



Cite this: *RSC Adv.*, 2018, 8, 27654

Preparation and application synthesis of magnetic nanocomposite using waste toner for the removal of Cr(vi)[†]

Hong Zhu,^a Yucheng Zhou,^a Shengsen Wang,^{ab} Xiaoge Wu,^a Jianhua Hou,^{ab} Weiqin Yin,^a Ke Feng,^{ab} Xiaozhi Wang ^{*abc} and Jie Yang^d

In this study, a novel magnetic nanocomposite was prepared using waste toner (WT) through high temperature decomposition, and calcination was conducted in different atmospheres (air, ammonia, and vacuum). WT calcined in ammonia (WT(NH₃)), and it was then utilized as an efficient absorbent for the reduction of Cr(vi) in aqueous solutions; a batch experiment with different conditions was performed to investigate its Cr(vi) removal ability. The effects of two pH-regulating acid (HCl and H₂SO₄) treatments were also studied. It was found that WT(NH₃) could remove about 99% Cr(vi) at pH 2 under H₂SO₄ treatment. The XRD and TEM results coupled with VSM results confirmed that WT(NH₃) is an Fe₃O₄/Fe₂N nanohybrid, which possesses excellent water-dispersibility and remarkable magnetic properties. XPS analysis showed the presence of Cr(vi) and Cr(III) on the surface of WT(NH₃), which indicated that Cr(vi) was reduced to Cr(III). Furthermore, H₂SO₄ regulation also promoted the reduction of Cr(vi) by WT(NH₃), and this reduction was higher than that obtained by HCl regulation.

Received 20th June 2018

Accepted 18th July 2018

DOI: 10.1039/c8ra05291c

rsc.li/rsc-advances

1. Introduction

Chromium (Cr) is a very common and toxic heavy metal, and it is usually present in water and soil. Most Cr in the environment is released from industrial activities including tanning, textile preservation, electroplating, and sewage sludge.^{1,2} The hazards of chromium to the human body mainly include skin and respiratory ulcers, meningitis, and lung cancer.³ Thus, the US Environmental Protection Agency has recommended that the Cr concentration should be below 50 μg L⁻¹ (ref. 4) in most natural waters. Cr is most frequently present in the hexavalent (Cr(vi)) or trivalent (Cr(III)) forms in aqueous solutions.⁵ These two common oxidation states of Cr are extremely different in charge, physicochemical properties and chemical and biochemical reactivities.⁶ In fact, the toxicity of Cr(vi) is five hundred times greater than that of Cr(III).⁷ Cr(vi) ions exist as chromate ions (CrO₄²⁻) and dichromate ions (Cr₂O₇²⁻) in solution, and these ions are more soluble and mobile than Cr(vi) in ground water; they are also more capable of oxidizing other species.⁸ Therefore, several

treatment methods have been applied such as ion exchange, electrolytic removal, chemical precipitation,⁹ adsorption,¹⁰ and reduction.¹¹ Recently, reduction–sorption has been considered as an efficient and economically viable method for Cr(vi) removal. Moreover, Cr(vi) can be reduced to its less toxic form.¹² However, this method requires absorbents that contain abundant active sites or functional groups.

With the increasing demand for electronic and electrical equipment (EEE) in modern society, a significant amount of EEE waste (WEEE) is produced, and the major constituents of WEEE plastics include polypropylene (PP) and high-impact polystyrene (HIPS).¹³ To recycle WEEE, some researchers have prepared R-PP/R-HIPS blends by adding polystyrene-poly(ethylene/propylene) (SEP) or glycidyl methacrylate (GMA) to obtain functional polymer nanocomposites.¹⁴ Also, it has been reported that near-critical water can be used to recycle printed circuit boards for the collection of copper foils and fabrics.¹⁵ The overuse of printing equipment has led to a significant increase in the number of waste toner cartridges.¹⁶ Powdery waste toner (WT) accounts for 8.0 wt% of a toner cartridge. Also, the weight of WT has reached about 36 t per year in China. Furthermore, if WT is disposed without reasonable management, its toxic components, including hazardous organic components, will be released into the environment, which can pollute the soil, underground water and air; this will be a tremendous threat to the human health.¹⁷ In fact, WT is a mixture that includes about 7.0 wt% polyacrylate, 55.0 wt% polystyrene, 3.0 wt% SiO₂, and 35.0 wt% Fe₃O₄.^{18–20} In recent years, very few researchers have studied the recycling and

^aCollege of Environmental Science and Engineering, Yangzhou University, Jiangsu 225127, China. E-mail: xzwang@yzu.edu.cn

^bJiangsu Collaborative Innovation Center for Solid Organic Waste Resource Utilization, Nanjing 210095, China

^cInstitutes of Agricultural Science and Technology Development, Yangzhou University, Yangzhou 225127, Jiangsu, China

^dKey Laboratory of Crop and Livestock Integration, Ministry of Agriculture, Nanjing 210095, China

[†] Electronic supplementary information (ESI) available. See DOI: 10.1039/c8ra05291c



transformation of WT. The method of vacuum-gasification-condensation was used to treat WT since the organic matter in WT can be decomposed and gasified at high temperature. Meanwhile, SiO_2 and Fe_3O_4 in WT were transformed into nanosized Fe_3O_4 and nano- SiO_2 .¹⁹ A thermal transformation process was investigated to convert WT into 98% pure iron by using the gases collected during the thermal transformation as an *in situ* source of carbon.²¹

Recently, iron-bearing materials have been applied to various fields such as drug delivery, sensors, electromagnetic interference shielding, and adsorbents for environmental remediation.^{22–24} In the environmental field, iron-bearing materials such as magnetite,^{25,26} iron sulfides,^{27,28} goethite,²⁹ and zerovalent iron³⁰ can remediate $\text{Cr}(\text{vi})$. It has also been reported that fabric- and cellulose-derived magnetic carbon-iron nanomaterials possess high efficiency for $\text{Cr}(\text{vi})$ removal,^{31,32} where $\text{Cr}(\text{vi})$ is reduced to $\text{Cr}(\text{iii})$ by the reductive intermediates generated from ZVI such as $\text{Fe}(\text{ii})$. Moreover, the nanocarbon structure and different carbonization temperatures affect the $\text{Cr}(\text{vi})$ removal capacity.^{33,34} Also, Fe_3O_4 is a type of iron oxide, and it exhibits remarkable performance owing to its high efficiency of adsorbing $\text{Cr}(\text{vi})$ and reducing $\text{Cr}(\text{vi})$ to less toxic $\text{Cr}(\text{iii})$.³⁵ Furthermore, compared to traditional adsorbents, this compound can be separated and recovered rapidly using an external magnetic field. It is known that Fe_3O_4 nanoparticles can be obtained by calcining WT. Although the calcination of WT has been previously studied, its calcination under different atmospheres and $\text{Cr}(\text{vi})$ uptake using the resultant nanoparticles have not been well investigated.

In this study, we attempted to develop a novel low-cost magnetic nanocomposite by recycling waste toner through high-temperature decomposition. Calcination was conducted in air, ammonia and vacuum to explore the influence of atmosphere on WT modification. Due to the unique properties of the materials, we used different acids to adjust the pH value during the experiments, and the results showed significant difference. The aim of this study is to characterize the as-prepared WTs, evaluate their applicability for $\text{Cr}(\text{vi})$ adsorption and reduction, and explore the mechanisms of $\text{Cr}(\text{vi})$ removal.

2. Materials and methods

2.1 Reagents

$\text{Cr}(\text{vi})$ aqueous solution was prepared by dissolving potassium dichromate ($\text{K}_2\text{Cr}_2\text{O}_7$). pH adjustment was carried out using 0.1 M HCl, 1.1 M H_2SO_4 and 0.1 M NaOH. All the reagents were of analytical grade and purchased from the Sinopharm Chemical Reagent Co., Ltd. (Shanghai China). Deionized (DI) water was used to prepare the chemicals. WT was collected from HP toner cartridges in printing shops from Yangzhou University.

2.2 Preparation

To prepare the magnetic nanomaterials in this study, WT was calcined in a tube furnace at 600 °C at a heating rate of 5 °C min^{-1} for two hours in an atmosphere of air, vacuum, and ammonia. The material was then cooled to ambient

temperature, and the finally prepared WTs denoted as WT(Air), WT(Vac), and WT(NH_3) were obtained by ball milling. Among them, the ammonia gas flow rate was 300–400 mL min^{-1} . For comparison, the original WT without any treatment was also used in the experiments.

2.3 Characterization

The surface morphology of WTs was examined *via* scanning electron microscopy (SEM, S-4800II) and transmission electron microscopy (TEM, TF-20) coupled with energy dispersive X-ray spectroscopy (EDS, Vario EL cube). Brunauer–Emmett–Teller (BET) specific surface areas were measured using Micromeritics ASAP 2460. Elemental oxidation states and functional groups were determined *via* X-ray electron spectroscopy (XPS, ESCALAB 250Xi). The structure and phase of the as-prepared WTs were analyzed with an X-ray diffractometer (XRD, D8-ADVANCE).

2.4 $\text{Cr}(\text{vi})$ removal experiments

Batch experiments were performed to study the removal of $\text{Cr}(\text{vi})$ from aqueous solution using different WTs in 100 mL conical flasks containing 25 mL of solutions at the desired initial $\text{Cr}(\text{vi})$ concentrations and pH values. The most optimum dosage of WTs was determined by adding gradient concentrations ranging from 1 to 8 g L^{-1} during the experiment. The absorption and reduction of $\text{Cr}(\text{vi})$ (50 mg L^{-1}) by WTs were performed using 0.1 M NaOH and two different pH-regulating acids (0.1 M HCl and 1.1 M H_2SO_4) to adjust the pH value.

The mixture was placed on an Orbit shaker with continual mixing at 300 rpm. All experiments were carried out at the room temperature of 25 °C and were performed in duplicate. After reacting for a certain time, WTs were filtered through a 0.45 μm membrane, and the residual $\text{Cr}(\text{vi})$ concentration was determined using an ultraviolet visible (UV) spectrophotometer at $\lambda = 540$ nm, according to the 1,5-diphenylcarbazide colorimetric method.³⁶ To measure total Cr concentration, the 1,5-diphenylcarbazide colorimetric method (with KMnO_4) was used.³⁷ Also, the leached iron in the solution was determined using the 1,10-phenanthroline spectrophotometry method.³⁸

3. Results and discussion

3.1 Characteristics of as-prepared WTs

Fig. 1 shows the XRD patterns of the original WT, WT(Air), WT(Vac) and WT(NH_3). The diffraction peaks of the original WT at the 2θ values of 30.1°, 35.58°, 43.11°, 53.45°, 57.13°, and 62.59° could be ascribed to the reflections of the (220), (311), (400), (422), (511), and (440) planes, respectively, which matched well with the standard pattern of JCPDS 019-0629 of the cubic lattice of Fe_3O_4 . The characteristic peaks of Fe_3O_4 of WT(Air) and WT(Vac) were still observed. Based on the Debye–Scherrer equation, the average crystallite size of Fe_3O_4 nanoparticles was calculated from the (311) plane, and it was found that the crystals were about 43.5 nm in diameter.³⁹ Fig. 1(d) shows clear Fe_2N peaks, which are in accordance with PDF#50-0958,⁴⁰ confirming that WT(NH_3) is an $\text{Fe}_3\text{O}_4/\text{Fe}_2\text{N}$ nanohybrid. Furthermore, our results suggested that the chemical



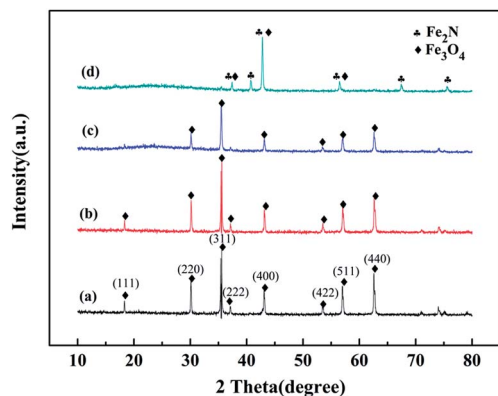


Fig. 1 XRD patterns of (a) original WT, (b) WT(Air), (c) WT(Vac), and (d) WT(NH₃).

composition of the material can be changed by annealing at 600 °C in an ammonia atmosphere.

As shown in Fig. 2, compared with the result for the spectrum of original WT, the organic groups disappeared in the spectra of WT(Air), WT(Vac) and WT(NH₃). The band at 578 cm⁻¹ was related to the vibration of the Fe–O functional group, which matched well with the characteristic peak of Fe₃O₄.⁴¹ This finding was in accordance with the XRD patterns. The weak peaks between 1101 and 1151 cm⁻¹ were assigned to the Si–O–Si group antisymmetric stretching vibration modes, indicating the existence of SiO₂ in the material.

It can be seen from the SEM image (Fig. 3(a)) that original WT is a granular mixture with a diameter of about 5 μm. After calcination, Fe₃O₄ and SiO₂ were retained in the residue. As shown in Fig. 3(b) and (c), WT(Vac) is found to be more aggregated than WT(NH₃), which was in accordance with the TEM results (Fig. 4(b)). The Fe₃O₄/Fe₂N nanoparticles displayed a spherical morphology with a mean diameter of 200 nm (Fig. 4(a)).

To further verify the distribution and structure of the material, TEM images of WT(NH₃) and WT(Vac) were obtained (Fig. 4). Both samples showed few tetragonal or spherical Fe₂N species, and spherical Fe₃O₄ species were also observed. Furthermore, WT(NH₃) had more phase interfaces of Fe₃O₄ and

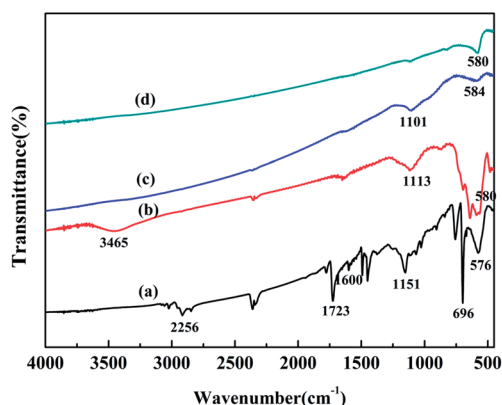


Fig. 2 FTIR spectra of (a) original WT, (b) WT(Air), (c) WT(NH₃), and (d) WT(Vac).

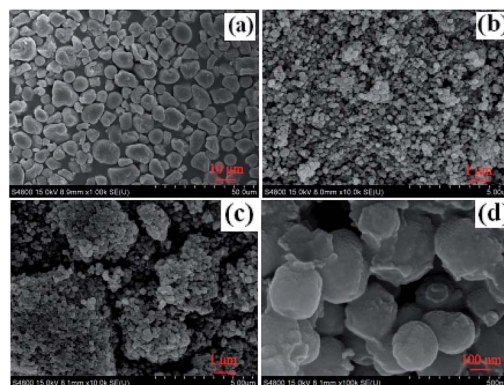


Fig. 3 SEM images of (a) original WT, (b) WT(NH₃), (c) WT(Vac), and (d) WT(NH₃).

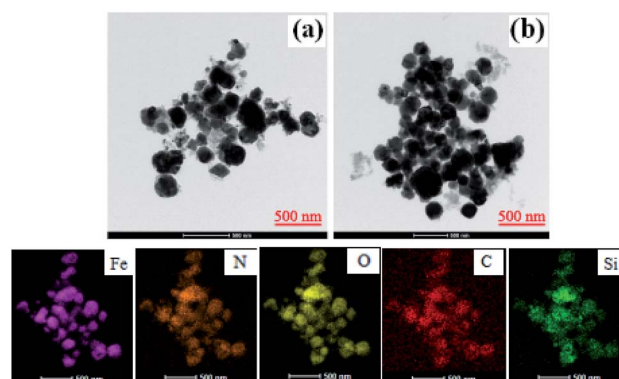


Fig. 4 TEM images of (a) WT(NH₃) and (b) WT(Vac) and the corresponding EDX elemental mappings of iron, nitrogen, oxygen, carbon, and silicon.

Fe₂N than WT(Vac). The EDS and TEM mapping analysis revealed the presence of Fe, Si, C and O in WTs and Fe, Si, C, N and O in WT(NH₃). Higher amount of N (8.6 wt%) was believed to be due to N-doping from annealing in ammonia (Table 1).

The Brunauer–Emmett–Teller (BET) surface areas of WT(Air), WT(Vac) and WT(NH₃) are shown in Table S1.† WT(NH₃) has a larger surface area (42.53 m² g⁻¹) than WT(Vac) (11.93 m² g⁻¹) and WT(Air) (11.75 m² g⁻¹), which can improve mass transfer and increase the number of active sites.⁴² The pore volume is in good agreement with the surface area.

The XPS spectra of WT(NH₃) depicted in Fig. 5 exhibit the presence of C, O, N, and Fe before the reaction. The C 1s spectra exhibit four peaks; the peaks at binding energies of 284.7 eV and

Table 1 Elemental compositions of the original WT, WT(Air), WT(Vac) and WT(NH₃) (%)

Sample	N	C	H	Fe
WT	0.049	52.66	5.232	32.07
WT(Air)	0.012	0.11	0.420	56.54
WT(Vac)	0.068	9.66	0.322	66.41
WT(NH ₃)	8.641	7.50	0.357	69.56



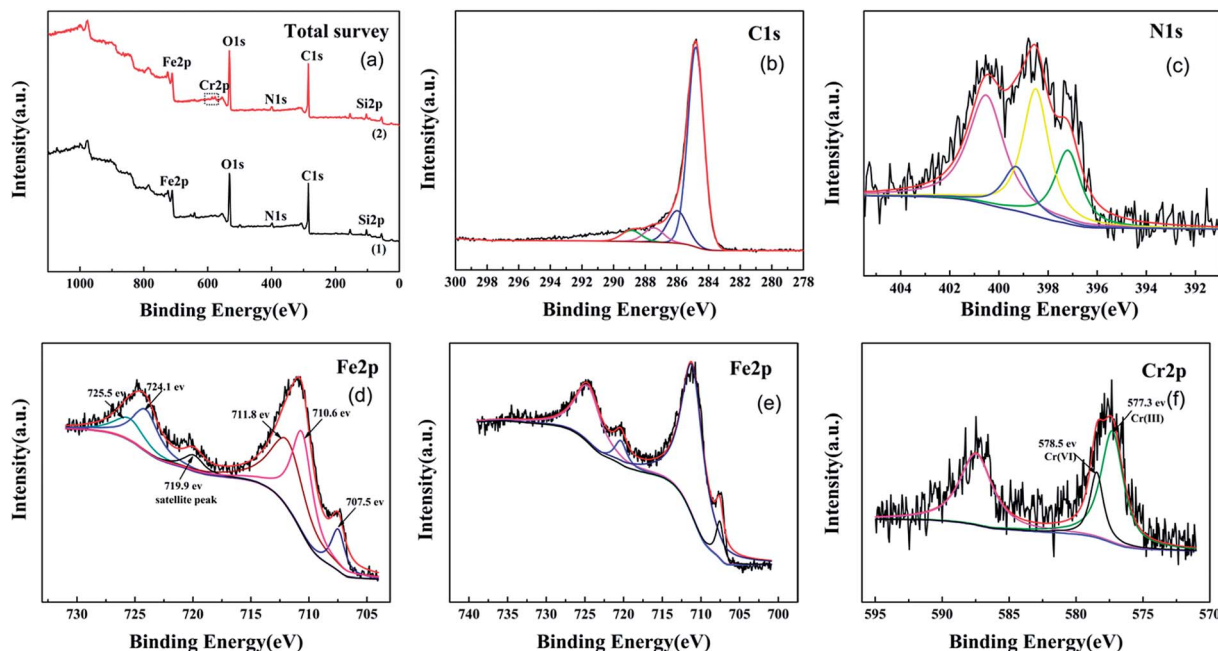


Fig. 5 XPS total survey (a), C 1s (b), N 1s (c), Cr-unloaded Fe 2p (d), Cr-loaded Fe 2p (e), and Cr 2p's (f) spectra.

285.9 eV corresponded to C–H, C–C, and hydroxyl (C–OH) bonds, and the peaks at 287.5 eV (C=O, O–C–O) and 288.9 eV corresponded to the carbonyl (O–C=O) bond (Fig. 5(b)).⁴³ Fig. 5(c) shows the high-resolution XPS spectra of the N peaks: pyrrolic-N at 400.5 eV, Fe–N at 399.3 eV, and pyridinic-N at 398.6 eV.⁴⁴ Nitrogen was successfully doped into Fe₃O₄ and reacted with iron, which was consistent with Fe₂N from XRD characterization. The XPS Fe 2p spectra of WT(NH₃) showed two peaks centered at 710.9 eV and 724.1 eV, which were assigned to Fe 2p_{3/2} and Fe 2p_{1/2}, respectively. For Fe 2p_{3/2}, the peak at BE of 710.6 eV was assigned to ferrous iron in Fe₃O₄ (FeO·Fe₂O₃) and Fe₂N, whereas that at 711.8 eV corresponded to Fe(III) in Fe₃O₄.⁴⁵ Metallic iron was also observed at the binding energy of 707.5 eV based on the Fe 2p_{3/2} spectra. The satellite peak at 719.9 eV was much weaker, which further proved the absence of Fe₃O₄ rather than γ -Fe₂O₃.⁴⁶ These results were consistent with the TEM and XRD results.

The magnetization curves of original WT and WT(NH₃) were studied using a vibrating sample magnetometer (VSM) at an ambient temperature. The corresponding hysteresis loops are shown in Fig. 6, and the saturation magnetization of original WT was determined to be about 37.9 emu g⁻¹, which decreased to 21.1 emu g⁻¹ (WT(NH₃)) after calcination in an ammonia atmosphere. This can guarantee its convenient separation from pollutants when used as an adsorbent.

3.2 Effect of sorbent on Cr(vi) removal

To compare the Cr(vi) removal efficiencies of WT(NH₃), WT(Vac), WT(Air) and original WT, the best reaction conditions were obtained: an initial Cr(vi) concentration of 50 mg L⁻¹ and sorbent dosage of 4 g L⁻¹ at pH = 2.0.

As shown in Fig. 7, Cr(vi) removal efficiencies decreased in the following order: WT(NH₃) > WT(Vac) > WT(Air) > original

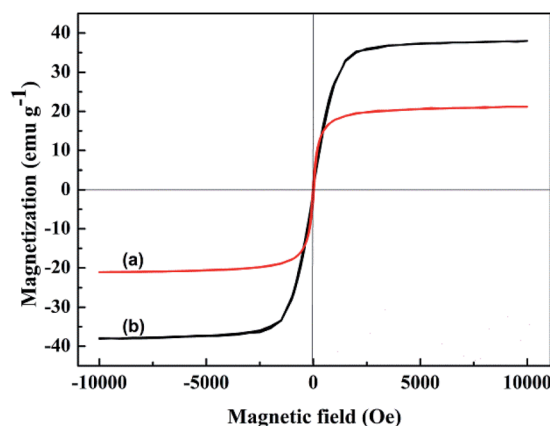


Fig. 6 Magnetic hysteresis loops of (a) original WT and (b) WT(NH₃).

WT. The effects of two pH-regulating acid (HCl and H₂SO₄) treatments were investigated. Under HCl treatment, WT(NH₃) removed 78.2% of Cr(vi) compared to WT(Vac) and WT(Air), which removed 29.9% and 5.3% of Cr(vi), respectively. Also, there was only a slight difference between WT(Vac) and WT(Air) under the two acid treatments. However, for WT(NH₃), its Cr(vi) removal efficiency was greatly affected. Within the first 1.5 h, its Cr(vi) removal efficiencies under HCl and H₂SO₄ treatments were almost identical. For HCl treatment, its efficiency showed only a gradually increasing trend after 2 h; on the contrary, its efficiency increased sharply with H₂SO₄ treatment, which resulted in 99.9% Cr(vi) removal after 7 h.

3.3 Cr(vi) removal kinetics and isotherms

The pseudo-second-order kinetic model was used to determine the potential rate-limiting step involved in the Cr(vi) removal process by WT(NH₃) and WT(Vac) (Fig. S1†). The kinetic



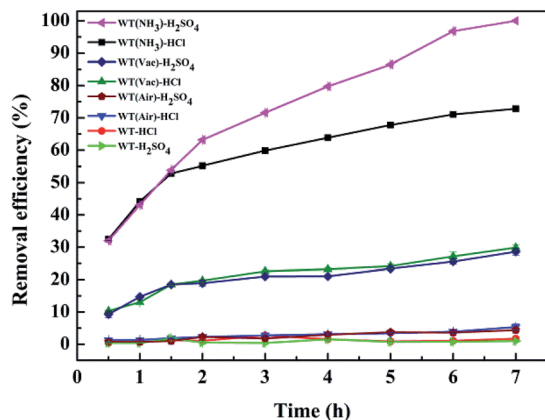


Fig. 7 Removal efficiency of Cr(vi) using different systems with reaction time. Reaction conditions: 4 g L⁻¹ as-prepared sorbents, 49.68 mg L⁻¹ Cr(vi) concentration, and pH = 2.0.

parameters and R^2 were determined by linear regression (Table S2†). The calculated q_e values of the pseudo-second-order kinetic model were close to the experimental data, indicating that the adsorption of Cr(vi) is mainly controlled by chemical adsorption. The k value of WT(Vac) was higher than that of WT(NH₃), which revealed that WT(Vac) exhibited faster adsorption of Cr(vi) and lower adsorption capacity than WT(NH₃).

The linearized modeled isotherms are plotted in Fig. S2.† As shown, high correlation coefficients and good R^2 values were obtained from the isotherm models described for the removal of Cr(vi) by WT(NH₃) in both HCl and H₂SO₄ systems (Table S3†). The adsorption of Cr(vi) was well fitted by the Langmuir model than the Freundlich model, indicating that Cr(vi) removal by WT(NH₃) is a monolayer adsorption process.

3.4 Effect of initial pH on Cr(vi) removal

The effect of initial pH on Cr(vi) removal by WT(NH₃) was investigated, and it was shown that the Cr(vi) adsorption is strongly dependent on the values of solution pH (Fig. 8). It can

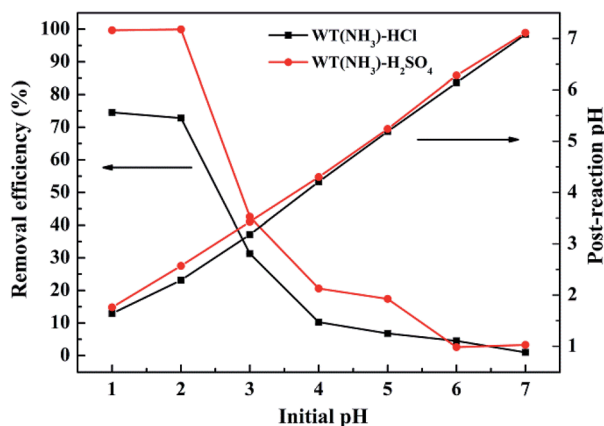


Fig. 8 Effect of initial pH on Cr(vi) removal efficiency by WT(NH₃). Reaction conditions: 4 g L⁻¹ WT(NH₃), 49.68 mg L⁻¹ Cr(vi) concentration, and 7 h reaction time.

be clearly observed that the removal efficiency of Cr(vi) decreased with an increase in the solution pH value. Specifically, the Cr(vi) removal efficiencies were 78.2% and 99.9% in the HCl and H₂SO₄ systems, respectively, at pH 2, and these values dropped sharply with an increase in pH to 3. Also, the pH increased after the reaction, which was possibly due to the consumption of H⁺ during the reduction of HCrO₄⁻ and Cr₂O₇²⁻ to Cr(III).^{47,48} At neutral pH, only 0.4–0.8% Cr(vi) was removed in the HCl and H₂SO₄ systems. With an increase in pH, the formation of Cr(vi) or Cr(III) precipitates was enhanced, which blocked the reactive sites.⁴⁹

To investigate the surface charge of WT(NH₃), we measured its zeta potential (Fig. S3†). The isoelectric point of WT(NH₃) was found to be about pH 2. Below this value, the surface of WT(NH₃) was positively charged, which was beneficial to bind with chromium anions through electrostatic attraction and ion exchange. At high pH values, the surface of WT(NH₃) became less positively charged, which might be unfavourable for Cr(vi) adsorption.⁵⁰

3.5 Cr(vi) removal mechanisms

To investigate the Cr(vi) removal mechanism of WTs, both the XPS and EDS techniques were used to characterize the surface chemical composition of WT(NH₃) before and after the reaction. After the reaction, some new peaks appeared, which coincided with those in the photoelectron spectra of Cr 2p. High resolution XPS spectra were used to clarify the state of Cr absorbed on the surface of WT(NH₃), and the results are shown in Fig. 5(f). The Cr 2p spectra exhibited two peaks, namely, Cr 2p_{3/2} (577.6 eV) and Cr 2p_{1/2} (587.5 eV). Using XPS-peak-differentiating analysis, the Cr 2p_{3/2} spectra were deconvoluted into two major components at binding energies of 578.5 eV and 577.3 eV, which were due to Cr(vi) and Cr(III), respectively.⁵¹ This demonstrated the coexistence of both Cr(vi) and Cr(III) on the surface of WT(NH₃) after reacting with Cr(vi), indicating that the Fe₃O₄/Fe₂N nanocomposite in WT(NH₃) is beneficial for the reduction of Cr(vi) to Cr(III). Moreover, the splitting of Cr 2p_{3/2} line peak into two peaks indicated the precipitation of Cr₂O₃ or Cr(OH)₃ during the adsorption process.⁵² Based on the analysis, the ratio of Cr(III)/Cr(vi) was found to be 2.8 : 1, implying that the reaction mechanisms might be sorption together with reduction processes. In addition, the concentrations of total Cr decreased by 14% and 46% in the H₂SO₄ and HCl systems, respectively. All the Cr(vi) ions were almost reduced to Cr(III) in the solution; thus, the H₂SO₄ system can promote the reduction of Cr(vi) by WT(NH₃) (Fig. S4(b)†).

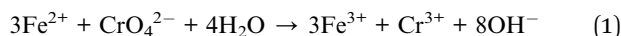
As shown in Fig. 5(e), the binding energies of Fe 2p after Cr adsorption at 724.6 eV, 720.4 eV, and 711.1 eV may be assigned to Fe(III), suggesting that Fe(II) on the surface of the material was oxidized to Fe(III) during the redox process. Hence, lower pH facilitated the transformation of Fe(III) to Fe(II) during the adsorption process. The measurements of the concentration of Fe(II) were performed in acidic solution (pH = 2) during the reaction (Fig. S4(a)†). Under H₂SO₄ treatment, the content of Fe(II) was considerably higher than that under HCl treatment



with and without Cr(vi), and the results showed that the content of Fe(II) ions reached about 108.3 mg L⁻¹ with no Cr(vi) in the H₂SO₄ system. When Cr(vi) was added, the concentration of Fe(II) ions decreased to about 9.24 mg L⁻¹ in 7 h.

These interesting findings indicate that Fe₃O₄/Fe₂N may have high reduction activity and can release a large amount of Fe(II) under H₂SO₄ treatment. As shown in Fig. 7, Fe₃O₄ alone has no effect on activation in the H₂SO₄ system. Also, both Fe₃O₄ and Fe₂N are inefficient in reducing Cr(vi) in the HCl system. Therefore, this suggests that the high efficiency of WT(NH₃) is due to the active sites at the interface of the two phases of Fe₂N and Fe₃O₄ or Fe₂N itself,⁵³ this is consistent with the TEM results, showing that WT(NH₃) has much more Fe₃O₄ and Fe₂N interfaces.

According to the above-mentioned analysis and discussion, a possible mechanism for Cr(vi) removal by WT(NH₃) is proposed (Fig. 9). First, Cr(vi) was partly absorbed by WT(NH₃) in both the HCl and H₂SO₄ systems. Second, WT(NH₃) released both Fe(III) and Fe(II). Meanwhile, some Fe(III) ions were converted to Fe(II) at a very low pH value. Third, the Cr(vi) ions were reduced to Cr(III) by Fe(II) (eqn (1)),³¹ which explained the presence of Cr(III) species in both the aqueous solution and on the surface of WT(NH₃). This was also supported by the decrease in the concentration of Fe(II) in the solution with Cr(vi). In addition, SO₄²⁻ could activate Fe₂N or the interface of Fe₂N and Fe₃O₄ and thus, it could accelerate the release of ferrous ions.



From the above-mentioned results, waste toner can be utilized to prepare functional nanomaterials under different conditions. As indicated in this study, the good sorptive and reductive capacities for CrO₄²⁻ are ascribed to the active sites and Fe₃O₄ and Fe₂N in the nanocomposites. Since functional magnetic Fe₃O₄ shows good affinity for many heavy metals such as lead,⁵⁴ arsenic⁵⁵ and copper,⁵⁶ the as-prepared toner-based sorbents can be used to remove a wide spectrum of heavy metals. Besides, the sorbents can be used to remediate wastewater contaminated by pharmaceuticals and other organic pollutants because Fe₂N may promote catalytic degradation of organic compounds.^{53,57} Thus, waste toner can be potentially used for the removal of heavy metals and organic contaminants.

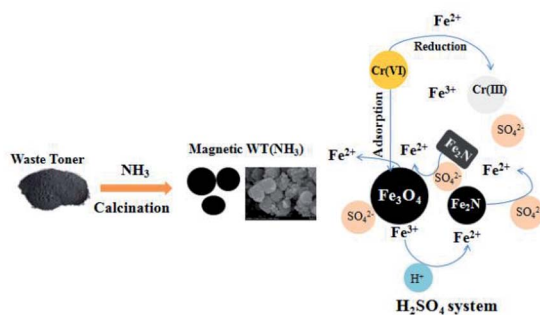


Fig. 9 Preparation route for magnetic WT(NH₃) and its mechanism for Cr(vi) removal in the H₂SO₄ system.

4. Conclusions

In this study, we prepared a novel magnetic nanocomposite using waste toner through high-temperature decomposition. The XRD analysis confirmed the existence of Fe₃O₄ and Fe₂N nanoparticles in WT(NH₃). Cr(vi) adsorption properties and magnetism characteristics of WT(NH₃) were investigated. WT(NH₃) exhibited high magnetization and good dispersibility. The enhanced efficiency for the removal of Cr(vi) can be related to its higher surface area and existence of Fe₂N. In fact, TEM results revealed that WT(NH₃) has more Fe₂N and Fe₃O₄ interfaces. The XPS analysis indicated that Cr(vi) was partly reduced to Cr(III) by Fe(II) during the process. Batch sorption experiments indicated that using H₂SO₄ to adjust pH could effectively help release a large amount of Fe(II) to reduce toxic Cr(vi) by WT(NH₃). Our study suggests that the facilely synthesized sorbent is efficient for the removal of Cr(vi) for environmental applications.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was financially supported by the Qing Lan Project, the National Science Foundation of China (31772394, 51602281), Social development project of Jiangsu Province (BE2015661), Six-talent peaks project in Jiangsu Province (2013-NY-017). We thank the Testing Center of Yangzhou University for Sample Characterization.

References

- 1 J. O. Nriagu and E. Nieboer, *J. Ecol.*, 1988, **71**, 1.
- 2 A. Broadway, M. R. Cave, J. Wragg, F. M. Fordyce, R. J. F. Bewley, M. C. Graham, B. T. Ngwenya and J. G. Farmer, *Sci. Total Environ.*, 2010, **409**, 267–277.
- 3 Y. T. Wang, E. M. Chirwa and H. Shen, *J. Environ. Eng.*, 2000, **126**, 300–306.
- 4 Agency, E. P. U. S., *Drinking Water Regulations and Health Advisories*, May 1995.
- 5 M. Dakiky, M. Khamis, A. Manassra and M. Mer'Eb, *Adv. Environ. Res.*, 2002, **6**, 533–540.
- 6 J. Kotaš and Z. Stasicka, *Environ. Pollut.*, 2000, **107**, 263–283.
- 7 M. K. Dinker and P. S. Kulkarni, *ChemInform*, 2015, **60**, 2521–2540.
- 8 E. A. Schmieman, J. N. Petersen, D. R. Yonge, D. L. Johnstone, Y. Bereded-Samuel, W. A. Apel and C. E. Turick, *Appl. Biochem. Biotechnol.*, 1997, **63**, 855–864.
- 9 A. E. Pagana and S. D. Sklari, *J. Membr. Sci.*, 2011, **367**, 319–324.
- 10 R. L. Goswamee, P. Sengupta, K. G. Bhattacharyya and D. K. Dutta, *Appl. Clay Sci.*, 1998, **13**, 21–34.
- 11 T. Ölmez, *J. Hazard. Mater.*, 2009, **162**, 1371–1378.
- 12 K. P. Singh, A. K. Singh, S. Gupta and S. Sinha, *Desalination*, 2011, **270**, 275–284.



- 13 Y. F. Kong, Y. C. Li, G. S. Hu, N. Cao, Y. Q. Ling, D. Pan, Q. Shao and Z. H. Guo, *Polym. Adv. Technol.*, 2018, 1–8.
- 14 Y. F. Kong, Y. C. Li, G. S. Hu, J. Lin, D. Pan, D. Y. Dong, E. Wujick, Q. Shao, M. J. Wu, J. Z. Zhao and Z. H. Guo, *Polymer*, 2018, **145**, 232–241.
- 15 H. J. Kang, Q. Shao, X. K. Guo, A. Galaska, Y. Y. Liu and Z. H. Guo, *Eng. Sci.*, 2018, **1**, 78–85.
- 16 J. Zheng, J. Ruan, L. Dong, T. Zhang, M. Huang and Z. Xu, *ACS Sustainable Chem. Eng.*, 2016, **5**, 161–167.
- 17 J. Ruan, J. Li and Z. Xu, *J. Hazard. Mater.*, 2011, **185**, 696–702.
- 18 J. Ruan, J. Li and Z. Xu, *Environ. Sci. Technol.*, 2013, **47**, 6457–6462.
- 19 J. Ruan, L. Dong, J. Huang, Z. Huang, K. Huang and H. Dong, *ACS Sustainable Chem. Eng.*, 2017, **5**, 4923–4929.
- 20 J. Ruan and Z. Xu, *Environ. Sci. Technol.*, 2012, **46**, 6214.
- 21 V. Gaikwad, U. Kumar, F. Pahlevani, A. Piadasa and V. Sahajwalla, *ACS Sustainable Chem. Eng.*, 2017, **5**, 11543–11550.
- 22 H. H. Hu, H. P. Liu, D. J. Zhang, J. J. Wang, G. W. Qin and X. F. Zhang, *Eng. Sci.*, 2018, **2**, 43–48.
- 23 H. K. Wu, Y. Zhang, R. Yin, W. Zhao, X. M. Li and L. Qian, *Adv. Compos Hybrid. Mater.*, 2018, **1**, 168–176.
- 24 Y. X. He, S. Yang, H. Liu, Q. Shao, Q. Y. Chen, C. Lu, Y. L. Jiang, C. T. Liu and Z. H. Guo, *J. Colloid Interface Sci.*, 2018, **517**, 40–51.
- 25 M. L. Peterson, A. F. White, G. E. Brown and G. A. Parks, *Environ. Sci. Technol.*, 1997, **31**, 1573–1576.
- 26 T. Kendelewicz, P. Liu, C. S. Doyle, G. E. Brown Jr, E. J. Nelson and S. A. Chambers, *Surf. Sci.*, 1999, **424**, 219–231.
- 27 Y. Gong, L. Gai, J. Tang, J. Fu, Q. Wang and E. Y. Zeng, *Sci. Total Environ.*, 2017, **595**, 743–751.
- 28 J. Pan, J. Jiang and R. K. Xu, *Chemosphere*, 2014, **101**, 71–76.
- 29 U. Schwertmann, U. Gasser and H. Sticher, *Geochim. Cosmochim. Acta*, 1989, **53**, 1293–1297.
- 30 B. A. Manning, J. R. Kiser, H. Kwon and S. R. Kanel, *Environ. Sci. Technol.*, 2007, **41**, 586–592.
- 31 B. Qiu, H. B. Gu, X. R. Yan, J. Guo, Y. R. Wang, D. Z. Sun, Q. Wang, M. Khan, X. Zhang, B. L. Weeks, D. P. Young, Z. H. Guo and S. Y. Wei, *J. Mater. Chem. A*, 2014, **2**, 17454–17462.
- 32 J. H. Zhu, H. B. Gu, J. Guo, M. J. Chen, H. G. Wei, Z. P. Luo, H. A. Colorado, N. Yerra, D. W. Ding, T. C. Ho, N. Haldolaarachchige, J. Hopper, D. P. Young, Z. H. Guo and S. Y. Wei, *J. Mater. Chem. A*, 2014, **2**, 2256–2265.
- 33 J. N. Huang, Y. H. Cao, Q. Shao, X. F. Peng and Z. H. Guo, *Ind. Eng. Chem. Res.*, 2017, **56**, 10689–10701.
- 34 B. Qiu, Y. R. Wang, D. Z. Sun, Q. Wang, X. Zhang, B. L. Weeks, R. O'Connor, X. H. Huang, S. Y. Wei and Z. H. Guo, *J. Mater. Chem. A*, 2015, **3**, 9817–9825.
- 35 G. Liu, Q. Deng, H. Wang, S. Kang, Y. Yang and D. H. L. Ng, *Chem.–Eur. J.*, 2012, **18**, 13418–13426.
- 36 S. C. Ponce, C. Prado, E. Pagano, F. E. Prado and M. Rosa, *Ecol. Eng.*, 2015, **74**, 33–41.
- 37 Q. Wang, X. Chen, K. Yu, Y. Zhang and Y. Cong, *J. Hazard. Mater.*, 2013, **246–247**, 135–144.
- 38 D. S. Sigman, *Acc. Chem. Res.*, 1989, **19**, 13–27.
- 39 A. Mirmohseni, M. S. S. Dorraji and M. G. Hosseini, *Electrochim. Acta*, 2012, **70**, 182–192.
- 40 X. Huang, Z. Yang, B. Dong, Y. Wang, T. Tang and Y. Hou, *Nanoscale*, 2017, **9**, 8102–8106.
- 41 L. Guo, P. Ye, J. Wang, F. Fu and Z. Wu, *J. Hazard. Mater.*, 2015, **298**, 28–35.
- 42 H. W. Liang, X. Zhuang, S. Brüller, X. Feng and K. Müllen, *Nat. Commun.*, 2014, **5**, 4973.
- 43 Z. Yue, S. E. Bender, J. Wang and J. Economy, *J. Hazard. Mater.*, 2009, **166**, 74–78.
- 44 L. Lin, Q. Zhu and A. W. Xu, *J. Am. Chem. Soc.*, 2014, **136**, 11027–11033.
- 45 D. Wilson and M. A. Langell, *Appl. Surf. Sci.*, 2014, **303**, 6–13.
- 46 Y. Tian, B. B. Yu, X. Li and K. Li, *J. Mater. Chem.*, 2011, **21**, 2476–2481.
- 47 F. Gao, H. Gu, H. Wang, X. Wang, B. Xiang and Z. Guo, *RSC Adv.*, 2015, **5**, 60208–60219.
- 48 S. Shi, J. Yang, S. Liang, M. Li, Q. Gan and K. Xiao, *Sci. Total Environ.*, 2018, **628–629**, 499–508.
- 49 T. Papadam, N. P. Xekoukoulotakis, I. Poullos and D. Mantzavinos, *J. Photochem. Photobiol., A*, 2007, **186**, 308–315.
- 50 J. Jiang, Q. Cai, W. Xu, M. W. Yang, Y. Cai, D. D. Dionysiou and K. E. O'Shea, *Environ. Sci. Technol.*, 2014, **48**, 8078–8085.
- 51 X. Tian, W. Wang, T. Na, C. Zhou, Y. Chao and S. Komarneni, *J. Hazard. Mater.*, 2016, **309**, 151–156.
- 52 Z. Ai, Y. Cheng, L. Zhang and J. Qiu, *Environ. Sci. Technol.*, 2008, **42**, 6955–6960.
- 53 J. Wang, C. Wang, S. Tong, J. Wang, C. Wang and S. Tong, *Catal. Commun.*, 2017, **103**, 105–109.
- 54 L. Tan, J. Xu, X. Xue, Z. Lou, J. Zhu and S. A. Baig, *RSC Adv.*, 2014, **4**, 45920–45929.
- 55 L. Y. Feng, M. H. Cao, X. Y. Ma, Y. S. Zhu and C. W. Hu, *J. Hazard. Mater.*, 2012, **217**, 439–446.
- 56 H. B. Hu, Z. H. Wang and L. J. Pan, *J. Alloys Compd.*, 2010, **492**, 656–661.
- 57 N. Zhang, H. Zhao, G. M. Zhang, S. Chong, Y. C. Liu, L. Y. Sun, H. Z. Chang and T. Huang, *J. Environ. Manage.*, 2017, **187**, 201–211.

