

## TEMPERATURE DEPENDENCE OF THE INFRARED SPECTROSCOPY OF OZONE IN SOLID OXYGEN<sup>a</sup>

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Irradiation with ultraviolet laser light (210 to 250 nm) generates ozone monomers and dimers in a solid oxygen matrix. For wavelengths longer than 240 nm, we observe a dominant doublet structure for each fundamental vibration of ozone,  $\nu_1$ ,  $\nu_2$ , and  $\nu_3$ , as observed in a lamp irradiation study<sup>b</sup>. In addition, we identify a doublet feature for the combination band,  $\nu_1 + \nu_3$ <sup>c</sup>. With higher excitation energy we observe even more features, especially in the  $\nu_3$  region and attribute them to ozone dimers in the oxygen solid. While the aforementioned doublet structures, for  $\nu_3$  at  $1037.8\text{ cm}^{-1}$  and  $1030.9\text{ cm}^{-1}$ , have been assigned to an ozone monomer and to a complex  $\text{O}\cdots\text{O}_3$ , respectively<sup>a</sup>, our investigations indicate the latter feature is due to an ozone monomer isolated in an unstable lattice site. This conclusion results from detailed temperature studies of ozone-doped oxygen matrices. Furthermore, from the temperature dependence of the line position of the ozone monomer feature we observe the  $\alpha$ - $\beta$  phase transition of the solid oxygen host. The potential of this temperature shift as a novel temperature sensor for cryogenic matrices will be discussed.

<sup>a</sup>The support for this work by the Air Force Office of Scientific Research High Energy Density Matter Program is gratefully acknowledged.

<sup>b</sup>L. Schriver-Mazzuoli, A. de Saxcé, C. Lugez, and A. Schriver, *J. Chem. Phys.* **102**, 690 (1995).

<sup>c</sup>M. J. Dyer, C. G. Bressler, R. A. Copeland, *Chem. Phys. Lett.* **266**, 548 (1997).