DYNAMICS OF LINEAR AND T-SHAPED ArI₂ DISSOCIATION UPON B→X EXCITATION: A DISPERSED INTRACAVITY LASER-INDUCED FLUORESCENCE STUDY

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We have recorded the dispersed fluorescence spectra of the linear and the previously well-studied T-shaped isomeric ArI₂ complexes following B→X excitation. The dispersed fluorescence features originate from the free I₂ B→X emission and reveal the I₂ product vibrational state distribution. The vibrational distribution of photofragment I₂(B) produced from dissociation of the T-shaped isomer shows a sharp Poisson distribution, beginning with \( \Delta v' = -3 \), which is nearly independent of excitation energy. The vibrational distribution of I₂(B) from the linear isomer is quite different. For excitation energies greater than that corresponding to I₂(B,v''=22), the product distribution is quite similar to that observed for excitation above the dissociation energy of the B state, i.e., it shows a broad, nearly gaussian distribution of I₂(B) vibrational states, well below the excitation energy. For excitation energies below that, the distribution becomes increasingly like the Poisson distribution observed for the T-shaped isomer. Bond dissociation energies for linear and T-shaped isomers are determined for the spectra. The dissociation time scale for the two isomers differs by about 3 orders of magnitude. To model the product state distribution of the linear isomer and therefore gain further insight into the Ar-I₂ interaction potential, a simple trajectory calculation based upon the classical dynamics of a compressed oscillator (I−I) ejecting a freely attached particle (Ar) is in progress. Results from the calculation are in good accord with the experiment.