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Photoanode and photocathode current matching toward tandem DSCs with higher photoconversion efficiency with respect to parent single-junction DSCs

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Photoanode and photocathode current matching toward tandem DSCs with higher photoconversion efficiency with respect to parent single-junction DSCs



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INTRODUCTION



One of the reasons to research on dye-sensitized solar cells (DSCs) is to realize a photovoltaic device with (partial) optical transparency, attractive for the smart fenestration of buildings, as well as for the development of indoor light-activated powering devices that require low levels of luminous input. DSCs present three different configurations: n-type, ptype and tandem. In the first two types of solar cells are based on the nature of the semiconducting electrodes with nanostructured features; for a tandem configuration (t-DSCs) can be assembled by coupling a

METHODOLOGY

Sintering process Spray deposition method /







photoanode (sensitized TiO_2) with a photocathode (sensitized NiO), both

photoelectrochemical active. Therefore, in t-DSCs both components of the redox couple are involved in the photoactivated processes of electron transfer in which the sensitizers act as mediators. Combining two sensitized photoelectrodes in the same cell represents a meaningful approach at the condition that the two sensitizers possess complementary absorption features and similar dynamics of excitation.

The NZNCs are actually solid solutions of NiO and ZrO₂ nanodomains, the use of spraydeposited nanostructured NiO/ZrO₂ composites as photocathodic material was extended also to t-DSCs when sensitized titania (TiO_2) was the photoanode. We report here the attainment of the maximum value for the overall efficiency of the t-DSC that employs VG10-C8 squaraine and P1 as anode and cathode sensitizers, respectively.

- Heat generation from inside
- High homogeneity (no DT)
- Lower Sintering Time

The dispersion to be sprayed had the mass concentration of 10 mg of NiO NPs precursor per mL of solvent. Dispersion was sprayed with a constant flow rate of 30 μ L/min at 0.55 mPa of operating pressure with nitrogen gas as atomizer. The scan rate of the nozzle was set at 10 mm/s while the distance between the nozzle and the substrate was 15 mm. During deposition, the substrate was heated up and its temperature was maintained at 70 °C.

RESULTS AND DISCUSSION

p-DSC:

n-DSC:

t-DSC:

0.15

TiO₂

TiO₂

1,⁻/l⁻

 $|_{3}^{-}/|^{-}$

 l_{ele}

<mark>₃⁻ /l⁻ Ni</mark>C

 l_{ele}

Physical properties of NZNCs



The EDX images show the localized presence of Zr through the cyano colored areas and Zr atoms increases in passing from lower to higher Zr/Ni ratio. The dispersion of ZrO_2 NPs results uniform on nanocomposite surface when Zr/Ni < 10%. Only at larger concentrations of ZrO_2 ($Zr/Ni \ge 10\%$) there is the verification of ZrO_2 macrostructures formation. The SD samples do not show the signal of Sn (from underlying FTO) in the EDX images. This implies that the coverage of the transparent conductive substrate through the SD procedure is much more uniform than in case of the coatings obtained via screen-printed NZNCs.

Characterization of DSC p-DSC n-DSC Current Density **P1** – **VG10-C8** sensitizer squaraine 0.0 0.5 0.1 0.2 0.3 0.4 Applied Voltage / V **Device**



CONCLUSION

The combined analysis of XRD patterns and SEM pictures demonstrated that spray-deposited NiO/ZrO₂ nanocomposites are solid solutions of ZrO_2 nanodomains embedded into a matrix of nanostructured NiO. Such an optimal composition of the nanocomposite cathode was employed

(photocathode material)	v _{oc} / mv	J _{sc} / mA cm ⁻²	FF / %	η / %
p-DSC (NiO)	128	2.738	0.33	0.12
p-DSC (NZNC)	124	3.523	0.34	0.15
n-DSC	580	3.621	0.68	1.43
t-DSC (NZNC) [#] spacer thickness: 15 mm	678	4.198	0.66	1.88
t-DSC (NZNC) ^{##} spacer thickness: 15 mm	682	2.800	0.66	1.26
t-DSC (NZNC) [#] spacer thickness: 25 mm	684	3.012	0.63	1.30
t-DSC (NZNC) ^{##} spacer thickness: 25 mm	688	2.352	0.59	0.95

The photoconversion efficiencies are 1.43% and 0.15% for the n-DSC and p-DSC, respectively. The t-DSC displayed better photoconversion properties in comparison to the parent n- and p-DSCs, for the open circuit voltage. The V_{OC} value is the result of the difference between the upper edge of the valence band of the p-type semiconductor (ZrO2/NiO nanocomposite) and the lower edge of the conduction band of the n-type semiconductor (TiO2). Current density is explained in terms of inter-electrodes thickness in the device with the different configurations considered (n-, p- and t-) DSC. In the experimental conditions, the spacer interposed between the DSC electrodes as a frame has a fixed value of thickness. The t-DSC has the larger sum of electrodes thickness, and this implies the thinnest electrolytic layer (l_{ele}) with respect to the others $[l_{ele}(t-DSC)]$ $< l_{ele}$ (n-DSC) $< l_{ele}$ (p-DSC)], given a higher current density.

REFERENCES

Mozaffari, S et al. Solar Cells. Renew. Sustain. Energy Rev. 2017, 71, 675–686; Liu, J. et al. Science China Materials, 2019, 62(2), 173-180.



Bonomo M. et al. J. Electrochem. Soc. 2019, 166 (8), D290–D300.



Mariotti, N. et al. *GreenChemistry*, **2020**, 22, 7168-7218