Preparation and Characterization of Fe₃O₄ Particles with Novel Nanosheets Morphology and Magnetochromatic Property by a Modified Solvothermal Method

Lin Zhuang¹, Wei Zhang², Yongxin Zhao¹-³, Hui Shen³, Han Lin⁴ & Jinhua Liang¹-³

¹State Key Laboratory of Optoelectronic Materials and Technologies, Guangdong Provincial Key Laboratory of Photovoltaics Technologies, School of Physics and Engineering, Sun Yat-Sen University, Guangzhou 510275, P. R. China, ²School of Physics and Optoelectronic Engineering, Guangdong University of Technology, Guangzhou 510006, China, ³School of Chemistry and Chemical Engineering, Sun Yat-Sen University, Guangzhou 510275, P. R. China, ⁴Department of Orthodontics, Guanghua School of Stomatology, Hospital of Stomatology, Sun Yat-sen University, Guangzhou 510055, P.R. China.

Novel-morphological Fe₃O₄ nanosheets with magnetochromatic property have been prepared by a modified solvothermal method. Such nanosheets could form one-dimension photonic crystal under an external magnetic field. The Fe₃O₄ nanosheets suspension could strongly diffract visible light and display varied colors with changing the intensity of the magnetic field. The photonic response is rapid, fully reversible and widely tunable in the entire visible spectrum. Excellent magnetic properties of these Fe₃O₄ nanosheets are exhibited with a high saturation magnetization (82.1 emu/g), low remanence (13.85 emu/g) and low coercive force (75.95 Oe). The amount of the solvent diethylene glycol (DEG) plays a key role in the formation of the sheet-shaped morphology. When the ratio of the DEG reaches 100%, the growing of the crystal plane (111) of Fe₃O₄ is inhibited and the sheet-like Fe₃O₄ crystals are formed.

Nanostructure magnetic Fe₃O₄ particles are of much interest because of their applications in a variety of areas such as ferrofluid, advanced magnetic materials, magnetic recording media and biomedical purpose, etc. With the advancement of magnetic materials, especially in the area of the magnetic responsive photonic crystal, considerable efforts have been attempted to improve their magnetic and optical properties. Different from other external stimuli systems of responsive photonic crystal, introducing magnetic components into the photonic crystal makes it more convenient and precise to control the properties through an external magnetic field. Such magnetic responsive photonic crystals have been demonstrated in the applications of different color display units including dynamic displays, rewritable photonic papers and full color printing systems.

Recently, a high-temperature hydrolysis method was reported to synthesize monodisperse superparamagnetic Fe₃O₄ particles with the nanosphere morphology. These Fe₃O₄ nanospheres could form one-dimension photonic crystal in aqueous and non-polar media, which could diffract the light of visible wavelengths. As known, the magnetic properties of nanoparticles are greatly dependent on their morphologies. Varied morphologies of Fe₃O₄ nanocrystals including nanotubes, hollow spheres, nanochains, nanoflowers and nanosheets have been fabricated. Compared to magnetite particles with other shapes, nanosheets have better crystallinity and smaller lattice distortion, which enable them possess better physical and magnetic properties. To the best of our knowledge, the synthesis of Fe₃O₄ nanosheets with the magnetochromatic property has hardly been reported.

Numerous chemical methods including microemulsions, sol-gel syntheses, sonochemical reactions, hydrothermal reactions, hydrolysis and thermolysis of precursors have been used to synthesize Fe₃O₄ magnetic nanoparticles. In the specific synthesis of nanosheets, the methods of solvothermal process, solid-state thermal decomposition route, supercritical fluid technique and bottom-up technique have been attempted. These synthesis processes are usually involved in one-step reaction, and the surface modification procedure is
incorporated with the synthesis of particles. In addition, in the conventional synthesis methods, deoxygenated protection and precisely control of the pH value are required.

Here we report a two-step solvothermal method to synthesize the Fe₃O₄ nanosheets with good magnetochromatic property without deoxygenated protection and controlling of the pH value. In the process, the surface modification of the Fe₃O₄ particles is performed after synthesizing the particles. The separation of these two steps simplifies our synthesis method and increases the reproducibility of the obtained nanosheets. In addition, there are more choices for capping agents since the modification step is independent. When an external magnetic field is introduced to the Fe₃O₄ nanosheets suspension, the particles suspension could diffract light with different wavelength and display various colors under magnetic fields with different intensity. The photonic response of the magnetic Fe₃O₄ nanosheets is reversible and widely tunable in the visible spectrum, indicating great potential applications in sensors and optical devices.

Figure 1 | XRD pattern of the Fe₃O₄ nanosheets.

Figure 2 | (a) Low magnitude TEM image of Fe₃O₄ nanosheets; (b) HRTEM of Fe₃O₄ nanosheets; (c) SAED pattern of Fe₃O₄ nanosheets. The scale bars are 0.2 μm, 5 nm, and 5 1/nm respectively.
Results and Discussion

The X-ray diffraction pattern of the Fe$_3$O$_4$ nanosheets is shown in Fig. 1, which reveals the formation of magnetite with well-defined crystallinity via the modified solvothermal process. All of the diffraction peaks match well with the normal characteristic diffractions of the Fe$_3$O$_4$ inverse spinel structure (PCPDFWIN v.2.02, PDF No. 89-0691). The calculated cell parameter (8.3915 Å) is comparable with the reported value. No obvious XRD peaks arising from impurities are found, which indicates no evidence that Fe$_3$O$_4$ particles are contaminated by foreign materials in the system.

In Fig. 2(a), the Fe$_3$O$_4$ nanosheets are of regular hexagonal or triangle structure with the edge length of 80–150 nm. The detailed crystalline information of the particles is shown in Fig. 2(b) and Fig. 2(c). The HRTEM image of selected area in the sample in Fig. 2(b) indicates that the nanosheets have well-defined crystallinity with an interplanar spacing of 0.296 nm. The diffraction points array tidily along a certain direction in the selected area electron diffraction (SAED) pattern as shown in Fig. 2(c). Crystal planes of (220) and (422) could be found in Fig. 2(c), which are consistent with the XRD pattern in Fig. 1. From Fig. 2(b) and Fig. 2(c), it’s clear that the growth rate of the crystal plane (111) is depressed, which is the key factor for the formation of the hexagonal or triangle shape of the nanosheets.

The nucleation of the Fe$_3$O$_4$ particles could be well controlled by changing the volume ratios of the solvent DEG and Ethylene Glycol (EG). The amount of the DEG plays a key role in the formation of the sheet-shaped Fe$_3$O$_4$ nanostructures, which can be confirmed by the comparison experiments with different concentrations of DEG. Fig. 3 shows that when the volume ratios of DEG/EG (v/v in mL) are 0/40, 26/14, 30/10 and 40/0, different sizes and shapes of Fe$_3$O$_4$ particles have been obtained. The diameters of the nanospheres are 180 nm (Fig. 3a), 100 nm (Fig. 3b) and 60 nm (Fig. 3c), and the shape of the nanosheet is shown in Fig. 3d. The particle size decreases as the ratio of DEG/EG increases. This size tunability may due to the different diffusion coefficient of the ions in the solution caused by the varying concentrations of the DEG. Since DEG has a stronger affinity to the Fe$^{3+}$ ions, the viscosity of the solvent increases as the portion of

Figure 3 | Low magnitude TEM images of Fe$_3$O$_4$ particles: (a) 180 nm particles (DEG/EG = 0/40); (b) 100 nm particles (DEG/EG = 26/14); (c) 60 nm particles (DEG/EG = 30/10); (d) nanosheets (DEG/EG = 40/0). All the scale bars are 200 nm.

Figure 4 | Low magnitude TEM images of Fe$_3$O$_4$ product under the conditions of different reaction time and percentage of DEG in the solvent: (a) 1 hour, DEG% = 65%; (b) 3 hours, DEG% = 65%; (c) 1 hour, DEG% = 100%; (d) 3 hours, DEG% = 100%.

Figure 5 | Photographs of Fe$_3$O$_4$ nanosheets colloidal solution formed in response to an external magnetic field. (a) and (b), the external magnetic field are both set beside the Fe$_3$O$_4$ nanosheets suspension.
the DEG increases and therefore the diffusion coefficient of the ions in the reaction system decreases, which leads to smaller grain size of the Fe$_3$O$_4$ particles and inhibits the formation of the secondary structure. In the solvothermal process, when the ratio of the DEG is rather low, the formation of the microstructure features after fast nucleation in solution relates to the aggregation growth process and the Ostwald ripening process. When the ratio of the DEG reaches 100%, the mechanism of the crystal growth is different from the former and the morphology of the final product becomes nanosheet.

To further understand the formation mechanism of the Fe$_3$O$_4$ nanosheets, a number of matrix precursor experiments were carried out under different reaction time and different concentrations of DEG. TEM images in Fig. 4(a) and Fig. 4(c) reveal that the samples consist of unconsolidated particles when the reaction time is 1 h. These unconsolidated particles only congregate but not form the crystal structure according to the electron diffraction patterns in Fig. 4(a) and Fig. 4(c). Fig. 4(b) reveals that the nanospheres preliminarily form after reaction for 3 h. In the reaction system, the Fe(III) compounds hydrolyze and deoxidize to Fe(II) compounds partially. Fe$_3$O$_4$ crystal nucleus are formed by the interaction of Fe(II) and Fe(III) hydrates. The growing of the crystal follows the process of homogeneous nucleation and the final product of Fe$_3$O$_4$ nanospheres is obtained. Fig. 4(d) shows the formation of the nanosheets. When the ratio of the DEG reaches 100%, DEG and Fe$^{3+}$ ions complex and form a steady solution. The growing of the crystal and aggregation of nanoparticles are slow under the strong complexation and the high viscosity of the solvent. The growth of the crystal plane (111) is limited and the shape of the nanosheets, but not the nanospheres, is formed.

The Fe$_3$O$_4$ sheet-like nanoparticles exhibit good magnetochromatic properties. The photographs and reflection spectra of the aqueous solution of Fe$_3$O$_4$ colloidal particles (ca. 15.00 mg mL$^{-1}$) in response to an external magnetic field are shown in Fig. 5 and Fig. 6. When a magnetic field is applied to the nanosheets suspension, the magnetic colloids displays clearly a gradient of colors. The particles in the suspension form a chain-like structure along with the direction of the external magnetic field, which is known as photonic crystal$^{21}$. The structure formed by the magnetic nanosheets could diffract visible light of different wavelength ranging from 480 to 640 nm as shown in Fig. 6. The reflection rate increases and the diffraction peaks blue shift as the magnetic field is increased from 0 to 327 G. The low reflection rate and broad peak width at a low magnetic field are due to the Brownian motion, which significantly interferes with the formation of the long-range order structure. When the magnetic field increases, the magnetic interaction between two magnetic nanoparticles surpasses the Brownian motion and the chain-like structure is enhanced, which makes the diffraction peak sharper and higher.

The magnetic properties of the modified nanosheets and the corresponding nanosheets suspension were investigated respectively. As shown in Fig. 7, the Fe$_3$O$_4$ nanosheets exhibit soft magnetism with a low remanence (13.85 emu/g) and low coercive force (75.95 Oe). The saturation magnetization of the magnetic nanosheets is 82.10 emu/g, which is much higher than that of the nanoparticles prepared by the common chemical coprecipitation method (about 60–65 emu/g). The high reaction temperature in our modified solvothermal method results in a higher crystallinity and consequently leads to a higher saturation magnetization. While in the observation of the hysteresis loop of the particles suspension as shown in Fig. 8, the colloid exhibits essentially no remanence or coercive force. This interesting result explores the previous conclusion that ferromagnetic particles cannot be dispersed in aqueous media. The remanence and the coercive force of the Fe$_3$O$_4$ nanosheets dispersed in aqueous solution are found to decrease to 0.042 emu/g and 1.94 Oe, respectively. This phenomenon indicates that if suitable surface modification is performed to decrease the magnetic force between the adjacent nanosheets, the ferromagnetic particles can rely on the Brownian motion to disrupt the chain-like structure induced by the magnetic moment, and then a transformation from ferromagnetism of the particles to a hysteresis loop with much small remanence and coercive force of the particles suspension appears. In this case, there will be no significant magnetic interactions remaining after removing the external magnetic field, which is important for achieving a reproducible and reversible optical response. The hysteresis loop of Fig. 8 displays irreversibility when the applied external magnetic field is between 100 and 900 G. When the external field is decreased from 900 G in the upper curve, the particles could still keep the saturation magnetization until the field is decreased to be about 700 G. And during the magnetic field is between 900 and 100 G, the magnetization value in the upper curve is clearly larger than that in the bottom curve. This is due to the complex effect of metastable state of the magnetic dynamic nanoparticles and the possible relaxation characteristic of the formed chain-like structure when decreasing the external magnetic field, the Brownian dynamics of particles, and the electrostatic interactions due to the PAA.

In summary, magnetic Fe$_3$O$_4$ nanosheets have been synthesized by a two-step solvothermal method. The Fe$_3$O$_4$ nanosheets suspension exhibits magnetochromatic property under an external magnetic
field. The formed one-dimension photonic crystals by the Fe₃O₄ nanosheets in aqueous solution under external magnetic field could strongly diffract visible light with different wavelength and display various colors. Diffraction peaks blue shift as the magnetic field increases which could cover the entire visible spectrum. The surface modification increases the dispersibility of the nanosheets and achieves a transformation of ferromagnetism of the particles to a hysteresis loop with much small remanence and coercive force of the particles suspension, which enables the optical response of the colloidal solution reproducible and reversible. Excellent magnetic properties of the nanosheets have been obtained with a high saturation magnetization (82.1 emu/g), low remanence (13.85 emu/g) and low coercive force (75.95 Oe). The amount of the solvent DEG plays a key role in the formation of the sheet-shaped Fe₃O₄ nanosstructures. The size of the Fe₃O₄ nanoparticles decreases as the ratio of DEG/EG increases in virtue of the strong affinity of DEG to Fe⁺⁺ ions and the higher viscosity of the solvent. A transformation of the shape from nanosphere to nanosheets appears when the ratio of DEG/EG is 40/0.

**Methods**

**Synthesis of polyacrylic acid.** 125 mL Deionized H₂O and 1.00 g ammonium persulfate (APS) were added into a three-neck bottle with a dropping funnel and reflux condensing tube. When the APS was dissolved under magnetic stirring. 5.00 g acrylic acid and 8.00 g isopropanol were further added. Then the mixture was heated to 65–70°C, following with the add of a solution containing 40.00 g acrylic acid, 2.50 g ammonium persulfate and 40 mL DI H₂O drop by drop through the dropping funnel. Due to the heat released in the polymerization reaction, the temperature of the system would rise and lead to reflux. After all the acrylic acid and APS solution reacted for 30 min, the mixture was kept refluxing at 94°C for 1 h and then cooled down.

**Synthesis and surface modification of Fe₃O₄ particles with sheet-like morphology.** Ferrous chloride (FeCl₂), ferric chloride (FeCl₃), Sodium Acetate (NaAc) and Polyethylene Glycol (PEG) were dissolved in 40 mL Diethyl glycol (DEG) under stirring and ultrasonic treatment. The molar ratio of FeCl₂:FeCl₃:NaAc:PEG was 1:1.9:22:1.25. The homogeneous yellow mixture was then transferred to a Teflon-lined stainless-steel autoclave and sealed to heat at 220°C for 7 h. The autoclave was cooled down to room temperature. The as-prepared Fe₃O₄ particles were washed several times with ethanol and DI H₂O before the surface modification step. 20 mL NaOH solution (1.00 mM) was added into the wet Fe₃O₄ precipitation, and then the mixture was ultrasonic treated to disperse the particles into the solution and form the magnetic nanoparticles (MNPs) suspension. 20 mL PAA solution (10.0 g/L) and 1 mL Fe₃O₄ (0.1 mol/L) were mixed to form an orange PAA-Fe⁺⁺ complex solution. The MNPs suspension was slowly added into the PAA-Fe⁺⁺ complex solution under mechanical stirring. The mixture was ultrasonic treated at 80°C for 15 min, and then the precipitate was collected by magnetic separation. After adding 20 mL NaOH (0.05 mol/L) solution into the precipitate and 10-s ultrasonic treatment of the mixture, the product was collected again by magnetic separation and washed by DI H₂O three times. Finally, the magnetic colloidal solution was obtained by dispersing the product in a certain amount of DI H₂O. It should be noted that the particles in the suspension are fairly stable in air and could be stored for months.

**Characterization.** X-ray diffraction (XRD) patterns were recorded on a Power X-ray Diffractometer (D/MAX-III) with Cu Kα radiation. Transmission electron microscopy (TEM) photographs were taken on a high-resolution transmission electron microscope (HRTEM, JEM-2010) at an accelerating voltage of 200 kV. The optical properties of the photonic crystals formed by the magnetic Fe₃O₄ nanosheets under external magnetic field, which was controlled by a home-made electromagnet, were collected by using a QE65000-ABS Scientific-grade Optic Fiber Spectrometer (Ocean Optic, Inc.). Room temperature magnetization measurements were performed with a LakeShore Model 7404 vibrating sample magnetometer (VSM).

Acknowledgments. This work was supported by the Natural Science Foundation of China (Grant No. 61376014 and No. 60906005), the Guangdong Provincial Natural Science Foundation of China (Grant No. S2011010106010 and No. 9451027501002848) and the Fundamental Research Funds for the Central Universities (Grant No. 11lgpy87 and No. 60ykpy66).

**Author contributions.** L.Z. wrote the manuscript and analyzed the experimental data, W.Z. provided crucial support for the experiments, Y.Z. and J.L. carried out the experiments, analyzed the experimental data, H.S. supervised the project and H.L. helped writing the main manuscript.

---

1. Sun, S., Murray, C., Weller, D., Folks, L. & Moser, A. Monodisperse FePt nanoparticles and ferromagnetic FePt nanocrystal superlattices. *Science* 287, 1899–1992 (2000).
2. Miller, M., Prinz, G., Cheng, S.-F. & Bounnak, S. Detection of a micron-sized magnetic sphere using a ring-shaped anisotropic magnetoresistance-based sensor: a model for a magnetoresistance-based biosensor. *Appl. Phys. Lett.* 81, 2211–2213 (2002).
3. Jain, T. K., Morales, M. A., Sahoo, S. K., Leslie-Pelecky, D. L. & Labhasetwar, V. Iron oxide nanoparticles for sustained delivery of anticancer agents. *Mol. Pharm.* 2, 194–205 (2005).
4. Babes, L., Denizot, B. T., Tanguy, G., Le Jeune, J. J. & Jallet, P. Synthesis of iron oxide nanoparticles used as MRI contrast agents: a parametric study. *J. Colloid Interface Sci.* 212, 474–482 (1999).
5. Chouria, I. et al. Molecular composition of iron oxide nanoparticles, precursors for magnetic drug targeting, as characterized by confocal Raman microscopy. * Analyst* 130, 1395–1403 (2005).
6. Moser, A. et al. Magnetic recording: advancing into the future. *J. Phys. D: Appl. Phys.* 35, R157 (2002).
7. Ge, J., Kwon, S. & Yin, Y. Niche applications of magnetically responsive photonic structures. *J. Mater. Chem.* 20, 5777–5784 (2010).
8. Jia, C.-J. et al. Iron oxide tube-in-tube nanostructures, *J. Phys. Chem. C* 111, 13022–13027 (2007).
9. Yang, M., Ma, J., Zhang, C., Yang, Z. & Lu, Y. General Synthetic Route toward Functional Hollow Spheres with Double-Shelled Structures. *Angew. Chem. Int. Ed.* 44, 6727–6730 (2005).
10. Wu, M. et al. Magnetic field-assisted hydrothermal growth of chain-like nanorods of metallic Fe. *Chem. Phys. Lett.* 401, 374–379 (2005).
11. Zhong, L.-S. et al. Self-assembled 3D flowerlike iron oxide nanostructures and the application in water treatment. *Adv. Mater.* 18, 2426–2431 (2006).
12. Xian-Ming, L. & Jang-Jyo, K. Solvothermal synthesis and magnetic properties of magneto ferrite nanoplatelets. *Mater. Lett.* 63, 428–430 (2009).
13. Chin, A. B. & Yacobi, I. I. Synthesis and characterization of magnetic iron oxide nanoparticles via w/o microemulsion and Massart’s procedure. *J. Mater. Process. Technol.* 191, 235–237 (2007).
14. Alborno, C. & Jacobo, S. E. Preparation of a biocompatible magnetic film from an aqueous ferrofluid. *J. Magn. Magn. Mater.* 305, 12–15 (2006).
15. Hee Kim, E., Sook Lee, H., Kook Kwak, B. & Kim, B.-K. Synthesis of ferrofluid with magnetic nanocrystals by sonochemical method for MRI contrast agent, *J. Magn. Mater.* 289, 328–330 (2005).
16. Wu, J., Chen, X., Wang, Z., Yang, X. & Qian, Y. A. soft-template-assisted hydrothermal approach to single-crystal Fe₃O₄ nanorods. *J. Cryst. Growth* 276, 571–576 (2005).
17. Kimata, M., Nakagawa, D. & Hasegawa, M. Preparation of monodisperse magnetic particles by hydrosynthesis of iron oxide. *Powder Technol.* 132, 112–118 (2003).
18. Chen, L., Zhao, C., Zhou, Y., Peng, H. & Zheng, Y. Controlled synthesis of Fe₃O₄ nanosheets via one-step pyrolysis of EDTA ferric sodium salt. *J. Alloys Compd.* 504, L46–L50 (2010).
19. Li, Z. et al. Supercritical Fluid Synthesis of Magnetic Hexagonal Nanoplatelets of Magnetite. *J. Am. Chem. Soc.* 132, 12540–12541 (2010).
20. Chen, K. C. et al. Large-Scale Synthesis of Fe₃O₄ Nanoplates at Low Temperature. *J. Phys. Chem. C* 111, 9136–9141 (2007).
21. Ge, J., Hu, Y., Zhang, T., Huyhn, T. & Yin, Y. Self-assembly and field-responsive optical diffractions of superparamagnetic colloids. *Langmuir* 24, 3671–3680 (2008).
Additional information
Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Zhuang, L. et al. Preparation and Characterization of Fe₃O₄ Particles with Novel Nanosheets Morphology and Magnetochromatic Property by a Modified Solvothermal Method. Sci. Rep. 5, 9320; DOI:10.1038/srep09320 (2015).