Magnetic Fe₃O₄@CoFe-LDH nanocomposite heterogeneously activated peroxymonosulfate for degradation of azo-dye AO7†

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In this study, a novel core@shell magnetic nanocomposite Fe₃O₄/CoFe-layered double hydroxide (Fe₃O₄@CoFe-LDH) was successfully synthesized by the co-precipitation method, and then employed as an efficient heterogeneous catalyst for activation of peroxymonosulfate (PMS) in removal of azo-dye acid orange 7 (AO7). The as-obtained nanocomposite was characterized by X-ray diffraction (XRD), Fourier transform infrared spectroscopy (FT-IR), scanning electron microscopy (SEM) and vibrating sample magnetometer (VSM). The results from these characterizations showed Fe₃O₄@CoFe-LDH to possess good ferromagnetism and a perfect crystalline structure with a typical core@shell morphology.

The system of Fe₃O₄@CoFe-LDH11/PMS (cobalt : iron molar ratio of 1 : 1) achieved 95.1% removal rate of AO7 (40 mg L⁻¹) within 15 min under the optimized conditions, which outperformed bare Fe₃O₄ and raw CoFe-LDH11. Meanwhile, Fe₃O₄@CoFe-LDH11 displayed good adaptability in a wide pH range from 4 to 9 and relatively low PMS activation energy (39.9 kJ mol⁻¹). The interference tests revealed HCO₃⁻/CO₃²⁻ to possess the strongest restriction effect. Only 57.7% AO7 was removed when 10 mM HCO₃⁻ was introduced, which was ascribed to HCO₃⁻ not only serving as a radical scavenger, but also increasing the pH of the system. The radical quenching tests demonstrated SO₄²⁻ as the dominant reactive species during the catalytic reaction. Based on X-ray photoelectron spectroscopy (XPS) analysis, the core structure of Fe₃O₄ served as an electron donor for accelerating the cycle of Co(II)/Co(III) at the active site of the LDH outer shell. Also, Fe₃O₄@CoFe-LDH exhibited outstanding stability and recyclability, and maintained high degradation efficiency of AO7 even after five cycles. In sum, the proposed magnetic Fe₃O₄@CoFe-LDH nanocomposite has great potential for remediation of wastewater contaminated with synthetic dyes.

1. Introductions

Synthetic dyes are chemical products widely used in the textile, leather, paper, cosmetics, and other industries. The increasing demand for dye production has led to the discharge of large quantities of wastewater, thereby contaminating natural water bodies and causing severe environmental problems. Besides, most of the dyes are toxic and resist natural degradation. Hence, the elimination of dyes from water bodies is of great importance for environmental remediation and public health. Numerous reports have so far been published on the removal of dyes from aquatic environments by means of adsorption, microbiological treatment, electrochemistry, photolysis, and so on. Unfortunately, these strategies do not meet the actual requirement due to complex operational management, low removal rates and high energy consumption. By comparison, advanced oxidation processes (AOPs) are ideal and appropriate for the treatment of non-biodegradable pollutants.

Recently, sulfate radical-based advanced oxidation processes (SR-AOPs) have attracted increasing attention due to their high redox potentials (2.5–3.1 V) and selectivity of sulfate radicals (SO₄⁻). Compared to HO· (1.9–2.7 V), the benchmark of radicals utilized for AOPs, SO₄⁻ possesses stronger oxidation capacity and longer half-life (30–40 µs, HO· less than 1 µs). Therefore, SR-AOPs exhibit better degradation and mineralization performances toward intractable contamination of water bodies. In general, SO₄²⁻ is produced by activation of peroxymonosulfate (PMS) or peroxodisulfate (PDS) as an oxidant. SR-AOPs can be divided into thermolysis, ultrasonic, photocatalysis, electrocatalysis, and transition metal activation. Among the above activation strategies, transition metal activation has extensively been employed owing to its lower energy-input and higher degradation rate. According to previous
studies, cobalt ion (Co²⁺) displayed the best catalytic properties in PMS activation. However, the lack of effective methods for recovering Co²⁺ from wastewater may cause secondary pollution and threaten both the ecological environment and human health. Hence, employing cobalt-based heterogeneous catalysts as PMS activators is a promising for the treatment of wastewater contaminated by dyes.

Due to the rich active sites and unique hierarchical architectures, two-dimensional (2D) materials, such as graphene and layered double hydroxides (LDH), have been considered as perfect PMS activator. LDH are multi-metal layered clay materials, consisting of positive metal cations (divalent metals and trivalent metals) and interlayer negative anions (NO₃⁻, Cl⁻, CO₃²⁻, etc.). The unique layered structure of LDH combined with its tuneable controllability of metal ions and structural stability made these materials promising for water splitting, energy storage and environmental remediation. For instance, Guo et al. successfully synthesized CoCu-LDH composites with great performance in PMS activation for removal of lomefloxacin. Zhao et al. reported that CoMn-LDH/PMS system achieved 52.2% TOC removal rate of acid orange G. Hence, cobalt-based LDH catalysts look efficient for the activation of PMS. However, the catalyst agglomeration might significantly impede the catalytic performance of LDH. Moreover, nanoparticle catalysts are difficult to separate and recover from the water bodies, thereby limiting their practical applications. Previous literature demonstrated Fe₃O₄ as an eco-friendly and non-toxic carrier with high magnetic separation force. For example, Jung et al. reported magnetic Fe₃O₄@Mgl-Al LDH adsorbent with large specific surface area, possessed much greater adsorption capacity for iodide. However, despite the extensive investigation of LDH coating on Fe₃O₄ with fast separation and recovery properties, little is still known about Fe₃O₄@LDH as a heterogeneous catalyst for PMS activation and wastewater purification.

Herein, a novel core@shell Fe₃O₄@CoFe-LDH was synthesized and applied for PMS activation in azo-dye AO7 degradation. The nanocomposite was prepared via a simple co-precipitation method and characterized by various analytical methods. The different reaction systems and effects of various parameters like Co/Fe molar ratio, pH, temperature, PMS dosage, and catalyst concentration were all systematic studied. The influences of common inorganic anions found in an aquatic environment on Fe₃O₄@CoFe-LDH/PMS performances were also tested to evaluate the interference capability. The as-obtained nanocomposite did not only exhibit great degradation rate of AO7, but also completely recovered from the aquatic environment through the application of an external magnetic field. Scavenging experiments were also carried out to identify the major involved radicals and possible reaction mechanisms. Overall, the developed magnetic nanocomposites showed good stability as PMS heterogeneous catalysts for dye wastewater treatment.

2. Experimental procedure

2.1. Materials

All chemical reagents used in this study were of analytical grade without further purification. Ferric chloride hexahydrate (FeCl₃·6H₂O, 99.0 wt%), sodium dihydrogen phosphate (NaH₂PO₄, 99.0 wt%), ethylene glycol (EG, 99.0 wt%), polyvinylpyrrolidone (PVP, MW = 40 000), acid orange 7 (AO7, 99.0 wt%) and peroxymonosulfate (2KHSO₅·K₂SO₄, 98.0 wt%) were purchased from Adamas-Beta Reagent Co., Ltd. Sodium hydride (NaOH, 98.0 wt%), sodium carbonate (Na₂CO₃, 99.8 wt%), sodium hydrogen carbonate (NaHCO₃, 99.5 wt%), sodium acetate (NaAc, 99.0 wt%), and tert-butanol (TBA, 99 wt%) were purchased from Shanghai Titan Scientific Co., Ltd. Cobalt chloride hexahydrate (CoCl₂·6H₂O, 98.0 wt%) and sulfuric acid (H₂SO₄) were purchased from Sinopharm Chemical Reagent Co., Ltd. All chemicals were commercially available. Deionized (DI) water was used to prepare solutions throughout the study.

2.2. Synthesis of magnetic Fe₃O₄@CoFe-LDH

Fe₃O₄ nanospheres were prepared via the solvothermal method with some modifications. Briefly, FeCl₃·6H₂O (10 mmol) and sodium acetate (52 mmol) were dissolved in 66 mL of ethylene glycol. After vigorously stirred for 30 min, the mixed solution was transferred to a 100 mL Teflon-lined stainless-steel autoclave and then heated at 473 K for 8 h. Finally, the obtained black precipitate was washed with DI and ethanol for several times, and dried at 333 K in an oven for 24 h.

As illustrated in Fig. 1, the magnetic Fe₃O₄@CoFe-LDH composites were synthesized via a simple co-precipitation method. FeCl₃·6H₂O (5 mmol) and CoCl₂·6H₂O (5 mmol) were dissolved in 100 mL DI water and stirred constantly for 30 min at ambient temperature. Then, the alkaline solution of NaOH (0.5 M) and Na₂CO₃ (0.2 M) was slowly added to the above solution with mechanical stirring (300 rpm) until pH reached 6.5. Subsequently, PVP (1.0 g) and as-prepared Fe₃O₄ (0.5 g) were added into the mixed solution under the ultrasound for 15 min at 298 K. The metal cation in mixed solution would be adsorbed on the surface of PVP modified Fe₃O₄ through the effect of electrostatic force of attraction. The pH of mixture was adjusted to 10 by the above-mentioned alkaline. The obtained suspension was aged at 338 K for 24 h. Finally, the products were collected by external magnetic field, washed with deionized water and ethanol, and dried overnight in oven at 333 K. The obtained red-brown composites were labeled as Fe₃O₄@CoFe-LDH11. Meanwhile, other materials with different Co/Fe ratios (4 : 1, 2 : 1, 1 : 2, 1 : 4, total metal concentration is 10 mmol) were synthesized based on the above method, and labeled as Fe₃O₄@CoFe-LDH14, Fe₃O₄@CoFe-LDH21, Fe₃O₄@CoFe-LDH12 and Fe₃O₄@CoFe-LDH14. Bare CoFe-LDH11 was also made in the absence of Fe₃O₄ for the control comparison.

Fig. 1 Schematic illustration of the synthetic route of Fe₃O₄@CoFe-LDH.

And, the detailed information of preparation method of CoFe₂O₄ was shown in the ESI (Text S1).

2.3. Characterizations

The phase structure of as-fabricated catalyst was identified by an X-ray diffraction (XRD, Ultima IV, Rigaku, Japan) using Cu Kα radiation (λ = 1.54056 Å, 40 kV, 20 mA) in the range of 5°–90° at a scan rate of 2° min⁻¹. The surface functional group of the composites was examined by Fourier transform infrared spectroscopy (FTIR, Thermo Nicolet IS10, Nicolet, USA) at room temperature. The morphology and chemical analysis were recorded on a scanning electron microscope (SEM, Gemini 300, ZEISS, Germany) equipped with an energy-dispersive X-ray spectrometer system (EDS, Quantax 400, Bruker, Germany). The specific surface area and porosity were analyzed using N₂ adsorption/desorption experiment (ASAP-2460, Micromeritics, USA). The magnetic properties of synthesized materials were measured at room temperature on a vibrating sample magnetometer (VSM, PPMS-9, Quantum Design, USA). The surface characteristics of the fresh and used samples were measured by the X-ray photoelectron spectroscopy (XPS, ESCALAB 250Xi, Thermo Fisher Scientific, USA). The binding energies of photoelectron peaks was corrected by referring to the binding energy of the C 1s neutral carbon peak at 284.6 eV.

2.4. Degradation experiments

All the degradation experiments were carried out in a 250 mL beaker containing 100 mL of AO7 (40 mg L⁻¹) at 298 K with mechanical stirring. In a typical procedure, the solid catalyst was added into the AO7 solution and mechanical stirred at 250 rpm for 5 min to dispersed completely. Subsequently, 0.12 mM of PMS was injected into the above solution. The initial pH of reaction system was adjusted to 6.0 by NaOH (0.1 M) and 0.12 mM of PMS was injected into the above solution. The initial 250 rpm for 5 min to dispersed completely. Subsequently, 100 mM of radical scavengers (TBA or ethanol) were mixed with AO7 solution before the catalyst and PMS were introduced.

3. Results and discussion

3.1. Characterization

As shown in Fig. 2a, the XRD peaks of bare Fe₃O₄ could be observed at 2θ of 30.2°, 35.6°, 43.2°, 57.2° and 62.8°, matched well with cubic iron oxide phase (JCPDS no. 19-0629). The main peaks of bare CoFe-LDH11 were located at 11.6°, 23.4°, 34.0°, 38.7°, 38.7°, 46.0°, 59.1° and 60.6°, indexed to (003), (006), (012), (015), (018), (110) and (113) facets of LDH (JCPDS no. 50-0235), respectively, which were consistent with the results in the previous works. Besides, the dominant peaks of magnetic Fe₃O₄@CoFe-LDH11 appeared at 2θ of 11.6°, 23.6°, 30.3°, 34.0°, 35.6°, 38.6°, 43.2°, 46.0°, 57.3°, 60.6°, 62.8° and 63.0°, which were familiar with both phase of pure Fe₃O₄ and LDH. Due to the unique structure of Fe₃O₄@CoFe-LDH11, it was observed that XRD peaks was weaker than bare Fe₃O₄ and CoFe-LDH11. These results suggested that the hybrids were successfully synthesized and consisted of Fe₃O₄ and LDH. The FT-IR spectra of Fe₃O₄, CoFe-LDH11 and Fe₃O₄@CoFe-LDH11 are shown in Fig. 2b. Clearly, one may observe that different samples have both three same peaks. The bands at 3418 cm⁻¹, 1384 cm⁻¹ were attributed to the O–H stretching vibration, bending vibration of hydroxyl deformation mode and –OH groups, respectively, which might be attribute to H₂O molecules adsorbed on the surface of composites interlayer water of LDH. It is worth noting that the hybrid has a characteristic peak of 1354 cm⁻¹ that disappeared in the Fe₃O₄, which was ascribed to the vibrational absorption of interlayer anions CO₃²⁻ of LDH. In addition, the absorption peaks at range of 763–471 cm⁻¹ were metal–oxygen (M–O) and metal–hydroxyl (M–OH) groups vibrations. The bands at 763 cm⁻¹ and 471 cm⁻¹ were associated with Co–OH and Co–O stretching vibration. The peaks located at 627 cm⁻¹ and 586 cm⁻¹ were corresponded to Fe–O stretch vibration, while the intensity of Fe₃O₄@CoFe-LDH11 around 586 cm⁻¹ was sharply decreased than bare Fe₃O₄ as a result of LDH encapsulated the Fe₃O₄ nanoparticles. Besides, the magnetic Fe₃O₄@CoFe-LDH11 showed a sharp band at 627 cm⁻¹, indicating the formation of Fe–O–Fe between Fe₃O₄ and LDH.

Fig. 2 XRD patterns (a) and FT-IR spectra (b) of bare Fe₃O₄, raw CoFe-LDH11 and Fe₃O₄@CoFe-LDH11.

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The magnetic properties of Fe₃O₄ and Fe₃O₄@CoFe-LDH11 were verified by the vibration sample magnetometry (VSM). As depicted in Fig. 3a, the saturation magnetization ($M_s$) value of Fe₃O₄ and Fe₃O₄@CoFe-LDH11 were 72.48 and 24.15 emu g⁻¹, respectively. Due to the non-magnetic substance LDH coating on the surface of Fe₃O₄, the magnetic properties of hybrid were weaker than the pristine Fe₃O₄. However, due to the non-magnetic substance LDH coating on the surface of Fe₃O₄, the magnetic properties of hybrid were weaker than the pristine Fe₃O₄. Despite this, the inset picture of Fig. 3a proved that Fe₃O₄@CoFe-LDH11 still could be completely separated from the solution within 1 min by external magnetic field, which was beneficial to the recycling of heterogeneous catalyst. Moreover, the Fe₃O₄@CoFe-LDH11 maintained 22.75 emu g⁻¹ of $M_s$ after PMS activation which indicated that it displayed outstanding stability and recyclability. The specific surface area and pore volume of samples were tested by Brunauer-Emmet-Teller method (Fig. 3b). N₂ adsorption–desorption isotherms of Fe₃O₄@CoFe-LDH11 was identified as a type IV curve, which indicated that Fe₃O₄@CoFe-LDH11 has both microporous and mesoporous pore structures, and the pore diameters of samples shown in the inset figure are at the range of 2–10 nm. Based on the results of Table 1, the specific surface area of Fe₃O₄@CoFe-LDH11 significantly increased from 46.22 to 122.65 m² g⁻¹ compared to bare Fe₃O₄ and was slightly less than that of CoFe-LDH11 which was reason of Fe₃O₄ core with lower surface area.²⁵ According to the literatures, it was reasonable to speculate that large surface area and rich pore structure of Fe₃O₄@CoFe-LDH11 might provide more active site for PMS activation and contaminants degradation.³⁷

SEM and EDS were utilized to determine the morphologies and element composition of different samples. As illustrated in the Fig. 4a, the pristine Fe₃O₄ possessed typical nanospherical structure with diameter of 200–500 nm and has an excellent dispersion. Fig. S3† suggested that CoFe-LDH11 exhibited nanosheet structure which was consistent with the results in the previous reports.²⁸ However, it should be noted that the detrimental effects of agglomeration make the size of LDH increased which might prevent the dye molecules mass diffusion. In the Fig. 4b, the shell of CoFe-LDH wrapped on the surface of Fe₃O₄ nanosphere, and because of relieving aggregation by Fe₃O₄ as a carrier, the dimensions of LDH were absolutely tinier than raw LDH. Then, the results of element mapping pictures (Fig. 4c–e) indicated that the Co elements were homogeneously distributed on the surface of Fe₃O₄. Overall, above results of characterization were compelling lines of evidence that Fe₃O₄@CoFe-LDH of core@shell structure was prepared successfully.

### 3.2. AO7 degradation of different catalysts

As shown in Fig. 5a, different reaction systems displayed variable performances toward the removal of target contaminants. The removal rate of AO7 by Fe₃O₄@CoFe-LDH11 through adsorption reached only 8.3%, while that of other samples was below 8.3%, and can be neglected. Using PMS without LDH and

| Samples                      | Specific surface area (m² g⁻¹) | Pore volume (cm³ g⁻¹) | Average pore size (nm) |
|------------------------------|-------------------------------|-----------------------|------------------------|
| Fe₃O₄                        | 46.22                         | 0.08                  | 7.37                   |
| CoFe-LDH11                   | 158.53                        | 0.43                  | 10.72                  |
| Fe₃O₄@CoFe-LDH11             | 122.65                        | 0.23                  | 8.22                   |

Fig. 4 SEM images of Fe₃O₄ (a) and Fe₃O₄@CoFe-LDH11 (b), EDS mapping of O, Fe, Co for Fe₃O₄@CoFe-LDH11 (c–e).
Where catalytic performances and a fast rate of PMS activation. The CoFe-LDH11/PMS system in this study had relatively high kinetics (eqn (1)).

\[
\ln \frac{C_t}{C_0} = -k_{\text{obs}} t
\]

where \( C_0 \) is the initial concentration of AO7, \( C_t \) is the concentration of AO7 at certain reaction time and \( k_{\text{obs}} \) (min\(^{-1}\)) represents the pseudo-first-order kinetics reaction rate constant for AO7 degradation. The rate constant \( k_{\text{obs}} \) of Fe\(_3\)O\(_4@\)CoFe-LDH11 calculated according to eqn (1) and shown in Fig. S5a\(^{+}\) was 0.549 min\(^{-1}\), a value 2.2-fold that of CoFe-LDH11 (\( k_{\text{obs}} \) = 0.255 min\(^{-1}\)) and 7.7-fold that of the CoFe\(_2\)O\(_4\) (\( k_{\text{obs}} \) = 0.071 min\(^{-1}\)). Among the catalytic systems, the excellent degradation efficiency of Fe\(_3\)O\(_4@\)CoFe-LDH11 was attributed to the synergistic effect between LDH and Fe\(_3\)O\(_4\). For further comparison, different reported catalysts used for the activation of PMS and degradation of AO7 are listed in Table 2. Fe\(_3\)O\(_4@\)CoFe-LDH11/PMS system in this study had relatively high catalytic performances and a fast rate of PMS activation.

The effects of Co/Fe molar ratio on AO7 degradation efficiency were also investigated and the results are depicted in Fig. 5b. All samples with different Co/Fe molar ratios displayed good capacity for activating PMS with a removal rate of AO7 reaching 90% within 15 min. As cobalt content increased from 1 : 4 to 1 : 1, the degradation efficiency of AO7 after 5 min varied from 75.9% to 91.7%. Therefore, within this range, the cobalt content played a key role in controlling the reaction rate. Further increase in cobalt content led to the \( k_{\text{obs}} \) values of Fe\(_3\)O\(_4@\)CoFe-LDH11, Fe\(_3\)O\(_4@\)CoFe-LDH21 and Fe\(_2\)O\(_4@\)CoFe-LDH41 reaching 0.549 min\(^{-1}\), 0.373 min\(^{-1}\) and 0.431 min\(^{-1}\), respectively (Fig. S5b\(^{+}\)). However, further raising the molar ratio of cobalt may cause excess divalent cobalt that could react with \( \text{SO}_4^{2-} \) to generate \( \text{SO}_4^{2-} \). This, in turn, resulted in a slightly declined AO7 degradation rate.\(^{49}\) Hence, a Co/Fe molar ratio of 1 : 1 was selected as the optimal value for subsequent catalytic experiments.

### 3.3. Effect of reaction conditions on catalytic activity

#### 3.3.1. Initial value of pH and temperature

The initial pH of the solution plays a key role in the catalytic degradation efficiency of the catalyst/PMS system.\(^{41}\) Thus, the influence of various pH values on AO7 degradation was studied at pH 4 to 10 and the results are presented in Fig. 6a. At pH of 4–9, the removal rate of AO7 reached over 90% after 15 min. The fastest degradation effect was recorded at pH 6 with 94% pollutants degraded in 5 min. According to the previous reports, the \( pK_{a1} \) and \( pK_{a2} \) of PMS are 0 and 9.4, respectively.\(^{42}\) Hence, at pH from 6–9, \( \text{HSO}_5^- \) as the main active species of PMS could easily be

![Fig. 5 AO7 degradation in different system (a) and effect of Co/Fe molar ration on AO7 degradation (b). Experiment conditions: pH = 6, T = 298 K, V = 0.1 L, [AO7] = 40 mg L\(^{-1}\), [catalyst] = 50 mg L\(^{-1}\), [PMS] = 0.12 mM.](image)

### Table 2 The comparison of AO7 removal efficiency by different catalyst in literatures

| Catalyst (g L\(^{-1}\)) | AO7 concentrations (mg L\(^{-1}\)) | PMS [mM] | Conditions | Removal efficiency | Ref. |
|-------------------------|-----------------------------------|----------|------------|-------------------|-----|
| CoFe\(_2\)O\(_4\) (0.50) | 20                                | 0.80     | pH = 6.3, T = 300 K | 96.0% (40 min) | 37  |
| β-FeOOH (5.70)          | 20                                | 1.00     | —          | 99.9% (180 min) | 38  |
| Porous CuO (0.05)       | 20                                | 0.10     | pH = 8.0, T = 298 K | 85.0% (60 min) | 39  |
| Co\(_3\)O\(_4@\)NF (——) | 35                                | 0.50     | pH = 6.7, T = 298 K | 98.0% (30 min) | 40  |
| CoCuAl-LDO (0.10)       | 20                                | 0.16     | —          | 97.6% (20 min) | 42  |
| MnFe-LDH (0.20)         | 20                                | 0.33     | pH = 6.1, T = 298 K | —              | —   |
| Fe\(_3\)O\(_4@\)CoFe-LDH11 (0.05) | 40                                | 0.12     | pH = 6.0, T = 298 K | 95.1% (15 min) | This work |
activated by Fe3O4@CoFe-LDH11 to generate SO42− radicals, leading to the fast AO7 degradation rate. As pH gradually decreased, the time required to achieve 90% AO7 removal rate was prolonged. This resulted from the increment in the concentration of H+, causing HSO5− to form more stable species of H2SO5. Simultaneously, the formation of CoOH+ could also inhibit catalyst activation of PMS to generate SO42−. At pH 10, large amounts of OH− could combine with active sites on the catalyst surface to form complexes, hindering the direct contact between Fe3O4@CoFe-LDH11 and PMS. This reduced

Fig. 6 Effects of (a) initial of pH, (b) temperature, (c) catalyst concentrations and (d) PMS dosages on AO7 removal efficiency. Experiment conditions: pH = 6 (except a), T = 298 K (except b), V = 0.1 L, [AO7] = 40 mg L−1, [catalyst] = 50 mg L−1 (except c), [PMS] = 0.12 mM (except d).

Fig. 7 Effects of anions on AO7 degradation (a) Cl−, (b) HCO3−, (c) H2PO4− and (d) NO3−. Experiment conditions: pH = 6, T = 298 K, V = 0.1 L, [AO7] = 40 mg L−1, [catalyst] = 50 mg L−1, [PMS] = 0.12 mM.
the removal rate of AO7 to 76%. Overall, Fe₃O₄@CoFe-LDH11/PMs exhibited excellent catalytic degradation of AO7 in a wide range of pH (4–9).

The effects of reaction temperature on AO7 degradation were also studied from 298 K to 318 K. As drawn in the illustration of constant and absolute temperature, respectively, kₐₚ value of Fe₃O₄@CoFe-LDH11/PMs system achieved 95% AO7 degradation in 7.5 min at room temperature (298 K). By comparison, while at 308 K and 318 K, the time required to achieve 95% AO7 removal were only 5 min and 3 min, respectively. Since PMS activation is an endothermic reaction, the kₐₚ of the reaction system would increase with the temperature. Furthermore, the Arrhenius formula (eqn (2)) was employed to further calculate and analyze the activation energy (Eₐ) of AO7 degradation.

\[ \ln k = \ln A - \frac{E_\text{a}}{RT} \]  

(2)

where k is the pseudo-first-order kinetics reaction rate constant, A is the preexponential factor, R and T was the ideal gas constant and absolute temperature, respectively. Eₐ is the activation energy. The Eₐ value of Fe₃O₄@CoFe-LDH11/PMs system calculated from Fig. S6b† was 31.99 kJ mol⁻¹. Hence, the degradation of AO7 in this system could be facilitated by raising the temperature.

3.3.2. Catalyst concentration and PMS dosage. The effects of catalyst concentration on the degradation rate of AO7 in Fe₃O₄@CoFe-LDH11/PMs system was investigated and the data are provided in Fig. 6c. The removal efficiency of AO7 within 3 min, as well as the reaction rate constant, both rose with the catalyst concentration. In detail, the degradation rate of AO7 after 5 min increased from 68.3% to 82.2%, 93.5% at catalyst concentration from 10 mg L⁻¹ to 50 mg L⁻¹, respectively. While the catalyst concentration was above 50 mg L⁻¹, the 5 min degradation rate of AO7 was ranged from 91.6% to 93.7%. Hence, a slight enhancement was noticed in AO7 removal efficiency. Increasing the catalyst dosage at lower catalyst concentrations (less than 50 mg L⁻¹) was equivalent to enhancing the number of catalytic active sites in the system. However, the catalyst concentration did not significantly affect the degradation rate of AO7 when the catalyst exceeded the upper limit of the range above 50 mg L⁻¹. In addition, the removal rate of AO7 after 15 min at 100 mg L⁻¹ catalyst concentration was only 91.5%, a value slightly lower than those obtained at 50 mg L⁻¹ and 75 mg L⁻¹ catalyst concentrations, since the excess of catalyst might compete with AO7 and consume SO₅²⁻ radicals generated during PMS activation. As a result, the AO7 degradation rate decreased, consistent with the literature. Therefore, 50 mg L⁻¹ was determined as the optimal catalyst concentration.

The effect of different PMS dosages on the removal rates of AO7 are presented in Fig. 6d. As PMS dosage increased from 0.06 to 0.12 mM, the degradation efficiency of AO7 increased significantly from 70.5% to 95.1% after 15 min. Besides, a linear growth relationship was recorded between the reaction rate constant k and PMS dosage (Fig. S6d†), indicated that the PMS concentration is the rate-controlling parameter of the catalytic degradation of AO7 under such conditions. Moreover, further increase PMS concentration led to no significant change in AO7 removal efficiency. This could be attributed to the excessive HSO₅⁻ which would react with the radicals to produce SO₅²⁻ with a weaker oxidizing ability (eqn (3) and (4)). Overall, considering the cost of PMS and the relationship between PMS dosage and degradation rate of AO7, 0.12 mM was selected as optimal PMS dosage for subsequent experiments.

\[
\text{HSO}_5^- + \text{SO}_4^{2-} \rightarrow \text{SO}_5^{2-} + \text{H}^+ 
\]  

(3)

\[
\text{HSO}_5^- + \text{HO}^- \rightarrow \text{SO}_5^{2-} + \text{H}^+ 
\]  

(4)

3.3.3. Effect of inorganic ions. Various inorganic ions exist in actual wastewater bodies, which may affect the degradation process of target contaminants. Hence, the influence of four inorganic anions (Cl⁻, HCO₃⁻, H₂PO₄⁻ and NO₃⁻) on the removal of AO7 was investigated and the results are displayed in Fig. 7. NO₃⁻ showed almost no obvious effect on AO7 degradation at different concentrations (Fig. 7d and S7,†). Compared to NO₃⁻, Cl⁻ exhibited an inhibitory effect on the catalytic system. After adding 10 mM Cl⁻, the AO7 removal rate reduced from 95.1% to 77.5% within 15 min. On the one hand, it is generally believed that Cl⁻ could directly react with PMS to generate HOCl in the presence of PMS and Cl⁻ in the system (eqn (4)). On the other hand, the generated SO₄²⁻ and HO⁻ by PMS activation could also deliver free electrons to Cl⁻ to form radicals of Cl²⁻ and ClO²⁻ (eqn (5) and (6)), k(SO₅²⁻ + Cl⁻) = 2.0 ×10⁶ M⁻¹ s⁻¹, k(HO⁻ + Cl⁻) = 4.3 ×10⁸ M⁻¹ s⁻¹. The redox potentials of newly generated free radicals Cl²⁻ and ClO²⁻ were estimated to 2.4 V and 1.5–1.8 V, respectively. As a result, the relative redox capacity of new generated radicals was the reason that led to a decrease in AO7 removal rate.

\[
\text{Cl}^- + \text{HSO}_5^- \rightarrow \text{SO}_4^{2-} + \text{HOCI} 
\]  

(5)

\[
\text{Cl}^- + \text{SO}_4^{2-} \rightarrow \text{SO}_2^{2-} + \text{Cl}^+ 
\]  

(6)

\[
\text{Cl}^- + \text{HO}^- \rightarrow \text{ClO}_2^- 
\]  

(7)

As shown in Fig. 7b and S7b,† the presence of HCO₃⁻ significantly inhibited the decomposition of AO7. The addition of 10 mM HCO₃⁻ led to only 57.7% removal of AO7. Note that HCO₃⁻ is a common free radical scavenger, quenching the generated SO₄²⁻ and HO⁻ in the catalytic/PMs system according to eqn (8) and (9), to form the less reactive HCO₃⁻. Furthermore, the introduction of HCO₃⁻ would cause the pH of the system rising. Simultaneously, HCO₃⁻ has strong buffer capability, which might keep the catalytic system under alkaline conditions. With addition of 1–10 mM H₂PO₄⁻, the decreasing trend of AO7 removal efficiency was similar to the system with Cl⁻ added. This result can be explained that H₂PO₄⁻ could transform into phosphate radicals with the assistance of SO₄²⁻ and HO⁻. 

\[
\text{HCO}_3^- + \text{SO}_4^{2-} \rightarrow \text{SO}_3^{2-} + \text{HCO}_3^- 
\]  

(8)

\[
\text{HCO}_3^- + \text{HO}^- \rightarrow \text{CO}_3^{2-} + \text{H}_2\text{O} 
\]  

(9)
3.4. Stability and reusability of Fe₃O₄@CoFe-LDH

Reusability and stability of catalysts are important when evaluating potential practical applications. Hence, five consecutive cycle experiments were conducted to measure the stability of Fe₃O₄@CoFe-LDH. As displayed in Fig. 8a, Fe₃O₄@CoFe-LDH exhibited outstanding reusability and stability after five cycles, with 98.1%, 94.9%, 94.4%, 94.3% and 88.7% AO7 degradation rate for each cycle. However, a slight reduction in AO7 removal rate occurred in the fifth catalytic degradation experiment, which might be explained by several reasons. (i) Loss of Fe₃O₄@CoFe-LDH active site during the catalytic degradation of AO7. (ii) The intermediate products of AO7 degradation adsorbed on Fe₃O₄@CoFe-LDH surface, that hindered the contact between the catalyst and PMS in the next run. In general, the stability and recyclability shown by Fe₃O₄@CoFe-LDH in the cycling experiments indicated that the magnetic samples could be used as an ideal heterogeneous catalyst in actual wastewater treatment.

3.5. Reaction mechanism

To verify the free radicals generated in Fe₃O₄@CoFe-LDH/PMS system, quenching tests were carried out by introducing two common free radical scavengers EtOH and TBA into the degradation system. The reaction rate constants of EtOH with sulfate radical and hydroxyl radical were determined as $3.51 \times 10^7$ M⁻¹ s⁻¹ and $9.1 \times 10^6$ M⁻¹ s⁻¹ respectively. EtOH is considered to capture both SO₄⁺ and HO⁺ at the same time. Meanwhile, the reaction rate constants of TBA/SO₄⁺ and TBA/HO⁺ were $4.0-9.1 \times 10^5$ M⁻¹ s⁻¹ and $3.8-7.6 \times 10^8$ M⁻¹ s⁻¹, thereby generally used as an effective scavenger for HO⁺. As illustrated in Fig. 8b, the addition of 10 mM and 100 mM TBA into the system led a mild decrease AO7 degradation efficiency within 10 min. By comparison, AO7 removal rate after 15 min showed almost no obvious change when compared to the control experiment, indicating that Fe₃O₄@CoFe-LDH/PMS system could generated HO⁺. As for EtOH, the introduction of 10 mM and 100 mM EtOH into AO7 solution, caused a remarkable declined in the degradation rate with only 77.4% and 35.9% AO7 removed within 15 min. In sum, Fe₃O₄@CoFe-LDH/PMS system could generate HO⁺ and SO₄⁺ during the catalytic degradation of AO7, while SO₄⁺ was determined as the dominant species decomposing the AO7 molecules.

To further interpret the reaction mechanism of Fe₃O₄@CoFe-LDH/PMS system, XPS was employed to analyze the valence states of Fe₃O₄@CoFe-LDH before and after the degradation reaction. In Fig. 9a, the Co 2p XPS spectrum of fresh Fe₃O₄@CoFe-LDH was divided into Co 2p₁/₂ and Co 2p₃/₂ peaks located at binding energies of 780.5 eV and 796.8 eV respectively. Two satellite peaks at 786.2 eV and 802.6 eV were

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**Fig. 8** Reusability of Fe₃O₄@CoFe-LDH for the degradation of AO7 (a), effect of TBA and EtOH on AO7 degradation (b). Experiment conditions: pH = 6, T = 298 K, V = 0.1 L, [AO7] = 40 mg L⁻¹, [catalyst] = 50 mg L⁻¹, [PMS] = 0.12 mM.

**Fig. 9** XPS spectra of (a) Co 2p and (b) Fe 2p of fresh and used Fe₃O₄@CoFe-LDH.
The mechanism of PMS activation was proposed for AO7 degradation sites in the outer shell structure of LDHs. From 70.2% before the reaction to 65.2% after reaction. Thus, a redox process of Co(II)/Co(III) took place on the samples surface during the catalytic reaction. Fe 2p XPS spectra of fresh and used samples are displayed in Fig. 8b. The peak of Fe 2p3/2 was divided into two peaks at 713.0 eV (Fe(II) 2p3/2) and 710.5 eV (Fe(II) 2p3/2), whereas the peak at 724.2 eV was related to Fe 2p1/2 of Fe(II). After catalytic reaction, no obvious change in the binding energy of Fe 2p was noticed. Additionally, the amount of Fe(II) increased from 60.7% to 69.3% after PMS activation, and the peaks related to Fe(II) were derived from ferrous on the surface of Fe3O4 in Fe3O4@CoFe-LDH. Therefore, the core structure of Fe3O4 would transfer free electrons to Co(II) active sites in the outer shell structure of LDHs.

Based on the above analysis and discussion, the reaction mechanism of PMS activation was proposed for AO7 degradation by Fe3O4@CoFe-LDH. Firstly, PMS was directly activated by Co(n) active sites on Fe3O4@CoFe-LDH surface to generate SO4(2-) (eqn (10)). Simultaneously, partial SO4(2-) reacted with H2O or OH- of the solution to form the HO• (eqn (11)). And then, these free radicals originated from the HO• would be degraded within 15 min by Fe3O4@CoFe-LDH/PMS system. The as-prepared magnetic nanocomposite exhibited excellent tolerance to pH (4-9) and relative activation energy (39.9 kJ mol⁻¹). The degradation efficiency of AO7 in Fe3O4@CoFe-LDH/PMS system was more or less affected by inorganic anions and the inhibiting order of anions was HCO3⁻ > Cl⁻ > H2PO4⁻ > NO3⁻ . The magnetic property of Fe3O4@CoFe-LDH could be conveniently recovered from an aqueous solution by external magnetic field for reuse. After five cycles, Fe3O4@CoFe-LDH maintained 88.7% AO7 degradation rate. Furthermore, quenching tests indicated that SO4(2-) was the dominant radical in Fe3O4@CoFe-LDH/PMS. The enhanced catalytic performances were mainly attributed to the synergistic effect between core structure of Fe3O4 and active sites of CoFe-LDH, resulting rapid transformation of Co(n)/Co(n). In sum, this work provides a new discovery of the potential of magnetic LDH for PMS activation in wastewater treatment.

**Conflicts of interest**

The authors declare that they have no conflict of interest.

**Acknowledgements**

This work was supported by the National Natural Science Foundation of China (Grant No. 41907270).

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