


 Cite this: *RSC Adv.*, 2019, 9, 41977

Synergetic effect of dual co-catalysts on the activity of BiVO₄ for photocatalytic carbamazepine degradation†

 Beibei Wang,^a Ping Li,^b Chunlei Du,^b Yan Wang,^b Daxin Gao,^b Songtao Li,^b Liying Zhang^b and Fuyu Wen^{ib*}

An efficient visible-light driven three components photocatalyst for carbamazepine (CBZ) degradation has been assembled by co-loading reduction cocatalyst Pt and oxidation cocatalyst Co₃O₄ (MnO_x) on BiVO₄. The apparent rate constant of the three components photocatalyst Pt/BiVO₄/Co₃O₄ for degradation of CBZ is 54 times that of Co₃O₄/BiVO₄ and 2.5 times that of Pt/BiVO₄, which shows a synergetic effect in the photocatalytic activity. The same synergetic effect is also observed for Pt/BiVO₄/MnO_x. The spatial separation of the reduction and oxidation cocatalysts could reduce the recombination of the photogenerated charges, which mainly accounts for the high photocatalytic activity of the three components photocatalyst. The photocatalytic intermediates of CBZ were detected by HPLC-ESI-MS, and a deductive degradation pathway of CBZ was proposed.

 Received 6th September 2019
 Accepted 13th December 2019

DOI: 10.1039/c9ra07152k

rsc.li/rsc-advances

1. Introduction

Pharmaceuticals and personal care products (PPCPs) are persistent organic pollutants which cannot be completely removed by conventional treatment plants.¹ For instance, the removal efficiency of carbamazepine (CBZ) through wastewater treatment plants (WWTPs) is mostly below 10%.² Drug residues in the natural environment eventually affect water quality, ecosystems and human health.³ For example CBZ is reported to be toxic even at concentrations below 100 mg L⁻¹.⁴ Therefore, it is urgent to seek more efficient technologies for the degradation of CBZ-containing waste-water.

Heterogeneous photocatalysis is one of the most ideal techniques because of its advantages of lower energy consumption, no secondary pollution, easy accessibility and so on.⁵ To date, the most widely employed semiconductor for CBZ degradation is TiO₂,⁶⁻⁹ however, it is severely limited as a sunlight-driven photocatalyst due to large band gap and low quantum yield.^{10,11} Therefore substantial research efforts have aimed to develop more efficient visible-light driven photocatalyst for CBZ degradation.¹²⁻¹⁶

Many semiconductors possess suitable band gap (e.g. 2.31 eV for BiVO₄ which corresponding an absorption edge at 535 nm) and sufficient band potentials for CBZ degradation but without or with low activity. The recombination of the photogenerated

electrons and holes may be the main reason. To enhance the separation efficiency of the photogenerated charges, cocatalysts which serve as the active sites for the photocatalytic reactions are usually necessary.¹⁷⁻²⁴ Moreover, when both the reduction and oxidation cocatalysts were co-loaded on the semiconductors, the synergistic effect on the photocatalytic activities were usually achieved.²⁵⁻²⁹ The strategy of dual cocatalysts has been successfully employed in photocatalytic water splitting,^{26,27,29,30} photocatalytic oxidation of some pollutants such as thiophene^{28,31} and some dyes.^{25,26,28} However, the synergistic effect of dual cocatalysts is far less investigated in the photocatalytic degradation of pharmaceuticals.

Herein we report an efficient visible-light driven three components photocatalyst for carbamazepine degradation. By co-loading reduction cocatalyst Pt and oxidation cocatalyst Co₃O₄ (MnO_x) on BiVO₄, the three components photocatalysts Pt/BiVO₄/Co₃O₄ (MnO_x) were assembled. The apparent rate constant of Pt/BiVO₄/Co₃O₄ for degradation of carbamazepine is 54 times of Co₃O₄/BiVO₄ and 2.5 times of Pt/BiVO₄, which shows a synergetic effect in the photocatalytic activity. The present work further demonstrates the universality of the strategy of dual-cocatalysts, which would be useful to construct highly efficient photocatalyst for pharmaceuticals degradation.

2. Experimental

2.1 Catalyst preparation

All of the reagents were of analytical grade, and were used without further purification. BiVO₄ was prepared by a hydrothermal process.³¹ The precursors NH₄VO₃ (12 mmol) and

^aEducational Technology Center, Chengde Medical University, Chengde 067000, China

^bHebei Key Laboratory of Research and Development for Traditional Chinese Medicine, Chengde Medical University, Chengde 067000, China. E-mail: wenyfuyu@vip.163.com

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c9ra07152k



$\text{Bi}(\text{NO}_3)_3 \cdot 5\text{H}_2\text{O}$ (12 mmol) were dissolved in 45 mL and 15 mL of 2.0 M nitric acid solutions, respectively. Then the above solutions were mixed to form a yellow homogeneous solution. The pH value of the solution was then adjusted to 2.0 with ammonia solution under stirring. The orange mixture was stirred for 0.5 h and aged for an additional 2 h. The orange precipitate at the bottom of the beaker was transferred to a 100 mL Teflon-lined stainless steel autoclave (70% capacity) and hydrothermal treated at 200 °C for 24 h. After the autoclave was cooled to room temperature, a vivid yellow powder was separated by filtration, washed with deionized water for more than 3 times, and then dried at 60 °C for overnight.

Co_3O_4 (MnO_x)/ BiVO_4 was prepared by photo-deposition method.²⁶ $\text{Co}(\text{NO}_3)_2$ solutions (0.5 g L^{-1}) or MnSO_4 solutions (0.5 g L^{-1}) was used as the precursor and NaIO_3 was employed as the electron acceptor. Typically, 0.5 g BiVO_4 powder was suspended in 100 mL NaIO_3 solutions (0.01 M), then 1.0 mL $\text{Co}(\text{NO}_3)_2$ solution or MnSO_4 solution was added and the suspension was then irradiated by a 300 W Xe lamp ($\lambda > 420 \text{ nm}$) under continuous stirring. After 5 h photo-deposition, the suspension was filtered, washed with deionized water for more than 3 times, and finally dried at 60 °C for overnight. The preparation of Pt/BiVO_4 was similar to Co_3O_4 (MnO_x)/ BiVO_4 but using H_2PtCl_6 solution (0.374 g L^{-1}) as the precursor and water as the hole scavenger.

$\text{Pt}/\text{BiVO}_4/\text{Co}_3\text{O}_4$ (MnO_x) was prepared by simultaneous photo-deposition method.²⁷ 0.5 g BiVO_4 powder was suspended in 100 mL deionized water, then 6.0 mL H_2PtCl_6 solution and 1.0 mL $\text{Co}(\text{NO}_3)_2$ solution (MnSO_4 solution) were added. After 5 h photo-deposition, the suspension was filtered, washed and finally dried at 60 °C for overnight.

The electrodes of BiVO_4 , $\text{Pt}/\text{BiVO}_4/\text{Co}_3\text{O}_4$ and $\text{Pt}/\text{BiVO}_4/\text{MnO}_x$ were prepared by electrophoretic deposition of corresponding powder on FTO substrate ($1 \times 2 \text{ cm}^2$),³² followed by drying in air and calcination at 573 K for 1 h.

2.2. Catalyst characterization

The prepared samples were characterized by X-ray powder diffraction (XRD) on a Rigaku D/Max-2500/PC powder diffractometer. Each sample powder was scanned using $\text{Cu K}\alpha$ radiation with an operating voltage of 40 kV and an operating current of 200 mA. The scan rate of 5° min^{-1} was applied to record the patterns in the range of 8–80° at a step of 0.02°. Brunauer–Emmett–Teller (BET) specific surface areas of the samples were determined by nitrogen adsorption at 77 K on Micromeritics ASAP 2420 system.

UV-Vis diffuse reflectance spectra (UV-Vis DRS) were recorded on a TU-1950 UV-Vis spectrophotometer (Beijing Purkinje General Instrument Co., Ltd.) equipped with an integrating sphere. The morphologies and particle sizes were examined by a Quanta 200 FEG scanning electron microscope (SEM). X-ray photoelectron spectroscopy (XPS) measurements, using a VG ESCALAB MK2 spectrometer with monochromatized Al-K α excitation.

Electron spin resonance (ESR) signals of radicals trapped by DMPO were recorded at ambient temperature on a Bruker

ESR A200 spectrometer. The samples were introduced into the home-made quartz cup inside the microwave cavity and illuminated with a 300 W Xe lamp (CERAMAX LX-300). The settings for the ESR spectrometer were as follows: sweep width, 140 G; microwave frequency, 9.82 GHz; modulation amplitude, 1 G.

Photoelectrochemical performances of the photoanodes were measured in a three-electrode setup, where Pt electrode and saturated mercury electrode were employed as counter and reference electrode, respectively. Electrolyte was 0.5 M Na_2SO_4 solution. A shutter was used to record both the dark and photocurrent during a single scan. A 300 W Xe lamp (Ushio-CERAMAX LX-300) and optical cutoff filter (kenko, L-42; $\lambda > 420 \text{ nm}$) was used as light source. Electrochemical impedance spectroscopy (EIS) were performed on a Ivium electrochemical workstation by applying an AC voltage of 10 mV amplitude in the frequency range of 10^5 Hz to 10^{-1} Hz in 0.5 M Na_2SO_4 .

2.3. Photocatalytic reaction

The photocatalytic degradation of carbamazepine (CBZ) was carried out using a 300 W Xe lamp and optical cut-off filter ($\lambda > 420 \text{ nm}$). Normally, 100 mg photocatalyst was dispersed in 100 mL 10 mg L^{-1} CBZ solutions. Prior to irradiation, the suspensions were magnetically stirred in dark for 30 min to establish adsorption/desorption equilibrium between CBZ and the photocatalysts. Top irradiation was used and after different irradiation time, 3 mL suspensions were centrifuged and used for UV-Vis characterization. Calibration based on the Lambert–Beer law was used to quantify the concentration of CBZ.

Total organic carbon (TOC) was measured using TOC-VCPN of Shimadzu Corporation of Japan. A combustion catalytic oxidation method of 680 °C is used. The measurement range is from $4 \mu\text{g L}^{-1}$ to $30\,000 \mu\text{g L}^{-1}$. The degradation intermediates were detected by mass spectrometry on Agilent HPLC-6500 Q-TOF in electrospray positive ion (ESI^+) mode with acetonitrile (solvent A) and ultrapure water with 0.1% V formic acid (solvent B) as mobile phase. A zorbax eclipse plus C18 column ($1.8 \mu\text{m}$, $2.1 \times 50 \text{ mm}$) was used as separation column.

3. Results and discussion

3.1 Characterization of the photocatalysts

Fig. 1 shows the UV-Vis diffuse reflectance spectra of BiVO_4 , Pt/BiVO_4 , $\text{MnO}_x/\text{BiVO}_4$, $\text{Co}_3\text{O}_4/\text{BiVO}_4$, $\text{Pt}/\text{BiVO}_4/\text{MnO}_x$ and $\text{Pt}/\text{BiVO}_4/\text{Co}_3\text{O}_4$. The absorption edge of the prepared BiVO_4 is about 535 nm, demonstrating a strong absorption in the visible light region. No obvious shift of the absorption edge is observed when different cocatalysts were loaded on BiVO_4 . The XRD patterns of the as-prepared BiVO_4 , $\text{Pt}/\text{BiVO}_4/\text{MnO}_x$ and $\text{Pt}/\text{BiVO}_4/\text{Co}_3\text{O}_4$ are shown in Fig. 2. The diffraction peaks of all of the samples are identical to the peaks for the monoclinic scheelite phase of BiVO_4 (JCPDS 14-0688). The crystal structures of the co-loaded BiVO_4 show no changes compared with that of pure BiVO_4 , implying that the





Fig. 1 UV-Vis DRS of different photocatalysts. The contents of the deposited Pt, MnO_x and Co_3O_4 are 0.2 wt%, 0.1 wt% and 0.1 wt%, respectively.

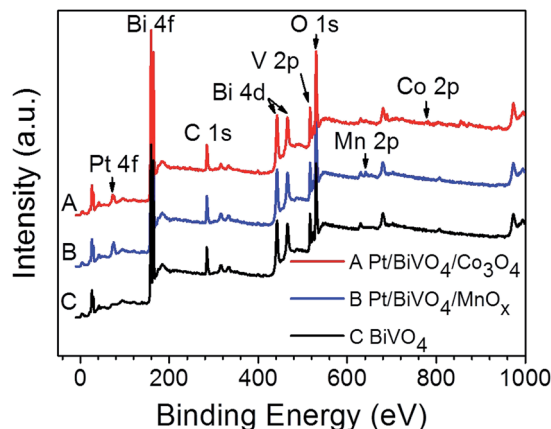


Fig. 3 XPS spectra of BiVO_4 , $\text{Pt/BiVO}_4/\text{Co}_3\text{O}_4$ and $\text{Pt/BiVO}_4/\text{MnO}_x$. The contents of the deposited Pt is 0.2 wt%, Co_3O_4 and MnO_x are all 0.1 wt%.

cocatalysts are loaded onto the surface of the BiVO_4 . N_2 adsorption–desorption isotherms (Fig. S1†) show that the co-loading of Pt and Co_3O_4 (MnO_x) did not change the framework of BiVO_4 and the BET surface areas of the three components photocatalysts $\text{Pt/BiVO}_4/\text{Co}_3\text{O}_4$ (MnO_x) are comparable with BiVO_4 .

SEM images of $\text{Pt/BiVO}_4/\text{Co}_3\text{O}_4$ further demonstrate that the cocatalysts are deposited on the surface of the BiVO_4 (Fig. S2†). Furthermore, as reported by Li *et al.*,^{26,27} the reduction cocatalyst Pt and oxidation cocatalyst Co_3O_4 may be deposited on the different facets of BiVO_4 , which will facilitate the charge separation of electrons and holes.

The chemical states of the loaded cocatalysts were examined by XPS characterization. The Pt 4f peak located at binding energy of 70.7 eV was observed for both three components photocatalysts, indicating the existence of the metallic Pt (0) on BiVO_4 (Fig. 3). The Co 2p peaks at binding energies of 777.9 eV and 795.0 eV demonstrating that Co_3O_4 was

deposited on BiVO_4 (Fig. 4a).²⁶ The Mn 2p peaks are located at binding energies of 653.5 eV and 641.9 eV, which are between those of Mn_2O_3 and MnO_2 (Fig. 4b). Based on this, the deposited Mn species can be ascribed to MnO_x , where x is between 1.5 and 2.0.²⁷

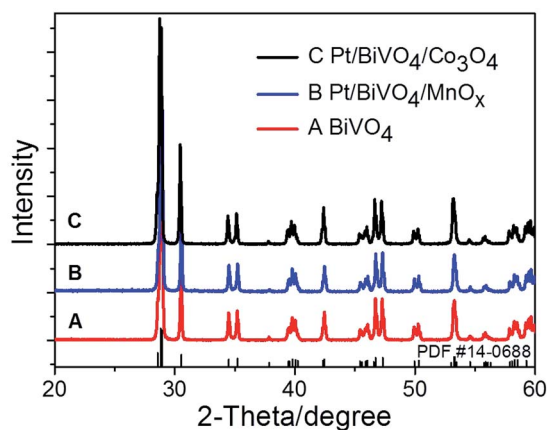


Fig. 2 XRD patterns of BiVO_4 , $\text{Pt/BiVO}_4/\text{MnO}_x$ and $\text{Pt/BiVO}_4/\text{Co}_3\text{O}_4$. The contents of the deposited Pt is 0.2 wt%, Co_3O_4 and MnO_x are all 0.1 wt%.



Fig. 4 The Co 2p XPS spectra of Co_3O_4 (a) and Mn 2p XPS spectra of MnO_x (b) deposited on BiVO_4 . The contents of the deposited Co_3O_4 and MnO_x are all 0.1 wt%.





Fig. 5 Photocatalytic degradation of CBZ on different photocatalysts. Reaction conditions: 100 mL CBZ solution with initial concentration of 10 mg L^{-1} , catalyst dosage 1 g L^{-1} , the contents of the deposited Pt is 0.2 wt%, Co_3O_4 and MnO_x are all 0.1 wt% and 300 W Xe lamp ($\lambda > 420 \text{ nm}$), top irradiation.

3.2 The effect of cocatalyst on photocatalytic degradation of carbamazepine

Fig. 5 shows the kinetic data of the photocatalytic degradation of CBZ on different photocatalysts. BiVO_4 without loaded cocatalyst exhibited a low removal (ca. 2%) in 120 min, and the removal was slightly enhanced when the oxidation cocatalyst MnO_x or Co_3O_4 was loaded on BiVO_4 . When reduction cocatalyst Pt was deposited on BiVO_4 , the removal of CBZ was enhanced to about 28% in 120 min. However, when Pt and Co_3O_4 were co-loaded on BiVO_4 , the removal was significantly improved to ca. 56%, much higher than those of Pt/BiVO_4 and $\text{Co}_3\text{O}_4/\text{BiVO}_4$. Same trend was also observed when Pt and MnO_x were co-loaded on BiVO_4 , yielding a removal of ca. 53% in 120 min, which is much higher than those of Pt/BiVO_4 and $\text{MnO}_x/\text{BiVO}_4$.

TOC analysis was carried out to evaluate the mineralization extent of CBZ on BiVO_4 and $\text{Pt/BiVO}_4/\text{Co}_3\text{O}_4$ photocatalysts (Fig. S3†). As shown in Fig. S3,† the TOC removal of CBZ in the presence of $\text{Pt/BiVO}_4/\text{Co}_3\text{O}_4$ is also much higher than that of BiVO_4 . Long-term circulation measurement was carried out to check the stability of the photocatalyst $\text{Pt/BiVO}_4/\text{Co}_3\text{O}_4$. As shown in Fig. S4,† the photocatalytic activity of $\text{Pt/BiVO}_4/\text{Co}_3\text{O}_4$ can be well maintained even after three runs, only less than 5% decreasing, indicating the excellent stability and durability of the three components photocatalyst.

The plots of $\ln(C_0/C)$ versus the reaction time (t) is presented in Fig. 6. All of the fitted curves show high corresponding apparent coefficients, revealing that the photocatalytic degradation of CBZ over different photocatalysts is in line with pseudo-first-order kinetic. The values of the rate constant k for different photocatalysts are listed in Table 1. The single loading of either oxidation cocatalyst (entry 2 and 3) or reduction cocatalyst (entry 4) can somewhat enhance the activity of BiVO_4 , demonstrating the roles of cocatalysts played in the facilitating of charge transfer of photogenerated holes or electrons; however, the

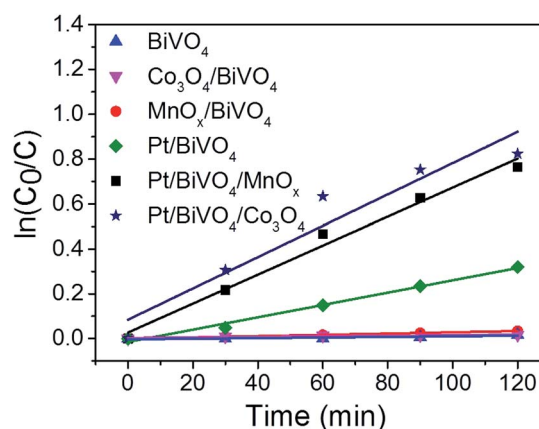


Fig. 6 Kinetic fitting of the CBZ degradation on different photocatalysts. Reaction conditions: 100 mL CBZ solution with initial concentration of 10 mg L^{-1} , catalyst dosage 1 g L^{-1} , the contents of the deposited Pt is 0.2 wt%, Co_3O_4 and MnO_x are all 0.1 wt% and 300 W Xe lamp ($\lambda > 420 \text{ nm}$), top irradiation.

enhancement is limited. The promotion caused by the reduction cocatalyst loading is more obvious than the oxidation cocatalyst loading, indicating that the reduction half-reaction may be the rate-determining step.²⁶ When reduction and oxidation cocatalysts were co-loaded on BiVO_4 , the rate constants (entry 5 and 6) were significantly improved. The k for $\text{Pt/BiVO}_4/\text{MnO}_x$ is 23 times of $\text{MnO}_x/\text{BiVO}_4$ and 2.3 times of Pt/BiVO_4 . Similarly, the k for $\text{Pt/BiVO}_4/\text{Co}_3\text{O}_4$ is 54 times of $\text{Co}_3\text{O}_4/\text{BiVO}_4$ and 2.5 times of Pt/BiVO_4 . Both of the three components photocatalysts show obvious synergetic effect of the dual cocatalysts ($k_5 > k_3 + k_4$, $k_6 > k_2 + k_4$). This demonstrates that only when the reduction and oxidation reactions were accelerated simultaneously we can get the much higher reaction rate than any other one.

The roles of cocatalysts played in facilitating photo-generated charges transfer can be further demonstrated by the photoelectrochemical measurements. As shown in Fig. S5,† the photocurrent density of the three components photocatalysts $\text{Pt/BiVO}_4/\text{Co}_3\text{O}_4$ and $\text{Pt/BiVO}_4/\text{MnO}_x$ are higher

Table 1 Apparent first-order rate constant k for degradation of CBZ over different photocatalysts

Entry	Photocatalysts	k^a ($\times 10^{-3} \text{ min}^{-1}$)
1	BiVO_4	0.12
2	$\text{Co}_3\text{O}_4/\text{BiVO}_4$	0.13
3	$\text{MnO}_x/\text{BiVO}_4$	0.28
4	Pt/BiVO_4	2.75
5	$\text{Pt/BiVO}_4/\text{MnO}_x$	6.46
6	$\text{Pt/BiVO}_4/\text{Co}_3\text{O}_4$	6.98

^a The apparent first-order rate constant k is calculated from the equation: $\ln(C_0/C) = kt$, where C_0 is the initial concentration and C is the concentration at certain time t . Reaction conditions: 100 mL CBZ solution with initial concentration of 10 mg L^{-1} , catalyst dosage 1 g L^{-1} , the contents of the deposited Pt is 0.2 wt%, Co_3O_4 and MnO_x are all 0.1 wt% and 300 W Xe lamp ($\lambda > 420 \text{ nm}$), top irradiation.



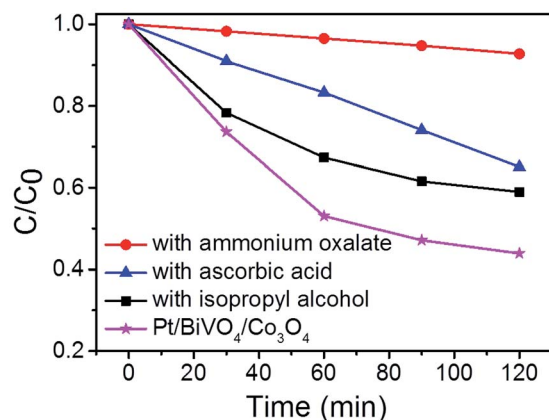


Fig. 7 Effect of scavengers on photocatalytic degradation of CBZ over Pt/BiVO₄/Co₃O₄ photocatalyst. Reaction conditions: 100 mL CBZ solution with initial concentration of 10 mg L⁻¹, scavenger concentration 50 mmol L⁻¹, catalyst dosage 1 g L⁻¹, the contents of the deposited Pt and Co₃O₄ are 0.2 wt% and 0.1 wt% respectively and 300 W Xe lamp ($\lambda > 420$ nm), top irradiation.

than that of bare BiVO₄. This demonstrating that due to the co-loading of cocatalysts, the photo-generated charges can be more efficiently trapped, which makes more photo-generated holes survived on the three components photocatalysts. Moreover, the photocurrent of Pt/BiVO₄/Co₃O₄ is higher than Pt/BiVO₄/MnO_x, demonstrating that the photogenerated holes (h⁺) would transfer easier from BiVO₄ to Co₃O₄ than to MnO_x, which cause the superior performance of Pt/BiVO₄/Co₃O₄ than Pt/BiVO₄/MnO_x on the photocatalytic CBZ degradation. The electrochemical impedance spectra (EIS) of BiVO₄ and Pt/BiVO₄/Co₃O₄ further demonstrate the role of cocatalysts played in facilitating photo-generated charges transfer. As shown in Fig. S6,† the arc radius on the EIS plot of Pt/BiVO₄/Co₃O₄ is smaller than that of BiVO₄, implying that the co-loading of Pt and Co₃O₄ make photo-generated charges transfer easier.

3.3 Identification of main reactive species

To identify the main reactive species in the photocatalytic degradation process, trapping experiments were conducted on Pt/BiVO₄/Co₃O₄ photocatalyst and the results are shown in Fig. 7. The addition of ammonium oxalate (AO, scavengers of h⁺ radical) has significant inhibition on CBZ degradation, indicating that h⁺ is crucial active species. The addition of ascorbic acid (VC, scavengers of ·O₂⁻ radical) and isopropyl alcohol (IPA, scavengers of ·OH radical) have feeble inhibition, which reveals that ·O₂⁻ and ·OH serve as synergistic active species but not as the most crucial active species. Similar results were observed for Pt/BiVO₄/MnO_x photocatalyst (Fig. S7†).

In order to gain direct evidence for the reactive radicals involved in the photocatalytic process, ESR analysis was employed using DMPO as trapping agent. As shown in Fig. 8, the sextet ESR signal centered at $g = 2.0065$ can be assigned to DMPO·O₂⁻. These results provide evidence of ·O₂⁻ formed in the presence of the photocatalyst BiVO₄ and Pt/BiVO₄/Co₃O₄.

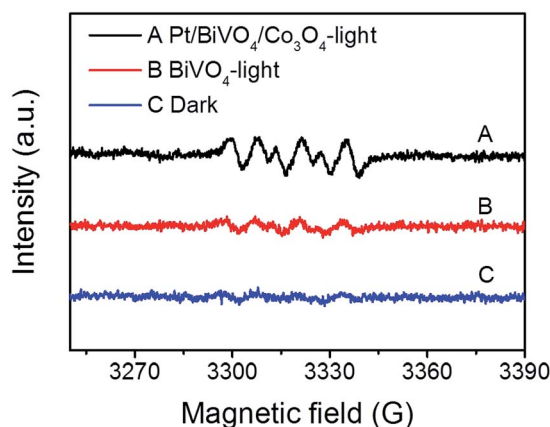


Fig. 8 *In situ* ESR spectra of DMPO·O₂⁻ generated by BiVO₄ and Pt/BiVO₄/Co₃O₄ after 10 min of visible-light irradiation ($\lambda > 420$ nm). The signal obtained in the presence of photocatalyst but without light irradiation is denoted as "Dark".

Moreover, the signal of ·O₂⁻ generated after 10 min of illumination on Pt/BiVO₄/Co₃O₄ is more obvious than those for BiVO₄, demonstrating the roles of cocatalysts played in facilitating charge separation and transfer.

3.4 Degradation pathway of CBZ photocatalyzed by Pt/BiVO₄/Co₃O₄

Based on the LC-ESI-MS results (Table S1 and Fig. S8†), the possible degradation pathway of CBZ photocatalyzed by Pt/BiVO₄/Co₃O₄ was proposed in Fig. 9. First of all, alkene double bond of CBZ was attack by ·OH radical and transferred into intermediate **P6** and **P9**.³³⁻³⁵ Intermediate **P6** was further hydroxylated to generate **P2** which might be further oxidized to generate **P8**.³⁵ As for intermediate **P9**, it might be further oxidized by ·O₂⁻ and photo-generated h⁺ to convert into **P4** and **P7**.³⁴ After aldehyde oxidation reaction in **P4**, **P5** was generated and further transferred to **P8**.³⁵ Product **P7** might undergo several successive steps including deamination and decarboxylation to produce **P1** and **P3**.^{7,34,36} Further oxidized by ·OH, h⁺ and ·O₂⁻, **P3** and **P8** could be finally mineralized into CO₂ and H₂O *via* ring-rupturing reaction.

The proposed mechanism of photocatalytic degradation of CBZ on Pt/BiVO₄/Co₃O₄ (MnO_x) is shown in Fig. 10. Under visible light irradiation, photo-generated electrons would transfer from the conduction band (CB) of BiVO₄ to the reduction cocatalyst Pt, on which the adsorbed O₂ is reduced to ·O₂⁻ radical. Meanwhile, the holes would migrate from the valence band (VB) of BiVO₄ to the oxidation cocatalyst Co₃O₄ (MnO_x), where the adsorbed CBZ or H₂O are oxidized to CO₂ and ·OH respectively. The produced ·O₂⁻ and ·OH also participate in the degradation of CBZ, which is consistent with the trapping experiment results. The co-loading of both the reduction cocatalyst Pt and oxidation cocatalyst Co₃O₄ (MnO_x) is beneficial for the efficient separation and transfer of the photo-generated electrons and holes, accounts for the high photocatalytic activity of CBZ degradation.



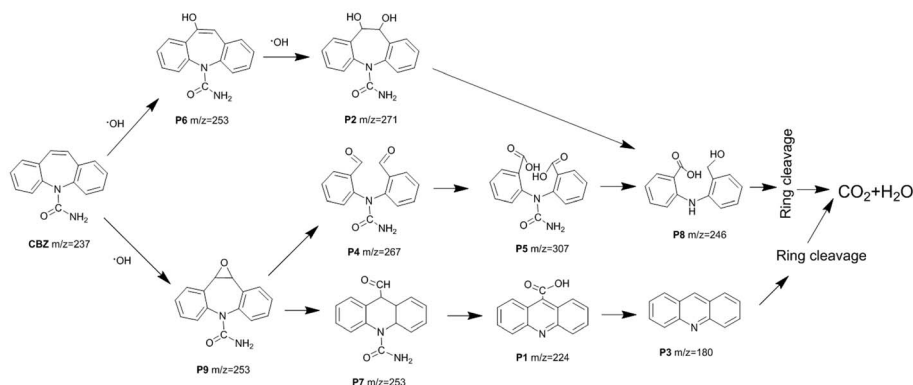


Fig. 9 Proposed degradation pathway of CBZ by Pt/BiVO₄/Co₃O₄ photocatalyst.

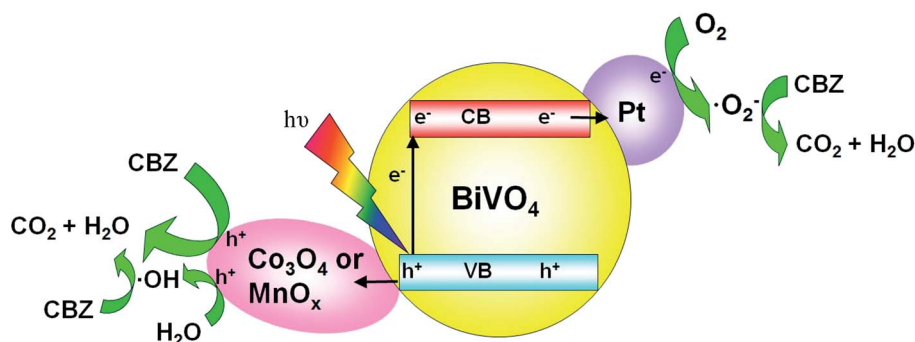


Fig. 10 Schematic description of the mechanism for the photocatalytic degradation of CBZ on Pt/BiVO₄/Co₃O₄ (MnO_x) photocatalyst under visible light irradiation.

4. Conclusion

A synergetic effect in photocatalytic degradation of CBZ was achieved when reduction cocatalyst Pt and oxidation cocatalyst Co₃O₄ (MnO_x) were co-loaded on BiVO₄. The co-loading of both the reduction cocatalyst and oxidation cocatalyst can facilitate the separation and transfer of the photo-generated electrons and holes, which is responsible for the high photocatalytic activity of the three components photocatalysts. This work further demonstrates the universality of the strategy of dual-cocatalysts, which would be useful to construct highly efficient photocatalyst for pharmaceuticals degradation.

Conflicts of interest

The authors declare no competing financial interest.

Acknowledgements

This work was financially supported by Scientific Research Fund of the Hebei Education Department (No. BJ2014002), Natural Science Foundation of Hebei Province, China (No. B2016406012 and H2017406022), National Natural Science Foundation of China (No. 81473101 and 81001401) and Undergraduate Innovation and Entrepreneurship Training Program of Chengde Medical University (No. 201424 and 201603). The authors thank

Prof. Can Li, Prof. Xu Zong, Dr Jingfeng Han, Dr Sheng Ye and Xianwen Zhang for the help on characterizations of the photocatalysts.

References

- 1 J. Rivera-Utrilla, M. Sanchez-Polo, M. A. Ferro-Garcia, G. Prados-Joya and R. Ocampo-Perez, *Chemosphere*, 2013, **93**, 1268–1287.
- 2 Y. J. Zhang, S. U. Geissen and C. Gal, *Chemosphere*, 2008, **73**, 1151–1161.
- 3 T. Heberer, *Toxicol. Lett.*, 2002, **131**, 5–17.
- 4 K. Fent, A. A. Weston and D. Caminada, *Aquat. Toxicol.*, 2006, **76**, 122–159.
- 5 P. A. K. Reddy, P. V. L. Reddy, E. Kwon, K. H. Kim, T. Akter and S. Kalagara, *Environ. Int.*, 2016, **91**, 94–103.
- 6 S. Murgolo, V. Yargeau, R. Gerbasi, F. Visentin, N. El Habra, G. Ricco, I. Lacchetti, M. Carere, M. L. Curri and G. Mascolo, *Chem. Eng. J.*, 2017, **318**, 103–111.
- 7 M. Nawaz, W. Miran, J. Jang and D. S. Lee, *Appl. Catal., B*, 2017, **203**, 85–95.
- 8 A. Carabin, P. Drogui and D. Robert, *J. Taiwan Inst. Chem. Eng.*, 2015, **54**, 109–117.
- 9 A. Y. C. Tong, R. Braund, D. S. Warren and B. M. Peake, *Cent. Eur. J. Chem.*, 2012, **10**, 989–1027.



- 10 A. Cincinelli, T. Martellini, E. Coppini, D. Fibbi and A. Katsoyiannis, *J. Nanosci. Nanotechnol.*, 2015, **15**, 3333–3347.
- 11 D. Kanakaraju, B. D. Glass and M. Oelgemoller, *Environ. Chem. Lett.*, 2014, **12**, 27–47.
- 12 Y. X. Gao, G. Yu, K. Liu, S. B. Deng, B. Wang, J. Huang and Y. J. Wang, *Chem. Eng. J.*, 2017, **330**, 157–165.
- 13 X. Chen, W. Y. Lu, T. F. Xu, N. Li, Z. X. Zhu, G. Q. Wang and W. X. Chen, *Chem. Eng. J.*, 2017, **328**, 853–861.
- 14 L. Lin, H. Y. Wang and P. Xu, *Chem. Eng. J.*, 2017, **310**, 389–398.
- 15 L. Lin, H. Y. Wang, W. B. Jiang, A. R. Mkaouar and P. Xu, *J. Hazard. Mater.*, 2017, **333**, 162–168.
- 16 Y. B. Ding, G. L. Zhang, X. R. Wang, L. H. Zhu and H. Q. Tang, *Appl. Catal., B*, 2017, **202**, 528–538.
- 17 J. H. Yang, D. G. Wang, H. X. Han and C. Li, *Acc. Chem. Res.*, 2013, **46**, 1900–1909.
- 18 F. Y. Wen and C. Li, *Acc. Chem. Res.*, 2013, **46**, 2355–2364.
- 19 K. Wenderich and G. Mul, *Chem. Rev.*, 2016, **116**, 14587–14619.
- 20 T. D. Nguyen, Q. T. P. Bui, T. B. Le, T. M. Altahtamouni, K. B. Vu, D. V. N. Vo, N. T. H. Le, T. D. Luu, S. S. Hong and K. T. Lim, *RSC Adv.*, 2019, **9**, 23526–23534.
- 21 D. Gao, W. Liu, Y. Xu, P. Wang, J. Fan and H. Yu, *Appl. Catal., B*, 2020, **260**, 118190.
- 22 S. C. Sun, Y. C. Zhang, G. Q. Shen, Y. T. Wang, X. L. Liu, Z. W. Duan, L. Pan, X. W. Zhang and J. J. Zou, *Appl. Catal., B*, 2019, **243**, 253–261.
- 23 H. G. Yu, R. R. Yuan, D. D. Gao, Y. Xu and J. G. Yu, *Chem. Eng. J.*, 2019, **375**, 121934.
- 24 M. H. Ai, J. W. Zhang, R. J. Gao, L. Pan, X. W. Zhang and J. J. Zou, *Appl. Catal., B*, 2019, **256**, 117805.
- 25 R. G. Li, X. P. Tao, R. T. Chen, F. T. Fan and C. Li, *Chem.–Eur. J.*, 2015, **21**, 14337–14341.
- 26 R. G. Li, H. X. Han, F. X. Zhang, D. G. Wang and C. Li, *Energy Environ. Sci.*, 2014, **7**, 1369–1376.
- 27 R. G. Li, F. X. Zhang, D. G. Wang, J. X. Yang, M. R. Li, J. Zhu, X. Zhou, H. X. Han and C. Li, *Nat. Commun.*, 2013, **4**, 7.
- 28 F. Lin, Y. N. Zhang, L. Wang, Y. L. Zhang, D. G. Wang, M. Yang, J. H. Yang, B. Y. Zhang, Z. X. Jiang and C. Li, *Appl. Catal., B*, 2012, **127**, 363–370.
- 29 B. J. Ma, F. Y. Wen, H. F. Jiang, J. H. Yang, P. L. Ying and C. Li, *Catal. Lett.*, 2010, **134**, 78–86.
- 30 L. C. Mu, Y. Zhao, A. L. Li, S. Y. Wang, Z. L. Wang, J. X. Yang, Y. Wang, T. F. Liu, R. T. Chen, J. Zhu, F. T. Fan, R. G. Li and C. Li, *Energy Environ. Sci.*, 2016, **9**, 2463–2469.
- 31 F. Lin, D. Wang, Z. Jiang, Y. Ma, J. Li, R. Li and C. Li, *Energy Environ. Sci.*, 2012, **5**, 6400–6406.
- 32 D. E. Wang, R. G. Li, J. Zhu, J. Y. Shi, J. F. Han, X. Zong and C. Li, *J. Phys. Chem. C*, 2012, **116**, 5082–5089.
- 33 X. Y. Gao, X. C. Zhang, Y. W. Wang, S. Q. Peng, B. Yue and C. M. Fan, *Chem. Eng. J.*, 2015, **273**, 156–165.
- 34 Y. Guo, P. F. Wang, J. Qian, Y. H. Ao, C. Wang and J. Hou, *Appl. Catal., B*, 2018, **234**, 90–99.
- 35 J. Cao, W. S. Nie, L. Huang, Y. B. Ding, K. L. Lv and H. Q. Tang, *Appl. Catal., B*, 2019, **241**, 18–27.
- 36 J. Xu, L. Li, C. S. Guo, Y. Zhang and W. Meng, *Appl. Catal., B*, 2013, **130**, 285–292.

