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One-Step Synthesis of Core(Cr)/Shell(\gamma\text{-Fe}_2\text{O}_3) Nanoparticles

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Interest in nanoparticles is stimulated from their technological importance, as they exhibit unique electrical, optical, and magnetic properties, which differ from their respective bulk.1–5 Magnetic nanoparticles with the size of 2–10 nm are particularly important because they are potentially useful in terabit magnetic storage, as carriers for biochemical complexes, as MRI contrast enhancing agents, etc.6 Core/shell nanoparticles, produced by various methods, can be applied for sensing, for high-density information storage, etc.7,8 Here we present the one-step synthesis of core/shell magnetic nanoparticles and their characterization.

The nanoparticles were prepared by adding 0.21 mL of Fe(CO)₅ and 39.98 mg pf of Cr(CO)₆ to 100 mL of degassed mesitylene (preheated to 70 °C). The molar ratio surfactant:carbonyls:solvent was 1:100:40000, and 9:1 for Fe(CO)₅:Cr(CO)₆. After stirring for 5 min, 226 mg of the polymer surfactant, Pluronic F127, was added. The temperature was then raised to reflux (~165 °C) for 24 h under reducing environment (5% H₂; 95% Ar). After cooling to room temperature, the resulting precipitate was collected by centrifugation, washed with toluene, and dried under vacuum.

The TEM image (Figure 1) shows the particles have spherical shape. The insert histogram shows particles size distribution of 13 ± 4 nm. The narrow size distribution is in good agreement with the magnetic measurements (Supporting Information). A small group of particles with the diameter 4–6 nm might be the result of decomposition at lower temperature.† An interesting feature in the image (see arrows) is a small dot inside of each particle core/shell structure. This suggests that the particles are not homogeneous and the material of the core differs from that of the shell.

The synchrotron powder XRD pattern (Supporting Information) of nanoparticles was fitted to more than one index, which supports finding heterogeneous particles from the TEM image. Apparently, the majority of the sample matches the γ-Fe₂O₃ crystal structure. However, since both Fe and Cr have the same crystal structure with different lattice constants, it is hard to determine what is the metal pattern at this point.

Figure 1. TEM image of the core/shell nanoparticles. (Inset) Histogram prepared by measuring ca. 200 nanoparticles.

The Mössbauer spectra (Supporting Information) are in good agreement with the results from SQUID (Supporting Information). They are characterized by a doublet at room temperature and by a sextet at 4.2 K. The change is due to the fact that the magnetic moment in the particles is random at room temperature, and they therefore behave like paramagnetic while there is an internal field at the blocked state at low temperatures, which generates a Zeeman splitting of the levels. At 295 K one observes two components: Fe³⁺ (96.57%) exhibiting superparamagnetic behavior and a small impurity of Fe²⁺ (3.43%). At 77 K we also observe Fe¹⁺ and Fe²⁺, but in lower amounts. Since almost all of the Fe atoms are trivalent, the crystal structure can be confirmed as γ-Fe₂O₃. Notice that no Fe⁰ could be detected, which indicates that the metal component from the XRD pattern is Cr. To ascertain the exact structure of the core/shell nanoparticles, we have carried out HR-TEM, EELS, and iron mapping.

Nanoparticles of approximately 12 nm in diameter were imaged by the HR-TEM, showing a high contrast core of about 4.5 nm in diameter (Figure 2a). The electron energy loss spectrum (EELS) (Figure 2d) clearly shows three edges: the O K, Cr L₂, and Fe L₂. Quantification of the EELS, from such particles, revealed an Fe:Cr ratio of about 9:1, which is in good agreement with the stoichiometry of the reaction. However, there is more than one source of oxygen, polymer, and oxide layers, so that the amount of oxygen varied at different spots of the sample. This result suggests that the sample is not homogeneous which is in good

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agreement with the TEM and XRD results. The EELS analysis of a single core/shell particle was performed in nanoprobe mode. One measurement on the entire particle resulted in an O:Cr:Fe ratio of 10:1.1:9.3, while a second measurement performed with the electron probe on the shell resulted in an O:Cr:Fe ratio of 10:0.3:6.3. This suggests that the shell in this particle is Fe₂O₃, which is in good agreement with the XRD pattern and Mössbauer spectra, so that the core should be Cr.

Lattice fringe could be observed both in the core and in the shell of the particles. The shell seems to consist of randomly oriented crystalline grains. Fast Fourier transform performed on such images gave several characteristic spots corresponding approximately to the d spacing: 2.97, 2.56, 2.10, and 1.67 Å. However, not all four spots appear in every particle analyzed. For example, in Figure 2a, the d spacings measured in the two designated areas are 2.54 and 2.08 Å that can be assigned to Fe₂O₃ or Fe₃Cr₂O₄. Mössbauer spectra (Supporting Information) also suggest the existence of Fe₂O₃. The possibility of Fe₃Cr₂O₄ is excluded since these lattice fringes were observed in the shell where Cr does not reside.

Energy-filtered imaging was performed by using a 40 eV energy window (708—748 eV) around the iron edge, to form an iron map. The regions containing Fe are bright, while other regions are dark in such an image. Thus, in the Fe mapping performed on the core/shell nanoparticles, shown in Figure 2, b and c, the donut-shaped bright structures indicate that the Fe constitutes the shell, which supports the finding from EELS analysis, while the dark region (particle core) in the core is probably the Cr.

On the basis of the result of EELS, Mössbauer spectra, and HR-TEM, the materials balance calculation (Supporting Information) was performed to estimate the density of particle core. The result shows there is no oxide in the particles core. Since Mössbauer spectra show no traceable amount of Fe⁰, we can conclude the particle core is Cr.

The decomposition temperature of Fe(CO)₅ (~60 °C) is much lower than that of Cr₂O₃ (~130 °C), thus, the decomposition reaction rate of Fe(CO)₅ should be faster than that of Cr₂O₃ at ~165 °C. Therefore, Fe should be formed before Cr, and the resulting particles should be Fe core and Cr shell. However, on the basis of the above results, they are confirmed to be Cr core and γ-Fe₂O₃ shell. To fully understand this result, the reaction mechanism (Supporting Information) is crucial. These nanoparticles were prepared in a one-step reaction from the carbonyl precursors. Due to the much lower decomposition temperature, Fe(CO)₅ was decomposed faster than Cr₂O₃ and forms a small amount of Fe clusters, which were then used as the catalysts to accelerate the decomposition of Cr₂O₃. Combine this with the consideration that the amount of Cr₂O₃ is only 10%, and thus, the formation of a Cr core presumably results from accelerated decomposition of Cr₂O₃ catalyzed by the Fe clusters. The Cr core is then used as the nucleation seed to form an Fe shell layer. Since particles of this size cannot stay stable as metal, the Fe shell oxidized to γ-Fe₂O₃, which protects the core from oxidation; in addition Cr has strong oxidation resistance. Therefore, the particle core stayed as Cr.

In conclusion, by mixing Fe(CO)₅ and Cr₂O₃ in the 9:1 ratio, we have successfully synthesized core/shell nanoparticles 13.5 nm in diameter, with uniform spherical shape and narrow distribution. The core/shell structure is formed by Fe clusters catalyzing the decomposition of Cr₂O₃. TEM image reveals the heterogeneous nature (core/shell structure), which is in good agreement with the synchrotron powder XRD pattern. The pattern shows more than one crystal structure in the materials. One was identified as γ-Fe₂O₃ and the other as a metal crystal structure. Mössbauer spectra, which support the superparamagnetic behavior determined by H—M measurement, do not show any traceable amount of Fe⁰. This suggests that the metal component might be Cr. EELS analysis and iron mapping suggest controlled stoichiometry and confirm a core made of Cr and a shell made of γ-Fe₂O₃.

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Note Added after ASAP Publication: In the version published on the Internet April 5, 2005, the current address for one of the authors was incorrect. In the final version, published April 6, 2005, and in the print version this is correct.

Supporting Information Available: Details of characterization techniques, magnetic measurements, and materials balance calculation. This material is available free of charge via the Internet at http://pubs.acs.org.

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