Measurement of Exposures to Air Pollutants, Metals and Pesticides

Halûk Özkaynak US EPA, National Exposure Research Laboratory, RTP, NC

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# Measuring Exposures in a Community Health Study

- Typically, there is no single exposure method that can be used to accurately characterize an individual exposures in conducting a community health study
- Measurements methods, include :
  - environmental
  - personal
  - biologic samples
  - questionnaires
  - time activity diaries
  - Source-oriented classifications (e.g., GIS, proximity to roads/agricultural areas)
- Need to consider methods strengths and weaknesses:
  - method difficulties, burden, invasiveness, cost, measurement frequency, ability to archive, exposure period represented, etc.

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## A White Paper on Measurement and Analysis of Exposures for the National Children's Study (NCS)

- A recent White Paper developed by an interdisciplinary team of scientists from EPA, CDC and academia summarizes and interprets environmental and personal exposure and biomonitoring sampling and analysis information for each critical life stage of a child.
- Potential exposure measurements, questionnaires and biological matrices which may be collected by the NCS are summarized in a series of tables by media, route and chemical class
- Can be obtained from: http://nationalchildrensstudy.gov/research/methods\_studies/finalwhite-paper-113004.cfm

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# Likely Agents of Interest

General Classes of Chemical Contaminants	Example Chemicals or Chemical Groups		
Criteria Air Pollutants	PM2.5, PM10, ozone, NO2, SO2, CO		
Environmental Tobacco Smoke	Criteria pollutants above and cotinine		
Biological Agents	housedust mite, rodent, arthropod, pet, fungi, pollen, endotoxins, virus, food allergens, parasites		
Induced Body Products	Hormones (Pregnancy/Adolescence), Histamine (Exercise & Injury), C-reactive Protein (Obesity)		
Metals	mercury (total and methyl-), lead, manganese, tin		
Organophosphorus Pesticides	chlorpyrifos, diazinon, malathion		
Pyrethroid Pesticides	cis- and trans-permethrin, cypermethrin, cyfluthrin, allethrin, bifenthrin, deltamethrin, esfenvalerate, cyhalothrin		
Phthalates	di-2-ethylhexyl phthalate, di-isononyl phthalate, diethyl phthalate, dibutyl phthalate, butyl benzyl phthalate		
VOCs	aromatic hydrocarbons, acrylamide, aldehydes, aliphatic hydrocarbons, halogenated hydrocarbons		
Persistent Organohalogens	PCBs, PCDFs, PBDEs, Dioxins		
Organochlorine and Triazine Pesticides	DDE/DDT, chlordane, heptachlor, lindane, dieldrin, atrazine		

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# Life Stages Of Interest for Exposure Measurement







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# **Object-to-Mouth Contact**



# **Dermal Contact**



# **Dietary and Non-Dietary Ingestion**



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## **Ingestion and Dermal Contact**







Modeling Indirect and Dietary Ingestion

\*1) Slide adapted from: Adgate (2004)
2) Use of product names do not constitute an endorsement

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### 3M Personal Organic Vapor Monitor (OVM)\*



\*1) Source Adgate (2004)
2) Use of product names do not constitute an endorsement



Source: Adgate (2004)

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# Carpet Residue/Dust Sampling



# EPA Baltimore PM Panel Study





Outdoors / Indoors







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# Personal Exposure Monitor (PEM) for Sampling PM<sub>2.5</sub>



#### Examples of media and instruments for evaluation pesticide exposures

#### **Biologic**

#### Environmental

Urine **Cord blood** Blood Soil Saliva **Breast milk** Surface residues **Amniotic fluid Clothing** Meconium Other Other

Air (personal, indoor) Questionnaires Dust Water Food

#### Instruments

Time-activities **Macro activities Micro** activities

Ecologic **Pesticide use** Land use



#### **Personal monitoring**

**Questionnaires** 

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# Criteria for Selecting an Exposure Method in a Community Health Study

- Assess importance of route/pathway by chemical type/class and life-stage
- Evaluate variability and uncertainty in implicit method for biomonitoring or personal or environmental measurements
- Determine suitability of method for testing key field study hypotheses
- Determine whether the method is appropriate for the entire study or on a subset

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# **Evaluation of When to Use Only Questionnaires**

- Identify chemical(s) and associated media, routes, and pathways of exposure and biologic matrix to be measured for study hypothesis as main effect, potential effect modifier, or confounder
- Identify life stage(s) for which the exposure needs to be measured including any critical windows of susceptibility
- Determine whether exposure to the chemical at the critical life stage(s) can be reliably estimated through the use of questionnaire data or other indirect measures (e.g., ambient monitoring and historic use data, time-activity logs, etc.) alone

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# Decision to Use Only Environmental Monitoring

- Biologic measure is not available, or
- Knowledge regarding the route of exposure is critical to testing the study hypothesis or for evaluating exposure mitigation options, and/or
- Exposures must by quantified during critical windows and this is more reliably done using environmental than biologic sampling, and
- Knowledge of target organ dose is not important, or toxicokinetic data is available for estimating target organ dose, and
- Exposures can be more reliably assessed or as reliably assessed using an environmental rather than a biological sample (especially when there is only one critical route of exposure), but the environmental sample is cheaper or participant burden lower.

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# Decision to Use Environmental Monitoring: Examples

- Metals (e.g., manganese) by inhalation
- VOCs with passive diffusion badge
- Criteria pollutants or some organic compounds for which biomarker is either not available or specifically linked
- Exposures to non-persistent compounds (e.g., pesticides). Likely to require repeat multimedia measures design when exposures are intermittent

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# Decision to Use Only Biologic Monitoring

- Knowledge regarding route of exposure is not critical for testing hypothesis, or biomarker reflects critical route of exposure, and
- Biomarker reflects exposure over critical life stage(s), or life stage is not important, and
- Biomarker reflects exposure to target tissue, or knowledge of target dose is not important, and
- Exposures can be more reliably assessed using a biomarker than by using an environmental sample, or
- Exposures can be as reliably assessed using a biomarker as by using an environmental sample, but assessment using the biomarker is cheaper, or the participant burden is lower.

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# Decision to Use Biologic Monitoring: Examples

- Persistent organic pollutants or metals such as lead and mercury or other compounds that are persistent in both the environment and biologic samples
- Plasma or urinary cotinine (as dosimeters of cigarette smoke exposure) or other compounds that are non-persistent in biologic sample but for which environmental exposure is constant
- Chemicals that are non-persistent in biologic sample and environmental exposure is not constant and/or exposures vary across the populations or temporally but for which exposures can be reliably estimated using a biomarker and participant burden and/or cost is lower than for environmental sampling. Likely to require repeat measures design.
- Multimedia exposures that can be characterized at lower cost and participant burden using an internal dosimeter rather than multiple route environmental sampling.

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# Decision to Use Both Environmental and Biologic Monitoring with Questionnaires

- Information regarding exposure route is critical, but exposures can't be reliably assessed with only environmental sampling, or
- Exposures must by quantified during critical windows and is more reliably done with environmental, and biologic sampling, or
- Biologic sampling is adequate to quantify internal dose, but environmental sampling is needed to characterize exposure route, or
- Environmental sampling is adequate to characterize exposure route, but biologic sampling is needed as internal dosimeter, or
- Exposures cannot reliably be assessed using either biologic or environmental sampling alone (e.g., pesticides or other non-persistent organic compounds may require intensive and repeat sampling depending on the scenario.

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### **Disclaimer**

Although this work was reviewed by EPA and approved for publication, it may not necessarily reflect official Agency policy.









# ISEA 2005 the 15th Annual Conference in Tucson, Arizona



# OCT 30<sup>TH</sup> - NOV 3<sup>RD</sup>, 2005

at the Westin La Paloma Resort & Spa



# Conference Deadlines

April 30, 2005	July 15, 2005	Up to October 1, 2005	Early June	Accepted through September, 2005
Abstract	Early Conference	Vendor and Sponsor	Notification of Paper	Late Breaking
Submissions*	Registration	Registration	Acceptance/Rejection	Posters

\*Visit www.conferencemanagement.net/isea for more information on submitting abstracts.

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& Mary Kay O'Rourke

All workshops are on a first come first serve basis. When the seats are filled, the workshop is closed. All fees due upon registration. ISEA reserves the option to cancel any workshop and fully refund the fees at the time of registration.

MINI-SYMPOSIA: Additional Symposia will be forthcoming on the ISEA conference website.

- · Air Pollution and Asthma Children's Centers Symposium (Chris Saint)
- Children's Exposure to Toxics Children's Centers Symposium (Chris Saint)
- Potential Exposures to Chemical or Biological Agents: Risk Evaluation, Mitigation and Planning (Brett Singer & Richard Sextro)
  - Environmental Justice Issues (Sarah Elwood)
- Exposures Along the US-Mexico Border (Shaibal Mukerjee, Rick Van Schoik & SCERP Investigators)
  - Exposure Assessment Needs for Environmental Policy, and Validity (David MacIntosh)
- Reliability of Bioaerosol Exposure Assessment Methods (David MacIntosh)
- Assessment of Environmental Exposures to Microbiological Agents (John Scott Meschke & Kristina D. Mena)
  - Exposures to Arsenic (Robin Harris & Séumas Rogan)
- Breast Milk Monitoring for Environmental Contaminants: A Useful Bio-fluid as a Measure of Maternal Body Burden and Infant Dose (Judith Schrieber)
- Revisions to the Exposure Assessment Guidelines of 1992: Proposed Changes and Panel Discussions (Gary Bangs)
  - Development of State and Regional Laboratory Capacity (Larry Needham)
- Understanding Dietary Exposure to Chemical Contaminants (Lisa Melnyk)
- The Detroit and Windsor PM and Air Toxics Research Studies Study Designs and Early Findings (Ron Williams & Amanda Wheeler)
  - Exposure Modeling for Outdoor and Indoor Air Pollution: Methodology, Applications and Evaluation of Models (Ted Palma & Stephen Graham)

# CONFERENCE SESSIONS

- Biological Aerosols
- Biomarkers & Exposure
- Bioterrorism & Other Disasters
  - Children's Exposure
- Climate Change & Exposure
  - Chillate Change & Exposure
  - Community Exposure
- Disparities in Exposure & Health
   Exposure Assessment
  - (Aggregate & Cumulative)
    - Exposures on Borders
- · Exposure & Genetics
- Emerging Assessment Technologies
  - GIS & Exposure
- Exposure Modeling

- · Exposure & Health Surveillance
  - · Gene / Environment Interactions
    - Pesticides
- Population Studies & Exposure
  - · Pick Accessment
  - Risk Assessment
     Scoles of Evenent
    - Scales of Exposure
      - Selected Toxins
        - Water
- Workplace Exposures
- Proposed sessions (watch website)



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