How Much Can We Learn About Nitrous Oxide Emissions from Background Sites and Simple Box Models?

 $\underline{J.W.\ Elkins}^{\scriptscriptstyle 1}$, G. Dutton², D. Nance², B. Hall¹, D.J. Mondeel², J.H. Butler¹, E. Dlugokencky¹, S. Wofsy³ and M. Rigby⁴

¹NOAA Earth System Research Laboratory, 325 Broadway, Boulder, CO 80305; 303-497-6224, E-mail: James.W.Elkins@noaa.gov

²Cooperative Institute for Research in Environmental Sciences (CIRES), University of Colorado, Boulder, CO 80309

³Harvard University, Cambridge, MA 02138

⁴University of Bristol, Bristol, United Kingdom

Atmospheric Nitrous Oxide (N₂O) is an important ozone-depleting gas that continues to rise in concentration even as chlorofluorocarbon emissions have virtually ceased. It is also a potent greenhouse gas with a global warming potential of 298 times per molecule that of carbon dioxide with 100 years time horizon. NOAA has been monitoring background concentrations of N₂O from weekly flask sampling since 1977, starting with five remote stations over a broad latitudinal coverage from Pt. Barrow, Alaska to South Pole, Antarctica. This network has expanded to thirteen flask sites and six in situ instrument sites. We have combined data from the collocated, ground-based sites using three different independent instruments all linked to the World Meteorological Organization N₂O calibration scale, primarily to assist in quantifying the global burden of atmospheric N₂O for international assessments of the state of the science in climate and stratospheric ozone depletion. The growth rate of atmospheric N₂O has been constant at 0.78±0.01(3σ) parts per billion (ppb) per year over the period (see figure), but with important deviations related to El Niño Southern Oscillation, transport, and changes in patterns of emissions. Recent studies used a combination of multiple atmospheric networks and different Global Climate Models to calculate emissions, even down to emission values for individual countries and sources. Slight calibration differences between networks of a few tenths of a ppb can have significant effects on the emissions calculated by these methods. Our approach is to use one calibration scale for our flask and in situ networks within different Global Monitoring Division (GMD) groups with simple box models to examine the locations of the emissions.

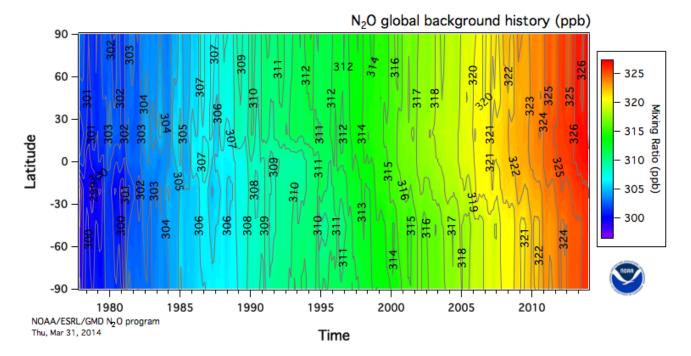


Figure 1. Global history of atmospheric N₂O (in ppb) from the NOAA GMD background observatories.