THE ROTATIONAL SPECTRUM OF CHLORINE NITRATE (CIONO₂) IN THE THREE LOWEST $n\nu_9$ POLYADS

Z. KISIEL, E. BIAŁKOWSKA-JAWORSKA, Institute of Physics, Polish Academy of Sciences, Al. Lotników 32/46, 02-668 Warszawa, Poland; R. A. H. BUTLER, Department of Physics, Pittsburg State University, Pittsburg, KS 66726; D. T. PETKIE, Department of Physics, Wright State University, Dayton, OH 45435; P. HELMINGER, Department of Physics, University of South Alabama, Mobile, AL 36688; F. C. DE LUCIA, Department of Physics, The Ohio State University, Columbus, OH 43210.

The rotational spectrum of the stratospherically important ClONO₂ molecule has recently been subjected to an extended analysis,^{*a*} which covered four states ranging from the ground state to the $\nu_5/\nu_6\nu_9$ dyad at ≈ 560 cm⁻¹. The analysis was based on a newly recorded broadband FASSST spectrum with an almost complete coverage of the 118-378 GHz region.

The remaining vibrational states below 560 cm⁻¹ are the three polyads associated with successively higher excitation of the ν_9 mode. These are the $(2\nu_9/\nu_7)$ dyad, the $(3\nu_9/\nu_7\nu_9)$ dyad, and the $(4\nu_9/\nu_72\nu_9/2\nu_7)$ triad. Rotational transitions in the triad were assigned for the first time and data sets for the two dyads were considerably improved over those obtained previously.^b Transition frequencies in all three polyads, for both ³⁵ClONO₂ and ³⁷ClONO₂, were fitted to within experimental accuracy by using a new coupling scheme which accounted in a unified manner for the pertinent Fermi and Coriolis interactions. Various tests used to ensure that the 'true' solution has been reached are discussed.

^aR.A.H.Butler et al., J. Mol. Spectrosc., 243, 1-9 (2007); 244, 113-116 (2007).

^bR.A.H.Butler et al., J. Mol. Spectrosc., 213, 8-14 (2002); 220, 150-152 (2003).