

# Abstract 746: Pediatric Bioavailability of Lead in Soil and Dust:

## Estimates from Soil, House Dust, and Blood Lead at the Bunker Hill Superfund Site

M. Stifelman, U.S. EPA, Region 10 Seattle, WA; S. Spalinger and I. von Lindern, TerraGraphics, Inc., Moscow, ID, USA.

Contact: stifelman.marc@epa.gov spalings@tgenviro.com vonlindi@tgenviro.com

### Abstract

More than a century of mining and smelting at Bunker Hill contaminated soil and interior dust with lead and other metals. Following closure of the smelter in 1981, lead in soil and dust were identified as primary exposure pathways for children. Soil and dust have been sampled annually as contaminated soil was replaced with clean soil at schools, parks, businesses, and approximately 2,200 residences. Annual voluntary blood lead screening has recruited over 50% of exposed children from 1988 through 2002. Blood lead levels have decreased as soil and dust lead levels have declined as the soil cleanup progressed. Bioavailability was estimated on an annual basis by comparing lead uptake from over 5,000 blood lead measurements paired with estimates of lead intake from thousands of soil and dust measurements. Lead intakes were estimated using a range of assumptions of the relative importance of lead levels in 1) house dust, 2) residential soil from a child's own yard, 3) soil from neighboring yards, and 4) the mean concentration from all yards in a town. The impact of various exposure assumptions on the estimated intake was minimal because of correlations among the four sources of exposure. From 1988 to 2002, the estimated aggregate geometric mean bioavailability of soil and dust averaged 18% over a range of 12-23%. Higher bioavailability coincided with increased intakes of lead in dust relative to soil, suggesting that dust is more bioavailable than soil. Attempts to separate soil and dust bioavailability are sensitive to soil and dust ingestion rate assumptions. However, assuming 10% soil and 25% dust bioavailability explains the annual variation in aggregate soil/dust bioavailability observed from 1988 to 2002 using IEUBK soil/dust ingestion rates. Greater bioavailability of dust may be caused by smaller particles in dust relative to soil which may account for an increase in bioavailability (due to greater surface area) and ingestion rate (smaller particles are more likely to cling to hands and fingers).

### Background

In risk assessment, bioavailability refers to the fraction of an external dose that enters systemic circulation (National Academy of Sciences, 2003). If the concentration of lead in environmental media available for exposure is the external dose, then bioavailability can include ingestion rates as well as the uptake fraction. Lead exposure from soil or dust is modeled as the product of concentration ( $\mu\text{g/g}$ ), ingestion rate ( $\text{g/day}$ ), and the unitless fraction available for uptake (i.e., bioavailability). See Figure 1 for predicted and observed blood lead levels using an aggregate soil/dust bioavailability of 18%.

### Methods

Aggregate bioavailability of lead in soil and dust has been estimated annually by comparing blood lead levels in resident children (internal dose) to lead concentrations in soil and dust (external dose) using age-specific default soil/dust ingestion rates of the EPA IEUBK Lead Model (U.S. Environmental Protection Agency, 2001).

### Methods cont.

Blood lead levels were converted to lead uptake using age-specific biokinetic slope factors which are the basis of the IEUBK Lead Model (Harley & Kneip, 1985; U.S. Environmental Protection Agency, 2001).

Bioavailability (unitless) = Internal dose ( $\mu\text{g/day}$ ) per external dose ( $\mu\text{g/day}$ )  
 Internal dose ( $\text{day/dL}$ ) = Blood lead level ( $\mu\text{g/dL}$ ) per biokinetic slope factor  
 Biokinetic slope factor = Blood lead level ( $\mu\text{g/dL}$ ) per lead uptake ( $\mu\text{g/day}$ )  
 External dose ( $\mu\text{g/day}$ ) = Sum of (soil+dust+ air+water+diet) intakes  
 Lead biokinetic slope factors at an uptake of 20  $\mu\text{g/day}$  (Harley & Kneip, 1985)

Age (months)	Biokinetic Slope Factor (day/dL)
0-12	0.295
13-24	0.405
25-36	0.365
37-48	0.350
49-60	0.365
61-72	0.345
72-84	0.325
85-96	0.250
97-108	0.230

#### Separating soil and dust bioavailability

Annual aggregate soil and dust bioavailability and the percent of lead intake from soil versus dust were tabulated from 1988 through 2002 (see Figure 2). Discrete bioavailability for soil and dust were calculated using the Solver function in Microsoft Excel 2000<sup>®</sup> as follows (see Solver report below):

- 1 Assume initial values for discrete soil and dust bioavailability are equal to the aggregate soil/dust bioavailability (i.e., a geometric mean of 18%).
- 2 Define the error of the initial value as:  
= sum of (observed-predicted aggregate bioavailability)<sup>2</sup>
- 3 Predicted aggregate bioavailability  
= (initial soil bioavailability\*weighting factor<sub>soil</sub>) + (initial dust bioavailability\*weighting factor<sub>dust</sub>)
- 4 Use Excel Solver to minimize the sum of errors with the following constraints:
  - 1) dust bioavailability > soil bioavailability
  - 2) dust bioavailability < 1.0
  - 3) soil bioavailability > 0.1

Microsoft Excel 9.0 Answer Report  
 Worksheet: [mls-request\_02-17-04.xls]Bio estimates  
 Report Created: 2/27/2004 2:12:45 PM

Cell	Name	Original Value	Final Value
\$L\$20	sum of errors (predict-obs) squared	0.015147361	0.009764467

Cell	Name	Original Value	Final Value
\$J\$4	GeoMean DUST Bioavailability	18.0%	25.4%
\$K\$4	GeoMean SOIL Bioavailability	18.0%	10.0%

Cell	Name	Cell Value	Formula	Status	Slack
\$J\$4	GeoMean DUST Bioavailability	25.4%	$\$J\$4 \geq \$K\$4$	Not Binding	15.4%
\$J\$4	GeoMean DUST Bioavailability	25.4%	$\$J\$4 \leq 1$	Not Binding	0.74593252
\$K\$4	GeoMean SOIL Bioavailability	10.0%	$\$K\$4 \geq 0.1$	Binding	0.0%

### Results

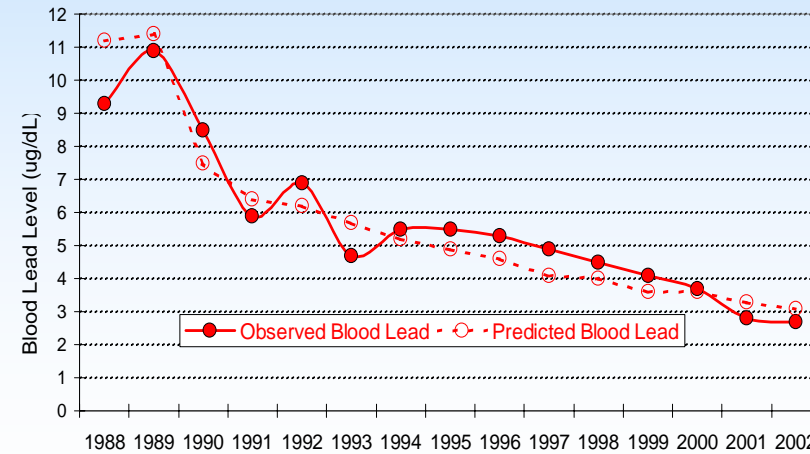


Figure 1 Observed and Predicted Blood Lead Using 18% Aggregate Bioavailability for Soil and Dust

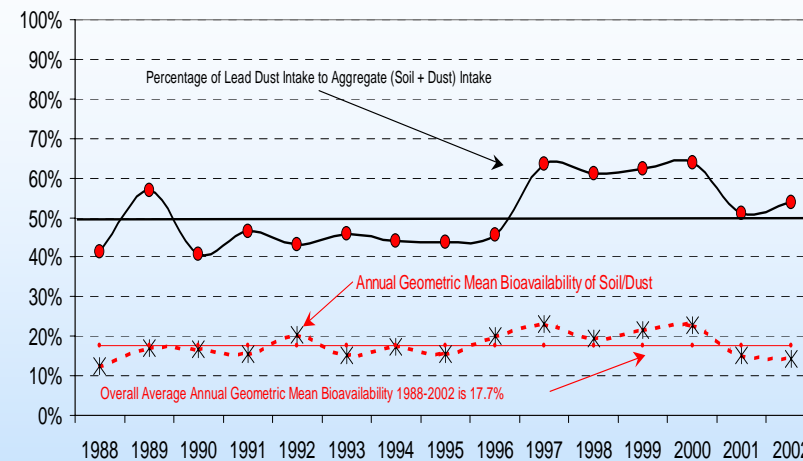


Figure 2 Aggregate Soil and Dust Bioavailability and Percentage of Lead Intake from Dust

### Discussion

Although measurements of lead bioavailability in soil media are available, similar measurements for lead in dust are lacking (National Academy of Sciences, 2003). Relative bioavailability of lead in soil and dust can be inferred from slope factors which relate concentrations in environmental media to blood lead concentrations in children, but this necessarily combines absorption with ingestion and other measures of exposure. Although soil and dust slope factors are highly variable, values soil slope factors are generally less than dust slope factors (Succop et al., 1998). Direct comparison between soil and dust bioavailability are impeded by differences in units. Soil lead is reported as a mass based concentration while dust lead is often reported as a mass per area basis (i.e., loading) which is a better predictor of blood lead levels than a mass based concentration (Lanphear et al., 1995; Lanphear et al., 1998; Succop et al., 1998).

### Conclusion

Analysis of estimated annual aggregate bioavailability and the relative soil and dust intakes suggest geometric mean bioavailability of 10% and 25%, respectively (average values are 36% and 11%). These results are consistent with observed increases in aggregate bioavailability during years when dust lead intakes were greater than soil lead intakes and predicted blood lead levels were below observed levels. Greater effective bioavailability of dust relative to soil could be caused in part by enhanced dermal adherence, ingestion, and absorption due to smaller particle size (Kissel et al., 1996; Steele et al., 1990; U.S. EPA Technical Review Workgroup for Lead, 1999). Differential assumptions of soil and dust bioavailability could be used to refine risk assessments and predictive blood lead modeling.

### References

Harley, N.H. & Kneip, T.H. (1985). An integrated metabolic model for lead in humans of all ages.  
 Kissel, J.C., Richter, K.Y. & Fenske, R.A. (1996). Factors affecting soil adherence to skin in hand-press trials. Bull Environ Contam Toxicol, 56, 722-8.  
 Lanphear, B.P., Emond, M., Jacobs, D.E., Weitzman, M., Tanner, M., Winter, N.L., Yakir, B. & Eberly, S. (1995). A side-by-side comparison of dust collection methods for sampling lead-contaminated house dust. Environ Res, 68, 114-23.  
 Lanphear, B.P., Matte, T.D., Rogers, J., Clickner, R.P., Dietz, B., Bornschein, R.L., Succop, P., Mahaffey, K.R., Dixon, S., Galke, W., Rabinowitz, M., Farfel, M., Rohde, C., Schwartz, J., Ashley, P. & Jacobs, D.E. (1998). The contribution of lead-contaminated house dust and residential soil to children's blood lead levels. A pooled analysis of 12 epidemiologic studies. Environ Res, 79, 51-68.  
 National Academy of Sciences. (2003). Bioavailability of Contaminants in Soils and Sediments Processes, Tools, and Applications. National Research Council, Committee of Contaminants in Soils and Sediments, Water Science and Technology Board, Division on Earth and Life Studies: Washington D.C.  
 Steele, M.J., Beck, B.D., Murphy, B.L. & Strauss, H.S. (1990). Assessing the contribution from lead in mining wastes to blood lead. Regul Toxicol Pharmacol, 11, 158-90.  
 Succop, P., Bornschein, R., Brown, K. & Tseng, C.Y. (1998). An empirical comparison of lead exposure pathway models. Environ Health Perspect, 106 Suppl 6, 1577-83.  
 U.S. Environmental Protection Agency. (2001). Integrated Exposure Uptake Biokinetic Model for Lead in Children (IEUBKwin v1.0). Environmental Protection Agency OERR Office of Emergency and Remedial Response: Washington, DC.  
 U.S. EPA Technical Review Workgroup for Lead. (1999). Short Sheet: IEUBK Model Bioavailability Variable.