Identifying & Regulating Environmental Impacts of Nanomaterials

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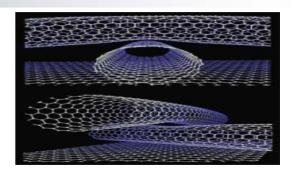
Regulatory Gaps

Nano-products likely to fall through regulatory gaps due to classification and nomenclature issues

- Which regulatory gaps apply to which nano-products
- To what extent, and where in the product life cycle
- Weight the risks that arise from regulatory gaps

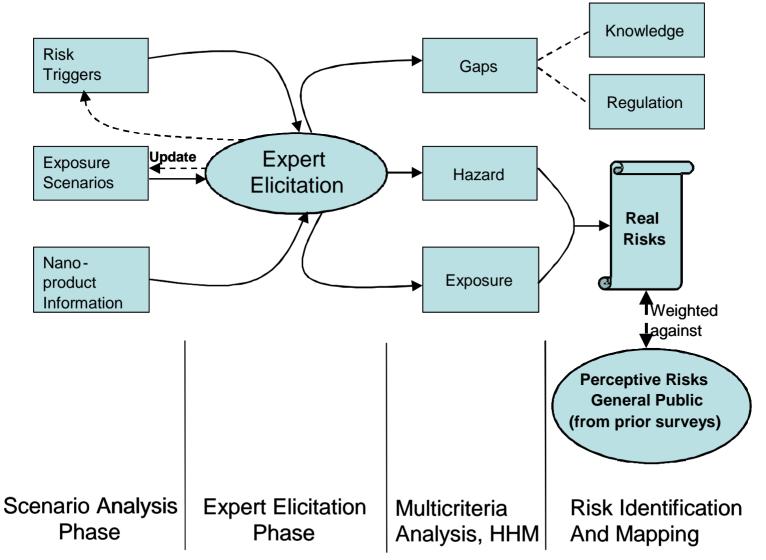
Life Cycle Stage	Relevant Risk Triggers	Possible products
R & D, Manufacturing and Application stage		
TSCA: R&D, Test marketing, Low volume exemption		Products that contain carbon nanotubes (Field Emission Displays), perhaps tennis rackets
FDA: Regulatory centers	FDA Nomenclature: Device or a Drug?	Sunscreen, Toothpaste, Food Supplement, MRI contrast agent
FIFRA: Pesticidal or antibacterial effects?	Antibacterial - harm useful bacteria in the environment or the human body?	Toothpaste, Air freshener
End-of-Life & Disposal stage		
RCRA: classification under RCRA?		Air freshener spray, Battery, MRI contrast agent, Field emission display
Transporting stage		
HIMLA		Battery, MRI contrast agent, Field emission display
Long Term Environmental Site Implications stage		
CERCLA (Superfund): Long term environmental, health risk		Air Freshener, Battery, Racquet, Field Emission display

Case Study: Carbon Nanotubes and the Low Volume Exemption



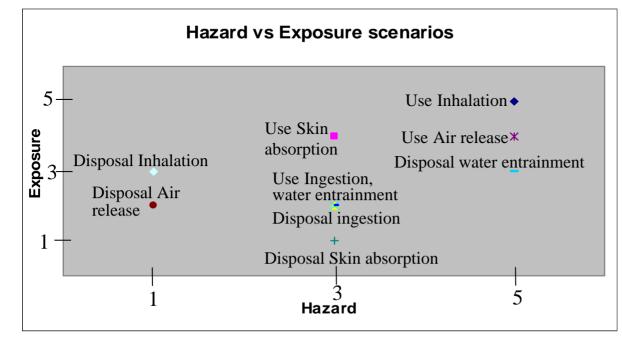
- Life Cycle Stage: Manufacture
- Regulatory Gap: Under TSCA, LVE limits regulation of facilities producing <10,000 kg/year
- Scenario:
 - About 44 carbon nanotube producers (Cientifica), and about 100 metric tons produced per year (UK Royal Society)
 - "A 40 inch computer display uses one-sixth of a gram of carbon nanotube powder (roughly 10,000 nanotubes) (Mann 2004). If all 25 metric tons of carbon nanotubes going to electron emission applications (estimated above) are used for computer displays, they would enter into 150 million displays."
- Implications:
 - □ Potential environmental impacts are widespread with no attention to risk.
 - Nanomaterials change the risk assessment paradigm. Mass does not correlate risk or exposure.

Methodology: Scenario Analysis & Expert Elicitation



Risk Identification for Air Fresheners

<u>Product Information</u>: 20-50 nm silver nanoparticles in polymer matrix with antibacterial properties (used as spray) <u>High-risk scenarios</u>: Inhalation and air release during use, water entrainment during disposal <u>Nano-property risk triggers</u>: Easy bio-availability, anti-bacterial property, ease of dispersion, affect on susceptible populations

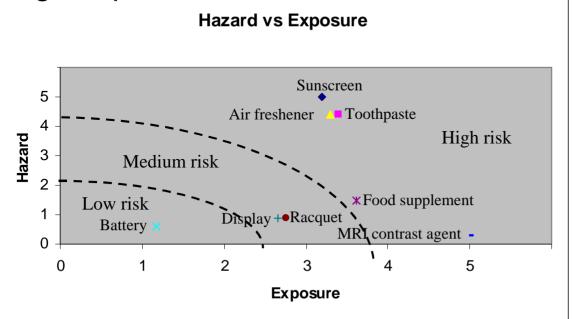


Intersection Scenarios:

- Wider bio-availability during air-release or water entrainment
- Anti-bacterial effects outside product cycle
 Susceptible Population (respiratory problems)

Risks from cross-section of products

Relative risks higher for sunscreens, toothpastes, and air fresheners based on number and weights of both 'high hazard' and 'high exposure' scenarios
Food supplements and MRI contrast agents pose some significant scenarios for high-hazard, but few scenarios for high exposure



Risk higher for products where Nanoparticles can be disengaged from the matrix or composite to which they are bound

Challenges & Opportunities

Risk:

- Including regulatory and knowledge gaps in framework
- Criteria to weigh risks arising from various sources

Regulatory:

- □ Statutes regulate products not technologies
- Dispersed through multiple agencies

Toxicology:

- Need standardized methods to collect toxicology data for application within risk context
- Risk versus benefit analysis
- Hazard communication to decrease exposure

Publications

- (In Press). Environmental Regulatory Implications for Nanomaterials under the Toxic Substances Control Act (TSCA), IEEE Technology & Society
- The Product Life Cycle and Challenges to Nanotechnology Regulation, Nanotechnology Law and Buiness, Vol 3, Issue 4.
- (Submitted). Identification of Risks in the Life-Cycle of Nanotechnology-Based Products, Journal of Industrial Ecology.
- Conference Presentations at MRS Fall 2005, NSTI Summer 2007

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