

121.

Accession number:20112714115306

Title:Electron mobility and injection dynamics in mesoporous ZnO, SnO<sub>2</sub>, and TiO<sub>2</sub> films used in dye-sensitized solar cells

Authors:Tiwana, Priti (1); Docampo, Pablo (1); Johnston, Michael B. (1); Snaith, Henry J. (1); Herz, Laura M. (1)

Author affiliation:(1) Department of Physics, Clarendon Laboratory, University of Oxford, Parks Road, Oxford, OX1 3PU, United Kingdom

Corresponding author:Snaith, H.J.(h.snaith@physics.ox.ac.uk)

Source title:ACS Nano

Abbreviated source title:ACS Nano

Volume:5

Issue:6

Issue date:June 28, 2011

Publication year:2011

Pages:5158-5166

Language:English

ISSN:19360851

E-ISSN:1936086X

Document type:Journal article (JA)

Publisher:American Chemical Society, 2540 Olentangy River Road, P.O. Box 3337, Columbus, OH 43210-3337, United States

Abstract:High-performance dye-sensitized solar cells are usually fabricated using nanostructured TiO<sub>2</sub> as a thin-film electron-collecting material. However, alternative metal-oxides are currently being explored that may offer advantages through ease of processing, higher electron mobility, or interface band energetics. We present here a comparative study of electron mobility and injection dynamics in thin films of TiO<sub>2</sub>, ZnO, and SnO<sub>2</sub> nanoparticles sensitized with Z907 ruthenium dye. Using time-resolved terahertz photoconductivity measurements, we show that, for ZnO and SnO<sub>2</sub> nanoporous films, electron injection from the sensitizer has substantial slow components lasting over tens to hundreds of picoseconds, while for TiO<sub>2</sub>, the process is predominantly concluded within a few picoseconds. These results correlate well with the overall electron injection efficiencies we determine from photovoltaic cells fabricated from identical nanoporous films, suggesting that such slow components limit the overall photocurrent generated by the solar cell. We conclude that these injection dynamics are not substantially influenced by bulk energy level offsets but rather by the local environment of the dye-nanoparticle interface that is governed by dye binding modes and densities of states available for injection, both of which may vary from site to site. In addition, we have extracted the electron mobility in the three nanoporous metal-oxide films at early time after excitation from terahertz conductivity measurements and compared these with the time-averaged, long-range mobility determined for devices based on identical films. Comparison with established values for single-crystal Hall mobilities of the three materials shows that, while electron mobility values for nanoporous TiO<sub>2</sub> films are approaching theoretical maximum values, both early time, short distance and interparticle electron mobility in nanoporous ZnO or SnO<sub>2</sub> films offer considerable scope for improvement. &copy; 2011 American Chemical Society.

