

MONITORING INDIVIDUAL POLLUTANT PARTICLE BEHAVIOUR ON IN-TACT LIVE AIRWAYS USING SYNCHROTRON X-RAY IMAGING

& Children's

Hospital

Government of South Australia

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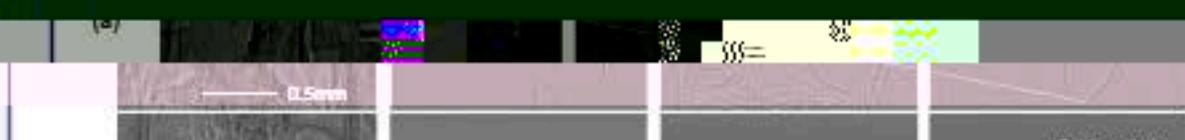
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Background

Non-biological particles small enough to be suspended in the air are continually inhaled as we breathe, and deposit on airway surfaces where they can remain and affect lung health. Pollutant particles from vehicles, buildings and industrial dusts have the potential to cause immediate and delayed health problems. Due to their small size, it has not been possible to non-invasively examine how individual particles move on the airway surface after deposition. Using live intact mouse airways we have begun to examine particle behavior after deposit on the airway wall, dynamically, non-invasively, using synchrotron phase contrast X-ray imaging.

In-vivo Results

The two panels on the left show the original X-ray image and its corresponding motion-detected frame that revealed the moving object on the airway. The panels on the right mark the nasal airways and show the same sequence of images each separated by 5 seconds.

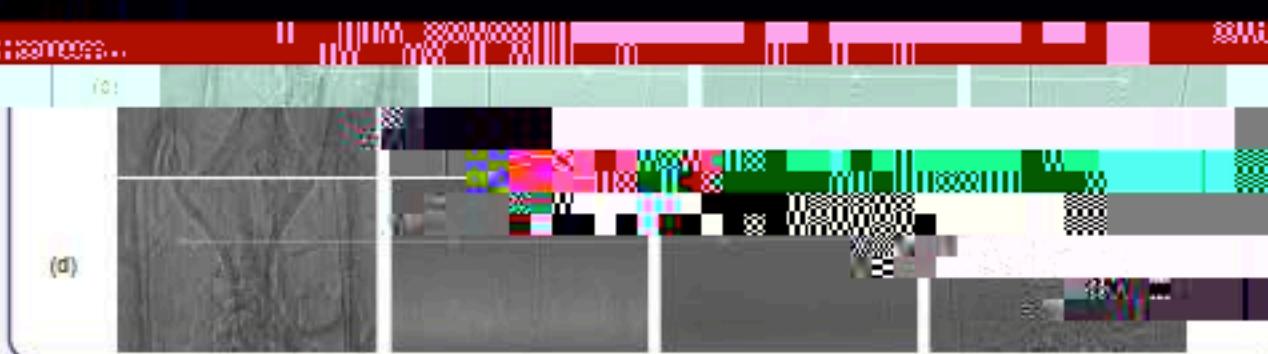


Materials and Methods

Experiments were performed on the RI 20Y11 beamline at the SPRU, University of Southampton.

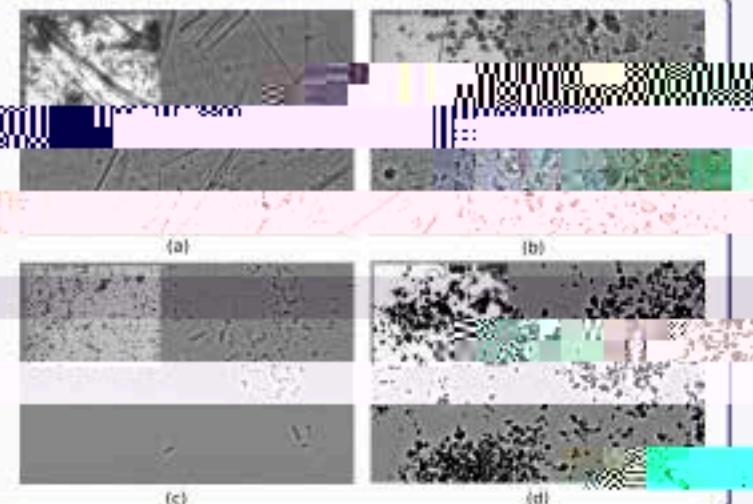
Asbestos particulates was determined. Asbestos

particulates were obtained from the National Institute for Occupational Safety and Health (CD1-Foxn1-nu). Mice were secured head-high on an imaging board, and the X-ray beam (dimensions 10 x 6mm) was directed ventro-dorsally through the mouse to image the nasal airways (where the ciliated epithelium is used as a model for the conducting airways of the human lung).



In-vitro Results

In-vitro dry particulate samples under PCXI (main pictures) and light microscopy (insets) (a) asbestos, (b) quarry dust, (c) fine PM10, (d) toner. The morphology of each of the particulates is clearly very different. The largely carbon-based particulates — combusted diesel, PM10 and laser printer toner — were not sufficiently visible to warrant *in-vivo* testing.



Conclusion

Individual particulates can be tracked non-invasively in live airways using PCXI. Further refinement of particle size and delivery techniques PCXI should provide a novel approach monitoring the behaviour of particles on airway walls.

Acknowledgements

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