



Measurements and modelling of OH and HO₂ radicals over a South-East Asian tropical rainforest during the OP3 field campaign

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Forests are the dominant source of volatile organic compounds (VOCs) into the atmosphere, with isoprene being the most significant species. The oxidation chemistry of these compounds is a significant driver of local, regional and global atmospheric composition. Recent evidence suggests a significant failing in our understanding of VOC oxidation chemistry under low NO_x conditions, with important consequences for modelling of OH and the climate gases methane and ozone.

During the Oxidant and Particle Photochemical Processes (OP3) project over Borneo in July 2008 aircraft observations of OH and HO₂ were made using FAGE (Fluorescence Assay by Gas Expansion). We investigate these observations with a constrained box model using the Master Chemical Mechanism (MCM). In line with previous work we significantly underestimate the observed OH and HO₂ concentrations, with mean observed to modelled ratios of OH and HO₂ in airmasses impacted with isoprene of (5.32 ± 8.74) and (1.18 ± 0.31) respectively. A variety of mechanistic changes are made to the model, and their impact on the model success will be described. None of the suggested mechanistic changes allow for a good simulation of both OH and HO₂ concentrations in the model.