## 5.143 Recent aircraft- and ground-based observations of longer-lived VOCs and comparisons to CAM-chem model simulations .

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## Abstract:

Volatile Organic Compounds (VOCs) play an important role in tropospheric chemistry and ozone formation. A primary sink for these species is through reaction with the hydroxyl radical (OH) and as such these species help determine the OH budget. The recent NSF/NCAR DC3, TORERO, CONTRAST and ORCAS campaigns have provided aircraft observations of longer-lived VOC species such as ethane, propane, acetone, 2-butanone, and methanol as well as other shorter-lived alkanes over a relatively large spatial extent. Observations are from the Trace Organic Gas Analyzer (TOGA) and the Advanced Whole Air Sampler (AWAS). These trace gases have long enough lifetimes to be transported over large distances, providing the opportunity to test our understanding and representation of global emissions, transport, and chemistry in the global chemistry climate model CAMchem (Community Atmosphere Model with Chemistry, a component of the Community Earth System Model). The model was run with specified dynamics to represent the specific dates of the campaigns allowing for guantitative comparison between model results and observations. We also compare the aircraft data, where possible, to groundbased networks such as the NOAA Global Greenhouse Gas Reference Network and the UCI Global Monitoring network. We find that current emissions inventories used in global models underestimate hydrocarbon emissions in the Northern Hemisphere and new insights are gained on VOC distributions in the much less well studied Southern Hemisphere.