

4.060 Atmospheric Chemistry of E- and Z-CF₃CH=CHCF₃.

Early Career Scientist

Presenting Author:

Freja F. Oesterstroem, Copenhagen Center for Atmospheric Research, Department of Chemistry, University of Copenhagen, Universitetsparken 5, DK-2100 Copenhagen Ø, Denmark, freja.oesterstroem@gmail.com

Co-Authors:

Simone Thirstrup Andersen, Copenhagen Center for Atmospheric Research, Department of Chemistry, University of Copenhagen, Universitetsparken 5, DK-2100 Copenhagen Ø, Denmark

Ole John Nielsen, Copenhagen Center for Atmospheric Research, Department of Chemistry, University of Copenhagen, Universitetsparken 5, DK-2100 Copenhagen Ø, Denmark

Abstract:

FTIR smog chamber experiments were performed to investigate the atmospheric fate of E- and Z-CF₃CH=CHCF₃ (1,1,1,4,4,4-hexafluoro-2-butene). The experiments were performed to study reactions of E-CF₃CH=CHCF₃ or Z-CF₃CH=CHCF₃ with Cl atoms, OH radicals, and O₃ in 700 Torr of N₂/O₂ diluents at 296 ± 2 K. The study determined the Cl atom, OH radical, and O₃ kinetics and the mechanism of the atmospheric oxidation of E- and Z-CF₃CH=CHCF₃. The main atmospheric fate for both compounds is reaction with OH radicals. Atmospheric chemistry of the reaction of Z-CF₃CH=CHCF₃ with OH and OD radicals were investigated by Baasandorj et al. in 2011 [1], no other previous studies are available. The results of the present study are compared to the findings of Baasandorj et al. assessing the atmospheric impact of E- and Z-CF₃CH=CHCF₃. Infrared spectra were recorded, and the atmospheric lifetimes, the radiative forcings and global warming potentials (GWP) of the two butenes were calculated. This study provides a comprehensive description of the atmospheric fate of E- and Z-CF₃CH=CHCF₃. We present here the first results on the atmospheric chemistry of E-CF₃CH=CHCF₃ and of the Cl- and O₃-initiated chemistry of Z-CF₃CH=CHCF₃.

References

[1] M. Baasandorj, A.R. Ravishankara, J.B. Burkholder, Atmospheric Chemistry of (Z)-CF₃CH=CHCF₃: OH Radical Reaction Rate Coefficient and Global Warming Potential, The Journal of Physical Chemistry A 115 (2011) 10539-10549.