Preparation and properties of poly(acrylic acid) oligomer stabilized superparamagnetic ferrofluid

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Abstract
Ferrofluids, which are stable dispersions of magnetic particles, behave as liquids that have strong magnetic properties. Nanoparticles of magnetite with a mean diameter of 10–15 nm, which are in the range of superparamagnetism, are usually prepared by the traditional method of co-precipitation from ferrous and ferric electrolyte solution. When diluted, the ferrofluid dispersions are not stable if anionic or cationic surfactants are used as the stabilizer. This work presents an efficient way to prepare a stable aqueous nanomagnetite dispersion. A stable ferrofluid containing Fe₃O₄ nanoparticles was synthesized via co-precipitation in the presence of poly(acrylic acid) oligomer. The mechanism, microstructure, and properties of the ferrofluid were investigated. The results indicate that the PAA oligomers promoted the nucleation and inhibited the growth of the magnetic iron oxide, and the average diameter of each individual Fe₃O₄ particle was smaller than 10 nm. In addition, the PAA oligomers provided both electrostatic and steric repulsion against particle aggregation, and the stability of dispersions could be controlled by adjusting the pH value of solution. A small amount of Fe₂O₃ was found in the nanoparticles but the superparamagnetic behavior of the nanoparticles was not affected.

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1. Introduction
Ferrofluids are colloidal dispersions of small single-domain magnetic particles suspended in a carrier fluid. Ferrofluids characteristically have both magnetic and fluid properties and have found a diverse range of applications, such as in audio devices, inertia dampers, stepper motors, sensors, vacuum seals, electromagnetic shielding, and high-density digital storage [1,2]. In general, finer particles are more surface active; i.e., they have a higher specific surface area. On the other hand, it is often difficult to handle fine particles such as colloids. For example, the filtration will be time-consuming or impractical, or else a centrifuge has to be used. But unlike other ultrafine particles, magnetic particles can be handled by an external magnetic field. Therefore, ferrofluids are widely studied for their applications in various fields in biology and medicine such as enzyme and protein immobilization, genes, radiopharmaceuticals, magnetic resonance imaging MRI, diagnostics, immunoassays, purification, separation, and controlled drug release [3–6]. However, the nanoparticles in the ferrofluids are likely to aggregate. Consequently, how to prevent aggregation between the nanoparticles during synthesis and treatment has received considerable attention when the applications need well-dispersed magnetic nanoparticles. Although well-dispersed magnetic nanoparticles can be obtained by ball milling [7], high-energy requirements and unavoidable contamination of the product have necessitated the development of more economical and reliable ways to produce magnetic particles by chemical coprecipitation.

Iron oxide (Fe₃O₄), the dominant magnetic material in magnetic fluid preparations, can be synthesized through the coprecipitation of Fe(II) and Fe(III) salts by addition of a
base. Stable ferrofluids can be prepared by adsorption of stabilizing agents on the surface of magnetite particles after a synthesis process. The feasibility of preparing magnetic fluids using anionic or nonionic surfactants as dispersing agents has been examined [8–10]. Massart obtained stable aqueous alkaline and acidic magnetic liquids by free precipitation [11,12]. The nature of the counterions and the pH of the suspensions played an important role in stabilizing the charged magnetic particles through interactions between their electrical double layers. The applications of magnetic fluids, which are stabilized entirely by electrostatic repulsion, are somewhat restricted, since this system is overly sensitive to conditions such as pH and ionic strength and offers little flexibility for changing the surface properties of the particles. Stabilization of magnetic fluid suspensions can be achieved by coating the particle surfaces with organic surfactants. When two such particles approach each other, the surfactants interpenetrate and the particles are subjected to steric repulsion due to an increase in osmotic pressure [8,13].

The traditional methods of preparing ferrofluids involve long-term grinding of magnetic materials in a carrier medium containing oleic acid, which is commonly used as a dispersing agent to stabilize dispersions in kerosene and other hydrocarbon dispersion media. This method has prompted the application of the same technique to prepare a water-based magnetic fluid. The latter preparation was anticipated to be much simpler, since one usually starts with aqueous solutions of ferrous and ferric salts. Shimoizaka et al. proposed the principle of bilayer stabilization in the early 1980s [14]. They first precipitated oleic acid-coated particles and then redispersed them in aqueous solutions of sodium dodecylbenzene sulfonate, poly(oxyethylene) nonylphenyl ethers, and di(2-ethylhexyl)adipate. The colloidal particles are thus doubly coated, first with a chemisorbed monolayer of oleic acid on top of which a second layer of the surfactant is physically adsorbed. Following this work, Khalafla and Reimers [7] and Wooding et al. [15] produced stable aqueous magnetic fluids using various saturated and unsaturated fatty acids as primary and secondary surfactants. In our previous studies [16,17], we have also synthesized ferrofluids by adsorbing stabilizing agents, lauric acid, after the co-precipitation process [7,16–18]. However, the ferrofluids thus prepared were not stable when being diluted. Aggregation of particles occurred when excess quantity of water was added. Applications of magnetite dispersions composed of electrostatic stabilizers and bilayer surfactants are restricted due to pH sensitivity and dispersion stability, respectively [7].

Recently, aqueous magnetic fluids have been reported using polymers as steric stabilizers and polymer templates [19–24]. But it should be noted that many of the reported polymeric stabilizers have not been designed with functional groups to bind to the surface of Fe₃O₄ particles, and thus, stability of ferrofluid has been limited [24]. This study presents a novel and simple procedure to synthesize an extremely stable and water-based Fe₃O₄ ferrofluid via a co-precipitation method in the presence of poly(acrylic acid) oligomers. The PAA oligomers were chemically bound on the surface of Fe₃O₄ nanoparticles to suspend the Fe₃O₄ particles in a fine state of dispersion, and the PAA oligomers provided both electrostatic and steric repulsion against particle aggregation.

2. Experimental

2.1. Materials

Fe(II) chloride tetrahydrate (99%), Fe(III) chloride hexahydrate (97%), and ammonium hydroxide (28% NH₃ in water, w/w) were purchased from ACROS and used as supplied. Poly(acrylic acid) (PAA oligomer: M₆₀ ∼ 2000) was purchased from Aldrich. Other chemicals were of analytical grade and used without further purification. Distilled and deionized water was used throughout the work.

2.2. Synthesis of ferrofluid

The stable Fe₃O₄ ferrofluid was synthesized using the method of coprecipitation of Fe(II) and Fe(III) salts in the presence of PAA oligomer (M₆₀ ∼ 2000). In this process, 1 g of PAA oligomer (M₆₀ ∼ 2000), 4.75 g of FeCl₃·6H₂O, and 1.75 g of FeCl₂·4H₂O were dissolved in 80 ml of water with vigorous stirring at 60°C under N₂. Then 7.5 ml of NH₄OH (28% (w/w)) was quickly added to the reactor and stirring was continued at 400 rpm for 30 min. Once the ammonium solution was added into the reactor, the color of the mixture turned from orange to black immediately. The ferrofluid thus obtained was designed by the sample code of PAA/MAG = 1/2, in which the weight ratio of PAA oligomers to Fe₃O₄ particles was theoretically calculated as 1/2 from the feed composition.

2.3. Observation of Fe₃O₄ nanoparticles

To observe the morphology of PAA oligomer-stabilized Fe₃O₄ particles, the ferrofluid was diluted in deionized water and observed using a TEOL JSM-1200 EX II transmission electron microscope (TEM). In addition, the ferrofluid was coated on a glass plate and dried naturally, and then was observed using a Hitachi S-800 scanning electron microscope (SEM).

2.4. Measurement of FTIR

To prepare the samples for the FTIR measurements (Bio-Rad FTS-3000 spectrometer), the PAA oligomers which were used as supplied were grind-blended with KBr and then compressed to form a pellet. In addition, the ferrofluid was dialyzed (molecular weight cut off MWCO = 6000–8000) against deionized water until the conductivity of the solution was stable and then was freeze-dried to form the PAA
oligomers stabilized Fe$_3$O$_4$ particles. In the same way, the sample of PAA oligomers stabilized Fe$_3$O$_4$ particles were grind-blended with KBr and then compressed to form a pellet for the measurement of FTIR. The transmission spectra were obtained for 64 scans at a resolution of 4 cm$^{-1}$.

2.5. Measurement of particle size distribution and zeta potential of Fe$_3$O$_4$ nanoparticles

The zeta potential and the size distribution of the PAA oligomers stabilized Fe$_3$O$_4$ particles were measured by using laser light scattering (Malvern Zeta Sizer 3000H) at 25$^\circ$C. Before the measurements, the ferrofluids were dia-layed for 7 days and diluted with distilled and deionized water.

2.6. Stability of ferrofluid

To investigate the effect of pH value on the stability of ferrofluid, the pH value of the ferrofluid was measured during the process of titration using 28% (w/w) ammonium solution as a titrant. When the pH value of solution was stable for several minutes, the stability of the particles suspended in the fluid was observed visually. Furthermore, the effect of acid medium on the stability of the PAA oligomers stabilized Fe$_3$O$_4$ particles was studied by using methacrylic acid as a titrant to titrate the ferrofluid, which contained 2 ml 28% (w/w) ammonium solution, and the pH value and stability of the Fe$_3$O$_4$ particles during the process of titration were observed.

2.7. TGA and XRD analysis

The ferrofluid was dialyzed until the conductivity of the solution was stable. Then the ferrofluid was freeze-dried to produce the PAA oligomer-stabilized Fe$_3$O$_4$ particles. The Fe$_3$O$_4$ particles were subjected to TGA (Perkin–Elmer TGA-7). The temperature was kept at 100$^\circ$C for 10 min and then was raised to 800$^\circ$C at a heating rate of 10$^\circ$C/min. Additionally, X-ray diffraction measurements with monochromatic CuK$_\alpha$ radiation were taken to investigate the crystal structure of the particles.

2.8. Measurement of magnetization of Fe$_3$O$_4$ particles

The magnetization of the PAA oligomer-stabilized nanoparticles were measured by SQUID magnetometer (superconducting quantum interface device, Quantum Design MPMS5) at 298 K and ±10,000 G applied magnetic field. Due to the applied magnetic field ($H$), the magnetic induction ($B$) could be measured according to Faraday’s law, and the magnetization ($M$) was then obtained by the relationship

\[ B = H + 4\pi M. \]  

The applied magnetic field was increased until the saturation magnetization was achieved. Then the applied field was reduced to zero to measure the remanence (residual magnetization). When the applied field was further reversed, the coercivity (the applied field that reduces magnetization to zero) and the saturation magnetization in the reverse direction could be obtained. This increasing and decreasing applied field process was repeated five times to get the magnetization curve (magnetization ($M$) versus applied magnetic field ($H$)) and examine the magnetic properties of the Fe$_3$O$_4$ particles.

In addition, the important parameters used to characterize the magnetic properties of solids, magnetic susceptibility ($\kappa$), and magnetic permeability ($\mu$) were also calculated using the following equations:

\[ \kappa = M / H, \]  

\[ \mu = B / M. \]

3. Results and discussion

3.1. Synthesis of ferrofluid

The traditional method of preparing Fe$_3$O$_4$ nanoparticles was to use the chemical co-precipitation of ferric and ferrous salts in alkaline medium:

\[ \text{Fe}^{2+} + 2\text{Fe}^{3+} + 8\text{OH}^- \leftrightarrow \text{Fe(OH)}_2 + 2\text{Fe(OH)}_3 \rightarrow \text{Fe}_3\text{O}_4 + \text{H}_2\text{O}. \]  

However, the Fe$_3$O$_4$ nanoparticles synthesized by this traditional method would aggregate easily, as shown in Fig. 1a. To prevent aggregation, the PAA oligomers were used to modify the traditional chemical co-precipitation method to synthesize the stable superparamagnetic ferrofluid. The ferrofluid containing the Fe$_3$O$_4$ nanoparticles could be suspended homogeneously and stably in water or monomer solution as shown in Fig. 1b. The proposed mechanism was shown in Fig. 2.

At first the Fe$^{2+}$ and Fe$^{3+}$ ions and the carboxylic acid groups of PAA oligomers formed a complex structure. After the ammonium solution was added, carboxylic acid groups on the PAA oligomer promoted the nucleation of Fe$_3$O$_4$ particles. Because the PAA oligomers acted as the template for Fe$_3$O$_4$ nucleation, the growth of particles would be hindered by the oligomers [25,26] and the size of the particles was about 5–10 nm, which was smaller than the traditional product. Also, a lot of carboxylic acid groups were bound on the surface of Fe$_3$O$_4$ nanoparticles to suspend the particles in a fine state of dispersion as discussed later. The Fe$_3$O$_4$ nanoparticles could be synthesized in situ and form a hyperbranch structure as shown in Figs. 3a and 3b. But there is no evidence to explain the formation of a hyperbranch structure.

From the chemical equation of Fe$_3$O$_4$ precipitation, Eq. (4), the suitable mole concentration ratio of Fe(II) to Fe(III) should be 1/2 for the synthesis of Fe$_3$O$_4$ nanoparticles. In addition, the suitable pH value for the synthesis of Fe$_3$O$_4$ nanoparticles should be higher than 6.93, which
could be calculated using (4) for $K_{sp}$ values of Fe(OH)$_2$ and Fe(OH)$_3$ at 25 °C of $8 \times 10^{-16}$ and $4 \times 10^{-38}$. If the mole concentration ratio of Fe(II) to Fe(III) was larger than 1/2, not only Fe$_3$O$_4$ but also Fe(OH)$_2$ would be produced. Moreover, as shown in Eq. (5), Fe(OH)$_3$ would be in excess and then converted to Fe$_2$O$_3$ if the mole concentration ratio of Fe(II) to Fe(III) were smaller than 1/2 or the pH value were lower than 6.93. In the reaction system we used, the mole concentrations of Fe(II) and Fe(III) were 0.1105 and 0.221 M and the pH value of the ferrofluid after synthesis was 8.8. Therefore, these conditions of reaction were suitable to form Fe$_3$O$_4$ nanoparticles:

$$2\text{Fe(OH)}_3 \rightarrow \text{Fe}_2\text{O}_3 + 3\text{H}_2\text{O}. \quad (5)$$

3.2. FTIR analysis

Since there are a large surface-to-volume atomic ratio, high surface activity, and a large amount of dangling bonds on the nanoparticle surface, the atoms on the surface are apt to adsorb ions or molecules in solution. In the system of Fe$_3$O$_4$ nanoparticles dispersed in an alkaline solution of PAA oligomers, the exposed atoms of Fe on the surface of Fe$_3$O$_4$ particles would adsorb OH$^-$ and COO$^-$ groups of...
PAA oligomers, and the atoms of O on the particle surface would adsorb H\(^+\). In addition, the OH-rich surface of Fe\(_3\)O\(_4\) nanoparticles could also react with the residual carboxylic acid groups of PAA oligomers.

Fig. 4 shows a comparison between the FTIR spectra of the Fe\(_3\)O\(_4\) nanoparticles, the pure PAA oligomers, and the ferrofluid prepared by co-precipitation in the presence of PAA oligomers. Previously, it was reported that the characteristic absorption bands of the Fe–O bond of bulk Fe\(_3\)O\(_4\) were at 570 and 375 cm\(^{-1}\) [27]. However, when the size of Fe\(_3\)O\(_4\) particles was reduced to nanoscale dimensions, the surface bond force constant increased due to the effect of finite size of nanoparticles, in which the breaking of a large number of bonds for surface atoms resulted in the rearrangement of nonlocalized electrons on the particle surface [28]. Therefore, the FTIR spectrum of Fe\(_3\)O\(_4\) nanoparticles would exhibit a blue shift and the characteristic absorption bands of the Fe–O bond were shifted to high wavenumbers of about 580 and 434 cm\(^{-1}\), as shown in Fig. 4. The strong band at ca. 1710 cm\(^{-1}\) as shown in the FTIR spectrum of PAA oligomer is attributed to the carboxyl group stretch of the PAA oligomer [29], and the bands at ca. 1400–1450 cm\(^{-1}\) are attributed to the stretch of C=O and the deformation vibration of OH [29]. In addition, the bands at ca. 1150–1280 cm\(^{-1}\) are attributed to the aliphatic acid [29]. The FTIR spectrum of the ferrofluid shows that the peak at 1710 cm\(^{-1}\) shrank and two new peaks at ca. 1540 and 1405 cm\(^{-1}\) appeared due to the binding of the carboxylic acid groups to the surface of the nanoparticles to form carboxylate groups [30,31]. The new peaks correspond to the COO\(^-\) antisymmetric vibration and the COO\(^-\) symmetric vibration, which indicate the bidentate bonding of the carboxyl groups to the surface Fe atoms [32,33]. Consequently, the interaction between PAA oligomers and Fe\(_3\)O\(_4\) nanoparticles was through chemical bonding and the mechanism shown in Fig. 2 was evidenced.

3.3. Zeta potential and light scattering analysis

The negative zeta potential indicates that the PAA oligomers are bound to the particle surface. Fig. 5a shows the zeta potentials of sample PAA/MAG = 1/2 in the solutions with different pH values. When the solution became more alkaline, the zeta potential became more negative that may due to the ionization of PAA oligomers. And the ionization of PAA oligomers would provide the electrostatic repulsion against the aggregation between nanoparticles. So Fig. 6a
shows that the sample in the solution with higher pH value would show small and narrow size distribution.

The zeta potentials of the samples with different PAA/MAG ratio were not changed drastically if the pH value of solutions were the same, as shown in Fig. 5b. But Fig. 6b shows that the sample with a higher concentration of PAA oligomer showed a small and narrow size distribution. The reason was due to the PAA oligomers that acted as templates and steric stabilizers, forming a wall to protect the particles. So the addition of the PAA oligomers in the reaction provides steric repulsion to improve the quality of the dispersion. These effects of polymeric stabilizers in metallic nanoparticle preparations were also discussed in previous studies [19–22]. Therefore, the PAA oligomers, which bonded to the surfaces of Fe₃O₄ particles, not only reduce the intrinsic particle size during their forming but also provide both factors of electrostatic and steric repulsion against the aggregation between Fe₃O₄ particles.

In addition, it is worth mentioning that when there was a high concentration of PAA oligomers in the ferrofluid, the effect of physical crosslinking due to the hydrogen bonding among COOH groups or the entanglement of PAA oligomers would cause a pseudo-aggregation of nanoparticles several weeks after the ferrofluid was synthesized, and therefore the size distribution of Fe₃O₄ measured by light scattering was much larger than its intrinsic size if ultrasonification was not applied before the measurement. Fig. 7 compared the size distribution of the Fe₃O₄ ferrofluid aggregates with or without ultrasonification before the light scattering measurement. It indicates that the Fe₃O₄ particles aggregated as clusters with a size range of 150–450 nm in the presence of high concentrations of PAA oligomers. The ultrasonification then destroyed the pseudo-aggregation in clusters and the size distribution of ferrofluid aggregates was reduced to about 10–40 nm.

3.4. Stability of ferrofluid

Because the ferrofluid was stabilized by PAA oligomers, the stability of Fe₃O₄ nanoparticles in a dispersing medium could be controlled by the ionization of carboxylic acid groups of PAA oligomers. The adjustment of the pH value of the medium solution could control the ionization degree of PAA oligomers and thus the stability of the ferrofluid.

The pKₐ value of PAA oligomers (MW ~ 2000) could be calculated to be 4.09 by Eqs. (6) and (7). This value is close to the pKₐ value of the aliphatic acid (4.0–4.5).

The concentration of carboxylic acid groups is

\[
C_0 = \frac{m}{MV}
\]

and

\[
pK_a = -\log \left( \frac{[\text{COO}^-][\text{H}^+]}{[\text{COOH}]} \right) = -\log \left( \frac{10^{-pK_a}}{C_0 - 10^{-pH}} \right)
\]

in which the pH value of the solution was measured to be 2.43 by dissolving 1 g PAA oligomer (m) in 80 g (≈80 L) of deionic water (V), and M is the molecular weight of the repeating unit of the PAA oligomer (72 g/mol). From further calculation, the relationship between the pH value and
the degree of ionization of carboxylic acid groups of PAA oligomer could be found as follows:

\[ pK_d = - \log \left( \frac{C_0(d)(H^+)}{C_0(1-d)} \right) = \text{pH} - \log \left( \frac{d}{1-d} \right) \]  

The symbol \( d \) used here is the ionization degree of acid groups. To simplify the calculation, it was assumed that more than 99.99% of carboxylic acid groups were ionized to provide electrostatic repulsion to form the most stable ferrofluid. The pH value under this condition was found to be higher than 8.1. In other words, the carboxylic acid groups of PAA oligomers were almost completely ionized and the ferrofluid was stable when the pH value was higher than 8.1. In our experiments, after the addition of ammonium solution into the reaction system to synthesize the ferrofluid, the final pH value of the ferrofluid was 8.8 which assured the formation of stable ferrofluids, and the ferrofluids thus synthesized would remain stable for several months. If the ammonium solution was added continuously as shown in Fig. 8, the ferrofluid was still stable due to the acid groups being almost ionized, and not only steric repulsion but also electrostatic repulsion were provided against particle aggregation. However, Fig. 9 reveals that the pH value of the ferrofluid decreased gradually during the process of titration by using methacrylic acid as a titrant. Once the pH value of the ferrofluid was lower than 5, the ferrofluid became unstable within several seconds and the aggregation of particles could be observed visually. It was because the ionization degree of acid groups would be lower than 90% when the pH value of the ferrofluid was lower than 5, and this condition could not provide sufficient electrostatic repulsion to preserve the stability of Fe$_3$O$_4$ nanoparticle suspension in ferrofluid. Consequently, the stability of ferrofluid could be controlled by adjusting the pH value of the solution, and therefore the ferrofluid could be applied properly.

3.5. XRD analysis

To investigate the effect of PAA oligomers on the preparation of Fe$_3$O$_4$ nanoparticles, the Fe$_3$O$_4$ nanoparticles prepared in the absence of PAA oligomers and the ferrofluid prepared in the presence of PAA oligomers were analyzed by XRD and the results are shown in Fig. 10. The results show
that the Fe$_3$O$_4$ nanoparticles prepared in the absence of PAA oligomers had six diffraction peaks at 2θ = 30.2°, 35.7°, 43.4°, 53.7°, 57.4°, and 62.9°, which were the characteristic peaks of standard Fe$_3$O$_4$ crystal (isometric-hexoctahedral crystal system). However, the ferrofluid prepared in the presence of PAA oligomers had an additional diffraction peak at ca. 2θ = 32.8°, which was the characteristic peak of Fe$_2$O$_3$ crystal (trigonal–hexagonal scalenohedral crystal system). These results indicate that some Fe$_2$O$_3$ particles consisted in the ferrofluid because the particles were synthesized in the initial condition of low pH value (in PAA oligomer solution pH 2.43), although the pH value of the final ferrofluid was 8.8 after the addition of ammonium solution. At the beginning of the precipitation reaction, the acidic condition would oxidize the Fe$_3$O$_4$ or oxidize the Fe$^{2+}$ to form Fe$^{3+}$ partially so that an initial mole concentration ratio of 1/2 in these ions would end up with excess quantities of Fe$^{3+}$ and form Fe$_2$O$_3$ as mentioned in Eq. (5). In addition, the broad diffraction peaks shown in Fig. 10 indicate that the particle size of magnetite was very small according to the Debye–Scherrer equation [34,35],

$$D = \frac{0.9\lambda}{\beta \cos \theta},$$

(9)

where $D$ is the average crystallite size (Å), $\lambda$ is the wavelength of X-rays (Cu$K_\alpha$: $\lambda = 1.5418$ Å), $\theta$ is the Bragg diffraction angle, and $\beta$ is the full width at half maximum (FWHM) (in radians). By using Eq. (9) and the FWHM of the diffraction peak at 2θ = 35.7°, the crystallite size of the Fe$_3$O$_4$ precipitated in the presence of PAA oligomer could be calculated to be 5.2 nm, which was smaller than the crystallite size of Fe$_3$O$_4$ (8.3 nm) synthesized in the absence of PAA oligomers. This result also shows that the presence of PAA oligomers reduced the particle size of Fe$_3$O$_4$ dramatically due to the fact that the carboxylic acid groups of PAA oligomers not only promoted nucleation but also inhibited growth of the Fe$_3$O$_4$ particles. Furthermore, the broad diffraction peak appeared in the range from 10° to 40° indicates the existence of amorphous PAA oligomers coated on the surface of Fe$_3$O$_4$ particles.

3.6. TGA analysis

Fig. 11 shows the TGA analysis of the pure Fe$_3$O$_4$ nanoparticles, the PAA oligomer ($M_w \sim 2000$), and the ferrofluid particles synthesized in the presence of PAA oligomers. The pure Fe$_3$O$_4$ nanoparticles showed insignificant weight loss from 100 to 800°C. But there were three stages during degradation shown in the TGA curve of PAA oligomers. The initial two stages of degradation (250–300°C) was due to the dehydration (Eq. (10)) [36–38] and decarboxylation (Eq. (11)) [36–38] of the carboxylic acid groups of the PAA oligomers, while the main degradation temperature of the PAA oligomer was near 400°C:

$$\text{HO} - \text{COO}^- + \text{H}_2\text{O} \rightarrow \text{HOCH}_2\text{COO}^-$$

(10)

The TGA curve of ferrofluid particles also showed three-stage decomposition. The first-stage decomposition, which occurred near 200°C, was due to dehydration (Eq. (12)), which was caused by the destruction of polyelectrolyte complex between COO$^-$ and NH$_4^+$ and the formation of amide bonds

$$\text{R} - \text{C} - \text{O} - \text{NH}_4^+ \rightarrow \text{R} - \text{C} - \text{NH}_2 + \text{H}_2\text{O}$$

(12)

Then the temperature was raised to 400°C to enable the main degradation of PAA oligomer to occur. The interesting decomposition which occurred at 700°C was due to the deoxidization of the Fe$_2$O$_3$ particles, which consisted in the ferrofluid as mentioned above, to form Fe$_3$O$_4$ at 700°C under N$_2$ atmosphere. Therefore, both the decomposition of PAA oligomers and the deoxidization of Fe$_2$O$_3$ contributed to the weight loss of the TGA analysis of ferrofluid particles. The residual weight percentage of ferrofluid particles after the TGA analysis (about 60%) was mainly the contribution of the Fe$_3$O$_4$ nanoparticles and a small amount of char yield induced from PAA oligomers. The results indicate that the weight ratio of PAA oligomers (~40%) to Fe$_3$O$_4$ nanoparticles (~60%) was nearly 1:2, where the ferrofluid was dialyzed before the TGA measurement. Theoretically, from the feed composition the weight ratio of PAA oligomers to Fe$_3$O$_4$ nanoparticles in the ferrofluid was cal-
Fe$_3$O$_4$ nanoparticles (50 emu/g), which was lower than the saturation magnetization of pure Fe$_2$O$_3$ nanoparticles was determined by using TGA analysis), the amount of nanoparticles in the ferrofluid, where the amount of Fe$_2$O$_3$ nanoparticles was lower than that of Fe$_3$O$_4$ nanoparticles. In addition, a lot of carboxylic acid groups of PAA oligomers were bound on the surface of Fe$_2$O$_3$ particles formed in the ferrofluid in the presence of PAA oligomers, the remanence and coercivity of the ferrofluid particles were zero and the magnetic hysteresis loop was not shown. It was concluded that the nanoparticles synthesized in this work were superparamagnetic.

3.7. Squid analysis

It is well known that the Fe$_3$O$_4$ nanoparticles show a superparamagnetic property. But when there was a quantity of Fe$_2$O$_3$ in the ferrofluid and the PAA oligomers were chemically bonded to nanoparticles, the magnetic property of the ferrofluid particles should be examined. The result of the SQUID analysis is shown in Fig. 12. When a magnetic field was applied, the dipolar particles aligned themselves with the applied magnetic field, and resulted in a measurable magnetization. The saturation magnetization of the ferrofluid was about 35 emu/g (emu per gram of nanoparticles) in the ferrofluid, where the amount of nanoparticles was determined by using TGA analysis), which was lower than the saturation magnetization of pure Fe$_3$O$_4$ nanoparticles (50 emu/g), due to the fact that the saturation magnetization reduced significantly when the particle size was smaller than 10 nm [39]. Also, the saturation magnetization of Fe$_2$O$_3$ nanoparticles was lower than that of Fe$_3$O$_4$ nanoparticles [40]. So the saturation magnetization of nanoparticles in the ferrofluid, which consisted of a small amount of Fe$_2$O$_3$ nanoparticles, was lower than that of pure Fe$_3$O$_4$ nanoparticles. Furthermore, the remanence and coercivity were zero, and there was no magnetic hysteresis loop as shown in Fig. 12. The magnetic susceptibility and permeability were calculated to be 0.024 and 1.302 (in the applied magnetic field from 0 to 100 G), which reveals superparamagnetic behavior [18,41]. These results indicate that the magnetic nanoparticles synthesized in the presence of PAA oligomers were superparamagnetic.

4. Conclusion

In this work, the method of co-precipitation was used to synthesize a water-based, stable ferrofluid that contained Fe$_3$O$_4$ nanoparticles in the presence of PAA oligomers. The PAA oligomers promoted the nucleation and inhibited the growth of the magnetic iron oxide so as to produce the smaller Fe$_3$O$_4$ nanoparticles. In addition, a lot of carboxylic acid groups of PAA oligomers were bound on the surface of Fe$_3$O$_4$ nanoparticles to suspend the Fe$_3$O$_4$ particles in a fine state of dispersion. The PAA oligomers provided both electrostatic and steric repulsion against particle aggregation, and the stability of dispersions could be controlled by adjusting the pH value of solution. Although there were some Fe$_2$O$_3$ particles formed in the ferrofluid in the presence of PAA oligomers, the remanence and coercivity of the ferrofluid particles were zero and the magnetic hysteresis loop was not shown. It was concluded that the nanoparticles synthesized in this work were superparamagnetic.

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