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Supporting Information

Controllable coverage of Bi_2S_3 quantum dots on one-dimensional

TiO₂ nanorod arrays by pulsed laser deposition technique for high

photoelectrochemical properties

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Fig. S1 Cross-section view FESEM images of (a) BS(900)/TiO₂, (b) BS(1800)/TiO₂, (c) BS(2700)/TiO₂ and (d) BS(3600)/TiO₂.

Fig. S1 shows the cross-section view FESEM images of $BS(n)/TiO_2$ photoelectrode. As the laser ablation pulse (*n*) increases, the coverage of QDs on the nanorods increases. When the QDs are deposited with low value of *n*, there are few QDs on the surface of TiO₂ nanorods (Fig. S1a and b). With the increase of *n*, the size and quantity of QDs increases and the top of TiO₂ nanorods become domed particles (Fig. S1c). While further increasing the value of *n*, the nanorods thicken obviously and the porosity between nanorods decreases (Fig. S1d). This result is consistent with the SEM images of the surface for $BS(n)/TiO_2$ photoelectrodes in Fig. 1, which indicates that the coverage of QDs on the surface of nanorods is able to be adjusted by the PLD technique.



Fig. S2 (a) Top-view and (b) cross-section view FESEM images of TiO_2 nanorod arrays. (c) TEM and (d) HRTEM images of TiO_2 nanorod arrays, the inset in (c) shows the SAED pattern of TiO_2 nanorod arrays.

Fig. S2a shows the top-view FESEM image of plain TiO₂ nanorod arrays. With an obvious porosity between them, TiO₂ nanorods grow uniformly all over the FTO substrate. From the cross-sectional view, it can be clearly seen that the TiO₂ nanorods are vertically aligned and the length of TiO₂ nanorods is about 2 μ m. Moreover, the diameter of plain TiO₂ nanorod is about 110 nm (Fig. S2c). The SAED pattern and the HRTEM image confirm the single-crystalline nature of TiO₂ nanorods. The lattice spacing of 0.35 nm can be indexed to the (110) plane, which indicates that the nanorods grow along the (110) crystal plane with a preferred (001) orientation. The XRD spectrum presented in Figure 3a also confirms that the TiO₂ nanorods are single crystalline and can be classified as tetragonal rutile phase (JCPDS file no. 21-1276) since all the diffraction peaks well match rutile phase.