

Abstract Submitted  
for the DAMOP06 Meeting of  
The American Physical Society

**Dissociation and ionization of  $\text{HD}^+$  in intense few-cycle laser pulses**<sup>1</sup> P. Q. WANG, A. M. SAYLER, V. ROUDNEV, B. GAIRE, NORA G. JOHNSON, M. LEONARD, E. PARKE, K.D. CARNES, B. D. ESRY, I. BEN-ITZHAK, J. R. Macdonald Laboratory, Physics Department, Kansas State University, Manhattan KS 66506 — All the dissociation and ionization channels of an  $\text{HD}^+$  molecular ion beam exposed to intense laser fields, namely  $\text{H}^+ + \text{D}$ ,  $\text{D}^+ + \text{H}$  and  $\text{H}^+ + \text{D}^+$ , have been studied simultaneously using a 3D coincidence momentum imaging method. These breakup channels are experimentally separated from each other. The laser pulse durations are from about 8 fs up to those comparable to the dissociation time scale of the molecule, with a wavelength of 790 nm and an intensity range of  $10^{13}$ - $10^{15}$  W/cm<sup>2</sup>. The focus of this study is on the phenomena in few-cycle laser pulses. The 3D momentum of each fragment is retrieved from its position and time signals, which provides angular and kinetic energy release spectra for each breakup channel. The dissociation of  $\text{HD}^+$  is found to be governed by bond-softening and above threshold dissociation, depending on the laser intensity. The ionization of  $\text{HD}^+$  is strongly aligned along the laser polarization and has a broad kinetic energy distribution which shifts to higher values at higher intensity.

<sup>1</sup>Supported by the Chemical Sciences, Geosciences and Biosciences Division, Office of Basic Energy Sciences, Office of Science, U.S. Department of Energy.

P. Q. Wang

Date submitted: 27 Jan 2006

Electronic form version 1.4