



# Non-target analysis of microplastics in indoor dust using liquid chromatography high-resolution mass spectrometry

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## INTRODUCTION

- Microplastics have become pervasive in the environment, including dust. These particles pose potential health risks due to their ability to be inhaled and accumulate in humans.
- Although microplastics are commonly analyzed using pyrolysis-gas chromatography-mass spectrometry, liquid chromatography-mass spectrometry [LC-MS] provides sensitive detection of a wide range of analytes, including thermolabile plastic additives and plasticizers.
- Indoor dust samples were collected in residential and industrial settings and analyzed using high-resolution LC-mass spectrometry following depolymerization treatment.
- Non-target analysis with MS/MS library matching using the NIST 2023 library and Kendrick Mass Defect analysis was applied to identify polymer homologous series.

## SAMPLE COLLECTION AND PREPARATION

### Sampling of indoor dust at residential and industrial settings



- Dust samples were collected using a vacuum cleaner from three bedrooms.
- Laundry lint was collected directly from the laundry dryer.



- Dust samples were collected using a Kimwipe from the floors of two laboratory rooms.

### Pretreatment

- All the samples were sieved through 1 mm sieve.
- Hydrogen peroxide was applied to remove organic substances from the surface of microplastics.
- The samples were dried in the oven.

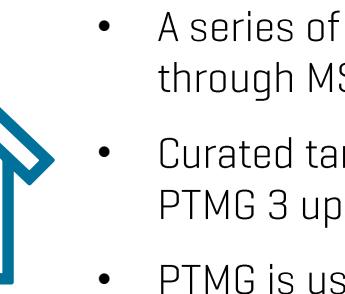
### Depolymerization

- 100 mg of samples were heated in 10 mL of LCMS grade methanol with 200 mg of potassium hydroxide.

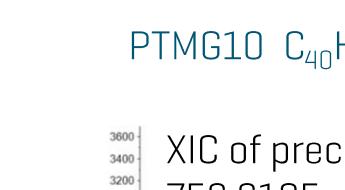
### Filtration and Dilution

- The extracts were filtered with 0.45  $\mu$ m PTFE syringe filters and then diluted 1:100 in acetonitrile.

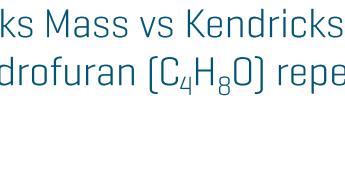
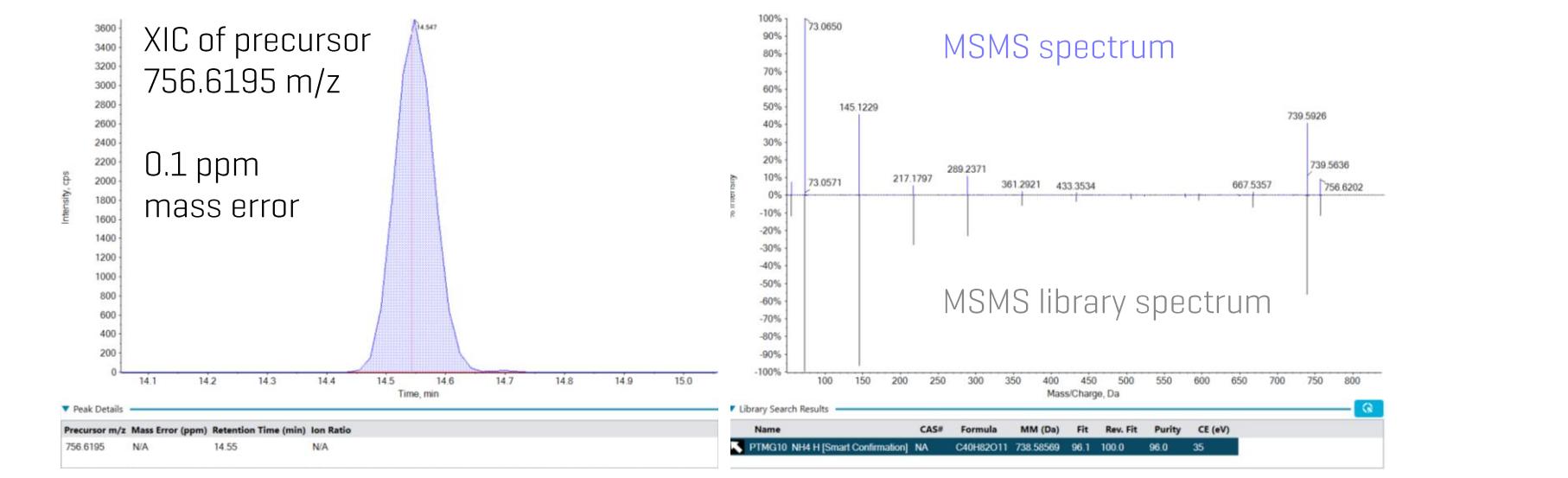
## UNKNOWN SCREENING OF POLYMERS IN DUST SAMPLES



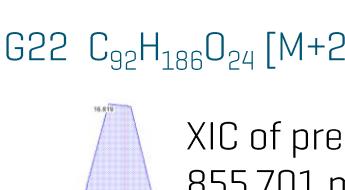
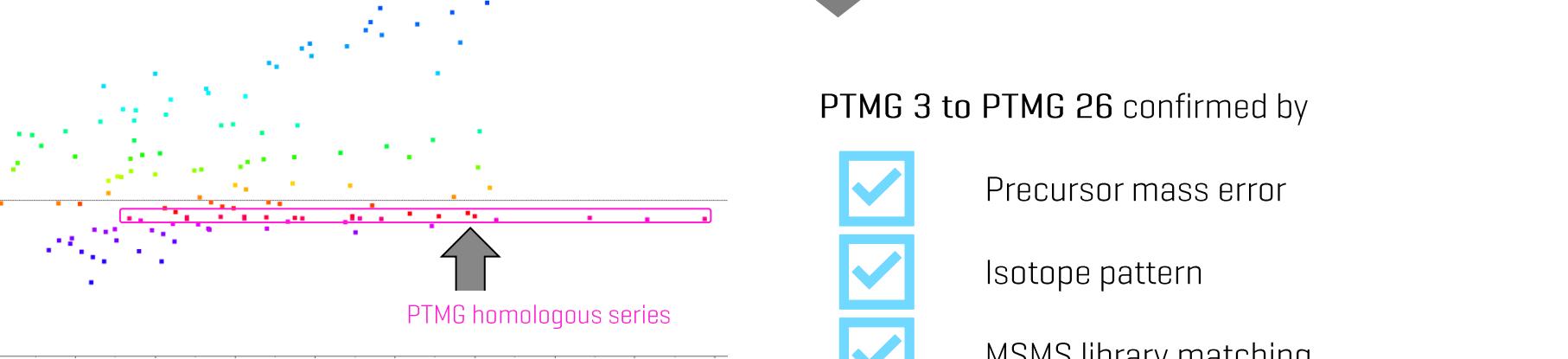
- A series of PTMG [Poly(tetramethylene ether) glycol] was found in residential samples through MSMS library matching.
- Curated targeted suspect screening of PTMGs revealed the detection of polymers from PTMG 3 up to PTMG 26.
- PTMG is used in flexible materials such as spandex and shoe soles.



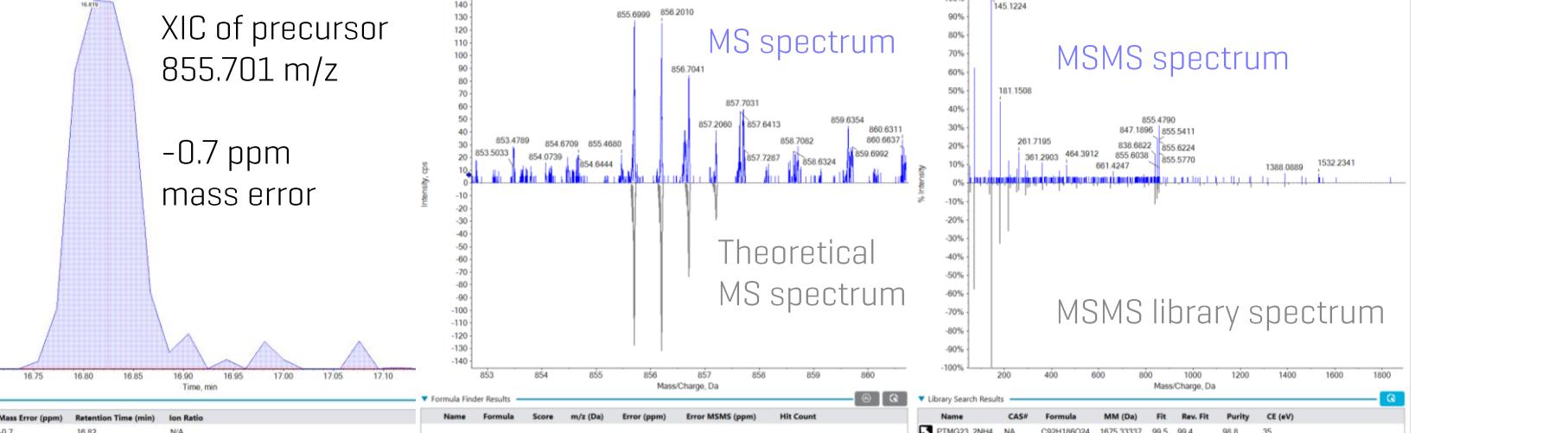
### PTMG10 $C_{40}H_{82}O_{11}$ $[M+NH_3]^+$ in bedroom dust found in non-target analysis



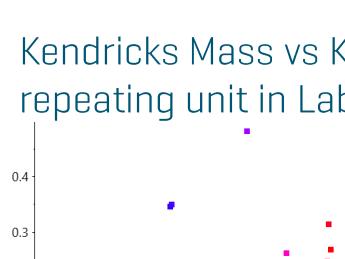
### Kendricks Mass vs Kendricks Mass Defect Plot with Tetrahydrofuran $[C_4H_8O]$ repeating unit in Laundry Lint



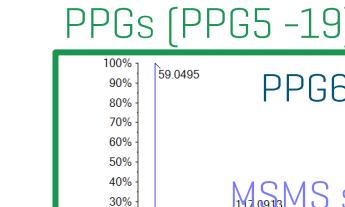
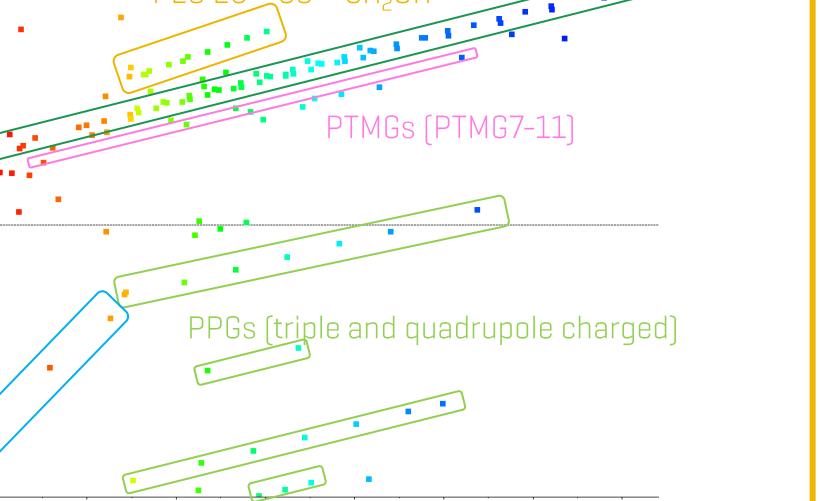
### PTMG22 $C_{92}H_{186}O_{24}$ $[M+2NH_3]^{2+}$ in bedroom dust found in suspect screening



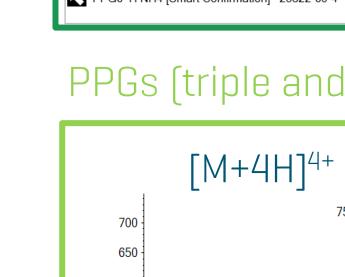
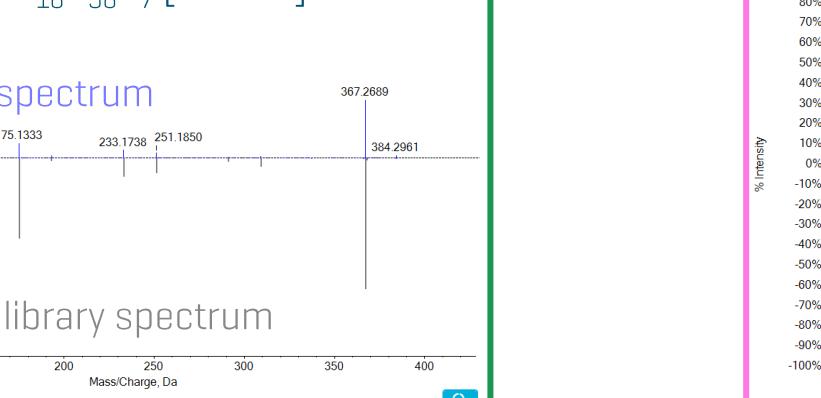
- PEGs, PPGs, and PTMGs dominated the dust sample collected in a non-residential setting.
- A KM vs KMD plot captured an overview of the polymers in the samples.



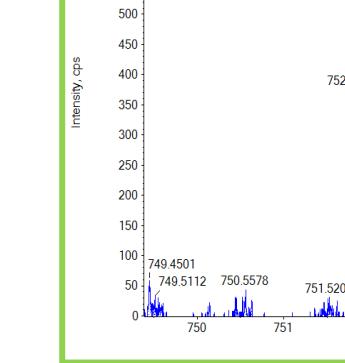
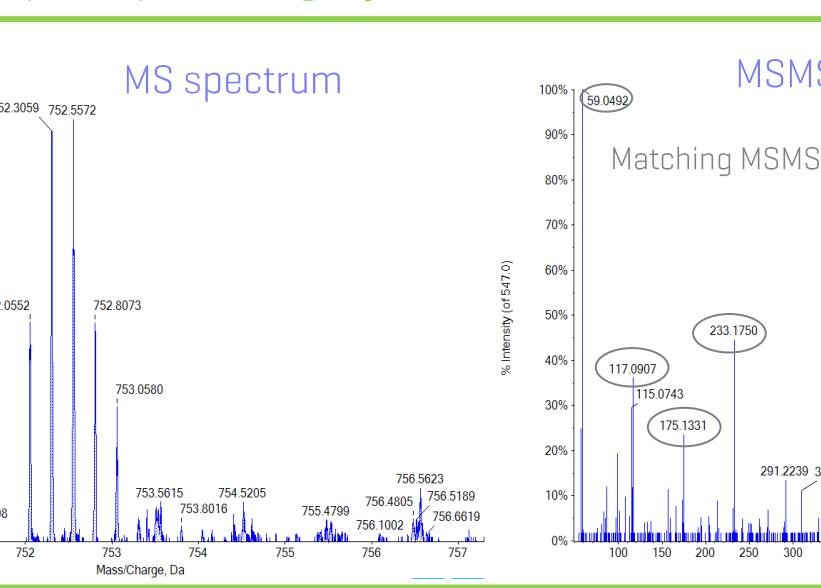
### Kendricks Mass vs Kendricks Mass Defect Plot with $CH_2O$ repeating unit in Laboratory dust



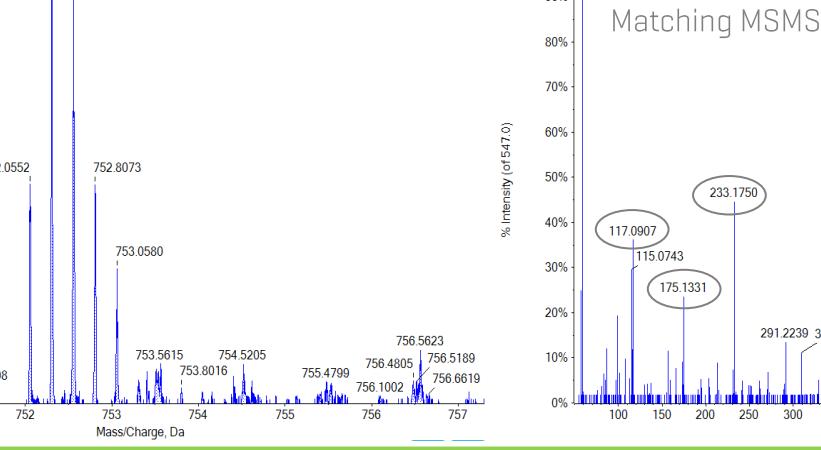
### PPGs [PPG5 -19]



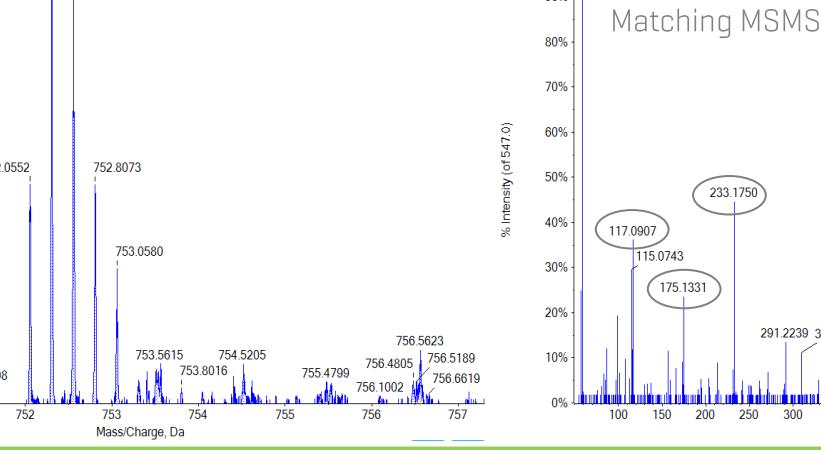
### PPGs [triple and quadrupole charged]



### PTMGs [PTMG7-11]



### MS spectrum



## ANALYSIS DETAILS

**Chromatography:** Chromatographic separation was performed using an Agilent E system with Phenomenex's Luna Omega Polar C18 [2 x 150 mm, 3  $\mu$ m] column, using 0.1% formic acid and 5 mM ammonium formate in ultra-pure water as mobile phase A and 0.1% formic acid and 5 mM ammonium formate in methanol as mobile phase B. The column temperature was maintained at 40°C, and the injection volume was 5  $\mu$ L.

**Mass Spectrometry:** Extracted samples were analyzed using the ZenoTOF7060 system with a Turbo Ion source with Twin Spray ESI probe. Analysis was conducted in data-dependent acquisition mode. The instrument parameters are in the following tables.

### MS source and gas parameters

Ionspray Voltage (IS)	5500 V
Heater Temperature (TEM)	550 °C
Gas 1	50 psi
Gas 2	40 psi
Curtain gas	35
CAD gas	10



TOF parameters	MS	MS/MS
TOF start - stop mass	50 - 3000 m/z	50 - 3000 m/z
TOF accumulation time	0.1 s	0.05 s
Declustering potential	60 psi	60 psi
Collision Energy	10 V	35 ± 15 V

**Data Analysis:** Non-targeted and suspect screening analyses were conducted using SCIEX Version 3.4.5 with NIST 2023 MSMS library. Kendrick Mass and Kendrick Mass Defect were calculated as follows:

Kendricks Mass [KM] = precursor mass  $\times$  [nominal mass of repeating unit /mono isotopic mass of repeating unit]  
Kendricks mass defect [KMD] = Nominal KM - exact KM

### TRADEMARKS/LICENSING

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