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Ultralight aerogel based on molecular-modified poly(*m*-phenylenediamine) crosslinking with polyvinyl alcohol/graphene oxide for flow adsorption†

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Poly(*m*-phenylenediamine) is an excellent adsorbent material. Nevertheless, it is difficult to recover and can even generate secondary pollution due to its powder-like form. Aerogels, monolithic three-dimensional structured materials, that are ultralight and have porous properties are efficient at adsorbing contaminants from water and can solve these problems. Here, we synthesized an aerogel based on molecular-modified poly(*m*-phenylenediamine) (mPmPD) crosslinking with polyvinyl alcohol (PVA) and graphene oxide (GO) (GO/mPmPD/PVA). Of note is that 3-aminophenylboronic acid was introduced into the polymer structure to induce a crosslinking reaction between boric acid units and PVA to constrain poly(*m*-phenylenediamine) in the aerogel. The GO/mPmPD/PVA aerogel shows stable mechanical properties in aqueous solution and an effective adsorption capacity for Ag(I); the maximum Ag(I) adsorption capacity is 917.41 mg g⁻¹. The mechanism of Ag(I) adsorption and reduction was clarified in that Ag(I) chelated with imine units, and a redox reaction occurred between Ag(I) and the benzenoid amine units. Furthermore, the GO/mPmPD/PVA aerogel also shows excellent adsorption ability toward methyl orange and Congo red dyes. This GO/mPmPD/PVA aerogel shows enormous potential for application to silver recovery and dye removal due to its low cost, effective adsorption capacity and facile separation with aqueous solution.

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1 Introduction

Silver, a precious metal, has been widely used in photography, coinage, and electroplating, and much wastewater containing Ag(I) has been discharged as result of these uses.^{1–4} Considering its scarcity, widespread application, and environmental pollution, it is necessary to treat and recover Ag(I) from wastewater. Compared with other technologies, adsorption is considered to be one of the most appropriate ways to treat metal ions, including Ag(I), in wastewater.^{5,6} In recent years, many absorbent materials, including clay minerals,^{7,8} activated carbons,⁹ and conducting polymers,^{5,10–12} have been developed. In particular, conducting polymers are considered to be promising materials for treating metal ions due to their strong chelating ability and redox reversibility.^{12,13}

One type of conducting polymer, polyphenylenediamines, have aroused a great deal of attention due to their superior

conductivity, multifunctionality, high capacity for removal of contaminants and low cost.^{5,14,15} Recently, some reports have suggested that polyphenylenediamines have excellent adsorption ability for Hg(II), Pb(II), Cr(VI), Ag(I), and dyes.^{5,12,14–16} Compared with other polyphenylenediamines, poly(*m*-phenylenediamine) (PmPD) is considered to be an excellent adsorbent because of its simple synthesis process. The polyreaction of poly(*o*-phenylenediamine) is only initiated under harsh conditions, where the reaction temperature is above 100 °C, and poly(*p*-phenylenediamine) dissolves in many common solvents, which limits its application in adsorption. However, PmPD has a small specific surface area, which influences the adsorption capacity, and is difficult to separate from water in the practical environment, even causing secondary pollution because of its powdered form. Hence, developing an environmentally friendly, low cost, and efficient method to solve these problems is needed.

Graphene oxide (GO) has attracted much attention in recent years due to its superior specific surface area.^{17,18} Graphene oxide and reduced graphene oxide (rGO) have been used in many fields, such as catalysis,¹⁹ electrochemical energy storage,^{20,21} and adsorption.⁵ Recently, lots of materials composed of graphene oxide and conducting polymers have

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been synthesized. Poly(*m*-phenylenediamine)/reduced graphene oxide was synthesized by redox reaction for the electro-sorption of Cu(II)²² and for adsorbing dyes,⁵ and polypyrrole/graphene oxide²³ was prepared to remove Cr(VI) from aqueous solution. Furthermore, three-dimensional (3D) structures of graphene oxide and reduced graphene oxide are considered to have more significance in practical applications. Recently, some reports have suggested that aerogels self-assembled by poly-aniline and graphene oxide nanosheets possess significant performances in electrochemical energy storage and catalysis.^{21,24,25} It is noteworthy that the advantages of ultralight weight, high porosity and high specific surface area are also of benefit for adsorption. Therefore, the aerogel is considered to be one of the most promising materials for recovery of metal ions from wastewater and to solve the water pollution problem. However, reduced graphene oxide-based aerogels are not hydrophilic and do not have good enough mechanical properties for application in aqueous solution.

Here, we synthesized an aerogel based on molecular-modified poly(*m*-phenylenediamine) (mPmPD) crosslinking with polyvinyl alcohol (PVA) and graphene oxide (GO). PVA, a flexible and tough polymer material, has already been used in many fields; for instance, in pharmaceuticals and as a flexible capacitor.^{26,27} Therefore, we chose PVA to strengthen the mechanical properties and hydrophilicity of the aerogel, a GO sheet to be the skeleton frame, and mPmPD to be the functional section. Inspired by the crosslinking reaction between boric acid and PVA, we introduced 3-aminophenylboronic acid (ABA) into the polymerization to modify PmPD at a molecular level to induce crosslinking with PVA.²⁸ A typical B 1s peak appears in the wide X-ray photoelectron spectrum of the GO/mPmPD/PVA aerogel, suggesting that boric acid groups exist in the aerogel. The GO/mPmPD/PVA aerogel shows stable mechanical properties in aqueous solution and high adsorption capacity for Ag(I). The maximum Ag(I) adsorption capacity is 917.41 mg g⁻¹. The mechanism of Ag(I) adsorption and reduction is explained by chelation between Ag(I) and imine units and the redox reaction between Ag(I) and benzenoid amine units. Furthermore, the GO/mPmPD/PVA aerogel also shows excellent adsorption ability for methyl orange and Congo red dyes. In a word, this GO/mPmPD/PVA supramolecular-structure aerogel shows

enormous potential in applications for silver recovery and dye removal due to its low cost, high adsorption capacity and facile separation with aqueous solution.

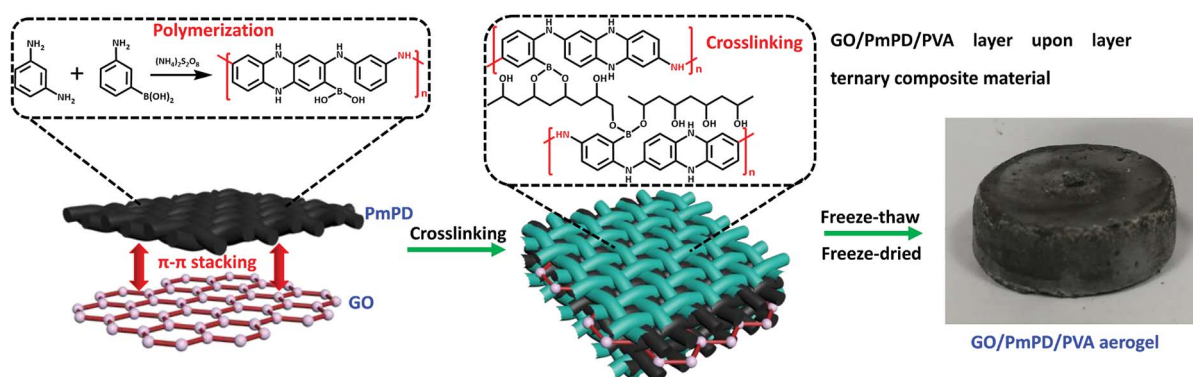
2 Experimental

2.1 Materials

Ammonium persulfate (APS), polyvinyl alcohol (PVA) (MW 1750), graphite powder, hydrogen peroxide (H₂O₂), potassium permanganate (KMnO₄), sulfuric acid (H₂SO₄), phosphoric acid (H₃PO₄), and silver nitrate (AgNO₃) were analytical pure grade and purchased from Shanghai Sinopharm Chemical Reagent Corporation. Methyl orange, Congo red, *m*-phenylenediamine (mPD), and 3-aminophenylboronic acid (ABA) were purchased from Aladdin Industrial Corporation and were analytical pure grade. Ultrapure water (18.2 MΩ cm) was used to prepare aqueous solutions for all experiments.

2.2 Preparation of GO/mPmPD/PVA supramolecular-structure aerogel

The GO/mPmPD/PVA supramolecular-structure aerogel was prepared through a one-step, *in situ* synthesis process. GO was prepared according to the improved Hummers' method,²⁹ and dispersed in ultrapure water and shaken ultrasonically for 3 hours to obtain a 2.5 mg mL⁻¹ uniform GO suspension. A certain amount of mPD and ABA were added to the GO suspension and vigorously stirred for 2 hours at 25 °C. Then, a certain amount of PVA solution (50 mg mL⁻¹) was added into the aqueous solution and stirred for 1 hour at 25 °C. After that, oxidant solution with APS (oxidant : monomer, ~1 : 1) was added to the system drop by drop to initiate the polymerization reaction, and the reaction solution was steadily stirred for 4 hours at 25 °C. The PmPD and GO sheets would be attracted together *via* strong π - π stacking interactions. The mix suspension was transferred to an oven for crosslinking for 2 hours at 65 °C. The mechanism of polymerization and crosslinking is shown in Scheme 1, where the boric acid units in PmPD can induce crosslinking between PmPD and PVA at the molecular scale. After the polymerization reaction and crosslinking, GO/mPmPD/PVA hydrogel samples were formed by freeze-thaw. To investigate the effect of ABA in the process,



Scheme 1 Schematic diagram of the synthesis mechanism of the GO/mPmPD/PVA aerogel.



system reactions with different ABA content were carried out. Photographs of samples after freeze–thaw are shown in Fig. S1,† and the optimal ratio of ABA : mPD is 1 : 9. GO/mPmPD/PVA hydrogel samples were washed with plenty of ultrapure water to remove impurities and then freeze-dried for 48 hours to obtain the final GO/mPmPD/PVA supramolecular-structure aerogels. Aerogels of various GO : mPD/ABA : PVA ratios were synthesized to find the most appropriate ratio. It is noteworthy that the sample cannot form an aerogel when the ratio between mPD/ABA and PVA exceeds 4 : 6. Considering PmPD as the main functional polymer, the aerogel of ratio GO : mPD/ABA : PVA = 1 : 4 : 6 was used in the following experiments.

2.3 Adsorption experiment

A series of Ag(I) adsorption experiments were designed. In brief, 25 mg GO/mPmPD/PVA aerogel was immersed in 20 mL Ag(I) solutions with a series of different concentrations. Tweezers were used to take out the GO/mPmPD/PVA aerogel after the system had been shaken for enough time at a constant temperature. The effect of initial pH, temperature, and adsorption time were also investigated. The adsorbed amount was calculated *via* the following formula:

$$q_e = \frac{(C_0 - C_e)V}{m} \quad (1)$$

where q_e (mg g^{−1}) is the amount of adsorbed Ag(I) at equilibrium, C_0 and C_e (mg L^{−1}) are the concentrations of the Ag(I) solution at the start time and equilibrium time, V (mL) is the volume of the solution and m (mg) is the dry mass of the adsorbent. The concentration of Ag(I) was determined by an inductively coupled plasma emission spectrometer.

GO/mPmPD/PVA aerogel was also used to remove methyl orange and Congo red dyes, and the adsorption properties for methyl orange and Congo red were investigated. Briefly, 10 mg GO/mPmPD/PVA aerogel was put into 20 mL methyl orange solutions with a series of different concentrations. The concentrations of methyl orange in the solutions were determined using a UV-vis spectrophotometer at 464 nm after shaking for 36 hours. The adsorption properties for Congo red

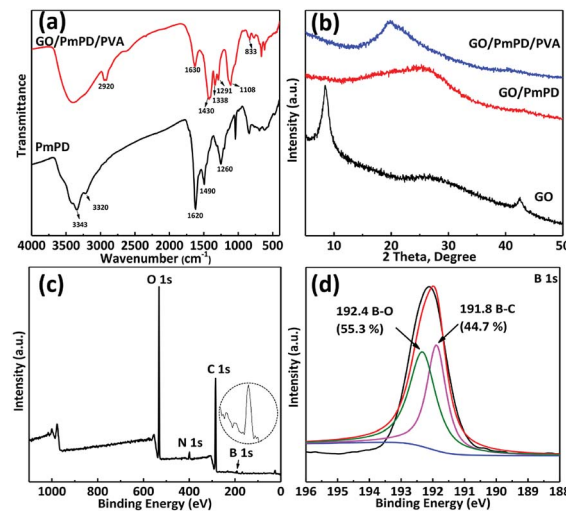


Fig. 2 (a) FT-IR spectra of PmPD and the GO/mPmPD/PVA aerogel. (b) XRD patterns of GO, GO/mPmPD and the GO/mPmPD/PVA aerogel. (c) Wide XPS spectrum of the GO/mPmPD/PVA aerogel. (d) High-resolution spectrum of B 1s.

were determined using the same method as for methyl orange, with the concentrations determined at 497 nm.

2.4 Characterization

The morphology of the GO/mPmPD/PVA aerogel was obtained by scanning electron microscopy (SEM, Sirion 200, FEI Co., USA). The concentration of Ag(I) was determined by inductively coupled plasma emission spectrometry (ICP-ES, Optima 7300 DV, PerkinElmer Inc., Electron Co., USA). The concentration of methyl orange and Congo red were determined using a UV-vis spectrophotometer (UV-2550, Shimadzu Co., Japan). The X-ray diffraction patterns of GO, GO/mPmPD, and GO/mPmPD/PVA aerogel were determined using CuK α radiation and an X-ray diffractometer (XRD, X'Pert PRO, PANalytical B.V., The Netherlands). Fourier-transform-infrared (FT-IR) spectra were measured using an infrared spectrophotometer (FT-IR, Nicolet 8700, Thermo Scientific Instrument Co., USA). X-ray photoelectron spectra were recorded using an X-ray photoelectron

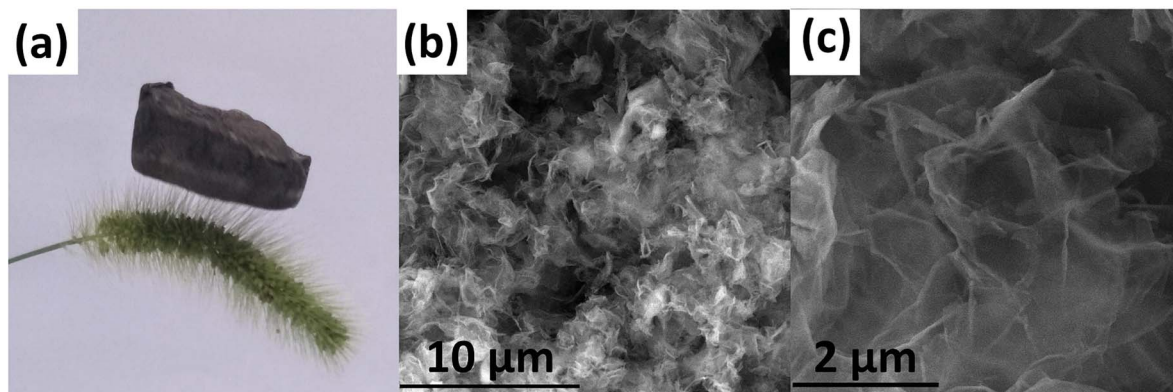


Fig. 1 (a) Photograph of GO/mPmPD/PVA aerogel, (b) and (c) SEM images of the GO/mPmPD/PVA aerogel.



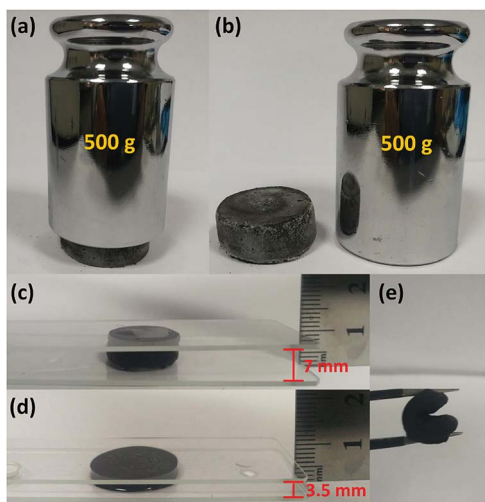


Fig. 3 (a and b) Photographs of the strong GO/PmPD/PVA aerogel with supporting weight. (c–e) Photographs of the GO/PmPD/PVA aerogel with stable mechanical properties after shaking for 36 hours in aqueous solution.

spectrometer (XPS, ESCALAB 250, Thermo-VG Scientific Co., USA). The pore size, pore volume and surface area were measured by accelerated surface area and porosimetry (Tristar II 3020M, Micromeritics Instrument Co., USA). Thermogravimetric analysis (TGA) was carried out using a thermoanalyser at a heating rate of $10\text{ }^{\circ}\text{C min}^{-1}$ under an air atmosphere flow (TGA Q5000IR, TA Instrument Co., USA).

3 Results and discussion

3.1 Characterization of GO/mPmPD/PVA aerogel

As shown in Fig. 1(a), the prepared ultralight GO/mPmPD/PVA aerogel can be supported on the slim fibers of *Setaria viridis*, and the fibers do not even deform under the pressure. SEM images of the aerogel with a ratio of 1 : 4 : 6 between GO, mPD/ABA, and PVA are presented in Fig. 1(b) and (c). The morphologies of other samples with various ratios of GO, mPD/ABA, and

PVA are shown in Fig. S2.† In the SEM images in Fig. 1(b) and (c), a lamellar structure is evident with many spaces in the GO/mPmPD/PVA aerogel. This may be attributed to the GO skeleton and crosslinking between PVA and PmPD. The pore size, pore volume, and surface area of PmPD and GO/mPmPD/PVA aerogels are given in Table S1.† The surface area of the GO/mPmPD/PVA aerogel is $23.49\text{ m}^2\text{ g}^{-1}$, which is much higher than the surface area of PmPD ($8.36\text{ m}^2\text{ g}^{-1}$). This higher surface area is beneficial for adsorption.³⁰

The FT-IR spectra of PmPD and the GO/mPmPD/PVA aerogel are shown in Fig. 2(a). For comparison, the spectra of GO sheets, PVA and GO/PmPD are presented in Fig. S3.† As shown in the spectrum of PmPD, the broad bands at 3343 cm^{-1} and 3220 cm^{-1} correspond to the $-\text{NH}-$ stretching vibration. The sharp peak at 1620 cm^{-1} can be associated with the quinoid imine and the peak at 1490 cm^{-1} can be associated with the benzenoid amine. The peak at 1257 cm^{-1} can be assigned to the stretching vibration of C–N in benzenoid amine groups.³¹ Quinoid imine and benzenoid amine groups are the most important functional groups for adsorption. In the spectrum of the GO/mPmPD/PVA aerogel, the peak at 1430 cm^{-1} can be associated with the amine of PmPD. The peak at 1291 cm^{-1} can be assigned to B–O.³² The peak around 2920 cm^{-1} can be assigned to the C–H of PVA.²⁶ The peak at 1108 cm^{-1} can be assigned to the C–O of GO and PVA, and the peak at 1630 cm^{-1} is associated with the C=O of GO and the quinoid imine of PmPD.^{26,31,33}

The XRD patterns of GO, GO/PmPD and GO/mPmPD/PVA aerogel are shown in Fig. 2(b). In the pattern for GO, the typical diffraction peak at about 10° can be assigned to the interlayer space in GO.³⁴ In the pattern for GO/PmPD, this diffraction peak at 10° disappeared and a new broad diffraction peak appeared at about 25° , suggesting the amorphous nature of PmPD.⁵ Compared with GO/PmPD, the broad diffraction peak in the GO/mPmPD/PVA aerogel shifted to about 20° , after crosslinking between boric acid groups and PVA. As noted in some previous reports,^{14,31} a shift in the diffraction peak of polymers to a low angle signifies an increase in their amorphous nature. Due to the disordered and loose-knit piles of the

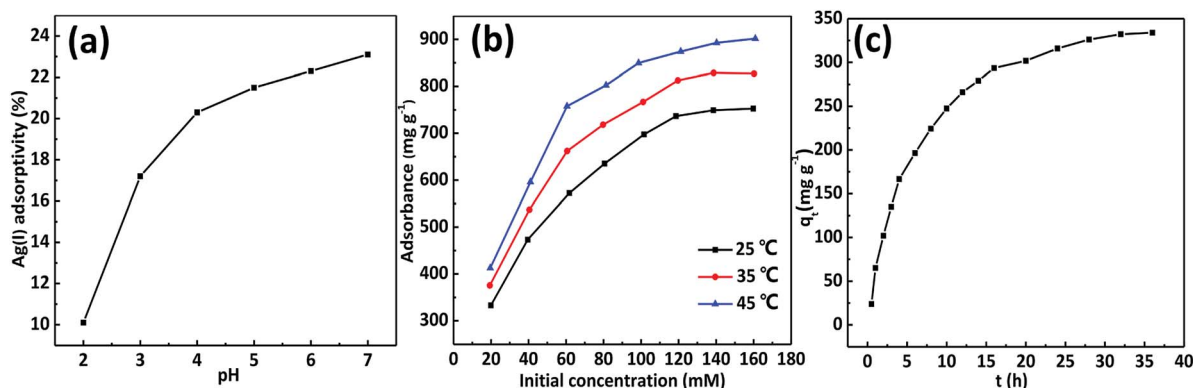


Fig. 4 (a) Effect of pH on Ag(I) adsorption by the GO/PmPD/PVA aerogel (temperature, $25\text{ }^{\circ}\text{C}$; initial concentration, 20 mM ; dose, 25 mg ; volume, 20 mL ; adsorption time, 36 h). (b) Effects of initial concentration and temperature on Ag(I) adsorption (dose, 25 mg ; volume, 20 mL pH, 7.0 ; adsorption time, 36 h). (c) Effect of contact time on Ag(I) adsorption (dose, 25 mg ; volume, 20 mL ; pH, 7.0 , temperature, $25\text{ }^{\circ}\text{C}$; concentration, 20 mM).



Table 1 Isotherm model equations for Ag(I) adsorption

Temperature, T (°C)	Langmuir model			Freundlich model		
	q_{\max} (mg \times g $^{-1}$)	K (L \times mg $^{-1}$)	R^2	K_f	n	R^2
25	917.413	0.00031	0.997	21.657	2.678	0.976

supramolecular chains, this amorphous structure is suitable for permeation and adsorption of ions onto the GO/mPmPD/PVA aerogel.

In order to further confirm the existence of boric acid groups, XPS was used to characterize the GO/mPmPD/PVA aerogel. As shown in the XPS results (Fig. 2(c)), the typical B 1s peak appearing in the wide spectrum of the GO/mPmPD/PVA aerogel means that ABA has successfully participated in the reaction system. In the high-resolution B 1s spectrum (Fig. 2(d)), the peak at 191.8 eV corresponds to B–C and the peak at 192.4 eV can be assigned to B–O.³² This suggests that the boric acid group plays a bridge-like role between PmPD and PVA. The TGA curve of for the GO/mPmPD/PVA aerogel is shown in Fig. S4,† and the result suggests that the GO/mPmPD/PVA aerogel has good thermostability for practical applications.

As shown in Fig. 3(a) and (b), the GO/mPmPD/PVA aerogel can support a weight of 500 g, which is about 2500 times its own weight. It is obvious that GO/mPmPD/PVA aerogel has strong mechanical properties. Fig. 3(c)–(e) show the stable mechanical properties after shaking at 300 rpm for 36 hours in aqueous solution. In Fig. 3(c) and (d), the GO/mPmPD/PVA aerogel is still intact and a lot of water was squeezed out after hard pressing. We found that the GO/mPmPD/PVA aerogel can bend and fold without any breakages (see Fig. 3(e)), which suggests that the aerogel is very tough. In a word, GO/mPmPD/PVA aerogel has excellent mechanical properties and hydrophilicity for practical applications in aqueous solution.

3.2 Properties of Ag(I) adsorption

As presented in Fig. 4(a), the pH was controlled from 2.0 to 7.0 by HCl to investigate the effect of pH for Ag(I) adsorption by the GO/mPmPD/PVA aerogel. Obviously, the adsorptivity of Ag(I) in a strong acid environment is much lower than in a neutral environment. More specifically, the adsorptivity of Ag(I) decreases rapidly, from 20.3% to 10.1%, when the pH is reduced from 4.0 to 2.0. On the other hand, the adsorptivity of Ag(I) only increases by 2.8% with a rise in pH from 4.0 to 7.0. This means that a weak acid or neutral environment is more suitable for adsorbing Ag(I) using the GO/mPmPD/PVA aerogel.

The effect of the initial Ag(I) concentration and temperature for adsorption of Ag(I) are shown in Fig. 4(b). As the initial Ag(I)

concentration increases, the adsorption capacity of Ag(I) increases, as a result of the increase in contact between Ag(I) and active sites in the GO/mPmPD/PVA aerogel. The adsorption capacity tends toward a stabilized value when the initial Ag(I) concentration is higher than 120 mM. Meanwhile, the adsorption capacity of Ag(I) rises from 752.67 mg g $^{-1}$ to 901.44 mg g $^{-1}$ as the temperature rises from 25 °C to 45 °C, indicating that high temperature is beneficial for Ag(I) adsorption. In order to achieve the maximum adsorption capacity of Ag(I) at room temperature and to figure out how the adsorbates interact with the adsorbents, a Langmuir model and a Freundlich model were used to analyze the adsorption isotherm, and these are shown in Fig. S5.† The relative parameters of the models are listed in Table 1. The experimental data fit better with the Langmuir model through comparing the correlation coefficients (R^2), 0.997 for the Langmuir model and 0.976 for the Freundlich model, which indicates adsorption of Ag(I) in a monolayer. To prove the effective adsorption capacity GO/mPmPD/PVA aerogel for adsorption of Ag(I), a comparison between GO/mPmPD/PVA aerogel and other adsorbents is shown in Table S2.†

The effect of contact time for Ag(I) adsorption is shown in Fig. 4(c). The adsorption capacity of Ag(I) increases rapidly in the initial 15 hours, then gradually slows down and tends to become stable. Moreover, the behavior of Ag(I) adsorption on the GO/mPmPD/PVA aerogel was analyzed using two kinetic equations and is presented in Fig. S6.† The fitting results for the two equations are shown in Table 2. The correlation coefficient of the pseudo-second-order kinetic equation (0.996) is higher than the correlation coefficient of the pseudo-first-order equation (0.960), suggesting that the Ag(I) ions are adsorbed mainly by chemisorption.

3.3 Adsorption mechanism of Ag(I) with the GO/mPmPD/PVA aerogel

In order to clarify the adsorption mechanism of the Ag(I), XPS was used to characterize the GO/mPmPD/PVA aerogel before and after adsorption of Ag(I), and the spectra are presented in Fig. 5. After adsorption, an Ag 3d peak appeared in the wide spectrum verifying that Ag has been adsorbed onto the GO/mPmPD/PVA aerogel. Furthermore, we found two peaks for Ag

Table 2 Summary of kinetic parameters for Ag(I) adsorption

Temperature, T (°C)	Pseudo-first-order			Pseudo-second-order		
	q_e (mg \times g $^{-1}$)	K_1 (g \times mg $^{-1}$ \times L $^{-1}$)	R^2	q_e (mg \times g $^{-1}$)	K_1 (g \times mg $^{-1}$ \times L $^{-1}$)	R^2
25	336.127	0.00231	0.960	395.257	6.4×10^{-6}	0.996



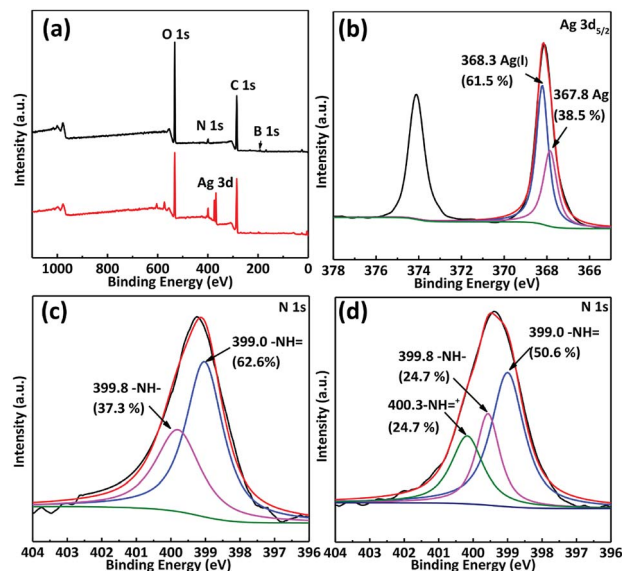
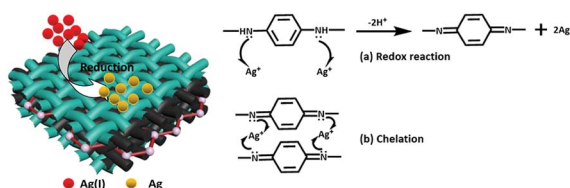


Fig. 5 (a) Wide XPS spectra of the GO/mPmPD/PVA aerogel before and after adsorption of Ag(I). (b) Ag 3d_{5/2} high-resolution spectrum. (c) N 1s high-resolution spectrum before adsorption of Ag(I). (d) N 1s high-resolution spectrum after adsorption of Ag(I).

at 367.8 eV and 368.3 eV in the high-resolution Ag 3d_{5/2} spectrum, which are assigned to Ag (38.5%) and Ag(I) (61.5%), respectively. This means that a portion of the Ag(I) ions has been reduced to Ag by the GO/mPmPD/PVA aerogel. Correspondingly, the proportion of benzenoid amine (–NH–) at 399.8 eV decreased from 37.3% to 24.7% after adsorption of Ag(I), as shown in Fig. 5(c) and (d). Simultaneously, a new peak (–NH=+) emerged at 400.3 eV, which can be attributed to chelation between the quinoid imine (–NH=) and Ag(I).³¹ Therefore, the mechanism of Ag(I) adsorption may be explained by the following three features. (1) Part of the Ag(I) is reduced to Ag and part of the benzenoid amine (–NH–) is transformed to the quinoid imine (–NH=) during a redox reaction. (2) A number of Ag(I) ions chelate with newly emerging and pre-existing quinoid imine (–NH=) to form –NH=+.³¹ (3) A small amount of the Ag(I) ions are adsorbed onto the GO/mPmPD/PVA aerogel through physical adsorption and through oxygen-containing functional groups such as hydroxyl and carboxyl.^{26,33} The redox reaction and chelation are shown in Scheme 2.

3.4 Properties of dye adsorption

The adsorption properties of GO/mPmPD/PVA aerogel with respect to dyes are shown in Fig. 6. The concentrations of



Scheme 2 Schematic illustration of the redox reaction and chelation during Ag(I) adsorption.

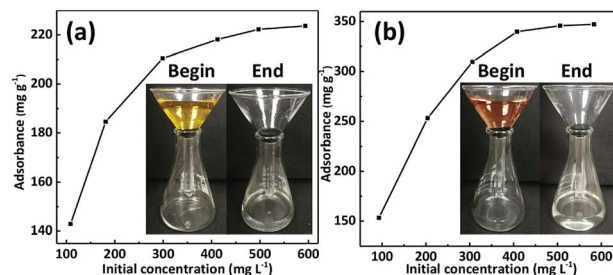


Fig. 6 Adsorption properties of the GO/mPmPD/PVA aerogel for (a) methyl orange and (b) Congo red (temperature, 25 °C; dose, 10 mg; volume, 20 mL; pH, 7.0; adsorption time, 36 h). Insets: methyl orange solution and Congo red solution changed into clean water after flowing through the GO/mPmPD/PVA aerogel (pH, 7.0; initial concentration, 10 mg L^{−1}).

methyl orange solution and Congo red solution were determined using a UV-vis spectrophotometer at 464 nm and 497 nm, respectively. The adsorbed amounts for both methyl orange and Congo red increase rapidly with increase in the initial concentration, when the initial concentrations are below 300 mg L^{−1}. When the initial concentrations exceed 300 mg L^{−1}, the amounts adsorbed increase slowly and tend to become stable. The adsorption capacities of methyl orange and Congo red can reach 223.66 mg g^{−1} and 347.19 mg g^{−1}, respectively. Practical experiments for filtration/adsorption of dyes were carried out and are shown in the insets in Fig. 6. We crammed the GO/mPmPD/PVA aerogels into the bottoms of the funnels and added solutions containing the dyes into the funnels. The colors of the solutions disappeared and became clean water after flowing through the GO/mPmPD/PVA aerogel, which suggests that the dyes were adsorbed by the aerogel. In summary, GO/mPmPD/PVA aerogel is not only able to easily recover Ag(I), but also able to remove dyes.

4 Conclusions

In this study, we synthesized GO/mPmPD/PVA, a supramolecular-structure aerogel, through *in situ* polymerization and chemical crosslinking. Inspired by the crosslinking reaction between boric acid and PVA, 3-aminophenylboronic acid was introduced into the system for chemical crosslinking. A typical B 1s peak appeared in the wide XPS spectrum of the GO/mPmPD/PVA aerogel suggesting that boric acid groups existed in the aerogel. The GO/mPmPD/PVA aerogel showed excellent hydrophilicity and great mechanical properties, which are of benefit for practical applications. The GO/mPmPD/PVA aerogel can adsorb Ag(I) with high efficiency, and the maximum Ag(I) adsorption capacity is 917.41 mg g^{−1}. The Ag(I) adsorption isotherm fits better with the Langmuir isotherm model and the kinetics agree with the pseudo-second-order model. Chelation between Ag(I) and imine units and redox reactions between Ag(I) and benzenoid amine units are the main methods of Ag(I) adsorption. Furthermore, the GO/mPmPD/PVA aerogel also shows excellent adsorption ability for methyl orange and Congo red dyes. Practical experiments for the filtration/adsorption of



dyes showed that solutions containing dyes become colorless after flowing through the GO/mPmPD/PVA aerogel. The GO/mPmPD/PVA supramolecular-structure aerogel shows enormous potential in applications for silver recovery and dye removal due to its effective adsorption capacity, low cost and facile separation with aqueous solution.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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