

Spatial and temporal variations of the size distribution of the atmospheric particulate persistent organic pollutants (POPs) and Persistent Bioaccumulative Substances (PBTs) in the Czech Republic

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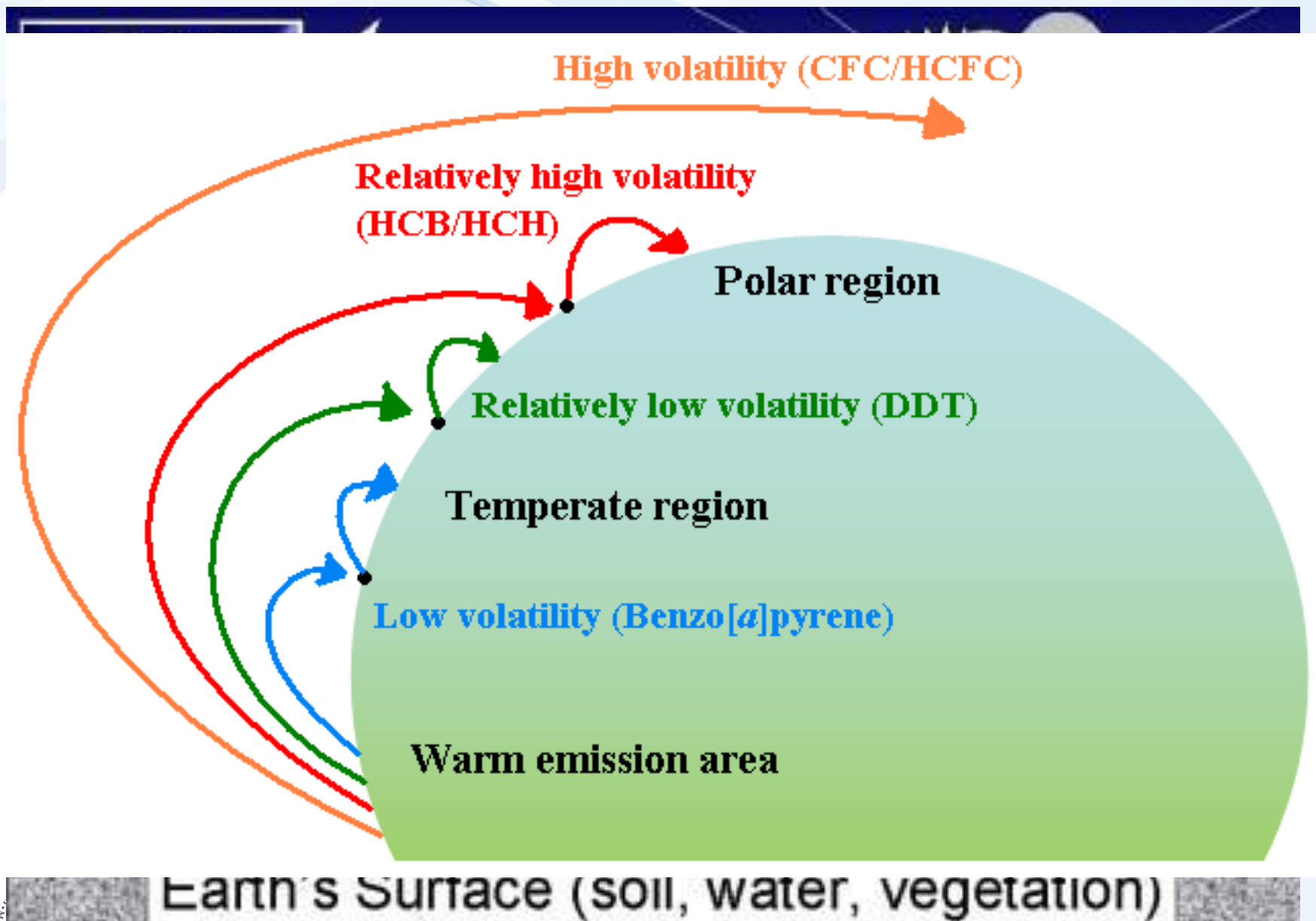
**Research Centre for Toxic Compounds in the Environment (RECETOX),
Brno, Czech Republic**

8th Network Conference in POPs, Birmingham, UK
8th May 2014

Outline

- Introduction
- Methodology
 - Sampling sites
 - Air sampler (cascade impactor)
 - Extraction and Clean-up
 - Separation and detection
- Results
 - PM₁₀ profiles
 - Combustion-related compounds (PAHs, PCDD/Fs, dl-PCBs)
 - Flame retardants (PBDEs, nBFRs and HBCDs)
 - Pesticides (OCPs and CUPs)
- Conclusion

Introduction

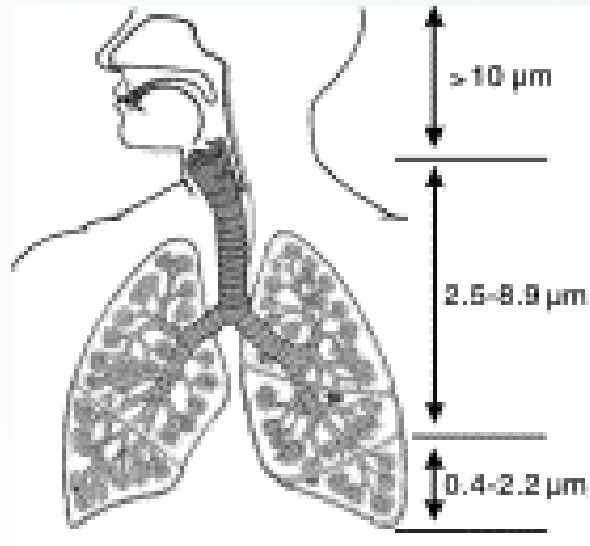


Introduction

Size of particles

Size of particles will determine:

- Residence time in the atmosphere
- Potential for LRAT
- Deposition efficiency
- Human health exposure from inhalation



Methodology

Sampling sites

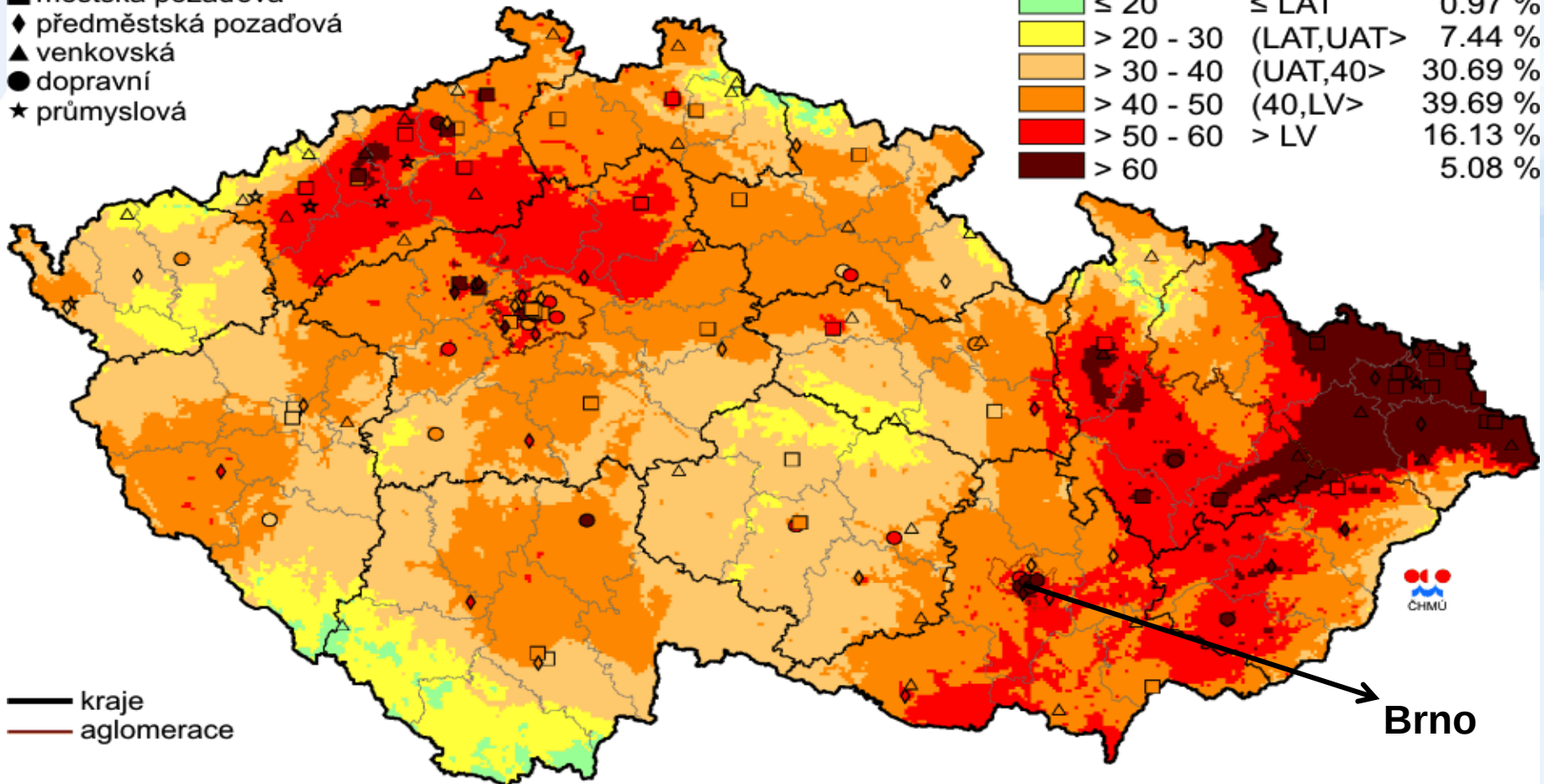
Data from the Czech Hydrometeorological Institute, 2010 www.chmi.cz

klasifikace stanic

- městská pozadová
- ◆ předměstská pozadová
- ▲ venkovská
- dopravní
- ★ průmyslová

koncentrace [$\mu\text{g}\cdot\text{m}^{-3}$]

≤ 20	≤ LAT	0.97 %
> 20 - 30	(LAT,UAT>	7.44 %
> 30 - 40	(UAT,40>	30.69 %
> 40 - 50	(40,LV>	39.69 %
> 50 - 60	> LV	16.13 %
> 60		5.08 %



Pole 36. nejvyšší 24hod. koncentrace PM_{10} v roce 2010

Methodology

Sampling sites

Telnice (15km from Brno)

Rural site



Kotlařska (Brno city centre)

Kerbside/Urban site



Methodology

Air sampling

High volume air sampler equipped with a cascade impactor



Fractions collected on QFFs:

- < 0.49 μm
- 0.49-0.95 μm
- 0.95-1.5 μm
- 1.5-3.0 μm
- 3.0-7.2 μm
- 7.2-10.0 μm

1 year sampling (Oct 2009-Oct 2010)

Seasonal data
corresponding of 3
weeks of sampling

Methodology

Extraction



Methodology

Clean-up



Methodology

GC-MS/LC-MS



- PAHs: GC-MS Agilent 6890/Agilent 5972
- PCDD/Fs, dl-PCBs, PBDEs and nBFRs: GC-HRMS Agilent 7890/Micromass AutoSpec Premier (sector MS)
- CUPs and HBCDs: LC-MS Agilent 1290-ABSCIEX5500

RESULTS

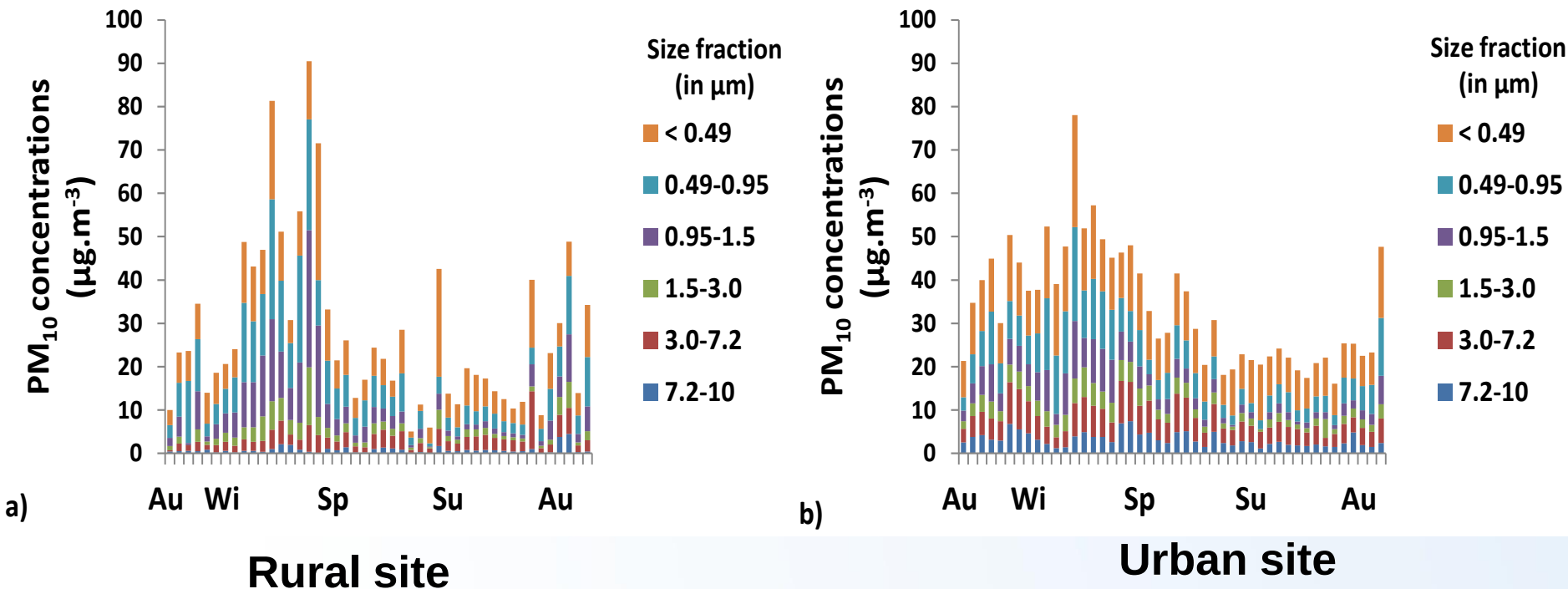


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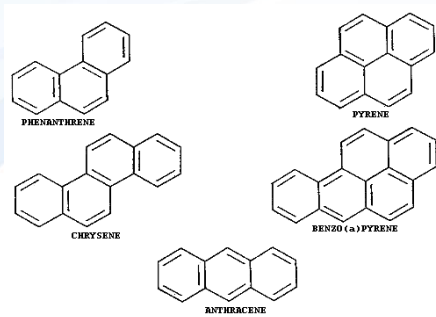
PM concentrations



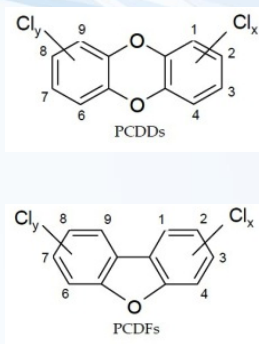
- Higher PM₁₀ concentrations at the urban site
- At both sites, higher concentrations were observed in winter (about 50 µg.m⁻³) than in summer (20 µg.m⁻³).
- **54-59%** of particles have a diameter < 0.95 µm

Combustion-related compounds

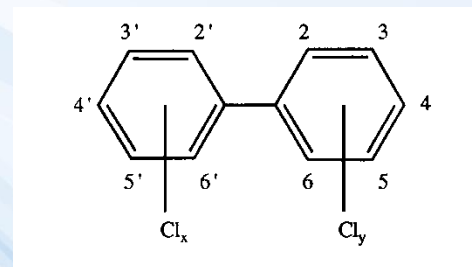
PAHs



PCDD/Fs



dl-PCBs



- **Sources:**

- Natural sources (forest fires, volcanic eruptions, products of biosynthesis of phytoplankton and bacteria)
- Anthropogenic sources (fossil fuel combustion, residential heating, waste incineration, traffic, oil refining, steel and iron manufacturing, coke and asphalt production and metal reclamation processes)

- **Long-half-life in the air, soil, sediment or biota**

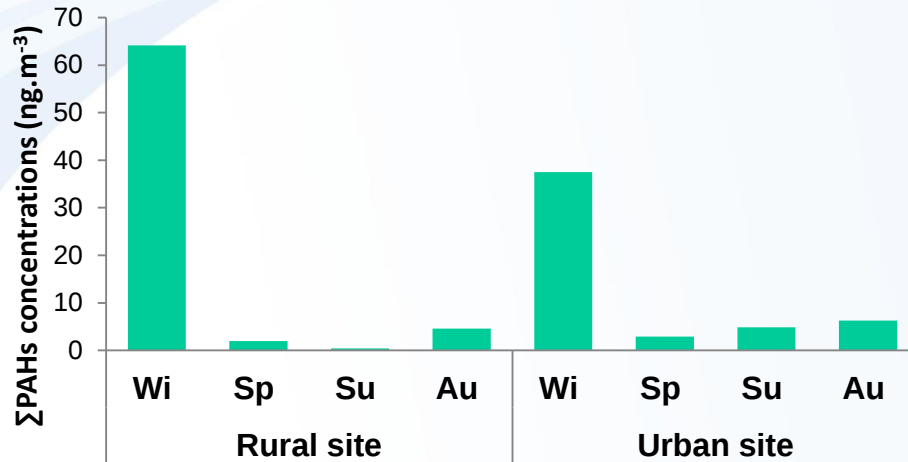
- **Health effects:**

- Carcinogenic and/or mutagenic properties for PAHs and high toxicity for PCDD/Fs and dl-PCBs

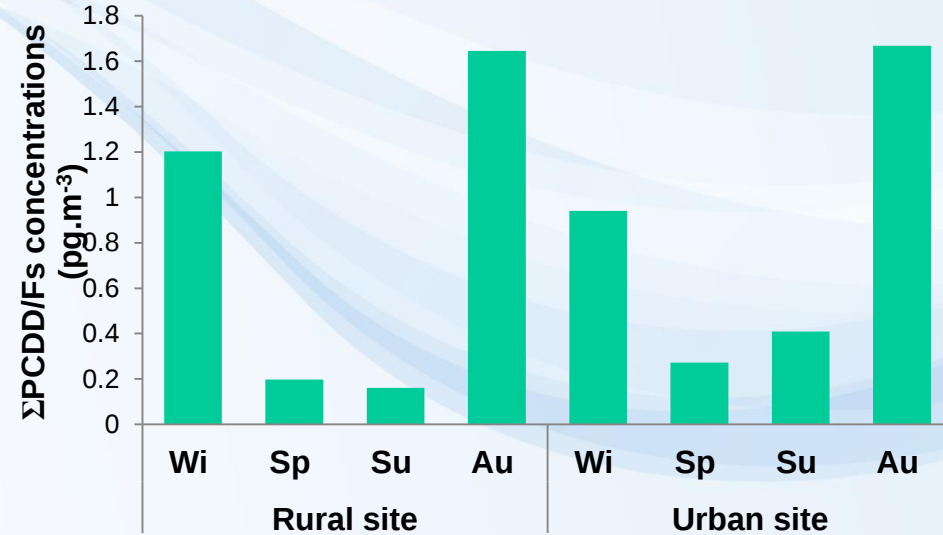
PAHs, PCDD/Fs and dl-PCBs

Total particulate concentrations

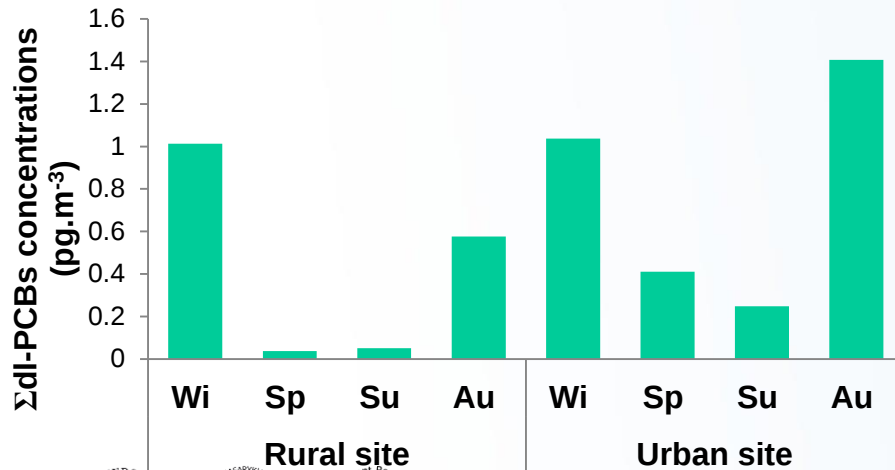
PAHs



PCDD/Fs



dl-PCBs



- Higher concentrations in cold seasons for all of the compounds at both sites
- Higher concentrations at the rural site for PAHs and PCDD/Fs in winter, which might reflect the spatial difference in domestic heating

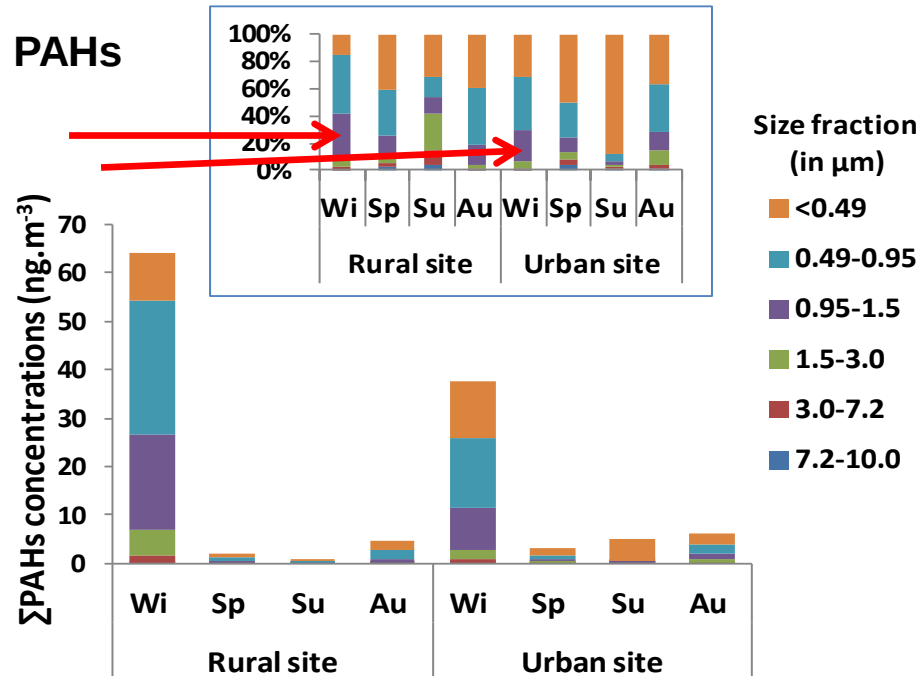
PAHs, PCDD/Fs and dl-PCBs

71% of PAHs < 0.95 μm

Size distribution

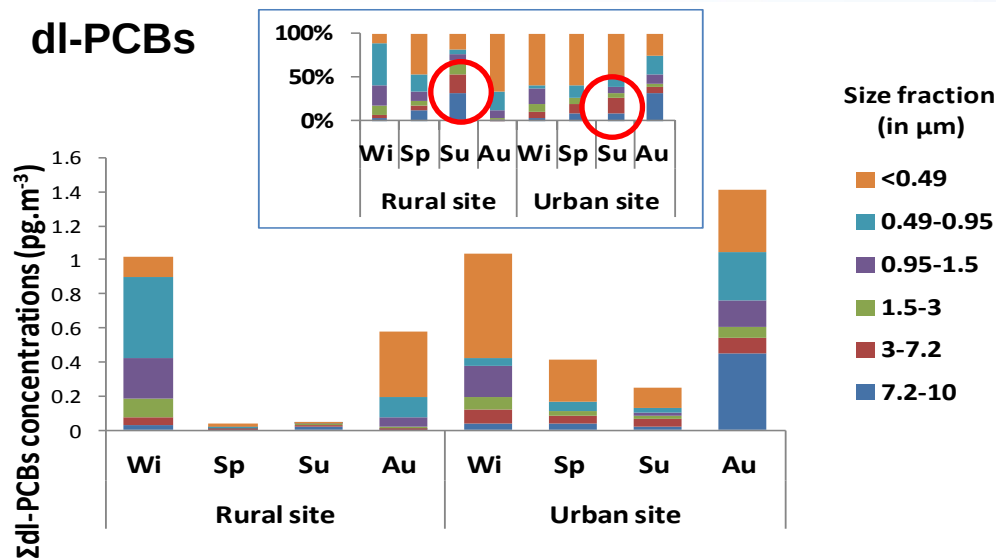
73% of PCDD/Fs < 0.95 μm

PAHs

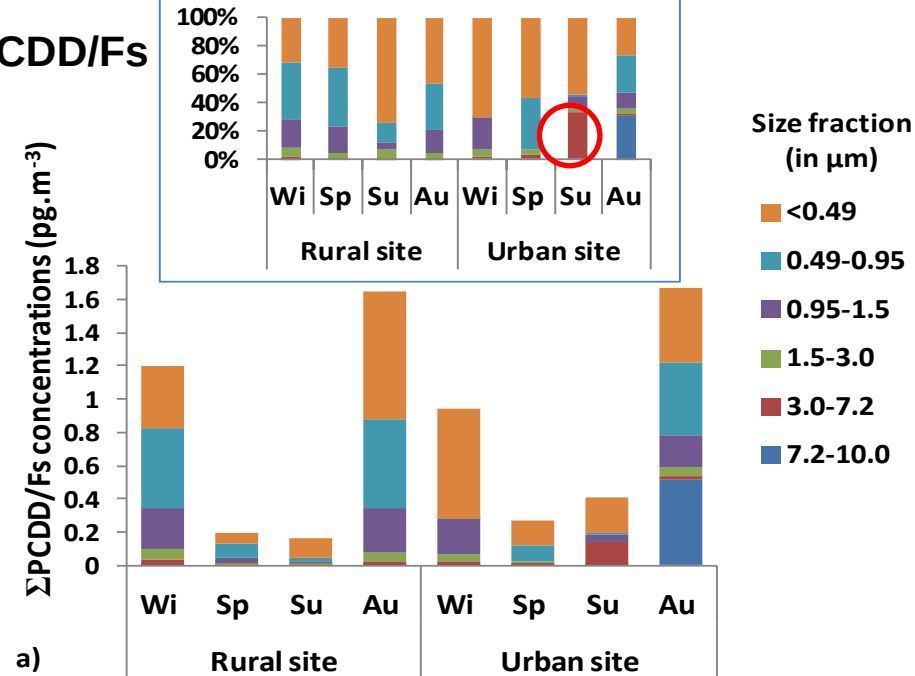


60% of dl-PCBs < 0.95 μm

dl-PCBs

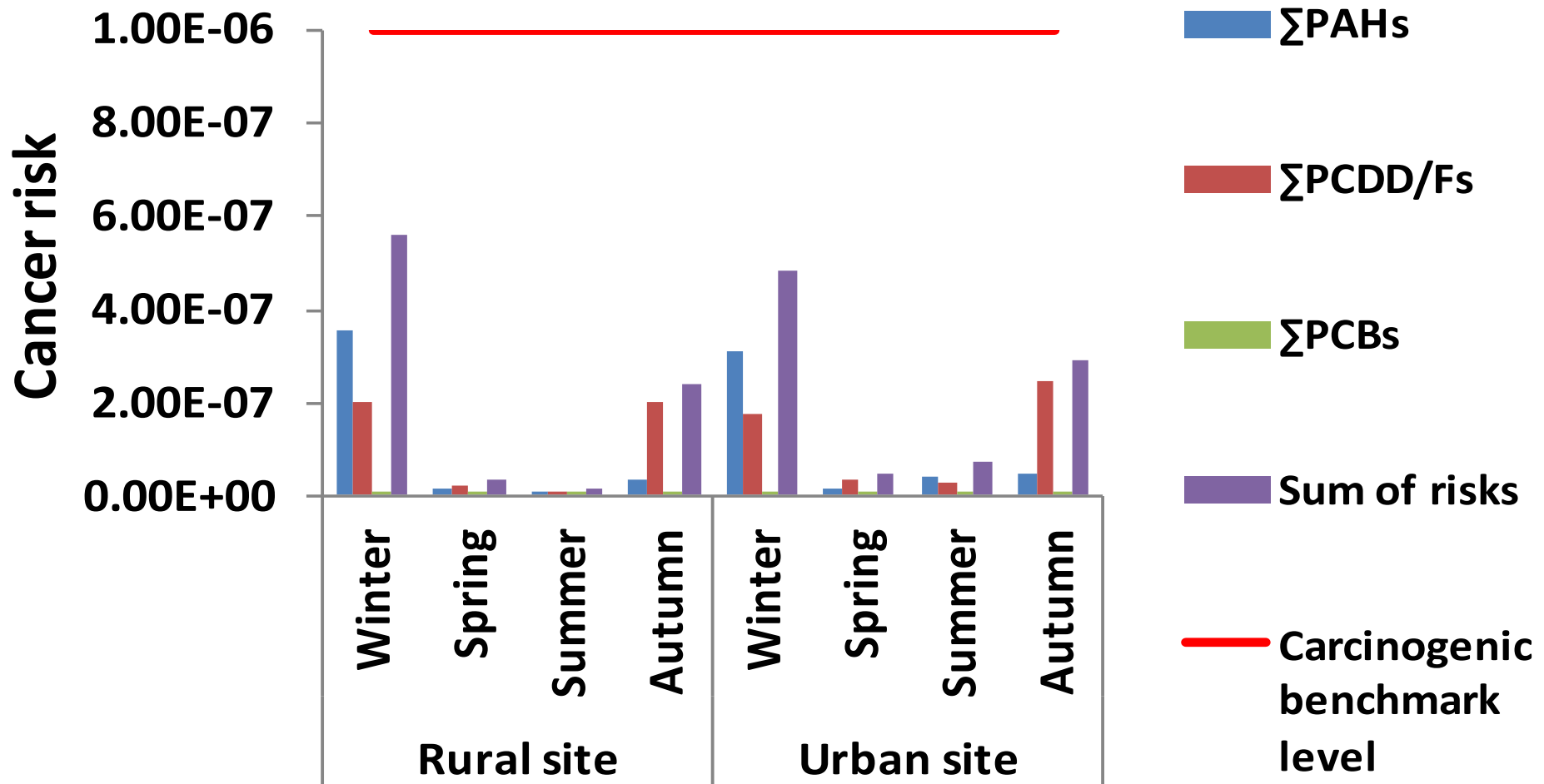


PCDD/Fs



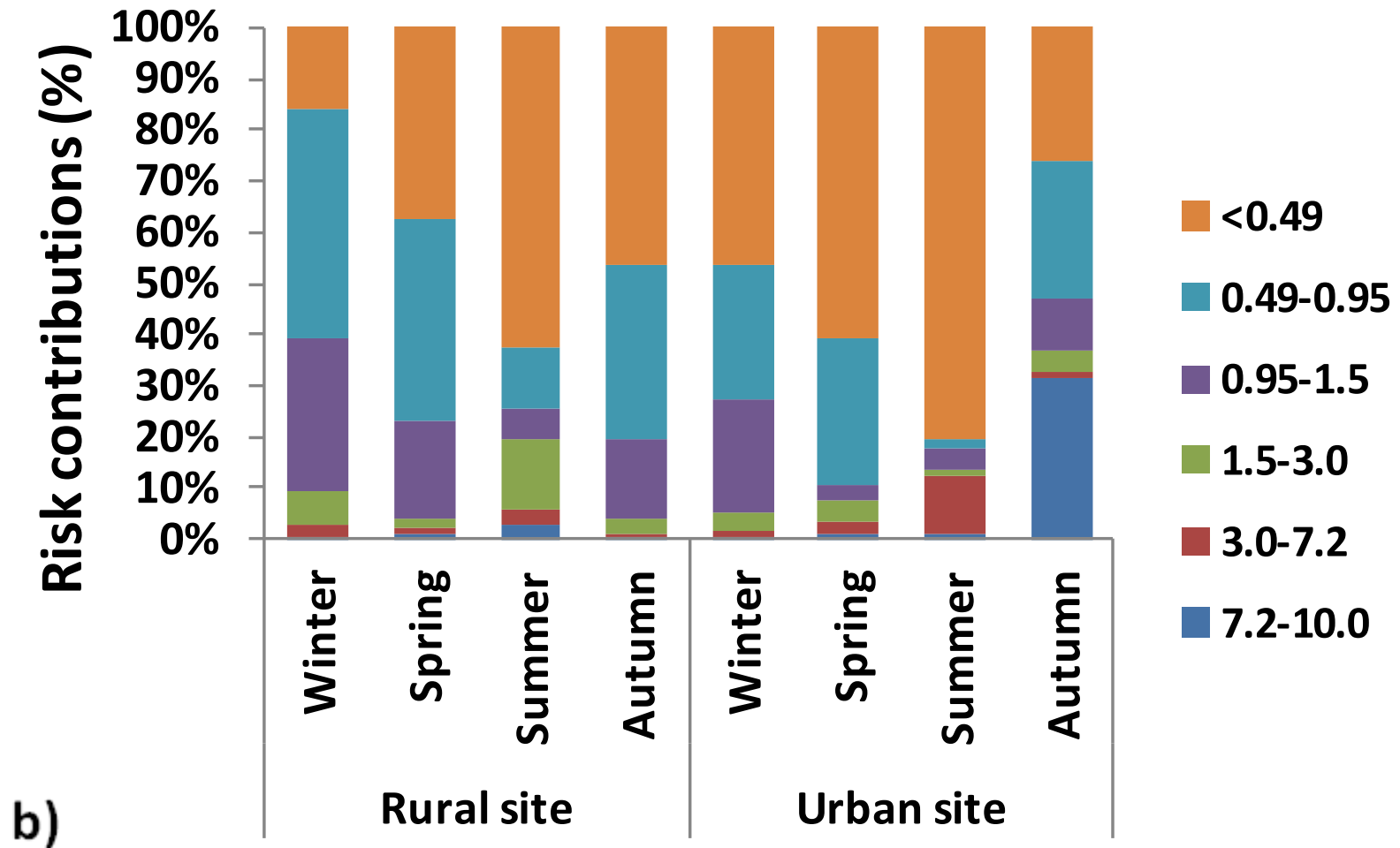
- Strong accumulation in the fine fraction for all compounds regardless the site or the season
- Higher contribution of the 0.95-1.5 μm fraction for PAHs in winter
- Increase of coarse particles in summer for PCDD/Fs and dl-PCBs

Cancer risks from inhalation exposure



- Cancer risks were lower than the carcinogenic benchmark level
- Higher risks in winter

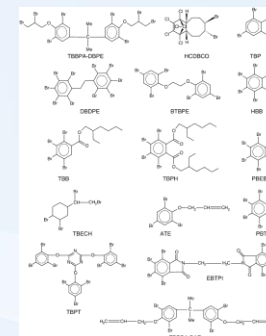
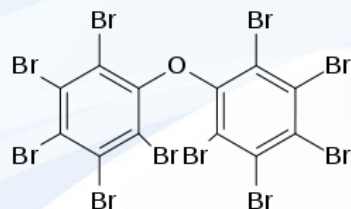
Cancer risks from inhalation exposure



- Most of the risks are due to fine particles
- However, the cancer risks can be overestimated by up to a factor of 2 in situations where there is a large amount of coarse particles.

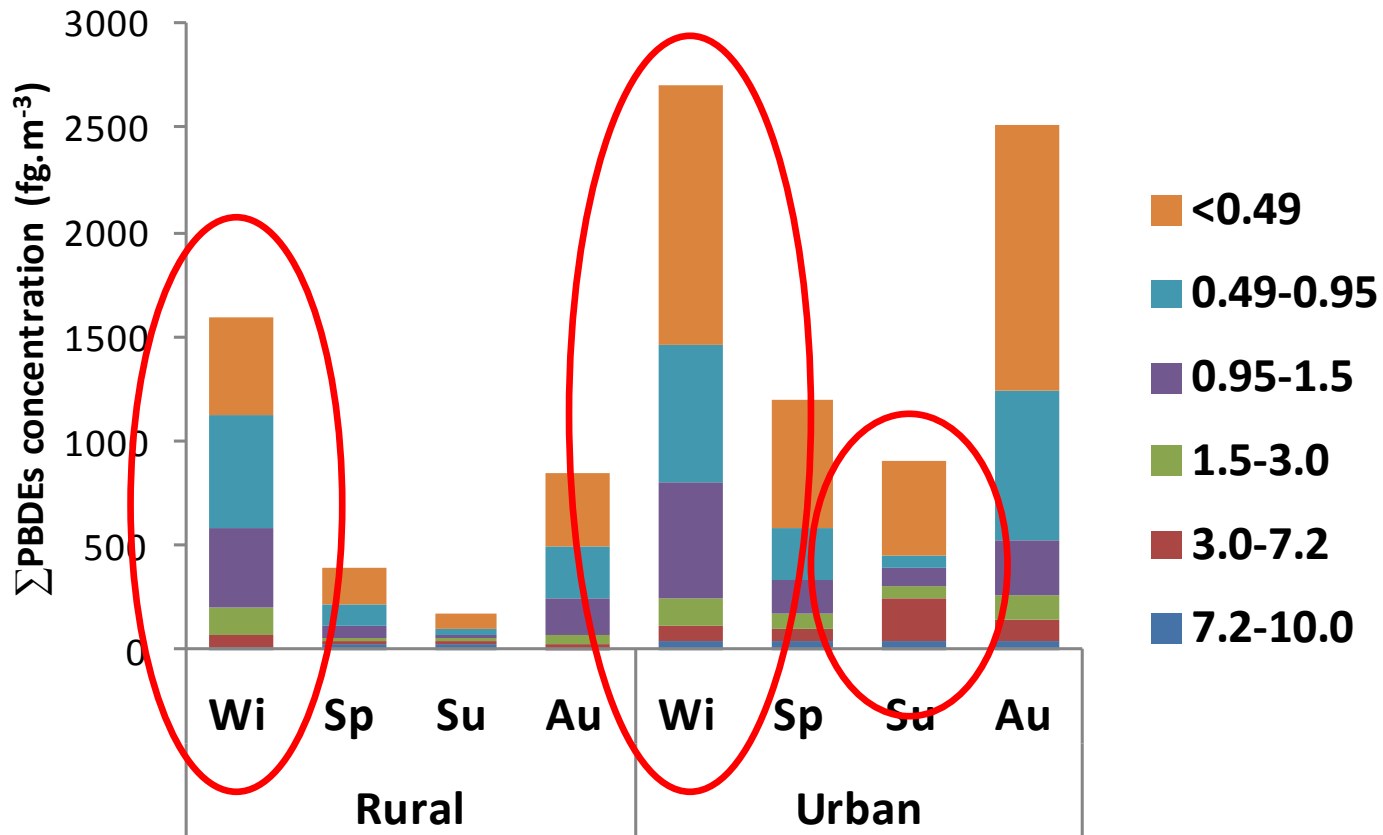
PBDEs, nBFRs and HBCDs

nBFRs



- Flame retardant additives used in electronics (computers, mobile phones, kitchen appliances), textiles, building materials, plastics, paints.
- Due to the restriction of some of PBDEs, nBFRs use has increased consistently for the last decade.
- Sources in the air: Release during production, volatilization during the use of products and inappropriate disposal.
- Bio-accumulative, persistent and toxic

PBDEs (without BDE-209)



- Higher concentration at the urban site
- 4x higher winter concentrations compared to summer concentrations
- 55% of PBDEs were associated to fine particles
- Increase of coarse fraction in summer due to volatilization and condensation onto coarse particles

nBFRs

10 target compounds

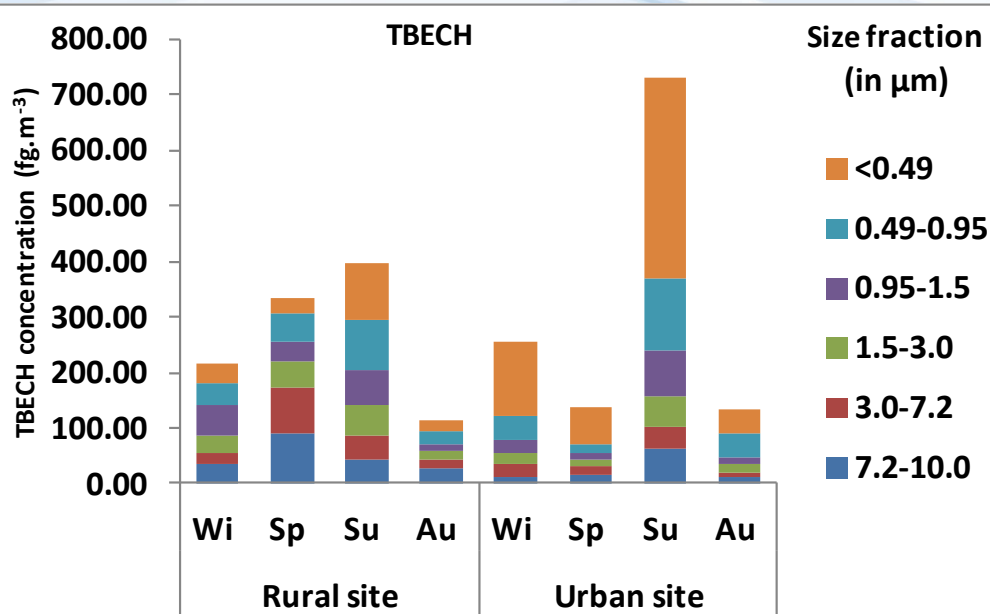
3 were not detected at all (TBCO, p-TBX and HCDBCO)

In general, higher summer concentrations due to indoor emissions

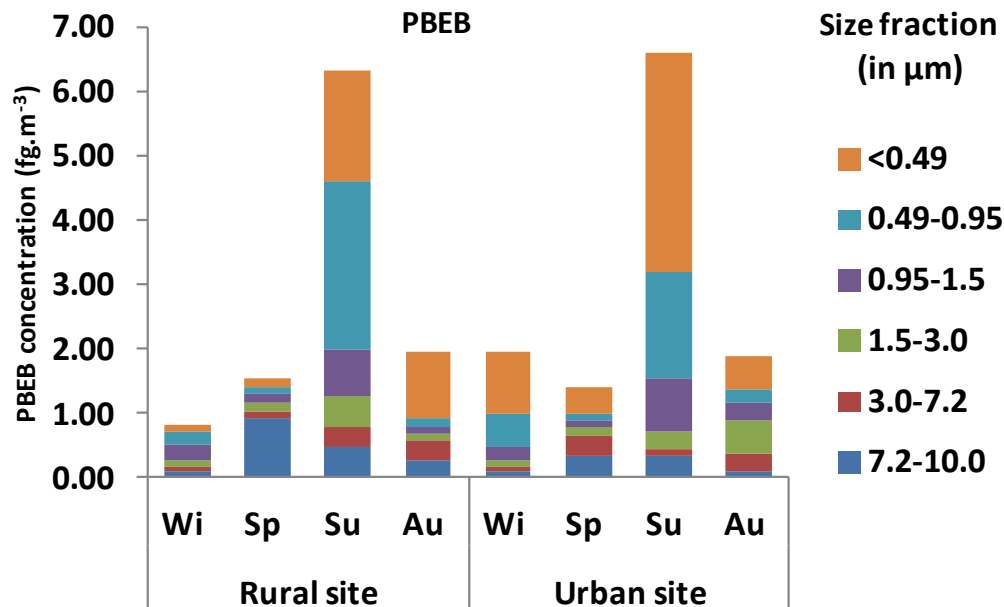
Higher concentrations at the urban site

In some cases, higher n-BFRs concentrations than PBDEs reflecting the growing use of alternative flame retardants

n-BFRs size distribution

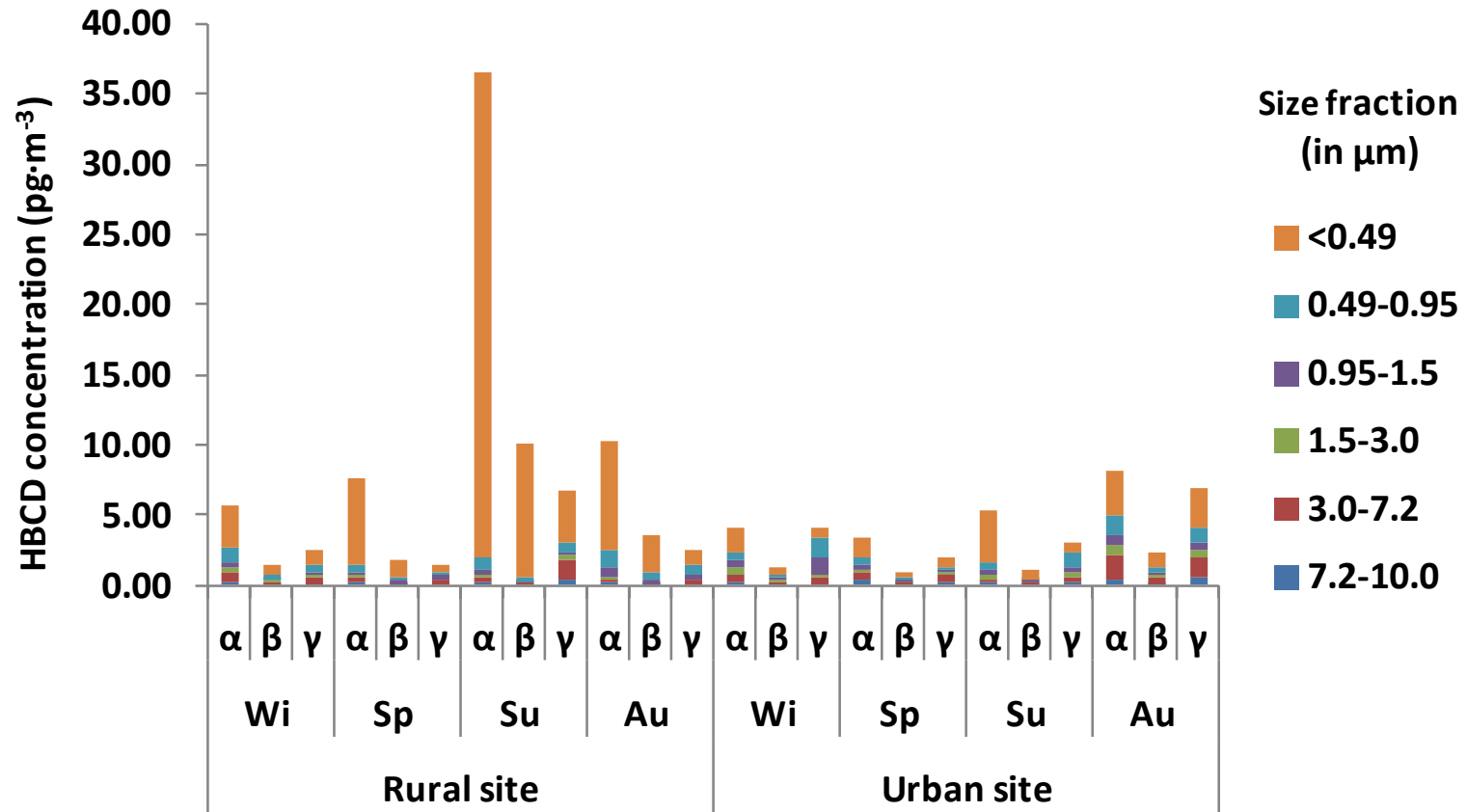


4/7 n-BFRs (TBECH, PBT, HBB and BTBPE) showed strong affinity toward fine particles



3/7 n-BFRs (s-DP, a-DP and PBEB) showed similar concentrations across size fractions

HBCDs

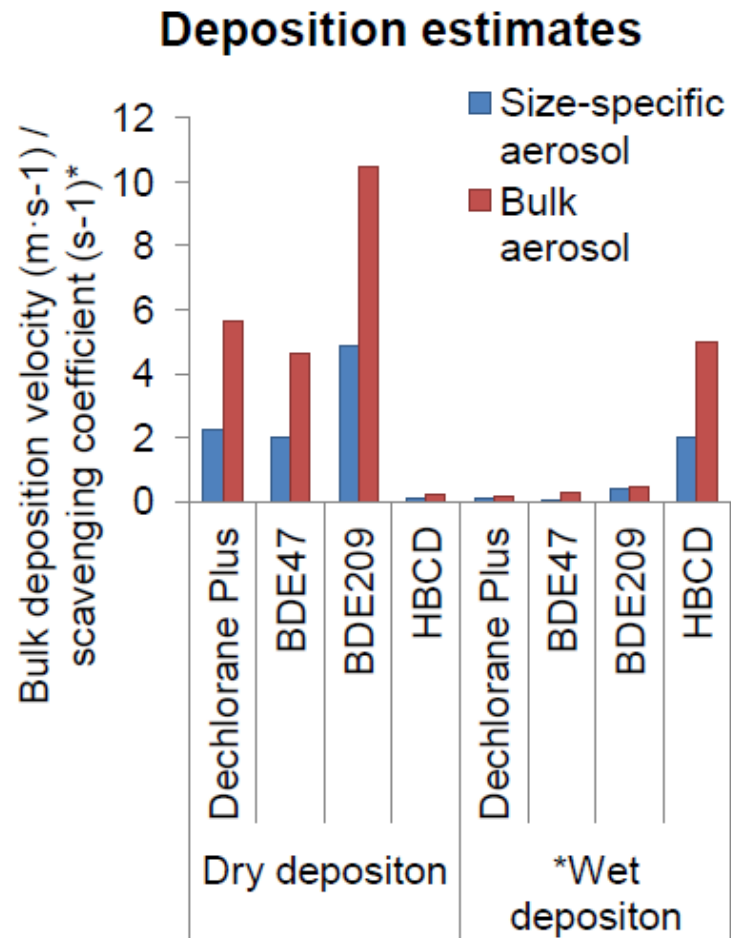


Higher concentrations in summer

Higher concentrations at the rural site (-> influence of the construction of new houses?)

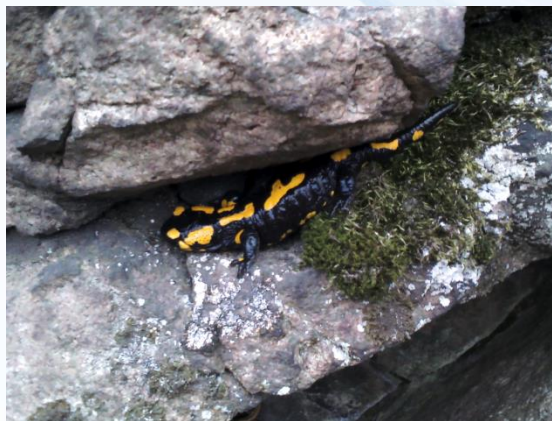
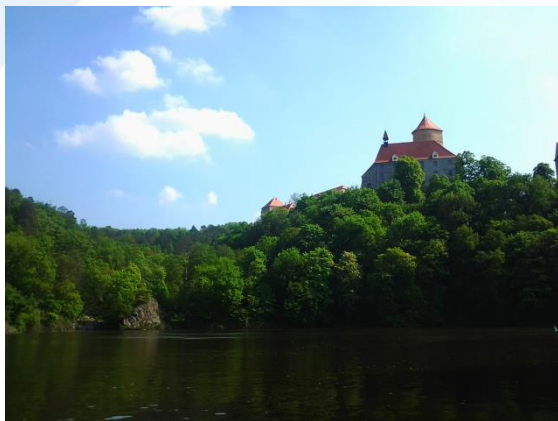
Strong accumulation in the fine fraction (57-74%)

Estimation of the wet and dry deposition



Thanks for your attention

Any questions?



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