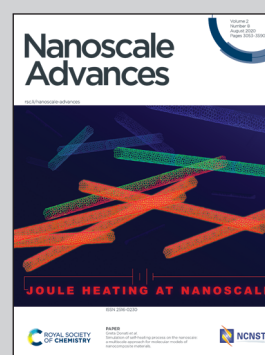


Showcasing research from Prof. Cannas, Department of Chemical and Geological Science, University of Cagliari, Italy, and Prof. Vejpravova, Department of Condensed Matter Physics, Charles University, Czech Republic.

Coupled hard-soft spinel ferrite-based core-shell nanoarchitectures: magnetic properties and heating abilities

Magnetic coupling between hard and soft spinel ferrites in a core-shell architecture permits to enhance the heat released by the nanoparticles under high-frequency magnetic fields. This work proposes a systematic study, on different sets of samples, of the core size, the chemical nature, and the thickness of the shell on the heating abilities. The correlation between the performances and the static/dynamic magnetic properties is possible thanks to the comprehensive study of chemical composition, in terms of stoichiometry and nanoscale mapping, oxidation state, spin canting, and cation distribution.

As featured in:



See Carla Cannas and Jana Vejpravova *et al.*, *Nanoscale Adv.*, 2020, 2, 3191.

Cite this: *Nanoscale Adv.*, 2020, 2, 3191

Coupled hard–soft spinel ferrite-based core–shell nanoarchitectures: magnetic properties and heating abilities†

Marco Sanna Angotzi, ^{‡ab} Valentina Mameli, ^{‡ab} Claudio Cara, ^{ab}
Anna Musinu, ^{ab} Claudio Sangregorio, ^{bcd} Daniel Niznansky, ^e
Huolin L. Xin, ^{fg} Jana Vejpravova ^{*eh} and Carla Cannas ^{*abi}

Bi-magnetic core–shell spinel ferrite-based nanoparticles with different CoFe_2O_4 core size, chemical nature of the shell (MnFe_2O_4 and spinel iron oxide), and shell thickness were prepared using an efficient solvothermal approach to exploit the magnetic coupling between a hard and a soft ferrimagnetic phase for magnetic heat induction. The magnetic behavior, together with morphology, stoichiometry, cation distribution, and spin canting, were investigated to identify the key parameters affecting the heat release. General trends in the heating abilities, as a function of the core size, the nature and the thickness of the shell, were hypothesized based on this systematic fundamental study and confirmed by experiments conducted on the water-based ferrofluids.

Received 18th February 2020
Accepted 5th May 2020

DOI: 10.1039/d0na00134a

rsc.li/nanoscale-advances

Introduction

Coupled bi-magnetic nanoparticles (NPs) exhibit outstanding properties, strongly dependent on the extent and nature of the interface.^{1–4} Antiferromagnet (AFM)/ferromagnet (FM) and AFM/ferrimagnet (FiM) interfaces are the most frequently studied systems. Their magnetic behaviour is characterized by exchange bias, which is manifested by an increase and asymmetry in the coercivity, giving rise to a shift of the hysteresis loop along the magnetic field axis.^{5–9} Since the pioneering work by Kneller and Hawig,¹⁰ many studies have focused on the investigation at the nanoscale of the coupling phenomena between hard and soft FM or FiM phases.¹¹ In particular, the magnetic behaviour of thin films has been deeply studied and described as a specific superposition of the intrinsic parameters of the *hard* and *soft* phases depending on the coupling strength

between the phases. Indeed, *hard* phases feature large magnetic anisotropy (expressed in terms of anisotropy constant, K) and moderate saturation magnetization (M_s), while *soft* phases present low K and large M_s . In thin films, the magnetization switching behaviour (*i.e.*, the hysteresis loop) depends on the relative dimension (thickness of the layer) of the *hard* and *soft* phases. When the *soft* phase thickness is lower than the double of the domain wall thickness of the *hard* one ($t_s \ll 2 \delta_h$), the phases are rigidly coupled and result in a rectangular stage-loop since the magnetization is reversed at the same nucleation field (H_N). In the case of thicker *soft* phases, the system behaves like a “spring-magnet,” showing different switching of the magnetization and, therefore a two-step loop.¹¹ Strongly coupled *hard*–*soft* heterostructures are desired for several applications. For example, they can be employed to replace rare-earth-based magnets, ideally combining the attractive properties of the

^aDepartment of Chemical and Geological Sciences, University of Cagliari, S.S. 554 bivio per Sestu, 09042 Monserrato (CA), Italy. E-mail: ccannas@unica.it

^bConsorzio Interuniversitario Nazionale per la Scienza e Tecnologia dei Materiali (INSTM), Via Giuseppe Giusti 9, 50121 Firenze (FI), Italy

^cIstituto di Chimica dei Composti OrganoMetallici - Consiglio Nazionale delle Ricerche (ICCOM-CNR), Via Madonna del Piano 10, 50019 Sesto Fiorentino (FI), Italy

^dDepartment of Chemistry “U. Schiff”, University of Florence, Via della Lastruccia 3-13, 50019, Sesto Fiorentino (FI), Italy

^eDepartment of Inorganic Chemistry, Charles University, Hlavova 8, 12800 Prague 2, Czech Republic. E-mail: jana@mag.mff.cuni.cz

^fCenter for Functional Nanomaterials, Brookhaven National Laboratory, 735 Brookhaven Ave, Upton, NY 11973, USA

^gDepartment of Physics and Astronomy, University of California, Irvine, CA 92697, USA

^hDepartment of Condensed Matter Physics, Charles University, Ke Karlovu 5, 12116 Prague 2, Czech Republic

ⁱConsorzio per la Promozione di Attività Universitarie Sulcis-Iglesiente (AUSI), Centro di Ricerca per l'Energia, l'Ambiente e il Territorio (CREATE), Palazzo Bellavista Monteponi, 09016 Iglesias (CI), Italy

† Electronic supplementary information (ESI) available: Samples characterization; low-temperature Mössbauer spectroscopy; DC magnetometry; details on magnetic fluid hyperthermia; AC magnetometry on powdered samples; intercalation process; magnetometry and calorimetric measurements on hydrophilic ferrofluids; literature MFH studies of bimagnetic spinel ferrite-based core–shell nanoparticles; experimental synthesis conditions. See DOI: 10.1039/d0na00134a

‡ These authors contributed equally.



hard (large K) and *soft* (high M_s) phases.^{12–16} The thermal stability of the *hard* component, combined with the ability of the *soft* part to reduce the switching field may be exploited in recording media, where to increase the areal bit density it is necessary to avoid the superparamagnetic limit without having such extensive magnetic anisotropy that would prevent the write-head to write the information.^{15,17,18} Again, the possibility to tune magnetic anisotropy and saturation magnetization has recently found usage in applications based on magnetic heat generation, such as catalysis^{19–22} or magnetic fluid hyperthermia (MFH).^{23–38} In this context, the engineering of bi-magnetic core-shell nanoarchitectures with sharp interfaces, homogeneous coating, and low size dispersity for a uniform magnetic response is crucial for maximizing the coupling between the *hard* and *soft* phases (*i.e.*, the interface) and requests proper synthesis methods able to guarantee a strict control over the composition, structure, and morphology of the particles.^{39–43} Isostructural phases with similar cell parameters should ensure epitaxial growth and can be considered the best candidates to build core-shell architectures. From this point of view, spinel ferrites are particularly appealing as they offer a broad spectrum of *hard* and *soft* magnetic behaviours, all sharing the same crystal structure. However, a systematic study of the magnetic parameters influencing the coupling, as already performed for thin films,⁴⁴ is still lacking for core-shell NPs. Some studies^{23–38} have reported the heating abilities of these systems since Lee *et al.* work in 2011 (Table 7S†).³³ Nevertheless, only a few of them discuss the data based on a large set of samples,^{28,31,35–37} and rarely the variation of a single parameter is studied keeping the other unchanged.^{23,24,30,31,33,35,37,38} More importantly, the undoubted demonstration of the core-shell architecture by nanoscale chemical mapping,^{26,32,33} and the actual chemical composition^{23,28,32,35,38} are hardly ever provided. All the above-cited aspects are essential as a solid starting point to ascribe the increased efficiency of systems to the exchange-coupled spinel ferrite phases. Generally, the heat release efficiency is explained in terms of optimized size, saturation magnetization, and effective magnetic anisotropy caused by exchange interaction. Nonetheless, some other aspects are usually ignored, as the cation distribution and the spin canting, even though their influence on the magnetic properties has been largely demonstrated.^{45–48} Notwithstanding important results that have been achieved on these promising systems in terms of heat release, the available literature data are still limited, and systematic studies are necessary to better understand the correlation between materials' feature and heat dissipation mechanism.

Recently, we set up an alternative synthesis method able to produce homogeneous core-shell NPs with high crystallinity, low dispersity, and precise control of the shell growth, as proven by a detailed characterization by XRD, STEM-EELS, STEM-EDX, FTIR, TGA, and room temperature ⁵⁷Fe Mössbauer spectroscopy.⁴³

Here we propose a systematic study of the DC/AC magnetic properties of three *hard* cobalt ferrite (CoFe₂O₄) cores covered with *soft* spinel iron oxide (magnetite Fe₃O₄/maghemite γ -Fe₂O₃) and manganese ferrite (MnFe₂O₄), characterized for their actual chemical composition (ICP-AES), their nanoscale

architecture and morphology (STEM-EDX), cation distribution, and spin canting (low-temperature ⁵⁷Fe Mössbauer spectroscopy). The evaluation of magnetic parameters provides hypotheses of general trends in the heating abilities as a function of the core size, the nature and the thickness of the shell, that have been then compared with experimental heat abilities obtained from aqueous ferrofluids.

Experimental

Methods

Three samples of CoFe₂O₄ NPs of different sizes, labelled as CoA, CoB, and CoC, were prepared by the solvothermal hydrolysis of mixed cobalt-iron oleates in a mixture of organic solvents with different polarities and water contents. The samples were used as seeds to produce core-shell nanostructures using a second solvothermal treatment (seed-mediated growth), with a shell of spinel iron oxide (maghemite/magnetite) and manganese ferrite, indicated as Co_x@Fe and Co_x@Mn (where $x = A, B, C$), respectively.⁴³ The samples CoA, CoA@Fe, CoA@Mn, CoC, CoC@Fe, and CoC@Mn correspond to the samples Co1, Co1@Fe, Co1@Mn, Co2, Co2@Fe, and Co2@Mn, respectively, described in previous work.⁴³ The experimental synthesis conditions for the samples CoB, CoB@Fe, and CoB@Mn are reported in the section “*Experimental conditions*” of the ESI (Tables 8S–10S†).

Equipment

The samples were characterized by X-ray diffraction (XRD), using a PANalytical X'Pert PRO with Cu K α radiation (1.5418 Å), a secondary monochromator, and a PIXcel position-sensitive detector. The peak position and instrumental width were calibrated 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 experimental diffraction patterns were fitted using the FullProf program.⁴⁹ The microstructural effects were treated using the Voigt approximation as implemented in the code: both instrumental and sample intrinsic profiles are supposed to be described by a convolution of Lorentzian and Gaussian components. The Gaussian and Lorentzian components of the instrumental function were determined using the diffraction pattern of the LaB₆ standard recorded for the identical instrumental set up, and the sample broadening was approximated by the Gaussian component only. In the structural model, the effects of spinel inversion and the core-shell nature approximated as a two-phase model were tested.

Transmission electron microscopy (TEM) images were obtained using a JEOL 200CX operating at 160 kV. The particle size distribution was obtained by measuring in the automatic mode over 1000 particles through the software Pebbles and adopting a spherical shape.⁵⁰ The mean particle diameter was calculated as the average value and the dispersity as the percentage ratio between the standard deviation and the average value.

High resolution TEM (HRTEM) images and EDX measurements were carried out in the STEM mode using an 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. The single line profiles were calculated using the Matlab command “improfile” for different sections all over a particle (*i.e.* over 360° with a step of 0.1°), and the results averaged.

⁵⁷Fe Mössbauer spectroscopy was done on a Wissel spectrometer using transmission arrangement and proportional detector LND-45431. An α -Fe foil was used as a standard, and the fitting procedure was done by the NORMOS program. The in-field measurements were done in a perpendicular arrangement of the external magnetic field with respect to the γ -beam and were used to get information about the cationic distribution and the canting phenomena in the spinel structure (see paragraph “Low-temperature Mössbauer spectroscopy” in the ESI, eqn (1S)†).

The chemical composition was studied by inductively coupled plasma-atomic emission spectroscopy (ICP-AES). The dried samples were digested 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 allowed to cool down, filtered, and diluted using 1% v/v HNO₃ solution. The ICP-AES measurements were performed on 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 were performed twice on different portions of the samples. The chemical formulas were calculated by assuming the absence of anion vacancies.

The magnetic property measurements were carried out using a SQUID magnetometer (MPMS7XL, Quantum Design). The temperature dependencies of magnetization in the zero field cooled (ZFC) and field cooled (FC) regimes were measured as follows: first, the sample was cooled down to 10 K in the zero external magnetic field. Next, the field of 10 mT was applied, and the temperature dependence of magnetization was measured on heating. Afterward, the sample was cooled down

in the applied field of 10 mT, and the temperature dependence of magnetization was measured again. The magnetization isotherms were recorded up to 7 T at selected temperatures in both polarities of the applied magnetic field. All data were corrected according to the organic content. AC susceptibility measurements were recorded with the amplitude of 0.3 mT and in the frequency range of 0.1–1000 Hz between temperatures of 10–400 K. The approach for the evaluation of magnetic parameters, including exact formulas, is reported in the ESI† (see paragraphs “DC Magnetometry”, “AC Magnetometry on powdered samples”, and “AC/DC Magnetometry on hydrophilic ferrofluids”).

Calorimetric estimation of specific absorption rate (SAR) was carried out using a non-adiabatic experimental set-up built at the Laboratorio di Magnetismo Molecolare (L.A.M.M.) using a power supply CELEs MP6/400 (FIVES CELES), a water-cooled heating station connected to the power supply, and an induction coil. Heating curves were recorded under a magnetic field of 17 kA m⁻¹ and 183 kHz for 300 s on water colloidal dispersions of the magnetic NPs. The hydrophobic NPs were made hydrophilic by an intercalation process with cetyltrimethylammonium bromide (CTAB, (C₁₆H₃₃N(CH₃)₃Br).⁴⁵ The concentration of the colloidal dispersion was 3.4 mg mL⁻¹ for all samples. The temperature of the sample was monitored by an optical fiber probe (OPTOCON-FOTEMP) dipped into the solution. The sample holder was surrounded by polystyrene and hosted in a glass Dewar, equipped by an ethylene glycol thermostat, to ensure the proper thermal insulation. The SAR, *i.e.*, the thermal power per mass unit, values have been estimated by a linear curve fitting in the first 20 s of the heating curves (initial slope method).

Results and discussion

Three samples of CoFe₂O₄ NPs of different sizes (CoA, CoB, and CoC) and their corresponding core-shell nanostructures (Cox@Mn and Cox@Fe, where Mn = MnFe₂O₄ and Fe = γ -Fe₂O₃/Fe₃O₄; x = A, B, C) were synthesized by solvothermal

Table 1 Cell parameter (*a*), volume-weighted particle size ($\langle D_{\text{TEM}_V} \rangle$) and standard deviation (SD_ D_{TEM_V}), shell thickness (Δ_{TEM}) calculated as the difference between $\langle D_{\text{TEM}_V} \rangle$ of the core-shell and core, magnetic size ($\langle D_{\text{MAG}} \rangle$), and M : Fe ratio determined by ICP-AES of the core-shell samples and respective core

Sample	<i>a</i> (Å)	$\langle D_{\text{TEM}_V} \rangle$ (nm)	SD_ D_{TEM_V} (nm)	Shell thickness Δ_{TEM} (nm)	$\langle D_{\text{MAG}} \rangle$ (nm)	M : Fe ^a
CoA	8.39(1)	5.9	1.0	—	4.4	0.49
CoA@Mn	8.40(2)	9.7	1.1	1.9	4.8	0.41
CoA@Fe	8.36(1)	10.9	1.2	2.5	8.0	—
CoB	8.38(1)	7.5	1.1	—	5.1	0.45
CoB@Mn	8.41(1)	13.2	1.6	2.9	6.3	0.43
CoB@Fe	8.35(1)	12.8	1.7	2.7	6.5	—
CoC	8.38(1)	9.0	1.3	—	5.3	0.55
CoC@Mn1	8.44(9)	13.3	1.7	2.1	6.9	0.41
CoC@Mn2	8.43(1)	14.9	1.6	3.0	7.6	0.45
CoC@Fe1	8.36(9)	11.7	1.5	1.4	8.4	—
CoC@Fe2	8.38(1)	12.8	1.7	1.9	8.2	—

^a Referred to the shell fraction in case of core-shell samples.



hydrolysis of metal-oleates in a mixture of organic solvents and water.^{43,51–53} Two different shell thicknesses, termed as CoC@Mn1/CoC@Mn2 and CoC@Fe1/CoC@Fe2, were prepared for CoC-based core-shell samples. The samples CoA, CoC, and respective core-shell NPs (except CoC@Mn1 and CoC@Fe1) have been described in a previous work⁴³ (where CoA was Co1 and CoC was Co2) and the results of their characterization, together with those of the sample CoB and core shells, are reported in Table 1. A complete list of the sizes and size distribution calculated by different techniques is reported in Table 1S.†

As for the CoA and CoC sets of core and core-shell samples,⁴³ the other samples (CoB, CoB@Fe, CoB@Mn, CoC@Mn1, CoC@Mn2) showed a XRD pattern typical of a spinel ferrite with no other phases (Fig. 1S and 2S†). The core-shell samples showed larger crystalline size with respect to the core and different cell parameter (a), higher for Cox@Mn and lower for Cox@Fe (Table 1). The TEM bright-field images (Fig. 1S and 2S†) revealed spherical NPs with narrow and monomodal size distribution (Table 1 and Fig. 1). HRTEM images (Fig. 1) revealed the high crystallinity of the NPs and epitaxial growth of the shell on the pre-existing cores without lattice mismatches. The formation of the core-shell architecture was proven through nanoscale chemical mapping by STEM-EELS and EDX.⁴³ Fig. 2 shows the STEM-EDX nanoscale chemical mapping of CoB, CoB@Mn, and CoB@Fe

with the corresponding line profile that unambiguously indicated the formation of a core-shell heterostructure through a homogeneous coating of the isostructural spinel phase around the core.

As stoichiometry, degree of inversion, and spin canting are crucial in determining the magnetic properties of spinel ferrite NPs, ICP-AES measurements (M : Fe ratios reported in Table 1) and low-temperature (4 K) ⁵⁷Fe Mössbauer spectroscopy in the absence and presence of a magnetic field (6 T) were performed on some selected samples (Fig. 3S, 4S and Table 2S†).

Concerning the cores, almost stoichiometric cobalt ferrite with a degree of inversion of approximately 0.7 was revealed for all samples, in agreement with previous studies.^{45,54–58} A sub-stoichiometry with a Mn : Fe ratio in the range of 0.40–0.45 and a degree of inversion of approximately 0.45 was found for all Cox@Mn samples, in agreement with the trend observed for single manganese ferrite NPs prepared by the same solvothermal method, suggesting the absence of Mn^{III} and indicating a slight preference of Mn^{II} for tetrahedral coordination in all samples.^{43,52,53,59} In the case of Cox@Fe samples, direct information on Fe^{II} : Fe^{III} was not achievable, but through the $[\text{Fe}]_{\text{oh}}/(\text{Fe})_{\text{Td}}$ ratio, it was possible to determine the nature of the iron-based shell (see paragraph entitled “Low-temperature Mössbauer Spectroscopy” in ESI for details†). In particular, this ratio should theoretically be 1.67 for maghemite and 2 for magnetite.⁶⁰ The core-shell samples showed values close to 1.67 with higher

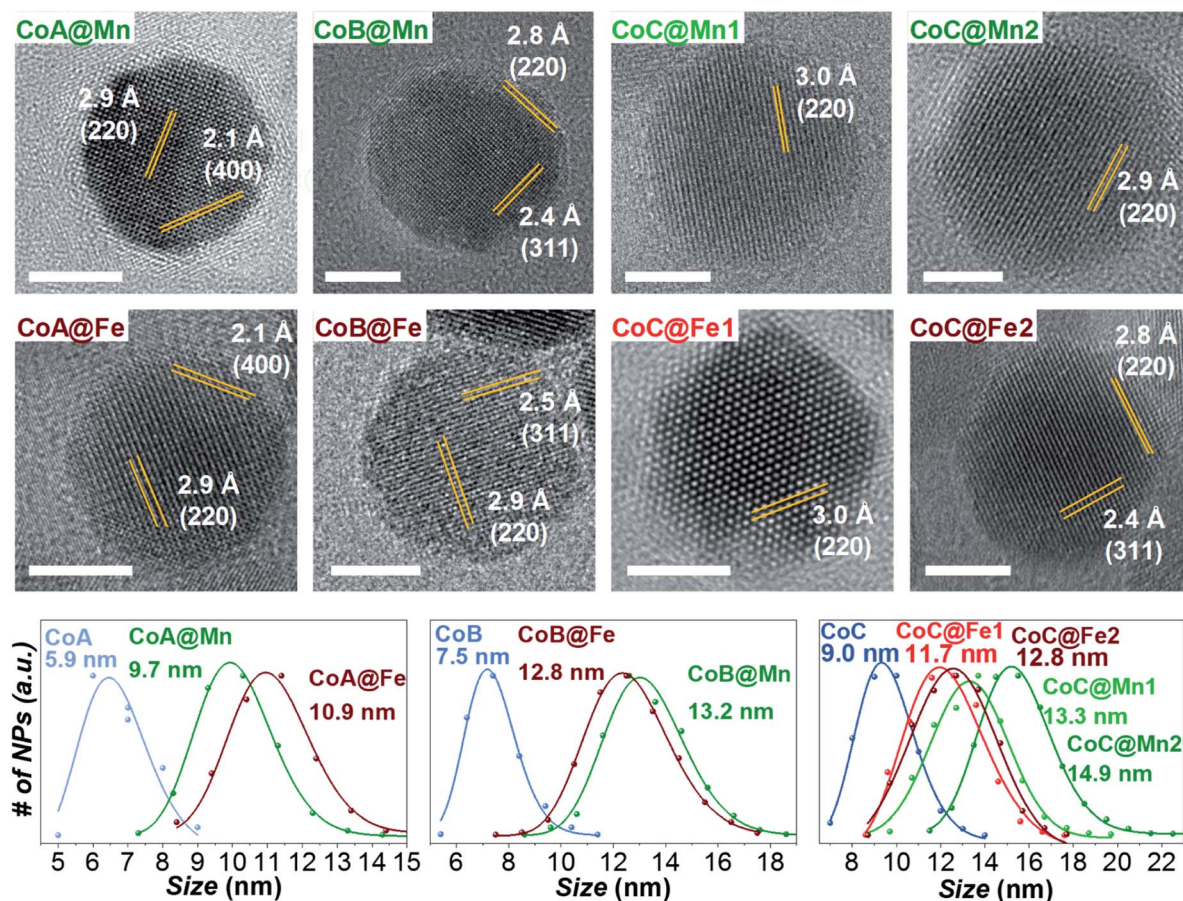


Fig. 1 HRTEM images and particle size distributions of the samples.



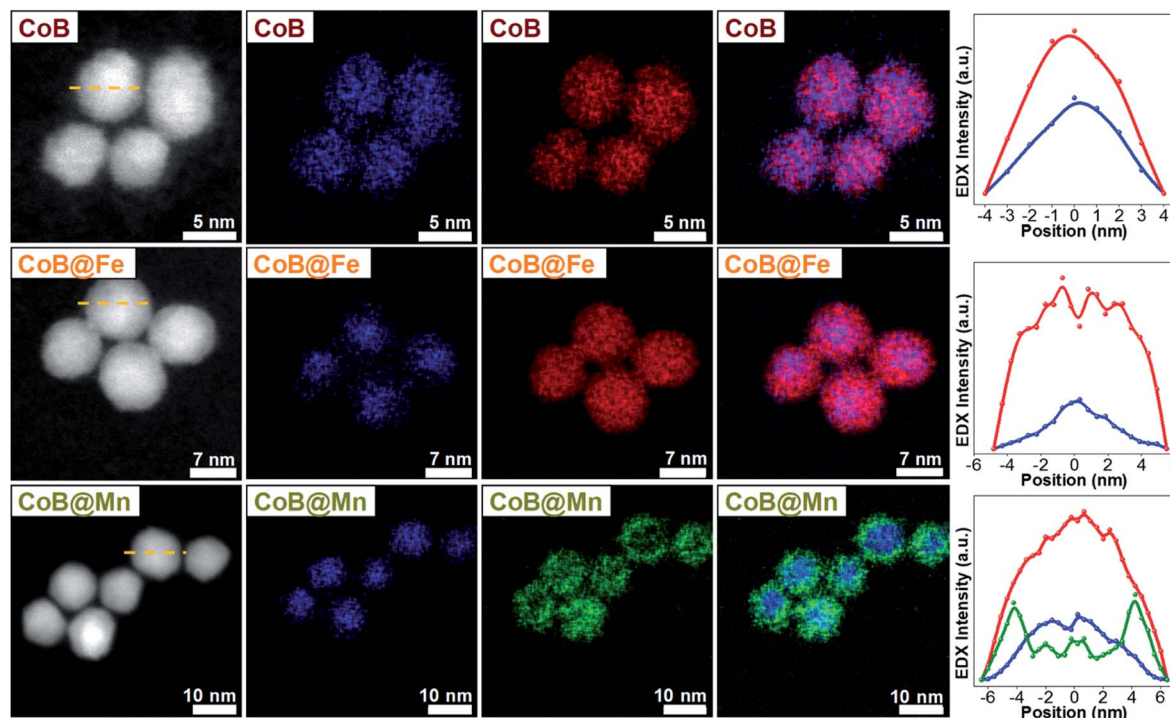


Fig. 2 STEM-EDX maps and line profiles across the NPs along the yellow dotted line of the samples CoB, CoB@Fe, and CoB@Mn. Cobalt is represented in blue, manganese in green, iron in red.

values for CoB@Fe and CoC@Fe, suggesting that oxidation phenomena took place at the surface (Table 2S[†]). These findings allowed the shell to be described as primarily composed of maghemite, even though magnetite is also present, especially for the larger core-shell samples (CoB@Fe and CoC@Fe). Moreover, spin canting (Table 2S[†]) was not revealed in both core and core-shell samples, supporting the evidence from HRTEM data of a homogeneous epitaxial growth of the shell around the core with a formation of a single crystalline domain, as also corroborated by XRD analysis (Table 1S[†]).

In light of these results, the magnetic properties can be discussed beyond any stoichiometric or structural variability.

The samples were characterized by DC and AC magnetometry measurements. ZFC-FC protocols and magnetization isotherms at 10 K and 300 K (Fig. 3, 5S and 6S[†]) were carried out on the cores and core-shell NPs to study the magnetic coupling between the *hard* and *soft* FiM phases. The rigid coupling between the two FiM phases in the core-shell systems was highlighted by (i) the presence of a single-stage hysteresis loop at 10 K (Fig. 3); (ii) a decrease in the coercive and anisotropy fields with respect to the corresponding cores at 10 K (Fig. 3 and Table 2); (iii) the presence of a single dominating maximum in ZFC-FC curves shifted toward higher temperatures, suggesting an increase in effective magnetic anisotropy and/or magnetic diameter (Fig. 5S[†] and Table 2); (iv) an increase in the magnetic diameter ($\langle D_{\text{MAG}} \rangle$) estimated by eqn (5S)[†] at 300 K as reported in the “DC magnetometry (calculations)” paragraph in ESI,[†] Table 1).

Furthermore, the comparison between the core-shell systems with a physical reference mixture of cobalt ferrite and

manganese ferrite of approximately 8 nm ($\langle D_{\text{XRD}} \rangle$) in a 1 : 1 mass ratio corroborates this hypothesis. For this sample, as expected, the hysteresis loop and the ZFC-FC curves (Fig. 3 and 4) display the contribution of the two individual spinel ferrite phases. Due to the low thickness of the *soft* shell, the two phases are expected to be rigidly coupled and reverse at the same nucleation field, H_n (which, as a first approximation is assumed to correspond to the anisotropy field H_K , listed in Table 2),⁶¹ which depends on the anisotropy constant, saturation magnetization, and volume fraction of the *soft* and *hard* phases.¹¹ Once the single-phase magnetic behaviour for all samples and the magnetic coupling in the core-shell samples were ascertained, the DC magnetic measurements together with the temperature dependence of in-phase and out-of-phase susceptibilities (on powders) were investigated to evaluate the critical magnetic parameters known as affecting the heating ability, such as saturation magnetization, magnetic volume, magnetic anisotropy, and Néel relaxation time (see “Details on magnetic fluid hyperthermia” paragraph in the ESI, Fig. 7S and 8S[†]).^{62,63}

At 10 K (Fig. 3, right side), the hysteretic behaviour is characterised by saturation magnetization values similar for all samples, in the range 90–100 A m² kg⁻¹. The reduced remanent magnetization (M_r/M_s) is around 0.7 for manganese ferrite coated core-shells and 0.8 for spinel iron oxide ones, suggesting a moderate prevalence of cubic to uniaxial anisotropy. Anisotropy field H_K is around 4 T for cobalt ferrite samples and in the range of 1.8–2.6 T for the core-shell samples, due to the rigid coupling of the *soft*-shell. On the contrary, magnetization isotherms at 300 K (Fig. 3, left side) display a superparamagnetic behaviour for all samples



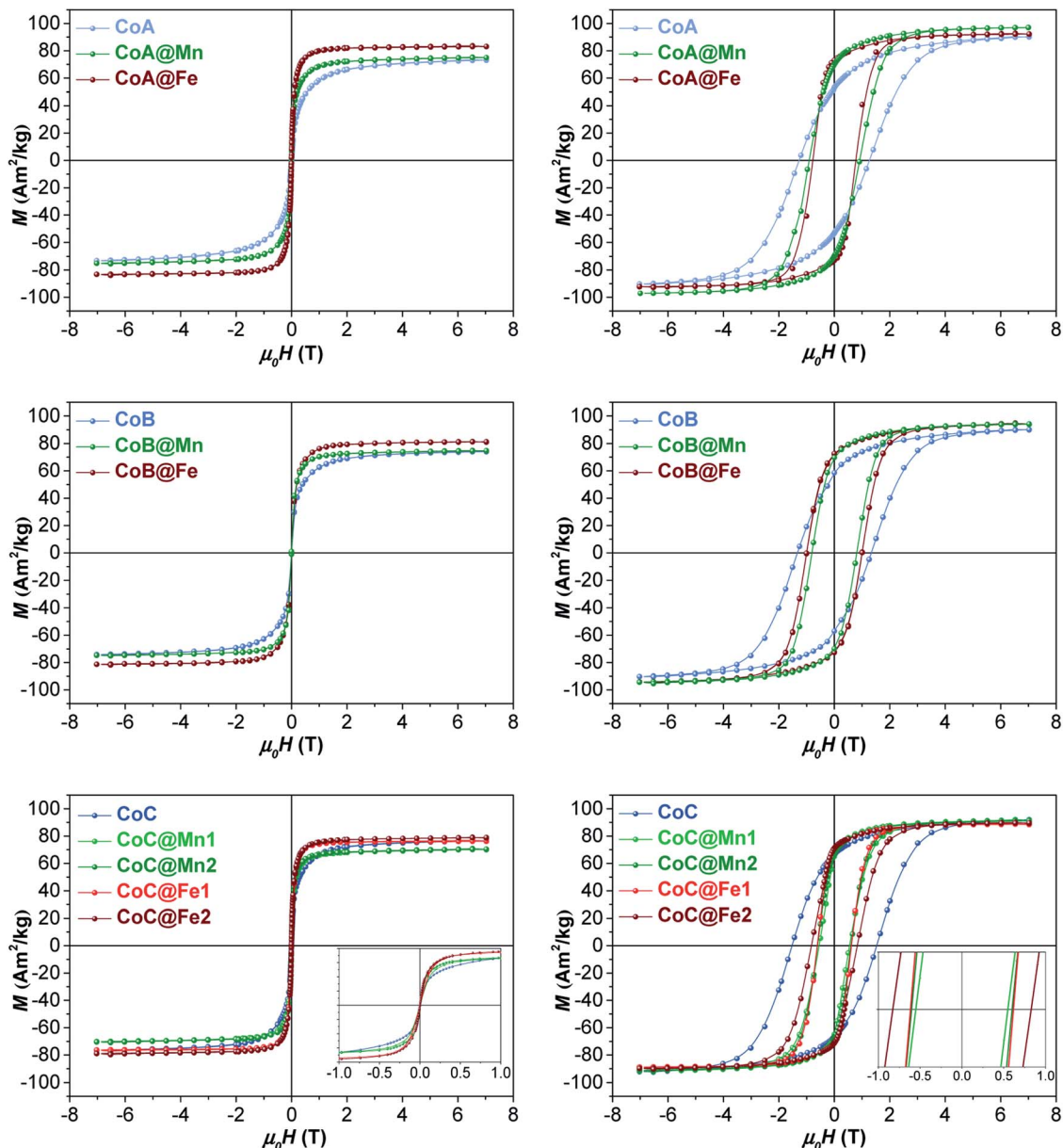


Fig. 3 Magnetization isotherms of core-shell samples and respective cores recorded at 300 K (left) and 10 K (right).

with generally higher saturation magnetization values (M_s^{300} , Table 2) for the core-shell NPs with respect to the cores, in agreement with the literature for similar systems.^{25,34,35} The highest saturation magnetization is reached for spinel iron oxide coated core-shell NPs. This is apparently in contrast with the results reported in the literature for similar systems prepared through a different synthesis strategy,³³ where higher saturation magnetization values were found for manganese ferrite shells than iron-oxide ones. Such a discrepancy could be related to differences in the formation mechanism of the nanoparticles, strictly dependent on the synthesis method, which may influence stoichiometry of the constituents, oxidation state of the metal(s), degree of inversion, spin canting phenomena, and as a consequence the magnetic properties. Moreover, within each set of samples, the magnetization at a low field (<1 T) and the magnetic

energy dissipation are always higher in the $\text{Co}_x\text{@Fe}$ with respect to $\text{Co}_x\text{@Mn}$, as shown in the insets of Fig. 3. The median magnetic moment (μ_m , Table 2) of the NPs and the magnetic diameter ($\langle D_{\text{MAG}} \rangle$, Tables 1 and 3S†) were estimated. $\langle D_{\text{MAG}} \rangle$ was found to increase moving from CoA to CoC, in agreement with the increased crystalline and particle sizes. As mentioned before, also an increase of $\langle D_{\text{MAG}} \rangle$ was observed in the core-shell samples as expected since a homogenous growth of magnetically coupled phases took place. Moreover, higher values were found for $\text{Co}_x\text{@Fe}$ samples with respect to the $\text{Co}_x\text{@Mn}$, and for CoC@Mn2 and CoC@Fe2 compared to CoC@Mn1 and CoC@Fe1 , respectively, in line with the observed $\langle D_{\text{XRD}} \rangle$ and $\langle D_{\text{TEM}} \rangle$.

Effective anisotropy constants (K) were calculated in different ways, as reported in the “DC magnetometry” paragraph of ESI (Table 4S†) and, besides the computation method, in all



Table 2 Basic parameters determined from the ZFC-FC curves, magnetization isotherms and temperature dependence of χ' and χ'' : maximum ZFC temperature (T_{\max}), furcation point of the ZFC-FC curves (2% difference, T_{diff}), blocking temperature (T_b), coercive field at 10 K (H_c^{10}), anisotropy field at 10 K (H_K^{10}), saturation magnetization at 10 and 300 K (M_s^{10} , M_s^{300}), remnant magnetization at 10 K (M_r^{10}), magnetic moment (μ_m , median), Néel relaxation time at 300 K (τ_N). T_b has been calculated as the maximum of the energy barrier distribution ($-d(M_{\text{FC}} - M_{\text{ZFC}})/dT$). H_K has been calculated by considering 2% of the difference between the magnetization and demagnetization curves in the magnetization isotherm at 10 K

Sample	T_{\max} (K)	T_{diff} (K)	T_b (K)	H_c^{10} (T)	H_K^{10} (T)	M_s^{10} (A m ² kg ⁻¹)	M_r^{10} (A m ² kg ⁻¹)	M_r/M_s	M_s^{300} (A m ² kg ⁻¹)	μ_m (10 ³ μ_B)	τ_N (s)
CoA	195(3)	270(9)	126(2)	1.28(1)	4.2(1)	90(3)	53(2)	0.55	73(2)	2.6	4×10^{-7}
CoA@Mn	246(3)	275(3)	185(5)	0.92(1)	2.6(1)	97(4)	67(2)	0.72	75(2)	3.7	4×10^{-6}
CoA@Fe	294(1)	262(3)	199(3)	0.76(1)	2.0(1)	92(1)	72(2)	0.78	83(3)	12.5	6×10^{-5}
CoB	241(3)	266(3)	163(2)	1.32(2)	3.8(1)	90(4)	58(3)	0.62	74(3)	3.9	6×10^{-7}
CoB@Mn	314(3)	312(3)	233(2)	0.81(1)	2.3(1)	94(3)	74(3)	0.73	75(3)	8.3	7×10^{-4}
CoB@Fe	337(3)	333(3)	237(4)	1.02(1)	2.5(2)	94(3)	72(3)	0.76	81(2)	7.6	5×10^{-3}
CoC	274(3)	313(3)	206(2)	1.54(1)	4.1(1)	92(1)	67(1)	0.67	77(1)	4.4	4×10^{-6}
CoC@Mn1	292(2)	295(1)	216(1)	0.56(2)	2.5(1)	92(1)	66(2)	0.69	70(1)	10.8	5×10^{-5}
CoC@Mn2	348(3)	>380	251(5)	0.60(1)	1.9(1)	91(2)	67(1)	0.72	71(1)	14.5	1×10^{-3}
CoC@Fe1	278(5)	270(5)	190(1)	0.60(1)	1.8(1)	89(3)	71(2)	0.79	77(3)	15.8	7×10^{-6}
CoC@Fe2	352(4)	>380	246(4)	0.83(1)	2.6(1)	90(2)	71(1)	0.79	79(2)	14.7	4×10^{-3}

cases, core-shell systems feature lower anisotropy constant values than those of the respective cores.

Since both anisotropy constants and magnetic volume affect the Néel relaxation time (τ_N), AC magnetometry was used to determine the temperature dependence of the in-phase (χ') and out-of-phase (χ'') component of the magnetic susceptibility at different frequencies (0.1–1000 Hz) (Fig. 9S and 10S†) and τ_N at 300 K estimated by the Vogel-Fulcher equation (Fig. 11S and eqn (17S)†)⁶⁴ is reported in Tables 2 and 5S.† τ_N are equal to 4×10^{-7} s, 6×10^{-7} s, and 4×10^{-6} s for CoA, CoB, and CoC, respectively, in line with the increased particles' size of the samples. In the same way, τ_N of core-shell samples is slower than those of the respective cores, due to the increased magnetic volume that dominates the overall decrease of effective anisotropy, as already observed. Generally, spinel iron oxide coated samples feature slower τ_N with respect to the manganese ferrite coated ones, as well as the thicker shell samples compared to the thinner ones.

Based on the values of M_s^{300} , $\langle D_{\text{MAG}} \rangle$, K , and τ_N , better heating performances should be expected, in order: (i) for the largest core particles, (ii) for the core-shell samples with respect to the

corresponding cores, (iii) and for the spinel iron oxide-coated samples with respect to the manganese ferrite ones, (iv) for the samples having the thicker shell than the thinner ones.

In this context, the samples were tested for magnetic heat release in experimental conditions close to those used for clinical applications of magnetic fluid hyperthermia (MFH) in terms of solvent and magnetic field parameters: the measurements were performed on aqueous colloidal dispersions obtained by intercalation with cetyltrimethylammonium bromide (see "Intercalation process" paragraph in ESI, Fig. 12S†) keeping the amplitude and frequency of the alternate magnetic field below the clinical threshold.⁶⁵ The heating curves are reported in Fig. 5.

While the CoA sample did not heat up, the larger CoB and CoC NPs provided a sizeable heat release corresponding to SAR values of 21 ± 1 and 32 ± 3 W g_{ox}⁻¹, respectively (Table 3). In all core-shell systems, a remarkable increase in SAR with respect to the cores alone was observed with values ranging between 20 and 59 W g_{ox}⁻¹, depending on the chemical nature and thickness of the coating (Table 3).

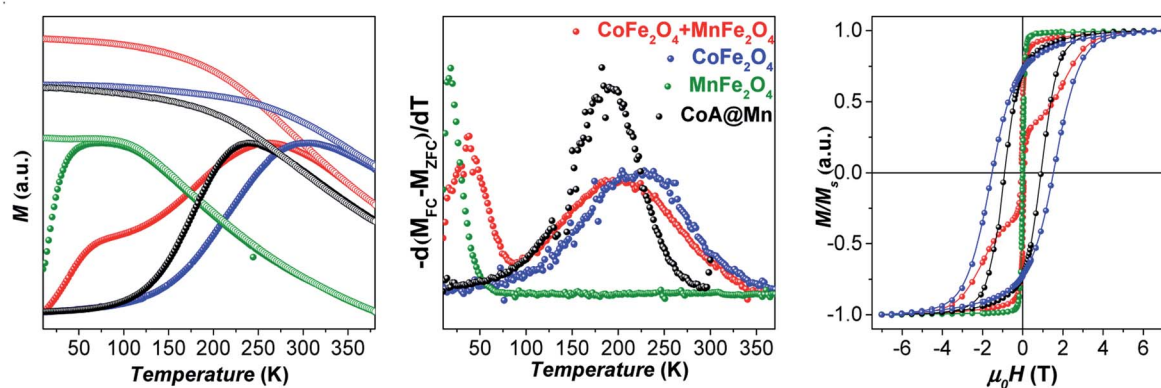


Fig. 4 ZFC (full circles) and FC (empty circles) curves, normalized for the magnetization at T_{\max} of the ZFC curve, recorded at a low external magnetic field (10 mT) of cobalt ferrite of ca. 8 nm, manganese ferrite of ca. 8 nm, a 1 : 1 w/w mixture of them, and CoA@Mn (left); anisotropy energy barrier distributions estimated by the first derivative $-d(M_{\text{FC}} - M_{\text{ZFC}})/dT$ (middle); magnetization isotherms recorded at 10 K (right).



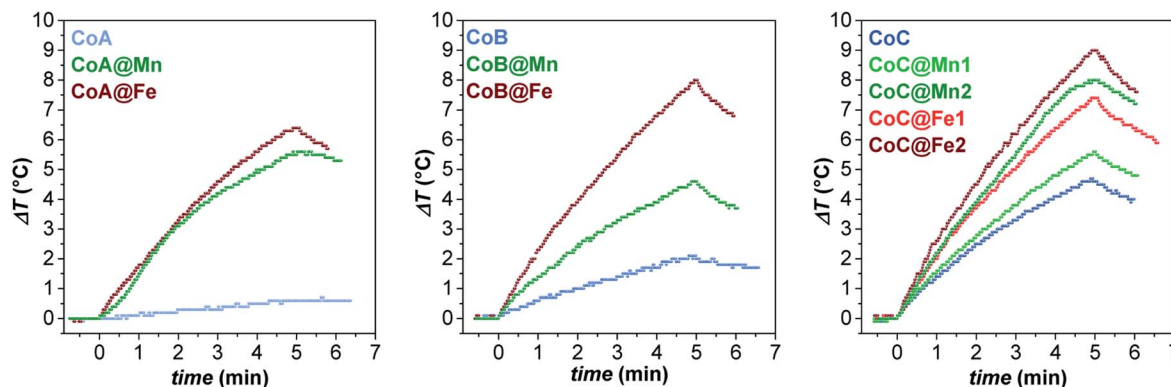


Fig. 5 Heating curves of the aqueous colloidal dispersions ($C_{\text{magn}} = 3.4 \text{ mg mL}^{-1}$) of all samples at $30 \text{ }^{\circ}\text{C}$, obtained under a magnetic field of 183 kHz and 17 kA m^{-1} .

Table 3 Specific absorption rate (SAR) and intrinsic loss power (ILP) values of the core–shell samples and respective core. SAR and ILP are given as the watt per gram of the spinel ferrite phase. * These values indicate a negligible heating release for the sample CoA

Sample	SAR ($\text{W g}_{\text{ox}}^{-1}$)	ILP ($\text{nH m}^2 \text{ kg}_{\text{ox}}^{-1}$)
CoA	0*	0*
CoA@Mn	20(1)	0.38(2)
CoA@Fe	42(2)	0.80(4)
CoB	21(1)	0.39(1)
CoB@Mn	27(2)	0.52(4)
CoB@Fe	48(1)	0.92(2)
CoC	32(2)	0.60(4)
CoC@Mn1	43(3)	0.81(6)
CoC@Mn2	47(2)	0.89(4)
CoC@Fe1	46(4)	0.88(8)
CoC@Fe2	59(2)	1.12(4)

The experimental hyperthermic data collected on these three different core–shell series revealed some general aspects:

- (1) Core–shell samples heated up more than respective cores;
- (2) In all sets, spinel iron oxide-coated core–shell NPs performed better than manganese ferrite-coated ones;
- (3) In the CoC series, both systems (CoC@Fe1/CoC@Fe2 and CoC@Mn1/CoC@Mn2) showed that an increase in the shell thickness induces an improvement in the heating abilities.

These results are totally consistent with the previous hypothesis based on the size and magnetic parameters of powdered samples. Nevertheless, to get close to the experimental conditions in which the heating abilities were evaluated, a set of AC magnetic measurements was carried out on the hydrophilic ferrofluids of the CoB series (Fig. 13S†). The temperature dependences of the in-phase and out-of-phase susceptibilities (χ' , χ'') showed a single maximum ($T_{\text{max}}^{\text{AC}}$): below the melting point for the CoB sample and above it for the core–shell ones. A noticeable asymmetry in the curves is visible for all samples indicating different behaviour below and above the melting points, making the estimation of Néel relaxation times more complicated. Indeed, when the solution melts, Brownian motions may occur and also the interparticle

interactions may change. Moreover, the $T_{\text{max}}^{\text{AC}}$ values of the ferrofluids result to be shifted towards higher temperatures with respect to those of the powdered samples. This behaviour is also observed in the ZFC-FC curves of the CoB dispersion if compared with the CoB powder (Fig. 14S†), in which other features are also visible: a decrease in the broadening of the ZFC peak and an enhanced flatness of the FC curve at temperatures below the T_{max} . Both the T_{max} shift and the enhancement in the flatness of the FC curve are hints of stronger interparticle interactions, which are independent of the ferrofluid's concentration (Fig. 14S†). These findings, even though appear awkward, suggest renormalization of interparticle interactions in the fluid in comparison with the powder and therefore the occurrence of agglomeration/aggregation phenomena which are independent of the concentration. Dipolar interactions most likely occur and cause the formation of secondary entities, whose size (number of primary NPs) and shape (random or controlled clustering such as chain-like alignment) affect the resulting magnetic behaviour and heating ability under an applied static or dynamic magnetic field. In the literature, specific studies on the effect on the heating abilities of the interparticle interactions for spinel ferrite-based core–shell nanoparticles are not available, probably due to the complexity of the systems in which many parameters may affect the magnetic and hyperthermic properties. Concerning single-phase systems, some experimental studies reported the enhancement or reduction of SAR as a function of the dipolar interactions,^{66–72} but other authors⁷³ provided a general theoretical model able to explain the heating release behaviour of NPs in the blocked-state, based on their intrinsic magnetic properties (anisotropy, magnetization) and experimental conditions (concentration and magnetic field amplitude). In our case, no changes occur in the strength of dipolar interactions, in the dispersions, in the concentration range $0.7\text{--}3.4 \text{ mg mL}^{-1}$ (Fig. 14S†). Moreover, also SAR values are independent of interparticle interactions, as reported in Fig. 15S and Table 6S† for a cobalt ferrite sample measured at different concentrations and different capping molecules. Unfortunately, besides this concentration and capping agent independences, it is not possible to speculate on the differences of magnetic properties



between powder and colloidal dispersion, the mechanism of formation of secondary entities, and their role in the magnetic properties and in the heating efficiency. Indeed, it has to be taken into account that ferrofluids are dynamic hybrid organic–inorganic systems in which the capping agents might be involved in different processes occurring in liquid phase and feature different physical properties (e.g. hydrophobicity for oleate and hydrophilicity for CTAB molecules). For all the above reasons, the evaluation of magnetic parameters from ferrofluids appears to be complicated and not strictly reliable, while the previous discussion of M_s^{300} , $\langle D_{MAG} \rangle$, K , and τ_N , extracted from measurements on powdered samples, although being a simplification of the system under study, helped in understanding the effect on the heat release performances of the magnetic features of the sole inorganic counterpart, making negligible the influences of the liquid-phase processes.

On the one hand, all the above findings seem to depict a more complex scenario behind the heating abilities of bi-magnetic core–shell systems than simple relationships with single magnetic or microstructural parameter(s). On the other hand, they suggest that a DC/AC magnetic characterization can be helpful in driving the engineering of the heat mediators and they also proved the enhancement of the heating abilities of these heterostructures, especially with a shell of spinel iron oxide, with respect to their corresponding cores in water colloidal dispersions, in principle making these systems biocompatible and promising for application in magnetic heat generation.

Conclusions

A seed-mediated growth strategy in a solvothermal condition made available three sets of bi-magnetic spinel ferrite core–shell NPs with low size dispersity, allowing the tuning of the core size, chemical nature of the shell, and shell thickness. The effects of these three parameters on the heating abilities were studied for the first time together, revealing unforeseen results on the shell influence. Direct proof of the core–shell structure formation was successfully provided by chemical mapping at the nanoscale using STEM-EDX. The chemical composition, cation distribution, and spin canting were also studied through ICP-AES and low-temperature ^{57}Fe Mössbauer spectroscopy, leaving out these features from their influence on the magnetic properties. DC magnetometry demonstrated the rigid coupling between the hard and the soft phases and provided, in combination with the AC magnetometry, the main magnetic parameters theoretically responsible for the heat generation. The availability of several samples allowed the identification of suitable comparisons from which a single parameter-effect emerged.

The heating abilities of the aqueous colloidal dispersions of all samples were tested under mild experimental conditions. For all sets of samples, spinel iron oxide shells featured higher heat release than those of manganese ferrite ones. Moreover, for the first time for hard core-based core–shell NPs, it was observed that the thicker the soft shell, the better the performances. These results were justified in agreement with the

hypothesized behaviour based on the magnetic properties, such as saturation magnetization at 300 K, magnetic volume, anisotropy and Néel relaxation time.

The study thus demonstrated the importance of a sophisticated approach based on the synergy of chemical, structural, and magnetic probes down to a single-particle level. Finally, considering the biocompatibility of the iron oxides, this systematic fundamental study proved that a proper design of cobalt ferrite cores coated with a homogenous crystalline shell of spinel iron oxide in principle should lead to biocompatible heat mediators with a net improvement in the heating abilities in comparison with the corresponding cores.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The University of Cagliari and V-Fase Project - RAS Piano Sulcis are acknowledged for the Ph.D. and PostDoc grants, respectively, of Marco Sanna Angotzi. Marco Sanna Angotzi also acknowledges CESA Project-RAS Piano Sulcis, which finances his current postdoctoral position. CESA Project-RAS Piano Sulcis and PON AIM (PON Ricerca e Innovazione 2014–2020-Azione I.2-D.D. n.407 del 27 febbraio 2018 “Attraction and International Mobility”, CULT-GEOCHIM Project, AIM1890410-3) are gratefully acknowledged for financing the fellowship of Valentina Mameli. This research used resources of the Center for Functional Nanomaterials, which is a U.S. DOE Office of Science Facility, at Brookhaven National Laboratory under Contract No. DE-SC0012704. Claudio Sangregorio thanks INFN HADROCOMBI and HADROMAG projects for partly funding the work. Jana Vejpravova acknowledges support of EC (ERC Starting grant TSuNAMI, no. 716265) and Materials Growth and Measurement Laboratory financed by the MEYS (LM2018096). The financial support of Fondazione Banco di Sardegna and Regione Autonoma della Sardegna Progetti Biennali di Ateneo Annualità 2016, Fondazione Sardegna CUP F72F16003070002 is gratefully acknowledged.

Notes and references

- 1 Y. Jun, J. Choi and J. Cheon, *Chem. Commun.*, 2007, 1203–1214.
- 2 R. G. Chaudhuri and S. Paria, *Chem. Rev.*, 2012, **112**, 2373–2433.
- 3 L. Carbone and P. D. Cozzoli, *Nano Today*, 2010, **5**, 449–493.
- 4 P. Mélinon, S. Begin-Colin, J. L. Duvail, F. Gauffre, N. H. Boime, G. Ledoux, J. Plain, P. Reiss, F. Silly and B. Warot-Fonrose, *Phys. Rep.*, 2014, **543**, 163–197.
- 5 R. L. Stamps, *J. Phys. D*, 2000, **3**, 444.
- 6 X. Sun, N. Frey Huls, A. Sigdel and S. Sun, *Nano Lett.*, 2012, **12**, 246–251.
- 7 J. Nogués, J. Sort, V. Langlais, V. Skumryev, S. Suriñach, J. S. Muñoz and M. D. Baró, *Phys. Rep.*, 2005, **422**, 65–117.



- 8 N. F. Troitino, B. Rivas-Murias, B. Rodríguez-González and V. Salgueiriño, *Chem. Mater.*, 2014, **26**, 5566–5575.
- 9 V. Mameli, A. Musinu, D. Niznansky, D. Peddis, G. Ennas, A. Ardu, C. Lugliè and C. Cannas, *J. Phys. Chem. C*, 2016, **120**, 27635–27645.
- 10 E. F. Kneller and R. Hawig, *IEEE Trans. Magn.*, 1991, **27**, 3560–3588.
- 11 A. López-Ortega, M. Estrader, G. Salazar-Alvarez, A. G. Roca and J. Nogués, *Phys. Rep.*, 2015, **553**, 1–32.
- 12 Virginia Commonwealth University, *US Pat.*, 2015 0001437A1, 2015.
- 13 V. G. Harris, *US Pat.*, 2012O16867OA1, 2012.
- 14 Industry-University Cooperation Foundation, *US Pat.*, 20140286817A1, Hanyang University Erica Campus, LG Electronics INC., 2014.
- 15 Fujifilm Corporation, *US Pat.*, 20110027588A1, Tohoku University, 2011.
- 16 S. Liu, *US Pat.*, 2006OOO5898A1, 2006.
- 17 BASF Aktiengesellschaft, *US Pat.*, 4770903, 1988.
- 18 University of Louisiana at Lafayette, USOO7964O13B2, 2011.
- 19 A. Kirschning, L. Kupracz and J. Hartwig, *Chem. Lett.*, 2012, **41**, 562–570.
- 20 R. B. N. Baig and R. S. Varma, *Chem. Commun.*, 2013, **49**, 752–770.
- 21 V. Polshettiwar, R. Luque, A. Fihri, H. Zhu, M. Bouhrara and J. M. Basset, *Chem. Rev.*, 2011, **111**, 3036–3075.
- 22 D. Zhang, C. Zhou, Z. Sun, L.-Z. Wu, C.-H. Tung and T. Zhang, *Nanoscale*, 2012, **4**, 6244.
- 23 S. L. Viñas, K. Simeonidis, Z. A. Li, Z. Ma, E. Myrovali, A. Makridis, D. Sakellari, M. Angelakeris, U. Wiedwald, M. Spasova and M. Farle, *J. Magn. Magn. Mater.*, 2016, **415**, 20–23.
- 24 H. Choi, M. An, W. Eom, S. W. Lim, I. B. Shim, C. S. Kim and S. J. Kim, *J. Korean Phys. Soc.*, 2017, **70**, 173–176.
- 25 M. S. Kim, C. S. Kim, H. J. Kim, K.-H. H. Yoo and E. J. Hahn, *J. Korean Phys. Soc.*, 2013, **63**, 2175–2178.
- 26 S. Noh, W. Na, J. Jang, J.-H. Lee, E. J. Lee, S. H. Moon, Y. Lim, J.-S. Shin and J. Cheon, *Nano Lett.*, 2012, **12**, 3716–3721.
- 27 J. Wang, Z. Zhou, L. Wang, J. Wei, H. Yang, S. Yang and J. Zhao, *RSC Adv.*, 2015, **5**, 7349–7355.
- 28 V. Pilati, R. Cabreira Gomes, G. Gomide, P. Coppola, F. G. Silva, F. L. O. Paula, R. Perzynski, G. F. Goya, R. Aquino and J. Depeyrot, *J. Phys. Chem. C*, 2018, **122**, 3028–3038.
- 29 L. Wang, Y. Yan, M. Wang, H. Yang, Z. Zhou, C. Peng and S. Yang, *J. Mater. Chem. B*, 2016, **4**, 1908–1914.
- 30 S. O. Solopan, N. Nedelko, S. Lewińska, A. Ślowska-Waniewska, V. O. Zamorskyi, A. I. Tovstolytkin and A. G. Belous, *J. Alloys Compd.*, 2019, **788**, 1203–1210.
- 31 F. Fabris, E. Lima, E. De Biasi, H. E. Troiani, M. Vásquez Mansilla, T. E. Torres, R. Fernández Pacheco, M. R. Ibarra, G. F. Goya, R. D. Zysler and E. L. Winkler, *Nanoscale*, 2019, **11**, 3164–3172.
- 32 M. Angelakeris, Z.-A. A. Li, M. Hilgendorff, K. Simeonidis, D. Sakellari, M. Filippousi, H. Tian, G. Van Tendeloo, M. Spasova, M. Acet and M. Farle, *J. Magn. Magn. Mater.*, 2015, **381**, 179–187.
- 33 J.-H. Lee, J.-T. Jang, J.-S. Choi, S. H. Moon, S.-H. Noh, J.-W. Kim, J.-G. Kim, I.-S. Kim, K. I. Park and J. Cheon, *Nat. Nanotechnol.*, 2011, **6**, 418–422.
- 34 S. Liébana-Viñas, K. Simeonidis, U. Wiedwald, Z.-A. Li, Z. Ma, E. Myrovali, A. Makridis, D. Sakellari, G. Vourlias, M. Spasova, M. Farle and M. Angelakeris, *RSC Adv.*, 2016, **6**, 72918–72925.
- 35 Q. Zhang, I. Castellanos-Rubio, R. Munshi, I. Orue, B. Pelaz, K. I. Gries, W. J. Parak, P. Del Pino and A. Pralle, *Chem. Mater.*, 2015, **27**, 7380–7387.
- 36 J. Robles, R. Das, M. Glassell, M. H. Phan and H. Srikanth, *AIP Adv.*, 2018, **8**, 2–8.
- 37 M. Hammad, V. Nica and R. Hempelmann, *IEEE Trans. Magn.*, 2017, **53**, 1–6.
- 38 O. V. Yelenich, S. O. Solopan, J. M. Greneche and A. G. Belous, *Solid State Sci.*, 2015, **46**, 19–26.
- 39 S. Laurent, D. Forge, M. Port, A. Roch, C. Robic, L. Vander Elst and R. N. Muller, *Chem. Rev.*, 2008, **108**, 2064–2110.
- 40 L. H. Reddy, J. L. Arias, J. Nicolas and P. Couvreur, *Chem. Rev.*, 2012, **112**, 5818–5878.
- 41 C. Cara, A. Musinu, V. Mameli, A. Ardu, D. Niznansky, J. Bursik, M. A. Scorciapino, G. Manzo and C. Cannas, *Cryst. Growth Des.*, 2015, **15**, 2364–2372.
- 42 C. Cannas, A. Ardu, A. Musinu, L. Suber, G. Ciasca, H. Amenitsch and G. Campi, *ACS Nano*, 2015, **9**, 7277–7286.
- 43 M. Sanna Angotzi, A. Musinu, V. Mameli, A. Ardu, C. Cara, D. Niznansky, H. L. Xin and C. Cannas, *ACS Nano*, 2017, **11**, 7889–7900.
- 44 E. E. Fullerton, J. Jiang and S. Bader, *J. Magn. Magn. Mater.*, 1999, **200**, 392–404.
- 45 V. Mameli, A. Musinu, A. Ardu, G. Ennas, D. Peddis, D. Niznansky, C. Sangregorio, C. Innocenti, N. T. K. Thanh and C. Cannas, *Nanoscale*, 2016, **8**, 10124–10137.
- 46 B. Pacakova, S. Kubickova, G. Salas, A. R. Mantlikova, M. Marciello, M. P. Morales, D. Niznansky and J. Vejpravova, *Nanoscale*, 2017, **9**, 5129–5140.
- 47 C. Pereira, A. M. Pereira, C. Fernandes, M. Rocha, R. Mendes, M. P. Fernández-García, A. Guedes, P. B. Tavares, J. M. Grenèche, J. P. Araújo and C. Freire, *Chem. Mater.*, 2012, **24**, 1496–1504.
- 48 J.-M. Greneche, *Hyperfine Interact.*, 2003, **148/149**, 79–89.
- 49 J. Rodríguez-Carvajal and T. Roisnel, *Int. Union Crystallogr.*, 1998, **20**, 35–36.
- 50 S. Mondini, a. M. Ferretti, a. Puglisi and a. Ponti, *Nanoscale*, 2012, **4**, 5356.
- 51 V. Mameli, M. S. Angotzi, C. Cara and C. Cannas, *J. Nanosci. Nanotechnol.*, 2019, **19**, 4857–4887.
- 52 M. S. Angotzi, V. Mameli, A. Musinu and D. Nizňanský, *J. Nanosci. Nanotechnol.*, 2019, **19**, 5008–5013.
- 53 M. Sanna Angotzi, V. Mameli, C. Cara, A. Ardu, D. Nizňanský and A. Musinu, *J. Nanosci. Nanotechnol.*, 2019, **19**, 4954–4963.
- 54 D. Peddis, N. Yaacoub, M. Ferretti, A. Martinelli, G. Piccaluga, A. Musinu, C. Cannas, G. Navarra, J. M. Greneche and D. Fiorani, *J. Phys. Condens. Matter*, 2011, **23**, 426004.



- 55 E. Fantechi, G. Campo, D. Carta, A. Corrias, C. de Julián Fernández, D. Gatteschi, C. Innocenti, F. Pineider, F. Ruggi and C. Sangregorio, *J. Phys. Chem. C*, 2012, **116**, 8261–8270.
- 56 D. Carta, M. F. Casula, A. Falqui, D. Loche, G. Mountjoy, C. Sangregorio and A. Corrias, *J. Phys. Chem. C*, 2009, **113**, 8606–8615.
- 57 V. Blanco-Gutiérrez, J. A. Gallastegui, P. Bonville, M. J. Torralvo-Fernández and R. Sáez-Puche, *J. Phys. Chem. C*, 2012, **116**, 24331–24339.
- 58 A. Repko, J. Vejpravová, T. Vacková, D. Zákutná and D. Nižňanský, *J. Magn. Magn. Mater.*, 2015, **390**, 142–151.
- 59 K. Vamvakidis, M. Katsikini, D. Sakellari, E. C. Paloura, O. Kalogirou and C. Dendrinou-Samara, *Dalton Trans.*, 2014, **43**, 12754–12765.
- 60 J. M. D. Coey, *Magnetism and magnetic materials*, Cambridge University Press, Cambridge, 2009.
- 61 G. Muscas, G. Concas, S. Laureti, A. M. Testa, R. Mathieu, J. A. De Toro, C. Cannas, A. Musinu, M. A. Novak, C. Sangregorio, S. S. Lee and D. Peddis, *Phys. Chem. Chem. Phys.*, 2018, **20**, 28634–28643.
- 62 P. de la Presa, Y. Luengo, M. Multigner, R. Costo, M. P. Morales, G. Rivero and A. Hernando, *J. Phys. Chem. C*, 2012, **116**, 25602–25610.
- 63 R. E. Rosensweig, *J. Magn. Magn. Mater.*, 2002, **252**, 370–374.
- 64 B. Aslibeiki, P. Kameli, H. Salamati, G. Concas, M. Salvador Fernandez, A. Talone, G. Muscas and D. Peddis, *Beilstein J. Nanotechnol.*, 2019, **10**, 856–865.
- 65 R. Hergt, S. Dutz and M. Zeisberger, *Nanotechnology*, 2010, **21**, 015706.
- 66 D. Niculaes, A. Lak, G. C. Anyfantis, S. Marras, O. Laslett, S. K. Avugadda, M. Cassani, D. Serantes, O. Hovorka, R. Chantrell and T. Pellegrino, *ACS Nano*, 2017, **11**, 12121–12133.
- 67 L. L. Gutiérrez, L. de la Cueva, M. Moros, E. Mazarío, S. de Bernardo, J. M. de la Fuente, M. P. Morales and G. Salas, *Nanotechnology*, 2019, **30**, 112001.
- 68 D. Serantes, K. Simeonidis, M. Angelakeris, O. Chubykalo-Fesenko, M. Marciello, M. Del Puerto Morales, D. Baldomir and C. Martinez-Boubeta, *J. Phys. Chem. C*, 2014, **118**, 5927–5934.
- 69 P. De La Presa, Y. Luengo, V. Velasco, M. P. Morales, M. Iglesias, S. Veintemillas-Verdaguer, P. Crespo and A. Hernando, *J. Phys. Chem. C*, 2015, **119**, 11022–11030.
- 70 C. Martinez-Boubeta, K. Simeonidis, A. Makridis, M. Angelakeris, O. Iglesias, P. Guardia, A. Cabot, L. Yedra, S. Estradé, F. Peiró, Z. Saggi, P. A. Midgley, I. Conde-Leborán, D. Serantes and D. Baldomir, *Sci. Rep.*, 2013, **3**, 1–8.
- 71 C. Guibert, V. Dupuis, V. Peyre and J. Fresnais, *J. Phys. Chem. C*, 2015, **119**, 28148–28154.
- 72 D. F. Coral, P. Mendoza Zélis, M. Marciello, M. D. P. Morales, A. Craievich, F. H. Sánchez and M. B. Fernández Van Raap, *Langmuir*, 2016, **32**, 1201–1213.
- 73 I. Conde-Leboran, D. Baldomir, C. Martinez-Boubeta, O. Chubykalo-Fesenko, M. Del Puerto Morales, G. Salas, D. Cabrera, J. Camarero, F. J. Teran and D. Serantes, *J. Phys. Chem. C*, 2015, **119**, 15698–15706.

