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Topical Review

Progress and prospects of hard hexaferrites for permanent magnet applications

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Abstract

Permanent magnets based on hard hexaferrite represent the largest family of magnets being used today by volume. They generate moderate remanence induction, but present crucial advantages in terms of availability, cost, resistance to demagnetization and corrosion and absence of eddy current losses. As a consequence, ferrites are the most logical candidate for substitution of rare-earths in selected applications that do not demand the best performing magnets. If the remanence of ferrite-based magnets was to be improved, even mildly, the door to a larger scale substitution could be opened. In this framework, we review here current strategies to improve the properties of hexaferrites for permanent magnet applications. We first discuss the potential of exploring the nanoscale. Second, progress related to controllably doping hexaferrites is revised. Third, results achieved by fabricating hard-soft magnetic composites using ferrites as the hard phase are presented. Finally, future prospects and new potential end applications for ferrite magnets are discussed.

Keywords: permanent magnets, hexaferrites, review, rare-earth substitution, doping, exchange-coupling

(Some figures may appear in colour only in the online journal)

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1. Introduction

1.1. The importance of hexaferrite magnets

Permanent magnets, in the form of lodestone [1], and later of magnetized steel, have been used by humankind for several millennia. During most of history, their practical use was limited to the compass, and they remained not much more than a kind of magician trick for most people. It is only with the industrial revolution that magnets began to be used on a large scale. Nowadays, permanent magnet materials are employed in many devices, specially in electrical to mechanical (and the converse mechanical to electrical) transducer devices, for information storage and in sensors. Since the beginning of the 20th century materials have been developed with the specific goal to improve those historic magnetic materials [2], thanks to the improved knowledge of material science and the parallel technological evolution. This path [3] provided a series of metallic alloys culminating in the Alnico family discovered by Mishima in the 30s, continuing with the hard ferrites developed by Phillips in the 50s, and reaching the SmCo and NdFeB rare-earth magnets developed in the 70s and 80s, respectively. This evolution has been accompanied by a corresponding increase in the tuning of the properties required for a hard magnet, specially the magnetic anisotropy.

Nowadays, the current market of ferrite magnets is dominated by the hard ferrites: the M-type hexaferrites $\text{BaFe}_{12}\text{O}_{19}$ (BFO), $\text{SrFe}_{12}\text{O}_{19}$ (SFO) and related materials [4]. Thus, the story of industrial age use of ferrites can be considered to start with the research of Takeshi Takei in Japan in the first half of the 20th century, in collaboration with Professor Kato [5]. While BFO was already known in the first part of the 20th century, it was only after the second World War that at Philips Laboratories, under the direction of Snoek, it was shown to have the same structure of magnetoplumbite ($\text{PbFe}_{12}\text{O}_{19}$). Phillips moved on to develop a series of ferrites with the magnetoplumbite structure. During the 1950s all the main hexaferrite phases were studied, as shown in the classic Smit and Wijn's book [4]. It was in 1952 when the barium hexaferrite material was commercialized.

Ferrites exhibit relatively large coercivity (H_C) up to 380 kA m^{-1} , but modest saturation magnetization (M_S or magnetization induction $B_S = \mu_0 M_S$), specific saturation (σ_S) and remanence (M_R or remanence induction, $B_R = \mu_0 M_R$). A useful parameter to empirically establish the suitability of a given material to realize compact permanent magnet with no shape constraints is the magnetic hardness, κ . This is defined by the ratio of anisotropy, K_1 , to magnetostatic energy, $\kappa = (K_1/m_0 M_S^2)^{1/2}$, and it is assumed that $\kappa > 1$ is the criterion for a material to be classified as hard. M-type hexaferrite have hardness higher than 1 (for example for BFO $\kappa = 1.34$), which explains why, despite the relatively low M_S and moderate anisotropy, they are largely used to realize permanent magnets. The highest grade ferrite magnets in the market (Hitachi Metals NMF-15 series) reach maximum values of remanence induction of 480 mT. The highest grade ferrite magnets in the market (Hitachi Metals NMF-15 series) reach maximum values of remanence induction of 480 mT.

Their maximal energy product (BH_{max}), figure of merit for the strength of magnets, reaches 44 kJ m^{-3} . These values are low when compared to that of rare-earth based materials, such as Nd-Fe-B, who typically present (Hitachi Metals NEOMAX Series) remanence inductions of 1.3 T, coercivities of 1500 kA m^{-1} ($\mu_0 H_C = 1.85 \text{ T}$) and energy products of 350 kJ m^{-3} [6].

However, ferrites do have some important advantages. The main one is definitely the availability and low cost of raw material powders, of approximately $1.5 \text{ \$ kg}^{-1}$, while NdFeB powders approximately cost $60 \text{ \$ kg}^{-1}$. They also exhibit excellent chemical stability and high electrical resistivity. For these reasons, ferrite magnets are widely used in mass applications that do not require large energy products and/or where increasing the weight of the device is not an issue—since the low flux density can be compensated by increasing the volume of magnetic material. These include inductors, transformers and electromagnets due to the low eddy current losses they entail. As a consequence of these advantages, ferrite magnets constitute by far the most widely used permanent magnet by weight [7].

Hexaferrites are the subject of a continuing research effort, both from fundamental and applied perspectives [8, 9]. In the last two decades the research on ferrites has been, however, mainly focused on ferrite nanomaterials. The systematic research devoted to ferrite nanoparticles lead to novel applications, ranging from engineering (e.g. magnetic recording media, magnetic seals, ferrofluids, magnetic separation, etc) [10–18] to medicine (e.g. magnetic hyperthermia, magnetic resonance imaging, drug delivery, etc) [19–28]. Based on ferrite nanoparticles new attractive magneto-responsive materials were developed, including ferromagnetic fluids (i.e. liquids exhibiting a spontaneous magnetic ordering, in contrast to superparamagnetic (SPM) ferrofluids) [29–32], magneto-optical suspensions and solid composites [33–35], soft magneto-electric [36], if we mention just a few. The acquired knowledge related to nanomagnetism, synthesis, and processing of the nanoparticles can also be crucial for the renaissance of most traditional applications of ferrites, i.e. for permanent magnets.

1.2. Structure and properties

The family of hard-ferrites is based on the magnetoplumbite structure, $\text{PbFe}_{12}\text{O}_{19}$, which belongs to the hexagonal space group $P6_3/mmc$. Thus, four indices are usually employed to indicate the planes, with the last one indicating the c -axis and the first three indicating the in-plane orientation of the unit vectors at 120° from each other. The crystal structure [37] is based on a close-packed arrangement of oxygen cations with the divalent (Ba^{2+} , Sr^{2+} or Pb^{2+}), and Fe^{3+} cations occupying the voids. A way to visualize the structure is to consider it to be built from two different building blocks. The first is the two layer S (for spinel) block. The S block presents the same occupancy of tetrahedral and octahedral sites as the spinel crystal structure, and thus it has the $\text{Fe}_6\text{O}_8^{2+}$ composition. The other block, R, is built from a three-layer hcp sequence, with

Figure 1. Schematic of the M-hexaferrite structure of BFO. Polyhedrals are colored: dodecahedral Ba^{2+} (green), and the Fe^{3+} sites: octahedral 2a (dark red), tetrahedral $4f_1$ (medium blue), bipyramidal 2b (dark blue), distorted octahedral $4f_2$ (gray), octahedral 12k (light red). The spin direction is parallel to the net direction for cations inside polyhedral colored red, and opposite for atoms inside polyhedral colored blue-gray (drawing made with VESTA [41]).

a quarter of the oxygen anions substituted in the intermediate hexagonal layer by the large divalent cation (Ba^{2+} , Sr^{2+}). Considering the interstitial Fe atoms, the composition of this block is $\text{BaFe}_6\text{O}_{11}^{2-}$. The full unit cell is obtained by ordering the blocks along the c direction, following the sequence RSR^*S^* , where the star indicates a rotation of 180° , giving a repeat distance of 2.2–2.3 nm. The in-plane unit cell constant is 0.56 nm, and corresponds to about four times the radius of the oxygen anion [38–40]. The iron cations thus occupy five unique interstitial sites, indicated by the Wyckoff positions 2a, 2b, $4f_1$, $4f_2$ and 12k. The five-fold coordinated (2b) has been found to correspond to two statistically half-occupied pseudo tetragonal 4 sites [37]. The other sites are the tetrahedral and octahedral sites of the S block ($4f_1$ and 2a), distorted octahedral sites in the R block ($4f_2$) and distorted octahedral sites at the boundary between R and S blocks (which we assign to the R block). A drawing is shown in figure 1.

The magnetic structure of the M-hexaferrites can be described by the Néel model of antiferromagnetism. The spin orientation of each iron cation is coupled by superexchange interaction through the oxygen anions. The Fe^{3+} cations have a spin-only magnetic moment. In the S block, the octahedral sites point in the net magnetization direction (i.e. the spin up direction), while the tetrahedral ones point in the opposite direction (spin down). Two octahedral sites of the R-block can be better described as distorted tetrahedral, and thus point in the

spin down. The bipyramidal site is along with the three octahedral ones, giving a net of another two magnetic moments. Considering the magnetic moment at each temperature of each site x , m_x , and the site occupancy (n_x), the magnetic moment per cell is:

$$m_c = n_{12k}m_{12k} + n_{2a}m_{2a} + n_{2b}m_{2b} - (n_{4f1}m_{4f1} + n_{4f2}m_{4f2})$$

Considering the occupancy of the M-type ferrites ($n_{12k} = 6$, $n_{2a} = 1$, $n_{2b} = 1$, $n_{4f1} = 2$, $n_{4f2} = 2$) and a spin-only magnetic moment at 0 K of Fe^{3+} ions of $5 \mu_B$, a magnetic moment per formula unit of $20 \mu_B$ is expected, which is close to the experimental value measured at 77 K ($19.9 \mu_B$, $100 \text{ A m}^2 \text{ kg}^{-1}$) [42].

The saturation magnetization at room temperature of the M-hexaferrites is not too low if compared to a ferrimagnetic oxide (for example magnetite, which is a high-saturation specific magnetization oxide and has $\sigma_s \sim 90\text{--}95 \text{ A m}^2 \text{ kg}^{-1}$), but it is very low when compared to the saturation magnetization of most materials used for permanent magnets. The reason for the use of hexaferrites actually lies in their large uniaxial magnetocrystalline anisotropy field (1.8 T), which yields large coercive fields. As a consequence, the coercive field can be larger than the remanence induction, which leads to robust resistance to demagnetization and large energy products. In this case the working point of the ferrite magnets depends weakly on their morphology.

The origin of the high magnetocrystalline anisotropy relies on the strong contribution to the single-ion term from Fe^{3+} ions placed in the trigonal bipyramidal 2b cavities, and to a minor extent in tetrahedral and octahedral distorted sites. On the other hand, the hexagonal structure makes the anisotropy to lie along the c -axis. The magnetocrystalline anisotropy is indeed well described, even at low temperature, as an uniaxial type with a second order term K_1 anisotropy constant. K_1 takes at room temperature a value of $3.5 \times 10^5 \text{ J m}^{-3}$ for SFO and $3.25 \times 10^5 \text{ J m}^{-3}$ for BFO [42]. Considering that magnetocrystalline anisotropy is uniaxial, the anisotropy field, H_A which is equal to $2K_1/\mu_0 M_s$ is 1.35 MA m^{-1} (1.7 T) and 1.43 MA m^{-1} (1.8 T), for BFO and SFO, respectively [4, 42].

A common problem when considering the feasibility of a new material for potential applications as permanent magnet material arises from the way its properties are often described in the literature. For example, the measurement of the magnetization saturation can be performed under different maximum applied magnetic fields, often at values that are not high enough to fully saturate the material or to overcome the irreversibility field (minor loop). In most of the studies, the changes of the magnetization are reported in terms of the specific magnetization, σ in $\text{A m}^2 \text{ kg}^{-1}$ or emu g^{-1} , since this quantity is not affected by powder packing or porosity in sintered samples. However for evaluating the performances of a ferrite as a magnet the proper magnitude is the magnetization (M_s in A m^{-1}) or the saturation induction (B_s in T). σ_s is preferable because uncertainty for mass is much lower than for volume measurement required to determine M_s . On the other hand, there is also the uncertainty that the change of σ_s can be due to modification of the M_s and/or the mass density.

Another key issue to be considered is how to determine the changes in H_A and of the symmetry of anisotropy with respect to the bulk. The measurement of these properties requires Singular Point detection technique [43, 44] in powders, nuclear magnetic resonances or ferromagnetic resonance experiments [42, 45]. However, often published works rely on the ‘approximation law to the saturation’ to the magnetization curves to estimate the anisotropy, a procedure that can present problems [46] to investigate nanomaterials and hybrid composites.

1.3. The potential and the challenges to improve ferrite magnets

Rare-earth elements are critical raw materials that present supply risk, price volatility and environmental problems. For these reasons, considerable efforts are being dedicated to reduce our dependence on them, which directly affect the permanent magnet value chain. The strategies include improving the recyclability of rare-earth materials and devices, fabricating competitive devices that are rare-earth-free and finding alternative magnets that could replace NdFeB and SmCo magnets. As the most widely used magnet by volume in the world, ferrites constitute an interesting candidate to attempt substitution with. However, to enable a wider replacement in more applications, the magnetic properties of ferrites need to be improved. Of course this substitution is generally realistic in applications where magnets with moderate $(BH)_{\max}$ ($50\text{--}200\text{ kJ m}^{-3}$) are needed, but it is important to remark that substitution by ferrites can be attempted in important applications that are expected to drive the permanent magnet market in the next few years and increase the NdFeB demand. Generators for wind turbines and automotive applications fall within this category, and the development of competitive devices based on high-grade/improved ferrites could entail large scale substitution. In light of this need, the importance of ferrite based magnets is being reinvigorated.

The $(BH)_{\max}$ value of hexaferrites in sintered ceramics is around 40 kJ m^{-3} . Most of the H_C values used to estimate $(BH)_{\max}$ have been frozen for more than 40 years. Considering that the large magnetic anisotropy field of the BFO and SFO, 1.7 T and 1.8 T, respectively is much larger than H_C and B_S , both sintered and bonded commercial magnets have coercive fields similar to the remanence induction and near of the saturation induction (0.4 T). In such a case, the $(BH)_{\max}$ is near the maximum achievable value, B_S^2/μ_0 . Hence, the first strategy to increase the $(BH)_{\max}$ requires the improvement of the saturation induction. However, for permanent magnets it should be considered that H_C should be larger than $N_{\text{ef}}M_S$ to prevent demagnetization, where N_{ef} represents the demagnetizing factor due to the shape of the magnet. Thus, a second target is to improve the coercive field to keep $H_C > N_{\text{ef}}M_S$. Because the H_C is proportional to the anisotropy field and hence to K_1/μ_0M_S , the targets of improvement both H_C and M_S by improving a single phase material require that a further increment of the magnetic anisotropy, K_1 , should take place.

On the other hand, the modification of the magnetic structure can lead to other changes in the features of the ceramics that should be taken into account. First is the change of the

exchange interactions and hence of the Curie temperature, T_C . In the particular case of the decreasing of the Curie temperature, the magnetization could decrease faster than in standard ceramics. This can impact negatively the use of magnets working in high temperature environments.

Another factor to take into account is that the negative thermal coefficient of hexaferrites makes the coercive field increase with temperature. This makes ferrite magnets perform often better up to $100\text{ }^\circ\text{C}$ than at room temperature. The temperature dependence of H_C depends on the relative temperature dependences of both K_1 and M_S [42] that should vary with the doping due to the change of the T_C . Any changes to each component might thus negatively affect the resulting useful temperature range.

In the framework of the current need for better ferrite magnets, we review here the most promising strategies, as well as recent results, to improve the performance of the hexaferrite based materials for their use in permanent magnets. We present three main pathways for future improvements. First, we consider how engineering the shape and size of hexaferrite nanoparticles can improve the magnetic properties of the commercial ones. In the second section, we consider how substituting one or more of the cations of the hexaferrites can improve the magnetic properties, considering three main themes: the improvement of the well-know LaCo-doped BFO, doping changes oriented towards improving the saturation magnetization, and changes oriented towards improving the coercivity. In the third section, we consider first exchange coupled composites, either with metal or oxide components. We comment on promising results of highly coherent magnetite-hexaferrite platelets and discuss the possibilities of exploiting the internal magnetostatic interactions for improving composites properties. Finally, the conclusions and future prospects of ferrite magnets are addressed.

2. Reducing the size to the nanoscale

2.1. The effect of the size

Among all, the reduction of the grain size down to the nanoscale is generally recognized as a useful route to improve the coercivity of a magnetic material. Indeed, one of the best examples of permanent magnet, Alnico alloys, is actually a nanocomposite. Alnico alloys, which 90 years after their introduction still today play a relevant role in the permanent magnet market, indeed, consist of elongated FeCo-rich ferromagnetic nanograins dispersed in a weakly magnetic Al–Ni matrix. The energy product of the material can be further enhanced through the application of an external magnetic field during the annealing process [47].

In a bulk ferro(ferrite)-magnetic material the process of reversal of magnetization may occur through different mechanisms such as nucleation and propagation of reverse domains and domain walls pinning. However, on squeezing the volume of the material a critical size is reached below which the formation of domain walls is no longer energetically favored, and a single domain (SD) spin configuration is attained. The magnetic behavior of the SD particle can then

Figure 2. Scheme of the room temperature variation of coercivity with the grain size from SPM to SD and MD states. PSD denotes the pseudo-single domain region (PSD) while the light grey zone indicates the size range where incoherent reversal processes might occur. Reproduced from [60] with kind permission of Società Italiana di Fisica.

be described by a giant magnetic moment $\mu = M_S V$, where V is the volume, arising from the combination of the individual atomic spins, which reverses its orientation through the coherent rotation of the single spins. For a given material the critical diameter, d_C , for the SD spin configuration state depends on the competition between three energy terms, magneto-static energy, magnetic anisotropy constant, K , and exchange energy, A . For a spherical nanocrystal, d_C , is expressed as [48]:

$$d_C = (18E_\sigma) / (\mu_0 M_S^2) = (36\sqrt{AK}) / (\mu_0 M_S^2) \quad (2.1)$$

where μ_0 is the vacuum permeability and $E_\sigma = 2(AK)^{1/2}$ is the surface energy of a domain. Typical values of d_C for most common magnetic materials, range from few tens up to several hundreds of nm. For Ba-ferrite theoretical and experimental estimates suggest d_C is ca. 500 nm [49, 50], although higher values up to 1 μm have been often observed [49, 51]. Critical diameters in the micron scale, up to 1.5 μm , have been also reported for SFO [52, 53]. González and coworkers estimated the activation volume, i.e. the size of the nucleus that gives rise to magnetization reverse, of a series of Ba-ferrite particles with sizes between 65 nm and the micrometric range. A jump in the coercive field from 200 kA m^{-1} to 320 kA m^{-1} was observed when the particle size decreased below 800 nm, while the activation size decreased from 40 nm to 25 nm along the series. The authors concluded that the reversal process in the particles above 800 nm occurs by domain wall propagation, while the coherent rotation is the basic reversal process for the smaller particles [54].

The transition from the multidomain (MD) to the SD state is not sharp and several non-uniform reversal modes as curling

and buckling can occur, where unlikely coherent reversal, the exchange energy does not remain constant [55, 56]. For example Chan *et al* measured by magnetic force microscopy the switching field of a series of isolated nanoparticles of doped barium ferrite and found the Stoner–Wohlfarth model was followed only for average size up to 50 nm; on increasing the size above this limit incoherent reversal modes produced a decrease of the switching field [57]. Moreover, an investigation by electron holography of the magnetic domain structure on isolated Ba-ferrite particles obtained by heat treatment at 1100 °C of a mixture of $\text{BaO} \cdot \text{Fe}_2\text{O}_3$ and an additive of Bi_2O_3 , demonstrated the coexistence of single-domain and two-domain magnetic states over the whole investigated range 0.1–2 μm [51].

As a result of the size dependence of the reversal process, the coercivity increases with decreasing the grain size, as schematically shown in figure 2, as long as the SD threshold is reached. On further shrinking the size, in fact the nanoparticles become unstable against thermal demagnetization till magnetic irreversibility completely disappears and the SPM state is attained. The threshold size below which superparamagnetism can be observed for a given measuring time, depends on the shape and crystal quality (presence of defects and impurities) of the grains. For Ba-ferrite and Sr-ferrite it is generally assumed SPM relaxation becomes dominant for size below 40–60 nm [58, 59]. However, it should be considered that these limits refer to time scale characteristics of laboratory experiments (typically from 10^2 to 10^4 s), which are much shorter than the shelf life required for a permanent magnet (the loss of magnetization must be less than 5% over 100 years).

The expected trend of the coercivity with the particle size was nicely confirmed by a systematic study performed on a

Figure 3. Trend of the coercive field (left) and of the reciprocal grain size (right) as a function of the annealing temperature for a series of BFO particles prepared by conventional ceramic route. Reprinted from [61], Copyright 2005, with permission from Elsevier.

family of polycrystalline BFO samples prepared through conventional solid-state reaction [61]. The size of the particles was tuned between 500 nm and several microns by increasing the calcination temperature from 1000 °C to 1400 °C, while the coercive field was observed to decrease with the grain size from 318 kA m⁻¹ for the smallest samples down to only 100 A m⁻¹ for the largest one, as shown in figure 3. The authors argued that at the small size, coercivity is determined by pinning of the magnetization at grain boundaries, which extent scales with the reciprocal of the size. Conversely, they suggested that in the larger multi-domain grain, the coercivity originated by nucleation and domain wall pinning is small. This study on one hand demonstrated how the coercivity of Ba-ferrite can be easily controlled by the grains size, and on the other confirm the critical single domain size is close to 500 nm.

2.2. The effect of the shape

In addition to the size, also the shape of the particles plays a relevant role in determining the coercive field. According to the Stoner–Wohlfarth model [65], in fact, for non-interacting single domain spheroidal particles, the anisotropy field, H_A , can be expressed as $H_A = H_K + D_N M_S$ where $H_K = 2K_1/\mu_0 M_S$ is the magnetocrystalline anisotropy and D_N is the demagnetizing term, which assumes positive or negative value depending whether the easy axis is parallel or perpendicular to the main axis of the spheroid. M-type ferrites naturally tend to assume a regular hexagonal thin platelet shape with the c -axis, which coincides with the magneto-crystalline easy axis, along the short dimension. Such a shape corresponds to the growth of low energy 001 crystallographic faces. However, often acicular platelets are formed, depending on the synthesis conditions ([66] and Refs. therein). In this situation, a strong demagnetizing field appears when the magnetization is along the easy axis. The decrease of the magnetic anisotropy due to the shape contribution is given by $H_A = H_K - D_N M_S$. D_N is a

Figure 4. Coercivity as a function of the diameter to thickness ratio in Ti-doped barium ferrite nanoparticles. The solid line represents the theoretical curve calculated according to the Stoner–Wohlfarth model assuming the particle shape as an ellipsoid of revolution. Reprinted from [56], with the permission of AIP Publishing.

tensor, but here for simplicity, can be treated as a scalar. The D_N term depends on the diameter (D) to thickness (t) ratio: $D_N = 1$ when $t \ll D$, as is the case of narrow platelets, then it decreases with D/t , reaching zero for equiaxial morphology ($D/t = 1$). For $D/t < 1$, D_N changes sign and H_A increases with respect to H_K . This discussion highlights how the proper design of the particle shape is critical to optimize the effective anisotropy of the ferrite.

The relevance of the shape has been indeed experimentally verified by several authors. For example, Kubo *et al* [56] investigated the effect of the particles shape on the coercivity of Ba-ferrite nanoparticles pure and doped with a small amount of Co and Ti prepared by the glass-crystallization method. They found that H_C decreased with increasing the diameter (d) to thickness (t) ratio down to ca. 32 kA m⁻¹ for d/t ca. 6 (figure 4).

On the other hand the platelet morphology offers the advantage of making small particles (i.e. below 100 nm) more easily orientable, either magnetically or mechanically. This is a fundamental feature since grain orientation along the c -axis allows for increasing the reduced remanence from half the saturation magnetization, which is the value obtained for isotropic magnets fabricated from powders composed of single domain, randomly oriented particles (0.5 is the theoretical value predicted by the Stoner–Wohlfarth model and it corresponds to $(BH)_{\max}$ of approximately 8 kJ m⁻³) up to values close to 1. The Stoner–Wohlfarth model also predicts that the

coercive field increases upon orientation: in fact, while in non-interacting randomly oriented particles $H_C = 0.49 H_A$, for oriented particles H_C coincides with H_A . Thanks to the platelet morphology, particles can be thus easily oriented and coercive field and remanence can be maximized, making the processing from powders to bulk magnets simpler. A similar result can be reached by piling up platelets. In such a situation interparticle interactions determine the decreasing of platelets' internal field that can improve the magnetic properties.

2.3. Experimental methods to prepare nanosized hexagonal ferrite nanoparticles

The control of the shape and of particle size down to the nano-scale is thus crucial for the fabrication of high-performance permanent magnets. Accordingly, attempts to prepare M-hexaferrite nanoparticles, nanorods and nanofibers, appeared in the literature since the 80s, at the very beginning of the nanomaterial era. However, it should be stressed that most of the recent literature is focused on the fabrication of nanosized M-hexaferrites with properties optimized for applications other than permanent magnets. M-type hexagonal ferrites, indeed, can be employed in several technologies such as microwave devices, magnetic recording or electromagnetic wave absorber, just to mention a few, where the nanostructure can have a significant impact [66]. The many techniques developed so far include solid-state reaction [72, 73], sol-gel [74, 75], co-precipitation [76–78], milling [79, 80], microemulsion [81, 82], aerosol pyrolysis [83], combustion synthesis [84], glass crystallization [85, 86], and hydrothermal techniques [87–91]. Particularly, low temperature chemical routes demonstrated to be effective to prepare M-hexaferrite nanostructures with controlled size and shape and good magnetic properties. Sol-gel has been largely employed to prepare nanoparticles of pure and substituted Ba- and Sr-ferrite [74, 92–95]. Importantly, the main efforts have been spent in the rationalization of the role of each parameter of the synthesis. Indeed, it has been shown that a fine control over the several parameters is required to avoid the formation of secondary phases as hematite or goethite, to control the size and to optimize the magnetic properties. Nga and coworkers studied the influence of pH, Fe/Sr molar ratio and calcination temperature and time on the structure and magnetic properties of SFO particles prepared by a sol-gel method using metal nitrates in citric acid as precursors [96]. They found that pure nanometric particles with hexagonal morphology could be obtained with Fe/Sr molar ratio of 10.5 and pH = 1 and after calcination at 850 °C. The particles had M_S of 56 A m² kg⁻¹ and $H_C = 525$ kA m⁻¹. A Ba/Fe molar ratios of 11.5 was instead found has the best one to obtain well-crystalline M-type Ba-ferrite powder with crystallite size 22 nm, and better magnetic properties ($H_C = 420$ kA m⁻¹) by the sol-gel auto combustion technique [97]. Similarly, Yu and Huang investigated the role of pH and citric acid/metal ion molar ratio in the sol-gel synthesis of BFO. Single domain particles with magnetization at 2.39 MA m⁻¹ of 61.7 A m² kg⁻¹, remanence of 30.8 A m² kg⁻¹ and coercivity of 406 kA m⁻¹ were obtained at pH = 3 and calcination at 900 °C [98]. In a previous work

it was suggested that adding a pre-decomposition step of the gel at 450 °C, in a standard sol-gel method with Fe:Ba = 11.5 and sintering at 900 °C, Ba-ferrite nanoparticles with average size ca. 100 nm, saturation magnetization of as high as 70 A m² kg⁻¹ and coercivity of 473 kA m⁻¹ could be obtained [74].

A tight control on the synthesis parameter is required also when nanosized M-type hexaferrites are prepared by the hydrothermal technique. This synthesis route has been largely proved to be effective to this goal [23, 89–91, 99, 100]. Moreover, as is the case of the sol-gel method, hydrothermal technique is simple and cost-effective, and can be easily scaled to industrial production. Despite many advantages, several studies have demonstrated how the properties of the final materials depend on many, often intertwined, synthesis parameters such as temperature, time, precursor molar ratio, nature of the base and its concentration, just to name the most important. This aspect, in one hand expands the potentiality of the technique in designing hexaferrites with tailored properties for each given application, on the other hand can make rather tricky the determination of the optimal preparation conditions to get the desired result. In this framework, some interesting results were recently reported by Saura-Muzquiz and coworkers, who demonstrated the fundamental role of the concentration of the metal ion solution, when adding the precipitating NaOH solution in the precursor preparation, in the synthesis of Sr-ferrite nanoparticles by a hydrothermal route [62]. By changing the iron concentration from 0.05 to 0.75 M, hexagonal platelet-like particles with mean diameters from 48 nm to 254 nm, whose magnetic properties were strongly dependent on the average size (specific remanence, σ_R , from 20 to 31 A m² kg⁻¹ and H_C from 68 to 175 kA m⁻¹) could be obtained. Notably, it was also shown that, in contrast to the bulk, in the thinnest nanoplatelets the termination layer was the Sr-containing one. Interestingly, the authors addressed the critical issue of processing the as-prepared nanopowders to obtain bulk magnets. By using spark plasma sintering (SPS) they managed to produce high-density (more than 95% of the bulk density) SFO magnets. X-ray pole figure measurements demonstrated that, thanks to the anisotropic shape, self-induced alignment of the nanoplatelets occurred during SPS (figure 5). This texturing, which was accompanied by the increase of the platelet thickness, provided high-performance permanent magnets with square hysteresis loop and $(BH)_{max}$ of 30 kJ m⁻³ (figure 5). The $(BH)_{max}$ could be further enhanced up to 36 kJ m⁻³ by annealing at 850 °C. This energy product was claimed by the authors to exceed those of the best dry-processed commercial ferrites.

In another work, Raghuram and coworkers used the hydrothermal method to prepare a series of Ba_{1-x}Sr_xFe₁₂O₁₉ nanoparticles with variable composition ($x = 0-0.8$) [101]. The particles exhibited average size between 15 and 20 nm and their shape evolved from plate-shaped grains to nanorods as a function of dopant. A marked decrease of M_S with x and an increase of both coercivity (from 159.1 to 189.5 kA m⁻¹) and critical temperature (from 703 K to 753 K) were observed as a function of x . However, the cation replacement was not found to be the only factor responsible for the observed trend. In

Figure 5. Properties of the pellet obtained by SPS of Sr-ferrite nanoplatelets SPS_XX, where XX denotes the initial iron molar concentration: (a) reconstructed (001) x-ray pole figures and texture index of SPS_005, SPS_015 and SPS_075; the texture index of a randomly oriented sample is 1.0; (b) fraction of oriented crystallites as a function of the α angle; the inset depicts the dependence of the texture index on the aspect ratio of the starting nanopowders; (c) hysteresis loop of SPS_005 pellet before (orange line) and after annealing at 850 °C for 4 h (red) and 16 h (black); (d) saturation magnetization, M_s , remanence, M_r , and coercivity, H_c , variation with the annealing time. Reprinted (adapted) with permission from [62]. Copyright 2018 American Chemical Society.

all cases the magnetic properties were far from those required for the realization of permanent magnets. The same observation was reported by Al-Hwaita and coworkers who systematically investigated micrometric $Ba_xSr_{1-x}Fe_{12}O_{19}$ ferrites prepared by solid state reaction and sol-gel methods and found that replacing Ba with Sr was not the only critical parameter for improving the magnetic properties, but other parameters as grain size and phase purity were similarly crucial [102].

Water-in-oil microemulsions have also been applied to prepare single domain M-type hexaferrite particles. Koutzarova and coworkers used a reverse microemulsion containing cetyltrimethylammonium bromide as cationic surfactant, n-butanol as co-surfactant, n-hexanol as a continuous oil phase, and water to obtain, after a calcination step at 900 °C, pure Ba-ferrite nanoparticles with average size 280 nm [103]. Most of the particles were found to have an almost perfect hexagonal shape with average aspect ratio diagonal/thickness of 7.7, although the smallest ones (ca. 100 nm) exhibited spheroidal or irregular shape, indicating that the growth process towards the most energetically favored hexagonal shape is incomplete. However, the particles exhibited low saturation magnetization of 48.9 A m² kg⁻¹ and coercivity, 240 kA m⁻¹ for both temperatures, which was attributed to the presence of small nanoparticles and the incomplete filling of the bipyramidal 2b sites by Fe³⁺ ions.

Low temperature solid state reaction represents a simple and efficient alternative method for the preparation of

strontium ferrite nanoparticles. Kiani *et al* [72] reported that sintering at 750 °C a powder obtained by mixing and grinding the corresponding metal chlorides and sodium hydroxide, yields pure, highly crystalline, hexagonally shaped SFO nanoparticles, with size ranging from 30 to 150 nm. The control of the Na⁺/Sr²⁺ and Fe³⁺/Sr²⁺ molar ratios was found to be essential to tune the size and morphology of the particles and avoid the formation of impurity phases as haematite and SrFe₂O₄. The as obtained nanoparticles exhibited high coercivity (412 kA m⁻¹; 0.52 T), saturation magnetization of 58.7 A m² kg⁻¹ and reduced remnant magnetization, $M_r/M_s = 0.49$, close to the theoretical value for randomly oriented single domain nanoparticles. Following a similar procedure some of us obtained SFO nanoparticles with a platelet morphology and average diameter ca. 70 nm. The nanoparticles displayed extremely good magnetic properties with $\sigma_s = 68$ A m² kg⁻¹ and a coercive field of 525 kA m⁻¹, as shown in figure 6 [104]. Since the proposed method is very simple, solvent-free, uses low cost reactants, and leads to high yield, it is very promising for the large-scale industrial production of nanoscale M-hexaferrite particles.

2.4. 1D nanostructures

One-dimensional magnetic nanostructures also received considerable attention since the high aspect ratio and the large surface may provide a remarkable contribution to the total

Figure 6. Representative bright field TEM image (left) and hysteresis loop (right) of M-type Sr-ferrite nanoparticles prepared according to the procedure described in [72]. The inset is the enlargement of the low field loop in the II quadrant. Reproduced from [104]. © IOP Publishing Ltd. All rights reserved.

Figure 7. Left: schematic diagram of the mechanism of formation of Sr-ferrite nanoribbons by the polymer-sol assisted single-spinneret electrospinning route followed by heat treatment in air at 800 °C for 2 h; Right: representative field emission scanning electron microscopy (first and second columns) and transmission electron microscopy (third column) images of three samples prepared from spinning solutions with PVP concentrations of 8.5% (a)–(c), 10.4% (e)–(j) and 12.3% (i)–(k), respectively. Reproduced from [63]. CC BY 4.0.

magnetic anisotropy, strongly modifying the coercivity. For example, an infinitely long chain of stacked platelets theoretically can have enough shape anisotropy to increase coercivity up to 1000 kA m⁻¹ [105]. In this context, several attempts have been carried out to realize mono-dimensional hexagonal M-hexaferrite structures and, among the methods developed so far electrospinning has been established as one of the most promising. SFO ribbons with average width in the submicrometric range, from 484 ± 15 to 1099 ± 18 nm, were successfully fabricated via a polyvinylpyrrolidone sol-gel assisted electrospinning method followed by heat treatment in air [63]. The nanoribbons, shown in figure 7 were formed by the assembling of smaller, strongly interacting ferrite nanoparticles whose average size decreased with increasing the width of the nanoribbon (from 131 ± 3 nm to 76 ± 3 nm). Both the width of the ribbon and the size of the constituent nanoparticles could be simply tuned through the polymer concentration in the spinning solution. The peculiar ribbons structure together with the high crystallinity of the nano-

particles granted for high coercive fields and good M_S and M_R values ($H_C = 582$ kA m⁻¹, $\sigma_S = 67.9$ A m² kg⁻¹ and $\sigma_R = 37.3$ A m² kg⁻¹ for the best sample).

BFO nanofibers of average length 1.5 μm consisting of randomly oriented, single nanoparticles of average diameter 70 nm, stacked along the nanofiber axis, were also prepared by electrospinning a homogeneous PVP/barium nitrate/iron nitrate precursor sol-gel solution, followed by calcination at 800 °C [64]. The authors investigated the growth mechanism and proposed a model which involves a series of phase transformations occurring at increasing temperatures, with Fe₂O₃, BaO, BaCO₃ and BaFe₂O₄ as intermediate species (figure 8). The resulting nanofibers showed at room temperature $\sigma_S = 71.5$ A m² kg⁻¹, reduced remanence of ca. 0.5 and $H_C = 473$ kA m⁻¹, lower than the theoretical expected value. Interestingly, the magnetization reversal mechanism of the BFO single-particle-chain nanofibers was theoretically analyzed and found to fit a curling model.

Figure 8. Schematic description of the reversal mechanism ((A), top) and of the formation of the BFO single-particle-chain nanofibers ((B), bottom). Reprinted with permission from [64]. Copyright 2012 American Chemical Society.

On the other hand, Liu and coworkers used a combined approach based on electrostatic spinning and high-temperature heating treatment, to prepare, by properly varying the experimental conditions, BFO hexagonal ferrites with different geometry (nanowire, nanoribbon and nanotube) [106]. The comparison of the magnetic properties of the samples demonstrated that while the nanostructure had similar saturation magnetization ($46\text{--}49 \text{ A m}^2 \text{ kg}^{-1}$), the coercivity was largely dependent on the morphology, increasing from nanowires (190 kA m^{-1}) to nanoribbons (224 kA m^{-1}) and nanotubes (259 kA m^{-1}). Despite the obtained coercivities are lower than those of many other hexagonal Ba-ferrites, this work nicely demonstrates the versatility of the technique for the fabrication of material with controlled properties for different applications.

3. Controlling doping

The structure of the M-hexaferrites suggests a direct way to improve their magnetic properties by substituting some of the cations of specific sites of the magnetoplumbite structure. The progress in this subject up to the early 80s was reviewed by Kojima [42]. However, there was a lack of reviews in the field between that work until the recent overviews by Pullar [66], Mahmood [9], and Lisjak *et al* [18]. Considering this large number of studies, here we will select and discuss the works reporting larger values of the magnetization, of the coercive field, or of the anisotropy field, than those of the best commercial hexaferrite magnets ($\sigma_S = 70 \text{ A m}^2 \text{ kg}^{-1}$, $H_C \geq 398 \text{ kA m}^{-1}$, $\mu_0 H_C = 0.5 \text{ T}$, and $H_A = 1.43 \text{ MA m}^{-1}$, $\mu_0 H_A = 1.8 \text{ T}$).

The doping of the M-type ferrites is in general designed to affect the intrinsic magnetic structure of this magnetoplumbite crystal in which magnetic properties are determined by the occupancy of five sites and their magnetic coupling through oxygen-driven superexchange interactions. Such design should take into account that the spin and orbital moments of the cations depend on the site and charge distribution [107], through the spin configuration (high spin, low or intermediate spin) that determines the net magnetic moment and the local anisotropy of an ion. In this framework, different scenarios of substitution can be considered. One is to change the total magnetization. Another is to enhance the magnetocrystalline anisotropy by increasing locally the magnetic anisotropy due to the contribution of the orbital moment or the change of the crystal and chemical structures. In general, the substitution of all the ions that are present in the structure has been explored [9, 42, 66]. However, the most common approach considers the substitution of cations, although changing even the oxygen anions have been tried [42]. We briefly enumerate the rationale for changing the cations:

- Substitution of Fe with non magnetic cations. An increase of the magnetization could be obtained if non-magnetic atoms like Al^{3+} , Ti^{2+} substitute Fe in the spin-down sites giving an increase of the net moment. However this also leads to a weakening of the exchange interactions that can reduce the Curie temperature, reducing the room temperature magnetization, or weakening or modifying the anisotropy symmetry [42, 108–110].
- Substitution of Fe with magnetic ions. For the first goal, Fe^{3+} and Mn^{2+} are the 3d transition metal ions with the largest magnetic moments (figure 2), and heavy rare earth ions [107, 111] have even larger magnetic moments. Hence the doping with other 3d metals could increase the total magnetization if ions occupy the spin down sites. Regarding the increase of the magnetocrystalline anisotropy, the substitution of ions with spin orbit contribution could be the main strategy. However, except in some cases [107, 112], the high symmetry of the octahedral and tetrahedral sites quenches the orbital moment. Hence the occupancy in the bipyramidal 2b site is the target. In addition, the anisotropy that finally results in the occupancy of any site [113], analysed in terms of the single ion model, depends on the spin orbit coupling and electronic configuration of the cation and the link in the charge distribution of the site. For example, Co^{2+} occupancy in CoTi-SFO changes the anisotropy from uniaxial to planar giving a ferrite with soft magnetic properties. In general, the contribution of the higher symmetry sites to the anisotropy cannot be neglected [108].
- Substitution of Ba^{2+} or Sr^{2+} ions. Non magnetic La^{2+} , Ca^{2+} , or Ag^{+} ions, or rare earth ions RE^{3+} [114] or others modify the crystal structure, thus modifying the exchange interactions and the magnetic anisotropy. For example, the slightly larger magnetic anisotropy of SFO ($K_1 = 35 \text{ kJ m}^{-3}$) compared with BFO ($K_1 = 32.5 \text{ kJ m}^{-3}$) is correlated with the smaller size of Sr^{2+} ($1.13 \text{ \AA}$) compared with Ba^{2+} ($1.35 \text{ \AA}$). This is explained because the smaller cell size implies a decreased Fe–O distance in the

2b sites, increasing the crystal field and zero-field splitting of Fe^{3+} and the magnetic anisotropy. As can be seen in figure 1(a), the $4f_2$ and 2b sites have three Sr^{2+} ions in their vicinity. Thus the substitution of $\text{Ba}^{2+}/\text{Sr}^{2+}$ ions by others induces a change of both the symmetry and the charge density around these sites [115].

Often the substitution of the ions is analyzed in terms of their charge, by their nominal oxidation state: divalent, trivalent or others [9, 66]. The selected ion can affect the synthesis and sintering processes, but the main influence is due to the required charge neutrality of the resulting hexaferrite. The doping with divalent cations or other multivalence ions can lead to the presence of Fe^{2+} instead of Fe^{3+} to maintain the charge neutrality. Fe^{2+} ions, in addition to contributing to the anisotropy, provide a possible charge transfer mechanism that can lead to an undesired increase of the electrical and thermal conductivity that are key features of the hexaferrites for many magnetic applications.

The main result can be obtained from the literature is that the changes of the properties induced by the doping are extremely difficult to predict and understand. In the following, the properties of LaCo hexaferrites, the only doped ferrite present in the market, are discussed first. Then the recent results are split in works that aim to increase the saturation magnetization and ones that try to increase H_C with features above the commercial ones.

3.1. LaCo–SFO magnets

The only successful story of doping hard hexaferrites is that of lanthanum and cobalt doping of SFO. Commercial magnets are produced by Hitachi [6] and TDK [116]. Considering the TDK catalogue, strontium ferrite magnets exhibit a maximum B_R up to 0.45 T and intrinsic H_C up to 319 kA m⁻¹ while LaCo-doped ferrites reach a B_R of 0.47 T and an intrinsic H_C of 343 kA m⁻¹, being the maximum $(BH)_{\max}$ of 38 kJ m⁻³ for the first and 44 kJ m⁻³ for the doped ferrites. Moreover, they keep low electrical and thermal conductivities, positive temperature H_C coefficient (0.1–0.4% K⁻¹), and T_C (around 430 °C–460 °C). Early work was performed by Yamamoto *et al* [117] but their properties were identified at the end of XX's century [118, 119]. The commercial LaCo-ferrites exhibit a weak improvement of the magnetic properties, a 10% in remanence and in the $(BH)_{\max}$, respect to those of the state of art SFO ferrites, demonstrating the high potential in the research of alternatives to the conventional hard ferrites.

Research has focused in the composition range $0 \leq x \leq 0.4$ of $(\text{Sr}_x\text{La}_{1-x})(\text{Fe}_{12-x}\text{Co}_x)\text{O}_{19}$ where La^{3+} and Co^{2+} substitute for Sr^{2+} and Fe^{3+} . Most of the studies report that a single phase magnetoplumbite structure is present in this composition range with a decreasing c -parameter while the a -parameter does not vary. In general, as it can be seen in figure 9, the M_S and M_R values increase slightly while H_C (H_C from the J versus H curves) exhibit a maximum value for a x between 0.2 and 0.4 depending on the preparation method [117, 118, 120–123]. Early studies of Iida *et al* [121], reported that LaCo–SFO sintered at 1200 °C present an increase of

H_C from 319 kA m⁻¹ of 383 kA m⁻¹ when doping from $x = 0$ to $x = 0.3$. The remanence increases weakly reaching a maximum at $x = 0.4$. For larger x both M_R and H_C decrease. Nishio *et al* [123] indicate that the doping that optimizes H_C depends on the thermal treatment. Nowadays, the work of Kikuchi *et al* [124] reports the maximum H_C of 0.8 T but with a small decrease in M_S . Likewise, H_A is also larger than for SFO and BFO and increases almost linearly with x up to $x = 0.4$. However, the H_A values depend on the preparation route and they go from 1.35 MA m⁻¹ [124] to 1.91 MA m⁻¹ [125]. The anisotropy is described in terms of a uniaxial anisotropy with a single second order constant K_1 as in SFO. At lower temperatures, higher order anisotropy constants need to be considered. At higher temperatures, a continuous increase of the H_A is correlated to the faster temperature dependent decrease of the magnetization rather than changes in anisotropy [125].

Mössbauer spectroscopy [122, 127], and NMR [45] and x-ray magnetic dichroism [128] studies indicate that the Co^{2+} ions occupy both 2a (spin up) and $4f_2$ (spin down) sites. The weak increase of the magnetization should indicate an almost equipartition of the Co cations between the two locations. On the other hand, the increase of the anisotropy field, even if the magnetization remains constant, is an indication of the increase of the magnetocrystalline anisotropy as reported in [126]. It can be explained by the contribution of the orbital moment of the Co^{2+} cations. In the case of Co–Ti hexaferrites, Co cations occupy other sites and the result is the change of the symmetry of the anisotropy and the decrease of the H_C [118]. The decrease of volume and the increase of the charge due to La^{3+} also stabilizes the Co^{2+} in the 2a and $4f_2$ sites. On the other hand, Morel *et al* [122] showed by Mössbauer and resistivity studies that the Fe^{2+} is present in powders obtained by the ceramic method. The imperfect distribution of La and Co ions could give rise to the Fe^{+3} cations replacement by Fe^{2+} ions, mainly in 2a site [126]. Also, the presence of Fe^{2+} cations has been demonstrated recently in single crystals [129, 130]. However, increasing the population of Fe^{2+} decreases the electrical resistivity.

However, the changes in M_R and H_C have been discussed in terms of the influence of the microstructure and morphology in the reversal process [118, 119, 131, 132]. Shape morphology of the particles determine intra and interparticle demagnetizing fields [133]. Typical platelet morphology of SFO leads to internal demagnetizing fields (i.e. shape anisotropy) that decreases the anisotropy field and thus the coercive field. Grossinger *et al* [120] explain in this way the drop of H_C above $x = 2$ in this way, even if H_A increases. Another factor is the particle size distribution, with the presence of single domain vs. multi domain particles. The reversal process of single domain particles is determined by the coherent rotation modes and gives rise to a high coercivity. Nishio *et al* [123] calculated the single domain critical size for these oxides that is around 1 micron, similar to that of SFO. In multi domain particles, the domain reversal is dominated by domain wall motion and lower switching fields are obtained. Hence the magnetic properties of these new magnets are determined both by the changes of the magnetic structure and by microstructural features.

Figure 9. (Left) intrinsic coercivity (H_c) and remanent induction (B_r) as a function of doping content (Reprinted from [120], Copyright 2004, with permission from Elsevier) and (Right) composition dependence of the anisotropy field H_a and H_c of $\text{Sr}_{1-x}\text{La}_x\text{Fe}_{12-x}\text{Co}_x\text{O}_{19}$ ferrite magnets measured at room temperature (Reprinted from [126], Copyright 2004, with permission from Elsevier).

Table 1. Data of doped-ferrites with specific magnetization larger than $70 \text{ A m}^2 \text{ kg}^{-1}$. x corresponds to the doping content, H_c , the coercive field, and σ_s the specific saturation magnetization.

Doping	Formula	Synthesis route	x	H_c (kA m^{-1})	σ_s ($\text{A m}^2 \text{ kg}^{-1}$)	Ref.
Ni	$\text{BaFe}_{12-x}\text{Ni}_x\text{O}_{19}$	Sol-gel	0.8	24	101	[139]
Zn, Ti	$\text{BaFe}_{12-2x}\text{Zn}_x\text{Ti}_x\text{O}_{19}$	Chemical route	0.4		110 ^a	[140]
Zn-Zr	$\text{SrFe}_{12-2x}\text{Zn}_x\text{Zr}_x\text{O}_{19}$	Mechanochemistry	1	24	108	[141]
Zn-Sn	$\text{SrFe}_{12-2x}\text{Sn}_x\text{Zn}_x\text{O}_{19}$	Sol-gel	2.5		97	[142]
Zn-Ni	$\text{SrFe}_{12-2x}\text{Ni}_x\text{Zn}_x\text{O}_{19}$	Sol-gel auto combustion method		160	99	[143]
Zn-Sb,V,Nb	$\text{SrFe}_{12-2x}\text{Y}_x\text{Zn}_x\text{O}_{19}$	Ceramics	0.2	238.7	75	[144]
Zn,La	ZnLa-SFO	Ceramics		191	74	[145]
Al,Cr	$\text{BaFe}_{12-2x}\text{Al}_x\text{Cr}_x\text{O}_{19}$	Sol-gel auto combustion method	4	318	88	[146]

^aStudy performed at low temperature.

Nowadays, several investigations consider the improvement of these LaCo-hexaferrites, mainly using Ca-doping using different synthesis methods [134–137]. Chen *et al* [136] obtained Ca doped LaCo-ferrites employing microwave calcination with H_c up to 397.9 kA m^{-1} and B_r of 0.44 T . Du *et al* [138] used doping with Al^{3+} . In these ferrites the magnetization decreases and the H_c increases linearly with the doping, reaching a H_c of 716.2 kA m^{-1} and specific magnetization of around $50 \text{ A m}^2 \text{ kg}^{-1}$.

3.2. Doping for increasing the magnetization

Several approaches have been attempted to increase the magnetization of M-hexaferrites by doping: substitution with a single non-magnetic element, substitution with a single magnetic element, or substitution with several (magnetic and non-magnetic elements). Table 1 summarizes the studies in which the specific magnetizations above the threshold of $70 \text{ A m}^2 \text{ kg}^{-1}$ of the high quality SFO and BFO.

Ab initio calculations by Dixit *et al* [147] predicted that Bi, Sb, Sn, and Sc ions can increase the magnetization. However, until now, no experimental result has obtained an increase in the magnetization. The case of Al^{3+} has been extensively studied, but without success. For a single magnetic cation the only work that reports an improvement is Wang *et al* [139], who studied the synthesis of magnetic Ni-doped BFO grown by sol-gel. They found a specific magnetization of $101 \text{ A m}^2 \text{ kg}^{-1}$ for $x = 0.8$. The authors suggest the origin

is the occupancy of $4f_2$ spin down sites with Ni^{2+} coupled to the presence of Fe^{2+} in $4f_1$ and $4f_2$ sites.

More promising is the approach of doping with multiple cations with a general formula $(\text{Sr})\text{BaFe}_{12-2x}\text{Y}_x\text{Z}_x\text{O}_{19}$ where Y, Z, are different cations. Often the substitution is performed with non magnetic ions (paramagnetic or diamagnetic) and there is substitution of Fe^{3+} by equal amounts of divalent and tetravalent cations to assure charge neutrality. Larger specific magnetizations were observed by Zn^{2+} doping combined with other diamagnetic ions, Ti [140], Zr [141], Sn [142], Nb [144], V [144], Sb [144], La [145] or with paramagnetic Ni cations [139] that gives rise to ferrites with σ_s between 75 and $110 \text{ A m}^2 \text{ kg}^{-1}$. Also the double doping of BFO with diamagnetic Al^{3+} and paramagnetic Cr^{3+} cations [146] gives rise to a ferrite with a maximum specific magnetization of $88 \text{ A m}^2 \text{ kg}^{-1}$ for $x = 0.4$.

In general, it was observed that the net magnetization and the specific magnetization do not exhibit monotonic trends with the doping. Wartewig *et al* [140] show that in Zn^{2+} and Ti^{4+} doped BFO synthesized by coprecipitation the magnetization at low temperatures first increases and then decreases above a doping of 0.4 as can be seen in figure 10 (left). The authors suggest that for small doping, Zn^{2+} occupies the tetragonal $4f_1$ sites while Ti cations go to the octahedral spin down $4f_2$ and spin up $12k$ sites and after $x > 1.2$ a small amount of Zn is placed at the 2b site and of Ti at the 2a site. At room temperature, Abdellhi *et al* [141] observed the similar doping dependence in mechanical milling Zn-Zr-SFO micro-particles, with a maximum magnetization of $108 \text{ A m}^2 \text{ kg}^{-1}$

for $x = 1$ ($\text{SrFe}_{10}\text{ZnZrO}_{19}$). Li *et al* [148] also reports the presence of a maximum in the magnetization but for lower Zr–Zn content and only at low temperatures. Ghasemi *et al* [142] have doped BFO with Sn^{4+} and Zn^{2+} synthesized by sol–gel ($\text{SrFe}_{12-2x}\text{Sn}_x\text{Zn}_x\text{O}_{19}$). An increasing doping dependence of the magnetization is found, with a maximum of $87 \text{ A m}^2 \text{ kg}^{-1}$ at $x = 2.5$, as can be seen in figure 10 (right). Mössbauer studies indicate that both Sn and Zn cations occupy spin down $4f_2$ and spin up 2b sites. In contrast, Fang *et al* [149] observed a continuous decrease of the magnetization using co-precipitation. Widyastuti *et al* [143] report the combination of Ni and Zn substituted BFO synthesized by sol–gel auto combustion method and further annealing. They obtain $94 \text{ A m}^2 \text{ kg}^{-1}$ in doped BFO with 2%wt Ni and 4%wt of Zn. Also, Sapoleva *et al* [144] report the same trend in ZSb, ZnNb and ZnV doped SFO.

The enhancement of the magnetization has been observed also in other compositions. Trudel *et al* [146] investigated the double substitution of Al^{3+} and Cr^{3+} cations in $\text{BaFe}_{12-2x}\text{Al}_x\text{Cr}_x\text{O}_{19}$ obtained by auto combustion sol–gel. For $x = 0.4$ the doped ferrite exhibits a specific magnetization of $88 \text{ A m}^2 \text{ kg}^{-1}$, larger than the corresponding BFO ($81 \text{ A m}^2 \text{ kg}^{-1}$) and similar mass density. The improvement of the magnetization does not follow a linear dependence. The authors propose that the increase of the magnetization takes place when Cr^{3+} begins to occupy the $4f_2$ site. Bsoul *et al* investigated doping with non magnetic Ti^{2+} and Ru^{4+} cations of BFO [150] and SFO [151]. In the two oxides a broad maximum in the magnetization versus the doping is observed. The specific magnetizations with substitution of $x = 0.2$ reach up to 10% respect to values of around $60 \text{ A m}^2 \text{ kg}^{-1}$ for both undoped oxides. Mössbauer spectroscopy studies reveal that Ti^{2+} and Ru^{4+} ions substitute Fe^{3+} ions at both the spin down $4f_2$ and $4f_1$ sites and the spin up 2a and 2b sites, for x values up to 0.2 and at the 12k site for $x > 0.2$.

Considering the mentioned results it appears that two-ion doping, mostly performed with non magnetic ions, allows to increase the magnetization in a certain doping range above which the magnetization decreases. In most of the discussed cases, cations occupy the two spin-down sites before reaching the maximum in the magnetization, while for higher doping the cations begin to occupy the spin-up sites giving rise to the reduction of the magnetization. The decrease of the magnetization is also related to the weakening of the superexchange interactions between the different sites due the presence of a large number of nonmagnetic cations in the spin down sites. This is also expected to be correlated with a decrease of the Curie temperature, although this point has not been investigated in the mentioned studies. However, as a rule the cation site distribution depends on the chemical process used to grow the material, as demonstrated for example by the magnetic properties of ZnTi [140, 152] and ZnZr [141, 148] doped hexaferrites.

A crucial point is that in all the previous cases reported H_C is reduced by doping. This is explained as in most cases the doping involves non-magnetic ions that weaken the magnetic order. As can be seen in table 1 ferrites with

$\sigma_s > 100 \text{ A m}^2 \text{ kg}^{-1}$ cannot be considered for hard magnets. The ferrites with $\sigma_s < 100 \text{ A m}^2 \text{ kg}^{-1}$ (22–24) exhibit $H_C > 159 \text{ kA m}^{-1}$ similar to those of standard hard hexaferrites. The only exception is the work of Trudel *et al* [146] in AlCr–BFO. For $x = 0.4$ the doped ferrite exhibits both a larger specific magnetization ($88 \text{ A m}^2 \text{ kg}^{-1}$) and a coercive field (318 kA m^{-1}) than the corresponding magnetization ($81 \text{ A m}^2 \text{ kg}^{-1}$) and H_C (127 kA m^{-1}) of BFO. The authors explain this increase due to the smaller particle size of the doped ferrites and the consequent increase of the coercivity by the single domain behavior of these particles.

3.3. Doping ferrites with high coercive field and the anisotropy

Given that most works that attempted to increase the magnetization reduced the anisotropy, we consider now works that have increased the latter. There is a large number of studies that report increases of H_C of M-hexaferrites by comparing the effect of different cations [108, 153, 154]. In this review, the discussion will be focused on selected studies that report materials with a coercive field larger than those of the best commercial SFO and BFO magnets, i.e. above of 400 kA m^{-1} ($\sim 0.5 \text{ T}$). The main properties of the selected works are summarized in the table 2.

We first consider single doping. Theoretical calculation performed by Dixit *et al* [147] conclude that doping with P, Co, Al, Ga, and Ti cations should increase the anisotropy constant and/or anisotropy field. The Al-doped ferrites are particularly interesting as in several cases [155–157], they exhibit H_C and induction magnetizations larger than those of NdFeB ($H_C \sim 1.19 \text{ MA m}^{-1}$, 1.5 T , and $\mu_0 M_S = 1.6 \text{ T}$). Hence these materials will not be demagnetized by the stray field generated by the NdFeB magnets and can be included in magnetic designs in which RE magnets are present. In addition, as can be seen in figure 11, the doping dependence of H_C exhibits peak variation which maximum H_C corresponds to different Al-doping, x , depending on the synthesis and treatment processes [155, 157, 159]. However all the works in table 2 and others [108, 154, 169–171] report that Al–SFO exhibits common features with the doping. First, the structure volume decreases as the Al content increases remaining c/a ratio almost unchanged. This is expected considering the small ionic radius of Al^{3+} ($0.5 \text{ \AA}$) in comparison with the Fe^{3+} ($0.64 \text{ \AA}$). Second, the doping gives rise to a corresponding decreasing of the magnetization as well the T_C . Most of the works discuss the decreasing of the magnetization in terms of the occupancy of the spin up 2a and 12k sites by non magnetic Al^{3+} cations [108, 159, 160] even though it could be due also to the presence of secondary phases. Theoretical studies by Dixit *et al* [171] confirm the preferential occupancy of Al in those sites. The occupancy of the spin down 4f site has been also reported [154]. The occupancy by non magnetic ions also weakens the super-exchange interactions between the Fe ions that cause the decreasing of T_C . This could cause the spin canting or the change of the anisotropy structure or strength. Regarding the anisotropy, the H_A of Al-doped ferrites was measured in single crystal [172]. A maximum H_A of

Figure 10. Left. Magnetization as a function of applied field at temperature of 5 K for $\text{BaFe}_{12-2x}\text{Zn}_x\text{TiO}_{19}$ (Reprinted from [140], Copyright 1999, with permission from Elsevier). Right. Dependence of the M_S from substitution amount x for $\text{SrFe}_{12-2x}(\text{Sb}_{0.5}\text{Zn}_{0.5})_x\text{O}_{19}$ (Reprinted from [142], with the permission of AIP Publishing).

Table 2. Data of doped-ferrites with $H_C > 0.5$ T. x corresponds to the doping content with the maximum specific magnetization, H_C , the coercive field, and σ_S the specific saturation magnetization, with the maximum specific magnetization.

Doping	Formula	Synthesis route	x	H_C (MA m ⁻¹)	σ_S (A m ² kg ⁻¹)	Ref.
Al	$\text{SrFe}_{12-x}\text{Al}_x\text{O}_{19}$	Auto-combustion	1.5	1.43	9	[155]
		Glycin-nitrate method	4	1.39	7	[156]
		Glycin-nitrate method	3	1.34	28	[157]
		Mechanochemical activation + calcination	4	0.77	23	[158]
		Microwave assisted combustion	1	0.59	35	[159]
		Glass vetrification	1.4	0.48	36	[160]
Mn	$\text{SrFe}_{12-x}\text{Mn}_x\text{O}_{19}$	Ceramics	5	0.77	35	[80]
		Mechanosynthesis	0.5	0.41	58	[161]
Cr	$\text{SrFe}_{12-x}\text{Cr}_x\text{O}_{19}$	Microwave -hydrothermal	0.3	0.58	30	[162]
Sm	$\text{SrFe}_{12-x}\text{Cr}_x\text{Sm}_x\text{O}_{19}$	Sol-gel	0.1	0.48	10	[163]
Al	$\text{Ca}_{0.6}\text{Sr}_{0.1}\text{La}_{0.3}\text{Fe}_{12-x}\text{Al}_x\text{O}_{19}$	Ceramics	1.4	0.48	36	[164]
Ca,Al	$\text{Sr}_{1-x}\text{Ca}_{x/12}\text{Fe}_{12-x}\text{Al}_x\text{O}_{19}$	Citrate melt autoignition method	4	1.67	12.7	[165]
			5	2.86	2	[166]
Al,Cr	$\text{Ca}_{0.33}\text{Sr}_{0.66}\text{FeFe}_9\text{Al}_x\text{Cr}_{4-x}\text{O}_{19}$	Sol-gel	3.5	1.11	35	[167]
CrSnAl	$\text{SrAl}_4(\text{Cr}_{0.5}\text{Sn}_{0.5})_x\text{Fe}_{8-x}\text{O}_{19}$	Sol-gel auto-combustion	0	1.43	12	[168]

2.62 MA m⁻¹ is obtained in $\text{SrFe}_{10}\text{Al}_2\text{O}_{19}$, 1.8 times larger than the corresponding value of SFO. However, the $\mu_0 M_S$ of this doped ferrite is 0.185 T, less than the half of the SFO, what suggests a decreasing of the K_1 . On the other hand, Behhera *et al* [170] report non monotonic but weak changes in K_1 as function of doping for $x < 1$ while Dixit *et al* [171], through *ab-initio* calculations, also suggest small changes in K_1 . Hence the large values of H_C and of H_A are explained from decreases in the magnetization (see figure 11).

Also, doping with magnetic Mn^{3+} [80, 161] or Cr^{3+} [153, 154, 162] cations has demonstrated to increase the H_C respect to the un-doped hexaferrites, which is correlated to the decrease of the magnetization. In the case of Mn cations (Mn^{3+}), early work of Obradors *et al* [173] show that the magnetic anisotropy field at low temperatures increases from 1.43 MA m⁻¹ of the SFO to 4.77 MA m⁻¹ for high doping levels of $x = 9$. Mn cations [80, 161] occupy first the spin up 12k and 2a sites and later the 4f₂ sites. Tenorio *et al*

[80] outlined that the Mn^{3+} ions occupy preferentially the 2a sites, up to the point of completely replacing Fe in those sites. If the Mn content increases further, Mn occupies first the 12k position and finally the 4f₁ and 4f₂ ones. Hence the main occupancy of spin-up sites is with Mn^{3+} which has a smaller magnetic moment (3.5 μ_B) than Fe^{3+} that explains the decrease in magnetization. On the other hand, the same study reports that $\mu_0 H_C$ reaches 0.97 T for $x = 5$ and a $\sigma_S = 35$ A m² kg⁻¹ ($\mu_0 M_S = 0.18$ T). The authors propose that structural microstrains lead to the increase in the magnetic anisotropy. Cr^{3+} doping shows similar phenomenology to Mn^{3+} doping. Katlakunta *et al* [162] reported a maximum $H_C = 580$ kA m⁻¹ with a $\sigma_S = 30$ A m² kg⁻¹ for a doping $x = 0.3$ in sintered materials obtained by microwave-hydrothermal method. In the low ($x < 0.5$) doping regime the reduction of the magnetization is determined by the substitution of Fe at the 12k, 2a and 4f₁ but for large doping ($x > 0.4$) also by the presence of non-magnetic impurities.

Figure 11. (Top left) The coercivity (H_c), specific magnetization at the field of 1.2 T and remanent magnetization for $\text{SrFe}_{12-x}\text{Al}_x\text{O}_{19}$ (Reprinted from [159], Copyright 2012, with permission from Elsevier). (Bottom left) Magnetic moment for unit formula and H_c of $\text{SrFe}_{12-x}\text{Mn}_x\text{O}_{19}$ (Reprinted from [80], Copyright 2017, with permission from Elsevier). (Right) Secondary electron microscopy micrographs of Al hexaferrite powder: typical platelet particles (a) and equiaxed particles (b) (Reprinted from [160], Copyright 2008, with permission from Elsevier).

The substitution of Sr with RE can give rise to improvements in the coercivity. Mangai *et al* [163] report the substitution of Sr by Sm. They observe the increase of H_C up to values around 477 kA m^{-1} but also the decrease of the magnetization.

Different doping approaches were investigated to improve the results obtained in Al doped hexaferrites [146, 164, 165, 167, 168] Trudel *et al* [146] investigated the Cr doping of Al-SFO. This magnetic cation is slightly smaller than Fe^{3+} in fact the CrAl-ferrite structure is contracted under the substitution. However, the magnetization and coercive field exhibit a not linear dependence of the doping: the σ_s is large (over $80 \text{ A m}^2 \text{ kg}^{-1}$) and H_C smaller than the corresponding values of the SFO. Trusov *et al* [165] and Gorbachev *et al* [166] investigate the simultaneous substitution of Sr for Ca and Fe for Al ($\text{Sr}_{1-x/12}\text{Ca}_{x/12}\text{Fe}_{12-x}\text{Al}_x\text{O}_{19}$) and obtain the largest H_C of 2.86 MA m^{-1} for $x = 5$ up to now published. However, the σ_s was very small, $2 \text{ A m}^2 \text{ kg}^{-1}$. These studies consider that the decrease of the magnetization is correlated to the main occupancy of 12k and 2a sites by the Al^{3+} , while the Ca doping enhances the contraction of the lattice structure, that leads to an increase of the anisotropy field due to the enhancement of the crystal field. Stingaciu *et al* [167] perform similar studies by doping Ca Al-doped ferrites with the magnetic Cr to increase the magnetization

($\text{Ca}_{0.33}\text{Sr}_{0.66}\text{Fe}_9\text{Al}_{3-x}\text{Cr}_x\text{O}_{19}$). They found that the Cr doping produces an increment of the magnetization but a decreasing of the coercive field. The study supports the statement that the doping does not modify significantly the anisotropy of these ferrites. Shekhawat *et al* [168] consider the Fe substitution with Cr^{3+} and Sn^{4+} producing $\text{SrAl}_4(\text{Cr}_{0.5}\text{Sn}_{0.5})_x\text{Fe}_{8-x}\text{O}_{19}$ ferrites by sol-gel auto-combustion method. The doping with these two elements decreases the H_C values but the H_C are always above 796 kA m^{-1} (1 T). These oxides exhibit unprecedented high saturation magnetizations, larger than 0.5 T, even in Al-SFO, much larger than previously reported in Al-doped ferrites and even in bulk SFO.

Finally, we should consider the influence of other factors in the hysteresis loop. As it can be deduced from table 2, different synthesis routes give rise to different doping-dependent magnetic properties. On one hand, this might be due to differences in the doping and the cation occupancy. Furthermore, the resulting material is affected by the nucleation and growth conditions that depend on the synthesis conditions or of the physico-chemical changes induced by the post-annealing treatments. On the other hand, as discussed in the previous section, the particle size also determines the magnetic properties. For example, for Al-doped powders, the single domain length is around 600 nm [155]. We note that in most works reported in

table 2 the H_C is high, likely related to particles in the sub-micrometric regime. Finally, most of these studies show that ferrites particles exhibit a platelet morphology (see figure 11), and hence H_C and H_A are determined both by the composition-dependent magnetocrystalline anisotropy and the shape anisotropy term coming from the morphology and magnetization.

In summary, the strategy of ferrite doping allows obtaining advances in the properties of the M-type hexaferrites, increasing magnetization or coercive field. Zn and Ni based doped ferrites exhibit larger magnetizations due to the cationic occupancy of the spin-down sites. In contrast, Al based ferrites exhibit large H_C due to their low magnetization as the cations occupy spin up sites. Commercial LaCo doped ferrites are an intermediate case in which Co cations occupy both spin-up and down sites giving a weak change of the magnetization but the La induced contraction of the crystal structure and the orbital moment of the Co^{2+} induce an increase of the anisotropy and hence of the coercive field. In addition, the submicrometric dimension of the powders and the platelet morphology determine the final hysteresis of the magnets. The fact that LaCo ferrites, that exhibit a weak improvement of the B_R and H_C with respect to undoped ferrites, have found their way into commercial applications demonstrates the important impact that mild improvements in the properties of hexaferrites can have.

4. Hard-soft composites

Given the limits of single phase improvements of hexaferrites, another venue to get better performance from ferrite-based magnets consists of fabricating composites of the hard hexaferrite with a soft magnetic material [174]. Hard magnetic materials exhibit a robust resistance to demagnetization due to their large magnetocrystalline anisotropy and are the base for permanent magnets. However, their saturation magnetization is often moderate, as is the case for ferrites which are ferrimagnetic materials. On the other hand, soft magnets such as metals (Fe, Co) or soft ferrites (Fe_3O_4 , MnFe_2O_4 , etc) present larger magnetization values than hexaferrites but their often cubic structure only allows for reduced magnetocrystalline anisotropies that lead to small coercivity and remanence values. The underlying idea is that as long as both materials are exchange-coupled, the spins of the soft phase align with that of the hard at remanence, therefore exploiting the larger magnetization of the soft phase to improve the remanence of the composite. The drawback is that the overall coercivity decreases as a consequence of the inclusion of a soft phase, but the model predicted an effective enhancement of the energy product even for lowered coercivities. A plethora of experimental efforts over the last three decades have exploited this effect in order to improve the magnetic properties of composites based on all the hard magnets [175], including of course ferrites. We will review in section 4.2 the main results obtained in systems based on hexaferrites as the hard phase.

In the absence of exchange-coupling, the possibility of exploiting the magnetodipolar interactions inside the magnetic material to align the soft spins with the hard phase has been discussed for a number of years as well [176–178], with a

Figure 12. Hysteresis loops corresponding to rigid and spring type magnetic coupling. (b) Scheme of the spring magnet. Reprinted figure with permission from [67], Copyright 2004 by the American Physical Society.

recent interest in this approach fueled by the European project AMPHIBIAN [179]. Several strategies and the main results obtained thus far using this approach will be reviewed in section 4.2.

Finally, in section 5, the future prospects for hard-soft magnetic composites for use in novel permanent magnets will be discussed.

4.1. Exchange-coupling

Achieving an efficient and robust exchange-coupling at the interface has represented the main challenge. This is due to the fact that orbital overlap and thus structural coherency at the interfaces is required on one hand; and on the other the soft phase size has to be below a certain threshold, indicated by the exchange length of the hard phase. This is typically of the order of a few nanometers for systems based on ferrites as a hard phase, which entails that these strict structural requirements are not straightforward to fulfill, especially in bulk systems on which industrial magnet production relies.

Based on the strength of the exchange interaction at the dissimilar interfaces, three types or regimes of coupling can be defined [67, 180–187]: Rigid, Spring and Decoupled. Figure 12 portrays the resulting hysteresis loops of each regime.

In the rigid coupling regime, the spins of the soft phase are completely and rigidly coupled to the hard spins and magnetization reversal occurs as a collective whole. It is equivalent to having a single magnetic phase with properties averaged between both phases.

In the spring-magnet case, both phases are effectively exchange-coupled at the interface but some of the spins of the soft, generally located further from the interface, are not rigidly coupled to the hard spins. This results in a magnetization reversal where low fields reverse those ‘loose’ soft phase spins, while higher negative fields are needed to reverse

Figure 13. Scheme of the synthesis process of a SFO/Co composite with a microstructure involving a soft nanosized Co phase. Reprinted by permission from Springer Nature Customer Service Centre GmbH: Springer Nature, Journal of Materials Science, [68], Copyright 2019.

the hard phase and rigidly coupled soft spins. In this regime, remanence is increased with the presence of the soft phase as negative fields are required to partially misalign the soft spins. In fact, if a negative field between the switching field of the hard and the soft is applied and subsequently removed, the soft ‘loose’ spins first reverse and then, upon field removal, return back to being aligned with the hard phase. This behavior resembles that of a spring, hence the name of the regime.

The third regime corresponds to the situation where there is absence of exchange-coupling between hard and soft constituents. In this case, magnetization reversal occurs as two completely independent phases. It is important to remark that from a single hysteresis loop, as in figure 12, it is difficult to discern between the spring and uncoupled regimes. In principle, in a spring magnet, an increased remanence M_{comp} with respect to that of the hard phase M_H should be observed; whereas in a decoupled system, the remanence value should comply with $M_H < M_{\text{comp}} < M_S$. However, differences in the alignment of the hard and soft particles could yield misleading observations, such as increases in M_{comp} in an uncoupled system, and additional means of verification should be implemented. One option is to measure the soft switching field $H_{\text{sw-s}}$ for different soft contents or thicknesses. If the system is in the decoupled/spring regime, $H_{\text{sw-s}}$ should be independent/dependent on the soft concentration, respectively. In the following, we will review the main results and developments obtained in recent years in hard-soft composites based on hexaferrites as the hard phase. The section will be divided in non-oxide soft phases and oxide soft phases.

4.1.1. Non-oxide-oxide composites. The use of metals as the soft compound presents the advantage of potentially reaching much higher magnetization values, due to the large magnetization of Fe, Co and their alloys. However, in order to stay within the rigid coupling regime, the soft phase must have a

size below a certain threshold, given by twice the domain wall thickness of the ferrite. This quantity is equal to the exchange length L_{ex} of the hard phase, in this case SFO or BFO. This is a relatively strict requirement for metals, as the ferrite’s exchange stiffness and magnetocrystalline anisotropy values leads to threshold sizes of the order of $2L_{\text{ex}} = 21 \text{ nm}$ [68, 185]. The challenge from a synthesis and processing point of view is considerable, especially taking into account that the permanent magnet industry produces thousands of tons of material per year [7]. First of all, a soft phase in the nanoscale has to be produced and supplied. Second, an intimate and coherent contact at the interface has to be achieved, as exchange-coupling demands the presence of chemical bonds and orbital overlap between hard and soft phases. In this framework, novel materials and innovative design, together with sophisticated synthesis and processing methods are needed.

Volodchenkov *et al* [68] have contributed to the field by designing and synthesizing the microstructures shown in figure 13, consisting of platelet shaped SFO particles surrounded by a thin metal layer, where the SFO platelet-shaped particles have an average size of approximately $1 \mu\text{m}$. The process involves the precipitation of a Co precursor, Co–O–OH, on the SFO surfaces and the subsequent reduction to metallic Co.

Based on these structures, their work develops a dense specimen where the Co is indeed observed to be located at the grain boundaries acting as a shell of thickness in the tens of nanometers range, as shown in figure 14.

The precipitated composite presents a substantial increase in remanence while keeping coercivity very close to that of the pure SFO control sample. The authors quantify the improvement in $(BH)_{\text{max}}$ as 70%, although the dense samples have not been magnetically oriented. Interestingly, they compare the magnetic properties of their novel microstructure to those of a composite of the same composition but fabricated by simply mixing hard and soft powders, with a soft Co phase that is

Figure 14. Scheme of the synthesis process of a SFO/Co composite with a microstructure involving a soft nanosized Co phase. Reprinted by permission from Springer Nature Customer Service Centre GmbH: Springer Nature, Journal of Materials Science, [68], Copyright 2019.

not in the nanoscale and not in close contact with the SFO. The marked decrease in remanence and coercivity compared to both the precipitated composite and the pure SFO sample is a clear indication of the threshold size effect of the soft phase.

Several authors have reported on hard-soft composite systems that reverse as a single magnetic phase [188–190]. Single-step reversal or the absence of discontinuities in the demagnetization curve is one of the fingerprints of rigid exchange-coupling in hard-soft composites. Li *et al* [189] report on exchange-coupled SFO/Fe–B composites. A marked increase in magnetization is observed, along with a substantial decrease in coercivity reaching $H_C = 49.1 \text{ kA m}^{-1}$, which brings forward one of the fundamental issues associated with robust exchange-coupling: the collapse of coercivity [191]. Zhang *et al* [190] fabricate structures based on SFO particles between 100 and 200 nm surrounded by FeN nanoparticles smaller than 20 nm. In their case, $H_C > 320 \text{ kA m}^{-1}$ for the exchange-coupled composites, probably due to the small size of the soft phase. In addition, they report mild enhancements of remanence. Xu *et al* [188] study a very similar microstructure although consisting of FeCo nanoparticles as the soft phase. They also observe slightly lower H_C and a mild increase in remanence. Although results are in all these three cases promising, all studies are performed in non-oriented specimens—i.e. no magnetic field was applied to orient the particles prior to the measurement. It is therefore difficult to exactly quantify what the increase in real remanence and energy product is.

An interesting strategy towards achieving structural coherence and small particle sizes is to produce core–shell hard-soft nanostructures. Several authors have studied these systems using SFO as the hard phase and FeCo as the soft shell. Xu *et al* [192, 193] confirm that the core–shell structures present better properties than simple mixtures of both types of particles, confirming the importance of small soft particle sizes, close contact between phases and avoiding agglomerates. However, no clear improvement in remanence with respect to pure SFO samples is observed. Dong *et al* [194] focus on core–shell microfibers where the robust exchange-coupling is again seen to lead to a relatively drastic decay in H_C . The samples studied in these articles are not magnetically oriented either.

It is important to keep in mind as well that relatively broad distributions of coercivities can exist, for both phases,

in polycrystalline powder samples, as a consequence of the particle size distributions. A broad distribution of switching events can lead to seemingly smooth demagnetization curves that are single-phase like, in systems that are in fact exchange-decoupled [195]. Henkel plots are often employed to further discern the existence of exchange-coupling in these systems [196]. Unfortunately, the study of the magnetic interactions in Henkel plots are only really applicable to Stoner–Wohlfart systems consisting of single-domain isolated particles with a very narrow size distribution [197]. In this framework, careful analysis of the dM/dH derivative can help reveal the switching events in the system.

4.1.2. All-oxide composites. All oxide hard-soft composites, based on SFO and a soft ferrite, have the advantage of being slightly easier to process and densify as a magnet since no oxidation vs reduction issues are to be faced. Sintering can be performed in air atmosphere, contrary to the case of metal-oxide composites. On the other hand, the best magnetization values of soft ferrites $\sigma_S = 100 \text{ A m}^2 \text{ kg}^{-1}$ is approximately half the one of Fe and FeCo $\sigma_S = 220 \text{ A m}^2 \text{ kg}^{-1}$, which makes it challenging to improve the magnetization of SFO by adding moderate amounts of soft phase.

Work has focused on a powder sample of different morphologies and dense sintered specimens. In the case of the former, Afshar *et al* [198] recently studied SFO/ $\text{Ni}_{0.6}\text{Zn}_{0.4}\text{Fe}_2\text{O}_4$ composite samples annealed at 700°C – 800°C . The demagnetization curves show two-step reversal, strongly suggesting that the magnetic phases are not exchange-coupled. As relatively high temperature is applied, there is a good chance that chemical bonds between hard and soft particles exist, but the particle sizes observed in their SEM images show particles that are clearly above the rigid coupling threshold size of 20 nm [68]. As a consequence, coercivity drops from $H_C = 358 \text{ kA m}^{-1}$ for pure SFO to $H_C = 88.1 \text{ kA m}^{-1}$ for 30 wt% $\text{Ni}_{0.6}\text{Zn}_{0.4}\text{Fe}_2\text{O}_4$; while magnetization negligibly improves. Dahal *et al* [199] synthesize SFO/ LaSrMnO_3 nanoparticulated composites with single-step reversal curves and supposedly good exchange-coupling between the phases, which may be the consequence of the small particle size. Although improvements in the remanence-to-saturation ratio

Figure 15. TEM images of the as-synthesized SFO particles (left) and Co-ferrite particles (right). [69] John Wiley & Sons. © 2016 The American Ceramic Society.

Figure 16. TEM and HAADF STEM images of bi-magnetic M/BHF/M sandwich composite nanoplatelets with 1.9 nm ((a) and (c)) and 2.5 nm ((b) and (d)) thick M layer. Adapted from [70] with permission of The Royal Society of Chemistry.

are observed, unfortunately the magnetization of the soft phase is lower than that of SFO, leading to decreases in both coercivity and remanence in the composites.

More sophisticated nanostructures have been investigated as well. Song *et al* [200] studied SFO/Ni_{0.5}Zn_{0.5}Fe₂O₄ hollow microfibers. Although small crystallites were obtained, no

Figure 17. HAADF STEM image of composite M/BFO/M sandwich nanoplatelet. The projected structures along $[10\bar{1}0]$ HF and maghemite along $[101]$ M) are superimposed over the images to illustrate the position of Ba^{2+} and Fe^{3+} ions. Different Fe lattice sites and the HF structure (trigonal 2b, tetrahedral 4f1, octahedral 12k, 2a and 4f2) are marked. Adapted from [70] with permission of The Royal Society of Chemistry.

clear improvements in remanence are reported and the composites appear to be decoupled based on the demagnetization curves. Core-shell BFO/ Fe_3O_4 core-shell-like particles were the focus of Mohseni *et al* work [201]. Larger BFO particles are seen to have their surface covered by small magnetite nanoparticles. The samples show promising results in the sense of a significant increase in overall remanence while coercivity remains roughly unchanged. The authors claim there is hard-soft exchange-coupling but two of the three composite studies clearly show two peaks in the dM/dH derivative while the third one, at a 9:1 hard:soft ratio seems to show a kink in the demagnetization curve. An increase in the energy product is claimed as well, although the samples are not magnetically oriented. Volodchenkov *et al* [202] replicated the interesting microstructure discussed in section 4.1.1. and shown in figure 13, this time using Fe_3O_4 instead of Co as the soft phase. A 37% improvement in $(\text{BH})_{\text{max}}$ is reported, which is indeed supported by a remarkable increase in remanence and an almost negligible decrease in coercivity. Unfortunately once again, the samples are not magnetically oriented which prevents a full assessment of their improved properties.

As far as sintered samples are concerned, Jenus *et al* [69] report on SFO/ CoFe_2O_4 composites. This system cannot be strictly considered a hard-soft composite since CoFe_2O_4 is magnetically hard, although its magnetization value $\sigma_s = 82 \text{ A m}^2 \text{ kg}^{-1}$ is indeed higher than that of SFO. By using SPS, the particle size of both phases, shown in

figure 15, is maintained small while achieving a competitive relative density of 93%. In addition, the samples of this study are magnetically oriented, which allow accurately establishing a remarkable 22% increase in energy product with respect to the pure SFO specimen: from $(\text{BH})_{\text{max}} = 21.9 \text{ kJ m}^{-3}$ to $(\text{BH})_{\text{max}} = 26.1 \text{ kJ m}^{-3}$.

4.1.3. Hexaferrite/maghemite bi-magnetic platelets. The research group (Department for Materials Synthesis) from Jožef Stefan Institute, Slovenia, used a simple water-based method for synthesis of the hard-soft exchange coupled nanoparticles composed of soft-magnetic spinel ferrite (maghemite—M) layers deposited on hard-magnetic BFO platelet cores [70, 71]. Even the composite nanoplatelets were developed for advanced medical applications; they can also be relevant as building blocks for production of permanent magnets. The synthesis is performed at relatively low temperature (60°C), it is environmentally friendly, inexpensive, relies on abundant raw materials and is also appropriate for scaling up to an industrial scale. Nevertheless, it enables excellent control of the composite structure. The spinel ferrite is deposited with a controlled co-precipitation of the ions from aqueous suspension containing the core nanoparticles [203]. For example, the M was deposited with the co-precipitation of $\text{Fe}^{2+}/\text{Fe}^{3+}$ ions using weak hydroxide under an inert atmosphere. The deposition process is controlled by the supersaturation of the precipitating iron species, which is enabled by immobilization of the highly reactive Fe^{3+} ions into the coordination compound with urea $[(\text{Fe}(\text{CO}(\text{NH}_2)_2)_6)(\text{NO}_3)_3]$ [204] and their controlled release with thermal decomposition of the complex. The precipitating iron species heterogeneously nucleates and the M grows exclusively on the basal surfaces of core BFO nanoplatelets present in the reaction suspension. Thus, the product nanoplatelets exhibited an incredibly uniform M/BFO/M sandwich-type structure (figure 16). The thickness of the M layers was very uniform, equal on the both sides of the cores. For lower layer thicknesses, the thickness can be controlled simply by adjusting the ratio between the iron ions and the surface area of the BFO core nanoplatelets (Fe/BFO ratio). For example, by adjusting the Fe/BFO ratio the thickness of deposited M surface layers on the BFO core nanoplatelets, 42 nm wide and $\sim 3 \text{ nm}$ thick, was successfully controlled up to $\sim 2 \text{ nm}$ (see figures 16(a) and (c)). For even thicker M layers the deposition had to be conducted in the two consecutive steps. Namely, with the increased Fe/BFO ratio, the supersaturation of the precipitating iron species increases until individual separated M nanoparticles are formed. The composite nanoplatelets with layer thickness of 2.5 nm were obtained by the deposition in the two steps. The product was fairly homogeneous (see figures 16(b) and (d)), even sometimes the larger composite platelets with much thicker layer ($>10 \text{ nm}$) grown on one side of the core were also present in a very low concentration.

Importantly, the co-precipitation method is not limited to co-precipitation of Fe^{2+} and Fe^{3+} ions. With some adaptation to the process, the Fe^{2+} ions can be exchanged with some other 2-valent ions constituting the spinel ferrites, for example Zn^{2+}

Figure 18. Magnetic hysteresis loops for randomly oriented Sc substituted BFO and composite nanoplatelets with different thicknesses of the M layers (the M/BFO/M platelets are referred to as CNPX.X, where ‘X.X’ marks the thickness of M layer in nm). Adapted from [70] with permission of The Royal Society of Chemistry.

Figure 19. (a) TEM image of composite nanoplatelet synthesized by deposition of maghemite layers onto the larger hexagonal BFO core nanoplatelets, and (b) HREM image with corresponding electron diffraction pattern. Adapted from [71] with permission of The Royal Society of Chemistry.

and Mn^{2+} . For example, zinc-ferrite (ZnFe_2O_4) layers were deposited just by exchanging Fe^{2+} with Zn^{2+} [70].

Details of the unique structure of composite nanoplatelets were revealed based on direct atomic-resolution high-angle annular dark field (HAADF) imaging with a probe aberration corrected STEM [70]. The M was deposited onto the Sc-substituted BFO core nanoplatelets, with relatively narrow distribution of widths, 42 ± 12 nm. Remarkably, the inspection of a large number of the sandwich M/BFO/M composite nanoplatelets revealed that the vast majority of the BFO cores exhibited equal structure across the nanoplatelet. In terms of two structural blocks usually used to describe the BFO structure: a cubic spinel ($(\text{Fe}_6\text{O}_8)^{2+}$) ‘S’ block and a hexagonal ($(\text{BaFe}_6\text{O}_{11})^{2-}$) ‘R’ block, the structure of a vast majority of the BFO core nanoplatelets can be presented with the $\text{SRS}^*\text{R}^*\text{S}$ stacking sequence across the nanoplatelet (figure 17). The structure closely defines the thickness of the nanoplatelets at 3.0 nm. As the BFO nanoplatelets terminate at the basal surfaces with the Fe-only S block, their composition

is Fe-rich corresponding to $\text{BaFe}_{15}\text{O}_{23}$ theoretical chemical formula. Only seldom the core nanoplatelets contained three R blocks (figure 16(c)); they were 4.2 nm thick and exhibited the $\text{BaFe}_{14}\text{O}_{22}$ theoretical composition. Later, systematic study showed that during hydrothermal growth, the BFO nanoplatelets crystalline structure of nanoplatelets always thicken with in a step-wise addition of the RS segments to their initial SRS^* structure [39].

Given the cubic S block termination, the structure of core nanoplatelets closely match with the spinel structure of M layers at the $(0001)_{\text{HF}}\parallel(111)_{\text{M}}$ interface. The S block of BFO structure is actually a slice of the cubic spinel structure. Formation of the low-energy coherent interface promotes heterogeneous nucleation and epitaxial growth of the M exclusively at the basal surfaces of the core nanoplatelets. The M was never deposited at the side surfaces of nanoplatelets, where the two structures do not match. Despite good matching between the two structures, the interface was always clearly visible on the STEM images, suggesting presence of strains. The M layer

Figure 20. Comparison of the magnetic hysteresis for BHF core nanoplatelets (~ 100 nm wide and ~ 8 nm thick, BaM 100 and corresponding composite nanoplatelets (S@BaM 100 measured PER and PAR to their basal surfaces. Adapted from [71] with permission of The Royal Society of Chemistry.

Figure 21. Simplified scheme of the internal fields of a BFO nanoplatelet placed inside a Fe particle. [69] John Wiley & Sons. © 2016 The American Ceramic Society.

was always slightly shifted along the interface relative to the structure of HF core.

The BFO nanoplatelets used as the cores of composite nanoplatelets display strongly anisotropic magnetic properties. Because of the strong uniaxial magnetocrystalline anisotropy, the BFO retains the easy axis oriented parallel to the crystallographic c -axis even when in the form of thin nanoplatelets, that is perpendicular to the nanoplatelet. Actually, the HF nanoplatelets are the only accessible platelet magnetic nanoparticles exhibiting such perpendicular easy axes [18, 66]. The ~ 50 nm wide and 3 nm thick Sc-substituted BFO nanoplatelets exhibited the M_S over $30 \text{ A m}^2 \text{ kg}^{-1}$ and H_C of $\sim 150 \text{ kA m}^{-1}$ [73, 205]. The properties of the nanoplatelets are actually substantial, when we consider the very small thickness of the nanoplatelets, which is just over one-unit cell in the corresponding c -direction of the BFO structure. After

the deposition of M layers, the M/BFO/M composite nanoplatelets retained the single-phase hysteresis loops without any constriction or kink, indicating rigid exchange coupling of the two magnetically-different phases [174, 206, 207]. Figure 18 shows the hysteresis loops for the BFO core nanoplatelets and the composite nanoplatelets with different thicknesses of the M layers [70]. A slightly constricted loop was only observed for the composites with the thickest M layer (CNP2.5). However, the two-phase behaviour was ascribed to the presence of a small amount of the nanoplatelets, which showed much thicker M layers; too thick to maintain the complete coupling with the core. As expected, the σ_S of the composite nanoplatelets increased proportionally to the M layer thickness, from $30 \text{ A m}^2 \text{ kg}^{-1}$ for the BFO core nanoplatelets to $52 \text{ A m}^2 \text{ kg}^{-1}$ for the thickest layer of the CNP2.5 composite. However, the H_C strongly decreased with the deposition of epitaxial M layers. The decrease of H_C was non-proportionally large, much larger than expected according to theory presented by Kneller in 1991 [174]. The H_C decreased for $\approx 80\%$ already for the composite with the thinnest M layer (CNP1.1), where the content of the soft-magnetic phase was only $\sim 33\%$. The decrease in H_C was ascribed to changes in the magnetic properties of BHF cores with the deposition of epitaxial M layers [70]. The changes in the properties of the BFO cores were first suggested from high-temperature measurements of the magnetization. The nanoplatelets were coated with a silica shell prior to the measurement to minimize interparticle interactions and block particle growth. The measurements showed that the Curie temperature (T_C) of the BFO nanoplatelets significantly decreased after deposition of the 1.1 nm thick M layers (T_C before and after the deposition was $\sim 450^\circ \text{C}$ and $\sim 310^\circ \text{C}$, respectively). Since the T_C is an intrinsic property reflecting the magnetic structure of the material, it was concluded that the structure changed after the deposition because of internal stresses induced in the cores by the epitaxial M layers. The influence of internal stresses was confirmed by deposition of weakly magnetic zinc-ferrite layers onto the BFO cores [70]. Expectedly, the deposition of the zinc ferrite decreased the M_S almost proportionally to its content, whereas the H_C of composite strongly decreased, confirming that the internal stresses induced in the core nanoplatelets by the epitaxial layers are very probable reason for the non-proportionally-high decrease in H_C of the M/BFO/M composites.

The large decrease in H_C was not observed when the epitaxial M layers (approximately 5 nm thick) were deposited onto larger BFO core nanoplatelets (approximately 100 nm wide and 8 nm thick) (figure 19). The nanoplatelets were measured after they were aligned with exposing their suspension in a liquid wax to a magnetic field, followed by solidification of the wax with cooling. Figure 20 shows the hysteresis loops for the core and the composite nanoplatelets measured with the magnetic field applied perpendicular (PER) and parallel (PAR) to the nanoplatelets' basal surfaces. The composite nanoplatelets exhibited significantly increased the M_S and M_R , while the H_C was only slightly lower than the H_C of cores. We assume that due to the larger thickness the structure of larger cores was more rigid and therefore less susceptible to stresses compared to the structure of smaller BFO nanoplatelets. Moreover, due

Figure 22. (Left) SEM image showing the SFO particles and the FeCo nanowire and (right) magnetization curves of an oriented ferrite bonded magnet (black curve) and a 70 wt% ferrite—30 wt% FeCo composite bonded magnet (red). Reprinted with permission from [209]. Copyright 2020 American Chemical Society.

to the larger area the nucleation of the M at the larger cores started at several spots on the basal surfaces. With growth the continuous M layers formed, however, they contained a high concentration of structural defects (figure 19(a)). In such a structure it is likely that the stresses due to the mismatch on the interface will be relaxed in the M layers, not significantly influencing the core. The improved M_S and M_R of the bi-magnetic composite nanoplatelets resulted in a large increase of the $(BH)_{\max}$, which was more than two times larger compared to that of the BFO cores (the $(BH)_{\max}$ was 7.2 kJ m^{-3} and 15.2 kJ m^{-3} for the cores and the composites, respectively) [71]. The result is very encouraging from the perspective of possible use of the composite nanoparticles as building blocks of permanent magnets. With optimization of the size of BFO core platelets and the composition and thickness of the spinel-ferrite shell the properties of the composite can be further improved.

4.2. Dipolar coupling

The third coupling discussed in the introduction corresponds to the absence of exchange-coupling between hard and soft phases. In this scenario, the internal dipolar interactions and the demagnetizing fields, especially of the high magnetization soft phases, normally lead to a substantial decrease of both remanence and coercivity. However, in certain conditions, the interphase dipolar interactions have been reported to lead to a shift in the hysteresis loop that entails an increase in remanence and energy product [176]. The idea is that mixing hard and soft with appropriate material's parameters, particle sizes, shapes and distribution, a certain fraction of the soft spins align with the effective H field created by the aligned SFO particles. A competition between the self-demagnetizing field of the soft and the effective 'external' field of the hard occurs.

This strategy has been scarcely pursued, although the idea pioneered by González, Montero and collaborators

[176, 177, 208] has led to the recently finished European project AMPHIBIAN further studying its potential. In their work from 1998, Montero *et al* explain how a BFO platelet inside a bigger Fe particle would create an internal field that, after being magnetized, would oppose the self-demagnetizing field of Fe. Figure 21 schematizes the situation. The simplified picture allows bringing forward the fact that the dipolar energy of the system is clearly lowered if the hard and soft magnetic moments are aligned. As a consequence, the demagnetization is shifted in an exchange-bias-like type of effect. Of course, the magnitude of this effect strongly depends on the size and shape of the particles involved, as the stray field lines distribution will be greatly affected by it, but also on the specific geometrical arrangement of hard and soft particles inside the system [176, 177, 208].

Recent unpublished work by our group has led to the observation of an increase in remanence in perfectly decoupled SFO/Fe magnetically oriented composite powders and injection molded magnets. With the aid of micromagnetic simulations, the results are interpreted as a consequence of a certain degree of alignment of the soft moments in the center of the Fe particles with the internal magnetic field lines of the hard. Unfortunately, the absence of exchange-coupling produces a loss of the squareness in the demagnetization curves that leads to a worse energy product even if remanence is increased.

An indirect way of exploiting the internal magnetodipolar interactions, in this case within the soft phase itself, is to fabricate composites based on high aspect-ratio soft nanostructures. Shape anisotropy can help stabilize large remanence values that can be exploited to fabricate permanent magnets. Recently accepted work [209] by the authors demonstrates a 40% increase in the energy product of SFO/FeCo bonded magnets (with respect to pure SFO bonded magnets), based on nearly spherical SFO particles and a 30 wt% addition of FeCo nanowires. Figure 22 summarizes the main result.

Hard and 'soft' phases are completely decoupled in such a system, with the improvement of properties relying on

the competitive properties of the FeCo nanowires themselves. This work constitutes an extremely promising strategy towards the development of rare-earth-free bonded magnets with improved properties.

5. Conclusions and prospects

In this review, we have gone over recent work dedicated to improving what is arguably the most popular material for permanent magnets, the M-type hexaferrites. Among the different methods that have been proposed, we have first considered the control of the nanoscale size and shape of the constituent particles. Engineering of both holds the promise of increasing coercivity. We believe that the main obstacles concern the realization of bulk magnets while preserving the advantage provided by nanostructuring and the need to find cost-efficient methods for the required size and shape control. So far, the promise of this approach remains to be fulfilled for real world materials.

Apparently the simpler route to improve the properties of M-hexaferrites for use in permanent magnet applications is to substitute cations that replace specific sites of the magnetoplumbite structure. This has produced the current best materials commercially available, the LaCo-doped ferrites, even if the increase of the $(BH)_{\max}$ is small (10%). While further improvements are sought, the results are somewhat disappointing in terms of the published effort and reproducibility. This can be attributed to the difficulty in populating specific cation sites, a problem which is common to many other doped-oxides. Even if such a goal is achieved, the doping approach is hampered by how interrelated the magnetic properties are. Increasing the magnetization or the coercivity by substituting some cations in specific sites usually decreases the anisotropy or the magnetization, giving rise in both cases to modest improvements in the $(BH)_{\max}$ energy factor. The positive results obtained so far in terms of M_S and H_C enhancement leave room for improvement. This will probably require considering the structural and magnetic design with multi-cationic oxides, controlling the preparation route to get materials with suitable morphology to play with the shape anisotropy, and reducing the particle size in the submicrometric region to obtain higher coercivity. This will be helped by the ability to perform first principles calculations to guide the experiments. It is clear that the developed doped materials such as the Al-doped hexaferrites with high coercivity should be considered in application design to substitute rare earth magnets. We note that hybrid sintered magnets of Al-SFO and SFO have been recently obtained [210].

Concerning exchange-coupled hard/soft magnetic systems, promising results have been obtained both based on metals and oxides as the soft phase. Unfortunately, and this is a factor shared also by doping experiments, most of the reports focus on non-oriented, often not dense samples. In this scenario, a real assessment of the potential of these materials is lacking at the moment. The nature of the interphase magnetic coupling strongly depends on particle size, shape and bonding to the dissimilar phase, and thus the processing of the specimens

to obtain dense oriented magnets is a critical step that needs to be addressed urgently.

As is the case for many other hard/soft systems based on other hard phases, it can be stated that results have been slightly disappointing thus far. It is the opinion of the authors of this review that the fact that interphase exchange-coupling can lead to a substantial decay of the coercivity [5, 37, 38] may explain part of the situation. The soft phase acts as an extremely effective source of domain walls in the reversal process that rapidly propagate and produce this collapse. In this sense, strategies towards implementing lower degrees of exchange-coupling that do not facilitate domain wall propagation from the soft to the hard particles are very interesting. These observations contradict many of the initial theories developed for hard-soft composites, which brings forward the challenge associated with properly modelling these systems. Up to now, models have been either too simplistic or too complicated to effectively predict behavior. The future may lie in the hands of analytical models in this sense.

Another reason for the disappointing results may come from the extremely severe structural and geometrical requirements that have to be met in order to obtain effective and robust exchange-coupling. In hexaferrite based composites, the soft phase has to have a size below 20 nm and be coherently bonded to the ferrite particles. In addition, the advances in synthesis techniques of nanostructures has led to beautiful work and sophisticated systems that hold promise for the future. In this sense, the synthesis and processing challenge to realize such structures at thousands of tons per year rate is huge. In general, scaling up the synthesis of the next generation ferrite-based systems may be the single most important challenge to address, and collaborations involving both research and industrial players are crucial in order to be successful.

The development of improved ferrite-based magnets needs to take into account the importance of fabricating materials that are orientable by a magnetic field as well. The raw materials powders have to be in an de-aggregated state and, ideally, the individual particles have to be single-crystalline or at least host a stable single-domain with uniaxial anisotropy. This will enable effective orientation under an external magnetic field before/during consolidation onto a dense body, which is crucial to ensure large energy products.

Also, the consolidation/densification step needs to be considered carefully. This is especially relevant for multiphase systems such as the composites discussed in section 3, as the processing needs to preserve the different phases. For instance, in the case of metal-oxide composites, novel sintering processes (cold sintering, pressure assisted methods) that prevent metal oxidation have to be implemented.

A hot topic is the design of 3D magnetic materials and magnets in particular [211, 212]. Additive manufacturing will allow the development of novel magnets and implement them in different applications for electronics, sensing, medicine and automotive. Polymer-bonded technology is employed in some of these applications, but the 3D printing is being considered to get better near-net features and it could allow creating complex molded structures coupled to other properties like shape memory [213]. In addition, additive manufacturing is focused

also on the development of 3D sintered ones. Advanced ferrites, as those here discussed, will allow the replacement of RE-compounds but also the design of the structures with different properties (higher remanence, higher coercivity or different magnetic orientation) to be incorporated in the 3D structures. Actually, early in-lab tests employ bulk ferrite [214] but optimized rheological properties of inks and precursor fibres could be obtained with the nanoparticles.

As a general prospect from the application point of view, it is worth noting that due to the promising results obtained in recent years related to the development of ferrite-based technologies, even moderate improvements in ferrite magnets could lead to substantial substitution of rare-earths in the market. In motors and generators, mild remanence enhancements can lead to significantly increased power output. In this sense, joint efforts both from the magnet side and the application side are important, as the ultimate goal of substitution will be achieved by fabricating a competitive final device.

Permanent magnets play a crucial role in low carbon technologies such as renewable energy generators and electric car motors and components. Thus, the pressing transition to a greener, more sustainable, future is expected to boost the demand in the next 10 years. This transition will be incomplete if we solely rely on rare-earth magnets due to the environmental impact associated with their mining. For these reasons, we expect ferrite-based motors for future mobility and generators for renewable energies to proliferate in the next few years.

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