

Spinel Ferrite Core-Shell nanostructures by a versatile solvothermal seed-mediated growth approach and study of their nanointerfaces.

*Marco Sanna Angotzi,^{ab} Anna Musinu,^{ab} Valentina Mameli,^{ab} Andrea Ardu,^{abc} Claudio Cara,^{abc} Daniel Niznansky,^d Huolin L. Xin,^{*e} Carla Cannas^{*abc}*

^aDepartment of Chemical and Geological Sciences, Università di Cagliari, S.S. 554 bivio per Sestu, 09042, Monserrato (CA), Italy

^bINSTM, Cagliari Unit

^cConsorzio AUSI, Palazzo Bellavista Monteponi, 09016 Iglesias, CI, Italy

^dDepartment of Inorganic Chemistry, Charles University of Prague, Czech Republic

^eCenter for Functional Nanomaterials, Brookhaven National Laboratory, Upton, New York, United States

ABSTRACT

An easy, low-cost, repeatable seed-mediated growth approach in solvothermal condition has been proposed to synthesize bimagnetic spinel ferrite core-shell heterostructures in the 10-20 nm particle size range. Cobalt ferrite and manganese ferrite nanoparticles (CoFe_2O_4 and MnFe_2O_4) have been coated with isostructural spinel ferrites like maghemite/magnetite, MnFe_2O_4 and CoFe_2O_4 with similar cell parameters to create different heterostructures. The conventional study of the structure, morphology and composition has been combined with advanced techniques in

order to achieve details on the interface at the nanoscale level. Clear evidences of the heterostructure formation have been obtained (i) indirectly by comparing the ^{57}Fe Mössbauer spectra of the core-shell samples and an *ad-hoc* mechanical mixture and (ii) directly by mapping the nanoparticles' chemical composition by electron energy loss spectroscopy (EELS) and energy-dispersive X-ray spectroscopy (EDX) in the scanning transmission electron microscopy mode (STEM). In addition, chemical-sensitive electron tomography in STEM-EDX mode has been applied in order to get detailed 3D images with an unprecedented sub-nanometer spatial resolution.

KEYWORDS

core-shell, ferrite, EELS, EDX, Mössbauer, solvothermal, tomography

TEXT

In recent years, magnetic nanoparticles with a core-shell heterostructure have been studied increasingly to combine the physical and chemical properties of different components to obtain multifunctional materials with a wide range of applications or to improve their performances.¹⁻⁴ In this framework, bi-magnetic core-shell nanoparticles play a fundamental role because of the interactions arising from the contact of antiferromagnetic (AF), ferromagnetic (FM) and ferrimagnetic (FiM) phases. Since the discovery of the exchange bias phenomena by Meiklejohn and Bean,⁵ who prepared Co/CoO core-shell nanoparticles by surface treatment, FM(or FiM)/AFM and AFM/FM(or FiM) couplings have been extensively studied.⁶⁻⁹ On the contrary, although the combination of magnetically hard and soft FM and FiM phases is of great interest for different applications (magnetic recording, permanent magnets, microwave absorption, biomedicine), these systems have been less explored.¹ In this context, ideal crystalline phases to study this type of magnetic interaction are spinel ferrites ($\text{M}^{\text{II}}\text{Fe}_2\text{O}_4$, $\text{M}^{\text{II}}=\text{Fe}^{2+}$, Co^{2+} , Mn^{2+} , Ni^{2+} , etc.) which may

have hard or soft magnetic nature depending on the type of divalent ion^{1,10-12} and can be easily prepared by a great variety of methods.^{10,13-17} For instance, cobalt ferrite, with a high magnetocrystalline anisotropy, shows a hard magnetic behaviour, while the isostructural manganese ferrite and spinel iron oxides (magnetite and maghemite) are magnetically soft phases. The main advantage of the use of isostructural phases is the possibility of an epitaxial growth of the shell around pre-existing seeds, with the so-called seed-mediated growth method.¹⁸⁻²¹ This two-step synthetic strategy represents a versatile approach to achieve well-defined heterostructures with controllable interfaces by changing size, shape, composition, and structure of the core and the precursors of the shell. In the literature, spinel ferrite core-shell nanoparticles^{18,22-28} with a low dispersity and high crystallinity have been mainly synthesized by the powerful high-temperature decomposition of acetylacetonates in the presence of surfactants.¹³ The necessity of more and more eco-friendly strategies, that use less amount of toxic organic solvents, less expensive precursors and lower temperatures, has moved the interest towards alternative synthetic approaches. Inexpensive and easy methods as coprecipitation²⁹ and metal complex polymerization^{30,31} have been proposed in some specific cases but they do not allow a good control of the shell growth. Hydrothermal/solvothermal methods³² have in principle several advantages as the use of low-boiling and inexpensive solvents, the ease of the synthesis and the possibility to monitor pressure and temperature and a good repeatability³³⁻³⁵. To the best of our knowledge, only a few examples are present in the literature for the synthesis of spinel ferrite core-shell heterostructures by these methods ($\text{NiFe}_2\text{O}_4@\text{CoFe}_2\text{O}_4$ ³⁶ and $\text{CoFe}_2\text{O}_4@\text{NiFe}_2\text{O}_4$ ³⁷), but with results, in terms of products quality, far from those obtained in thermal decomposition conditions.

In this work, a seed-mediated growth approach in solvothermal conditions has been developed to synthesize core-shell nanoparticles made up of a hard magnetic phase (CoFe_2O_4) and a soft one

($\text{Fe}_3\text{O}_4/\gamma\text{-Fe}_2\text{O}_3$, MnFe_2O_4) with the same spinel structure, similar cell parameter and phase-contrast imaging. Clear evidences of the effective production of the core-shell structure have been obtained combining conventional techniques such as powder X-ray diffraction (XRD), transmission electron microscopy (TEM), induced coupled-plasma (ICP) and advanced ones (STEM-EELS, STEM-EDX, and STEM-EDX tomography) to map the chemical composition at the nanoscale. An indirect approach based on ^{57}Fe Mössbauer spectroscopy has been also used, allowing to demonstrate the potentiality of this technique for the study of coupled bimagnetic phases in heterostructures.

RESULTS AND DISCUSSION

Two samples of CoFe_2O_4 nanoparticles of different sizes, labelled as Co1 and Co2, and one sample of MnFe_2O_4 , labelled as Mn1, were prepared by solvothermal hydrolysis of mixed cobalt-iron or manganese-iron oleates in a mixture of organic solvents with different polarities and water contents. The Co1, Co2 and Mn1 nanoparticles were used as seeds to produce core-shell nanostructures by means of a second solvothermal treatment (seed-mediated growth). In particular, for both Co1 and Co2, two core-shell samples were prepared with a shell of spinel iron oxide (maghemite/magnetite) and manganese ferrite, indicated as $\text{Co}_x@\text{Fe}$ and $\text{Co}_x@\text{Mn}$ (where $x=1, 2$), respectively. Moreover, two core-shell samples with a shell of cobalt ferrite and spinel iron oxide were prepared for sample Mn1, labelled as $\text{Mn1}@\text{Co}$ and $\text{Mn1}@\text{Fe}$, respectively. The details about the synthesis are reported in the METHODS section.

Composition, structure and morphology. XRD patterns of the cores as well as the core-shell systems (**Figure 1**) show the typical reflections of a spinel phase. The cell parameters, a , for the samples Co1, Co2 and Mn1 are in good agreement with the value for the CoFe_2O_4 and MnFe_2O_4 reported in the literature (8.3919 Å, PDF Card: 022-1086 and 8.4990 Å, PDF Card: 010-0319,

respectively). From the profile analysis, the average crystallite sizes have been estimated and are reported in **Table 1**.

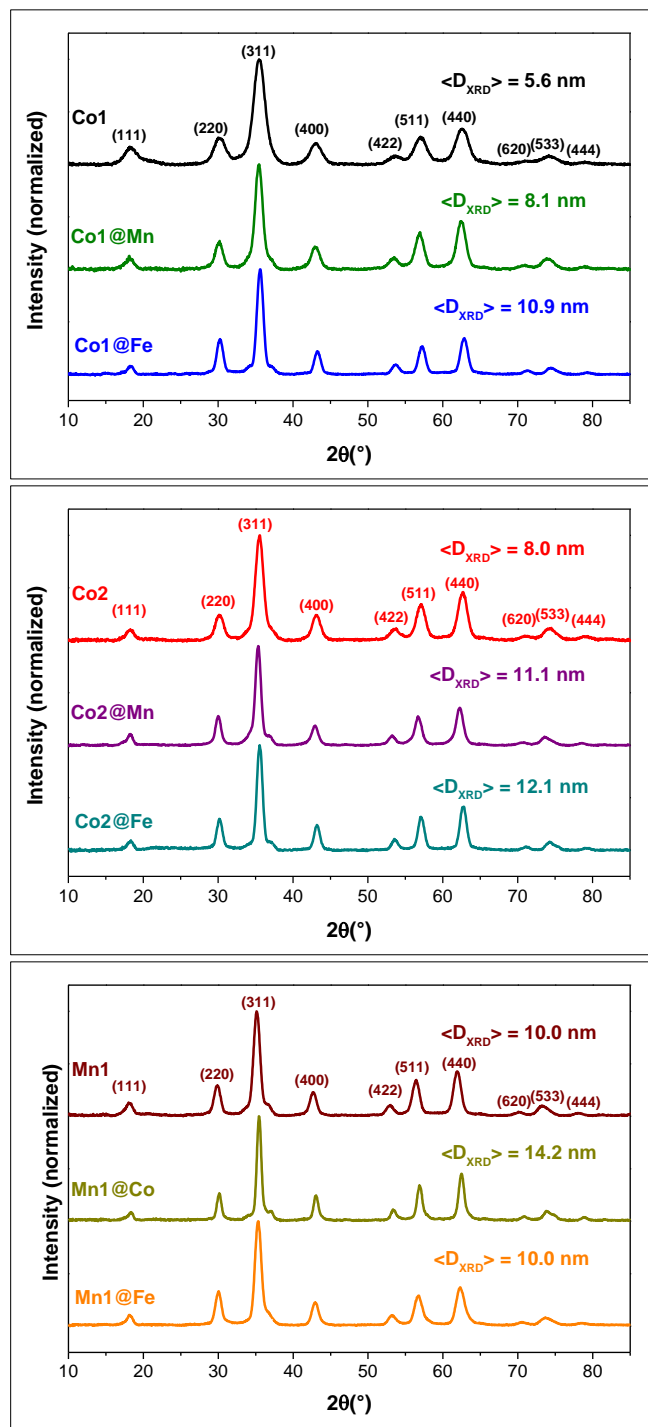


Figure 1. XRD Pattern of the samples.

Sample	a (Å)	<D _{XRD} > (nm)	<D _{TEM} > (nm)	Inorganic phase (%)
Co1	8.39±0.01	5.6±0.2	5.4±0.9	73
Co1@Mn	8.40±0.02	8.1±0.2	9.4±1.0	84
Co1@Fe	8.36±0.01	10.9±0.4	10.5±1.2	88
Co2	8.38±0.01	8.0±0.2	8.5±1.2	80
Co2@Mn	8.43±0.01	11.1±0.7	14.4±1.6	92
Co2@Fe	8.38±0.01	12.2±0.8	12.1±1.6	90
Mn1	8.47±0.01	10.0±0.2	10.8±1.4	85
Mn1@Co	8.40±0.01	14.2±0.8	17.9±2.1	92
Mn1@Fe	8.42±0.01	10.0±0.8	11.1±1.4	91

Table 1. Cell parameter (a), Crystallite size (<D_{XRD}>), Particle size (<D_{TEM}>) and amount of Inorganic phase calculated from TGA of the samples.

A slight variation of the lattice parameter has been observed for some core-shell systems. An increase of the lattice parameter is observed in the case of Co1@Mn and Co2@Mn, in agreement with the higher value of cell parameter for manganese ferrite than that of cobalt ferrite. Inversely, a decrease of the lattice parameter is detected when the shell is made of spinel iron oxide. The core-shell samples Mn1@Co and Mn1@Fe are subjected to a decrease of the lattice parameter due to the presence of cobalt ferrite or spinel iron oxide, respectively. Concerning the crystallite size obtained from the experimental profile, all the core-shell samples are characterised by higher values than the cores (**Table 1**), suggesting that a growth process took place, except for the sample Mn1@Fe, where the crystallite size is the same as the core. This can be explained by the formation of an amorphous shell of spinel iron oxide around the pre-formed core or partial dissolution of the core surface.

The TEM Bright Field images of the cores (**Figure 2**) show well-separated spheroidal nanoparticles with particle size similar to crystallite size, suggesting a high crystallinity of the particles (**Table 1**). Moreover, all the samples are characterised by narrow unimodal particles size distributions, with a dispersity in a 13-17 % range. TEM images of the core-shell samples, reported in **Figure 2**, show spherical particles with the presence of a unimodal size distribution and mean particle size higher than the core, suggesting that there has not been nucleation of new particles but the growing of the new phase around the pre-existing seeds, forming a core-shell heterostructure. This is also supported by the significant decrease of the dispersity with respect to the original core.

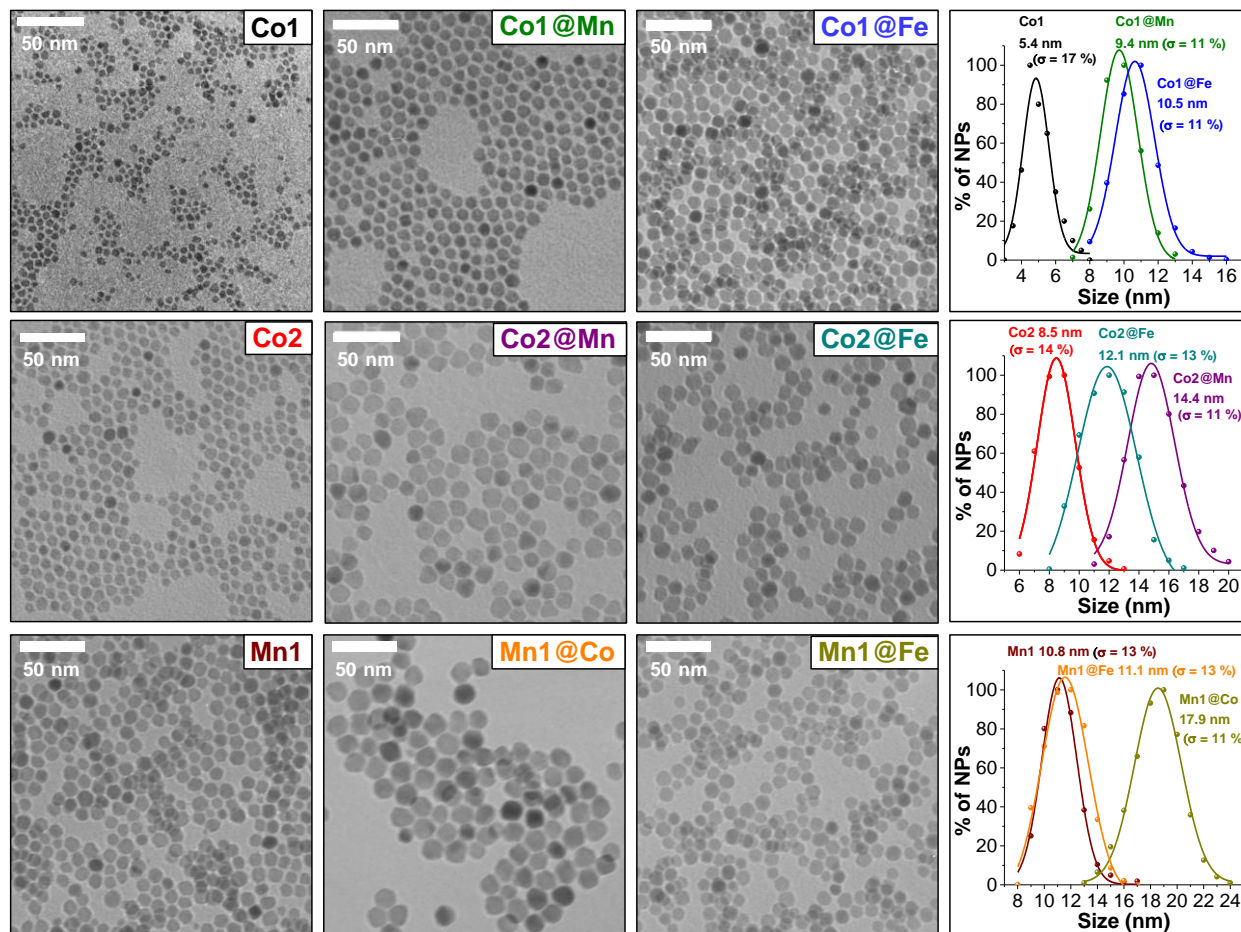


Figure 2. TEM Bright Field images and particle size distribution of the samples.

A careful study by means of high resolution microscopy (HRTEM and HRSTEM) has been conducted on all the samples, showing highly crystalline particles with no evidences of any amorphous part or heterojunctions. Here, as an example, the images of Co1@Mn and Co2@Fe samples are shown in **Figure 3**. The interlayer distances confirm the presence of the spinel oxide phase in agreement with the XRD data. The core-shell samples show no lattice mismatch, suggesting an epitaxial coating on the respective cores. Unfortunately, due to the very similar phase-contrast imaging, even high resolution microscopy cannot distinguish the core from the shell.

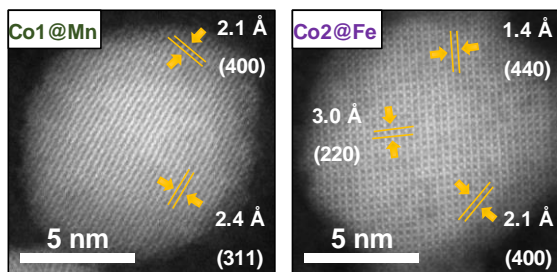


Figure 3. HRSTEM images of the Co1@Mn and Co2@Fe samples.

ICP analyses have been carried out on both core and core-shell systems. The $M^{II}:Fe^{III}$ ($M^{II} = Co^{2+}$ or Mn^{2+}) molar ratio for Co1, Co2 and Mn1 samples have been found equal to 0.49, 0.55 and 0.50, respectively, suggesting the formation of almost stoichiometric spinel ferrites. Taking into account the $M^{II}:Fe^{III}$ molar ratio found in the cores, it has been possible to roughly estimate the $M^{II}:Fe^{III}$ molar ratio in the shell, in the case of cobalt or manganese ferrite coating. Indeed, Co1@Mn, Co2@Mn and Mn1@Co samples present a $M^{II}:Fe^{III}$ molar ratio in the shell equal to 0.48, 0.45 and 0.64, which indicates the formation of an almost stoichiometric spinel ferrite shell.

The capping by oleate molecules has been demonstrated by FTIR (**Figure 1S**) and TGA (**Figure 2S**). FTIR spectra show the main vibrational modes associated with the oleate molecules linked at

the surface of the nanoparticles, as the COO⁻ vibrational modes (ν_{as} (COO⁻), ν_s (COO⁻)) and those ones related to the hydrocarbon chain.^{15,12} The complete assignment for all the samples is reported in **Table 4S**.

TGA curves, recorded under an oxygen atmosphere are shown in **Figure 2S**. The weight losses percentages obtained for the samples are reported in **Table 1**. These weight percentages correspond to a monolayer of oleate molecules surrounding the nanoparticle surface, as reported in the supporting information **Table 5S**. These values are also really close to those reported by other authors for oleic acid-coated spinel iron oxides nanoparticles and suggest the presence of a close-packed monolayer of the capping agent.³⁸⁻⁴⁰

Evidence of heterostructure by RT Mössbauer spectroscopy. As it has been said so far, the understanding of the core-shell heterostructure is not trivial, due to the same spinel crystallographic structure, similar cell parameter and phase-contrast imaging of the two counterparts. In this work, room temperature ⁵⁷Fe Mössbauer spectroscopy is proposed as an alternative tool to study bimagnetic core-shell heterostructures. In detail, room temperature (RT) ⁵⁷Fe Mössbauer measurements have been carried out for all the samples (**Figure 4**) and the hyperfine parameters are shown in **Table 2**.

The sample Co1 ($\langle D_{XRD} \rangle = 5.2$ nm) shows a broad singlet associated with particles having a relaxation time near the Mössbauer measurement time window (τ_M) accompanied with a sharper one related to the particles in the superparamagnetic state. The spectrum of the sample Co2 ($\langle D_{XRD} \rangle = 8.0$ nm) is fitted by one sextet deriving from the overlapping at room temperature accounting for Fe^{III} in the octahedral and tetrahedral sites of cobalt ferrite. The sample Mn1 ($\langle D_{XRD} \rangle = 10.0$ nm) shows a sharp singlet, which indicates the presence of nanoparticles in superparamagnetic state.

Clear differences can be evidenced by the comparison of the cores with the correspondent core-shell systems: the appearance of two sextets in place of singlets, indicating the formation of nanoparticles in the blocked state (Co1@Mn, Co1@Fe, Mn1@Co) and a separation of the two sextets (Co2@Mn, Co2@Fe). The two sextets can be due to two different spinel phases (CoFe₂O₄ and MnFe₂O₄ or maghemite/magnetite) or to two different sublattices (octahedral and tetrahedral) in a spinel phase due to the formation of a unique coherent structural domain, as observed by HRSTEM (**Figure 3**). The sample Mn1@Fe shows a collapsed sextet, due to the presence of nanoparticles in the edge between the superparamagnetic and the blocked state.

The isomer shift values for all the samples but Co2@Fe are in the range of Fe^{III} (0.28-0.39 mm/s), whereas Co2@Fe presents a higher isomer shift for one sextet (IS=0.50 mm/s), due to the presence of Fe^{II}, suggesting its not complete oxidation to Fe^{III}. The different behaviour with respect to the Co1@Fe, in which only Fe^{III} is present, is probably due to the reduced nanoparticle size that leads to a complete oxidation of magnetite to maghemite.

The hyperfine field values can be affected by the nature of the coating (manganese ferrite or spinel iron oxide) and the thickness of the shell (and consequently the final crystallite size). Specifically, Co1@Fe (<D_{XRD}> = 10.9 nm) shows hyperfine fields for both sextets higher than Co1@Mn (<D_{XRD}> = 8.1 nm), as manganese ferrite and maghemite are expected to have anisotropy constants of the same magnitude (for bulk materials: 3·10³ J/m³ and 5·10³ J/m³, respectively), the bigger crystallite size of the particles of Co1@Fe (10.9 nm) seems to be primarily responsible for the higher hyperfine field values. On the contrary, Co2@Mn (<D_{XRD}> = 11.1 nm) and Co2@Fe (<D_{XRD}>=12.2 nm) have similar crystallite sizes and the increase in the hyperfine field values for the sample Co2@Fe can be due principally to the higher anisotropy constant of magnetite, that for bulk material is 1.3·10⁴ J/m³.⁴¹ The sample Mn1@Co (<D_{XRD}> = 14.2 nm)

shows the highest hyperfine field values for both octahedral and tetrahedral sites, due to the big crystallite size and the presence of cobalt ferrite in the shell, which forms the 78% of the total nanoparticle volume. The sample Mn1@Fe shows lower hyperfine field values for both sites in comparison with Mn1@Co, being manganese ferrite and maghemite/magnetite two magnetically soft phases with respect to cobalt ferrite (which magnetocrystalline anisotropy in the bulk is $2.9 \cdot 10^5 \text{ J/m}^3$). It is known that the O_h - T_d exchange interactions (J_{AB}) are stronger than T_d - T_d ones (J_{AA}) or O_h - O_h ones (J_{BB}). Therefore, the hyperfine field of the tetrahedral sublattice will be more affected by the cations located in the octahedral sublattice.⁴² In particular, for the Co2@Fe sample, the highest difference in the hyperfine field, with respect to the Co2@Mn sample, has been detected for T_d sites, probably due to the presence of Fe^{II} only in the O_h sites.

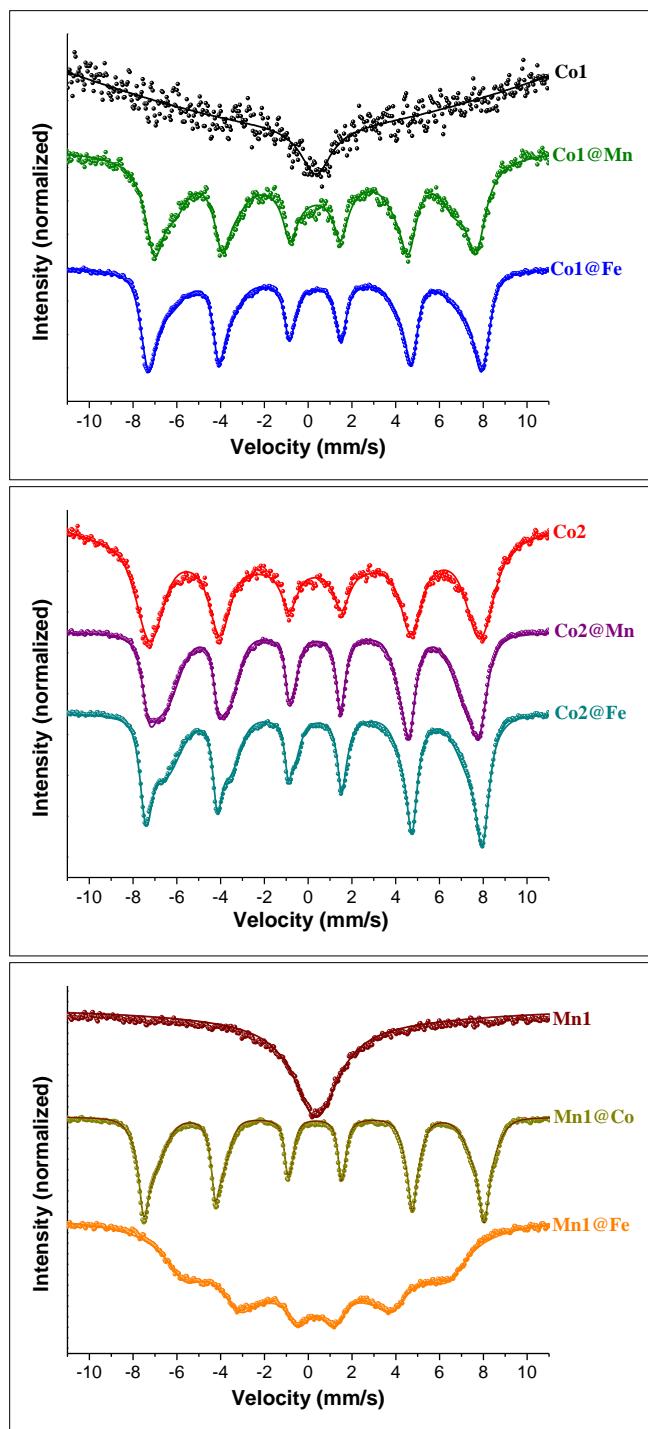


Figure 4. Room Temperature ^{57}Fe Mössbauer spectra of the samples.

	Type	Size (nm)	Signal	ISO (mm/s)	QUA (mm/s)	BHF (T)	FWHM (mm/s)
Co1	CoFe ₂ O ₄	5.6±0.2	Singlet	0.2(1)	-	-	12(1)
			Singlet	0.44(6)	-	-	1.1 (2)
Co1@Mn	CoFe ₂ O ₄ @MnFe ₂ O ₄	8.1±0.2	Sextet	0.34(2)	- 0.10(1)	39.7(2)	1.56(5)
			Sextet	0.32(1)	0.00(1)	45.8(1)	0.67(2)
Co1@Fe	CoFe ₂ O ₄ @ γ -Fe ₂ O ₃ /Fe ₃ O ₄	10.9±0.4	Sextet	0.35(1)	- 0.02(1)	42.4(2)	0.64(1)
			Sextet	0.32(1)	- 0.01(1)	47.3(1)	0.47(1)
Co2	CoFe ₂ O ₄	8.0±0.2	Sextet	0.32(1)	- 0.02(1)	46.9(1)	0.87(2)
Co2@Mn	CoFe ₂ O ₄ @MnFe ₂ O ₄	11.1±0.7	Sextet	0.39(1)	- 0.02(1)	43.4(1)	0.55(1)
			Sextet	0.28(1)	0.01(1)	46.9(1)	0.28(1)
Co2@Fe	CoFe ₂ O ₄ @ γ -Fe ₂ O ₃ /Fe ₃ O ₄	12.2±0.8	Sextet	0.50(1)	0.00(1)	43.9(1)	0.64(1)
			Sextet	0.29(1)	- 0.02(1)	47.8(1)	0.34(1)
Mn1	MnFe ₂ O ₄	10.0±0.2	Singlet	0.37(1)	-	-	2.8(1)
Mn1@Co	MnFe ₂ O ₄ @CoFe ₂ O ₄	14.2±0.8	Sextet	0.37(2)	- 0.01(2)	47.1(1)	0.41(2)
			Sextet	0.26(1)	0.01(1)	48.3(1)	0.29(1)
Mn1@Fe	MnFe ₂ O ₄ @ γ -Fe ₂ O ₃ /Fe ₃ O ₄	10.0±0.8	Sextet	0.39(1)	0.03(1)	24.8(1)	1.00(1)
			Sextet	0.35(2)	0.01(2)	37.5(2)	1.04(2)

Table 2. Room Temperature ⁵⁷Fe Mössbauer parameters of the samples.

In order to verify if a core-shell structure was obtained, a reference sample (CoMn_R) made from a physical mixture of cobalt ferrite (Co_R) and manganese ferrite (Mn_R) nanoparticles in a

mass ratio 1:1 was prepared and analysed. The nanoparticles were synthesised by the same procedure reported in the METHODS section, resulting in a crystallite size of about 8.7 nm for both Co_R and Mn_R samples, similar to the Co1@Mn (8.1 nm) sample. In the physical mixture, the oleate molecules that surround the particles should hinder the direct contact of the two magnetic phases. **Figure 5** shows RT Mössbauer spectra of two different cores (Co_R, Mn_R), the physical mixture (CoMn_R) and the core-shell nanoparticles (Co1@Mn). **Table 3** shows Mössbauer parameters for all these samples. The spectrum of the mixture (CoMn_R) is the sum of the two subspectra (one singlet and one sextet) having the same parameters as the pure phases. On the contrary, the spectrum of core-shell nanoparticles shows two sextets with values of hyperfine fields equal to 39.7 T and 45.8 T, which differ from the values found for cobalt ferrite or the physical mixture (47.3 T). This comparison permits to exclude the independent formation of two phases (CoFe₂O₄ and MnFe₂O₄) with similar size and can be considered as an indirect proof that the two magnetic phases in the sample Co1@Mn are in a close contact. This is in agreement with TEM data that evidence the formation of a monomodal distribution of the particles and HRSTEM that evidence a single crystal structure (**Figure 3**).

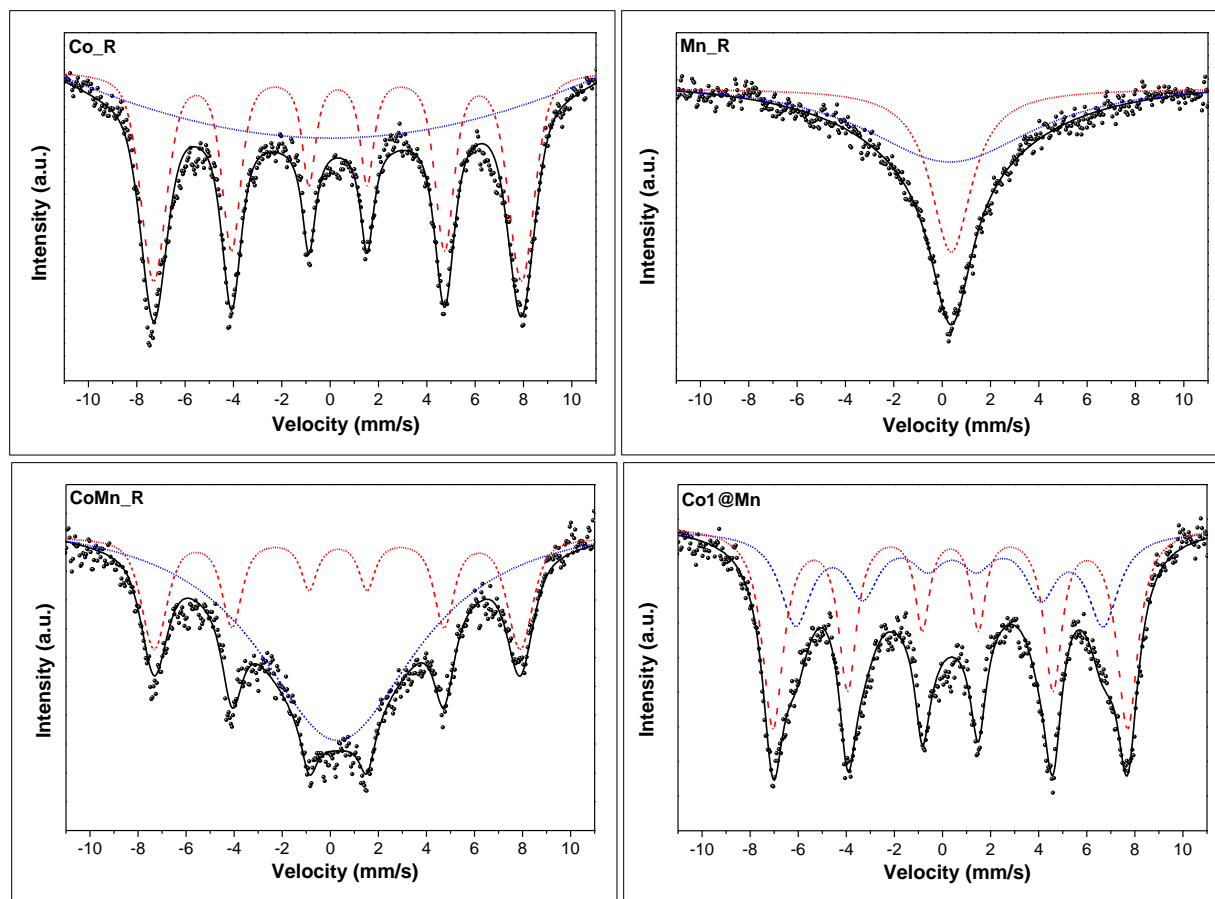


Figure 5. Room Temperature ^{57}Fe Mössbauer spectra of Co_R, Mn_R, CoMn_R and Co1@Mn.

Sample	Type	Size (nm)	Signal	ISO (mm/s)	QUA (mm/s)	BHF (T)	FWHM (mm/s)
Co_R	CoFe_2O_4	8.7 ± 0.3	Sextet	0.32(1)	-0.02(1)	47.3(1)	0.65(2)
Mn_R	MnFe_2O_4	8.7 ± 0.5	Singlet	0.32(6)	-	-	8(1)
			Singlet	0.38(2)	-	-	2.1(1)
CoMn_R	$\text{CoFe}_2\text{O}_4 + \text{MnFe}_2\text{O}_4$	8.0 ± 0.5	Singlet	0.31(3)	-	-	8.1(1)
			Sextet	0.30(1)	-0.04(2)	47.3(1)	0.80(2)
Co1@Mn	$\text{CoFe}_2\text{O}_4 @ \text{MnFe}_2\text{O}_4$	8.1 ± 0.2	Sextet	0.34(2)	-0.10(1)	39.7(2)	1.56(5)
			Sextet	0.32(1)	0.00(1)	45.8(1)	0.67(2)

Table 3. Room Temperature ^{57}Fe Mössbauer parameters of Co_R, Mn_R, CoMn_R and Co1@Mn.

Chemical mapping at the nanoscale. It is worth noting that in order to have a direct and unambiguous prove of the formation of core-shell heterostructures, chemical mapping at the nanoscale is mandatory.⁴³⁻⁴⁵ Consequently, STEM-EDX, STEM-EDX tomography, STEM-EELS (for the Cox@Me samples) mapping of Fe, Co and Mn elements have been carried out. In the literature, only EELS^{18,25,24,28,36} is generally adopted to make chemical mapping at the nanoscale and there are no examples of the combined use of the two techniques. Thanks to the recent substantial technological progresses of energy-dispersive X-ray spectroscopy, in particular the development of ultrasensitive detectors (Super-XTM detector with multiple silicon drift detectors),⁴⁶⁻⁴⁸ high-quality EDX maps⁴⁹ and a detailed comparison between EDX and EELS is here presented in order to strength our findings about the creation of a core-shell heterostructure.

Figure 6 shows STEM-EDX and STEM-EELS chemical mapping, the corresponding line profiles of the samples and, as an example, an EDX and EELS spectrum for the Co2@Mn sample. All the cores (Co1, Co2 and Mn1) show homogenous distribution of Co (or Mn) and Fe throughout the particles, with a $M^{II}:Fe^{III}$ ($M^{II} = Co^{2+}$ or Mn^{2+}) ratio, calculated from EDX data, equal to 0.56, 0.59 and 0.48, respectively. These values are in good agreement with the ICP data, suggesting the formation of almost stoichiometric ferrites and highlighting the reliability of the quantitative elemental mapping by STEM-EDX at the nanoscale.⁵⁰ STEM-EDX chemical mapping of the core-shell samples unambiguously indicates the formation of a core-shell heterostructure. The single particle line profile calculated from EDX data reveals that the amount of Co (in case of Co1@Fe, Co1@Mn, Co2@Fe and Co2@Mn) or Mn (in case of Mn1@Fe and Mn1@Co) decreases gradually up to the surface. This behaviour suggests that there had been a slight dissolution of the core surface during the seed-mediated growth treatment. This is consistent with the literature data that the high temperature induces a partial dissolution of the core surface, leading to a mixed

chemical composition at the interface.^{18,25,28,29,51} To validate the EDX results, STEM-EELS analyses have been carried out on the cobalt ferrite-based core-shell samples (Co1@Fe, Co1@Mn, Co2@Fe and Co2@Mn). The chemical mapping and the single particle line profile show the same tendency and demonstrate how STEM-EDX can be successfully employed to study core-shell heterostructures with a sub-nanometer spatial resolution.⁴⁹

STEM-EDX tomography has been performed on the Co2@Fe sample and the three-dimensional reconstruction of the Co and Fe distributions are shown in **Figure 7** and **Movie 1**. In agreement with the above-findings, these results have permitted to visualize the core-shell heterostructure, pointing out the 3D details: spheroidal shape of the particle and homogenous coating of the shell around the well-centred core, *i.e.* with a uniform shell thickness. To the best of our knowledge, this is the first example of chemical tomography performed on this type of systems at this length scale, allowing to demonstrate the power of this technique.

On the basis of above findings, thanks to the direct (chemical mapping at the nanoscale) and indirect approach (multitechnique analyses by the comparison with a mechanical mixture), the achievement of core-shell heterostructures has been proven.

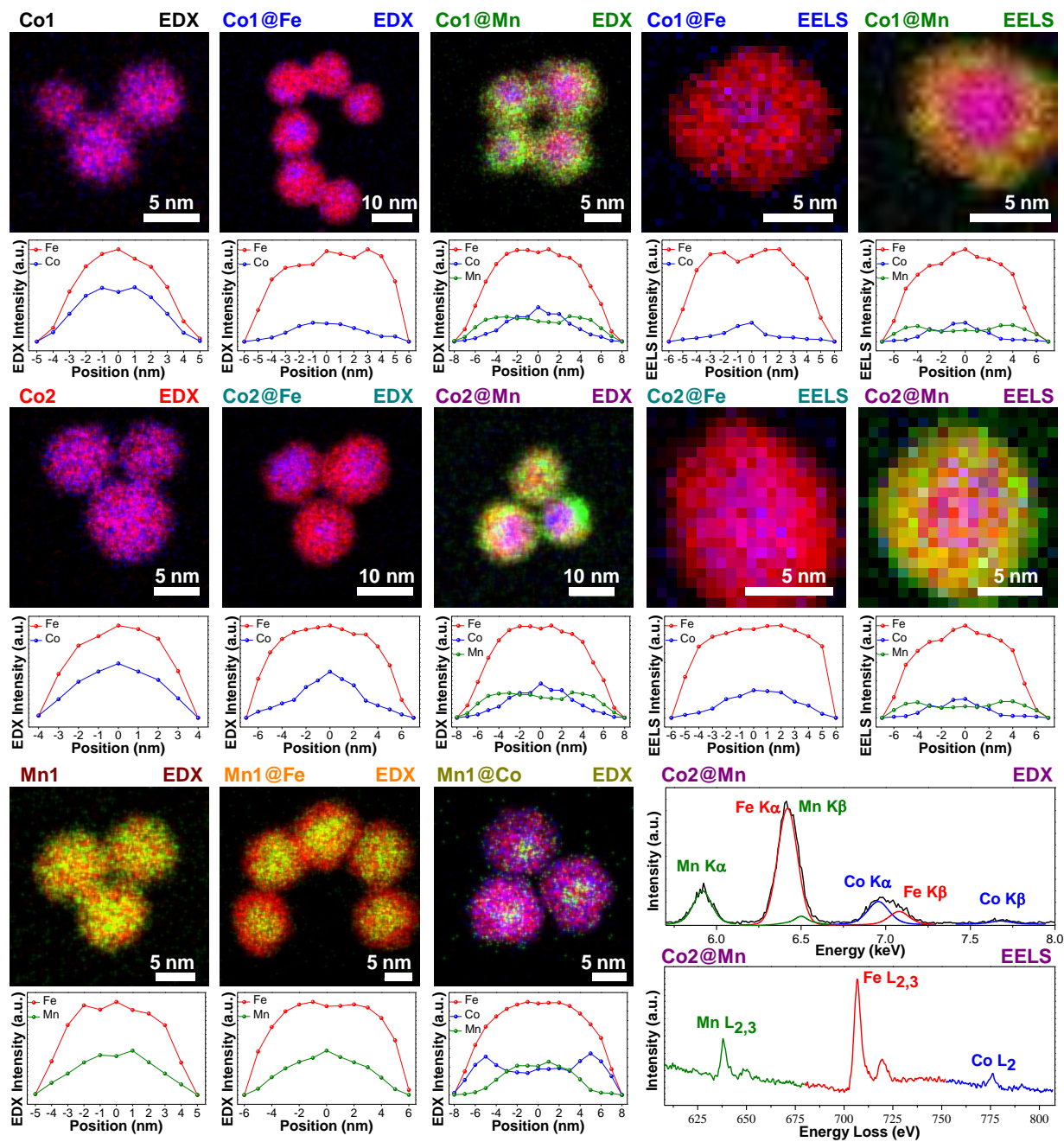


Figure 6. STEM-EDX and STEM-EELS mapping of the particles. EDX and EELS images are shown on top of the line profile across the samples. Cobalt is represented in blue, manganese in green, iron in red. Bottom right, two examples of EDX and EELS spectra collected for the Co₂@Mn sample.

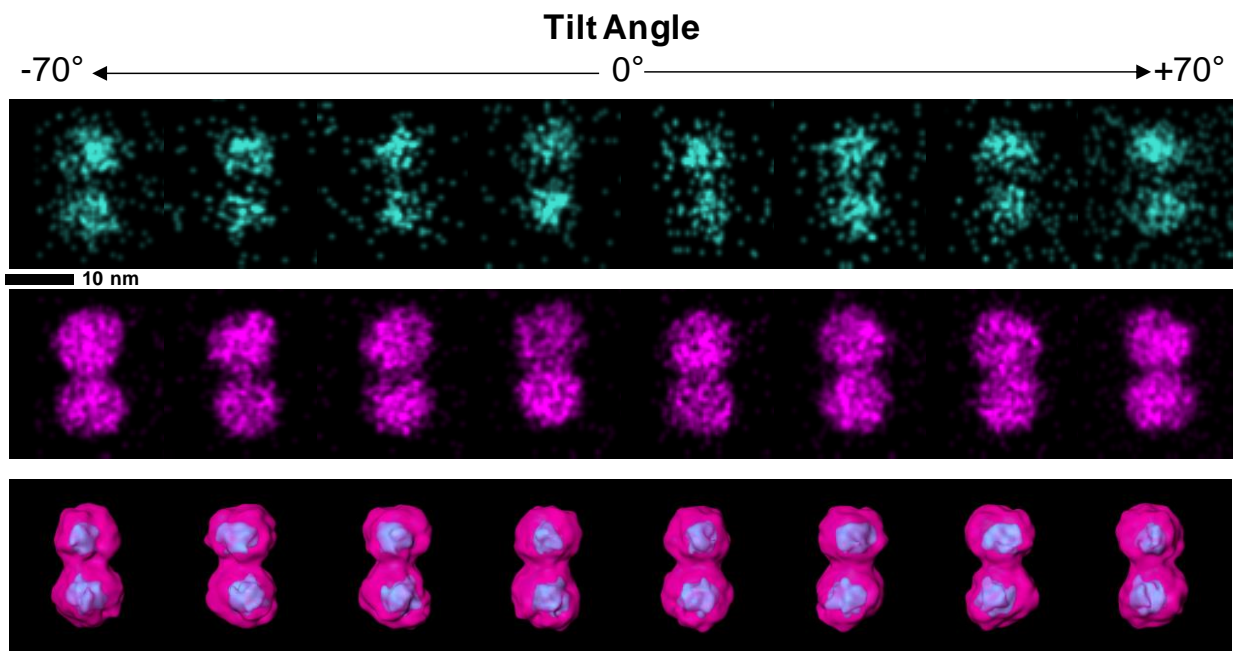


Figure 7. STEM-EDX tomography of the Co₂@Fe sample. Top two rows show the projected distribution of Co and Fe mapped by STEM-EDX and the third row shows the isosurface rendition of the 3D distributions of Co and Fe in two touching Co₂@Fe particles.

CONCLUSIONS

A simple, low-cost seed-mediated growth strategy in solvothermal condition is proposed to synthesize hydrophobic bimagnetic spinel ferrite core-shell nanoparticles. Conventional techniques (XRD, TEM, HRTEM/HRSTEM, TGA, FTIR, ICP) show for all the samples a unique crystal domain, spherical shape, low dispersity and a monolayer of oleate molecules as the nanoparticles' capping. The higher particle size and lower dispersity than the core are indirect clues of the homogeneous growing of the shell around the pre-formed seeds. A further indication supporting this idea is given by ⁵⁷Fe Mössbauer spectroscopy, suitably applied to coupled core-shell nanoparticles and not-coupled *ad-hoc* mechanical mixtures. Direct proof of the core-shell structure formation is successfully provided by chemical mapping at the nanoscale, by the combination of STEM-EELS, STEM-EDX and STEM-EDX tomography. In particular, STEM-EDX is revealed to be a powerful technique to obtain high-quality compositional maps and 3D

reconstructions of the core-shell heterostructure with sub-nanometer spatial resolution. All the above-findings suggest that the solvothermal seed-mediated growth can be proposed as a powerful and versatile approach to create other classes of heterostructures properly combining magnetic and not-magnetic materials with diverse chemical-physical properties and diverse architectures.

METHODS

Chemicals. Oleic acid (>99.99%), 1-pentanol (99.89%), hexane (84.67%) and toluene (99.26%) were purchased from Lach-Ner; 1-octanol (>99.99%) and $\text{Mn}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$ (>97.0%) from Sigma-Aldrich; absolute ethanol and $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ (99.0%) from Penta; NaOH (>98.0%) from Fluka; $\text{Fe}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$ (98.0%) from Lachema; $\text{FeCl}_2 \cdot 4\text{H}_2\text{O}$ (99%) from Merck.

Synthesis of $\text{M}^{\text{II}}\text{-Fe}^{\text{III}}$ oleate in 1-pentanol. The mixed $\text{M}^{\text{II}}\text{-Fe}^{\text{III}}$ oleates ($\text{M}^{\text{II}} = \text{Co}^{2+}, \text{Mn}^{2+}$) and the Fe^{II} -oleate were used as metal precursors for the synthesis of CoFe_2O_4 or MnFe_2O_4 and $\gamma\text{-Fe}_2\text{O}_3$, respectively. They were synthesized and collected following the procedure set up by A. Repko *et al.*³⁴ The molar ratios among the reactants are given in **Table 1S** for the different metal oleates. First, a pale-yellow sodium oleate solution was prepared in a 250 mL round-bottom flask dissolving the sodium hydroxide in 10 mL of distilled water and adding 20 mL of ethanol together with the oleic acid. Secondly, the iron(III) nitrate and $\text{Me}(\text{II})$ nitrate ($\text{M}^{\text{II}} = \text{Co}^{2+}, \text{Mn}^{2+}$) or only iron(II) chloride were dissolved in 10 mL of distilled water producing an orange solution. This was then added to the sodium oleate one and mechanically stirred in order to obtain the $\text{M}^{\text{II}}\text{-Fe}^{\text{III}}$ oleate or the Fe^{II} -oleate. After the addition, the formation of the metals oleate complexes made the solution black and viscous. The successive addition of 20 mL of hexane led to a liquid biphasic system, the upper one is the metals oleate containing organic phase while the lower is the water phase. This mixture was boiled under reflux for 60 min to complete the formation of metals oleate.

The as-described procedure was carried out under an inert atmosphere in the case of the Fe^{II} oleate synthesis. Then, the system was left to cool down to room temperature. The water phase was removed by Pasteur pipette, while 20 mL of water, 5 mL of ethanol and 5 mL of hexane were added to the organic phase and stirred in order to wash it from the inorganic residuals. The mixture was boiled under reflux for 30 min. This washing step was done twice. Finally, the water phase was completely removed and 15 mL of 1-pentanol were added to the flask. The mixture was heated for about 30 min to induce the complete evaporation of hexane. The obtained product as a viscous black liquid (M^{II}-Fe^{III} oleate or Fe^{II} oleate in pentanol) was moved into a 40 mL glass vial with Teflon cup with the help of 5 mL of 1-pentanol. The composition of the product, i.e. the final amount of pentanol and consequently the concentration of metals oleate was estimated from its weight, assuming quantitative yield from metals salts.

Core: solvothermal preparation of colloidal CoFe₂O₄ and MnFe₂O₄ nanoparticles. Two samples of CoFe₂O₄ nanoparticles of different sizes, labelled as Co1 and Co2, and one sample of MnFe₂O₄, labelled as Mn1, were prepared by solvothermal hydrolysis of mixed cobalt-iron or manganese-iron oleates in a mixture of organic solvents with different polarities and water contents. The metal oleate was prepared separately, in order to avoid the formation of by-products.³³ The formation mechanism of the nanoparticles has been already described by Repko et al.^{33,52,34} In solvothermal condition, water causes the hydrolysis of the oleate and the formation of nanoparticles takes place after nucleation and growth in the organic phase until they reach a critical diameter, which bring them to precipitate. At the end of the treatment, in fact, a black precipitate is found at the bottom of the teflon liner, while two liquid phases are present above, an aqueous and an organic one. The water phase is always colourless indicating, as expected, that no particles are present in it. On the contrary, the colour of the organic phase becomes darker with

decreasing solvent polarity due to the presence of hydrophobic oleate-capped nanoparticles with a size below the critical diameter necessary to precipitate. The appropriate amount of M^{II}-Fe oleates (M^{II}= Co²⁺ or Mn²⁺, in a 1:2 M^{II}:Fe^{III} ratio to produce M^{II}Fe₂O₄) in 1-pentanol was moved into a 50 mL teflon liner and a further amount of a mixture of other solvents was added to reach a total volume of 20 mL, as described in **Table 2S**. Then, 10 or 5 mL of water, depending on the reaction temperature, was added. The solvents and their relative ratios have been chosen according to the study by Repko *et al.*³⁴ in order to prepare particles of different sizes. The liner free space was flushed with nitrogen and was enclosed in a stainless-steel autoclave (Berghof DAB-2), briefly shaken and put vertically into a pre-heated (180° or 220 °C) oven. The reaction time was 10 hours. After the heat treatment, the autoclave was left to cool down to room temperature, and the as-prepared magnetic nanoparticles were magnetically separated from the supernatant that was discarded. A purification process was conducted twice as follows. First, the particles were dispersed in 10 mL of hexane (with the help of sonication), then 10 mL of ethanol were used in order to wash and precipitate the nanoparticles that were finally separated by a magnet. At the end of this step, the nanoparticles were dispersed in 5 mL of hexane and centrifuged at 3000 rpm for 5 min. In this case the supernatant, which is the desired product, was saved and the precipitate was discarded. The nanoparticles concentration was estimated by sampling an aliquot of the colloidal dispersion, drying it and weighing it. A summary of the synthesis conditions for each sample is reported in **Table 2S**. It is worthy of note that both the solvent polarity (by using 1-octanol or toluene together with 1-pentanol) and the reaction temperature (180 °C or 220 °C) play the main role in the nanoparticles growth and their final size.

Core-shell: solvothermal preparation by seed-mediated growth. The Co1, Co2 and Mn1 nanoparticles were used as seeds to produce core-shell nanostructures by means of a second

solvothermal treatment (seed-mediated growth). In particular, for both Co1 and Co2, two core-shell samples were prepared with a shell of spinel iron oxide and manganese ferrite, indicated as $\text{Co}_x@Fe$ and $\text{Co}_x@Mn$ (where $x=1, 2$), respectively. Moreover, two core-shell samples with a shell of cobalt ferrite and spinel iron oxide were prepared for sample Mn1, labelled as $\text{Mn1}@Co$ and $\text{Mn1}@Fe$ respectively. Different attempts have been carried out in order to achieve the best experimental conditions, in terms of ratio seeds/shell precursor, solvent, concentration of the precursors, allowing the production of bigger nanoparticles with a narrow size distribution. For example, we found out that 1-octanol is not a suitable solvent because it leads to a high dispersity of the product. This can be probably ascribed to the low colloidal stability of the seed nanoparticles dispersion. Being toluene a good solvent to disperse nanoparticles, it has been chosen instead of 1-octanol. We used a mixture of M^{II} -Fe oleates ($M^{II}=\text{Co}^{2+}$ or Mn^{2+} , in ratio $M^{II}:\text{Fe}^{III}=1:2$) in order to create a shell of cobalt or manganese ferrite, or Fe^{II} -oleate in order to create a shell of spinel iron oxide. Indeed, in these experimental conditions, the use of Fe^{III} or a mixture of Fe^{II} and Fe^{III} -oleates lead to hematite. The amount of hexane dispersion containing the appropriate quantity of particles (seeds) was added in a teflon liner. The particles were precipitated by adding ethanol in a 1:1 volume ratio with respect to the hexane dispersion, held by a magnet and the liquid was discharged. Then, the particles were dissolved in 10 mL of toluene (with the help of sonication) and 10 mL of a 1-pentanol solution of metals oleate was added. Finally, after the addition of 5 mL of distilled water, the liner was enclosed into the autoclave and treated at 220°C for 10 hours. After the heat treatment, the purification steps were the same as for the seeds. **Table 3S** summarises the synthesis conditions for the core-shell nanostructures.

Reference sample: the physical mixture. A reference sample made up of CoFe_2O_4 - MnFe_2O_4 physical mixture (weight ratio 1:1) was prepared. Both the phases were synthesised as described

in the previous paragraph about the cores and the synthesis conditions are shown in **Table 2S**. Specifically, a proper amount of a hexane dispersion of CoFe₂O₄ nanoparticles (Co_mix) was added to the MnFe₂O₄ one (Mn_mix), dried and characterized.

Characterization techniques. The chemical composition was studied by Inductively Coupled Plasma - Atomic Emission Spectrometry (ICP-AES). The dried samples were digested by using HNO₃. The digested sample solutions were stirred at room temperature for 1 h, then heated up to ~50 °C for 2 h. The solutions were left to cool down, filtered and diluted by using 1% v/v HNO₃ solution. The ICP measurements were made by means of a Liberty 200 ICP Varian spectrometer under the following conditions: Fe line: 259.940 nm, Co line: 238.892 nm, Mn line: 257.610 nm; Fe, Co and Mn concentration range: (0.1÷1.5) ppm; Fe detection range: (0.015÷750) ppm, Co detection range: (0.050÷2500) ppm, Mn detection range: (0.003÷150) ppm. The analyses have been repeated two times on different portions of the samples. The chemical formulas were calculated by assuming the absence of anions vacancies.

The samples were characterized by X-ray Diffraction (XRD), using a PANalytical X'Pert PRO with Cu K α radiation (1.5418Å), secondary monochromator and PIXcel position sensitive detector. Calibration of peak position and instrumental width was done using powder LaB₆ from NIST. The hexane dispersions were dried on a glass plate and measured in the angular range 10°-90° with step 0.039°. The analyses of the XRD patterns (identification of the crystalline phase, background subtraction) were done by the PANalytical X'Pert HighScore software. The most intense X-ray peaks ((220), (311), (400), (422), (511), (440)) were fitted with Origin Software by PseudoVoigt function, using a 1:1 gaussian:lorentzian ratio ($m_u = 0.5$):

$$y = y_0 + A \left[m_u \frac{2}{\pi} \frac{w}{4(x - x_c)^2 + w^2} + (1 - m_u) \frac{\sqrt{4 \ln 2}}{\sqrt{\pi} w} e^{-\frac{4 \ln 2}{w^2} (x - x_c)^2} \right]$$

In reciprocal space

$$q = \frac{2\sin\theta}{\lambda} = \frac{\sqrt{h^2 + k^2 + l^2}}{a}$$

The crystallite size (diameter) was obtained from full width at half maximum of q (denoted as Γ_q) by

$$d = \frac{1.1034}{\Gamma_q}$$

Room temperature ^{57}Fe Mössbauer spectroscopy was done on Wissel spectrometer using transmission arrangement and proportional detector LND-45431. A α -Fe foil was used as a standard, and fitting procedure was done by NORMOS program.

TEM images were obtained by using a JEOL 200CX operating at 200 kV. The particle size distribution was obtained by measuring in the automatic mode over 1000 particles by means of the software Pebbles and adopting a spherical shape.⁵³ The mean particle diameter was calculated as the average value and the dispersity as the percent ratio between the standard deviation and the average value.

HRTEM images were carried out using JEOL JEM 2010 UHR equipped with a Gatan imaging filter (GIF) and a 794-slow scan CCD camera.

EDX measurements were carried out in the STEM mode using a FEI Talos F200X with a field-emission gun operating at 200 kV equipped with a four-quadrant 0.9-sr energy dispersive X-ray spectrometer. 3D tomography was acquired in STEM-EDX mode with tilt series from -70° to 70° and 10° intervals.

EELS and HRSTEM measurements were carried out using an aberration-corrected scanning transmission electron microscope (Hitachi HD2700C) with a cold field emission gun operating at 200 kV equipped with a parallel EELS spectrometer (high resolution Enfina).

Transform Infrared Spectroscopy (FT-IR) spectra were recorded in the region from 350 to 4000 cm^{-1} by using a Bruker Vertex 70 spectrophotometer on the colloidal dispersions by means of a Platinum ATR Unit A 225, (standard ATR crystal material: diamond).

TGA curves were obtained on powders by using a Mettler-Toledo TGA/SDTA 851 in the 25-1000°C range, with a heating rate of 10°C/min under 50 mL/min O_2 flow.

AUTHOR INFORMATION

Corresponding Authors

*E-mail (CC) ccannas@unica.it

*E-mail (HLX) hxin@bnl.gov

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ASSOCIATED CONTENT

The Supporting Information is available free of charge on the ACS Publications website. Synthesis conditions for different metal oleates; synthesis condition for the pure spinel ferrite nanoparticles; synthesis condition for the core-shell nanoparticles; FT-IR spectra of the samples; assignment of FT-IR bands of the samples; TGA curves of the samples; computation of capping molecule packing on the nanoparticles surface.

REFERENCES

- (1) López-Ortega, A.; Estrader, M.; Salazar-Alvarez, G.; Roca, A. G.; Nogués, J. Applications of Exchange Coupled Bi-Magnetic Hard/soft and Soft/hard Magnetic Core/shell Nanoparticles. *Phys. Rep.* **2015**, *553*, 1–32.
- (2) Chaudhuri, R. G.; Paria, S. Core/Shell Nanoparticles: Classes, Properties, Synthesis Mechanisms, Characterization, and Applications. *Chem. Rev.* **2012**, *112*, 2373–2433.
- (3) Liu, F.; Hou, Y.; Gao, S. Exchange-Coupled Nanocomposites: Chemical Synthesis, Characterization and Applications. *Chem. Soc. Rev.* **2014**, *43*, 8098–8113.
- (4) *Magnetic Nanoparticles: From Fabrication and Clinical Applications*; Thanh, N. T. K., Ed.; CRC Press,.; 2012.
- (5) Meiklejohn, W. H.; Bean, C. P. New Magnetic Anisotropy. *Phys. Rev.* **1957**, *105*, 904–913.
- (6) Lo, A.; Tobia, D.; Winkler, E.; Golosovsky, I. V; Gonza, M. A.; Surin, S.; Arbiol, J.; Peiro, F.; Zysler, R. D.; Baro, M. D.; *et al.* Size-Dependent Passivation Shell and Magnetic Properties in Antiferromagnetic/Ferrimagnetic Core/Shell MnO Nanoparticles. *J. Am. Chem. Soc.* **2002**, *132*, 9398–9407.
- (7) Kavich, D. W.; Dickerson, J. H.; Mahajan, S. V.; Hasan, S. A.; Park, J. H. Exchange Bias of Singly Inverted FeO/Fe₃O₄ Core-Shell Nanocrystals. *Phys. Rev. B - Condens. Matter Mater. Phys.* **2008**, *78*, 1–6.
- (8) Winkler, E. L.; Lima, E.; Tobia, D.; Saleta, M. E.; Troiani, H. E.; Agostinelli, E.; Fiorani, D.; Zysler, R. D. Origin of Magnetic Anisotropy in ZnO/CoFe₂O₄ and CoO/CoFe₂O₄ Core/shell Nanoparticle Systems. *Appl. Phys. Lett.* **2012**, *101*, 2–6.
- (9) Lottini, E.; López-Ortega, A.; Bertoni, G.; Turner, S.; Meledina, M.; Tendeloo, G. Van; Julián Fernández, C.; Sangregorio, C. Strongly Exchange Coupled Core|Shell Nanoparticles with High Magnetic Anisotropy: A Strategy toward Rare-Earth-Free Permanent Magnets.

- Chem. Mater.* **2016**, *28*, 4214–4222.
- (10) Cannas, C.; Musinu, A.; Peddis, D.; Piccaluga, G. Synthesis and Characterization of CoFe₂O₄ Nanoparticles Dispersed in a Silica Matrix by a Sol–Gel Autocombustion Method. *Chem. Mater.* **2006**, *18*, 3835–3842.
- (11) Fantechi, E.; Innocenti, C.; Albino, M.; Lottini, E.; Sangregorio, C. Influence of Cobalt Doping on the Hyperthermic Efficiency of Magnetite Nanoparticles. *J. Magn. Magn. Mater.* **2015**, *380*, 365–371.
- (12) Mamei, V.; Musinu, A.; Ardu, A.; Ennas, G.; Peddis, D.; Niznansky, D.; Sangregorio, C.; Innocenti, C.; Thanh, N. T. K.; Cannas, C. Studying the Effect of Zn-Substitution on the Magnetic and Hyperthermic Properties of Cobalt Ferrite Nanoparticles. *Nanoscale* **2016**, *8*, 10124–10137.
- (13) Sun, S.; Zeng, H.; Robinson, D. B.; Raoux, S.; Rice, P. M.; Wang, S. X.; Li, G. Monodisperse MFe₂O₄ (M=Fe, Co, Mn) Nanoparticles. *J. Am. Chem. Soc.* **2004**, *126*, 273–279.
- (14) Niederberger, M.; Pinna, N. *Metal Oxide Nanoparticles in Organic Solvents. Synthesis, Formation, Assembly and Application*; Springer, 2009.
- (15) Cannas, C.; Musinu, A.; Ardu, A.; Orrù, F.; Peddis, D.; Casu, M.; Sanna, R.; Angius, F.; Diaz, G.; Piccaluga, G. CoFe₂O₄ and CoFe₂O₄/SiO₂ Core/Shell Nanoparticles: Magnetic and Spectroscopic Study. *Chem. Mater.* **2010**, *22*, 3353–3361.
- (16) Cannas, C.; Ardu, A.; Peddis, D.; Sangregorio, C.; Piccaluga, G.; Musinu, A. Surfactant-Assisted Route to Fabricate CoFe₂O₄ Individual Nanoparticles and Spherical Assemblies. *J. Colloid Interface Sci.* **2010**, *343*, 415–422.
- (17) Cannas, C.; Ardu, A.; Musinu, A.; Suber, L.; Ciasca, G.; Amenitsch, H.; Campi, G.

- Hierarchical Formation Mechanism of CoFe₂O₄ Mesoporous Assemblies. *ACS Nano* **2015**, *9*, 7277–7286.
- (18) Lee, J.-H.; Jang, J.-T.; Choi, J.-S.; Moon, S. H.; Noh, S.-H.; Kim, J.; Kim, J.-G.; Kim, I.-S.; Park, K. I.; Cheon, J. Exchange-Coupled Magnetic Nanoparticles for Efficient Heat Induction. *Nat. Nanotechnol.* **2011**, *6*, 418–422.
- (19) Song, Q.; Zhang, Z. J. Controlled Synthesis and Magnetic Properties of Bimagnetic Spinel Ferrite CoFe₂O₄ and MnFe₂O₄ Nanocrystals with Core-Shell Architecture. *J. Am. Chem. Soc.* **2012**, *134*, 10182–10190.
- (20) Thanh, N. T. K.; Maclean, N.; Mahiddine, S. Mechanisms of Nucleation and Growth of Nanoparticles in Solution. *Chem. Rev.* **2014**, *114*, 7610–7630.
- (21) Xia, Y.; Gilroy, K. D.; Peng, H.-C.; Xia, X. Seed-Mediated Growth of Colloidal Metal Nanocrystals. *Angew. Chemie Int. Ed.* **2017**, *56*, 60–95.
- (22) Zeng, H.; Sun, S.; Li, J.; Wang, Z. L.; Liu, J. P. Bimagnetic Core/Shell FePt/Fe₃O₄ Nanoparticles. *Nano Lett.* **2004**, *4*, 187–190.
- (23) Masala, O.; Hoffman, D.; Sundaram, N.; Page, K.; Proffen, T.; Lawes, G.; Seshadri, R. Preparation of Magnetic Spinel Ferrite Core/shell Nanoparticles: Soft Ferrites on Hard Ferrites and Vice Versa. *Solid State Sci.* **2006**, *8*, 1015–1022.
- (24) Noh, S.-H.; Na, W.; Jang, J.-T.; Lee, J.-H.; Lee, E. J.; Moon, S. H.; Lim, Y.; Shin, J.; Cheon, J. Nanoscale Magnetism Control via Surface and Exchange Anisotropy for Optimized Ferrimagnetic Hysteresis. *Nano Lett.* **2012**, *12*, 3716–3721.
- (25) López-Ortega, A.; Estrader, M.; Salazar-Alvarez, G.; Estradé, S.; Golosovsky, I. V.; Dumas, R. K.; Keavney, D. J.; Vasilakaki, M.; Trohidou, K. N.; Sort, J.; *et al.* Strongly Exchange Coupled Inverse Ferrimagnetic Soft/hard, Mn_xFe_{3-x}O₄/FexMn_{3-x}O₄, Core/shell

- Heterostructured Nanoparticles. *Nanoscale* **2012**, *4*, 5138.
- (26) Chen, J.; Ye, X.; Oh, S. J.; Kikkawa, J. M.; Kagan, C. R.; Murray, C. B. Bistable Magnetoresistance Switching in Exchange-Coupled CoFe₂O₄-Fe₃O₄ Binary Nanocrystal Superlattices by Self-Assembly and Thermal Annealing. *ACS Nano* **2013**, *7*, 1478–1486.
- (27) Estrader, M.; López-Ortega, A.; Estradé, S.; Golosovsky, I. V.; Salazar-Alvarez, G.; Vasilakaki, M.; Trohidou, K. N.; Varela, M.; Stanley, D. C.; Sinko, M.; *et al.* Robust Antiferromagnetic Coupling in Hard-Soft Bi-Magnetic Core/shell Nanoparticles. *Nat. Commun.* **2013**, *4*, 2960.
- (28) Gavrilov-Isaac, V.; Neveu, S.; Dupuis, V.; Taverna, D.; Gloter, A.; Cabuil, V. Synthesis of Trimagnetic Multishell MnFe₂O₄@CoFe₂O₄@NiFe₂O₄ Nanoparticles. *Small* **2015**, *11*, 2614–2618.
- (29) Gomes, J. D. A.; Sousa, M. H.; Tourinho, F. A.; Aquino, R.; da Silva, G. J.; Depeyrot, J.; Dubois, E.; Perzynski, R. Synthesis of Core–Shell Ferrite Nanoparticles for Ferrofluids: Chemical and Magnetic Analysis. *J. Phys. Chem. C* **2008**, *112*, 6220–6227.
- (30) Honarbakhsh-Raouf, A.; Emamian, H. R.; Yourdkhani, A.; Ataie, A. Synthesis and Characterization Of CoFe₂O₄/Ni_{0.5}Zn_{0.5}Fe₂O₄ Core/Shell Magnetic Nanocomposite By The Wet Chemical Route. *Int. J. Mod. Phys. B* **2010**, *24*, 5807–5814.
- (31) Wang, J.; Zhou, Z.; Wang, L.; Wei, J.; Yang, H.; Yang, S.; Zhao, J. CoFe₂O₄@MnFe₂O₄/polypyrrole Nanocomposites for in Vitro Photothermal/magnetothermal Combined Therapy. *RSC Adv.* **2015**, *5*, 7349–7355.
- (32) *Handbook of Nanoparticles*; Aliofkhazraei, M., Ed.; Springer International Publishing: Cham, 2016.
- (33) Repko, A.; Nižňanský, D.; Poltírová-Vejpravová, J. A Study of Oleic Acid-Based

- Hydrothermal Preparation of CoFe₂O₄ Nanoparticles. *J. Nanoparticle Res.* **2011**, *13*, 5021–5031.
- (34) Repko, A.; Vejpravová, J.; Vacková, T.; Zákutná, D.; Nižňanský, D. Oleate-Based Hydrothermal Preparation of CoFe₂O₄ Nanoparticles, and Their Magnetic Properties with Respect to Particle Size and Surface Coating. *J. Magn. Magn. Mater.* **2015**, *390*, 142–151.
- (35) Cara, C.; Musinu, A.; Mamei, V.; Ardu, A.; Niznansky, D.; Bursik, J.; Scorciapino, M. A.; Manzo, G.; Cannas, C. Dialkylamide as Both Capping Agent and Surfactant in a Direct Solvothermal Synthesis of Magnetite and Titania Nanoparticles. *Cryst. Growth Des.* **2015**, *15*, 2364–2372.
- (36) Almeida, T. P.; Moro, F.; Fay, M. W.; Zhu, Y.; Brown, P. D. Tuneable Magnetic Properties of Hydrothermally Synthesised Core/shell CoFe₂O₄/NiFe₂O₄ and NiFe₂O₄/CoFe₂O₄ Nanoparticles. *J. Nanoparticle Res.* **2014**, *16*, 1–13.
- (37) Sattar, A. A.; EL-Sayed, H. M.; ALSuqia, I. Structural and Magnetic Properties of CoFe₂O₄/NiFe₂O₄ Core/shell Nanocomposite Prepared by the Hydrothermal Method. *J. Magn. Magn. Mater.* **2015**, *395*, 89–96.
- (38) van Ewijk, G. A.; Vroege, G. J.; Philipse, A. P. Convenient Preparation Methods for Magnetic Colloids. *J. Magn. Magn. Mater.* **1999**, *201*, 31–33.
- (39) Gnanaprakash, G.; Philip, J.; Jayakumar, T.; Raj, B. Effect of Digestion Time and Alkali Addition Rate on Physical Properties of Magnetite Nanoparticles. *J. Phys. Chem. B* **2007**, *111*, 7978–7986.
- (40) Ayyappan, S.; Gnanaprakash, G.; Panneerselvam, G.; Antony, M. P.; Philip, J. Effect of Surfactant Monolayer on Reduction of Fe₃O₄ Nanoparticles under Vacuum. *J. Phys. Chem. C* **2008**, *112*, 18376–18383.

- (41) Coey, J. M. D. *Magnetism and Magnetic Materials*; 2010.
- (42) Zuo, X.; He, Y.; Yang, A.; Bernardo, B.; Harris, V. G.; Vittoria, C. Calculation of Exchange Constants in Spinel Ferrites with Magnetic S-State Ions. *J. Appl. Phys.* **2005**, *97*, 10F104.
- (43) Han, L.; Meng, Q.; Wang, D.; Zhu, Y.; Wang, J.; Du, X.; Stach, E. A.; Xin, H. L. Interrogation of Bimetallic Particle Oxidation in Three Dimensions at the Nanoscale. *Nat. Commun.* **2016**, *7*, 13335.
- (44) Wang, D.; Liu, S.; Wang, J.; Lin, R.; Kawasaki, M.; Rus, E.; Silberstein, K. E.; Lowe, M. A.; Lin, F.; Nordlund, D.; *et al.* Spontaneous Incorporation of Gold in Palladium-Based Ternary Nanoparticles Makes Durable Electrocatalysts for Oxygen Reduction Reaction. *Nat. Commun.* **2016**, *7*, 11941.
- (45) Wang, D.; He, H.; Han, L.; Lin, R.; Wang, J.; Wu, Z.; Liu, H.; Xin, H. L. Three-Dimensional Hollow-Structured Binary Oxide Particles as an Advanced Anode Material for High-Rate and Long Cycle Life Lithium-Ion Batteries. *Nano Energy* **2016**, *20*, 212–220.
- (46) Schlossmacher, P.; Klenov, D. O.; Freitag, B.; Harrach, S. Von; Steinbach, A. Nanoscale Chemical Compositional Analysis with an Innovative S/TEM-EDX System. *Microsc. Anal. Nanotechnol. Suppl.* **2010**, *24*, S5–S8.
- (47) Schlossmacher, P.; Klenov, D. O.; Freitag, B.; von Harrach, H. S. Enhanced Detection Sensitivity with a New Windowless XEDS System for AEM Based on Silicon Drift Detector Technology. *Microsc. Today* **2010**, *18*, 14–20.
- (48) Harrach, H. S. Von; Dona, P.; Freitag, B.; Soltau, H.; Niculae, A.; Rohde, M. An Integrated Silicon Drift Detector System for FEI Schottky Field. *Microsc. Microanal.* **2009**, *15*, 8–9.
- (49) Allen, L. J.; D'Alfonso, A. J.; Freitag, B.; Klenov, D. O. Chemical Mapping at Atomic Resolution Using Energy-Dispersive X-Ray Spectroscopy. *MRS Bull.* **2012**, *37*, 47–52.

- (50) Kothleitner, G.; Neish, M. J.; Lugg, N. R.; Findlay, S. D.; Grogger, W.; Hofer, F.; Allen, L. J. Quantitative Elemental Mapping at Atomic Resolution Using X-Ray Spectroscopy. *Phys. Rev. Lett.* **2014**, *112*, 1–5.
- (51) Gomes, J. A.; Azevedo, G. M.; Depeyrot, J.; Mestnik-Filho, J.; Paula, F. L. O.; Tourinho, F. A.; Perzynski, R. Structural, Chemical, and Magnetic Investigations of Core–Shell Zinc Ferrite Nanoparticles. *J. Phys. Chem. C* **2012**, *116*, 24281–24291.
- (52) Repko, A.; Nižňanský, D.; Matulková, I.; Kalbáč, M.; Vejpravová, J. Hydrothermal Preparation of Hydrophobic and Hydrophilic Nanoparticles of Iron Oxide and a Modification with CM-Dextran. *J. Nanoparticle Res.* **2013**, *15*, 1767.
- (53) Mondini, S.; Ferretti, A. M.; Puglisi, A.; Ponti, A. Pebbles and PebbleJuggler: Software for Accurate, Unbiased, and Fast Measurement and Analysis of Nanoparticle Morphology from Transmission Electron Microscopy (TEM) Micrographs. *Nanoscale* **2012**, *4*, 5356–5372.

