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Defected ZnWO₄-decorated WO₃ nanorod arrays for efficient photoelectrochemical water splitting†

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The utilization of solar energy in photoelectrochemical water splitting is a popular approach to store solar energy and minimize the dependence on fossil fuels. Herein, defected ZnWO₄-decorated WO₃ nanorod arrays with type II heterojunction structures were synthesized *via* a two-step solvothermal method. By controlling the amount of Zn precursor, WO₃ nanorods were decorated *in situ* with tunable amounts of ZnWO₄ nanoparticles. Characterization confirmed the presence of abundant W⁵⁺ species in the defected ZnWO₄-decorated WO₃ samples, leading to enhanced light absorption and charge-separation efficiency. Therefore, the decorated WO₃ nanorod arrays show much higher photoelectrochemical (PEC) activity than pure WO₃ nanorod arrays. Specifically, the sample with optimal ZnWO₄ decoration and surface defects exhibits a current density of 1.87 mA cm⁻² in water splitting at 1.23 V vs. RHE under 1 sun irradiation (almost 2.36 times higher than that of pure WO₃), a high incident photon-to-current efficiency of nearly 40% at 350 nm, and a relatively high photostability. However, the decoration of WO₃ with too much ZnWO₄ blocks the light absorption of WO₃, inhibiting the PEC performance, even when many defects are present. This work provides a promising approach to rationally construct defected heterojunctions as highly active PEC anodes for practical applications.

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1. Introduction

As a promising method to convert solar energy into storable chemical energy, photoelectrochemical (PEC) water splitting into H₂ and O₂ has attracted wide scientific and technological interest.^{1–4} Since the first discovery of PEC water splitting on TiO₂ by Fujishima *et al.*, many studies focused on fabricating efficient photoanodes have been conducted.^{5–8} Among the various semiconductor materials, n-type tungstic oxide (WO₃) has emerged as a promising photoanode material for PEC water splitting because of its suitable bandgap (*ca.* 2.7 eV), which can utilize a portion of visible light, and its valence band (VB) edge (*ca.* 3 V vs. RHE), which is positive enough to provide a sufficient driving force for oxygen production.^{9,10}

However, one of the greatest challenges hindering the photoactivity of WO₃ is its high rate of electron–hole recombination.¹¹ To solve this problem, the construction of

heterostructures with matching band structures on WO₃ has been shown to be an efficient strategy. Type II heterojunctions, in which the both VB and conduction band (CB) of one semiconductor are either lower or higher than those of the other, are the most common composites. The formation of the proper band offset on the interface can drive photo-induced holes to the semiconductor with higher VB potential, while the electrons are driven to the semiconductor with lower CB potential. Thus, the recombination of charges is alleviated to some degree, and the spatial separation of charge carriers is enhanced.^{12–16} In recent years, several semiconductors have been applied to fabricate type II heterojunctions with WO₃, such as BiVO₄,¹⁷ BiOCl,¹⁸ and tungstate materials (*e.g.*, NiWO₄,¹⁹ CoWO₄,¹⁴ and ZnWO₄ (ref. 20)). Among them, tungstate materials have drawn considerable attention and been widely studied due to their superior compatibility with WO₃ crystals. Using a modified scanning electrochemical microscope method, Leonard *et al.*²⁰ found that the addition of Zn to WO₃ resulted in the best photo-performance among 25 types of metals. Furthermore, several studies on the construction of zinc tungstate and WO₃ have also been carried out.^{13,21,22} However, the PEC activity of WO₃ requires further improvement. The introduction of W⁵⁺ species/oxygen vacancies is a promising approach because the dopant and shallow donors could prevent the recombination of photo-induced electron–hole pairs^{23,24} and create defect levels below the CB, inducing new channels for light absorption and thus improving the PEC performance.^{25,26}

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† Electronic supplementary information (ESI) available: Lattice parameters, unit cell volumes, fitted values of *R_s* and *R_{ct}*, XPS survey spectra of WO₃ and WZ-*x*, fitting of PL data for WZ-*x*, and PEC stabilities of WO₃ and WZ-2. See DOI: 10.1039/c8ra10060h



Herein, we develop a hydrothermal method with two steps, the formation of WO_3 nanorod arrays (NAs) and the reaction of Zn precursor with WO_3 NAs to form ZnWO_4 nanoparticles (NPs), to finally synthesize ZnWO_4 -decorated WO_3 NAs ($\text{WO}_3/\text{ZnWO}_4$, see Scheme 1). Importantly, the insertion of Zn atoms into the WO_3 NA crystals leads to distortion and forms abundant W^{5+} species. By controlling the amount of Zn precursor, tunable amounts of ZnWO_4 NPs were crystallized *in situ* on the surfaces of WO_3 nanorods (named WZ- x , where x reflects the content of ZnWO_4 NPs). Characterization confirmed the presence of abundant defects on the surface of ZnWO_4 . Therefore, with its continuous interface, the $\text{WO}_3/\text{defected ZnWO}_4$ NA shows a much higher PEC activity than that of pure WO_3 NA. In particular, sample WZ-2 exhibits a current density as high as 1.87 mA cm^{-2} in PEC water splitting at 1.23 V vs. RHE under 1 sun irradiation (almost 2.36 times higher than that of pure WO_3), a high incident photon-to-current efficiency (IPCE) of nearly 40% at 350 nm, and a relatively high photostability.

2. Experimental

2.1. Materials

Ammonium metatungstate, $(\text{NH}_4)_6(\text{H}_2\text{W}_{12}\text{O}_{40}) \cdot x\text{H}_2\text{O}$, was obtained from Innochem Chemicals. Hydrochloric acid (36–38%), H_2O_2 (30%), acetone and absolute ethanol were obtained from Tianjin Yuanli Chemical Institute. Zinc acetate ($\text{Zn}(\text{CH}_3\text{COO})_2$) was obtained from J&K Scientific. All reagents were analytical grade and used without further purification. Milli-Q ultrapure water ($>18 \text{ m}\Omega \text{ cm}$) was used in all experiments.

2.2. Sample preparation

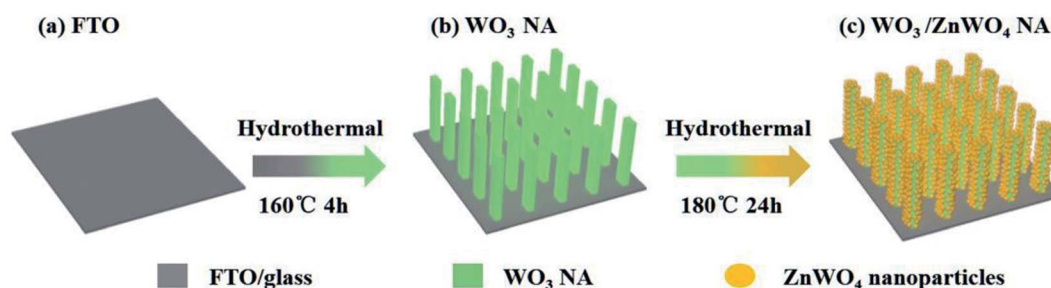
2.2.1 Synthesis of WO_3 nanorod arrays. Glass substrates coated with fluorine-doped tin oxide (FTO) were cut into slides with dimensions of $1 \text{ cm} \times 5 \text{ cm}$ and washed sequentially with acetone, absolute ethanol and deionized water for 30 min each while under ultrasonication. In a typical preparation process, 1 g of $(\text{NH}_4)_6(\text{H}_2\text{W}_{12}\text{O}_{40}) \cdot x\text{H}_2\text{O}$ was dissolved in 95 mL ultrapure water followed by the addition of 3 mL hydrochloric acid (36–38%) and stirring for 5 min. Subsequently, 2 mL of H_2O_2 (30%) was added followed by vigorous stirring for 1 h. The as-prepared solution (20 mL) was then transferred into a Teflon-lined autoclave (50 mL) in which the FTO substrate had been

placed at an angle against the wall of the Teflon liner with the conducting side facing down. Hydrothermal synthesis was conducted at 160°C for 4 h. The resulting WO_3 NA/FTO was rinsed with ultrapure water and dried at 60°C for 12 h. Finally, the obtained sample was annealed at 500°C in air for 1 h at a heating rate of $5^\circ \text{C min}^{-1}$.

2.2.2 Synthesis of $\text{WO}_3/\text{defected ZnWO}_4$ nanorod arrays (NAs). $\text{WO}_3/\text{defected ZnWO}_4$ NAs were synthesized from the above WO_3 NA/FTO *via* a hydrothermal growth process using only Zn precursor. Briefly, WO_3 NA/FTO was added to a solution of 20 mL ethanol containing zinc acetate at a concentration of $c_x \text{ mM}$ (c_1, c_2, c_3). The solution was then solvothermally treated in a Teflon-lined autoclave at 180°C for 24 h. Subsequently, the obtained film was washed with absolute ethanol, dried at 60°C overnight, and annealed at 500°C in air for 1 h (at a heating rate of $5^\circ \text{C min}^{-1}$). The molar concentration of $\text{Zn}(\text{CH}_3\text{COO})_2$ in the prepared solution was fixed as $c_1 = 3.125 \text{ mM}$, $c_2 = 6.25 \text{ mM}$ or $c_3 = 12.5 \text{ mM}$, and the obtained films were named WZ- x (WZ-1, WZ-2, or WZ-3, respectively).

2.3. Characterization

Crystal structures were characterized by X-ray diffraction (XRD) using a Rigaku D/MAX-2500 diffractometer equipped with $\text{Cu K}\alpha$ radiation at 40 kV and 140 mA at a scanning rate of 5° min^{-1} . Raman spectra were obtained using a Raman spectrometer (DXR Microscope) with a green semiconductor laser (532 nm) as the excitation source. Scanning electron microscopy (SEM) images were recorded using a field-emission scanning electron microscope (Hitachi, S-4800). High-resolution transmission electron microscopy (TEM) observations were made with a Tecnai G² F-20 transmission electron microscope. X-ray spectroscopy (EDX) elemental maps were obtained using an EDX system attached to the TEM instrument. X-ray photoelectron spectroscopy (XPS) was carried out using a PHI-1600 X-ray photoelectron spectrometer equipped with $\text{Al K}\alpha$ radiation. The binding energy was calibrated by the C1s peak (284.8 eV). Ultraviolet-visible (UV-Vis) diffuse reflectance spectroscopy (DRS) was conducted with a Shimadzu UV-2600 spectrometer equipped with a 60 mm-diameter integrating sphere using BaSO_4 as the reflectance. Steady-state photoluminescence (PL) spectra were recorded using a Hitachi F-4600 instrument with excitation at 325 nm.



Scheme 1 Synthetic process of $\text{WO}_3/\text{defected ZnWO}_4$ nanorod arrays (NAs).



2.4. Measurement of photoanode PEC performance

The PEC properties of the as-prepared WZ-*x* samples were evaluated in a typical three-electrode quartz cell with a Pt wire counter electrode and a saturated Ag/AgCl reference electrode in 0.5 M Na₂SO₄ electrolyte. The illumination source was a 300 W Xe arc lamp (100 mW cm⁻², PLS-SXE300, Beijing Trusttech. Co. Ltd). Linear sweep voltammetry (LSV) was conducted by sweeping the potential to the positive direction at a scan rate of 10 mV s⁻¹. Based on the Nernst equation, E (vs. RHE) = E (vs. Ag/AgCl) + 0.059 × pH + 0.197,²⁷ the potential vs. Ag/AgCl reference electrode was converted to the potential vs. RHE.

The PEC stability (I - t curve) was evaluated at a potential of 0.4 V vs. Ag/AgCl under AM 1.5 G simulated sunlight irradiation. Electrochemical impedance spectroscopy (EIS) measurements were recorded with a sinusoidal ac perturbation of 10 mV applied over the frequency range of 1–10⁵ Hz.

The IPCE was calculated according to eqn (1):²⁸

$$\text{IPCE}(\lambda) = (1240 \times I)/(\lambda \times J_{\text{light}}) \times 100\%, \quad (1)$$

where λ is the wavelength (nm), I is the photocurrent density (mA cm⁻²), and J_{light} is the incident power illumination density of monochromatic light (mW cm⁻²).

Supposing 100% faradaic efficiency, the applied bias photon-to-current efficiency (ABPE) was calculated by eqn (2):²

$$\text{ABPE} = I \times (1.23 - V_{\text{bias}})/J_{\text{light}} \times 100\%, \quad (2)$$

where V_{bias} is the applied potential, and J_{light} is the incident illumination power density (100 mW cm⁻²).

3. Results and discussion

3.1. Crystal structure and morphology

As shown in the XRD patterns (Fig. 1a), the pure WO₃ NA film shows sharp and well-defined diffraction peaks corresponding to a typical monoclinic WO₃ phase (JCPDS no. 43-1035).²⁹ Compared to the pure WO₃ NAs, the WZ-*x* samples show new diffraction peaks at 18.87°, 30.53° and 36.28°, respectively corresponding to the (100), (111) and (021) crystal planes of pure monoclinic sanmartinite ZnWO₄, which has a space group of $P2_1/c$ with C_4 symmetry and two molecules per unit cell ($Z = 2$;

JCPDS no. 15-0774).³⁰ As shown in Fig. 1b, the diffraction peaks of ZnWO₄ were slightly shifted to higher 2θ values as the content of Zn increased, resulting in smaller interplanar distances in ZnWO₄. This may be attributed to the existence of defects formed during the synthetic process with the insertion/substitution of Zn atoms on the surface of WO₃. The results in Table S1 (ESI†) indicate slight variations in the lattice parameters and cell volumes of ZnWO₄ (compared with JCPDS card no. 15-0774). This suggests the distortion of the octahedral [ZnO₆] and [WO₆] clusters, generating crystal defects in the ZnWO₄ lattice.³¹

The crystal structures of the as-prepared photoanodes were also verified by Raman spectroscopy (Fig. 1c). Pure WO₃ exhibits six main Raman peaks located at *ca.* 74, 136, 274, 328, 718 and 808 cm⁻¹, which are attributed to the active Raman scattering modes of monoclinic WO₃.^{23,32} The strong Raman peaks at 718 and 808 cm⁻¹ are assigned to the W–O stretching modes, while the two other peaks at 274 and 328 cm⁻¹ are related to bending vibrations (W–O–W).²⁸ An extra peak at 906 cm⁻¹ is observed for WZ-*x* samples and can be ascribed to symmetric stretching ($\leftarrow \text{O} \leftarrow \text{W} \rightarrow \text{O} \rightarrow$), which is contributed to by the newly generated ZnWO₄ crystals.³³ The intensity of the peak at 906 cm⁻¹ increases gradually with increasing Zn amount (from WZ-1 to WZ-3). Furthermore, the synthesized crystals exhibit a relatively broad vibrational mode, indicating short-range structural disorder.³⁴

The surface morphologies of the WZ-*x* samples were characterized by SEM and TEM, as shown in Fig. 2. WO₃ exhibits a uniform rod-like morphology. The rods are cuboid shaped with lengths of 500 nm to 1.5 μm, widths of 200–400 nm, and heights of *ca.* 2.4 μm (Fig. 2a and b). After secondary solvothermal crystallization and thermal calcination (WZ-*x* samples), the surface layers of the WO₃ arrays are converted to ZnWO₄ NPs, forming a well-contacted heterojunction structure (Fig. 2c–h). Compared to the pristine WO₃ NAs, the WZ-*x* samples show much rougher surfaces after the formation of ZnWO₄ NP layers. Compared to the other WZ-*x* samples, the lowest concentration of Zn precursor (3.125 mM; WZ-1) resulted in relatively fewer ZnWO₄ NPs with crystal sizes of *ca.* 20–40 nm. When the Zn precursor concentration was increased to 6.25 mM (WZ-2), the amount of ZnWO₄ NPs increased along with the crystal size (*ca.* 50 nm). Further increasing the concentration of

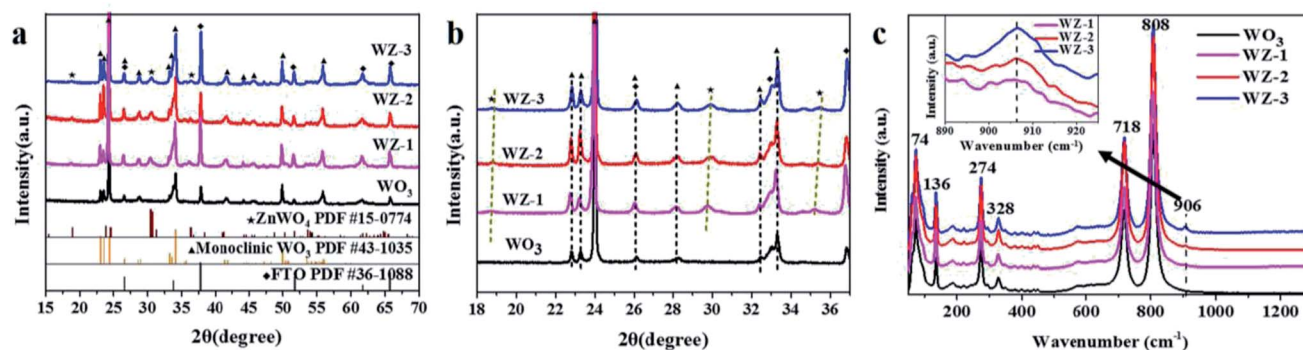
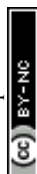


Fig. 1 XRD patterns (a and b) and Raman spectra (c) of pure WO₃ and WZ-*x* NAs.



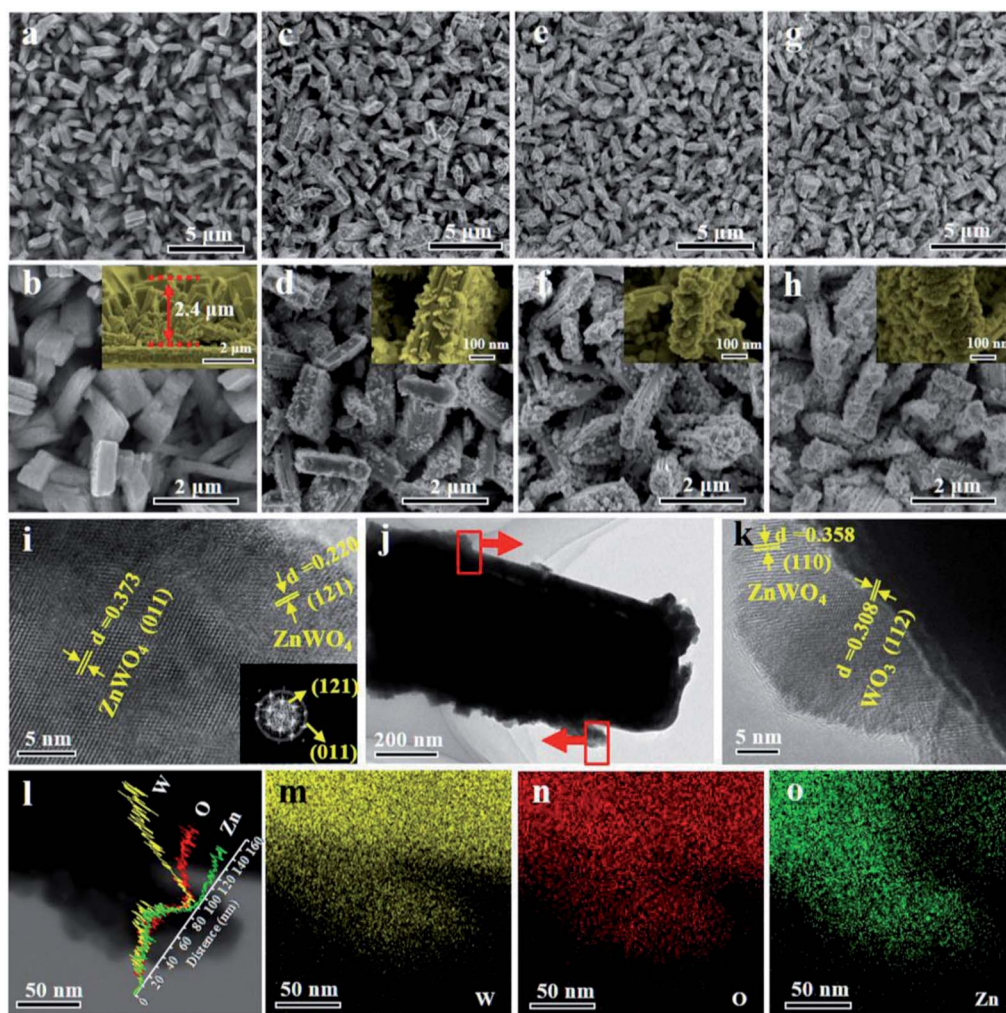


Fig. 2 SEM images of WO_3 (a and b), WZ-1 (c and d), WZ-2 (e and f) and WZ-3 (g and h). TEM images (i–k) and EDX element line scans (l) and maps (m–o) of WZ-2. The inset in (b) is the cross-sectional view of a WO_3 NA.

Zn precursor to 12.5 mM (WZ-3) caused the ZnWO_4 NPs to become much larger and denser, almost covering the surface of WO_3 completely. This dense coverage might inhibit optical absorption and mass transfer for the PEC reaction.

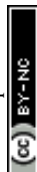
The moderate coverage of ZnWO_4 on WO_3 in sample WZ-2 should promote optical absorption and facilitate the separation of generated electron–hole pairs for the constructed heterojunction. Moreover, this hierarchically structured film can provide sufficient surface active sites to effectively enable reactions at the interface between the photoanode and electrolyte, thus improving the PEC performance.³⁵

The TEM images (Fig. 2i–k) confirm the formation of $\text{WO}_3/\text{ZnWO}_4$ NAs. The spacing of 0.308 nm is in good agreement with the interplanar spacing of the WO_3 (112) plane,³⁶ while the spacings of 0.358, 0.373 and 0.220 nm can be indexed to the (110), (011) and (121) planes of monoclinic ZnWO_4 .³⁷ The corresponding fast Fourier transform (FFT) pattern (inset in Fig. 2i) clearly reveals that the ZnWO_4 NPs are highly crystalline. It is worth noting that a compact interface in the $\text{WO}_3/\text{ZnWO}_4$ heterojunction was formed (Fig. 2k). This can be attributed to

the *in situ* reaction of the Zn precursor with WO_3 NAs, which is beneficial for the migration of photogenerated carriers between the two composites. Meanwhile, the elemental line scanning profiles (Fig. 2l) confirm that W and O are dispersed in the nanorods and branched ZnWO_4 regions, while Zn exists only in the surrounding ZnWO_4 regions. This further verifies that WO_3 was decorated with well-defined defected ZnWO_4 in sample WZ-2. Similarly, the EDX maps (Fig. 2m–o) also confirm the primary distribution of W (and O) and Zn in the rod core and on the surface, respectively.^{11,38}

3.2. Chemical state

XPS was conducted to detect the surface chemical states of the as-prepared photoanodes. The survey spectra (Fig. S1a, ESI†) show the presence of W, O and Zn in the pure WO_3 and WZ-*x* samples without any impurities. In the high-resolution O 1s XPS spectra (Fig. S1b, ESI†), the peak with a binding energy of 530.5 eV is attributed to O–W and O–Zn bonds.^{39,40} The Zn 2p XPS spectra (Fig. 3a) of the WZ-*x* samples show similar characteristic Zn 2p_{3/2} and Zn 2p_{1/2} peaks centered at 1021.9 and



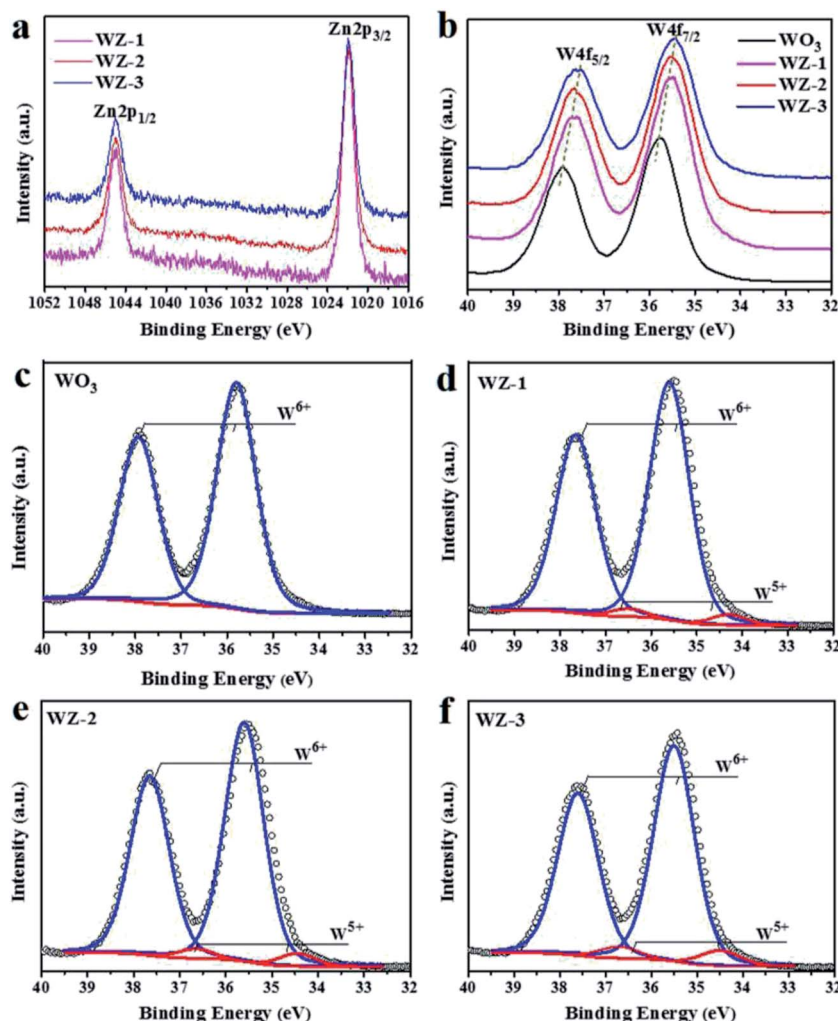


Fig. 3 XPS spectra of WO_3 , WZ-1, WZ-2 and WZ-3: Zn2p (a) and W4f (b), and the fitted W4f XPS spectra of WO_3 (c), WZ-1 (d), WZ-2 (e) and WZ-3 (f).

1045.0 eV, respectively. These energies are slightly higher than those reported in the literature (1021.6 and 1044.7 eV, respectively),⁴⁰ indicating that fewer electrons locate in the Zn atoms of WZ-*x* compared to in pristine ZnWO_4 . Because Zn has a lower electronegativity (1.65) than W (2.36), the shift in binding energy confirms the close interaction between WO_3 and ZnWO_4 . As shown in Fig. 3b, the WZ-*x* samples exhibit obvious low-energy shifts in W 4f binding energy compared to pure WO_3 . For samples WZ-2 and WZ-3, a 0.3 eV negative shift in the W 4f binding energy is observed. This indicates that the formation of WZ-*x* heterojunctions increases the outer electron cloud density of the W atom, which results from W–O–Zn bonding and/or the presence of surface defects.⁴¹

Pristine WO_3 shows only two symmetrical sharp peaks of W 4f (Fig. 3c), which is typical of W^{6+} , while the W 4f peak of WZ-*x* can be divided into four peaks (Fig. 3d–f). The strong peaks at 35.4 and 37.6 eV correspond to W^{6+} , while the weak peaks at 34.5 and 36.7 eV correspond to W^{5+} .²⁶ By fitting the XPS peak of W 4f, the contents of W^{5+} in samples WZ-1, WZ-2 and WZ-3 are estimated to be 3.9%, 4.9% and 7.2% (atomic percentage),

respectively, and the content of W^{5+} is positively correlated with the amount of ZnWO_4 .

3.3. Optical properties

The UV-Vis DRS spectra of the WO_3 and WZ-*x* photoanodes are shown in Fig. 4a. For pristine WO_3 , the absorption edge appears at *ca.* 460 nm, consistent with its intrinsic band-gap absorption.⁴² After coating with ZnWO_4 NPs, the absorption intensity is obviously enhanced in the ultraviolet region compared to pure WO_3 . Among the WZ-*x* samples, WZ-2 achieves the best absorption due to the moderate decoration of defected ZnWO_4 on WO_3 in this sample. The enhanced optical absorption should be related to the enhanced light scattering by the rough surface. Moreover, the W^{5+} species in ZnWO_4 create defect levels below the CB, giving rise to absorption in the far-visible and/or infrared regions and producing tailing at the onset of absorption.²¹ The high-resolution VB XPS spectra of WO_3 and WZ-*x* were recorded to determine the VB edge. As shown in Fig. 4b, compared to pure WO_3 (2.8 eV),⁴³ the VB positions of WZ-*x* are clearly shifted toward a lower energy (0.1–0.2 eV),



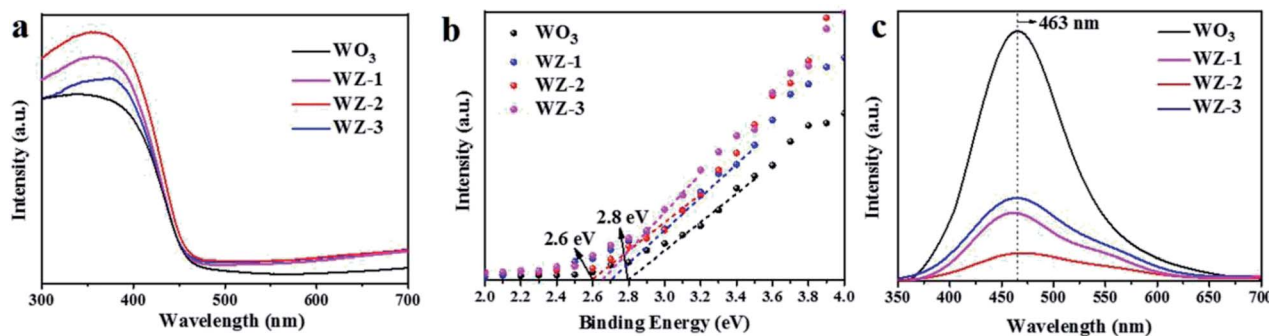


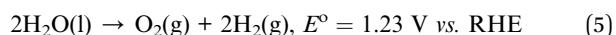
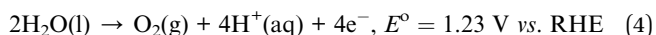
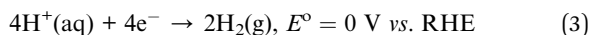
Fig. 4 UV-Vis DRS spectra (a), high-resolution VB XPS spectra (b), and photoluminescence (PL) spectra (c) of WO₃ and WZ-*x*.

indicating that the ZnWO₄ NPs have a lower VB edge (and lower CB on the basis of bandgap data¹⁶) than WO₃, which benefits the formation of type II heterojunctions.²⁶

The recombination of photogenerated charge carriers was investigated through PL emission measurements. Generally, a higher PL intensity indicates a higher recombination rate of photoexcited electrons and holes, while a lower PL intensity indicates a lower recombination rate.¹⁴ Fig. 4c shows that pure WO₃ and WZ-*x* exhibit a major PL emission peak centered at around 463 nm due to the intrinsic feature of WO₃ and/or ZnWO₄. This peak is ascribed to the radiative decay of self-trapped excitons in the crystals.^{35,44} However, the PL intensities of WZ-*x* are clearly lower than that of pristine WO₃, indicating a much greater charge-separation efficiency of photoexcited electron-hole pairs. For surface ZnWO₄, the PL peak at *ca.* 460 nm (Fig. S2, ESI†) is mainly caused by the charge transition between the O 2p orbitals and the empty d orbitals of the central W⁶⁺ ions in the WO₆²⁻ complex, while the yellow-red emission is extrinsic and may be related to W⁵⁺.⁴⁴ The defects induce the new energy states in the bandgap, which may benefit the separation efficiency of e⁻-h⁺ pairs.³¹ Fitting the PL peak indicated that the amount of W⁵⁺ species in WZ-*x* gradually increased with increasing Zn precursor concentration, consistent with the XPS results. However, for sample WZ-3, the total coverage of ZnWO₄ on the WO₃ surface will block light absorption and significantly inhibit the activity of WO₃, even when a large number of defects are present. Thus, WZ-2 with abundant defects and a heterojunction structure exhibits the highest separation efficiency of photogenerated charge carriers.⁴⁵

3.4. PEC performances of photoanodes

PEC catalysis is emerging as a promising method for solar water splitting (hydrogen and oxygen generation).⁹ In PEC water splitting, the oxidation and reduction processes are separated into two half-cell reactions [eqn (3) and (4)], with eqn (5) showing the overall reaction:



To identify the PEC performances of the pristine WO₃ and WZ-*x* photoanodes, PEC measurements were performed under simulated sun light illumination (100 mW cm⁻²) in 0.5 M Na₂SO₄ electrolyte. As shown in Fig. 5a, no current was detected for pure WO₃ in the dark (WO₃-dark). Under light illumination, the photocurrent density of pure WO₃ reaches 0.79 mA cm⁻² at an applied bias 1.23 V vs. RHE. The ZnWO₄-coated samples WZ-1 and WZ-3 show enhanced photocurrent densities of 1.44 and 1.24 mA cm⁻², respectively, while WZ-2 exhibits the highest photocurrent density of *ca.* 1.87 mA cm⁻² at 1.23 V vs. RHE, nearly 2.36 times higher than that of pure WO₃. The PEC performances of recently reported WO₃-based photoanodes are summarized in Table S2 (ESI†). Notably, WZ-2 shows a higher PEC current than the previously reported photoanodes. Moreover, the on-set potentials of the WZ-*x* photoanodes (0.70 V vs. RHE for WZ-1, 0.68 V for WZ-2, and 0.75 V for WZ-3) are also lower than that of pure WO₃ (0.77 V vs. RHE). Based on the LSV results in Fig. 5a, WZ-2 achieved the maximum ABPE in this study (0.18% at 1.05 V vs. RHE; Fig. 5b), compared to only 0.07% for pure WO₃.

The IPCE for PEC water splitting was carried out at 1.2 V vs. RHE, and the results are shown in Fig. 5c. The wavelengths of the initial light response are below 470 nm for the pure WO₃ and WZ-*x* photoanodes and IPCE increases gradually as the irradiation wavelength decreases. Based on the IPCE values, the PEC decreases in the following order: WZ-2 > WZ-1 > WZ-3 > pure WO₃. The IPCE value of WZ-2 reaches 40% at 350 nm. Sample WZ-2 also possesses relatively good photostability, with only a slight loss after long-term continuous illumination (Fig. S3, ESI†). After continuous PEC testing (over 3 h), WZ-2 shows nearly the same XRD peaks (Fig. S4a, ESI†) and surface morphology (Fig. S4b, ESI†) as those of a fresh sample, again confirming the relatively good photostability of WZ-2.

For the WO₃ photoanode, the occurrence of hole-capture reactions (*e.g.*, 2H₂O + 2h⁺ → H₂O₂ + 2H⁺) on the surface is responsible for the corrosion of WO₃ *via* the formation of peroxotungstates, which hinders charge transfer at the WO₃/electrolyte interface. In WZ-*x* samples, the decoration of WO₃ with defected ZnWO₄ forms a type II heterogeneous structure. Since ZnWO₄ possesses higher VB positions than WO₃, the photo-induced holes will transfer rapidly to the ZnWO₄ NPs, preventing the accumulation of holes on WO₃ and preventing the photocorrosion of WO₃. In addition, ZnWO₄ has better



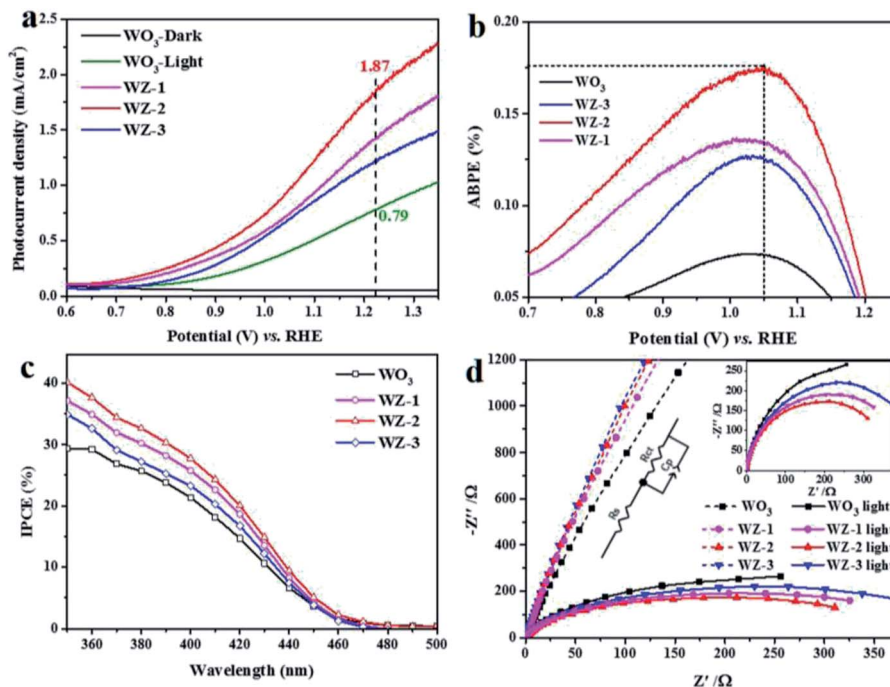
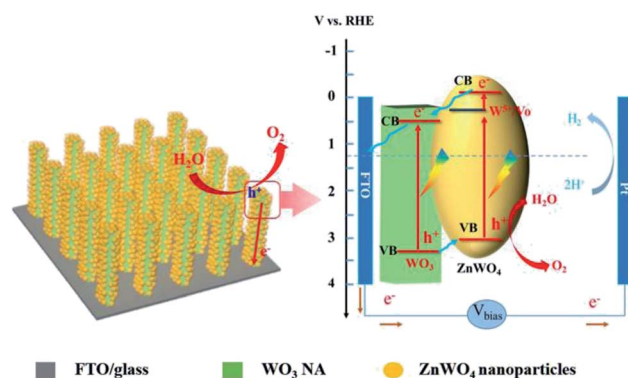


Fig. 5 PEC properties of WO_3 and WZ- x photoanodes in a 0.5 M Na_2SO_4 aqueous solution at pH = 6.8 under light illumination (100 mW cm^{-2}). J - V curves (a), applied bias photon-to-current efficiency (ABPE) (b), incident photon to current efficiency (IPCE) at 1.2 V vs. RHE (c), and electrochemical impedance spectroscopy (EIS) spectra recorded in the dark and under light irradiation (d). The inset in (d) shows the EIS fitting model. R_s , series resistance; R_{ct} , charge-transfer resistance; C_p , capacitive reactance.

chemical stability and photocorrosion resistance than WO_3 .^{20,31} Therefore, the decoration of WO_3 with ZnWO_4 benefits the stability of the photoanode.

To obtain more insight into the charge-transfer kinetics of the photoanodes, EIS measurements were carried out. The arc radii of the Nyquist plots can be used to evaluate the charge-transfer resistance at the semiconductor/electrolyte interface, with a smaller arc radius implying a smaller charge-transfer resistance.⁴⁶ Based on the EIS plots (Fig. 5d), the light irradiation significantly decreases the arc radius of the photoanode, and the effect is more obvious for WZ- x than for pure WO_3 . This suggests that WZ- x exhibits more efficient light absorption and conversion than pure WO_3 . The charge-transfer resistances of the photoanodes were calculated by fitting the EIS spectra (see the inset in Fig. 5d), where R_s and R_{ct} represent the series resistance and interfacial charge-transfer resistance across the electrode/electrolyte interface, respectively.⁴⁷ The fitted values of each component are listed in Table S3 (ESI†). The R_s values for all photoanodes are similar, indicating that the series resistance effect is negligible. In the dark, R_{ct} decreases as the amount of ZnWO_4 NPs decreases ($R_{ct} = 35\,552 \, \Omega$ for WZ-3, $34\,638 \, \Omega$ for WZ-2, $33\,858 \, \Omega$ for WZ-1, and $31\,347 \, \Omega$ for pure WO_3). However, under light irradiation, the R_{ct} values of all of the photoanodes are obviously decreased, and the order is also altered ($R_{ct} = 592 \, \Omega$ for WO_3 , $488.5 \, \Omega$ for WZ-3, $422.4 \, \Omega$ for WZ-1, and $391.3 \, \Omega$ for WZ-2, consistent with the PEC performances of the photoanodes). Notably, the construction of WZ- x heterojunctions benefits charge transfer under light irradiation, unless the coating of defected ZnWO_4 is too thick (WZ-3).

Scheme 2 shows the proposed process of charge separation and transfer for WO_3 /defected ZnWO_4 NAs. As reported, the CB and VB energies of WO_3 are 0.64 and 3.3 V, respectively, while the corresponding values for ZnWO_4 are -0.1 and 3.05 V, respectively.^{13,20} As the shallow donors, the surface W^{5+} species in ZnWO_4 will introduce a new band level below the CB (*ca.* 0.30 V, calculated based on the peak emission wavelength in the PL spectrum). Therefore, the matched band structures benefit the formation of interface heterojunctions between WO_3 and defected ZnWO_4 in the WZ- x samples. Under light illumination, the electrons are excited from the VBs of WO_3 and ZnWO_4 to their own CBs, leaving positive holes in the VBs. The photo-generated electrons in the CB of the ZnWO_4 NPs would first



Scheme 2 Schematic diagram of the WO_3 @defected ZnWO_4 photoanode and the proposed charge-transfer processes.



migrate to the band of W^{5+} (or related oxygen vacancies) and then to the CB of WO_3 . The accumulated electrons will rapidly transfer from WO_3 to the FTO substrate *via* back contact and finally reach the Pt counter electrode for hydrogen generation. Meanwhile, the photogenerated holes in the VB of WO_3 have a higher probability of transferring to the surface of $ZnWO_4$ because of the internal electric field of the heterojunction. Thus, the holes have a higher probability of reacting with water molecules to generate oxygen gas.

The good PEC performance of sample WZ-2 in this study should be closely related to the heterojunction and surface W^{5+} species. The heterojunction will force the spatial separation of electrons and holes in WO_3 and $ZnWO_4$, respectively, while the surface W^{5+} species will inhibit charge recombination.

4. Conclusions

Using a two-step solvothermal method, we successfully fabricated WO_3 /defected $ZnWO_4$ as a photoanode for efficient PEC water splitting. By controlling the amount of Zn precursor, tunable amounts of $ZnWO_4$ nanoparticles were decorated on WO_3 nanorods. The type II heterojunction with abundant defects enhances the light absorption and charge-separation efficiency. Thus, the WZ-x samples show much higher PEC activities than those of pure WO_3 NA. Among the samples, WZ-2 exhibits the highest photocurrent, with a current density of 1.87 $mA\ cm^{-2}$ in PEC water splitting at 1.23 V *vs.* RHE (almost 2.36 times higher than that of pure WO_3), a high IPCE of *ca.* 40% at 350 nm, and a relatively high photostability. This work paves the way for fabricating highly active PEC anodes with defected heterojunction structures for practical applications.

Conflicts of interest

There are no conflicts to declare.

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