



Global modeling of aerosol transport and optical properties with ORISAM-TM4 model in sectional framework including organics, inorganics, dust and sea-salts

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Black carbon (BC) and primary organic carbon (OC_p) are directly emitted by combustion sources whereas organic carbon aerosols further involve photooxidation reactions (OC_{sec}). This complexity together with size effects, both linked to emissions and secondary organic formation due to organic gaseous precursors, is poorly represented in models.

To explore this complexity, a global aerosol model ORISAM-TM4 [Guillaume et al., 2007] has been developed by coupling a global chemistry-transport model TM4 [Van Velthoven et al., 1996] and the aerosol sectional model ORISAM (Organic and Inorganic Spectral Aerosol Model) [Bessagnet et al., 2004; Lioussé et al., 2005; Cousin et al., 2005]. This modeling system allows us to take into account a sectional size-distribution (8-20 bins between 0.04 μm and 10 μm), updated emission inventories for gases and primary particles and detailed aerosol chemical composition (BC and primary OC being the particle cores on which 6 organic/inorganic chemical species are absorbed/adsorbed). More recently, additional aerosol components have been incorporated in particle cores (sea-salts, dust), giving major contribution in the coarse particle mode. A simple model of coarse nitrate formation from heterogeneous processes [Hodzic et al., 2005] is also implemented. Simulations of global aerosol transport and optical properties have been performed with ORISAM-TM4 endowed with the RAD module [Mallet et al., 2005] for the complex internally-mixed aerosol (dust/organics/inorganics) over the detailed full size-spectrum. Sensitivity tests have

been run on external vs. internal mixing state models. A second series of tests were made on the impact of explicit vs. simply parametrized OC. Tentative validation of these results is performed against AERONET network and MODIS satellite data.