


 Cite this: *RSC Adv.*, 2021, **11**, 21642

# Facile preparation of $Zn_xCd_{1-x}S/ZnS$ heterostructures with enhanced photocatalytic hydrogen evolution under visible light

 Jing Dong,<sup>a</sup> Wenjian Fang,<sup>b</sup> Weiwei Xia,<sup>c</sup> Qihong Lu<sup>c</sup> and Xianghua Zeng<sup>\*bc</sup>

Hydrogen evolution from water using solar energy is regarded as a most promising process, thus, exploring efficient photocatalysts for water splitting is highly desirable. To avoid the rapid recombination of photogenerated electrons and holes in CdZnS semiconductors,  $Zn_xCd_{1-x}S/ZnS$  composites were synthesized via a one-step hydrothermal method and then annealed at 400 °C for 60 min under argon flow.  $Zn_xCd_{1-x}S/ZnS$  composites are composed of ZnS nanosheets decorated with  $Zn_xCd_{1-x}S$  nanorods, and TEM and UV-vis absorption spectra confirm the formation of the heterostructure between  $Zn_xCd_{1-x}S$  nanorods and ZnS nanosheets. Because of the well-matched band alignment, stronger optical absorption and larger carrier density,  $Zn_{0.2}Cd_{0.8}S/ZnS$  has the highest hydrogen production, with a photocatalytic hydrogen production rate up to 16.7 mmol g<sup>-1</sup> h<sup>-1</sup> under visible light irradiation. Moreover, the photocatalyst also exhibits high stability and good reusability for hydrogen production reaction. The facile and efficient approach for ZnS based heterostructures could be extended to other metal compound materials.

 Received 24th April 2021  
 Accepted 13th June 2021

DOI: 10.1039/d1ra03195c

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## Introduction

In consideration of the global energy crisis and environmental issues, much effort has been made to replace exhaustible fossil fuels with clean energy sources. Hydrogen, with a fuel value (~143 kJ g<sup>-1</sup>) three times higher than that of gasoline and only non-polluted water as a by-product, has been regarded as one of the most important green energy carriers to replace fossil fuels. Among various strategies for hydrogen evolution, photocatalytic hydrogen evolution is regarded as the most promising way from water splitting using solar energy, because of the direct utilization of solar energy to achieve H<sub>2</sub> generation from water splitting, as well as its much simpler and more economically competitive systems.<sup>1</sup>

Ternary ZnCdS nanomaterials were regarded as a promising substitute for noble-metal co-catalysts in photocatalytic reactions because of their tunable band gap energy and lattice constant between 0.331 ((100) of ZnS) and 0.359 nm (<(100) of CdS) in comparison with ZnS or CdS materials.<sup>2-6</sup> Recently some studies on ZnCdS-related photocatalytic hydrogen reactions have been reported. Such as, Jin *et al.*<sup>7</sup> reported that ZnCdS catalysts with a (Zn + Cd):S molar ratio of 1:3.5 has much higher

photocatalytic activity and exhibited an excellent stability after six cycles. And the H<sub>2</sub> production of (CdS-ZnS)-TiO<sub>2</sub> supported photocatalytic system was studied by Tambwekar *et al.*<sup>8</sup> The hydrogen evolution rate of ZnCdS-CdS heterostructure was obtained from 25.46 to 72.82 and 52.82 μmol h<sup>-1</sup>, and the value increased to 192.28 μmol h<sup>-1</sup> after the adoption of VS<sub>2</sub> to the heterostructure surface.<sup>9</sup> GQDs/ZnCdS/PdS exhibited a H<sub>2</sub> evolution rate of 517 μmol h<sup>-1</sup>, which is 15, 7 and 1.7 times higher than that of pure ZnCdS, GQDs/ZnCdS, and ZnCdS/PdS, respectively.<sup>10</sup> α-Fe<sub>2</sub>O<sub>3</sub>/Zn<sub>0.4</sub>Cd<sub>0.6</sub>S heterostructure has visible light photocatalytic H<sub>2</sub> production of 536.8 μmol h<sup>-1</sup>.<sup>11</sup> The ZnCdS-CH solid solution without noble metal loading achieved a superior photocatalytic H<sub>2</sub> activity rate of 0.971 mmol h<sup>-1</sup> under visible light irradiation (λ ≥ 420 nm), which exceeds those of coprecipitated Zn<sub>0.5</sub>Cd<sub>0.5</sub>S samples by more than 13 times.<sup>12</sup> The hydrogen production rate of Fe<sub>1-x</sub>Pt<sub>x</sub>-ZnCdS NPs had a significant enhancement over the pure ZnCdS (740 μmol g<sup>-1</sup> h<sup>-1</sup>). The hydrogen production activity of ZnCdS-NiCoP composite catalyst was improved greatly, which reached 5.2 times that of pure ZnCdS.<sup>13</sup> The highest hydrogen production rate of 2.265 mmol g<sup>-1</sup> h<sup>-1</sup> was achieved by the 0.5 wt% Fe<sub>0.3</sub>Pt<sub>0.7</sub>-ZnCdS nanocomposites, which was even better than that of 0.5 wt% Pt-ZnCdS (1.626 mmol g<sup>-1</sup> h<sup>-1</sup>) under the same condition.<sup>14</sup>

The photocatalytic H<sub>2</sub> evolution rate of Zn<sub>0.3</sub>Cd<sub>0.7</sub>S nanorods was improved from 517.4 to 3310.1 μmol g<sup>-1</sup> h<sup>-1</sup> by loading a suitable amount of Ni<sub>3</sub>C NPs as co-catalyst under visible light irradiation.<sup>15</sup> Hollow Zn<sub>0.6</sub>Cd<sub>0.4</sub>S cage material exhibited the highest hydrogen production rate of 5.68 mmol h<sup>-1</sup> g<sup>-1</sup> under cocatalyst-free and visible-light irradiation conditions.<sup>16</sup> The

<sup>a</sup>College of Chemistry and Chemical Engineering, Yangzhou University, Yangzhou 225002, P. R. China

<sup>b</sup>College of Electrical, Energy and Power Engineering, Yangzhou University, Yangzhou 225127, P. R. China. E-mail: xhzeng@yzu.edu.cn

<sup>c</sup>College of Physics Science and Technology & Institute of Optoelectronic Technology, Yangzhou University, Yangzhou 225002, P. R. China


addition of NiB greatly improved the photocatalytic performance of CdZnS, and the hydrogen production of NiB/CdZnS catalysts reached  $8.137 \text{ mmol g}^{-1} \text{ h}^{-1}$ , which is 17 times that of pure CdZnS.<sup>17</sup> A typical “type II” band alignment forms at  $\text{Zn}_{0.5}\text{Cd}_{0.5}\text{S}$  hybridized with carbon materials lead to the improved  $\text{H}_2$  generation rate of  $10.8 \text{ mmol g}^{-1} \text{ h}^{-1}$ , exceeding the  $\text{Zn}_{0.5}\text{Cd}_{0.5}\text{S}$  nanosheets ( $6.4 \text{ mmol g}^{-1} \text{ h}^{-1}$ ) and QDs ( $3.4 \text{ mmol g}^{-1} \text{ h}^{-1}$ ).<sup>18</sup> A controlled ZnCdS QDs developed by a simple ZIF-8 templating method exhibited significantly enhanced photocatalysis performance with a  $\text{H}_2$  production rate of  $3.70 \text{ mmol h}^{-1} \text{ g}^{-1}$  ( $\text{Zn}_{0.5}\text{Cd}_{0.5}\text{S}$  QDs),<sup>19</sup> and the highest hydrogen production rate of  $\text{Zn}_{0.5}\text{Cd}_{0.5}\text{S}$ /dodecahedron ZIF-67 composite reached  $23.264 \text{ mmol g}^{-1} \text{ h}^{-1}$  under visible light irradiation,<sup>20</sup> which was explained as the matched valence band position for ZIF-67 and ZCS and the exposure of rich active sites. The  $\text{WO}_3/\text{ZnCdS}$  compound catalysts showed the hydrogen production activity of the 35 wt%  $\text{WO}_3/\text{ZnCdS}$  to  $98.68 \mu\text{mol mg}^{-1}$ , about 9.6 times that of pure ZnCdS ( $10.28 \mu\text{mol mg}^{-1}$ ).<sup>21</sup>

Although many efforts have been done to develop efficient photocatalytic systems and photocatalysts by using ZnCdS materials and their composites, it is still an urgent task to develop low-cost co-catalysts for advancing photocatalytic  $\text{H}_2$  production. In this paper, simple hydrothermal and annealing reaction methods were developed for preparation of pure ZnS nanosheets, CdS nanorods and  $\text{Zn}_x\text{Cd}_{1-x}\text{S}/\text{ZnS}$  composites. The photocatalytic hydrogen production rate of  $\text{Zn}_{0.2}\text{Cd}_{0.8}\text{S}/\text{ZnS}$  (#3) up to  $16.7 \text{ mmol g}^{-1} \text{ h}^{-1}$  has been obtained under visible light irradiation with a good stability. This value is larger than the reported pure  $\text{Zn}_x\text{Cd}_{1-x}\text{S}/\text{ZnS}$  composites. Furthermore, the preparation method is simple. The outstanding photocatalytic performance benefited from the efficient charge transfer between ZnCdS nanorods and ZnS nanosheets in the  $\text{Zn}_x\text{Cd}_{1-x}\text{S}/\text{ZnS}$  heterojunctions, as well as more active sites on the surface of nanosheets. This study will be helpful to obtain catalysts for photocatalytic  $\text{H}_2$  production based on low-priced metal composite materials.

## Experimental section

### Materials and chemicals

All chemicals were analytical-grade and used without further purification. Zinc acetate [ $\text{Zn}(\text{OAc})_2 \cdot 2\text{H}_2\text{O}$ ], cadmium acetate [ $\text{Cd}(\text{OAc})_2 \cdot 2\text{H}_2\text{O}$ ], thiourea [ $(\text{NH}_2)_2\text{CS}$ ] and ethanol were purchased from Sinopharm Chemical Reagent Co., Ltd (Shanghai China). Ethylenediamine was purchased from Aladdin Chemical Reagent Co., Ltd (Shanghai).

### Sample preparation

ZnS nanosheets, CdS nanorods and  $\text{Zn}_x\text{Cd}_{1-x}\text{S}/\text{ZnS}$  heterostructure were synthesized according to the following methods. ZnS nanosheets were prepared according to the previous reports with some modifications.<sup>22</sup> Typically, 60 mM  $\text{Zn}(\text{OAc})_2 \cdot 2\text{H}_2\text{O}$  was dissolved in 6 mL deionized water under constant magnetic stirring for approximately 10 min, then, 24 mL ethylenediamine was added to the solution. After the mixed solution was stirred continuously and cooled to room temperature, 0.18 M  $(\text{NH}_2)_2\text{CS}$

was added into the above solution. After stirring for another 30 min, the mixture was transferred into a 50 mL Teflon-lined stainless steel autoclave and keeping sealed under  $160 \text{ }^\circ\text{C}$  for 6 h. The product was collected and washed with water and ethanol for several times and dried at  $60 \text{ }^\circ\text{C}$  for 24 h. Furthermore, the precursor was annealed in the quartz tube furnace at  $400 \text{ }^\circ\text{C}$  for 60 min under argon (Ar) flow. After heat treatment, ZnS nanosheets were achieved, labeled as sample #1. CdS nanorods were prepared with the same procedure, except that 60 mM  $\text{Cd}(\text{OAc})_2 \cdot 2\text{H}_2\text{O}$  was used as cationic precursor (so called as sample #4). For  $\text{Zn}_x\text{Cd}_{1-x}\text{S}/\text{ZnS}$  heterostructures, 48 mM  $\text{Zn}(\text{OAc})_2 \cdot 2\text{H}_2\text{O}$  and 12 mM  $\text{Cd}(\text{OAc})_2 \cdot 2\text{H}_2\text{O}$  were used as cationic precursor, labeled as sample #2; 36 mM  $\text{Zn}(\text{OAc})_2 \cdot 2\text{H}_2\text{O}$  and 24 mM  $\text{Cd}(\text{OAc})_2 \cdot 2\text{H}_2\text{O}$  were used as cationic precursor, named as sample #3.

### Characterization

The crystal structures of the samples were analyzed by powder XRD (D8 Advance, Bruker-AXS) using  $\text{Cu K}\alpha$  ( $\lambda = 0.154056 \text{ nm}$ ), and their patterns were collected in the  $2\theta$  range from  $10^\circ$  to  $80^\circ$  using a continuous scanning method at a scanning speed of  $2^\circ (2\theta) \text{ min}^{-1}$ . The morphology and microstructure of the as-synthesized samples were characterized using Hitachi S-4800 field emission SEM and Tecnai G2 F30 field emission TEM, operated at an accelerating voltage of 300 kV, the composition of the products was measured by energy dispersive X-ray spectroscopy (EDX). Brunauer–Emmett–Teller (BET) surface area measurements were performed with  $\text{N}_2$  adsorption at 77 K on a BSD-PS (M) instrument. Before analyses, the samples were degassed at  $80 \text{ }^\circ\text{C}$ . Absorption measurements were carried out using a UV-vis-NIR spectrophotometer (UV-vis, Cary-5000, Varian). Photoluminescence (PL) measurements were performed using a Britain Renishaw InVia spectrophotometer, with a 325 nm line of a He–Cd laser as the excitation light source in a closed-cycle He cryostat. In addition, the chemical states of Zn, Cd and S elements were analyzed by X-ray photoelectron spectroscopy (XPS, Thermo ESCALAB250Xi) which was equipped with a standard monochromatic Al-K $\alpha$  source ( $h\nu = 1486.6 \text{ eV}$ ), the binding energies were referred to C 1s peak (284.8 eV).

The PEC analysis was carried out on Zahner CIMPS electrochemical workstation (Germany) in a quartz cell, using a three-electrode cell with a Pt wire as the counter electrode, Ag/AgCl electrode as the reference electrode and FTO glass covered with photocatalyst as the working electrode. A 300 W Xe lamp (CEL-HXF 300, Beijing Au-light, China) was employed as an incident light source to study the PEC response of the samples, and  $\text{Na}_2\text{SO}_4$  (0.2 M) solution was used as the electrolyte.

### Photocatalytic $\text{H}_2$ production

The photocatalytic reactions were carried out in a Pyrex reaction cell connected to a closed gas circulation and evacuation system. Firstly, 50 mg prepared samples and  $\text{H}_2\text{PtCl}_6$  solution ( $0.004 \text{ g mL}^{-1}$ ) containing 0.5 mg Pt were dispersed in 50 mL of mixed aqueous solution containing 40 mL deionized water and 10 mL sacrificial agent (lactic acid) in a Pyrex reaction cell with constant stirring. Then, the suspension was thoroughly



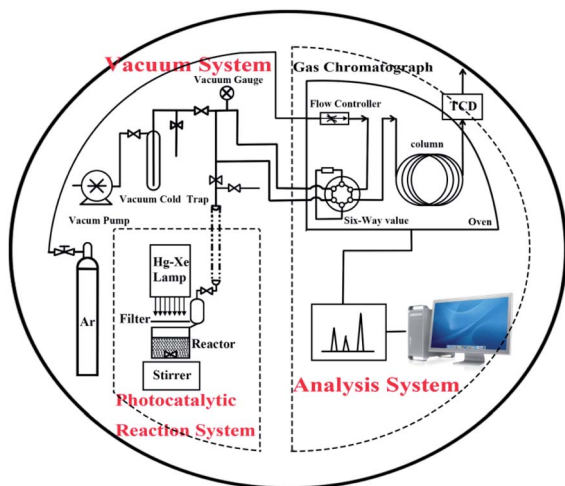


Fig. 1 Scheme of photocatalytic reactor and closed gas circulation and evacuation system made of Pyrex glass.

degassed and irradiated by a Xe lamp (300 W) with a cutoff filter ( $\lambda$  greater than 420 nm). The amount of  $H_2$  was analyzed every 30 min using an online gas chromatography. The setup was designed by Liu *et al.*,<sup>23</sup> as shown in Fig. 1.

## Results and discussion

The XRD patterns of the prepared samples are shown in Fig. 2a, where the diffraction peaks can be indexed as the hexagonal

ZnS (PDF#36-1450) and hexagonal CdS (PDF#41-1049) for samples #1 and #4, respectively. While for samples #2 and #3, the diffraction peaks are composed with both hexagonal ZnS and hexagonal  $Zn_xCd_{1-x}S$  (PDF#40-0836). With the increase of Cd precursor, the diffraction peaks belonging to  $Zn_xCd_{1-x}S$  become stronger and the peak positions shift to low-angle side continuously. At the same time, the peak intensity assigned to hexagonal ZnS decreases gradually while its positions is nearly unchanged, indicating that the content of ZnS decreases constantly as shown in the magnified XRD pattern (Fig. 2b). Moreover, from the typical peak positions and Vegard's law, the lattice parameters of  $Zn_xCd_{1-x}S$  solid solutions varies linearly with composition at constant temperature,<sup>24,25</sup> from the parameters of sample #2 and #3, we further estimate that  $x$  is approximately to 0.3 in #2, and 0.2 in #3. The main lattice parameters were displayed in Table 1. The structure and the chemical composition of sample #3 were characterized using

Table 1 The main lattice parameters for samples #2 and #3

	$d$	$(h\ k\ l)$	$2\theta$
#2	3.5005	(100)	25.424
	3.2940	(002)	27.047
	3.0912	(101)	28.858
#3	3.5239	(100)	25.252
	3.3105	(002)	26.910
	3.1107	(101)	28.674

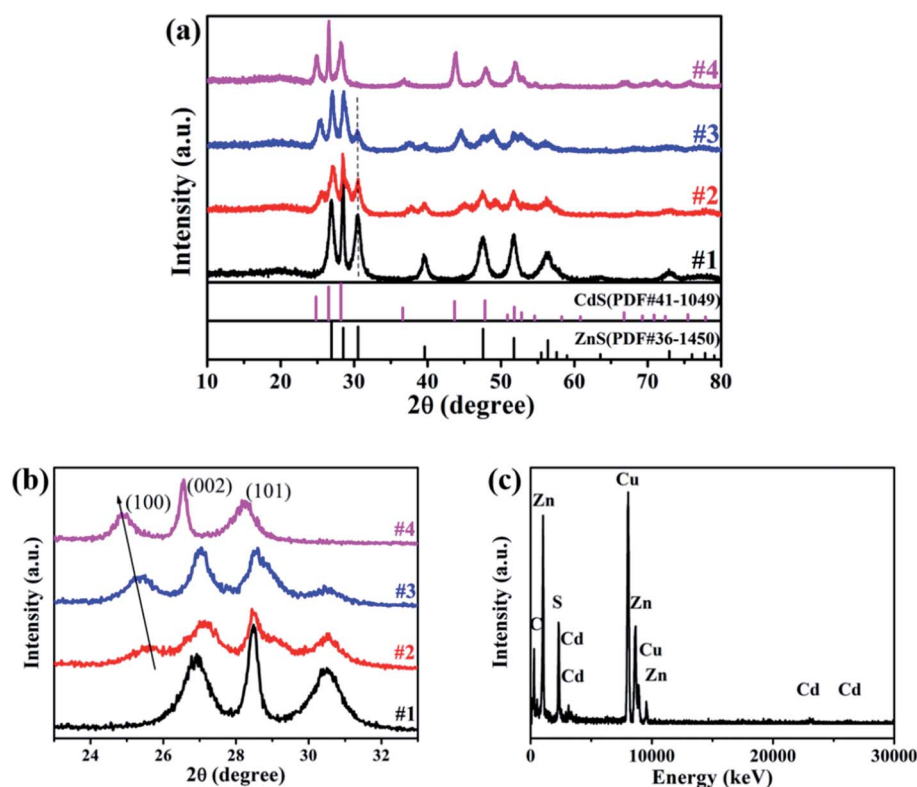


Fig. 2 (a) and (b) XRD patterns of samples ZnS (#1),  $Zn_xCd_{1-x}S/ZnS$  (#2, #3) and CdS (#4); (c) EDX spectrum for sample #3.



Table 2 Elemental percentage in sample #3 (EDX) data

Element	Atomic percentage %
S	40.15
Zn	54.83
Cd	5.00

EDX spectroscopy, and the EDX spectrum clearly reveals the existence of Zn, Cd and Cu elements in the prepared samples, as shown in Fig. 2c.

The atomic weight percentages of the constituents are given in Table 2, we find that the metal element Zn is dominated in sample #3, and the compositional percentages of  $(\text{Zn}^{2+} + \text{Cd}^{2+})$  and  $\text{S}^{2-}$  are in the required stoichiometric ratio, but larger than 1. So, there are a lots of sulfur vacancies in the prepared samples.

The morphologies of samples #1, #3 and #4 were investigated with SEM images as shown in Fig. 3. From Fig. 3a, sample #1 is composed with nanosheets, each nanosheet has a thickness of about 50 nm (inset of Fig. 3a), and sample #4 is composed with nanorods, each individual nanorod has a diameter of several tens nm and length of several hundred nm (Fig. 3b). Low- and high-magnification images of sample #3 were displayed in Fig. 3c and d, which show that after replacing Zn precursor with Cd, the products are mainly composed with nanosheets with some nanorods sparsely growing on the surface of ZnS nanosheets. Sample #2 has the same morphology as sample #3, but it is thicker and narrower than sample #3.

TEM and HRTEM measurements were carried out to investigate the microstructure features, and HAADF-STEM to determine the elements distribution. TEM images of selected

$\text{Zn}_x\text{Cd}_{1-x}\text{S}/\text{ZnS}$  are displayed in Fig. 4a, the nanorods with a diameter of  $\sim 20$  nm were grown on the nanosheets, which is consistent with the result of SEM. Furthermore, from the HRTEM image (Fig. 4b), one can find that the well-resolved lattice fringes with separations of 3.53 and 3.31 Å matched well with the interplanar spacing of (100) and (002) of  $\text{Zn}_{0.2}\text{Cd}_{0.8}\text{S}$  with a high crystallinity, indicating that the  $\text{Zn}_{0.2}\text{Cd}_{0.8}\text{S}$  nanorods grows along the [001] direction. At the same time, the lattice spacing of 3.13 Å in the nanosheet matched well with the (002) plane of hexagonal ZnS. The result is in good agreement with XRD. From elemental mappings in Fig. 4c–f, it is found that Zn and S were well-distributed, while Cd mainly gathered in nanorod area, the results confirm that #2 and #3 were composed of both ZnS nanosheets and  $\text{Zn}_x\text{Cd}_{1-x}\text{S}$  nanorods, and  $\text{Zn}_x\text{Cd}_{1-x}\text{S}$  nanorods were sparsely distributed on the surface of ZnS nanosheets.

Therefore, from the XRD patterns, SEM and TEM images show that samples #2 and #3 were composed with  $\text{ZnS}/\text{Zn}_x\text{Cd}_{1-x}\text{S}$  heterostructures with ZnS nanosheets decorated with  $\text{Zn}_x\text{Cd}_{1-x}\text{S}$  nanorods.

The PL spectra were carried out to study the properties of electrons' recombination light emission. As displayed in Fig. 5a, a wide peak from 400 to 700 nm was observed for all the samples, while ZnS (sample #1) has the strongest PL intensity, which was ascribed to the sulfur-related, zinc-related vacancies and the surface states.<sup>26,27</sup> A decrease in the fluorescence intensity in samples #2 and #3 implies the suppressed recombination of photogenic electron–hole pairs upon attaching  $\text{Zn}_x\text{Cd}_{1-x}\text{S}$  nanorods on ZnS nanosheets, and an increased photogenerated electron–hole separation, and the shift of peak position probably attributed to the transition at the interface.

UV-vis-NIR absorption spectra (Fig. 5b) reveal that samples #1 and #4 have only a single typical absorption edge, while two

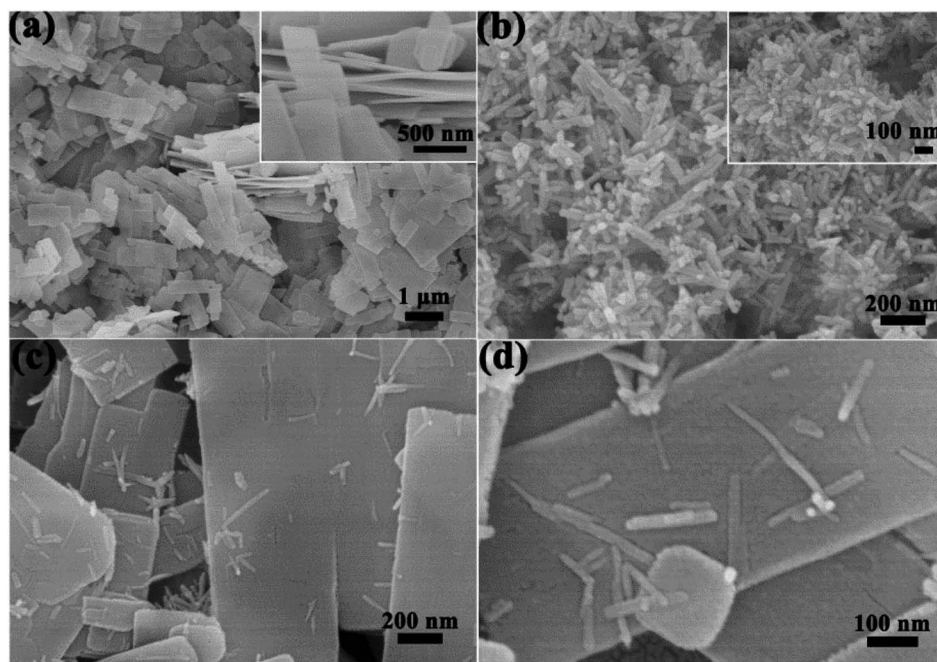


Fig. 3 SEM images of samples #1 (a), #4 (b) and #3 (c and d).



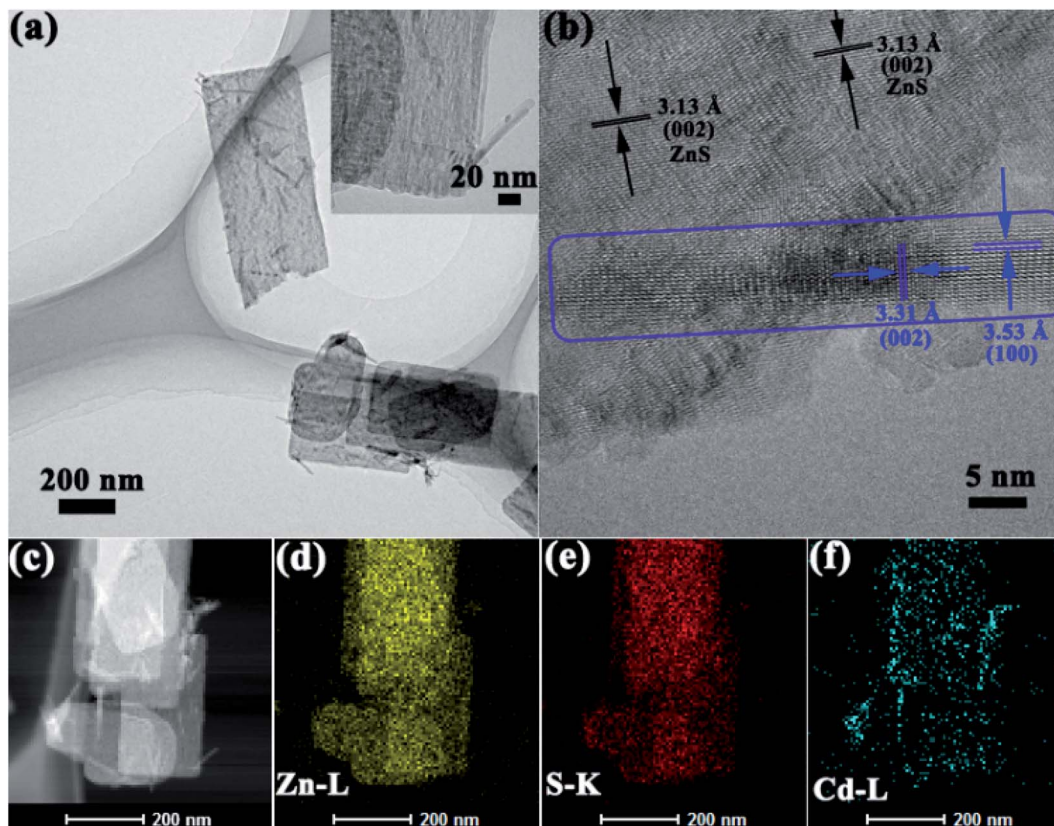


Fig. 4 (a) TEM and (b) HRTEM images, (c–f) elemental mappings of sample #3.

absorption edges can be easily found in samples #2 and #3, one at  $\sim 358$  nm is ascribed to ZnS, the other at  $\sim 500$  nm for sample #2 and  $\sim 525$  nm for sample #3 can be attributed to the intrinsic bandgap absorption of  $\text{Zn}_x\text{Cd}_{1-x}\text{S}$ . The observed two absorption edges in samples #2 and #3 reveal the formation of the  $\text{Zn}_x\text{Cd}_{1-x}\text{S}/\text{ZnS}$  heterostructures. Sample #3 has a stronger absorption and bigger absorption edge than sample #2. Then, the band gap energy is obtained as 3.46, 2.48, 2.35 and 2.19 eV for samples #1–#4, respectively.

The hollow porous structures of the as-prepared samples were elucidated by nitrogen adsorption/desorption measurements to obtain their BET surface areas. As shown in Fig. 5c, the BET surface area was calculated to be  $149.3 \text{ m}^2 \text{ g}^{-1}$  for pure ZnS

nanosheets (#1), and  $103.2$ ,  $82.3$  and  $20.9 \text{ m}^2 \text{ g}^{-1}$  for #2, #3 and #4, respectively. From the results, one can find that pure CdS nanorods possess a low BET surface area, and sample #3 has larger BET surface than sample #2, which is favorable to adsorb molecules.

XPS measurements were used to investigate the surface composition of the four samples, the binding energies have been calibrated using the carbon C 1s peak (285.0 eV) as reference. Zn 2p spectra exhibited two peaks at  $\sim 1020.9$  eV and  $1043.8$  eV, which were assigned to Zn  $2p_{3/2}$  and  $2p_{1/2}$  of  $\text{Zn}^{2+}$ , as shown in Fig. 6a. In comparison with sample #1, the Zn 2p binding energy in  $\text{Zn}_x\text{Cd}_{1-x}\text{S}/\text{ZnS}$  heterostructures (#2, #3) has a slightly shift to higher energy, indicating the decreased

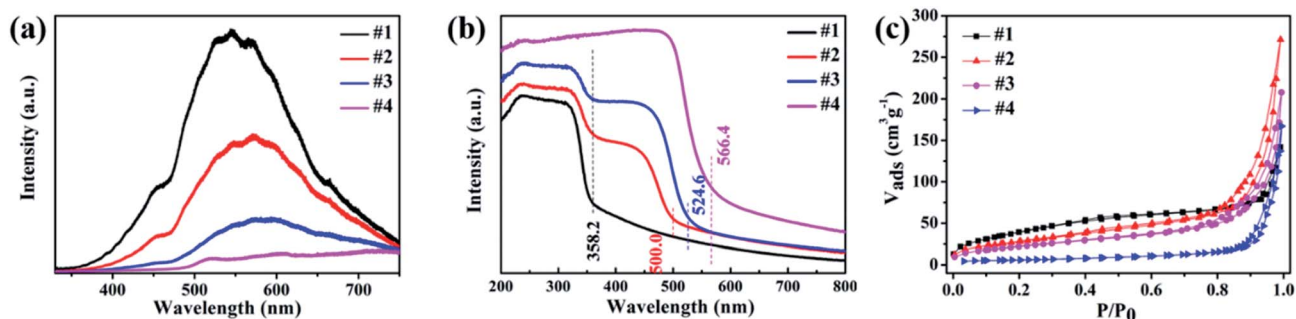


Fig. 5 (a) PL spectra, (b) UV-vis absorption spectra and (c) BET surface areas for the prepared samples ZnS (#1),  $\text{Zn}_x\text{Cd}_{1-x}\text{S}$  (#2, #3) and CdS (#4).



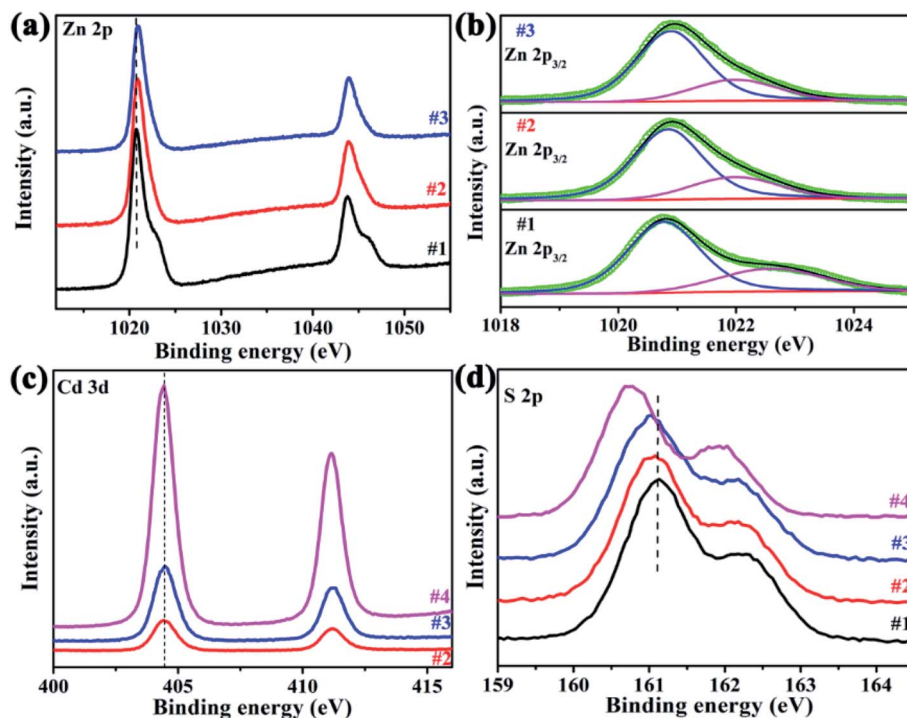


Fig. 6 XPS spectra. (a) Zn 2p, (b) deconvoluted Zn  $2p_{3/2}$  peaks, (c) Cd 3d and (d) S 2p.

electron density around Zn atoms.<sup>28</sup> The Zn  $2p_{3/2}$  core level spectra can be deconvoluted into two peaks, as shown in Fig. 6b, where the lower one is ascribed to zinc in the lattice, while the higher one was related to the Zn<sup>2+</sup> in the sulfur deficient

regions.<sup>29</sup> The relative intensity ratios between the higher one and total zinc are equal to 0.298, 0.280 and 0.285 for sample #1, #2 and #3, respectively. That means, many sulfur vacancies exist in sample #2 and #3, and the concentration is approximate,

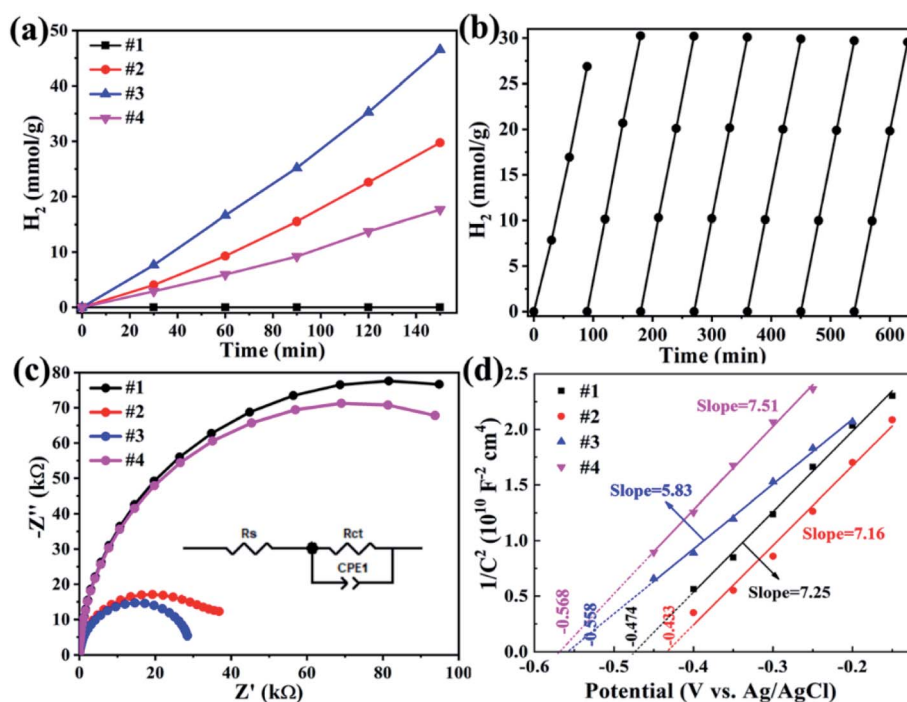


Fig. 7 (a) Hydrogen production of pure ZnS CdS and Zn<sub>x</sub>Cd<sub>1-x</sub>S/ZnS under visible light irradiation (with a 420 nm cut filter), (b) hydrogen production cycle of sample #3; electrochemical performances of all samples, (c) EIS Nyquist plots in a frequency range from 1 to 100 kHz, (d) Mott-Schottky plots at 1000 Hz.



consistent with the EDX results. The Cd 3d XPS spectra are shown in Fig. 6c, in which two peaks at 404.4 eV and 411.1 eV correspond to Cd 3d<sub>5/2</sub> and 3d<sub>3/2</sub>, respectively, and there is no obvious peak shift between pure CdS nanorods and Zn<sub>x</sub>Cd<sub>1-x</sub>S/ZnS heterostructures. For S 2p, the peaks at about 161.0 eV and 162.2 eV are belong to S 2p<sub>3/2</sub> and S 2p<sub>1/2</sub>, as shown in Fig. 6d. With the introduction of Cd, the peaks of S shift to a lower binding energy, implying that the introduction of Cd atoms leads to the formation of Zn<sub>x</sub>Cd<sub>1-x</sub>S. The above XPS analyses demonstrated that the electron transformation from Zn and Cd to S in Zn<sub>x</sub>Cd<sub>1-x</sub>S/ZnS heterostructures can increase electron density of active sites and beneficial to the various reactivity.<sup>30,31</sup>

The apparent quantum efficiency was measured using the 300 W xenon lamp with a 420 nm band pass filters and calculated according to eqn (1):

$$\begin{aligned} \text{AQE}(\%) &= \frac{\text{number of reacted electrons}}{\text{number of incident photons}} \times 100\% \\ &= \frac{\text{number of evolved H}_2 \text{ molecules} \times 2}{\text{number of incident photons}} 100\% \quad (1) \end{aligned}$$

The photocatalytic H<sub>2</sub> production was carried out under visible light irradiation (with a 420 nm cut filter) for samples ZnS, CdS and Zn<sub>x</sub>Cd<sub>1-x</sub>S/ZnS, as shown in Fig. 7a. From Fig. 7a, we found that pure ZnS nanosheets exhibit no photocatalytic activity due to the poor visible light absorbability, while ZnS nanosheets decorated with Zn<sub>x</sub>Cd<sub>1-x</sub>S nanorods reveal an enhanced photocatalytic activity. Sample Zn<sub>0.3</sub>Cd<sub>0.7</sub>S/ZnS has a hydrogen production rate of 9.2 mmol g<sup>-1</sup> h<sup>-1</sup>, while Zn<sub>0.2</sub>Cd<sub>0.8</sub>S/ZnS (#3) achieved a H<sub>2</sub> production rate up to 16.7 mmol g<sup>-1</sup> h<sup>-1</sup> with an apparent quantum efficiency of 10% at 420 nm, approximately 1.5 times of the former. The value is nearly three times of pure CdS nanorods (5.9 mmol h<sup>-1</sup> g<sup>-1</sup>).<sup>32</sup> This result is larger than recently reported 10.8 mmol g<sup>-1</sup> h<sup>-1</sup> with Zn<sub>0.5</sub>Cd<sub>0.5</sub>S hybridized with carbon materials as catalysts,<sup>18</sup> but smaller than the value of 23.264 mmol g<sup>-1</sup> h<sup>-1</sup> with Zn<sub>0.5</sub>Cd<sub>0.5</sub>S/dodecahedron ZIF-67 composite.<sup>20</sup> Moreover, different from the reported severe photocorrosion happened in CdS materials,<sup>33-35</sup> the as-prepared Zn<sub>0.2</sub>Cd<sub>0.8</sub>S/ZnS heterostructures have a relatively stable H<sub>2</sub> reproduction with seven consecutive cycles (Fig. 7b), and the initial low rate of hydrogen production was mainly attributed to the activation process of the catalyst.

EIS measurements was carried out to understand the charge transfer process and the electron-hole separation efficiency occurring at the interface between the photoelectrode and electrolyte, as presented in Fig. 7c, the semicircle in the Nyquist plot of each sample corresponds to the parallel combination of R<sub>ct</sub> and CPE1.<sup>36</sup> The resultant values for the components of the EEC model corresponding to the best fit are presented in Table

Table 3 The EEC impedance parameters

Sample	#4	#1	#2	#3
R <sub>s</sub> (kΩ)	22.31	25.32	25.83	23.16
R <sub>ct</sub> (MΩ)	150.8	164.6	38.73	30.50

3. The smaller diameter corresponds to the lower charge transfer resistance. One can find that sample #3 has smaller charge transfer resistance.

Mott-Schottky (M-S) plots were explored to determine the flatband potential (V<sub>fb</sub>) for the four sample, as depicted in Fig. 7d. All curves show positive slopes, corroborating the n-type characteristics. M-S plots can also be used to estimate the electron carrier concentration using the following equation.<sup>37</sup>

$$N_d = \left( \frac{2}{e_0 \epsilon \epsilon_0 A^2} \right) [d(1/C_s^2)/dV]^{-1}, \quad (2)$$

here e<sub>0</sub>, ε and ε<sub>0</sub> are the electron charge with a value of 1.6 × 10<sup>19</sup> C, dielectric constant with a value of 10,<sup>38</sup> and permittivity of vacuum with a value of 8.85 × 10<sup>-12</sup> F m<sup>-1</sup>, respectively. A, V and C<sub>s</sub> are the area of the film electrode, the bias applied on the electrode and surface capacitance, respectively. From Fig. 7d, Zn<sub>0.2</sub>Cd<sub>0.8</sub>S/ZnS (#3) has the smallest value of slope, so it has the highest carrier concentration up to 2.42 × 10<sup>19</sup>, larger than the value of 1.95 × 10<sup>19</sup>, 1.98 × 10<sup>19</sup> and 1.88 × 10<sup>19</sup> cm<sup>-3</sup> for sample #1, #2 and #4, respectively. The higher carrier concentration results from the sulfur vacancy. Therefore, the enhanced H<sub>2</sub> production rate of sample #3 can be ascribed to the improved charge transfer activity. Besides, the flat-band potentials derived from the M-S plots are equal to -0.474, -0.433, -0.558 and -0.568 V versus Ag/AgCl electrode at neutral solution for #1, #2, #3 and #4, respectively, that correspond to the related conduction band versus Ag/AgCl electrode.<sup>39</sup>

In consideration of band gap energy and the conduction band potential versus Ag/AgCl electrode, the energy band alignment can be obtained for sample #3, as shown in Fig. 8. The figure shows that it is type-II heterojunction for Zn<sub>0.2</sub>Cd<sub>0.8</sub>S/ZnS heterostructure. For Zn<sub>0.2</sub>Cd<sub>0.8</sub>S/ZnS heterostructure, under visible-light irradiation (λ > 420 nm), electrons will be excited from the conduction band of ZnCdS nanorods to the valence band due to the lower band gap energy. As the conduction potential of ZnCdS is more negative than that of ZnS, the excited electrons will transit from the conduction band of ZnCdS nanorods to that of

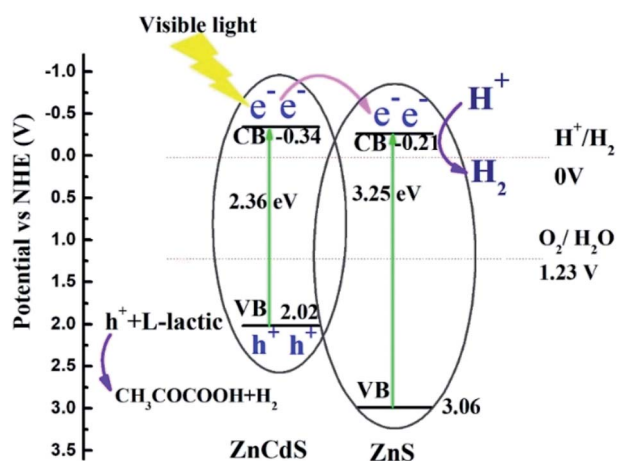


Fig. 8 The schematic illustration for electron charge transfer and H<sub>2</sub> evolution mechanism for the Zn<sub>0.2</sub>Cd<sub>0.8</sub>S/ZnS nanocomposites.



ZnS nanosheets, resulting in the separation of electron-hole pair. Due to the existence of internal fields, the potential barrier at the interface of the heterostructure prevents photogenerated back to ZnCdS nanorods, the photogenerated electrons have a longer lifetime. At the same time, sample #3 has higher electron carriers, meaning that the electrons could be rapidly transmitted to start reduction reaction with H<sub>2</sub>O, producing hydrogen, while the hole sacrificial agent L-lactic acid in the system consumed electrons, forming pyruvic acid and hydrogen (h<sup>+</sup> + L-lactic → CH<sub>3</sub>COCOOH + H<sub>2</sub>). These phenomena effectively promoted the photoinduced electron-hole separation and resulted in the excellent photocatalytic performance. The better photocatalytic H<sub>2</sub> reproduction of sample #3 than sample #2 can be ascribed to the lower charge transfer resistance and the stronger optical absorption, the lower charge transfer resistance will accelerate electrons' transition to participate the reduction reaction with H<sub>2</sub>O; and the stronger visible light absorption of sample #3 can provide more photogenerated electrons for hydrogen production (2H<sup>+</sup> + 2e<sup>-</sup> → H<sub>2</sub>).

## Conclusion

In summary, pure ZnS nanosheets, CdS nanorods and Zn<sub>x</sub>-Cd<sub>1-x</sub>S/ZnS composites were synthesized *via* simple hydrothermal and annealing reaction methods. Zn<sub>x</sub>Cd<sub>1-x</sub>S/ZnS heterostructures were composed with ZnS nanosheets decorated with some Zn<sub>x</sub>Cd<sub>1-x</sub>S nanorods. Type-II Zn<sub>0.2</sub>Cd<sub>0.8</sub>S/ZnS heterostructures have a lower charge transfer resistance, and stronger optical absorption, as well as a well-matched band energy alignment, hence they achieved a H<sub>2</sub> production rate up to 16.7 mmol g<sup>-1</sup> h<sup>-1</sup> under visible light irradiation (λ > 420 nm), approximately 1.5 and 3 times higher than that of Zn<sub>0.3</sub>Cd<sub>0.7</sub>S/ZnS (#2) and pure CdS nanorods (5.9 mmol h<sup>-1</sup> g<sup>-1</sup>) respectively. The strategy for the construction of heterojunctions can provide more perspective for the preparation of photocatalyst in different application field with a simple preparation method.

## Conflicts of interest

The authors declare no competing financial interest.

## Acknowledgements

This work was supported by the National Natural Science Foundation of China (Grant 61604127, 61474096).

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