BIOAVAILABILITY OF BROMINATED FLAME RETARDANTS IN HOUSEHOLD **DUST**

Sonia García Alcega

INTRODUCTION

Dust

- Very ubiquitous and can be found easily in different range of environments (houses, offices, cars).
- Formed by a complex heterogeneous mixture between semi volatile organic compounds and particle-bound matter derived from biological materials (skin, cells, plant pollen, human and animal hairs, textile fibbers, fungal spores,...)
- Calculations by Harrad et al have suggested that exposure of young children to PBDE via this pathway may be significant.

INTRODUCTION

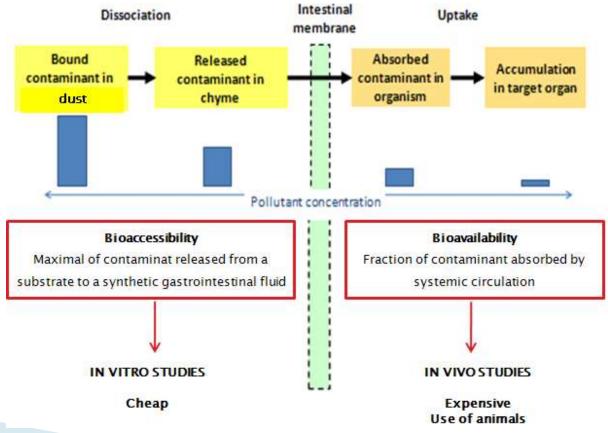
- Available data from researches realized by different countries indicate that BDE-209 is the dominating PBDE congener recovered in dust samples in Europe.
- There is a lack of information about the exact mechanism of exposure to PBDEs, toxicological effects and its implications in humans.

AIM OF THE STUDY

- This project is related to the interaction between brominated flame retardants in home dust and human health.
 - In vitro gastric model
 - → analyze the bioaaccessibility of PBDEs in home dust.
 - Caco-2 cell cultures
 - → predict the absorption, transepithelial transport and cytotoxicity of those chemical compounds.
 - Objective: to obtain more knowledge and potentially answer questions about bioavailability as we have now crossed the membrane.

IN VITRO GASTRIC MODEL

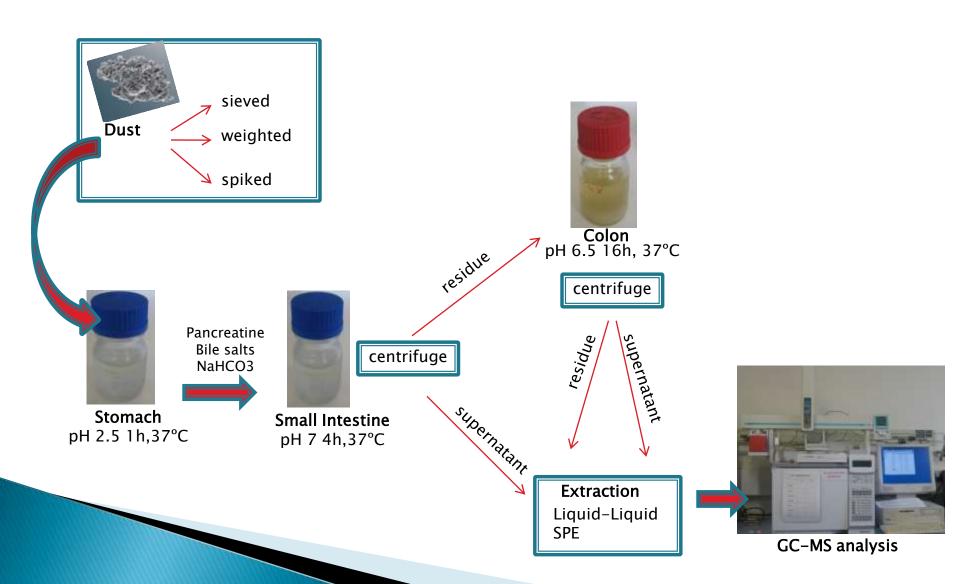
The **bioaccessibility** and **bioavailability** of contaminants in humans can be determined using *in vitro* and *in vivo* analysis.



IN VITRO GASTRIC MODEL

- Gut model consists of a laboratory digestion procedure designed to reproduce gastro intestinal tract chemistry and function.
- It consists on a **three-step** model: **acidic gastric phase** followed by a neutral or slightly **alkaline intestinal phase** with a third **colon phase**.
- According to the literature, measurement of bioaccessibility with in vitro methods offer the potential to provide a reasonable estimation of bioavailability.

IN VITRO GASTRIC MODEL



GC-ECD, GC-MS

• Designing a good method to analyze BDE-209 in my samples

GC-ECD

Column: 30m
Oven program:
100 °C, 1min
200 °C @ 15°C/min, hold 4min
290 °C @ 10°C/min, hold 10min

Transfer line: 280°C

Carrier gas flow: 3-4 ml/min

Splitless injection

GC-MS

Column: 15m Oven program: 100 °C, 2min

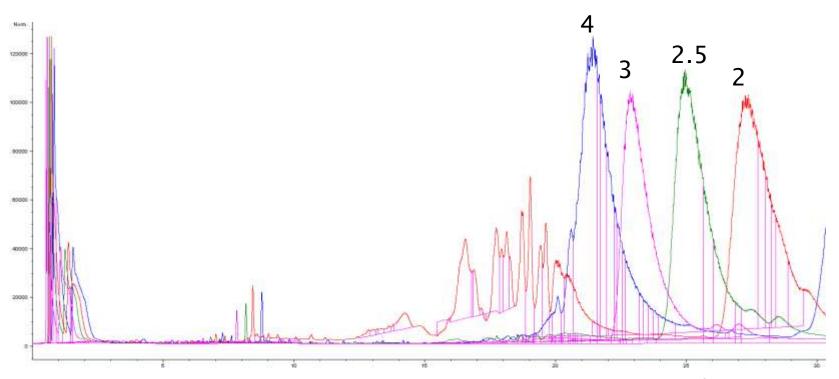
295 °C @ 15°C/min, hold 15min

Transfer line: 280°C

Carrier gas flow: 1.5-2 ml/min

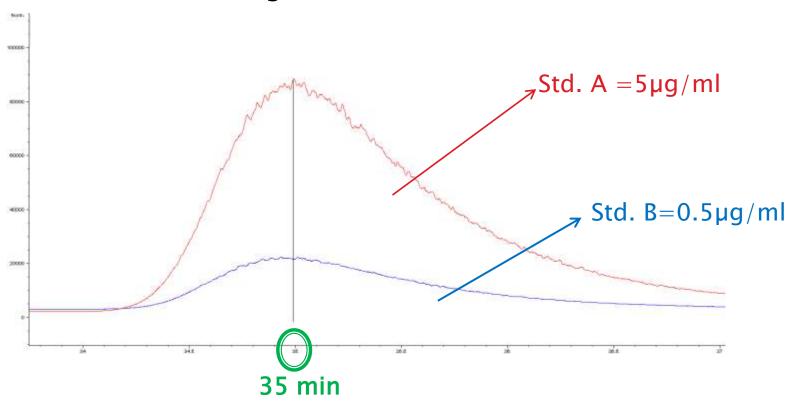
Electron Capture Negative Ionizaton

Mode m/z=496.7 Splitless injection



- BDE-209 appears earlier and with sharper peak using 4 ml/min flow.
- We have tried 5ml/min flow but this was not possible as too much pressure was needed.
- This was using a 30m column which was only one available until recently.

Chromatogram of the BDE-209 standards



We have tried to analyze less concentrated standards (0.05 μ g/ml - 0.0005 μ g/ml but we did not see any peak.

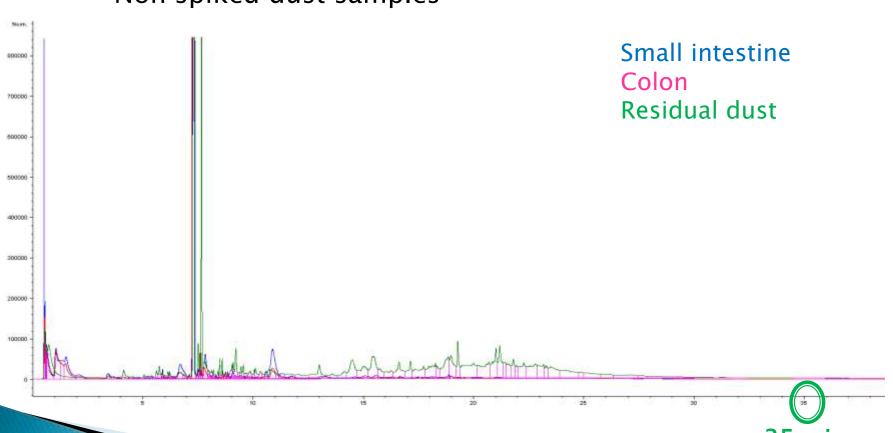
Small intestine, stomach and colon solutions were spiked with 500µl of Std. A (5 µg BDE/ml) and processed as normal samples.

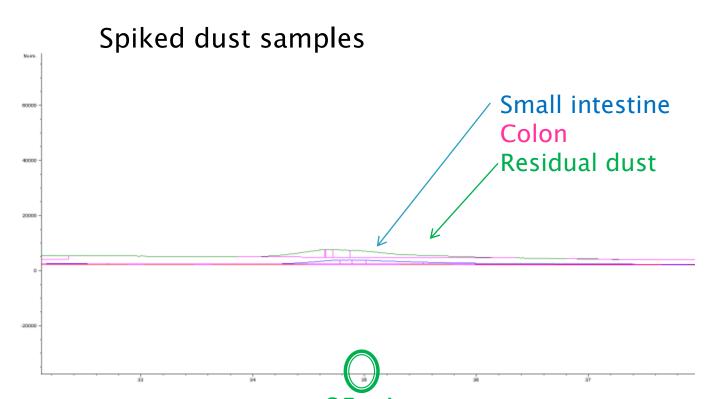
Small intestine Stomach Colon

The concentration of BDE-209 in the small intestine and stomach is equivalent to the Std. A peak.

By contrast, the concentration found in the colon is like the Std. B peak.

Non spiked dust samples





We didn't detect BDE-209 in colon samples.

CONCLUSION

- The protocols for the bioaccessibility determination have been learned and tested successfully using PAH.
- Initial experiments have been undertaken with a house dust but these results were not conclusive.
- We need to develop a reliable method of PBDE determination in the lab.

FUTURE WORK

- Analyze the same dust samples with the GC-MS (15m column) and compare BDE-209 results with the GC-ECD.
- Find other compounds in dust samples (BDE- 28, 47, 99, 100, 153, 154, 183).
- Study and comparison of different dust properties (size, elemental components, optical properties).

