



Supplement of

Gaseous products and Secondary Organic Aerosol formation during long term oxidation of isoprene and methacrolein

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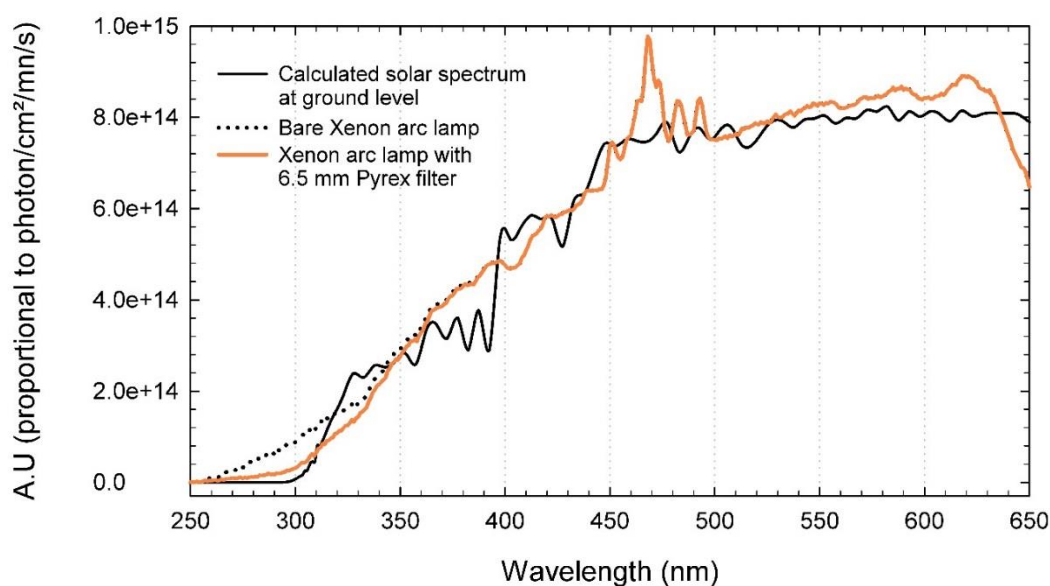


Figure S1 Solar irradiation spectrum (solid black curve) calculated from TUV NCAR, 12 h, Equator, 21 June compared to irradiation spectra of xenon arc lamp without filter (dotted black curve) and with a 6.5 mm thicknesses Pyrex[®] filter (orange curve). The lamps spectra are scaled to the solar spectrum to facilitate comparison of their shapes.

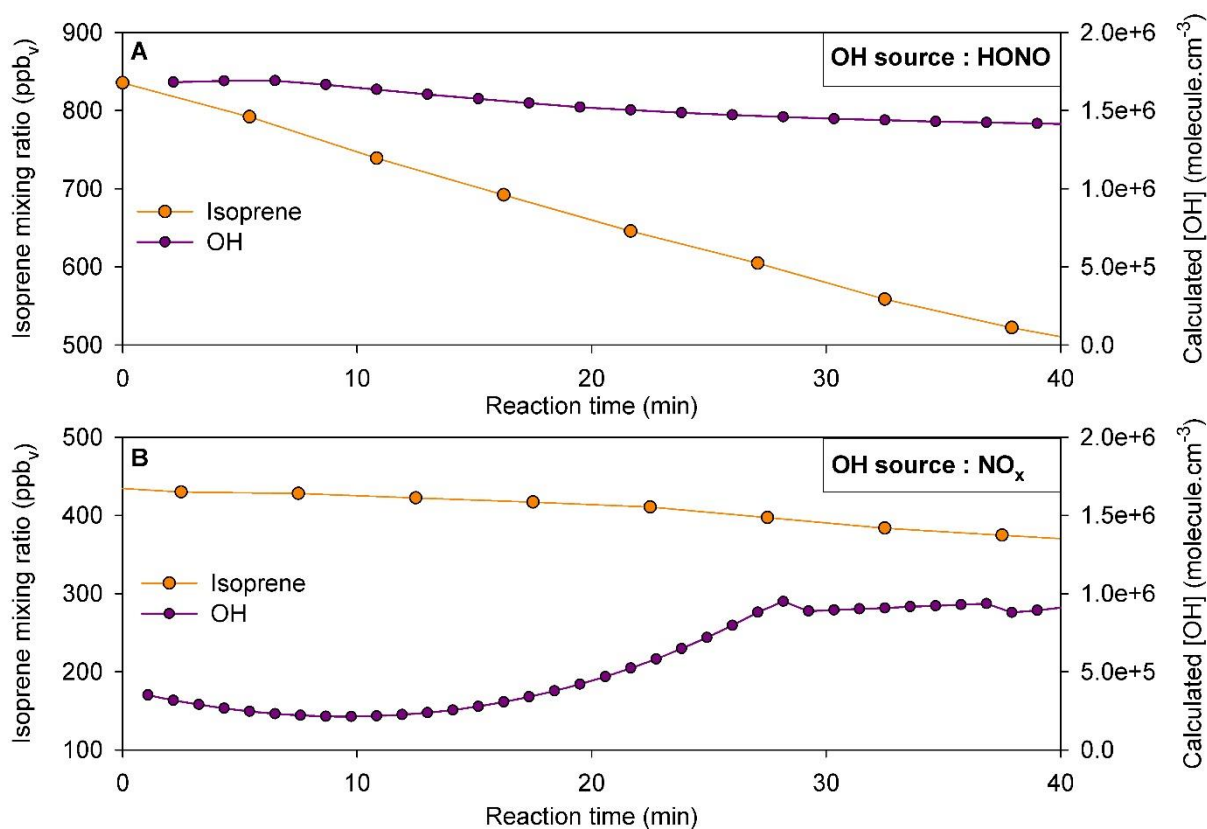


Figure S2 Time profiles of isoprene mixing ratios and calculated OH concentrations during isoprene photooxidation experiments performed after manual cleaning with (A) HONO (I280113) and (B) NO_x (I210512) as OH source.

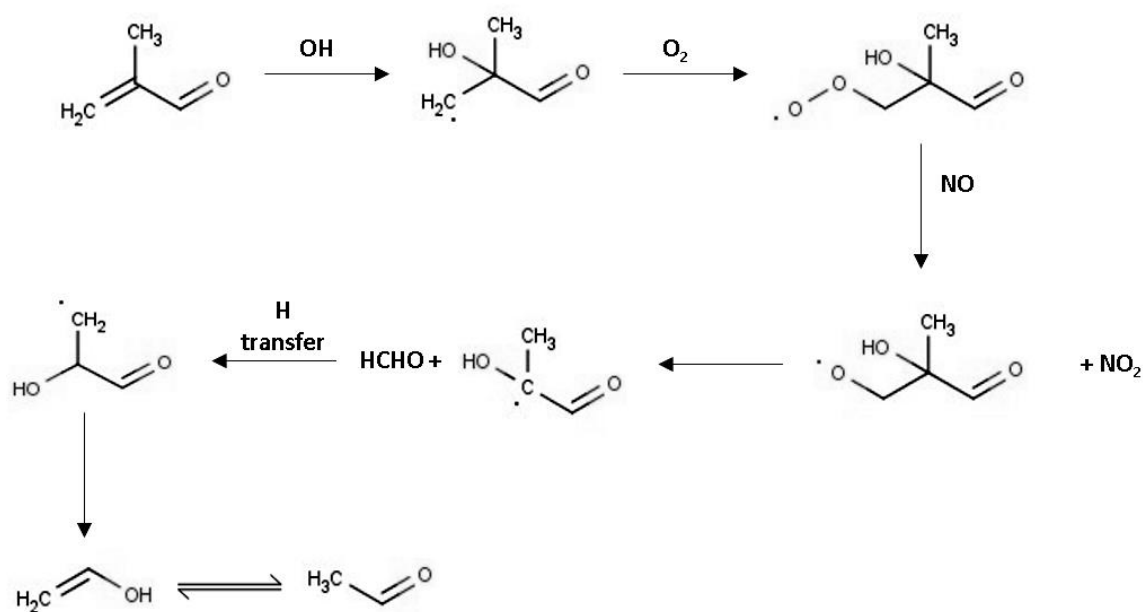


Figure S3 Suggested primary acetaldehyde formation from MACR which implies an H transfer in gaseous phase.

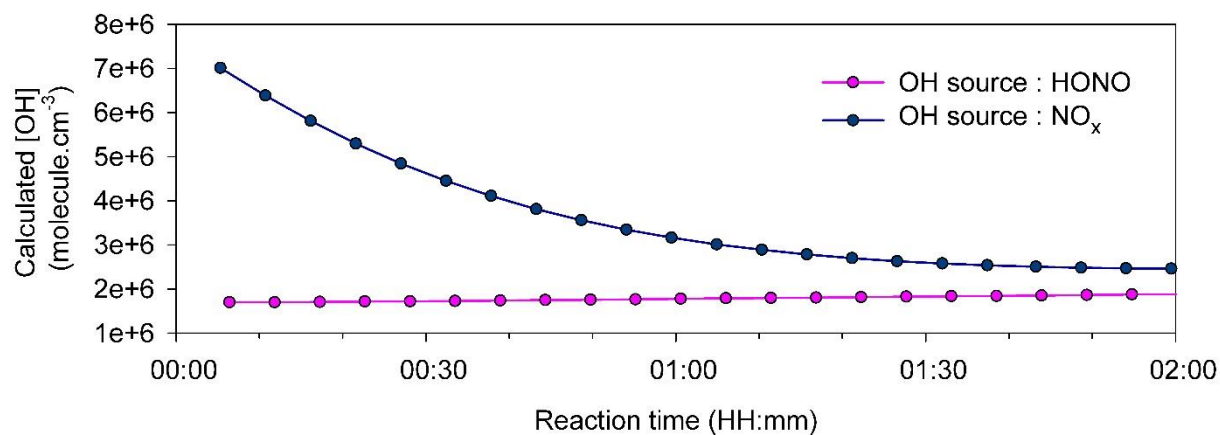


Figure S4 Time profiles of calculated OH concentrations during MACR photooxidation experiments performed with HONO (pink curve; M230113) and NO_x (blue curve; M240512) as OH source.

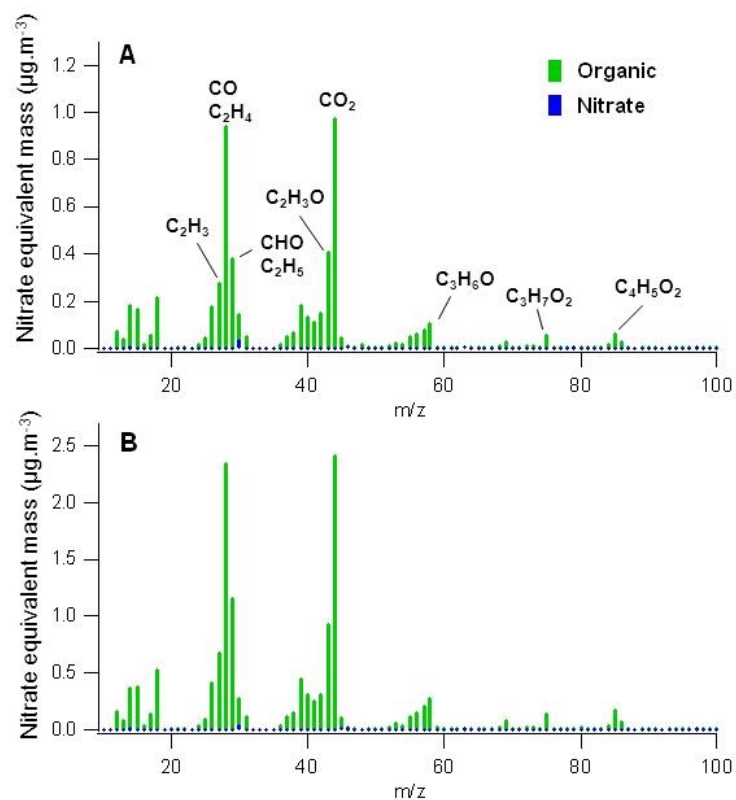


Figure S5 High-resolution mass spectra of SOA from (A) isoprene (I110411) and (B) MACR (M120411) photooxidation in the presence of NO_x . Spectra were taken at the maximum of SOA growth. The contribution of CO^+ to the total signal was estimated from the CO_2^+ organic signal ($\text{CO}_2^+ = \text{CO}^+$) like it was proposed by Chhabra et al. (2010) for isoprene experiments.