

Abstract Submitted
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Preparation of Vibrationally Excited H₂ in a Coherent Superposition of M -States Using Stark Induced Adiabatic Raman Passage (SARP)¹
NANDINI MUKHERJEE, WENRUI DONG, WILLIAM PERREAULT, RICHARD ZARE, Stanford Univ — We prepare a large ensemble of rovibrationally excited ($v = 1, J = 2$) H₂ molecules in a coherent superposition of M -states using Stark-induced adiabatic Raman passage (SARP) with linearly polarized single mode pump (532 nm) and Stokes (699 nm) laser pulses of duration 6 ns and 4 ns. A biaxial superposition state, $|\psi\rangle = 1/\sqrt{2} [|v = 1, J = 2, M = -2\rangle - |v = 1, J = 2, M = +2\rangle]$, is prepared using SARP with a sequence of a pump laser pulse partially overlapping with a cross polarized Stokes laser pulse co-propagating along the quantization z -axis. The degree of phase coherence is measured by recording interference fringes in the ion signal produced using the O(2) line of 2+1 resonance enhanced multiphoton ionization (REMPI) from the rovibrationally excited ($v = 1, J = 2$) level as a function of REMPI laser polarization angle. The ion signal is measured using a time-of-flight mass spectrometer. Nearly 60% population transfer from H₂ ($v = 0, J = 0$) ground state to the superposition state in H₂ ($v = 1, J = 2$) is measured from the depletion of Q(0) REMPI signal of the ($v = 0, J = 0$) ground state. The M -state superposition behaves much like a multi-slit interferometer where the number of slits, i.e. the number of M -states, and their separations, i.e. the relative phase, can be varied experimentally.

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