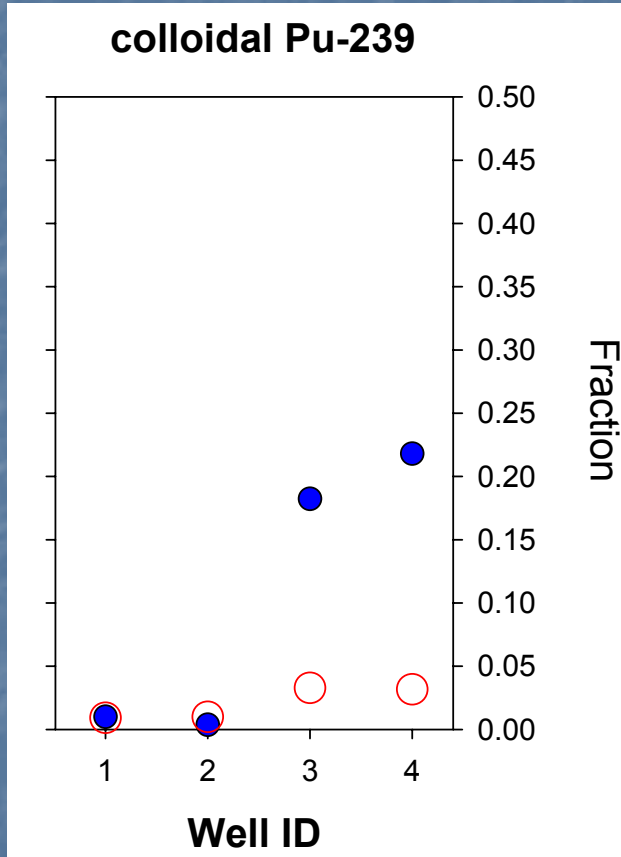
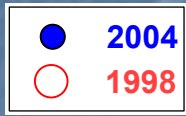
^{244}Cm concentrations 1-2 orders of magnitude higher than ^{239}Pu
Increasing 240/239 ratio due to ^{240}Pu production from ^{244}Cm decay
Less ^{244}Cm in 2004 than 1998-

$$K_D = 40 \text{ mL/g @ pH=4}; K_D = 15,000 \text{ mL/g @ pH=6.7}$$





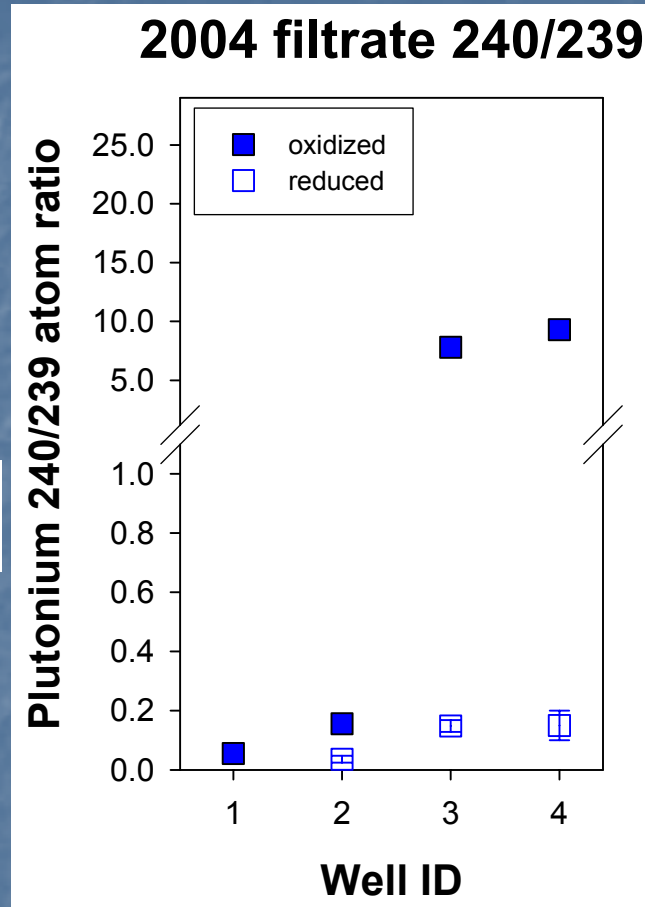
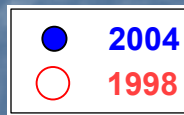
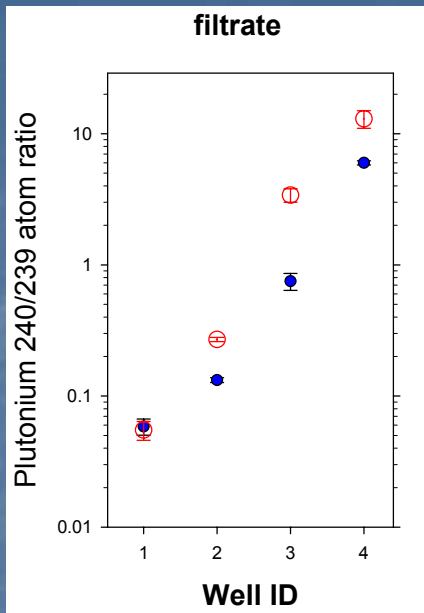
Source dependent controls on ^{240}Pu speciation



Less colloidal ^{240}Pu
 Less reduced- 1-25% in 1998; 5-55% in 2004



Source dependent controls on ^{240}Pu speciation



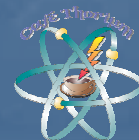
^{240}Pu produced from ^{244}Cm decay results in more oxidized forms
Isotope specific differences in apparent Pu mobility
Prior work w/ $^{239,240}\text{Pu}$? Other sites- Oak Ridge?



Summary

Variability in Pu isotope ratios, redox state and colloid associations attributable to Pu source effects and groundwater chemistry

- **Methods matter**
 - Considerable effort devoted to improving gw Pu speciation methods
- **Are colloids important in F-area groundwater for Pu?**
 - low ^{239}Pu colloidal abundances (1-20%)
 - rapid decrease in ^{239}Pu concentration downstream from source (<1km)
- **Impacts of natural groundwater variability and remediation are seen in Pu concentrations and speciation**
 - 1998 "Freshening Event" - more oxidized/lower colloidal ^{239}Pu
 - remediation changes to pH result in lower ^{244}Cm , ^{239}Pu
- **^{240}Pu differs from ^{239}Pu**
 - ^{240}Pu found further downstream from source than ^{239}Pu
 - ^{244}Cm source results in more oxidized/mobile forms of ^{240}Pu



Future Needs

Field studies provide important insights into processes that impact Pu speciation and transport in the subsurface environment

- Continued use/development of reliable field methods for in-situ speciation
- Need time-series sampling to capture seasonal and episodic variability
- Consider groundwater methods intercomparison, multi-lab "Colloid cookout"?
- Improve parameterization for colloid reactive transport models

Recent references (used for this presentation)

Methods

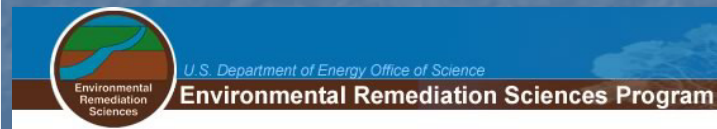
- Dai et al., 2001 (J. Envir. Rad. v53)
- Buesseler et al., 2003 (ES&T v37)
- Hasselov et al., 2006 (Sci. Total Envir., sub.)

Field

- Dai et al., 2002 (ES&T, v36)
- Dai et al., 2005 (J. Cont. Hydro., v76)

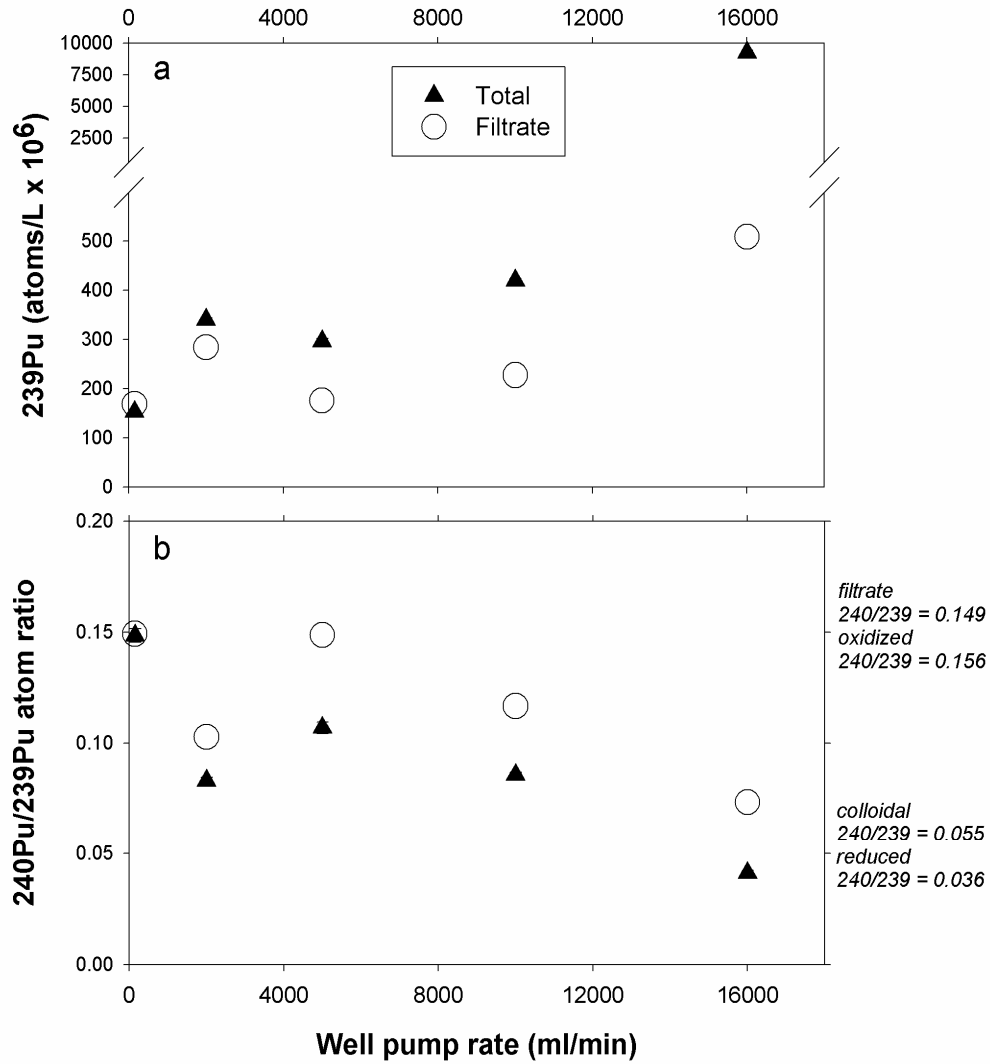
Lab

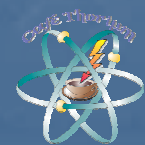
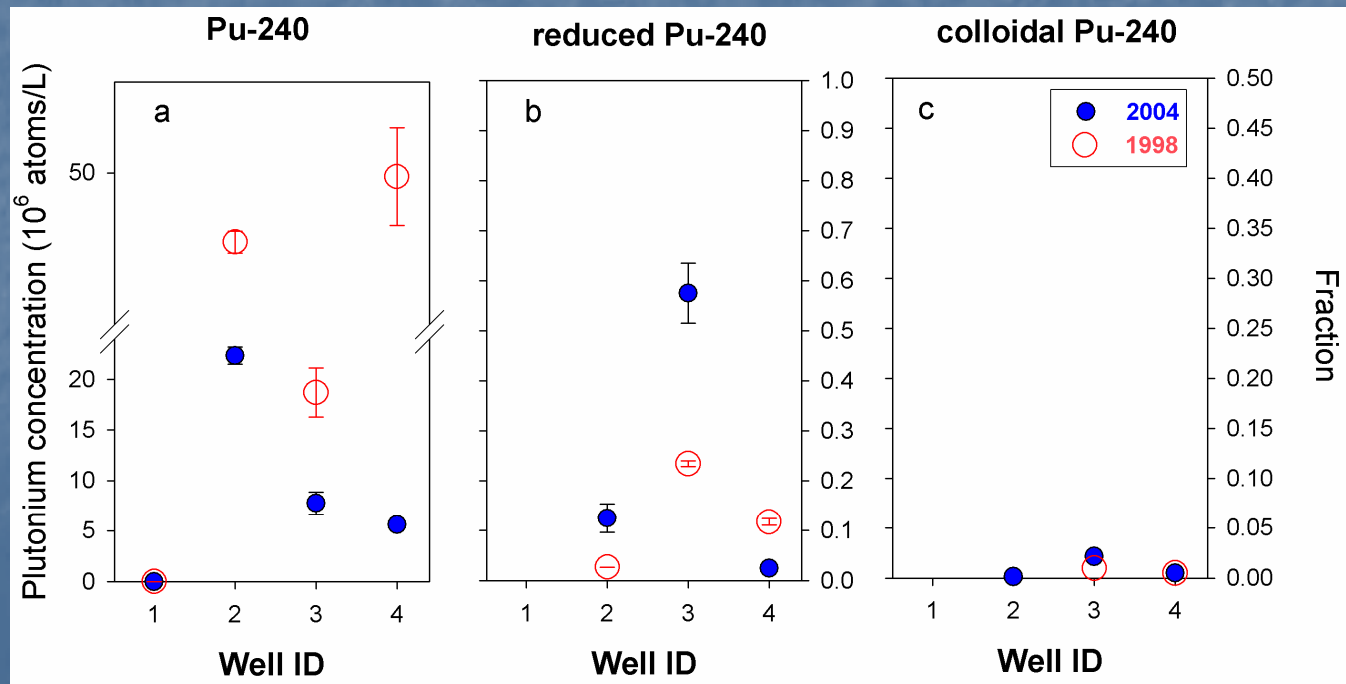
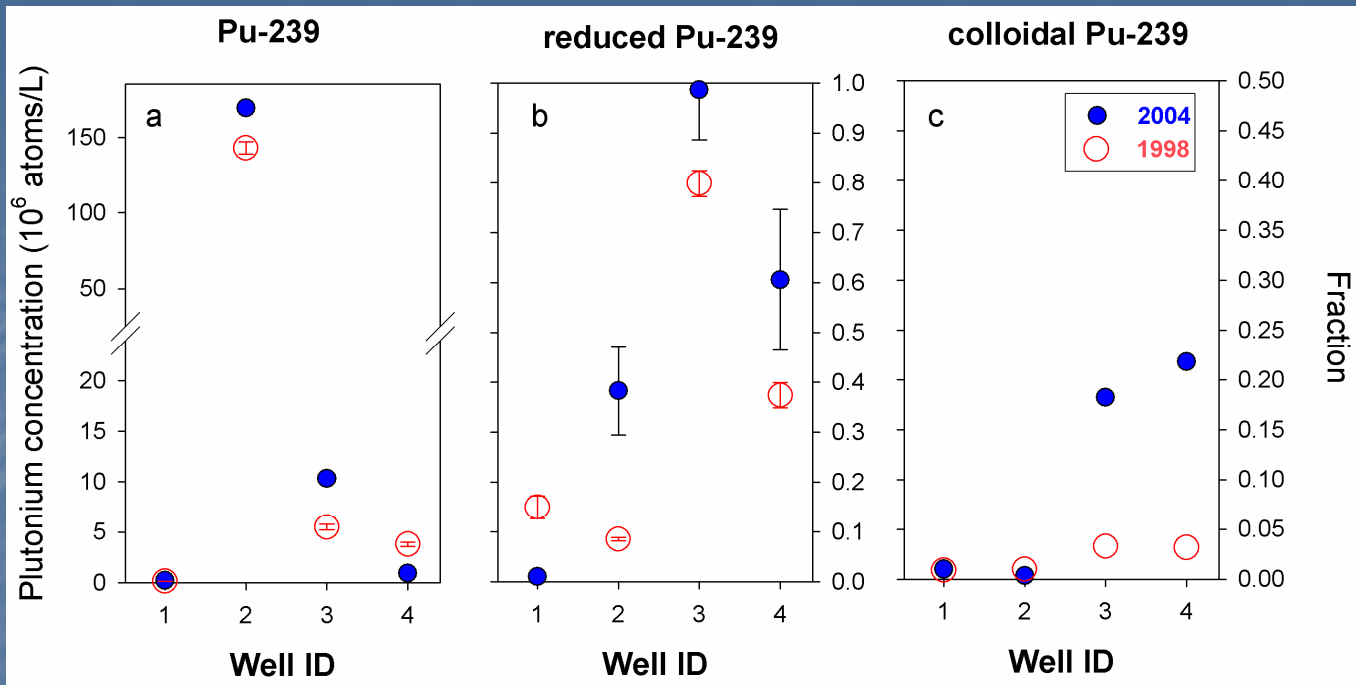
- Kaplan et al. 2004, 2006a,b (ES&T, v38, v40)
- Powell et al. 2004, 2005, 2006 (ES&T, v28, v39, v40)





Well 92D Flow Rate Experiment





Groundwater sampling and processing diagram

