

4.068 PROFILES OF NO_x AND O₃ IN AN AMAZONIAN RAINFOREST: COMPARISON OF MEASURED PROFILES WITH A MULTI-LAYER CANOPY CHEMICAL EXCHANGE MODEL.

Early Career Scientist

Presenting Author:

Stefan Aiko Wolff, Biogeochemistry Department, Max Planck Institute for Chemistry – Mainz, Germany, aikos-box@gmx.de

Co-Authors:

Laurens Ganzeveld, Environmental Sciences Department, Wageningen UR – Wageningen, The Netherlands

Anywhere Tsokankunku, Biogeochemistry Department, Max Planck Institute for Chemistry – Mainz, Germany

Christopher Poehiker, Biogeochemistry Department, Max Planck Institute for Chemistry – Mainz, Germany

Leonardo Deane de Abreu Sá, Instituto Nacional de Pesquisas Espaciais/ INPE, Centro Regional da Amazônia/CRA – Belém, PA, Brazil

Antonio Ocimar Manzi, Instituto Nacional de Pesquisas da Amazônia/ INPA – Manaus, AM, Brazil

Rodrigo Souza, Instituto Nacional de Pesquisas da Amazônia/ INPA – Manaus, AM, Brazil

Ivonne Trebs, Luxembourg Institute of Science and Technology, Environmental Research and Innovation (ERIN) Department, L-4422 Belvaux, Luxembourg

Matthias Soergel, Biogeochemistry Department, Max Planck Institute for Chemistry – Mainz, Germany

Abstract:

In 2011, an 80 m high walk up tower for atmospheric research was erected at the ATTO (Amazon Tall Tower Observatory) site (02°08'38.8"S, 58°59'59.5"W) in the remote Amazonian rainforest. The nearly pristine environment allows biosphere-atmosphere studies within an ecosystem far away from large anthropogenic emission sources. Since April 2012 vertical mixing ratio profiles of H₂O, CO₂ and O₃ were measured at 8 different heights between 0.05 m and 79.3 m. During five intensive campaigns (Oct-Dec 2012, Oct-Nov 2013, Mar 2014, Aug-Sep 2014, Oct-Dec 2015) nitric oxide (NO) and nitrogen dioxide (NO₂) were also measured. Since the end of 2015 NO_x measurements are performed continuously.

We applied the Multi-layer Canopy Chemical Exchange Model – MLC-CHEM to support the analysis of the observed profiles of NO_x and O₃. This includes inferring bi-directional surface-atmosphere exchange fluxes as well as the role of the canopy interactions between the emissions, dry deposition, chemistry and turbulent transport of trace gases. During our investigation of diurnal and seasonal differences between model and

measurements, we conducted a set of sensitivity studies to analyse the effects of changes in NO_x -soil emissions, in-canopy turbulence and resistances for O_3 and NO_2 uptake on wet surfaces. These analyses suggest some modification in the representation of some of the poorly constrained canopy processes resulting in a significantly better agreement between the simulated and measured exchange fluxes and concentrations. Furthermore we have compared different NO_x -soil emission flux-scenarios with NO_x -fluxes measured by different techniques.