



Cite this: *RSC Adv.*, 2018, 8, 5180

Novel magnetic g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composite for the very effective visible-light-Fenton degradation of Orange II

Zhuliang Wang,^{ab} Yiping Fan,^a Rong Wu,^a Yaoxing Huo,^a Hao Wu,^{id} *^{ab} Fang Wang^{ab} and Xiaohong Xu^{id} *^{ab}

A novel magnetic heterogeneous g-C₃N₄/α-Fe₂O₃/Fe₃O₄ catalyst was successfully synthesized through a simple hydrothermal method. The structure, morphology, and optical properties of the catalyst were characterized. The photocatalytic activity of the heterogeneous g-C₃N₄/α-Fe₂O₃/Fe₃O₄ catalyst for the photo-Fenton degradation of Orange II in the presence of H₂O₂ irradiated with visible light (λ > 420 nm) at neutral pH was evaluated. The g-C₃N₄/α-Fe₂O₃/Fe₃O₄ photocatalyst was found to be an excellent catalyst for the degradation of Orange II and offers great advantages over the traditional Fenton system (Fe(II/III)/H₂O₂). The results indicated that successfully combining monodispersed Fe₃O₄ nanoparticles and g-C₃N₄/α-Fe₂O₃ enhanced light harvesting, retarded photogenerated electron–hole recombination, and significantly enhanced the photocatalytic activity of the system. The g-C₃N₄/α-Fe₂O₃/Fe₃O₄ (30%) sample gave the highest degradation rate constant, 0.091 min⁻¹, which was almost 4.01 times higher than the degradation rate constant for α-Fe₂O₃ and 2.65 times higher than the degradation rate constant for g-C₃N₄/α-Fe₂O₃ under the same conditions. A reasonable mechanism for catalysis by the g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composite was developed. The g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composite was found to be stable and recyclable, meaning it has great potential for use as a photo-Fenton catalyst for effectively degrading organic pollutants in wastewater.

Received 13th December 2017
 Accepted 20th January 2018

DOI: 10.1039/c7ra13291c

rsc.li/rsc-advances

1. Introduction

Increasing amounts of synthetic organic dye pollutants are being released into the environment after synthetic dyes have been used in the textile, leather, pharmaceutical, food, and other industries.¹ Advanced oxidation processes including ozonation, Fenton reactions, photo-Fenton reactions, photocatalysis, ultrasonication, and other processes can potentially be used to degrade recalcitrant organic contaminants because total mineralization is more likely to be achieved using advanced oxidation processes than using classical wastewater treatment processes.^{2–7} The Fenton reaction, an advanced oxidation process, involves producing radicals using homogeneous Fe(II/III) and H₂O₂. The Fenton reaction has been shown to be an effective treatment for wastewater produced by various industries because it is fast, simple to perform, and does not produce toxic products.^{8,9} However, there are some drawbacks to

homogeneous processes such as the Fenton reaction in that sludge is generated, the pH must be adjusted before and after the reaction, and the catalyst is lost in the effluent. These drawbacks limit the applications of the Fenton reaction.¹⁰ Using a solid catalyst in a “heterogeneous Fenton oxidation reaction” may solve these problems.^{11,12}

Hematite (α-Fe₂O₃), a common semiconductor with a narrow band gap of 1.9–2.1 eV, has been used as a heterogeneous photo-Fenton catalyst. α-Fe₂O₃ is easy and cheap to produce, nontoxic, and can be used over a wide pH range.¹³ The photocatalytic degradation efficiency achieved using α-Fe₂O₃ can be markedly increased by coupling the α-Fe₂O₃ with other oxides or semiconductors.^{14–20} Graphitic carbon nitride (g-C₃N₄), which has been studied in recent years, is a typical metal-free polymeric semiconductor that is very chemically stable and environmentally benign, and it has been found to have a band gap of 2.7 eV, making it suitable for absorbing visible light. g-C₃N₄ is therefore a promising candidate for coupling with α-Fe₂O₃ to form a heterogeneous photo-Fenton catalyst. Many attempts have been made to fabricate α-Fe₂O₃/g-C₃N₄ composites for photodegrading organic dyes using visible light.^{21,22} However, the g-C₃N₄/α-Fe₂O₃ heterogeneous catalyst is not appropriate for practical use because it is difficult to remove from suspension and reuse. Magnetically separable photocatalysts have

^aSchool of Chemistry and Materials Science of Shanxi Normal University, Key Laboratory of Magnetic Molecules and Magnetic Information Materials of Ministry of Education, Linfen, 041004, China. E-mail: xuxh@sxnu.edu.cn; wuhao@sxnu.edu.cn

^bResearch Institute of Materials Science of Shanxi Normal University, Collaborative Innovation Center for Shanxi Advanced Permanent Magnetic Materials and Technology, Linfen, 041004, China



attracted considerable interest because they can be removed from suspensions easily.

Many magnetic materials are available (*e.g.*, Fe_3O_4 , $\gamma\text{-Fe}_2\text{O}_3$, and MFe_2O_4 , where M is Ba^{2+} , Co^{2+} , Mg^{2+} , Mn^{2+} , Ni^{2+} , or Zn^{2+}). Magnetite (Fe_3O_4) is one of the most popular materials for producing magnetically recoverable catalysts because it is cheap, has a desirable degree of magnetism, and is nontoxic.²³ However, Fe_3O_4 is very conductive, so it can transfer photo-generated electrons rapidly, improving charge carrier separation efficiency.²⁴ It has recently been found that immobilizing a photocatalyst using Fe_3O_4 nanoparticles can effectively increase the cyclic utilization rate and improve the photodegradation efficiency of the catalyst.^{25–29} The preparation of magnetically separable visible-light-driven photocatalysts based on Fe_3O_4 is an active research field. However, to the best of our knowledge, $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composites have not previously been prepared or their photo-Fenton catalytic properties investigated.

In the work presented here a novel magnetic $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composite was synthesized using a simple hydrothermal method. The heterogeneous photo-Fenton activity of the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composite was investigated using the composite to degrade Orange II, a typical recalcitrant organic contaminant. The effects of the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$ to Fe_3O_4 weight ratio, scavengers of reactive species, and the role of hydrogen peroxide (H_2O_2) on the photocatalytic activity of the composite were investigated in detail. The results demonstrated that successfully combining monodispersed Fe_3O_4 nanoparticles and $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$ enhanced light harvesting, retarded photo-generated electron-hole recombination, and significantly enhanced the photocatalytic activity of the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$. The magnetic recovery and reusability of the composite were also investigated, and the results indicated that the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composite will be a valuable photocatalyst that could be used to organic dye wastewater treatment.

2. Experimental

2.1. Chemical reagents and materials

Ferric chloride ($\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ 99%) and polyethylene glycol 4000 (PEG-4000) were purchased from Shanghai Chemical Reagents Company (Shanghai, China). Urea (99%) was purchased from Kermel (Tianjin, China) and NaBH_4 (AR) was purchased from Da mao (Tianjin, China). TCNE (98%), Orange II (99.9%, $\text{C}_{16}\text{H}_{11}\text{N}_2\text{NaO}_4\text{S}$) and sodium acetate (NaAc 99%) were provided by Acros Organics (New Jersey, USA). Ethylene glycol (EG \geq 99%) was purchased from Sigma-Aldrich. Coumarin (COU 99%) was purchased from J&K Scientific Ltd. (Beijing China). All chemicals were analytical grade and used as received without further purification.

2.2. Characterization

The crystal phases of the sample were analyzed by X-ray diffraction (XRD: Rigaku Ultima IV, Japan). A scanning electron microscope (SEM, JSM-7500F, JEOL, Japan) was used to characterize the morphology of the obtained products. The

composite of the sample was investigated by the energy dispersive spectroscopy (EDS) attached to the SEM. The structure and morphology of the products were examined by transmission electron microscopy (TEM: JEOL2100, Japan). Fourier transform infrared spectra (FT-IR) were measured using Spectrum One infrared spectrometer with KBr as the reference. The photoluminescence spectra (PL) of samples were measured with a fluorescence spectrophotometer (LSP920, UK) using a Xe lamp as an excitation source with optical filters. UV-vis diffuse reflectance spectroscopy was recorded with a spectrophotometer (UV-2700, Shimadzu Japan). The PL spectra of hydroxyl radical ($\cdot\text{OH}$) detection were measured by a fluorescence spectrophotometer (Cray Eclipse, Agilent Technologies Australia). The specific surface area and pore size distributions were measured with liquid nitrogen at 77 K using a Micromeritics ASAP 2020 instrument *via* the Brunauer–Emmett–Teller (BET) method (Quantachrome, USA). The magnetic property was measured by Physical Property Measurement System (PPMS, XL-5 Quantum Design USA) at room temperature.

2.3. Preparation of the samples

2.3.1. Synthesis of $g\text{-C}_3\text{N}_4$. Graphitic carbon nitride was prepared using a thermo-induced copolymerization method using urea as the precursor.³⁰ Urea (30 g) and tetracyanoethylene (40 mg) were mixed carefully, then the mixture was placed in a crucible with a loose cover. The crucible was placed in a muffle furnace, and the temperature was increased at $20\text{ }^\circ\text{C min}^{-1}$ to $550\text{ }^\circ\text{C}$, then held for 4 h. The crucible was cooled to room temperature, then the $g\text{-C}_3\text{N}_4$ produced was ground to a powder to prepare it for further use.

2.3.2. Synthesis of Fe_3O_4 . The Fe_3O_4 was synthesized using a solvothermal method in our previous report.³¹ $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ (0.6757 g) was dissolved in ethylene glycol (20 mL) in a sonication bath to give a yellow transparent solution. Anhydrous sodium acetate (1.6404 g) was added, and the mixture was heated until the reactants were fully dissolved, then polyethylene glycol 4000 (0.3 g) was added. The mixture was transferred to a Teflon-lined stainless steel autoclave (50 mL capacity), which was heated to $200\text{ }^\circ\text{C}$ and kept at that temperature for 12 h. The autoclave was allowed to cool naturally to room temperature. The products were separated using a magnet, rinsed several times with deionized water and then ethanol, and then dried at $60\text{ }^\circ\text{C}$ for 24 h.

2.3.3. Preparation of $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composites. The $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composites were prepared using borohydride (NaBH_4) reduction method and hydrothermal method. First, as-prepared Fe_3O_4 (an appropriate amount for the desired product) and $g\text{-C}_3\text{N}_4$ (0.3 g) were dispersed in deionized water (12.5 mL) with ultrasonication, then $\text{FeCl}_3 \cdot 9\text{H}_2\text{O}$ (0.6757 g) was dissolved in the suspension with stirring. Second, a NaBH_4 solution which pre-prepared by dissolving solid NaBH_4 (0.1892 g) in deionized water (12.5 mL) was quickly added to the suspension described above, and the mixture was vigorously stirred for 1 h. In this process, the ferric ion (Fe^{3+}) was reduced to zero-valent iron (Fe^0).³² Third, the above mixture was then transferred to a Teflon-lined stainless steel autoclave



(50 mL capacity), which was heated to 120 °C for 12 h to allow the Fe⁰ oxidized to α -Fe₂O₃,³³ then the desired g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ composite was formed. Last, the suspension produced was washed several times with deionized water and ethanol, then dried at 60 °C. Composite samples with different Fe₃O₄ mass fractions were produced, and they are later labeled g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ (*x*), where *x* is the Fe₃O₄ mass percent in the g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ composite. Samples with Fe₃O₄ mass percentages of 10%, 20%, 30%, 40%, and 45% were prepared using the procedure described above. A schematic diagram of the growth process of the g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ composites is shown in Fig. 1.

2.4. Photodegradation of Orange II

The photocatalytic efficiencies of the samples that were prepared were evaluated using the samples to degrade Orange II in the presence of H₂O₂ and irradiated with visible light produced using a Xe light (500 W) with a 420 nm cut-off filter. For a degradation experiment, a 50 mg L⁻¹ (the initial concentration, C₀) stock solution (100 mL) of Orange II was placed in a glass beaker, then the catalyst being tested was added and the mixture was put in a dark place and constantly stirred using a magnetic stirrer for 30 min to allow absorption-desorption equilibrium to be reached. H₂O₂ (0.1 mL) was then added and the mixture was immediately irradiated with visible light. All the tests were performed at a constant temperature of 25 ± 0.2 °C, achieved using a water bath. A 3.0 mL aliquot of the solution was removed at each of a series of specified times, and absorbance at 484 nm was measured using a UV-visible spectrophotometer.

2.5. Measurement of ·OH

The formation of ·OH was detected by the PL technique using COU as a probe molecule. Experimental procedures were as follows: 0.1 g of g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ composites was dispersed in 100 mL of 1 × 10⁻³ M COU aqueous solution in a glass beaker, then the solution was constantly stirred using a magnetic stirrer for 30 min to allow absorption-desorption equilibrium to be reached. H₂O₂ (0.1 mL) was then added and

the mixture was immediately irradiated with visible light using a Xe light (500 W) with a 420 nm cut-off filter. All the tests were performed at a constant temperature of 25 ± 0.2 °C, achieved using a water bath. A 3.0 mL aliquot of the solution was removed at each of 15 min, and the reaction solution was filtrated to measure the PL intensity at 456 nm by an Agilent Technologies Cary Eclipse Fluorescence Spectrophotometer. The excitation wavelength was 332 nm, the scanning speed was 120 nm min⁻¹. The width of excitation and emission slit was set to be both 5 nm.

3. Results and discussion

The X-ray diffraction patterns of the g-C₃N₄, α -Fe₂O₃, Fe₃O₄, and g-C₃N₄/ α -Fe₂O₃ materials and the g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ composites with different Fe₃O₄ contents are shown in Fig. 2a. The g-C₃N₄ gave a pronounced peak at 27.5° indexed as a (002) diffraction plane corresponding to the characteristic interlayer stacking peak of a conjugated aromatic system (JCPDS files no. 87-1526).³⁴ The positions and relative intensities of all the diffraction peaks for the Fe₃O₄ sample matched the data for pure face-centered cubic Fe₃O₄, and the diffraction peaks in the range 30.6–63.2° were indexed as (220), (311), (400), (422), (511), and (440) reflections (JCPDS files no. 19-0629).³⁵ Diffraction peaks were found for α -Fe₂O₃ at 24.2°, 33.3°, 35.7°, 40.9°, and 49.5°, and these were indexed as (012), (104), (110), (113), and (024) planes of α -Fe₂O₃, respectively (JCPDS files no. 33-0664).¹⁶ The g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ composite X-ray diffraction patterns contained the diffraction peaks for g-C₃N₄, α -Fe₂O₃, and Fe₃O₄, and the intensity of the Fe₃O₄ peak increased as the Fe₃O₄ weight percent increased. This suggested that the desired g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ composites were successfully prepared using the hydrothermal method.

Fourier-transform infrared spectra were acquired to characterize the structural properties of the samples, and the results are shown in Fig. 2b. The g-C₃N₄ sample gave a broad band at 3400 cm⁻¹ indicative of N–H stretching vibrations and a band in the range 1214–1634 cm⁻¹ corresponding to heterocycle C–N and C=N stretching vibrations.³⁶ A breathing mode at 806 cm⁻¹ was related to typical *s*-triazine unit bending vibrations.³⁷ The composites containing Fe₃O₄ and α -Fe₂O₃ had characteristic peaks at 470 and 560 cm⁻¹ attributed to Fe–O stretching.^{38,39} It can be seen that the main characteristic peaks of g-C₃N₄, Fe₃O₄, and α -Fe₂O₃ were found in the g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ composite spectra, further demonstrating that α -Fe₂O₃ and Fe₃O₄ had been deposited on the surfaces of the g-C₃N₄ sheets.

The morphologies of the samples and the ways in which the samples formed were investigated using scanning electron microscopy and transmission electron microscopy. Scanning electron microscopy images for g-C₃N₄, Fe₃O₄, and g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ (30%) are shown in Fig. 3a–c, respectively. The pristine g-C₃N₄ sample had a sheet-like structure. The Fe₃O₄ nanoparticles were nearly spherical. In contrast, the g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ composite produced using the 12 h hydrothermal reaction process was irregular and rough in shape because of the Fe₃O₄ and α -Fe₂O₃ deposited on the g-C₃N₄ surfaces. It can be seen from the transmission electron microscopy images

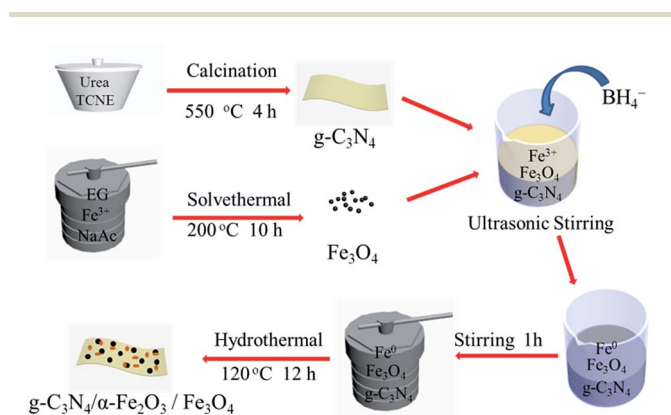


Fig. 1 The schematic diagram for growth process of the g-C₃N₄/ α -Fe₂O₃/Fe₃O₄ composites.



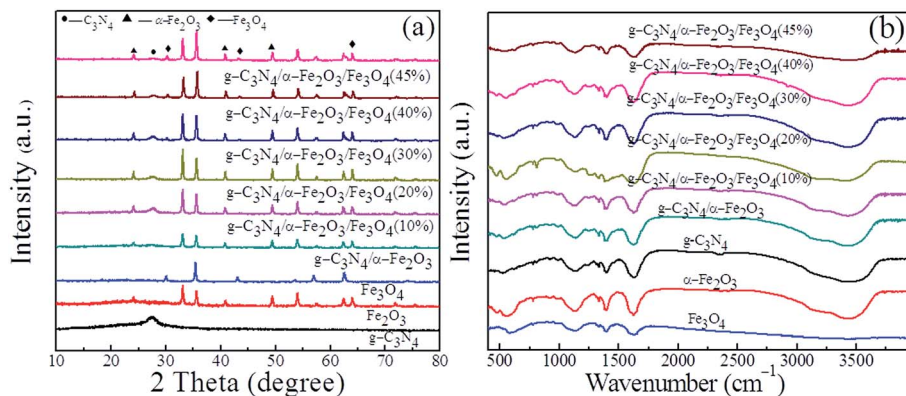


Fig. 2 (a) XRD patterns and (b) FT-IR spectra of $g\text{-C}_3\text{N}_4$, Fe_3O_4 , $\alpha\text{-Fe}_2\text{O}_3$ and $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ samples.

shown in Fig. 3d and e that the pristine $g\text{-C}_3\text{N}_4$ had typical two-dimensional sheet-like nanosheets and that the Fe_3O_4 were very dispersible. It can clearly be seen from the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ transmission electron microscopy image shown in Fig. 3f that Fe_3O_4 and $\alpha\text{-Fe}_2\text{O}_3$ particles were uniformly distributed over the $g\text{-C}_3\text{N}_4$ sheet surfaces and a sandwich-like structure was formed. This unique architecture can greatly promote separation efficiency of electron-hole pairs. High resolution

transmission electron microscopy (HRTEM) image of $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ is shown in Fig. 3g. It can be noted that there are two different lattice fringes on the surface of $g\text{-C}_3\text{N}_4$, the lattice fringe of 0.271 nm and 0.253 nm in the observed nanocrystallites agrees well with the (104) lattice planes of $\alpha\text{-Fe}_2\text{O}_3$ and the (311) lattice planes of Fe_3O_4 , respectively, which indicates the existence of the heterojunction structure between $g\text{-C}_3\text{N}_4$, $\alpha\text{-Fe}_2\text{O}_3$ and Fe_3O_4 .^{40,41}

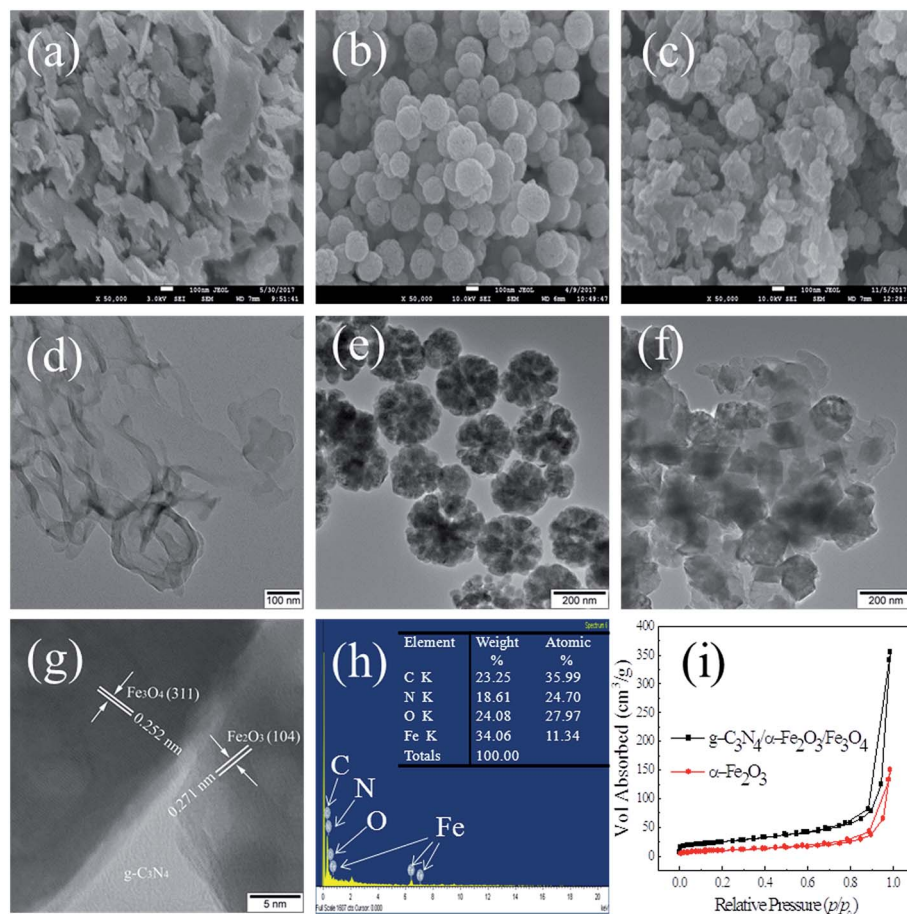


Fig. 3 SEM (a–c), TEM (d–f), HRTEM (g), EDX (h), and N_2 adsorption–desorption isotherm (i) of the samples.



Energy dispersive spectroscopy was performed to determine the elemental composition of the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ (30%) sample, and the results are shown in Fig. 3d. The peaks were ascribed to the elements C, N, O, and Fe and agreed with the X-ray diffraction patterns, further confirming the presence of $g\text{-C}_3\text{N}_4$, Fe_3O_4 , and $\alpha\text{-Fe}_2\text{O}_3$ particles in the composite. The specific surface areas of the $\alpha\text{-Fe}_2\text{O}_3$ and the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ (30%) composite were determined by acquiring N_2 adsorption/desorption isotherms (Fig. 2h). Both samples gave type IV adsorption-desorption isotherms according to the Brunauer-Deming-Deeming-Teller classification. Type IV adsorption-desorption isotherms are typical of materials with mesoporous domains. The BET surface area of $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ (30%) was $88.37\text{ m}^2\text{ g}^{-1}$, which was much higher than the surface area of $\alpha\text{-Fe}_2\text{O}_3$ ($38.54\text{ m}^2\text{ g}^{-1}$). Importantly, the pore volume was also higher for $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ (30%), at $0.55\text{ cm}^3\text{ g}^{-1}$, than for $\alpha\text{-Fe}_2\text{O}_3$ ($0.23\text{ cm}^3\text{ g}^{-1}$). The high pore volume of $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ (30%) would have facilitated charge separation in the composite system and made the composite suitable for heterogeneous photo-Fenton applications.²⁴

It is well known that enhancing absorption in the visible-light region improves the activity of a photocatalyst. The UV-vis diffuse reflection spectra of $\alpha\text{-Fe}_2\text{O}_3$, $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$, and $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ with different Fe_3O_4 weight percentages were acquired and are shown in Fig. 4a. $\alpha\text{-Fe}_2\text{O}_3$ had an absorption threshold at 657 nm associated with a band gap energy of 1.9 eV, which agreed with the previously reported band gap energy.⁴² $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$ had an absorption threshold at 617 nm associated with a band gap energy of 2.01 eV, which was lower than the band gap energy of pure $g\text{-C}_3\text{N}_4$ ($\sim 2.7\text{ eV}$). The $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ (10%), $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ (30%), and $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ (45%) absorption thresholds were at 627, 661, and 669 nm, respectively, and the band gap energies were 1.98, 1.88, and 1.85 eV, respectively. These results indicate that the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composite absorption edge shifted to a longer wavelength and the ability to absorb visible light improved as the Fe_3O_4 to $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$ weight ratio increased. The increase in visible light absorption may have been caused by heterogeneous structures forming. The results imply that the response of the photocatalyst to visible light was improved by loading Fe_3O_4 and $\alpha\text{-Fe}_2\text{O}_3$ onto the $g\text{-C}_3\text{N}_4$ sheets, which may

have increased the photocatalytic activity by allowing more photogenerated charge carriers to be formed.

The migration, transfer, and recombination processes for the photogenerated electron-hole pairs in the samples were investigated by acquiring room temperature PL spectra of $\alpha\text{-Fe}_2\text{O}_3$, $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$, and the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composites. The spectra are shown in Fig. 4b. No obvious $\alpha\text{-Fe}_2\text{O}_3$ emission peak was found because the $\alpha\text{-Fe}_2\text{O}_3$ peak intensity was much weaker than the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$ and $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composite peak intensities.²¹ The main $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$ and $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composite emission bands were centered on 480 nm, and the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composite emission peaks were much weaker than the pure $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$ emission peak, indicating that the recombination of photogenerated electron-hole pairs was efficiently hampered. The emission peak became weaker as the amount of Fe_3O_4 present increased, meaning the lifetimes of the photogenerated charge carriers would have increased as the amount of Fe_3O_4 present increased.

The photo-Fenton photocatalytic activities of $\alpha\text{-Fe}_2\text{O}_3$, $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$, and $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ composites were determined using the catalysts to decompose Orange II when irradiated with visible light ($\lambda > 420\text{ nm}$). As shown in Fig. 5a, Orange II was degraded by each sample, and the photocatalytic activities of the composites were closely related to the compositions of the composites. All the $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ samples had stronger photocatalytic activities than did $\alpha\text{-Fe}_2\text{O}_3$ and $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$. The enhanced photocatalytic activities of the composites could have been caused by the composites having higher surface areas, higher visible-light absorption capacities, and higher electron-hole separation efficiencies.

The experimental data were analyzed to investigate the Orange II degradation kinetics, and it was found that a pseudo-first-order model fitted the data. The formula for the pseudo-first-order model is

$$\ln(c_0/c) = kt,$$

where c_0 is the initial Orange II concentration in the aqueous solution, c is the actual Orange II concentration at the specified reaction time, k is the degradation rate constant, and t is the reaction time. The rate constants k for the different samples are

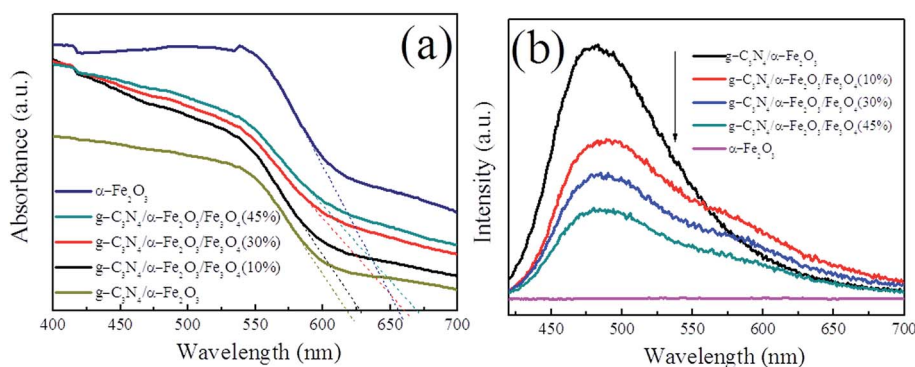


Fig. 4 UV-vis DRS (a) and PL spectra (b) of $\alpha\text{-Fe}_2\text{O}_3$, $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3$, and $g\text{-C}_3\text{N}_4/\alpha\text{-Fe}_2\text{O}_3/\text{Fe}_3\text{O}_4$ samples.



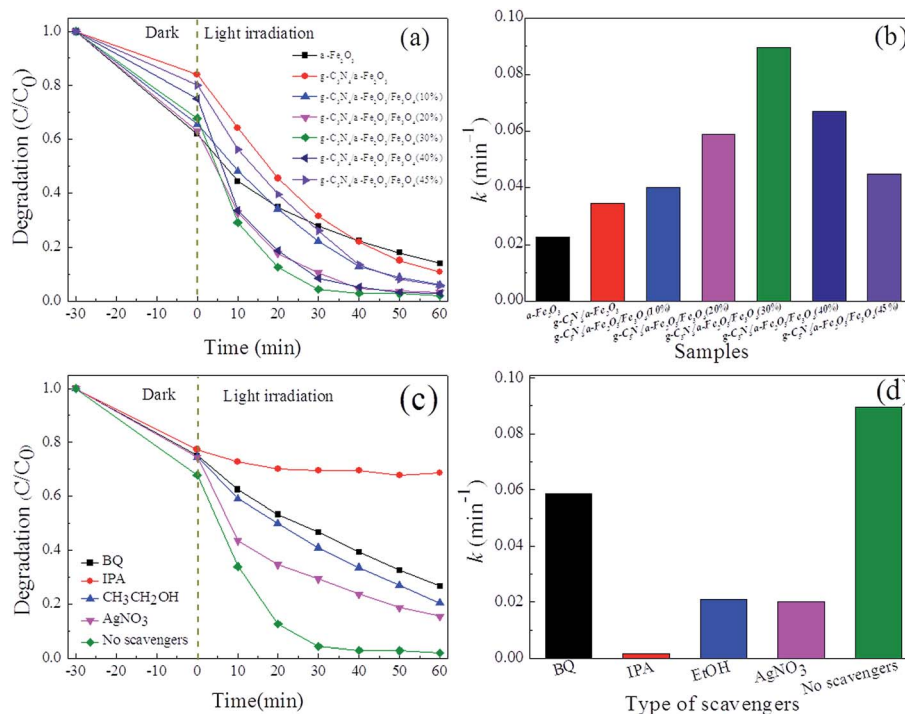


Fig. 5 (a) Degradation rate and (b) kinetic constant of all samples, (c) degradation rate and (d) kinetic constant of g-C₃N₄/α-Fe₂O₃/Fe₃O₄ (30%) samples with different scavengers.

shown in Fig. 5b. It is worth noting that the photocatalytic activities of the samples first increased and then decreased as the amount of Fe₃O₄ present increased. The g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composite with a Fe₃O₄ mass content of 30% had the highest rate constant, 0.091 min⁻¹, which was almost 4.01 times higher than the rate constant for α-Fe₂O₃ and 2.65 times higher than the rate constant for g-C₃N₄/α-Fe₂O₃. This may have been because combining a little Fe₃O₄ with α-Fe₂O₃ and g-C₃N₄ could allow heterojunctions to form, giving the composite a lower rate of electron-hole pair recombination. This would have meant that more h⁺ and e⁻ would have been generated to participate in the degradation process, enhancing the photocatalytic activity.⁴³ However, increasing the amount of Fe₃O₄ present further would have allowed some Fe₃O₄ to act as a recombination center for photoinduced electrons and holes,³⁴ and may have caused the active sites on the g-C₃N₄ surfaces to be covered, decreasing the photocatalytic activity of the composite. It is therefore very important that the most appropriate degree of doping is achieved to improve photocatalytic activity.

The reactive species trapping experiments were performed to evaluate role of the active species in the photocatalytic process. 1,4-Benzoquinone (BQ), isopropanol (IPA), ethanol (EtOH), and silver nitrate (AgNO₃) were used to capture superoxide radicals ([•]O₂⁻), hydroxyl radicals ([•]OH), holes (h⁺), and electrons (e⁻), respectively.^{44,45} The results of degradation rate and kinetic constant of g-C₃N₄/α-Fe₂O₃/Fe₃O₄ (30%) samples with different scavengers are shown in Fig. 5c and d. It can be seen that using IPA as a radical scavenger markedly decreased the degradation rate, indicating that [•]OH radicals were involved in the

photocatalytic process. Likewise, using EtOH, BQ, and AgNO₃ decreased the degradation rate to some extent, indicating that [•]O₂⁻, h⁺, and e⁻ also played roles in the photocatalytic process.

To further detect the formation of [•]OH in our visible-light-Fenton system, the PL technique was performed using COU as a probe molecule according to the published literatures.^{46,47} Because of COU can readily react with [•]OH to produce highly fluorescent product, 7-hydroxycoumarin (7HC), we can detect the formation of [•]OH by the PL spectra of 7HC in an aqueous solution containing g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composite, COU and H₂O₂ under visible light irradiation. As shown in Fig. 6a, a gradual increase in the PL intensity at about 456 nm was observed with increasing irradiation time, which indicated that fluorescent product 7HC was formed during visible-light-Fenton process due to the specific reaction between [•]OH and COU. Thus, it could be inferred that the [•]OH is producing at the g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composites surface, which is consistent with the results of reactive species trapping experiments.

The results described above led us to propose the catalytic mechanism for the g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composite shown in Fig. 6b. There are five main possible aspects to the Orange II catalytic degradation process. (i) Both α-Fe₂O₃ and g-C₃N₄ can efficiently absorb visible light and form photogenerated electron-hole pairs (eqn (1)) because the g-C₃N₄ and α-Fe₂O₃ band gap energies are about 2.70 and 1.90 eV, respectively. (ii) The Fe(III) on the α-Fe₂O₃ surfaces may be reduced to Fe(II) by excited electrons, then Fe(II) could react with H₂O₂ to form the Fenton system, generating [•]HO radicals (eqn (3)). (iii) The g-C₃N₄ conduction band bottom level is more negative than the O₂/[•]O₂⁻ redox potential (-0.16 eV),⁴⁸ so the photogenerated



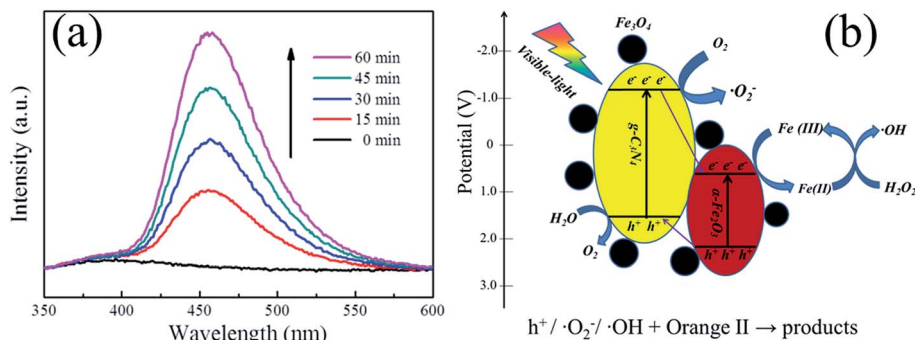
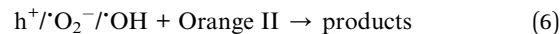
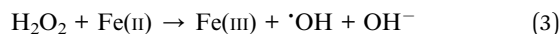
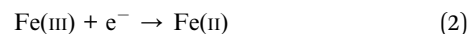
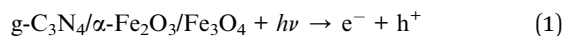


Fig. 6 (a) PL spectra of COU solution with g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composite in the presence of H₂O₂ under visible light irradiation. (b) Possible mechanism for the enhanced activity of g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composite.

electrons in the g-C₃N₄ conduction band could react with O₂ on the photocatalyst surfaces to form $\cdot\text{O}_2^-$ radicals (eqn (4)). (iv) The presence of heterojunctions between α-Fe₂O₃ and g-C₃N₄ may allow the holes generated in the α-Fe₂O₃ valence band to be transferred to the g-C₃N₄ valence band. The photogenerated holes could react with H₂O to produce O₂ (eqn (5)), and the O₂ generated could react with electrons to generate $\cdot\text{O}_2^-$. Likewise, the electrons generated in the g-C₃N₄ conduction band could be transferred to the α-Fe₂O₃ conduction band, reducing more Fe(III) to Fe(II) on the α-Fe₂O₃ surfaces, directly contributing to the production of $\cdot\text{HO}$ radicals.

The g-C₃N₄/α-Fe₂O₃/Fe₃O₄ catalyst also had a high surface area to which Orange II could adsorb, benefiting the degradation process. It is worth noting that the high conductivity of Fe₃O₄ will mean that some Fe₃O₄ between the α-Fe₂O₃ and g-C₃N₄ nanosheet interfaces could act as a medium for the transmission of electrons and holes between g-C₃N₄ and α-Fe₂O₃, strongly decreasing the probability that photo-generated electron-hole pairs in the g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composite system will recombine. This will cause more electrons and holes to form active species that could be involved in the photodegradation of Orange II. This photo-Fenton system works through $\cdot\text{OH}$, $\cdot\text{O}_2^-$, and holes that all acting as reactive species (eqn (6)), significantly increasing the photocatalytic activity relative to a traditional Fenton system.



The magnetic properties of Fe₂O₃, Fe₃O₄ and the g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composites were measured at room temperature, and the results are shown in Fig. 7a. The α-Fe₂O₃ magnetization curve showed that α-Fe₂O₃ was almost non-magnetic. Saturation magnetization for Fe₃O₄ and g-C₃N₄/α-Fe₂O₃/Fe₃O₄ was found at about 83.58 and 17.98 emu g⁻¹, respectively. The saturated magnetization was lower for g-C₃N₄/α-Fe₂O₃/Fe₃O₄ than for the Fe₃O₄ nanoparticles because of the presence of nonmagnetic g-C₃N₄ and α-Fe₂O₃. Separation of the magnetic photocatalyst from a treated solution using a magnet is shown in the inset in Fig. 7a. We concluded that the nanocomposite was magnetic enough to allow the composite to be magnetically separated from a treated solution.

The reusability of a photocatalyst is an important parameter from the economic viewpoint. Recycling tests were conducted to

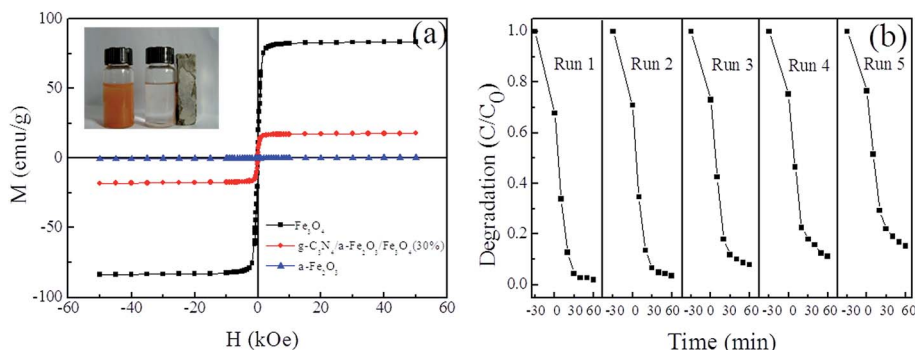


Fig. 7 (a) Magnetization curves for the α-Fe₂O₃, Fe₃O₄ nanoparticles and g-C₃N₄/α-Fe₂O₃/Fe₃O₄ (30%) composite. Inset of the figure shows separation of the composite from the treated solution using a magnet. (b) Reusability of the nanocomposite after five successive runs.



investigate the stability of g-C₃N₄/α-Fe₂O₃/Fe₃O₄, and the results are shown in Fig. 7b. More than 85% of the original photocatalytic activity of g-C₃N₄/α-Fe₂O₃/Fe₃O₄ (30%) was retained after five successive experimental runs using the same experimental conditions. The photocatalytic activity may have decreased with repeated use because of photocorrosion of the nanocomposite. These results indicated that the g-C₃N₄/α-Fe₂O₃/Fe₃O₄ catalyst is quite stable and could be reused.

4. Conclusions

Magnetic heterogeneous g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composites with different Fe₃O₄ mass percentages were successfully fabricated via a facile hydrothermal synthesis route. Combining the Fenton and photocatalytic processes in one reaction system gave the composite excellent catalytic activity. Of the samples that were prepared, g-C₃N₄/α-Fe₂O₃/Fe₃O₄ (30%) had the best catalytic activity. The enhanced photocatalytic activities could be ascribed to increases in the surface area, visible light absorption capacity, and electron-hole separation efficiency, leading to more h⁺ and e⁻ being generated to participate in the degradation process. A possible catalytic mechanism for the g-C₃N₄/α-Fe₂O₃/Fe₃O₄ composite was proposed. This photo-Fenton system works through ·OH, ·O₂⁻, and holes that all acting as reactive species, significantly increasing the photocatalytic activity relative to a traditional Fenton system. The g-C₃N₄/α-Fe₂O₃/Fe₃O₄ catalyst could be reused and was magnetic enough to be easily separated from a treated solution. The simply prepared magnetic heterogeneous composite was found to be an effective visible-light-driven photo-Fenton catalyst that may be useful in environmental applications.

Conflicts of interest

The authors declare no competing financial interest.

Acknowledgements

This work was supported by the National Science Foundation of China (Grant No. 61434002), the Special Funds of Sanjin Scholars Program.

References

- H. Lin, H. Zhang, X. Wang, L. Wang and J. Wu, *Sep. Purif. Technol.*, 2014, **122**, 533–540.
- M. A. Oturan and J.-J. Aaron, *Crit. Rev. Environ. Sci. Technol.*, 2014, **44**, 2577–2641.
- L. Clarizia, D. Russo, I. Di Somma, R. Marotta and R. Andreozzi, *Appl. Catal., B*, 2017, **209**, 358–371.
- S. Rahim Pouran, A. R. Abdul Aziz and W. M. A. Wan Daud, *J. Ind. Eng. Chem.*, 2015, **21**, 53–69.
- M. Sayed, J. A. Khan, L. A. Shah, N. S. Shah, F. Shah, H. M. Khan, P. Zhang and H. Arandiyani, *J. Phys. Chem. C*, 2017, **122**, 406–421.
- M. Sayed, L. A. Shah, J. A. Khan, N. S. Shah, J. Nisar, H. M. Khan, P. Zhang and A. R. Khan, *J. Phys. Chem. A*, 2016, **120**, 9916–9931.
- A. M. Khan, A. Mehmood, M. Sayed, M. F. Nazar, B. Ismail, R. A. Khan, H. Ullah, H. M. Abdur Rehman, A. Y. Khan and A. R. Khan, *J. Mol. Liq.*, 2017, **236**, 395–403.
- H. Li, Y. Li, L. Xiang, Q. Huang, J. Qiu, H. Zhang, M. V. Sivaiah, F. Baron, J. Barrault, S. Petit and S. Valange, *J. Hazard. Mater.*, 2015, **287**, 32–41.
- J. M. Monteagudo, A. Durán and C. López-Almodóvar, *Appl. Catal., B*, 2008, **83**, 46–55.
- A. Mirzaei, Z. Chen, F. Haghghat and L. Yerushalmi, *Chemosphere*, 2017, **174**, 665–688.
- X. Zhang, Y. Ding, H. Tang, X. Han, L. Zhu and N. Wang, *Chem. Eng. J.*, 2014, **236**, 251–262.
- J. A. Khan, X. He, N. S. Shah, M. Sayed, H. M. Khan and D. D. Dionysiou, *Chem. Eng. J.*, 2017, **325**, 485–494.
- D. B. Jiang, X. Liu, X. Xu and Y. X. Zhang, *J. Phys. Chem. Solids*, 2018, **112**, 209–215.
- L. Zhou, J. Lei, L. Wang, Y. Liu and J. Zhang, *Appl. Catal., B*, 2017, DOI: 10.1016/j.apcatb.2017.08.039.
- J. Zhang, T. Yao, C. Guan, N. Zhang, H. Zhang, X. Zhang and J. Wu, *J. Colloid Interface Sci.*, 2017, **505**, 130–138.
- Q. Tian, W. Wu, L. Sun, S. Yang, M. Lei, J. Zhou, Y. Liu, X. Xiao, F. Ren, C. Jiang and V. A. L. Roy, *ACS Appl. Mater. Interfaces*, 2014, **6**, 13088–13097.
- L. Qin, M. Liu, Y. Wu, Z. Xu, X. Guo and G. Zhang, *Appl. Catal., B*, 2016, **194**, 50–60.
- B. Liu, L. Tian, R. Wang, J. Yang, R. Guan and X. Chen, *Appl. Surf. Sci.*, 2017, **422**, 607–615.
- S. Guo, G. Zhang, Y. Guo and J. C. Yu, *Carbon*, 2013, **60**, 437–444.
- D. Bi and Y. Xu, *Langmuir*, 2011, **27**, 9359–9366.
- D. Xiao, K. Dai, Y. Qu, Y. Yin and H. Chen, *Appl. Surf. Sci.*, 2015, **358**, 181–187.
- S. Hu, R. Jin, G. Lu, D. Liu and J. Gui, *RSC Adv.*, 2014, **4**, 24863.
- A. Akhundi and A. Habibi-Yangjeh, *Ceram. Int.*, 2015, **41**, 5634–5643.
- S. Kumar, S. T. B. Kumar, A. Baruah and V. Shanker, *J. Phys. Chem. C*, 2013, **117**, 26135–26143.
- D. Zhu, S. Liu, M. Chen, J. Zhang and X. Wang, *Colloids Surf., A*, 2018, **537**, 372–382.
- D. Zhang, S. Cui and J. Yang, *J. Alloys Compd.*, 2017, **708**, 1141–1149.
- J. Wang, Y. Chen, G. Liu and Y. Cao, *Composites, Part B*, 2017, **114**, 211–222.
- R. Nagarjuna, S. Challagulla, R. Ganesan and S. Roy, *Chem. Eng. J.*, 2017, **308**, 59–66.
- S. Hu, W. Ouyang, L. Guo, Z. Lin, X. Jiang, B. Qiu and G. Chen, *Biosens. Bioelectron.*, 2017, **92**, 718–723.
- M. Zhang and X. Wang, *Energy Environ. Sci.*, 2014, **7**, 1902.
- N. Gao, H. Wu, Y. Chang, X. Guo, L. Zhang, L. Du and Y. Fu, *Spectrochim. Acta, Part A*, 2015, **134**, 10–16.
- Y.-P. Sun, X.-q. Li, J. Cao, W.-x. Zhang and H. P. Wang, *Adv. Colloid Interface Sci.*, 2006, **120**, 47–56.



- 33 C. Cai, Z. Zhang, J. Liu, N. Shan, H. Zhang and D. D. Dionysiou, *Appl. Catal., B*, 2016, **182**, 456–468.
- 34 X. Zhou, B. Jin, R. Chen, F. Peng and Y. Fang, *Mater. Res. Bull.*, 2013, **48**, 1447–1452.
- 35 X. Li, X. Huang, D. Liu, X. Wang, S. Song, L. Zhou and H. Zhang, *J. Phys. Chem. C*, 2011, **115**, 21567–21573.
- 36 E. S. Baeissa, *Frontiers in Nanoscience and Nanotechnology*, 2016, **2**, 100–106.
- 37 M. Mousavi and A. Habibi-Yangjeh, *J. Colloid Interface Sci.*, 2016, **465**, 83–92.
- 38 X. Zhang, S. Lin, Z. Chen, M. Megharaj and R. Naidu, *Water Res.*, 2011, **45**, 3481–3488.
- 39 T. Wang, X. Jin, Z. Chen, M. Megharaj and R. Naidu, *Sci. Total Environ.*, 2014, **466–467**, 210–213.
- 40 B. Jia, L. Gao and J. Sun, *J. Am. Ceram. Soc.*, 2007, **90**, 1315–1318.
- 41 P. Liu, Q. Xu, D. Gao, S. Shi and B. Xia, *Journal of Advances in Nanomaterials*, 2017, DOI: 10.22606/jan.2017.21002.
- 42 Y. Cui, J. Briscoe, Y. Wang, N. V. Tarakina and S. Dunn, *ACS Appl. Mater. Interfaces*, 2017, **9**, 24518–24526.
- 43 S. Chi, C. Ji, S. Sun, H. Jiang, R. Qu and C. Sun, *Ind. Eng. Chem. Res.*, 2016, **55**, 12060–12067.
- 44 E. Cheng, S. Zhou, M. Li and Z. Li, *Appl. Surf. Sci.*, 2017, **410**, 383–392.
- 45 W. J. Kim, E. Jang and T. J. Park, *Appl. Surf. Sci.*, 2017, **419**, 159–164.
- 46 K.-i. Ishibashi, A. Fujishima, T. Watanabe and K. Hashimoto, *Electrochem. Commun.*, 2000, **2**, 207–210.
- 47 Q. Xiang, J. Yu and P. K. Wong, *J. Colloid Interface Sci.*, 2011, **357**, 163–167.
- 48 F. Dong, L. Wu, Y. Sun, M. Fu, Z. Wu and S. C. Lee, *J. Mater. Chem.*, 2011, **21**, 15171–15174.

