

Biomonitoring Chemicals with Short Half-lives in Humans: Issues in Interpreting and Communicating Individual Results

Materials for: July 26, 2012 Meeting of Scientific Guidance Panel (SGP)
Biomonitoring California¹

For chemicals with short half-lives, biomonitoring measurements can be very variable over short periods of time. An individual result does not necessarily characterize a person's exposure to a chemical over time; even a short time, like a day or week. Biomonitoring California is in the process of developing results return materials for a broad range of chemicals, including those with short half-lives. Part of this process involves usability testing of draft materials with a small set of participants. As another part of this process, we are seeking SGP input on the following discussion questions:

- What additional context, if any, might be important to provide to participants on interpreting their individual results for chemicals with short half-lives, beyond the standard template? What basic messages do you suggest we try to convey?
(See this link for an example of the template:
<http://www.oehha.ca.gov/multimedia/biomon/pdf/03162012FOXMockResultsPacket.pdf>)
- One approach to providing context would be to give information to participants about how the half-life of a chemical could affect their individual results. If we chose this approach, what type of information would be most important to include on half-life and how do you suggest we frame such information?
- Half-life is only one of many factors that affect an individual's results for a given chemical. Which other relevant factors do you think would be important to explain to participants?
For example:
 - Repeated exposures, such as via routine product use
 - Timing of when a biological sample is taken, such as after a meal
- Do you have other comments on interpreting and/or communicating biomonitoring results for chemicals with short half-lives, from your background reading or your own experience?

Selected Background References (those marked with an asterisk [*] were sent to the SGP):

Aylward L. et al. (2012). Interpreting variability in population biomonitoring data: Role of elimination kinetics. *J Expo Sci Environ Epidemiol*. doi: 10.1038/jes.2012.35. [Epub ahead of print]

Braun J. et al. (2012). Variability of urinary phthalate metabolite and bisphenol A concentrations before and during pregnancy. *Environmental Health Perspectives* 120 (5):739-745.

¹California Environmental Contaminant Biomonitoring Program, codified at Health and Safety Code section 105440 et seq.

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Hildebrand S et al. (2009). Biomonitoring of the di(2-ethylhexyl) phthalate metabolites mono(2-ethyl-5-hydroxyhexyl) phthalate and mono(2-ethyl-5-oxohexyl) phthalate in children and adults during the course of time and seasons. *Int J Hyg Environ Health* 212(6):679-684.

*Li Z. et al. (2010). Variability of urinary concentrations of polycyclic aromatic hydrocarbon metabolite in general population and comparison of spot, first-morning, and 24-h void sampling. *J Expo Sci Environ Epidemiol* 20:526–535.

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Peck J et al. (2012). Intra- and inter-individual variability of urinary phthalate metabolite concentrations in Hmong women of reproductive age. *J Expo Sci Environ Epidemiol* 20(1):90–100.

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Teitelbaum S. et al. (2008). Temporal variability in urinary concentrations of phthalate metabolites, phytoestrogens and phenols among minority children in the United States. *Environ Res* 106:257–269.

*Ye X et al. (2011). Variability of urinary concentrations of bisphenol A in spot samples, first morning voids, and 24-hour collections. *Environmental Health Perspectives*, 119(7):983-988.