The interesting low-frequency ring-pucker vibration in the far infrared region of trimethylene oxide (oxetane) has been the subject of previous, extensive investigations. Ueda and Shimanouchi\(^a\) found that by studying difference bands in the near-IR originating from C-H stretching and ring-puckering modes, values for the puckering energy levels could be obtained that agree well with the far-IR data. Recently, we have obtained a high resolution, diode laser, mid-IR spectrum of trimethylene oxide seeded in a pulsed-jet expansion with argon, (The rotational and vibrational temperatures are approximately 5 K and 50 K, respectively.) in the expectation that the effect of the ring mode on the puckering vibration could be determined. Preliminary analysis of the spectrum yields the following, previously undetermined upper-state rotational constants for this mode, \(A' = 0.4009 \text{ cm}^{-1}\), \(B' = 0.3909 \text{ cm}^{-1}\), and \(C' = 0.2204 \text{ cm}^{-1}\). The band origin \(v_0\) is determined to be 908.427 \text{ cm}^{-1}, which represents greater accuracy and precision than the previously available value of 908.7 \text{ cm}^{-1}.\(^b\)