Geophysical Research Abstracts Vol. 21, EGU2019-7099-1, 2019 EGU General Assembly 2019 © Author(s) 2019. CC Attribution 4.0 license.



Processing of HCl in the Antarctic stratospheric polar vortex

Wuhu Feng (1,2), Martyn Chipperfield (1,2), Sandip Dhomse (2), Graham Mann (1,2), John Plane (3), Luke Arabham (4), and Michelle Santee (5)

(1) National Centre for Atmospheric Science, University of Leeds, Leeds, United Kingdom, (2) School of Earth and Environment, University of Leeds, Leeds, United Kingdom, (3) School of Chemistry, University of Leeds, Leeds, United Kingdom, (4) National Centre for Atmospheric Science, Department of Chemistry, University of Cambridge, Cambridge, United Kingdom, (5) Jet Propulsion Laboratory, California Institute of Technology, California, USA

Since the discovery of Antarctic ozone hole in 1985, significant progress has been made in understanding how different processes (e.g. dynamics, transport, microphysics, photochemistry, radiation, etc.) control the extent of the observed ozone depletion. It is well accepted that this depletion is primarily caused by efficient catalytic loss through chemical cycles containing the ClO and BrO radicals. Stratospheric halogens exist in this form through heterogeneous processing on polar stratospheric clouds, which converts the reservoir species HCl and ClONO₂ to active chlorine. Although the main processes that lead to ozone depletion are generally well understood, there remain open questions such as the duration of chlorine activation and its dependence on the abundance of nitrogen oxides (Kühl et al., 2004; Feng et al., 2011). It is still problematic to fully parameterise PSC-related processes in global models (SPARC, 2015). Furthermore, satellite observations (e.g., MLS) show that HCl is essentially completely depleted inside the dark, midwinter Antarctic polar vortex but most models significantly overestimate HCl in this region (e.g., SPARC 2010; Groo β et al., ACP, 2018).

Here we have used two state-of-the-art global models (the TOMCAT/SLIMCAT chemical-transport model (Chipperfield et al., 2018) and the UM-UKCA chemistry-climate model (Morgenstern et al., 2009)) to revisit the processing of the stratospheric HCl in the wintertime Antarctic polar vortex. The models are forced (or nudged) with ECMWF ERA-Interim and ERA5 reanalysis datasets and have detailed stratospheric chemistry. Both models have been run from 1980 to 2018. We compare the model results with MLS measurements and assess the extent to which the models capture the observed processing of HCl under different meteorological conditions. The year-to-year variation of polar ozone depletion will also be discussed, in particular for the recent years of decreasing stratospheric chlorine loading.

References:

Chipperfield, M.P., et al., On the Cause of Recent Variations in Lower Stratospheric Ozone, Geophys. Res. Lett., 45, 5718-5726, doi:10.1029/2018GL078071, 2018.

Feng, W., et al.,: Modelling the effect of denitrification on polar ozone depletion for Arctic winter 2004/2005, Atmos. Chem. Phys., 11, 6559-6573, doi:10.5194/acp-11-6559-2011, 2011.

 $Groo\beta$, J.-U., et al., On the discrepancy of HCl processing in the core of the wintertime polar vortices, Atmos. Chem. Phys., 18, 8647-8666, doi:10.5194/acp-18-8647-2018, 2018.

Kühl, S., et al.,: Stratospheric chlorine activation in the Arctic winters 1995/96–2001/02 derived from GOME OCIO measurements, Adv. Space Res., 34, 798–803, 2004.

Morgenstern, O., et al..: Evaluation of the new UKCA climate-composition model – Part 1: The stratosphere, Geosci. Model Dev., 2, 43-57, https://doi.org/10.5194/gmd-2-43-2009, 2009.

SPARC report: Chemistry-Climate Model validation, 2010.

SPARC report, https://www.sparc-climate.org/activities/polar-stratospheric-clouds/, 2015.