Geophysical Research Abstracts Vol. 13, EGU2011-6785, 2011 EGU General Assembly 2011 © Author(s) 2011



Global and regional emissions of HFC-143a (CH3CF3) and HFC-32 (CH2F2) from in situ and air archive atmospheric observations

Simon O'Doherty (1), Matt Rigby (2), Alistair Manning (3), Jens Mühle (4), Ben Miller (5), Ray Wang (6), Peter Simmonds (1), Dickon Young (1), Brian Greally (1), Stefan Reimann (7), and the S. O'Doherty Team (1) University of Bristol, Atmospheric Chemistry Research Group (ACRG), Chemistry, Bristol, United Kingdom (s.odoherty@bristol.ac.uk, 00441179252990), (2) Department of Earth, Atmospheric and Planetary Sciences, Massachusetts Institute of Technology, Cambridge, Massachusetts, USA., (3) Met Office, Exeter, UK, (4) Scripps Institution of Oceanography, University of California, San Diego, La Jolla, California, USA., (5) 5Global Monitoring Division, NOAA Earth System Research Laboratory, Boulder, Colorado and the Cooperative Institute for Research in Environmental Sciences, University of Colorado, Boulder, Colorado, USA., (6) School of Earth and Atmospheric Sciences, Georgia Institute of Technology, Atlanta, Georgia, USA., (7) Laboratory for Air Pollution and Environmental Technology, Swiss Federal Laboratories for Materials Testing and Research (Empa), Dübendorf, Switzerland., (8) Centre for Australian Weather and Climate Research, CSIRO Marine and Atmospheric Research, Aspendale, Victoria, Australia., (9) School of Earth and Environmental Sciences, Seoul National University, Seoul, South Korea., (10) Research Institute of Oceanography, Seoul National University, Seoul, South Korea.

High frequency, in situ observations from the Advanced Global Atmospheric Gases Experiment (AGAGE), and AGAGE associated stations, Gosan (South Korea), and Hateruma (Japan) for the period 2003 to 2010, combined with archive flask measurements dating back to 1978, have been used to capture the rapid growth of HFC-143a (CH3CF3) and HFC-32 (CH2F2) in the atmosphere. These are the first reported in situ global measurements of these two gases. HFC-143a is the third and HFC-32 is the sixth most abundant HFC. At the beginning of 2010 the global average for HFC-143a was 9.6 ppt in the lower troposphere and the growth rate was 13%/yr, and HFC-32 was 3.3 ppt with a growth rate was 23%/yr. The extensive observations have been combined with the AGAGE 2-D 12-box atmospheric transport model to simulate global atmospheric abundances and derive global emission estimates. It is estimated that \sim 18 Gg/yr of HFC-143a and \sim 11 Gg/yr of HFC-32 were emitted globally in 2009, and the emissions are estimated to have increased 15%/yr for HFC-143a and 26%/yr for HFC-32 since 2000. Observations of polluted air masses at selected individual AGAGE sites have been used to produce regional emission estimates for Europe (the EU15 countries), China, and Australia, respectively. Comparisons between these top-down estimates to bottom-up estimates based on reports by individual countries to the UNFCCC show a range of approximately four in the differences. This process of independent verification of emissions, and an understanding of the differences, is vital for assessing the effectiveness of international treaties, such as the Kyoto Protocol.