

Experimental study on the capture/desorption of gaseous methyl iodide on sea salt aerosols

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Iodine-131, when released into the environment during severe nuclear power plant accident can have a high radiological impact on the population at short term [1]. Interaction between gaseous Iodine compounds and aerosols was not considered by the current post-accident management. In this context, this work was focused on investigating the influence of sea salt aerosols on the transport of gaseous methyl iodide (CH₃I). The identification of uptake processes as well as the formation of new products at the particle surfaces was the main objectives.

We have studied the interaction between NaCl particles as surrogate of sea salt particles and CH₃I in various humidity conditions to reproduce the atmospheric conditions.

The nature of this interaction was investigated by Infrared Spectroscopy (DRIFTS, Diffuse Reflectance Infrared Fourier Spectroscopy). Solid NaCl was exposed to CH₃I (1000 and 500 ppm) with a relative humidity (RH) ranging between 0 and 80%.

DRIFTS results clearly evidenced adsorbed CH₃I on NaCl particles surface under both dry and humid conditions. The kinetic results exhibited very low uptake coefficients in the order of 10⁻¹⁴ in all the experimental conditions. The adsorption process follows three regimes that can be fitted with First-order Langmuir adsorption isotherm models.

Additionally, to the CH₃I absorption bands, the DRIFT spectrum evidenced typical absorption bands that could be assigned either to the CH₂ deformation of CH₂I₂ or to CH₃ degenerate rocking of CH₃Cl. The formation of new bands appears only when CH₃I is in presence of halogenated salts [2]. However, at RH=80%, the water layer at the particle surface inhibits the interaction between gaseous CH₃I and NaCl surface due to the low solubility of CH₃I in water.

Although the uptake coefficients of CH₃I are quite low, a coverage of particle surface with CH₃I-derived compounds may affect the reactivity of the particles and in term the cycling life of Iodine in the atmosphere.

Références :

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[2] Chebbi, M.; Azambre, B.; Cantrel, L.; Koch, A. J. Phys. Chem. C 2016, 120 (33), 18694–18706.