## 4.086 OH and HO2 concentration observations in the upper troposphere inside and outside of Asian monsoon influenced air..

Early Career Scientist

**Presenting Author:** 

Daniel Marno, Max-Planck Institut fur Chemie, daniel.marno@mpic.de

## Co-Authors:

Christopher Künstler, Institute of Energy and Climate Research, IEK-8: Troposphere, Forschungszentrum Jülich Korbinian Hens, Max-Planck Institut fur Chemie Cheryl Ernest, Max-Planck Institut fur Chemie Monica Matinez, Max-Planck Institut fur Chemie Hendrik Fuchs, Institute of Energy and Climate Research, IEK-8: Troposphere, Forschungszentrum Jülich Efstratios Bourtsoukidis. Max-Planck Institut fur Chemie Jonathan Williams, Max-Planck Institut fur Chemie Frank Holland, Institute of Energy and Climate Research, IEK-8: Troposphere, Forschungszentrum Jülich Andreas Hofzumahaus, Institute of Energy and Climate Research, IEK-8: Troposphere, Forschungszentrum Jülich Laura Tomsche, Max-Planck Institut fur Chemie Bettina Hottmann. Max-Planck Institut fur Chemie Horst Fischer, Max-Planck Institut fur Chemie Theresa Klausner, Deutsches Zentrum für Luft- und Raumfahrt (DLR) Hans Schlager, Deutsches Zentrum für Luft- und Raumfahrt (DLR) Greta Stratmann, Deutsches Zentrum für Luft- und Raumfahrt (DLR) Paul Stocks, Deutsches Zentrum für Luft- und Raumfahrt (DLR) Helmut Ziereis, Deutsches Zentrum für Luft- und Raumfahrt (DLR) Birger Bohn, Institute of Energy and Climate Research, IEK-8: Troposphere, Forschungszentrum Jülich Andreas Zahn, Karlsruhe Institute of Technology IMK-ASF Andreas Wahner, Institute of Energy and Climate Research, IEK-8: Troposphere, Forschungszentrum Jülich Jos Lelieveld, Max-Planck Institut fur Chemie Harder Hartwig, Max-Planck Institut fur Chemie

## Abstract:

The Asian monsoon convectively transports pollutants like volatile organic compounds (VOCs),  $NO_X$ , and  $SO_2$  from the boundary layer over South Asia into the upper

troposphere where they can potentially enter the stratosphere, or be dispersed globally throughout the troposphere. Therefore, it is crucial to understand the oxidizing capacity of this system and its impact on pollutant degradation and aerosol formation. The Hydroxyl radical (OH) plays a central role and is the most important oxidizing molecule in the atmosphere. During the OMO-ASIA campaign in the summer of 2015,  $HO_X$  (OH and HO 2) was measured onboard the High Altitude Long-Range Research Aircraft (HALO). Two laser-induced fluorescence instruments based on the fluorescence assay by gas expansion technique (LIF-FAGE) were installed, the AIR-LIF instrument from Forschungszentrum Jülich GmbH and the HORUS instrument from the Max Planck Institute for Chemistry, Mainz. To measure the chemical background of OH potentially produced inside HORUS from highly oxidized VOCs, an Inlet Pre-injector (IPI) system was used. This was the first time an IPI system was implemented within an airborne LIF-FAGE instrument measuring HO<sub> $\chi$ </sub>. Inside the Asian monsoon outflow total HO<sub> $\chi$ </sub> concentrations did not increase significantly compared to outside. However, OH concentrations were on average 37% higher inside the anticyclone. This is mainly attributable to increased  $NO_X$  levels within the anticyclone with a significant NO source from lightning. This strong shift of  $HO_X$ towards OH results in accelerated oxidation rates of pollutants, implying that the Asian monsoon anticyclone acts like an atmospheric purifier in addition to a pollution pump. The CAABA-MECCA box model was used to test our current understanding of HO<sub>X</sub> chemistry in the upper troposphere. Constraining the model with measurements from OMO-Asia has provided insight into the extent of differences in the contribution and composition of  $HO_X$  sources, sinks and cycling in and outside the anticyclone.