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A novel Z-scheme WO$_3$/CdWO$_4$ photocatalyst has been fabricated with sheet-like tungsten trioxide (WO$_3$) hybridized by rods-like cadmium tungstate (CdWO$_4$) via hydrothermal and chemisorptions method. The as-synthesized WO$_3$/CdWO$_4$ photocatalyst exhibited enhanced photocatalytic efficiency for the degradation of different organic dyes under visible light irradiation. It was found that the photocatalytic performance of the composite WO$_3$/CdWO$_4$ was much higher than that of either WO$_3$ or CdWO$_4$ for the degradation of each organic dye. The highest activity of the composite was recorded for the degradation of MB which was about 7 times greater than pure CdWO$_4$ and 2.3 times than that of pure WO$_3$. The enhanced performance of the photocatalyst was mainly attributed to the increased surface area and introduction of WO$_3$ into the composite sample, which can induce higher adsorption activity for organic dyes and the increased electron-hole separation at the interface of two semiconductors by establishing an inner electric field.

Introduction

Since the discovery of photocatalytic splitting of water by TiO$_2$ in 1972 [1], photocatalysis has become one of the promising water decontamination techniques, in which a semiconductor material is used as a catalyst to degrade the organic dyes or environmental pollutants. Unlike the traditional methods of purification, photocatalysis technology has received significant attention because of its effective removal of structurally stable refractory pollutants and excellent degradation efficiency [2]. However, it has been noticed the practical applications of single-phase photocatalysts are usually restricted due to limited visible-light response and high recombination rate of photoexcited electron-hole pairs which play important role in photocatalytic reaction [3]. Hence, exploring the novel semiconductors with excellent photocatalytic activity and irreplaceable merits for the decomposition of organic pollutants present in water or air has become a research hotspot nowadays, which is necessary for solving many problems of mankind regarding the environmental pollution [4].

To design the efficient photocatalysts is a critical issue that involves light absorption, transfer/separation of photogenerated charge carriers and the subsequent catalysis, etc. During the last few years, various methods such as doping [5], coupling plasmonic resonance [6], size and crystal facet of semiconductors and controlling morphology [7, 8] have been adopted to enhance the photocatalytic performance of photocatalysts. Among these strategies, the construction of heterostructures has been proved more effective for improving photocatalytic efficiency. Therefore, coupling of two kinds of semiconductors with visible light response has recently become very famous as a Z-scheme process [9]. This scheme provides some advantages: (1) efficient charge separation, (2) simultaneously high reduction power and deep oxidation power even with visible light of less energy, and (3) different kinds of photocatalysts can be developed [3, 10]. Further, the synergistic effect of heterojunctions can excellently slow down the recombination and speed up the separation rate of photogenerated electron-hole pairs which results in the enhancement of photocatalytic activity [11]. Thus, Z-scheme/heterojunction photocatalysts [12] have attracted much attention among all other photocatalysts. In the recent years, it has been observed that W-based metal oxides (tungstates) exhibit very good physicochemical properties because of their self-trapped excitons [13] and also it has been confirmed that the tungstate materials show good photocatalytic activity for degradation of contaminants [14]. CdWO$_4$, a layered compound crystallizing in a monoclinic wolframite structure is considered to be an attractive material because of its optical, chemical and structural properties [13, 15]. It has wide band gap larger than the conventional photocatalyst TiO$_2$ [16] thus it seems to indicate limited photo-absorption, however, it could be still excited by light energy higher than its band gap to induce energy-rich electron–hole pairs that consequently degrade the toxic pollutants into nontoxic CO$_2$ and H$_2$O components during the photocatalytic process [17]. But the fast recombination ratios of photogenerated charge carriers and poor response to visible light damage the practical applications of CdWO$_4$. Therefore, significant efforts have been taken into account to overcome such problems. The photocatalytic efficiency of Eu$^{3+}$-doped CdWO$_4$ was found higher than that of undoped sample for degradation of MO [18].
Similarly, the photocatalytic H₂ production rate of CdS-modified CdWO₄ was found about 34 times higher than the pure CdWO₄ [19]. Very recently, Xu et al. [20] fabricated CdWO₄-deposited reduced graphene oxide and studied its enhanced photocatalytic properties by degrading methylene blue under UV light; the results were much higher than single CdWO₄. Generally, the photocatalytic performance of the semiconductors can be enhanced by increasing the separation efficiency of photogenerated charge carriers. Therefore, the combination of two photocatalysts would be proved an ideal system in extending the absorption spectra to visible light region to attain a high separation efficiency of photogenerated charge carriers. However, to date, there is no report on the coupling ofWO₃ with CdWO₄ and effect of this coupling on the photocatalytic efficiency of resulting photocatalyst.

In this article, we fabricated a novel Z-scheme heterostructured WO₃/CdWO₄ photocatalyst for the first time by simple mixing and heating method and studied its enhanced photocatalytic activities for the degradation of organic pollutants under visible-light irradiation.

All the chemicals used for synthesis in this work, were reagent grade and were commercially available and were used without further purification. First the rodlike CdWO₄ was synthesized by hydrothermal method. Typically 5 mmol of Na₂WO₄.2H₂O and 5 mmol of Cd(NO₃)₂.4H₂O were dissolved in 15 ml distilled water and stirred for 30 min, during the stirring aqueous ammonia was added dropwise to the solution to attain a pH value of 9.30. The clear white solution was taken into stainless steel autoclave and heated at 180°C for 24 hours. After washing several times with absolute ethanol and deionized water, a white colour powder was obtained. WO₃ sheets like structure was synthesized by CVD method: 0.5 g tungsten powder was taken into alumina ceramic boat and placed in the furnace. It was heated at 600°C with ramp rate 10°C/min in air flow for 3 hours. The tube was then allowed to cool at room temperature and yellow colour powder was collected. WO₃/CdWO₄ photocatalyst was fabricated as follows; appropriate amounts of WO₃ and CdWO₄ (1:1) were dispersed in 10 ml of ethanol in separate glass beakers and sonificated for 1 hour. The dispersed solutions were then mixed and magnetically stirred for 1 hour and then heated at 80°C for 12 hours. The obtained powder was then calcined at 550°C for 2 hours. The final product was preserved for characterizations and measurements. The phase characterization of as-prepared photocatalyst WO₃/CdWO₄ was done by X-ray diffraction (XRD; Philips X’Pert Pro MPD), using a Cu Kα radiation source (λ = 0.15418 nm) with 2θ from 10° to 80°. The field emission scanning electron microscopy (FESEM) and transmission electron microscope (TEM, H-600-II, and Hitachi) high resolution transmission electron microscopy (HRTEM) and selected area electron diffraction (SAED) pattern were employed to characterize the morphology and composition of the as-prepared sample. The chemical composition of the samples was determined by energy dispersive X-ray (EDX) analysis (Hitachi S-3500). Fourier transform infrared (FTIR) spectra of samples were measured by using Nicolet Avatar-370 spectrometer. The Brunauer–Emmett–Teller (BET) specific surface areas of the sample materials were measured in a Micromeritics nitrogen adsorption instrument HYA2010C2 (Beijing ZhongKe Hui Yu Technology Co., Ltd.). The optical absorption and energy band gap spectra of the samples were examined by UV-VIS-NIR (Hitachi-U4100) spectrophotometer whereas the photoluminescence (PL) spectra were measured with Hitachi FL-4500 fluorescence spectrometer at room temperature.

Experimental

Fabrication of WO₃/CdWO₄ photocatalyst

In order to evaluate the photocatalytic activity of the fabricated photocatalyst, an aqueous solution of each dye MO, RhB and MB (100 mL, 10 mg L⁻¹) was prepared in glass beaker and then 100 mg of the photocatalyst were added in this solution. The glass beaker was covered and kept in darkness under constant magnetic stirring for 30 minutes to obtain adsorption–desorption equilibrium between the photocatalysts and dye solution, and then this solution was exposed to 500 W xenon lamp which was used as a visible light source. After 10 min time intervals, 5 mL of solution was taken and centrifuged to remove the photocatalyst particles before measuring the UV absorption spectra. The filtrates were then analysed by measuring the variations in absorption band maximum (MO 508 nm, RhB 553 nm and MB 663 nm) using a UV-VIS-NIR (Hitachi-U4100) spectrophotometer. To examine the photostability of the photocatalyst, the same materials were centrifuged and dried and used again for degradation by repeating the above mentioned steps for second and third time.

Results and discussion

Phase structures and morphology

X-Ray diffraction (XRD) analysis was employed to characterize the crystal structure and to identify the phase of the as-synthesized photocatalyst WO₃/CdWO₄ as well as pure CdWO₄ and WO₃ as shown in Fig. 1. Fig. 1 (a) shows the typical XRD pattern of CdWO₄ microrods, the well-defined diffraction peaks with relative diffraction intensities can be clearly indexed to a pure monoclinic phase of cadmium tungstate (JCPDS Card No. 87-1114, with lattice constants a = 5.0289 Å, b = 5.8596 Å, c = 5.0715 Å and α = γ = 90°, β = 91.5190°) with distinctive peaks centred at 23.3°, 29.02°, 29.58°, 30.49°, 35.34°, 35.76° and 47.7° match well with (110), (−111), (111), (020), (002), (200) and (220) respectively. Fig. 1 (b) presents the XRD of the WO₃ sheetslike structure. All the diffraction peaks are well matched with the orthorhombic crystalline phase of WO₃, which was in good agreement with the standard card (JCPDS Card No. 89-4479, with lattice parameters a = 7.3612 Å, b = 7.5739 Å, c = 7.7620 Å and α = β = γ = 90°) with distinctive peaks at 23.02°, 23.52°, 24.25°, 33.14° and 34.12° corresponding to (002), (020), (200), (022) and (220) planes respectively. Fig. 1 (c) represents the
XRD pattern of the as-prepared composite WO₃/CdWO₄. It can be noted that the composite sample contains the peaks of both materials with slightly different positions that could be expected after mixing and heating process. The composite sample is purely composed of CdWO₄ and WO₃ that can be further confirmed from the EDX spectrum as shown in Fig. 1 (d). The EDX spectrum shows that the as-synthesized product contains only W, O and Cd elements and no other element or impurity found in the fabricated WO₃/CdWO₄ photocatalyst.

![Fig. 1 (a) XRD patterns of CdWO₄, (b) XRD patterns of WO₃, and (c) XRD patterns of WO₃/CdWO₄ photocatalyst (d) The energy dispersive X-ray (EDX) spectrum of WO₃/CdWO₄ photocatalyst.](image)

The morphology and structure of the as-synthesized materials were characterized by SEM as shown in Fig. 2. The Fig. 2 (a) represents the typical SEM image of WO₃ microsheets that are uniformly distributed over the entire surface at different orientations. Fig. 2 (b) shows the rodlike structure of CdWO₄. It can be seen that the rods are assembled in such a way that it presents a view more like to the leaf of a palm tree. Fig. 2 (c and d) depicts the SEM image of the as-prepared WO₃/CdWO₄ photocatalyst at different magnifications. It can noted that WO₃ microsheets were shrunk when combined with CdWO₄ rods and some of them attached/stuck with the rods as can be seen in a close view in Fig. 2 (d). Although by mixing the two materials, there is no big change observed in their structures but this mixing plays an obvious effect on the optical and photocatalytic properties of the obtained product/photocatalyst.

![Fig. 2 (a) SEM image of sheetslike WO₃, (b) SEM image of rodlike CdWO₄, and (c, d) SEM images of WO₃/CdWO₄ photocatalyst at different magnifications.](image)

![Fig. 3 (a-d) FESEM images of the as-prepared WO₃/CdWO₄ photocatalyst at different magnifications.](image)

![Fig. 4 (a and b) represents the TEM characterization of the composite sample at different magnifications. The rodlike shape of CdWO₄ can be clearly seen here and we can also note in Fig. 4 (a and b) that there are sheets of WO₃ attached with this rod. Fig. 4 (c) shows the HRTEM image of the as-synthesized WO₃/CdWO₄ photocatalyst. It can be clearly noticed that there are two kinds of lattice fringes of two different materials which confirm the interfaces between WO₃ and CdWO₄ (see also Fig. S1 ESI†). From the HRTEM image (Fig. 4c), the spacing between two adjacent lattice fringes was found to be 0.307 nm that corresponds to the (-111)
plane of CdWO₄. Fig. 4 (d) represents the SAED image of the sample.

**FTIR analysis and BET surface area measurements**

To further confirm the composition information of WO₃/CdWO₄ composite, Fourier transform infrared (FTIR) spectra of the samples were measured as displayed in Fig. 5. It shows typical FTIR spectra of pure CdWO₄, pure WO₃ and WO₃/CdWO₄ Z-scheme photocatalyst respectively. The broad absorption peak around 3428 cm⁻¹ in Fig. 5 (a) is associated with H–O vibration for surface hydration layers, and the weak peak around 1633 cm⁻¹ is attributed to H–O–H bonding of the water molecules [22]. The two peaks around 1395 and 1568 cm⁻¹ may be assigned to the symmetrical and asymmetrical vibrations of carboxylate groups respectively. The two strong peaks at 1060 cm⁻¹ and 1140 cm⁻¹ are the characteristics of stretching vibrations of C–O of citric species [23], which confirms the presence of CdWO₄. A set of peaks below 1000 cm⁻¹ shown in Fig. 5 (b) can be ascribed to O–W–O stretching modes of WO₃, while the broad bands in the range 3200-3550 cm⁻¹ could be assigned to stretching vibrations of O–H for WO₃ [24].

![FTIR spectra](image)

**Fig. 5** FTIR spectra of CdWO₄, WO₃ and WO₃/CdWO₄ (a) in full range, and (b) in short range.

To gain the knowledge about the surface area of the as-prepared photocatalyst, the N₂ adsorption/desorption isotherms of all the samples were measured as indicated in Fig. 6. It is well known that high surface area of a photocatalyst is favorable to adsorb organic pollutants from the aqueous solution and increases the contact area between the catalyst and pollutants during the photocatalytic reaction. Therefore, for a material to be a good photocatalyst, its specific surface area is given much importance as it plays significant role during photocatalysis providing more active sites [25]. Fig. 6 shows the typical nitrogen adsorption–desorption curves for CdWO₄, WO₃ and WO₃/CdWO₄. The BET specific surface area of all the samples was calculated from the data points in the pressure range 0 to 1 by the Brunauer–Emmett–Teller (BET) equation [26]. According to these calculations, the BET specific areas of CdWO₄, WO₃ and WO₃/CdWO₄ were estimated as 6.844, 7.950 and 15.803 m²/g respectively. In addition, the BET analysis shows that the WO₃/CdWO₄ photocatalyst has larger specific surface area than both CdWO₄ and WO₃ that possibly makes it beneficial for the decontamination of organic dyes.

![BET analysis](image)

**Fig. 6** Nitrogen adsorption–desorption curves for CdWO₄, WO₃ and WO₃/CdWO₄ respectively.

**Optical absorption properties**

To investigate the optical properties of the fabricated photocatalyst WO₃/CdWO₄, the UV spectra were measured at room temperature. As indicated in Fig. 7 (a), the absorption band edges of CdWO₄ and WO₃ are around 390 and 420 nm while the absorption band edge of WO₃/CdWO₄ is located around 460 nm. It can be seen that the absorption edge of WO₃/CdWO₄ composite sample has shifted significantly towards longer wavelengths as compared to CdWO₄ and WO₃ which means that the absorption edge of WO₃/CdWO₄ photocatalyst is shifted to the lower energy region. The optical band gaps, E₉, of all the three samples were calculated by the well-known equation used in ref. (24). The band gaps of CdWO₄, WO₃ and WO₃/CdWO₄ photocatalyst were estimated as 3.30, 2.77 and 2.97 eV respectively as can be seen from Fig. 7 (b). The band gap of the composite WO₃/CdWO₄ sample exists between CdWO₄ and WO₃. This result reveals that the formation of tightly chemically bonded interfaces between CdWO₄ and WO₃ phases can make WO₃/CdWO₄ composite photocatalyst shift to the lower energy region.
photocatalytic activity of the WO$_3$/CdWO$_4$ under visible light irradiation were calculated as 0.07221 m$^{-1}$, 0.07614 m$^{-1}$ and 0.07982 m$^{-1}$ respectively. It can be seen that WO$_3$/CdWO$_4$ photocatalyst showed the enhanced photocatalytic activity for the degradation of pollutants compared to pure CdWO$_4$ and pure WO$_3$. In the WO$_3$/CdWO$_4$ system, the photocatalyst exhibited the photocatalytic performance about 7 times greater than the bare CdWO$_4$ (0.01155 m$^{-1}$) and about 2.3 times greater than bare WO$_3$ (0.03459 m$^{-1}$) for the case of MB degradation. The enhancement could be ascribed to the introduction of WO$_3$ into the composite sample, which can induce higher adsorption activity for organic dyes and higher migration efficiency of photo-generated charge carriers. Fig. 8 (c) shows the stability WO$_3$/CdWO$_4$, WO$_3$ and CdWO$_4$ for the case of MB degradation. For the recycling process, the catalyst samples were centrifuged, dried at 80°C for 12 hours and then reused for degradation of the same concentration of the dye. The samples were recycled for 3 times and the results showed excellent stability and recycling capability (Fig. 8 c). The materials took almost same time for degradation without any significant loss of efficiency even after 3rd cycle. The stability of the samples for the degradation of RhB and MO are given in Fig.S3 (ESI†). The apparent rate constants of all the samples and relative coefficients are summarized in Table 1. The photocatalytic performance of the as-synthesized photocatalyst in comparison to CdWO$_4$ and WO$_3$ for the degradation of MO and RhB can also be studied from this table.

**Table 1** The values of rate constants CdWO$_4$, WO$_3$, and WO$_3$/CdWO$_4$ for the degradation of MB, RhB and MO respectively.

<table>
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<th>Dye</th>
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In order to express the reaction kinetics of the organic dyes, we have to deal with the following first order equation:

$$-\ln(C/C_0)=kt$$

where $C_0$ and $C$ represents the concentrations of dye in the solution at times $t$ and $t$, respectively, and $k$ is the apparent first-order rate constant. The photodegradation kinetics for CdWO$_4$, WO$_3$ and WO$_3$/CdWO$_4$ over the organic pollutants is shown in Fig. 8 (a) by plotting a graph between $\ln(C/C_0)$ and irradiation time $t$ that are almost straight lines indicating that the photodegradation reaction obeys the first-order kinetics. From Fig. 8 (b), the pseudo-first-order rate constants ($k$) for the degradation of MO, RhB and MB with WO$_3$/CdWO$_4$ under visible light irradiation were calculated as 0.07221 m$^{-1}$, 0.07614 m$^{-1}$ and 0.07982 m$^{-1}$ respectively. It can be seen that WO$_3$/CdWO$_4$ photocatalyst showed the enhanced photocatalytic activity for the degradation of pollutants compared to pure CdWO$_4$ and pure WO$_3$. In the WO$_3$/CdWO$_4$ system, the photocatalyst exhibited the photocatalytic performance about 7 times greater than the bare CdWO$_4$ (0.01155 m$^{-1}$) and about 2.3 times greater than bare WO$_3$ (0.03459 m$^{-1}$) for the case of MB degradation. The enhancement could be ascribed to the introduction of WO$_3$ into the composite sample, which can induce higher adsorption activity for organic dyes and higher migration efficiency of photo-generated charge carriers. Fig. 8 (c) shows the stability WO$_3$/CdWO$_4$, WO$_3$ and CdWO$_4$ for the case of MB degradation. For the recycling process, the catalyst samples were centrifuged, dried at 80°C for 12 hours and then reused for degradation of the same concentration of the dye. The samples were recycled for 3 times and the results showed excellent stability and recycling capability (Fig. 8 c). The materials took almost same time for degradation without any significant loss of efficiency even after 3rd cycle. The stability of the samples for the degradation of RhB and MO are given in Fig.S3 (ESI†). The apparent rate constants of all the samples and relative coefficients are summarized in Table 1. The photocatalytic performance of the as-synthesized photocatalyst in comparison to CdWO$_4$ and WO$_3$ for the degradation of MO and RhB can also be studied from this table.

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Possible mechanism for enhanced photocatalytic activity of Z-scheme WO₃/CdWO₄ photocatalyst

It is well known that there are some crucial factors like adsorption ability, optical absorption, phase structure and separation efficiency of photo-generated charge carriers that influence the photocatalytic activity [27]. However, to fully understand the photocatalytic reaction mechanism occurring during the photodegradation of as-prepared WO₃/CdWO₄ photocatalyst, the band edge positions of the valence band (VB) and conduction band (CB) of both CdWO₄ and WO₃ are necessary to be determined because they have strong relation with the photocatalytic oxidation process of organic compounds. The valence band potentials of CdWO₄ and WO₃ were calculated according to the following empirical equations [28]:

\[ E_{VB} = \chi - E^e + \frac{1}{2}E_g \]  
\[ E_{CB} = E_{VB} - E_g \]

where \( E_{VB} \) and \( E_{CB} \) are the valence band and conduction band edge potentials respectively, \( \chi \) is the electronegativity of the semiconductor, which is the geometric mean of the electronegativity of the constituent atoms. The values of \( \chi \) for CdWO₄ and WO₃ have been calculated as 6.28 eV and 6.59 eV according to literature [29]. \( E^e \) is the energy of free electrons on the hydrogen scale (about 4.5 eV) and \( E_g \) is the band gap energy of the semiconductor. On the basis of band gap positions, the band edge potentials of VB and CB for WO₃ [29] are 3.48 eV and 0.71 eV while for CdWO₄ are 3.43 eV and 0.13 eV respectively. The schematic illustration of energy band structure of WO₃ and CdWO₄ is presented in Fig. 9.

After bringing the two materials CdWO₄ and WO₃ in contact, the electrons from the CB of CdWO₄ started to diffuse into the CB of WO₃ as the CB potential of CdWO₄ is at lower potential than that of WO₃. As a result, negative charges will accumulate in WO₃ near the interface. Simultaneously, the holes from VB of WO₃ move into the VB of CdWO₄ because the VB of WO₃ is at higher potential than that of CdWO₄. This movement of the holes gives rise to a layer of positive charges in CdWO₄ near the interface between two semiconductors. So an electric field is established between the interfaces as shown in Fig. 9. The equilibrium between CdWO₄ and WO₃ is established when the Fermi levels are reached and the further diffusion of electrons from CdWO₄ to WO₃ is stopped. In the presence of visible light irradiation, both CdWO₄ and WO₃ absorb photons of energy equal to or greater than the corresponding band gap energy, that cause to excite the electrons in the VB to the CB and leave holes in the VB of both materials. The excited electrons present in the CB of CdWO₃ are then transferred to the CB of WO₃, and the holes are left in the VB of CdWO₄. The inner established electric field (\( E \)) at the heterojunction interfaces promotes this transfer of photo-induced charge carriers. The photo-induced electron-hole pairs are thus effectively separated because of a junction formed between CdWO₄ and WO₃ interface, resulting in the slow-down recombination photo-generated electron-hole pairs. As a result, the photocatalytic activity of the Z-scheme photocatalyst WO₃/CdWO₄ was improved. This conclusion can be verified by the PL spectra of all the three samples.

![Fig. 9 Schematic diagram of the (a) band energy of CdWO₄ and WO₃ before contact and (b) formation of a p–n junction and the proposed charge separation process of WO₃/CdWO₄ heterostructures under visible-light irradiation.](image)

![Fig. 10 Photoluminescence (PL) spectra of CdWO₄, WO₃ and WO₃/CdWO₄ at room temperature.](image)

In order to investigate the fact of photo-generated electron–hole pairs in the product and to understand the possible mechanism, the photoluminescence emission spectra were measured, since the PL emission spectra originate from the recombination of free charge carriers in a semiconducting material and there is a strong link between PL intensity and the photocatalytic efficiency. Generally, a higher PL intensity shows a higher recombination rate of photo-generated charge carriers which leads to a lower photocatalytic performance. On the other hand, a lower PL intensity shows a lower recombination rate of photo-generated electron–hole pairs, indicating that more photo-generated electrons and holes can participate in the oxidation and reduction reactions and as a result the photocatalytic performance is improve [31]. Fig. 10 shows the PL spectra of rodslike CdWO₄ microstructures, WO₃ microsheets and WO₃/CdWO₄ composite at room temperature in the wavelength range of 270-800 nm with an excitation wavelength of 235 nm. The PL emission peak for pure CdWO₄ centred at 463 nm whereas for pure WO₃ at 427 nm. It has been noticed that the coupling of WO₃ microsheets with rodslike CdWO₄ had significant effect on the PL
intensity of the obtained composite WO$_3$/CdWO$_4$. It can be that the PL intensity of composite dramatically decreased by the introduction of WO$_3$. The intensity peak for WO$_3$/CdWO$_4$ appeared at 476 nm which may be ascribed to the band gap recombination of electron-hole pairs. It can be noted that the addition of WO$_3$ to CdWO$_4$ not only alters the spectral position of the peak but also decreased the relative PL intensity of the as-prepared composite sample. These findings reveal that the heterojunction/synergistic effect between WO$_3$ and CdWO$_4$ plays an important role in the effective electron-hole separation that might be a reason for the superior photocatalytic activity of Z-scheme WO$_3$/CdWO$_4$ photocatalyst under visible light irradiation environment. The efficient separation of photo-generated electron-hole pairs in WO$_3$/CdWO$_4$ photocatalyst was confirmed by the PL emission spectra as it is strongly correlated with the transfer behaviour of photo-generated electron-hole pairs in semiconductors [32]. On the addition of WO$_3$ to CdWO$_4$, the surface recombination of electrons and holes was inhibited, therefore the PL intensity of the final product is decreased which shows that the recombination rate of photo-generated charge carriers in the composite sample has slowed down and as a result higher photocatalytic activity is achieved. The decreased PL intensity of WO$_3$/CdWO$_4$ photocatalyst is thus an obvious evident of high quantum efficiency and high photocatalytic activity. Finally, the increased surface area (15.803 m$^2$ g$^{-1}$) of the composite WO$_3$/CdWO$_4$ as compared to CdWO$_4$ and WO$_3$ played an important role in the enhancement of its catalytic activity. The larger surface area of the catalyst provides more active sites for photocatalytic reaction, resulting in higher catalytic activity.

Conclusion

A novel Z-scheme WO$_3$/CdWO$_4$ photocatalyst has been fabricated by a simple mixing and heating method. The as-prepared WO$_3$/CdWO$_4$ photocatalyst exhibited high photocatalytic efficiency for the degradation of different organic pollutants under visible light irradiation. It has been found that the photocatalytic performance of the composite WO$_3$/CdWO$_4$ was much higher than that of either WO$_3$ or CdWO$_4$ for the degradation of each organic dye. The highest activity of the composite was recorded for the degradation of MB which was about 7 times greater than pure CdWO$_4$ and 2.3 times than that of pure WO$_3$. The enhanced performance of the photocatalyst was mainly attributed to the increased surface area and introduction of WO$_3$ into the composite sample, which can induce higher adsorption activity for organic dyes and the increased electron-hole separation at the interface of two semiconductors by establishing an inner electric field.

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Notes and references

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† Footnotes should appear here. These might include comments relevant to but not central to the matter under discussion, limited experimental and spectral data, and crystallographic data.

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