



Cite this: *CrystEngComm*, 2017, **19**, 6694

Effect of precursor concentration on size evolution of iron oxide nanoparticles†

Hamed Sharifi Dehsari, ^a Anielen Halda Ribeiro, ^a Bora Ersöz, ^a Wolfgang Tremel, ^b Gerhard Jakob ^c and Kamal Asadi ^{*a}

Thermal decomposition is a promising route for the synthesis of magnetic nanoparticles. The simplicity of the synthesis method is counterbalanced by the complex chemistry of the system such as precursor decomposition and surfactant-reducing agent interactions. Control over nanoparticle size is achieved by adjusting the reaction parameters, namely, the precursor concentration. The results, however, are conflicting as both an increase and a decrease in nanoparticle size, as a function of increasing concentration, have been reported. Here, we address the issue of size-controlled synthesis *via* the precursor concentration. We synthesized iron oxide nanoparticles with sizes from 6 nm to 24 nm with narrow size distributions. We show that the size does not monotonically increase with increasing precursor concentration. After an initial increase, the size reaches a maximum and then shows a decrease with increasing precursor concentration. We argue that the observation of two different size regimes is closely related to the critical role of the amount of surfactant. We confirm the effect of surfactant amount on nucleation and growth and explain the observed trend. Furthermore, we show that the nanoparticles show size-dependent but superior superparamagnetic properties at room temperature.

Received 1st August 2017,
Accepted 16th October 2017

DOI: 10.1039/c7ce01406f

rsc.li/crystengcomm

Introduction

Iron oxide nanoparticles have been the focus of intense research because of their magnetic properties,^{1,2} biocompatibility^{3,4} and bright application prospect in catalysis,^{5,6} magnetic fluidity,⁷ biomedicine^{3,8–10} biotechnology,^{11–14} and data storage.^{15,16} Various approaches such as coprecipitation,¹⁷ hydrothermal and solvothermal chemistry,¹⁸ polyol synthesis,¹⁹ reverse micelle,²⁰ sol-gel,²¹ and thermal decomposition^{22–24} techniques have been developed for the synthesis of iron oxide nanoparticles. Thermal decomposition is considered to be a promising and reproducible route for highly magnetic nanoparticles with uniform morphology, narrow size distribution and high crystallinity.^{22,23} In thermal decomposition, an iron complex, typically iron(III) acetylacetone (Fe(acac)₃), is dissolved in a high boiling point solvent, and oleic acid and oleylamine are added as surfactants. Upon heating, Fe(acac)₃ decomposes in the presence of the surfactants.²² Precise control over the size and dispersity of magnetic nanoparticles is crucial for different applications like biotechnology.^{9–11} The

nanoparticle size, morphology, dispersity, and crystallinity are controlled, among other factors, by the precursor,^{25,26} solvent,^{27,28} surfactant^{26,29} and their respective concentrations.^{30–32} Typically, the LaMer model³³ is used to explain the nucleation and growth of nanoparticles. Under identical heating and synthetic conditions for different reactions, the nucleation rate depends on supersaturation and thereby on the initial precursor concentration. Therefore, tuning the precursor concentration has been used to control the size of nanoparticles.^{26,27} Concentration tuning can be achieved in different ways, *e.g.* changing the amount of solvent or precursor. Concentration tuning by changing the amount of solvent is straightforward because the solvent has only a dilution effect. By changing the precursor concentration, the ratio between the surfactant and the precursor also varies. Several studies have shown that an increase in concentration may lead to both an increase^{25,27,34,35} and a decrease^{26,36} of the particle size. A conclusive explanation is still lacking.

Here, we address the effect of concentration on the final size (distribution) and magnetic properties of nanoparticles. First, as a reference, a nanoparticle batch was prepared and characterized under “standard conditions”. Size tuning by controlling the precursor concentration is achieved in two ways: by changing (1) the amount of solvent or (2) the amount of the precursor. We show that the particle size monotonically decreases with increasing amount of solvent,

^a Max Planck Institute for Polymer Research, Ackermannweg 10, Mainz, Germany.
E-mail: asadi@mpip-mainz.mpg.de

^b Institute of Inorganic Chemistry and Analytical Chemistry, Johannes Gutenberg University, Mainz, Mainz, Germany

^c Institute of Physics, Johannes Gutenberg-University Mainz, Mainz, Germany

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c7ce01406f



whereas for changing the precursor amount, two size regimes are observed, first an increase and then a decrease in size as the amount of precursor increases. We show that the ratio of the surfactant to the precursor plays a crucial role in tuning the size by changing the concentration, and we explain the observed opposite increasing/decreasing trend reported in the literature.^{25–27,34,35}

Iron oxide nanoparticles are superparamagnetic whose magnetic properties depend on their size. Magnetic measurements show that the nanoparticles possess room-temperature magnetizations close to the theoretically predicted values.

Experimental section

Iron(III) acetylacetone (97%), oleylamine (OAM, >70%), benzyl ether (BE, technical grade 99%), oleic acid (OAC, technical grade 90%), hexane, ethanol and acetone were all purchased from Sigma Aldrich. 1,2-Hexadecanediol (99%) was purchased from TCI. All chemicals were used as received.

The standard sample was prepared under “standard conditions”, wherein a three-necked round-bottom flask was charged with 2 mmol of iron acetylacetone, 6 mmol of OAC and 6 mmol of OAM, 10 mmol of 1,2-hexadecanediol, and 20 mL of benzyl ether. The mixture was heated to 110 °C, and maintained at that temperature for 60 min under vacuum. The temperature was then raised to 180 °C under an N₂ blanket with a heating rate of 6.5 °C min⁻¹ and kept at that temperature for 120 min to fully decompose the precursor. Subsequently, the temperature was raised to ~300 °C with a constant heating rate of 3.3 °C min⁻¹ and refluxed for 1 hour. Upon cooling of the solution, the nanoparticles were precipitated by addition of ethanol. The nanoparticles were washed three times with a mixture of toluene/ethanol/acetone followed by centrifugation (6000 rpm, 10 min), and finally stored under argon in toluene or hexane.

From standard conditions, the amount of solvent (series A), the total amount of precursor (B) and the total amount of surfactant (C) were varied. Other synthesis and post synthesis parameters/processes were kept unchanged with respect to the standard conditions. A summary of the experimental conditions is given in Table 1.

The particle size was characterized by transmission electron microscopy (TEM) and high resolution TEM (HRTEM) with an accelerating voltage of 120 kV using a JEOL JEM1400 and 200 kV using a FEI Tecnai F20 200 kV, respectively. The size distribution was obtained from statistical size

Table 1 Summary of experimental conditions. A, B and C are the variable parameters, where A = 28 mL (A₁), 24 mL (A₂), 16 mL (A₃), 12 mL (A₄); B = 0.6 mmol (B₁), 1.2 mmol (B₂), 1.6 mmol (B₃), 3 mmol (B₄), 4.5 mmol (B₅), 6 mmol (B₆); C = 2 mmol (C₁), 4 mmol (C₂), 8 mmol (C₃), 10 mmol (C₄)

Reactants	Standard conditions	Series A	Series B	Series C
Benzyl ether (ml)	20	A ₁ –A ₄	20	20
Fe(acac) ₃ (mmol)	2	2	B ₁ –B ₆	2
OAC (mmol)	6	6	6	C ₁ –C ₄
OAM (mmol)	6	6	6	C ₁ –C ₄

analysis of more than 2000 particles.¹⁵ The X-ray diffraction (XRD) pattern was recorded at room temperature using a diffractometer equipped with a monochromatic copper radiation source CuK α (λ = 1.5406 Å). The pattern was collected in the 15–65° (2 θ) range with a scan step of 0.03°. The mean size and lattice parameter of the crystal domains were calculated from the XRD pattern by using the Scherrer³⁷ and Bragg³⁸ equations. Magnetization measurements of powder samples were performed using a VSM (Cryogenic Ltd) magnetometer. Hysteresis loops $M(H)$ were measured under a maximum applied field of 50 kOe at 2 K and at room temperature. Zero-field-cooling (ZFC) and field-cooling (FC) magnetization curves were also measured at 100 Oe in the temperature range from 5 to 300 K. Thermogravimetric measurements (TGA) were performed on dried powder samples from 20 to 800 °C at a heating rate of 10 °C min⁻¹ under N₂. The number of ligands on the surface of the nanoparticle (grafting density) was extracted from the TGA/DTG data.

Results and discussion

Under standard conditions, nanoparticles with regular polyhedral morphologies were obtained, as shown in Fig. 1a. The size histogram can be fitted well with a normal Gaussian distribution. It gives a mean size of 13.00 nm with a standard deviation of 1.1 nm and hence a polydispersity of ~8.5% (Fig. 1b). The reference nanoparticles are monodisperse with a narrow size distribution. Particle aggregation is effectively hindered by the surfactant, and well dispersed colloidal solutions in hexane or toluene were obtained. Due to the size monodispersity and regular morphology, the nanoparticles have a strong tendency to self-assemble in a closely packed arrangement (Fig. S1†).

The high resolution TEM (HRTEM) image of the nanoparticles, shown in the inset of Fig. 1a, demonstrates that the particles have a highly crystalline structure. The crystalline structure is extended up to the edge of the nanoparticles, thereby minimizing the thickness of the disordered shell^{24,39} at the particle surface. It also shows different crystal planes depending on the orientation of the NPs with respect to the direction of the electron beam. To further investigate the crystallinity, the selected area electron diffraction (SAED) pattern of standard nanoparticles is shown in Fig. 1c. The *d* spacings in the diffraction patterns were calculated, and they matched well with the reported values for stoichiometric Fe₃O₄, as shown in the inset of Fig. 1c.^{22,28}

Fig. 1d shows the XRD patterns of the standard sample. The peak positions and relative intensities of the nanoparticles are in good agreement with the reported diffractograms of magnetite.⁴⁰ The narrow sharp peaks reveal the high purity of the synthesized nanoparticles. The calculated mean value of inter-planar distances using Bragg's law³⁸ (ESI†) amounts to 8.3778 Å in accordance with the literature value.^{22,28,41} The crystal structure is indexed to the fcc inverse cubic spinel structure of magnetite.^{22,40–42} The crystallite size of the nanoparticles was calculated to be 11.8 ±



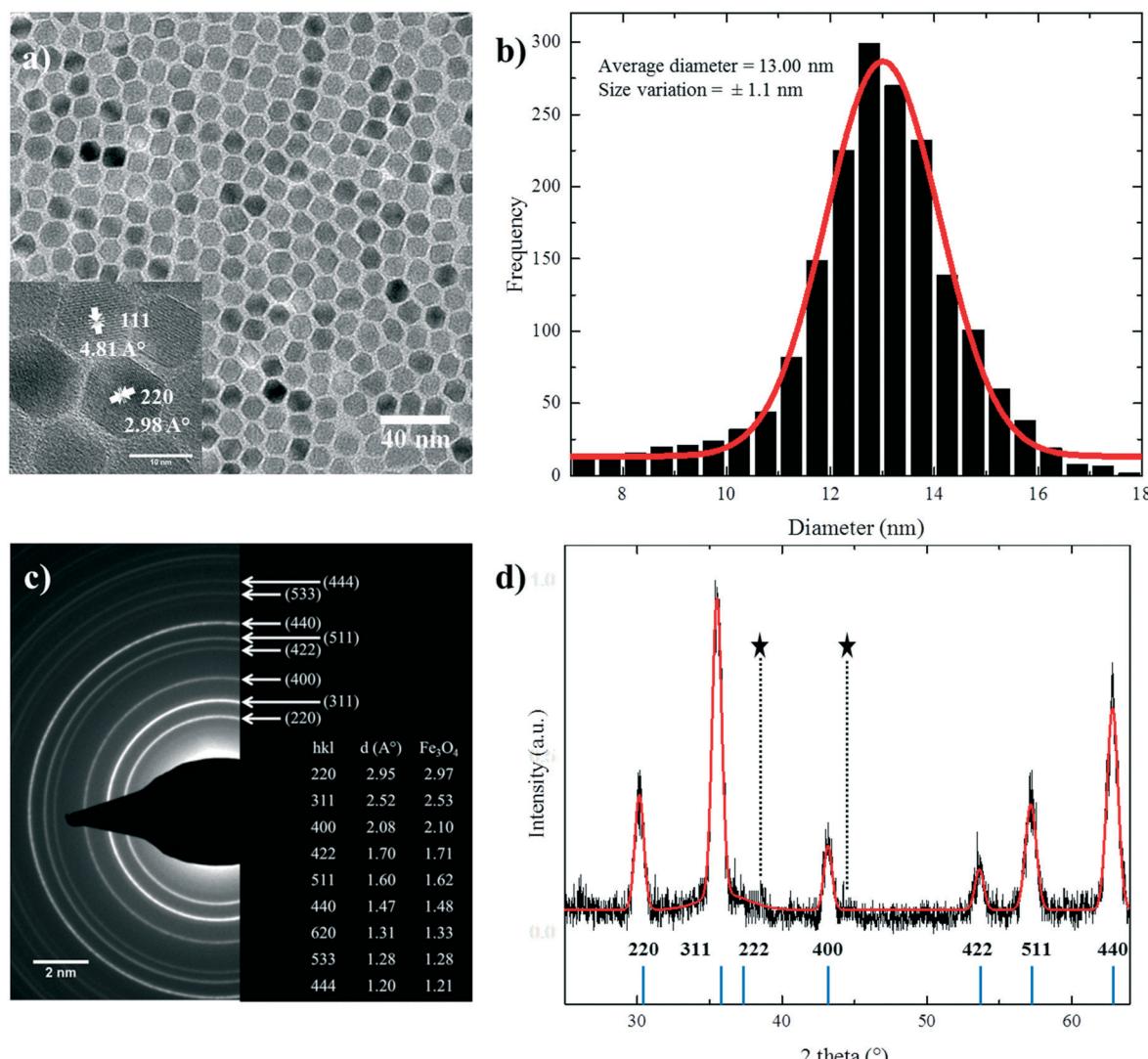


Fig. 1 (a) TEM image of nanoparticles prepared under standard conditions. The inset shows a typical HRTEM image where crystalline planes are indicated. (b) Size distribution obtained for more than 2000 particles. (c) Selected area electron diffraction (SAED) pattern of the standard reference nanoparticle. The lattice spacing, d (Å), was determined using the diffraction rings and their respective hkl indexes. (d) XRD patterns of the standard reference nanoparticle (black line) and the corresponding modeling result (solid red line). The positions of the Bragg reflections are represented by vertical blue lines. Holder reflections are indicated by stars.

0.17 nm from the XRD diffractograms using Scherrer's formula^{37,43} (ESI†), which is in good agreement with the size determined by statistical analysis of the TEM images.

The ligand shell around the nanoparticles consists predominantly of oleate, as determined from the FTIR spectra of the nanoparticles (Fig. S2a†). The bands at around 3000 and 2800 cm⁻¹ are for the methyl stretch on the surface of the nanoparticles. The bands at 1300–1650 cm⁻¹ are due to the asymmetric and symmetric COO⁻ bands of oleate, and the bands found at around 700 and 400 cm⁻¹ are characteristic Fe–O absorption bands of iron oxide.^{44,45}

The surfactant coverage was determined by TGA (Fig. S2b†). The graphs show three weight loss plateaus.^{46,47} The first plateau below 200 °C is attributed to the evaporation of adsorbed water and/or solvent molecules from the powder. The second and the third plateaus between 200–400 °C and

500–750 °C correspond to desorption from the surface and the decomposition of the surfactant molecules, respectively.

In the next step, we investigate the influences of the amount of *A*) solvent, *B*) precursor, and *C*) surfactant on the size evolution (dispersity) of the nanoparticles.

Amount of solvent

Different reactions were performed where only the amount of solvent (*A*) was systematically varied from 12 ml (A_1), 16 ml (A_2), 20 ml (standard sample), 24 ml (A_3) to 28 ml (A_4). The TEM images of the resulting nanoparticles are shown in Fig. 2a–d. The XRD diffractograms (Fig. S3†) show a comparison of crystallinity with the standard reference nanoparticles. The TGA traces for all the nanoparticles (A_1 – A_4) show (Fig. S4†) that the grafting densities were almost ~3 molecules per

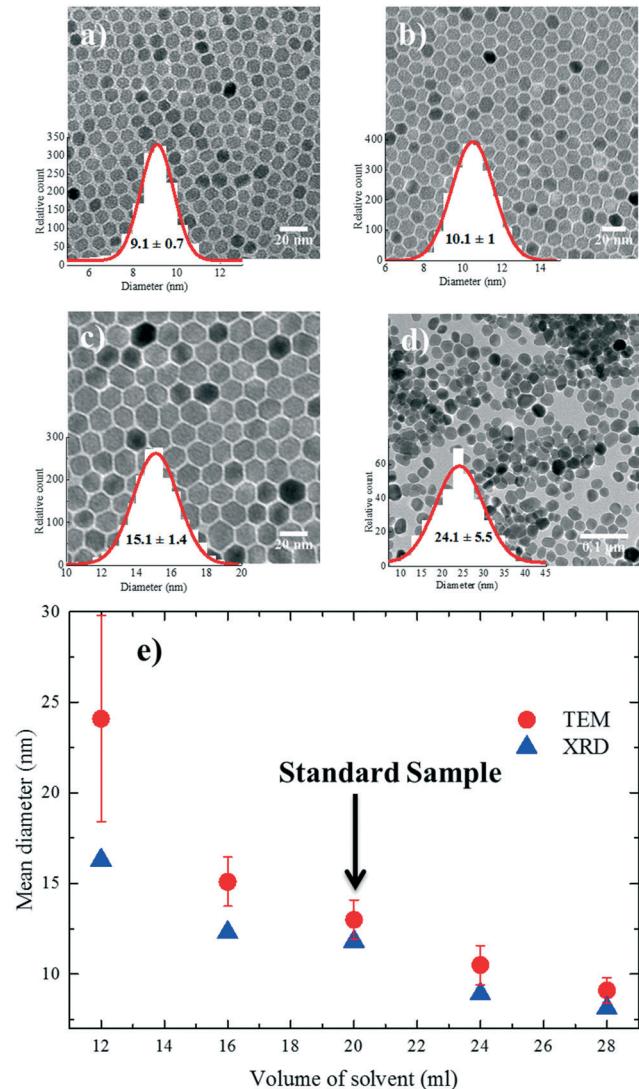


Fig. 2 TEM images and particle size distribution of samples (a) A_1 , (b) A_2 , (c) A_3 and (d) A_4 . (e) Evolution of nanoparticle size as a function of solvent volume. Nanoparticle diameters were determined from both TEM and XRD analyses.

nm^2 compared to the standard reference (Table S1†). The nanoparticle size extracted from both TEM and XRD is given in Fig. 2e. The nanoparticles A_1 – A_3 are truly monodisperse with polydispersities well below 10%. With an increasing amount of solvent (from 12 ml to 28 mL), *i.e.* decreasing concentration, the average size of the nanoparticles drops from 24.1 ± 5.5 nm to 9.1 ± 0.7 nm.

The observed drop in size with the increase in solvent volume can be explained by the decrease in the concentration of growth species (monomer) in the reaction medium. At a low solvent volume, the concentration of the monomer in the solution is high. The concentration of the available monomers at the interface of the nuclei (the crystal growth front) is close to that of the bulk solution. Hence, the diffusion distance for the monomers is shorter, which leads to a higher mass transfer and therefore a higher growth rate.²⁷ As a result, larger nanoparticles are formed during the same growth period

compared to the standard conditions. We note that for low solvent volumes (high concentrations), nanoparticle growth is reaction-controlled, as the growth rate is mainly governed by reactions on the surface of the nanoparticles. As the amount of solvent increases (low concentration), the concentration of the precursor, and therefore the monomer, is lowered which increases the diffusion constant and reduces the growth rate due to less mass transfer in the reaction

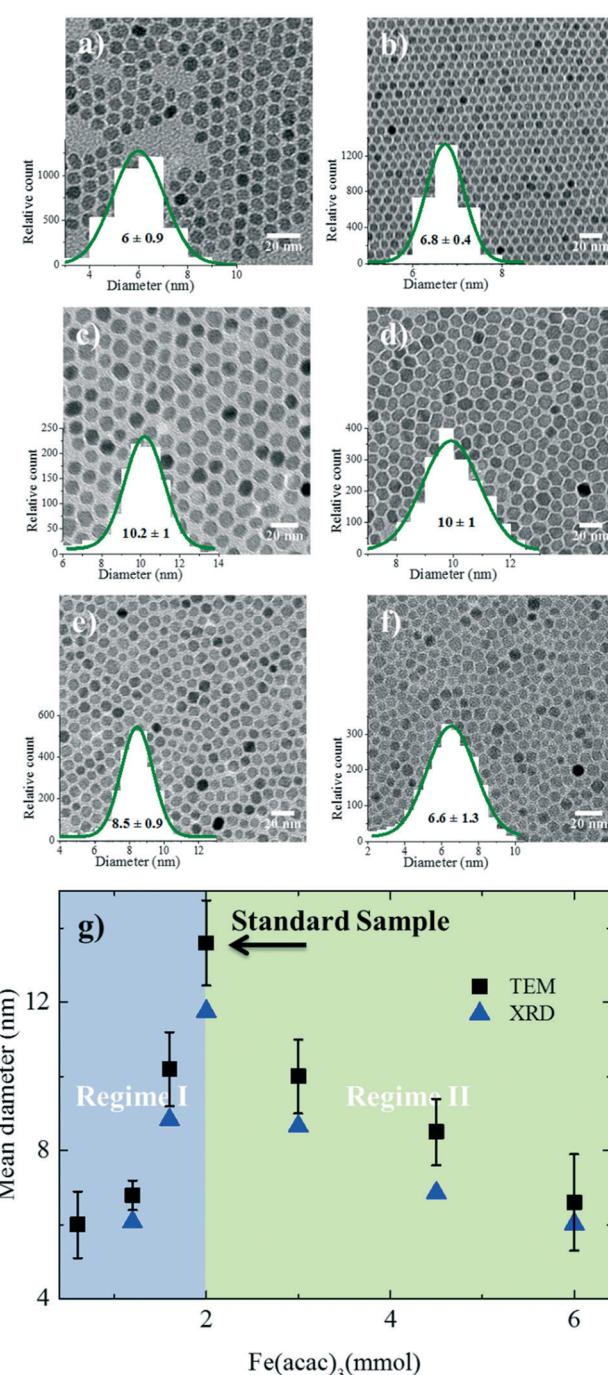


Fig. 3 TEM images of samples (a) B_1 , (b) B_2 , (c) B_3 , (d) B_4 , (e) B_5 and (f) B_6 . Particle size distributions obtained by fitting TEM histograms are also shown. (g) Evolution of the nanoparticle size (TEM and XRD) as a function of precursor amount. The dashed line is a guide to the eye.

medium. Nanoparticle growth is therefore diffusion-limited; as a result, smaller nanoparticles are formed during the same growth time compared to the reference conditions.

Amount of precursor

The amount of $\text{Fe}(\text{acac})_3$ (*B*) was systematically varied from 0.6 to 6 mmol (B_1 – B_6). The TEM images of the resulting nanoparticles with the corresponding size distribution are shown in Fig. 3a–f. XRD diffractograms are given in Fig. S5a.† The nanoparticle size evolution is given in Fig. 3g, and two different size regimes were observed. In the first regime, the size of the nanoparticles increases from 6 ± 0.9 nm to 13.00 ± 1.1 nm by increasing the amount of the precursor from 0.6 to 2 mmol. The increase in size can be understood based on the increased concentration of the monomer (growth species) in a fixed reaction volume, as discussed for the solvent case. To show that the governing mechanism is the same, we have plotted the size evolution for both series *A* and *B* as a function of the precursor concentration in Fig. S6a.† The trend observed for the case of the solvent variation describes well the first regime in Fig. 3g. The trend of increasing size is held up to a precursor concentration of 2 mmol, beyond which the nanoparticle size continuously drops. We note that as the concentration of the precursor increases in a fixed reaction volume, two parameters are changed simultaneously: the precursor concentration and the ratio between the surfactants and $\text{Fe}(\text{acac})_3$. The drop in size can be attributed to the decreasing amount of the surfactants available for the stabilization of the monomers. Therefore, the saturation concentration of the monomer (growth species) increases and more nuclei are formed. As a result, less monomer is available for the growth and hence smaller nanoparticles are formed.^{30,31,48} To substantiate the role of the surfactant to precursor ratio, we systematically varied the amount of surfactant in the next step.

The role of surfactant amount

Several studies have investigated the effect of surfactant concentration. By changing the amount, both trends of either increasing^{26,28,30,31,49–51} or decreasing size^{28,29,46,52} have been observed. To elucidate the role of precursor/surfactant ratio, different syntheses were performed wherein only the amount of surfactant ($\text{OAC} = \text{OAM} = \text{C}$) was systematically varied from 2 mmol to 10 mmol (C_1 – C_4). The TEM images of the resulting nanoparticles and the XRD diffractograms are shown in Fig. 4a–d and S5b,† respectively. The mean nanoparticle size (Fig. 4e) continuously increases from 5.8 ± 0.8 nm to around 16.3 ± 2 nm as the precursor/surfactant ratio decreases. As the amount of surfactant increases, more oleate molecules react with the precursor and more stable monomers with reduced reactivity are formed.^{28,30,31,48,49,53} According to LaMer's model³³ for nucleation and growth, a reduced active monomer concentration reduces the nucleation rate and hinders the formation of a large number of nuclei and hence favors the growth of larger nanoparticles. We have also plotted the size evolution for both series *B* and *C* as a function of

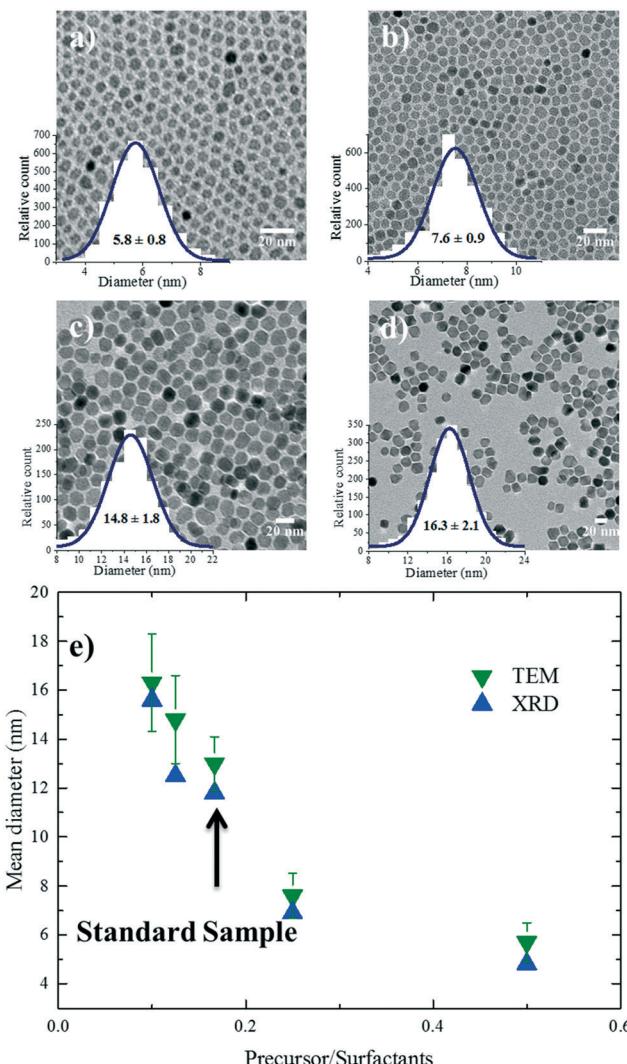


Fig. 4 TEM images of samples (a) C_1 , (b) C_2 , (c) C_3 and (d) C_4 . Particle size distributions obtained by fitting TEM histograms are also shown. (e) Size (based on TEM and XRD) evolution of the nanoparticles as a function of precursor/surfactant ratio. The dashed line is a guide to the eye.

precursor/surfactant ratio in Fig. S6b.† The trend observed for the case of the surfactant variation describes well the second regime in Fig. 3g. It has also been shown that an excess of surfactants prevents nanoparticle growth by blocking the growth sites and stabilizing the growth species,^{28,48} and manifests a dramatic drop in the mass reaction yield of the nanoparticles.

The magnetic properties of particles depend on their size.^{2,29} Representative magnetization hysteresis loops of samples A_1 – A_4 as a function of applied field measured at 300 and 2 K are given in Fig. 5a and b, respectively. At 300 K, no hysteresis loops were observed for samples A_1 – A_4 , confirming that the nanoparticles are superparamagnetic. However, sample A_4 (24.1 ± 5.7 nm) showed a hysteresis loop with a small coercivity, H_C (~ 10 Oe), and a remnant magnetization, indicating that the critical nanoparticle size is reached.^{4,50,54} The hysteresis loops at 2 K confirm that all the samples are ferri-ferromagnetic at 2 K.



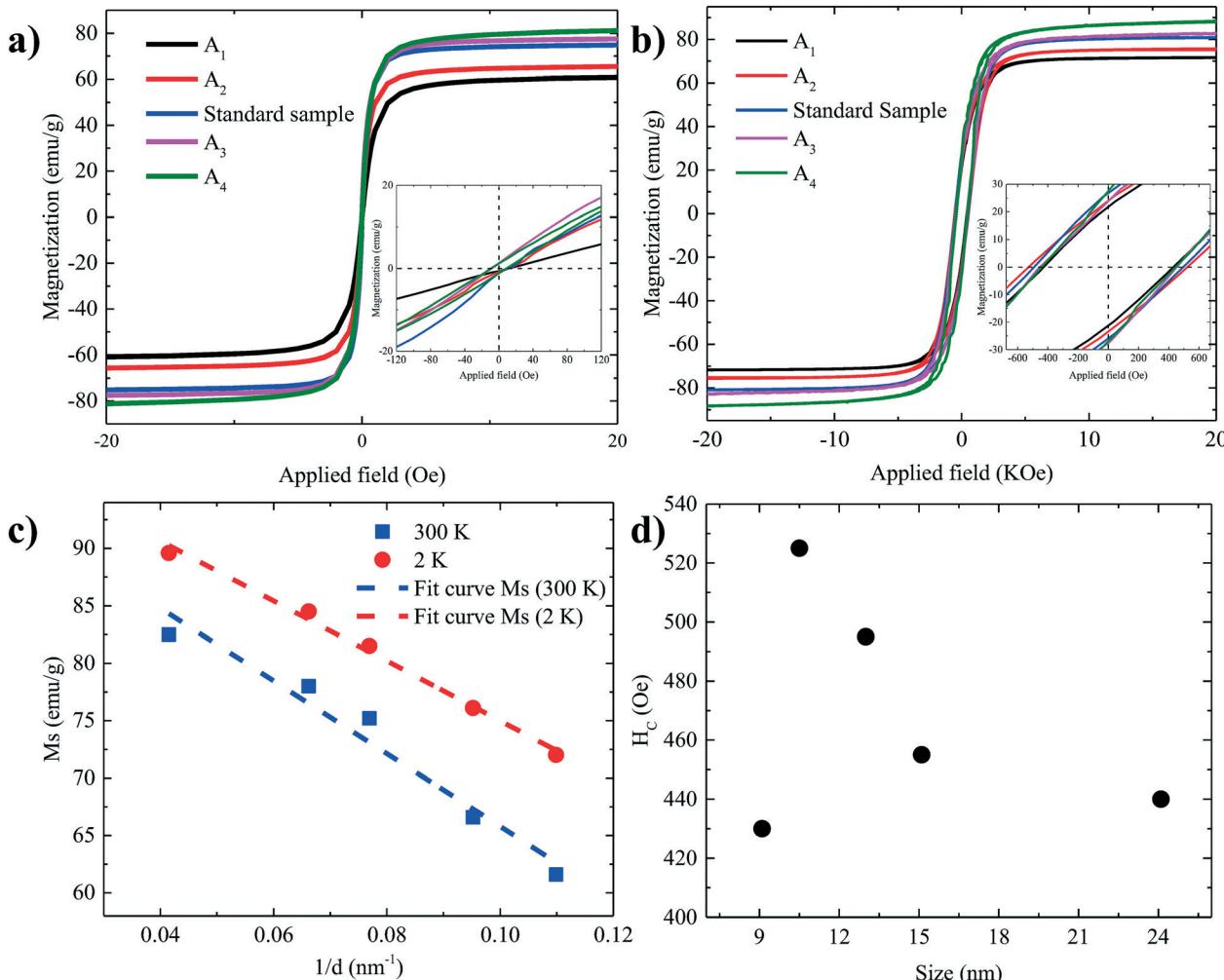


Fig. 5 Hysteresis loops of all samples at (a) 300 K and (b) 2 K. (c) The values of M_s as a function of $1/d$ in both 300 K and 2 K. (d) The values of H_c as a function of size.

Table 2 Summary of the magnetic properties of all samples at room temperature and 2 K. SPM and ferri-ferro are superparamagnetic and ferri-ferromagnetic

Samples	D_{TEM} (nm)	M_s (300 K) (emu g ⁻¹)	M_s (2 K) (emu g ⁻¹)	H_c (300 K) (Oe)	H_c (2 K) (Oe)	M_r (2 K) (emu g ⁻¹)	T_b (K)	K (10 ⁵ erg cm ⁻³)
A ₁	9.1	61.6	72	SPM	430	21.1	103 (±13)	9.314 (±2.25)
A ₂	10.5	66.6	76.1	SPM	525	23.9	170 (±20)	8.42 (±2.48)
Standard	13.0	75.2	81.5	SPM	495	26.3	222 (±20)	6.66 (±1.73)
A ₃	15.1	78	84.5	SPM	455	25.6	252 (±19)	4.83 (±1.38)
A ₄	24	82.5	89.6	10	440	26	ferri-ferro	ferri-ferro

The values of the saturation magnetization at 300 K and 2 K are listed in Table 2. As the size of the nanoparticles increases, M_s approaches the values reported for the bulk phase magnetite (84 emu g⁻¹) and the theoretically predicted value (98 emu g⁻¹) at low temperature.^{22,39,55-58}

The nanoparticles are covered with a non-magnetic shell that is composed of the surfactant layer and a disordered spin layer (spin canting layer).^{59,60} The thickness of the shell e can be calculated using the following relation:^{58,61,62}

$$M_s = M_s(\text{bulk}) \left(1 - \frac{6e}{d} \right) \quad (1)$$

Fig. 5c shows M_s plotted as a function of inverse nanoparticle diameter, $1/d$, at both 300 K and 2 K, which can be fitted well with eqn (1). The calculated values of the shell thickness amount to 0.54 nm and 0.43 nm at 300 K and 2 K, respectively.

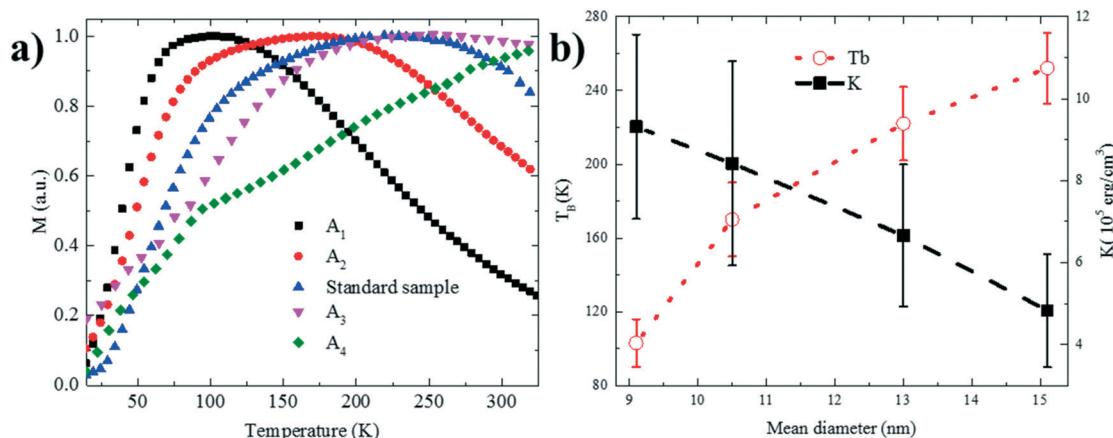


Fig. 6 (a) Temperature dependence of magnetization measured after zero-field cooling (ZFC) at 100 Oe. The magnetization data are normalized with respect to the value at the maximum of the ZFC magnetization $M(T_B)$ for each sample. (b) Values of T_B (blocking temperature) and K (anisotropy constant) as a function of size.

The coercivity, H_C , at 2 K increases with increasing nanoparticle size, as shown in Fig. 5d, from 430 Oe for 9.1 nm to 525 Oe for 10.5 nm. A further increase in the size of the nanoparticle reduces H_C by shifting from a single to a multi-domain regime.^{2,4,28,54} The flip of magnetic moments in each domain is controlled by magnetic anisotropy energy, domain wall motion and thermal energy. When the nanoparticle diameter approaches the regime of a single domain, domain wall motion does not exist. By increasing the size, the magnetic anisotropy energy increases. As a result, H_C increases. However, by further increasing the size, multidomain particles are formed. Due to the existence of the domain wall motion, the coercivity is reduced.^{2,4}

The transition temperature from superparamagnetic to ferro-ferrimagnetic and the blocking temperature T_B were determined with zero-field cooling (ZFC) and field cooling (FC) in an applied magnetic field of 100 Oe between 5–320 K (Fig. 6a). With increasing temperature, the thermal energy increases, and the ZFC curve starts to plunge as T_B is reached. The values of T_B are given in Fig. 6b as a function of size. With increasing size, the volume of the nanoparticle increases, and T_B moves to higher temperatures.^{23,25,56} The increase in T_B with size is caused by the increased magnetocrystalline energy. Consequently, higher thermal energies are required to unblock the magnetic moment of larger nanoparticles. Thermal anisotropy constants were calculated based on the equation:^{23,25}

$$K = \frac{25k_B T_B}{V} \quad (2)$$

where V is the mean volume of the iron core, T_B is the blocking temperature, and k_B is the Boltzmann constant. The anisotropy constant K increases because of the increased surface-to-volume ratio as the nanoparticle size decreases.^{23,25,31,58,63} A summary of the magnetic properties of the nanoparticles is given in Table 2.

Conclusion

In summary, we have shown that size-controlled synthesis of iron oxide nanoparticles can be achieved by changing the concentration *via* (1) the amount of solvent or (2) the amount of precursor. Two competing mechanisms were identified which influence the nucleation and growth of the nanoparticles: (i) concentration-dependent monomer diffusion and (ii) monomer stabilization by an excessive amount of surfactant. The former controls the size when the amount of solvent (to some extent, the precursor concentration) is changed. At a fixed solvent amount, lowering the precursor amount changes the precursor/surfactant ratio which leads to monomer stabilization, less nucleation and hence growth of larger nanoparticles. The resulting nanoparticles show superparamagnetic behavior at room temperature and ferro-ferrimagnetic behavior at low temperature with a high M_s close to the theoretical value for magnetite.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

K. A. acknowledges the Alexander von Humboldt Foundation for the funding provided in the framework of the Sofja Kovalevskaja Award, endowed by the Federal Ministry of Education and Research, Germany. The authors acknowledge the support from the Max-Planck Institute for Polymer Research (Mainz, Germany) and the technical help of Dr. Ingo Lieberwirth, Michael Steiert, Verona Maus, Michelle Beuchel, Ann-Kathrin Schönbein and Elham khodabakhshi. Open Access funding provided by the Max Planck Society.

References

- 1 C. Yang, J. Wu and Y. Hou, *Chem. Commun.*, 2011, **47**, 5130–5141.
- 2 A. G. Kolhatkar, A. C. Jamison, D. Litvinov, R. C. Willson and T. R. Lee, *Int. J. Mol. Sci.*, 2013, **14**, 15977–16009.
- 3 N. A. Frey, S. Peng, K. Cheng and S. Sun, *Chem. Soc. Rev.*, 2009, **38**, 2532–2542.
- 4 L. Wu, A. Mendoza-Garcia, Q. Li and S. Sun, *Chem. Rev.*, 2016, **116**, 10473–10512.
- 5 B. Sahoo, S. K. Sahu, S. Nayak, D. Dhara and P. Pramanik, *Catal. Sci. Technol.*, 2012, **2**, 1367–1374.
- 6 X. Mou, X. Wei, Y. Li and W. Shen, *CrystEngComm*, 2012, **14**, 5107–5120.
- 7 M. López-López, J. Durán, A. Delgado and F. González-Caballero, *J. Colloid Interface Sci.*, 2005, **291**, 144–151.
- 8 M. De, P. S. Ghosh and V. M. Rotello, *Adv. Mater.*, 2008, **20**, 4225–4241.
- 9 L. H. Reddy, J. L. Arias, J. Nicolas and P. Couvreur, *Chem. Rev.*, 2012, **112**, 5818–5878.
- 10 R. Dinali, A. Ebrahiminezhad, M. Manley-Harris, Y. Ghasemi and A. Berenjian, *Crit. Rev. Microbiol.*, 2017, **43**, 493–507.
- 11 F. Assa, H. Jafarizadeh-Malmiri, H. Ajamein, N. Anarjan, H. Vaghari, Z. Sayyar and A. Berenjian, *Nano Res.*, 2016, **9**, 2203–2225.
- 12 K. Ulbrich, K. i. Holá, V. Šubr, A. Bakandritsos, J. Tucek and R. Zboril, *Chem. Rev.*, 2016, **116**, 5338–5431.
- 13 W. Zhang, Z.-L. Yu, M. Wu, J.-G. Ren, H.-F. Xia, G.-L. Sa, J.-Y. Zhu, D.-W. Pang, Y.-F. Zhao and G. Chen, *ACS Nano*, 2017, **11**, 277–290.
- 14 M. Mahmoudi, S. Sant, B. Wang, S. Laurent and T. Sen, *Adv. Drug Delivery Rev.*, 2011, **63**, 24–46.
- 15 L. Wu, P.-O. Jubert, D. Berman, W. Imai, A. Nelson, H. Zhu, S. Zhang and S. Sun, *Nano Lett.*, 2014, **14**, 3395–3399.
- 16 Q. Dai, D. Berman, K. Virwani, J. Frommer, P.-O. Jubert, M. Lam, T. Topuria, W. Imai and A. Nelson, *Nano Lett.*, 2010, **10**, 3216–3221.
- 17 Y. S. Kang, S. Risbud, J. F. Rabolt and P. Stroeven, *Chem. Mater.*, 1996, **8**, 2209–2211.
- 18 T. Daou, G. Pourroy, S. Begin-Colin, J. Greeneche, C. Ulhaq-Bouillet, P. Legaré, P. Bernhardt, C. Leuvrey and G. Rogez, *Chem. Mater.*, 2006, **18**, 4399–4404.
- 19 W. Cai and J. Wan, *J. Colloid Interface Sci.*, 2007, **305**, 366–370.
- 20 Y. Lee, J. Lee, C. J. Bae, J. G. Park, H. J. Noh, J. H. Park and T. Hyeon, *Adv. Funct. Mater.*, 2005, **15**, 503–509.
- 21 N. Tang, W. Zhong, H. Jiang, X. Wu, W. Liu and Y. Du, *J. Magn. Magn. Mater.*, 2004, **282**, 92–95.
- 22 S. Sun, H. Zeng, D. B. Robinson, S. Raoux, P. M. Rice, S. X. Wang and G. Li, *J. Am. Chem. Soc.*, 2004, **126**, 273–279.
- 23 J. Park, K. An, Y. Hwang, J.-G. Park, H.-J. Noh, J.-Y. Kim, J.-H. Park, N.-M. Hwang and T. Hyeon, *Nat. Mater.*, 2004, **3**, 891–895.
- 24 M. Unni, A. M. Uhl, S. Savliwala, B. H. Savitzky, R. Dhavalikar, N. Garraud, D. P. Arnold, L. F. Kourkoutis, J. S. Andrew and C. Rinaldi, *ACS Nano*, 2017, **11**, 2284–2303.
- 25 J.-H. Huang, H. J. Parab, R.-S. Liu, T.-C. Lai, M. Hsiao, C.-H. Chen, H.-S. Sheu, J.-M. Chen, D.-P. Tsai and Y.-K. Hwu, *J. Phys. Chem. C*, 2008, **112**, 15684–15690.
- 26 R. Hufschmid, H. Arami, R. M. Ferguson, M. Gonzales, E. Teeman, L. N. Brush, N. D. Browning and K. M. Krishnan, *Nanoscale*, 2015, **7**, 11142–11154.
- 27 H. Zeng, P. M. Rice, S. X. Wang and S. Sun, *J. Am. Chem. Soc.*, 2004, **126**, 11458–11459.
- 28 W. Baaziz, B. P. Pichon, S. Fleutot, Y. Liu, C. Lefevre, J.-M. Greeneche, M. Toumi, T. Mhiri and S. Begin-Colin, *J. Phys. Chem. C*, 2014, **118**, 3795–3810.
- 29 P. Guardia, N. Pérez, A. Labarta and X. Batlle, *Langmuir*, 2009, **26**, 5843–5847.
- 30 C. J. Meledandri, J. K. Stolarczyk, S. Ghosh and D. F. Brougham, *Langmuir*, 2008, **24**, 14159–14165.
- 31 A. Demortiere, P. Panissod, B. Pichon, G. Pourroy, D. Guillou, B. Donnio and S. Begin-Colin, *Nanoscale*, 2011, **3**, 225–232.
- 32 F. B. Effenberger, R. A. Couto, P. K. Kiyohara, G. Machado, S. H. Masunaga, R. F. Jardim and L. M. Rossi, *Nanotechnology*, 2017, **28**, 115603–115610.
- 33 V. K. LaMer and R. H. Dinegar, *J. Am. Chem. Soc.*, 1950, **72**, 4847–4854.
- 34 L. I. Cabrera, Á. Somoza, J. F. Marco, C. J. Serna and M. P. Morales, *J. Nanopart. Res.*, 2012, **14**, 1–14.
- 35 N. Miguel-Sancho, O. Bomati-Miguel, A. G. Roca, G. Martinez, M. Arruebo and J. Santamaria, *Ind. Eng. Chem. Res.*, 2012, **51**, 8348–8357.
- 36 Y. Li, J. Liu, Y. Wang and Z. L. Wang, *Chem. Mater.*, 2001, **13**, 1008–1014.
- 37 G. Arrhenius, *J. Chem. Educ.*, 1955, **32**, 228–229.
- 38 K. Venkatesan, D. R. Babu, M. P. K. Bai, R. Supriya, R. Vidya, S. Madeswaran, P. Anandan, M. Arivanandhan and Y. Hayakawa, *Int. J. Nanomed.*, 2015, **10**, 189–198.
- 39 J. Mohapatra, A. Mitra, D. Bahadur and M. Aslam, *CrystEngComm*, 2013, **15**, 524–532.
- 40 R. M. Cornell and U. Schwertmann, *The iron oxides: structure, properties, reactions, occurrences and uses*, John Wiley & Sons, 2003.
- 41 S. Belaïd, S. Laurent, M. Vermeersch, L. Vander Elst, D. Perez-Morga and R. N. Muller, *Nanotechnology*, 2013, **24**, 055705–055712.
- 42 X. Teng and H. Yang, *J. Mater. Chem.*, 2004, **14**, 774–779.
- 43 Y. Eom, M. Abbas, H. Noh and C. Kim, *RSC Adv.*, 2016, **6**, 15861–15867.
- 44 M. Klokkenburg, J. Hilhorst and B. Erne, *Vib. Spectrosc.*, 2007, **43**, 243–248.
- 45 K. Yang, H. Peng, Y. Wen and N. Li, *Appl. Surf. Sci.*, 2010, **256**, 3093–3097.
- 46 C. Moya, X. Batlle and A. Labarta, *Phys. Chem. Chem. Phys.*, 2015, **17**, 27373–27379.
- 47 I. Castellanos-Rubio, M. Insausti, E. Garaio, I. G. de Muro, F. Plazaola, T. Rojo and L. Lezama, *Nanoscale*, 2014, **6**, 7542–7552.
- 48 M. Yin, A. Willis, F. Redl, N. J. Turro and S. P. O'Brien, *J. Mater. Res.*, 2004, **19**, 1208–1215.



49 W. Y. William, J. C. Falkner, C. T. Yavuz and V. L. Colvin, *Chem. Commun.*, 2004, 2306–2307.

50 Y. Zhu, F. Jiang, K. Chen, F. Kang and Z. Tang, *J. Alloys Compd.*, 2011, **509**, 8549–8553.

51 Z. Chen, *Synth. React. Inorg., Met.-Org., Nano-Met. Chem.*, 2012, **42**, 1040–1046.

52 J. Vargas and R. Zysler, *Nanotechnology*, 2005, **16**, 1474–1479.

53 W. W. Yu and X. Peng, *Angew. Chem., Int. Ed.*, 2002, **41**, 2368–2371.

54 D. L. Leslie-Pelecky and R. D. Rieke, *Chem. Mater.*, 1996, **8**, 1770–1783.

55 K. O'grady and A. Bradbury, *J. Magn. Magn. Mater.*, 1983, **39**, 91–94.

56 A. Roca, M. Morales, K. O'Grady and C. Serna, *Nanotechnology*, 2006, **17**, 2783–2788.

57 D. Jiles, *Introduction to magnetism and magnetic materials*, CRC press, 2015.

58 D. Caruntu, G. Caruntu and C. J. O'Connor, *J. Phys. D: Appl. Phys.*, 2007, **40**, 5801–5809.

59 R. Kodama, A. Berkowitz, E. McNiff Jr and S. Foner, *J. Appl. Phys.*, 1997, **81**, 5552–5557.

60 S. Schwaminger, D. Bauer, P. Fraga-García, F. Wagner and S. Berensmeier, *CrystEngComm*, 2017, **19**, 246–255.

61 M. Zheng, X. Wu, B. Zou and Y. Wang, *J. Magn. Magn. Mater.*, 1998, **183**, 152–156.

62 C. Liu and Z. J. Zhang, *Chem. Mater.*, 2001, **13**, 2092–2096.

63 S. Mørup and E. Tronc, *Phys. Rev. Lett.*, 1994, **72**, 3278–3281.

