

Probing water vapor-induced modulation of charge carrier dynamics in mesoporous TiO₂ using ultrafast transient absorption spectroscopy

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Área temática: C. Propiedades de nanomateriales

Mesoporous titanium dioxide (TiO₂) thin films exhibit unique properties due to their high surface area and porous structure, making them highly sensitive to environmental conditions. In this work, we investigate the influence of water vapor on the ultrafast charge carrier dynamics of mesoporous TiO₂ films using transient absorption spectroscopy in the femtosecond–picosecond regime. Measurements were performed at an excitation wavelength of 325 nm under controlled relative humidity (RH) conditions ranging from low (~5–6%) to near-saturation (~94–95%). The transient absorption response shows a pronounced dependence on RH, with clear modifications in both spectral features and decay kinetics as humidity increases. These changes indicate that water adsorption within the mesoporous network strongly affects charge carrier lifetimes and recombination pathways. The observed behavior is attributed to the interaction of water molecules with the TiO₂ surface, which can introduce or modify surface trap states, influence charge separation, and alter recombination dynamics. The high internal surface area of mesoporous films enhances these effects, leading to a marked sensitivity to humidity variations. These results demonstrate that water vapor is a key factor governing the charge carrier dynamics in mesoporous TiO₂, highlighting the importance of conducting characterization under operando conditions in applications such as photoelectrocatalysis.