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Talbot, Nicholas
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PARTICLE SIZE, MASS AND CHEMICAL TRANSITIONS FROM AN OUTDOOR TO INDOOR ENVIRONMENT IN PRAGUE, CZECH REPUBLIC WITH ATTENTION TO NITRATE

Nick TALBOT^{1,2}, Lucie KUBELOVÁ^{1,2}, Otakar MAKEŠ^{1,2}, Michael CUSACK¹, Jaroslav SCHWARZ¹, Petr VODIČKA¹, Jakub ONDRÁČEK¹, Vladimír ŽDÍMAL¹

¹Institute for Chemical Process Fundamentals of the ASCR, v.v.i., Prague 6, Czech Republic, talbot@icpf.cas.cz

²Department of Environmental Studies, Faculty of Science, Charles University, Prague, Czech Republic.

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INTRODUCTION

In the absence of internal sources, particle number concentrations indoors are largely dependent on concentrations outdoors (Hussein *et al.*, 2006). Once indoors, particles are subjected to changes in temperature, relative humidity and different organic compounds, changing the particles physico-chemical state. (Lunden *et al.*, 2003).

As people spend most of their time indoors it's important to understand how outdoor originating particulate can migrate indoors, their relative abundance indoors and how physico-chemical processes acting upon particulate can change the overall aerosol composition. Species volatility can complicate indoor data, change mass size distribution profiles, and implant artifacts on data measurements, enhancing uncertainty on results. This research describes the findings from high temporal resolution I/O, offline and online measurements, identifies outdoor aerosol loading dynamics and describes proportional compositions indoors relative to calculated I/O ratios.

EXPERIMENTAL SETUP

Data was collected from a ground floor flat in Suchbát, Prague during summer 2014. The flat is located in a residential area in northwestern Prague about 6 km from the city centre and is recognized as a suburban background site. Particle number size distribution profiles were collected using a Scanning Mobility Particle Sizer (SMPS) model TSI 3936L consisting of a long DMA 3081 and CPC 3775. The SMPS scanned 64 channels per decade, through sizes 14.8-724nm with 3-minute up-scan, 30 second down scan and 90 second purge. An Aerodyne Compact Time of Flight Aerosol Mass Spectrometer (C-ToF AMS) ran alongside the SMPS sampling every minute, providing real-time chemical composition data (Drewnack *et al.*, 2005).

Sequential sampling was conducted using an automated switching valve which allowed two-5 minute SMPS scans outdoors followed by two scans indoors. Two Berner Low Pressure Impactors (BLPI) were used for a size resolved chemical composition analysis with one BLPI located indoors and one outside. Substrate filters were changed every 24 hours. Meteorological data was collected from a permanent air research station located within the same ICPF compound where all measurements were collected.

RESULTS AND CONCLUSIONS

A New Particle Formation (NPF) event on the afternoon of Saturday 30th August shows a clearly defined, size resolved, outdoor-indoors infiltration [Fig 1]. This allowed for the calculation of an I/O ratio using time/size subsets during the NPF event. Ultrafine particles <40nm in diameter had the lowest I/O ratio at 0.45 whilst particles >80nm in diameter gave a consistent ratio between 0.65-0.7. Apparent growth rate defining particle growth between 20-50nm during the same NPF event were 4.4nm/hr outdoors.

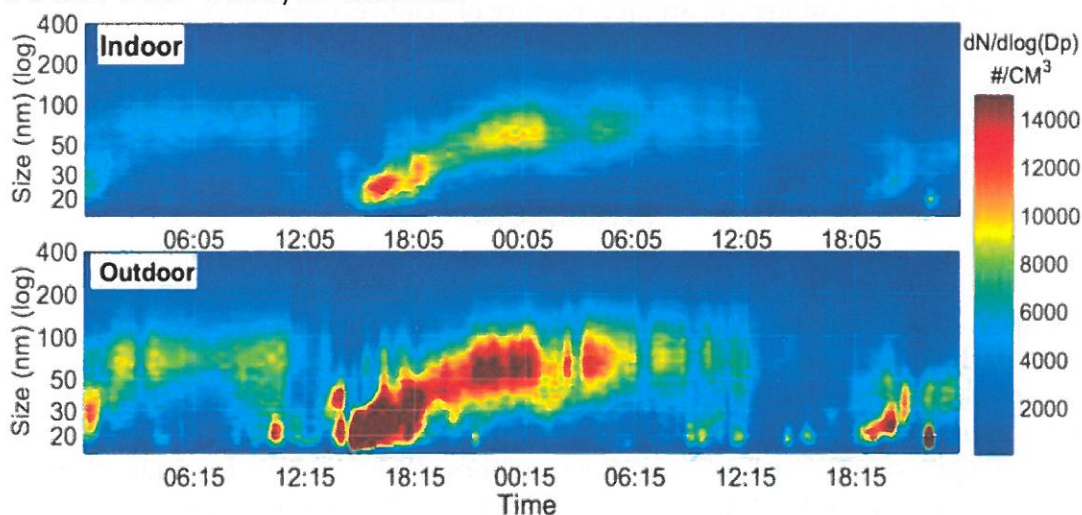


Fig 1: NPF event outdoors seen to infiltrate indoors with changing size and concentration.

Size characterization of NR-PM₁ from AMS measurements gives outdoor and indoor modal diameters for Organic matter, NH₄, NO₃ and SO₄ between 300-500nm. Further intercomparison of these non-refractory species shows indoor NO₃ concentrations were proportionally far lower than expected when considering the I/O ratio. The known volatility of NH₄NO₃ was identified as the most likely cause. Larger NO₃ mass concentration losses were observed on substrate from concurrent BLPI measurements. This difference highlights the time dependent factor when accounting for semi-volatile species.

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