



 Cite this: *RSC Adv.*, 2023, 13, 33403

# Facile fabrication of $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ magnetic composites and their application in $\text{Cu}(\text{II})$ ion removal

 Jiachen Zhu, Ping Li, Bowen Yang, Shengjie Lan,\* Weiyuan Chen and Donghai Zhu \*

In this study, we fabricated magnetic  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites through the seed deposition technique to achieve  $\text{Cu}(\text{II})$  ion removal from aqueous solutions. As indicated by the characterization results, three-dimensional flower-like spheres composed of external  $\text{Mg}(\text{OH})_2$  were formed, with nano- $\text{Fe}_3\text{O}_4$  particles uniformly embedded in the “flower petals” of the spheres. The efficacy of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 in  $\text{Cu}(\text{II})$  ion removal was examined through batch experiments. The impact of solution pH on removal efficiency was examined, and the pseudo-second-order model and the Langmuir model provided good fits to the adsorption kinetics and isotherm data, respectively. Remarkably,  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 exhibited a significant removal capacity of  $1051.65 \text{ mg g}^{-1}$  for  $\text{Cu}(\text{II})$  ions. Additionally, the composite displayed a notable saturation magnetization value of  $17.3 \text{ emu g}^{-1}$ , facilitating isolation from sample solutions through external magnetic fields after  $\text{Cu}(\text{II})$  ion absorption. At the solid–liquid interface, a mechanism involving ion exchange between  $\text{Mg}(\text{II})$  and  $\text{Cu}(\text{II})$  cations was realized as the mode of  $\text{Cu}(\text{II})$  ion removal. The composites’ effective adsorption properties and rapid magnetic separation highlighted their suitability for use in treating copper-contaminated water.

 Received 1st September 2023  
 Accepted 8th November 2023

DOI: 10.1039/d3ra05961h

[rsc.li/rsc-advances](https://rsc.li/rsc-advances)

## 1. Introduction

Heavy metal ions (HMIs) are extensively used in industrial applications, such as pigments, batteries, and mineral processing, owing to their unique physical and chemical properties. However, rapid industrialization has led to increasingly severe water pollution caused by HMIs, which are both cumulative and non-biodegradable.<sup>1</sup> When these HMIs accumulate to a certain level, they pose significant risks to both human health and the ecological environment.<sup>2</sup> Hence, it is essential to treat wastewater by removing HMIs. Numerous methods have been employed for this purpose, such as ion exchange,<sup>3</sup> adsorption,<sup>4</sup> chemical precipitation,<sup>5</sup> and membrane separation.<sup>6</sup> Due to its high adsorption effectiveness and simplicity of operation, adsorption is an excellent option for eliminating low-concentration HMIs from aqueous solutions.

A survey of the literature revealed that many adsorption materials have been employed to remove HMIs; these include activated calcium carbonate,<sup>7</sup> activated carbon,<sup>8</sup> alumina,<sup>9</sup> graphene,<sup>10</sup> and carbon nanotubes.<sup>11</sup> Nonetheless, these materials are unsuitable for large-scale industrial wastewater treatment because of their low adsorption capacities for HMIs or high costs. Magnesium hydroxide ( $\text{Mg}(\text{OH})_2$ ), and especially its nano-

form, has recently garnered considerable attention because of its high adsorption capability, low cost, and low adverse impact on the environment.<sup>12,13</sup> For example, Zhu *et al.* demonstrated the efficient adsorption of  $\text{Pb}(\text{II})$  and  $\text{Cd}(\text{II})$  ions from wastewater utilizing coral-like hierarchical porous  $\text{Mg}(\text{OH})_2$ .<sup>14</sup> Jiang *et al.* reported a 92.6% removal rate of  $\text{Ni}(\text{II})$  ions from wastewater using flower-like globular  $\text{Mg}(\text{OH})_2$ .<sup>15</sup> However, obtaining  $\text{Mg}(\text{OH})_2$  with a specific morphological structure and a sizable specific surface area often necessitates the use of structure-directing molecules or crystal modifiers, thereby increasing costs.<sup>16</sup> Furthermore, the small size of  $\text{Mg}(\text{OH})_2$  particles makes them susceptible to loss during operations. Additionally, the challenge of separating  $\text{Mg}(\text{OH})_2$  particles from aqueous solutions after adsorbing target ions hinders its practical application and wider adoption.<sup>17</sup>

Composite materials consisting of a multifunctional shell and a magnetic core offer the combined advantages of the shell’s functionality and the core’s magnetic separation capabilities.<sup>18</sup> Numerous recent studies have explored this concept.<sup>19,20</sup> Huang *et al.* fabricated  $\text{Fe}_3\text{O}_4@\text{Cu}/\text{C}$  and  $\text{Fe}_3\text{O}_4@\text{CuO}$  composites through a self-sacrificing template method for catalytic dye removal using  $\text{H}_2\text{O}_2$  as an oxidant.<sup>21</sup> Liu *et al.* synthesized an 18-crown-6 ether-modified  $\text{Fe}_3\text{O}_4@\text{R}-\text{COOH}$  adsorbent through an amidation reaction for  $\text{Cs}^+$  removal.<sup>22</sup> Xiong *et al.* fabricated multifunctional  $\text{Ag}_3\text{PO}_4/\text{Fe}_3\text{O}_4/\text{diatomite}$  composites through a hydrothermal approach.<sup>23</sup> These resulting composites not only exhibit

State Key Laboratory of Plateau Ecology and Agriculture, School of Chemical Engineering, Qinghai University, Xining 810016, PR China. E-mail: lanshengjie@126.com; zhudonghai-2001@163.com



improved multifunctional activity involving catalysis, adsorption, and sterilization but can also be separated from aqueous solutions using an external magnet. Therefore, incorporating magnetic particles into  $\text{Mg}(\text{OH})_2$  could facilitate the separation and recovery of such composite adsorbents from aqueous solutions.

In the current research,  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  magnetic composites were synthesized through a straightforward and efficient method without the use of structure-directing molecules or crystal modifiers. Thereafter, these composites were used in aqueous solutions to eliminate  $\text{Cu}(\text{II})$  ions. The structure and morphology of the magnetic composites were characterized. The outer layer, with a flower-like globular morphology of  $\text{Mg}(\text{OH})_2$ , exhibited a remarkable ability to adsorb heavy metal ions. Moreover, embedding  $\text{Fe}_3\text{O}_4$  magnetic particles facilitates the rapid separation of the composites from aqueous solutions after target ions have been adsorbed under the action of an external magnetic field. Additionally, the impacts of the solution pH values on the  $\text{Cu}(\text{II})$  ion isolation were evaluated. Furthermore, the mechanism of  $\text{Cu}(\text{II})$  adsorption on  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  magnetic composites was determined by analyzing the adsorption isotherms, adsorption kinetics, and the characteristics of the adsorbed products.

## 2. Experimental

### 2.1 Materials

Every reagent that was used in this investigation, including magnesium sulfate heptahydrate ( $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$ ), nitric acid ( $\text{HNO}_3$ ), sodium hydroxide ( $\text{NaOH}$ ), nano- $\text{Fe}_3\text{O}_4$  ( $\geq 99.5\%$ , 20 nm), copper nitrate trihydrate ( $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$ ), ammonia solution ( $\text{NH}_3 \cdot \text{H}_2\text{O}$ : 25%), ethanol ( $\text{CH}_3\text{CH}_2\text{OH}$ ), were of analytical grade. The ammonia solution and nitric acid were procured from Shuangshuang Chemical Co., Ltd (Yantai, China), and the Sinopharm Chemical Reagent Co., Ltd (Shanghai, China) supplied other reagents.

### 2.2 Synthesis of $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ composites

A seed deposition technique was applied to synthesize composites of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ . First, 24.65 g of  $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$  was dissolved in 50 mL of deionized water, followed by dispersing a certain amount of nano- $\text{Fe}_3\text{O}_4$  particles into the magnesium salt solution through ultrasonic processing. Next, the suspension was heated and maintained at a constant 60 °C. Thereafter, the suspension received a drop-by-drop addition of 150 mL ammonia while being agitated at 850 rpm. After the drop-wise addition, the reaction solution was left undisturbed to sit for 2 h. The resulting precipitate was separated by centrifugation and rinsed with deionized water. Final  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites were obtained after freeze-drying for 8 h. According to the additive amounts of nano- $\text{Fe}_3\text{O}_4$  (0.10 g, 0.20 g and 0.30 g), the synthesized composites were labeled as  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -1,  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -2 and  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3, respectively. For comparison, pure  $\text{Mg}(\text{OH})_2$  without adding nano- $\text{Fe}_3\text{O}_4$  particles was also synthesized under the same conditions.

### 2.3 Characterization

The composite specimens were characterized employing a D5000 X-ray diffractometer (XRD, Karlsruhe, Germany). Employing a JSM 6700 field emission scanning electron microscope (FESEM, JEOL, Japan) and an HT-7700 transmission electron microscope (TEM, Hitachi, Japan), the morphology and dimensions of the specimens were analyzed. The specific SA and porosity structure parameters of the specimens were analyzed through the Brunauer–Emmett–Teller (BET) method utilizing an ASAP 2020 analyzer (Micromeritics Instrument Corp., USA). The magnetic characteristics of the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites were assessed employing a PPMS-9T vibrating sample magnetometer (VSM, Quantum Design, USA).

### 2.4 Adsorption experiments

The  $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$  was used to prepare solutions with varying concentrations of  $\text{Cu}(\text{II})$ . For the kinetic experiments, 12 mg of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites were individually introduced into 160 mL of 25.4  $\text{mg L}^{-1}$   $\text{Cu}(\text{II})$  solution and shaken using an orbital shaker. At specific intervals, a polyether sulfone membrane with a pore size of 0.45 micrometers was used to filter aliquots of the samples. The atomic absorption spectrometer model ZEEnit 700 P (Analytik Jena, Germany) was employed to determine the concentration of  $\text{Cu}(\text{II})$  that was still present in the solution. The adsorption isotherms were analyzed through the same procedure but with varying concentrations of  $\text{Cu}(\text{II})$  solutions. We conducted all experiments in duplicate. The eqn (1) and (2) were applied to determine the removal efficiency ( $E\%$ ) of  $\text{Cu}(\text{II})$  ions and the adsorption capacity ( $q_t$ ) of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ , respectively.

$$E = \frac{(C_0 - C_t)}{C_0} \times 100\% \quad (1)$$

$$q_t = \frac{(C_0 - C_t)V}{m} \quad (2)$$

where  $C_0$  ( $\text{mg L}^{-1}$ ) and  $C_t$  ( $\text{mg L}^{-1}$ ) denote the initial  $\text{Cu}(\text{II})$  ions concentration and concentration at time  $t$ , respectively. In addition, the volume of the  $\text{Cu}(\text{II})$  ions solution is shown by  $V$  (L), and  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composite weight is indicated by  $m$  (g).

## 3. Results and discussion

### 3.1 Magnetic property and remove ability

Through VSM measurements at room temperature,  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites were evaluated for their magnetic properties. Fig. 1a shows all of the synthesized composites exhibited S-like curves in their magnetic hysteresis loops, characterized with low magnetic remanence and coercivity, indicating that these composites exhibited superparamagnetic behavior. The saturation magnetization ( $M_s$ ) values of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -1,  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -2 and  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 were 7.4, 11.6 and 17.3  $\text{emu g}^{-1}$ , respectively.  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites have lower  $M_s$  values. This may result from the embedding of  $\text{Fe}_3\text{O}_4$

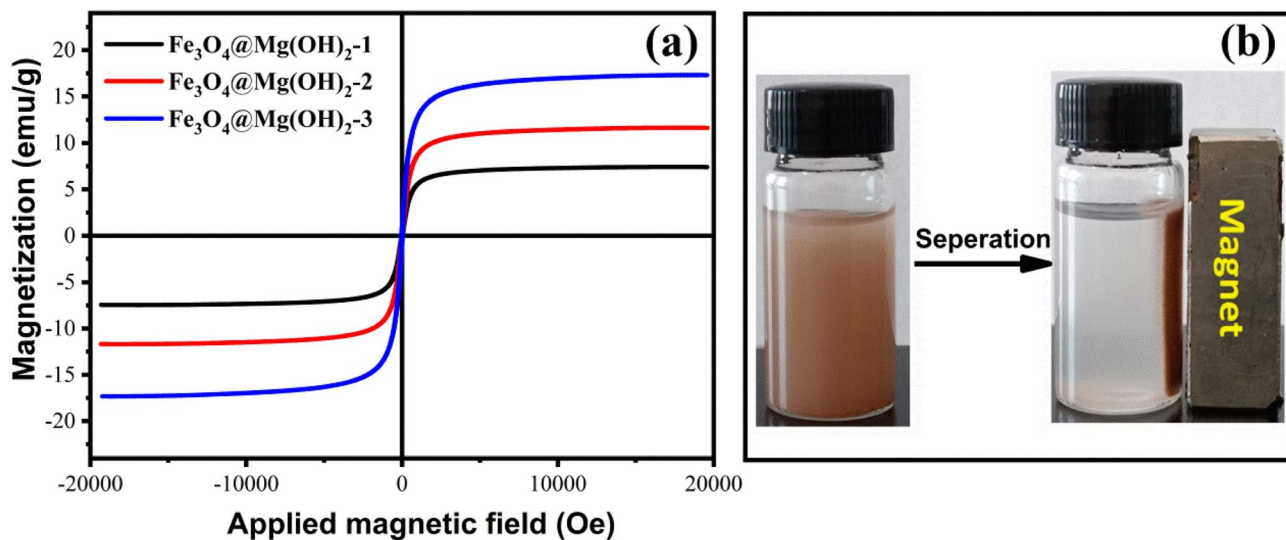


Fig. 1 (a) Magnetic hysteresis loops of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites obtained at room temperature, (b) magnetic separation process of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composite.

in non-magnetic  $\text{Mg}(\text{OH})_2$  nanosheets and the low levels of  $\text{Fe}_3\text{O}_4$  in the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composite.<sup>24</sup> Some studies have found that an  $M_s$  value of  $16.3 \text{ emu g}^{-1}$  is sufficient for magnetic separation with a conventional magnet.<sup>25,26</sup> According to this standard, only the  $M_s$  value of achieved with  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 meet the requirements of magnetic separation. Moreover, uniformly dispersed  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 quickly coalesced under the influence of an external magnetic field, as shown in Fig. 1b. This further confirm that the  $M_s$  value for this configuration of the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composite is sufficient for efficient magnetic separation.

To investigate the remove ability of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites, comparative experiments with an initial concentration of  $25.4 \text{ mg L}^{-1}$  at pH 5 were carried out. The removal capacities of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -1,  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -2 and  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 for  $\text{Cu}(\text{II})$  were 335.86, 329.52 and  $322.14 \text{ mg g}^{-1}$ , respectively. Under the same conditions, the removal capacity of pure  $\text{Mg}(\text{OH})_2$  was  $357.15 \text{ mg g}^{-1}$ . Apparently, with the increase of  $\text{Fe}_3\text{O}_4$  loading amount, the removal capacity of the composites decreases gradually. However, the amount of  $\text{Fe}_3\text{O}_4$  added is too small to meet the requirements of magnetic separation. Considering comprehensively,  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 was selected for subsequent experiments because of its magnetic properties to meet the magnetic separation, while it still has a high removal capacity for  $\text{Cu}(\text{II})$ .

### 3.2 Characterization of $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ composite

As depicted in Fig. 2, XRD was employed to characterize the composition and phase of the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composite. The diffraction peaks at approximately  $30.32^\circ$ ,  $35.65^\circ$ ,  $43.24^\circ$ ,  $57.43^\circ$ , and  $63.06^\circ$  could be attributed to the diffraction patterns of the  $\text{Fe}_3\text{O}_4$  crystal with a face-centered cubic spinel structure (JCPDS #88-0866).<sup>27</sup> Meanwhile, the residual diffraction peaks corresponding to the as-prepared composites, at approximately  $18.53^\circ$ ,  $32.88^\circ$ ,  $37.94^\circ$ ,  $50.84^\circ$ , and  $58.68^\circ$ ,

corresponded to the (001), (100), (101), (102), and (110) planes, respectively, of hexagonal  $\text{Mg}(\text{OH})_2$  (JCPDS #44-1482),<sup>12</sup> confirming the successful generation of the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites.

The  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composite' morphology was examined through SEM and TEM analyses. As demonstrated in Fig. 3a, the resulting composites exhibited a consistent flower-like spherical structure with a diameter of 1–2  $\mu\text{m}$ . The enlarged SEM image (Fig. 3b) showed that each flower sphere was composed of interwoven nanosheets with a multi-layered porous structure. Such a porous architecture is potentially conducive to enhanced adsorption phenomena. Notably,  $\text{Fe}_3\text{O}_4$  nanoparticles were not observed in the enlarged SEM image. Nevertheless, a signal corresponding to Fe was visible in the EDS spectrum (Fig. 3c),

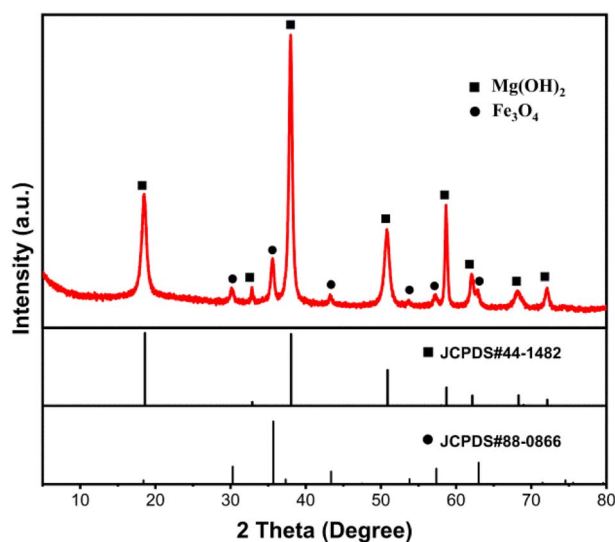


Fig. 2 XRD pattern of the as-prepared  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composite.



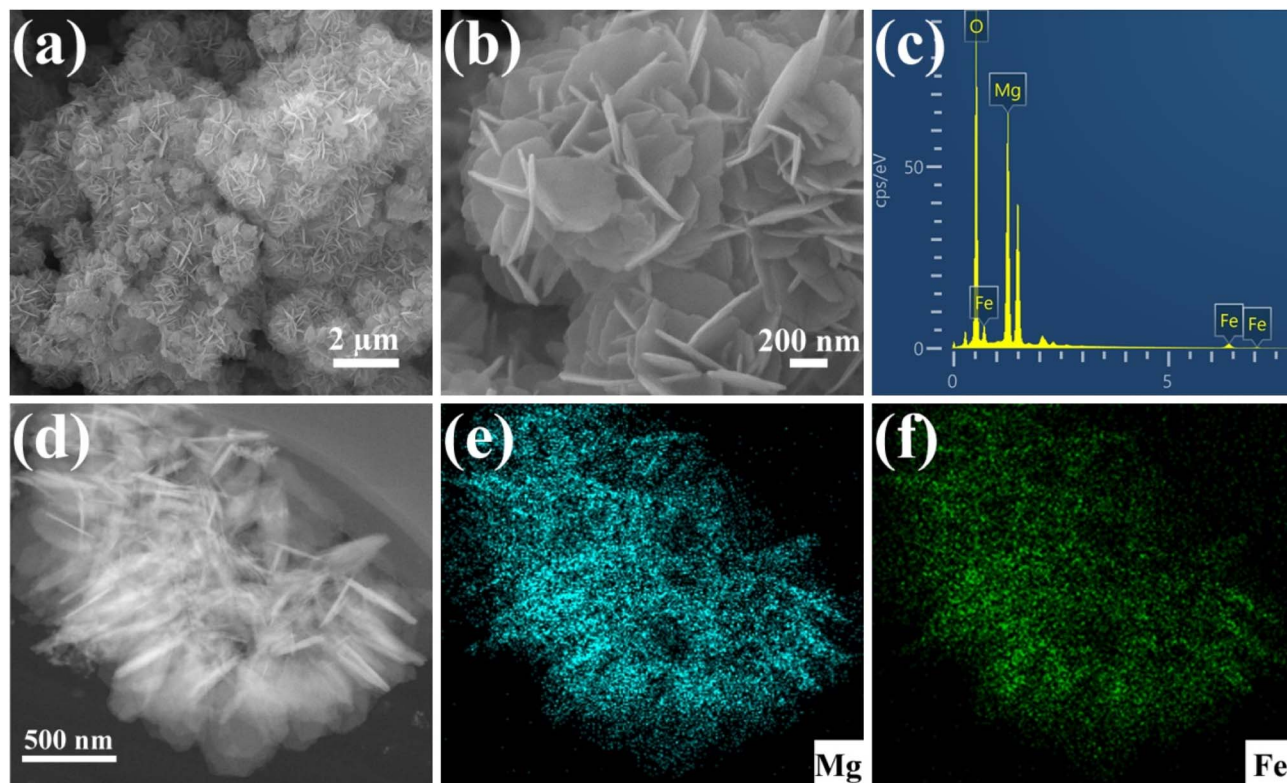


Fig. 3 (a and b) SEM images, (c) EDS spectra, (d) TEM image, and (e and f) the corresponding EDS mapping of the as-prepared  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2\text{-3}$  composite.

suggesting that  $\text{Fe}_3\text{O}_4$  nanoparticles were embedded and enveloped within  $\text{Mg}(\text{OH})_2$  nanosheets. Further insights were gained from the TEM-EDS mapping of the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2\text{-3}$  composite, as depicted in Fig. 3e and f. This mapping revealed a co-distribution of Mg and Fe elements within the same spatial region, demonstrating the even dispersion of  $\text{Fe}_3\text{O}_4$  nanoparticles within the  $\text{Mg}(\text{OH})_2$  nanosheets.

The porous characteristics of the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2\text{-3}$  composite was evaluated *via* the nitrogen adsorption-desorption

approach. According to the IUPAC classification, the nitrogen absorption-desorption curve (Fig. 4a) featured H3 hysteresis loops and type-IV isotherms.<sup>14</sup> The non-coincidence of the two branches of the hysteresis ring at relative pressures ( $P/P_0$ ) between 0.5 and 1 is a primary characteristic of mesoporous materials. This finding confirmed the presence of mesoporous structures within the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2\text{-3}$  composite. These pore structure characteristics aligned with the micro-morphology observed through SEM. The  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2\text{-3}$  composite's

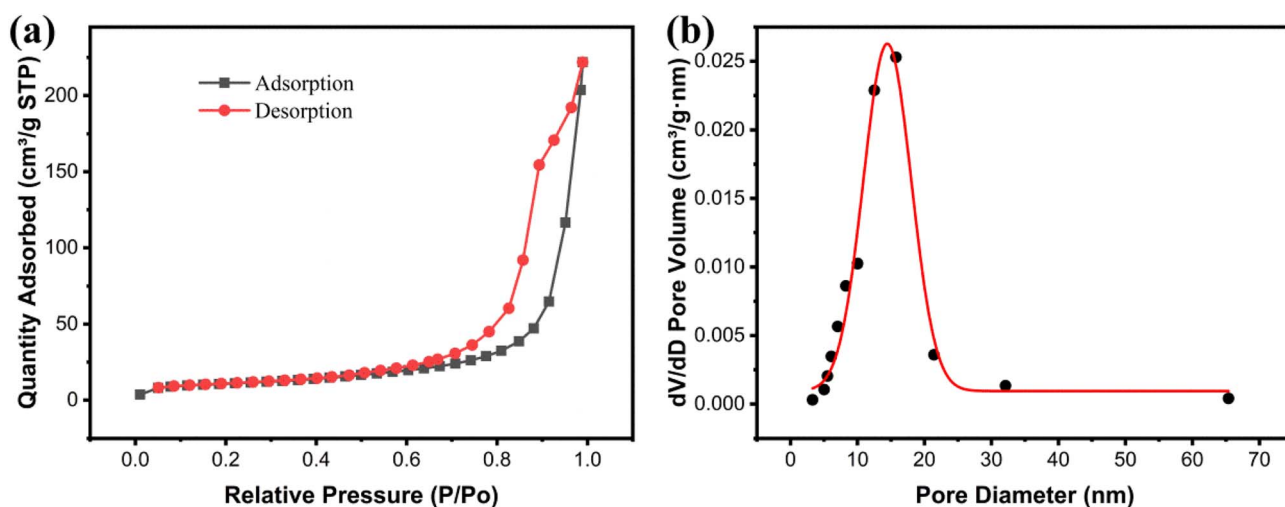


Fig. 4 (a) Nitrogen adsorption-desorption isotherms and (b) pore size distributions of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2\text{-3}$ .

surface area calculated using the BET model was  $39.78 \text{ m}^2 \text{ g}^{-1}$ . The corresponding pore size distribution curve derived from the nitrogen desorption data (Fig. 4b) confirmed the mesoporous nature of the composites consisting of pores averaging  $16.10 \text{ nm}$  in size, predominantly distributed within the  $5\text{--}25 \text{ nm}$  range and  $0.35 \text{ cm}^3 \text{ g}^{-1}$  pore volume. Notably, the substantial specific surface area and mesoporous structure can be attributed to the spherical structure resembling a flower of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 comprising an assembly of  $\text{Mg}(\text{OH})_2$  nanosheets. This unique structure provides a multitude of active adsorption sites and facilitates the rapid diffusion of ions, thereby enhancing the adsorption capacity.

### 3.3 Adsorption studies

**3.3.1 Effect of pH values.** The removal efficiency of  $\text{Cu}(\text{II})$  by  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 was influenced by the initial pH of the solution. Fig. 5 displays the results of this study, which were in an initial pH range of  $2.0\text{--}6.0$ . Notably,  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 had a low efficiency in removing  $\text{Cu}^{2+}$  at a pH value of  $2.0$ . This may occur because of the abundance of  $\text{H}^+$  ions when the pH values of the solution are low. These  $\text{H}^+$  ions readily neutralize the outer layer of  $\text{Mg}(\text{OH})_2$ , which otherwise possesses excellent adsorption capabilities. With the increase in pH value, the concentration of  $\text{H}^+$  ions decreased rapidly, leading to an increase in the removal efficiency of  $\text{Cu}^{2+}$  by  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3. When the pH of the solution was within the range of  $4.0$  to  $6.0$ ,  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 exhibited a high removal efficiency for  $\text{Cu}^{2+}$ . At a pH higher than  $6.0$ , an elevated  $\text{OH}^-$  ion concentration resulted in the precipitation of  $\text{Cu}(\text{II})$  ions or their transition into  $\text{Cu}(\text{OH})^+$  ions. This alteration notably influenced the adsorption process, causing the influence of pH to stabilize after reaching  $6.0$ . Consequently, the pH value of the solution was adjusted to  $5.0$  for subsequent experiments.

**3.3.2 Adsorption kinetics.** A series of adsorption kinetics tests were conducted to evaluate the adsorption rate of  $\text{Cu}(\text{II})$  on  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composites. The evolution of the amount of

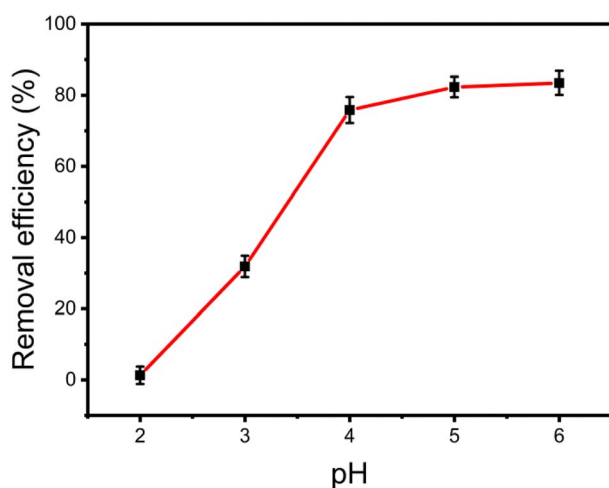


Fig. 5 Effect of solution pH values on the  $\text{Cu}(\text{II})$  adsorption.

$\text{Cu}(\text{II})$  adsorbed on the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composites over time is depicted in Fig. 6a. Notably,  $\text{Cu}(\text{II})$  was rapidly adsorbed on the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites within the first  $60 \text{ min}$  since many adsorption sites were available during this phase. The adsorption rate reduced gradually as time passed throughout the process of adsorption, leading to equilibrium within  $120 \text{ min}$  because of the progressive occupation of available adsorbent sites on the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composites by  $\text{Cu}(\text{II})$  ions during the latter stages.

Further investigation into the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composite's adsorption behavior for  $\text{Cu}(\text{II})$  was done by fitting experimental data to the pseudo-first-order kinetic model, pseudo-second-order kinetic model, and the in-particle diffusion model<sup>12</sup> as demonstrated below:

$$\ln(q_e - q_t) = \ln q_e - k_1 t \quad (3)$$

$$\frac{t}{q_t} = \frac{1}{k_2 q_e^2} + \frac{t}{q_e} \quad (4)$$

$$q_t = k_{id} t^{0.5} + C_{id} \quad (5)$$

where  $q_t$  ( $\text{mg g}^{-1}$ ) and  $q_e$  ( $\text{mg g}^{-1}$ ) denote the adsorption capacities of the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composites at time  $t$  (min) and at equilibrium,  $k_1$  ( $\text{min}^{-1}$ ) and  $k_2$  ( $\text{g mg}^{-1} \text{ min}^{-1}$ ) denote the equilibrium rate constants in the pseudo-first-order and pseudo-second-order kinetic models, respectively.  $k_{id}$  denotes the adsorption rate constant ( $\text{mg g}^{-1} \text{ min}^{-0.5}$ ), and  $C_{id}$  is the constant of the in-particle diffusion model. The fitting results are shown in Fig. 6.

Table 1 displays the correlation coefficients ( $R^2$ ) that were calculated using the fitted kinetic data. The outcomes presented in Fig. 6 and Table 1 suggested that the  $R^2$  value linked to the pseudo-first-order kinetic model was less compared to the value associated with the pseudo-second-order kinetic model. Furthermore, the computed adsorption capacity ( $q_{e,\text{cal}}$ ) obtained through the pseudo-first-order kinetic model was substantially different from the experimental value ( $q_{e,\text{exp}}$ ), indicating that the adsorption of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 to adsorb  $\text{Cu}(\text{II})$  ions was not consistent with that of the pseudo-first-order kinetic model. In comparison, the pseudo-second-order kinetic model achieved an  $R^2$  of  $0.9960$  when fitting the data, and the pseudo-second-order kinetic modeling yielded a  $q_{e,\text{cal}}$  of  $350.88 \text{ mg g}^{-1}$ , which was close to the  $q_{e,\text{exp}}$  of  $322.14 \text{ mg g}^{-1}$ . These results indicated the occurrence of chemisorption during the process of adsorption. Additionally, according to Fig. 6d, the adsorption process was analyzed using the in-particle diffusion model. The fitted curve does not intersect at the origin suggesting that speed is limited by more than just particle diffusion.

**3.3.3 Adsorption isotherms.** The maximum adsorption capacity, which can be determined based on the adsorption isotherm, is a key parameter for evaluating the adsorbent's performance and exploring the adsorption mechanism. Consequently, the experimental data of  $\text{Cu}(\text{II})$  adsorption on the composite were fitted utilizing the Langmuir (eqn (6)) and Freundlich isotherm (eqn (7)) models.<sup>24</sup>

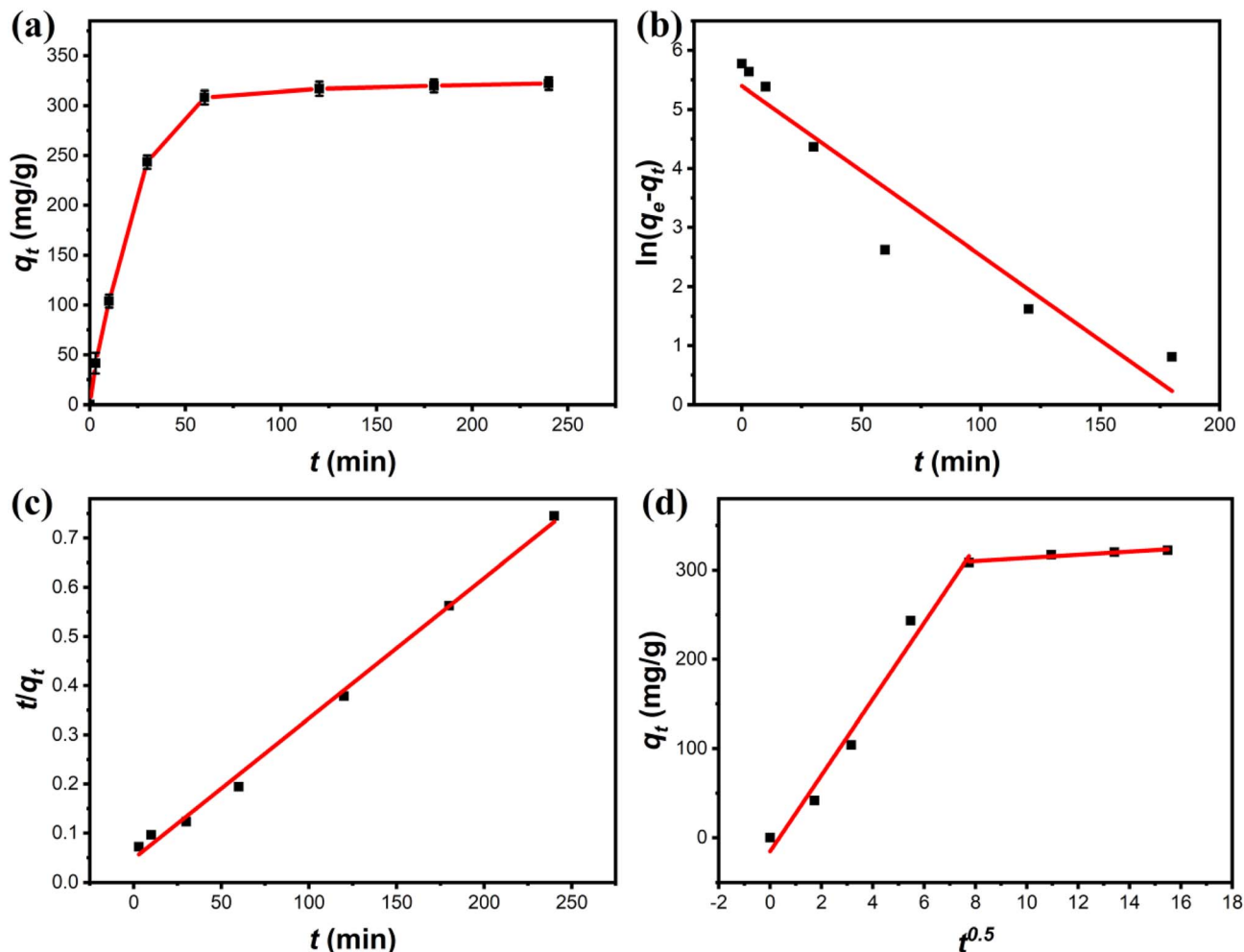


Fig. 6 (a) Adsorption kinetic of Cu(II) onto  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2\text{-3}$  and fitting using the (b) pseudo-first-order, (c) pseudo-second-order, and (d) intra-particle diffusion models.

Table 1 Parameters of the kinetic models for Cu(II) adsorption on  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2\text{-3}$

Models	Parameters	Values
Pseudo-first-order kinetic model	$q_{e,\text{cal}}$ ( $\text{mg g}^{-1}$ )	220.99
	$k_1$ ( $\text{min}^{-1}$ )	0.0287
	$R^2$	0.9242
Pseudo-second-order kinetic model	$q_{e,\text{cal}}$ ( $\text{mg g}^{-1}$ )	350.88
	$k_2$ ( $\text{g mg}^{-1} \text{min}^{-1}$ )	$1.68 \times 10^{-4}$
	$R^2$	0.9960
Intra-particle diffusion model	$k_{\text{id1}}$ ( $\text{g mg}^{-1} \text{min}^{-0.5}$ )	1.7582
	$R_1^2$	0.9436
	$k_{\text{id2}}$ ( $\text{g mg}^{-1} \text{min}^{-0.5}$ )	42.7244
	$R_2^2$	0.9793
Experiment	$q_{e,\text{exp}}$ ( $\text{mg g}^{-1}$ )	322.14

$$\frac{c_e}{q_e} = \frac{c_e}{q_m} + \frac{1}{q_m k_L} \quad (6)$$

$$\ln q_e = \ln k_F + \frac{1}{n} \ln c_e \quad (7)$$

where the Cu(II) equilibrium concentration is represented by  $c_e$  ( $\text{mg L}^{-1}$ ),  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2\text{-3}$  composite maximum adsorption capacity is illustrated by  $q_m$  ( $\text{mg g}^{-1}$ ), and  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2\text{-3}$  composite's equilibrium adsorption capacity is illustrated by  $q_e$  ( $\text{mg g}^{-1}$ ). In addition, the Langmuir constant is represented by  $k_L$  ( $\text{L mg}^{-1}$ ), and the Freundlich constant is indicated by  $k_F$  ( $\text{mg g}^{-1}$ ). The Freundlich exponent  $n$  indicates the adsorption strength.

Fig. 7 and Table 2 present the fitting outcomes. Based on the obtained correlation coefficient ( $R^2$ ), In contrast to the Freundlich model, the Langmuir model exhibited better alignment with the adsorption of Cu(II) by  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites. The results indicated that  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites adsorption of Cu(II) predominantly involved monolayer adsorption onto homogeneously active centers.  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites had a maximum adsorption capacity for Cu(II) of  $1115.97 \text{ mg g}^{-1}$ , as predicted by the Langmuir model; this value was impressively similar to the actual value ( $1051.65 \text{ mg g}^{-1}$ ). The affinity between the adsorbent and the adsorbed ion is denoted by the strength index  $n$  in the Freundlich model. Furthermore, the value of  $1/n$  in our

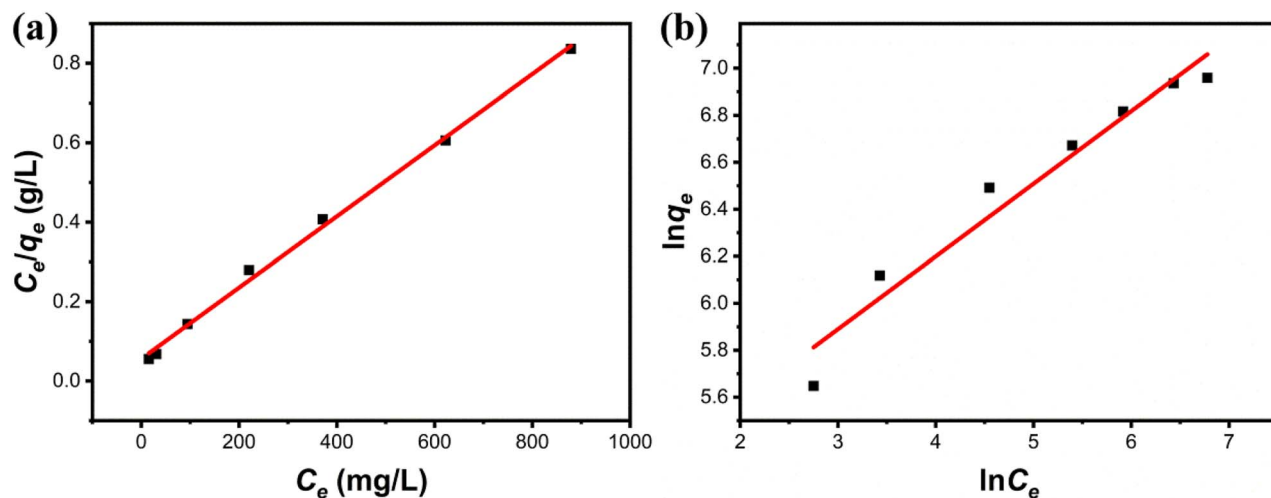


Fig. 7 Adsorption isotherms of  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composites for  $\text{Cu}(\text{II})$  fitted using the (a) Langmuir model and (b) Freundlich model.

Table 2 Isotherm model parameters for adsorption of  $\text{Cu}(\text{II})$  by  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites

Models	Parameters	Value
Langmuir	$q_m$ ( $\text{mg g}^{-1}$ )	1115.97
	$k_L$ ( $\text{L mg}^{-1}$ )	0.016
	$R^2$	0.9969
Freundlich	$k_F$ ( $\text{mg g}^{-1}$ )	142.81
	$n$	3.2321
	$R^2$	0.9543

experiment was 0.309, indicating that  $\text{Cu}(\text{II})$  exhibited a high propensity for adsorption onto the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composite. In addition, comparisons were made between the adsorption capabilities of a variety of typical adsorbents for  $\text{Cu}(\text{II})$  that were reported in the research literature (Table 3). Accordingly, the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composites that were produced for this investigation proved to be excellent adsorbents for removing  $\text{Cu}(\text{II})$  since they were able to facilitate the rapid and efficient removal of  $\text{Cu}(\text{II})$  from an aqueous solution.

**3.3.4 Effect of coexisting cations.** Industrial wastewater often contains multiple coexisting cations, resulting in competitive adsorption of binding sites on the adsorbent surface.<sup>35</sup> In this work, the effect of different coexisting cations including  $\text{Na}^+$ ,  $\text{K}^+$ ,  $\text{Co}^{2+}$ ,  $\text{Ni}^{2+}$  and  $\text{Zn}^{2+}$  with initial concentration of  $25.4 \text{ mg L}^{-1}$

Table 3 Different adsorbents and their maximum  $\text{Cu}(\text{II})$  adsorption capacities

Adsorbent	Time (min)	$q_{\text{max}}$ ( $\text{mg g}^{-1}$ )	References
$\text{Ni}@\text{Mg}(\text{OH})_2$	50	40.18	28
$\text{Mg-Al-D2EHPA}$	60	68.66	29
$\text{MgAl-LDH}@\text{RHB}$	1440	104.34	30
$\text{Mg}(\text{OH})_2/\text{GO}$	1440	216.0	31
$\text{MCs}@\text{Mg/Fe-LDHs}$	1440	338.98	32
$\text{CaAl-LDH}$	600	122.7	33
Phosphate	1440	50.84	34
$\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$	120	1051.65	This work

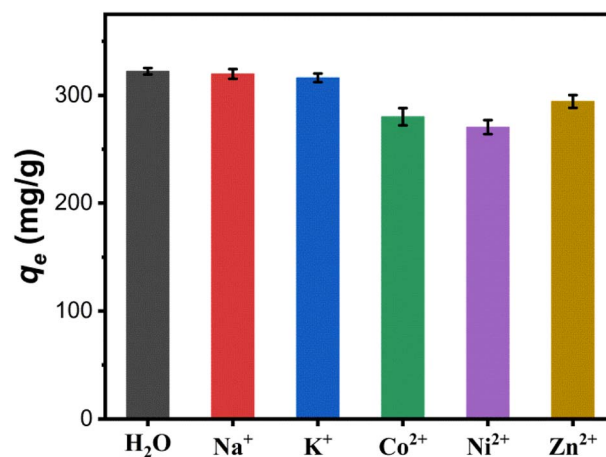


Fig. 8 Effect of coexisting cations on the adsorption capacity of the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composite for  $\text{Cu}(\text{II})$ .

on the adsorption capacity of the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$ -3 composite for  $\text{Cu}(\text{II})$  was investigated. As shown in Fig. 8, its adsorption capacity for  $\text{Cu}(\text{II})$  decreased from  $322.14 \text{ mg g}^{-1}$  in aqueous solution without these coexisting cations to 319.66, 316.08, 280.12, 270.53 and  $294.4 \text{ mg g}^{-1}$  in the presence of  $\text{Na}^+$ ,  $\text{K}^+$ ,  $\text{Co}^{2+}$ ,  $\text{Ni}^{2+}$  and  $\text{Zn}^{2+}$ , respectively. Through comparative analysis, it can be found that the coexistence cations of  $\text{Na}^+$  and  $\text{K}^+$  have little effect on the adsorption capacity of the composite. While, the coexistence cations of  $\text{Co}^{2+}$ ,  $\text{Ni}^{2+}$  and  $\text{Zn}^{2+}$  have a certain inhibitory effect on the remove of  $\text{Cu}(\text{II})$ . This can be attributed to the fact that the coexisting ions of  $\text{Co}^{2+}$ ,  $\text{Ni}^{2+}$  and  $\text{Zn}^{2+}$  can also combine with the hydroxyl on the surface of  $\text{Mg}(\text{OH})_2$  to form precipitation and deposit *in situ*,<sup>14</sup> which hinders the ion exchange between  $\text{Cu}^{2+}$  and  $\text{Mg}^{2+}$ .

### 3.4 Mechanism

The mechanism of  $\text{Cu}(\text{II})$  removal by the  $\text{Fe}_3\text{O}_4@\text{Mg}(\text{OH})_2$  composite was evaluated by analyzing the adsorption products



at different time intervals. Fig. 9 shows the SEM image and the corresponding element mapping of the composites after Cu(II) adsorption for 10 min. Fig. 9a displays an abundance of small spherical particles on the petal-like Mg(OH)<sub>2</sub> nanosheets, indicating the formation of novel substances post-adsorption. Furthermore, the EDS element mapping analysis revealed a uniform dispersion of Cu within the adsorbed product, indicating that the composites furnish a multitude of active centers conducive to the binding of Cu(II) ions. Within this short span of 10 min allotted for adsorption, strong Cu signals were detected for the adsorption products, indicating that the as-prepared composites adsorbed Cu(II) at a relatively high rate. Notably, no signal corresponding to N was detected for the original composites; however, a prominent N signal was observed after adsorption, indicating the formation of N-containing compounds in the adsorption products.<sup>36</sup> Notably, in the adsorption process, the sole source of the N element was the nitrate used during the preparation of the Cu(II) solution.

To further confirm the composition of products following Cu(II) adsorption, the composites' products were characterized using XRD after a 120 min adsorption duration, as depicted in Fig. 10. Similar to the original composites, the distinct diffraction peaks of Fe<sub>3</sub>O<sub>4</sub> (JCPDS #88-0866) were also identified in the spectrum of the adsorption product. This confirmed the magnetic nature of the adsorption product, facilitating its separation from aqueous solutions through magnetic means. Concurrently, other residual diffraction peaks corresponding to the monoclinic Cu<sub>2</sub>(NO<sub>3</sub>)(OH)<sub>3</sub> (JCPDS #75-1779)<sup>37</sup> were observed, indicating the

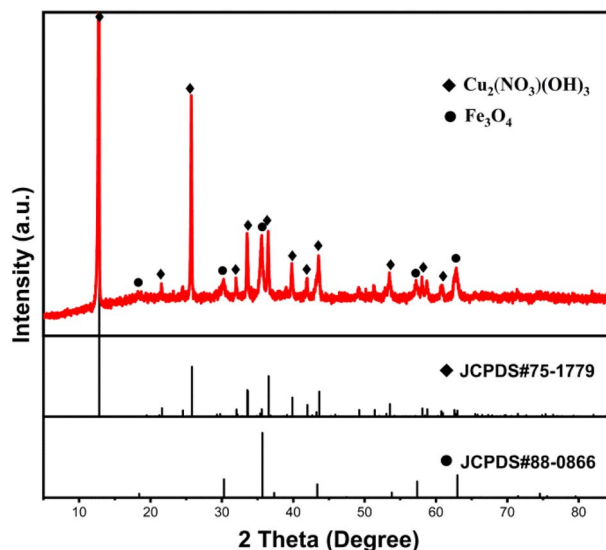


Fig. 10 XRD pattern of the Fe<sub>3</sub>O<sub>4</sub>@Mg(OH)<sub>2</sub> after adsorbing Cu(II) for 120 min.

involvement of NO<sub>3</sub><sup>-</sup> ions from the solution in the adsorption reaction. Moreover, the diffraction peaks corresponding to the magnesium-containing substance were hardly detected, suggesting that most of the Mg(II) ions were exchanged. To confirm this, the concentrations of Mg(II) ions that are present in the solution at different adsorption time intervals were measured. After

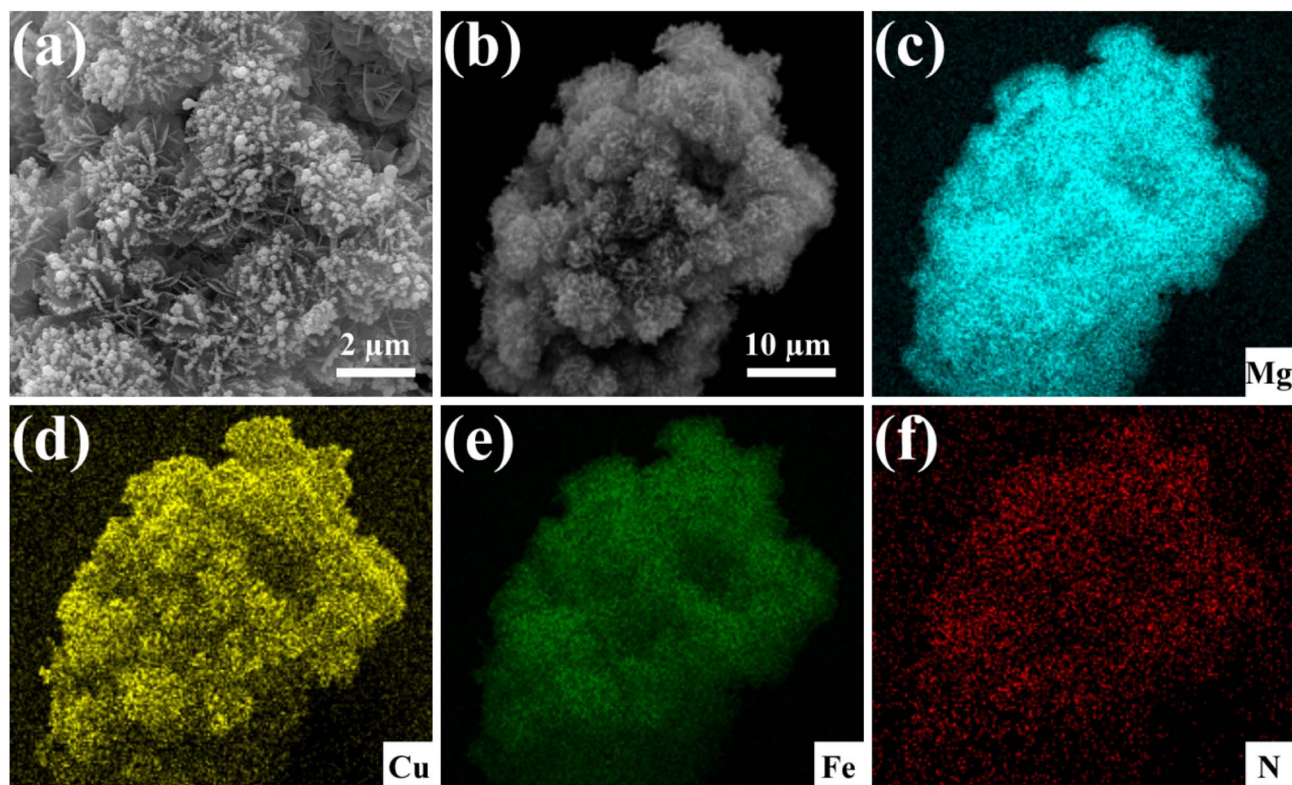


Fig. 9 SEM image of the Fe<sub>3</sub>O<sub>4</sub>@Mg(OH)<sub>2</sub> composites after adsorbing Cu(II) for 10 min (a), along with the corresponding elemental mapping of a selected area (b–f).



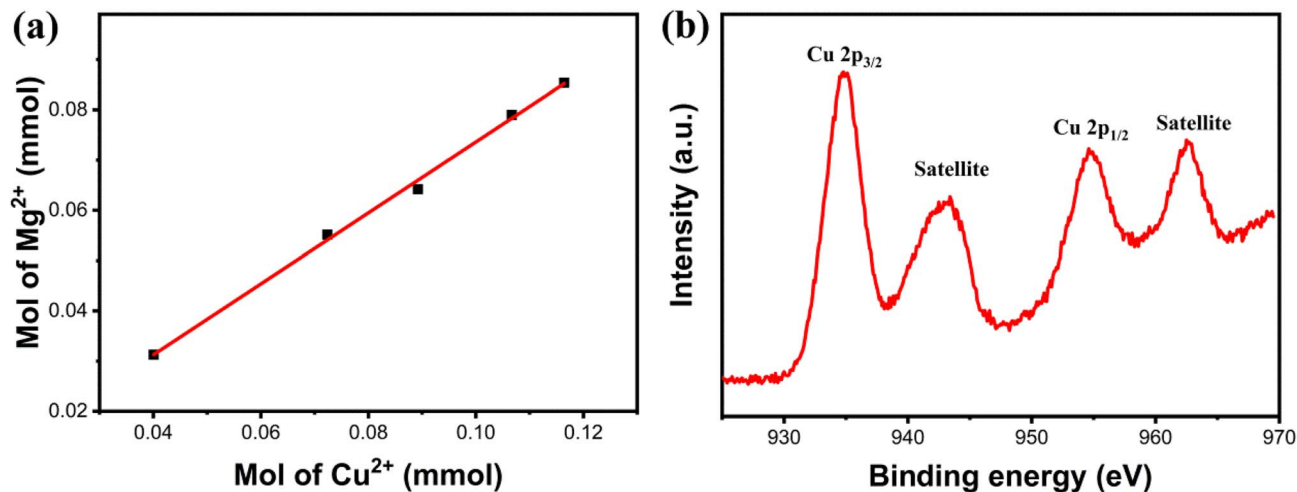
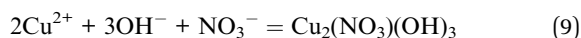


Fig. 11 (a) Quantitative measurements of the molar amount of Mg(II) released and the molar amount of Cu(II) removed, (b) high resolution XPS spectra of Cu 2p after adsorption.

adsorption, the findings showed that Mg(II) ions appeared in the solution, and the concentration of Mg(II) ions increased with the extension of the adsorption time. Interestingly, an almost linear relationship was observed between the molar release of Mg(II) ions and the number of adsorbed Cu(II) ions (as shown in Fig. 11a). The valence state of the Cu compound on the surface of the Fe<sub>3</sub>O<sub>4</sub>@Mg(OH)<sub>2</sub> composite was determined by high resolution XPS. Fig. 11b shows the binding energy of Cu 2p orbital to be 934.8 eV (Cu 2p<sub>3/2</sub>) and 954.7 eV (Cu 2p<sub>1/2</sub>). Furthermore, two satellite bands (938–948 eV and 958–966 eV) were also shown in the adsorption product. These confirmed that Cu exists in divalent state after adsorption<sup>38</sup> and there is no reduction coupled adsorption on the surface of Fe<sub>3</sub>O<sub>4</sub>@Mg(OH)<sub>2</sub> composite.

The above investigation led to the following cation exchange reaction summary of the adsorption process:



First, an ionization equilibrium was observed between Mg(OH)<sub>2</sub> and Mg(II) alongside OH<sup>−</sup> ions, and the concentration of OH<sup>−</sup> ions surrounding the Mg(OH)<sub>2</sub> nanosheets was greater relative to that in the solution.<sup>39</sup> When Cu(II) ions moved close to the Mg(OH)<sub>2</sub> nanosheets, they bonded with OH<sup>−</sup> ions and NO<sub>3</sub><sup>−</sup> ions from the solution, resulting in Cu<sub>2</sub>(NO<sub>3</sub>)(OH)<sub>3</sub> deposition on the nanosheet surface *in situ*. As OH<sup>−</sup> ions were consumed, the ionization equilibrium of Mg(OH)<sub>2</sub> shifted toward the right, increasing the influx of Mg(II) ions into the solution.

## 4. Conclusions

This study applied the seed deposition approach to synthesize a novel magnetic Fe<sub>3</sub>O<sub>4</sub>@Mg(OH)<sub>2</sub> composite, which was used for removing Cu(II) ions from aqueous solutions. Analysis of the phase and morphology revealed that nano-Fe<sub>3</sub>O<sub>4</sub> particles were uniformly embedded within the “flower petals” of globular

Mg(OH)<sub>2</sub>. The composite demonstrated the advantages of flower globular Mg(OH)<sub>2</sub> and magnetic nano-Fe<sub>3</sub>O<sub>4</sub> particles, providing benefits such as effective adsorption toward heavy metal ions and rapid magnetic separation from sample solutions. The Fe<sub>3</sub>O<sub>4</sub>@Mg(OH)<sub>2</sub> composite exhibited a high adsorption capacity of 1115.97 mg g<sup>−1</sup> toward Cu(II) ions, as revealed by calculations using the Langmuir equation. The adsorption isotherm data were consistent with the Langmuir model, and the kinetic data were consistent with the pseudo-second-order model. Furthermore, the removal mechanism of Cu(II) ions was analyzed in this study. The results illustrated that Cu(II) ions precipitated in the form of Cu<sub>2</sub>(NO<sub>3</sub>)(OH)<sub>3</sub> particles on the surface of the flowered globular Fe<sub>3</sub>O<sub>4</sub>@Mg(OH)<sub>2</sub> magnetic composite during the adsorption process.

## Author contributions

Jiachen Zhu: investigation, writing – original draft. Ping Li: data curation. Bowen Yang: investigation. Shengjie Lan: methodology, supervision and writing – review & editing. Weiyan Chen: investigation. Donghai Zhu: conceptualization and writing – review & editing.

## Conflicts of interest

There are no conflicts of interest to declare.

## Acknowledgements

This work received financial support from the Natural Science Foundation of Qinghai Province (No. 2020-ZJ-934Q).

## References

- 1 S. Arooj, S. Naz, A. Qaisar, S. Satti, A. Razzaq, A. F. Alrefaei, M. F. Albeshr, M. Durrani, S. F. Dai and R. U. Khan, *J. Appl. Anim. Res.*, 2023, **51**, 530–539.

- 2 W. Yang, W. Cai, J. Zhou, C. Dang, X. Peng, Y. Chen, X. Wei, S. Bo, S. Liang and Z. Luo, *J. Environ. Chem. Eng.*, 2021, **9**, 106634.
- 3 L. Ulloa, M. Martinez-Mincherro, E. Bringas, A. Cobo and M. F. San-Roman, *Sep. Purif. Technol.*, 2020, **253**, 117516.
- 4 Y. Fei and Y. H. Hu, *Chemosphere*, 2023, **335**, 139077.
- 5 M. H. El Awady, M. A. Ahmed and A. Dahaba, *Egypt. J. Chem.*, 2020, **63**, 1697–1712.
- 6 X. X. Cheng, Y. R. Zhang, S. L. Shao, C. X. Lai, D. J. Wu, J. T. Xu, X. S. Luo, D. L. Xu, H. Liang and X. W. Zhu, *Desalination*, 2023, **548**, 116266.
- 7 X. W. Song, Y. W. Cao, X. Z. Bu and X. P. Luo, *Appl. Surf. Sci.*, 2021, **536**, 147958.
- 8 X. A. Bai, L. Y. Xing, N. Liu, N. A. Ma, K. X. Huang, D. P. Wu, M. M. Yin and K. Jiang, *Nanomaterials*, 2021, **11**, 3255.
- 9 Y. J. O. Asencios and M. R. Sun-Kou, *Appl. Surf. Sci.*, 2012, **258**, 10002–10011.
- 10 M. Sivakumar, J. Widakdo, W. S. Hung, C. F. Wang, C. C. Hu, K. R. Lee and J. Y. Lai, *Sep. Purif. Technol.*, 2022, **288**, 120485.
- 11 J. F. Jin, J. S. Sun, K. H. Lv, X. B. Huang, J. T. Wang, J. P. Liu, Y. R. Bai, X. Guo, J. W. Zhao, J. Liu and Q. L. Hou, *J. Mol. Liq.*, 2021, **334**, 116087.
- 12 H. Amrulloh, Y. S. Kurniawan, C. Ichsan, J. Jelita, W. Simanjuntak, R. T. M. Situmeang and P. A. Krisbiantoro, *Colloids Surf., A*, 2021, **631**, 127687.
- 13 P. P. Wang, Y. X. Ye, D. W. Liang, H. M. Sun, J. Liu, Z. F. Tian and C. H. Liang, *RSC Adv.*, 2016, **6**, 26977–26983.
- 14 H. Zhu, L. Li, W. D. Chen, Y. J. Tong and X. S. Wang, *J. Hazard. Mater.*, 2021, **416**, 125922.
- 15 D. M. Jiang, Y. H. Yang, C. T. Huang, M. Y. Huang, J. J. Chen, T. D. Rao and X. Y. Ran, *J. Hazard. Mater.*, 2019, **373**, 131–140.
- 16 X. Y. Shen, Y. X. Huang, H. M. Shao, Y. Wang, Q. Han, J. S. Chen, B. C. Li and Y. C. Zhai, *Trans. Nonferrous Met. Soc. China*, 2022, **32**, 3149–3162.
- 17 Q. Sun, G. C. Zhu, J. Wu, J. Lu and Z. H. Zhang, *Environ. Technol.*, 2021, **42**, 894–904.
- 18 S. Melhi, *Environ. Pollut. Bioavailability*, 2023, **35**, 2163922.
- 19 A. Iqbal, M. R. Jan, J. S. Shah and M. N. Sarwar, *J. Chem. Technol. Biotechnol.*, 2023, **98**, 967–977.
- 20 E. S. Behbahani, K. Dashtian and M. Ghaedi, *J. Hazard. Mater.*, 2021, **410**, 124560.
- 21 Y. F. Huang, L. Zhang, L. Ma, Y. Li and C. L. Zhong, *Catal. Lett.*, 2020, **150**, 815–825.
- 22 Z. Liu, Y. Q. Zhou, M. Guo, B. L. Lv, Z. J. Wu and W. Z. Zhou, *J. Hazard. Mater.*, 2019, **371**, 712–720.
- 23 C. Y. Xiong, Q. F. Ren, S. H. Chen, X. Y. Liu, Z. Jin and Y. Ding, *Mater. Today Commun.*, 2021, **28**, 102695.
- 24 B. Peng, T. T. Song, T. Wang, L. Y. Chai, W. C. Yang, X. R. Li, C. F. Li and H. Y. Wang, *Chem. Eng. J.*, 2016, **299**, 15–22.
- 25 Z. Y. Ma, Y. P. Guan and H. Z. Liu, *J. Polym. Sci., Part A: Polym. Chem.*, 2005, **43**, 3433–3439.
- 26 X. L. Zhao, J. M. Wang, F. C. Wu, T. Wang, Y. Q. Cai, Y. L. Shi and G. B. Jiang, *J. Hazard. Mater.*, 2010, **173**, 102–109.
- 27 C. P. Chen, P. Gunawan and R. Xu, *J. Mater. Chem.*, 2011, **21**, 1218–1225.
- 28 M. Zhang, W. Q. Song, Q. L. Chen, B. J. Miao and W. C. He, *ACS Appl. Mater. Interfaces*, 2015, **7**, 1533–1540.
- 29 N. Rouahna, D. Barkat, A. Ouakouak and E. Srasra, *J. Environ. Chem. Eng.*, 2018, **6**, 1226–1232.
- 30 A. Y. Li, Y. Zhang, W. Z. Ge, Y. T. Zhang, L. H. Liu and G. H. Qiu, *Bioresour. Technol.*, 2022, **347**, 126425.
- 31 P. P. Wang, Y. X. Ye, D. W. Liang, H. M. Sun, J. Liu, Z. F. Tian and C. H. Liang, *RSC Adv.*, 2016, **6**, 26977–26983.
- 32 Y. Y. Xie, X. Z. Yuan, Z. B. Wu, G. M. Zeng, L. B. Jiang, X. Peng and H. Li, *J. Colloid Interface Sci.*, 2019, **536**, 440–455.
- 33 W. Yao, J. Wang, P. Y. Wang, X. X. Wang, S. J. Yu, Y. D. Zou, J. Hou, T. Hayat, A. Alsaedi and X. K. Wang, *Environ. Pollut.*, 2017, **229**, 827–836.
- 34 A. Ivanets, N. Kitikova, I. Shashkova, Y. Matrunchik, L. Kul'bitskaya and M. Sillanpaa, *Environ. Technol. Innovation*, 2016, **6**, 152–164.
- 35 U. Upadhyay, I. Sreedhar, S. A. Singh, C. M. Patel and K. L. Anitha, *Carbohydr. Polym.*, 2021, **251**, 117000.
- 36 P. S. Das, S. Bakuli, A. Samanta, A. K. Mandal, J. Ghosh, A. Deyand and A. K. Mukhopadhyay, *Mater. Res. Express*, 2017, **4**, 025025.
- 37 T. T. Dai and Y. Q. Mao, *Chem. Phys. Lett.*, 2020, **739**, 137018.
- 38 M. C. di Gregorio, P. Ranjan, L. Houben, L. J. W. Shimon, K. Rechav, M. Lahav and M. E. van der Boom, *J. Am. Chem. Soc.*, 2018, **140**, 9132–9139.
- 39 C. R. Li, Z. Y. Zhuang, F. Huang, Z. C. Wu, Y. P. Hong and Z. Lin, *ACS Appl. Mater. Interfaces*, 2013, **5**, 9719–9725.