HOW ULTRAFINE PARTICLES MAY INTERACT WITH PULMONARY CELLS

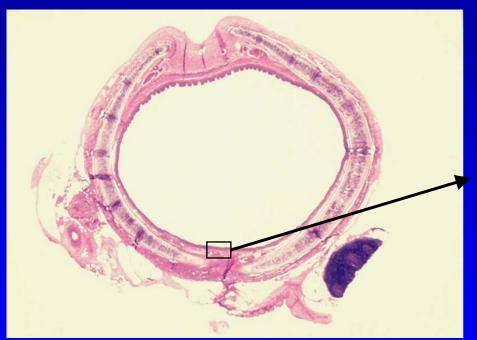
P. Gehr, B. Rothen, S. Schürch*, N. Kapp, M. Geiser,

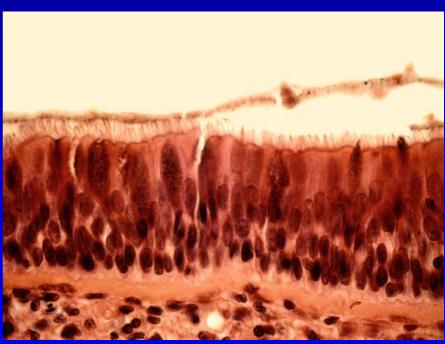
Institute of Anatomy, University of Bern, Bern, Switzerland

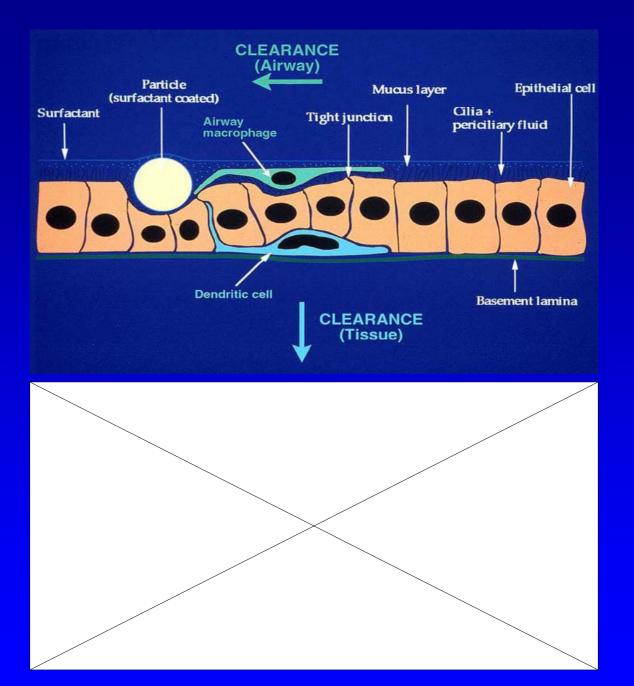
*Health Sciences Center, University of Calgary, Calgary, Alberta, Canada

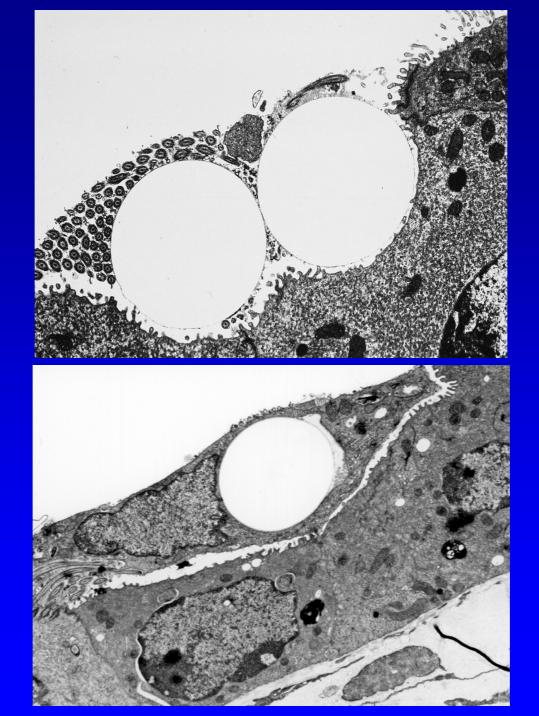
Trachea

Airway epithelium

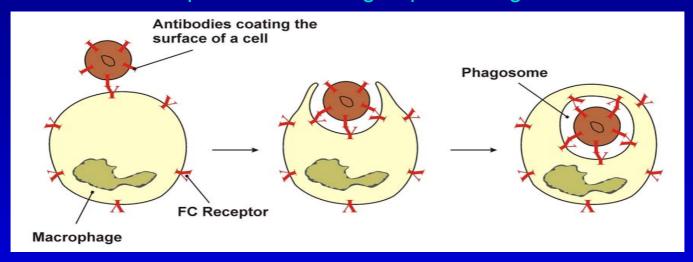




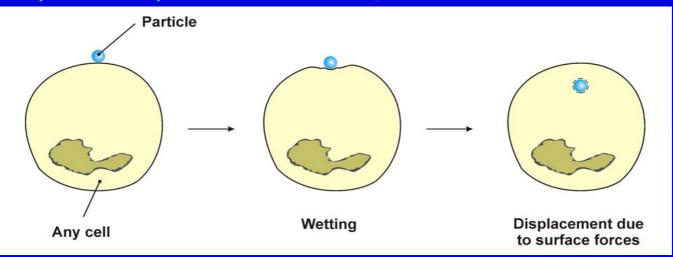




Phagocytosis: • professional phagocytes: macrophages • professional antigen-presenting cells: dendritic cells



Non-specific uptake: • macrophages, dendritic cells, epithelial cells etc.



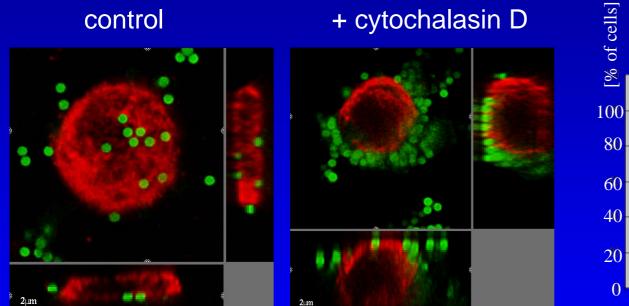
Models

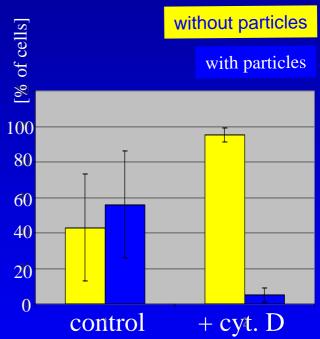
- phagocytic cells: macrophages
- non-phagocytic cells: red blood cells

Incubation with FITC labeled polystyrene particles of different sizes

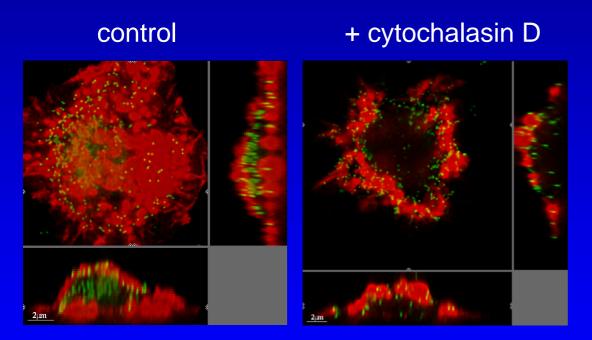
Preparation for confocal laser scanning microscopy

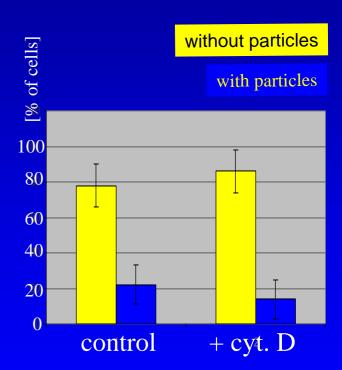
■ F-Actin 1-μm particles



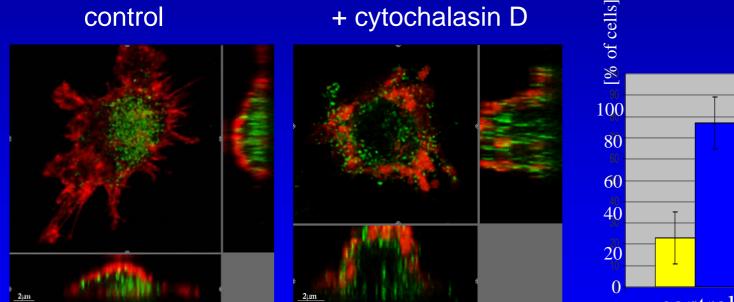


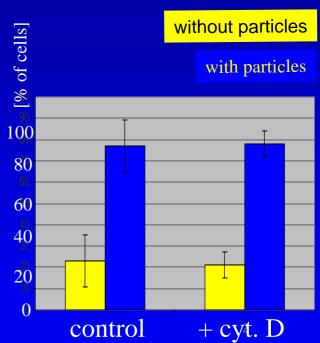
■ F-Actin 0.2-μm particles (neutral)



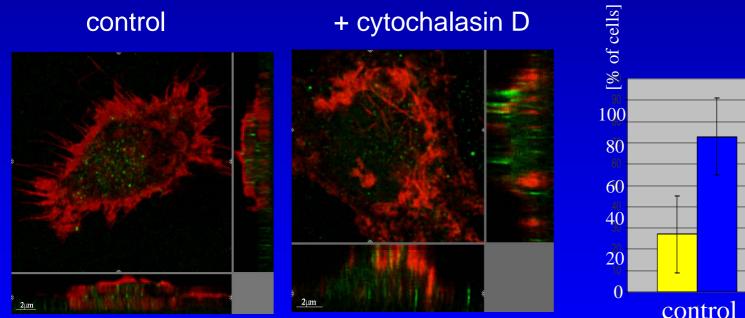


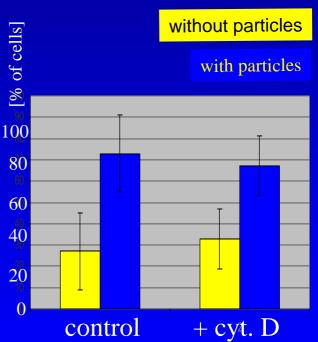
F-Actin 78-nm particles (neutral)



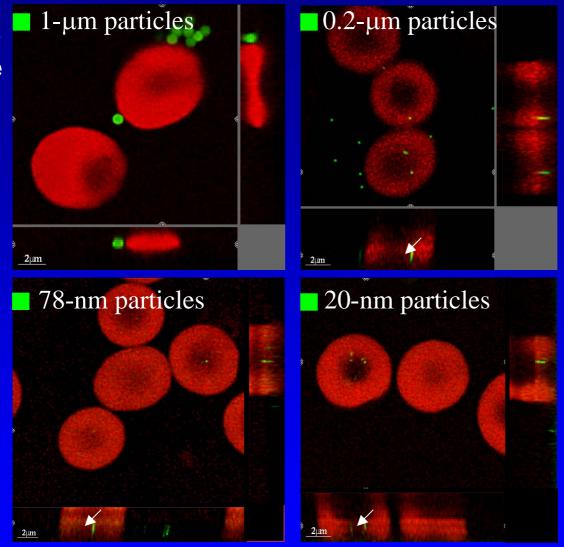


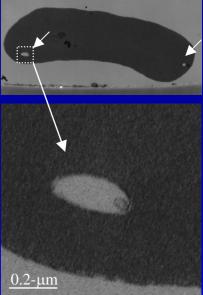
F-Actin 20-nm particles (negatively charged)





Autofluorescence





Summary

1-μm particles

- taken up by phagocytosis
- no uptake by red blood cells

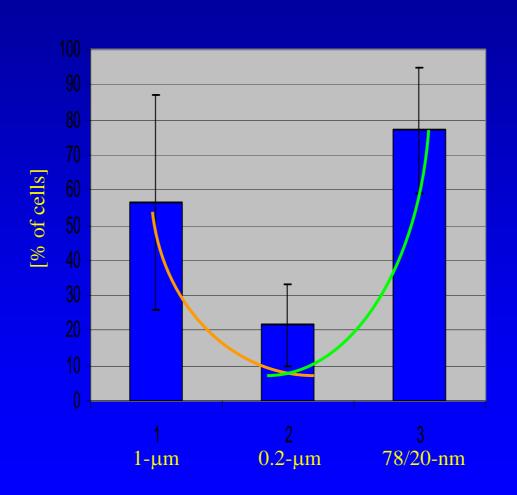
0.2-μm particles

- reduced uptake, mechanism not clear
- uptake by red blood cells

78-nm and 20-nm particles

- increased uptake by adhesive interaction
- uptake by red blood cells

with particles



non-specific uptake

Phagocytosis

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