

Original Research

The Efficiency Adsorption of Ammonia Nitrogen, Phosphate and Basic Blue 3 by Fulvic Acid Decorated Fe₃O₄ Magnetic Nanocomposites

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Abstract

Magnetic nanocomposites have attracted much attention in the field of wastewater treatment due to their great advantages of fast and efficient removal and separation. Therefore, Fe₃O₄ magnetic nanocomposites coated with fulvic acid (Fe₃O₄/FA) were applied to remove ammonia nitrogen, phosphate and basic blue 3 (BB3) from the effluent. The performance of the adsorbents towards the pollutant removal under different contact time, pH, ionic strength and temperatures were investigated. The whole process with adsorption kinetics and adsorption thermodynamics were also studied. The results showed that the adsorption equilibrium could be achieved in 180 min. The high susceptibility of ammonia nitrogen, phosphorus and BB3 to the alternation of pH, ionic strength and temperatures indicated that electrostatic interaction played a dominant role in the process. Fe₃O₄/FA exhibited excellent adsorption performance for ammonia nitrogen, phosphorus and BB3, and the q_{max} obtained by the Langmuir model were 106.21 mg/g, 116.81 mg/g and 203.54 mg/g, respectively. Its adsorption performance remained constant in the treatment of ammonia nitrogen, phosphate and BB3 after consecutive 6 cycles. In practical wastewater treatment, Fe₃O₄/FA improved the removal of ammonia nitrogen and phosphate. Therefore, Fe₃O₄/FA magnetic nanocomposites have a promising application for wastewater treatment.

Keywords: magnetic nanocomposites, fulvic acid, ammonia nitrogen, phosphate, basic blue 3

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Introduction

Wastewater containing nitrogen, phosphorus and dye from urban living and industrial development has put forward a great challenge to the environment. Once surplus nitrogen and phosphorus were discharged into wastewater, the aquatic organism would suffer breathing difficulties, weakened immunity and even mass mortality [1]. What's more, the high concentration of ammonia nitrogen enriched in organisms can be further converted into strong oxidation and carcinogenic nitrite, resulting in greater harm [2]. Excessive phosphorus content will lead to eutrophication of waters, rapid overgrowth of aquatic algae and the sharp decrease of water dissolved oxygen content. At the same time, eutrophication can also cause water odor, deteriorate the water quality, and increase the difficulty and cost of water treatment [3].

Dyeing wastewater with deep chromaticity and complex composition is harmful to the environment. Dyes own the characteristics of absorbing light, reducing the transparency of water, consuming a large number of dissolved oxygen, affecting aquatic organisms and microbial growth, and destroying the self-purification capacity of water. The aromatic polyphenyl ring derivatives produced by substituting hydrogen on the dye benzene ring to amine, nitro and halogen groups own high biotoxicity, and some of them even have malign effects as teratogenic, carcinogenic and mutagenic, etc. Besides, the dyeing wastewater may contain heavy metals, such as cuprum, chromium, lead and so on. These heavy metals with high toxicity are difficult to be degraded by organisms, which further aggravate heavy metal pollution and increase plants, animals, and human health risks [4, 5].

So far, different kinds of physical and chemical methods have been applied to removing nitrogen, phosphorus and dye in the wastewater treatment, such as adsorption, coagulation, oxidation, electrolysis, and biological treatment [6, 7]. Compared with these methods, adsorption is a high-efficiency, economical and convenient method with good selectivity. Particularly, it presents practical application value in removing low concentration pollution. Many titanate nanotubes and polymer-based nanomaterials have excellent adsorption properties for nitrogen, phosphorus, dyes and organic pollutants [8-13]. However, the difficulty of recycling these materials after adsorption limited its application in water purification and remediation.

Magnetite particles (Fe_3O_4), as a kind of highly magnetic nanomaterial, perfectly solved the problem of adsorbent recycling. Fe_3O_4 has been widely used in removing various pollutants for its good saturation magnetic strength and easiness of being re-collected with a magnet from water [14, 15]. However, the pure Fe_3O_4 can be oxidized in air and easily aggregate in an aqueous solution. To solve the problem, fulvic acid (FA) was introduced to the surface of the magnetic particles as a protective agent. FA is an organic substance in the

soil and water environment with many active sites on its surface for the complexation of pollutants. Hence, $\text{Fe}_3\text{O}_4/\text{FA}$ composites were made as a novel stable core-shell composite material. The composites not simply have the magnetic separation property but also have the multi-functional group characteristic of shell organics, showing good removal performance for pollutants [16-19]. According to our previous study, the agglomeration of Fe_3O_4 nanoparticle could be significantly reduced and the dispersity of the composite was able to be improved in the water solution with the participation of FA [15].

Therefore, the performance of $\text{Fe}_3\text{O}_4/\text{FA}$ composites in removing ammonia nitrogen, phosphate and basic blue 3 (BB3) under different environmental conditions, including ion strength, pH and temperature were discussed. Meanwhile, the reuse performance of $\text{Fe}_3\text{O}_4/\text{FA}$ was studied for practical application.

Material and Methods

Chemicals

$\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$, $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$, ammonium hydroxide, NH_4Cl , NaCl , HCl , NaOH and KH_2PO_4 were provided by China Chemical Reagent Co., LTD. The fulvic acid (FA) was purchased from Aldrich (Sigma-Aldrich, Steinheim, Germany). Basic blue 3 (BB3) was provided by Sinopharm Chemical Reagent Co. Ltd. (Shanghai, China). Ammonia nitrogen solution was prepared by dissolving NH_4Cl with appropriate amounts of deionized water. Phosphate solution was prepared by dissolving KH_2PO_4 with appropriate amounts of deionized water. All of the chemical reagents used in experiments were analytical grade reagents.

Preparation of Adsorbents

$\text{Fe}_3\text{O}_4/\text{FA}$ magnetic composites were prepared by the co-precipitation method [17,20-21]. The detailed procedures were as follows: dissolving 10.0 g of $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ and 8.0 g of $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ into 200 mL of distilled water and heated to 90°C, then adding 20 mL of ammonium hydroxide and 1.0 g of FA immediately. The mixture would react for 1 h under the protection of nitrogen at 90°C and cooled to room temperature. The black precipitate at the bottom of the beaker was collected and washed by distilled water many times till pH reached neutral. Finally, colloidal samples were left in a vacuum drier to dry and then ground for subsequent use.

Adsorption Procedures

All adsorption experiments were conducted in 10 mL polyethylene centrifuge tubes. In detail, the suspension liquid of $\text{Fe}_3\text{O}_4/\text{FA}$ adsorbents and background electrolyte solution NaCl were mixed in advance. The stock solution of ammonia

nitrogen, phosphate and BB3 was added to make the concentration of each component reach the preset value, respectively. pH was adjusted to the desired value with the trace amount of HCl and NaOH solution at the concentration of 0.1 mol/L. The centrifuge tube with suspension was oscillated at 150 rpm to make the adsorption process reach equilibrium and then the supernatant was obtained by solid-liquid separation with the magnet, and the residual concentration of the pollutant was measured by spectrophotometer. In the desorption experiments, 500 mL of solution containing 0.01 mol/L NaCl with 50 mg/L ammonia nitrogen, phosphorus and BB3 were respectively placed into three beakers and adding 0.5 g/L $\text{Fe}_3\text{O}_4/\text{FA}$ into them. The suspension was mechanically stirred for 24 hours before magnetic separation. The solid part was repeatedly rinsed with 0.1 mol/L HCl or NaOH solution until no contaminants were detected in the eluent. Last, the residual $\text{Fe}_3\text{O}_4/\text{FA}$ was collected and dried at 60°C. Six times adsorption-desorption experiments were carried out and the adsorption rates of pollutants on magnetic material were subsequently measured.

To study the adsorption properties of the composites in mixed effluent, the industrial wastewater in the Shaoxing area was referred to prepare the simulated wastewater. It contains 10 mg/L Ni(II), 10 mg/L Pb(II), 50 mg/L ammonia nitrogen, 50 mg/L phosphate, 50 mg/L BB3 and 0.01 mol/L NaCl (as background electrolyte). 250 mL simulated wastewater was added into a 500 mL beaker following the above-mentioned methods at pH 5.0 to carry out static batch experiment.

Analytical Methods

The sorption rate and adsorption capacity of the contaminants on the magnetic composites were calculated by (1) and (2):

$$\text{Sorption rate (\%)} = (C_0 - C_{\text{eq}}) / C_0 \times 100\% \quad (1)$$

$$\text{Adsorption capacity } q_e \text{ (mg/g)} = (C_0 - C_{\text{eq}}) \cdot V / m \quad (2)$$

...where C_0 (mg/L) was the initial concentration of the pollutants, C_{eq} (mg/L) was the equilibrium concentration of the pollutants, V (L) was the volume of solution, and m (g) is the weight of the adsorbents used. All experiments were duplicated and the average standard deviation of the duplicated experiments was less than $\pm 5.0\%$.

Results and Discussion

Effect of Contact Time

Fig. 1 described the adsorption kinetics curve of ammonia nitrogen, phosphate and BB3 on $\text{Fe}_3\text{O}_4/\text{FA}$ composites. With the contact time increased from 10 min to 180 min, the sorption rate (%) of

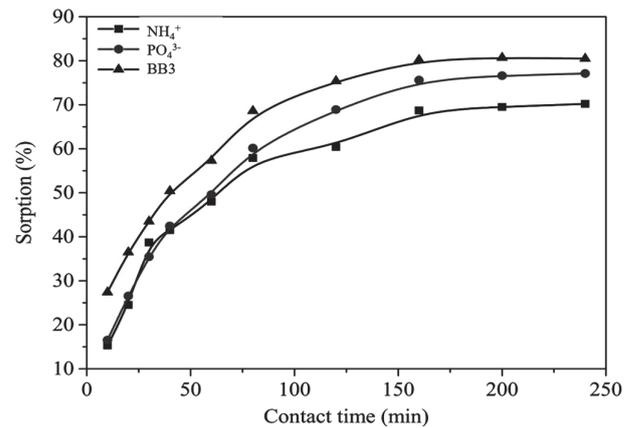


Fig. 1. Effect of contact time on the adsorption of ammonia nitrogen, phosphate and BB3 by $\text{Fe}_3\text{O}_4/\text{FA}$ (pH = 5.0, T = 298K, m/V = 0.5g/L, the initial concentration of ammonia nitrogen, phosphate and BB3 were 50 mg/L).

ammonia nitrogen rise from 15% to 70% and attained adsorption equilibrium. The adsorption tendency of phosphate and BB3 were analogous to ammonia nitrogen's curve, reaching the peak value at 180 min. The adsorption kinetics of ammonia nitrogen, phosphate and BB3 achieved equilibrium in just 180 min, which confirmed that the removal mechanism of the three pollutants on $\text{Fe}_3\text{O}_4/\text{FA}$ was essentially chemisorption.

Effect of pH and Ionic Strength

The adsorption of ammonia nitrogen, phosphate and BB3 on $\text{Fe}_3\text{O}_4/\text{FA}$ composites in different pH and ionic strengths were respectively shown in Fig. 2. As shown in Fig. 2a) and Fig. 2b), the sorption rate (%) of ammonia nitrogen, phosphate gradually decreased with the increase of pH value, indicating that the alternative of the solution from acidic to alkaline was not conducive to the removal of ammonia nitrogen and phosphate. The zero charge point of the composite was 2.2 in our previous study [15]. At pH > 3.0, the functional group sites on its surface were negatively charged due to deprotonation $\text{SOH} \leftrightarrow \text{SO}^- + \text{H}^+$, and the electronegativity gradually increased with the increase of pH value. Ammonia nitrogen and phosphorus are usually electronegative which makes it hard to bond to the surface of the composites for the same electrical properties and leads to the dropping adsorption rate as the pH value goes up. The adsorption of BB3 has the opposite trend comparing with ammonia nitrogen and phosphorus (Fig. 2c). BB3 is positively charged in a wide pH range. With the increase of pH, the charge attraction of these forms and the surface sites of $\text{Fe}_3\text{O}_4/\text{FA}$ composite gradually increased, leading to the increase of adsorption rate. These experimental phenomena indicated that the electrostatic interaction was the dominant adsorption mechanism [22-25].

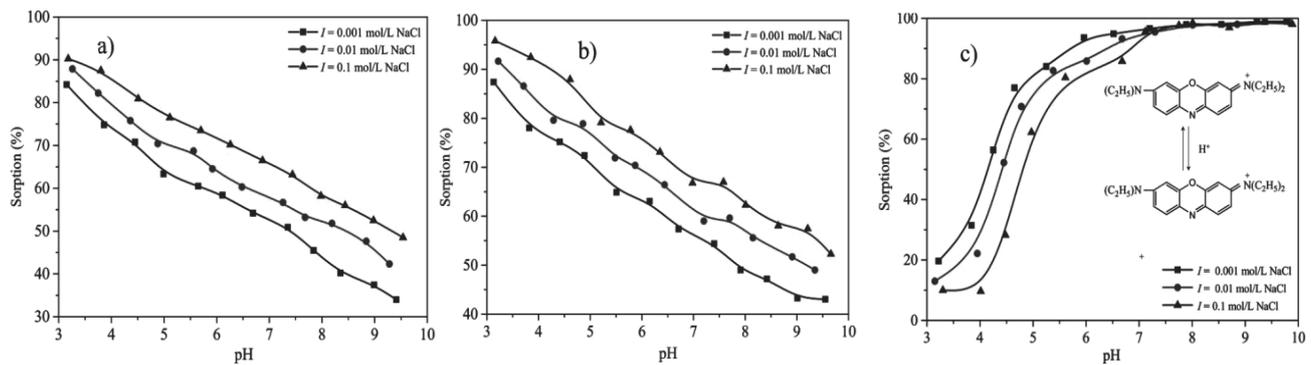


Fig. 2. Effect of pH and ionic strength on the adsorption of a) nitrogen; b) phosphorus; c) BB3 by $\text{Fe}_3\text{O}_4/\text{FA}$ ($T = 298 \text{ K}$, $m/V = 0.5 \text{ g/L}$, $C_{\text{NH}_4^+}$, $C_{\text{phosphorus}}$ and $C_{\text{BB}_3} = 50 \text{ mg/L}$).

Sorption Isotherms

To further investigate the adsorption mechanism, sorption isotherms of $\text{Fe}_3\text{O}_4/\text{FA}$ composites on ammonia nitrogen, phosphorus and BB3 were depicted in Fig. 3. We can see that the sorption of ammonia nitrogen and phosphorus increased with the increasing temperature, indicating the whole adsorption process of ammonia nitrogen and phosphorus was endothermic. The sorption of BB3 decreased with the increasing temperature, suggesting the adsorption process of BB3 was exothermic. The Langmuir model and Freundlich model were applied to simulate the process. All of the parameters were listed in Table 1-3.

Compared with the correlation coefficient (R^2), the Langmuir model fitted better than the Freundlich model. The results proved that the surface of $\text{Fe}_3\text{O}_4/\text{FA}$ composites was relatively uniform, and chemisorption was the main removal mechanism of ammonia nitrogen, phosphorus and BB3 pollutants. The q_{max} obtained by the Langmuir model for ammonia nitrogen and phosphorus at 328 K were 106.21 mg/g and 116.81 mg/g, respectively. The q_{max} obtained by the Langmuir model for BB3 at 298 K were 203.54 mg/g. Under the condition of the above three temperature, the

q_e in the experiment were smaller than the q_{max} obtained by the Langmuir model fitting. This suggested that the unsaturated monolayer adsorption of ammonia nitrogen, phosphorus and BB3 on $\text{Fe}_3\text{O}_4/\text{FA}$. The n value of the experimental results of the Freundlich model was between 0 and 1, indicating that the adsorption of ammonia nitrogen, phosphorus and BB3 on $\text{Fe}_3\text{O}_4/\text{FA}$ composite was a spontaneous nonlinear process.

Regeneration

The reuse performance of the adsorbents is a key indicator to evaluate the potential application of the adsorbents in practical wastewater treatment. Therefore, it is imperative to explore the reuse performance of $\text{Fe}_3\text{O}_4/\text{FA}$ composites. Considering the pH effect on the removal of the three pollutants, we adopted 0.1 M of NaOH solution as the eluent of ammonia nitrogen and phosphorus, and 0.1 M of HCl solution as the eluent of BB3. After being regenerated, the composites were used to finish the subsequent removal experiment. The sorption rates (%) of the three pollutants on $\text{Fe}_3\text{O}_4/\text{FA}$ composites almost stayed the same after six successive adsorption-desorption cycles (Fig. 4). The result

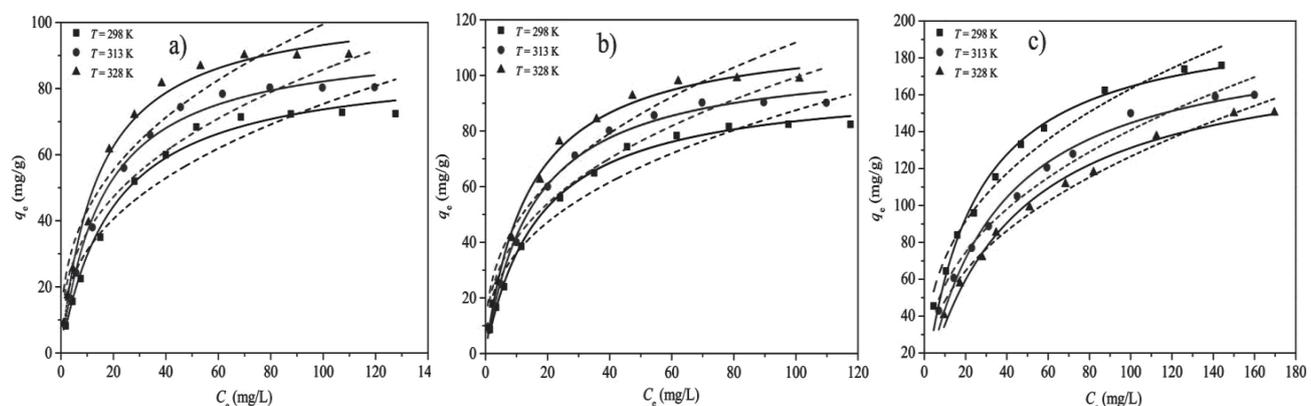


Fig. 3. Adsorption isotherms of a) ammonia nitrogen; b) phosphorus; c) BB3 on $\text{Fe}_3\text{O}_4/\text{FA}$ (solid lines denote the fit curves of Langmuir model and dash lines denote the fit curves of Freundlich model, $\text{pH} = 5.0$, $m/V = 0.5 \text{ g/L}$, $I = 0.01 \text{ mol/L NaCl}$).

Table 1. Parameters of the Langmuir and Freundlich models for the adsorption of ammonia nitrogen on Fe₃O₄/FA.

Correlation parameters		T = 298 K	T = 313 K	T = 328 K
Langmuir	q_{max} (mg/g)	88.30	95.32	106.21
	b (L/mg)	0.050	0.062	0.072
	R^2	0.990	0.993	0.990
Freundlich	K_f (mg ¹⁻ⁿ ·L ⁿ /g)	12.78	15.58	18.61
	n	0.385	0.370	0.418
	R^2	0.914	0.917	0.364

Table 2 Parameters of the Langmuir and Freundlich models for the adsorption of phosphorus on Fe₃O₄/FA.

Correlation parameters		T = 298 K	T = 313 K	T = 328 K
Langmuir	q_{max} (mg/g)	97.87	107.39	116.81
	b (L/mg)	0.058	0.065	0.071
	R^2	0.996	0.994	0.995
Freundlich	K_f (mg ¹⁻ⁿ ·L ⁿ /g)	14.98	17.30	19.73
	n	0.383	0.379	0.377
	R^2	0.934	0.921	0.933

Table 3 Parameters of the Langmuir and Freundlich models for the adsorption of BB3 on Fe₃O₄/FA.

Correlation parameters		T = 298 K	T = 313 K	T = 328 K
Langmuir	q_{max} (mg/g)	203.54	194.82	187.28
	b (L/mg)	0.042	0.029	0.023
	R^2	0.989	0.984	0.986
Freundlich	K_f (mg ¹⁻ⁿ ·L ⁿ /g)	31.06	22.84	18.43
	n	0.360	0.395	0.418
	R^2	0.950	0.948	0.954

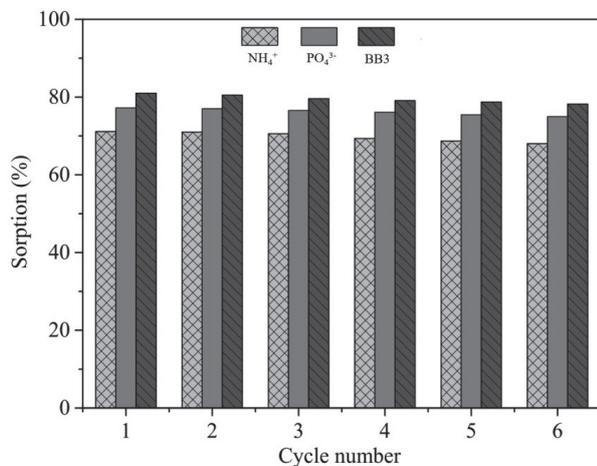


Fig. 4. The reuse performance of Fe₃O₄/FA for ammonia nitrogen, phosphorus and BB3 removal (pH = 5.0, T = 298K, m/V = 0.5 g/L, I = 0.01 mol/L NaCl, the concentration of NH₄⁺, PO₄³⁻ and BB3 were 50 mg/L).

showed excellent reused performance of the magnetic composites.

Removal Performance in Mixed Wastewater

In order to assess the practical potential of Fe₃O₄/FA composites in sewage treatment, the simulated wastewater was prepared for further study on the removal performance of the adsorbents. We made a comparison between the sorption rate (%) of each contaminated component in the multi-solute system and the corresponding single-solute system to obtain the removal efficiency of the composites. As shown in Fig. 5, we found that the sorption rates (%) of Ni(II), Pb(II) and BB3 in simulated wastewater by Fe₃O₄/FA were lower than those of the corresponding single-component system, while the removal of ammonia

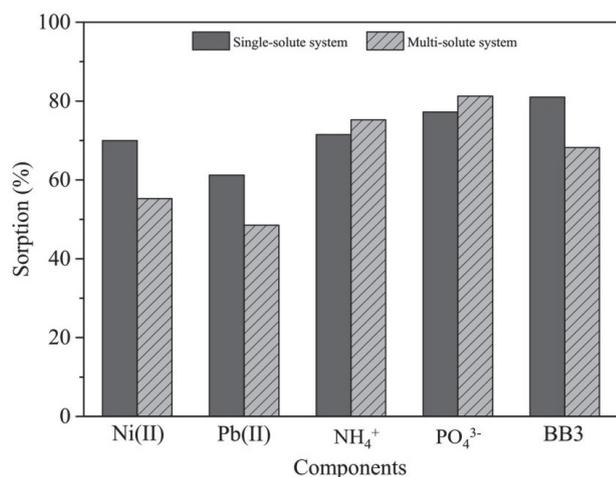


Fig. 5. The adsorption performance of Fe₃O₄/FA in simulation wastewater with multiple pollutant components (pH = 5.0, T = 298K, m/V = 0.5 g/L, 10 mg/L Ni(II), 10 mg/L Pb(II), I = 0.01 mol/L NaCl, the concentration of NH₄⁺, PO₄³⁻ and BB3 were 50 mg/L).

nitrogen and phosphorus were higher than those of the corresponding single-component system. The results were possibly derived from interaction among multiple components, which influenced electrostatic and complexation between pollutants and adsorbents [26]. Generally, Fe₃O₄/FA composites showed strong ability in the removal of the five pollutants in mixed wastewater.

Conclusions

In the study, we found that the adsorption of Fe₃O₄/FA on ammonia nitrogen, phosphorus and BB3 could achieve equilibrium in 180 min. As the ionic strength increased, the adsorbability of ammonia nitrogen and phosphorus was improved, but the adsorption of BB3 was inhibited. The results indicated that electrostatic binding was the dominant adsorption mechanism in the process. The q_{max} obtained by the Langmuir model for ammonia nitrogen, phosphorus and BB3 were 106.21 mg/g, 116.81 mg/g and 203.54 mg/g, respectively. The adsorption capacity for ammonia nitrogen and phosphorus increased with the increasing temperature, indicating that the removal process was an endothermic process. The opposite phenomenon was found for the adsorption of BB3, suggesting the removal process of BB3 was an exothermic process. The adsorption process matched better with the Langmuir model which revealed that the surface of the composites was relatively uniform and chemisorption was the major removal mechanism. Ammonia nitrogen, phosphorus and BB3 occurred unsaturated monolayer adsorption in the surface of the composites, which belonged to spontaneous process. The removal of ammonia nitrogen and phosphorus were higher than those of the corresponding single-component system,

while the removal of BB3 was lower than those of the corresponding single-component system. The experimental phenomenon probably resulted from the interaction among multiple components and affected electrostatic interaction and complexation between those pollutants and adsorbents. Furthermore, the adsorption of ammonia nitrogen, phosphorus and BB3 on Fe₃O₄/FA remained constant after six adsorption-desorption cycles, showing excellent regeneration capacity.

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Conflict of Interest

The authors declare no conflict of interest.

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