

PHOTOELECTRON SPECTROSCOPY OF ICN^- : CHARACTERIZATION OF A CONICAL INTERSECTION IN ICN

ELISA M. MILLER, LEONID SHEPS,^a YU-JU LU, *JILA, Department of Chemistry and Biochemistry, University of Colorado at Boulder, Boulder, CO 80309*; ANNE B. McCOY, *Department of Chemistry, The Ohio State University, Columbus, OH, 43210*; and W. CARL LINEBERGER, *JILA, Department of Chemistry and Biochemistry, University of Colorado at Boulder, Boulder, CO 80309*.

We report the photoelectron spectrum of ICN^- to probe transitions to the ground state ($X^1\Sigma^+$) and first 5 excited states ($^3\Pi_2$, $^3\Pi_1$, $^3\Pi_0-$, $^3\Pi_0+$, and $^1\Pi_1$) of neutral ICN . We spectroscopically resolve the first 3 excited states and a conical intersection region between the $^3\Pi_0+$ and $^1\Pi_1$ states for the first time. The spectra are assigned with the aid of previously published high-level calculations by Morokuma and coauthors^b. Our assignments are further verified by comparison to the photoelectron spectra of the dihalides I_2^- and IBr^- . The poor Franck-Condon overlap between the ground states of the anion and neutral precludes direct observation of the adiabatic electron affinity, $\text{EA}(\text{ICN})$. However, through thermochemical cycles involving narrow transitions to excited states, we determine the $\text{EA}(\text{ICN})$ to be 1.7 ± 0.1 eV and the dissociation energy, $D_0(\text{ICN}^-)$, to be 0.9 ± 0.1 eV. To our knowledge, the $\text{EA}(\text{ICN})$ has not been previously reported in experiment or theory; therefore, this is the first $\text{EA}(\text{ICN})$ determination. In addition, we observe at least four spectral peaks with kinetic energies of ≤ 5 , 45, 70, and 160 meV that are independent of the photon energy over the 2.6 - 4.1 eV energy range. It appears that these peaks are related to autodetachment from excited anions to the ground electronic state of the neutral. We use a combination of previous calculations by Morokuma et al. and two-dimensional SO-MR-CI scans with a fixed CN distance of the anion potential energy surfaces to aid in an autodetachment mechanism. Support from NSF and AFOSR is gratefully acknowledged.

^aPresent address: Sandia National Laboratories, Livermore, CA 94551

^bS. Yabushita and K. Morokuma, *Chem. Phys. Lett.* 175, 518 (1990); Y. Amatatsu, S. Yabushita, and K. Morokuma, *J. Chem. Phys.* 100, 4894 (1994).