

Nitryl chloride as a ‘new’ radical source and its role in production of ozone in polluted troposphere: an overview of the results from four field campaigns in China

Tao Wang (1), Yee Jun Tham (1), Likun Xue (2), Zhe Wang (1), Xinfeng Wang (2), Weihao Wang (1), Hao Wang (2), Hui Yun (1), Keding Lu (3), Min Shao (3), Peter K. K. Louie (4), Donald R. Blake (5), Steven S. Brown (6,7), and Yuanhang Zhang (3)

(1) Department of Civil and Environmental Engineering, Hong Kong Polytechnic University, Hong Kong, China (cetwang@polyu.edu.hk), (2) Environment Research Institute, Shandong University, Jinan, Shandong, China, (3) State Key Joint Laboratory of Environmental Simulation and Pollution Control, College of Environmental Sciences and Engineering, Peking University, Beijing, China, (4) Environmental Protection Department, Government of the Hong Kong Special Administrative Region, Hong Kong, China, (5) Department of Chemistry, University of California at Irvine, Irvine, CA, USA, (6) NOAA Earth System Research Laboratory, Chemical Sciences Division, Boulder, CO, USA, (7) Department of Chemistry and Biochemistry, University of Colorado, Boulder, CO, USA

Nitryl chloride (ClNO_2) - a trace gas produced from heterogeneous reactions of dinitrogen pentoxide (N_2O_5) on aerosols containing chlorine - can significantly affect radical budget and concentrations of ozone and other secondary pollutants. However, the abundance, formation kinetics, and impact of ClNO_2 are not fully understood under different environmental conditions. This presentation gives an overview of recent field campaigns of ClNO_2 and related chemical constituents in China, including one at a mountain top (957 m a.s.l) in Hong Kong of South China in winter 2013 and three in North China (urban Ji’nan, semi-rural Wangdu, and Mt Tai (1534 m a.s.l)) in summer 2014. ClNO_2 and N_2O_5 were measured with a chemical ionization mass spectrometry (CIMS) system with iodide as the primary ions. Ambient concentrations of several hundreds ppt and up to 4.7 ppbv of ClNO_2 were observed in these locations, suggesting existence of elevated ClNO_2 in both coastal and inland atmospheres of China. Measurements in North China exhibited generally low concentrations of N_2O_5 , indicative of its fast uptake of on aerosols under aerosol and humid conditions. Indications of anthropogenic sources of chloride were observed at all these sites. The impact of photolysis of ClNO_2 on radical budget and ozone enhancement was assessed with a MCM model which was updated with detailed chlorine chemistry and constrained by measurement data for the southern and a northern site. The results show that the ClNO_2 could increase ozone production by 2-16% in the following day. Overall, our study re-affirms the need to include ClNO_2 related reactions in photochemical models for prediction of ground-level ozone in polluted environments.