

INFRARED RESONANCE ENHANCED PHOTODISSOCIATION SPECTROSCOPY OF  $\text{Si}^+(\text{CO}_2)_n$ ,  $\text{Si}^+(\text{CO}_2)_n\text{Ar}$ , AND  $\text{Si}^+(\text{O}_2)(\text{CO}_2)_n$  CLUSTERS

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Weakly bound  $\text{Si}^+(\text{CO}_2)_n$ ,  $\text{Si}^+(\text{CO}_2)_n\text{Ar}$ , and  $\text{Si}^+(\text{O}_2)(\text{CO}_2)_n$  complexes are produced by laser vaporization in a pulsed supersonic expansion. The ions are then mass selected in a reflectron time-of-flight mass spectrometer where they are photodissociated by a tunable infrared OPO/OPA laser system near the asymmetric stretch ( $\nu_3$ ) of  $\text{CO}_2$ . Dissociation is more efficient on resonance, thus monitoring the fragmentation channel as a function of infrared laser frequency produces the IR absorption spectrum for the complex of interest.  $\text{Si}^+(\text{CO}_2)_n$  and  $\text{Si}^+(\text{O}_2)(\text{CO}_2)_n$  clusters fragment by the loss of whole  $\text{CO}_2$  units while  $\text{Si}^+(\text{CO}_2)_n\text{Ar}$  complexes fragment by the loss of argon. Interpretation of the observed infrared spectra will be discussed.