

Characterization of particulate matter in Kraków, Poland.

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Airborne particulate matter poses a significant problem in South Poland. Kraków is classified at the eighth place among 575 most PM_{2.5}-polluted cities and at the 145th place among 1100 most PM₁₀-polluted cities listed in the WHO ranking (WHO, 2013). According to its basin location the city often suffers under temperature inversion causing high pollution periods. Continuous air quality monitoring and several studies conducted already in the region showed that residential solid fuel combustion and traffic have a dominant influence on particulate matter burden (e.g., Larsen *et al.*, 2007), being responsible for increased concentration of toxic and noxious chemicals. However, the available information is not persuasive enough to become a catalyst for efficient policies reducing the emissions and lowering the human exposure to hazardous substances.

The current study combines two sampling campaigns (February 2011 – PM₁₀ and February 2013 – PM₁₀ and PM_{2.5}). Particulate matter was collected in the centre of Krakow (Aleja Mickiewicza, AGH, University of Science and Technology) with low volume samplers operated with daily (24h) intervals. Quartz fibre filters (Pall, 47mm) were used for sampling, field blanks were collected.

Particulate matter mass was measured gravimetrically under stable humidity and temperature conditions. Circular aliquots were punched out of the filters in order to conduct chemical analyses. Analytes and the respective aerosol sources are listed in Table 1.

Table 1. Characteristic sources their chemical fingerprints and analytical methods.

| Source | Component group | Method |
|----------------------|-------------------|-----------------|
| Combustion | EC, OC | Thermal-optical |
| RWC, BA | Saccharides | HPAE PAD |
| SA | Inorganic anions | HPAE CD |
| SA | Inorganic cations | HPCE CD |
| Combustion | PAHs | GC-MS |
| Combustion, Industry | Selected elements | XRF |
| Combustion, Industry | Particulate Hg | CV-AAS |

RWC = residential wood combustion, BA = biological aerosol, SA = secondary aerosol, EC = elemental carbon, OC = organic carbon, PAHs = polycyclic aromatic hydrocarbons

The aim of the study is to chemically characterize PM₁₀ and PM_{2.5} as well as to quantify aerosol sources

being locally relevant. The comparison of two winter periods will help to assess the pollutants which were not reduced over a period of two years.

The source apportionment method used for Kraków is based on the macro-tracer model derived in Austria (Puxbaum *et al.*, 2005). This model is taking advantage of the existence of source specific compounds and stability of their relations in emission and ambient air.

The winter PM₁₀ concentrations observed during the sampling period in winter 2011 exceeded constantly the admissible threshold of 50 µg/m³. The similar trend is expected for 2013, as no powerful reduction measures have been applied in the city since that time.

In 2011 the highest mass contribution was found for carbonaceous aerosols (50%), followed by inorganic secondary aerosols (30%). Levoglucosan (anhydrous saccharide used to assess residential wood burning impact) concentrations did not exceed 1% of PM₁₀. Analysed PAHs contributed only 0.1% to PM₁₀ mass, but benzo(a)pyrene (BaP) concentrations were around 10 ng/m³ (being 10 times higher than the annual limit value). Residential coal combustion is expected to be the most prominent BaP source.

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