

Supplementary Information.

Kinetic competition in liquid electrolyte and solid state Cyanine Dye Sensitized Solar Cell.

Sergio Tatay^a, Saif A. Haque^b, Brian O'Regan^b, James R. Durrant^b, W.J.H. Verhess^c, J.M. Kroon^c, Anton Vidal^d and Emilio Palomares,^{*d}

Figure 1S

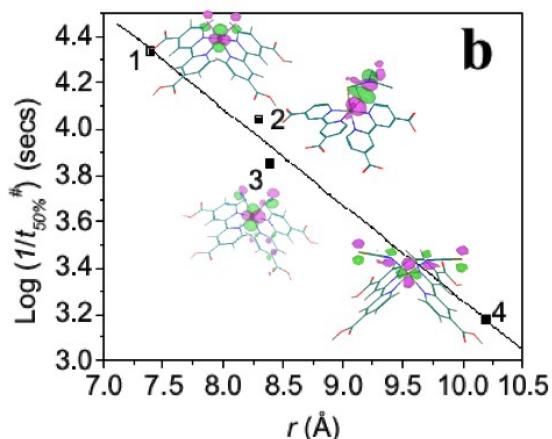
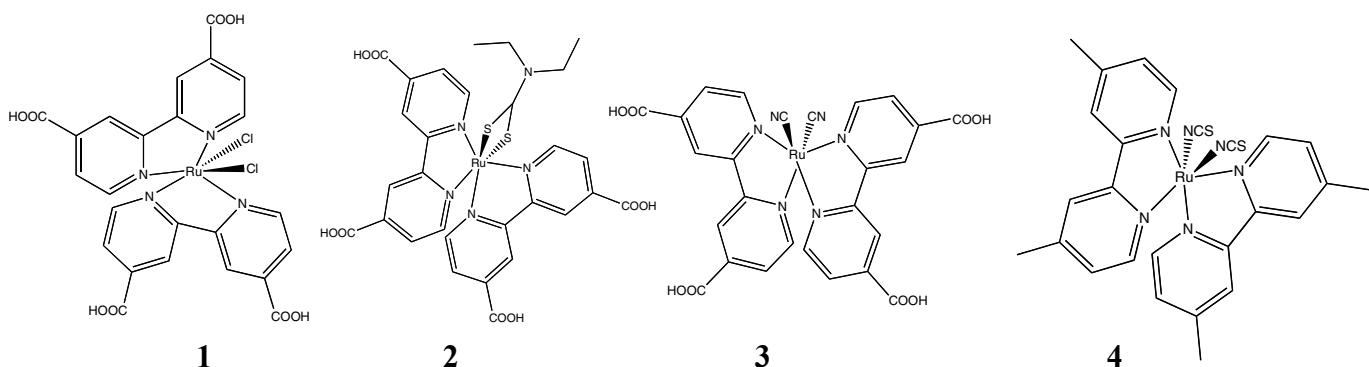


Figure 1S. Molecular structure of Ru(dcbpy)₂ (**1**), Ru(dcbpy)₂DTC (**2**), Ru(dcbpy)₂ (CN)₂ (**3**), Ru(dcbpy)₂(NCS)₂ (**4**). The **Figure 1S (b)** shows the linear correlation between the distance of the HOMO (D^+ / D^*) of the dyes to the TiO_2 nanoparticle surface (X axis) and the estimated value ($1/t_{50\%}$) of the back electron transfer reaction .

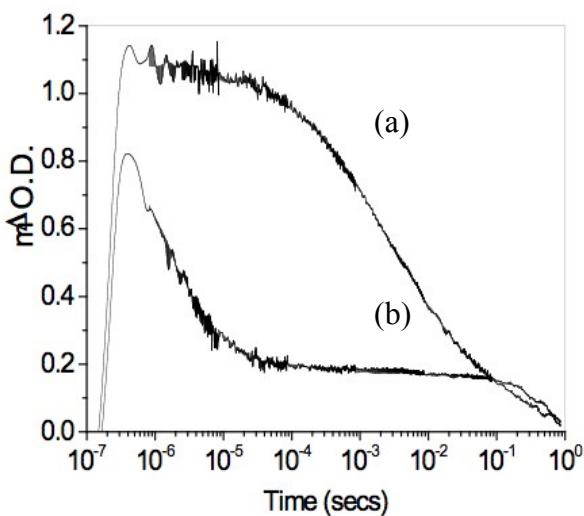


Figure 2S. Transient absorption kinetics observed for a complete N719 (dye 4 Figure 1S) sensitised TiO_2 (a) in absence of electrolyte and (b) in the presence of electrolyte. The electrolyte solution comprised 0.05M I_2 and 0.1M LiI in propylene carbonate. The monitored wavelength was 800nm.

As can be observed a biphasic decay is registered in the presence of electrolyte for N719 sensitised devices. The slow phase (0.1-1 s) at this wavelength is assigned to the absorption of I_2 species generated by the following reaction:

