Unravelling the growth mechanism of the co-precipitation of iron oxide nanoparticles with the aid of synchrotron X-Ray diffraction in solution†

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Co-precipitation is the most ubiquitous method for forming iron oxide nanoparticles. For a typical co-precipitation synthesis, the pH of a ferrous and/or ferric ion solution is increased via the addition of a base. The latter can be added either slowly (a steady addition over either minutes or hours) or fast (a one-time addition) resulting in an abrupt increase in the pH. However, understanding the mechanism of particle formation is still lacking, which limits the reproducibility of the co-precipitation reaction due to intermediate phases still being present in the final product. In this work, we study in detail a co-precipitation synthesis with an abrupt increase in pH via the addition of sodium carbonate. Fast and reproducible mixing at defined precursor and base solution temperatures was achieved utilising a flow reactor. Transmission electron microscopy, electron diffraction and room temperature 57Fe Mössbauer spectroscopy showed a distinct transition from an amorphous ferrihydrite phase to a mixture of magnetite-maghemite (Fe3O4/γ-Fe2O3). Synchrotron X-ray diffraction revealed the initial formation of crystalline iron hydroxide carbonate (green rust) plates occurring before the Fe3O4/γ-Fe2O3 appeared. The ferrihydrite particles increase in size over time as the proportion of iron hydroxide carbonate plates are re-dissolved into solution, until the ferrihydrite particles crystallise into Fe3O4/γ-Fe2O3.

Introduction

Iron oxide nanoparticles (IONPs) are used extensively in applications in biomedicine, magnetic storage media, water treatment and catalysis.1–7 The most ubiquitous synthetic method used to date to form IONPs is the co-precipitation method, which is carried out in aqueous solutions containing ferric (Fe3+) and ferrous (Fe2+) salts as precursors,8 to which a base is added at moderate temperatures (<100 °C). While IONPs mostly contain magnetite and maghemite phases, there are many other iron phases that can form during synthesis; each of them has very different magnetic properties. Even though the co-precipitation reaction is widely used, the mechanism of the reaction is still poorly understood, due to a severe lack of information on how intermediates are formed. This often leads to difficulties in obtaining reproducible syntheses, as the intermediates can remain present to a varying degree products.

Proposed mechanisms for IONP formation originate from studies involving a slow transition in pH (from ~2 to above 8) and investigating the phases formed during the transition through multiple pH regimes.9,10 These studies show intermediate formation of iron oxyhydroxide species9–15 and iron hydroxide2,10,12,16,17 species, all of which transition to magnetite or maghemite (Fe3O4, and γ-Fe2O3 respectively) over time. From the iron oxyhydroxide intermediates the most commonly reported are goethite,9–11 and a poorly crystalline or amorphous ferrihydrite.14,15 For example, Baumgartner et al. showed with cryo-transmission electron microscopy (TEM) the formation of amorphous ferrihydrite which attaches to the surface of the growing IONP.15

However, most co-precipitation syntheses proceed via a single and abrupt increase in pH, which is also the most common approach for large-scale and continuous reactor systems14,18–21 due to the shorter process times. Hence, the understanding of the abrupt pH transitions is extremely important. Ahn et al. showed that abrupt pH changes result in different particle formation routes than slow pH additions,
showing the precipitation of ferrous hydroxide [Fe(OH)₂] and its transition to several intermediate phases (also due to pH inhomogeneities in the solution after mixing) including lepidocrocite.²² Milosevic et al. continuous flow reactors to study more abrupt pH transitions and in situ magnetic characterisation to study the growth mechanism. From the magnetic measurements they hypothesised that the particles originated as a matrix of primary ferrihydrite of around 3 nm in size, and grew over time to form 7 nm iron oxide particles.¹⁴

In this work, we have used a millifluidic flow reactor to facilitate the fast mixing of the base and the precursor solution at a well-controlled reaction temperature to study the growth mechanism of Fe₃O₄/γ-Fe₂O₃ NPs. The flow reactor allowed for an abrupt transition of the pH with minimal inhomogeneities of either the pH or temperature of the solution. Characterisation was performed with transmission electron microscopy (TEM), synchrotron X-ray diffraction (XRD) in solution, and room temperature ⁵⁷Fe Mössbauer spectroscopy. The abrupt pH transition caused the formation of two intermediate phases, i.e., amorphous ferrihydrite and iron hydroxide carbonate (green rust). The ferrihydrite grew at the expense of the iron hydroxide carbonate and crystallised into magnetite/maghemite.

Experimental procedure

Chemicals

Iron(II) chloride hexahydrate (96%) was obtained from Chem Cruz, USA. Iron(II) chloride tetrahydrate (99.0%) and sodium carbonate decahydrate (99.0%) were from Honeywell, Fluka, USA. All chemicals were used as received. For all experiments deionised water was used.

Synthesis of iron oxide nanoparticles

In a typical experiment a 0.033 M FeCl₂·4H₂O and 0.066 M FeCl₃·6H₂O solution was made up in deionised water. In a separate flask a 0.34 M solution of Na₂CO₃·10H₂O was prepared in deionised water. These two solutions were then loaded into 25 mL glass syringes (Hamilton 1000 series Gastight; Hamilton, Switzerland) and pumped with a dual syringe pump (Legato210; KDScientific, USA) at 5 ml min⁻¹ for both syringes through 6 m of PTFE tubing with an 1/16 inch (= 1.58 mm) outer diameter, and an inner diameter of 1 mm. This tubing was submerged in a water bath heated to 60 °C. Tubing of 6 m was sufficient to preheat the reagents to the final reaction temperature of 60 °C before mixing at the used flow rates.²² Mixing happened in a T-mixer made of ETFE (P-632; IDEX Health & Science, USA) with a channel diameter of 0.5 mm which was kept at 60 °C under stirring. The solution was collected for 30 s (after steady-state operation of the flow reactor was ensured), to fill the aging vessel with 5 ml solution. Subsequently, flow to the aging vessel was stopped and reaction was allowed to continue for up to 10 min. The procedure described above for the synthesis of iron oxide NPs is shown in Fig. 1A.

The experiments with only Fe²⁺ or Fe³⁺ were carried out identically to the procedure above, except they were either carried out with 0.1 M FeCl₂·4H₂O only or 0.1 M FeCl₃·6H₂O only.

Transmission electron microscopy

The transmission electron microscopy (TEM) images and electron diffraction were captured using a JEOL 1200 EX microscope with a tungsten filament operating at a 120 kV acceleration voltage. The high-resolution TEM (HRTEM) imaging was carried out on a FEI Titan G2 60–300 equipped with a Cs image corrector operating at 80 kV. The TEM samples were prepared by withdrawing 20 µL of solution from the stirred reaction flask every min (with the first sample withdrawn immediately after sampling at 30 s) and dropping it onto a carbon coated 200 mesh copper grid (EM Resolutions Ltd) sitting on a filter paper. The filter paper absorbed the excess liquid causing the solution to dry immediately and effectively quenching the reaction.

Synchrotron X-ray diffraction

The synchrotron X-ray diffraction was carried out at Elettra Synchrotron in Trieste, Italy using a wavelength of 1 Å for all measurements. Diffraction patterns were collected for 1 min in transmission mode at room temperature (~24 °C) through the quartz capillary (~1 mm inner diameter and <0.2 mm wall
thickness) of a flow cell mounted in the beamline. This made it possible to characterise the NPs in solution via injection into the quartz capillary after the specified amount of reaction time (1, 2, 3, 4 or 5 min). After these times, a volume of the nanoparticle solution was withdrawn from the aging vessel with a syringe through a thin PTFE tubing (inner diameter 0.5 mm, for rapid cooling to room temperature), and then fed into the quartz capillary, as shown in Fig. 1B. The process of filling the capillary typically took 25 s, and the measurement was started within the next minute. The detector position and angle was calibrated with a boron lanthanide (LaB6) reference powder in the same capillary setup. The resulting pattern was radially averaged and the background of the capillary and solution were subtracted as described in section 2 of the ESI (Fig. S1 and S2†). It should be noted that the phases formed are stable upon cooling in the capillary for at least 10 min (Fig. S3†).

$^{57}$Fe Mössbauer spectroscopy

Samples for room temperature $^{57}$Fe Mössbauer spectroscopy were withdrawn from the flask at the specified reaction time (1, 3 and 10 min), and pumped through 2 m of 1/16 inch tubing in an ice bath to quench the reaction. The samples were frozen with liquid nitrogen and then freeze dried to remove the water, before storage in a glovebox prior to measurements. Samples were further processed for the measurement by mixing the dried powder with sucrose to form a solid dispersion and mounted in a 2.1 cm coin shaped absorber.

A SeeCo W302 spectrometer (SeeCo Inc., UCA) was operated in transmission geometry and in constant acceleration mode. A $^{57}$Co in Rh foil was the source of the 14.4 keV γ-rays, with velocity calibration performed by recording a reference spectrum from a 10 μm thick foil of α-Fe at room temperature. All spectra were folded and baseline corrected using a cubic spline correction derived from the α-Fe calibration spectrum and following a protocol implemented in the Recoil analysis program. All spectra obtained for the 1 min samples were fitted using pairs of Lorentzian doublets, with the isomer shift and quadrupole splitting allowed to vary as free parameters; the best fit to the spectrum was determined by the lowest value of $\chi^2$. The 3 min and 10 min samples were fitted using the ‘center of gravity’ method. This method allows for the area weighted mean isomer shift of a magnetically split $^{57}$Fe Mössbauer spectrum to be determined without the need to assume any specific underlying model for the microenvironment of the Fe atoms, provided that the data is obtained from thin, texture free absorbers and possesses a flat, well defined background and that the sample comprises magnetite/maghemite only. It is for this reason we were unable to deduce an α-value of the t = 3 min sample, since it appeared to contain an appreciable amount of ferrihydrate (ca. 20%).

Results and discussion

The co-precipitation synthesis was carried out with sodium carbonate, which has been previously reported to result in slower growth kinetics than the strong bases, and thus would allow the kinetics to be readily studied. The described semi-flow procedure for the synthesis of iron oxide NPs, facilitated fast mixing (mixing times <50 ms) to rapidly obtain the reaction pH, hence minimise local pH variations, and a controlled reaction temperature of 60 °C throughout the synthesis (including the mixing step). Furthermore, it allowed for the co-precipitation synthesis to be studied multiple times under identical processing conditions, and thus guaranteeing reproducibility needed for a detailed study into the kinetics of nanoparticle synthesis. The samples synthesised were characterised by TEM, synchrotron XRD and room temperature $^{57}$Fe Mössbauer spectroscopy, carried out at the reaction times described above.

TEM characterisation

The initial particles formed immediately after mixing, as apparent from the change in colour to a dark black-reddish solution when exiting the T-Mixer (Fig. S4†). The first TEM sample was taken immediately after transfer to the aging vessel (i.e., 30 s) and showed particles of 2.2 ± 0.5 nm in size which formed aggregates (Fig. 2A). By 1 min the particles increased to 3.0 ± 0.7 nm. The particles continued to increase approximately linearly in size as the size distribution broadened at the same time (Fig. 2B). This continued until the particles reached a maximum size after 5 min of ~7 nm. With longer times (up to 10 min) the average particle size stayed the same, although the polydispersity increased (Fig. 2C).

The crystal structure of the TEM samples was studied with electron diffraction (Fig. S5†). Each diffraction pattern was radially averaged to allow it to be studied as a two-dimensional plot, and normalised (Fig. 2D). The initial sample-formed after 30 s had two broad peaks in the electron diffraction pattern at ~2.5 Å and ~1.5 Å. At 3 min the particles are crystalline; the pattern for iron oxide is clearly observed and stays as the only component observed till the end of the reaction. Electron diffraction on the TEM samples indicate that a transition between the initial phase to crystalline magnetite or maghemite (Fe₃O₄/Fe₂O₃) occurred between 2 to 3 min, which corresponds to a particle size of 3.6 ± 0.8 nm to 4.7 ± 1.1 nm.

Aberration corrected TEM was carried out at 80 kV to further study the pre-transition sample after 2 min of reaction. Two distinct structures are seen in TEM showing a smaller proportion of faceted plates of a roughly hexagonal shape and smaller nanoparticles 2–4 nm in size (Fig. 3A). The high-resolution imaging of the 2–4 nm nanoparticles did not show distinct crystallinity, with a few diffraction spots in the Fourier transform of the image at ~2.4 Å (Fig. 3B) and the high-resolution images showing a partially crystalline structure (Fig. 3D). High resolution imaging of part of the plate and the Fourier transform of the image showed hexagonal symmetry and spacings of ~2.5 Å and ~1.5 Å (Fig. 3C and E), indicating that the plate is highly crystalline. The spacings and symmetry seen in the Fourier transform is almost identical to characterisation of iron hydroxide carbonate (green rust) plates reported by McGill et al. The spacings of ~2.5 Å and ~1.5 Å are in the same position as the peaks in the electron diffraction of...
Fig. 2D at 30 s, 1 min and 2 min, and thus would originate from these plates. Green rust structures have been shown to adopt hexagonal plate-like structures with the CO$_3^{2-}$ or SO$_4^{2-}$ counter-ions.

Control experiments were carried out using either solely FeCl$_3$·6H$_2$O or FeCl$_2$·4H$_2$O. For the reaction with FeCl$_3$·6H$_2$O alone, only small poorly crystalline particles were formed (Fig. S6†), which did not grow further when the reaction was continued for up to 10 min. For FeCl$_2$·4H$_2$O alone, plates were formed which had the same electron diffraction patterns as observed in Fig. 2D at 30 s, 1 min and 2 min (Fig. S6†). The FeCl$_2$·4H$_2$O alone sample oxidised over time to form a brown solution. This indicates that the electron diffraction patterns from the mixed samples shown in Fig. 2D originated from the plate structures with the small spherical particles being amorphous, and moreover that the spherical particles formed would not grow further without Fe$^{2+}$ present in the reaction mixture.

**Synchrotron X-Ray diffraction studies**

Metal hydroxides can be unstable and the crystal structure decomposes when dried from solution or studied with an electron beam.$^{15,29,30}$ To avoid drying of sample, XRD was carried out through a capillary in which the nanoparticle solution was injected and studied with a synchrotron light source to achieve prompt measurements during the synthesis. The synchrotron XRD measurements were taken at set time points to retain the benefits of the controlled mixing times and temperature homogeneity of the flow reactor. In contrast, dedicated in situ cells normally have much smaller reaction volumes without the possibility for mixing.$^{31,32}$ The XRD patterns of samples are shown in Fig. 4A. The presence of Fe$_3$O$_4$/$\gamma$-Fe$_2$O$_3$ was first observed at 3 min as shown by XRD, and in agreement with the electron diffraction data (Fig. 2D). At 1 min and 2 min the XRD shows the formation of a crystalline phase which corresponds to iron hydroxide carbonate [Fe$_6$(OH)$_{12}$CO$_3$], which is formed of 4 Fe$^{2+}$ centres and 2 Fe$^{3+}$ centres. At 3 min, the peaks for the iron hydroxide carbonate and iron oxide co-exist, with the latter having a Scherrer size of 7 nm at 3 min, which increases to $\sim$9 nm at 5 min. The iron hydroxide carbonate makes up a small proportion of the volume of diffraacting objects that are eventually seen, surmised by comparing the XRD intensities after 5 min (Fig. 4B). The peak area of the iron hydroxide carbonate linearly decreases until peaks of magnetite (and/or maghemite) are observed, and these increase also linearly in peak area after the iron hydroxide carbonate peaks disappear (Fig. 4B).
57Fe Mössbauer spectroscopy studies

Room temperature 57Fe Mössbauer spectroscopy measurements were performed on the 1 min sample (i.e., before transition), the 3 min sample (at the transition) as well as the final product at 10 min. For comparison we also measured the 1 min sample of the Fe3+ sample alone, prepared in the absence of Fe2+. All mixtures were freeze dried to give a fine powder.

The spectrum of the product formed at 1 min (Fig. 5) corresponds to that of a quadrupole ferric doublet, with the best fit obtained using two Lorentzian doublets (each with an isomer shift ca. 0.35 mm s⁻¹) and which is reminiscent of ferrihydrite.33 There are very small differences between the samples prepared with and without Fe2+, namely a small increase in the sub-component with higher quadrupole splitting, which could be due to the increase of amorphous material formed when Fe2+ is omitted from the reaction (Fig. S7†). There is no sign of any Fe2+ from the iron hydroxide carbonate plates in either of these samples, which is likely to be due to their oxidation during isolation and the freeze-drying process.

The room temperature spectrum of the product at 3 min (Fig. 5) is a superposition of a doublet and broad sextet, indicative of a range of Fe environments within the sample. Best fitting of the spectrum was performed using Voigtian lines (to represent a Gaussian distribution of Lorentzian lines) and indicates that ca. 20% of the sample is the original ferrihydrite.
drite, with the remainder of the sample comprising of a poorly magnetic iron oxide. The presence of very broad absorption lines, which were best fit using a very broad distribution of hyperfine fields are likely to arise from a non-stoichiometric solid solution of magnetite/maghemite. The spectrum of the final product (10 min), comprises a partially collapsed magnetically split sextet pattern with very broad lines, and is indicative of magnetically blocked particles on the Mössbauer timescale, with a broad size distribution. The spectrum was fitted using the model independent ‘centre of gravity’ method from which we surmise that the numerical proportion of Fe atoms in a magnetite environment, \( \alpha \), was 0.40 ± 0.17 i.e., corresponding to a mixture of magnetite and maghemite.\(^{27}\) We did not observe any oxidation ageing of the material during the acquisition of any of the spectra, which took place over at least 3 d for each sample.

**Proposed mechanism of iron oxide nanoparticle formation**

Ferrihydrite is a poorly crystalline iron oxyhydroxide material with a general formula of \( \text{Fe}_{10}\text{O}_{14}(\text{OH})_2 \),\(^{34}\) formed from \( \text{Fe}_{13} \) clusters of \( \text{Fe}^{3+} \) known as Keggin clusters.\(^{35,36}\) The formation of the initial ferrihydrite, occurs by olation of the hydroxylated iron and oxolation.\(^{37}\) The ferrihydrite structure is formed of a central Keggin cluster with an iron depleted shell.\(^{38}\) These Keggin clusters that are central to ferrihydrite formation also have a similar crystal structure to the central magnetite unit.\(^{35}\) The formation of amorphous ferrihydrite primary particles or pre-nucleation clusters have been reported in numerous previous reports as an intermediate phase in iron oxide formation.\(^{10,14,15,36,37}\)

Iron hydroxide carbonate \( [\text{Fe}_6(\text{OH})_{12}\text{CO}_3] \) is a type of green rust. Green rust is a mixed valence 2-dimensional structure which is structurally similar to the amorphous \( \text{Fe(OH)}_2 \) sheets,\(^{11}\) and can be stabilised by a range of counter ions such as \( \text{CO}_3^{2−}, \text{SO}_4^{2−}, \text{Cl}^{−}, \text{Br}^{−} \) in either a hexagonal or rhombohedral crystal structure.\(^{17,28}\) The structure is formed via the olation of \( \text{Fe}^{2+} \) hydroxide tetrahedra\(^{11}\) and some incorporation of \( \text{Fe}^{3+} \) hydroxide tetrahedral centres, either by incorporation or oxidation.

From our data we observe a linear increase in particle size of the ferrihydrite particles via TEM and room temperature \(^{57}\)Fe Mössbauer spectroscopy with time, starting from initially formed 2.2 ± 0.5 nm particles. We observe a size dependent transition between the ferrihydrite and the magnetite/maghemite which occurs when the particles reach ~4 nm. This transition corresponds to the disappearance of peaks from the iron hydroxide carbonate phase in the X-ray diffraction pattern. The intensity of the iron hydroxide carbonate pattern decreases over time before and after the formation of iron oxide occurs. This indicates that as the reaction progresses the iron hydroxide carbonate structure is re-dissolved and the ions are incorporated into the ferrihydrite, providing a source of \( \text{Fe}^{2+/3+} \), causing it to grow and transition into magnetite/maghemite. The iron hydroxide carbonate acts as a reservoir of \( \text{Fe}^{2+} \) and \( \text{Fe}^{3+} \) ions leading to slow continuous growth first of the ferrihydrite and then subsequently of the magnetite/maghemite. A schematic of this growth mechanism, which differs from classical nucleation and growth concepts, is shown in Fig. 6. In reaction systems without carbonate ions present, a reservoir containing \( \text{Fe}^{2+} \) ions could be formed as a green rust structure with another counter ion \( \text{SO}_4^{2−}, \text{Cl}^{−} \text{or Br}^{−} \),\(^{17,28}\) or as amorphous \( \text{Fe(OH)}_2 \) sheets.\(^{11}\)

The incorporation of \( \text{Fe}^{2+} \) into the structure of magnetite has been previously predicted to be a rate limiting step.\(^{38–41}\) As \( \text{Fe}^{2+} \) has a higher solubility in water,\(^{42}\) it becomes the primary additive causing the primarily \( \text{Fe}^{3+} \) ferrihydrite particles to grow. Without the \( \text{Fe}^{2+} \) present, the ferrihydrite would not grow to magnetite (as in the control experiment, see Fig. S6†). Therefore, in a common co-precipitation synthesis, if there was an oxidation of the \( \text{Fe}^{2+} \), this would lead to its deficiency and hinder the conversion from ferrihydrite to \( \text{Fe}_3\text{O}_4/\gamma\text{Fe}_2\text{O}_3 \). This potentially leads to considerable problems with reproducibility of the synthesis of IONPs due to the presence of residual ferrihydrite particles, which are weakly magnetic and amorphous.
The amount of residual ferrihydrite in the final product can therefore be reduced by protecting the Fe$^{2+}$ ions from oxidation or increasing the amount of Fe$^{2+}$ in the reaction.

### Summary and conclusions

For this work a flow reactor was utilised to facilitate fast mixing (<50 ms) and well-controlled, hence reproducible, process conditions during a co-precipitation synthesis. Synchrotron X-ray diffraction (in solution, using a flow cell mounted in the beamline), as well as TEM and room temperature 57Fe Mössbauer spectroscopy studies were performed to unravel the particle formation mechanism.

These studies revealed that two initial phases were formed upon the direct mixing of the iron chloride precursor solution with the pH 12 solution of sodium carbonate. The phases were a poorly crystalline ferrihydrite phase formed primarily of Fe$^{3+}$ and crystalline iron hydroxide carbonate plates formed primarily of Fe$^{2+}$. The ferrihydrite was characterised by means of room temperature 57Fe Mössbauer spectroscopy and TEM, while the dynamic changes of the (metastable) iron hydroxide carbonate plates could only be monitored via synchrotron X-ray diffraction in solution. The iron hydroxide carbonate concentration decreased over time until it disappeared after 4 min of the reaction. The ferrihydrite phase grew with time from 2 nm particles until it transitioned to Fe$\gamma$O$_4$/$\gamma$-Fe$_2$O$_3$ when the particles reached ~4 nm in size, which happened after 3–4 min of the reaction. The ferrihydrite phase acted as the seeds that grew into the eventual Fe$\gamma$O$_4$/$\gamma$-Fe$_2$O$_3$ particles, while the iron carbonate plates acted as a feedstock to supply the iron ions for the particles to grow and the phase change to occur. Without the presence of Fe$^{2+}$ in solution, the formation and growth of the Fe$\gamma$O$_4$/$\gamma$-Fe$_2$O$_3$ NPs did not occur.

A new particle formation mechanism was detailed, differing drastically from the most commonly discussed classical nucleation and growth process. Such comprehensive understanding of the mechanism is essential not only for robust syntheses, but also for the tuning of particle properties for the targeted applications. In particular, the growth mechanism shows a separation of the nucleation and growth stages in the reaction which will allow chemists greater control over the size of the nanoparticles that can be readily synthesised via co-precipitation.

### Conflicts of interest

There are no conflicts of interest to declare.

### Acknowledgements

APL, MOB, AG, NTKT would like to thank EPSRC for funding (grant EP/M015157/1). AS thanks the EPSRC CDT for the Advanced Characterisation of Materials (grant EP/L015277/1) for his studentship. NTKT thanks AOARD (FA2386-17-1-4042 award). AH thanks the CERIC ERIC internal research project: Nano Analytics in Pharmaceutics for support. We would like to thank Giorgio Bais and Maurizio Polentarutti for support with the synchrotron XRD studies.

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