EXPERIMENTAL CHARACTERIZATION OF THE He + $I^{35}Cl(E,v^{\dagger}=11,12)$ AND He + $I^{35}Cl(\beta,v^{\dagger}=0-2)$ INTER-MOLECULAR POTENTIAL ENERGY SURFACES

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Two-photon excitation of the T-shaped and linear He \cdots I³⁵Cl($X^{1}\Sigma^{+}, \nu''=0$) complexes are used to access multiple intermolecular vibrational levels within the He + I³⁵Cl($E0^{+3}P_2, \nu^{\dagger}=11,12$) and He + I³⁵Cl($\beta 1^{3}P_2, \nu^{\dagger}=0-2$) ion-pair state intermolecular potentials. The excitation utilizes different metastable intermolecular levels within the He + I³⁵Cl($B^{3}\Pi_{0+}, \nu'=2,3$) and He + I³⁵Cl($A^{3}\Pi_{1}, \nu'=15$) potentials, and thus varying Franck-Condon windows, to access levels with varying amounts of vibrational excitation within the ion-pair state intermolecular potential. The He + ICl($E, \nu^{\dagger}=11,12$) and He + ICl($\beta, \nu^{\dagger}=0-2$) intermolecular potentials are found to be nearly identical with an overall minimum in the near T-shaped orientation and binding energies $D_{0}^{\dagger}=40 \text{ cm}^{-1}$. Intermolecular stretching and bending frequencies are measured to be 25 and 13 cm⁻¹, respectively. Since common intermolecular levels are accessed by transitions from both the T-shaped and linear He \cdots ICl($X, \nu''=0$) ground state conformers, the relative binding energies of the conformers can be directly measured; the linear conformer is found to be 5.4(2) cm⁻¹ more strongly bound than the T-shaped conformer.