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Dr. Mohammad K. Nazeeruddin



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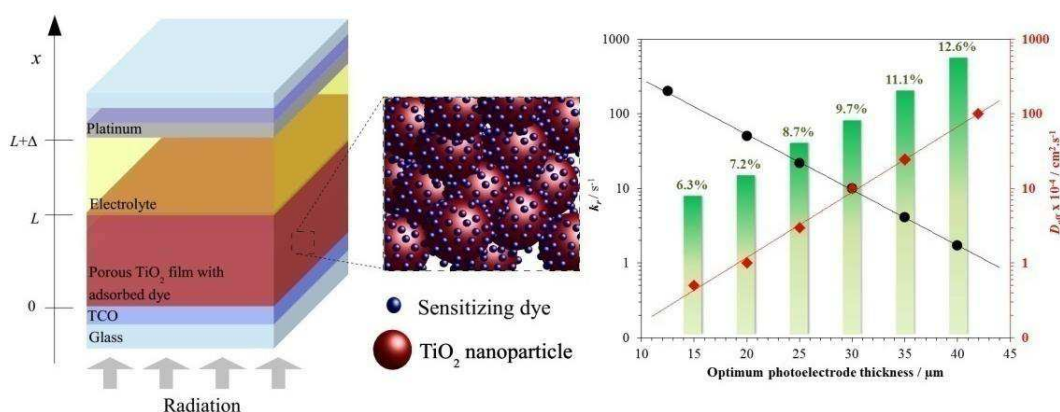
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## P167 Optimization of photoelectrode features in Cobalt (II/III) based electrolyte DSCs

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Dye-sensitized solar cells (DSSCs) have been considered an effective alternative to conventional p-n junction devices. Their low production cost, ability to harvest diffuse light and versatile applications, makes them a very interesting PV technology<sup>1-3</sup>. However, for the DSSC technology to become a competitive alternative, higher efficiencies are required. Up to now the major drawback in increasing the DSSCs efficiency has been the excessive loss of voltage during the dye-regeneration reaction; the use of iodine/triiodide redox electrolyte limits the open-circuit potential to 0.8V<sup>5</sup>. With the current state-of-art sensitizers, such as N719, and high surface area TiO<sub>2</sub>, I<sup>-</sup>/I<sub>3</sub><sup>-</sup> electrolytes achieve an efficiency maximum of 11.1%<sup>6</sup>. However, by replacing I<sup>-</sup>/I<sub>3</sub><sup>-</sup> by cobalt based redox couples new opportunities for higher efficiency DSSCs have been unlocked. The best DSSCs result achieved so far is 12.3%, accomplished by the complementary use porphyrin dyes with Co<sup>(II/III)</sup> electrolytes<sup>7</sup>. These have emerged as the future electrolytes for liquid state DSSC<sup>7-13</sup>. In comparison to I<sup>-</sup>/I<sub>3</sub><sup>-</sup>, Co<sup>(II/III)</sup> complexes possess higher reduction potentials, originating  $V_{oc}$  close to 1V; they also have several advantages such as non-corrosiveness and negligible visible absorption<sup>7</sup>. Even though the remarkable 12.3% world record, higher efficiency is desired to commercialize the DSSC technology<sup>14</sup>. The drawbacks hampering the efficiency in Co<sup>(II/III)</sup> devices are mass-transfer limitations due to electrolyte's high viscosity and very fast electron recombination with the Co<sup>(II/III)</sup> species<sup>12</sup>. In fact in these devices, the photoanode's thickness and porosity was found to be crucial in their performance<sup>13</sup>. In this work we use phenomenological modeling to optimize the photoelectrode thickness, considering the slower Co<sup>(II/III)</sup> ionic diffusion coefficient ( $1.4 \times 10^{-6}$  vs  $5 \times 10^{-6}$  cm<sup>2</sup>·s<sup>-1</sup> found in I<sup>-</sup>/I<sub>3</sub><sup>-</sup>)<sup>15</sup>. It is shown that a photoanode with higher electronic mobility should allow increasing its optimal thickness in current Co<sup>(II/III)</sup> DSSCs. New photoanode architecture is proposed, that tackles the shortcomings of cobalt-based electrolytes. Our approach uses high surface area particles with enhanced electron mobility due to the exposed (001) facets of TiO<sub>2</sub> crystals. By increasing the crystals size in (001) direction it is possible to create high electrical conductivity material capable of adsorbing large amounts of dye; these unique structures show higher electron diffusion with lower recombination rates even in the presence of Co<sup>(II/III)</sup>. This work shows that careful optimization of the TiO<sub>2</sub> crystal features is a promising route for increasing the thickness of photoanodes without increasing recombination and mass transfer resistance; and thus opening a new route for a higher efficient DSSC.



**Figure 1** Figure 1. Influence of recombination and transport are crucial in optimizing the photoelectrodes thickness

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