

HIGH-RESOLUTION SPECTROSCOPY OF MgKr^+ IN ITS GROUND AND LOW-LYING ELECTRONICALLY EXCITED STATES

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Diatomic molecules RgM consisting of a rare-gas atom Rg and an alkaline-earth-metal atom M and their singly and doubly-charged cations RgM^+ and RgM^{2+} have unusual chemical properties that are related to the low first and second ionization energies of M and the high ionization energy of Rg . In MgAr the second ionization energy of Mg is lower than the first ionization energy of Ar . Consequently, MgAr^{2+} is thermodynamically stable and Rydberg series of MgAr^+ can be observed that converge on the $X^{2+} \ ^1\Sigma^+$ ground state of MgAr^{2+} .^a In this contribution, we present the results of spectroscopic investigations of MgKr^+ in its ground and low-lying electronically excited states that complement earlier studies of this cation.^{b,c} Pulsed-field-ionization zero-kinetic-energy (PFI-ZEKE) photoelectron spectra of the $X^+ \ ^2\Sigma^+$ ground state of MgKr^+ were recorded following single-photon excitation from the $a \ ^3\Pi_0$ metastable state of MgKr . Vibrational channel interactions enabled the observation of the lowest vibrational levels of MgKr^+ and the determination of an accurate value of the adiabatic ionization energy of metastable MgKr ($38183 \pm 2 \text{cm}^{-1}$). Using isolated-core multiphoton Rydberg dissociation (ICMRD) spectroscopy,^d spectra of several low-lying electronically excited states of MgKr^+ were observed that are associated with the $\text{Kr} + \text{Mg}^+(nl)$ dissociation limits with $n = 3, 4$ and $l = s, p$ and d . These states may be regarded as the lowest members of Rydberg series converging on the ground state of MgKr^{2+} . These studies represent first steps towards studying the doubly charged cation MgKr^{2+} .

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