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Non-equilibrium dynamics driven by 1.5-cycle strong light field in strongly-correlated organic conductor

Optical responses of strongly correlated organic compounds has attracted much attention, because they exhibit ultrafast changes in the conducting and/or dielectric properties upon photo-excitations [1-4]. Recent progress in several fs optical pulse enables us to drive various new aspects of the strongly correlated system [5-7]. Dynamical localization[8,9], i.e., reduction of the intersite electronic transfer integral t by an AC electric field, $E(\omega)$, is a promising strategy for controlling strongly correlated systems with a competing energy balance between t and the Coulomb repulsion energy[10]. Here we describe a charge localization induced by the 9.3 MV/cm instantaneous electric field of a 1.5 cycle (7 fs) infrared pulse in an organic conductor α -(ET)2I3 [7]. A large reflectivity change of $> 25\%$ and a coherent charge oscillation along the time axis reflect the opening of the charge ordering gap in the metallic phase. This optical freezing of charges, which is the reverse of the photoinduced melting of electronic orders, is attributed to the $\sim 10\%$ reduction of t driven by the strong, high-frequency ($\omega \sim t/\hbar$) electric field.

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