

## Improved open-circuit voltage in ZnO/PbS quantum dot heterojunction solar cells through the surface passivation of ZnO nanowires

<sup>o</sup>Jin Chang,<sup>1</sup> Yuki Kuga,<sup>1</sup> Yuhei Ogomi,<sup>2,4</sup> Shuzi Hayase,<sup>2,4</sup> Kenji Yoshino,<sup>3,4</sup> Taro Toyoda,<sup>1,4</sup>  
Qing Shen,<sup>1,4</sup>

Univ. Electro-Commun.<sup>1</sup>, Kyushu Inst. Tech.<sup>2</sup>, Miyazaki Univ.<sup>3</sup>, CREST JST<sup>4</sup>

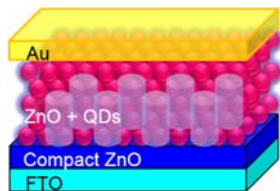
Email: shen@pc.uec.ac.jp

### Introduction

Solid state quantum dot solar cells (QDSCs), such as bulk heterojunction PbS/ZnO QDSCs, are attracting much attention owing to their broadband light-harvesting and the potential of high efficiency and low cost. Recent progress has pushed the efficiency of PbS/ZnO QDSCs above 8% through the QD surface passivation.<sup>1</sup> However, the open-circuit voltages ( $V_{oc}$ ) are still lower than theoretical expectations calculated from the corresponding energy levels.<sup>2</sup> One reason could be due to the defects located near the conduction band of ZnO. Here, we proposed a method that an ultra-thin TiO<sub>2</sub> layer grown on ZnO surfaces could eliminate the effect of ZnO defects and improve the cell performances.

### Experimental Method

ZnO-PbS bulk heterojunction solar cells (as illustrated in Fig. 1) were fabricated by solution-processed methods.<sup>3</sup> Firstly, ZnO nanowires (ZnO-NWs) were grown by a wet-chemical method on fluorine-doped tin oxide glasses coated with a ZnO compact layer. Then, thin TiO<sub>2</sub> layers were coated on ZnO-NWs by a chemical bath deposition (CBD) method. Finally, ZnO-NWs were filled and coated with PbS QDs, followed by the deposition of a gold contact layer.

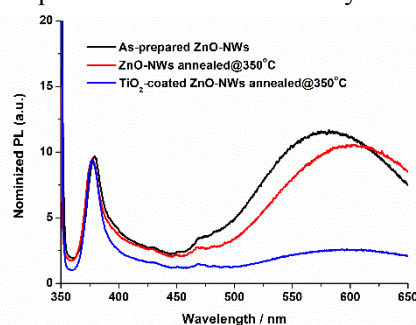


**Fig. 1** Schematic illustration of the structure of ZnO-PbS bulk heterojunction solar cells.

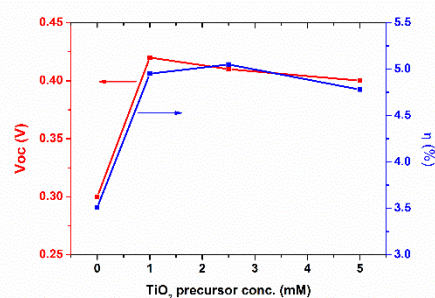
### Results and Discussion

Fig. 2 shows the fluorescence spectra of ZnO-NWs with and without TiO<sub>2</sub> coating. The emission intensity at around 600 nm significantly dropped after TiO<sub>2</sub> treatment, which was attributed to the decrease of surface defects in ZnO. As shown in Fig. 3, the  $V_{oc}$  and efficiency of ZnO/PbS cells were significantly

improved by the TiO<sub>2</sub> treatment. When higher concentration of TiO<sub>2</sub> precursors was applied, the  $V_{oc}$  slightly decreased, which could be due to the intrinsic defects within TiO<sub>2</sub> layer. Our results indicate the surface passivation can suppress recombination through removing ZnO defects and thus significantly improve the  $V_{oc}$  and power conversion efficiency.



**Fig. 2** Fluorescence spectra of as-prepared ZnO-NWs, and annealed ZnO-NWs with and without TiO<sub>2</sub> passivation.



**Fig. 3** The effect of TiO<sub>2</sub> passivation on the  $V_{oc}$  and efficiency of ZnO/PbS solar cells.

### References

1. C.-H. M. Chuang, P. R. Brown, V. Bulović and M. G. Bawendi, *Nat. Mater.*, **2014**, doi: 10.1038/nmat3984.
2. R. L. Z. Hoyer, B. Ehrler, M. L. Böhm, D. Muñoz-Rojas, R. M. Altamimi, A. Y. Alyamani, Y. Vaynzof, A. Sadhanala, G. Ercolano, N. C. Greenham, R. H. Friend, J. L. MacManus-Driscoll and K. P. Musselman, *Adv. Energy Mater.*, **2014**, doi: 10.1002/aenm.201301544.
3. H. Wang, T. Kubo, J. Nakazaki, T. Kinoshita and H. Segawa, *J. Phys. Chem. Lett.*, **2013**, *4*, 2455-2460.