

Donnerstag, 18.10.2018

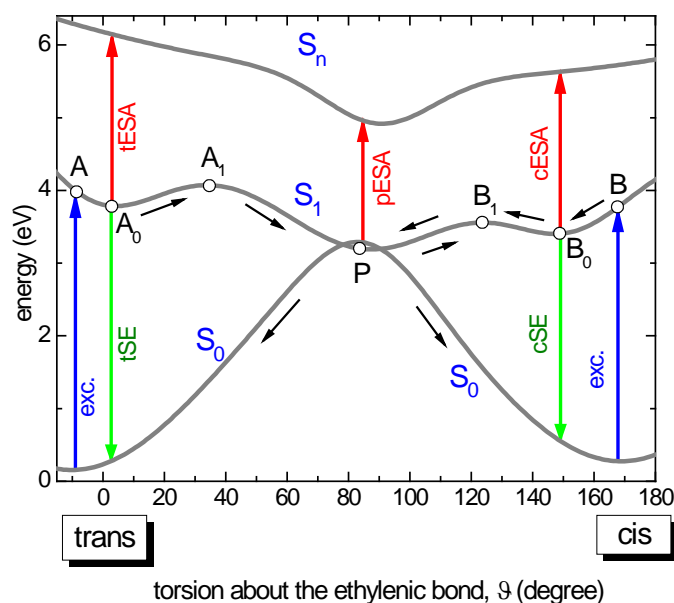
Hörsaal C, Chemie Zentralbau, 17:15 Uhr

**Sprecher:** **Nikolaus Ernsting**  
*Humboldt-Universität zu Berlin*

**Thema:** **New views through old holes:  
looking for the perpendicular  
“phantom” state of  
photoisomerizing stilbene**

**Abstract:**

In the photoisomerization path of stilbene, a perpendicular state P on the  $S_1$  potential energy surface is expected just before internal conversion through a conical intersection  $S_1/S_0$ . The situation is depicted schematically below:



For decades the observation of P was impossible because of a short lifetime  $\tau_P$  in combination with slow population flow over a barrier ( $A_1$  or  $B_1$  in the figure). But these limitations can be overcome by ethylenic substitution.

Comparing stilbene and 1,2-dimethyl-stilbene with precision transient absorption spectroscopy in the range 250-800 nm, we determine the electronic properties of P.

By substitutions – also at the phenyl moieties - the excited *trans*, *cis*, and *perpendicular* forms are stabilized on a >10 ps timescale. Their excited-state absorption bands (ESA in the figure) are used for resonance in fs stimulated Raman spectroscopy. In this way we find several new phenomena during the photoreaction.

**Organisation: C. Lambert**

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