Electronic predissociation of CN($\text{A}^3\Pi$)-H$_2$/D$_2$ was examined by characterizing action spectra and product state distributions. Both spin-orbit (CN($\text{A}^3\Pi_{1/2}$)-H$_2$→CN($\text{A}^3\Pi_{3/2}$)+H$_2$) and internal conversion (CN($\text{A}^3\Pi_{3/2}$)-H$_2$→CN($\text{X}^2\Sigma^+$)+H$_2$) decay channels were observed. For comparison with the predissociation data, CN(A)+H$_2$ collisional energy transfer was examined at low temperatures (near 10 K). The product state distributions resulting from collisions showed symmetry propensities that were not evident in the distributions resulting from predissociation. Furthermore, the symmetry propensity exhibited by spin-orbit transfer was the opposite of that seen in A→X transfer. We argue that the lack of symmetry preference in the predissociation dynamics is indicative of resonant scattering processes. The results may be explained by considering the symmetry properties of the intermolecular potential energy surfaces, and the regions of these surfaces that are sampled by predissociation and collisional transfer events.

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