

Abstract Submitted
for the DAMOP13 Meeting of
The American Physical Society

**Angle-Resolved Strong Field Ionization of Polyatomic Molecules:
More than the Orbitals Matters** OUMAROU NJOYA, SUNY Stony Brook,
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Stony Brook — We compare the time and angle-dependent strong field ionization
yields of three molecules with very similar electronic structure. A pump pulse in
the deep ultraviolet excites the molecules to their first bright excited state (HOMO-
LUMO excitation). We then measure the strong field ionization yield due to a near
infrared probe pulse as a function of delay and angle between pump and probe po-
larization vectors. *ab initio* electronic structure calculations allow us to associate
the parent ion yields with removal of an electron from the LUMO orbital (occupied
after excitation by the pump). Despite the fact that the orbitals are very similar,
the yields are very different, indicating that it is not the orbital shape alone which
determines angle-dependent ionization yields.

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Date submitted: 24 Jan 2013

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