

Research Brief 202: Size-resolved Chemical Characterization of Atmospheric Aerosols

Release Date: 10/05/2011



Background



Wind erosion of a mine tailings pile in a typical semi-arid environment.
(Photo by: Blenda Machado)

Crushing, grinding, smelting, refining, and tailings management operations at mines are important sources of airborne particulates that can contain metalloids such as arsenic and lead. Human exposures to this dust can occur through inhalation and ingestion. Children are more likely than adults to ingest dust, and are more vulnerable to the effects of toxic contaminants.

Characterization of airborne dust in regions influenced by mining activity can provide information about sources, fate and transport, and the potential for human exposures.

Particle size affects residence times (how long a particle stays suspended in the atmosphere); smaller particles stay airborne longer. This influences fate and transport, leading to clues about contaminant sources. Coarse particles (>2.5

μm), which are generally created through mechanical action, such as crushing, grinding, or by strong winds, and have residence times of a few hours. By contrast, the fine particles (<2.5 μm) that result from smelting and refining operations have residence times as long as 10 days and can disperse more widely into the environment.

Particle diameter affects aerosol deposition in the human respiratory system. Coarse particles are mainly deposited in the upper respiratory tract (nasal cavity, pharynx, and larynx). They are ultimately swallowed and eliminated through the digestive system. In contrast, fine particles penetrate more deeply into the lungs and can be transferred directly to the blood stream. Thus, determining the distribution of toxic metals in dust fractions also has important implications for human exposure and public health.

At the University of Arizona's Superfund Research Program ([UA SRP](#)), scientists led by Dr. Eric A. Betterton and Dr. A. Eduardo Sáez are performing size-resolved chemical characterization of atmospheric aerosols near mining sites in Arizona. By determining the chemical components of mine tailings dust as a function of particle diameter, their research will generate source apportionment data, improve fate and transport models of airborne particulates, and better assess potential health impacts of contaminated dust.

Advances

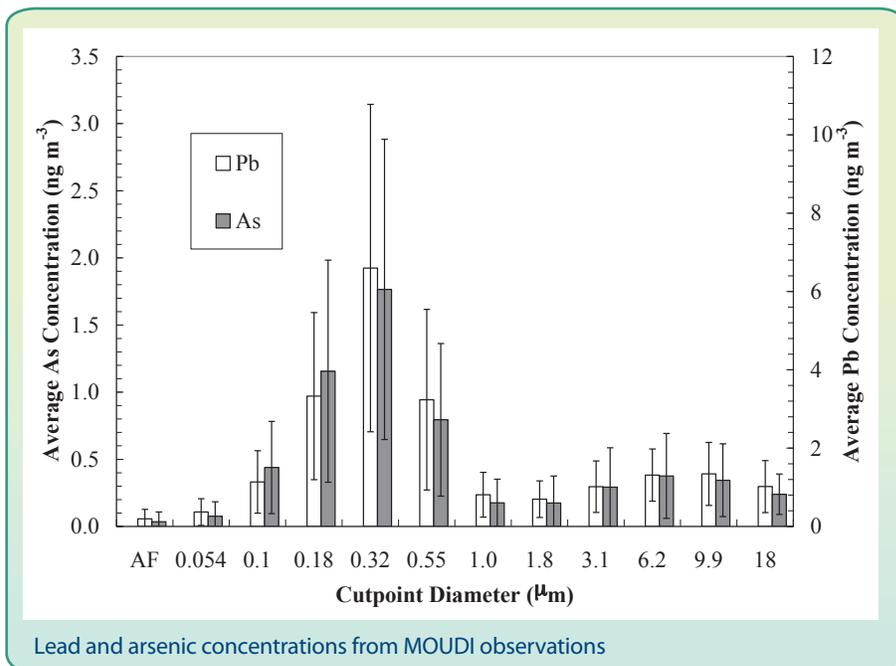
Dr. Betterton and Dr. Sáez's group performed size-resolved chemical characterization of atmospheric aerosols sampled over a period of a year near an active copper mine and smelter site in Hayden, Arizona. In addition to standard measurements using a total suspended particulate (TSP) collector, aerosols were characterized with a 10-stage (0.054 to 18 μm aerodynamic diameter) multiple orifice uniform deposit impactor (MOUDI) and a scanning mobility particle sizer (SMPS).

The MOUDI yields mass concentration of particulates by particle diameter, while the SMPS yields number and volume concentration by particle diameter. The collected particulates were analyzed for metal content by inductively-coupled plasma mass spectroscopy.

TSP samples collected at the active mining and smelting site in Hayden were compared to other sites in southeast Arizona: a clean mountain-top site; a metal-free mine tailings pile; and a typical urban environment. These comparisons demonstrated that airborne arsenic and lead were at least 9 and 4 times higher, respectively, at the Hayden site than any of the other sites sampled.

Scanning electron microscopy and electron dispersive spectroscopy revealed the presence of globular arsenic- and lead-containing particulates, suggesting a history of high-temperature processing in fine-fraction samples collected in Hayden.

The MOUDI samples that showed both arsenic and lead concentrations followed a bimodal distribution, with maxima centered at approximately 0.32 μm and 7.0 μm particle diameter. (Click on the Figure for a larger view). The irregular fractured nature of particles containing arsenic and lead in the coarse-size fraction ($>2.5\mu\text{m}$) suggest that they originate as windblown dust from mine tailings and other sources. The globular fine fraction ($<2.5\mu\text{m}$) contained approximately 70% of the total arsenic and lead, which is likely the product of condensation and coagulation of smelting vapors. In support of this, SMPS data showed the highest readings of ultrafine particle number concentration when the wind came from the general direction of the smelter. Unfortunately, particles detected by the SMPS cannot be chemically analyzed and therefore cannot be assigned to a source. Thus, the ultrafine particles observed with the SMPS could be associated with arsenic or lead vapor, or sulfuric acid production at the smelter.



Significance

More detailed observations of dust fluxes at a Superfund site in [Dewey-Humboldt, AZ](#), will be used to develop a model for the horizontal and vertical flux of particles, and to further characterize contaminant transport from mining and smelting operations. Additionally, Drs. Betterton and Sáez are working with other UA SRP investigators: first, to see if vegetation cover at this site will reduce dust; and second, to supply actual size-fractionated dust samples for analysis of arsenic solid-phase speciation and for inhaled arsenic toxicity studies. Together, these UA SRP projects will provide insights into the source, fate and transport, and health effects of metal-contaminated airborne particulates.

To our knowledge, this is the first demonstration of size-resolved atmospheric aerosols associated with arsenic and lead in the vicinity of mining operations, information necessary for source apportionment and exposure assessment. The location of many mining operations in dry climates, in addition to the warmer, drier conditions predicted for the Southwestern US, may make contaminated atmospheric dust and aerosols an increasingly important exposure route that could result in adverse effects on human health and ecology.

For more information, contact:

Eric A. Betterton

Department of Atmospheric Sciences
 Physics & Atmospheric Sciences Building, Room 542
 1118 E. 4th Street
 P.O. Box 210081
 Tucson, AZ 85721-0081
 Phone: (520) 621-6832
 Email: betterton@atmo.arizona.edu

To learn more about this research, please refer to the following source:

Csavina, Janae, Andrea Landázuri, Anna Wonaschütz, Kyle Rine, Paul Rheinheimer, Brian Barbaris, William Conant, Avelino Eduardo Sáez, and Eric A. Betterton. 2011. Metal and Metalloid Contaminants in Atmospheric Aerosols from Mining Operations *Water Air Soil Pollution* (2011) 221:145–157.
[DOI: 10.1007/s11270-011-0777-x](https://doi.org/10.1007/s11270-011-0777-x)

