

Fossil and non-fossil source contributions to atmospheric carbonaceous aerosols during grassland fires

V. Ulevicius¹, S. Byčėnkiėnė¹, C. Bozzetti², A. Vlachou², K. Plauškaitė¹, G. Mordas¹, V. Dudoitis¹, G. Abbaszade³, J. Blee², R. Fröhlich², K.R. Dällenbach², F. Canonaco², J.G. Slowik², J. Dommen², R. Zimmermann³, J. Schnelle-Kreis³, G.A. Salazar², K. Agrios^{4,5}, S. Szidat⁴, I. El Haddad², and A.S.H. Prévôt².

¹Department of Environmental Research, SRI Center for Physical Sciences and Technology, 02300, Vilnius, Lithuania

²Laboratory of Atmospheric Chemistry, Paul Scherrer Institute (PSI), 5232, Villigen-PSI, Switzerland

³Helmholtz Zentrum München, German Research Center for Environmental Health (GmbH), Joint Mass Spectrometry Centre, Cooperation Group Comprehensive Molecular Analytics and Helmholtz Virtual Institute of Complex Molecular Systems in Environmental Health - Aerosol and Health, 85764 Neuherberg, Germany

⁴Department of Chemistry and Biochemistry & Oeschger Centre for Climate Change Research, University of Bern, 3012 Bern, Switzerland

⁵Laboratory of Radiochemistry and Environmental Chemistry, PSI, 5232 Villigen, Switzerland

Keywords: aerosol, source apportionment, carbonaceous aerosol.

Presenting author email: ulevicv@ktl.mii.lt

Wood and grass burning are the major sources of biomass-burning aerosols which are composed mainly of organic matter (Crutzen *et al.*, 1979). Approximately 90% of vegetation burning is caused by human-induced fires and only a minor fraction derives from natural processes such as lightning. The main type of biomass burning in Lithuania and surrounding countries in early spring is illegal last year's [grass burning](#) for land clearing (Ulevicius *et al.*, 2010, Byčėnkiėnė *et al.* 2013).

Continuous air monitoring and time integrated particulate matter sampling were carried out on March 2014 in Preila, Lithuania (55° 55' N, 21° 04' E 5 m above sea level) – a representative coastal background site, an ideal location for studying the long-range transport of air pollutants in the South-eastern Baltic region due to the absence of local sources.

An ACSM (Aerosol Chemical Speciation Monitor, Aerodyne Research, Inc., Billerica, MA, USA) was deployed at site. 1-h time resolution ACSM data were analysed to elucidate the PM₁ chemical composition. A graphical user interface SoFi (Canonaco *et al.*, 2013), developed at PSI was used to perform a PMF source apportionment of the non-refractory OA mass spectra collected during March 2014. For ¹⁴C analysis, the accelerator mass spectrometer MICADAS was used, which was equipped with a gas-capable ion source (Szidat *et al.*, 2014). ¹⁴C was determined after combustion of filter punches in an elemental analyser, which was directly coupled to the MICADAS. An estimate of fossil and non-fossil primary (POC_f, POC_{nf}) and secondary organic carbon (SOC_f, SOC_{nf}) was achieved by coupling ACSM-PMF results, ¹⁴C data and gas chromatography – mass spectrometry organic markers measurements using a CMB approach.

During the event, when grass burning was most intense, the levoglucosan concentration increased up to 683 ng m⁻³. Concentrations of mannosan varied from 3.10 to 68.0 ng m⁻³ and galactosan 1.02 – 12.0 ng m⁻³. Organics (46 %, 3.2 μg m⁻³ (σ = 4.8 μg m⁻³)) composed the major fraction of the NR-PM₁ aerosol concentration composition with lower contributions from sulphate

(17 %, 1.2 μg m⁻³ (σ = 1.1 μg m⁻³)), nitrate (20 %, 1.4 μg m⁻³ (σ = 1.8 μg m⁻³)), ammonium (15 %, 1.00 μg m⁻³ (σ = 0.9 μg m⁻³)), and chloride (2 %, 0.1 μg m⁻³ (σ = 0.3 μg m⁻³)). A further separation between fossil and non-fossil primary and secondary contributions to total carbon (POC_f, POC_{nf}, SOC_f, SOC_{nf}) was obtained (Fig. 1).

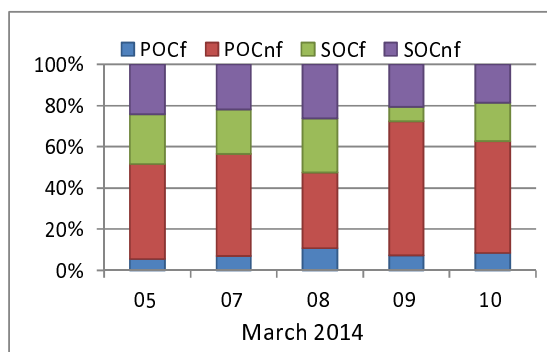


Figure 1. Median percentage contributions of different sources to total carbon.

This work was supported by the Lithuanian-Swiss Cooperation Programme “Research and Development” project AEROLIT (No. CH-3-ŠMM-01/08).

- Byčėnkiėnė, S., Ulevicius V., Dudoitis V., Pauraitė J. (2013). *Adv. Met.* **2013**, Article ID 380614.
- Canonaco, F., Crippa, M., Slowik, J. G., Baltensperger, U., and Prévôt, A.S.H. (2013). *Atmos. Meas. Tech.* **6**, 3649-3661.
- Crutzen, P.J., Heidt, L.E., Krasnec, J.P., Pollock, W.H., Seiler, W. (1979). *Nature* **282**, 253-256.
- Szidat, S., Salazar, G. A., Vogel, E., Battaglia, M., Wacker, L., Synal, H.-A., and Türler, A. (2014). *Radiocarbon*, **56**, 561-566.
- Ulevicius, V., Byčėnkiėnė, S., Remeikis, V., Garbaras, A., Kecorius, S., Andriejauskienė, J., Jasinevičienė, D., Mocnik, G. (2010). *Atmos. Res.* **98**, 190-200.