Cite this: *RSC Adv.*, 2021, 11, 23679

Preparation, characterization and application of CS@PDA@Fe₃O₄ nanocomposite as a new magnetic nano-adsorber for the removal of metals and dyes in wastewater†

Xingjing Zhang, Baohe Li, Xiaoqian Han * and Nong Wang

This study aimed to develop a novel magnetic chitosan/dopamine/Fe₃O₄ nano-adsorber (CS@PDA@Fe₃O₄) for the removal of heavy metal ions and organic dye molecules from aqueous solution. CS@PDA@Fe₃O₄ was prepared by surface modification of PDA/Fe₃O₄ nanoparticles with chitosan using IPTES as the cross-linker. The surface structure, composition, and properties of the CS@PDA@Fe₃O₄ nano-adsorber were characterized by elemental (EDS), spectroscopic (XRD, XPS, and FT-IR), magnetic intensity (VSM), surface and morphological (TEM and SEM) analyses. In order to study its adsorption behavior, equilibrium and kinetics studies were carried out through batch experiments. Additionally, the influences of the pH value, initial concentration, adsorbent dose, and contact time were also evaluated. The CS@PDA@Fe₃O₄ nano-adsorber exhibited high adsorption capacity especially for Cu(II), with a maximum adsorption capacity of 419.6 mg g⁻¹. The experimental data were well described by the Langmuir isotherm kinetic models.

Received 14th May 2021
Accepted 29th June 2021

DOI: 10.1039/d1ra03779j

rsc.li/rsc-advances

1. Introduction

A growing social and economic concern is water contamination due to the discharge of many dyes and heavy metal ions into the environment, which causes a potential threat to human health and the environment.^{1,2} As everyone knows, heavy metal ions and most of the dyes are very poisonous or strongly oncogenic, and may cause bad effects to people's brains, lungs, and kidneys.³ Therefore, it is particularly important to treat wastewater with a feasible and appropriate method.⁴ The traditional ways such as electrochemical precipitation,⁵ sorption,⁶ reduction,⁷ advanced oxidation,⁸ *etc.* have been used to remove pollutants from water. In all technologies, the adsorption method has the advantages of low cost, and being green and environmentally friendly for the elimination of pollutants in water.⁹ Various adsorbents have been reported for the treatment of metal ions and dye-contaminated water: activated carbon, zeolites, and clay minerals, which have disadvantages such as low adsorption abilities, narrow application scope, and difficult recovery problem.¹⁰ New materials modified by natural adsorbents for easy adsorbent recovery have received extensive attention in recent years.¹¹

Chitosan (CS), as the second most abundant natural polysaccharide after cellulose, has a large number of hydroxyl and amino functional groups in its chain moiety and exerts outstanding ability to interact with a wide variety of molecules by physical and chemical forces.¹² Therefore, it can be used for wastewater treatment.^{13–15} However, the solubility of chitosan in an acidic environment limits its further application.¹⁶ Due to using glutaraldehyde as a crosslinking agent in the traditional method, the amino content of chitosan decreased sharply, which affected the adsorption property. Therefore, we introduce dopamine as a new cross-linking agent for surface modification or modification.^{17,18}

Dopamine (DA), a protein-based biomaterial that is found in the tentacles of mussel sea creatures, has abundant functional groups such as catechol groups, amine groups, and aromatic moieties. Hence, it can easily adsorb heavy metal ions and toxic organic pollutants through chelation and π – π stacking interactions.^{19,20} Furthermore, it has some advantages of easy processing, in a one-pot preparation method and chemical versatility, which proved to be a powerful tool for the removal of environmental water pollutants.^{21,22}

In this study, a new, low-cost, nano-adsorber with the increased number of adsorption sites was developed. This new biopolymer adsorbent is a chitosan derivative, CS@PDA@Fe₃O₄ which was synthesized by assembling biomimetic polymer (PDA) and chitosan onto magnetic nanoparticles based on esterification reaction. The structure and properties of nano-adsorber were investigated by means of FTIR, SEM, TEM,

School of Chemical and Chemical Engineering, Lanzhou Jiaotong University, Lanzhou 730070, P. R. China. E-mail: hxqnv2011nv@163.com

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d1ra03779j

XRD, VSM, and elemental analysis. The various studies in terms of isotherms and kinetics of adsorption of heavy metal ions and organic dye molecules were carried out. The effects of solution pH, dosage, initial concentration, and contact time on the adsorption process were optimized.

2. Experimental

2.1. Instruments and materials

$\text{FeCl}_2 \cdot 4\text{H}_2\text{O}$, $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$, sodium hydroxide (NaOH) and dopamine (DA) were purchased from Kaixin Chemical Industry. Trometamol was purchased from Zhongqin Chemical Reagent (Shanghai, China). Chitosan and γ -isocyanatopropyltriethoxysilane (IPTES) were purchased from Odofuni Biological Technology (Nanjing, China). Hydrochloric acid (HCl) was purchased from Chemical Factory (Beijing, China). Methylene blue (MB), Fuchsin basic (FB), Rhodamine B (RhB) were cationic dyes gained from Damao Chemical Reagent Factory (Tianjin, China). All other reagents were analytical grade and were used without any extra purification. Aqueous solutions at various concentrations were prepared from HgCl_2 , were used as sources for $\text{Hg}(\text{II})$, CuCl_2 , PbCl_2 were used as sources for $\text{Cu}(\text{II})$, $\text{Pb}(\text{II})$, respectively. The morphology was analyzed by scanning electron microscopy (SEM) (MIRN3, TESCAN, China) and (TEM) (Philip Tecnai20, Netherlands). Infrared spectroscopy was recorded by a Fourier transform infrared spectrometer (FTIR) (Vector22, Bruker, Germany). Vibrating-sample magnetometry (VSM) (Lake Shore7410, America). The types and contents of the elements were analyzed by an energy dispersive spectrometer (EDS) (FE SEM-4100, Hitachi Limited, Japan).

2.2. Synthesis of magnetite Fe_3O_4 nanoparticles

Magnetite Fe_3O_4 nanoparticles were synthesized by the method of coprecipitation according to reported investigation.²³ Briefly, FeCl_3 (8.15 g) and FeCl_2 (3.98 g) were weighed and dissolved in 25 mL deionized water respectively, the obtained mixture was mixed further stirred for 30 min at 60 °C, followed by NaOH (100 mL, 2 mol L^{-1}) for another 2 h. The produced mixture was washed in deionized water until neutral and dried at 70 °C for 12 h to obtain Fe_3O_4 nanoparticles.

2.3. Preparation of $\text{PDA@Fe}_3\text{O}_4$ nanocomposite

In a round-bottom flask, a solution of 300 mL of Tris-HCl buffer solution (pH = 8.5) was configured and then a certain amount of dopamine and magnetic Fe_3O_4 nanoparticles (1 g) was added. After sonication for 30 min, the reaction temperature was rose to 100 °C for 12 h to obtain the $\text{PDA@Fe}_3\text{O}_4$ nanocomposite.

2.4. Synthesis of $\text{CS@PDA@Fe}_3\text{O}_4$ nanocomposite

Chitosan (0.5 g) was added to anhydrous pyridine solution at 90 °C for 12 h. Add 0.5 mL IPTES in constant temperature stirring for 12 h. Then, $\text{PDA@Fe}_3\text{O}_4$ (0.5 g) was added, the stirring of mixture continued for 12 h. The black $\text{CS@PDA@Fe}_3\text{O}_4$ nanocomposite was separated using the magnetic decantation, washed with distilled water and alcohol thoroughly. The product was dried at 70 °C under the vacuum for 12 h.

2.5. Adsorption experiments

The adsorption was studied by batch methods that were carried out by placing $\text{CS@PDA@Fe}_3\text{O}_4$ (0.01 g) in series of flasks containing heavy metal ions or dyes in an aqueous solution (30 mL) with the desired initial concentration and pH. Then, the flasks were shaken in a thermostat oscillator at a certain temperature with the constant rate for a given time. The effects of operating variables such as pH value, contact time, initial concentration, and amount of adsorbent on the degree of adsorption were investigated. To investigate the effect of different solution pH on adsorption of heavy metal and dye, the desired pH range of the heavy metal and dye was adjusted from 1 to 7. The working concentration of heavy metal ions or dyes was 5.4 g L^{-1} for kinetic studies and 1.35–8.1 g L^{-1} for isotherm studies. Separate the magnetic microsphere $\text{CS@PDA@Fe}_3\text{O}_4$ from the solution and measure the absorbance of the solution before and after adsorption.

The amount of pollutant adsorbed per gram of the adsorber was calculated based on the following equation.²⁴

$$\eta = \frac{C_0 - C_e}{C_0} \times 100\% \quad (1)$$

$$Q = \frac{V(C - C_e)}{M} \quad (2)$$

where η is the adsorption capacity (mg g^{-1}); C_0 and C_e are initial and equilibrium concentrations of the heavy metal ions or dyes (mg L^{-1}) in the testing solution (mg L^{-1}); V is the volume of the solution (L), and M is the weight of resin beads (g).

2.6. Recycling of $\text{CS@PDA@Fe}_3\text{O}_4$ nanocomposite

For the desorption experiment, the $\text{CS@PDA@Fe}_3\text{O}_4$ nanocomposite was separated, washed several times with absolute ethanol solution and distilled water, and used for the subsequent adsorption after drying at 70 °C under the vacuum for 12 h. In order to test the reusability of the $\text{CS@PDA@Fe}_3\text{O}_4$ nanocomposite, this adsorption–desorption cycle was repeated 6 times by using the same adsorbent.

3. Results and discussion

3.1. Characterization of $\text{CS@PDA@Fe}_3\text{O}_4$ biopolymer adsorbent

The nanoparticles of $\text{CS@PDA@Fe}_3\text{O}_4$ were synthesized in a three-step procedure. The $\text{CS@PDA@Fe}_3\text{O}_4$ was prepared using coating layer upon layer of Fe_3O_4 , PDA and CS (Fig. 1). The adsorption principle of $\text{CS@PDA@Fe}_3\text{O}_4$ biopolymer adsorbent was that its chitosan layer molecule contains a large number of amino and hydroxyl groups, which formed mesh cage molecules by hydrogen bond, chelate many heavy metal ions and dye molecules. The prepared $\text{CS@PDA@Fe}_3\text{O}_4$ nanocomposite was characterized by various techniques such as FT-IR, vibration sample magnetometer (VSM), X-ray diffraction (XRD), transmission electron microscopy (TEM), scanning electron microscope (SEM), energy dispersive X-ray spectrometer (EDS), and X-ray photoelectron spectroscopy (XPS) analyses.

Fig. 1 Synthesis of CS@PDA@Fe₃O₄ nanocomposite and heavy metal ion absorption.

3.1.1. TEM, SEM, VSM, FT-IR and XRD of CS@PDA@Fe₃O₄. The FT-IR spectra of Fe₃O₄, PDA@Fe₃O₄ and CS@PDA@Fe₃O₄ were shown in Fig. S1.† The peaks at 585 cm⁻¹ and 623 cm⁻¹ in these three spectra are attributed to the Fe₃O₄ characteristic absorption. In the spectrum of PDA@Fe₃O₄, the peaks at 1610 cm⁻¹ could be attributed to the vibrations of the aromatic ring in PDA, which is consistent with the reference reported.²² In CS@PDA@Fe₃O₄, the peaks absorption at 2914 cm⁻¹ and 3453 cm⁻¹ correspond to -OH stretching vibrations and -NH₂ stretching vibrations, respectively. Compared with the pure Fe₃O₄, the new peak at 1097 cm⁻¹ is attributed to Si-O stretching vibrations of the crosslinking agent IPTES, and all results indicate that CS@PDA@Fe₃O₄ nano-adsorber has been successfully synthesized.

The magnetic hysteresis loops of Fe₃O₄, PDA@Fe₃O₄ and CS@PDA@Fe₃O₄, at 300 K, are shown in Fig. 1b. The magnetization saturation (MS) value of the above materials was 80.5, 68.5 and 38.2 emu g⁻¹, respectively. Although the magnetic intensity decreases obviously after the coating of dopamine and chitosan layer by layer, the CS@PDA@Fe₃O₄ nano-adsorber can still be easily separated by using magnetic field, and facilitated collection, regeneration, and reutilization (shown in the insets in Fig. S2†).²⁵

The XRD patterns of Fe₃O₄, PDA@Fe₃O₄, CS@PDA@Fe₃O₄ are displayed in Fig. S3.† The six peaks of Fe₃O₄ at 2θ = 30.5°, 35.6°, 43.2°, 54°, 57.4°, and 62.9° highly match the characteristic peaks of magnetite (Fe₃O₄, PDF NO. 01-071-6336).²⁶ There is no obvious characteristic peak of dopamine in PDA@Fe₃O₄, indicating that the structure of dopamine coating Fe₃O₄ is amorphous. In the spectrum of CS@PDA@Fe₃O₄, an obvious crystal characteristic diffraction peaks at 2θ = 20.6°, which belongs to the characteristic crystal peak of chitosan, further proving the successful introduction with chitosan to the surface of Fe₃O₄ nanoparticle. Moreover, the diffraction peaks of Fe₃O₄ crystal are well consistent with the characteristic diffraction peaks of CS@PDA@Fe₃O₄. These results indicated the Fe₃O₄ was successfully introduced. The surface morphologies of the pure Fe₃O₄ and the CS@PDA@Fe₃O₄ were shown in Fig. S4.† By comparing Fig. S4d and e,† it could be clearly observed that the surfaces of the CS@PDA@Fe₃O₄ (Fig. S4e†) are smoother and the volume increased significantly than Fe₃O₄ (Fig. S4d†). They remained basically consistent, which were spherically granular

and partially barreled. It can be seen from Fig. 1c and d that the agglomeration phenomenon of CS@PDA@Fe₃O₄ (Fig. S4g†) is much weaker than Fe₃O₄ (Fig. S4f†) due to the surface modification by coated of the chitosan. Thus, it proved that Fe₃O₄ as a magnetic core was coated.

3.1.2. EDS of CS@PDA@Fe₃O₄. We also compared the proportion of N, C, H, Fe, Si in Fe₃O₄, PDA@Fe₃O₄ and CS@PDA@Fe₃O₄ performed with EDS data, the results were listed in Table S1.† The significant improvement of N and C element content about PDA@Fe₃O₄ strongly demonstrated the successful introduction of dopamine, formed a coating layer on Fe₃O₄. The Si content in CS@PDA@Fe₃O₄ was provided by linker IPTES in the synthesis process, thus it can be seen that CS@PDA@Fe₃O₄ nano-adsorber was successfully synthesized.

3.2 Adsorption mechanism of the CS@PDA@Fe₃O₄ nano-adsorber

The XPS analysis was further carried out to study the chemical elements on the surface of the nano-adsorber and the adsorption mechanism of nano-adsorber on heavy metal ions and dye molecules. Taking the adsorption of Cu(II) as an example, virgin and Cu(II)-loaded CS@PDA@Fe₃O₄ was performed. The wide scan spectra of the CS@PDA@Fe₃O₄ before and after Cu(II) adsorption showed the same peak components of C, N, O, Fe, and Si elements (Fig. S5a†). After adsorption, new peaks appear at 931 and 74 eV, which can be attributed to Cu 2p and Cu 3p, the peak of Cl 3p at 200 eV. Based on the comparison and analysis, it is reasonable to believe that the Cu(II) was loaded onto the nano-adsorber.

By comprising the N 1s spectrum of the virgin and Cu(II)-loaded CS@PDA@Fe₃O₄ in Fig. S5b,† the peak for the species of -NH₂ (396.6 eV) was increased to 396.8 eV and the peak of -NH- (397.1 eV) increased to 398.1 eV after Cu(II) adsorption.²⁷ Fig. S5c† shows the O 1s spectrum of PDA/CS/Fe₃O₄ before and after adsorption. After Cu(II) adsorption, the peak for the species of -OH (527.1 eV) was increased to 527.3 eV and the peak of C-O (397.1 eV) increased to 529.5 eV after Cu(II) adsorption. It is further demonstrated that the amino group and hydroxyl group play an important role in the adsorption process of CS@PDA@Fe₃O₄.²⁸

3.3 Effect of contact time, pH, adsorbent dose and initial concentration on properties of the CS@PDA@Fe₃O₄ nano-adsorber

The effect of contact time for the adsorption of heavy metals ions and organic dyes molecules by CS@PDA@Fe₃O₄ nano-adsorber at various intervals of time ranging from 10 to 120 min was investigated in the ambient experimental background. The results obtained were presented in Fig. 2a. The results confirmed that adsorption of metal ions and dyes has a consistent change rule: the initial stage of the rate of adsorption was faster and then gradually reduced and reached saturation.¹³ Thus, 30 min, 70 min, 80 min, 100 min, 60 min, 60 min of contact time were respectively optimized for the adsorptive removal of Cu(II), Hg(II), Pb(II), MB, FB and RhB.

The solution pH, as a critical operational variable, has a significant effect on the surface charge of the adsorption and the metal ion chemistry. For that, we investigated different processes when the pH ranged from 1 to 12 for heavy metal ions and dyes. The Cu(II) cations tend to precipitate in the form of hydroxyl salts of Cu(II) ions and FB has a large color change in alkaline solution, which has a detrimental effect on adsorption research. Therefore, the pH range of both of them was adjusted to 1–7. It can be seen from Fig. 2b that the removal rate of adsorbent on metal ions and dye molecules increases gradually with the increasing pH of the solution under acidic conditions. When the solution was neutral, the adsorption reaches saturation. The strongly acidic and alkaline conditions are not

conductive to the effect of metal ions and dyes. The strong acidic solution containing a large number of H⁺ will form a competitive relationship with metal ions for adsorption reduce the active sites, and there will be the partial dissolution of chitosan, thus reducing the adsorption performance. Besides, metal ions may be partially precipitated under basic conditions. In consequence, the optimum pH of heavy metal ions and dye molecules adsorption onto CS@PDA@Fe₃O₄ nano-adsorber was 7.

The adsorbent dosage is a key parameter for the sorption capacity of adsorbents. The dosage of CS@PDA@Fe₃O₄ nano-adsorber was varied in the range of 40–240 mg, in a fixed initial dyes and heavy metal ions concentration of 50 mg L⁻¹, the pH value of the solution was 7, at ambient experimental conditions. The effect of the adsorbent dosage on the adsorption of heavy metal ions and dye molecules were investigated, and the results were shown in Fig. 2c. With the increase of adsorption dose, the adsorption efficiency of heavy metal ions and dye molecules increased, which could be described based on the availability of active binding sites for the adsorbates on the surface of the adsorbent. When the amount of adsorbent increases to a certain value, the removal rate decreases after it remains stable. The reason could be that after the adsorption reaches equilibrium, part of the adsorbent will fall off from the adsorbent. Therefore, 100 mg of the CS@PDA@Fe₃O₄ nano-adsorber was chosen as the ideal dose for the following adsorption experiment.

The effects of different initial concentrations of dye molecules and heavy metal ions on their adsorption at optimal

Fig. 2 Effect of (a) shaking time; (b) initial pH and (c) adsorbent dosage on the adsorption with the pH value of the solution was 7 at ambient experimental conditions.

contact time and pH were investigated. The relationship between adsorption amount and initial concentration was presented in Fig. 3. The adsorption of heavy metal ions and dye molecules showed a common change rule. As the initial concentration increases, the collision probability between the magnetic microspheres and the adsorbent increases, but when the initial concentration is large enough, the adsorption sites of the nano-adsorber are filled and saturated, and the removal rate reaches equilibrium or even decreases.

3.4 Adsorption kinetics

The adsorption kinetics are the crucial factors for determining the rate of the adsorption process, which provides essential data about the mechanism and rate of removal. Table 1 shows the kinetic profiles of heavy metals ions and organic dyes adsorption operation using CS@PDA@Fe₃O₄ as adsorbents, with time increasing. In order to investigate the removal mechanism of the adsorption of Cu(II), Hg(II), Pb(II) heavy metal ions and MB, FB, RhB dye molecules on CS@PDA@Fe₃O₄ nano-adsorber, the pseudo-first-order (3), pseudo-second-order (4) and intra-particle diffusion kinetic models (5) were employed to explore the experimental data.²⁹

$$\log(Q - Q_t) = \log Q - \frac{K_1 t}{2.303} \quad (3)$$

$$\frac{t}{Q_t} = \frac{t}{Q} + \frac{1}{K_2 Q^2} \quad (4)$$

$$Q_t = K_{\text{int}} t^{0.5} + C \quad (5)$$

where K_1 , K_2 and K_{int} are pseudo-first-order rate constant (min^{-1}), pseudo-second-order rate constant ($\text{g mg}^{-1} \text{min}^{-1}$) and intraparticle diffusion rate constant ($\text{mg g}^{-1} \text{min}^{-1/2}$) of adsorption, respectively. Q and Q_t are the adsorption capacity (mg g^{-1}) at equilibrium time and at time t (min), respectively. C (mg g^{-1}) is a constant of the intra-particle diffusion model.

The non-linear form of the reaction and diffusion-based kinetic parameters and their results were summarized in Table 1. Through the comparison and analysis of R^2 values in the table, the kinetic model. With the higher R^2 values of the pseudo-second-order kinetics model, the conclusion is drawn that the pseudo-second-order kinetics model was well simulated, which signifying that chemisorption dominated the adsorption process.

3.5 Adsorption isotherms

The adsorption isotherms are the crucial factors for optimizing the application of adsorbent, which explain the interaction between adsorbate molecules and adsorbent.

For the adsorption isotherm studies, the adsorption data of heavy metal ions and dye molecules were fitted to Langmuir (6) and Freundlich (7) equations, respectively.³⁰ The results were shown in Table 2.

$$\frac{C_e}{Q} = \frac{1}{K_L Q_{\text{max}}} + \frac{C_e}{Q_{\text{max}}} \quad (6)$$

Fig. 3 Effect of initial concentration on adsorption at 30 °C.

Table 1 Parameters of kinetics model for adsorption of pollutants on CS@PDA@Fe₃O₄

Pollutants	Pseudo-first-order			Pseudo-second-order			Intraparticle diffusion	
	<i>Q</i>	<i>K</i> ₁	<i>R</i> ²	<i>Q</i>	<i>K</i> ₂	<i>R</i> ²	<i>K</i> _{int}	<i>R</i> ²
Cu ²⁺	784.15	0.145	0.8746	416.67	0.0032	0.9999	57.404	0.9255
Hg ²⁺	9.16	0.032	0.9622	42.90	0.0035	1.0000	0.2349	0.9758
Pb ²⁺	2.63	0.048	0.8657	12.48	0.0400	0.9999	0.1563	0.9909
MB	2.07	0.061	0.6893	2.99	0.1020	0.9993	0.0464	0.986
FB	1.24	0.043	0.9626	2.48	0.0550	0.9974	0.0789	0.9091
RhB	0.58	0.055	0.9130	3.08	0.1500	0.9997	0.0149	0.9531

$$\ln Q = \ln K_f + \frac{\ln C_e}{n} \quad (7)$$

where *C_e* is the equilibrium concentration of metal ions in solution (mg L⁻¹), and *Q* is the equilibrium adsorption capacity (mg g⁻¹), *Q_{max}* (mg g⁻¹) and *K_L* (L mg⁻¹) are the Langmuir constant which is related to the adsorption capacity and energy of adsorption, respectively. *K_f* is the Freundlich constant related to adsorption capacity.

The *R*² of both fitting models is relatively high, and the *R*² of the Langmuir model is generally higher than that of the Freundlich model. This result indicates that the adsorption of pollutants on PDA/CS/Fe₃O₄ nano-sorbents occurred as a single monolayer, and the interaction force between metal ions can be ignored. Based on the theoretical analysis of the obtained data, the theoretical *Q_{max}* value of the nano adsorbent for Cu(II), Hg(II), Pb(II), MB, FB and RhB were respectively 909.0 mg g⁻¹, 94.0 mg g⁻¹, 20.2 mg g⁻¹, 3.6 mg g⁻¹, 2.4 mg g⁻¹, 2.9 mg g⁻¹, which was very close to the experimental data.

3.6 Desorption and reusability

Considering the cost in practical applications, it is necessary to study the analysis and regeneration of nano-adsorber. Desorption tests were studied to assess the regeneration and recycling of the CS@PDA@Fe₃O₄ nanocomposite regarding removing heavy metal ions and dye molecules *via* performing six adsorption/desorption cycles. It can be seen from Fig. 4 that the removal rate of nano-adsorber decreased slightly after the analysis operation with an ethanol solution, but it still has high removal efficiency.

Fig. 4 Desorption and reusability of CS@PDA@Fe₃O₄.

4. Conclusions

In this paper, a simple method is used to synthesize chitosan-grafted CS@PDA@Fe₃O₄ magnetic nano-adsorber with Fe₃O₄ as the magnetic core and PDA as the intermediate layer. These nano-adsorber membranes were utilized to eliminate the heavy metal ions (Cu(II), Hg(II), Pb(II)) and dye molecules (MB, FB, RhB) from aqueous solutions. The effects of static adsorption parameters such as pH, adsorbent concentration, metal initial concentration and time on adsorption performance were investigated. The results showed that the magnetic nano-adsorber had good adsorption performance and magnetic responsiveness. Besides, the experiment of desorption and regeneration proves that CS@PDA@Fe₃O₄ magnetic nano-adsorber can be recycled. This study has practical guiding significance for the treatment of pollutants in adsorbed water.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the Innovation Fund of Small and Medium-sized Enterprises of Gansu province (Grant No. 1407GCCA013), the Key Technology and Industrial Application Demonstration Project of High Quality and High Purity Nano Calcium Carbonate of Guangxi province (Grant No. 17202030-2).

Table 2 Parameters of isotherm model for adsorption of pollutants on CS@PDA@Fe₃O₄

Pollutants	Langmuir parameters			Freundlich parameters		
	<i>K_L</i>	<i>Q_{max}</i>	<i>R</i> ²	$\ln K_f$	<i>N</i>	<i>R</i> ²
Cu(II)	22.0	909.0	0.9759	9.2199	1.015	0.9713
Hg(II)	35.4	94.0	0.9868	5.246	2.70	0.9883
Pb(II)	165.0	20.2	0.9955	3.4434	5.50	0.9490
MB	0.4	3.6	0.9833	0.0565	2.416	0.9267
FB	12.3	2.4	0.9905	0.3238	5.214	0.9501
RhB	13.1	2.9	0.9287	0.4508	3.948	0.9465

References

- 1 M. A. Albakri, T. A. Saleh, Y. Mankour, T. F. Garrison and O. C. S. Al Hamouz, *J. Colloid Interface Sci.*, 2021, **582**, 428–438.
- 2 N. Ali, S. Uddin, A. Khan, S. Khan, S. Khan, N. Ali, H. Khan, H. Khan and M. Bilal, *Int. J. Biol. Macromol.*, 2020, **161**, 1305–1317.
- 3 V. Ball, D. Del Frari, M. Michel, M. J. Buehler, V. Toniazio, M. K. Singh, J. Gracio and D. Ruch, *BioNanoScience*, 2011, **2**, 16–34.
- 4 A. R. Cestari, E. F. Vieira, A. A. Pinto and E. C. Lopes, *J. Colloid Interface Sci.*, 2005, **292**, 363–372.
- 5 C. Dumitriu, S. I. Voicu, A. Muhulet, G. Nechifor, S. Popescu, C. Ungureanu, A. Carja, F. Miculescu, R. Trusca and C. Pirvu, *Carbohydr. Polym.*, 2018, **181**, 215–223.
- 6 Y. Guan, H. Y. Yu, S. Y. H. Abdalkarim, C. Wang, F. Tang, J. Marek, W. L. Chen, J. Militky and J. M. Yao, *Int. J. Biol. Macromol.*, 2019, **132**, 51–62.
- 7 V. K. Gupta, P. J. M. Carrott, M. M. L. Ribeiro Carrott and Suhas, *Crit. Rev. Environ. Sci. Technol.*, 2009, **39**, 783–842.
- 8 C. Hou, H. Zhu, Y. Li, Y. Li, X. Wang, W. Zhu and R. Zhou, *Appl. Microbiol. Biotechnol.*, 2015, **99**, 1249–1259.
- 9 A. Khan, S. Badshah and C. Airoidi, *Colloids Surf., B*, 2011, **87**, 88–95.
- 10 A. Khan, S. Begum, N. Ali, S. Khan, S. Hussain and M. Sotomayor, *Water Sci. Technol.*, 2017, **75**, 2034–2046.
- 11 A. Khan, F. Wahid, N. Ali, S. Badshah and C. Airoidi, *Desalin. Water Treat.*, 2014, **56**, 1099–1109.
- 12 C. Lee and S. Y. Lee, *Colloids Surf., B*, 2015, **127**, 89–95.
- 13 Y. Liu, K. Ai and L. Lu, *Chem. Rev.*, 2014, **114**, 5057–5115.
- 14 Y. Liu and Y.-J. Liu, *Sep. Purif. Technol.*, 2008, **61**, 229–242.
- 15 C. Lobo, J. Castellari, J. Colman Lerner, N. Bertola and N. Zaritzky, *Int. J. Biol. Macromol.*, 2020, **164**, 1575–1583.
- 16 A. L. Popovic, J. D. Rusmirovic, Z. Velickovic, T. Kovacevic, A. Jovanovic, I. Cvijetic and A. D. Marinkovic, *J. Ind. Eng. Chem.*, 2021, **93**, 302–314.
- 17 A. K. Sharma, Priya, B. S. Kaith, N. Sharma, J. K. Bhatia, V. Tanwar, S. Panchal and S. Bajaj, *Int. J. Biol. Macromol.*, 2019, **124**, 331–345.
- 18 Y. Xie, C. Chen, X. Lu, F. Luo, C. Wang, A. Alsaedi and T. Hayat, *J. Hazard. Mater.*, 2019, **379**, 120786.
- 19 L. Yang, Y. Peng, C. Qian, G. Xing, J. He, C. Zhao and B. Lai, *Chemosphere*, 2021, **263**, 128120.
- 20 S. Yang, Y. Hou, S. Xiong, F. Chen, Y. Jiang, C. Pan, J. Tang and G. Yu, *Sep. Purif. Technol.*, 2021, **258**, 117986.
- 21 J. Ye, X. Cong, P. Zhang, E. Hoffmann, G. Zeng, Y. Liu, W. Fang, Y. Wu and H. Zhang, *Appl. Surf. Sci.*, 2015, **356**, 128–134.
- 22 S. Zhang, Y. Zhang, G. Bi, J. Liu, Z. Wang, Q. Xu, H. Xu and X. Li, *J. Hazard. Mater.*, 2014, **270**, 27–34.
- 23 X. Xin, Q. Wei, J. Yang, L. Yan, R. Feng, G. Chen, B. Du and H. Li, *Chem. Eng. J.*, 2012, **184**, 132–140.
- 24 X. Li, Y. Qi, Y. Li, Y. Zhang, X. He and Y. Wang, *Bioresour. Technol.*, 2013, **142**, 611–619.
- 25 X. Guo, B. Du, Q. Wei, J. Yang, L. Hu, L. Yan and W. Xu, *J. Hazard. Mater.*, 2014, **278**, 211–220.
- 26 L. Zhou, Y. Wang, Z. Liu and Q. Huang, *J. Hazard. Mater.*, 2009, **161**, 995–1002.
- 27 W. Lee, J. U. Lee, B. M. Jung, J.-H. Byun, J.-W. Yi, S.-B. Lee and B.-S. Kim, *Carbon*, 2013, **65**, 296–304.
- 28 Y. Wang, Y. Zhang, C. Hou and M. Liu, *J. Taiwan Inst. Chem. Eng.*, 2016, **61**, 292–298.
- 29 X. Liu and D. J. Lee, *Bioresour. Technol.*, 2014, **160**, 24–31.
- 30 L. Hadjittofi, M. Prodromou and I. Pashalidis, *Bioresour. Technol.*, 2014, **159**, 460–464.