Injected Electrons Populate an Electronic State at the Interface of the SnO₂ Core and TiO₂ Shell

Absorbance

Scientific Achievement

- Spectro-electrochemical analysis showed electrons injected into core/shell SnO₂/TiO₂ mesoporous thin films were not present in the SnO_2 core or the TiO₂ shell.
- An electronic state at the interface between the core and shell was proposed with an energetic position 200 mV more positive than either SnO_2 or TiO_2 .

Significance and Impact

Control of the energetic position and physical location of injected electrons enables control of unwanted charge recombination reactions and catalysis.

Research Details

- About 2.5 nm thick rutile TiO₂ shells were formed on 15 nm diameter SnO₂ nanocrystals present in a mesoporous thin films via atomic layer deposition, followed by sintering.
- Reduction of TiO₂ and SnO₂/TiO₂ films resulted in the appearance of one unique absorption spectra consistent with the formation of one electronic state.
- Standard addition of the SnO₂ and TiO₂ absorption ٠ spectra did not simulate the SnO₂/TiO₂ data, indicating that the core/shell has an electronic state that is distinct from SnO₂ and TiO₂.
- Core/shell SnO₂/TiO₂ thin films have a higher capacitance than SnO_2 , but comparable to TiO_2 .

James, E. M.; Barr, T. J.; Meyer, G. J. ACS Appl. Energy Mater. 2018, ASAP, DOI: 10.1021/acsaem.7b00274.

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Α. Β. 2.5 nm SnO₂ TiO₂ 10 nm **C.** 0.02 D. -0.6 0.00 SnO₂/TiO SnO₂/TiO₂ TiO. TiO₂ SnO. SnO₂ at 50e⁻/particle 0.2 -0.0 40 30 20 500 400 600 700 800 900 1000 Wavelength (nm) Ε. TEM image of a SnO₂/TiO₂ film. 0.2 Scheme showing idealized conduction B) band (CB) offsets and electrons 0.1 Potential (V vs NHE) localized at the core/shell interface. 0.0 UV-Vis absorption spectra of the thin C) -0.1 films with ~50 electrons present in each nanocrystallite. -0.2 Density of states of the oxide D) -0.3 semiconductors. Absorbance: 0.1

Reduction potential onset vs. pH with E) an overlaid linear fit.

Work was performed at University of North Carolina at Chapel Hill.





The University of Texas



