

Electron kinetics study of dye-sensitized solar cells made from MgO coated SnO₂ nanocrystalline films

V P Susira Perera*, L S G Liyanage, R Chinthaka L De Silva
Department of Physics, The Open University of Sri Lanka, Nawala, Nugegoda.

The studies on Dye-sensitized nanocrystalline photovoltaic solar cells (DSNCs) widen the horizons of the low cost alternative photovoltaics and arouse the fundamental research interests of the investigators. A DSNC fabricated by SnO₂ crystallites coated with ultra thin layer of MgO (MgO/SnO₂) has been reported to deliver a high open circuit photovoltage comparable to the DSNCs fabricated only with SnO₂ crystallites. The photo excited dye molecules adsorb on the MgO outer layer suppose to inject electrons into the conduction band of SnO₂ via tunneling through the MgO barrier. Suppression of recombination of electrons in the conduction band with the dye cations and the acceptors at the electrolytic interface build up the quasi-fermi level of SnO₂ resulting increase in the photovoltage. The competition between the back ward reactions and the collection of electrons at the substrate is determine by the electron diffusion length $L_n=(D_n \tau_n)^{1/2}$ where D_n is the electron diffusion coefficient and τ_n is the carrier lifetime. Though the increment of photovoltage and the efficiency is interpreted as a consequence of the blocking layer, no previous research has been done to study the electron kinetics of this cell. We have conducted some studies to analyze D_n and τ_n of the electrons injected to MgO/SnO₂ DSNC using intensity modulated photocurrent (IMPS) and intensity modulated photo voltage (IMVS) spectroscopy.

MgO/SnO₂ film was deposited on a conducting tin oxide (CTO) glass by the following method. 7 mg of MgO was grinded in a mortar with few drops of acetic acid and 2 ml of Tin oxide Colloid. This solution was diluted with ethyl alcohol and spray onto the CTO glass at 120 °C. After depositing the film it was sintered in a furnace at 500 °C for 30 minutes. These films were kept immersed in a ruthenium bipyridyl dye in alcohol for one hour. The cell was fabricated by placing a CTO glass coated with Pt on the film and filling the capillary space with an electrolyte containing I⁻/I³⁻ redox couple. A similar procedure is followed to construct DSNCs of bare SnO₂, except the addition of MgO in the preparation. Intensity modulated photovoltage spectroscopy (IMVS) and Intensity modulated photocurrent spectroscopy (IMPS) measurements were carried out using a blue light emitting diode ($\lambda_{\text{max}}=470\text{nm}$) driven by a frequency response analyzer. The LED provided both the dc and ac components of the illumination. D_n and τ_n were calculated from the minima of the IMVS and IMPS curves respectively at different intensities.

The lifetime of electrons decreased with the increment of light intensity for both SnO₂ and SnO₂/MgO cells but the lifetime of the electrons in SnO₂/MgO film is always higher compared to bare SnO₂ cell. The diffusion coefficient increases as the light intensity increases where the diffusion coefficient is also higher at any particular light intensity for SnO₂/MgO cell. Since both the D_n and τ_n are greater, diffusion length of electrons in SnO₂/MgO cells is large. Nevertheless electrons transport efficiently and recombination losses are minimal in the DSNC constructed with MgO coated SnO₂ film that claim for the high photovoltage and efficiency. This is a different approach to interpret the enhancement of high voltage and efficiency previously reported on the SnO₂/MgO dye-sensitized solar cells.

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