

## 2.107 Heterogeneous uptake of N<sub>2</sub>O<sub>5</sub> in urban and sand dust plumes observed in spring in Beijing, China: implications for parameterizations and particulate nitrate formation.

Early Career Scientist

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Abstract:

Dinitrogen pentoxide (N<sub>2</sub>O<sub>5</sub>) and its heterogeneous product on aerosol, nitryl chloride (ClNO<sub>2</sub>), contribute to nocturnal nitrate formation and impact daytime oxidative capacity. However, ambient observations of N<sub>2</sub>O<sub>5</sub> and ClNO<sub>2</sub> are still limited, which precludes full understanding of reactive nitrogen chemistry in various conditions. Here we present observations of N<sub>2</sub>O<sub>5</sub> and ClNO<sub>2</sub> at a ground site in Beijing in 2017, focusing on the intercepted urban and sand dust plumes with abundant N<sub>2</sub>O<sub>5</sub> and/or ClNO<sub>2</sub>. High levels of N<sub>2</sub>O<sub>5</sub> (up to 2.2 ppbv) were observed on May 1<sup>st</sup> night due to insignificant heterogeneous loss. In comparison, ClNO<sub>2</sub> mixing ratios of up to 3.3 ppbv were frequently observed in late May in humid and chemically processed urban plumes. Significant levels of ClNO<sub>2</sub> (up to 0.7 ppbv) characterized with very fast heterogeneous loss of N<sub>2</sub>O<sub>5</sub> ( $k(\text{N}_2\text{O}_5)$  up to 0.02 s<sup>-1</sup>) were observed in a sand storm event. N<sub>2</sub>O<sub>5</sub> uptake coefficient ( $\gamma$ ) is calculated for various air masses and found more variable than that suggested by parameterizations. The observed  $\gamma$  is mostly linked to the ratio of aerosol volume to surface area density ( $V_a/S_a$ ) but less dependent on [H<sub>2</sub>O] or water-soluble ions. Utilizing the derived uptake coefficient ( $\gamma=0.027\pm0.010$ ) along with related data, nocturnal nitrate production rates are calculated and found correlated to observed nitrate increasing rates. In the ClNO<sub>2</sub>-rich urban plumes, heterogeneous uptake of N<sub>2</sub>O<sub>5</sub> results in comparable or

higher nitrate formation potential than daytime  $\text{OH} + \text{NO}_2$  reaction. Higher  $\text{NO}_3$  production coupled with larger proportion of  $\text{N}_2\text{O}_5$  lost in heterogeneous uptake is responsible for more significant nocturnal nitrate production. Overall, our results indicate notable nighttime chemistry of  $\text{N}_2\text{O}_5$  in spring time of urban Beijing and its significant contribution to particulate nitrate formation.