

Taking the step from bulk to size-segregated aerosol description: Modelling of size distributions with the EMEP/MSC-W model

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Knowledge of the size distribution of atmospheric aerosols in the submicron size range is essential for estimation of aerosol climate effects and also health implications. Concerning effects on Earth's radiative balance, aerosol extinction of short and long wave radiation and interaction with clouds depends on their size. On the other hand, growing evidences suggest increased health effect from transport-related particles, in particular ultrafine ones (e.g. Oberdörster, 2001; Krzyzanowski et al., 2005). Ultrafine particles contribute negligibly in currently regulated PM₁₀ and PM_{2.5} mass and should be described in terms of number density.

The EMEP/MSC-W model (Simpson et al., 2012) is an atmospheric chemistry transport model, calculating photo-chemistry and PM on regional to global scale. It is flexible with respect to the choice of projection and horizontal resolution (tested for between 4x4 km and 1x1° grid), and has 20 vertical layers up to 100 hPa. The standard model deals with bulk aerosol mass, distinguishing fine and coarse particles.

Table 1. Median particle number concentrations in the size range 30 - 50 nm and 50 - 500 nm. Observed values adopted from Asmi et al. (2011).

Site	N ₃₀₋₅₀ obs. (#/cm ³)	N ₃₀₋₅₀ model (#/cm ³)	N ₅₀₋₅₀₀ obs. (#/cm ³)	N ₅₀₋₅₀₀ model (#/cm ³)
Birkenes	156	296	511	121
Aspvreten	284	241	1081	132
SMEAR	223	239	878	115
Pallas	89	97	205	37
K-Pusztza	697	592	3120	375
Melpitz	860	434	2327	275
Kosetice	700	460	3194	265
Waldhof	878	413	2434	263
Cabauw	1914	889	3387	554
Finokalia	220	660	1345	625
Ispra	1341	1032	4448	598
Mace Head	105	364	241	199

Recently, aerosol dynamics of the sectional aerosol model MAFOR (Karl et al., 2011; <http://mafor.nilu.no>) have been implemented in the EMEP model to simulate size-resolved particle number and mass concentrations. In this work, 16 size sections were used to represent the aerosol size distribution from 1 nm to 10 μm. The new approach consistently solves the time evolution of the particle number and mass concentration distribution using the fixed sectional method in a robust manner on a 3-D atmospheric grid.

We present European maps of size-resolved aerosol and evaluate the model ability to reproduce observed particle size distributions. First comparison of modelled number concentration in the sizes of 30 - 50 nm (N₃₀₋₅₀) and 50 - 500 nm (N₅₀₋₅₀₀) to observational data (2008-2009) published by Asmi et al. (2011), summarized in Table 1, revealed reasonable agreement for most remote sites and esp. for the ultrafine fraction, whereas N₅₀₋₅₀₀ was underestimated by the model. The discrepancies are partly associated with uncertainties in the particle emission data, but are also due to currently deficient treatment of processes: better descriptions of particle deposition and of particle growth due to condensation of inorganic and organic vapours, are expected to improve the model's predictive capabilities.

Modelled size distribution at Birkenes, Ispra and K-Pusztza for winter and summer are shown in Figure 1. The shape of the size distributions has little variability among the sites, probably due to the use of a uniform particle emission spectrum for entire Europe. Increased nucleation of sulphuric acid led to higher numbers of nucleation mode particles in summer.

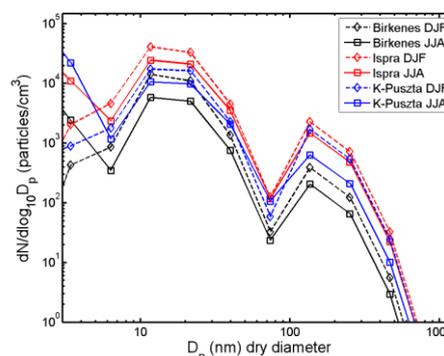


Figure 1. Modelled number size distribution in winter (DJF) and summer (JJA) at various sites in Europe.

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