

The background of the slide is a collage of images related to electronic waste. It includes a large pile of discarded mobile phones, a truck dumping a load of e-waste into a container, and various other electronic components and devices scattered around.

Migration Pathways to the environment

"Horizon Scanning" for FRs present in e-waste

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- Analysis of Bisphenol A in urine samples (SPE, LC-QQQ MS), collaboration with a Medical Spanish Research Center in Catalonia (CREAL)
- Development of extraction/clean-up methods for acrylamide/ELISA kits (ABRAXIS LLC, Pennsylvania)
- Development of extraction/clean-up methods for PFASs in food (PERFOOD project) Supervision Dr. Stefan van Leeuwen. Summer 2009, VU University
- Review articles (e.g. Ballesteros Gómez, Rubio Bravo, Soledad, *Recent Advanced in Environmental Analysis*, Analytical Chemistry, 83 (2011) 4579-4613)



e-waste background

- Most polymer electronic materials contain flame retardants, often with halogens
- Globally e-waste growing by about 40 million tons a year
- > 1 billion mobile phones sold worldwide in 2007
- e-waste can end up in landfills
 - >4.6 million tons in US landfills in 2000
 - Hong Kong, 10-20 percent of discarded computers go to landfill
- Regulations:
 - Convention of Basel to control hazardous waste between countries, e.g. to prevent electronic waste being dumped in landfills due to its hazardous content
 - In practice dumping continues in many countries



Objectives ER1

- To study the mechanisms via which FRs migrate from e-waste
 - Develop analytical screening methods for “established” (PBDEs, TBBPA, HBCD), novel (BDBPE, DBDPE, TBB, TBPH, PBT), alternative (PFRs) and unknown FRs (e.g. degradation products)
 - Horizon scanning e-waste
 - Occurrence, leaching (field and lab)
 - Hazard of e-waste
 - Toxicity profiling (e.g. microsomes)

ANALYTICAL METHODS DEVELOPMENT FOR SCANNING FRs

EXTRACTION

Simplifying sample
treatment: dispersive
SPE, QUECHERS,
surfactant-based
extraction

REPORTED EXTRACTION/ CLEANUP METHODS

- Different extraction methods
 - Soxhlet
 - ASE
 - Ultrasonic extraction
 - SPE
- Wide range of solvent mixtures
 - Petroleum ether
 - Toluene
 - Dichloromethane
 - Hexane
 - Acetone
- Cleanup methods for abiotic and biotic samples
 - Sulphuric washing
 - Deactivated or sulphuric acid impregnated silica column
 - Florisil column
 - SPE cartridges
 - Alumina column
 - Sulphur removal (activated copper, AgNO₃ on silica, TBA reagents and GPC)

- Developing comprehensive separation/detection methods for scanning FRs

- GCXGC (Agilent)-HRTOF(Brukers)

Why?

GCxGC has not been coupled before to HRTOF to combine enhanced separation and high mass accuracy for identification

How?

High scan rate needed (20-30Hz) due to narrow peaks coming from GCxGC thus leading to a decrease in sensitivity, so optimization of TOF parameters affecting sensitivity will be carried in depth

Compatible sources commercially available will be checked

APCI [some recent articles propose it as suitable for BRFs, Zhou et al. Anal. Bioanal. Chem 396 (2010); Mascolo et al. J. Chromatogr.A 1217 (2010) 4601]

APLI (possible collaboration)

APPI (coupling development needed, only commercially available for direct probe injection)

- **Developing comprehensive separation/detection limits for scanning FRs: Other options**

- GC-QTOF
- LC (Agilent)-ESI/APCI /APPI-HRTOF(Bruckers)
- LC-QTOF (University of Antwerp)
- APPI , APCI direct probe-HRTOF
- LC-QTRAP for full scan/MRM/MSⁿ combined experiments (University of Cordoba)



THANK YOU FOR
YOUR ATTENTION!