



## Measurement of uptake coefficients of HO<sub>2</sub> radical with aerosol particles sampled in the ambient air

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HO<sub>x</sub>(=OH+ HO<sub>2</sub>) radicals play a central role in the tropospheric chemistry. The total concentration of HO<sub>x</sub> radicals is determined by the balance of sources and sinks. Recent field measurements of HO<sub>x</sub> radicals suggested that the heterogeneous loss of HO<sub>2</sub> by aerosol particles was potentially important HO<sub>x</sub> sink in the troposphere. However, there are a few data to assess the contribution to HO<sub>2</sub> loss by aerosols. In this study, the HO<sub>2</sub> uptake coefficients for aerosol particles sampled by filter in ambient air were measured using an aerosol flow tube (AFT) coupled with a chemical conversion/laser-induced fluorescence (CC/LIF) technique at 760Torr and 298K. Aerosol particles were regenerated by atomizer using the water extract of PM2.5 sampled on filter at three sites(Taishan and Mangshan in China, and Yokosuka in Japan). The CC/LIF technique enabled experiments at HO<sub>2</sub> radical concentrations similar to those in the atmosphere (-10<sup>8</sup> molecules/cm<sup>3</sup>). The uptake coefficients of HO<sub>2</sub> by sampled aerosol particles were in the range of 0.1-0.4 at 75% of relative humidity. To assess the impact of the heterogeneous loss for HO<sub>2</sub>, we carried out box model calculations for the diurnal variation of HO<sub>2</sub> concentration.