

1D05

## Trial to Observe Wavepacket Motions near the Potential “Funnel” in Stilbene Photoisomerization by Pump-dump-probe Spectroscopy

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### [Introduction]

In ultrafast reactions, it is considered that the excited-state molecule relaxes rapidly to ground state through a potential “funnel”, where the excited- and ground-state potential energy surfaces (PES) get close to each other. This potential region is expected to provide an efficient channel for rapid relaxation and to play a key role in ultrafast reactions. However, our knowledge is very limited on how the molecular structure (or nuclear wavepacket) evolves in this important potential region. Therefore, it is highly desirable to obtain quantitative information about the wavepacket motion near the funnel region. With this fundamental interest in mind, we tried to observe the wavepacket motion approaching the funnel region in a model system, photoisomerization of cis-stilbene, using pump-IR dump-probe spectroscopy.

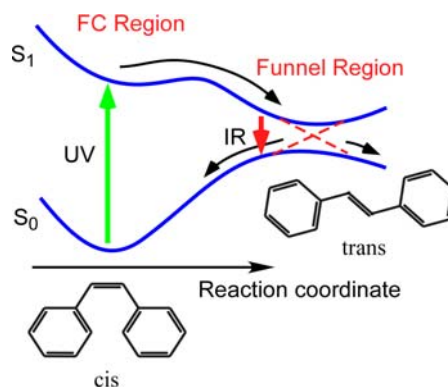


Fig. 1 Schematic illustration of the potential funnel region in cis-stilbene photoisomerization.

### [Experiment]

The output of a Ti: sapphire amplifier was divided into three beams for generating the pump, dump and probe pulses, respectively. The pump pulse at 266 nm was produced through 3<sup>rd</sup>-harmonic generation of the 800 nm fundamental pulse. The dump pulse at various infrared wavelengths was generated by a commercial OPA system pumped by the 800 nm pulse. The white continuum probe pulse was obtained by focusing the fundamental pulse on a sapphire plate.

### [Results and Discussion]

As shown in Figure 1, it is suggested that the photogenerated S<sub>1</sub> state of cis-stilbene initially undergoes a twisting deformation and forms the relaxed S<sub>1</sub> state, corresponding to a shallow

local minimum of the  $S_1$  potential. This relaxed  $S_1$  state, which exhibits a strong  $S_n \leftarrow S_1$  absorption at 640 nm, is a precursor of the isomerization and is converted to the  $S_0$  state (trans and cis) in a picosecond through the funnel region. To investigate the wavepacket motion that moves toward the funnel region, we introduced an infrared dump pulse at 0.5 ps after photoexcitation and monitored the transient absorption by the probe at 1 ps. As shown in figure 2, the  $S_n \leftarrow S_1$  transient absorption at 640 nm is decreased significantly by the dump pulse at 1.5  $\mu\text{m}$  ( $6500\text{ cm}^{-1}$ ). This result strongly indicates that the  $S_1$  population is depleted by the dump pulse, implying that the  $S_1$  population is resonantly driven back to the  $S_0$  state by the stimulated emission dumping. We evaluated the following dump efficiency,

$$\text{dump efficiency} = 1 - \frac{\Delta\text{Abs}(\text{dump-on})}{\Delta\text{Abs}(\text{dump-off})}, \quad (1)$$

and plotted it against the pump-dump delay. As shows in figure 3 (b), the dump efficiency shows almost instantaneous rise as short as 11 fs, which was evaluated by an analysis taking account of the instrumental response (FWHM=133 fs). The rise time of the dump efficiency corresponds to the time that the wavepacket needs to reach the potential region, where the  $S_1$ - $S_0$  energy difference matches the  $6500\text{-cm}^{-1}$  dump photon energy. To monitor the wavepacket motion at different PES regions, we carried out experiments with changing the dump photon energy, as shown in figure 3. It was found that the rise is almost identical in the range of 10-20 fs for the dump photon energy from  $8500\text{ cm}^{-1}$  to  $3500\text{ cm}^{-1}$ , while it is substantially longer (76 fs) for the  $2470\text{ cm}^{-1}$  dump pulse (we note that the temporal response of the experiment does not change significantly with change of the dumping wavelengths).

The pump-dump-probe result clearly indicates the wavepacket motion toward the PES region, where the  $S_0$  and  $S_1$  potentials get as close to as  $2470\text{ cm}^{-1}$ , and provides its quantitative characterization near the funnel region.

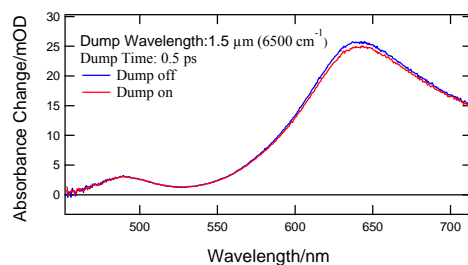


Fig. 2 The transient absorption spectrum of cis-stilbene with/without dump pulse (pump-probe delay time: 1 ps; pump-dump delay time: 0.5 ps).

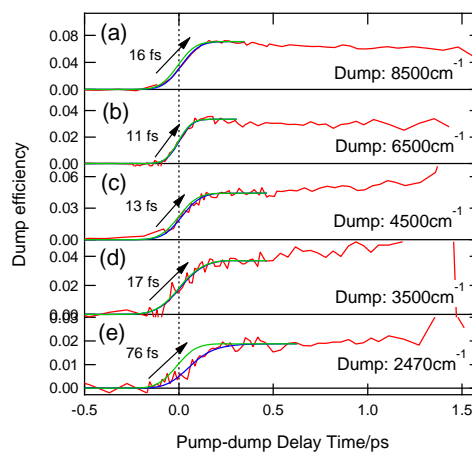


Fig. 3 Dump efficiencies as a function of pump-dump delay time for various dump photon energy.