



Comparison of measured reactive trace gas profiles with a multi-layer canopy chemical exchange model in an Amazonian rainforest

Stefan Wolff (2), Laurens Ganzeveld (4), Anywhere Tsokankunku (2), Christopher Pöhlker (2), Leonardo Deane de Abreu Sá (3), Antonio Ocimar Manzi (1), Rodrigo Souza (1), Ivonne Trebs (2,5), and Matthias Sörgel (2)

(2) Biogeochemistry Department, Max Planck Institute for Chemistry – Mainz, Germany, (4) Environmental Sciences Department, Wageningen UR – Wageningen, The Netherlands, (3) Instituto Nacional de Pesquisas Espaciais/ INPE, Centro Regional da Amazônia/CRA – Belém, PA, Brazil, (1) Instituto Nacional de Pesquisas da Amazônia/ INPA – Manaus, AM, Brazil, (5) now at: Luxembourg Institute of Science and Technology, Environmental Research and Innovation (ERIN) Department, L-4422 Belvaux, Luxembourg

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.

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 O₃ and NO₂ 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 comparison between the simulated and measured exchange fluxes and concentrations.