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In situ TEM oxidation study of Fe thin-film transformation to single-crystal magnetite nanoparticles

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ABSTRACT

In this work, we present an in situ transmission electron microscopy (TEM) study of Fe thin films to Fe nanoparticle formation and their oxidation to single-crystal magnetite nanoparticles. Amorphous Fe thin films were prepared by sputtering on TEM carbon grids. The thin Fe films were continuously heated in situ from room temperature to 700 °C under vacuum (4×10^{-4} Pa). With the increase in temperature, the continuity of the thin film starts breaking, and Fe nanoparticle nucleation centers are formed. At 600 °C, the thin film transforms into metallic Fe nanoparticles (NPs) with a small presence of different Fe oxide NPs. Further increase in the temperature to 700 °C resulted in the full oxidation of the NPs (i.e., no core-shell were found). Zero-loss energy filtered diffraction and HRTEM analysis of the lattice spacing reveals that all NPs have fully transformed into single-phase magnetite NPs. The structural study of the magnetite NPs shows that magnetite NPs are free of antiphase domain boundary defects. This work demonstrates that under low partial pressure of oxygen at elevated temperatures a complete oxidation of Fe NPs into magnetite single-crystal nanoparticles can be achieved.

Introduction

The study of iron oxide nanoparticles (NPs) has burgeoned during the last decades [1–3] due to their range of applications that include water purification [4], ferrofluids [5], hyperthermia treatments of tumors [6], chemical sensing [7, 8], magnetic

resonance imaging contrast enhancement agents [9], and catalyst support [10–12].

There are both chemical [13–17] and physical methods to produce iron oxide NPs [16, 18]. Physical methods include post-growth oxidation of sputtered Fe NPs [19, 20] and laser-induced pyrolysis [21]. Independently from the preparation methods,

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obtaining iron oxides NPs with bulk-like structures has been shown to be very challenging, *e.g.*, core/shell structures Fe/Fe-oxides are a common outcome [19, 20, 22] as well as the presence of antiphase boundaries [23]. Such type of defects, commonly occurring in thin Fe oxide films [24, 25], are known to reduce the magnetization of the material [23], detrimental to some of their applications. Although methods for the production on large scale of free-standing iron oxide NPs are important, the fundamental understanding of iron crystals formation and evolution under controlled gas and temperature is a key for the development of potential industrial applications. In this work, we demonstrate that annealing in low O₂ pressure of pre-sputtered iron thin films results in single-crystal Fe₃O₄ (magnetite) NP formation. By performing in situ TEM studies, we were able to characterize the structural phase transformations from thin Fe film to single-crystal magnetite NPs.

Historically ample literature has been produced on the oxidation mechanism of iron and phase transformations of iron oxides from bulk materials to thin films down to nanoparticles [16, 17, 26–31]. Phase diagrams for the Fe–O system in bulk materials are well known [32], but they are not directly applicable to nanoscale objects. In situ heating and gas environmental electron microscopy [33] provide unique opportunities to investigate the oxidation down to the nanometer and atomic scale in quasi-equilibrium conditions. One of the challenges present during this type of studies is the effect of the electron beam and the residual gas molecules in the microscope column [34] on the oxidation dynamics [35], hence the final product of the oxidation. In this work, we utilize the residual O₂ in the microscope column to perform low-pressure (4.0×10^{-4} Pa) oxidation study during in situ heating of iron sputtered thin films as a function of temperature. To avoid phase transformations driven by the electron beam, heating of the sample was done with the beam off, and the TEM imaging was performed for very short time (in order of seconds) by continuously changing area of observation. Due to the close lattice spacing of the iron oxide phases of interest (Fe₃O₄ and γ -Fe₂O₃), standard TEM imaging techniques are not able to distinguish between these two phases; here we show that energy filtered electron diffraction can overcome such a limitation.

Materials and methods

Samples for TEM investigation were deposited directly on amorphous holey carbon films supported by TEM Cu grids (AGAR Holey Carbon Films on 200 Mesh Copper Grids, AGS147) by glow discharge sputter coating. The holey carbon support film has a nominal thickness of 10–15 nm; such a thickness has been chosen as a compromise between ultrathin carbon films (3 nm) and standard carbon film (28–30 nm), in order to ensure robustness during the heating experiment in the presence of residual gas. The TEM grids were put centrally in the bottom of the vacuum chamber of a high-resolution plasma sputter coater (model JFC-2300HR, JEOL Ltd) with the carbon film facing up toward an iron target (SuperVac® Fe target 99.9% purity, 0.1 mm thickness, Testbourne Ltd). Sputtering was performed for 60 s at a current setting of 40 mA using gas argon (99.999% Zero Grade Argon, BOC); the deposited film thickness, as measured by the quartz monitor of the sputter coater, was below 1 nm.

In situ electron microscopy was performed using an environmental transmission electron microscope (TEM) operating at 300 kV (Titan, FEI) and equipped with a third-order image aberration corrector (Cetcor, CEOS). Samples were loaded on a single tilt heating holder (Gatan, model 628). Sample temperature was controlled and monitored by using a *SmartSet* hot stage controller (Gatan, model 901).

Results and discussion

The as-sputtered thin film supported by the carbon film was transferred in air and loaded to the TEM microscope column with a vacuum basic pressure of 4.0×10^{-4} Pa, as measured by the microscope column vacuum gauge, corresponding to a residual oxygen partial pressure of 0.8×10^{-4} Pa (calculated assuming a 21% Oxygen fraction as in the atmosphere). As shown in Fig. 1, both HRTEM images (Fig. 1a) and electron diffraction patterns (Fig. 1b) do not reveal any atomic ordering (no atomic fringes in HRTEM) neither clear sharp diffraction rings.

The temperature was then ramped up to 600 °C from room temperature, with a linear rate in 30 min. Before exposing the sample to the electron beam for imaging, the specimen holder was left to stabilize thermally to reduce drift for additional 60 min. After

this annealing step, nanoparticles with round shape with an average diameter of 5.2 nm were formed, as shown in Fig. 2a, b.

Diffraction patterns show that annealing has started the crystallization process, which resulted in crystalline NP formation within the amorphous/semicrystalline thin film. This is concluded from the simultaneous presence of sharp and diffuse diffraction rings presented in Fig. 2c–e. Detailed analysis of these diffraction rings using JEMS software revealed that the crystalline NPs have spacings mostly corresponding to bulk iron- α phase as shown in Fig. 2c, as well as some diffraction spots and a minor ring compatible with the other major bulk iron oxide phases such as magnetite (Fe_3O_4) as shown in Fig. 2d, hematite ($\alpha\text{-Fe}_2\text{O}_3$), and maghemite ($\gamma\text{-Fe}_2\text{O}_3$). Calculated diffraction patterns for wustite (FeO) and iron carbide (Fe_3C) do not fit the experimental data. In fact, in bulk ferrite (α -iron), carbon solubility is as low as < 0.00005 wt% at room temperature which can increase up to 0.02 wt%, at about 720 °C, where Fe is supersaturated with C in the interstitial positions. Quenching the supersaturated Fe down to the range 20–300 °C leads to Fe_3C precipitate formation [36]. In this work, we have not found any evidence (both by HRTEM and electron diffraction) of the Fe_3C structure at the experimental temperatures used; hence, solubility of C into NPs if

present does not result in a distinctive structural phase.

With further increase in the temperature, raised to 700 °C, the film completely dewets from the support and the formation of nanoparticles with increased abundance, size, and pronounced morphology is obtained (Fig. 3). Faceting of the NPs is now evident (Fig. 3a, b) compared to the round nanoparticles obtained at 600 °C (Fig. 2). We note that crystallinity of the NPs is extending well to their surfaces, in contrast to observed disorder at surfaces in metallic NPs [3]. The drastic phase transformation due to the increase in temperature is also evident from the diffraction patterns taken at 700 °C, as shown in Fig. 3. Diffraction ring analysis (as discussed later in Fig. 5a) confirms atomic planes spacing compatible with both magnetite (Fe_3O_4) and maghemite ($\gamma\text{-Fe}_2\text{O}_3$), indicating that the low pressure of oxygen in the microscope column is sufficient, at 700 °C, for a full oxidation of the formed nanoparticles. In fact, no evidences of core/shell structures or metallic nanoparticles were found by imaging or electron diffraction.

HRTEM imaging of single nanoparticles show lattice spacings corresponding to orientations of NPs along the major crystallographic axes [111], [100], and [2-1-1], Figure 4. Atomic spacing measurements from the HRTEM images are compatible with both maghemite and magnetite phases.

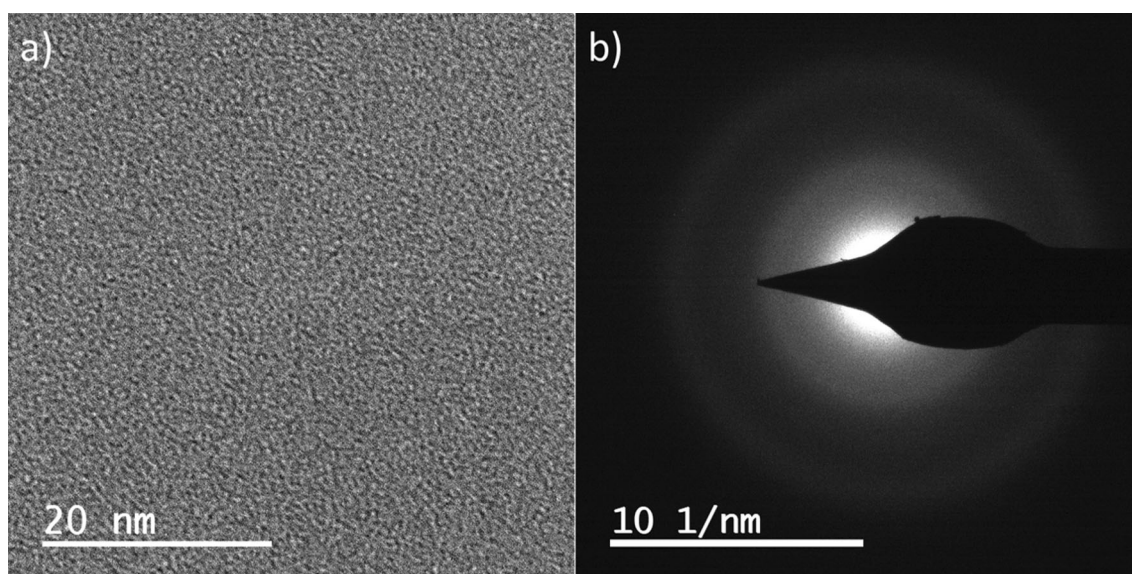
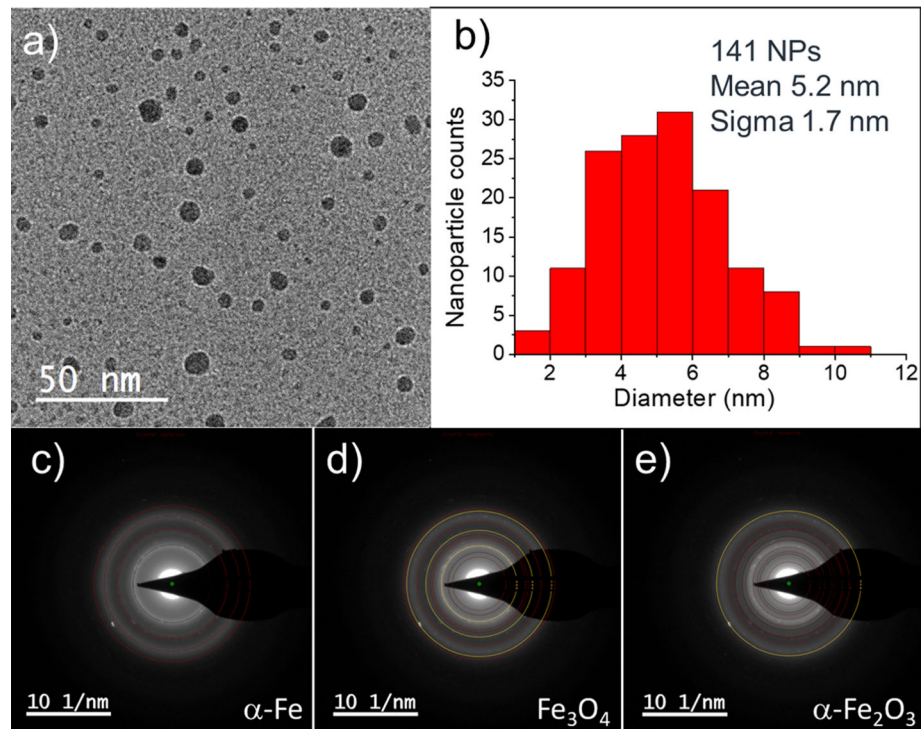


Figure 1 As sputtered Fe thin film before annealing. **a** Bright field TEM image showing the support and sputtered material; **b** diffraction pattern showing diffuse rings typical of amorphous materials.

Figure 2 Nanoparticle formation at 600 °C. **a** Bright field image showing nanoparticles formed onto the carbon film support; **b** nanoparticle statistic over 141 nanoparticles from several images showing a 5.2 nm mean NPs diameter (equivalent area circular diameter); diffraction pattern showing rings corresponding to the NP polycrystalline phase; DP analysis shows that the main phase is α -Fe (**c**), with extra rings compatible with the main iron oxide phases magnetite (**d**), hematite (**e**), and maghemite (not shown here).

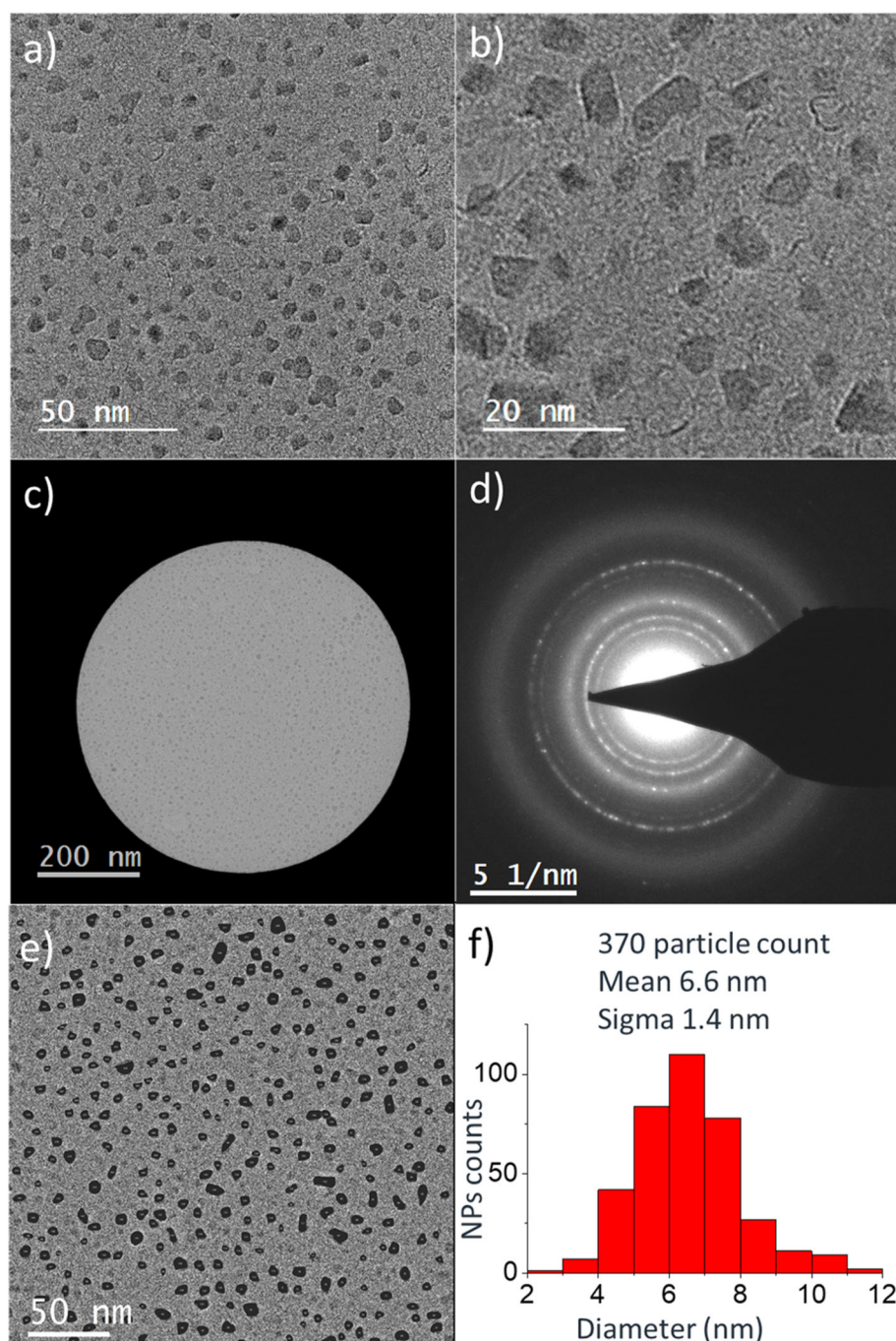


We note that the accuracy of spacing measurements using the HRTEM ($\sim 2\%$ error) does not allow to distinguish between the maghemite and magnetite NPs, since the difference of their lattice spacings is only $\sim 0.5\%$. This problem can be overcome by utilizing the super-reflections and kinematically allowed diffraction reflections for maghemite. Even though magnetite and maghemite share virtually almost the same crystallographic spacings (lattice constants are $a_{\text{magn}} = 8.397 \text{ \AA}$, $a_{\text{magh}} = 8.351 \text{ \AA}$), the broken symmetry along the c -axis of maghemite, i.e., $c = 3a$, provides unique characteristic super-reflections and kinematically allowed reflections of maghemite that are not present in magnetite diffraction patterns. As shown in Fig. 5b, the low index lattice spacings corresponding to (110), (210), (211), and (421) atomic planes should be present in the maghemite diffraction patterns while they are forbidden in magnetite. The most intense diffraction reflections of these, thus the easier to be identified, are due to (210) and (211) planes. The difficulty in imaging diffraction patterns at these low spatial frequencies is the proximity of transmitted (000) beam. The (000) beam contains the majority of the transmitted electron intensity, as well as diffusively scattered electrons which broaden the (000) beam; as a result, the low index plane reflections usually overlap

with the transmitted beam, as shown in Fig. 5a. We note that even with a beam stopper the low-angle Bragg scattered electrons will be superimposed with the spread of the (000) beam. In order to capture the low scattering vectors and separate them from the (000) beam, we used post-column electron energy filtering to reduce the background. Figure 5c shows a diffraction pattern taken at a longer camera length with a 5-eV energy window centered on the zero-loss peak (ZLP) to ensure the removal of inelastic scattered electron from the diffraction pattern. The energy filtered diffraction pattern clearly shows the (111) and (220) diffraction rings corresponding to Fe_3O_4 ; however, the (110), (210), and (211) representing the maghemite phase are missing. In addition, if the maghemite phase was present in the NPs, the (210) and (211) Bragg reflections would have produced rings with slightly higher intensity than the (111) reflections of magnetite observed in Fig. 5c). This clearly shows that the completely oxidized NPs have a magnetite structure.

The results obtained show that annealing at higher temperature (700 °C) under low background O_2 pressure promotes the formation of single-phase magnetite nanoparticles. These findings are in stark contrast with reports of formation of NPs under higher O_2 pressure which are usually core-shell

Figure 3 Nanoparticle structure and size at 700 °C; **a**, **b** Bright field image showing increased size and faceting of the nanoparticles; **c** selected-area image relative to diffraction pattern **d** showing rings compatible with both Fe_3O_4 and $\gamma\text{-Fe}_2\text{O}_3$; **e**, **f** nanoparticle size analysis showing a 6.6 nm mean NP diameter (equivalent area circular diameter).



[19, 20, 22] and have a significant number of structural defects including ant domain phase boundaries that originate at the core–shell interface, and are responsible for complex magnetic behaviors of the nanoparticles. It is interesting also to note that according to the calculated structural phase stability for iron oxides with respect to temperature and O_2 pressure [37] for bulk systems, at both 600 °C and 700 °C and at O_2 pressure of 10^{-4} Pa, only the $\alpha\text{-Fe}_2\text{O}_3$

phase is energetically favorable, while the Fe_3O_4 phase is expected above 800 °C. These calculations do not correspond to NPs; our observations clearly show that, at the nanoscale, the transformation into Fe_3O_4 happens at lower temperature (700 °C), and it is preceded not from $\alpha\text{-Fe}_2\text{O}_3$ NPs, as discussed above. We note that the quasi-equilibrium conditions during in situ TEM annealing will also contribute to the

Figure 4 HRTEM images of oxidized nanoparticles at 700 °C showing nanoparticles oriented along different zone axes, with clear faceting and no disorder at their surfaces. (The insets are digital diffractograms).

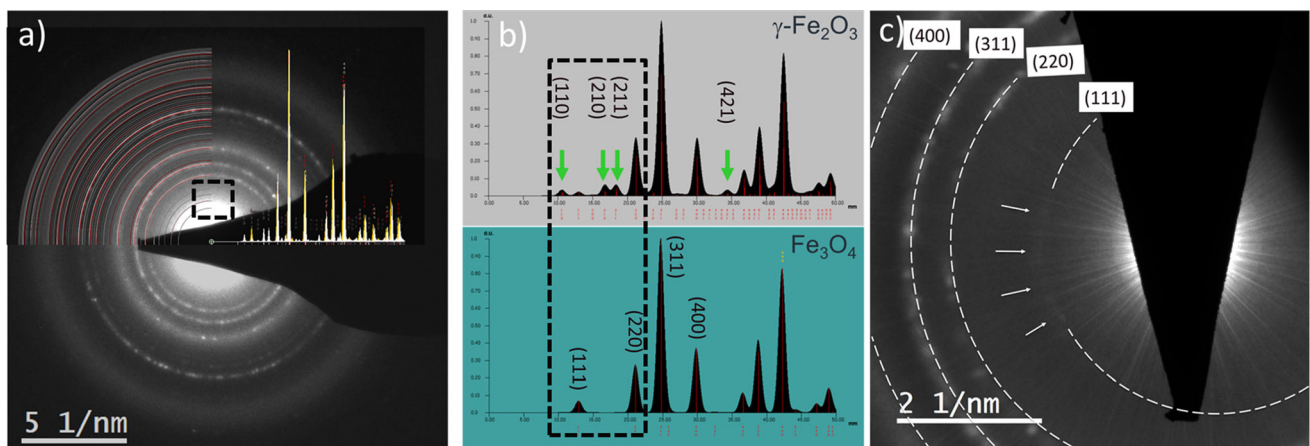
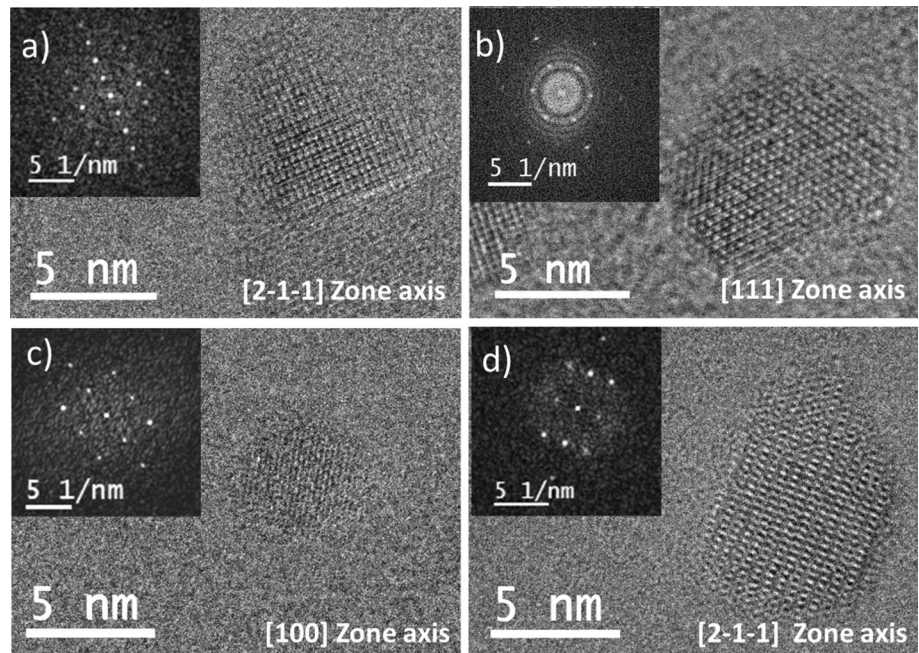


Figure 5 Analysis of diffraction pattern from nanoparticles at 700 °C. **a** DPs superimposed with Fe_3O_4 and $\gamma\text{-Fe}_2\text{O}_3$ calculated for powder of 20 nm NPs; the black dashed box highlights the area where the low index diffraction rings of Fe_3O_4 and $\gamma\text{-Fe}_2\text{O}_3$

should/can appear, as outlined also by black box in **b**. **c** Energy filtered DPs, with superimposed Fe_3O_4 calculated diffraction rings (dashed white lines).

discrepancy between the bulk and NP behaviors besides size effects.

Conclusions

In summary, in this work we studied the evolution of sputtered amorphous Fe thin films deposited on amorphous C as a function of temperature and low (background) O_2 pressure by using in situ transmission electron microscopy. We have shown that

amorphous Fe thin films dewet with increasing temperature and spherical shape nanoparticles are formed. Continuous annealing in the presence of residual oxygen O_2 under a microscope column vacuum of 4.0×10^{-4} Pa drives the crystallization and oxidation process. The structural phases of the nanoparticles at 600 °C are with mixed phases and structure. Electron diffraction measurements show that the majority of NPs have atomic lattice spacings corresponding to $\alpha\text{-Fe}$. Further annealing to 700 °C drastically changes both the morphology and

structure of the nanoparticles. Nanoparticle mean size increases from 5.2 to 6.6 nm, and from spherical they became faceted. Zero-loss energy filtered electron diffraction patterns revealed that all particles transform into magnetite nanoparticles, with no trace of other Fe oxide phases or core-shell structured nanoparticles. The atomic resolution TEM imaging confirms the single-crystal structure of the magnetite nanoparticles and their structural integrity. This work demonstrates that annealing in low pressure O₂ leads to creation of single-phase magnetite nanoparticles starting from amorphous Fe thin film and can provide a way to avoid core-shell or mixed Fe oxide phase nanoparticles.

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Compliance with ethical standards

Conflict of interest The authors declare that there is no conflict of interests regarding the publication of this article.

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