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Microstructure evolution and phase analysis of Sm60Ni40 alloy
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Abstract:	The paper investigates the microstructure evolution and phase analysis of the Sm 60 Ni 40 alloy. The arc melted sample retained the high temperature stable (> 600 °C) Sm 7 Ni 3 and Sm 3 Ni 2 phases along with the congruently melting SmNi phase. Annealing the as-cast sample at 630 o C for 100 hours stabilized the high temperature stable (600 °C - 630 °C) Sm 3 Ni 2 phase. Rietveld refinement was performed to resolve the crystal structure of Sm 3 Ni 2 phase and it was observed that Sm 3 Ni 2 phase stabilizes in monoclinic crystal structure (space group: C2/m) with a lattice parameter of a=13.49 Å, b=3.75 Å, c=9.68 Å and β=106.6°. The Curie temperature of the Sm 3 Ni 2 phase was determined to be ~110 K. The high squareness ratio of 82% along with high coercivity of 3 T indicates that the Sm 3 Ni 2 phase possess spontaneous uniaxial magnetic anisotropy.
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From
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To
Prof. H. S. Kim
Editor
Intermetallics

Dear Professor

I am submitting herewith a manuscript titled “**Microstructure evolution and phase analysis of Sm₆₀Ni₄₀ alloy**” by G.Vijayaragavan, D. Prabhu, M. B. Ponnuchamy, K. R. S. Preethi Meher, Ravi Gautam, Mainak Saha, R. Gopalan, K. G. Pradeep, to be considered for publication in Intermetallics.

Sm based binary alloy systems with low melting eutectics are studied with interest as they are considered potential bonding materials for consolidation of Sm-Fe-N magnets at low temperatures. Sm-Ni is one such system with one of the eutectic reactions at 630 °C. In this report, we present the experimental results on the microstructure and phase evolution of the Sm₆₀Ni₄₀ alloy both in as cast and annealed (630 °C for 100 hr) states in comparison with the thermodynamically predicted phase formation sequence at various temperatures. Interestingly, we were able to stabilize significant (80% area fraction) proportion of the high temperature stable Sm₃Ni₂ phase at room temperature. Though the crystal structure of this phase is reported in Open quantum material database OQMD to be rhombohedral based on DFT calculations with certain assumptions, in the present work our experimental observation is providing a **new information i.e. the structure of Sm₃Ni₂ being Monoclinic** in contrast from the one reported in OQMD. We have also resolved the crystal structural of Sm₃Ni₂ phase using XRD Rietveld analysis an information which was not available in literature prior to this work. Further, we have also reported the magnetic properties of this phase in this paper.

We believe the information provided in this paper is novel and would be most suitable to the journal exclusively dedicated to “intermetallics”, and we hope you will find it worthy of publication in your valued journal.

Thank you
Yours sincerely
On behalf of the team of authors,



(D PRABHU)

Highlights of the paper

- The paper reports the structural and magnetic properties of a high temperature stable Sm_3Ni_2 phase, not reported experimentally earlier in literature to the best of our knowledge.
- The structure is determined to be monoclinic (space group $C2/m$) with a Curie temperature of 110 K.

Microstructure evolution and phase analysis of Sm₆₀Ni₄₀ alloy

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Abstract:

The paper investigates the microstructure evolution and phase analysis of the $\text{Sm}_{60}\text{Ni}_{40}$ alloy. The arc melted sample retained the high temperature stable ($> 600\text{ }^\circ\text{C}$) Sm_7Ni_3 and Sm_3Ni_2 phases along with the congruently melting SmNi phase. Annealing the as-cast sample at $630\text{ }^\circ\text{C}$ for 100 hours stabilized the high temperature stable ($600\text{ }^\circ\text{C} - 630\text{ }^\circ\text{C}$) Sm_3Ni_2 phase. Rietveld refinement was performed to resolve the crystal structure of Sm_3Ni_2 phase and it was observed that Sm_3Ni_2 phase stabilizes in monoclinic crystal structure (space group: $C2/m$) with a lattice parameter of $a=13.49\text{ \AA}$, $b=3.75\text{ \AA}$, $c=9.68\text{ \AA}$ and $\beta=106.6^\circ$. The Curie temperature of the Sm_3Ni_2 phase was determined to be $\sim 110\text{ K}$. The high squareness ratio of 82% along with high coercivity of 3 T indicates that the Sm_3Ni_2 phase possess spontaneous uniaxial magnetic anisotropy.

Keywords: Sm-Ni alloy, Annealing, Microstructure, Magnetic property and Crystal structure

1. Introduction

Recently, Sm-based low melting eutectics are being explored in literature as a potential metal binder for consolidation of Sm-Fe-N due to its good wettability and strong reducing ability [1]. In particular, Sm-Cu and Sm-Ni based alloy systems with eutectic melting points below the decomposition temperature of $\text{Sm}_2\text{Fe}_{17}\text{N}_3$ are considered prospective materials.

Sm-Ni binary system consisting of multiple eutectic compositions could be explored for its suitability as an ideal binder alloy. Recently, a few reports emerged investigating the phase formation in Sm-Ni system. Pan et al. [2] studied the Sm-Ni system across the entire composition range and developed the phase diagram using differential thermal analysis and X-ray diffraction techniques. SmNi and SmNi_5 , Sm_3Ni were identified to be congruently melting phases while, SmNi_2 , SmNi_3 , Sm_2Ni_7 , $\text{Sm}_5\text{Ni}_{19}$ and $\text{Sm}_2\text{Ni}_{17}$ phases form by peritectic reaction.

1 The binary Sm-Ni phase diagram has further been evaluated by Xuping et al. [3] and the
2 observations were consistent with the reported literature. Independently, G. Borzone et al. [4]
3 reported the formation of two new metastable phases Sm_3Ni_2 and Sm_7Ni_3 by peritectic reaction
4 which decompose at temperature below 600 and 619 °C respectively. The crystal structure of
5 the Sm_7Ni_3 phase was reported based on the indexing of diffraction pattern to $\text{hp}20\text{-Fe}_3\text{Th}_7$
6 structure type. To the best of our knowledge, no experimental data has been reported for the
7 crystal structure of Sm_3Ni_2 apart from the one available in the open quantum data base which
8 is theoretically predicted structure using density function theory calculations [5][6]. However,
9 intermetallic phases with composition R_3Ni_2 form via peritectic reaction as reported in the
10 literature for $\text{R} = \text{Tb}, \text{Dy}, \text{Ho}, \text{Er}, \text{Y}$ and Gd . The crystallographic information is also available
11 for $\text{Tb}_3\text{Ni}_2, \text{Dy}_3\text{Ni}_2, \text{Ho}_3\text{Ni}_2, \text{Er}_3\text{Ni}_2$ and Gd_3Ni_2 compounds. Monoclinic structure for Tb_3Ni_2
12 and Dy_3Ni_2 [Dy_3Ni_2 type, $\text{mS}20, \text{C}2/\text{m}$] [7], hexagonal for Er_3Ni_2 [Er_3Ni_2 type, $\text{hR}45, \text{R}-3$] [8]
13 and tetragonal structure for Y_3Ni_2 [Y_3Ni_2 type, $\text{tP}80, \text{P}4_12_12$] [9] exist. The Ho_3Ni_2 exist at
14 low temperature [Dy_3Ni_2 type, $\text{mS}20, \text{C}2/\text{m}$] [7] as well as at high temperatures [Er_3Ni_2 type,
15 $\text{hR}45, \text{R}-3$] [8]. Recently, monoclinic structure was reported for Gd_3Ni_2 [Dy_3Ni_2 type, $\text{mS}20,$
16 $\text{C}2/\text{m}$] with lattice parameters [$a = 1.3418, b = 0.372, c = 0.9640 \text{ nm}$ and $\beta = 106.2^\circ$] [10]. In
17 this study, we have investigated the Sm-Ni system close to the eutectic reaction (E3) having
18 substantially lower reaction temperature (623 °C) as indicated in Fig. 1a. i.e. a fixed Ni content
19 of 40 at. % towards understanding the microstructure evolution, phase stability and their
20 magnetic properties.
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49 2. Experimental details

50 The $\text{Sm}_{60}\text{Ni}_{40}$ (at. %) composition was melted using vacuum arc melting technique under inert
51 Argon atmosphere. Elemental Samarium (Sm) and Nickel (Ni) with purity higher than 99.9%
52 was used for preparing the alloy ingots. To compensate for the losses during melting, 1 wt.%
53 excess Sm was added. In order to understand the phase formation as a function of temperature,
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1 Sm-Ni phase diagram was generated using CALPHAD (Calculation of Phase Diagram)
2 approach with the thermodynamic data available elsewhere [NIMS CPDDB
3 (<https://mits.nims.go.jp>)] [11], [12]. The annealing temperature of 630 °C for 100 hours just
4 above the corresponding eutectic reaction temperature (i.e. 623 °C for the composition under
5 study) was determined from the phase diagram in order to obtain a homogeneous
6 microstructure. The arc melted bulk alloy ingot was wrapped in a tantalum foil and sealed in a
7 quartz tube filled partially with argon for heat treatment.
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12 Phase analysis and crystal structure investigation were carried out by X-Ray diffraction (XRD)
13 using Cu- k_{α} radiation (Panalytical, X'pert pro, Netherlands). The XRD patterns were obtained
14 from fine powder samples by crushing the as-cast and annealed ingots. Phase identification of
15 the as-cast sample was performed utilizing crystallographic data from the Inorganic Materials
16 Database (Atom Work) of NIMS, Japan (<https://mits.nims.go.jp>) [13]. The XRD pattern of
17 annealed samples were subjected to Rietveld refinement using Fullprof software for discerning
18 the different phases. Crystal structure information of R_3Ni_2 stoichiometry phases with other
19 rare earths such as Tb, Dy, Ho, Er, Y, Gd is available in literature [10]. The XRD patterns
20 were fit using pseudo-Voigt profile with axial divergence asymmetry. Microstructure imaging
21 and composition analysis was performed using FEG-Scanning electron microscope (SEM)
22 (Zeiss, Merlin Compact, Germany) attached with an energy dispersive X-ray spectrometer
23 (EDS) (EDAX, Octane plus, USA). Three-dimensional elemental distribution analysis at near
24 atomic-scale was performed using a local electrode atom probe tomography (LEAP) (Cameca,
25 5000XR, USA). Site-specific atom probe tomography (APT) tips was prepared using a dual-
26 beam Focused ion beam (FIB) (Thermofisher scientific, Helios G4 UX) following the lift out
27 procedure described elsewhere [14],[15]. APT measurement was carried out in the laser pulsing
28 mode with laser pulse frequency of 250 kHz and 30 pJ pulse energy while the tips were
29 maintained at 60 K. Data reconstruction and analysis was performed using IVAS 3.8.10
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1 software provided by Cameca Inc. The magnetic properties were measured using Physical
2 property measurement system (PPMS) with vibrating sample magnetometer attachment
3 (Quantum Design, Dynacool, USA). The Magnetisation (M) vs Applied Field (H)
4 measurements were carried out from 300 K down to 20 K by applying a magnetic field between
5 +9 T to -9 T. Thermomagnetic (Zero field cooled and Field cooled) measurements were carried
6 out in the temperature range of 20 K to 300 K with an applied field of 0.05 T and 5 T.
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15 **3. Results and Discussion**

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18 Fig. 1a shows the generated equilibrium phase diagram of Sm-Ni. It can be noticed that 3
19 different eutectic reactions namely, E1 (1275 °C), E2 (821 °C) and E3 (623 °C) are possible,
20 and it can be observed that the reaction temperature decreases with increasing Sm content.
21 Table 1 summarizes all the possible reactions and the associated phases that are products of the
22 various reactions [12]. The enlarged portion (in fig. 1b) shows the phase field around E3 where
23 the composition under study lies. The schematic of microstructure evolution as a function of
24 temperature is shown for the desired composition of Sm₆₀Ni₄₀ in fig. 1b. Accordingly, the ideal
25 microstructure at room temperate should consist of SmNi and Sm₃Ni with phase fraction of 60
26 and 40 vol.% respectively. Fig. 2(a) shows the backscattered electron (BSE) image of the as
27 cast sample. Three different phases corresponding to varying contrast (dark, white, and grey
28 phase) are identified. To estimate the composition of the individual phases, EDS line scan was
29 performed along the red dotted line shown in fig. 2(a) cutting across the three phases. The
30 concentration profile obtained from EDS is shown in fig 2(b) and the three phases could be
31 identified as SmNi (dark), Sm₇Ni₃ (white) and Sm₃Ni₂ (grey). The intensity distribution of Sm
32 and Ni in the elemental map (fig. 2 (c & d)) is in agreement with the concentration profile. The
33 as-cast microstructure consisting of high temperature stable Sm₇Ni₃ and Sm₃Ni₂ phases
34 retained along with SmNi suggests non-equilibrium solidification of the molten liquid due to
35 the water-cooling of the copper mold used in the arc melting furnace. According to fig. 1b,
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1 SmNi should be the first solid phase emerging out of the Sm₆₀Ni₄₀ liquid phase at 850 °C as
2 indicated in the schematic. Upon further solidification, a peritectic reaction between the
3 remaining liquid and SmNi solid phase is expected at 630 °C. The observation of Sm₃Ni₂ being
4 spatially separated from the SmNi phase suggests the suppression of peritectic reaction [P1 in
5 table.1]. The remaining liquid therefore should have undergone eutectic solidification at 623
6 °C resulting in the formation of 17 Vol.% Sm₃Ni₂ and 43 Vol.% Sm₇Ni₃ phases. However, the
7 thermodynamically predicted phase fraction of Sm₃Ni₂ and Sm₇Ni₃ phases at 623 °C are 12
8 Vol.% and 46 Vol. % respectively, indicating no further phase change.
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20 Fig 3. shows the XRD powder diffraction pattern of the as-cast sample. Most of the peaks
21 correspond to the SmNi and Sm₇Ni₃ phases. Few unknown peaks were observed which did not
22 correspond to either of the phases. it is plausible that Sm₃Ni₂ phase (detected in SEM analysis)
23 could contribute to diffraction resulting in the observation of additional peaks. However, it
24 should be noted that, peak position was not in agreement with crystal structure predicted in
25 OQMD for Sm₃Ni₂ [5][6].
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35 Therefore, to obtain equilibrium microstructure, the sample was annealed at 630 °C for 100 h
36 based on fig. 1b followed by furnace cooling. Fig. 4 shows the Rietveld refined XRD pattern
37 for the annealed sample which indicates the presence of Sm₃Ni₂ and SmNi phases. The
38 microstructure of the annealed sample is shown in fig. 5a which contains three different regions
39 of varying contrast. The grey contrast region appears to be the major phase while the dark and
40 white regions are present as minor fractions. SEM-EDS line scan was carried out along the red
41 dotted line shown in fig. 5a cutting across all three contrasting regions to obtain their local
42 chemical composition. Fig. 5b shows the concentration profile and based on the contents of
43 Sm and Ni present, the three phases were identified to be Sm₃Ni₂ (~80 % area fraction, grey
44 phase), SmNi (~10 % area fraction, dark phase) and Sm₃Ni (~10% area fraction, white phase).
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60 The elemental maps of Sm and Ni corresponding to the BSE image in fig. 5a confirm the
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1 presence of three phases. The sporadic distribution of equilibrium SmNi and Sm₃Ni phases
2 along with their spatially separated nature suggest that the precipitation of these two phases are
3 due to the partial decomposition of the Sm₃Ni₂ phase below 600 °C. Further, close observation
4 of the Sm₃Ni precipitate (in fig. 5c) revealed an intragranular phase contrast which are distinct,
5 indicative of the coexistence of multiple phases within, possibly in their metastable states.
6 SEM-EDS line scan performed along the dotted line in fig. 5c does not show any significant
7 variation in chemical composition across the metastable region which was also confirmed by
8 the elemental mapping in fig. 5d. To further ascertain their phase constituents, site-specific
9 APT measurement was performed along the multiphase contrast (metastable) region. Fig. 6a
10 shows the elemental distribution map of Ni which appears to be inhomogeneous. 1D
11 concentration profile along a 10 nm diameter cylindrical region of interest indicates local
12 composition variations corresponding to Sm, SmNi, Sm₃Ni and Sm-Ni-O. To further visualize
13 the distribution of Oxygen rich regions, and to determine their local chemical composition, an
14 8 at.% isoconcentration surface was used to delineate the atom map of O as shown in inset of
15 Fig. 6b. The proximity histogram corresponding to one of the O rich regions (marked in
16 rectangle) is constituted with 24.9 ± 0.89 at.% O, 15.8 ± 0.75 at.% Ni and 59.1 ± 1.01 at.% Sm.
17 Significant amount of O seems to have been incorporated in the sample post-annealing even
18 though annealing was performed under controlled Ar atmosphere. Based on the APT
19 determined local chemical composition, it can be inferred that the Oxygen rich regions could
20 be metastable and may not be related to any stable oxide phases. However, the Sm₃Ni₂ is
21 resolved to be monoclinic with space group C2/m. The refined lattice parameter values are $a =$
22 13.487 \AA , $b = 3.754 \text{ \AA}$, and $c = 9.682 \text{ \AA}$ & $\beta = 106.6^\circ$ different from the crystal structure and
23 parameters reported in the OQMD for this structure [5][6]. However, it could be noted that the
24 reported data is similar to isostructural Gd₃Ni₂ reported by et al [10].
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1 Having stabilized significant fraction (80% area fraction) of high-temperature stable Sm_3Ni_2
2 phase in the annealed condition, detailed magnetic property evaluation has been performed.
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4 The temperature dependent magnetization measured with an applied field of 500 Oe for the as-
5 cast alloy is shown in fig. 7a which shows two magnetic transitions. The first transition
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7 identified as T_{C1} at 45 K correspond to the curie temperature of SmNi compound which was
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9 identified as T_{C1} at 45 K correspond to the curie temperature of SmNi compound which was
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11 found to be in good agreement with the value reported in literature [16],[17]. The second Curie
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13 transition identified as T_{C2} at 110 K should be that of either Sm_7Ni_3 or Sm_3Ni_2 , but it could not
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15 be ascertained as the magnetic properties of these high temperature phases are not yet reported.
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17 Fig.7b shows the hysteresis loop measured for the as-cast alloy at 20 K (below T_{C1}), 100 K
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19 (below T_{C2}) and 120 K (above T_{C2}). The hysteresis loop at 20 K exhibited a two-phase
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21 behaviour suggesting that two ferromagnetic phases are magnetically decoupled. At 100 K
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23 (above the T_C of SmNi phase), a well-defined hysteresis loop can be observed with a coercivity
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25 of 1 kOe corresponding to either of Sm_7Ni_3 or Sm_3Ni_2 phases. At 120 K, the hysteresis vanishes
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27 and a linear change in magnetization typical of a paramagnetic behaviour observed since it is
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29 well above the curie temperature of all the ferromagnetic phases present in the as-cast alloy.
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39 Figure 8a. shows the temperature dependent magnetization curve measured for the annealed
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41 sample which contains ~80% (area fraction) of Sm_3Ni_2 phase. The Curie transition at 45 K
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43 (T_{C1} i.e. of SmNi) was observed but only as a small slope change due to the low volume fraction
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45 of the SmNi phase present in the annealed sample. Whereas the T_{C2} remained similar to the as-
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47 cast sample at 110 K. The inset shows the derivative of the M vs T to clearly identify the two
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49 Curie transition temperatures. Based on the SEM microstructure obtained area fraction of
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51 phases and taking into account the thermomagnetic behaviour of the as-cast and annealed
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53 samples, the transition at 110 K should correspond to the Curie temperature of Sm_3Ni_2 phase.
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55 The Sm_7Ni_3 and Sm_3Ni phases present in the as-cast and annealed samples respectively may
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be diamagnetic or have a Curie temperature below 5 K. The hysteresis loop of the annealed sample shown in fig. 8b is similar to the as-cast sample for the 120 K and 100 K. The dominant Sm_3Ni_2 phase present in the annealed sample exhibits a very high coercivity ~ 3 T at 20 K. The high coercivity and squareness (82%) of the hysteresis loop clearly suggests that the Sm_3Ni_2 is a hard magnetic phase with high uniaxial anisotropy. The high magnetocrystalline anisotropy of the Sm_3Ni_2 phase could be due to the Sm ion having a positive Stevens coefficient ($\alpha_J = 4 \times 10^{-2}$) [18] with a prolate shell [19]. The large separation distance between the zero-field cooled and field cooled thermomagnetic measurements carried out at an applied field of 5 T for the annealed sample shown in fig. 8b as inset further confirms the strong anisotropy of the Sm_3Ni_2 phase [20]. Recently, it was reported that Sm based alloys could act as binders for the consolidation of Sm-Fe-N powders. The magnetic properties of the binder alloy which will be present in the intergranular/interparticle regions is crucial and is considered favourable if they are paramagnetic as they can decouple the ferromagnetic interactions. The paramagnetic nature of Sm_3Ni_2 phase at room temperature highlights the possibility of further exploration as a potential binder material towards consolidation of Sm-Fe-N based bulk magnets.

4. Summary

The microstructure evolution, phase formation and magnetic properties of $\text{Sm}_{60}\text{Ni}_{40}$ alloy both in as cast and annealed (630 °C for 100h) conditions were investigated. The major observations are summarized below,

1. Significant fractions (~80% area fraction) of high temperature stable Sm_3Ni_2 can be stabilized at room temperature by annealing the as-cast material slightly above the E3 temperature.
2. The crystal structure of Sm_3Ni_2 phase has been determined using XRD Rietveld analysis to be monoclinic.
3. At room temperature Sm_3Ni_2 phase is paramagnetic which can exhibit high uniaxial magnetocrystalline anisotropy below 110K.
4. The very high coercivity of ~ 3 T observed at 20 K combined with 82% squareness ratio confirms the hard magnetic nature of Sm_3Ni_2 phase.

Based on the above observations, it can be concluded that appropriate heat treatment protocols could be devised to synthesize phase pure Sm_3Ni_2 alloy which offers the potential of utilizing them as binder material for the synthesis of high-performance Sm-Fe-N based bulk magnets.

Table.1. shows reaction in enlarged portion of the Sm-Ni phase diagram [12]

Reaction	Type	Temperature (°C)	ID
$L \rightarrow \alpha - \text{Sm} + \text{Sm}_2\text{Ni}_{17}$	Eutectic	1275	E1
$L \rightarrow \text{SmNi} + \text{SmNi}_2$	Eutectic	821	E2
$L \rightarrow \text{Sm}_3\text{Ni}_2 + \text{Sm}_7\text{Ni}_3$	Eutectic	623	E3
$L + \text{SmNi} \rightarrow \text{Sm}_3\text{Ni}_2$	Peritectic	630	P1
$L + \text{Sm}_3\text{Ni} \rightarrow \text{Sm}_7\text{Ni}_3$	Peritectic	652	P2

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34 **Figure captions**

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38 **Fig. 1** a) Sm-Ni binary phase diagram b) Enlarged region around the $\text{Sm}_{60}\text{Ni}_{40}$ composition
39 with the inset schematic presenting the evolution of phases and microstructure as a function of
40 temperature.
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46 **Fig. 2** (a) SEM-BSE microstructure of the as cast $\text{Sm}_{60}\text{Ni}_{40}$ alloy; (b) 1D concentration profile
47 obtained along the dotted line (red color) in (a); EDX elemental map of (c) Sm and (d) Ni.
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52 **Fig. 3** X-ray diffraction pattern of the as cast sample obtained in powder form.
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55 **Fig. 4** X-ray diffraction pattern (red) of the annealed sample (630 °C for 100h) in powder form
56 overlaid with the Rietveld refined pattern (in black).
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Fig. 5 (a) SEM-BSE microstructure of the annealed sample (630 °C for 100h); (b) 1D concentration profile obtained along the dotted line (red color) in (a) along with the corresponding elemental maps of Sm and Ni. (c) Magnified SEM-BSE image of the Sm₃Ni precipitate showing the metastable intragranular region; (d) 1D concentration profile obtained along the dotted line (red color) in (c) along with the corresponding elemental maps of Sm and Ni.

Fig. 6 a) Elemental distribution map of Ni along with the 1D concentration profile obtained along a 10 nm diameter cylindrical region of interest with 0.5 nm bin width; (b) Oxygen rich regions in the elemental map (inset) delineated with 8 at.% isoconcentration surface and the representative proximity histogram obtained from the rectangular (red color) region with 0.1 nm bin width. Inset SEM-BSE micrograph also shows the region of interest (ROI) from where site-specific APT tips were prepared for composition analysis.

Fig. 7 a) Thermomagnetic curve of the as-cast sample showing two curie transitions at T_{C1} and T_{C2}; b) Hysteresis loop measured at various temperatures showing the magnetic property of the as-cast alloy.

Fig. 8 a) Thermomagnetic curve of the annealed sample showing two curie transitions at T_{C1} and T_{C2} and inset shows the derivative of the curve highlighting the two indicated transitions. b) Hysteresis loop measured at various temperatures showing the high coercivity of the Sm₃Ni₂ phase and inset shows the difference (vertical double headed arrow) between FC and ZFC measured for the annealed sample with an applied field of 5 T exhibiting the anisotropy of the Sm₃Ni₂ phase.

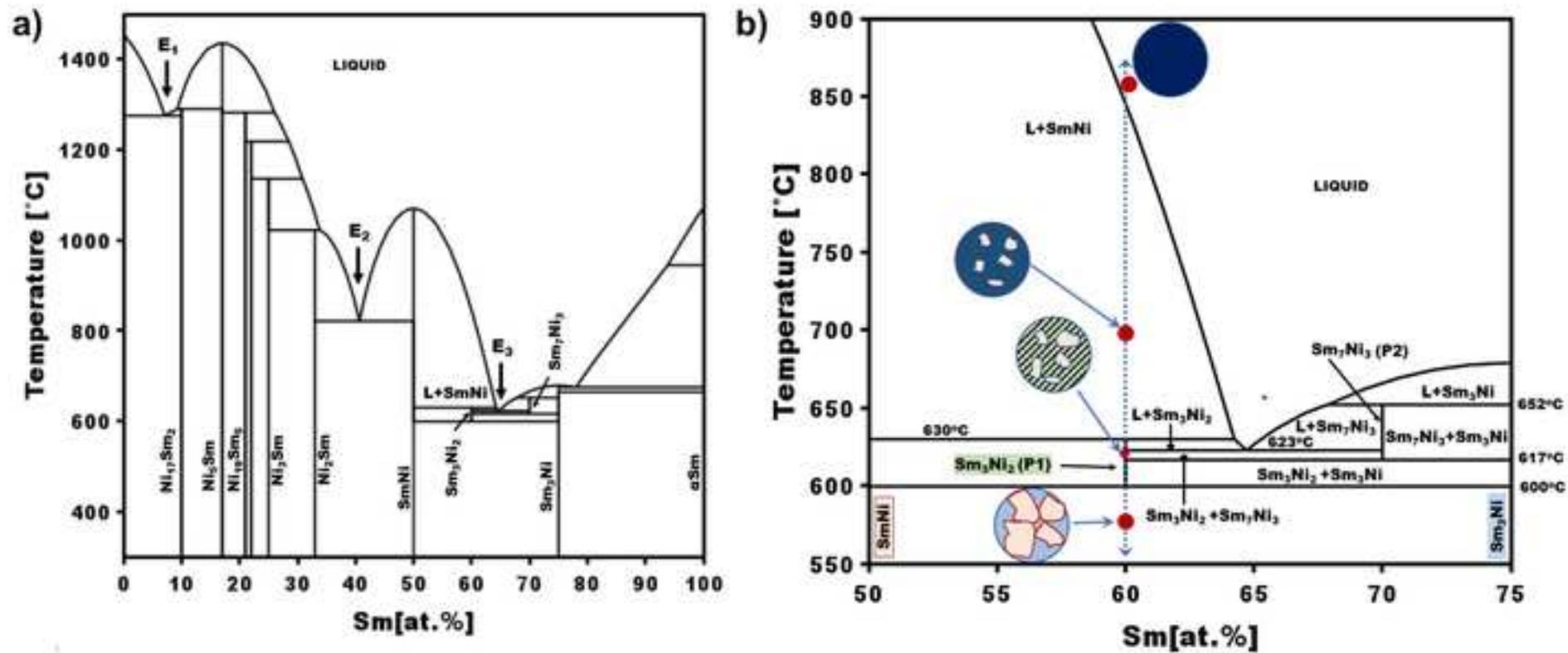
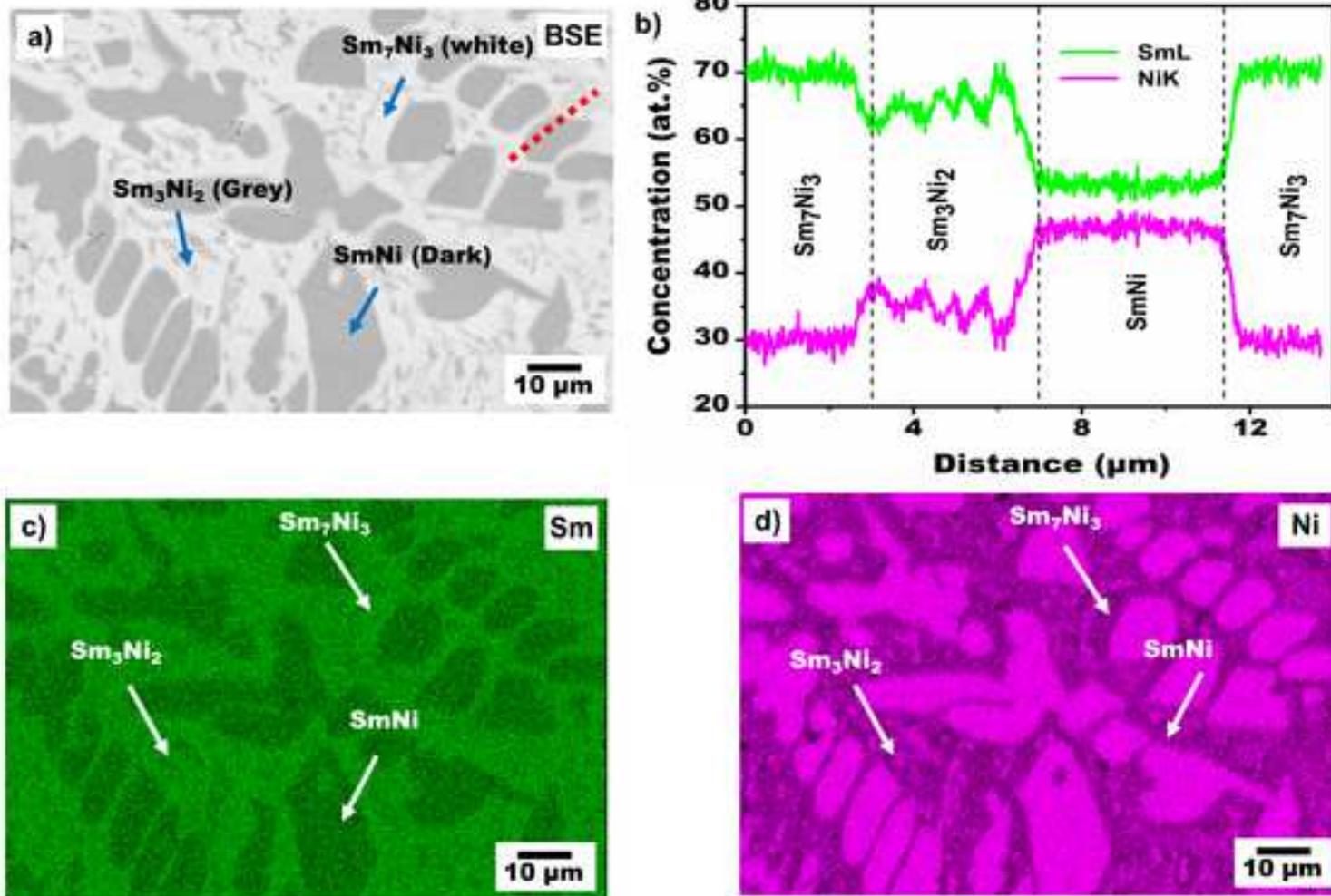


Fig. 1

**Fig. 2**

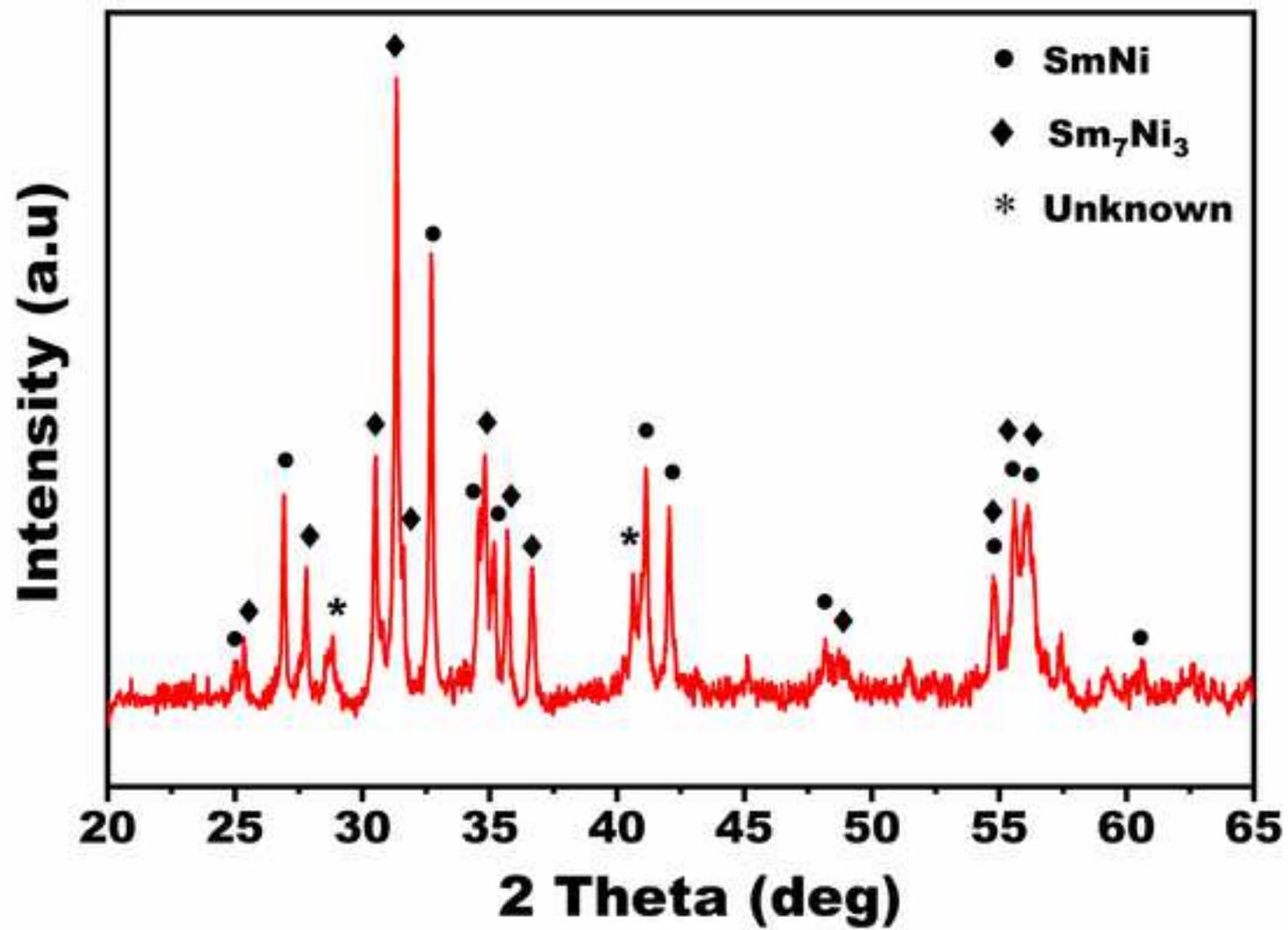
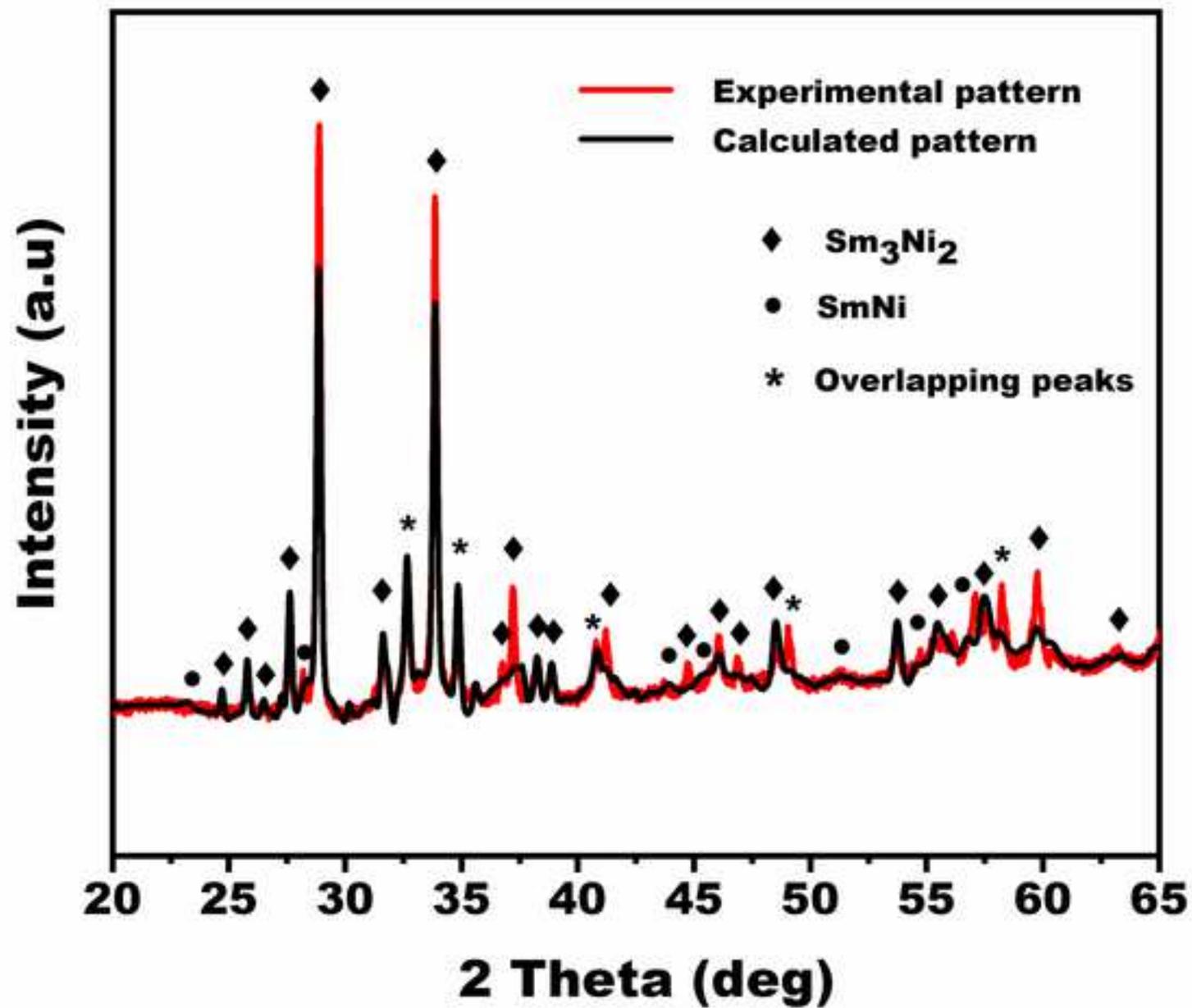
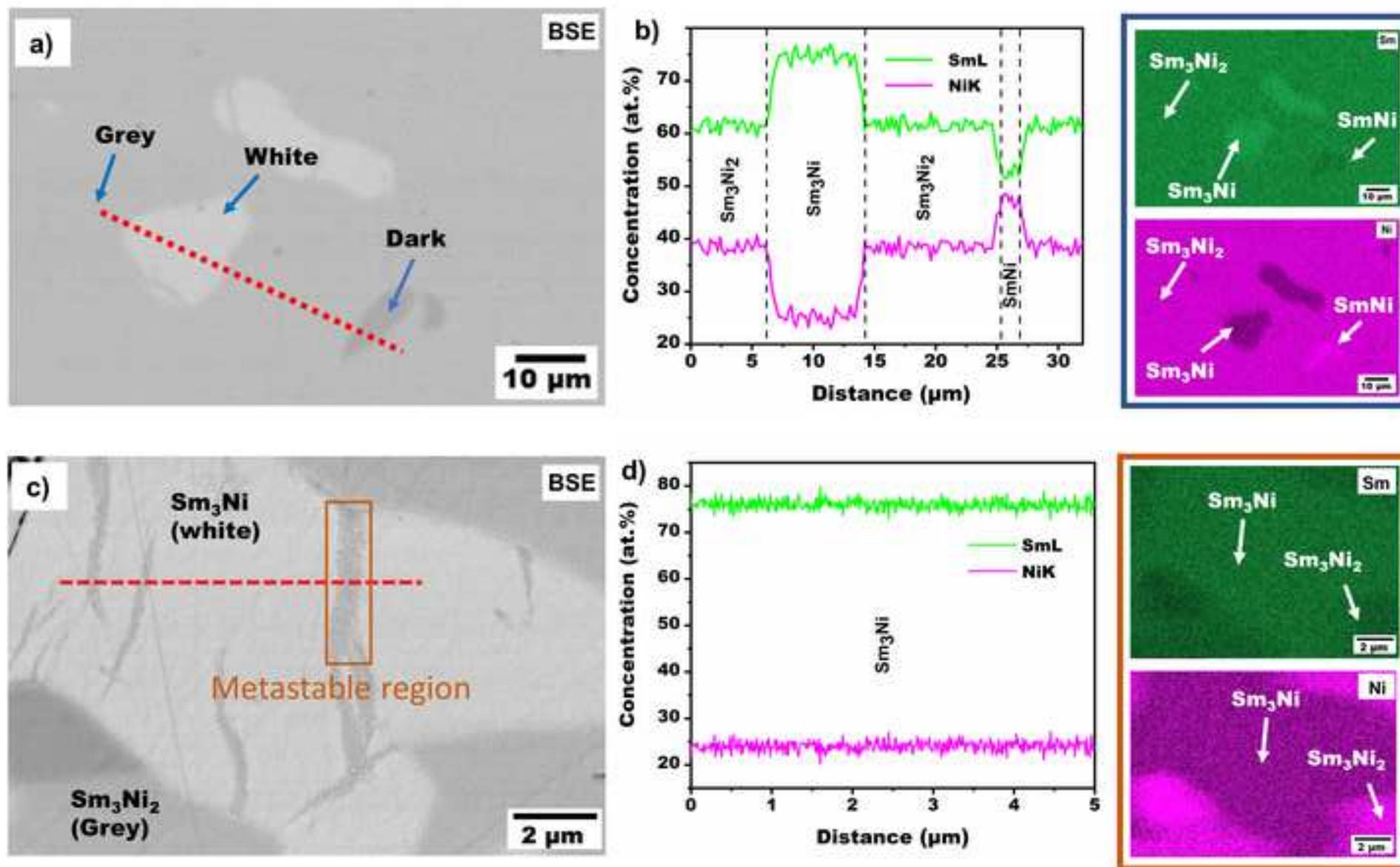
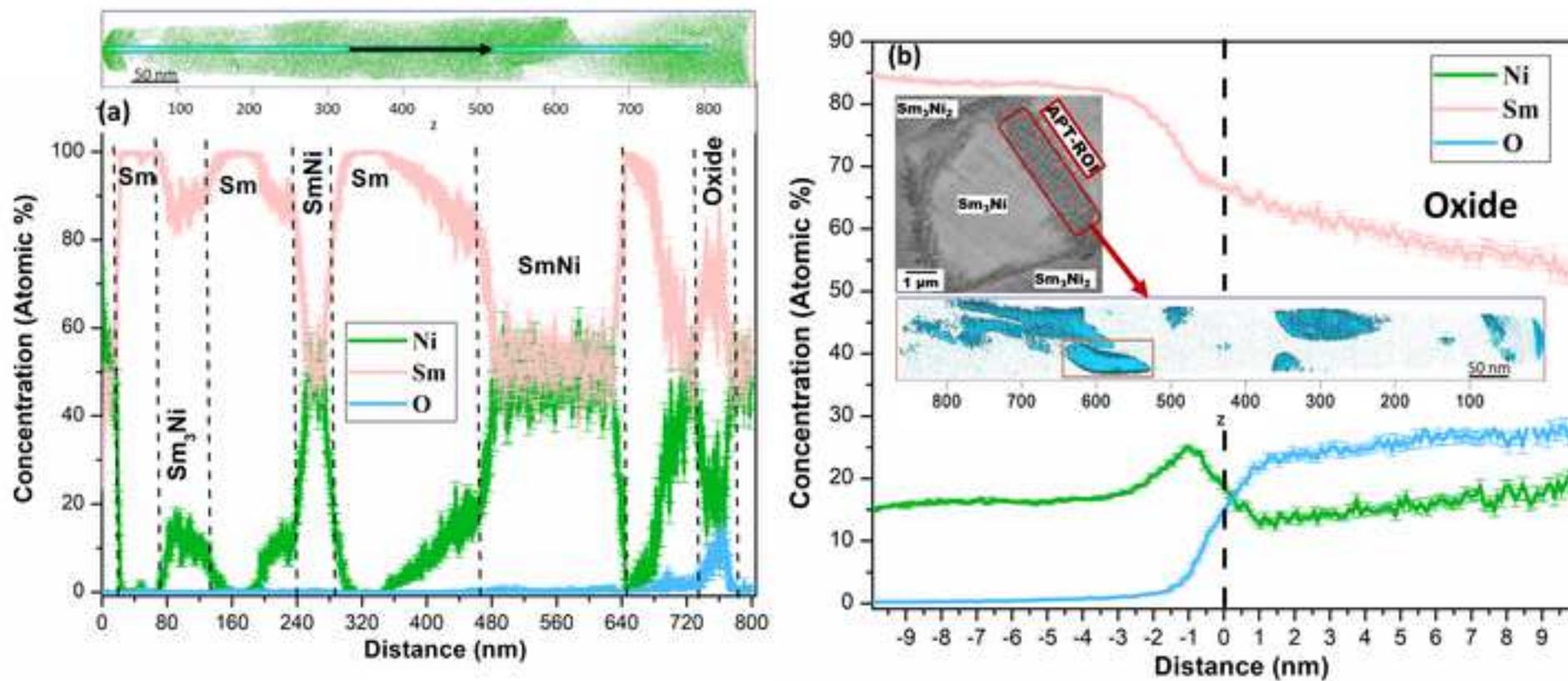
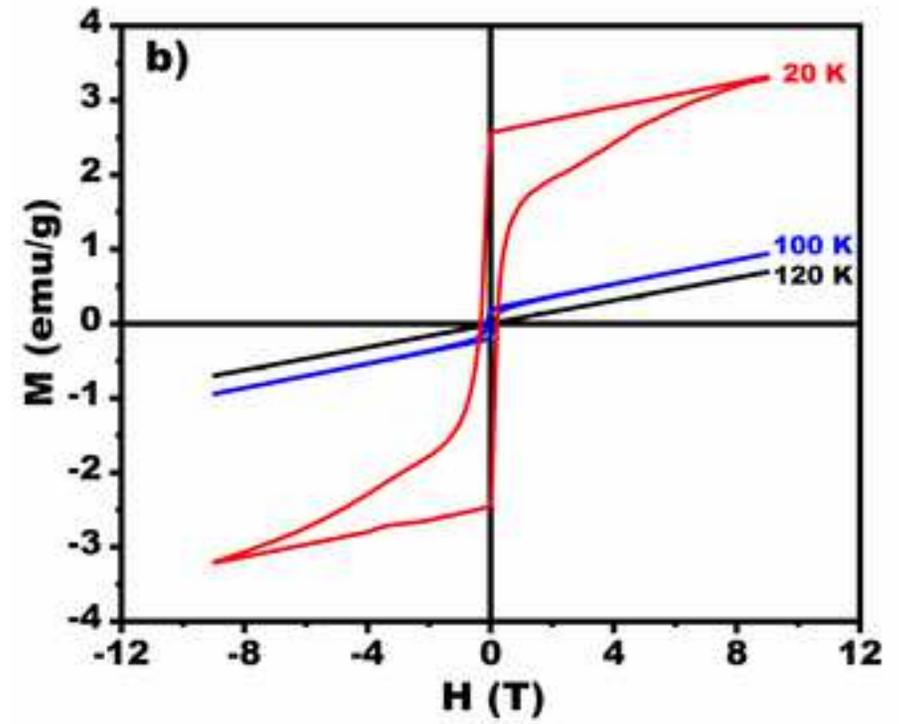
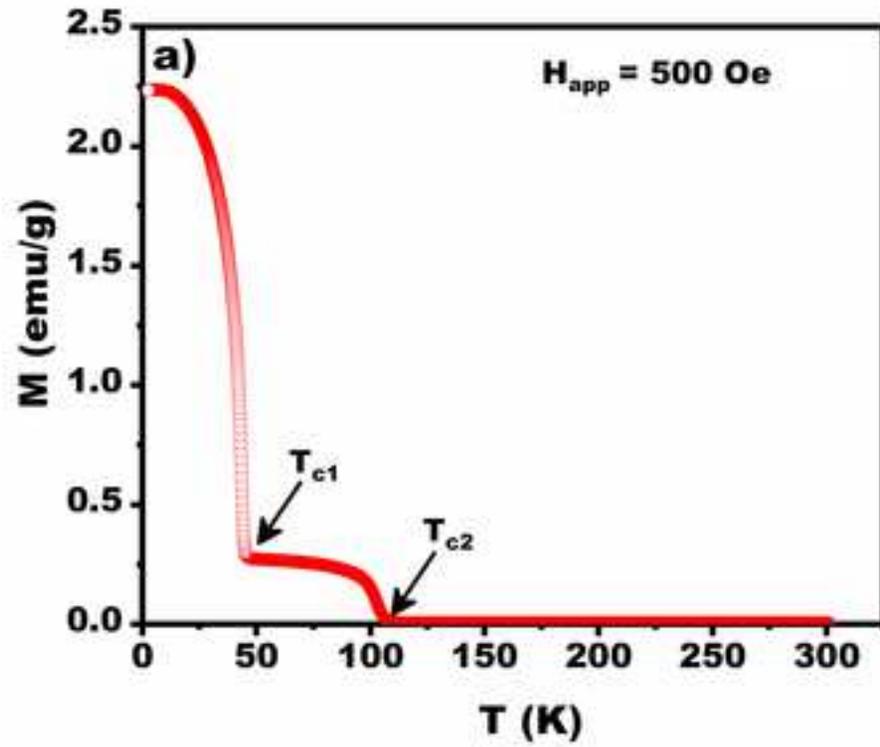


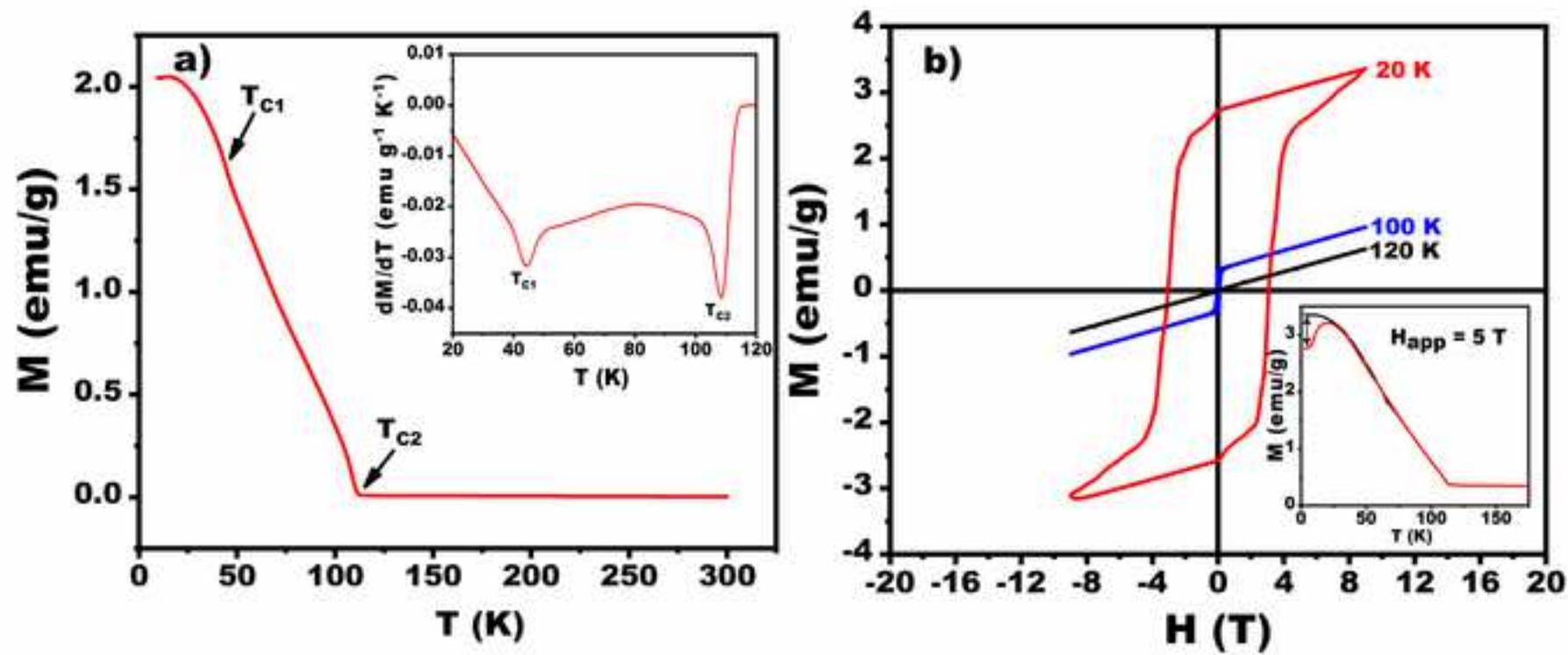
Fig. 3

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**Fig. 7**

**Fig. 8**

Declaration of interests

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

The authors declare the following financial interests/personal relationships which may be considered as potential competing interests: