## FAR INFRARED SYNCHROTRON SPECTRUM OF TRIMETHLYENE OXIDE

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Rotationally-resolved vibrational spectra of trimethlyene oxide (c-C<sub>3</sub>H6O) from 650 through 1200 cm<sup>-1</sup>were recorded using far infrared synchrotron radiation at the Canadian Light Source with better than 0.001 cm<sup>-1</sup>resolution. The observed bands correspond to at least eight different fundamental vibrations in this region. Due to the low frequency ring puckering motion,<sup>a</sup> the observed rovibrational pattern of each band is congested with hot-combination bands that originate in the first two excited ring puckering states (52.9 cm<sup>-1</sup>, 142.6 cm<sup>-1</sup>). The ongoing analysis of the strong *b-type* bands corresponding to asymmetric in-plane CO stretching ( $\nu_{10}$ :1008 cm<sup>-1</sup>)<sup>b</sup> will be discussed along with the identification of allowed Coriolis interactions arising from nearby energy levels related to in-plane CC stretching ( $\nu_{9}$ : 940 cm<sup>-1</sup>,  $\nu_{3}$ : 1033 cm<sup>-1</sup>).

<sup>&</sup>lt;sup>a</sup>G. Moruzzi et al., J. Mol. Spectrosc. **219**, **152** (**2003**).

<sup>&</sup>lt;sup>b</sup>Bánhegyi et al. Spectrochim. Acta. **39A, 761 (1983).**