



# **Bioavailability – Metals, Organics, and Use at Hazardous Waste Sites**

May 28, 2008 Session 1: "Metals"

Dr. Dominic Di Toro, University of Delaware Environmental Control of Metal Bioavailability

Dr. Nicholas Basta, Ohio State University Assessing Oral Contaminant Human (Bio)availability in Soil with *In Vitro* Gastrointestinal Methods: Uncertainties, Data Gaps, and Research Needs





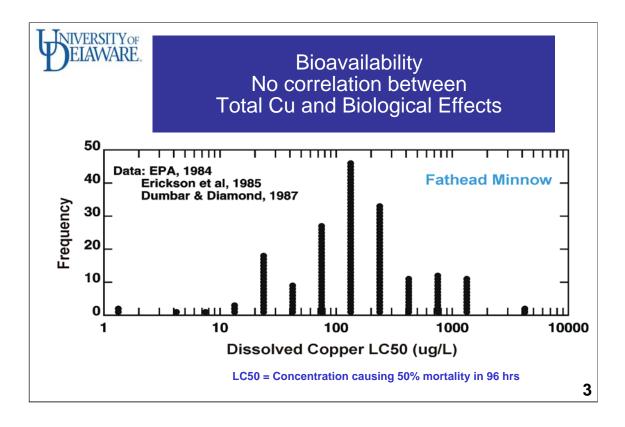


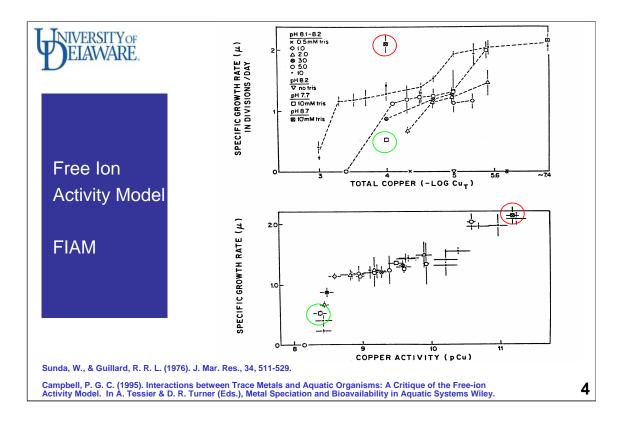
#### **Environmental Control of Metal Bioavailability**

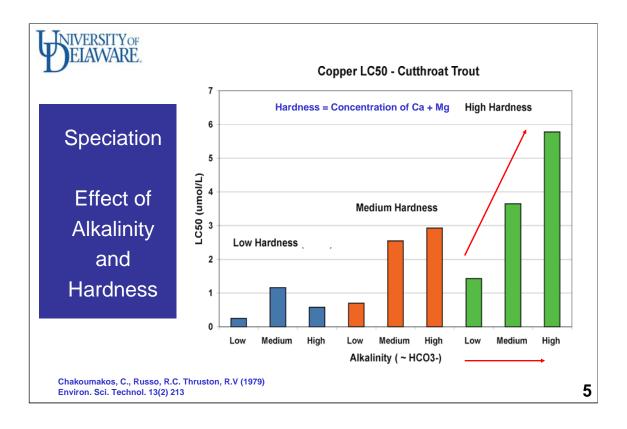
Dominic M. Di Toro

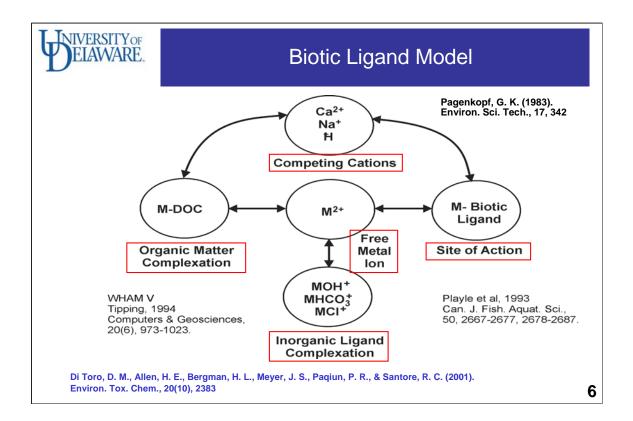
Edward C. Davis Professor of Civil and Environmental Engineering Center for the Study of Metals in the Environment Department of Civil and Environmental Engineering University of Delaware Newark, DE

> Superfund Basic Research Program Webinar EPA CLU-IN 28 May 2008









#### **Overview of Biotic Ligand Model Framework:**

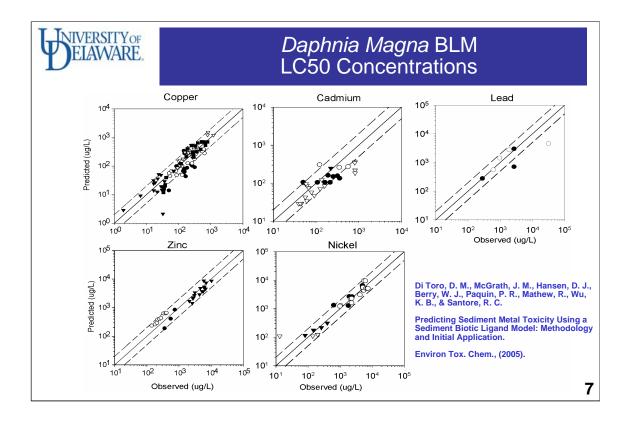
The essence of the overall BLM framework (above) was first proposed by Pagenkopf (1983) as the Gill Site Interaction Model (GSIM).

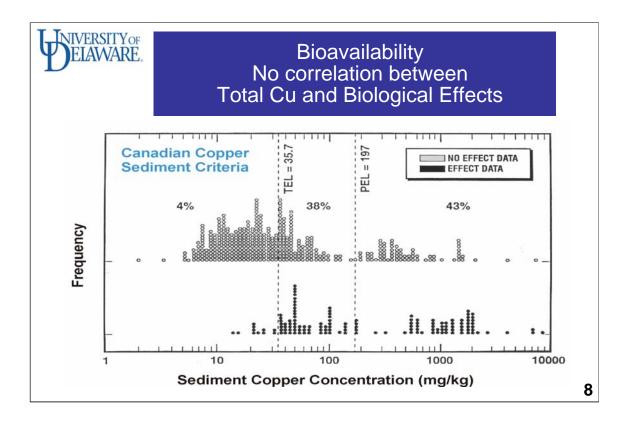
• Chemical equilibrium basics will not be discussed here. However, such models are not new and are generally well accepted by the scientific community.

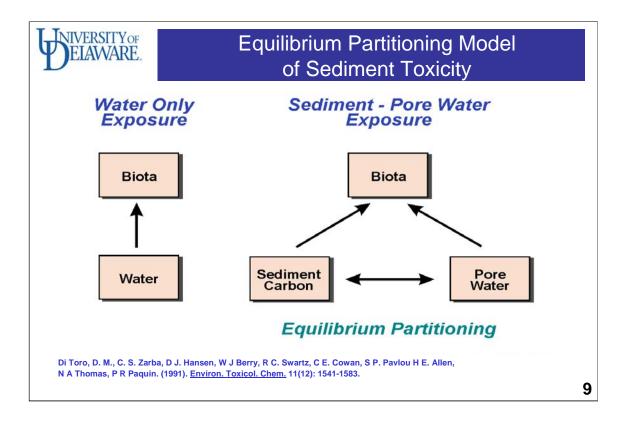
The BLM consists of 3 main types of interactions:

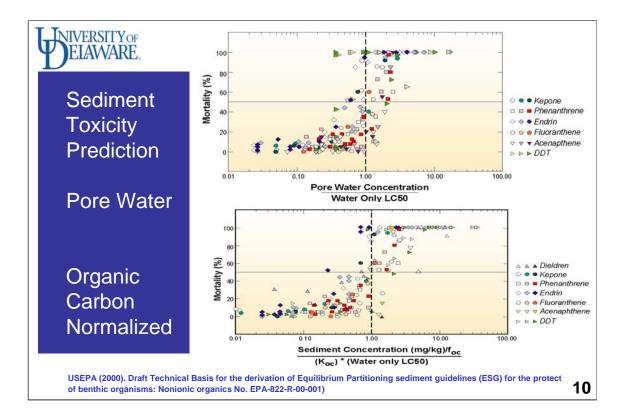
- Metal-Inorganic Ligand Interactions-
  - Chemical Equilibrium in Soils and Solutions (CHESS) (Santore and Driscoll, 1995) serves as the basis of the speciation computations and was adapted to include the metal-OM and metal-biotic ligand interactions described below.
- Metal-Organic Mater (OM) Interactions-
  - Based on the Windermere Humic Aqueous Model (WHAM), Version 5 (Tipping, 1994)
- Biotic Ligand Interactions-
  - Adapted from Playle et al., 1993a,b.

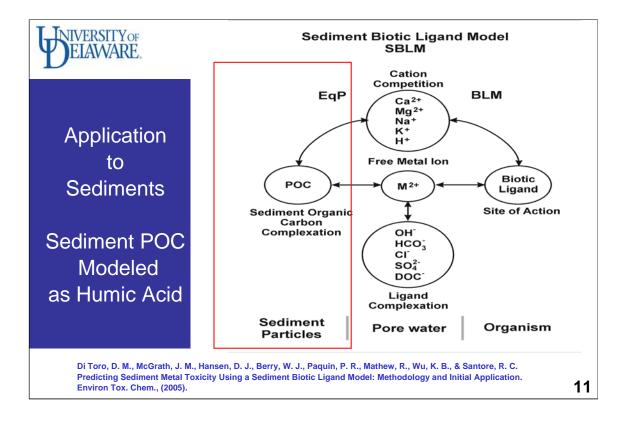
Each of these will be discussed in turn.

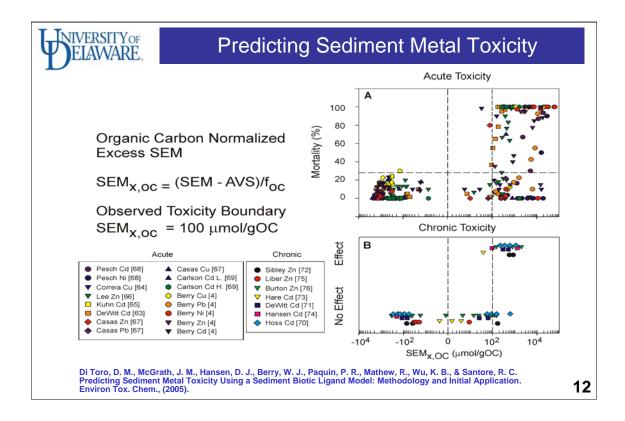


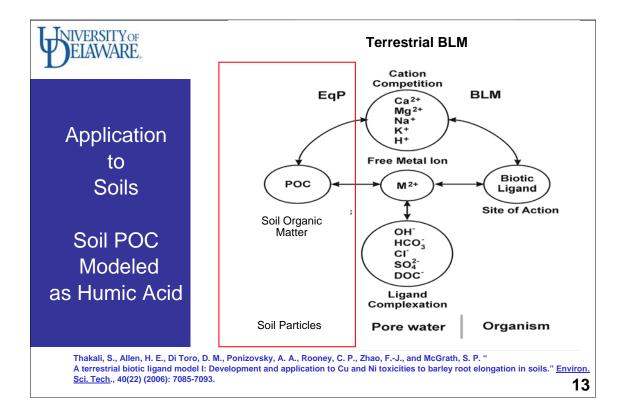


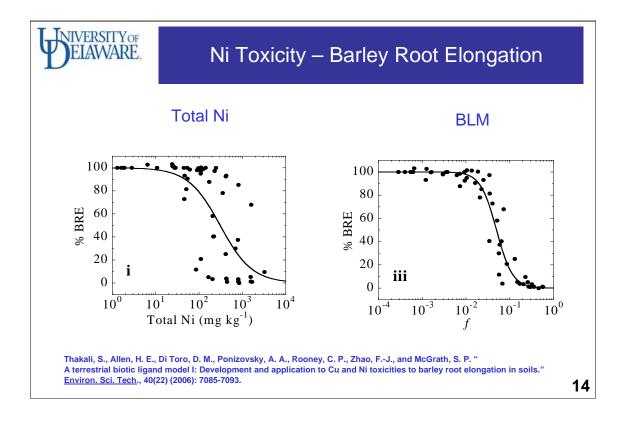


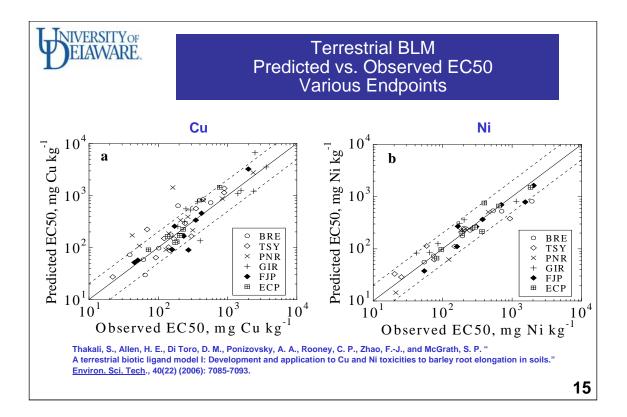


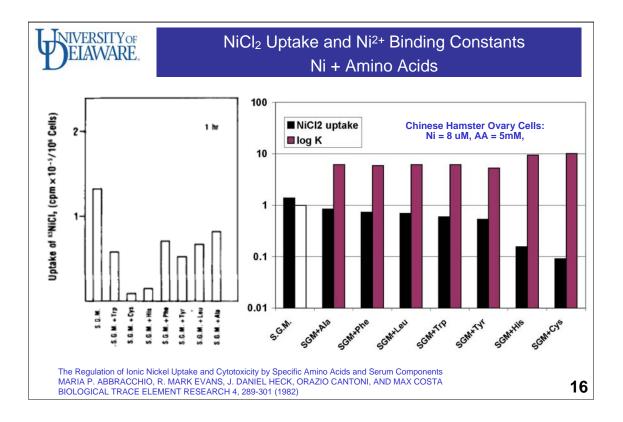


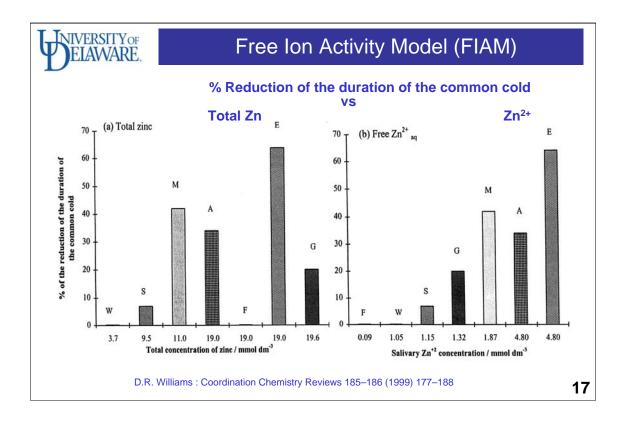


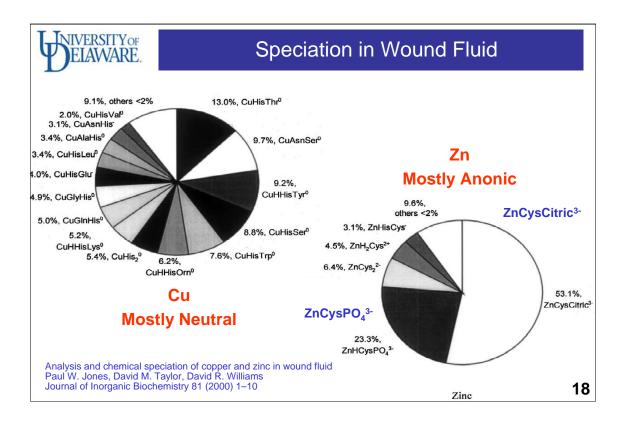












Assessing Oral Contaminant Human (Bio)availability in Soil with *In Vitro* Gastrointestinal Methods: Uncertainties, Data Gaps, and Research Needs

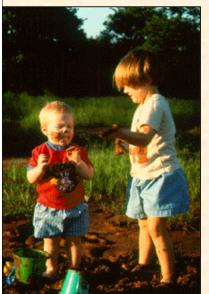
> Nick Basta Professor of Soil and Environmental Chemistry School of Environment and Natural Resources Ohio State University

Dr. Kirk Scheckel National Risk Management Research Laboratory U.S. EPA, Cincinnati, OH

Dr. Karen Bradham National Exposure Research Laboratory U.S. EPA, Research Triangle Park, NC

USEPA / Superfund Basic Research Program Webinar May 28, 2008

### Using Bioavailability to Adjust Risk in the Soil Ingestion Pathway "Soil Contaminant Oral Bioavailability"



Risk = [Soil] (EF) (ED) (IR) (BIO) (BW) (AT) How do we measure BIO for children? Animal model dosing trials costly, lengthy, not easily obtained data

#### *In Vitro* Gastrointestinal Methods An Inexpensive, Fast, Accessible Alternative



Sequential extraction, 37°C

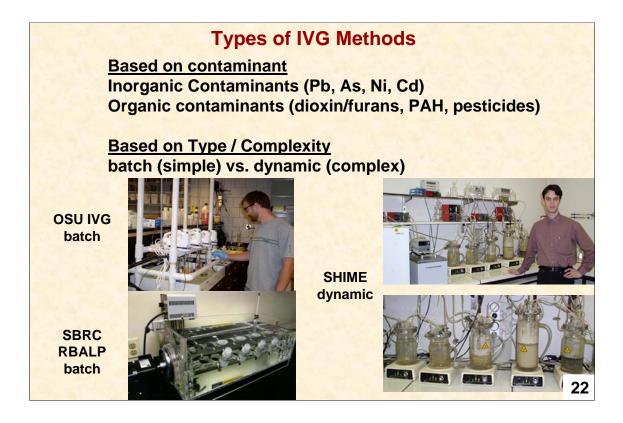
all have a stomach phase some have an intestinal phase

may have several intestinal simulations for duodenum, jejunum, colon, etc.

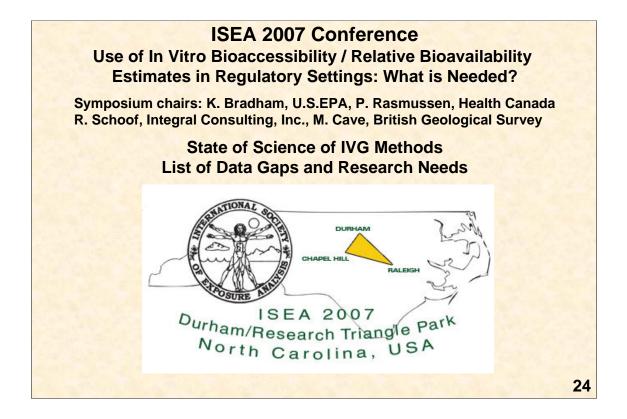
*in vitro* "(bio)availability"

dissolved contaminantbioaccessible contaminant

bioaccessibility > bioavailability, so in vitro assumes worst case



Select Types of IVG Methods		
Method	Туре	Main application(s)
PBET / RBALP (Ruby, Drexler)	Batch, fasting	Pb
OSU IVG (Basta, Rodriguez)	Batch, fasting	Pb, As, Cd
RIVM, (Oomen, Sips)	Batch, fed	PAH / Pb, As
SERDP (Lowney)	Batch, fasting	Pb, As
SHIME (Van de Wiele)	Dynamic, fed	PAH, As
TIM, tiny TIM	Dynamic, fed	PAH



## U.S. EPA

Guidance for Evaluating the Oral Bioavailability of Metals in Soils for Use in Human Health Risk Assessment OSWER 9285.7-80, May 2007

#### Recommended Criteria for Validation of Test Methods adapted from ICCVAM

"Data generated adequately measure or predict the toxic endpoint of interest and demonstrate a linkage between either the new test and effects in the target species."

> In vitro gastrointestinal (IVG) method must be correlated with an acceptable *in vivo* model IVG must be *predictive*

# Acceptable In Vivo Models



accurate bioavailability

unlikely model



acceptable model for Pb, As, other bioavailability

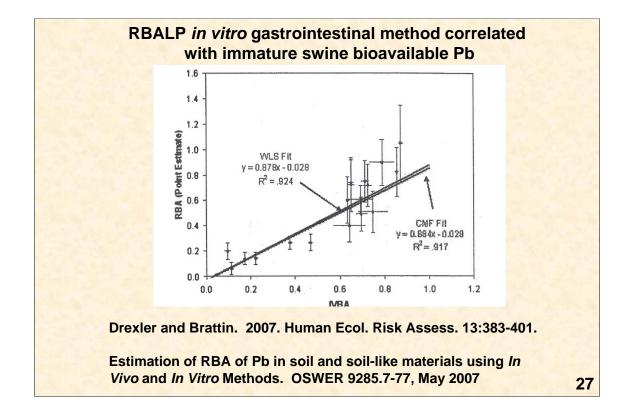
USEPA Pb OK; As?

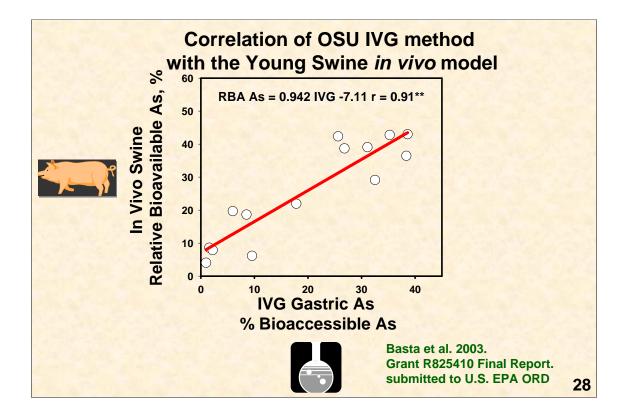


acceptable model for bioavailability

expensive ethical issues acceptable bioavailability? inexpensive recent developments Dave Thomas USEPA RTP (ISEA 2007) 26

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#### **OSU** In Vitro Gastrointestinal Method





#### Simulated GI extraction at 37°C

Gastric bioaccessibility and Intestinal bioaccessibility

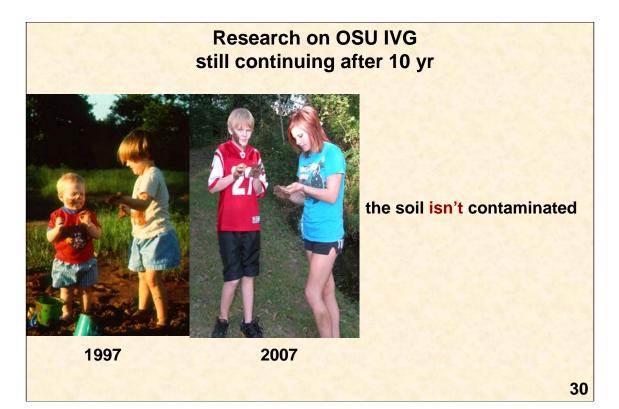
Development of Chemical Methods to Assess the Availability of Arsenic in Contaminated Media, R825410

U.S. EPA, Office of Research and Development National Center for Environmental Research

N.T. Basta, R.R. Rodriguez, and S.W. Casteel Nov 1996 to October 2000.

Rodriguez et al. 1999. ES&T 33:642-649.

Basta et al., 2007. J. Environ. Health Sci. Part A 42:1275-1181

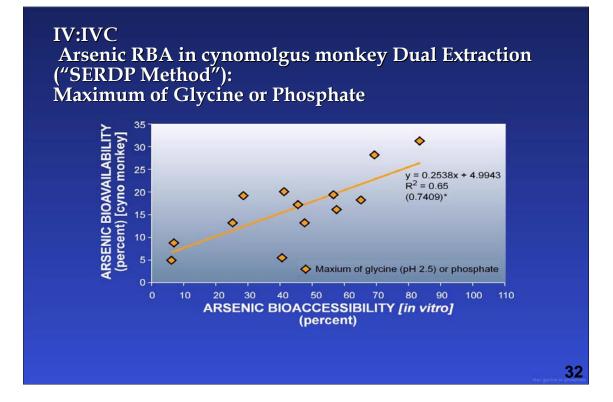


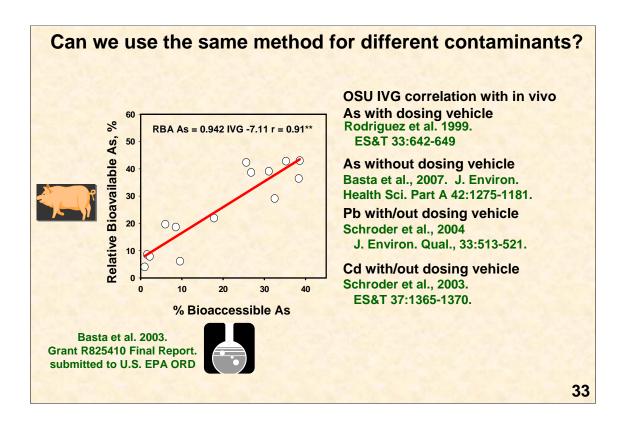
Correlation of "SERDP" method with Relative Bioavailable Arsenic

Lowney, presented at ISEA 2007 Primate (cynomolgus monkey) RBA As vs. "SERDP" As

"SEDRP" As: gastric bioaccessibility 0.4 M glycine/HCl pH 1.5 <u>OR</u> 0.4 M K<sub>2</sub>HPO<sub>4</sub>, pH 2.5

use larger bioaccessible As value of two methods





#### *IVG* Method Correlation Studies Will the method work for other contaminated soils?



Most correlation studies conducted on highly contaminated wastes

often > 2,000 mg/kg contaminant of concern

Estimating RBA of Pb in Soil and Soil-like materials (OSWER 9285.7-77, May 2007) Most of 19 solid waste materials from smelter origin Pb content: 1,590 to 14,200 mg/kg, median 7,225 mg/kg

Estimating RBA of Arsenic in Contaminated Soils and Solid Media (Rodriguez et al., 1999) As content: 233 to 17,500 mg/kg, median 1,460 mg/kg

#### *In Vitro* Models Method Validation Issues

Will the in vitro test work for all types contaminants/media?



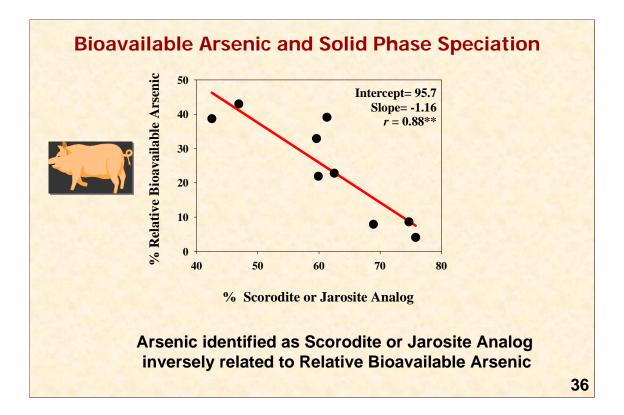
Do we have to conduct validation studies for all contaminant sources?

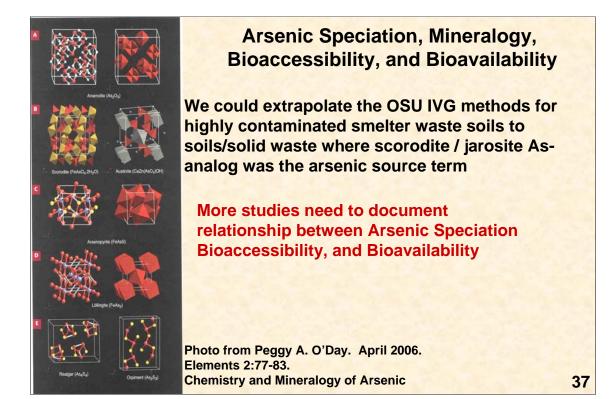
mining waste, battery waste, paint, coal ash, etc.?



Better approach – contaminant speciation SEM/EDX (J. Drexler); EXAFS (K. Scheckel)

Which species are bioavailable? Does the *in vitro* test measure them?





# **Contaminant Concentration in Soil / Solid Waste**

when will bioavailability adjustments be made?

Highly Contaminated unreasonable adjustment

> Moderately Contaminated

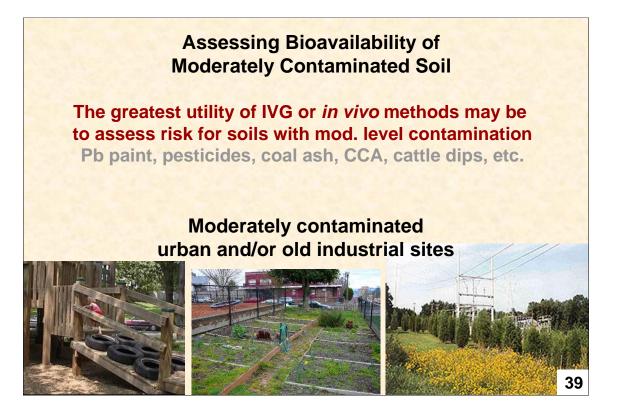
reasonable adjustment

Background

High level: 7,000 mg/kg total As or Pb Bioavailability has to be very very low unreasonable adjustment

Moderate level: 300 mg/kg As moderate bioavailability so reasonable adjustment

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Bioavailable (*in vivo*) vs. Bioaccessible (*in vitro*) Method Detection Limits and Contaminant Levels

most *in vivo* dosing studies require highly contaminated soil > 500 to 5,000 mg/kg contaminant

Moderately contaminated soil levels could be < 1000 mg/kg Pb; < 100 mg/kg As Below *in vivo* detection limits

Below *in vivo* working range but easily measured by IVG methods

<u>A Strong Advantage of IVG methods</u> is the ability to estimate (bio)availability at moderate levels Highly Contaminated in vivo and in vitro

> Moderately Contaminated only *in vitro*

Background

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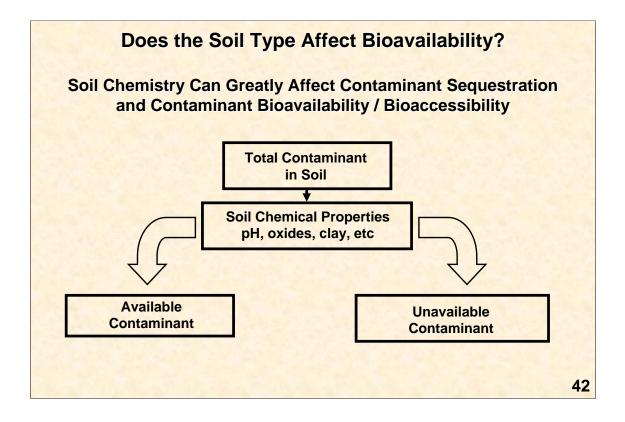
# Are we confident to use IVG methods to Estimate Contaminant Bioavailability in Soil for Moderately Contaminated Soils?

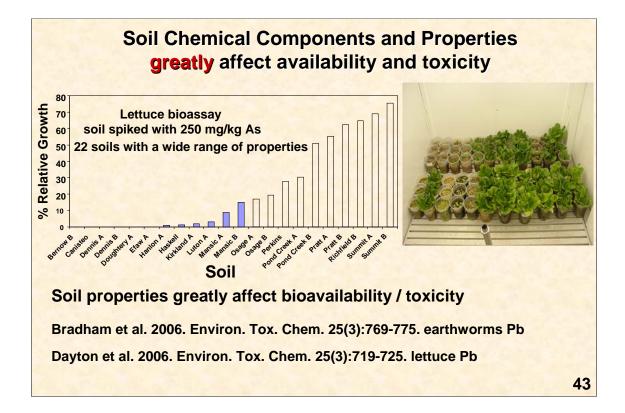
Knowledge of chemical speciation is essential!

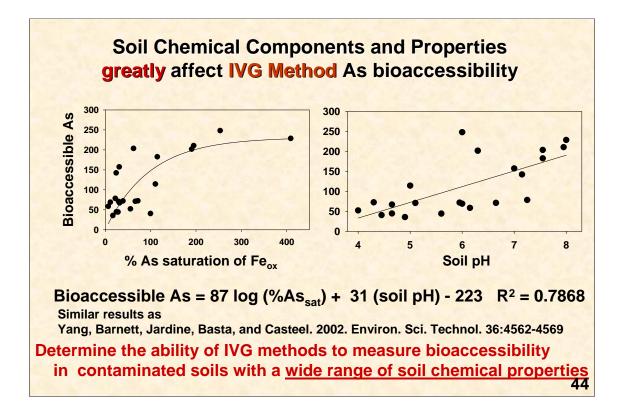
contaminant species in old orchard soil same as contaminant species in smelter soil (*in vivo* correlation study)?

Yes: then we are more confident to use the IVG (*in vitro*) method for the orchard soil









# U.S. EPA

## Guidance for Evaluating the Oral Bioavailability of Metals in Soils for Use in Human Health Risk Assessment OSWER 9285.7-80, May 2007

"A detailed protocol for the test method......, and a description of the known limitations of the test including a description of the classes of materials that the test can and cannot accurately assess."

- > Specify the contaminant chemical speciation and
- whether the IVG method has been correlated with *in vivo* for the contaminant species in the test material
- > Measure soil chemical parameters that affect bioavailability

### Summary

### Uncertainties, data gaps, research needs

\* Research leading to acceptance of existing / new *in vivo* models

- Document the relationship between arsenic speciation, bioaccessibility, and bioavailability
- Test the use of soil chemical / speciation methods to support IVG data when IVG is the only option
- Determine the ability of IVG methods to measure bioaccessibility in contaminated soils with a <u>wide range of soil chemical properties</u>

