PROBING THE MOLECULAR DYNAMICS OF A $\text{Cu}(\text{CD}_3\text{OD})$ CLUSTER WITH PHOTODETACHMENT-PHOTOIONIZATION SPECTROSCOPY

<u>J. BARBERA</u>, V. DRIBINSKI, and W.C. LINEBERGER, *JILA/Department of Chemistry, University of Colorado, Boulder, CO 80309*; S. HORVATH, and A.B. McCOY, *Department of Chemistry, Ohio State University, Columbus, OH 43210*.

We report the femtosecond nuclear dynamics of neutral $Cu(CD_3OD)$ van der Waals clusters, investigated using photodetachment-photoionization spectroscopy. Photodetachment of $Cu(CD_3OD)$ anion with a 150 fs 400 nm laser pulse produces a vibrationally excited neutral complex that undergoes ligand reorientation and dissociation. The time evolving neutral is interrogated by delayed femtosecond resonant two photon ionization. This study shows that the nascent $Cu(CD_3OD)$ complex dissociates on prompt (3 ps) and slower timescales (30 ps). The prompt component reflects direct dissociation upon photodetachment, while the slower dissociation arises from the coupling of CD_3OD molecular rotation into the Cu-(CD_3OD) dissociation coordinate. Theoretical investigations provide insight to the nature of the molecular dynamics which produce the observed dissociation characteristics. Supported by NSF and AFOSR