

Comprehensive Characterization Strategies for Ultrafine Particles: Lessons from Beryllium Health and Safety Studies.

A.B. Stefaniak¹, G. A. Day¹, R.C. Scripsick², and M.D. Hoover¹

¹ Division of Respiratory Disease Studies, National Institute for Occupational Safety and Health, Morgantown, WV, USA

² Health Safety and Radiation Protection Division, Los Alamos National Laboratory, Los Alamos, NM, USA

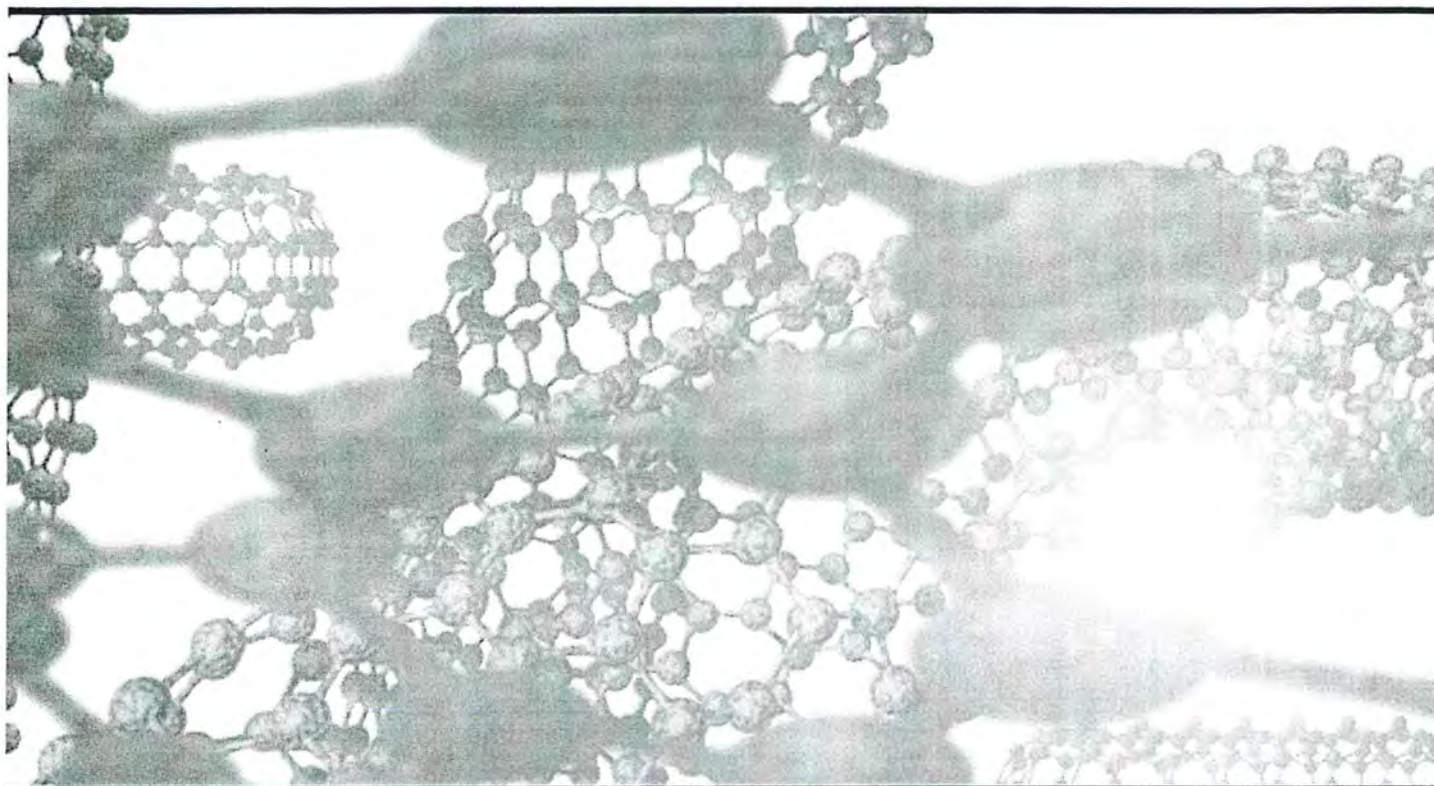
Information on the physicochemical and bioavailability properties of nanoparticles is needed to help identify appropriate metrics of exposure (e.g., airborne mass, particle size, surface area, and/or chemical composition) for protecting worker health. Knowledge of appropriate exposure metrics would aid in developing exposure limits based on bioavailability through the inhalation and dermal routes of exposure. Thus, a suite of standard analytical techniques for characterization of properties of nanoparticles would be of value. Experience from studies of ultrafine beryllium aerosols demonstrate the capabilities and limitations of physicochemical and bioavailability characterization techniques as well as practical considerations for analysis.

Traditional analytical techniques and methodologies for ultrafine particles (e.g., microscopy, electron and x-ray analyses, surface area, and spectroscopy) are applicable for nanoparticles. Care must be taken to ensure that particle properties measured in the laboratory are representative of the particle properties in their native environment. These considerations would include the type of test (destructive or nondestructive), and influence of sample preparation (heating, drying, etc.) and analysis (e.g., beam probe-particle interaction) on the particle sample.

Appropriate cell-free *in vitro* models are needed to study bioavailability of a material in a target biological compartment. Standard dissolution solvents include serum ultrafiltrate (SUF), a simulant of extracellular lung fluid having neutral pH; artificial sweat, a simulant of skin surface fluid having pH 6.5; and phagolysosomal simulant fluid (PSF), a simulant of the macrophage phagolysosome having pH 4.5. Traditional dissolution techniques, e.g., static, flow-through, etc. are applicable for nanoparticles. Note that use of membrane filters having pore sizes of 0.025 μm or greater to separate the particle sample from the dissolution solvent in these techniques may not always be feasible for nanoparticles. Thus, alternative approaches to preventing solid material from migrating into the solvent to bias dissolved mass measurements may be needed, e.g., isolating via centrifugation, ultrafiltration, or using density gradient separation.

Nanomaterials

a risk to health at work?



First International Symposium on Occupational
Health Implications of Nanomaterials

12-14 October 2004
Palace Hotel, Buxton, Derbyshire, UK

Report of Presentations at Plenary and Workshop Sessions and
Summary of Conclusions

David Mark
Chair of Symposium
Health and Safety Laboratory,
Buxton, SK17 9 JN



NANOMATERIALS – a risk to health at work?

First International Symposium on Occupational Health Implications of Nanomaterials

**Palace Hotel, Buxton, Derbyshire, UK
12 – 14 October 2004**

Report of Presentations at Plenary and Workshop Sessions and
Summary of Conclusions

David Mark,
Chair of Symposium
Health and Safety Laboratory,
Buxton, SK17 9JN
www.hsl.gov.uk