

## NEW INFRARED SPECTRA OF THE NITROUS OXIDE TRIMER

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Infrared spectra of N<sub>2</sub>O trimers are studied using a tunable diode laser to probe a pulsed supersonic slit-jet expansion. A previous observation by R.E. Miller and L. Pedersen [J. Chem. Phys. **108**, 436 (1998)] in the N<sub>2</sub>O  $\nu_1 + \nu_3$  combination band region ( 3480 cm<sup>-1</sup>) showed the trimer structure to be noncyclic, with three inequivalent N<sub>2</sub>O monomer units which could be thought of as an N<sub>2</sub>O dimer (slipped antiparallel configuration) plus a third monomer unit lying above the dimer plane. The present observations cover the N<sub>2</sub>O fundamental band regions  $\nu_3$  ( 1280 cm<sup>-1</sup>) and  $\nu_1$  ( 2230 cm<sup>-1</sup>). In the  $\nu_3$  region, two trimer bands are assigned with vibrational shifts and other characteristics similar to those in the  $\nu_1 + \nu_3$  region, but in the  $\nu_1$  region all three possible trimer bands are observed. Relationships among the various bands such as rotational intensity patterns, vibrational shifts, and the properties of the related N<sub>2</sub>O dimer, generally support the conclusions of Miller and Pedersen. Three trimer bands are also observed for the fully <sup>15</sup>N-substituted species in the  $\nu_1$  region, and these results should aid in detection of the as-yet-unobserved pure rotational microwave spectrum of the trimer. Finally, three combination bands involving the intermolecular van der Waals modes at 2253.7, 2255.5, and 2269.4 cm<sup>-1</sup> have been measured. The analyses of these bands and the identification of the nature of the intermolecular modes involved are currently underway.