



Evaluating Airborne Exposures to Microplastic Particles

**Inhaled Particles XIII – NanOEH Conference
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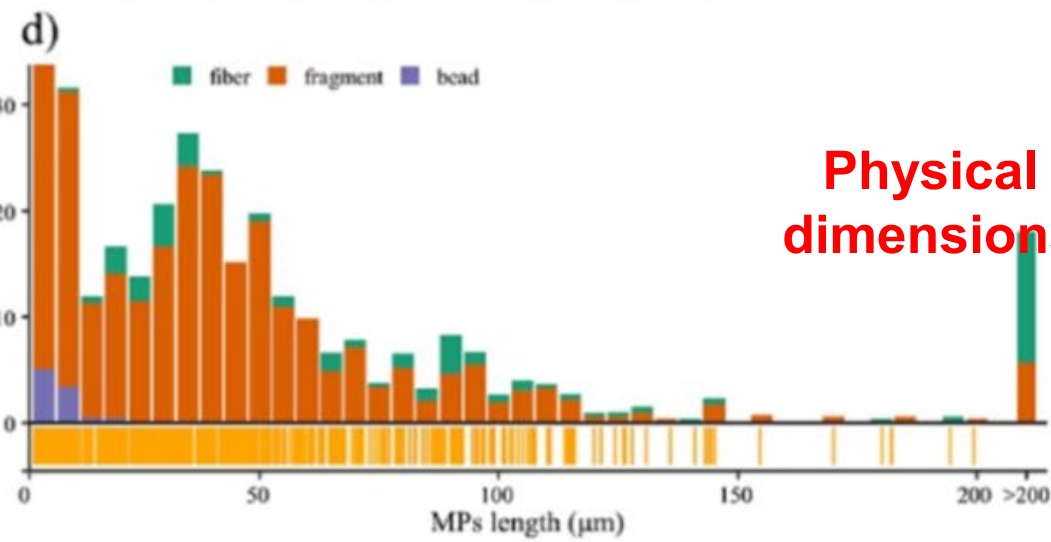
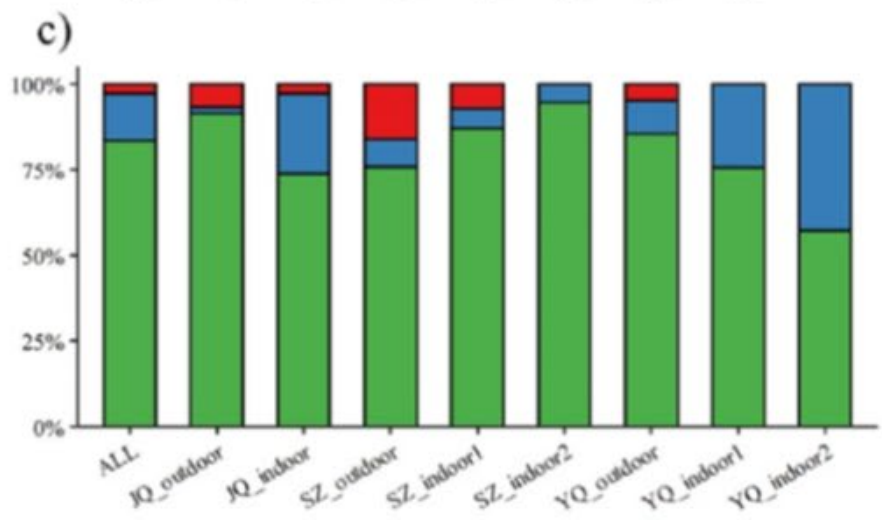
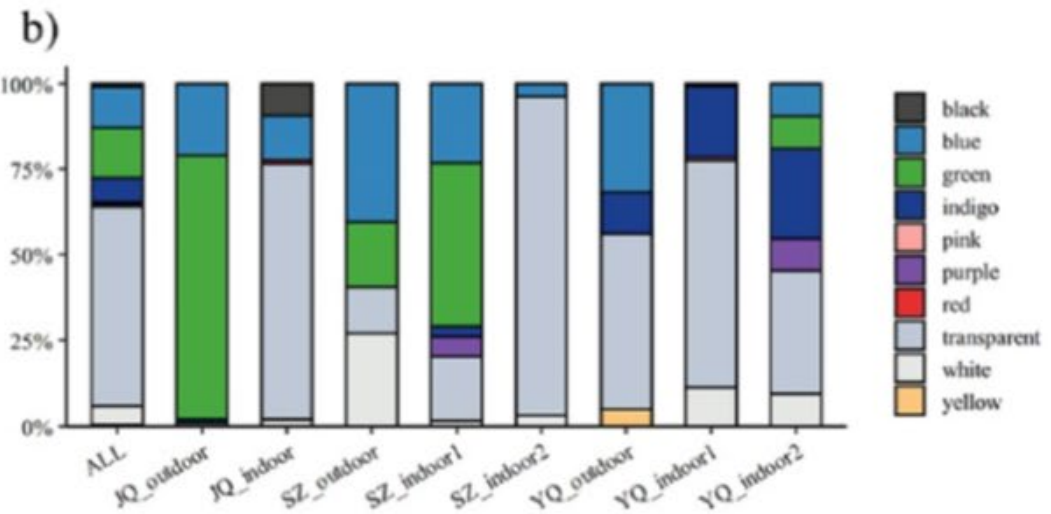
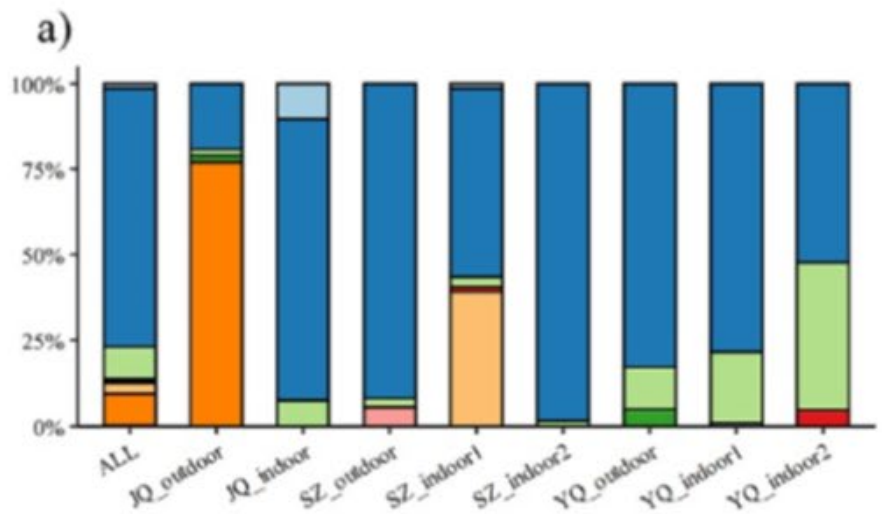
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Plastics are Ubiquitous in the Environment: Airborne Microplastics (MPs)

- Indoor and outdoor air sampling have demonstrated the presence of plastic particles that have a range of compositions, morphologies, and sizes
 - Submicrometer (incl. nanoscale) up to several mm in length and/or diameter (*Rahman et al., 2021; Wright et al., 2019; Zhang et al., 2020*)
 - Fibers, fragments, spheres (beads) (*Xie et al., 2022*)
 - Polypropylene, polyethylene, polyester, polyvinyl chloride, nylon (*Vianello et al., 2019; Xie et al., 2022*)
- Plastic particles can be distinguished from other airborne particulates
 - Available literature suggests that 4-50% by mass of sampled particles are composed of synthetic polymers (*Brahney et al., 2020, 2021; Dris et al., 2017; Prata et al., 2020*)
- Polymeric particles have been identified in digested excised human lung tissue (*Jenner et al., 2022*)

Broad Characteristics of Airborne MPs

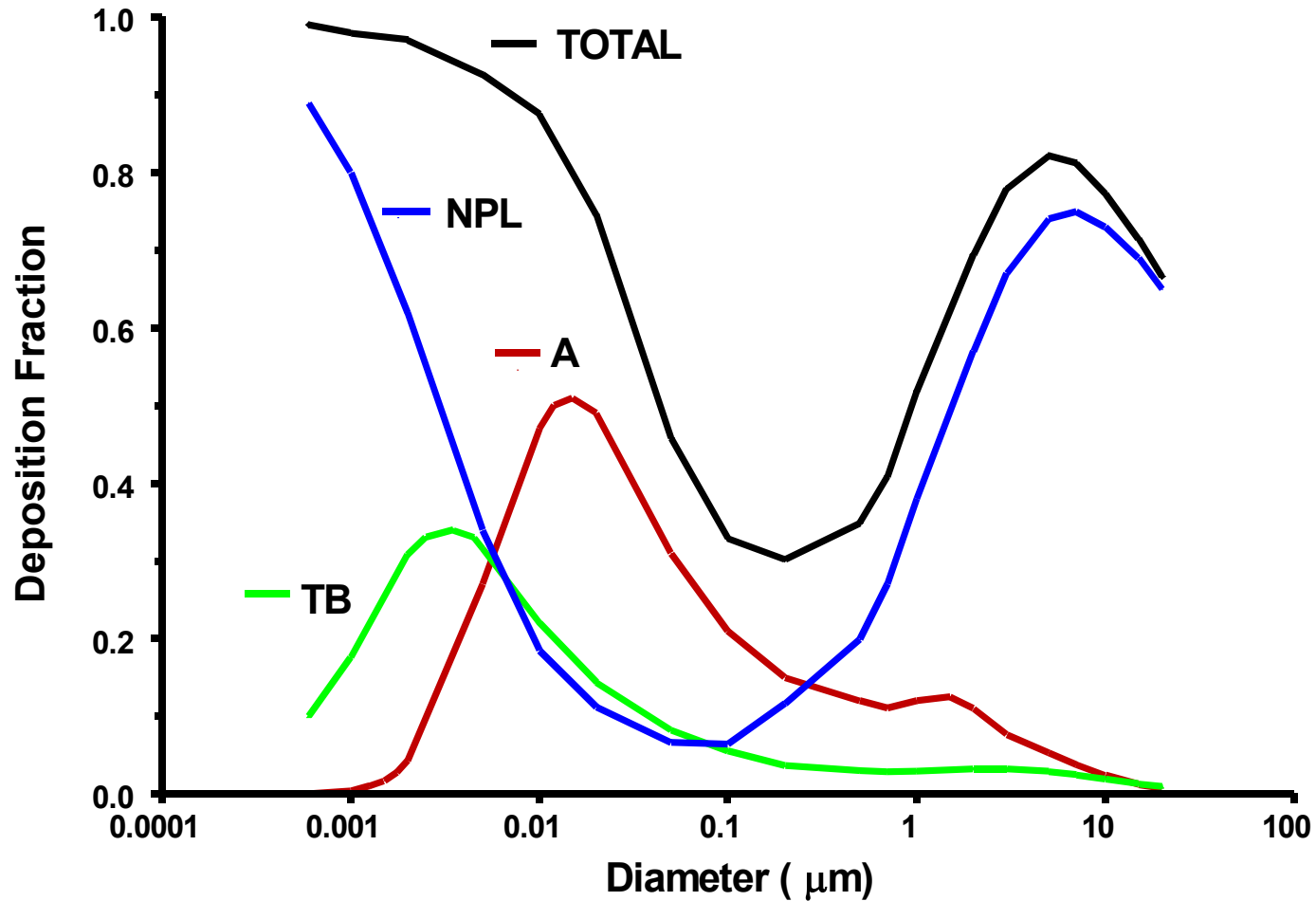
(indoor and outdoor)



Challenges in the Characterization of Airborne MPs Exposure

- Size detection limits of analytical tools used for imaging and identification
 - Raman, FTIR spectroscopies; GC-MS; thermogravimetric analysis; enhanced darkfield hyperspectral imaging (*Nel et al., 2021; Rahman et al., 2021*)
 - Can dyes (or coatings) interfere with identification? (*Xie et al., 2022*)
- Distinguishing polymeric from other sampled particles
 - What is their contribution to overall airborne PM mass and number concentrations?
 - Is this fraction constant across all particle size fractions?
 - Use of dyes (e.g., Nile red) is commonly used (*Nel et al., 2021; Xie et al., 2022*)
- Suite of tools is needed to fully characterize polymeric particles (chemical identity, morphology, size/size distribution)
- Distinction between physical dimensions and aerodynamic diameters*

Fractional Deposition of Inhaled Particles in the Human Respiratory Tract (*nose-breathing*)



Inhalable particulate matter (human)
<100 μm

Respirable particulate matter (human)
<4 μm

PM_{2.5} <2.5 μm

Conceptual Overview of Project

Indoor airborne particulates



Respirable particulates



Respirable microplastics

Morphology?
Concentration?
Composition?

Respirable Particle Sampling Devices

Cyclone type Cutoff



Impactor type Cutoff (2.5 μm)

CYCLONE OPERATION

- ▶ Air enters through a **slit** on the side of the sampler which creates cyclonic action.
- ▶ Large particles fall into the cap (grit pot) at the bottom and are discarded.
- ▶ Small particles are thrown up onto the filter for analysis.



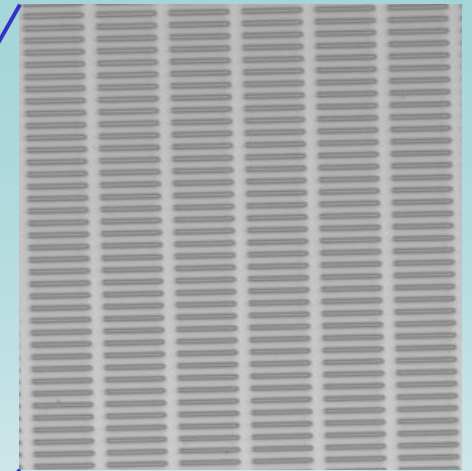
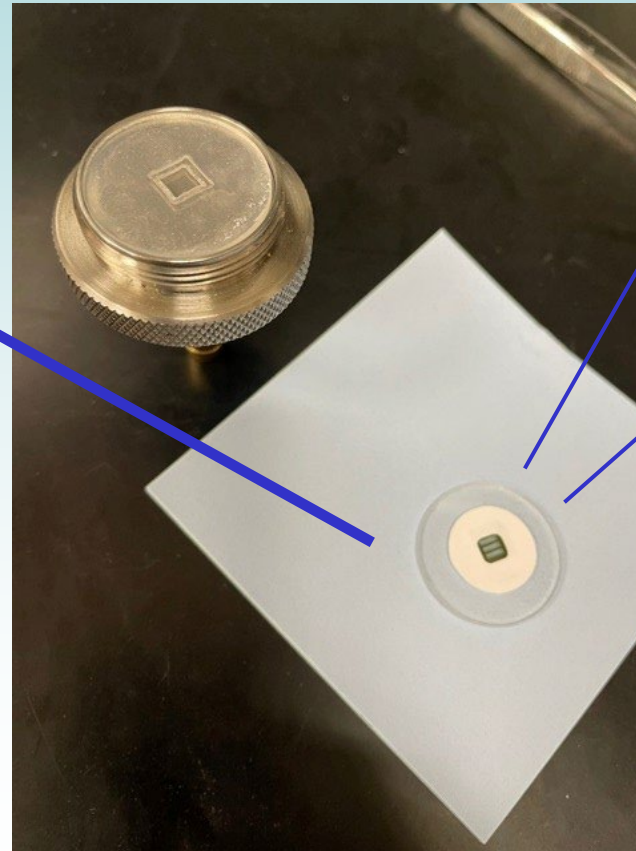
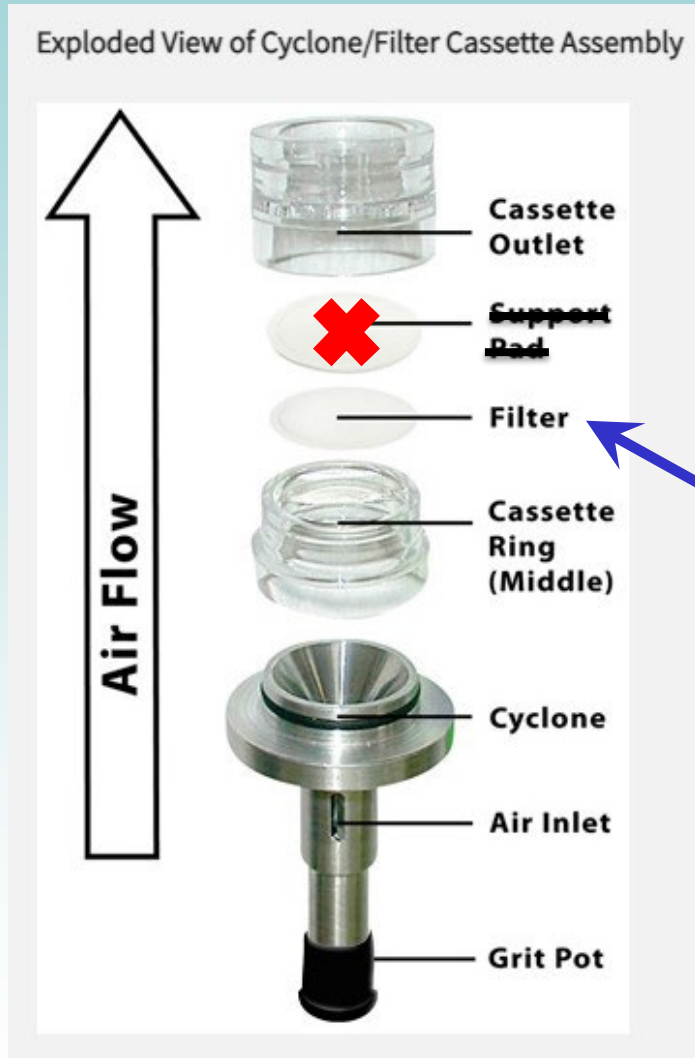
Cap (grit pot) must be in place during sampling.

TABLE 3. Respirable Fraction

Particle Aerodynamic Diameter (μm)	Respirable Particulate Matter (RPM) Fraction Collected (%)
0	100
1	97
2	91
3	74
4	50
5	30
6	17
7	9
8	5
10	1



Respirable Cartridge: SiMPore Substrate Substitution

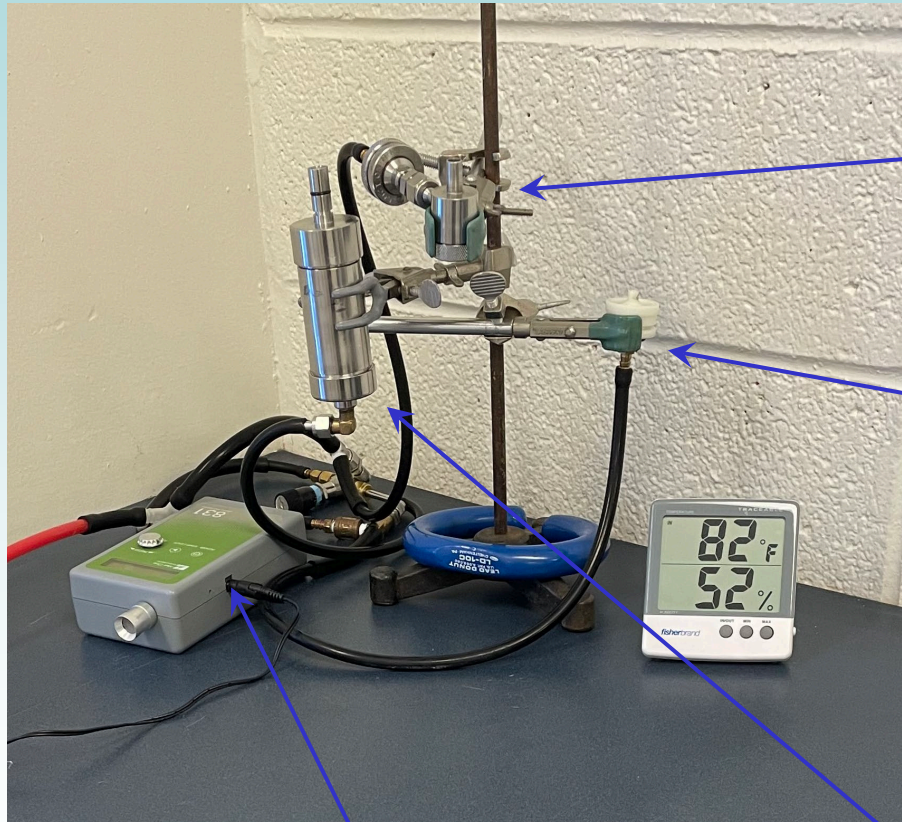


Ultrathin SiN membrane
Each pore = 1 x 50 μm
(8, 4, 2, 1 μm)
>3000 pores across 3
membrane windows,
highly permeable

Imaging and analysis take place
directly on the membrane

- No extraction
- Optically thin

Indoor Air Sampling for Respirable Microplastics



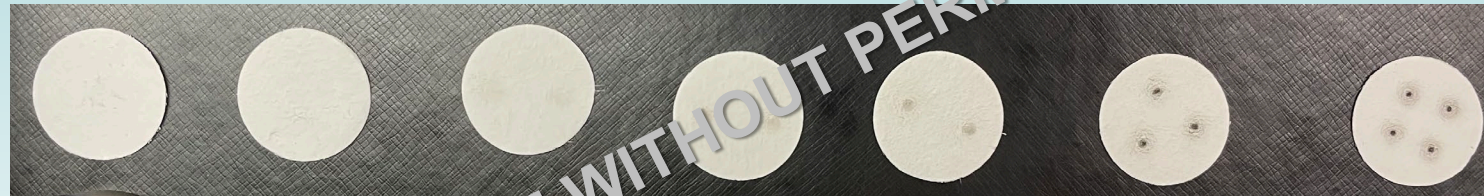
Sampler
(impactor) with
~2.5 μm cutoff

Respirable PM sampling
with SiMPore membrane
(pore size 1 μm x 50 μm)

Optical particle
counter

Cascade impactor with 7
stages (cutoff range from 10
 μm to 0.5 μm)

University laundromat air sampled with the cascade impactor (arranged from largest to smallest cutoff)



Effective
Cut-off
Diameters

10 µm

6.1 µm

3.7 µm

2.2 µm

1.4 µm

0.8 µm

0.5 µm

* 96% of the mass belonged to the particles found in the bottom four stages

MMAD: 0.44 µm

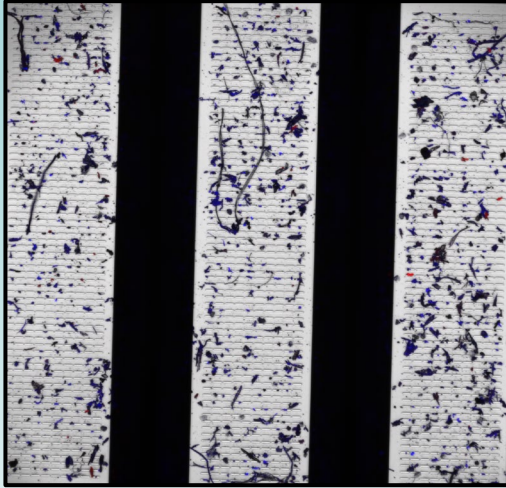
GSD: 4.59

TSP: 9.8 µg/m³

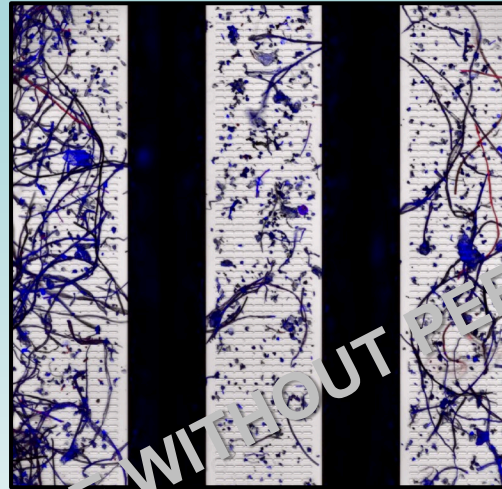
With PM_{2.5} Inlet: 11.6 µg/m³

Air and Surface Dust Samples

Machine Shop
(air)



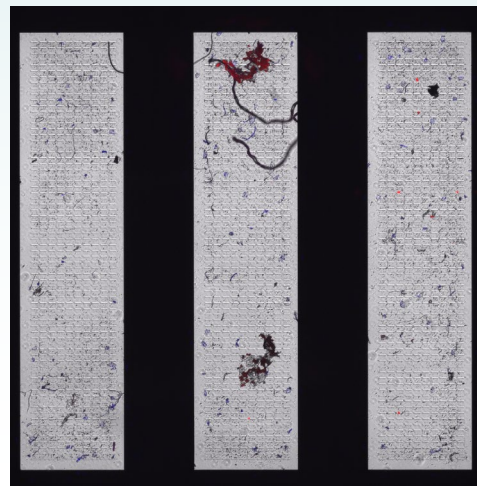
Campus Office
(surface dust)



Household Laundry
Room (open-face)



Household Laundry
Room (2.5 μm impactor)



- + Nile Red
- + Trypan Blue
- Plastic
- Cellulosic
- Proteinaceous

Air

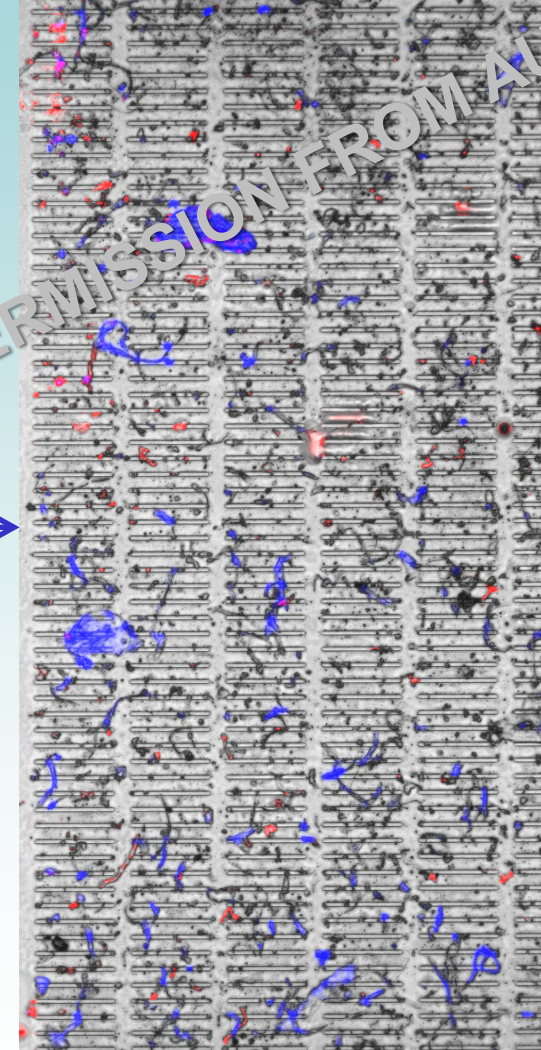
Staining and imaging done by
S.S. Romanick, G. Madejski,
unpublished

4-hr Indoor Air Respirable PM Sample:

SiMPore Membrane, 1 μm pore

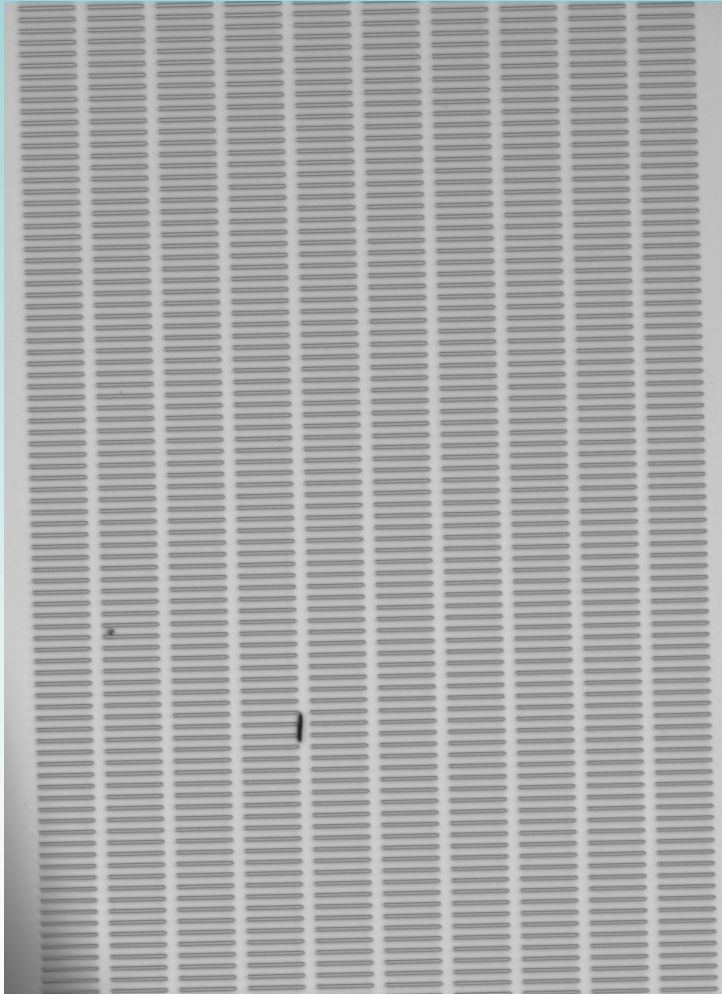


+ Nile Red
+ Trypan Blue

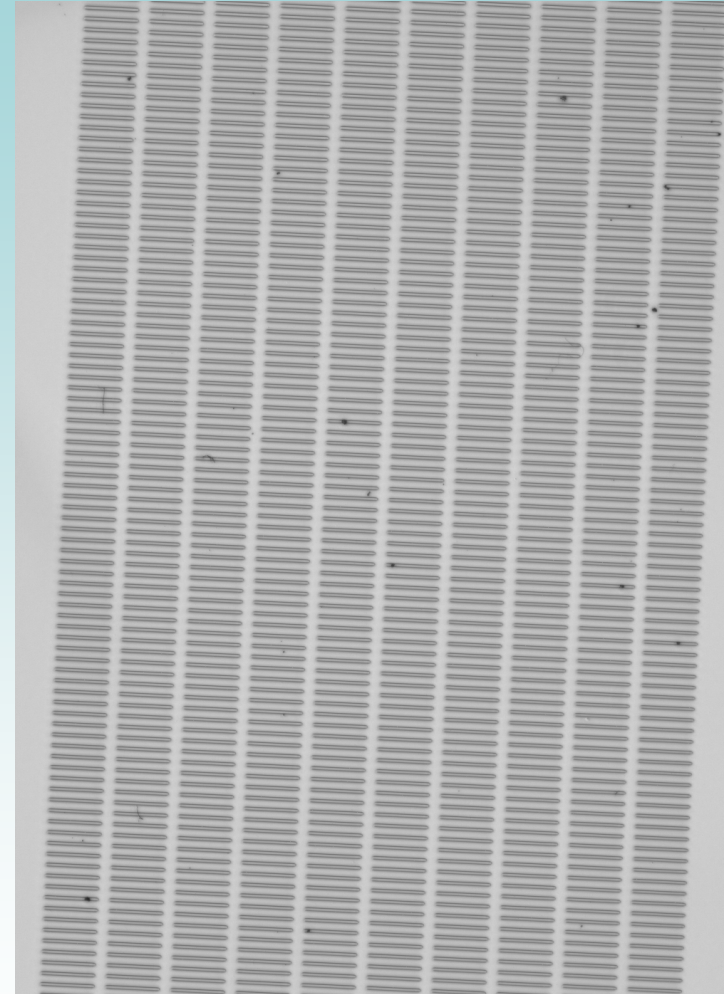


Plastic
Cellulosic
Proteinaceous

Sampling Controls

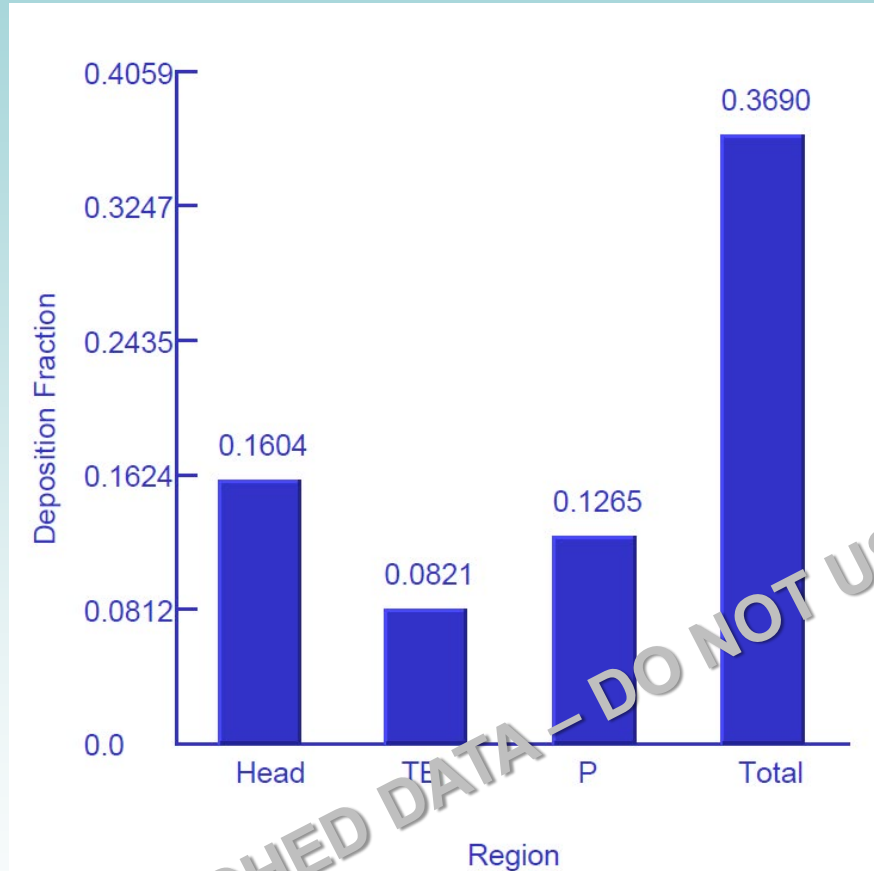


Field Blank
(SiMPore membrane in sampler)



Sample from Clean Room

Estimated Deposited Dose of Respirable Microplastic Particles



~1% of the particles were Nile Red-stained

Estimated deposited dose in the tracheobronchial and pulmonary regions is **0.26 μg** for a modeled 24-hour exposure

$$\text{Deposited Dose} = \underbrace{\text{airborne concentration} \times \text{exposure time}}_{\text{inhaled dose}} \times \underbrace{\text{tidal volume} \times \text{respiratory rate}}_{\text{minute ventilation}} \times \text{deposition fraction}$$

Multiple-Path Particle Deposition Model

Input parameters: human symmetric airway morphometry, oro-nasal normal augmented breathing mode, tidal volume = 625 mL, and breathing rate of 12 min⁻¹, MMAD = 0.44 μm , GSD = 4

Conclusions and Future Directions

1. Microplastics were present in all indoor air samples collected thus far (~1% of sampled PM was Nile red-positive).
 - a. Identification of specific polymers
 - b. Are the polymer types equally distributed across all particle size classes?
2. Sampling revealed the presence of respirable microplastics.
 - a. More work is needed to determine the broad applicability of these findings (is this true for all indoor environments?).
 - b. How do indoor and outdoor environments compare?
3. Using the sampling data from the university laundromat, we estimated a deposited mass dose in the tracheobronchial plus pulmonary regions of ~0.26 μg (for a modeled 24-hour exposure scenario).

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