

Most of the previous field studies of N₂O₅ uptake and ClNO₂ production have been conducted in the United States of America (US) and Europe (Brown et al., 2009; Chang et al., 2016). Direct field investigation of the N₂O₅ heterogeneous processes in China is very limited. Pathak et al. (2009, 2011) analyzed the aerosol composition and suggested that the accumulation of fine NO₃⁻ aerosol downwind of Beijing and Shanghai was due to significant N₂O₅ heterogeneous re-

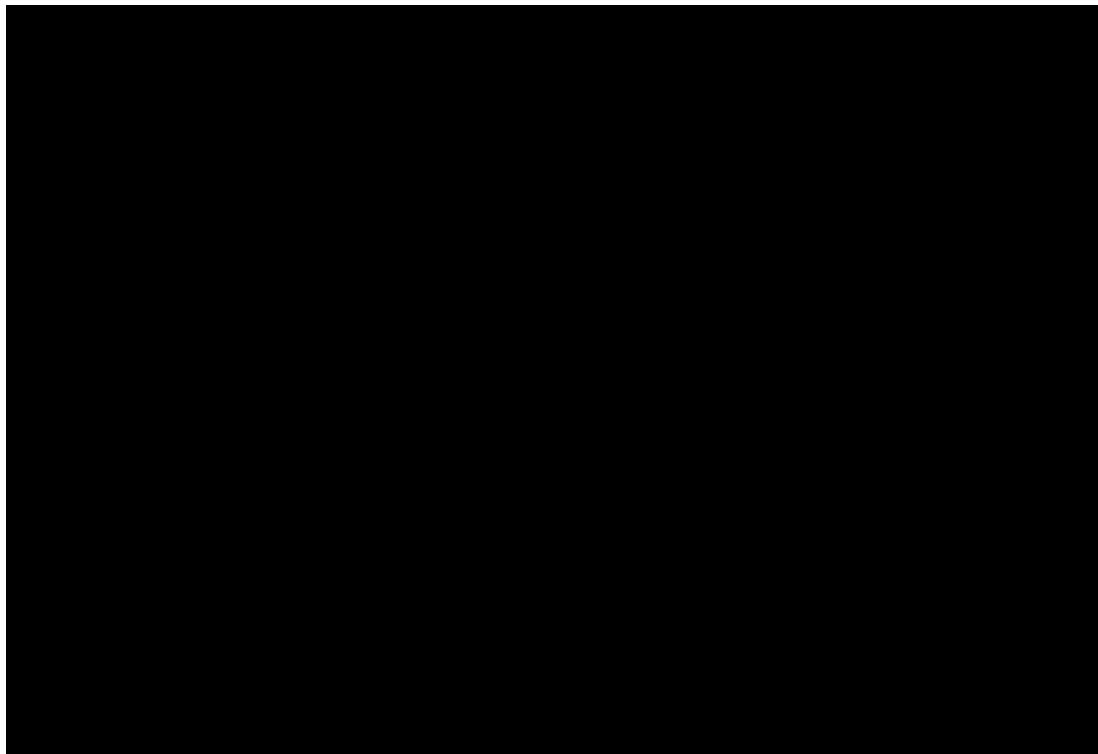


Figure 1. Time series of concentrations of N₂O₅, CINO₂, NO₃ production rates, the steady-state lifetimes of N₂O₅, and related gas and aerosol data at Wangdu from 21 June to 9 July 2014. N₂O₅ and CINO₂ are 1 min averaged data, whereas the NO_x, O₃, and NO₃ production rates and $\langle N_2O_5 \rangle$ are given as 5 min averages. The data for S_a and fine particulate NO₃ are in 10 and 30 min time resolutions, respectively. The data gaps were caused by technical problems, calibrations, or instrument maintenance.

inant source of particulate nitrate. Moreover, the production rate of HNO₃, as calculated from the gas-phase reactions of OH + NO₂ and NO₃ + VOC, shows a decreasing trend towards the night (Fig. 3c), and the combination of these rates on average is less than one-third of the average ρNO_3 , which was determined from the slope of nighttime particulate NO₃ in Fig. 3b. The increase in nighttime NO₃ is $6.436 \text{ Tf} / 9.9626 \text{ Tf} = 0.6436$.

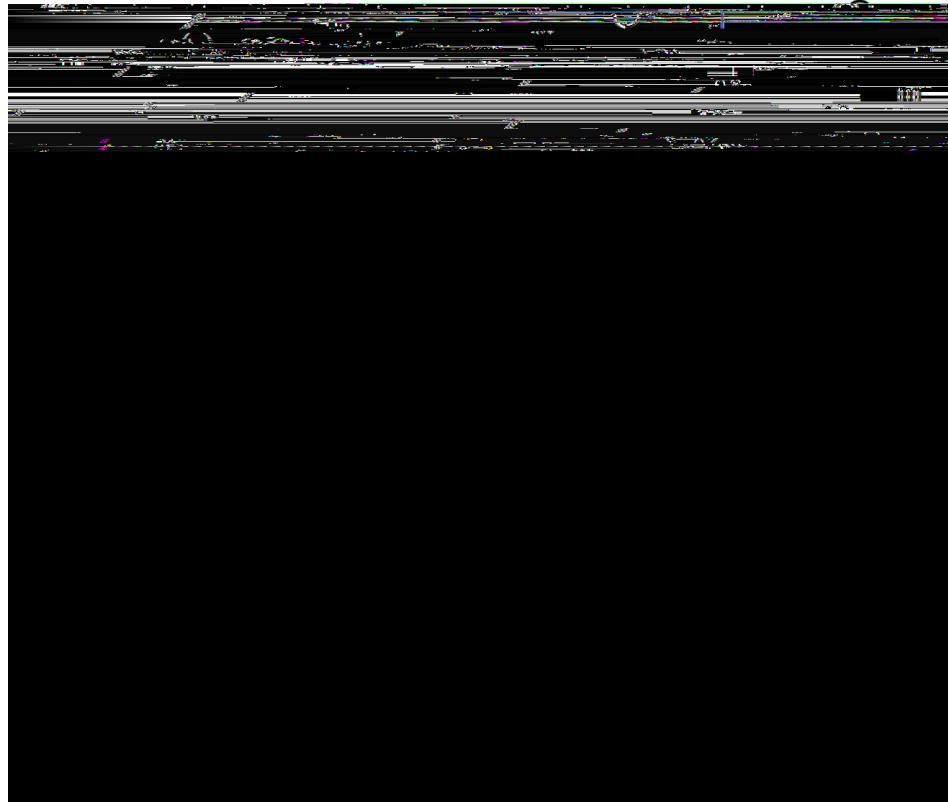


Figure 7. Scatter plots for (a) yield derived from the field versus yield calculated from the parameterization, using $k_{R4}=k_{R3}$ of 483 (recommended by Bertram and Thornton, 2009; solid circle) and 836 (recommended by Behnke et al., 1997; pink open circle). Error bars represent the uncertainty in field-derived , and the black dotted line represents the 1 : 1 ratio; (b) field-derived yield versus aerosol water content; (c) field-derived yield versus chloride; and (d) field-derived yield versus CH₃CN=

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