Investigation of Ferrimagnetic Sr-hexaferrite Nanocrystals for Clinical Hyperthermia

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Sr-hexaferrite samples were produced via the conventional ceramic method. X-ray diffractometry (XRD) patterns confirmed the single nanocrystal phase as Sr-hexaferrite (SHF) where any pattern peaks of unreacted Fe₂O₃ phase were not detected. The mean crystallite size values were determined to be 44 ± 3 nm and 41 ± 3 nm for SHF-O1 and SHF-O2, respectively. The chemical bonding peaks of our sample indicated that the structure of Sr-hexaferrite formation was confirmed by FTIR spectra result. Scanning electron microscopy (SEM) images indicated clearly observed porosity regions with relative densities as high as 94 % and 87 % for SHF-O1 and SHF-O2 samples. The vibrating sample magnetometry (VSM) of each sample at 2K and under a magnetic field of 10 kOe yielded saturation magnetizations, $M_s$ of 93.5 and 94.1 emu/g; remanence values, $M_r$ of 76.4 and 67.8 emu/g for SHF-O1 and SHF-O2, respectively. The magnetization loops of both samples indicated a soft ferrimagnetic behaviour in which the saturation magnetizations were higher than those measured at room temperature in the previous studies. The coercivities, $H_c$ were measured to be 150Oe for both samples. The squareness values, $SQR$ ($M_r/M_s$) were measured to be high, approximately 0.82 and 0.72 for SHF-O1 and SHF-O2, respectively. Depending on the adequate values of magnetization an coercivity along with small mean crystallite size and low porosity values of the obtained Sr-hexaferrite samples, we estimate that these samples are likely to be evaluated further for the potential use as thermoseeds in the field of clinical hyperthermia.

Keywords: hexaferrites, ferrimagnetic properties, magnetization, coercivity.

1. INTRODUCTION

Hexaferrites have been massively fabricated materials and today, their applications involve microwave and radar technology, magnetic recording and data storage, automotive industry and the others [1]. M-type hexaferrites have been examined more frequently than other six classes. 90% of the magnetic materials that are produced today are based on hexagonal ferrites [2].

The magnetic properties of Sr-hexaferrites such as the saturation magnetization, coercivity have been experimentally studied by replacing Sr$^{2+}$ and/or Fe$^{3+}$ sites with cationic dopants [3, 4]. The magnetic measurements at the cryogenic temperature range have initiated in the last twenty years, by the invention of required equipments which are able to work below 4.2K [5].

M-type hexaferrites are metal oxides with a formula of MFe$_{12}$O$_{19}$. They are frequently utilized as permanent magnets [6]. Apart from the high saturation magnetizations and magneto-crystalline anisotropies, also high intrinsic coercivities; they possess a number of electrical properties as well such as high electrical resistivity, low dielectric loss, low eddy current loss etc [7–9]. High frequency and microwave field i.e radar absorbing materials, is an example of their applications. Medical equipment applications also require hard hexaferrite magnets [10, 11].

Despite their very old research history, there are still undefined properties of especially M-type hexaferrites. In addition, new and interesting fields of application have been emerging upon the recent studies [12]. Two of the M-type hexaferrites, BaFe$_{12}$O$_{19}$ and SrFe$_{12}$O$_{19}$, exhibit a magnetoplumbite structure. Easy magnetization direction is the c-axis in both of them [13].

Many studies have focused on the electrical and magnetic properties of hard M-type hexaferrites, which have been produced by different synthesis methods such as chemical routes [14, 15] and solid oxide reaction methods. Ferrimagnetic materials are magnetically characterized by magnetometer, often at room temperature. High and low magnetic fields are applied in this characterization; high magnetic fields are utilized particularly for determining the saturation magnetization [16].

Clinical hyperthermia is a thermal treatment of the cancer using ferromagnetic seeds and the application of a magnetic field. Sr-hexaferrites exhibit some tunable magnetic properties, i.e. high saturation magnetization, to be used in this context [17]. Most previous researches have been focused on the room temperature magnetic behaviours of the hexaferrites except some of the very recent studies on the low temperature magnetization [18].

In order to improve the efficiency of the hyperthermia treatment, high saturation magnetization is often preferred [19]. Therefore, the aim of this paper is to conduct the magnetic characterization of Sr-hexaferrites at 2K for a higher saturation magnetization in order to enhance the ferrimagnetic properties at the low temperature region.

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2. EXPERIMENTAL DETAILS

2.1. Ferrite preparation

The polycrystalline Sr-hexaferrites were synthesized using conventional ceramic method. The raw materials of alfa-Fe$_2$O$_3$ (min 99 wt.%) and SrCO$_3$ (98 – 99 wt.%) were mixed in the appropriate proportion and calcined at 1000 °C for 1 h and 2 h to produce Sr-hexaferrite phase then coded with SHF-O1 and SHF-O2, respectively. Afterwards, the hexaferrite product was ball-milled multiple times inside a hardened steel container with steel balls as the grinding medium to obtain fine powders with mean size of 0.8 mm in order to achieve a single domain in each powder particle. After mixing with some water, obtained wet slurry was depressed under the pressure of 5 MPa and the external magnetic field of 10 kOe. Then green compacts were fired at 1250 °C for 1 h.

2.2. Characterization studies

2.2.1. Phase identification

X-ray diffraction (XRD) data from the sintered samples were collected at angular angles between 20 – 80° via Rigaku D-Max2200 diffractometer using CuKα ($\lambda = 1.54056$ nm) radiation to confirm the phase composition. The mean crystallite size values of the sintered samples were calculated through Debye-Scherrer’s formula using the full width at half maximum (FWHM) values obtained from XRD data as in Eq. 1:

$$L = \frac{k\lambda}{\beta \cos \Theta}$$  \hspace{1cm} (1)

where $L$ is the average crystallite size; $\beta$ is FWHM; $\Theta$ is the half of the Bragg angle; $\lambda$ is the X-ray wavelength; $k$ is a constant. Based on Bragg’s Law, the plane spacing of hexagonal crystal, $d$ is related to the lattice parameters (a and c) as in Eq. 2:

$$\frac{1}{d^2} = \frac{4}{3} \left( \frac{h^2 + h.k + k^2}{a^2} + \frac{l^2}{c^2} \right).$$  \hspace{1cm} (2)

The lattice parameters of both Sr-hexaferrites were also calculated by Bragg’s Law using $d$-spacings and ($hkl$) indices of the diffraction planes.

2.2.2. Microstructural characterization

The bulk densities of the Sr-hexaferrites were calculated by determining the dry weight of the samples after drying in oven for 30 min, and measuring the regular dimensions of both samples. The total porosity for the sintered samples was calculated based on the relative density. The porosities of the samples were also measured by bottle pycnometer method based on the Archimedes principle.

Sr-hexaferrite samples were demagnetized by heating at a rate of 5 °C/min up to 460 °C and were held for 2 h. before the analysis. The sintered samples were surface-finished by polishing with SiC papers. Scanning electron microscopy (SEM) technique was conducted using JEOL-JSM 6060 to examine the microstructural morphology.

2.2.3. FTIR analysis of hexaferrites

Fourier transform infrared (ATR-FTIR) spectroscopy results were recorded between the spectra range of 3600 – 650 cm$^{-1}$ using Perkin-Elmer Spectrum BX spectrometer in order to confirm the formation of the chemical bonds of the synthesized Sr-hexaferrites.

2.2.4. Magnetic behaviour of hexaferrites

To conduct the magnetic characterization of the synthesized Sr-hexaferrites, the magnetization vs. magnetic field curves were recorded at 2K under the applied field of 10kOe via vibrating sample magnetometer (VSM) using Cryogenic Limited PPMS.

3. RESULTS AND DISCUSSION

3.1. Microstructural and structural analysis results

Table 1 indicates the sintered densities of both samples. The theoretical density of SrFe$_{12}$O$_{19}$ was considered to be 5.1 g/cm$^3$ in order to calculate the percentages of porosity. The resultant relative densities can be derived as 94 % and 87 % for SHF-O1 and SHF-O2 samples.

<table>
<thead>
<tr>
<th>Sample</th>
<th>$\phi$, mm</th>
<th>$t$, mm</th>
<th>$\rho$, g/cm$^3$</th>
<th>$P$, %</th>
<th>$P_{ii}$, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>SHF-O1</td>
<td>10</td>
<td>5</td>
<td>4.78</td>
<td>6</td>
<td>2.94</td>
</tr>
<tr>
<td>SHF-O2</td>
<td>6</td>
<td>3</td>
<td>4.44</td>
<td>13</td>
<td>13</td>
</tr>
</tbody>
</table>

SEM images of SHF-O1 and SHF-O2 samples are given in Fig. 1. Grain boundaries were not evident even after high-magnification microscopic examination. Thus the grain morphologies were not observed. However, the porosity regions were clearly observed for both samples.

Fig. 2 indicates the XRD pattern peaks of sintered SHF-O1 and SHF-O2 samples. XRD results confirmed the single nanocrystal phase as magnetoplumbite Sr-hexaferrite (JCPDS card number 00-024-1207) in both samples. The pattern peaks of unreacted Fe$_2$O$_3$ phase were not detected.

The crystallite size values from the highest to lowest peak intensity were measured as 43.1 nm, 41.38 nm, 47.89 nm for SHF-O1 and 43.32 nm, 37.48 nm, 42.81 nm for SHF-O2, corresponding to the peaks of (107), (114) and (110). According to the results, the crystallites along <110> directions were higher than those along <114> directions, which indicate a platelet growth in the Sr-hexaferrite crystals [20].

The lattice parameters (a, c) of Sr-hexaferrite crystal were calculated as 6.16 Å°, 23.7 Å° for SHF-O1 and 6.16 Å°, 23.55 Å° for SHF-O2, respectively.

3.2. FTIR results

In order to confirm the formation of Sr-hexaferrite, FTIR spectrum of SHF-O1 sample was examined (Fig. 3). First, since the band range of 3451 – 3498 cm$^{-1}$ is related to the stretching of water molecules [21], the small band at 3492 cm$^{-1}$ in our sample corresponds to the band of O=H. The sharp peak at 2920 cm$^{-1}$ indicates the surface bending bond of the hydroxide (–OH) group due to the damped environment [22].
343

Fig. 1. SEM images of both samples: a – SHF-O1; b – SHF-O2

Fig. 2. XRD patterns of both samples: a – SHF-O1; b – SHF-O2

The small peaks around 1722 cm\(^{-1}\) show the stretching in the vibration of C=O bond, whereas the relatively stronger small band at 1065 cm\(^{-1}\) corresponds to the C-O bond stretching. Finally, the clear sharp peak at 797 cm\(^{-1}\) is related to metal oxide stretching of vibration [21].

Fig. 3. FTIR spectrum of the sample of SHF-O1

As a result, the chemical bonding peaks of our sample indicated that the structure of Sr-hexaferrite formation was confirmed by FTIR spectra result.

3.3. Magnetic behaviour results

According to magnetic measurements at 2K, it was confirmed that both samples exhibit typical soft ferrimagnetic behaviour as shown by the narrow hysteresis curves in Fig. 4. The magnetic alteration from hard to soft type in the ferrimagnetic hexaferrites, where spins are non-collinear is a function of measurement temperature [23].

Fig. 4. M-H hysteresis loops of the samples: a – SHF-O1; b – SHF-O2. Full arrow: saturation and blank arrow: remanence values

The magnetic parameters of two Sr-hexaferrite samples were calculated from the low temperature M-H curves. The saturation magnetizations, Ms were determined to be 93.5 and 94.1 emu/g for SHF-O1 and SHF-O2, respectively at 2K. These values were higher than those measured at the room temperature [12].
Because a less thermal energy causes a small fluctuation in the magnetic moments thus, the saturation magnetization rises [12]. Remanence values, Mr were determined to be 76.4 and 67.8 emu/g for SHF-O1 and SHF-O2, respectively. Remanence values of the Sr-hexaferrites at the room temperature in the previous studies were substantially lower, 13.2 emu/g in comparison [13] due to canted type spin structures in the ferrimagnetic hexaferrites. The coercivities, Hc at 2K were measured to be 150 Oe for both SHF-O1 and SHF-O2 samples which were significantly lower in comparison to those at room temperature [13], typical of M-type hexaferrites.

At 2K the squareness values, SQR (Mr/Ms) were measured to be high, approximately 0.82 and 0.72 for SHF-O1 and SHF-O2, respectively. These results can be attributed to the appropriate sintering treatment which permitted to obtain an oriented Sr-hexaferrite [24]. Instead, the room temperature SQR values were lower, i.e. 0.54 for Sr-hexaferrite [13] because the spin canting causes a reduced remanence in the hexaferrites [12]. The mean crystallite size values of 44 ± 3 nm and 41 ± 3 nm for SHF-O1 and SHF-O2, respectively were calculated to be smaller than the critical value for a single-domain (650 nm) of the Sr-hexaferrites [25].

Table 2 indicates the magnetic behaviour results of Sr-hexaferrites were given at 2K, 10K and 300K. The saturation magnetization value, Ms of our sample at 2K was close to that of Tm doped SHF measured at 10K but it was lower than BHF value measured at 2K. On the contrary, the coercivity, Hc of our sample at 2K was lower in comparison to the values obtained in other studies both at 300K and 10K, which can be attributed to the variations in the crystalline size. In order to examine the demagnetization behaviour of the samples, the second quadrant of the ferrimagnetic curve was converted into the magnetic induction (B) vs. field strength (H) graph as in Fig. 5 a. The remanent induction (Br) and the coercivity (Hc) values of the Sr-hexaferrites were shown on the B-H graph.

On the second quadrant of the ferrimagnetic curve, the largest rectangle was also drawn to determine the maximum energy product, BHmax of our sample as shown in Fig. 5 a. The parabola curve in Fig. 5 b indicates the maximum energy product vs field strength (H) graph. In order to draw this curve, the combinations of M-H values in the second quadrant were considered. The vertex of this curve was taken as the BHmax of our sample. BHmax values were calculated to be 1.6 and 1.2 kJ⋅m⁻³(0.19 and 0.16 MGOe) for SHF-O1 and SHF-O2, respectively.

Table 2. Magnetic behaviour of the Sr-hexaferrite samples

<table>
<thead>
<tr>
<th>Magnetic properties</th>
<th>Temperature</th>
<th>BHF(Ce, Zn) a,b</th>
<th>SHF(Tm)</th>
<th>SHF(Zr)</th>
<th>SHF-O1</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ms, emu/g</td>
<td>300K</td>
<td>65 (67)</td>
<td>70</td>
<td>52</td>
<td>46 *</td>
</tr>
<tr>
<td></td>
<td>2K</td>
<td>117 (117)</td>
<td>–</td>
<td>–</td>
<td>93</td>
</tr>
<tr>
<td></td>
<td>10K</td>
<td>–</td>
<td>97</td>
<td>75</td>
<td>–</td>
</tr>
<tr>
<td>Mr, emu/g</td>
<td>300K</td>
<td>18 (7)</td>
<td>41</td>
<td>25</td>
<td>22 *</td>
</tr>
<tr>
<td></td>
<td>2K</td>
<td>10 (6)</td>
<td>–</td>
<td>–</td>
<td>76</td>
</tr>
<tr>
<td></td>
<td>10K</td>
<td>–</td>
<td>54</td>
<td>37</td>
<td>–</td>
</tr>
<tr>
<td>Hc, kOe</td>
<td>300K</td>
<td>7.5 (3.1)</td>
<td>4.3</td>
<td>4.5</td>
<td>8.8 *</td>
</tr>
<tr>
<td></td>
<td>2K</td>
<td>2.3 (1.5)</td>
<td>–</td>
<td>–</td>
<td>0.2</td>
</tr>
<tr>
<td></td>
<td>10K</td>
<td>–</td>
<td>2.8</td>
<td>3.5</td>
<td>–</td>
</tr>
<tr>
<td>Mr/Ms</td>
<td>300K</td>
<td>0.3 (0.1)</td>
<td>0.6</td>
<td>0.5</td>
<td>0.5 *</td>
</tr>
<tr>
<td></td>
<td>2K</td>
<td>0.1 (0.1)</td>
<td>–</td>
<td>–</td>
<td>0.8</td>
</tr>
<tr>
<td></td>
<td>10K</td>
<td>–</td>
<td>0.6</td>
<td>0.5</td>
<td>–</td>
</tr>
</tbody>
</table>

Reference: [26] [27] [28] Present study, [18]

+: magnetic values at 300 K for a sample close to SHF-O1;
a,b: two different doping levels.
Heating effect generated by the ferromagnetism is a significant point in the clinical hyperthermia application. Specific absorption rate (SAR) is a measure of the heating effect and this parameter has been investigated by some of the researchers for Sr-hexaferrite. SAR value of 0.5 – 0.7 W/g has been found in Sr-hexaferrite composites at a frequency of 214 kHz and amplitude of 220e, which shows some potential usage in the clinical hyperthermia [29]. SAR values between couple of units and tens W/g have been measured in Sr-hexaferrite/maghemite nano-composites at a frequency of 108 kHz and amplitude of 188 – 855 Oe [30]. SAR values in the range of 50 – 750 W/g have been determined in Fe₃O₄ nano-particles at a frequency of 265 kHz and amplitudes of 184 – 625 Oe depending on the particle size [31]. Also, some nano-spinel ferrites of MgFe₂O₄, Zn₆Fe₁₀O₁₆, MnZnFe₂O₄ have also been investigated for their SAR values and they have been measured to be 3.9, 36 and 110 W/g, respectively for these ferrites [31]. As a result, it is estimated that our Sr-hexaferrite ferrimagnets may produce similar orders of heating effect to be used in the clinical hyperthermnia. Thus, it will be quite beneficial to test the heating effects of our Sr-hexaferrite samples for future studies.

4. CONCLUSIONS

Sr-hexaferrites were successfully synthesized via the conventional sintering. As-sintered densities and the porosities determined indicated a low porosity content. This was confirmed by SEM images of both samples. However, the grain boundaries could not be observed in the SEM microscopy due to nano-sized crystals. XRD was utilized to ensure the phase purity and the patterns indicated that both of the samples contained a single phase of Sr-hexaferrite. The crystallite size values calculated by Debye-Scherrer’s formula varied in the range of 37 – 47 nm.

Magnetic behaviour of the Sr-hexaferrite samples altered from hard to soft ferrimagnetism with the decreasing temperature in comparison to the previous studies where the room-temperature magnetization was determined. In contrast to the room-temperature studies, the collinear type of magnetic structure is present at 2K with an increase in the magnetization values. The saturation magnetization increased by 1.1 % when the samples of SHF-O1 and SHF-O2 were compared due to the higher XRD intensity observed in SHF-O2. The coercivities of both samples at 2K were measured to be 150Oe, which can be attributed to the close crystal size values in the samples.

Consequently, it is anticipated that the performance of our Sr-hexaferrite nanocrystals with a number of appreciable properties might be useful as ferrimagnetic thermoseeds for clinical hyperthermia applications, where a combination of high magnetization and low coercivity is required.

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