



# Speciation, Fate, and Cycling of Arsenic in Subsurface Environments

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Z=33

As

Atomic wt.  
74.922

## Mechanistic understanding of arsenic speciation can help predict its behavior in subsurface environments

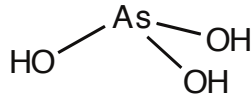
- Can we use geochemical scenarios to categorize potential As mobilization?
- How do we optimize kinetics of biogeochemical processes to enhance natural As attenuation?

### Geochemical Parameters:

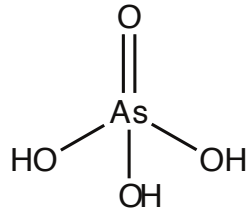
- Amount of labile iron
- Amount of sulfur available for reduction/oxidation
- pH & Eh (local and gradients)
- Role of nitrogen species?

# Arsenic Speciation in the Environment

## Inorganic Arsenic



Arsenious acid or  
Arsenite (As<sup>III</sup>(OH)<sub>3</sub>)  
pK<sub>a1,2,3</sub> = 9.23, 12.13, 13.40

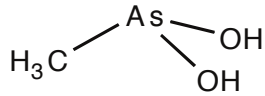


Arsenic acid or  
Arsenate (H<sub>3</sub>As<sup>V</sup>O<sub>4</sub>)  
pK<sub>a1,2,3</sub> = 2.20, 6.97, 11.53

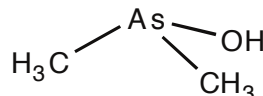
## Arsenic Minerals

Orpiment (As<sub>2</sub>S<sub>3</sub>)  
Realgar (As<sub>4</sub>S<sub>4</sub>)  
Arsenopyrite (FeAsS)  
Scorodite (FeAs<sup>V</sup>O<sub>4</sub>·2H<sub>2</sub>O)  
Oxides, Arsenites, Arsenates

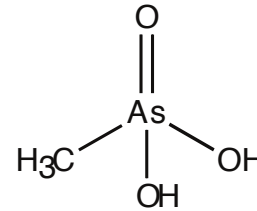
## Methylated Arsenic Compounds



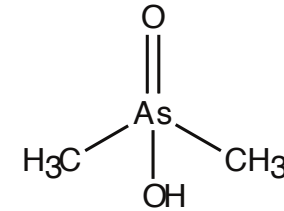
Monomethylarsonous  
acid or MMA<sup>III</sup>  
(As(OH)<sub>2</sub>CH<sub>3</sub>)



Dimethylarsinous  
acid or DMA<sup>III</sup>  
(As(OH)(CH<sub>3</sub>)<sub>2</sub>)

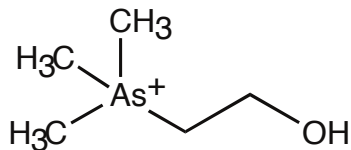


Monomethylarsonic  
acid or MMA<sup>V</sup>  
(AsO(OH)<sub>2</sub>CH<sub>3</sub>)

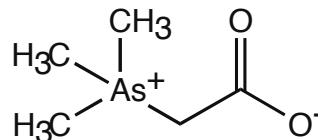


Dimethylarsinic  
acid or DMA<sup>V</sup>  
(AsO(OH)(CH<sub>3</sub>)<sub>2</sub>)

## Organoarsenic Compounds

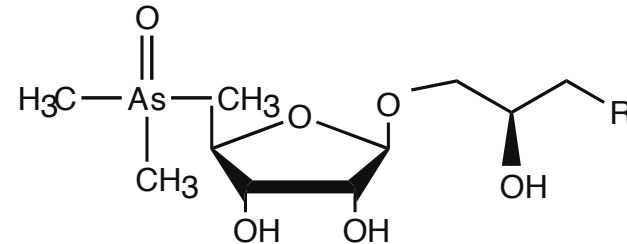


Arsenocholine  
(CH<sub>3</sub>)<sub>3</sub>As<sup>+</sup>CH<sub>2</sub>CH<sub>2</sub>OH)



Arsenobetaine  
(CH<sub>3</sub>)<sub>3</sub>As<sup>+</sup>CH<sub>2</sub>COO<sup>-</sup>)

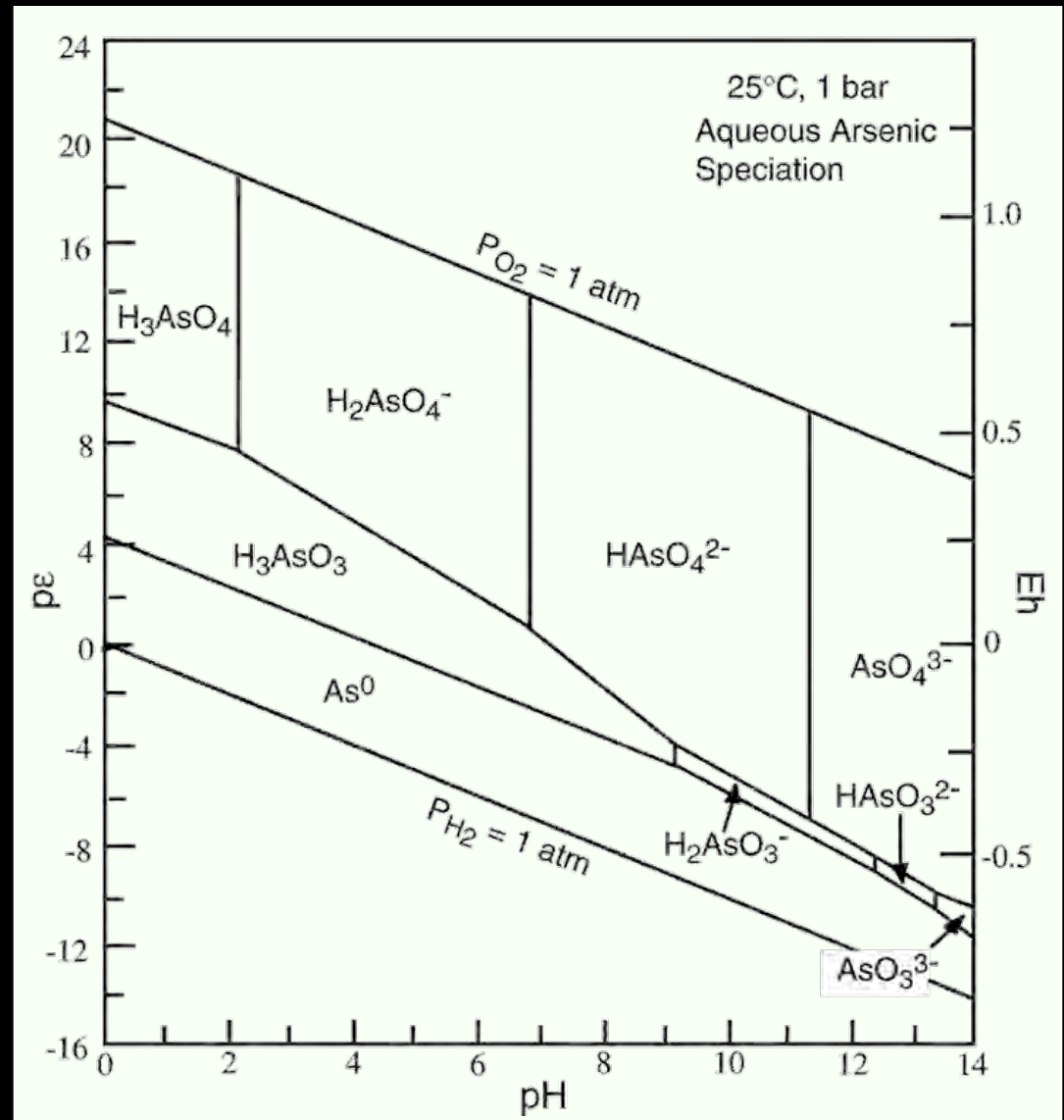
## Organoarsenic Lipids



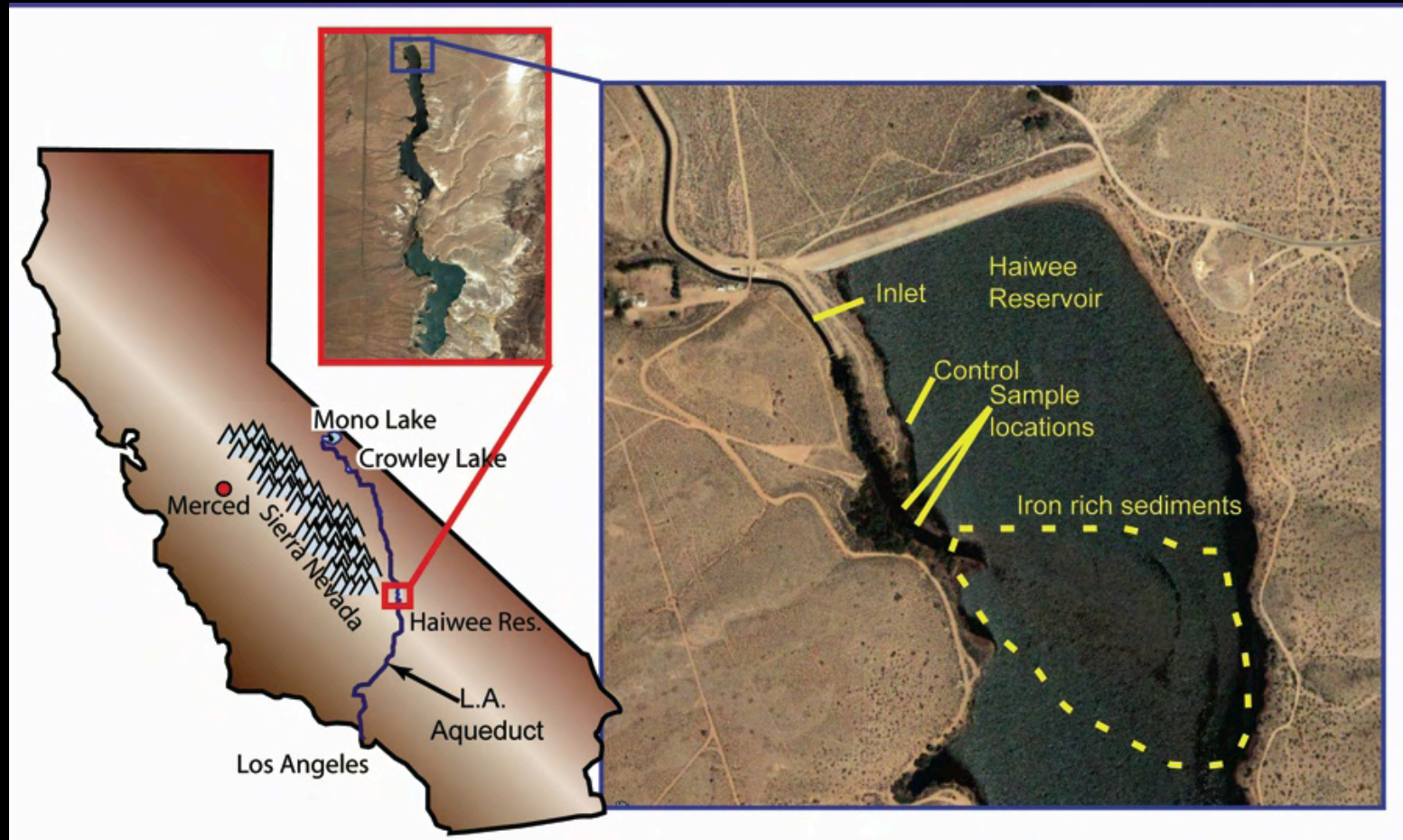
R = OH  
R = OP(O)(O<sup>-</sup>)OCH<sub>2</sub>CH(OH)CH<sub>2</sub>OH  
R = SO<sub>3</sub><sup>-</sup>  
R = OSO<sub>3</sub><sup>-</sup>

# Arsenic Speciation & Partitioning: Tied to Major Element Chemistry

- Precipitation important for sulfides and sulfates
- Adsorption: Strongly associates with Iron hydroxides/oxides; competitive sorbates?
- Organic carbon and microbial activity
- Microbial coupling/competition with Nitrogen species?



# Haiwee Reservoir, Owens Valley

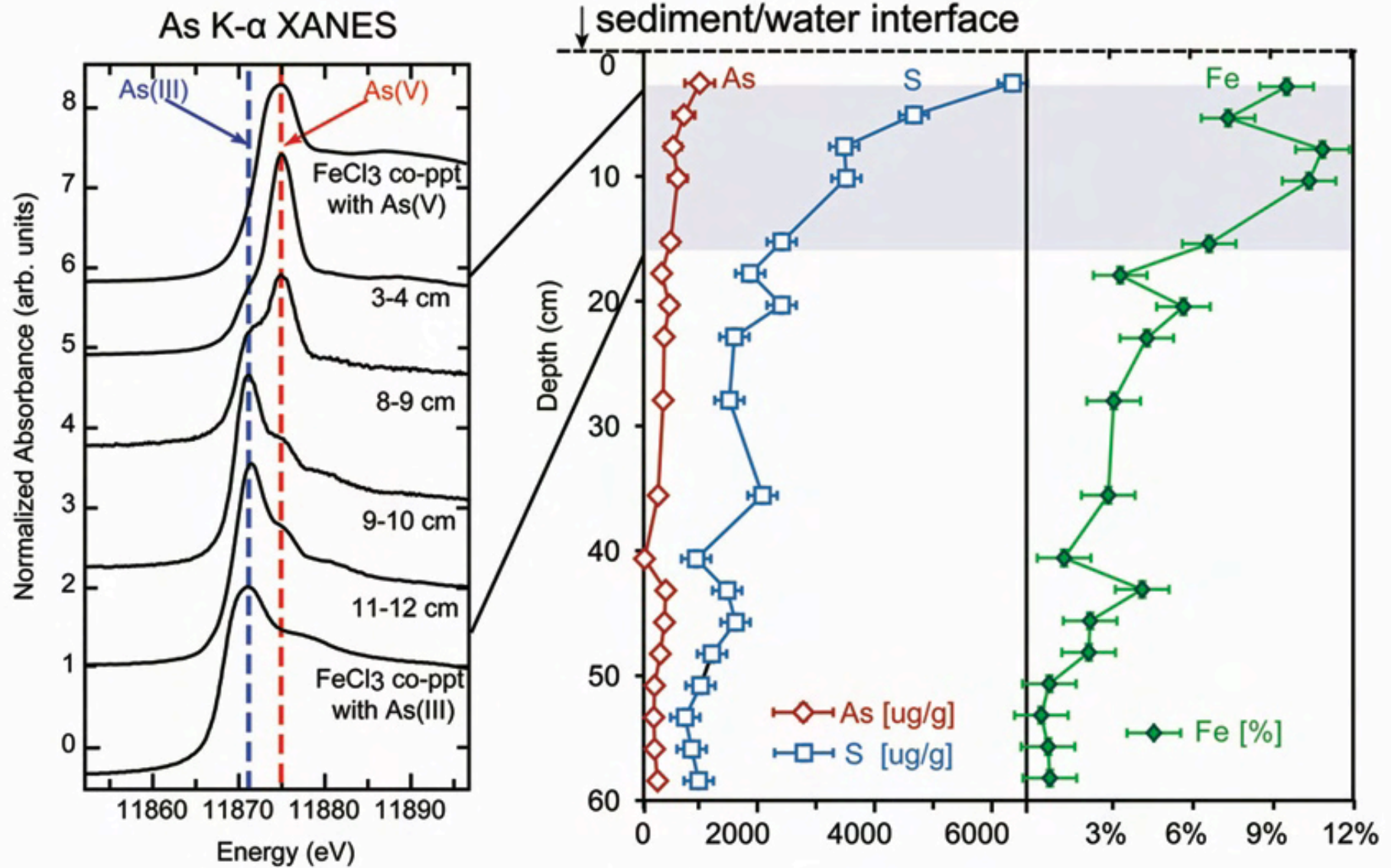


- Aqueduct water dosed with  $\text{FeCl}_3$  to remove As
- Deposition of high Fe, low S sediments with sorbed As(V)

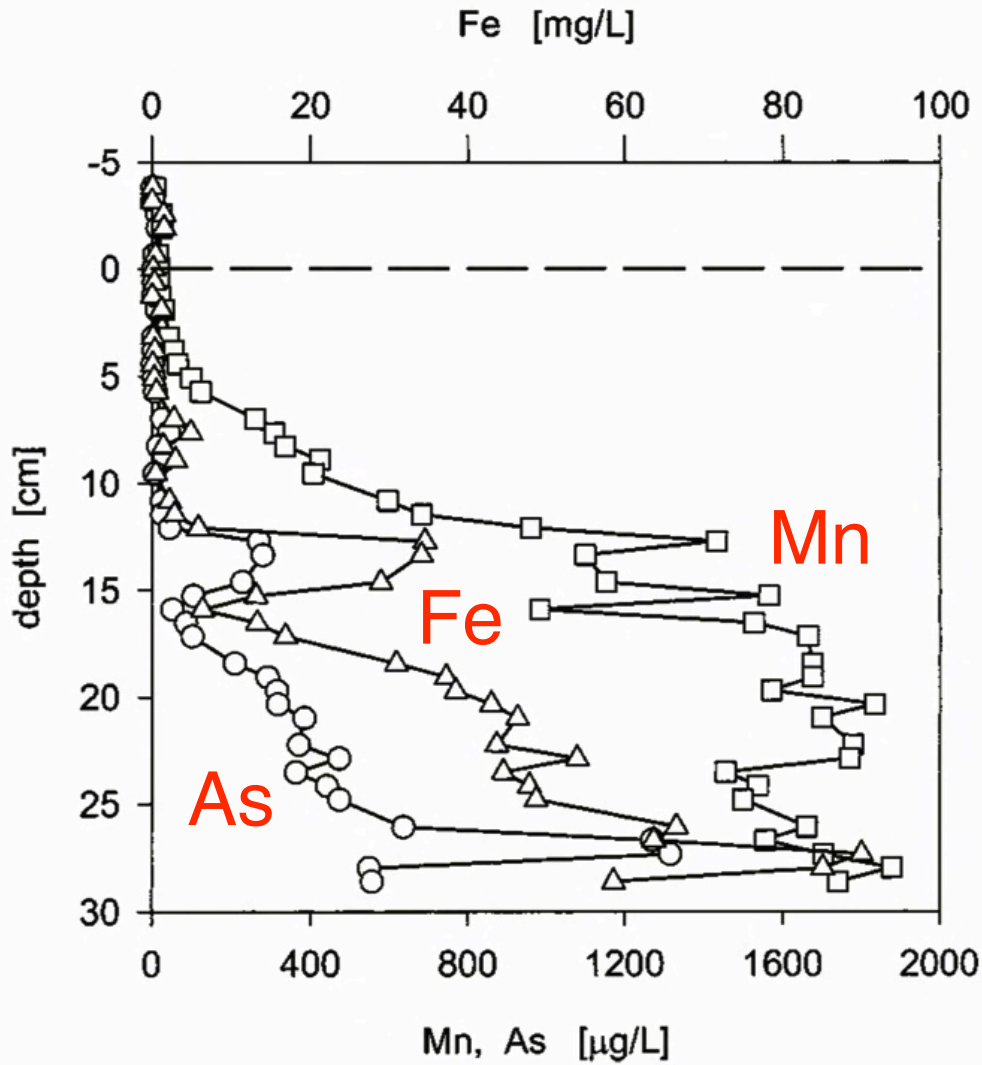
# Haiwee Reservoir, Owens Valley



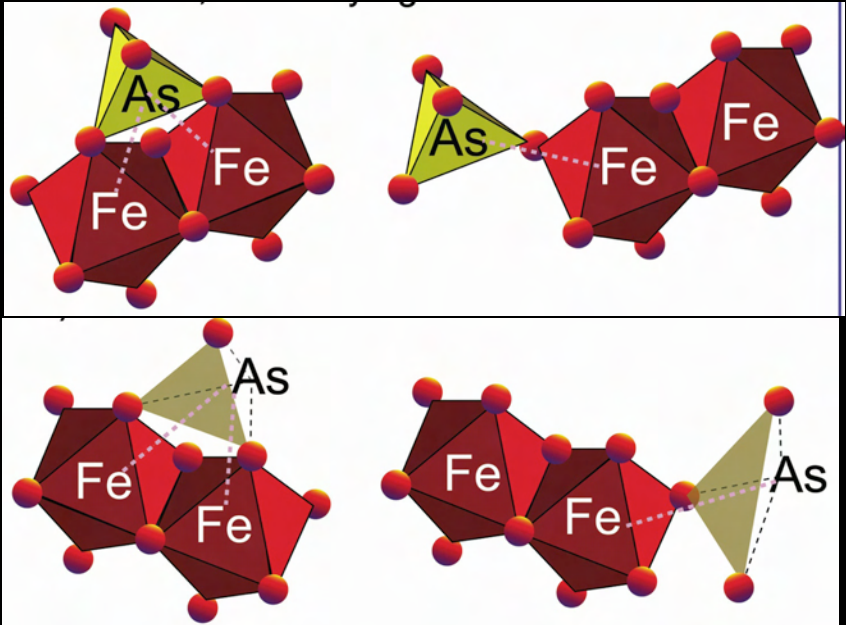
# Haiwee Reservoir: Core Sediments



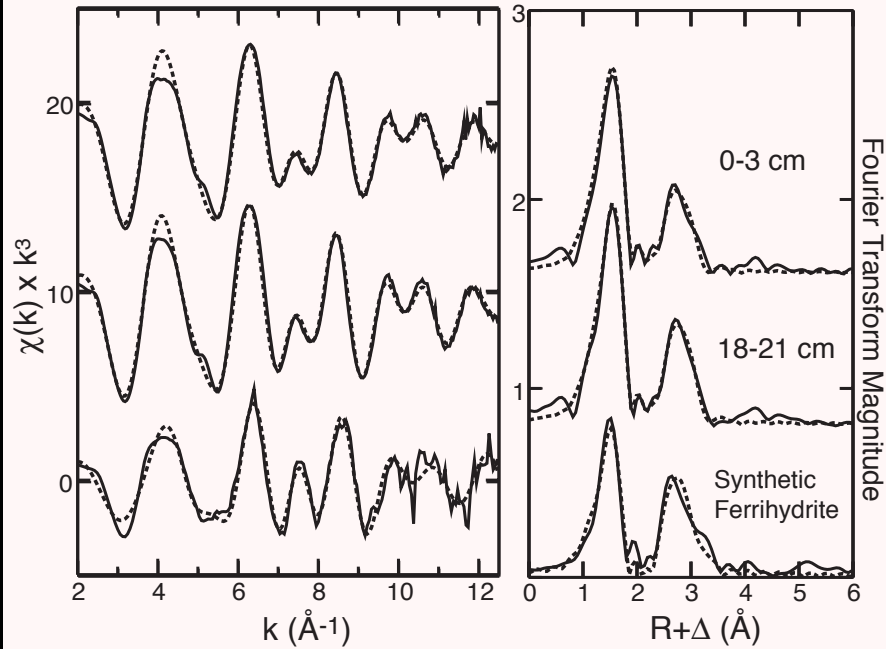
# Porewater Concentrations



# Sediment As & Fe Speciation



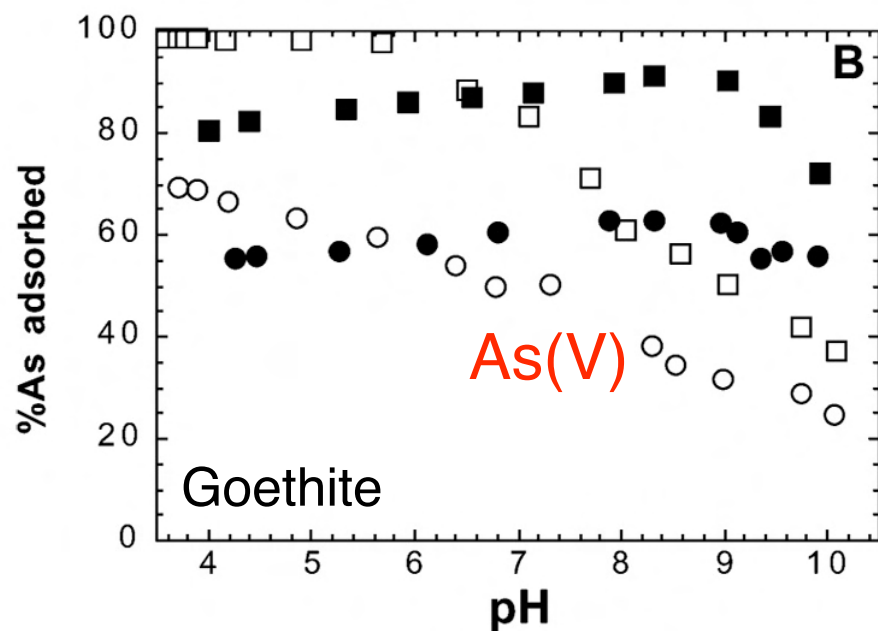
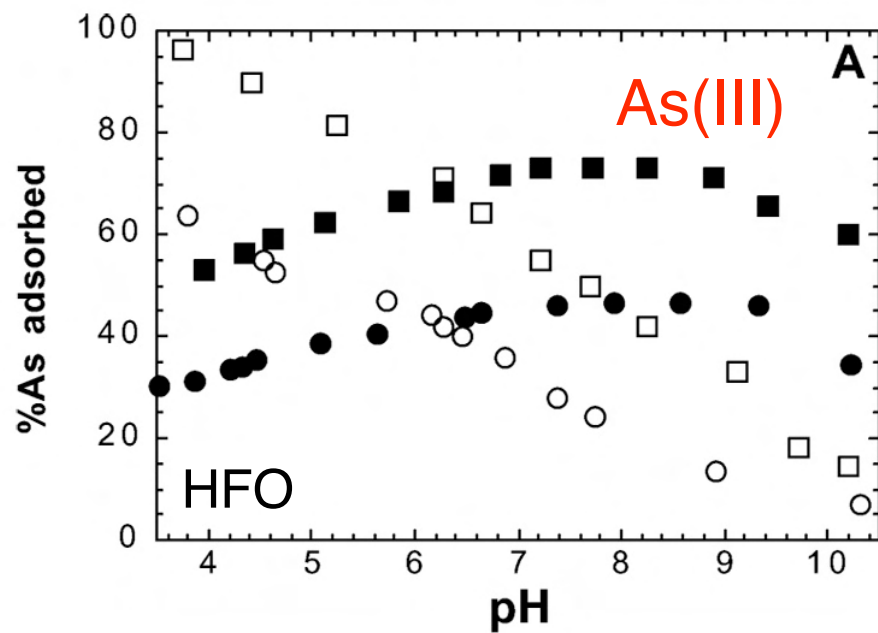
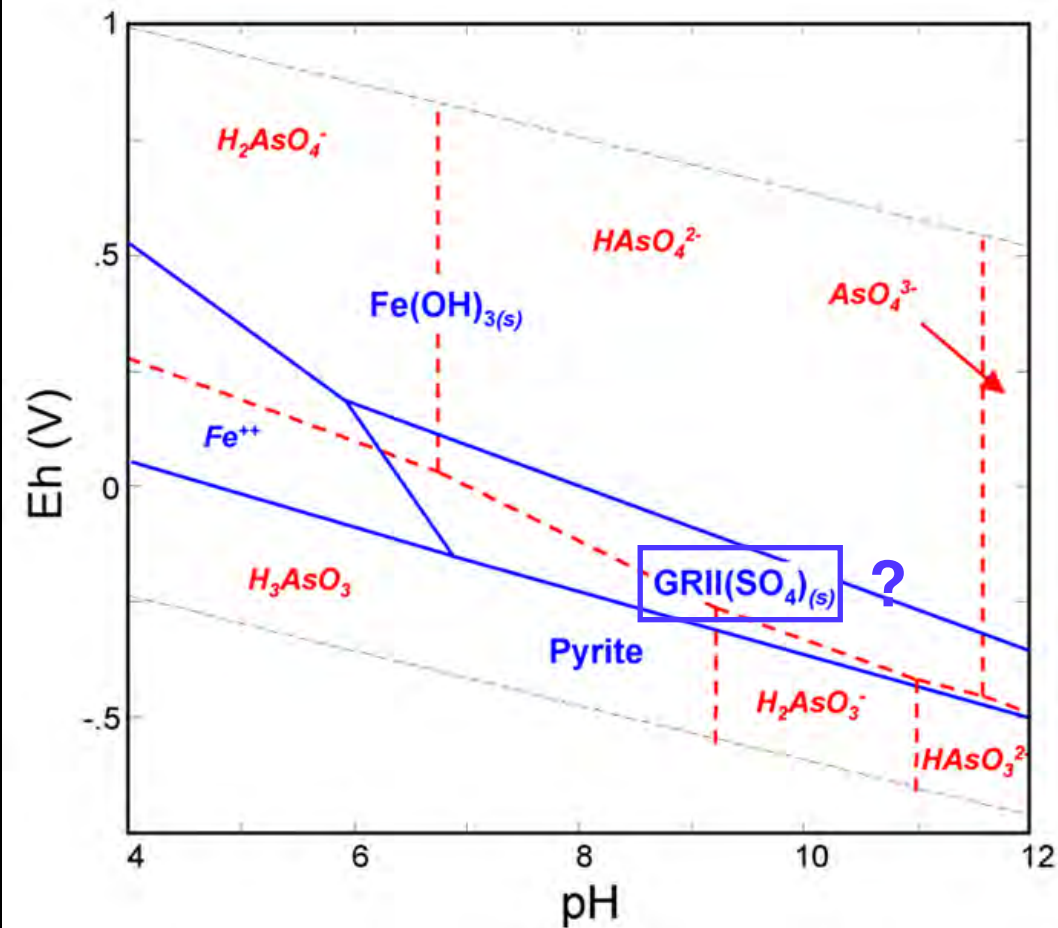
Fe EXAFS: Haiwee Sediments



Kneebone et al. (2002) ES&T 36, 381-386

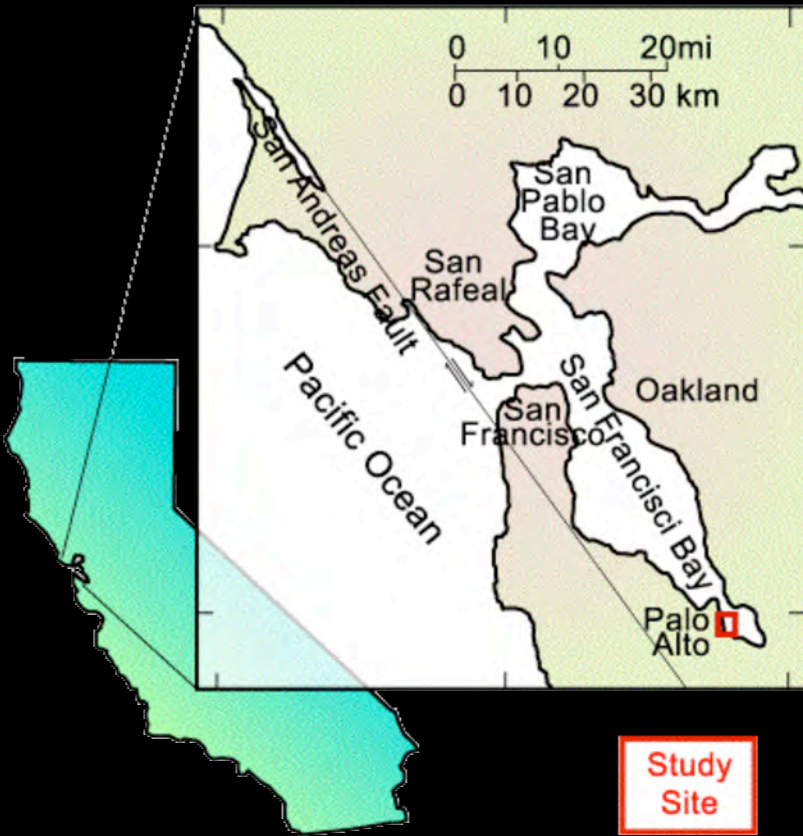


# High Iron, low Sulfur, Carbon System



## High Iron, low Sulfur, Carbon System

- Reductive dissolution of sorbent  $\text{Fe}(\text{OH})_3$  releases As
- Low potential for Sulfur reduction -- no removal by sulfides
- Reduction of As(V) to As(III) -- may remain sorbed
- As(III) sorption depends on pH, competitive sorbates, available sorbents



**Study Site**

- Tidal influence
- Sulfate reducing

## Bay Road Site East Palo Alto CA.



# Natural Arsenic Attenuation

## Bay Road Site, East Palo Alto (CA, USA):

Subsurface plume below former sodium arsenite herbicide & pesticide manufacturing facility (1926-71)

### Contaminant Plume: [As<sub>T</sub>] in Groundwater

Up to 100 mg l<sup>-1</sup>

### [As<sub>T</sub>] in Sediments

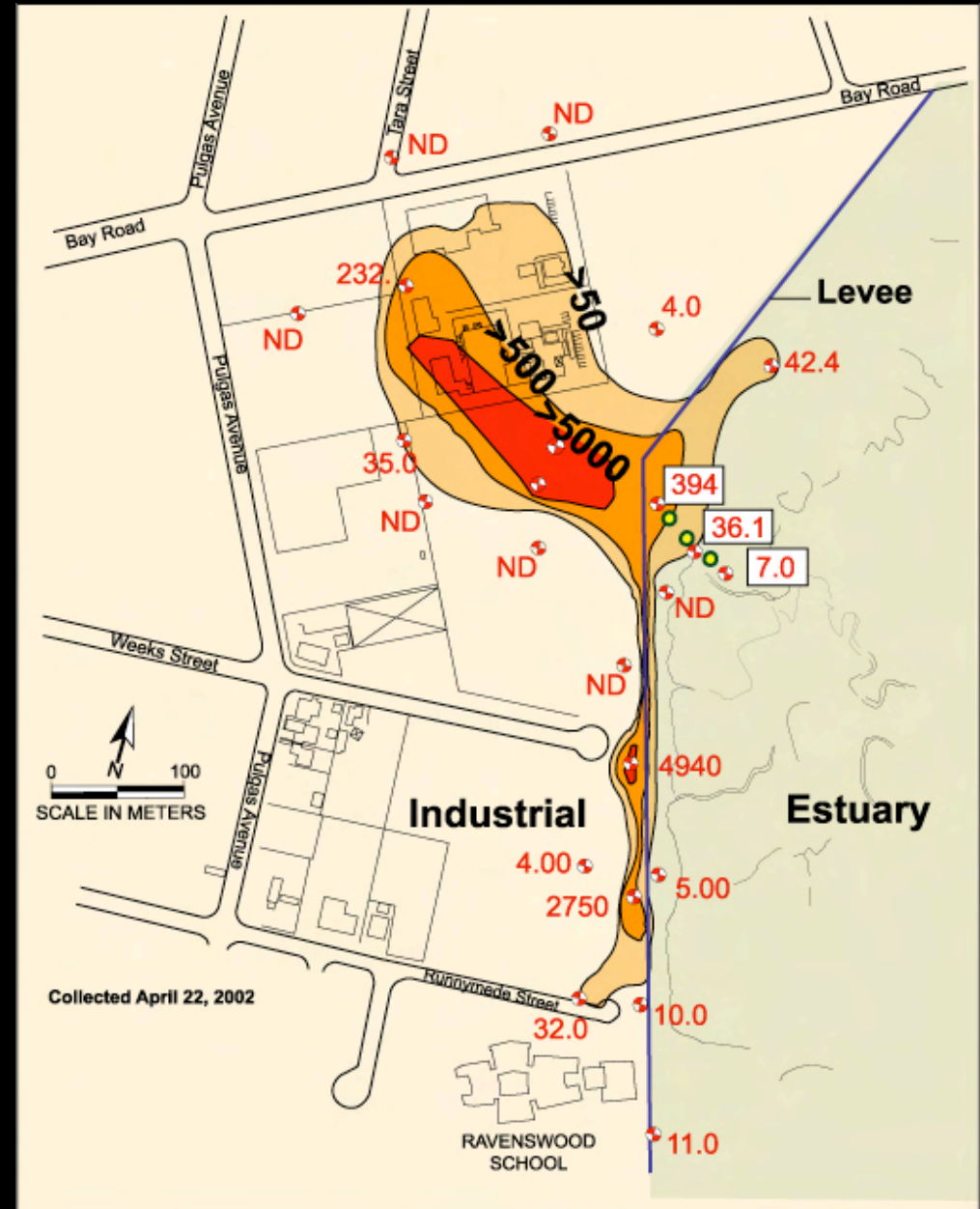
Up to 1000 mg kg<sup>-1</sup>

### Down-gradient of Plume: [As<sub>T</sub>] in Groundwater

<0.01 mg l<sup>-1</sup>

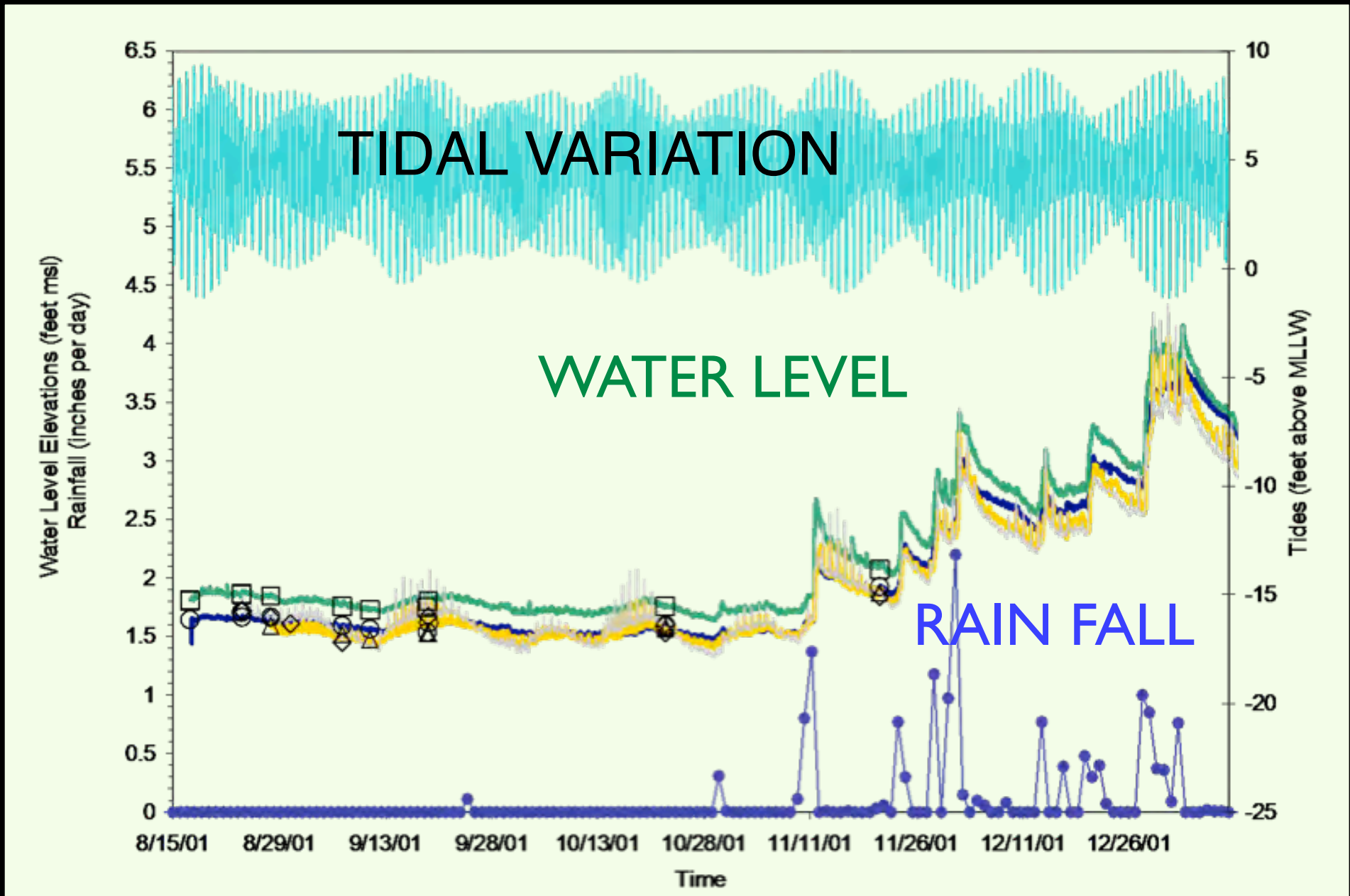
### [As<sub>T</sub>] in Sediments

Natural Background

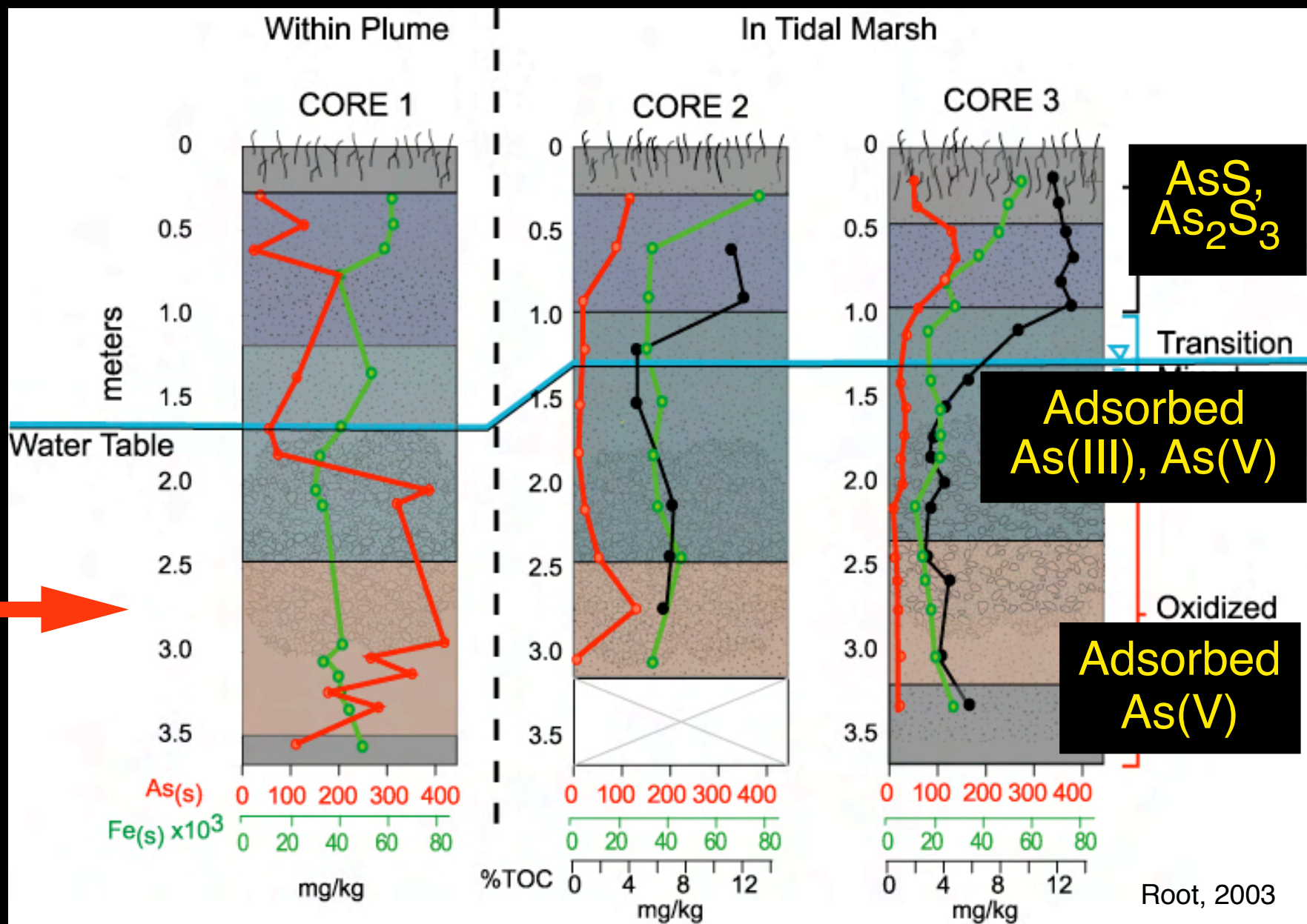




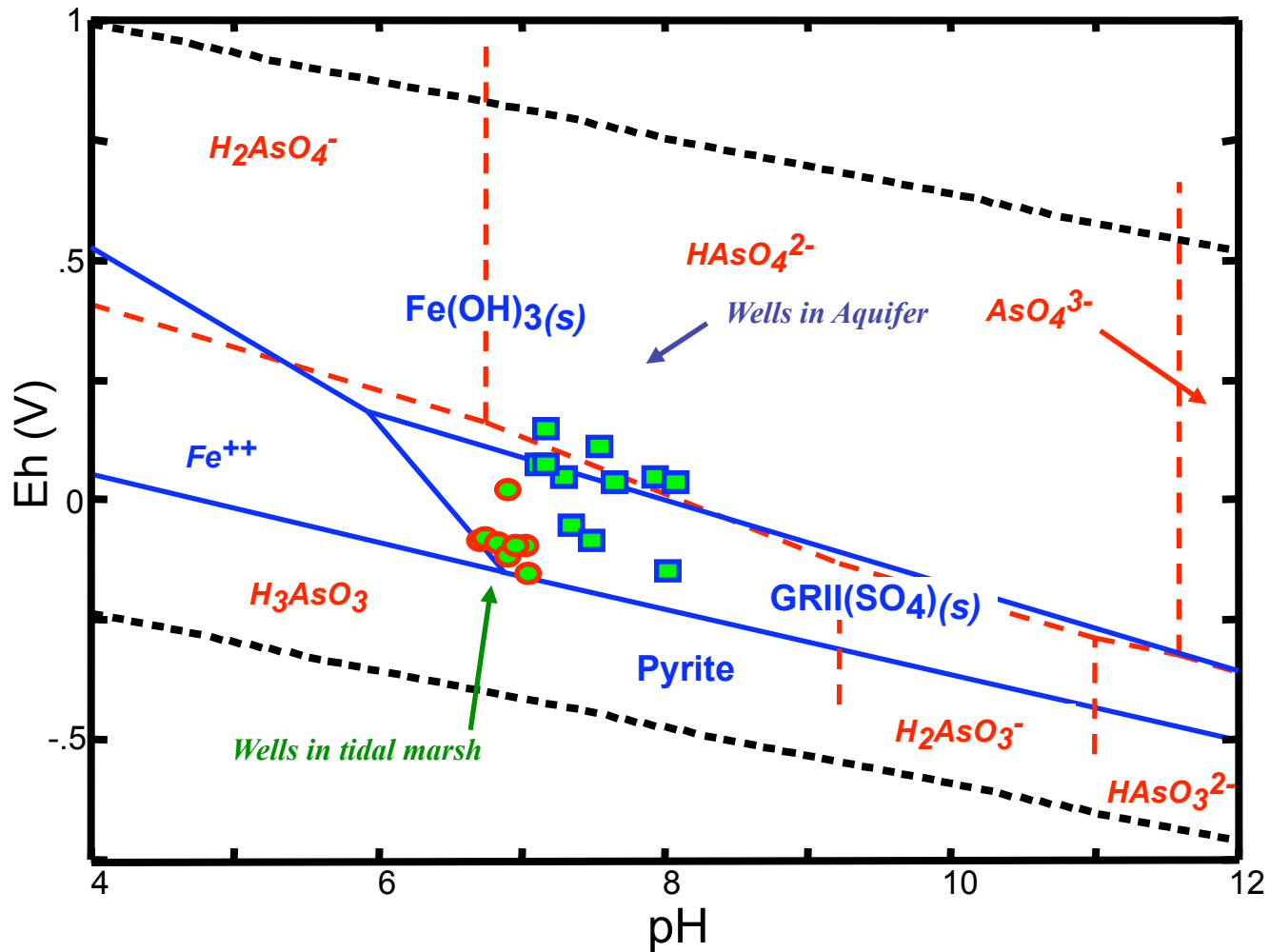
# Seasonal variation in well water level compared to tidal variation and rainfall (8/15/01-1/15/02)



# Sediment Arsenic, Iron, & Organic Carbon



# As-Fe-S Speciation



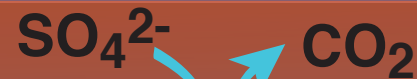
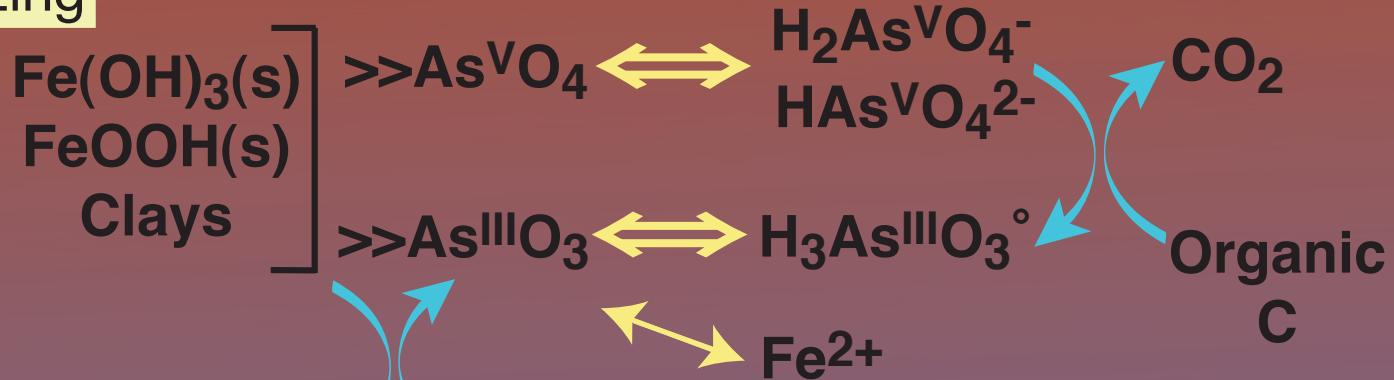
Green Rust:  $GR^{II}: Fe^{II}_6Fe^{III}_2(OH)_{16}(SO_4) \cdot 4H_2O$



**Solid & Adsorbed  
Species**

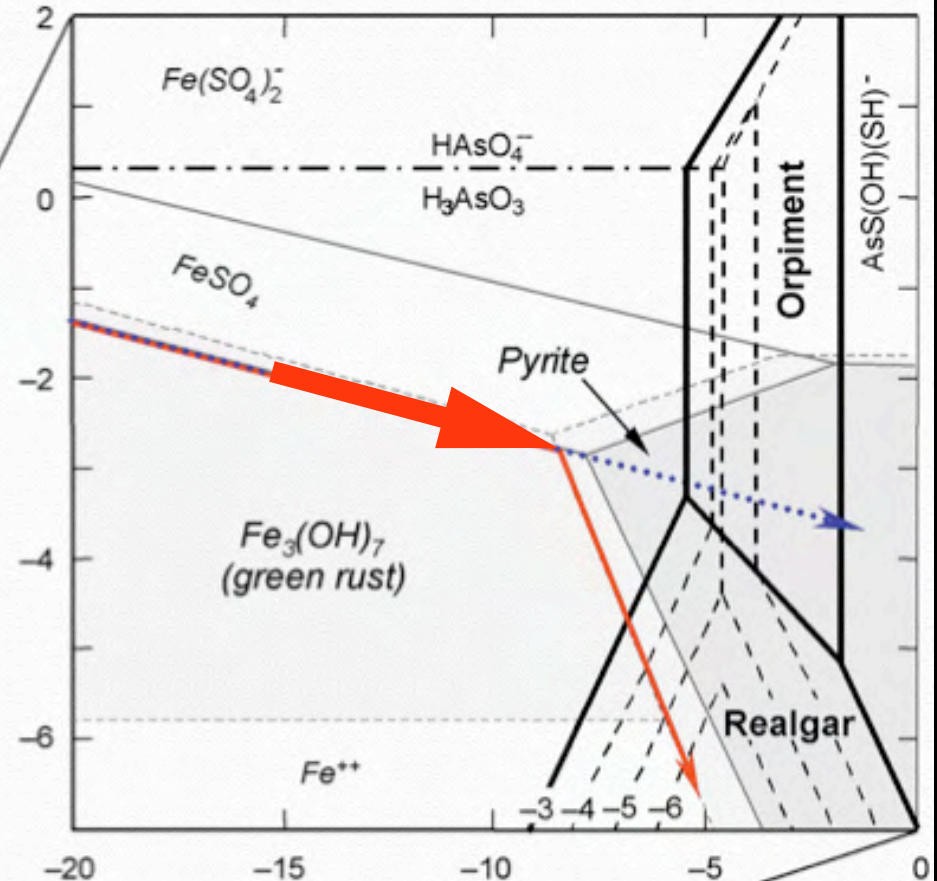
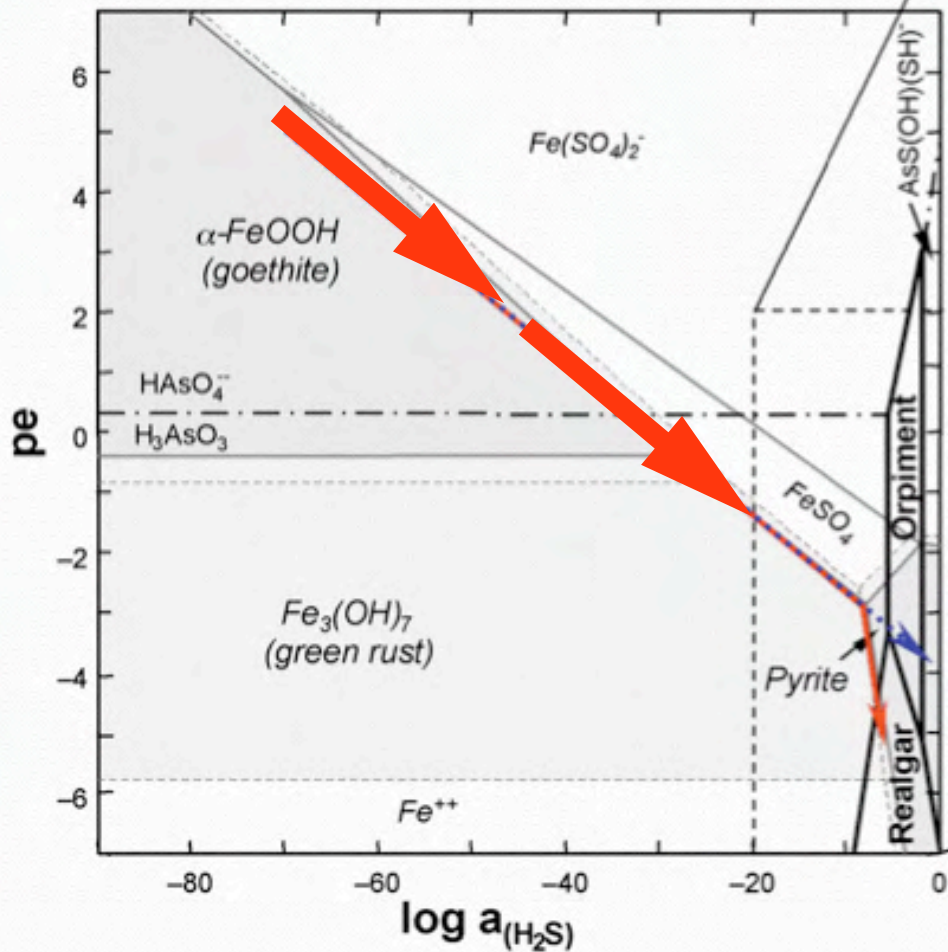
**Aqueous  
Species**

Oxidizing



Reducing

# Deciphering biotic vs. abiotic reaction rates



$$[As]_T = 100 \text{ } \mu\text{M}$$

$$[SO_4]_T = 28 \text{ mM}$$

$$\text{pH} = 7$$

O'Day et al. (2004) PNAS 101, 13703-13708

# Soil Amendments for As Stabilization Bay Road Site

## Amendments:

Ferrous sulfate (3% w/w)

Portland Cement (Type V, 10% w/w)

## As Concentrations:

500-5000 mg/kg

## Treatments:

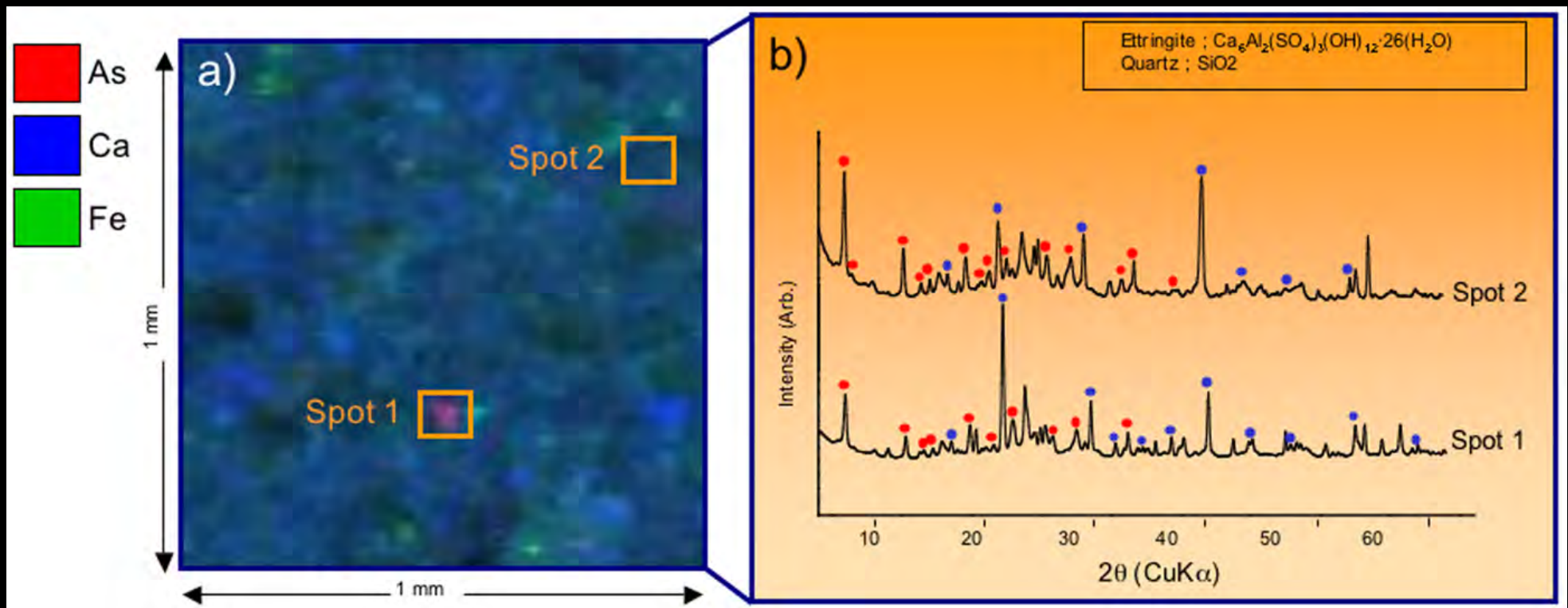
1992, 1996, 2000

1-9 m depth

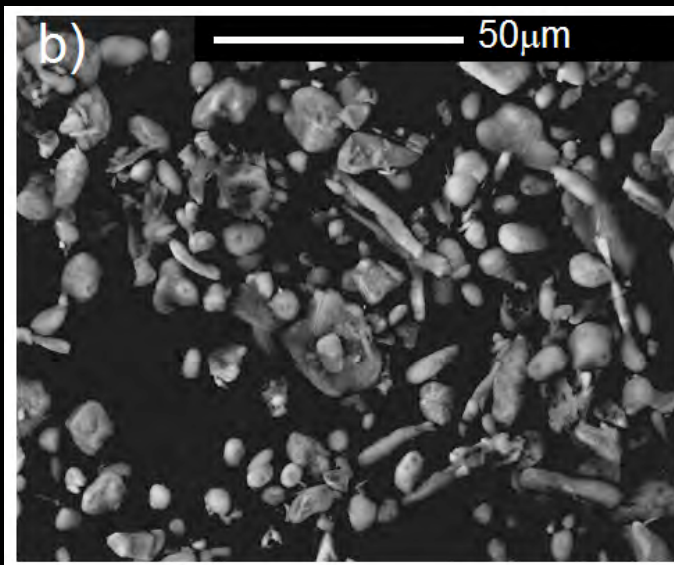
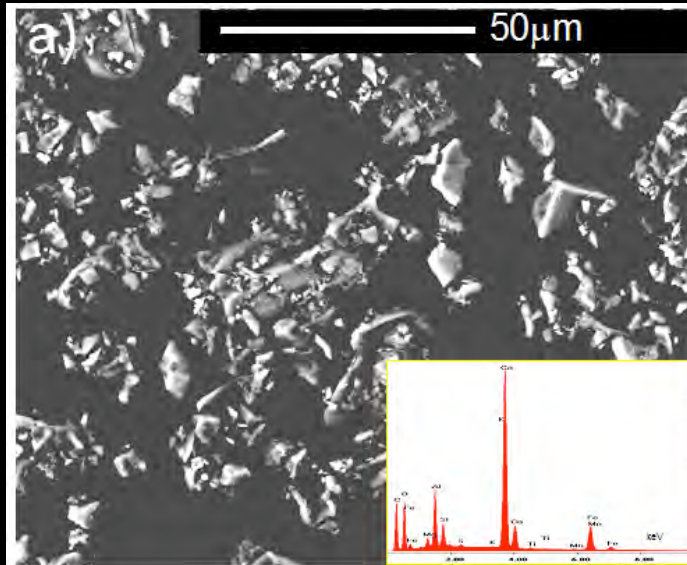
surface capped



# Microfocused Synchrotron XRD: Bay Road Field Samples

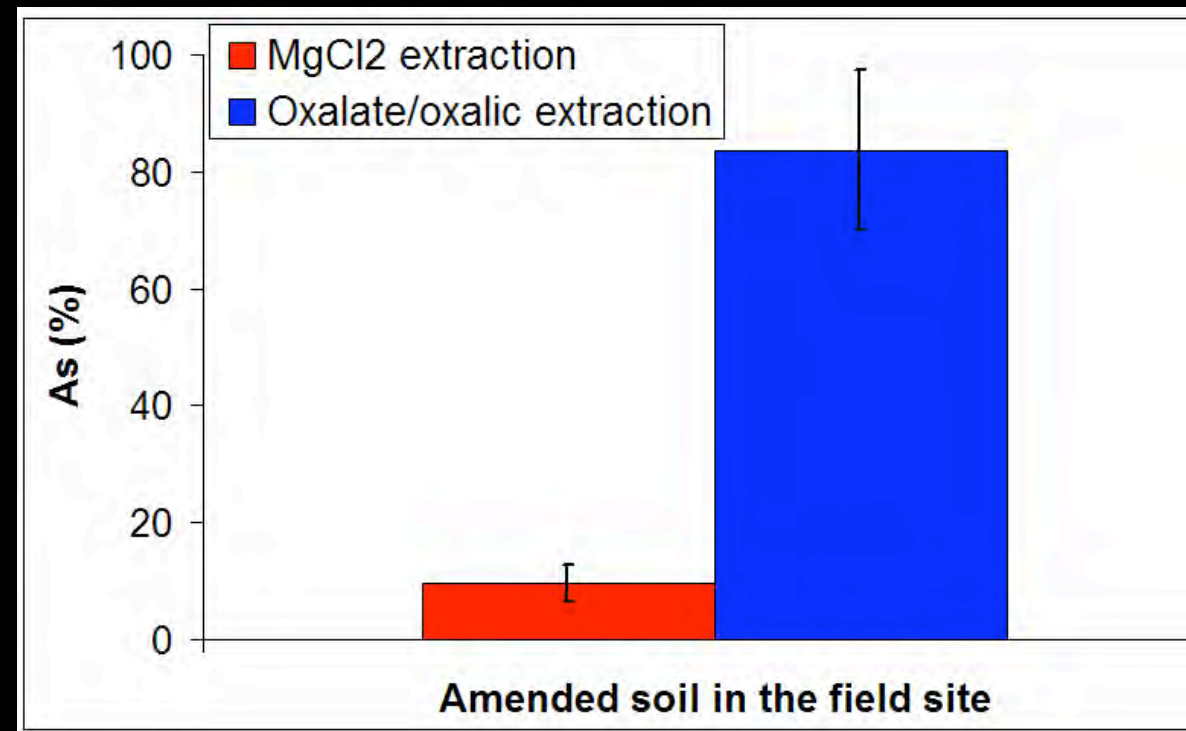


SEM



# Soil Amendments for As Stabilization

- Arsenate incorporated into crystalline sulfate phases
- No evidence for reduction to As(III) after 10+ years
- High pH stabilized
- Aging process relatively rapid -- weeks?  
(experiments in progress)

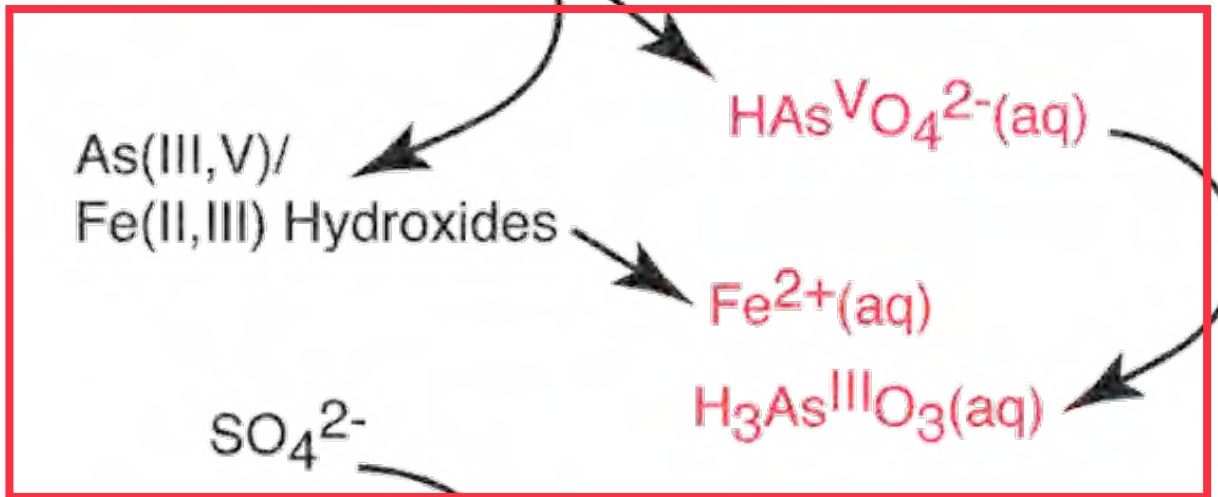


Oxidized

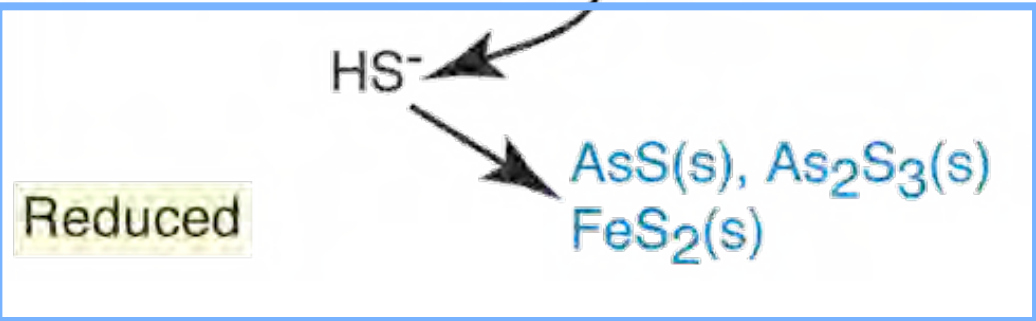
As-Fe-S

As(V)/Fe(OH)<sub>3</sub>  
As(V)/FeOOH

-- Rates?  
-- Sorption Capacity?



SO<sub>4</sub><sup>2-</sup>



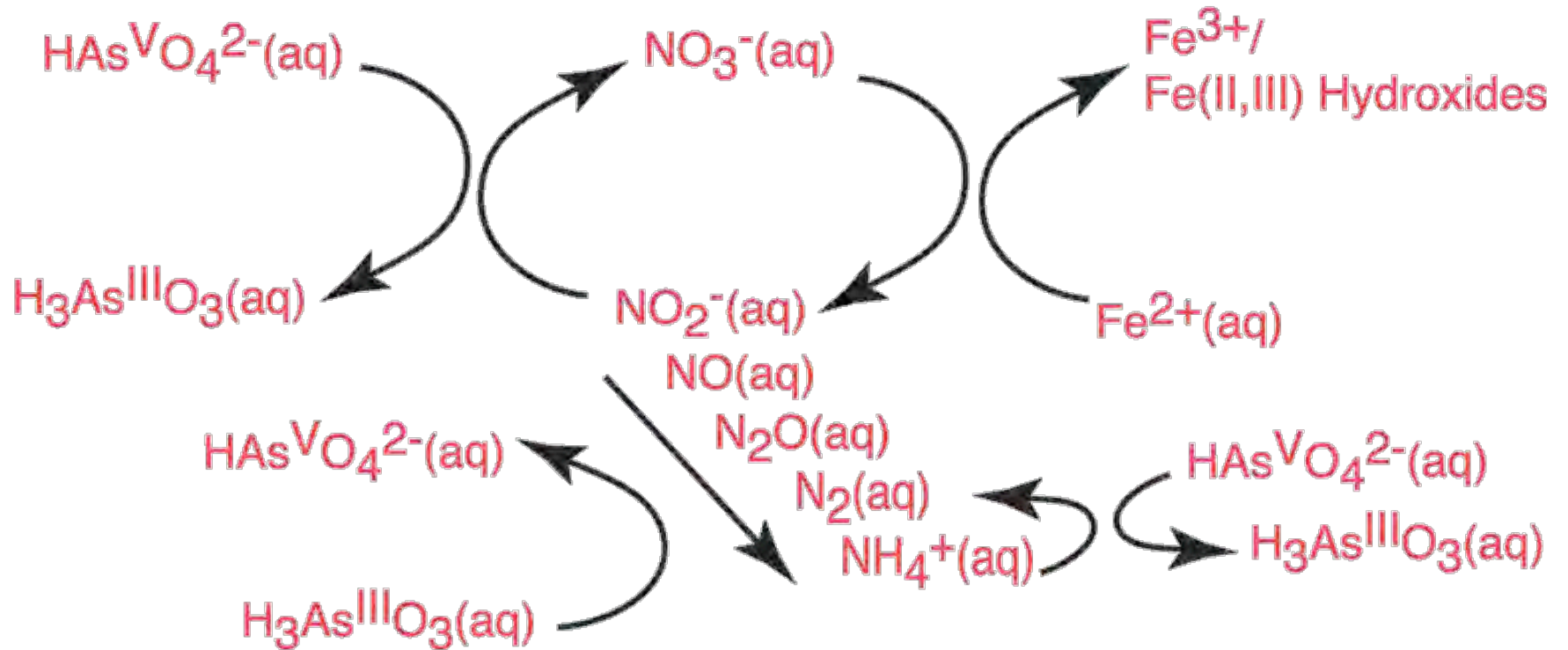
Reduced

CO<sub>2</sub>

Organic matter

-- Limited by rate of  
sulfate reduction

As-Fe-N



Z=33

As

Atomic wt.  
74.922

## Assessing the Potential for Arsenic Mobilization

- Rates of reductive dissolution of Fe(III) and Fe(II,III) (hydr)oxides and potential release of sorbed As
- pH-dependent desorption and competitive effects (phosphate, sulfate, silica)
- Rates of sulfate reduction and production of As-bearing sulfides; rates of re-oxidation
- Influence of N species on As-Fe-S redox rates
- Cost/benefit of amendment stabilization
- Validation of reactive transport models: accurate coupling of biogeochemical and hydrologic processes