

## CORRESPONDENCE

## The Assessment of Environmental and Occupational Exposure to Hazardous Substances by Biomonitoring

by PD Dr. rer. nat. Lygia T. Budnik, Univ.-Prof. Dr. med. Xaver Baur  
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### Sum of Many Factors

Although biomonitoring for lead toxicity is based on hemotoxicity, lead is known to act through many other pathological mechanisms. For example, lead is known to damage beta-receptors and the renin-angiotensin-aldosterone system, leading to increases in blood pressure (1); this is not included in biomonitoring. At any rate, lead's hypertensive activity has been known for 120 years and the American environmental agency considered it appropriate to publish a systematic review of cardiovascular damage from lead in March 2007, containing 130 references (2). This should be considered in Germany too.

It was repeatedly clear in this article that the toxicity of a substance is influenced by the sum of many different factors. This has also been pointed out by Emily F. Madden of the American Food and Drug Administration (FDA). At the end of a review of the interactions between multiple environmental or occupational exposures to heavy metals in the development of cancer, she concluded that the combined effects of different metals may be greater than the sum of their individual effects (3). Perhaps then the cumulative effects of different metals might also effect the development of disease, even when they are all within the normal ranges. In my opinion, these aspects of chronic and complex exposure to heavy metals and pollutants deserve more attention, as this could help to guarantee comprehensive protection of the population, particularly of children and of people who are already ill.

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### Epidemiological Benefits

It is generally accepted that industrial biomonitoring is a valuable tool as it helps to control workplace-related threshold limit values. It is also capable to determine a quantitative relationship between external and internal exposure and to support the diagnosis of diseases caused by acute effects of harmful substances. Biomonitoring may also be used in epidemiological studies to determine group differences between subjects with high and low exposure to improve the description of average pollutant concentrations from chronic exposure.

However, when assessing health risks of chronic diseases with long induction periods that arise from these exposures, the article is a bit short-spoken. Biomonitoring is less meaningful when studying these diseases, such as most types of cancer (1). Due to the short half-life of most of these agents, past exposures can hardly be estimated by biomonitoring procedures. In some cases risk estimation is only possible when complex mathematical models are applied.

Occupational epidemiologists use a variety of methods to evaluate these exposure scenarios, including historical ones, and the associated risks. Data sources, such as registry data, archive materials, expert evaluations and personal assessment by employees, as well as biological parameters or workplace measurements, may be for example combined in job exposure matrices which allow for automatic classification of historical exposures to evaluate health risks in epidemiological studies.

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### In Reply:

Extremely complex interactions may occur when subjects are exposed to multiple pollutants. There are many unsettled issues and unknowns here. It is not essential that individual pollutants (such as heavy metals) should exceed reference or limit values. Additive, superadditive or even antagonistic effects are possible here, although only initial studies have been performed. Detailed recent studies have shown that lead does indeed cause a variety of adverse effects besides its well known neuro-toxicity and hematotoxicity. Timely and sensitive biomonitoring

can help in clarifying the etiology, or in confirming or excluding the etiological role of lead.

Exposure biomonitoring alone is generally not helpful in risk assessment or in determining the etiology of a disease when the exposure is long past. As our article describes, biochemical and biological effect monitoring (such as DNA- or protein-adduct formation or chromosomal aberrations [1, 2]) are capable of recording long-term effects and can lead to important new information about dose-effect relationships at the molecular level. These validated toxicological data may be combined with other procedures, such as occupational medical assessments, and analyzed by mathematical models to give more precise risk assessments.

What is important is that biomonitoring results of this kind should be interpreted in the context of clinical findings. Evaluation of carcinogenic effects must include the long periods of latency, individual susceptibility and cocarcinogenic aspects. The medical advisory committee on occupational diseases has recently recommended that the interaction between two occupational agent groups, namely polycyclic aromatic hydrocarbons and cocarcinogenic asbestos fibers, should be included in the list of occupational diseases (3).

Classical exposure monitoring can be used in long term studies, for example, to investigate persistent biological substances which are absorbed from the environment and accumulated in the body. The American Environmental Protection Agency and the WHO are currently

performing large scale studies to investigate the background exposure of the population to these pollutants (4). DOI: 10.3238/arztebl.2009.0508a

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**Conflict of interest statement**

The authors of all contributions declare that no conflict of interest exists according to the guidelines of the International Committee of Medical Journal Editors.