RSC Advances



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Cite this: RSC Adv., 2017, 7, 30109

Received 6th May 2017 Accepted 30th May 2017 DOI: 10.1039/c7ra05114j

rsc.li/rsc-advances

1. Introduction

Protein is an important material basis of life and is closely linked to various forms of life activity.¹ As protein is the direct manifestation of life phenomena, the analysis of protein is of great significance in the field of life science. However, protein is usually present in a complex sample matrix, which causes significant interference in its detection.² Moreover, the disturbance of other different proteins is a challenge that particularly needs to be addressed. Consequently, sample clean-up, separation and enrichment are necessary before protein analysis.

Recently, magnetic separation has attracted widespread attention due to its unique advantages.^{3–5} Compared to conventional solid-phase extraction (SPE), the magnetic materials do not need to be packed into the cartridge, which avoids the problem of column clogging encountered in SPE. In addition, the phase separation of magnetic materials is much more conveniently performed by applying an external magnetic field,⁶ so that magnetic adsorbents can be easily recycled.^{7,8} Furthermore, the magnetic nanocomposites have a large specific surface area, so extraction and separation can be attained with only a small amount of adsorbents and a shorter equilibration time, improving the extraction capacity and efficiency.^{9,10}

Amino acid-modified graphene oxide magnetic nanocomposite for the magnetic separation of proteins[†]

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A novel amino acid-modified graphene oxide magnetic nanocomposite was successfully synthesized for the magnetic separation of proteins. The extraction capacity was influenced by several parameters including the type of amino acid, the solution pH value, the addition of electrolyte, the extraction time and the protein concentration. The nanocomposite showed excellent extraction capacity for bovine hemoglobin (868.3 mg g⁻¹). By adsorbing bovine hemoglobin, bovine serum albumin, ovalbumin and lysozyme, the nanocomposite had a better performance than magnetic graphene oxide in the extraction of bovine hemoglobin and lysozyme, owing to the intensive electrostatic interaction. The proposed nanocomposite could be easily recycled and reused more than three times. Moreover, the proposed method was successfully applied to selectively extract bovine hemoglobin in acid protein mixtures and real samples.

cost technique that requires no cumbersome operation and offers high extraction efficiency, and is widely used in sample pretreatment procedures.¹¹⁻¹³

Graphene (G) is a novel carbon-based material with twodimensional properties.¹⁴ It has attracted a great amount of research interest because of its physical and chemical properties such as extraordinary electrical conductivity, and excellent thermal and mechanical qualities.¹⁵⁻¹⁸ In addition, graphene has been widely used in sample pretreatment owing to its large specific surface area, good dispersibility and hydrophobicity.19,20 Graphene oxide (GO), the oxidation product of graphene, is considered to be an ideal adsorbent material in analytical chemistry.²¹ On one hand, GO still maintains the basic framework of graphene, which means it has a huge surface area. On the other hand, there are many oxygenated groups on the surface of GO, such as epoxy, hydroxyl and carboxyl, making GO more hydrophilic.22 Furthermore, the functional groups can form hydrogen bonding and electrostatic interactions with adsorbates, and can tune the adsorption properties of specific adsorbates via functional modification.23 Therefore, GO has potential for sample pretreatment. However, it is difficult to separate GO from aqueous solution, which limits its direct use in sample pretreatment. To overcome this drawback, graphene hybrid materials have become the research focus. Magnetic GO possesses high adsorption capacity and can be easily separated by an external magnetic field, which indicates good potential in magnetic separation applications.²⁴ Recently, magnetic GO has been applied as the sorbent for organic compounds,²⁵ heavy metal ions,²⁶ phosphopeptides²⁷ and proteins.²⁸ Moreover, Ding et al. modified the functional guanidinium ionic liquids on GO to synthesize a magnetic

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[†] Electronic supplementary information (ESI) available. See DOI: 10.1039/c7ra05114j

composite material, which improved the extraction capacity of proteins.²⁹ Modification of the oxygenated groups on GO seems to be a good approach to enhance the extraction efficiency of proteins.

Amino acids, the basic building units of protein, exhibit good biocompatibility.³⁰ In addition, they possess functional groups such as amino, carboxyl, phenyl and sulfhydryl groups which can form hydrogen bonding, electrostatic and π - π interactions with proteins. Modification of the amino acids on GO can improve the biocompatibility and adsorbability of the adsorbents. However, modification of amino acids on GO as the absorbent for proteins has rarely been reported.

In this paper, amino acid-modified graphene oxide magnetic nanocomposite (AMGO@Fe₃O₄) was synthesized for the magnetic separation of proteins. We have chosen four kinds of amino acid with different functional groups to modify the surface of GO. Bovine hemoglobin (BHb) was selected as the representative protein to explore the adsorption capacity. After adsorption, the concentration of protein in the supernatant was measured by a UV-vis spectrophotometer. The parameters influencing extraction efficiency, including pH value, addition of electrolyte, extraction time and protein concentration, were optimized in detail. AMGO@Fe₃O₄ has also been used to explore extraction selectivity, and the results indicated that it could selectively extract BHb in an acidic protein mixture and in real samples.

2. Materials and methods

2.1 Materials

Expanded graphite powder (100 mesh) was supplied by the Xinghe Graphite Co., Ltd (Qingdao, China). Amino acids were obtained from Beijing Chemical Reagent Co. (Beijing, China). 1-Ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride (EDC) was purchased from J&K Chemical Technology Co., Ltd (Beijing, China). Ferric chloride and ferrous chloride were bought from Xilong Chemical Co., Ltd (Guangdong, China). Ammonia solution (25%) was acquired from Beijing Chemical Works (Beijing, China). Proteins including bovine hemoglobin (BHb), bovine serum albumin (BSA) and lysozyme (Lyz) were obtained from Beijing Keao Biological Pharmaceutical Co., Ltd (Beijing, China). Ovalbumin (OVA) was from Beijing Century Hualin Biotechnology Co., Ltd (Beijing, China). Bovine plasma was purchased from Beijing Keao Biological Pharmaceutical Co., Ltd (Beijing, China). All other reagents used in this study were analytical grade, and deionized water was used throughout the work.

2.2 Preparation of GO

GO was synthesized from expanded graphite powder following a modified Hummers' method.³¹ About 18 g of potassium permanganate was added into a mixture of frozen concentrated H_2SO_4 (360 mL) and H_3PO_4 (40 mL), then 3 g of expanded graphite powder was slowly added with vigorous agitation to avoid a sudden increase in temperature. The reaction was maintained at 50 °C for 12 h. After this, the mixture was transferred into an ice–water mixture with the same volume, and $30\% H_2O_2$ was added dropwise until the color of the slurry turned to golden yellow. The slurry was washed with 10% HCl and deionized water and centrifuged to remove the supernatant. Then the product was purified by dialysis for 1 week to remove the remaining metal species. After the dialysis, the GO was ultrasonicated for 2 h to obtain an aqueous dispersion (10 mg mL⁻¹).

2.3 Preparation of AMGO

EDC (0.25 g) was added into 10 mL GO aqueous dispersion (10 mg mL⁻¹) under agitation. After stirring for 30 min, different amino acids (0.3 g each) including L-arginine (Arg), glutamic acid (Glu), phenylalanine (Phe) and cysteine (Cys) were added to separate GO/EDC solutions. The mixtures were stirred for 24 h at room temperature. Then the amino acid-modified GO materials (AMGO) were washed with deionized water and centrifuged 5 times to remove the unreacted amino acids. Finally, the materials were dried in oven at 60 °C for 12 h.

2.4 Preparation of AMGO@Fe₃O₄

AMGO@Fe₃O₄ was synthesized by the chemical coprecipitation method.²⁶ The synthesized AMGO was dispersed in deionized water and ultrasonicated for 6 h to obtain an aqueous dispersion of AMGO (1 mg mL⁻¹). Then, 250 mg of ferric chloride and 500 mg of ferrous chloride were added slowly to 50 mL of the AMGO dispersion at 50 °C under N₂. Afterwards, 2 mL of ammonia solution was added slowly for synthesis of magnetite particles, and the reaction was maintained under vigorous stirring for 30 min. Then the mixture was subjected to an external magnetic field to separate the AMGO@Fe₃O₄. Finally, the AMGO@Fe₃O₄ was washed with deionized water 5 times and dispersed in deionized water (2 mg mL⁻¹).

2.5 Magnetic separation procedures

Magnetic separation experiments were performed using the AMGO@Fe₃O₄ as the adsorbent and BHb as the target protein to examine the extraction capacity. A 1 mL portion of the homogenized adsorbent solution (2 mg mL⁻¹) was placed in a centrifuge tube. The adsorbent was isolated using an external magnetic field to remove the supernatant. Then 2 mL of aqueous protein solution was transferred into the centrifuge tube and the mixture was shaken at room temperature for 1 h. Afterwards, AMGO@Fe₃O₄ was magnetically separated by an external magnetic field. The concentration of protein in the supernatant was measured by a UV-vis spectrophotometer at the maximum absorption wavelength.

2.6 Characterization of graphene nanocomposites

Fourier transform infrared spectra were recorded in the range 4000–500 cm⁻¹ on a Bruker Fourier Transform Infrared Spectrometer (Horiba, Germany). The magnetic properties were measured with a 7307 Vibrating Sample Magnetometer (Lake-Shore, USA). X-ray photoelectron spectroscopy data were evaluated using a PHI-5300 ESCA X-ray photoelectron spectrometer

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(PHI, USA). Transmission electron microscopy images were taken with an H-7650B transmission electron microscope (Hitachi, Japan). Ultraviolet visible detection was carried out on a U-3010 UV-vis spectrophotometer (Hitachi, Japan). Zeta potential data were measured using a SZ-100 Nanoparticle Analyzer (Horiba, Japan).

3. Results and discussion

3.1 Characterization of AMGO@Fe₃O₄

The AMGO@Fe₃O₄ synthesis procedure is illustrated schematically in Fig. 1. First, GO was prepared by the modified Hummers' method, and then the oxygenated groups including epoxy, hydroxyl and carboxyl were introduced to the surface of graphene. Second, the amide reaction occurred between the amine groups of amino acids and the carboxyl groups of GO, introducing amino acids to the surface of GO. Elemental analysis results showed that the content of nitrogen increased from 0.58% to 8.21%, and the content of carbon decreased from 44.25% to 38.91% after modification. Third, the AMGO@Fe₃O₄ was synthesized by the chemical coprecipitation method. The iron ions (Fe²⁺ and Fe³⁺) were seized on the surface of the AMGO sheets and then the Fe₃O₄ precipitate formed in the presence of ammonia solution. Finally, the nanocomposites we required were obtained.



Fig. 1 Schematic illustration of the synthesis of AMGO@Fe $_3O_4$. AMGO, amino acid-modified graphene oxide; EDC, 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride.

We chose the glutamic acid modified magnetic composite AMGO-Glu@Fe $_3O_4$ as the model to study the physical and chemical properties.

3.1.1 Fourier transform infrared spectrometry. FTIR was carried out to identify the successful modification of amino acids and magnetic particles and the results are shown in Fig. S1.[†] For the FTIR spectrum of GO, the peaks at 1719.5 cm⁻¹ and 1618.2 cm^{-1} were attributed to C=O and C=C stretching vibrations, respectively, indicating the successful oxidation process.30 For the FTIR spectra of AMGO-Glu and AMGO-Glu@Fe₃O₄, the peaks at 1617.3 cm⁻¹ and 1380.9 cm⁻¹ were assigned to the C=O stretching vibration and C-N bending vibration of -NHCO-, respectively.²⁹ The peak at 1576.8 cm⁻¹ corresponded to the N-H bending vibration. The results indicated that the amino acids had been covalently grafted to the surface of GO. In addition, both the GO@Fe₃O₄ and AMGO-Glu@Fe₃O₄ presented a sharp peak at 578.6 cm⁻¹ attributed to the Fe-O stretching vibration. This proved the existence of Fe₃O₄.

3.1.2 Vibrating sample magnetometer (VSM). A VSM was used to study the magnetic properties of the synthesized materials at room temperature. The magnetization curves of $GO@Fe_3O_4$ and AMGO-Glu@Fe_3O_4 are shown in Fig. 2A. The saturation magnetization values of $GO@Fe_3O_4$ and AMGO-Glu@Fe_3O_4 and AMGO-Glu@Fe_3O_4 were 45.77 emu g⁻¹ and 60.56 emu g⁻¹, respectively. This may be because the negative charge of AMGO-Glu increased after modification, and more iron ions were seized on the surface of the AMGO-Glu sheets, which increased the load amount of Fe_3O_4. Both GO@Fe_3O_4 and AMGO-Glu@Fe_3O_4 exhibited superparamagnetic properties according to the S-like magnetic hysteresis loops. Fig. 2B illustrates that AMGO-Glu@Fe_3O_4 could be quickly separated from the dispersion when an external magnetic field was applied, which is important for further research.

3.1.3 X-ray photoelectron spectroscopy. XPS spectra were used to further investigate the presence of functional groups on AMGO@Fe₃O₄. As shown in Fig. 3A, all of the materials showed the same peaks at approximately 285 and 530 eV, which were attributed to C 1s and O 1s, respectively.⁵ The peaks at about 710 eV for GO@Fe₃O₄ and AMGO-Glu@Fe₃O₄ were consistent with the characteristic peaks of Fe 2p. In Fig. 3D, the peaks at 711.3 eV and 724.8 eV, which corresponded to Fe $2p_{2/3}$ and Fe



Fig. 2 (A) Magnetic hysteresis loops of GO@Fe₃O₄ (a) and AMGO-Glu@Fe₃O₄ (b), and (B) response of AMGO-Glu@Fe₃O₄ to an external magnetic field.



Fig. 3 XPS spectra of (A) wide scan of GO (a), GO@Fe₃O₄ (b) and AMGO-Glu@Fe₃O₄ (c), (B) C 1s, (C) N 1s, and (D) Fe 2p spectra of AMGO-Glu@Fe₃O₄.

 $2p_{1/2}$, indicated the formation of the Fe₃O₄ phase in the composites.²⁶ The peak at about 400 eV for AMGO-Glu@Fe₃O₄ (Fig. 3A and C) was attributed to N 1s, illustrating the existence of nitrogen from amino acids. The C 1s spectra of AMGO-Glu@Fe₃O₄ (Fig. 3B) could be deconvoluted into four fitting curves with peaks located at about 284.8, 285.6, 286.6, and 288.8 eV, assigned to C-C, C-N, C-O, and C=O, respectively.⁵ The appearance of C-N confirmed the successful modification of GO with amino acids.

3.1.4 Transmission electron microscopy. The morphologies of GO, AMGO and AMGO@Fe₃O₄ were characterized by transmission electron microscopy. From Fig. 4A, it can be seen that GO had a flake-like shape with some corrugations. After modification of amino acids, the morphology of AMGO-Glu was still a folded and wrinkled sheet (Fig. 4B), which meant that AMGO-Glu retained the characteristic of a large surface area. Fig. 4C and D show that the Fe₃O₄ nanoparticles were distributed on the surface of AMGO-Glu, which indicated the successful synthesis of Fe₃O₄.²⁵

3.2 Optimization of the magnetic separation conditions

Bovine hemoglobin (BHb) was chosen as the representative protein to study the extraction capacity of the prepared AMGO@Fe $_3O_4$. The extraction amount was calculated according to the following formula:²⁶

$$Q_{\rm e} = \frac{(C_{\rm i} - C_{\rm e})V}{m} \tag{1}$$

where $Q_e (\text{mg g}^{-1})$ is the mass of protein extracted by unit mass of AMGO@Fe₃O₄, $C_i (\text{mg mL}^{-1})$ and $C_e (\text{mg mL}^{-1})$ are the initial and equilibrium protein concentrations, V(mL) is the volume of the initial solution and m (g) is the weight of the adsorbent based on the dry weight of AMGO@Fe₃O₄.

Several factors including the type of amino acid, pH value, addition of electrolyte, extraction time, and protein concentration were optimized to attain the best adsorption efficiency.

3.2.1 Effect of the type of amino acid. As shown in Table S1,† four kinds of amino acid have been selected to modify the surface of GO in this experiment. All of the extraction experiments were carried out at room temperature. The extraction capacities of the four kinds of AMGO@Fe₃O₄ are shown in Fig. 5. The results indicated that the extraction amounts of the adsorbents were different (AMGO-Glu@Fe₃O₄: 868.3 mg g⁻¹, AMGO-Arg@Fe₃O₄: 647.5 mg g⁻¹, AMGO-Cys@Fe₃O₄: 769.9 mg g⁻¹, AMGO-Phe@Fe₃O₄: 600.6 mg g⁻¹) for the same protein BHb. This phenomenon might be explained by the multiple interactions between protein and adsorbent including electrostatic, hydrogen bonding and π - π interactions. When the four



Fig. 4 TEM images of (A) GO, (B) AMGO-Glu, (C) AMGO-Glu@Fe₃O₄, and (D) AMGO-Glu@Fe₃O₄.



Fig. 5 Effect of the type of amino acid on the extraction of bovine hemoglobin (BHb). Glu, glutamic acid; Arg, ∟-arginine; Cys, cysteine; Phe, phenylalanine.

kinds of amino acid were modified on GO, their different functional groups would have different interactions with protein. As phenylalanine was grafted on GO, the introduction of the benzene ring would enhance the π - π interaction on protein. On the other hand, L-arginine would strengthen hydrogen bonding interaction and change the electrostatic interaction owing to the existence of $-NH_2$, and cysteine and glutamic acid would increase the negative charge on the surface and enhance the electrostatic interaction between protein and adsorbent. As AMGO-Glu@Fe₃O₄ was extracted in the greatest quantity, electrostatic interaction may be the main driving force, and AMGO-Glu@Fe₃O₄ was selected for use in further experiments.

3.2.2 Effect of pH value. The pH value played an important role in the extraction capacity of protein in the magnetic separation procedure. It could influence the conformation of protein and the charges on both protein and adsorbent. The pH value was varied from 3.0 to 10.0 using PBS buffer solution (0.05 mol L^{-1}) and the results are shown in Fig. 6A. It can be seen that the amount extracted reached a maximum at pH 6.0. This was because, at pH 6.0, the AMGO-Glu@Fe₃O₄ was highly anionic, but BHb (pI = 6.8) was highly cationic, as shown in Fig. 7. The electrostatic attraction led to the high extraction capacity. However, with pH > 6.0, the amount of BHb extracted decreased sharply with increasing pH value. This could be attributed to the fact that the surfaces of BHb and AMGO-Glu@Fe₃O₄ were both negatively charged when the pH was greater than the pI of BHb. Although there were hydrogen bonding and π - π interactions between AMGO-Glu@Fe3O4 and BHb, the electrostatic repulsion interaction was much stronger and reduced the extraction performance of the adsorbent. In addition, a higher pH value could weaken hydrogen bonds between protein and adsorbent. As the extracted amount reached its maximum at pH 6.0, a solution pH value of 6.0 was chosen for further experiments.

3.2.3 Effect of the addition of electrolyte. The influence of the addition of electrolyte on extraction capacity was investigated using PBS buffer solutions with different concentrations of NaCl (0.0–0.5 mol L^{-1}). As shown in Fig. 6B, the extraction capacity of BHb decreased with the increase in concentration of NaCl. The negative impact might be ascribed to the electrostatic interaction between protein and adsorbent. Under the conditions of solution, BHb with positive charges on its surface attracted negative ions in the solution, which would reduce the electrostatic attraction between BHb and AMGO-Glu@Fe₃O₄. With the increase in concentration of NaCl, the salt ions strongly shielded the electrostatic attraction, making it difficult to adsorb and rearrange protein on the surface of the adsorbent, and decreasing the extraction efficiency. As the addition of NaCl could not increase the extraction capacity of protein, NaCl was not added in further experiments.

3.2.4 Effect of extraction time. In order to obtain an appropriate adsorption time, the experiment was carried out with different extraction times, and the results are presented in Fig. 6C. It can be seen that a large amount of BHb could be adsorbed within 5 minutes, which exhibited the superior adsorption capacity of AMGO-Glu@Fe₃O₄. In the following 35 min, protein adsorption increased gradually and after 40 min, the amount extracted reached equilibrium and did not change significantly with increasing time after that. This could be interpreted by the fact that AMGO-Glu@Fe₃O₄ had many adsorption sites on its surface at the start, so that the protein could be easily adsorbed. As time progressed, more and more adsorption sites had been occupied, so the adsorption process slowed down and eventually reached equilibrium. Since the extraction process reached equilibrium at 40 min, this was selected as the optimum extraction time.

3.2.5 Effect of protein concentration. The effect of protein concentration was studied using different initial BHb concentrations, with values in the range $0.1-2.0 \text{ mg mL}^{-1}$. In order to



Fig. 6 Effects of extraction conditions. (A) Effect of pH: protein concentration, 1 mg mL^{-1} ; concentration of NaCl, 0.0 mol L⁻¹; extraction time, 1 h. (B) Effect of the solution ionic strength: protein concentration, 1 mg mL^{-1} ; extraction time, 1 h; solution pH 6.0. (C) Effect of extraction time: protein concentration, 1 mg mL^{-1} ; solution pH 6.0. (D) Effect of protein concentration: concentration of NaCl, 0.0 mol L⁻¹; extraction time: protein concentration of NaCl, 0.0 mol L⁻¹; solution pH 6.0. (D) Effect of protein concentration: concentration of NaCl, 0.0 mol L⁻¹; extraction time, 1 h; solution pH 6.0.



Fig. 7 Zeta potentials of AMGO-Glu@Fe $_3O_4$ and BHb under different pH values.

increase in protein concentration, and almost all of the protein in the solution was adsorbed on the surface of the adsorbent when the concentration of protein was less than 0.5 mg mL⁻¹. This is because there were sufficient adsorption sites on AMGO-Glu@Fe₃O₄ to extract all of the protein at lower concentrations, and the extraction amount was only dependent on the amount of protein in the solution. When the initial concentration was in the range 0.5–1.0 mg mL⁻¹, the upward trend turned to gradual, and then the extraction capacities remained almost constant with concentrations greater than 1.0 mg mL⁻¹. The results indicated that the adsorption capacity of AMGO-Glu@Fe₃O₄ was limited, and the adsorption sites were saturated when the concentrations less than 1.0 mg mL⁻¹. Therefore, concentrations less than 1.0 mg mL⁻¹ could be selected as an appropriate value for the BHb concentration.

3.3 Comparison of GO@Fe₃O₄ and AMGO-Glu@Fe₃O₄

guarantee the attainment of adsorption equilibrium, the extraction time was chosen to be 1 h. As can be seen from Fig. 6D, the extraction amount increased linearly with the $GO@Fe_3O_4$ and AMGO-Glu@Fe_3O_4 were used to adsorb proteins under the same conditions to evaluate the advantage of the proposed AMGO-Glu@Fe_3O_4. It can be seen from Fig. 8 that Paper



Fig. 8 Comparison of protein extraction ability of GO@Fe_3O_4 and AMGO-Glu@Fe_3O_4.

both adsorbents had the ability to extract proteins, but their extraction capacities were different. Because of the conjugated structure and the existence of oxygenated groups on GO, the GO@Fe₃O₄ could form electrostatic, hydrogen bonding and π - π interactions with proteins. When glutamic acid was modified on the surface of GO, the extraction capacities of BHb and Lvz increased. This may be because the carboxyl groups in glutamic acid could increase the negative charge on the surface after modification. As shown in Fig. 9, the zeta potentials of GO@Fe₃O₄ and AMGO-Glu@Fe₃O₄ were -6.8 mV and -14.1 mV, respectively. In addition, the chosen pH value of extraction was 6.0, so the BHb (pI = 6.8) and Lyz (pI = 11.0) were positively charged. The increase in negative charge could enhance the electrostatic attraction between protein and the adsorbent, which improved the extraction capacities. However, under the same pH value, BSA (pI = 4.8) and OVA (pI = 4.7) were negatively charged, and the increase in negative charge could strengthen the electrostatic repulsion, so that the extraction capacities decreased. Moreover, the adsorption ratios of BHb/ BSA and BHb/OVA were 10.54 and 4.79, respectively, which

indicated that AMGO-Glu@Fe₃O₄ could selectively extract BHb from an acidic protein mixture.

3.4 Regeneration of the AMGO@Fe₃O₄

In order to explore the regeneration of AMGO-Glu@Fe₃O₄, a series of desorption experiments were performed. It can be seen in Sections 3.2.2 and 3.2.3 that an increase in pH value and concentration of NaCl could decrease the amount of BHb extracted. In addition, the hydrogen bonds between AMGO-Glu@Fe₃O₄ and BHb could be broken under alkaline conditions. Therefore, 0.05 mol L⁻¹ PBS buffer solution (pH 11.0) containing 0.5 mol L⁻¹ NaCl was chosen as the eluent. After desorption, the AMGO-Glu@Fe₃O₄ was recovered by an external magnetic field and washed 3 times with deionized water, then it was utilized to extract BHb again. Three consecutive extraction–desorption cycles were carried out and the results are shown in Fig. S2.† The results showed only a minor decrease in the extraction capacity after three consecutive cycles, indicating the good recyclability of the proposed AMGO-Glu@Fe₃O₄ in magnetic separation.

3.5 Extraction selectivity

In order to examine the possibility of selective extraction of BHb based on AMGO-Glu@Fe₃O₄, a protein mixture including BSA and BHb was used as a model. Meanwhile, the extraction selectivity of GO@Fe3O4 was explored under the same conditions for comparison. After adsorption and desorption, the supernatants were both collected for analysis by sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) assay. The results are shown in Fig. S3.[†] For GO@Fe₃O₄, the presence of protein was not observed in the supernatant after adsorption (lane 3), whereas both the BSA band around 66 kDa and the BHb band around 14 kDa were found in the eluent (lane 5). The results showed that both BHb and BSA were adsorbed by GO@Fe₃O₄, which indicated that the protein extraction of GO@Fe₃O₄ was nonspecific. On the other hand, after adsorption by AMGO-Glu@Fe₃O₄, only the BSA band was discovered in the supernatant (lane 4), while in the recovered eluent, only the



Fig. 9 Zeta potentials of $GO@Fe_3O_4$ and $AMGO-Glu@Fe_3O_4$.



BHb band was observed (lane 6). The results showed that AMGO-Glu@Fe $_3O_4$ could selectively extract BHb from BSA.

3.6 Analysis of real samples

To evaluate the performance of AMGO-Glu@Fe₃O₄ in real samples, the nanocomposite was used to extract BHb from bovine plasma. After adsorption and desorption, the supernatants were both collected for analysis by SDS-PAGE assay. As shown in Fig. S4,[†] some protein bands were recorded for the bovine plasma sample (lane 2), which may be attributed mainly to BSA, BHb, transferrin and so on. After adsorption by AMGO-Glu@Fe₃O₄, the bands were still observed in the supernatant (lane 3). This is because the amount of proteins in bovine plasma was excessive for the adsorbent; however, in the recovered eluent, lane 4 showed a major band corresponding to 14 kDa, which was identified as BHb. The results showed that AMGO-Glu@Fe₃O₄ could selectively extract BHb in complex real samples.

3.7 Validation of the proposed method

To evaluate the analytical performance of the proposed method, stability, repeatability and precision experiments were carried out. The stability experiment was performed by detecting a sample continuously over 3 days under the same conditions and the relative standard deviation (RSD) was 1.00% (n = 3). The repeatability experiment was performed by monitoring three copies of the same sample under the same conditions and the RSD was 0.76% (n = 3). The apparatus precision experiment was performed by measuring the same sample three times using a UV-vis spectrophotometer, and the value of RSD was 0.02% (n = 3). The results indicated the proposed method possessed excellent precision, repeatability, and stability.

4. Conclusions

A novel amino acid-modified GO magnetic nanocomposite AMGO@Fe₃O₄ was successfully synthesized for the magnetic separation of proteins. The material displayed good magnetic properties and proteins could be separated rapidly using an external magnet. The research results showed that the AMGO@Fe₃O₄ was superior to GO@Fe₃O₄ in the extraction of BHb and Lyz, and could selectively extract BHb from BSA in complex real samples. Moreover, the AMGO@Fe₃O₄ could be easily regenerated, and the extraction capacity decreased only slightly after being used three times. The proposed method possessed excellent precision, repeatability and stability, and could be used as a simple and efficient separation technique for proteins in real samples.

Conflict of interest

The authors have declared no conflict of interest.

Acknowledgements

This work was supported by the Natural Science Foundation of China (No. 21575076 and 21621003), the National Key Research

and Development Program of China (2016YFA0203101), and the Beijing Municipality Science and Technology Program (D161100002116001).

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