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**Probing Time-Dependent Electron Interactions in Double-Rydberg Wavepackets** X. ZHANG, R.R. JONES, Department of Physics, University of Virginia — Subpicosecond half-cycle electric field pulses (HCPs) have been used as a time-resolved probe of interactions between electrons in a 3-body Coulomb system. We produce double-Rydberg wavepackets (DRWs) using a multi-step, ultra-fast, isolated core excitation of ground-state barium atoms. Two laser pulses first create a 5dnd radial wavepacket. At some time within the first Kepler period of the nd Rydberg electron's motion, additional laser pulses excite the 5d core-electron into a superposition of  $Ba^+ Ng$  Rydberg states. Initially, the two electrons in the  $Ngnd$  DRW have well defined energies, momenta, and radial positions as they move in singly and doubly-charged Coulomb potentials, respectively. However, the electrons soon become highly correlated as they exchange energy and angular momentum. If left unaltered, the atom eventually autoionizes, creating a free electron and an  $N'L$  ionic wavepacket ( $N' < N$ ). We probe the evolution of the DRW using a HCP that has sufficient strength to impulsively ionize both the initial  $Ng$  (ionic) and nd (neutral) wavepackets individually, but insufficient amplitude to efficiently ionize the  $N'L$  ions that result from autoionization. We monitor the production of  $Ba^{++}$  ions as a function of the delay between the HCP and the launch of the  $Ng$  ionic wavepacket to probe the time-dependent energy-exchange between the two electrons. Classical trajectory Monte Carlo simulations aid in the interpretation of our experimental results. This work has been supported by the NSF.

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