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1 Introduction

With the rapid development of aquaculture, antibiotics are widely used to treat infectious diseases of fish. However, remnant antibiotic drugs have the potential to harm people and the environment.^{1,2} Levofloxacin hydrochloride is a quinolone with antibacterial activities.^{3,4} Residual levofloxacin hydrochloride in water can increase the resistance of microbes, which is harmful to human health and safety.⁵⁻⁷ Hence, it is important to develop a method to remove the levofloxacin hydrochloride in water.

Photocatalytic technology is an advanced oxidation method that is an outstanding achievement for water treatment because it is inexpensive and produces no secondary pollution.^{8,9} The photocatalytic reaction produces free radicals in a chain reaction until the end products of degradation are CO₂ and H₂O. For wastewater treatment, there is higher research value in the photocatalytic reaction for the treatment of pollutants because of its low selectivity, rapid reaction process, and simple operation compared with conventional oxidation technology.¹⁰

Synergetic effect of photocatalysis and peroxymonosulfate activated by MFe_2O_4 (M = Co, Mn, or Zn) for enhanced photocatalytic activity under visible light irradiation⁺

Mingyang Long,^a Di Li, ^b *^a Hongmiao Li,^a Xinguo Ma, ^c Qianqian Zhao,^a Qi Wen^a and Fang Song^b

Nanosized MFe₂O₄ (M = Co, Mn, or Zn) photocatalysts were synthesized *via* a simple sol-gel method. MFe₂O₄ photocatalysts exhibited lower photocatalytic activity for the degradation of levofloxacin hydrochloride under visible light irradiation. For enhancement of photocatalytic activity, MFe₂O₄ was used to activate peroxymonosulfate and degrade levofloxacin hydrochloride under visible light irradiation. The influences of peroxymonosulfate dosage, levofloxacin hydrochloride concentration, pH value, and temperature on peroxymonosulfate activation to degrade levofloxacin hydrochloride were investigated in detail. The mechanism of activation of peroxymonosulfate by MFe₂O₄ was proposed and proved by radical quenching experiments, electron spin resonance analysis, X-ray photoelectron spectroscopy, electrochemical impedance spectroscopy, and transient photocurrent responses. The combined activation effects of photogenerated e⁻/h⁺ and transition metals on peroxymonosulfate to produce sulfate radical clearly enhanced the degradation efficiency.

> Fe-based heterogeneous catalysts have been widely explored and studied because they are environmentally friendly, inexpensive, and non-toxic compared to other metals. In addition, spinel ferrite nanoparticles are magnetic semiconductors and have been used for activating peroxymonosulfate (PMS).11-13 Ferrite nanoparticles are a magnetic nanomaterial that can be simply recycled from solution by applying an external magnetic field. The presence of ferrite magnetic nanoparticles greatly increases the efficiency of pollutant removal. These materials directly activate PMS to produce sulfate radical (SO_4^{-}) . Previous conductivity measurements determined that the band gaps of CoFe₂O₄, MnFe₂O₄, and ZnFe₂O₄ were in the range of 0.5-0.6 eV.14-16 This indicates that CoFe2O4, MnFe2O4, and ZnFe₂O₄ can be considered as narrow band gap semiconductors that can absorb a greater amount of visible light, and their narrow band gaps allow sunlight to be fully utilized. Furthermore, CoFe₂O₄, MnFe₂O₄, and ZnFe₂O₄ nanoparticles increase the efficiency of removal and can be simply recycled from a heterogeneous suspension using an external magnetic field after the completion of photocatalytic reactions. Thus, secondary pollution created by the disposal of photocatalysts would not be produced.17-20

> In this work, nanosized MFe₂O₄ (M = Co, Mn, or Zn) photocatalysts were synthesized by a sol-gel method. To further increase the degradation efficiency of levofloxacin hydrochloride, MFe₂O₄ photocatalysts were used to activate PMS under visible light irradiation. In addition to photogenerated e^-/h^+ ,

[&]quot;School of Chemistry and Chemical Engineering, Xi'an University of Architecture and Technology, Xi'an, 710055, China. E-mail: di.li@xauat.edu.cn

^bInstrument Analysis Center, Xi'an University of Architecture and Technology, Xi'an, 710055, China

School of Science, Hubei University of Technology, Wuhan 430068, China

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Paper

Fe, Co, Mn, and Zn can also activate PMS to produce SO_4 .⁻, and thus, the combined activation effects of photogenerated e^-/h^+ and transition metals on PMS to produce SO_4 .⁻ can clearly enhance the degradation efficiency. The influences of PMS dosage, pH value, levofloxacin hydrochloride concentration, and temperature on PMS activation were investigated in detail. Moreover, the degradation mechanisms used by MFe₂O₄ photocatalysts to activate PMS were systematically studied.

2 Experimental

2.1 Materials

Levofloxacin hydrochloride (LVX) of analytical reagent grade quality was used without further purification. Potassium peroxymonosulfate ($2KHSO_5 \cdot 3KHSO_4 \cdot K_2SO_4$, Oxone®) (PMS) was purchased from Sigma-Aldrich. Other chemicals were analytical or reagent grade commercial products. All solutions were prepared with deionized water.

2.2 Preparation of MFe₂O₃

First, 4 mmol Fe(NO₃)₃·9H₂O, 2 mmol Co(NO₃)₂·6H₂O, and 12 mmol citric acid were dissolved in a mixture of deionized water (10 mL) and ethanol (20 mL), which was maintained at 70 °C until it entered into the gel state. The gel was dried at 80 °C until it formed a xerogel, which was calcined in a muffle furnace at 550 °C for 5 hours with a heating rate of 5 °C min⁻¹ to obtain CoFe₂O₄ powder. The ZnFe₂O₄ sample was prepared using the same protocol. However, the MnFe₂O₄ sample was calcined at 400 °C for 5 hours, although the other preparation steps were carried out in the same manner as those for CoFe₂O₄.

2.3 Characterizations

The UV-Vis diffuse reflectance spectra (UV-Vis DRS) of the samples were obtained on a UV-Vis spectrophotometer (Lambda 950, PerkinElmer) using an integrating sphere accessory, and BaSO₄ was used as a reflectance standard. X-ray diffraction (XRD) experiments were carried out using a Rigaku D/MAX 2500 diffractometer with Cu Ka radiation. The size and morphologies of MFe₂O₄ were characterized with the aid of a JSM-7000F field emission scanning electron microscope. Fourier transform infrared (FT-IR) spectroscopy was recorded on a Thermo Nicolet iS5 spectrometer with a KBr disk. The magnetic property was measured at room temperature with the Quantum Design MPMS-SQUID VSM-094. X-ray photoelectron spectroscopy (XPS) measurements were performed using a Kratos AXIS ULTRA DLD. Electron paramagnetic resonance (EPR) measurements of spin-trapped radicals with spin-trap reagent 5,5-dimethyl-1-pirroline-N-oxide (DMPO) (Sigma-Aldrich) were carried out at room temperature with a Bruker A300 spectrometer equipped with a high-pressure mercury lamp as the irradiation source. To minimize experimental errors, the same type of quartz capillary tube was used for all EPR measurements. An EPR spectrometer was coupled to a computer for data acquisition and instrument control.

2.4 Photo-electrochemical properties

Photo-electrochemical measurements were carried out using a conventional three-electrode, single-compartment glass cell fitted with a synthesized quartz window using a potentiostat. The quartz electrolytic cell was filled with 0.1 M Na₂SO₄. A total of 10 mg of synthesized MFe₂O₄ was dropped onto the ITO glass $(2.0 \times 4.0 \text{ cm})$, which was used as a working electrode for electrochemical impedance spectroscopy (EIS) and chronoamperometry experiments. A 500 W xenon lamp (Institute of Electric Light Source, Beijing) was used as the light source for the visible light irradiation in the photoelectrochemical analyses. A 420 nm cutoff filter was placed onto the window face of the cell to ensure the desired irradiation conditions. The counter and reference electrodes were a platinum black wire and Ag/AgCl electrode, respectively. The photoelectrochemical experiment was performed using an electrochemical system (Shanghai Chenhua-CHI660e).

2.5 Photocatalytic oxidative degradation

The photocatalytic activities of MFe_2O_4 were evaluated by LVX decomposition under visible light irradiation. In the case of visible light irradiation, a 300 W halogen lamp (Philips Plusline, Shanghai) was focused through a window. A 420 nm cutoff filter was placed onto the window face of the cell to ensure the desired irradiation conditions. The average light intensity was 65 mW cm⁻². The radiant flux was measured with a power meter (Beijing Normal University, Beijing).

A cylindrical double-layer glass photochemical reactor with internal diameter 70 mm, external diameter 80 mm, and height 100 mm was utilized for the photocatalysis reaction. A distance of approximately 17 cm between the lamp and reactor was maintained. Running water was piped into the layer to maintain a constant temperature.

The photocatalytic degradation of LVX in aqueous solution was studied using MFe₂O₄ as the photocatalyst at room temperature and under normal atmospheric pressure. MFe₂O₄ (50 mg) and 100 mL LVX (10 mg L⁻¹) aqueous solution were added to the reactor, and then stirred with a magnetic stirrer prior to irradiation by a halogen lamp at room temperature. Prior to irradiation, the solution was incubated in the dark for 30 min to ensure equilibrium of the working solution. After the reaction, the sample solution was centrifuged to remove MFe₂O₄, and the solution obtained this way was extracted into a quartz cell. The absorbance of the samples was measured using quartz cells every 10 min.

For the photocatalytic test of MFe_2O_4 activating PMS, MFe₂O₄ was dispersed into LVX solution and magnetically stirred in the dark for 30 min to ensure adsorption–desorption equilibrium. A certain amount of PMS was added to the reactor, and the concentration of LVX solution was monitored by measuring the absorbance as the initial concentration as C_0 . Then, the lamp was turned on to initiate the photodegradation reaction. After five min, 3 mL of the solution was removed and filtered using a 0.22 mm membrane to separate the catalyst powders from the solution, where the instantaneous concentration of LVX was measured every 10 min as C.

3 Results and discussion

3.1 Controlling the synthesis of MFe₂O₄

The XRD patterns were used to study the crystal structure of MFe₂O₄. The XRD patterns of the phase characteristics and crystalline structure information for MFe₂O₄ are shown in Fig. 1. For CoFe₂O₄ nanoparticles, the diffraction peaks of all the samples were easily indexed as $CoFe_2O_4$, which were in agreement with the standard card (JCPDS card number: 79-1744, $R\bar{3}m$ (166) Space Group) with major peaks at $2\theta = 18.28^{\circ}$, 30.08°, 35.43°, 37.06°, 43.05°, 56.04°, and 62.52° corresponding to the diffractions of (003), (104), (113), (006), (024), (125), and (119), respectively. For MnFe₂O₄, the diffraction peaks of all the samples were easily indexed as MnFe2O4, which were in agreement with the standard card (JCPDS card number: 73-1964, $Fd\bar{3}m$ (227) Space Group) with major peaks at $2\theta = 18.03^{\circ}$, 29.66°, 34.93°, 42.44°, 56.10°, and 61.59° corresponding to (111), (220), (311), (400), (333), and (440), respectively. For ZnFe₂O₄, the diffraction peaks of all the samples were easily indexed as ZnFe₂O₄, which were in agreement with the standard card (JCPDS card number: 73-1963, Fd3m (227) Space Group) with major peaks at $2\theta = 18.38^{\circ}$, 30.24° , 35.63° , 43.30° , 57.28° , and 62.91° corresponding to (111), (220), (311), (400), (511), and (440), respectively.

The microstructures of MFe₂O₄ were observed by scanning electron microscopy (SEM), as shown in Fig. 2a–d. The CoFe₂O₄, MnFe₂O₄, and ZnFe₂O₄ nanoparticles were agglomerated together. The CoFe₂O₄ result (Fig. 2a and b) shows agglomerated nanoparticles 10–20 nm in size. The MnFe₂O₄ (Fig. 2c) particle size was approximately 30 nm, and the ZnFe₂O₄ nanoparticles (Fig. 2d) were 20–30 nm in size.

3.2 FT-IR spectrum and the optical properties of MFe₂O₄

The FT-IR spectrum of MFe₂O₄ is shown in Fig. 3. For CoFe₂O₄, the absorption band at 1113 cm⁻¹ is characteristic of the cobalt ferrite system, and this may be due to the residual FeOOH. The absorption bands present at approximately 568 cm⁻¹ were due to the stretching vibrations of metal oxide in the octahedral group complex Co(π)-O²⁻ and Fe(π)-O²⁻ tetrahedral group complex of the cobalt ferrite phase, respectively, which proves the existence of spinel ferrite.²¹ The peaks at 1635 cm⁻¹, 2336 cm⁻¹, and 3373 cm⁻¹ were attributed to the vibrational stretching of the O-H bond of H₂O due to the bending of the



Fig. 1 The XRD pattern of MFe₂O₄ prepared by a sol-gel method.



Fig. 2 SEM images of MFe₂O₄: (a) CoFe₂O₄, (b) CoFe₂O₄, (c) MnFe₂O₄, and (d) ZnFe₂O₄.



Fig. 3 FT-IR spectra of MFe₂O₄.

absorbed water molecules.²² For $MnFe_2O_4$, the characteristic peaks at approximately 563 cm⁻¹ and 637 cm⁻¹ corresponded to the formation of Mn–O and Fe–O bonds at the octahedral sites of spinel-type compounds.²³ For $ZnFe_2O_4$, the characteristic absorption peaks at 537 cm⁻¹ and 452 cm⁻¹ denoted the stretching vibration of the Fe–O and Zn–O bonds, indicating the existence of $ZnFe_2O_4$.²⁴

UV-Vis diffuse reflectance spectroscopy (DRS) was used to research the optical property of the photocatalysts.²⁵ There was strong absorption for $CoFe_2O_4$ and $MnFe_2O_4$ in the range of 200–900 nm, and strong absorption for $ZnFe_2O_4$ in the range of 200–600 nm, as shown in Fig. 4. Therefore, the visible light was efficiently utilized. The steep shape of the spectra indicated that the visible light absorption was not caused by a transition from the impurity level, but rather, was caused by the band-gap transition.²⁶

3.3 Magnetic properties of MFe₂O₄ photocatalysts

Fig. 5 shows the H–M hysteresis loop of MFe₂O₄ samples prepared by a sol–gel method. There were excellent magnetic properties for CoFe₂O₄ and MnFe₂O₄, and the values of specific magnetization (M_s) were 90.8 and 52.3 emu g⁻¹, which thus indicated superparamagnetism. By comparison, ZnFe₂O₄ was 5.02 emu g⁻¹, which was much less than that of CoFe₂O₄ and This article is licensed under a Creative Commons Attribution-NonCommercial 3.0 Unported Licence.

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Fig. 4 UV-Vis diffuse reflectance spectra of MFe₂O₄.



Fig. 5 Magnetic hysteresis loops of MFe₂O₄. The inset shows photos of well-dispersed MFe₂O₄ in water and magnetic separation of MFe₂O₄.

MnFe₂O₄. Difficulty in recycling of the photocatalyst could significantly hinder its extensive application in wastewater treatment. Thus, by taking advantage of the magnetic properties of CoFe₂O₄ and MnFe₂O₄, the photocatalysts can be easily recycled for multiple usages. The inset of Fig. 5 shows that CoFe₂O₄ and MnFe₂O₄ can be readily dispersed in water to form a stable solution. In addition, the photocatalyst rapidly responds to the external magnet because of its excellent magnetic properties. After magnetic separation, most of the CoFe₂O₄ and MnFe₂O₄ photocatalyst particles were drawn to the bottle sidewall. As a result, this dispersion and separation process can be repeatedly applied with CoFe₂O₄ and MnFe₂O₄ using an external magnetic field, which is convenient for their reusability in water treatment and minimizes any secondary pollution.

3.4 Photocatalytic properties of MFe₂O₄

The photocatalytic activity of MFe₂O₄ was estimated by the removal of LVX. To further depict the photocatalytic reaction, the photocatalytic degradation process was also fitted to pseudo first-order kinetics, and the value of the rate constant *k* is equal to the corresponding slope of the fitting line, as shown in Fig. 6. The first-order linear relationship was revealed by the plots of $\ln(C/C_0)$ vs. irradiation time (*t*), where *C* denotes the concentration of LVX at irradiation time *t*, and *C*₀ denotes the concentration during the adsorption equilibrium of the photocatalysts before irradiation. *Via* the first order linear fit, the



Fig. 6 First-order plots for the photocatalytic degradation of LVX using $\mathsf{MFe}_2\mathsf{O}_4.$

determined reaction rate constants *k* were 0.00269, 0.01003, and 0.00691 min⁻¹, respectively, for CoFe₂O₄, MnFe₂O₄, and ZnFe₂O₄. The strongest photocatalytic activity was exhibited by the MnFe₂O₄ sample.

3.5 Activating PMS with MFe₂O₄ to enhance the degradation efficiency

Recently, there has been increased interest by researchers regarding the advanced oxidation processes (AOPs) of the sulfate radical (SO4 ·-).27,28 Because of a higher redox potential, the sulfate radical can degrade many organic pollutants.^{29,30} As a monopersulfate compound, PMS can be activated not only by a transition metal, but also by photogenerated e^{-}/h^{+} to generate strong oxidizing sulfate radicals. Therefore, activating PMS to enhance the degradation rate with MFe₂O₄ under visible light irradiation might be a good choice. The degradation rate of LVX in different systems is shown in Fig. 7a-c. The photocatalytic experiments showed that the photocatalytic activity of MFe₂O₄ was low, and its degradation rate was the slowest of all. The degradation rate for the MFe₂O₄/PMS system without visible light was much slower than that of the MFe₂O₄/Vis/PMS system. The degradation efficiency for activation of PMS with MFe₂O₄ under visible light irradiation was much higher than that for PMS only activated by MFe₂O₄ and MFe₂O₄ only under visible light irradiation. The activation of PMS with MFe₂O₄ resulted from photogenerated e^{-}/h^{+} , Fe, and M, and the degradation rate was much faster than that which occurred when PMS was only activated by MFe₂O₄.

A comparison between this work and other photocatalyst performances for the degradation of LVX was conducted under the same conditions. As shown in Fig. S1,[†] the photocatalytic experiments showed that the efficiency of LVX degradation in the FeWO₄/Vis/PMS and Fe₂(MoO₄)₃/Vis/PMS systems was lower. However, the removal rate increased to 92% in an hour in the CoFe₂O₄/Vis/PMS system. The removal rate also reached 70–80% in an hour in the MnFe₂O₄/Vis/PMS and ZnFe₂O₄/Vis/PMS systems. This indicated that the degradation efficiency for PMS activation with MFe₂O₄ under visible light irradiation was much higher than that with FeWO₄ or Fe₂(MoO₄)₃.

3.5.1 Effect of PMS dosage, pH value, temperature, and LVX concentration. The effect of PMS dosage on the performance of the MFe₂O₄/Vis/PMS system towards LVX degradation



Fig. 7 The degradation rate of MFe₂O₄ in different systems: (a) CoFe₂O₄, (b) MnFe₂O₄, (c) ZnFe₂O₄ ([photocatalyst] = 0.5 g L⁻¹, [LVX] = 10 mg L⁻¹, [PMS] = 2 mM, 0.5 mM, and 3 mM for CoFe₂O₄, MnFe₂O₄, and ZnFe₂O₄, respectively); effect of PMS dosages on the degradation of LVX with PMS activated by MFe₂O₄ photocatalysts: (d) CoFe₂O₄, (e) MnFe₂O₄, and (f) ZnFe₂O₄ ([photocatalyst] = 0.5 g L⁻¹, [LVX] = 10 mg L⁻¹); effect of pH value on the degradation of LVX: (g) CoFe₂O₄, (h) MnFe₂O₄, and (i) ZnFe₂O₄ ([photocatalyst] = 0.5 g L⁻¹, [LVX] = 10 mg L⁻¹); effect of pH value on the degradation of LVX: (g) CoFe₂O₄, (h) MnFe₂O₄, and (i) ZnFe₂O₄ ([photocatalyst] = 0.5 g L⁻¹, [LVX] = 10 mg L⁻¹, [PMS] = 2 mM, 0.5 mM, and 3 mM for CoFe₂O₄, MnFe₂O₄, and ZnFe₂O₄, respectively); effect of temperature on the degradation of LVX: (j) CoFe₂O₄, (k) MnFe₂O₄, and (l) ZnFe₂O₄ ([photocatalyst] = 0.5 g L⁻¹, [LVX] = 0.5 g L⁻¹, [LVX] = 0.5 g L⁻¹, [LVX] = 10 mg L⁻¹, [PMS] = 2 mM, 0.5 mM, and 3 mM for CoFe₂O₄, and ZnFe₂O₄, respectively); effect of LVX concentrations on the degradation of LVX: (m) CoFe₂O₄, (n) MnFe₂O₄, and (o) ZnFe₂O₄ ([photocatalyst] = 0.5 g L⁻¹, [LVX] = 0.5 g L⁻¹, [PMS] = 2 mM, 0.5 mM, and 3 mM for CoFe₂O₄, MnFe₂O₄, and ZnFe₂O₄, respectively).

was investigated (Fig. 7d–f). The LVX degradation rate increased with the increase in the PMS dosage in all systems, and thus, the more dosages of PMS that were added, the more SO_4 ⁻ that was produced. However, the degradation rate gradually decreased when the PMS dosage continuously increased, which was

ascribed to the self-quenching effect between the sulfate radicals and PMS (eqn (1)).^{31,32} The experimental results indicated that the optimal dosage of PMS was 0.2 mmol, 0.05 mmol, and 0.3 mmol, respectively, for $CoFe_2O_4$, $MnFe_2O_4$, and $ZnFe_2O_4$.

$$HSO_5^- + SO_4^{-} \to SO_5^{-} + SO_4^{2-} + H^+$$
 (1)



Fig. 8 Effects of *tert*-butyl alcohol and ethanol addition on the photocatalytic degradation of LVX ([photocatalyst] = 0.5 g L^{-1} , [LVX] = 10 mg L^{-1} , [PMS] = 2 mM, 0.5 mM, and 3 mM for CoFe₂O₄, MnFe₂O₄, and ZnFe₂O₄, respectively): (a) CoFe₂O₄, (b) MnFe₂O₄, (c) ZnFe₂O₄; (d) EPR spectra of CoFe₂O₄ under visible light irradiation (DMPO as the radical trapper); (e) electrochemical impedance spectroscopy of CoFe₂O₄ sample electrodes with and without PMS under visible light irradiation ($\lambda > 420 \text{ nm}$); (f) transient photocurrent responses of CoFe₂O₄ sample electrodes with and without PMS under visible light irradiation ($\lambda > 420 \text{ nm}$).

The pH value had an obvious effect on PMS activation, which was confirmed by a previous study.³³ Therefore, the effect of the pH value on the degradation rate was studied. As exhibited in Fig. 7g-i, the results showed that the pH value clearly influenced the degradation efficiency. According to published literature, it was mainly attributed to the fact that H^+ hindered the production of OH' and SO₄^{•-}, resulting in a decrease in the number of active radicals.³⁴ Therefore, the removal rate increased as the pH value increased.

The influence of the initial temperature on the removal of LVX by the MFe₂O₄ photocatalyst was further investigated. As shown in Fig. 7j–l, the degradation rate increased with increasing temperature, which was due to PMS activation in the endothermic reaction and a higher reactive oxygen species (ROS) production rate at higher temperatures, with LVX being degraded by $CoFe_2O_4$ in 10 min at 40 °C. This might be caused by the self-activation reaction of PMS under higher temperatures.³²

The effects of initial LVX concentration on the degradation behaviour are shown in Fig. 7m–o. As the initial levofloxacin concentration increased from 10 mg L^{-1} to 20 mg L^{-1} , the degradation kinetics decreased. This may have occurred due to the higher concentration of LVX that resulted in additional active ROS being produced.³² However, the limited photocatalysts and PMS could not produce sufficient radicals to degrade a solution with higher LVX concentration.

3.5.2 Degradation mechanism after PMS activation with MFe_2O_4 . To investigate the degradation mechanism that occurs after activation of PMS with MFe_2O_4 and confirm the role of the active species in the degradation process, *tert*-butanol and ethanol were introduced as radical scavengers. Ethanol was chosen as the radical scavenger for the sulfate radical and hydroxyl radical.³² However, *tert*-butanol was the only effective scavenger for the hydroxyl radical.^{35–37} After 1 mmol *tert*-butanol

was added to LVX solution, there was not an obvious inhibitory effect (Fig. 8a–c), which indicated that the hydroxyl radical was not the primary active specie in the $MFe_2O_4/PMS/Vis$ system. Obvious inhibition of the degradation rate was observed after excess ethanol was added to the LVX solution, in which the molar ratio of ethanol and PMS was 2000 : 1 (Fig. 8a–c). When the molar ratio was increased to 4000 : 1, the inhibition of the degradation rate was more obvious, which indicated that sulfate radical was the key active specie in the Vis/MFe₂O₄/PMS system.

To confirm the above results, electron paramagnetic resonance (EPR) spectroscopy was used to trace intermediate radical species that existed in the Vis/CoFe₂O₄/PMS system. The trapping agent 5,5-dimethylpyrrolineoxide (DMPO) was used to capture radicals 'OH and SO₅⁻⁻ in the Vis/CoFe₂O₄/PMS system.^{38,39} As shown in Fig. 8d, the characteristic peak signals of DMPO-'OH and DMPO-SO₄⁻⁻ adducts were found,⁴⁰ which not only proved the coexistence of 'OH and SO₄⁻⁻ in this system.

Electrochemical impedance spectroscopy (EIS) was used to characterize electrochemical interfacial reactions. The photocatalytic decomposition of LVX can be explained as an electrochemical oxidation reaction in which reactants supply electrons to an anode. A smaller arc radius of the EIS Nyquist plot represents a faster electron transfer rate and a more efficient electron-hole separation.⁴¹ Fig. 8e shows the EIS response of $CoFe_2O_4$ under visible light irradiation ($\lambda > 420$ nm). The radius of the arc on the EIS Nyquist plot reflects the reaction rate occurring at the surface of the electrode. The radii tended to significantly decrease after adding PMS to the solution, indicating that both charge-transfer resistance and capacitive reactance decreased. This suggested that there was an effective separation of photogenerated electron-hole pairs and that fast interfacial charge transfer to the electron donor/electron



Fig. 9 (a, d, g, and j) XPS spectra of Fe 2p and Co 2p for CoFe₂O₄ before and after photocatalytic oxidation of LVX. (b, e, h, and k) XPS spectra of Fe 2p and Mn 2p for MnFe₂O₄ before and after photocatalytic oxidation of LVX. (c, f, i, and l) XPS spectra of Fe 2p and Zn 2p for ZnFe₂O₄ before and after photocatalytic oxidation of LVX.

acceptor occurred, as suggested by Leng *et al.*⁴² With the existence of PMS, the interfacial electron transfer rate and the electron-hole separation efficiency of the Vis/CoFe₂O₄/PMS system greatly increased.

To further confirm that PMS efficiently captured the photogenerated electrons in the Vis/CoFe₂O₄/PMS system, the transient photocurrent responses of CoFe₂O₄ were also measured under visible light irradiation. As shown in Fig. 8f, the introduction of PMS significantly decreased the density of photocurrent in the Vis/CoFe₂O₄/PMS system. It is likely that the photogenerated electrons were efficiently trapped by the PMS, which was consistent with results from a previous report.⁴³

To investigate the role of iron, zinc, and manganese in Vis/ MFe_2O_4/PMS systems, the XPS spectrum was used to analyze the

samples before and after photocatalytic experiments (Fig. 9). In the XPS spectrum of $CoFe_2O_4$, the Fe 2p peaks at 707.95 eV and 721.53 eV were assigned to Fe $2p_{3/2}$ and Fe $2p_{1/2}$ (Fig. 9a and d), respectively, showing that a portion of Fe species existed in the form of Fe²⁺ in CoFe₂O₄ before the photocatalytic experiment. However, the area of Fe³⁺ peaks decreased, and the area of Fe²⁺ peaks increased after activation of PMS with CoFe₂O₄, which indicated that Fe³⁺ on the surface of CoFe₂O₄ was partially transformed to Fe²⁺.⁴⁴ This proved that the regeneration of Fe(m) and a cycle of Fe(m)/Fe(n) occurred in the Vis/CoFe₂O₄/ PMS systems.⁴⁵ The Co 2p peaks at 776.93.45 and 792.98 eV were assigned to the binding energies of Co $2p_{3/2}$ and Co $2p_{1/2}$ (Fig. 9g and j), respectively, indicating that a portion of Co species existed in the form of Co³⁺. However, the area of the



Scheme 1 Schematic drawing illustrating the mechanism of activation of PMS with MFe_2O_4 photocatalyst under visible light irradiation.

Co²⁺ peaks decreased, and the area of the Co³⁺ peaks increased after activation of PMS with CoFe₂O₄, which indicated that Co²⁺ on the surface of CoFe₂O₄ was partially transformed to Co^{3+,45} This proved that regeneration of Co(π) and a cycle of Co(π)/ Co(π) occurred in the Vis/CoFe₂O₄/PMS systems. Similar conclusions were drawn from the XPS spectra (Fig. 9) of MnFe₂O₄ and ZnFe₂O₄ before and after activation of PMS. It proved that cycles of Fe(π)/Fe(π), Co(π)/Co(π), Mn(π)/Mn(π), and Zn(π)/Zn(π) existed in the Vis/MFe₂O₄/PMS systems.

According to the above results, the mechanism of activating PMS with MFe₂O₄ is shown in Scheme 1 (eqn (2)-(20)). The MFe₂O₄ photocatalyst absorbed visible light and then produced photogenerated e⁻/h⁺ pairs.⁴⁶ PMS was activated by photogenerated electrons (e⁻) and then produced SO₄^{•-} radicals. SO_4 reacted with OH^- to form OH^- and SO_4^{2-} . SO_4^{2-} then reacted with h^+ to form SO₄[•]. The h^+ also activated PMS to produce SO5.-, and then, SO5.- reacted with each other to generate SO₄⁻⁻ (eqn (2)-(7)).^{47,48} PMS was also activated by iron to produce sulfate radicals. Because of the effective electron transfer from PMS to Fe(III) and Fe(II), which resulted in regeneration of Fe(m) and cycling between Fe(m) and Fe(m), enhanced degradation efficiency was achieved in activating PMS with CoFe₂O₄ under visible light irradiation (eqn (8)-(10)).⁴⁴ Similarly, there was also regeneration of Co(II) and cycling between Co(II) and Co(III) for CoFe₂O₄ (eqn (11)-(13)),⁴⁵ which indicated that cobalt also activates PMS to enhance the degradation efficiency, in addition to iron. MnFe₂O₄ and ZnFe₂O₄ activated PMS and regenerated in the same manner as that of CoFe₂O₄ (eqn (14)-(19)).⁴⁴ Finally, LVX was mainly degraded by sulfate radicals and hydroxyl radicals (eqn (20)).49-51

$$MFe_2O_4 + h\nu \rightarrow h^+ + e^-$$
(2)

$$HSO_5^- + e^- \rightarrow SO_4^{-} + OH^-$$
(3)

$$SO_4^{\cdot-} + OH^- \rightarrow SO_4^{2-} + OH$$
 (4)

$$\mathrm{SO_4}^{2-} + \mathrm{h^+} \to \mathrm{SO_4}^{\cdot-}$$
 (5)

$$\mathrm{HSO}_{5}^{-} + \mathrm{h}^{+} \to \mathrm{SO}_{5}^{\cdot -} + \mathrm{H}^{+} \tag{6}$$

$$\mathrm{SO}_5^{\bullet-} + \mathrm{SO}_5^{\bullet-} \to 2\mathrm{SO}_4^{\bullet-} + \mathrm{O}_2$$
 (7)

$$Fe(III) + HSO_5^- \rightarrow Fe(II) + SO_5^{-} + H^+$$
(8)

$$Fe(II) + HSO_5^- \rightarrow Fe(III) + SO_4^{-} + OH^-$$
(9)

$$\mathrm{SO}_5^{\bullet-} + \mathrm{SO}_5^{\bullet-} \to 2\mathrm{SO}_4^{\bullet-} + \mathrm{O}_2 \tag{10}$$

$$\operatorname{Co(II)} + \operatorname{HSO}_5^{-} \to \operatorname{Co(III)} + \operatorname{SO}_4^{-} + \operatorname{OH}^{-}$$
(11)

$$Co(III) + HSO_5^{-} \rightarrow Co(II) + SO_5^{-} + H^+$$
(12)

$$SO_5^{\bullet} + SO_5^{\bullet} \rightarrow 2SO_4^{\bullet} + O_2$$
 (13)

$$Mn(II) + H_{S}O_{5}^{-} \rightarrow Mn(III) + SO_{4}^{\cdot -} + OH^{-}$$
(14)

$$Mn(III) + HSO_5^- \rightarrow Mn(II) + SO_5^{\bullet-} + H^+$$
(15)

$$\mathrm{SO}_5^{\cdot-} + \mathrm{SO}_5^{\cdot-} \to 2\mathrm{SO}_4^{\cdot-} + \mathrm{O}_2$$
 (16)

$$Zn(II) + HSO_5^- \rightarrow Zn(III) + SO_4^{-} + OH^-$$
(17)

$$Zn(III) + HSO_5^- \rightarrow Zn(II) + SO_5^{\cdot -} + H^+$$
(18)

$$\mathrm{SO}_5^{\cdot -} + \mathrm{SO}_5^{\cdot -} \to 2\mathrm{SO}_4^{\cdot -} + \mathrm{O}_2 \tag{19}$$

$$LVX + SO_4^{-} + OH \rightarrow degradation$$
 (20)

3.6 The degradation pathway of LVX

During the process of CoFe₂O₄ activating PMS to enhance degradation efficiency under visible light irradiation, the SO₄.attacks LVX and then disassembles its molecular structure. The intermediate products then form, and the possible degradation pathways are shown in Scheme 2 by UPLC-MS analysis (Fig. S2[†]). As the weak electron donor, methyl groups facilitate the attack on LVX by electrophilic species (SO₄^{•-}) in the demethylation process, resulting in the formation of the intermediate product of m/z = 310. With sustained degradation, the SO_4 · · · cleaved the ring, and the intermediate products of m/z =279 were formed.⁵² With photodegradation proceeding, the nitrogen bonded with C7 was exfoliated to obtain the compound of m/z = 223. Then, the SO₄⁻⁻ continued to attack the intermediate products and disconnect the ring of the intermediate structure, and the intermediate compound of m/z178 formed. As photodegradation proceeded, SO_4 . continued to cleave the functional groups, and glycol was formed (m/z = 65). Finally, the above intermediate products were mineralized into H2O, CO2, and others.

3.7 Stability of MFe₂O₄

To study the reusability and stability of photocatalysts, recycling experiments were carried out (Fig. S3a–c†). When the degradation experiment was finished, the CoFe₂O₄ was filtered from the LVX solution. The CoFe₂O₄ was further dried at 60 °C and was used for the next degradation experiment. After three recycling experiments, there were no obvious changes in LVX degradation. Similarly, recycling experiments were conducted in Vis/ MnFe₂O₄/PMS and Vis/ZnFe₂O₄/PMS systems. It was found that

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Scheme 2 Possible degradation pathway of LVX.

there was satisfactory reusability and stability for the three photocatalysts.

4 Conclusions

Nanosized MFe₂O₄ was successfully synthesized by a simple sol-gel method. The photocatalytic properties of MFe₂O₄ were investigated under visible light irradiation, and they showed decreased photocatalytic activity for the degradation of levofloxacin hydrochloride under visible light irradiation. For enhanced photocatalytic activity, MFe₂O₄ was used to activate peroxymonosulfate to degrade levofloxacin hydrochloride under visible light irradiation. A series of systematic experiments proved that the PMS dosage conferred an obvious effect on the degradation rate. Furthermore, there was a noteworthy effect by the levofloxacin hydrochloride concentration and the pH value on the levofloxacin hydrochloride degradation. After activation of PMS with MFe₂O₄, a degradation mechanism was proposed and proved by EPR, EIS, transient photocurrent, and XPS. Kinetic studies using radical scavenger technologies suggested that sulfate radical controls the degradation process.

Author contributions

Di Li: project administration, Mingyang Long: investigation, Hongmiao Li: data curation, Xinguo Ma: methodology, Qianqian Zhao: formal analysis, and Qi Wen: writing – review and editing.

Conflicts of interest

There are no conflicts to declare.

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