



Testing of a global carbon cycle model that includes Carbonyl Sulfide (COS)

Adam Wolf (1), Ian Baker (2), J.A. Berry, N.J. Blake (3), D.R. Blake (3), E. Campbell (1), A.S. Denning (2) S.R. Kawa (4), S. Meinardi (3), S.A. Montzka (5), U. Seibt (1), S.A. Vay (6), Z. Zhu (4)

(1) Carnegie Inst. of Science, Dept. of Global Ecology, 260 Panama St., Stanford, CA 94305 USA (joeberry@stanford.edu, /Phone: +1 650 464 1047), (2) Dept of Atmos. Sci., Colorado State University, Fort Collins, CO, 0523-1371 USA (denning@colostate.edu, /Phone: +1 970 491 8359), (3) University of California 570 Rowland Hall Mail Code: 2025 Irvine, CA 92697 USA (DRBLAKE@uci.edu, /Phone: +1 949 824 4195) (4) NASA Goddard Space Flight Center, Code 613.3, Greenbelt, MD 20771 USA (kawa@maia.gsfc.nasa.gov, /Phone: +1 301 614 6004), (5) NOAA/ESRL/GMD, 325 Broadway, Boulder, CO 80305 USA (Stephen.A.Montzka@noaa.gov, /Phone: +1 303 497 6657) (6) NASA LaRC, Code E303 Hampton VA 23681 USA (Stephanie.a.vay@nasa.gov, /Phone: +1 757 864 1574)

Recent plant chamber and ambient atmospheric measurements suggest that atmospheric carbonyl sulfide (COS) may be a useful tracer for carbon cycle processes. We have included a mechanistic representation of COS exchange by leaves and soils in SiB3, a land-surface/carbon-cycle model. SiB generated COS and CO₂ fluxes were used together with other information on sources and sinks to simulate time series of the concentration of COS and CO₂ in the global atmosphere with chemical transport models. Here, we describe our parameterization of COS exchange in SiB3 and compare our meso-scale and global-scale simulations of COS and CO₂ concentrations to observations at 13 NOAA background atmospheric sites and many points in the free troposphere and the marine and continental boundary layers sampled by aircraft campaigns (NOAA, INTEX-NA and TC4).