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Highly efficient removal of uranium(vi) from wastewater by polyacrylic acid hydrogels†

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Polyacrylic acid (PAA) hydrogel prepared by radical polymerization in a clean and extremely simple way was used to adsorb U(vi) ions from aqueous solutions. Fourier transform infrared spectroscopy (FTIR) characterization demonstrated that the PAA hydrogels were successfully synthesized, FTIR and transmission electron microscope (TEM) characterization illustrated the adsorption mechanism, and the adherence of the U(vi) ions on PAA was confirmed by X-ray photoelectron spectroscopy (XPS) energy dispersive X-ray (EDX). Batch experiments with various pH, initial concentration and contact time were conducted to evaluate the adsorption performance. The adsorption kinetics of the PAA hydrogels followed a pseudo-second-order model and exhibited a 3-stage intraparticle diffusion mode. Equilibrium data were best described by a Langmuir model, and the estimated maximum adsorption capacity for the adsorbent was 445.11 mg g⁻¹. Moreover, the U(vi) loaded adsorbent could be regenerated using 0.1 M HNO₃ acid solution, and the regenerated adsorbent after five cycles could retain 90.91% of the adsorption capacity of the fresh adsorbent. The results suggested that the PAA hydrogels could be considered to be very effective and promising materials for U(vi) removal from waste water.

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1. Introduction

Over the years, an increase in some given pollutants such as industrial waste, have contaminated fresh water and caused deterioration in the quality of potable water. Waste water from electronic device manufacture, plating facilities and the production of electricity in nuclear power plants may contain considerable amounts of toxic and polluting heavy metals.^{1,2} Among the metal contaminants, particular concern has been given to uranium, an important actinide element.³ Uranium is a heavy metal having both radiochemical and toxicological effects.⁴ Excessive amounts of uranium have entered into the environment through the activities of the nuclear industry due to different applications including nuclear power plants, the manufacture of glass, pottery glaze, electron microscopes and nuclear weapons.⁵ Uranium in the environment can ultimately reach the top of the food chain and be ingested by humans, causing harmful diseases such as severe kidney or liver damage.⁶ Therefore, methods for the removal of uranium from wastewater are of great importance.

The removal of heavy metal ions from waste water can be performed by physical and chemical methods such as adsorption, chemical precipitation, membrane filtration, chromatographic extraction, ion exchange and electrochemical techniques.^{7,8} And the adsorption is considered to be

a promising and efficient method for removal of U(vi) and other metal ions.⁹ The adsorption mechanism involves the establishment of equilibria between the adsorbate and the adsorbent through processes such as ion exchange, chelation, complexation, electrostatic interactions and micro-precipitation even at extremely low concentrations, thus facilitates the removal of heavy metal ions from very dilute solutions.¹⁰ Adsorption of U(vi) ions on different functional groups containing adsorbents from aqueous solution had been widely studied. Results indicated that the adsorption of U(vi) ions was mostly dominated by complexation and/or ion exchange at acidic pH values.⁹ A number of adsorbents such as, polymers,¹¹ functionalized SBA-15,¹² chitosan¹³ and MOFs,¹⁴ have been investigated for the removal of U(vi) from aqueous solutions. Among them, polymer materials, with abundant adsorption sites, have attracted wide attention. Adsorption of U(vi) cations using polymers takes place by the attraction mechanism, counting on the difference in charge between the positively charged metal ion and the negatively charged active sites distributed everywhere along the backbone and branch of any polymer, where these active sites govern the whole uptake of the U(vi) cations by the polymers.

Hydrogel, one of the most widely used polymers, consists of a three-dimensional network of hydrophilic polymers which allows the diffusion of solutes into the interior network and can be prepared using a covalent or a non-covalent approach with different functional groups such as carboxylic acid, hydroxyl and amine groups.¹⁵ These groups attached onto the polymeric networks can be used easily for the removal of metal ions. Synthesis and characterization of poly(HEA/MALA) hydrogel and

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its application in removal of heavy metal ions from water were conducted by Wu.¹⁶ Wang and Li¹⁷ have used ion-imprinted composite hydrogels to investigate Cu²⁺ removal from waste water. Pb²⁺ and Cu²⁺ sorption properties of Gum Ghatti-grafted poly(acrylamide-co-acrylonitrile) biodegradable hydrogel were studied by Mittal.¹⁸ The most widely abundant function groups present in the structure of any hydrogel polymer are hydroxyl groups and carbonyl groups but although these hydroxyl and carbonyl groups are present in great abundance, they are not effective sites for removing and attracting the metal cations. Functional groups like carboxyl groups were found to have a very governing role in removing heavy metal cations, that they bind to the heavy metals through salt formation.¹¹ Polyacrylic acid (PAA), is remarkably stable and can deliver high binding capacity because of its high density of carboxyl-containing.¹⁹ Although there have been extensive academic and industrial interests in the studies of the hydrogel polymer recently,^{20,21} few of them have focused on the removal of uranium using PAA hydrogel materials.

In the present work, PAA hydrogel was prepared by free radical polymerization as a clean and extremely simple way, and was used for removal of U(vi) ions from aqueous solution. The synthesized PAA hydrogels are not water soluble and were characterized. In order to explore the adsorption ability and mechanism of the PAA polymer hydrogel, the influence of experimental conditions such as pH, contact time and initial concentration were investigated. The adsorption kinetics and isotherms of U(vi) cations were also evaluated. The application of PAA based hydrogel to remediate U(vi) cations as in our case has not been reported so far. From the results obtained in this work, we could establish a suitable strategy with promising application for efficient harmless removal of U(vi) ions.

2. Materials and methods

2.1 Materials

Acrylic acid (AA) and Methylene-Bis-Acrylamide (MBA) were supplied by Sinopharm Chemical Reagent Co., Ltd. Ammonium persulphate (APS) was provided by Shanghai SANGON Biological Engineering Co., Ltd, China. Sodium Hydrogen Sulfite (SHS) was purchased from Xilong Chemical Co., Ltd. All chemicals were of analytical grade and were used without further purification. All water mentioned in this paper was double deionized water (was prepared by Fuzhou University).

Stock solutions (1000 mg L⁻¹) of U(vi) ions were prepared by dissolving 1.859 g UO₂(NO₃)₂·6H₂O in 1000 mL distilled water. The solutions of different concentrations used in various experiments were obtained by dilution of the stock solutions.

2.2 Preparation of PAA hydrogel

PAA was prepared by radical polymerization with APS as an initiator and SHS as coagulant aid in room temperature.²² The acrylic monomer with a mass fraction of 30% and 0.5 g MBA were dissolved in a beaker by supersonic wave. About 0.21 g APS and 0.21 g SHS were added into this solution. Then the final solution volume was diluted to 50 mL under mechanical stirring. Finally, the obtained solution above stands at 25 °C for 2 h.

The product was washed with deionized water until the unreacted substance and fluid wax were completely removed, the hydrogel was dried in vacuum freeze-drying machine for 24 h and stored in vacuum drying oven at 50 °C.

2.3 Characterization of adsorbent

2.3.1 Fourier transform infrared spectroscopy (FTIR). FTIR spectra of the AA, PAA and PAA after adsorption were obtained by using a FTIR spectrophotometer. The FTIR spectrum was recorded with a spectrometer in the range of 4000–400 cm⁻¹ using an attenuated total reflectance technique.

2.3.2 X-ray photoelectron spectroscopy, Transmission electron microscopy and energy dispersive X-ray. The elemental compositions and surface species of the PAA before and after adsorption were analyzed by XPS (ESCALAB 250). Transmission electron microscopy (TEM) images were performed on a JEOL-1230 electron microscope operated at 100 kV. A scanning electron microscope (SEM, Nova Nano 230) was employed to perform EDX measurements. EDX measurements were obtained between 0 and 20 keV.

2.4 Batch adsorption experiments

The stock solution was diluted to give standard solutions of appropriate concentrations. The batch adsorption experiments were conducted in 50 mL conical flask with 10 mL of standard solutions. About 10 mg PAA was then added into this solution and was left shaking for a certain period of time. The conical flask were covered with rubber plug. After filtration, the concentration of U(vi) ions in the filtrate was analyzed at a wavelength of 650 nm using Arsenazo-III as the complexing agent by UV-vis spectrophotometer.²³

The adsorption capacity of U(vi) in the adsorption system, q_e , was calculated using the following equation:

$$\text{Adsorption capacity } (q_e) = \left(\frac{C_0 - C_e}{m} \right) V \quad (1)$$

where q_e is the equilibrium adsorption capacity per gram dry weight of the adsorbent (mg g⁻¹), C_0 and C_e are the initial and equilibrium concentration of U(vi) ions (mg L⁻¹), respectively. V is the volume of U(vi) ions solution (mL) and m is the mass of the adsorbent used (g).

2.4.1 Effect of pH. The effect of pH on the adsorption of U(vi) ions was studied at 25 °C in a pH range of 1–5.5 at metal ion concentrations of 60, 100 and 200 mg L⁻¹. The solution pH was adjusted to the desired value by adding either nitric acid or sodium hydroxide standardized solutions.

2.4.2 Effect of the initial concentration of the U(vi) ions. The effect of initial concentration of the U(vi) ions was carried out by soaking 10 mg of PAA in a series of conical flasks which contained 10 mL of U(vi) ions solution at definite concentrations (20–600 mg L⁻¹) and pH 4.0. The conical flasks were conditioned on the shaker at 150 rpm while keeping the temperature at 25 °C for about 3 h. After adsorption, solution was filtered and the residual concentration was determined.

2.4.3 Effect of contact time. The experiments were performed at pH 4.0 by shaking 30 mg PAA with 30 mL (60, 100 and



200 mg L⁻¹) U(VI) ions solution. The contents of the conical flask were agitated on a shaker at 150 rpm and 25 °C. One milliliters of solution after adsorption were taken at time intervals for the analysis of residual U(VI) ions concentration in solution.

2.4.4 Adsorption kinetic and isotherms studies of PAA.

Isotherms studies were performed by soaking 0.01 g PAA in a series of conical flasks containing 10 mL of U(VI) ions solution at different initial concentration (20, 50, 100, 200, 300, 400, 500 and 600 mg L⁻¹) shaking for 3 h at 25 °C. After adsorption, the residual concentration of U(VI) ions in the solution was determined. Sorption isotherms are plots of the equilibrium adsorption capacity (q_e) (according to eqn (1)) versus the equilibrium concentration of the residual U(VI) in the solution (C_e).

The kinetic studies were investigated by placing 0.01 g PAA in a series of conical flasks containing 10 mL of U(VI) ions solution with three different initial concentration (60, 100 and 200 mg L⁻¹), and the samples were shaken at 25 °C for designated time periods (5, 10, 20, 30, 60, 90, 120, 150, 180 min).

2.4.5 Desorption and regeneration experiments. After the adsorption experiment, the uranium loaded PAA (0.01 g) was left in contact with 10 mL 0.1 M HCl solution for 24 h at room temperature, and then washed with distilled water. Subsequently, it was dried in vacuum freeze-drying machine for 24 h before reuse. The adsorption-desorption procedure was conducted for five cycles under the same conditions.²⁴

3. Results and discussion

3.1 Characterization

The FTIR spectra of samples, including AA, PAA and PAA after adsorption, are presented in Fig. 1. The strong absorption peaks at 1695 cm⁻¹ are attributed to the C=O of carboxylic group from AA, PAA and U-PAA complexes.²⁵ In the spectrum of PAA, compared with that of AA, the peak of C=C at 1635 cm⁻¹ disappears, which indicated that the successful synthesis of PAA. In the spectrum of U-PAA complexes, two characteristic absorption bands lying at 1530 and 1450 cm⁻¹ are observed,

which are attributed to the symmetric COO⁻ and asymmetric COO⁻ of carboxylate, respectively. And the appearance of a peak at 929 cm⁻¹ is from the stretching vibration of O=U=O.²⁶ In addition, the weak absorption peak around 477 cm⁻¹ can be assigned to the stretching vibration of M-O bond.²⁷ These data indicate that PAA hydrogels were successfully chelated with UO₂²⁺ cations.

The coordination number of hexavalent uranium is often 5 or 6 in aqueous solution.²⁸ And according to the results of U-PAA complexes of FTIR characterization, each UO₂²⁺ is randomly combined with two carboxylates in the chain of the polymer (not necessarily in the same chain), water molecules can also participate the coordination to meet the saturated coordination.

The chemical states of the PAA before and after adsorption of uranium ions were investigated by XPS and results were shown in Fig. 2. It's obvious that PAA before adsorption showed three peaks of binding energy, C 1s (284 eV), N 1s (399 eV) and O 1s (532 eV). However, the peak at the binding energy of 382 eV (U 4f) appeared after adsorption of U(VI) ions. The elemental compositions of PAA calculated from the XPS spectra were summarized in Table 1. Compared with the pure PAA, the oxygen contents increased and the uranium element appeared on the PAA after adsorption, which indicated that the UO₂²⁺ was successfully grafted onto PAA.

The TEM micrograph of PAA is given in Fig. 3. Before adsorption, some holes and openings in the structure were observed in Fig. 3(c), which increases the contact area and can improve metal ion adsorption. And loosely structure can be

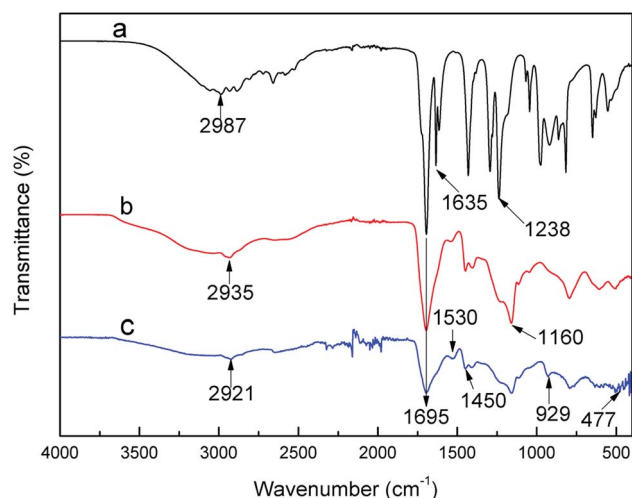


Fig. 1 Fourier transform-infrared (FT-IR) spectra of (a) AA, (b) PAA and (c) U-PAA complexes.

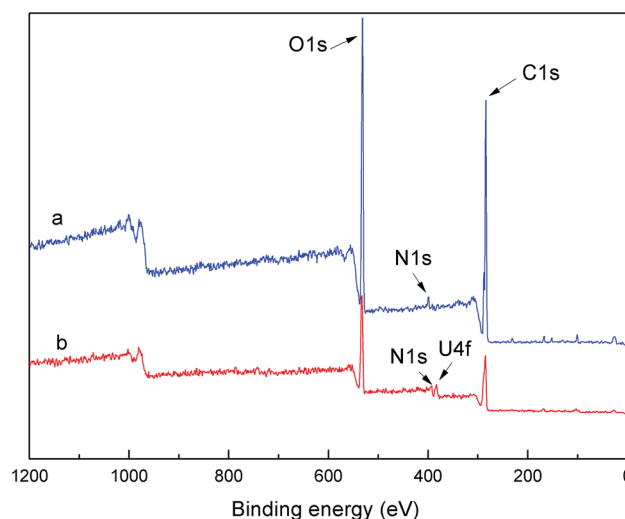


Fig. 2 XPS spectra for PAA. (a) Before adsorption and (b) after adsorption.

Table 1 Elemental composition of the PAA by XPS spectra

| Samples | C 1s (%) | N 1s (%) | O 1s (%) | U 4f (%) |
|-----------------------|----------|----------|----------|----------|
| PAA before adsorption | 68.68 | 7.47 | 23.85 | 0 |
| PAA after adsorption | 48.44 | 4.24 | 45.57 | 1.72 |



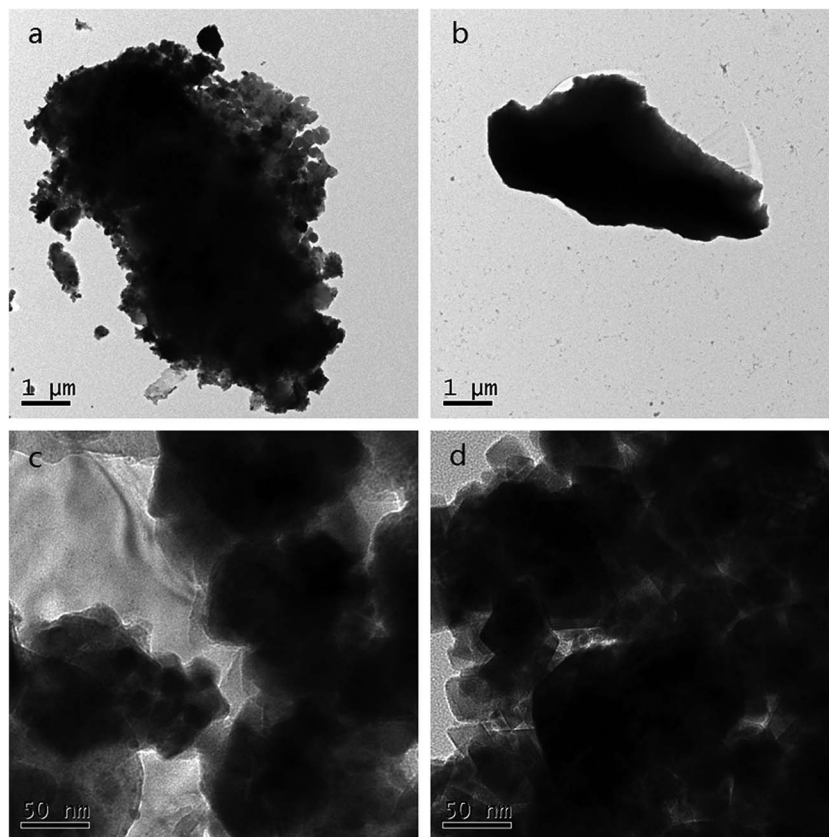


Fig. 3 TEM images of PAA. (a) and (c) Before adsorption, (b) and (d) after adsorption.

seen in Fig. 3(a), which may exist as a mass of carboxyl groups. After adsorption, Fig. 3(d) shows that the agglomeration became aggravated and dense microstructure can be seen in Fig. 3(b), demonstrating there may be strong chelation interactions between one UO_2^{2+} cation and two $-\text{COO}^-$ groups. Compared Fig. 3(c) and (d), it is observed that the interstitial spaces of PAA hydrogels were filled up after the application of the adsorbent to the uranium solution. These images prove that, $\text{U}(\text{vi})$ ion was adsorbed by PAA hydrogel into its pores and developed a dense structure.

EDX analysis of PAA before and after adsorption were carried out to confirm the adherence of $\text{U}(\text{vi})$ ions onto PAA surface and depicted in Fig. 4. After adsorption, the results showed that the element C, O and N decreased by 6.95 wt%, 2.76 wt% and 1.70 wt% respectively. In addition, Fig. 4(a) shows that there are not U elements existed in the PAA before adsorption, while it can be noticed the appearance of U peaks in EDX spectrum in Fig. 4(b). The U elements accounted for 11.41 wt% in the PAA after adsorption, which probably ascribed to the perfect adsorption capacity of PAA. The occurrence of uranium peak in EDX spectrum confirms the $\text{U}(\text{vi})$ adherence onto PAA.

3.2 Adsorption properties of PAA for the UO_2^{2+} ions

3.2.1 Effect of initial pH on the adsorption. The initial pH of the aqueous solution is an significant parameter in the adsorption process, since pH values affect not only the

speciation of heavy metal ions in solution, but also protonation of the functional groups of the adsorbent.²⁹ The selected pH values in the following experiments were set below 5.5 due to the probability of the hydrolysis and precipitation of UO_2^{2+} cations at higher pH values. Fig. 5 illustrates the effect of pH on the removal efficiency of UO_2^{2+} ions by PAA. As shown in the picture, it is found that the higher adsorption capacity for $\text{U}(\text{vi})$ ions was achieved at higher pH values. However, the adsorption capacity is relatively low for $\text{U}(\text{vi})$ ions at pH = 1.0, this may be for the reason of the leading role of the $-\text{COOH}$ at low pH, and a highly competitive adsorption between H^+ ions and UO_2^{2+} ions in the solution. The adsorption capacity increases with the increase of the solution pH values for the PAA hydrogels when pH range of 1.0–4.0, which attributed to the transformation from some $-\text{COOH}$ groups to $-\text{COO}^-$ groups, leading the electrostatic attraction between PAA hydrogels and UO_2^{2+} cations to strengthen, and accordingly, the adsorption capacity increases. Moreover, the PAA adsorbent can adsorb UO_2^{2+} quite efficiently in the pH range of 4.0–5.5, and the results indicated that the approximately maximum adsorption capacity of 45.89, 78.14, 142.79 mg g^{-1} occurred at an initial $\text{U}(\text{vi})$ concentration of 60, 100 and 200 mg L^{-1} , respectively.

3.2.2 Effect of initial concentration of the $\text{U}(\text{vi})$ ions. Fig. 6 illustrates the effect of the initial concentrations of $\text{U}(\text{vi})$ ions on the adsorption by PAA at 25 °C, which indicates that along with the increasing concentrations of the $\text{U}(\text{vi})$ ions, the adsorption capacities of PAA enhanced sharply at first, and



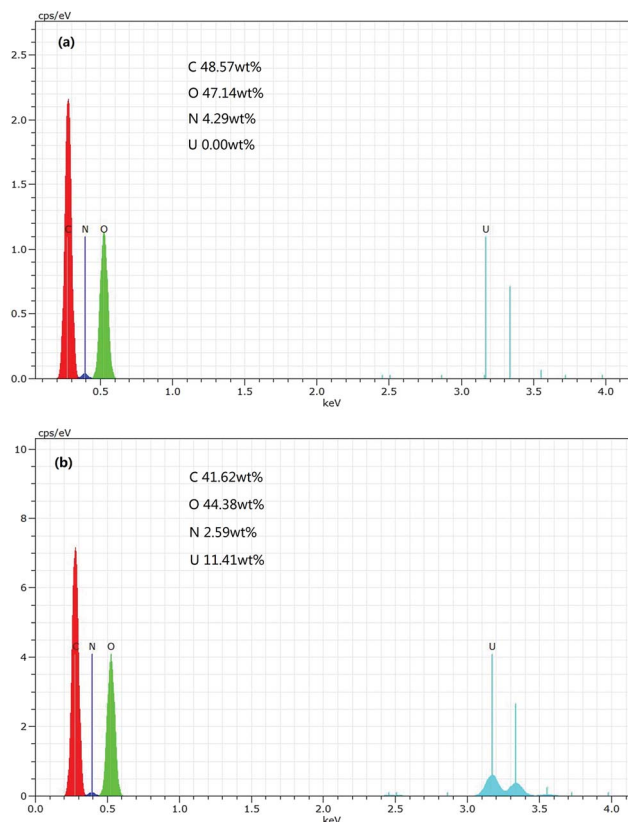


Fig. 4 EDX spectra of PAA. (a) Before adsorption (b) after adsorption.

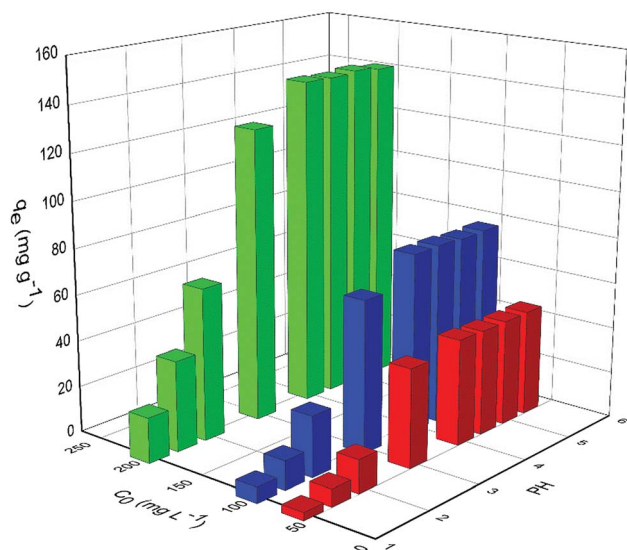


Fig. 5 The effect of initial solution pH on U(VI) removal by PAA hydrogels at various initial concentrations (60, 100 and 200 mg L⁻¹). (Volume, 10 mL; adsorbent dose, 0.01 g; pH value, 1, 1.5, 2, 3, 4, 4.5, 5, 5.5; contact time, 180 min; temperature, 25 °C; rotating speed, 150 rpm). Note: data are presented as the mean \pm standard deviation of triplicate experiments.

then gently rise to reach a plateau, which means PAA could be completely saturated with UO_2^{2+} at enough high initial concentrations.

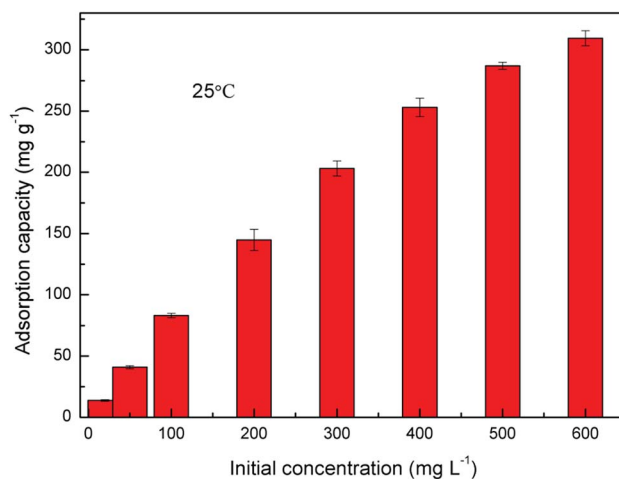


Fig. 6 The effect of initial concentrations (20, 50, 100, 200, 300, 400, 500, 600 mg L⁻¹) on U(VI) removal by PAA hydrogels. (Volume, 10 mL; adsorbent dose, 0.01 g; pH value, 4.0; contact time, 180 min; temperature, 25 °C; rotating speed, 150 rpm.)

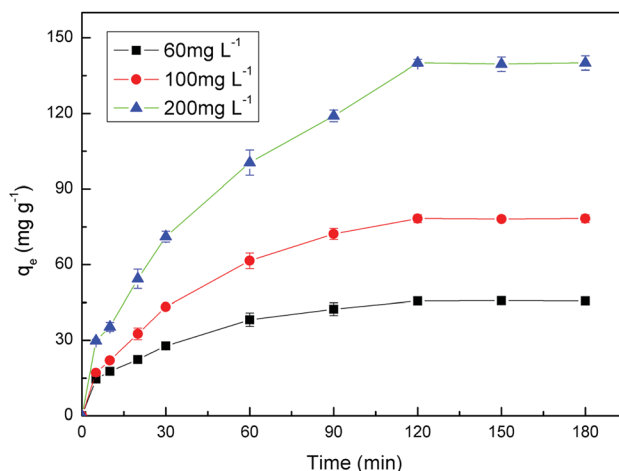


Fig. 7 Effect of contact time on U(VI) adsorption. (Volume, 30 mL; adsorbent dose, 0.03 g; initial concentration, 60, 100 and 200 mg L⁻¹; pH value, 4.0; contact time, 0, 5, 10, 20, 30, 60, 90, 120, 150, 180 min; temperature, 25 °C; rotating speed, 150 rpm.)

3.2.3 Effect of contact time on the adsorption and kinetic studies. U(VI) removal by adsorbent as a function of contact time with different initial concentration (60, 100 and 200 mg L⁻¹) is shown in Fig. 7. It can be seen clearly from the figure that adsorption was rapid originally, and slowed down thereafter, and finally reached equilibrium. The initial U(VI) adsorption was a rapid process and almost reached maximum uptake at 120 min and no more significant uptake of U(VI) was observed at longer time. Therefore, the contact time of 120 min is sufficient to reach equilibrium for PAA hydrogels in the adsorption of UO_2^{2+} ions, which was selected in the further experiments.

Sorption kinetics were investigated to evaluate both the rate of U(VI) sorption and the equilibrium time required for the sorption isotherm. Three adsorption kinetic models,

Table 2 Pseudo-first-order, pseudo-second-order and intra-particle diffusion kinetic model parameters

| Kinetic models | Parameters | 298 K | | |
|--------------------------|--|-----------------------|------------------------|------------------------|
| | | 60 mg L ⁻¹ | 100 mg L ⁻¹ | 200 mg L ⁻¹ |
| Pseudo-first-order model | $k_1 \times 10^{-2}$ | 2.646 | 2.602 | 1.997 |
| | R^2 | 0.991 | 0.981 | 0.982 |
| Pseudo-second-order | q_e (mg g ⁻¹) | 46.8165 | 79.3651 | 143.6782 |
| | $k_2 \times 10^{-3}$ (g mg ⁻¹ min ⁻¹) | 1.165 | 0.4851 | 0.2245 |
| | R^2 | 0.995 | 0.992 | 0.987 |
| Intra-particle diffusion | k_1 | 4.0989 | 8.0955 | 13.0555 |
| | k_2 | 2.3225 | 6.4599 | 12.377 |
| | $k_3 \times 10^{-3}$ | 2.08 | -2.13 | -6.35 |

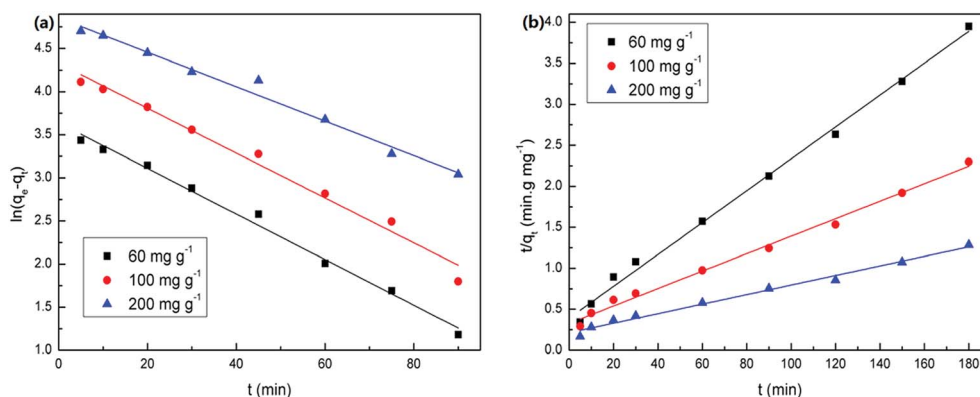


Fig. 8 Sorption kinetic plots of U(vi) ions onto PAA at various initial concentrations. (a) Pseudo-first-order and (b) pseudo-second-order sorption kinetics.

namely pseudo-first-order, pseudo-second-order and intra particle model have been chosen to describe the kinetic data. The pseudo-first-order model is expressed in the following equation:²⁴

$$\ln(q_e - q_t) = \ln q_e - k_1 t \quad (2)$$

The pseudo-second-order model is described in the following equation:³⁰

$$\frac{t}{q_t} = \frac{1}{k_2 q_e^2} + \frac{t}{q_e} \quad (3)$$

where q_t and q_e (mg g⁻¹) stand for the amount of U(vi) ions adsorbed on the adsorbents at time t (min) and equilibrium, respectively. k_1 (min⁻¹) is the rate constant of the pseudo-first-order model, while k_2 (g mg⁻¹ min⁻¹) is the pseudo-second-order rate constant.

The values of corresponding kinetic parameters calculated from the pseudo-first and the pseudo-second-order models are presented in Table 2, and the plots of kinetic model are shown in Fig. 8. The results from Fig. 8(a) and (b) indicate that the pseudo-second order model fits the experiment data better than the pseudo-first order model. Also, the theoretical calculated $q_{e,cal}$ values obtained from the pseudo-second order model are closer to the experimental ones $q_{e,exp}$ and this further confirms the best applicability of the pseudo-second-order model.

In order to investigate if pore or film diffusion was the controlling step in the adsorption, intra-particle diffusion model was further tested as follows:³¹

$$q_t = k_i t^{1/2} + C \quad (4)$$

where k_i (mg g⁻¹ min^{-1/2}) is the intra-particle diffusion rate constant and C is the thickness of the boundary layer and

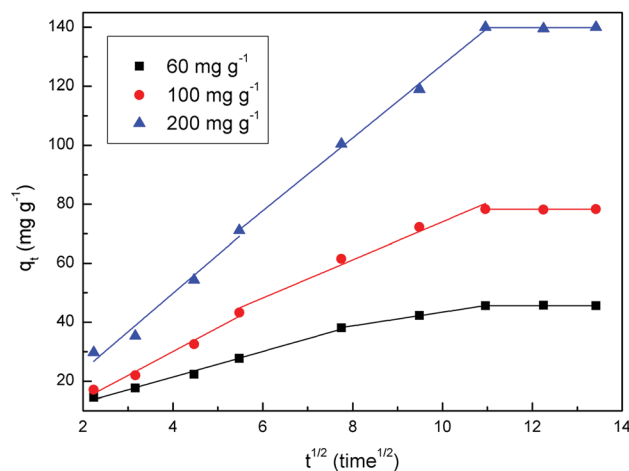


Fig. 9 Intraparticle diffusion kinetics of U(vi) onto PAA at various initial concentrations.



q_t (mg g^{-1}) stands for the amount of metal ions adsorbed on the adsorbents at time t (min). Fig. 9 shows the intra-particle diffusion plots for the U(VI) cations adsorption on PAA. It is apparent that more than one linear portion appear in all of the plots, which indicates that the adsorption is affected by two or more steps.³² Moreover, intra-particle diffusion rate constants in every step follow the order of $k_1 > k_2 > k_3$, which are presented in Table 1. This could be explained as follows: the first portion represents instantaneous adsorption or external surface adsorption stage, in which large numbers of U(VI) ions were adsorbed rapidly by COO^- of the PAA. After almost all the exterior COO^- were occupied, the U(VI) ions started to enter into the pores of the adsorbents and the second portion of the plot seemed to refer to the diffusion into mesopores. Moreover, the third portion of the plots was nearly parallel, suggesting that equilibrium state reached at last.³³

3.2.4 Adsorption isotherm studies. Adsorption isotherm studies are remarkably significant to determine the efficacy of adsorption. Several adsorption isotherms initially used for gas phase adsorption are available and readily adopted to correlate adsorption equilibria in heavy metals adsorption. Some well-known ones are Langmuir, Freundlich, Temkin, Redlich–Paterson and Sips equation.³⁴ In this work, the adsorption isotherms were investigated using three equilibrium models, which are Langmuir, Freundlich and Temkin isotherm models.

The Langmuir isotherm assumes that monolayer adsorption takes place on an adsorbent that have a structurally homogeneous surface, on which the binding sites have the same affinity for the adsorption, and no interaction occurs between adsorbates.³⁵ The Langmuir model can be represented as:

$$q_e = \frac{q_{\max} K_L C_e}{1 + K_L C_e} \quad (5)$$

where C_e is the equilibrium concentration of the adsorbate (mg L^{-1}), q_e is adsorption capacity at equilibrium (mg g^{-1}), q_{\max} is the maximum adsorption capacity of the adsorbent (mg g^{-1}) and K_L is the Langmuir adsorption constant related to the energy of adsorption (L mg^{-1}).

The well-known Freundlich isotherm is an empirical equation predicting that the adsorption occurs on an energetically heterogeneous surface, on which the adsorbed molecules are interactive and adsorption capacity is related to the concentration of adsorbate at equilibrium.³⁶ The Freundlich model is expressed as:

$$q_e = K_F C_e^{\frac{1}{n}} \quad (6)$$

where K_F and n are the Freundlich constants related to adsorption capacity and intensity, respectively, q_e and C_e are the adsorption capacity at equilibrium (mg g^{-1}) and equilibrium concentrations of adsorbate in liquid phases (mg L^{-1}), respectively.

The Temkin isotherm is based on the assumption that the free energy of adsorption is a function of the surface coverage and takes into account the interactions between adsorbents and adsorbates.³⁷ The isotherm model is written as

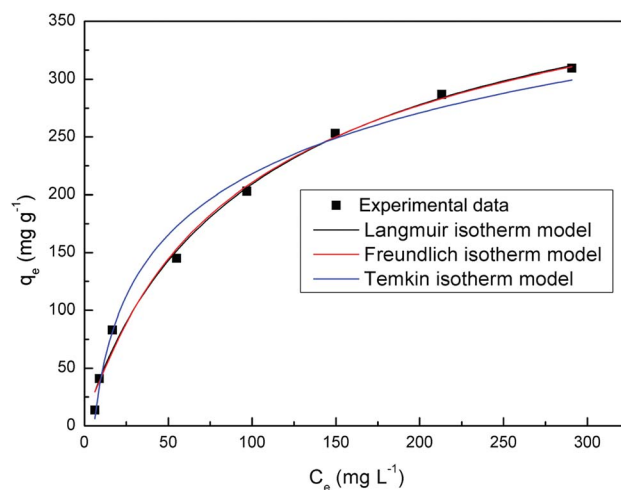


Fig. 10 Non-linear Langmuir, Freundlich and Temkin isotherm for the adsorption of U(VI) ions onto PAA hydrogels. (Volume, 10 mL; adsorbent dose, 0.01 g; initial concentration, 20, 50, 100, 200, 300, 400, 500 and 600 mg L^{-1} ; pH value, 4.0; contact time, 180 min; temperature, 25 °C; rotating speed, 150 rpm.)

$$q_e = \frac{RT}{b_T} \ln(a_T C_e) \quad (7)$$

where R is the ideal gas constant ($8.314 \text{ J mol}^{-1} \text{ K}^{-1}$), T is the absolute temperature (K), a_T is the equilibrium binding constant corresponding to the maximum binding energy and b_T is the Temkin isotherm constant.

Fig. 10 shows the equilibrium adsorption of UO_2^{2+} onto the PAA hydrogels and the fitting plot of the three isotherm models. For the studied system, the Langmuir isotherm correlates best ($R^2 = 0.991$) with the experimental data from adsorption equilibrium of UO_2^{2+} by PAA in these three models, suggesting a monolayer adsorption. Furthermore, the degree of suitability of adsorbent towards U(VI) ions was assessed from the values of separation factor constant (R_L), which has always been used to indicate the adsorption is favorable or not.³⁸ The value of R_L will indicate the type of isotherm is, irreversible ($R_L = 0$), favorable ($0 < R_L < 1$), linear ($R_L = 1$), or unfavorable ($R_L > 1$), which can be calculated from the following equation:

$$R_L = \frac{1}{1 + K_L C_0} \quad (8)$$

where K_L (L mg^{-1}) is the Langmuir constant, and C_0 (mg L^{-1}) is the initial concentration of U(VI) ions. The K_L value obtained from fitting plot of Langmuir isotherm model is 0.0136. And the R_L values of PAA, according to eqn (5), are greater than zero and less than unity, which suggests that the adsorption processes, between the adsorbent (PAA) and UO_2^{2+} , are favorable. The maximum adsorption capacity of PAA for UO_2^{2+} obtained by Langmuir isotherm model is 445.11 mg g^{-1} .

3.2.5 Desorption and reusability studies. From practical point of view, reusability is an very significant feature of an advanced adsorbent. The adsorption behavior of U(VI) onto PAA hydrogels is remarkably reliant on pH and both a lower and higher pH values negatively impact the uranium adsorption. At high pH, a stable precipitation product, $\text{UO}_2(\text{OH})_2$, can be



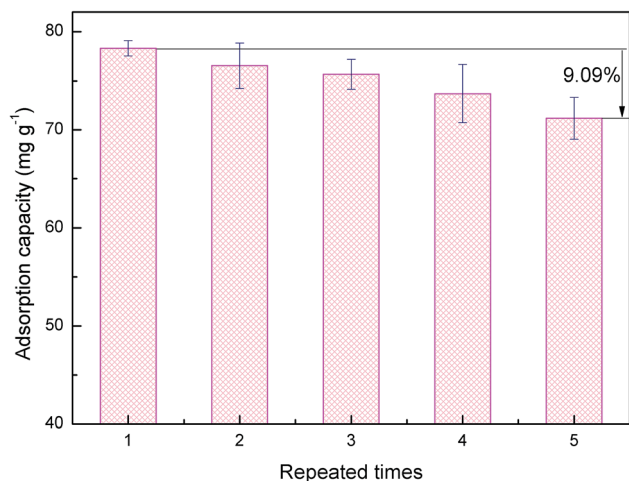


Fig. 11 Adsorption/desorption cycles (initial concentration: 100 mg L⁻¹).

Table 3 Comparison of adsorption capacities for U(vi) ion of various adsorbents

| Adsorbent | Adsorption capacity (mg g ⁻¹) | Reference |
|-------------------------------------|---|-----------|
| HNTs-Fe ₃ O ₄ | 88.32 | 40 |
| AgOH-MWCNTs | 140 | 41 |
| CMPEI/MSU-H | 124 | 42 |
| SA/CMC | 101.76 | 43 |
| SBA-15 | 170 | 44 |
| PAA hydrogels | 445.11 | This work |

formed by the desorbed uranium,³⁹ which was difficult to separate from the adsorbent. Therefore, desorption of the adsorbed U(vi) from PAA can only take place in acid solution (0.1 M HNO₃). The capacity of PAA for five adsorption-desorption cycles is shown in Fig. 11. The PAA hydrogels could be utilized repeatedly and the adsorption capacity only slightly declined (by 9.09%). The regeneration experiments in this study indicate that the PAA hydrogels has the potential of regeneration and can be used as an efficient adsorbents for wastewater treatment.

3.2.6 Comparison with other adsorbents. Table 3 presents a comparison of the U(vi) adsorption capacity of other adsorbents including organic, inorganic materials and composites of them. Among these materials, HNTs-Fe₃O₄, AgOH-MWCNTs, CMPEI/MSU-H and SA/CMC were relatively difficult to prepared due to their complex preparation process. And SBA-15 is low in chemical reactivity. Compared to these adsorbents, PAA hydrogels can be used as efficient adsorbents due to its simple synthesizing procedures and high chemical reactivity. Moreover, PAA hydrogels performed relatively excellent adsorption capacity.

4. Conclusions

In this work, we have demonstrated a clean and extremely simple strategy for preparing PAA hydrogels, which are

identified to be an excellent adsorbent for the removal of UO₂²⁺ ion from aqueous solutions. Structure characterization indicated that the PAA hydrogel was successfully synthesized, complexation took a dominant role in adsorption process and U(vi) ions were adhered onto PAA hydrogels. The adsorption kinetics of PAA hydrogels were well described by pseudo-second-order kinetic and intraparticle diffusion model. Langmuir model was most appropriate to fit the adsorption data, and the maximum sorption capacity of PAA for U(vi) was evaluated to be 445.11 mg g⁻¹ at pH 4.0. According to this paper, its good reusability and excellent adsorption capacity give PAA hydrogels a high potential for practical application in the removal of heavy metal ions from polluted water.

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