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## Fabrication and behaviors of CdS on Bi<sub>2</sub>MoO<sub>6</sub> thin film photoanodes

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Most Bi-based photoelectrodes have suitable band gaps and can effectively promote hydrogen evolution from water splitting, but there are few studies up to now for simple preparation methods for Bi-based binary metal oxides as photoanodes. Here, we prepared a novel Bi<sub>2</sub>MoO<sub>6</sub> thin film photo-anode without a template; our preparation methods of Bi-based binary metal oxides with controlled morphologies were conducted by growing the Bi<sub>2</sub>MoO<sub>6</sub> directly on an electrical substrate *via* an *in situ* growth process. The photoanodes show well-shaped thin film morphologies and exhibit impressive photoelectrochemical properties compared to the Bi-based photoanodes synthesized by conventional methods. A 2× enhanced photocurrent was obtained when the Bi<sub>2</sub>MoO<sub>6</sub> thin film photoanodes were modified with CdS in comparison with the primary Bi<sub>2</sub>MoO<sub>6</sub> (about 0.85 mA cm<sup>-2</sup>) under identical conditions. The enhanced photoelectrochemical properties were studied using several techniques including SEM, XRD, XPS, UV-vis diffuse reflectance, *etc.* and the results were in good agreement with each other. Moreover, the Bi<sub>2</sub>MoO<sub>6</sub> thin film photoanodes possess long-term stability under solar irradiation and show a considerable photocurrent.

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### 1. Introduction

Photoelectrochemical (PEC) water splitting, as a promising strategy for a renewable energy supplement and environmental protection, has attracted worldwide attention.<sup>1–3</sup> The determining factor for efficient water splitting is development of a semiconductor photoanode which has rapid charge transfer, a wide absorption spectrum, and excellent stability.<sup>4,5,6</sup> However, the traditional way of synthesizing photoanodes limits widespread application due to small specific surface area, low visible light responses, and serious charge recombination problems.<sup>7–10</sup> Therefore, developing a new material with high efficiency, durability, easy preparation, and low cost of manufacturing is extremely urgent. Cheng *et al.* developed various methods of preparing TiO<sub>2</sub> photoelectrodes and obtained excellent results.<sup>11,12</sup> Recently, Yu *et al.* synthesized a new type of TiO<sub>2</sub> photoelectrode which directly grows on fluorine-doped tin oxide (FTO) using TiCl<sub>3</sub> mediated surface treatment of TiO<sub>2</sub> nanorods that were designed and fabricated.<sup>10,13</sup> Furthermore, TiO<sub>2</sub> thin film electrodes were successfully synthesized by Zhang *et al.*<sup>14</sup> using a sol-gel method and the maximum photocurrent value was only 0.5 mA cm<sup>-2</sup>. We prepared well-aligned ZnO nanowire arrays, by adding other semiconductors, such as BiMoO<sub>6</sub> and Au, which significantly

enhanced the photoelectrochemical properties.<sup>15,16</sup> Attaching substrates increased surface area that was conducive to improved photocurrent due to better transportation of charge carriers and more reactive sites.

Compared with single metal oxides, very few simple and effective preparation methods of binary metal oxides have been developed. Binary metal oxides containing Bi(III) have been identified as promising semiconductor electrodes in solar energy conversion. For example, perovskite bismuth ferrite (BiFeO<sub>3</sub>) with a direct band-gap of approximately 2.2 eV is a promising multifunctional material that also exhibits photocatalytic properties.<sup>17,18</sup> Phase-pure BiFeO<sub>3</sub> films were grown directly *via* dual-source low-pressure from ligand-matched precursors and exhibited high activities.<sup>19</sup> In particular, Bi<sub>2</sub>WO<sub>6</sub>, BiVO<sub>4</sub>, and Bi<sub>2</sub>MoO<sub>6</sub> raised great concern due to their suitable band gaps that absorb visible light.<sup>20–25</sup> T. W. Kim and K. S. Choi synthesized nanoporous BiVO<sub>4</sub> electrodes which achieved a photocurrent density of 1.04 mA cm<sup>-2</sup> at 1.23 V *versus* RHE; the ZnFe<sub>2</sub>O<sub>4</sub> layer increased the photocurrent significantly with 2.84 mA cm<sup>-2</sup> and uniform conditions as well.<sup>26</sup> We fabricated a nanoporous BiVO<sub>4</sub> photoanode using a facile method for rational controlled pore-size and obtained the highest photocurrent density (3.5 mA cm<sup>-2</sup>) at a potential of 0.7 V *vs.* RHE for sulfite oxidation.<sup>27</sup> Correspondingly, using traditional methods such as spin-coating, dip-coating, or electrostatic self-assembly deposition, our synthesized Bi<sub>2</sub>WO<sub>6</sub> thin films are reported. As far as we know, Bi<sub>2</sub>MoO<sub>6</sub> as a thin film-type of electrode applied for photoelectrochemical purposes has rarely been reported.<sup>28–32</sup> Zhu *et al.* prepared Bi<sub>2</sub>MoO<sub>6</sub> thin

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film electrodes by means of hydrothermal methods.<sup>30</sup> Shortcomings of photoanodes with Bi<sub>2</sub>MoO<sub>6</sub> powder prepared by the above methods often caused poor attachment of particles to the electrical substrate, suppressed the transfer of photo-generated charge, and lowered the photocurrent. Recently, Gong *et al.* synthesized Bi<sub>2</sub>MoO<sub>6</sub> porous nanoflake photoanodes using BiOI as the template with a Bi source. The maximum photocurrent density was 120  $\mu\text{A cm}^{-2}$  under AM 1.5G irradiation at a constant applied bias of 1.0 V *vs.* RHE.<sup>1</sup> Based on the above study, we attempted to design and synthesize novel Bi<sub>2</sub>MoO<sub>6</sub> thin film electrodes with a simple method and high photocatalytic activities.

In this work, we exhibit a novel Bi<sub>2</sub>MoO<sub>6</sub> thin film photoanode prepared by means of a simple method. This preparation method grows Bi<sub>2</sub>MoO<sub>6</sub> on the electrical substrate directly *via* an *in situ* growth process without the aid of any template. We further raised the photocurrent on Bi<sub>2</sub>MoO<sub>6</sub> thin film photoanodes and modified with CdS. The photocurrent was improved by two times in comparison with the primary Bi<sub>2</sub>MoO<sub>6</sub> (about 0.85 mA cm<sup>-2</sup>) under identical conditions.

## 2. Experimental section

### 2.1 Preparation of composite photoanodes

All chemicals were analytical grade and used directly without any further purification. A precursor solution was prepared in which (NO<sub>3</sub>)<sub>3</sub>·5H<sub>2</sub>O and MoCl<sub>5</sub> with atomic ratios of 2 : 1 were dissolved in 8 mL ethylene glycol (EG). Then, 16 mL EG solution containing 0.544 g polyethylene glycol 600 (PEG-600) was added, followed by 60 mL ethanol and the solution was magnetically stirred and equally transferred into two 100 mL polytetrafluoroethylene stainless autoclaves. The fluorine-doped tin oxide (FTO) (1 × 5 cm<sup>2</sup>) substrate was put in the autoclave after washing with a cleaning agent, acetone solution, isopropyl alcohol, ethanol, and water, respectively. After that, the autoclave should be kept at 432 K for 24 h by a hydrothermal reaction. Finally, the materials were annealed in air at 772 K for 2.5 h after which a yellow-green film was obtained on the FTO.

Herein, the concentrations of Bi<sup>3+</sup> were controlled at 20 mM, 30 mM, and 40 mM, respectively, and the corresponding products were named as BM-20, BM-30, and BM-40. The CdS products prepared with a hydrothermal anion exchange method and 0.1234 g cadmium nitrate tetrahydrate and 0.03 g thioacetamide were separately dissolved in 10 mL deionized water. Then, the two prepared solutions were mixed and reacted in a water bath at 312 K for 15 min. The Bi<sub>2</sub>MoO<sub>6</sub> & CdS composite photoanode was synthesized by dripping with 100  $\mu\text{L}$  of the above solution and dried at 422 K.

### 2.2 Characterization of bismuth molybdate films

Morphology was characterized by a field-emission scanning electron microscope (JSM-6701F, JEOL) at an accelerating voltage of 5 kV. Transmission electron microscopy (TEM) measurements were employed using a FEI Tecnai TF20 microscope at 200 kV. The crystalline structure was identified by X-ray diffraction analysis (XRD, Rigaku RINT-2000) using Cu K $\alpha$

radiation at 40 keV and 40 mA. X-ray diffraction spectra (XRD) measurements were performed on a Rigaku RINT-2000 instrument utilizing Cu K $\alpha$  radiation (40 kV). XRD patterns were recorded from 10° to 90° with a scanning rate of 0.067° s<sup>-1</sup>. UV-vis diffuse reflectance spectra were taken on an UV-2550 (Shimadzu) spectrometer using BaSO<sub>4</sub> as the reference. The element composition was detected by X-ray photoelectron spectroscopy (XPS, ESCALAB 250Xi).

### 2.3 Photoelectrochemical measurements

All PEC measurements were conducted on an electrochemical workstation (CHI760E) in a three-electrode system. The prepared photoanode was the working electrode, a Pt plate was the counter electrode, and a saturated calomel electrode (SCE) was the reference electrode. A 300 W xenon lamp equipped with an AM 1.5 filter was used as the irradiation source with a light intensity of about 100 mW cm<sup>-2</sup> estimated with a radiometer (Newport, Models 1916C and 818-P). A 0.2 M Na<sub>2</sub>SO<sub>4</sub> aqueous solution was employed as the electrolyte. Photocurrent response tests of the photoanodes with on and off cycles were carried out at a fixed bias of 0.6 V *vs.* SCE.

## 3. Results and discussion

### 3.1 Crystal structure and morphology

As shown in Fig. 1A–F, the stacking Bi<sub>2</sub>MoO<sub>6</sub> nanoparticles of different precursor's concentration are irregularly grown in FTO. It can be clearly seen in Fig. 1A that the Bi<sub>2</sub>MoO<sub>6</sub> nanoparticles have regular nanoflower structures before annealing. After annealing, the morphology of Bi<sub>2</sub>MoO<sub>6</sub> nanoparticles was changed because the PEG-600 was eliminated. Fig. 1B shows the morphology of BM-20; here, the precursor Bi(NO<sub>3</sub>)<sub>3</sub> mole fraction is 20 mM and the Bi<sub>2</sub>MoO<sub>6</sub> nanoparticles show reduced growth in FTO, caused by a decreased illumination utilization rate. In addition, when precursor Bi(NO<sub>3</sub>)<sub>3</sub> mole content is 40 mM, the Bi<sub>2</sub>MoO<sub>6</sub> nanoparticles grew excessively in FTO, caused by a high combination of photogenerated electron-hole pairs. When the mole content of Bi<sub>2</sub>MoO<sub>6</sub> nanoparticles was 40 mM then it exposed more active crystal surfaces. Furthermore, growing irregular patterns increases specific surfaces and exposes more active sites, which contributes to absorption of light, decreases the recombination of photogenerated charge carriers, and improves photoelectric currents of the photoanodes. Fig. 1E and F show the morphologies of pure CdS and the Bi<sub>2</sub>MoO<sub>6</sub> & CdS photoanode. It can be seen that the pure CdS nanoparticles are regularly spaced and that some CdS was deposited onto the surface of Bi<sub>2</sub>MoO<sub>6</sub> nanoparticles in the Bi<sub>2</sub>MoO<sub>6</sub> & CdS composites.

With a view to further investigating CdS nanoparticles located in the composite, energy-dispersive spectroscopy (EDS) was employed and the results of different points of Bi<sub>2</sub>MoO<sub>6</sub> & CdS composite are shown in Fig. 2. The different elements and their contents can be clearly seen in Fig. 2. Because the contents of O and S were significantly decreased in the upper layer, it can be inferred that only a portion of CdS was deposited onto the surface of Bi<sub>2</sub>MoO<sub>6</sub>. Correspondingly, in the lower layer, no Bi



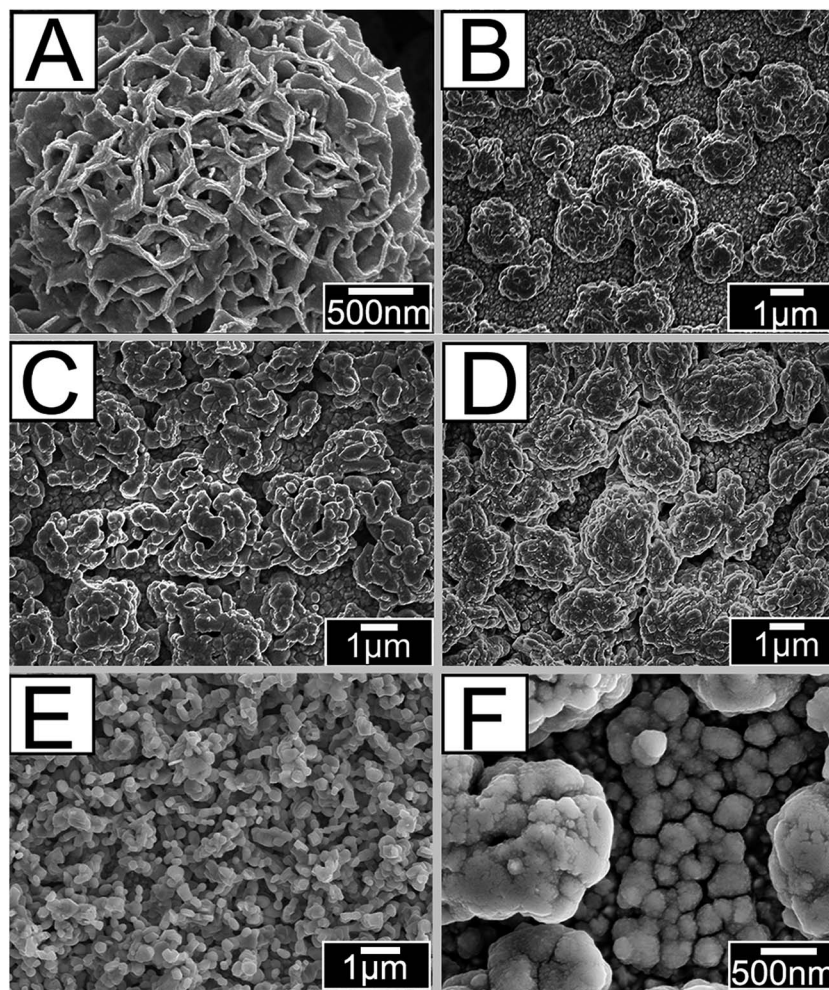


Fig. 1 SEM patterns of: (A) the  $\text{Bi}_2\text{MoO}_6$  before annealing, (B) BM-20, (C) BM-30, (D) BM-40, (E) CdS, and (F)  $\text{Bi}_2\text{MoO}_6$  & CdS composites.

and Mo elements were observed and the S and Cd element signals were significantly high, which clearly indicated that these were from CdS nanoparticles. The Si elements were from the FTO glass under high testing energy. As a result, it can be reasonably speculated that a portion of CdS was deposited onto the surface of  $\text{Bi}_2\text{MoO}_6$  and another portion of CdS was deposited onto the FTO.

The X-ray diffraction (XRD) patterns of the BM-20, BM-30, BM-40, and  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanodes are exhibited in Fig. 3. It is clearly shown that the diffraction peaks were well indexed to  $\text{Bi}_2\text{MoO}_6$  (JCPDS 21-0102); namely, the characteristic sharp peaks could be indexed to the (111), (131), (200), (202), and (133) crystal faces at 23.524, 28.264, 46.707, and 55.538. Furthermore, we observed the exposed (131) crystal faces were different with the different precursor contents; particularly, the BM-30 displayed maximal intensity of (131) crystal faces, exhibiting the best catalytic activity of current density at  $0.38 \text{ mA cm}^{-2}$ . The diffraction peaks of CdS were not observed; because of its low content in the composite, the CdS diffraction peaks are inconspicuous. In addition, the patterns have distinct diffraction peaks at 26.5, 37.7, 51.7, 61.7, and 65.7, which could be indexed to the  $\text{SnO}_2$  structure on the FTO.

### 3.2 The element composition of $\text{Bi}_2\text{MoO}_6$ & CdS composite photoanodes sample

To determine chemical composition and identify chemical states of the elements in the  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanode sample, X-ray photoelectron spectroscopy (XPS) spectra are also presented in Fig. 4. Specifically, Fig. 4A is a survey spectrum of the  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanode, which demonstrates that elements of Bi, Mo, O, Cd, and S exist in the  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanode. In Fig. 4B the binding energies 157.8 and 163.1 eV correspond to Bi  $4f_{7/2}$  and Bi  $4f_{5/2}$ , respectively.<sup>31–33</sup> For the XPS spectrum of S 2p in Fig. 4B, the peak located at 160.5 was assigned to S  $2p_{3/2}$  and another one located at 162.6 corresponded to S  $2p_{1/2}$ . In Fig. 4C, the binding energies of around 232.2 eV and 235.3 eV could be ascribed to Mo 3d.<sup>34</sup> The two peaks centered at 405.5 eV and 412.2 eV in the Cd 3d XPS spectrum (Fig. 4D) are ascribed to the Cd  $3d_{5/2}$  and Cd  $3d_{3/2}$ , respectively.<sup>35</sup>

### 3.3 UV-vis absorption spectra

Three samples, namely  $\text{Bi}_2\text{MoO}_6$ , CdS, and  $\text{Bi}_2\text{MoO}_6$  & CdS photoanodes, were employed. The sample of pure  $\text{Bi}_2\text{MoO}_6$  was



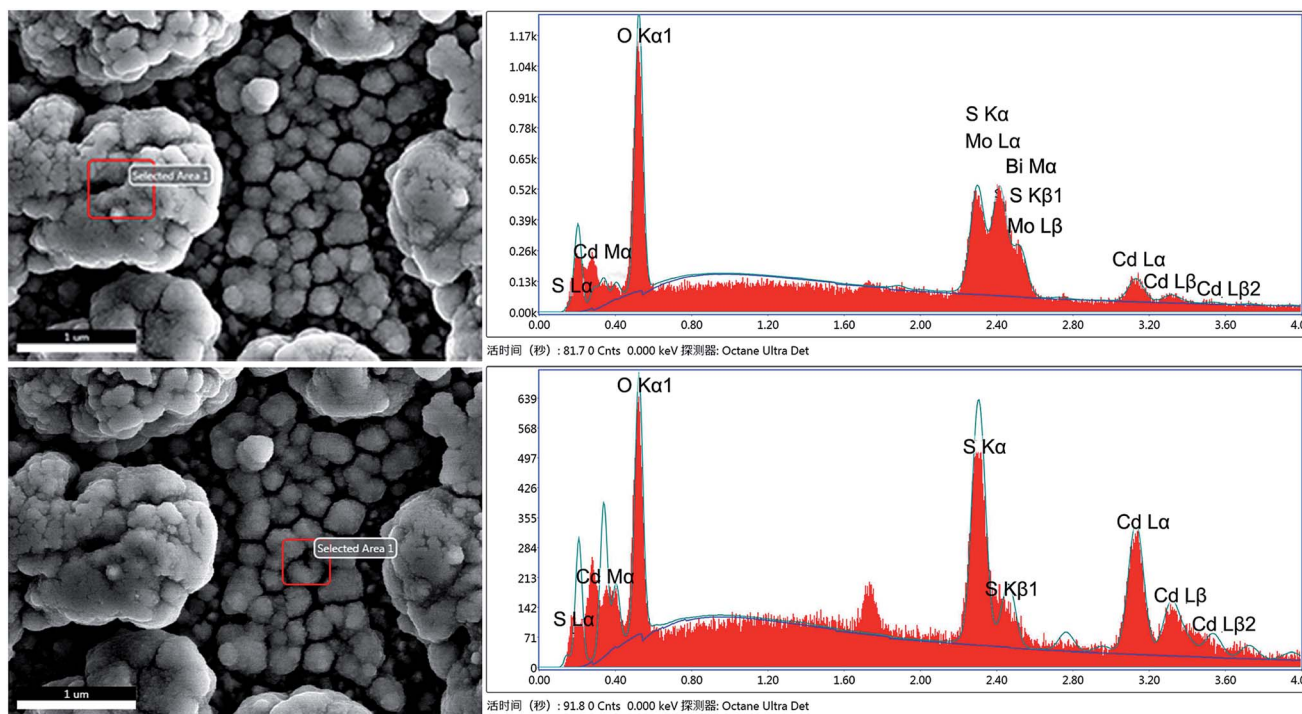


Fig. 2 The EDX patterns of  $\text{Bi}_2\text{MoO}_6$  & CdS composites.

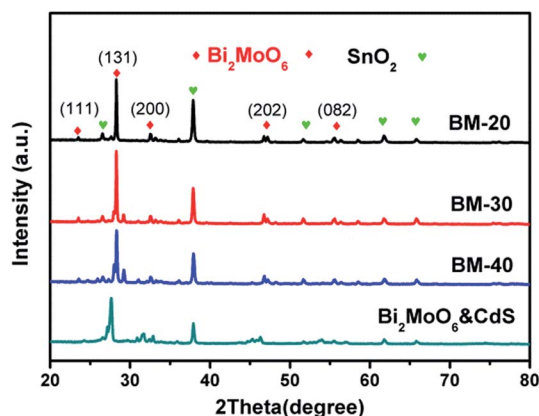


Fig. 3 XRD patterns of the BM-20, BM-30, BM-40, and  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanodes samples.

prepared with  $\text{Bi}(\text{NO}_3)_3$  as the precursor and the mole content is 30 mM. The  $\text{Bi}_2\text{MoO}_6$  & CdS photoanode agreed with the XPS sample's data.

UV-vis light absorption spectra of pure  $\text{Bi}_2\text{MoO}_6$ , CdS, and  $\text{Bi}_2\text{MoO}_6$  & CdS photoanodes are shown in Fig. 5A, respectively. It can be clearly seen that absorption of the pure  $\text{Bi}_2\text{MoO}_6$  photoelectrode was 450 nm, the pure CdS photoelectrode was 500 nm, and the  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoelectrode was 490 nm. The UV-vis DRS of the  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoelectrode has an obviously red shift compared to pure  $\text{Bi}_2\text{MoO}_6$ . In addition, the band gap of CdS and  $\text{Bi}_2\text{MoO}_6$  was calculated in Fig. 5B. Specifically, the band gap of CdS was 2.46 eV and  $\text{Bi}_2\text{MoO}_6$  was 2.73, which was beneficial to

photocatalytic excitation by visible light. This suggested that the as-fabricated  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoelectrode had a suitable band gap. Therefore, good PEC properties of the  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoelectrode under solar irradiation are presented.

### 3.4 The photoelectrochemical properties

The photoelectrochemical properties of BM-20, BM-30, BM-40, and  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanode were investigated. Specifically, an experiment was carried out in 0.2 M  $\text{Na}_2\text{SO}_4$  under illumination of  $100 \text{ mW cm}^{-2}$  coupled with an AM 1.5 filter to provide simulated solar irradiation. The linear-sweep voltammograms under transient illumination are presented in Fig. 6A, from which it can be seen that the composite electrodes of  $\text{Bi}_2\text{MoO}_6$  & CdS have a much lower photocurrent onset potential than the pure  $\text{Bi}_2\text{MoO}_6$ . This is mainly because of the extended absorption edges of the  $\text{Bi}_2\text{MoO}_6$  photoelectrode resulting from the loaded CdS. Furthermore, pure CdS presented an abnormal decline following potential augmenting because pure CdS has a serious light corrosion. Fig. 6B shows amperometric  $I-t$  curves of pure CdS, BM-20, BM-30, BM-40, and  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanodes. The current density of pure CdS rapidly descended with irradiation time while pure  $\text{Bi}_2\text{MoO}_6$  BM-40 displayed maximum photocurrent density ( $0.38 \text{ mA cm}^{-2}$ ) at a potential of 0.6 V vs. RHE for 0.2 M  $\text{Na}_2\text{SO}_4$ . The  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanode exhibited the highest photocurrent density ( $0.86 \text{ mA cm}^{-2}$ ) under the same conditions. When CdS was added to the pure  $\text{Bi}_2\text{MoO}_6$  photoanode, it extended the absorption edges as well as providing more active sites, which facilitated the photoanode



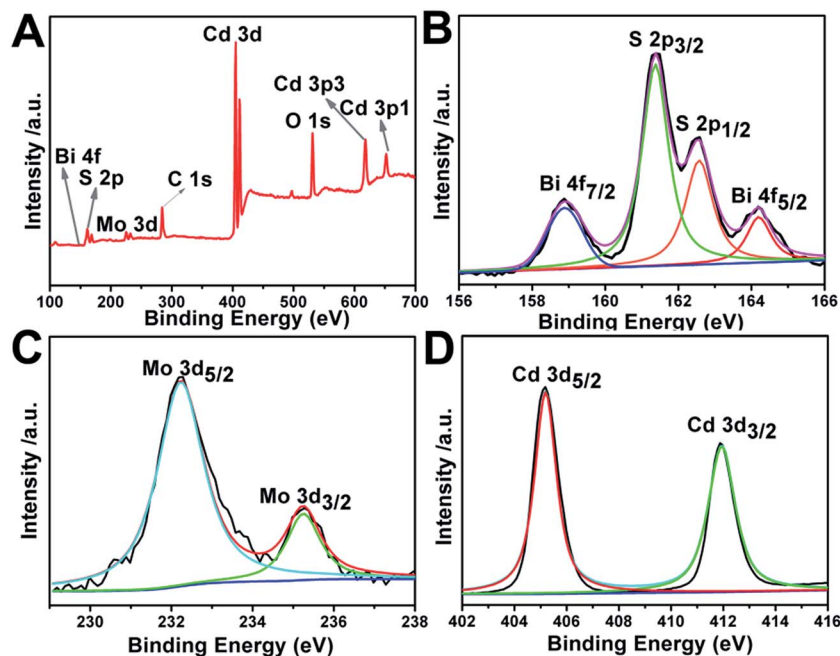


Fig. 4 The X-ray photoelectron spectroscopy (XPS) spectra of  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanode.

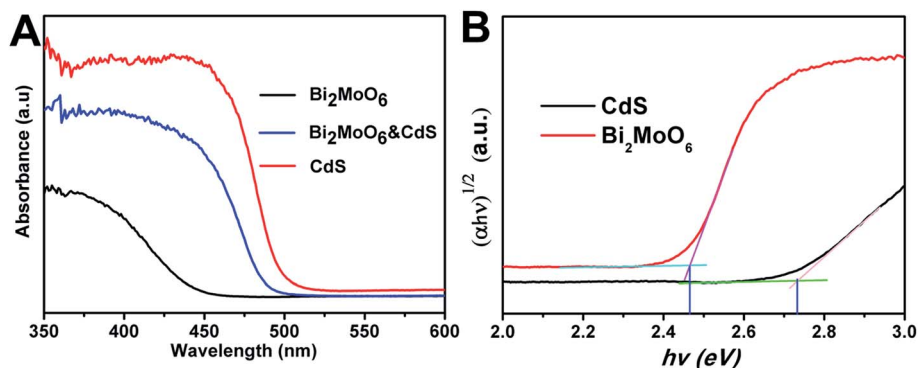


Fig. 5 (A) The UV-vis absorption spectra of the  $\text{Bi}_2\text{MoO}_6$ , CdS, and  $\text{Bi}_2\text{MoO}_6$  & CdS electrodes; (B) the band gap of CdS and  $\text{Bi}_2\text{MoO}_6$ .

taking advantage of the wider light irradiation and photo-generated charges carriers. Meanwhile,  $\text{Bi}_2\text{MoO}_6$  could carry photogenerated charges of CdS away over time so that CdS light

corrosion was inhibited. Therefore, the  $\text{Bi}_2\text{MoO}_6$  & CdS composite electrode remarkably enhanced the photocurrent property.

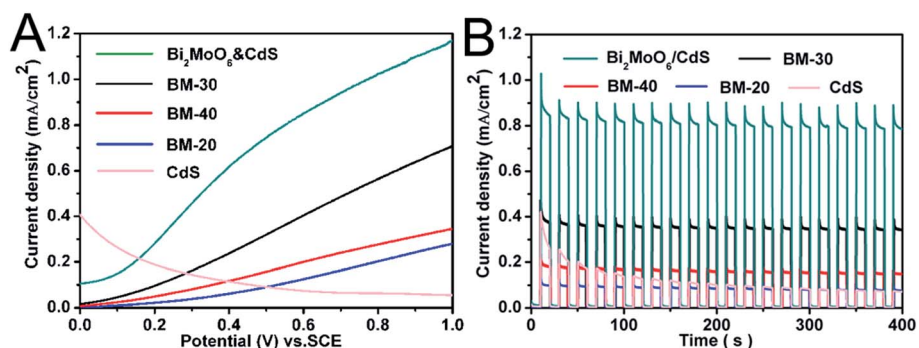


Fig. 6 (A) Linear sweep voltammograms of BM-20, BM-30, BM-40, and  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanode samples in visible light illumination ( $100 \text{ mW cm}^{-2}$ ); (B) amperometric  $I-t$  curves of BM-20, BM-30, BM-40, and  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanodes samples.



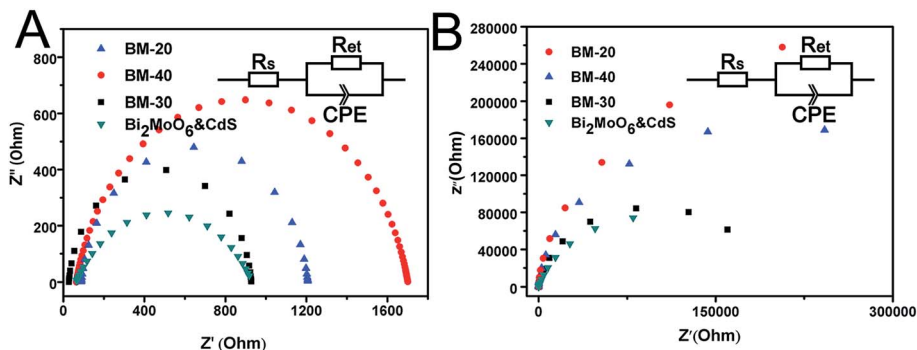


Fig. 7 Electrochemical impedance spectroscopy (EIS) Nyquist plots of BM-20, BM-30, BM-40, and  $\text{Bi}_2\text{MoO}_6$  & CdS composite photoanodes samples.

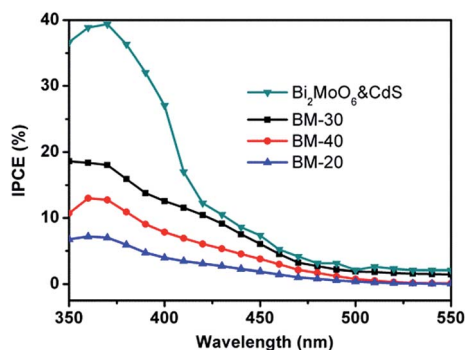


Fig. 8 The incident photon-to-current conversion efficiency (IPCE) of BM-20, BM-30, BM-40, and  $\text{Bi}_2\text{MoO}_6$  & CdS.

The interfacial charge transfer on the electrode can be characterized by Electrochemical Impedance Spectroscopy (EIS) shown in Fig. 7. The interfacial charge transfer resistance ( $R_{ct}$ ) can be represented by the diameter of the semicircle in a Nyquist plot. It can be clearly seen that the  $R_{ct}$  under illumination (Fig. 7A) is lower than that in the dark (Fig. 7B), which suggests a fast interfacial charge transfer upon illumination. The diameter of  $\text{Bi}_2\text{MoO}_6$  & CdS semicircle is smaller than that of pure  $\text{Bi}_2\text{MoO}_6$ , proving that the decoration of CdS facilitated the charge transfer process due to a type II band alignment between  $\text{Bi}_2\text{MoO}_6$  and CdS.

Fig. 8 shows measurements of the incident photon-to-current conversion efficiency (IPCE) of Z BM-20, BM-30, BM-40, and  $\text{Bi}_2\text{MoO}_6$  & CdS.

IPCE was calculated as follows:<sup>19,27,36</sup>

$$\text{IPCE (\%)} = \frac{1240 \times I \text{ (mA cm}^{-2}\text{)}}{P_{\text{light}} \text{ (mW cm}^{-2}\text{)} \times \lambda \text{ (nm)}} \times 100 \quad (1)$$

where  $I$  is the measured photocurrent density at a specific wavelength,  $\lambda$  is the wavelength of incident light, and  $P_{\text{light}}$  is the measured light power density at that wavelength. IPCE analysis of the photoanodes showed a similar distribution compared with current-potential characteristics; BM-20 has an exhibited maximum conversion efficiency of 18% in 350 nm and when CdS is introduced the maximum conversion efficiency is 40% at 370 nm, which is because CdS extended the absorption edges of the  $\text{Bi}_2\text{MoO}_6$  photoelectrode.

## 4. Conclusion

In summary, since most Bi-based photoelectrodes have suitable band gaps and can effectively promote hydrogen evolution from water splitting, a novel  $\text{Bi}_2\text{MoO}_6$  thin film photoanode was prepared by growing  $\text{Bi}_2\text{MoO}_6$  on an electrical substrate directly *via an in situ* process. The photoanodes show well-shaped thin film morphologies and exhibit impressive photoelectrochemical properties compared to Bi-based photoanodes synthesized by conventional methods. Particularly, the morphology of our thin film photoanodes can be controlled by precursor content. Crystal structure and morphology was carefully studied with SEM, XPS, UV-vis diffuse reflectance, and PEC performance *etc.*, and the results are in good agreement with each other.

The  $\text{Bi}_2\text{MoO}_6$  thin film photoanodes possess long-term stability under solar irradiation and show a considerable photocurrent. The results of our photoelectron studies proved that the precursor  $\text{Bi}(\text{NO}_3)_3$  mole fraction with 30 mM can reach a maximal photocurrent of  $0.38 \text{ mA cm}^{-2}$ . In addition, we further enhanced the  $\text{Bi}_2\text{MoO}_6$  thin film photoanodes photocurrent after CdS modifying the thin film electrodes, the photocurrent of which doubled in comparison with the primary  $\text{Bi}_2\text{MoO}_6$  (about  $0.85 \text{ mA cm}^{-2}$ ) under the identical conditions. We believe that this facile method may be suitable for synthesizing nanostructured  $\text{Bi}_2\text{MoO}_6$ -based materials for use in solar energy devices.

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