INFRARED SPECTROSCOPY OF TWO ISOMERS OF THE OCS-CS2 COMPLEX

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A second isomer of OCS-CS₂ complex, with a nonplanar cross-shaped structure, has been studied for the first time by analysing an infrared band in the region of the OCS stretching vibration (2062 cm⁻¹). This isomer has C_s symmetry and the observed band consists of purely c-type rotational transitions. The ground state rotational parameters are found to be A=0.07306 cm⁻¹, B=0.03325 cm⁻¹ and C=0.02879 cm⁻¹, in good agreement with a previous semi-empirical calculation.^{*a*} In addition, a hybrid band with a- and b-type rotational transitions has been assigned to be due to the known planar form of OCS-CS₂, as previously studied by microwave spectroscopy.^{*a*} The spectra were recorded using a rapid-scan tunable diode laser spectrometer to probe a pulsed supersonic jet expansion. Calculations indicate that the planar isomer of OCS-CS₂ is the lowest in energy, in contrast to CO₂-CS₂ where the crossed form is believed to be the lowest.^{*b*}

^aJ. J. Newby, M. M. Serafin, R. A. Peebles and S. A. Peebles, Phys. Chem. Chem. Phys. 7, 487 (2005).

^bC. C. Dutton, D. A. Dows, R. Eikey, S. Evans and R. A. Beaudet, J. Phys. Chem. A 102, 6904 (1998).