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Synthesis of mesoscopic particles of multi-component rare earth permanent magnet compounds

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ABSTRACT

Multielement rare earth (R)–transition metal (T) intermetallics are arguably the next generation of high-performance permanent magnetic materials for future applications in energy-saving and renewable energy technologies. Pseudobinary Sm₂Fe₁₇₋ₓNₓ and (R,Zr)(Fe,Co,Ti)₁₂ (R = Nd, Sm) compounds have the highest potential to meet current demands for rare-earth-element-lean permanent magnets (PMs) with ultra-large energy product and operating temperatures up to 200°C. However, the synthesis of these materials, especially in the mesoscopic scale for maximizing the maximum energy product ([BH]ₘₚₓ), remains a great challenge. Nonequilibrium processes are apparently used to overcome the phase-stabilization challenge in preparing the R–T intermetallics but have limited control of the material’s microstructure. More radical bottom-up nanoparticle approaches based on chemical synthesis have also been explored, owing to their potential to achieve the desired composition, structure, size, and shape. While a great achievement has been made for the Sm₂Fe₁₇₋ₓNₓ progress in the synthesis of (R,Zr)(Fe,Co,Ti)₁₂ magnetic mesoscopic particles (MMPs) and R–T/T exchange-coupled nanocomposites (NCMs) with substantial coercivity (Hₑ) and remanence (Mₑ), respectively, remains marginal.

1. Introduction

The current high-end permanent magnet, Nd₂Fe₁₄B (P4/mm/m) compound, has a relatively low Curie temperature Tₛ of 313°C [1,2], and, since its sintered magnet (Nd₂Fe₁₄B: 0.982 vol.%, O₂: 600 ppm, grain orientation: 0.991) has reached the room-temperature ([BH]ₘₚₓ) of 474 kJm⁻³ [3], approaching the theoretical limit of 520 kJm⁻³, high-performance permanent magnetic compounds that outperform the Nd₂Fe₁₄B and operate at elevated temperatures (typically 200°C) for highly efficient electric motors and generators are ever-increasing demand [4–8]. Uniaxial magnetocrystalline multielement R–T intermetallics are arguably the exclusive candidates that can process ultra-large intrinsic magnetic properties, where strong spin-orbit coupling (SOC) of 4f electrons of R sublattice originates large uniaxial magnetocrystalline anisotropy field, Hₐ, and large magnetic moment and strong exchange interactions of 3d electrons of T sublattice result in large saturation magnetization, Mₛ, and high Tₛ, respectively, [9,10]. Among them, R2T₇ (Th₂Zn₁₇-type, R = Sm) and RT₁₂ (ThMn₁₂-type, I₄/mmm) compounds have the potential to meet current demands for rare-earth-element-lean PMs, owing to their intrinsic magnetic properties superior to those of the Nd₂Fe₁₄B compound (Figure 1) [1,2,11–22].

Magnetism on the mesoscopic scale, which is known as micromagnetism, exhibits particularly rich extrinsic behavior. Hₑ is an extrinsic property of crucial importance in permanent magnetism and is governed by the real structure of materials under Brown’s paradox [23,24]: Hₑ is reduced to aHₑ by defects, where the factor α (0 ≤ α < 1) describes microstructural details [25–29]. The MMPs, especially magnetic nanoparticles (MNPs), are an important class of magnet building blocks that can be used to fabricate high-performance anisotropic PMs [30–32]. Their unique feature is the size-dependent coercivity: Hₑ of a single-domain grain increases beyond the superparamagnetic critical size (Dₛₚ) as Hₑ ≈ 1 − (Dₛₚ/D)³/₂, reaches the
maximum at the single-domain critical size ($D_{sd}$) given by $D_{sd} \approx 72\sqrt{A_{ex}K_{1}/\mu_0M_s^2}$ where $A_{ex}$ is the exchange stiffness and $K_1$ is the first anisotropy constant, and then decreases as $H_s \sim 1/D^3$, provided that the grain has a strong cubic anisotropy [33–36]. The grain-size dependent coercivities of representative $R_2\text{Ti}_1\text{Nd}$ and $RT_1\text{Nd}$ compounds are named a few and shown in Figure 2 [37–50]. Owing to the phase-stabilization challenges, control over the microstructure of $R$–$T$ multielement materials is still a non-trivial task, though the $R$–$T$ permanent magnetic materials have been established since 1960s [51] and the Nd$_2$Fe$_{14}$B compound has been utilized since its discovery in 1984 [52–55].

Synthesis of multielement $R$–$T$ intermetallics, especially the $RT_{12}$ compound, is very challenging due to their complex crystal structure, desired phases formed in narrow compositions and at very high temperatures (700–1200°C), and poor chemical stability in the air environment [56–60]. In general, the $R$–$T$ intermetallics with equilibrium phases can be synthesized by equilibrium processes under the framework of their equilibrium phase diagrams, such as cooling of the alloying liquid with a very low cooling rate and annealing of the as-casting ingots at elevated temperatures for a time as long as possible. The later process utilizing arc-melting or induction-melting and subsequent annealing is convenient to synthesize the intermetallics [12,13,16–19,22,37,58,59,61]. However, the equilibrium processes often lead to the formation of impurities because the strict equilibrium conditions to give pure phases are hardly realized, and/or the metals $R$ and $T$ can easily form several equilibrium phases, thus make the microstructure, especially the size, less controllable. In contrast, the nonequilibrium processes are appropriate for synthesizing not only metastable compounds but also the intermetallics with desired crystal structures free from impurities and a fine grain microstructure [61]. The most typical method is first to form the amorphous phase, followed by annealing at an appropriate temperature [14,40–50]. The annealing evolves the formation of metastable phases, which can be produced at various extreme nonequilibrium conditions, and the dynamical transformation between the metastable and the equilibrium phases, corresponding to the local free-energy minima. The differences between the crystallographic symmetries of the phases result in the differences in the local free-energy minima for the formation of the phases. Other factors, such as the composition and the atomic binding energy, of course, also play an important role in the formation and stability of the phases. The descending sequence of the symmetries for the structures of the $R$–$T$ compounds has been found as follows: Amorphous, CaCu$_5$-type ($P6/mmm$), TbCu$_7$-type ($P6/mmm$), Th$_2$ Ni$_{17}$-type ($P6_3/mmc$), Nd$_5$Fe$_{14}$B-type ($P4_3/mmm$), ThMn$_{12}$-type ($I4/mmm$), Th$_2$Zn$_{17}$-type ($R-3m$), and Nd$_3$(Fe, Ti)$_{29}$-type ($A_2/m$) [56,57,61]. The differences
between the free-energy minima for the formation of the last five equilibrium phases in the sequence may be quite small, depending on the composition of alloys and the condition of the synthetic process, and, thus only one equilibrium phase usually forms as the final one under a given condition of compositions and processes [61]. The CaCu$_2$-type ($P6/mmm$) structure with the highest symmetry among those of the R–T metastable and equilibrium phases is the basic one from which the structures of various R–T compounds can be derived by replacements of the R atoms with a pair of T atoms, which is known as dumbbell atoms, as follows [56,57]:

\[
\begin{align*}
2RT_5 & - R + 2T = RT_{13}(I4/mmm), \\
3RT_5 & - R + 2T = R_3T_{17}(R-3m \text{ or } P6_3/mmc), \\
5RT_5 & - 2R + 4T = R_3T_{20}(A2/m).
\end{align*}
\]

For most of the R–T compounds, the metastable CaCu$_2$-type phases may form in a narrow temperature range and, thus, are hardly observed experimentally. The formation of the CaCu$_2$-type phases can be observed by careful annealing with a very slow heating rate for a short time [46], whereas the metastable TbCu$_2$-type phases form in a sufficiently wide temperature range to be observed in various synthetic processes [14,40,43]. In practice, the metastable compounds usually crystallize at annealing temperatures slightly higher than the crystallization temperature of the amorphous phase, and subsequently can be dynamically transformed into more stable compounds at higher annealing temperatures [61]. Therefore, the choice of the appropriate annealing temperatures under a given condition of compositions and processes is essential to promote the formation of the desired intermetallic compounds. The following processes are those based on the method above: mechanical alloying; mechanical milling including high-energy ball milling (HEBM) and surfactant-assisted ball milling (SABM); rapid quenching/melt-spinning; hydrogenation disproportionation desorption recombination (HDDR) [61]. They are popularly adopted for the massive production of fine powders, although they have limited control of the microstructure of materials. DC magnetron sputtering nonequilibrium process is often used to fabricate metastable RT$_{12}$ films [15,20]. Solid-state and solid-gas reactions are the other two powerful nonequilibrium processes for the synthesis of the metastable and intermetallic compounds [19,37–39], and they are further discussed hereinafter. Among bottom-up synthetic approaches, chemical synthesis is the most versatile method for the preparation of MMPs with controllable composition, structure, size, and shape [30–32,62]. Recent advances in the solution-phase synthesis of MNPs followed by the solid-state reaction have overcome the phase stabilization challenges in preparing R–T intermetallics, leading to the successful synthesis of some binary R–T MMPs (e.g., SmCu$_5$, Sm$_2$Co$_{17}$) with ultra-large room-temperature $H_c$ [62,63]. Herein we present an overview of our ongoing research in the context of other recent developments in the chemical synthesis of the most challenging multielement pseudobinary R$_5$T$_{17}$ and RT$_{12}$ intermetallics with an emphasis on grain size and composition control. This review covers the basis behind the use of nanoparticles (NPs) as precursors for the microstructure control of MMPs and presents the most recent results of Sm$_2$Fe$_7$N$_3$ and (R,Zr)(Fe,Co,M)$_{12}$ (R = Nd, Sm; M = Ti, V, Cr, Mn, Co, Mo, W, Al, Si, Ga) MMPs. The review also summarizes the efforts in the chemical synthesis of magnetically hard/soft exchange-coupled R–T/T NCMs.

2. Synthesis of nanostructured precursors

The high negative reduction potentials of R cations, a large difference in reduction potentials of R and T cations, and high chemical instability of R metals make it impossible to directly synthesize R–T intermetallics by solution-phase chemical reactions. An alternative chemical synthetic approach is to first synthesize nanostructured precursors, which are chemically stable and readily synthesized by solution-phase reactions, followed by R–D reactions of the precursors. Monodisperse nanostructured precursors with controllable composition, size, and shape are an important key to determine the microstructure of MMPs, and they are advantageous depending on their structural fashions such as core@shell, encapsulated, doped, or mixed oxide NPs.

The core@shell nanostructures composed of T metal or T oxide (T–O) cores and R oxide (R–O) shells, namely T/T–O@R–O NPs hereinafter, are usually synthesized via a two-step reaction: the T or T–O NPs are firstly synthesized, followed by the deposition of R–O over the surface of the T or T–O NPs. The Co (Fm-3m) and amorphous Fe NPs with particle sizes of < 10 nm were readily synthesized by thermal decomposition of Co$_2$(CO)$_6$ and Fe(CO)$_5$, respectively, as reported by Sun and co-workers [64,65]. The Co (P6/mmm) nanorods with the length in the range of 200–300 nm and the average diameter of 20 nm could be synthesized by using a Ru-catalyzed solvothermal reaction of cobalt laureate in the presence of hexadecylamine in 1,2-butane diol [66]. Amorphous Fe nanospheres with a particle size of 200 nm were synthesized by Carpenter and co-workers [67], where FeSO$_4$ was reduced to Fe by NaBH$_4$ at room temperature in the presence of sodium citrate. A series of Fe–O NPs with tunable sizes in a wide range from ca. 10 nm to several hundred nanometers were also successfully synthesized by the following procedures. Wüstite FeO (Fm-3m) NPs were synthesized by reductive thermal decomposition of Fe(acac)$_3$ (acac = acetylacetone) with oleic acid (OA) and oleylamine (OAm); their sizes were tuned from 14 to 100 nm by controlling the reaction temperatures [68]. Magnetite Fe$_3$O$_4$ (Fd-3m)
nanocubes were synthesized by thermal decomposition of Fe(acac)_3 in the presence of OA in benzyl ether solvent at 290°C [69]; their sizes were tuned from 20 to 200 nm by varying the OA concentration, as shown in Figure 3 [70]. Hyeon and co-workers developed the ultra-large-scale synthesis of smaller Fe_3O_4 nanospheres with tunable sizes in the range of 5–20 nm by thermal decomposition of Fe(III) oleate in various solvents with different boiling points [71]. The reactivity of nanostructured precursors toward the following R–D reaction can be tailored by adopting NPs with different shapes as the surface energy can govern the interdiffusion of metal atoms. For this purpose, Fe_3O_4 (Fd-3m) NPs with rationally-controlled shapes from a cube, truncated cube, cuboctahedron, truncated octahedron, to octahedron were prepared by varying the concentration of CPC (CPC = cetylpyridinium chloride), as shown in Figure 4 [70]. Once the transition metal core like the Co core was prepared, amorphous Sm–O shell could be deposited by thermal decomposition of Sm(acac)_3 at 300°C in 1-octadecene solvent, as shown in Figure 5(a) [65,66]. The temperature rate was kept as low as 2°C min⁻¹ to avoid homogeneous nucleation of particulate Sm–O NPs. The method has been successfully extended to synthesize Fe_3O_4@Sm–O NPs, as shown in Figure 5(b) [72]. The molar ratio of Fe to Sm could be tuned by adjusting relative amounts of Sm(acac)_3 and Co or Fe_3O_4 NPs.

**Figure 3.** Size evolution of Fe_3O_4 (Fd-3m) NPs. (a,c,e,g) Transmission electron microscopy (TEM) images and (b,d,f,h) size distributions. Reprinted with permission from [70]. Copyright 2020 Wiley VCH.

**Figure 4.** Shape evolution of Fe_3O_4 (Fd-3m) NPs: (a–c) cubes, (d–f) truncated cubes, (g–i) cuboctahedra, (j–l) truncated octahedra, (m–o) octahedra with small size, and (p–r) octahedra with large size. (a,d,g,j,m,p) TEM images, (b,e,h,k,n,q) Scanning electron microscopy (SEM) images, and (c,f,i,l,o,r) size distributions. Insets show the (100) and (111) planes in red and blue, respectively. Reprinted with permission from [70]. Copyright 2020 Wiley VCH.

**Figure 5.** Chemically-synthesized core@shell nanostructured precursors. (a,b) TEM images of Co@Sm–O (Co core: Fm-3m, 8 nm; Sm:Co = 1:4.3 at%) and Fe_3O_4@Sm–O (Fe_3O_4 core: Fd-3m, 79 nm; Sm:Fe = 1:11.5 at%) (b). (a) Reprinted with permission from [65]. Copyright 2020 Wiley VCH. (b) Reproduced with permission from [72]. Copyright 2020 the Chemical Society of Japan.
The T/T–O@R–O nanostructures are highly desirable because their size can be fully tuned in the mesoscopic scale from a few nanometers to one micrometer by well-established solution chemical synthetic methods. However, this strategy has succeeded rather in binary R–T MMPs [65,66]; it may be inappropriate for higher-multielement R–T MMPs since the synthetic process involves multi-step reactions giving a very low overall yield and the R/T ratio is not well-controlled. For better control of the R–T composition and full access to the control of particle size, the encapsulated nanostructures, where R–O and T–O in crystalline and/or amorphous forms are co-precipitated within one nanostructure (namely RT–O NPs hereinafter), can be used as precursors. The SmCo–O (7 nm) NPs were synthesized by thermal decomposition of Sm(OAc)₃ and Co(OAc)₂ (OAc = acetate) by Sun and co-workers, as shown in Figure 6(a) [73]. We successfully extended this method to synthesize multielement NPs, e.g., Sm–O, Zr–O, and Ti–O encapsulated CoFe₂O₄ (Fd-3m) NPs (5 nm), which will be reported in the forthcoming paper. Besides these, the Sm(Co or Fe)–O NPs with tunable sizes in the range of 60–220 nm and different morphologies were synthesized directly from thermal decomposition of Sm(acac)₃ and Co/Fe(acac)₃ (Figure 6(b–i)) [74] or SmCo–olate complex (Figure 6(k–s)) [75]. The SmCo–O NPs with various morphologies, such as Sm(OH)₃–Co nanorods [76] or urchin-like [77], Sm(OH)₃–Co(OH)₂ nanoflakes [78,79], and SmCo–O nanofibers [80,81], could be successfully obtained by sonification [76], hydro/solvothermal reaction [77–79], and electrospinning [80,81]. Once the R amounts in these encapsulated nanostructures are under the critical concentration for the solid-solution formation, the R-doped T–O nanostructures can be synthesized by using procedures which are similar to the above [81–85]. A series of R-doped Fe₂O₃ or Fe₃O₄ (R = Sm, Eu, Gd, Tb, Ho, Er, Y) NPs with different shapes and particle sizes tuned in the range of 5 nm–1 μm could be obtained by thermal decomposition [82,84], hydrothermal reaction [85], and ultrasonication [86]. These precursors with R compositions

Figure 6. Chemically-synthesized nanostructured precursors. (a) TEM image of Sm–O encapsulated CoO (Fm-3m) NPs (7 nm, Sm: Co = 1:3.6). (b–d) TEM images of Sm–O encapsulated CoO (P6₃mc) multipods (Sm:Co = 1:4.5): (b) 60 ±10 ×10 ±3 nm, (c) 110 ±20 ×23 ±5 nm, and 220 ±40 ×45 ±5 nm. (e) HRTEM (high-resolution TEM) image of an enlarged part of a nanorod (inserted). (f) HAADF-STEM (high-angle annular dark-field scanning TEM) image and (g–i) elemental mapping images of Sm (g), Co (h), and O (i) of the multipods shown in (c). (j) TEM image of the multipods (shown in (c)) imbedded into a CaO matrix. (k–m) TEM images of Sm–O encapsulated CoO (Fm-3 m) NPs: (k) 110 ±8 nm, (l) 150 ±12 nm, (m) and 200 ±15 nm. (n) HRTEM image of a section of one 200 nm NP shown in (m), showing a mixture of smaller CoO and Sm–O NPs. (o) HAADF-STEM image and (p–r) elemental mapping images of Sm (p), Co (q), and O (r) of one representative 200 nm NP shown in (m). (s) TEM image of one 200 nm NP (shown in (m)) coated with a 10 nm layer of NGC (N-doped graphitic carbon). (t) SEM image of Fe₂O₃, NdFe₂O₄, and Fe₃(MoO₄)₃ mixed NPs (ca. 200 nm). (a) Reproduced with permission from [73]. Copyright 2020 The Royal Society of Chemistry. (b–j) Reprinted with permission from [74]. Copyright 2020 Wiley VCH. (k–t) Reproduced with permission from [75,98]. Copyright 2020 American Chemical Society.
varied below 20 at.% are suitable for the synthesis of RT$_{12}$MMPs.

The most simple and robust strategy for the control over the composition of the multielement R–T MMPs is to prepare mixed NPs of R–O and T–O (namely [T–O,R–O] NPs hereinafter): the precursors are synthesized via one-pot solution chemical synthesis; their stoichiometry is well determined by the feeding ratio of starting materials. Of Sm$_2$O$_3$/Sm(OH)$_3$, Co/Co–O/Co(OH)$_2$/CoOOH, and/or Fe/Fe–O NPs were usually prepared by reducive thermal decompositions [87], solvolothermal reaction [88], co-precipitation [89–96], and ultrasonication [97]. The sol-gel method is rather simple and efficiently produces a mixture of superfine oxide NPs, where a mixture of R–O and T–O gel is formed by using a poly-network gel process at elevated temperatures, followed by calcination to convert the gel to the superfine oxides [98–100]. By varying the concentration of the R and T ions in the gel, the size of oxide NPs could be controlled in a wide range from ca. 10 nm to several hundred nanometers, as shown in Figure 6(1) [98]. Monodisperse and size-tunable T/T–O@R–O and R–T nanostructured precursors with homogeneous composition have opened horizons for optimization of the microstructure of R–T MMPs since they could be embedded in dispersant matrices (e.g., CaO, graphite oxide GO), as shown in Figure 6(j) [74,78,79], or tightly coated with other layers intact (e.g., CaO, GO, nitrogen-doped graphitic carbon NGC) in core@shell structures, as shown in Figure 6(s) [73,75,76]. While mixed NPs strategy partially succeeded in only the embedding, leading to the limited kinetic control of the microstructure of R–T MMPs [88–92,100]. Although the mixed NPs exhibited poor control of particle size, they were capable to synthesize multielement R–T MMPs owing to feasible control of composition [98].

3. Synthesis of multielement R–T MMPs

Multielement R–T intermetallics, represented by Sm$_2$Fe$_{17}$N$_3$, (Nd,Zr)(Fe,Co,Ti)$_{12}$N, and (Sm,Zr)(Fe,Co,Ti)$_{12}$, with large $H_a$, large $M_s$, and high $T_c$ are the most promising candidates to replace Nd$_2$Fe$_{14}$B for modern permanent magnet applications (Figure 1). However, their chemical synthesis remains a great challenge due to the high negative reduction potentials of R cations (e.g., Sm$^{3+}$: $E_{\text{red}} = 2.304 \text{V}$, Nd$^{3+}$: $E_{\text{red}} = 2.323 \text{V}$) and very low chemical stability of R metals. Generally, the R and T cations are reduced by strong reducing agents (e.g., CaH$_2$, Ca) accompanied by diffusion of R and T atoms to form R–T intermetallics under high-temperature solid-state reaction conditions, which is known as R–D process. Before the R–D process, it may need to remove organics from nanostructured precursors by calcination to avoid any undesirable formation of carbides and/or C interstitial compounds in the following R–D reaction, and/or adopt H$_2$ pre-reduction of the calcined precursors to promote the R–D reaction. In this synthesis, the R–T MMPs encounter common issues of sintering in the R–D reactions at high temperatures and oxidation in air environments. A great strategy is the coating of nanostructured precursors with stable materials, which were mentioned in Section 2, to stabilize MMPs formed in high-temperature solid-state reaction conditions and against oxidation in air. The chemical synthetic approach has been successful to prepare rather simple binary R–T MMPs. Indeed, the SmCo$_5$ and Sm$_2$Co$_{17}$ MMPs with tunable sizes in a wide range from a few nanometers to a few micrometers have been successfully synthesized [65,66,73–78,80,81,87–93,97,99,100]. The resultant MMPs were dispersible in common solvents [74,75,91,100], possessed ultra-large $H_a$ reaching the highest yet reported room-temperature $H_a$ of 7.2 T [91] for any permanent magnetic materials, and were strongly stable against oxidation at elevated temperatures [75]. The chemical synthesis of SmCo$_5$ MMPs has been well documented in a previous review [63]. Thus, the nanostructured precursors are readily prepared by a wide range of solution chemical synthetic methods and the intriguing results of chemically-synthesized SmCo$_5$ MMPs are triggering the chemical synthesis of MMPs of Sm$_2$Fe$_{17}$N and (R,Zr)(Fe,Co,M)$_{12}$ ($M =$ Nd, Sm; $M =$ Ti, V, Cr, Mn, Co, Mo, W, Al, Si, Ga) compounds. The chemically synthesized MMPs of these compounds and their room-temperature magnetic properties are summarized in Table 1.

3.1. Sm$_2$Fe$_{17}$N MMPs

The Sm$_2$Fe$_{17}$ (Th$_2$Zn$_{17}$-type, R-3 m) compound, representative to the R$_2$T$_{17}$ intermetallic series, has a relatively low $\mu_0M_r =$ 1.03 T, very low $\mu_0H_C < 1$ T, and very low $T_c =$ 116°C [100–102]. The interstitial doping of the Sm$_2$Fe$_{17}$ compound with nitrogen atoms forms the Sm$_2$Fe$_{17}$N$_3$ (Th$_2$Zn$_{17}$-type, R-3 m) compound [19,101–103]; its intrinsic magnetic properties enormously increases to $\mu_0M_r =$ 1.57 T, $\mu_0H_C =$ 26 T, and $T_c =$ 473°C [19], superior to those of the Nd$_2$Fe$_{14}$B compound (Figure 1). Nitrogen atoms interstitially doped in the 9e octahedral sites (Figure 7) expand the unit-cell volume and increase the Fe–Fe exchange interactions, resulting in the increases in $M_s$ and $T_c$, respectively [102–104]. The hybridization of Sm–f states and N–p states changes magnetization direction from the easy-plane in the Sm$_2$Fe$_{17}$ to the easy-c axis in the Sm$_2$Fe$_{17}$N$_3$ and affects band energy, leading to a large uniaxial magnetocrystalline anisotropy [104]. The Sm$_2$Fe$_{17}$N$_3$ phase is metastable and, thus synthesized by nitridation of the Sm$_2$Fe$_{17}$ with N$_2$ or NH$_3$ in gas-solid reaction or with melamine ($C_6H$_4$N$_6$)
Table 1. Synthetic details and room-temperature magnetic properties of Sm\textsubscript{2}Fe\textsubscript{17}N and (R,Zr)(Fe,Cu,M)\textsubscript{12} (R = Nd, Sm; M = Ti, V, Cr, Mn, Co, Mo, W, Al, Si, Ga) MMPs synthesized by a reduction-diffusion (R–D) process. \( ^{\text{①}}\) T/T–O@R–O NPs: core@shell structure. RT–O NPs: encapsulation, and [T–O,R–O] NPs: mixture of oxide NPs. \( ^{\text{②}}\) Gas-solid nitridation. \( ^{\text{③}}\) High pressure. \( ^{\text{④}}\) Solid-state nitridation. \( ^{\text{⑤}}\) Dehydrogenation after the rinse with H\textsubscript{2}O. \( ^{\text{⑥}}\) Slow-oxidation of Ca residue before the rinse with H\textsubscript{2}O. A typical chemically-synthetic procedure for the synthesis of multielement R–T MMPs composed of calcination, pre-reduction by H\textsubscript{2}, and (R–D) process using Ca or CaH\textsubscript{2} as reductant.

| Materials | Structural fashion\( ^{\text{①-⑥}}\) | Size (nm) | Calcination temperature (°C) | H\textsubscript{2} reduction temperature (°C) | R–D reaction (Nitridation) temperature (°C) | MMPs Size (μm) | \( T_c (°C) \) | \( M_r \text{ (emu} \ g \text{) \text{ }^-1)\) | \( \mu_B H_m (T) \) | \( \mu_B H_m (T) \) | Ref. |
|-----------|---------------------------------|------------|-----------------|-----------------|---------------------------------|-------------|----------|-----------------|-------------|-------------|-----|
| R\textsubscript{8}Fe\textsubscript{17} (Th\textsubscript{2}Zn\textsubscript{17}, R–3 m) | \[Fe\textsubscript{8}O\textsubscript{16}Fe\textsubscript{16}SmFe\textsubscript{O} \textsubscript{36}\] | ca. 100 | 900 | 700 | 900 (430 \( ^{\text{②}} \)) | 0.7 \( \pm 0.0 \) | – | – | – | – | 2.32 | [39] |
| Sm\textsubscript{2}Fe\textsubscript{17}N | \[Fe\textsubscript{2}O\textsubscript{3}Fe\textsubscript{16}SmFe\textsubscript{O}\] | ca. 100 | 1000 | 900 | 1000 (430 \( ^{\text{②}} \)) | 3.5 \( \pm 1.3 \) | – | – | – | – | 1.3 | [72] |
| Fe\textsubscript{2}O\textsubscript{3}Sm–O NPs | ca. 30 | 1000 | 900 | 900 (430 \( ^{\text{②}} \)) | 1.9 \( \pm 1.0 \) | – | – | – | – | 1.3 | [74] |
| RFe\textsubscript{12} (Th\textsubscript{4}Mn\textsubscript{12}, R–3 m) | \[Fe\textsubscript{2}O\textsubscript{3}NdFe\textsubscript{O} \textsubscript{3}Fe\textsubscript{16}Mo\textsubscript{O} \textsubscript{36}\] | ca. 200 | 1000 | 900 | 900 (430 \( ^{\text{②}} \)) | 0.47 \( \pm 0.09 \) | – | 132 | – | 89 | 2.47 \( ^{\text{⑤}} \) | [95] |
| NdFe\textsubscript{14}O\textsubscript{12} | \[Fe\textsubscript{2}O\textsubscript{3}NdFe\textsubscript{O} \textsubscript{3}Fe\textsubscript{16}Mo\textsubscript{O} \textsubscript{36}\] | ca. 200 | 1000 | 900 | 900 (430 \( ^{\text{②}} \)) | 0.6 \( \pm 0.2 \) | – | 134 | – | 100 | 2.28 \( ^{\text{⑤}} \) | [96] |
| NdFe\textsubscript{14}O\textsubscript{12} | \[Fe\textsubscript{2}O\textsubscript{3}NdFe\textsubscript{O} \textsubscript{3}Fe\textsubscript{16}Mo\textsubscript{O} \textsubscript{36}\] | ca. 200 | 1000 | 900 | 900 (430 \( ^{\text{②}} \)) | 0.9 \( \pm 0.3 \) | – | 142 | – | 110 | 1.81 \( ^{\text{⑤}} \) | [97] |
| NdFe\textsubscript{14}O\textsubscript{12} | \[Fe\textsubscript{2}O\textsubscript{3}NdFe\textsubscript{O} \textsubscript{3}Fe\textsubscript{16}Mo\textsubscript{O} \textsubscript{36}\] | ca. 200 | 1000 | 900 | 900 (430 \( ^{\text{②}} \)) | 0.6 | – | – | – | – | 2.78 \( ^{\text{⑥}} \) | [96] |
| NdFe\textsubscript{14}O\textsubscript{12} | \[Fe\textsubscript{2}O\textsubscript{3}NdFe\textsubscript{O} \textsubscript{3}Fe\textsubscript{16}Mo\textsubscript{O} \textsubscript{36}\] | ca. 200 | 1000 | 900 | 900 (430 \( ^{\text{②}} \)) | 0.9 | – | – | – | – | 2.37 \( ^{\text{⑥}} \) | [96] |
| NdFe\textsubscript{14}O\textsubscript{12} | \[Fe\textsubscript{2}O\textsubscript{3}NdFe\textsubscript{O} \textsubscript{3}Fe\textsubscript{16}Mo\textsubscript{O} \textsubscript{36}\] | ca. 200 | 1000 | 900 | 900 (430 \( ^{\text{②}} \)) | 1.5 | – | – | – | – | 2.01 \( ^{\text{⑥}} \) | [96] |
in a solid-state reaction under high temperature and/or pressure conditions. In practice [72], the $\text{Fe}_3\text{O}_4 @ \text{Sm-O}$ NPs (ca. 30 nm) described in Section 2 (Figure 5(b)) were calcined at 1000°C for 1 h in air (Figure 8(a,b,g(i))), and subsequently reduced by $H_2$ at 900°C for 1 h (Figure 8(c,d,g(ii))). The resultant NPs were converted into $\text{Sm}_3\text{Fe}_{17}$ MMPs by the Ca R–D reaction at 900°C for 1 h in Ar, subsequently converted into $\text{Sm}_3\text{Fe}_{17}\text{N}_3$ MMPs by in situ nitridation at various temperatures for 10 h in $\text{N}_2$ (Figure 8(e,f,g(iii))). The $H_c$ of $\text{Sm}_3\text{Fe}_{17}\text{N}_3$ MMPs depended on the nitriding temperature, as shown in Figure 8. This is related to thermal transformations in the nitridation process: nitridation in a gas-solid reaction can only proceed at relatively high temperatures (400–500°C); nitrogen concentration in $\text{Sm}_3\text{Fe}_{17}\text{N}_3$ ($0 < x \leq 3$) is a function of nitriding temperature at low temperatures, while the $\text{Sm}_2\text{Fe}_{17}\text{N}_3$ decomposed into $\text{SmN}_3$ and Fe at high temperatures [19,105,106]. For example, the resultant $\text{Sm}_3\text{Fe}_{17}\text{N}_3$ MMPs formed by the nitridation at 435°C have an average particle size of $1.9 \pm 1.0$ μm and a room-temperature $\mu_r H_c$ of 1.3 T in an isotropic sample before the rinse with $\text{H}_2\text{O}$, as shown in Figures 8(e,f,g(iii)) and 9.

In the light of nanostructured precursors, the particle size of $\text{Sm}_3\text{Fe}_{17}\text{N}_3$ MMPs can be controlled.

**Figure 7.** Crystal structure of $\text{Sm}_2\text{Fe}_{17}\text{N}_3$ (Th$_2$Zn$_{17}$, R-3m) compound.

**Figure 8.** Structural analysis of chemically synthesized $\text{Sm}_3\text{Fe}_{17}\text{N}_3$ MMPs (R-3m, 1.9 ±1.0 μm), (a,c,e) SEM images, (b,d,f) energy-dispersive spectroscopy elemental maps (Fe K edge: purple, Sm L edge: yellow), and (g) Rietveld refinement XRD patterns of $\text{Fe}_3\text{O}_4 @ \text{Sm-O}$ NPs calcined at 1000°C for 1 h (a,b,g(i)), subsequently reduced by $H_2$ at 900°C for 1 h (c,d,g(ii)), and $\text{Sm}_3\text{Fe}_{17}\text{N}_3$ synthesized by Ca R–D at 900°C for 1 h in Ar and in situ nitridation at 435°C for 10 h in $\text{N}_2$ (e,f,g(iii)). Inset in (e) shows the grain size distribution of the $\text{Sm}_3\text{Fe}_{17}\text{N}_3$ MMPs. Reproduced with permission from [72]. Copyright 2020 the Chemical Society of Japan.

**Figure 9.** Room-temperature magnetic properties of $\text{Sm}_3\text{Fe}_{17}\text{N}_3$ MMPs (R-3m, 1.9 ±1.0 μm) formed by in situ nitridation of $\text{Sm}_2\text{Fe}_{17}$ MMPs at various temperatures for 10 h. (a,b) M–H curves before (a) and after (b) the rinse with $\text{H}_2\text{O}$. (c) Nitriding-temperature dependence of coercivity (red rectangles: before the rise with $\text{H}_2\text{O}$, blue circles: after the rinse with $\text{H}_2\text{O}$). Repotted with permission from [72]. Copyright 2020 the Chemical Society of Japan.
through thermodynamics based on careful observation of reaction temperature [39]. To this end, a synthesis using a mixture of Fe–O and Sm–O NPs prepared by a sol-gel method, as described in Section 2, was conducted via consecutive calcination, H₂ pre-reduction, and Ca R–D at various temperatures in the ranges of 500–1000°C, 700–900°C, and 900–1000°C, respectively, followed by in situ nitridation at 430 °C. The particle size of the resultant Sm₂Fe₁₇N₃ MMPs was largely dependent on the processing temperature. The first two processes conducted at low temperatures were crucially important to control the final MMP size at the following R–D process as their resultant small NPs facilitated the Ca R–D reaction at low temperatures to obtain small MMPs. The Ca R–D process, in its turn, effectively determined the particle size of the resultant Sm₂Fe₁₇N₃ MMPs as higher R–D reaction temperatures led to larger MMPs due to the sintering of the particles. As a result, the Sm₂Fe₁₇N₃ MMPs exhibited particle sizes tuned in the range of 0.7–3.5 µm and room-temperature μ₀Hₘ of 2.32 T was obtained for the smallest size of 0.69 µm (Figure 2(a), the blue rectangles). However, the syntheses above required an excessive amount of Sm by 25–30 at.% to the 2:17 stoichiometry to compensate for Sm evaporation during the Ca R–D reaction. The thermodynamic control could not proceed to obtain much smaller sizes as the calciothermic reduction must be conducted at temperatures far above the melting point of Ca (845°C). Recently, an efficient kinetic approach that was advanced by the engineering of nanostructured precursors could achieve the formation of Sm₂Fe₁₇N₃ MMPs with better-controlled size and composition by Sun and co-workers [74]. In this synthesis, SmFe–O nanocubes (110 nm) were tightly coated with CaO by thermal decomposition of Ca(acac)₂ at 200°C, calcined at 185°C for 5 h, and reduced by Ca at 850°C for 30 min in Ar. The resultant Sm₂Fe₁₇ MMPs were then mixed with C₆H₆N₆ and annealed at 600°C for 6 h in Ar to form 100 nm Sm₂Fe₁₇N₃ MMPs. The overall synthesis cost an excessive Sm composition of the SmFe–O nanocubes by only 2 at.% to the 2:17 stoichiometry to compensate for Sm loss. The resultant Sm₂Fe₁₇N₃ MMPs were well dispersible in conventional solvents and exhibited room-temperature μ₀Hₘ of 1.54 T in an anisotropic PEG–embedded Sm₂Fe₁₇N₃ (PEG = polyethylene glycol) sample. This Hₘ value was smaller than that reported above because their particle size (100 nm) was smaller than the single-domain critical diameter of Sm₂Fe₁₇N₃, Dₘₐₓ = ca. 390 nm [36]. This method would give the Sm₂Fe₁₇N₃ MMPs with larger sizes approaching the Dₘₐₓ by increasing the particle size of the precursors, leading to ultralarge Hₘ.

### 3.2. RT₁₂ MMPs

NdFe₂₃N (ThMn₁₂-type, I₄/mmm) and Sm(Fe,Co)$_{12}$ (ThMn₁₂-type, I₄/mmm) compounds are the most impressive candidates for rare-earth-element-lean permanent magnets: their thin films were found to possess significantly large $M_t$ and $H_{c}$ (Figure 1), $(BH)_{max}$ theoretical limits of 550 and 630 KJm$^{-3}$, respectively, and high $T_C$ of 550 and 586°C, respectively; all the properties surpass those of the Nd₂Fe₁₄B [15,20,107–109]. Recently, anisotropic Sm(Fe₀.₅Co₀.₅)$_{12}$–B films composed of columnar grains (40 nm) textured with amorphous B intergranular boundary have been realized. The films exhibited a large room-temperature μ₀Hₘ of 1.2 T, μ₀M_r of 1.5 T, and very small temperature-dependent Hₘ, promising excellent stability at $T \geq 150°C$ [110]. Unfortunately, they are metastable phases and, thus could not be realized in bulk for the fabrication of PMs. To this end, one can partially substitute Fe with stabilizing elements M (M = Ti, V, Cr, Mn, Co, Mo, W, Al, Si, Ga) in preferential 8f, 8i, and 8j sites, depending on the stabilizing elements, for example, Ti, V, and Mo in 8i; Co and Si in 8f and 8j; Ga in 8 j, as shown in Figure 10 [12,13,17,18,40–50,106,111–117]. The substitution with a large concentration of M leads to a significant reduction in $M_t$, except for Co, as shown in Figure 1 for Ti. The Co substitution results in the $M_t$ enhancement based on the Slater-Pauling curve, which is elucidated by an increase in the majority-spin state density below the Fermi level [114,117]. An effective strategy that stabilizes the low M-substituted concentration compounds and rationally enhances $M_t$ is to substitute Sm or Nd with other R elements of smaller atomic number (e.g., Zr, Y) in preferential 2a sites (Figure 10), though it reduces $H_{c}$ for Zr, as shown in Figure 1 [16,21,22,117–126]. The substituted compounds have been found to be stable at even very high temperatures [127]. The stabilization by the substitution can be understood as a decrease of local mismatches in interatomic distances in the structure unfavorable for the orbital hybridization [121],

![Figure 10](image-url)
resulting in low formation energies of the substituted compounds [114,126]. Interstitial nitrogen doping into 2h octahedral sites in Nd(Fe,M)_12 (Figure 10) is of crucial importance to enhance \( M_s \) and \( H_c \); it could be explained as the hybridization between N-\( p \) (2b) and Fe-\( d \) (8j) gives raises in the magnetic moment and the crystal field parameter \( r^2 A_k^0 \) [128,129].

Despite such impressive \( H_c \) and enormous research efforts, the RT\(_{12}\) MMPs with substantial \( H_c \) have not been realized yet, the reported \( H_c \) values being still less than 10% of the corresponding \( H_c \) [40–50,111,112,124]. The following critical issues make it extremely difficult to practically realize the hard magnetic potential of RT\(_{12}\) compounds. Firstly, the size and shape of RT\(_{12}\) MMPs have not been well optimized, especially in the sub-micrometer range, as shown in Figure 2(b) [40–50,111,112,124]. Secondly, since equilibrium phases formed along with the typical RT\(_{12}\) phase are ferromagnetic, the conventional methods cannot introduce appropriate non-magnetic intergranular boundary phases, unlike Nd\(_2\)Fe\(_{14}\)B where their intergranular boundary is greatly facilitated by the eutectic reaction with Nd phase [8,21,111,130]. Finally, the RT\(_{12}\) compounds usually melt at relatively high temperatures, making it difficult to fabricate anisotropic magnets through the liquid-phase sintering [8,21,111,130]. Chemically synthesized fine RT\(_{12}\) MMPs with optimum microstructure may become suitable for sintering into fully dense anisotropic magnets, but is greatly challenging to be obtained. In a typical synthesis, a mixture of Fe\(_2\)O\(_3\) (P4\(_3\)2\(_1\)2, R-3c), NdFeO\(_3\) (Pmma), and Fe\(_2\)(MoO\(_4\))\(_3\) (P\(_2\)/c, Pbnm) NPs (ca. 200 nm), as described in Section 2 (Figure 6(i)), reduced by H\(_2\) at 700°C to convert into Fe(Mo) alloyed (Im-3m) and Nd\(_2\)O\(_3\) (C2/m, P-3m1) NPs, and subsequently converted into NdFe\(_{10}\)Mo\(_2\) (I4/mmm) MMPs (ca. 3–8 μm) by CaH\(_2\) R–D at 1010°C for 4 h [Figure 11(a,b)] [98]. The particle size of the oxide NPs was crucial in controlling the particle size of NdFe\(_{10}\)Mo\(_2\) MMPs: the CaH\(_2\) R–D reaction could be carried out at lower temperatures (e.g., 960°C) and for a short time for the smaller oxide NPs (ca. 50 nm), resulting in smaller NdFe\(_{10}\)Mo\(_2\) MMPs [98,131]. The nitridation of NdFe\(_{11}\)Ti was conducted using the same procedure for the synthesis of Sm\(_2\)Fe\(_{17}\)N\(_5\), as described in Section 3.1, but at higher temperatures (550–600°C). As a result, the NdFe\(_{10}\)Mo\(_2\)N MMPs (ca. 3–8 μm) exhibited a room-temperature \( \mu_B H_c \) of 0.35 T [131]. The method has been successfully applied to synthesize a series of ternary, quaternary, and quinary R(Fe,M)\(_{12}\)X (R = Nd, Y, Nb, Tb, Er; M = Ti, Mo, W; Si; X = N, H) MMPs, as shown in Table 1 [131–136]. For the formation of the R(Fe,M)\(_{12}\)X phase almost free from impurities such as Fe and/or TiFe\(_2\) phases, these syntheses were required to compensate the Sm evaporation in Ca R–D process of 2–10 at.% exceeded to the 1:12 stoichiometry, depending on R elements. Since the formation of R(Fe,M)\(_{12}\)X phase required very high temperature (960–1100°C) for long reaction time (4–8 h), the particle sizes of resultant MMPs were in the range of 3–8 μm. The size control through thermodynamics, which is applicable for Sm\(_2\)Fe\(_{17}\)N\(_5\), is no longer effective for RT\(_{12}\) in the mesoscopic scale in such high-temperature reactions. The particle size may be further controlled to some extent through kinetics, in which the mixed oxide NPs are embedded in dispersant matrices (e.g., CaO, GO) to suppress the resultant MMPs from sintering. A more effective
approach to the kinetic control is to adopt encapsulated nanostructures as precursors (Figure 6(a–s)), where all elements are incorporated in a single NPs with homogeneous composition. We have examined the feasibility of this approach for the synthesis of quinary (Sm,Zr)(Fe,Co,Ti)_{12} MMPs using Sm–O, Zr–O, and Ti–O encapsulated CoFe_{2}O_{4} (Fd-3m) NPs (5 nm), which was mentioned in Section 2. As a result, we have successfully synthesized ultrafine (Sm,Zr)(Fe,Co,Ti)_{12} MMPs with partially controllable size. The resulting (Sm,Zr)(Fe,Co,Ti)_{12} MMPs exhibited a relatively large room-temperature $H_c$ and high susceptibility to magnetic-field alignment, resulting in anisotropic bulk magnets with a large $(BH)_{max}$, which will be reported in the forthcoming paper. Taking these potentials into consideration, synthetic prospects will move a step forward in the size control through kinetics, which is advanced by the use of encapsulated nanostructure as precursors, to obtain their particle size in the range below 1 μm.

### 3.3. Oxidation and Hydrogenation of R–T MMPs

The chemical synthesis of R–T MMPs involves calcothermic reduction, resulting in the formation of byproducts (e.g., CaO, CaCO_{3}) and residual reductants (e.g., CaH_{2} or Ca). Therefore, a post purifying process is required; however, it is still a challenge to achieve high purity and improved magnetic properties of R–T MMPs. The purification using water-based solutions as a rinsing agent has been found to form side products and, thus affect the magnetic properties of as-synthesized R–T MMPs to some extent [81]. Generally, water-rinsing induced oxidation and hydrogenation of R–T MMPs are inevitable side reactions. Owing to low ionization potentials, R elements can be easily oxidized to form surface oxides once R–T MMPs are exposed to air environment, especially in acidic solutions, leading to an attenuation of magnetic performance. Recently, a new washing process has been reported for highly effective purification and stabilization against oxidation of Sm–Co and Nd_{2}Fe_{14}B MMPs by Cho and co-workers [81]. In this procedure, a methanol solution of NH_{4}Cl was used as a rinsing agent, in which Ca and CaO react with NH_{4}Cl to form NH3 and methanol-soluble CaCl_{2}. As a result, neither surface oxidation nor any damage to the MMPs was observed by HRTEM characterization, resulting in a substantial improvement of $M_s$, even near a $M_s$ theoretical value for Sm–Co MMPs. For the nanoscale R–T MMPs sensitive to air, the situation is much severe at elevated temperatures. Their protection against long-term and severe oxidation is crucially important for practical applications, especially for high-temperature applications. Recently, Sun et al. demonstrated a new strategy for the chemical synthesis and stabilization of SmCo_{5} NPs for high-performance upnanomagnet applications in a broad temperature range [75]. In particular, the chemically synthesized SmCo–O NPs composed of SmO_{3} and CoO were coated with a layer of NGC (Figure 6(s)), embedded in CaO matrix, and reduced by Ca at 850°C to produce SmCo_{3}@NGC MMPs. The resultant SmCo_{3}@NGC MMPs showed efficient stability against oxidation: they could maintain 99.2% or 98.3% of magnetization after their exposure to air at room temperature for 5 days or 100°C for 48 h, respectively. The protection of R–T MMPs against oxidation is also necessary to prevent the oxidation induced decomposition. It has been found that the (Sm_{0.2}Zr_{0.2})(Fe_{0.75}Co_{0.25})_{11.5}Ti_{0.5} phase was stable at very high temperatures up to at least 1100°C in an almost oxygen-free atmosphere, but was decomposed above 427°C [127].

Hydrogenation of R–T MMPs through the exothermic reaction of residual Ca with H_{2}O in a rinsing process using water-based solutions has been found to considerably reduce $H_c$ of SmFe_{17}N_{3} MMPs [72,95,96]. As seen in Figure 9, the $H_c$ of SmFe_{17}N_{3} MMPs rinsed with distilled H_{2}O is reduced by 40–80% those of as-synthesized ones. This could be understood as H atoms interstitially occupy available 18g tetrahedral sites (Figure 7) to cause a decrease in $H_c$ [137,138]. It was also obviously observed that the dehydrogenation of hydrogen-doped SmFe_{17}N_{3} MMPs by means of annealing in vacuum was ineffective to recover their $H_c$ [95]. As a result, the SmFe_{17}N_{3} MMPs (0.6 μm) exhibited room-temperature $\mu_0H_c$ of 2.8 T, 1.56 T, and 2.28 T in the form of anisotropic samples for as-synthesis, rinse with distilled water, and a combination of the rinse and dehydrogenation in a reduced atmosphere at 200°C, respectively. Slow oxidation of residual Ca to CaO, prior to the rising process, was then applied to avoid the in situ formation of H_{2}, resulting in the $H_c$ preservation of as-synthesized SmFe_{17}N_{3} MMPs [96]. However, the oxidation also generated water-insoluble SmO_{3} and CaCO_{3} impurities, deteriorating $M_s$. The purification process using a NH_{4}Cl/methanol solution, as previously described [81], is greatly capable of selectively rinsing out impurities and avoiding the in situ generation H_{2}; it has been extended to successfully prepare dispersible SmFe_{17}N_{3} MMP [74]. Unlike what was observed in the SmFe_{17}N_{3} compound, hydrogen interstitial doping in preferential 2b octahedral sites in RT_{12} compound (Figure 10) has been found to enhance their $M_s$, $H_a$, and $T_c$ [132–135,139–147]. The increase in $H_a$ was attributed to an increase in the crystal field and a change in the local symmetry of 4f-electron shell along the c-axis, while the increases in $M_s$ and $T_c$ were attributed to an unit-cell-volume expansion and strong Fe–Fe exchange interactions, respectively, beyond the hydrogen interstitial doping [139–147].
4. R-T/T exchange-coupled NCMs

Magnetically hard/soft exchange-coupled NCMs have long been a potential candidate for high-performance permanent magnets since it can possess a large \((BH)_{max}\) which is deduced from large \(H_{c}\) and \(M_s\) of the corresponding hard and soft magnetic constituents, respectively [148,149]. Subject to the R elements crisis, they have the high potential to meet current demands for R-element lean PMs with large \((BH)_{max}\) and operating temperatures of 150–200°C. The effectiveness of exchange-coupled interaction, which is represented by the microstructure factor \(\alpha\) [25–29], is inversely proportional to the ratio \(D_s/\delta_w\), where \(D_s\) is the size of a soft magnetic phase and \(\delta_w = \pi \sqrt{A_{cw}/K_{e}}\) is the domain wall width of a hard magnetic phase [148,149]. An effective exchange-coupled interaction requires the size of the soft magnetic phase being small enough \((D_s < 2\delta_w)\) for anisotropic bulk NCMs) and the size of the hard magnetic phase approaching the exchange length \(l_{ex}\) of the soft magnetic phase [148–151]. These fundamentals have led to two following approaches to the development in the exchange-coupled NCMs based on the hard magnetic materials. One is to adopt materials with a large \(K_1\) but a consequently small \(\delta_w\), such as \(L_1\)-FePt [152], \(Nd_2Fe_14B\) [153,154], \(SmCo_5\) [87,155–158], and \(Sm_2Fe_{17}N_3\) [150], as their large \(H_e\) compensated for a small \(M_s\) corresponding to a small \(D_s\) \((D_s < ca. 10 \text{ nm})\) [148,149]. Another is to adopt materials with a moderate \(K_1\) and a consequently relatively large \(\delta_w\), such as \(L_1\)-FePd [159–161] and \(HfCo_7\) [162], as they had benefitted to \(\alpha\) and \(D_s\), and a large \(M_s\) corresponding to a large \(D_s\) \((D_s < ca. 20 \text{ nm})\) could be compensated for a moderate \(H_e\) [148,149,161,162]. The latter approach is not usually considered, since the gain in \(\alpha\) is overcompensated by the reduction of \(H_e\).

Apart from the microstructure issues above, the most obvious obstacle to the synthesis of R-T/T NCMs is the formation of undesirable phases as the R and T metals can form many equilibrium and metastable phases. The design of magnetically hard/soft exchange-coupled NCMs with the soft phase as an equilibrium phase of the hard phase, which is similar to the one that was proved to be so effective in \(L_1\)-FePd/Fe NCMs [159–161], is a solution. Chemical synthesis of R-T/T NCMs has had very limited success in obtaining large \((BH)_{max}\). The \(Sm_2Fe_{17}N_3\) has a large \(K_1\) of 16.2 MJm\(^{-3}\) [19]; the \(Sm_2Fe_{17}N_3\) (2.4 nm)/\(Fe_{65}Co_{35}\) (9 nm) alternatively multilayered anisotropic NCMs have been theoretically predicted to have a grant \((BH)_{max}\) of 1 MJm\(^{-3}\) (120 MG Oe) [149]. Nevertheless, large \((BH)_{max}\) of \(Sm_2Fe_{17}N_3\) NCMs has never been realized, because there still has no their bulk materials prepared with a grain size of the order of 10 nm, where large \(H_e\) and \(M_s\) could be achieved in their NCMs [150,151]. Since this size range is an unrealistic practice for metastable \(Sm_2Fe_{17}N_3\) little attention has been paid to the \(Sm_2Fe_{17}N_3\) NCMs so far. Owing to it’s a huge \(K_1\) of 17.2 MJm\(^{-3}\), a relatively small \(\mu_s\) of 1.07 T [36], and easy chemical synthesis among R-T intermetallics, the SmCo\(_5\) has drawn considerable attention as a hard magnetic phase [87,155–158]. Generally, the SmCo\(_5\)/Fe NCMs were synthesized by a simultaneous Ca R-D of mixed oxide NPs [87,155,156]; they exhibited very small \(M_s\), resulting in very small \((BH)_{max}\). The main reason for such the reported small \((BH)_{max}\) in the SmCo\(_5\)/Fe NCMs is a lack of an easy-axis alignment of the hard magnetic SmCo\(_5\) phase. It has been demonstrated that the anisotropic FePt/Fe\(_{0.8}\)Ni\(_{0.2}\) NCMs could enhance \((BH)_{max}\) by 224% of that of the corresponding isotropic NCMs [163]. Therefore, chemical synthesis of particulate NCMs, which is similar to the ones that were proved to be so effective in \(L_1\)-FePd/Fe NCMs [161] and \(HfCo_7/Fe_{65}Co_{35}\) NCMs [162], is of crucial importance to fabricate the anisotropic NCMs with ultra-large \((BH)_{max}\). In a typical synthesis [157], particulate SmCo\(_5\)/Co MMs (200 nm, Co-soft phase: 4 wt.% was synthesized by Ca R-D of Sm\(_{4}\)Co\(_{3}\)Ni\(_{3\cdot1}\)]-\(4H_2\) O\(\rightarrow \)GO MMs and Co(acac)\(_2\) additive at 960°C; their isotropic sample showed room-temperature \(\mu_s\)\(H_c\) of 2.07 T, \(M_s/M_{\text{sat}}\) of 0.75, and \((BH)_{max}\) of 80 kJm\(^{-3}\) (10 MG Oe). This small \((BH)_{max}\) resulted from a low fraction of the soft magnetic Co phase (4 wt.% ) and/or too large grain size of the hard magnetic SmCo\(_5\) phase (ca. 200 nm) to an effective exchange coupling [148–151]. It is highly perspective to gain a drastically enhanced \((BH)_{max}\) as the particulate SmCo\(_5\)/Co MMs can be magnetically aligned to form the anisotropic NCMs. In the line with the problem of R-T/T NCMs, the great challenge in the synthesis of SmCo\(_5\)/Fe(or Co) NCMs is to achieve a high fraction of the soft phase since other phases can be formed with the increase in the fraction [155,157]. An effective chemical synthesis avoiding this alloying issue is to disperse SiO\(_2\)-coated Fe NPs in SmCo\(_5\) matrix, which is described in the literature elsewhere [156]. Prospects are presumably better for RT\(_{12}\)/T NCMs, where T is the only equilibrium phase of the RT\(_{12}\) phase at their high fraction. There have unfortunately been no reports on the RT\(_{12}\)/T NCMs yet.

5. Conclusions and prospects

Multicomponent R-T permanent magnetic materials, which are multielement or multi-phase ones, are expected to shape the growth of \((BH)_{max}\) back to the ‘Moore’s law’ after decades of \(Nd_2Fe_14B\) magnets since the theoretical \((BH)_{max}\) limit is given by \(\mu_sM_s^2/4\) while the \(M_s\) can further increase (Figure 1). However, extraordinary achievements in searching for high-
performance magnetic materials with ultra-large intrinsic properties with the aid of theoretical calculations in recent years, especially \((R,\text{Zr})(\text{Fe,Co,Ti})_{12}\) \((R = \text{Nd, Sm})\) compounds (Figure 1), leave a large gap behind them to the fabrication of their corresponding magnets. This lag practically comes from challenges in the synthesis of these compounds with optimum microstructure in the mesoscopic range for maximizing practically relevant extrinsic properties and the introduction of appropriate grain boundaries for the fabrication of their anisotropic magnets. While conventional synthetic techniques that are popular with the production of magnetic material powders remain unsatisfactory, chemical synthetic approach with recent advances in the solution-phase synthesis of nanostructured precursors followed by solid-state reaction may become suitable to overcome the challenges. As nanostructured precursors are used, the chemical approach broadens microstructural control horizons given by thermodynamics and kinetics. The thermodynamics, unfortunately, addresses a limit in the size control of RT_{12} intermetallics, where their formation requires very high temperatures. Prospects seem to be only adopting the kinetic control, where appropriate engineering of nanostructured precursors toward the following solid-state reaction is a crucial key. The ideal nanostructured precursors are particulate NPs composed of all constituents, and they are monodisperse and homogenous in composition. In addition to the precedent challenges in the synthesis of R-T intermetallics, the synthesis of R-T/T NCs faces new problems of achieving the R-T phase with a particle size of the order of 10 nm for effective exchange interactions, and the formation of undesired phases with an increase in the T fraction. As a result, while monodisperse Sm_{2}Fe_{17}N_{3} MMPs with very well-controllable size and ultra-large \(H_{C}\) are ready for the fabrication of anisotropic magnets, progress in the development of \((R,\text{Zr})(\text{Fe,Co,Ti})_{12}\) MMPs and R-T/T NCs remains marginal. Once MMPs with primarily optimum microstructure are synthesized, their anisotropic magnets can be fabricated by a rapid low-temperature current sintering method \([164]\) or an infiltration treatment \([165]\), instead of conventional liquid-phase sintering.

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