Structural, Magnetic and Electrical Properties of Co-Doped Barium Monoferrite BaFe_2O_4

Samar I. Abdelsalam^a, Adel Maher Wahba^{a,*}, M.A. Amer^b, A.S. Doma^c, Bahaa Eddin M. Moharram^a

^aFaculty of Engineering, Department of Engineering Physics and Mathematics, Tanta University, Egypt

^bDepartment of Physics, Faculty of Science, Tanta University, Egypt

^c Polymeric Materials Research Dep., Advanced Technology and New Materials Research Institute (ATNMRI), City of Scientific Research and Technological Applications (SRTA-City), New Borg El-Arab City 21934, Alexandria, Egypt

* Corresponding author: adel.mousa@f-eng.tanta.edu.eg

Abstract- Cobalt-substituted barium monoferrite with the formula $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ ($x = 0.0 - 0.4$, step 0.1) has been prepared through the sol-gel autocombustion method. Phase purity for prepared samples sintered at 700 and 1050°C was analyzed by X-ray diffraction (XRD) patterns that indicated the formation of the typical orthorhombic phase for the samples. Fourier-transform infrared (FTIR) spectroscopy was used to investigate the effect of cobalt doping on the vibrational modes of the studied samples. Scherrer and Williamson-Hall formulae were used to define the microstrain and the crystallite size being in the range of 44.1 to 59.7 nm, which was further confirmed with transmission electron microscopy (TEM) images. Magnetic characteristics including magnetic saturation and coercivity were obtained from hysteresis loops traced using vibrating sample magnetometry (VSM). Magnetic properties were correlated with the composition of the prepared samples. Finally, dielectric measurements were operated to further confirm the structural and magnetic analysis.

I. INTRODUCTION

Along the past few decades, magnetic nanomaterials have been under intense investigation due to their versatile use in technological application including drug delivery [1], magneto-optic media [2], radar absorption [3] and modern microwave devices [4]. This is attributed to the distinctive properties exhibited by them such as excellent chemical stability, good dielectric properties, and the flexibility of tailoring its magnetic properties [5-7].

Spinel ferrites are assigned by the formula MFe_2O_4 , where M refers to a divalent metal cation ($\text{M} = \text{Ba}, \text{Sr}, \text{Ni}, \text{Co}$, etc.). The structure of these materials depends mainly on the cation M and/or its substituent. For example CoFe_2O_4 has cubic structure [8], that of SrFe_2O_4 is hexagonal [9] and that of BaFe_2O_4 is orthorhombic [10]. Generally, less symmetric structures are produced when the difference between the ionic radius of the cation M and that of Fe^{3+} cation is larger. For each of these structures M and Fe cations occupy two characterizing sites, namely octahedral and tetrahedral sites specified by the lattice made up by oxygen anions. Among the spinel ferrites, barium monoferrite (to distinguish it from hexagonal barium ferrite $\text{BaFe}_{12}\text{O}_{19}$) is characterized by the unique combination of moderate magnetization and large coercivity, which permits its use in quite specified applications, e.g. magnetic storage and permanent magnets [11]. Moreover, many works have reported its wide use as photocatalytic [12, 13] and pigments [14].

The magnetic properties of BaFe_2O_4 , like other spinel ferrites, could be controlled using the substitution technique of either Ba^{2+} [9] and/or Fe^{3+} cations. The substitution of Fe^{3+} could be by either trivalent ions (Al^{3+} , Cr^{3+}) [15, 16] and/or divalent ions (Zn^{2+} , Co^{2+}) [17, 18].

Several preparation methods have been reported by researchers for synthesizing ferrite materials with the desired composition stoichiometry, structure, and morphology to achieve the properties compatible with the specified application. Such methods include solid state or ceramic [19], co-precipitation [20], hydrothermal [21], and sol-gel autocombustion [22]. The latter presents some advantages among which, simple reagents, inexpensive equipment, doping control, low-temperature production, good stoichiometric control, homogenous mixing, and finally, the short preparation time [23-25].

In the present work, the substitution of Co^{2+} in $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ system has been investigated from the point of view of its effects on structure, magnetic and dielectric properties. XRD, TEM, and IR spectroscopy were used to analyze the structural properties including phase purity, crystallite size, lattice parameters and strain for prepared samples sintered at both 700 and 1050°C. VSM was used to obtain saturation magnetization, coercivity and remanence. The effect of Co^{2+} doping on the magnetic properties of BaFe_2O_4 has been investigated and correlated with the structural and electrical properties.

II. EXPERIMENTAL

A. Preparation of Nanocrystalline $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$

Nanocrystalline Co-doped barium monoferrite samples with the formula $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ ($x = 0.0 - 0.4$, step 0.1) were synthesized using sol-gel autocombustion technique. The starting materials were analytical grade nitrates of $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, $\text{Ba}(\text{NO}_3)_2$, and $\text{Fe}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$ and anhydrous citric acid $\text{C}_6\text{H}_8\text{O}_7$. Stoichiometric amounts of the metal nitrates were dissolved separately in minimum amounts of distilled water, and then mixed together. Citric acid solution was then added to the nitrates solution in a molar ratio of 1:1 (to chelate Ba^{2+} , Co^{2+} , Fe^{3+} ions) and then, a stirring process was operated for one hour. Meanwhile, ammonium hydroxide was added drop by drop to adjust the pH number from ~1 (for the acidic solution) to 7, after which the stirring process has been allowed to continue for further an hour. Heating at 100°C was then initiated on the hot plate

to enforce water evaporation till the formation of a dark vicious gel was observed. Continuing the heating process initiated the autocombustion process from the hottest point of the formed gel, which then propagates up like an erupting volcano, forming an ash-like powder. The latter was ground in an agate mortar to obtain a loose powder. For each composition, the fine powder was divided into two parts for being sintered in a programmable muffle furnace at 700 and 1050 °C for 2 h, separately, with a heating rate of 5 °C/min.

B. Characterization techniques

XRD diffraction patterns of the sintered samples with different compositions were recorded using a Philips diffractometer (X'pert MPD) with a goniometer using Cu-K α radiation ($\lambda = 1.54056$ Å). The diffracted intensities were collected in the step-scan mode (step size $2\theta = 0.03^\circ$ with step time 1.5 s). The angular range was 15 to 70°. The recorded patterns were used for performing the structural analysis for the studied samples, from which information such as phase structure, lattice parameters, crystallite size, and unit-cell volume were achieved. Infrared spectroscopy (FTIR) of the prepared samples was carried out on Bruker Tensor 27 FTIR Spectrometer in the range of 200 – 900 cm $^{-1}$ to confirm the phase structure of the prepared BaFe $_2$ O $_4$ sample and to analyze the changes induced on it upon doping with Co $^{2+}$. For this purpose, 0.2 mg of the ferrite was mixed with 200 mg of KBr and pressed into pellets. The powder morphology was recorded by TEM images (TEM, JEOL JEM-100CX with an accelerating voltage extending up to 100 kV) (HR-TEM, Tecnai G20, FEI, Netherland) with an accelerating voltage of 200 kV.. Magnetic characteristics of the studied samples, namely saturation magnetization, remanence, and the coercive field were deduced from M - H hysteresis loops traced at room temperature, using VSM (Lake Shore-7400) with an applied field extending up to 20KOe). The electrical properties including the ac conductivity and the dielectric constant (with its real, ϵ' , and imaginary, ϵ'' , parts) were achieved at room temperature using a IN 3536 LCR Meter in the frequency range of 1 MHz to 8MHz. The ac conductivity (σ_{ac}) was obtained using the relation $\sigma_{ac} = 2\pi f \epsilon_0 \epsilon' \tan \delta$, where f is the frequency (Hz), ϵ_0 is the electrical permittivity of free space, ϵ' is the dielectric constant and $\tan \delta = \epsilon''/\epsilon'$ is dielectric loss factor. The parameter ϵ' was determined from the formula $C d/\epsilon_0 A$ where C is the capacitance, d is the pellet thickness and A is its cross-sectional area. The imaginary part of the dielectric constant is depicted from the formula $\epsilon'' = d G/2\pi f \epsilon_0 A$, where G is the measured conductance ($1/R$).

III. RESULTS AND DISCUSSION

A. Structural analysis

The XRD patterns for the prepared BaCo $_x$ Fe $_{2-x}$ O $_4$ ($x = 0.0$ to 0.4, step 0.1) sintered at 700 °C are shown in Fig. 1(a). The XRD pattern quite matches that of the orthorhombic crystal structure of barium monoferrite (identified by JCPDS card 70-2468). The peak at $2\theta = 31.09^\circ$ (indicated with •), recorded for samples with $x \geq 0.2$, corresponds to the maximum-intensity peak, of the diffraction plane (110), for the cubic perovskite phase of BaFeO $_3$ (JCPDS card 75-0426). That perovskite-phase impurity increases with Co doping as indicated from the figure. Fig. 1(b) shows the diffraction

Figure 1. XRD pattern of BaCo $_x$ Fe $_{2-x}$ O $_4$ samples sintered at (a) 700 °C and (b) 1050 °C. • indicates BaFeO $_3$ and * indicates Ba $_2$ FeO $_4$.

patterns of the same samples sintered at 1050 °C. Extra feeble peak appeared at $2\theta = 29.2^\circ$ (indicated with *) corresponding to the orthorhombic Ba $_2$ FeO $_4$ oxide, with maximum-intensity diffraction plan (112) (JCPDS card 23-1022). Worth to mention is that sintering at 1050 °C enhanced the sharpness and intensity of the diffraction peaks for $x \leq 0.2$, indicating the enrichment of size and crystallinity. It is expected that BaFeO $_3$ and Ba $_2$ FeO $_4$ impurities (for $x = 0.3$ and 0.4 samples) impeded the crystal growth of the original orthorhombic structure of BaCo $_x$ Fe $_{2-x}$ O $_4$.

The crystallite size was estimated by two well-known methods. The first used the broadening of the peaks according to Scherrer's formula [26], $D = k\lambda/(\beta \cos \theta)$, Where D is the particle size, λ is X-ray wavelength, k is a

shape-factor constant taken as 0.94, β is the peak full-width at half-maximum (FWHM) intensity and 2θ is the diffraction angle. The second method involved the use of Williamson-Hall strain-size formula [27]:

$$(d\beta\cos\theta)^2 = \frac{k}{D} d^2 \beta \cos\theta + \left(\frac{\varepsilon}{2}\right)^2$$

Where d is the spacing between diffraction plans corresponding to (h, k, l) lattice indices and ε is the strain. The particle size is determined from the slope of the linearly fitted data of $(d\beta\cos\theta)^2$ on the y axis versus $(d^2\beta\cos\theta)$ on the x axis. The root of the y -intercept gives the strain. An example of the use of both methods is shown in Fig. 2a, b, for the sample with $x = 0.2$ sintered at 700°C. The crystallite size and microstrain for all samples and for both sintering temperatures are illustrated in Table 1. It is clear that for sintering temperature of 700 °C, the crystallite size hardly changed with Co^{2+} doping (40 – 44.2 nm), while for sintering at 1050 °C, the crystallite size and crystallinity (see Fig. 1b) decreased with dopant content, being attributed to phase impurities. This is confirmed when considering the significant growth of the undoped BaFe_2O_4 sample (from 40.9 to 57.9 nm) upon increasing the sintering temperature to 1050 °C.

The orthorhombic-phase lattice parameters of the studied samples (a , b and c) were calculated using (suggested) Miller indices (h , k , and l) corresponding to the XRD reflections and their d spacing according to lattice geometry equation [28]:

$$\frac{1}{d^2} = \frac{h^2}{a^2} + \frac{k^2}{b^2} + \frac{l^2}{c^2}$$

Using successive iterations, the deduced lattice parameters were used to attain theoretical values for the d -spacing of the diffraction planes. The maximum standard deviation of experimental and theoretical d -spacing values was less than 0.001. The unit cell volume was calculated using $V = abc$. The lattice parameters and unit cell volume for all samples are listed in Table 2. Variation of the unit cell volume with Co concentration for both sintering temperatures is illustrated in Fig.3. There is a slightly decrease in unit cell

Figure 2. (a) Scherrer plot and (b) W-H strain-size plot of $\text{BaCo}_{0.2}\text{Fe}_{1.8}\text{O}_4$ sintered at 700 °C.

TABLE 1

CRYSTALLITE SIZE (D) AND STRAIN (ε) OF $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ SINTERED AT 700 AND 1050 °C

x	D (nm)				$\varepsilon \times 10^{-3}$	
	Scherrer		W-H		700 °C	1050 °C
	700 °C	1050 °C	700 °C	1050 °C		
0.0	40.9	57.9	35.0	57.5	0.974	0.789
0.1	40.0	56.6	30.3	42.1	0.798	0.546
0.2	42.3	53.3	39.2	38.3	1.152	0.721
0.3	44.2	43.7	38.3	45.5	0.721	1.188
0.4	43.4	44.1	32.4	35.8	0.782	0.842

TABLE 2

LATTICE PARAMETERS AND UNIT CELL VOLUME OF $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ samples SINTERED AT 700 AND 1050 °C

Lattice Parameters (Å)							Unit-cell vol. (Å ³)	
x	700 °C			1050 °C			700 °C	1050 °C
	a	b	c	a	b	c	V	V
0.0	19.027	5.385	8.451	19.05	5.382	8.448	865.9	866.0
0.1	19.030	5.385	8.461	19.02	5.376	8.452	867.1	864.3
0.2	19.025	5.384	8.463	19.03	5.381	8.446	867.0	864.9
0.3	19.025	5.376	8.440	19.02	5.379	8.447	863.2	864.0
0.4	19.010	5.376	8.441	18.97	5.367	8.455	862.7	860.6

Figure 3. unit cell volume for $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ sintered at 700 and 1050 °C

volume with increasing the dopant content for $x \geq 0.2$. The monotonic decrease in volume cannot be explained on the basis of the difference between the ionic radii of Co^{2+} ion (0.74 Å) and Fe^{3+} ion (0.64 Å). As such difference is expected to induce lattice expansion as recorded for sample with $x = 0.1$ sintered at 700 °C. This behavior might be interpreted as follow. For sample with $x = 0.1$ sintered at 700 °C, Co^{2+} cation simply substitutes Fe^{3+} causing both the creation of electronic charge imbalance and an increase in the unit cell volume as compared with the undoped BaFe_2O_4 sample (due to the larger ionic radius of Co^{2+}). However, upon sintering to 1050 °C, enough energy is supplied for the oxidation of Co^{2+} to Co^{3+} (0.545 Å) to achieve the charge balance with a resulting reduction in the unit cell volume. Magnetic behavior of this sample supports this scenario, as will be discussed below. For higher concentration of Co^{2+} ($x \geq 0.2$) and for either sintering temperatures, the induced non stoichiometry is compensated by the occurrence of higher valence state of Fe^{3+} cations, i.e. Fe^{4+} with the smaller ionic radius (0.585 Å) resulting in the reduction of the unit cell volume. This could be confirmed by the appearance of traces of BaFeO_3 and/or Ba_2FeO_4 impurities in XRD patterns; either of those compounds has Fe^{4+} cations. The effect of

Figure 4. HRTEM images of (a) BaFe_2O_4 sintered at 700 °C; (b) $\text{BaCo}_{0.4}\text{Fe}_{1.6}\text{O}_4$ sintered at 700 °C; (c) the electron diffraction pattern for sample of (a).

replacing Fe^{3+} cation by Co^{2+} could be simply clarified by the following chemical equation $2\text{Fe}^{3+} \rightarrow \text{Co}^{2+} + \text{Fe}^{4+}$.

To attain a better view of the crystallite size and morphology HRTEM images of BaFe_2O_4 and $\text{BaCo}_{0.4}\text{Fe}_{1.6}\text{O}_4$ sintered at 700 °C was shown at Fig (4a and b). The powder morphology was recorded by TEM images (TEM, JEOL JEM-100CX with an accelerating voltage extending up to 100 kV). Fig.4c shows the diffraction pattern of BaFe_2O_4 sintered at 700 °C. The particles have a tendency to agglomerate probably due to the magnetic interaction between the particles [29]. The average particle size obtained from TEM images agrees quite well with that deduced from the XRD data analysis. The arrows in Fig. 4b clarify the formation of perfectly-packed orthorhombic crystallites. The crystallinity of undoped barium ferrite is a bit incomplete as compared to that of $\text{BaCo}_{0.4}\text{Fe}_{1.6}\text{O}_4$; the particles are less crystallized with distorted shapes and electron diffraction pattern is relatively dim. Barium monoferrite is reported to

Figure 5. FTIR absorption spectra of barium cobalt ferrites with the formula $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ with different substitutions sintered at (a) 700 °C and (b)

fully crystallizes at 900 °C [9], which has been assured from our XRD data (see Fig. 1a, b).

Figure (5a, b) shows FTIR spectra of $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ with different Co^{2+} ion content sintered at 700 and 1050 °C. As just explained by the above discussion, IR spectrum for undoped BaFe_2O_4 sintered at 700 °C did not show any vibrational modes (Fig a). However, spectra for the Co-doped samples showed well-identified behavior. Vibrational peaks at 302, 377, 503, 630 and 775 cm^{-1} were recorded for each sample for both sintering temperatures. All these peaks are characteristics for the orthorhombic phase of BaFe_2O_4 [13, 30-33]. With increasing Co content, the peak around 302 cm^{-1} becomes sharper which might attribute to the vibrational coupling between Fe (Ba)–O and Co–O. It could be inferred that Co^{2+} joined well the internal orthorhombic phase and the $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ was successfully prepared by autocombustion sol-gel method [34]. Worth to mention is that a vibrational shoulder was recorded at around 802 cm^{-1} . In $\text{La}_{1-x}\text{Ca}_x\text{FeO}_3$ pervoskite investigated by Barbero *et al.* [35], new band was observed at around 877 cm^{-1} , whose intensity increased with the increase of divalent Ca content. They assigned this band to the formation of Fe^{4+} . Moreover, Gholizadeh *et al* [36] reported that substituting Co in $\text{LaFe}_{1-y}\text{Co}_y\text{O}_3$ resulted in the oxidation of Co^{2+} and Fe^{3+} to Co^{3+} and Fe^{4+} respectively. Based on the findings in literature and according to the results of our structural analysis, substitution trivalent Fe^{3+} cation by divalent Co^{2+} one may enforce oxidation of Co^{2+} to Co^{3+} and/or Fe^{3+} to Fe^{4+} in order to maintain the electronic charge neutrality. The probability of this oxidation process increases with either degree of substitution (x value) or by increasing the sintering temperature. Accordingly, we can assign the shoulder at 802 cm^{-1} to the Fe^{4+} ions formation.

B. Magnetic properties

Magnetic hysteresis (M - H) loops for $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ samples ($x = 0.0 - 0.4$, step 0.1) sintered at 700 °C and 1050 °C are depicted in Fig. 6(a, b) respectively. The magnetic characteristics including magnetization at 20 kOe (M_{20}), coercivity (H_c), and remnant field (M_s) are listed in Table 3 and Table 4 for sintering temperature 700 and 1050 °C, respectively. The saturation magnetization (M_s) was deduced by plotting M versus $1/H$ and taking the limit as H approaches ∞ [37, 38]. For the barium monoferrite sample ($x = 0.0$) sintered at 700 °C, a typical M - H loop for barium

Figure 6. Hysteresis pattern of undoped and cobalt doped barium ferrites $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ sintered at (a) 700 °C and (b) 1050 °C

monoferrite is exhibited [39]. The necking appearing at relatively weak applied field could be attributed to the incomplete crystallinity that could appear in a nonuniform particle size distribution (a mixture of small and large particle) which agrees with TEM images for this sample. Small particles could exhibit antiferromagnetic behavior which induces additional character to a well-known ferromagnetic behavior of BaFe_2O_4 . This possibility is supported by the TEM image of that sample besides the absence of a clear IR spectrum for it (see Fig. 5a). In addition, previous works recorded similar observation, for $\text{SrFe}_{12}\text{O}_{19}$ and $\text{BaFe}_{12}\text{O}_{19}$, with no explanation [34, 40].

TABLE 3
MAGNETIC PROPERTIES AND PARTICLE SIZE D OF $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$
SAMPLES SINTERED AT 700°C

x	M_r (emu/g)	M_{20} (emu/g)	M_s (emu/g)	H_c	D (nm)
0.0	0.772	2.46	2.87	2880	40.9
0.1	0.147	1.03	1.33	315	40.0
0.2	0.013	0.231		342	42.3
0.3	0.024	0.387		485	44.2
0.4	0.033	0.532		453	43.4

MAGNETIC PROPERTIES AND PARTICLE SIZE D OF $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$
SAMPLES SINTERED AT 1050°C

x	M_r (emu/g)	M_{20} (emu/g)	M_s (emu/g)	H_c	D (nm)
0.0	1.153	2.44	2.56	5260	57.9
0.1	0.749	1.17	1.75	761.4	56.6
0.2	0.0247	0.200		436.9	53.3
0.3	0.0062	0.153		402.6	47.0
0.4	0.0149	0.188		578.5	44.1

Sintering at 1050 °C was sufficient to enhance the crystallinity and to increase the crystallite size from 40.9 to 57.9 nm, the fact which resulted in almost doubling the coercivity (2880 to 5260 Oe).

Low substitution of Co cations ($x = 0.1$) resulted in a clear reduction of magnetization and a collapse in coercivity, which could be explained as follows. First, the magnetic moment of Co^{2+} ($3\mu\text{B}$) cation is smaller than that of the substituted Fe^{3+} ($5\mu\text{B}$) one. Second, the substitution itself weakens the strength of the magnetic exchange interaction of $\text{Fe}^{3+}-\text{O}^{2-}-\text{Fe}^{3+}$ [41]. Finally, the reduction in M_s could be related to the existence of some oxygen vacancies that maintain the electron neutrality of the structure when Co^{2+} replaced Fe^{3+} [40]. Those oxygen vacancies, beside the substantial disruption of the nearest-neighbor magnetic surrounding of Fe^{3+} cations, could explain the steep reduction of coercivity, as compared to the undoped sample. On sintering the same sample to 1050 °C, the oxygen vacancies are compensated by the oxidation of Co^{2+} to Co^{3+} or Fe^{3+} to Fe^{4+} with a resulting reduction of the unit-cell volume, a subsequent enhancement of magnetic exchange interaction, and thus an increase in M_s . Moreover, the crystallite size has significantly increased (Table 4) producing a partial recovery of coercivity (an increase from 315 to 761 Oe).

With higher degrees of Co^{2+} substitution ($x \geq 0.2$), the magnetization was significantly diminished; the magnetic

behavior of the samples deviated strongly from being ferromagnetic and also saturation magnetization was absent for any of them. Samples sintered at 700 °C exhibited small differences in magnetization, while all showed almost identical behavior for sintering temperature of 1050 °C. Worth to mention is that, the crystallite size hardly increased with increasing sintering temperature for these samples, and so, no significant changes in H_c was observed. In addition, continuing nonstoichiometric substitution of Fe^{3+} by Co^{2+} drastically eliminated the magnetic super interaction between the magnetic moments of Fe^{3+} cations. In an attempt to provide electron neutrality, the oxidation of Co^{2+} and/or Fe^{3+} to nonmagnetic higher valence states is highly probable. This induced both an increase in the distance between nearest-neighbor Fe^{3+} ions and also a reduction of the unit-cell volume; both affect the super-exchange interaction oppositely [41, 42]. With increasing sintering temperature, the concentration of nonmagnetic (low spin state) Co^{3+} [43] and Fe^{4+} [44] increases and the cell-volume effect is minimized. Considering both factors, the M_{20} value increases with x ($x \geq 0.2$) for samples sintered at 700 °C (Table 3), when the effect of unit-cell volume dominates. For the same samples sintered at 1050 °C, the role of high-valence cations dominates that of the cell volume when M_{20} is almost constant with x (Table 4).

It is indeed beneficial that the controlled Co substitution enables the tailoring of the magnetic behavior of BaFe_2O_4 ; it transformed it from hard to soft magnetic material with Co^{2+} substitution ratio does not exceed 10%. This is a quite desirable characteristic for recording media with low noise level at high recording density [45].

C. Electrical properties

Ac conductivity

Fig.6 shows the frequency-dependent ac conductivity plot. From this figure, the conductivity shows an increase trend with an increase in frequency for all samples. The conductivity in ferrites may be explained by Verwey's hopping mechanism [46]. According to this mechanism, the electrical conductivity in ferrites is mainly due to hopping of electrons between ions of the same element present in more than one oxidation state. The hopping probability depends upon the separation between ions involved and the potential barrier that must be overcome, which is known as the activation energy for hopping [47]. In addition, the ac conductivity has been related to the hopping of electrons

Figure.6 Variation of ac conductivity for BaFe_2O_4 and $\text{BaCo}_{0.2}\text{Fe}_{1.8}\text{O}_4$ as a function of $\log(f)$

between Fe^{3+} and Fe^{2+} and the hopping rate increase with the increase in applied field [48]. As compared to grains, grain boundary is dominant at low frequency which resulting in slow hopping between Fe^{2+} and Fe^{3+} ions. While at high frequency, grains influence more that encourage the hope mechanism [49].

An increase in the sintering temperature increases the grain size [50, 51]. Larger size implies the reduction of the concentration of less conductance grain bounders as compared to that of more conductance grains, which enable easier flow for hopping electrons increased and therefore, a lower barrier height. Which could interprets the increase in Ac conductivity with increasing sintering temperature.

Dielectric properties

The variation of dielectric constant (ϵ') with frequency for pure barium ferrite (BaFe_2O_4) and cobalt doped barium ferrite $\text{BaCo}_{0.2}\text{Fe}_{1.8}\text{O}_4$ sintered at 700 °C and 1050 °C has been depicted in Fig. 7.

In all samples the dielectric constant decreases continuously with increasing frequency and becomes almost constant in high frequency region. The decrease in dielectric constant with increasing frequency of applied field is observed by other researchers in a number of ferrite [52]. According to Maxwell–Wagner and Koops theory [53, 54], ferrites consist of highly conducting grains separated by grain boundaries with poor conductance. For ultrafine nanoparticles, grain-boundaries-to-grains volume ratio is quite larger in comparison with their bulk counterparts. The effect of grain boundaries dominates at low frequencies. This observation is in agreement with other researchers [55–57]. At low frequency the space charge polarization is supposed to be responsible for high dielectric constant value. According to Koops model, the charge carriers cause an increase in both space charge and rotational polarizations at the grain boundaries for low-frequency regime. Particles in nano regime have larger surface grains area, which results in larger space-charge polarization and a subsequent increase in ϵ' . Furthermore, loosely bound electrons in oxygen anions constitute an additional source for polarization in dielectric materials through aligning themselves along the even small applied electric field [58]. That dielectric polarization with weak electric field is the reason for the recorded high ϵ' at low frequency. The dielectric polarization in ferrites matches with the mechanism of electrical conduction; it is equivalent to the electronic exchange between ferrous and ferric ions at the octahedral sites ($\text{Fe}^{2+} \leftrightarrow \text{Fe}^{3+} + e^{-}$). The exchange of electron in ferrite systems has been responsible for dielectric polarization i.e. the displacement of electrons in the direction of the applied electric field. The further reduction of ϵ' with f is attributed to the fact that beyond a certain frequency of external field, the electronic exchange between the ferrous and ferric ions cannot follow the applied electric field [53, 59]. With increasing the sintering temperature, the dielectric constant of a specified sample also increases. This could be attributed to the introduction of sufficient thermal energy for liberating localized dipoles that are capable of aligning themselves with the applied field direction [60]. Worth to noting is that, upon doping Co^{2+} (for $\text{BaCo}_{0.2}\text{Fe}_{1.8}\text{O}_4$) the dielectric constant increased which could be attributed to the increased in hopping sites between Fe^{3+} and Fe^{4+} and/or Co^{2+} and Co^{3+} besides the ferrous and ferric hopping.

Figure 7. Variation of dielectric constant for BaFe_2O_4 and $\text{BaCo}_{0.2}\text{Fe}_{1.8}\text{O}_4$ vs. Log (f)

Fig. 8 depicted the variation of dielectric loss (ϵ'') as a

Figure 8. Variation of dielectric loss factor for BaFe_2O_4 and $\text{BaCo}_{0.2}\text{Fe}_{1.8}\text{O}_4$ vs. log (f)

function of frequency for pure BaFe_2O_4 and $\text{BaCo}_{0.2}\text{Fe}_{1.8}\text{O}_4$ sintered at 700 and 1050 °C. At low frequencies, the dipoles interactions are high and decreases with increasing frequency; thereby a remarkable reduction in dielectric loss is observed [61]. Upon introducing Co cations, for $\text{BaCo}_{0.2}\text{Fe}_{1.8}\text{O}_4$, both dielectric constant and dielectric loss increase, which could be attributed to oxygen vacancies, grain boundary defects, valence variations and possibly interfacial polarization [62].

CONCLUSION

A series of Co-doped barium monoferrite compounds $\text{BaCo}_x\text{Fe}_{2-x}\text{O}_4$ ($0.0 \leq x \leq 0.4$) have been synthesized by sol-gel auto combustion method with the subsequent heat treatment at 700 and 1050 °C. XRD confirmed the formation of pure orthorhombic phase for $x = 0.0 - 0.2$, while few traces of BaFe_3O_4 and Ba_2FeO_4 appeared with increasing either x or sintering temperature. The average crystallite size and unit cell volume were calculated for each sample. The

unit cell volume are found to decrease with increasing Co content ($x \geq 0.2$) for both sintering temperature owing to the formation of high valence cations, Fe^{4+} and/or Co^{3+} . FTIR data has confirmed the phase structure and the presence of higher valence state of Fe^{3+} . The TEM images confirmed XRD calculations. The magnetic behavior of the samples was thoroughly correlated with their structural and compositional properties of the samples. A high degree of flexibility in tailoring M_s and H_C values is suggested in view of the presented discussion. Moreover, the substitution of non-stoichiometric Co^{2+} ions in mono-barium ferrite results in increasing ac conductivity, dielectric constant and dielectric loss.

REFERENCES

- [1] P. V. Baptista, "Cancer nanotechnology-prospects for cancer diagnostics and therapy," *Current Cancer Therapy Reviews*, vol. 5, pp. 80-88, 2009.
- [2] C. J. Murphy, T. K. Sau, A. M. Gole, C. J. Orendorff, J. Gao, L. Gou, S. E. Hunyadi, and T. Li, "Anisotropic metal nanoparticles: synthesis, assembly, and optical applications," ed: ACS Publications, 2005.
- [3] S. Harker, G. Stewart, W. Hutchison, A. Amiet, and D. Tucker, "Microwave absorption and 57 Fe Mössbauer properties of Ni-Ti doped barium hexaferrite," *Hyperfine Interactions*, vol. 230, pp. 205-211, 2015.
- [4] I. A. Auwal, A. Baykal, H. Güngüneş, and S. E. Shirsath, "Structural investigation and hyperfine interactions of $\text{BaBi}_x\text{La}_x\text{Fe}_{12-2x}\text{O}_{19}$ ($0.0 \leq x \leq 0.5$) hexaferrites," *Ceramics International*, vol. 42, pp. 3380-3387, 2016/02/01/ 2016.
- [5] L. Gonsalves and V. Verenkar, "Synthesis and characterization of nanosize nickel-doped cobalt ferrite obtained by precursor combustion method," *Journal of thermal analysis and calorimetry*, vol. 108, pp. 877-880, 2012.
- [6] S. Majetich, J. Scott, E. Kirkpatrick, K. Chowdary, K. Gallagher, and M. McHenry, "Magnetic nanoparticles and magnetocrystalline anisotropy," *Nanostructured Materials*, vol. 9, pp. 291-300, 1997.
- [7] A. K. Menon and B. Gupta, "Nanotechnology: a data storage perspective," *Nanostructured Materials*, vol. 11, pp. 965-986, 1999.
- [8] M. Rajendran, R. Pullar, A. Bhattacharya, D. Das, S. Chintalapudi, and C. Majumdar, "Magnetic properties of nanocrystalline CoFe_2O_4 powders prepared at room temperature: variation with crystallite size," *Journal of Magnetism and Magnetic Materials*, vol. 232, pp. 71-83, 2001.
- [9] A. Mulyawan, W. A. Adi, S. Mustofa, and A. Fisli, "The Phase Transformation and Crystal Structure Studies of Strontium Substituted Barium Monoferrite," in *IOP Conference Series: Earth and Environmental Science*, 2017, p. 012070.
- [10] R. Candeia, M. Souza, M. Bernardi, S. Maestrelli, I. Santos, A. Souza, and E. Longo, "Monoferrite BaFe_2O_4 applied as ceramic pigment," *Ceramics International*, vol. 33, pp. 521-525, 2007.
- [11] M. Sugimoto, "The past, present, and future of ferrites," *Journal of the American Ceramic Society*, vol. 82, pp. 269-280, 1999.
- [12] S. Da Dalt, B. B. Sousa, A. K. Alves, and C. P. Bergmann, "Structural and photocatalytic characterization of BaFe_2O_4 obtained at low temperatures," *Materials Research*, vol. 14, pp. 505-507, 2011.
- [13] S. Mandizadeh, M. Salavati-Niasari, and M. Sadri, "Hydrothermal synthesis, characterization and magnetic properties of BaFe_2O_4 nanostructure as a photocatalytic oxidative desulfurization of dibenzothiophene," *Separation and Purification Technology*, vol. 175, pp. 399-405, 2017.
- [14] R. Candeia, M. Souza, M. Bernardi, S. Maestrelli, I. Santos, A. Souza, and E. Longo, "Monoferrite BaFe_2O_4 applied as ceramic pigment," *Ceramics International*, vol. 33, pp. 521-525, 2007.
- [15] I. C. Nlebedim, N. Ranvah, Y. Melikhov, P. Williams, J. Snyder, A. Moses, and D. Jiles, "Effect of temperature variation on the magnetostrictive properties of $\text{CoAl}_x\text{Fe}_{2-x}\text{O}_4$," *Journal of Applied Physics*, vol. 107, p. 09A936, 2010.
- [16] S. Singhal and K. Chandra, "Cation distribution and magnetic properties in chromium-substituted nickel ferrites prepared using aerosol route," *Journal of solid state chemistry*, vol. 180, pp. 296-300, 2007/01/01/ 2007.
- [17] A. M. Wahba, M. B. Mohamed, and N. Imam, "Correlating structural, magnetic, and luminescence properties with the cation distribution of $\text{Co}_{0.5}\text{Zn}_{0.5+x}\text{Fe}_{2-x}\text{O}_4$ nanoferrite," *Journal of Magnetism and Magnetic Materials*, vol. 408, pp. 51-59, 2016.
- [18] A. M. Wahba and M. B. Mohamed, "Structural and magnetic characterization and cation distribution of nanocrystalline $\text{Co}_x\text{Fe}_{3-x}\text{O}_4$ ferrites," *Journal of Magnetism and Magnetic Materials*, vol. 378, pp. 246-252, 2015.
- [19] J. Dho, E. Lee, J. Park, and N. Hur, "Effects of the grain boundary on the coercivity of barium ferrite $\text{BaFe}_{12}\text{O}_{19}$," *Journal of Magnetism and Magnetic Materials*, vol. 285, pp. 164-168, 2005.
- [20] A. Ataie, M. Piramoon, I. Harris, and C. Ponton, "Effect of hydrothermal synthesis environment on the particle morphology, chemistry and magnetic properties of barium hexaferrite," *Journal of Materials Science*, vol. 30, pp. 5600-5606, 1995.
- [21] J. Wang, "Prepare highly crystalline NiFe_2O_4 nanoparticles with improved magnetic properties," *Materials Science and Engineering: B*, vol. 127, pp. 81-84, 2006.
- [22] M. George, A. M. John, S. S. Nair, P. Joy, and M. Anantharaman, "Finite size effects on the structural and magnetic properties of sol-gel synthesized NiFe_2O_4 powders," *Journal of Magnetism and Magnetic Materials*, vol. 302, pp. 190-195, 2006.
- [23] K. Khaja Mohaideen and P. A. Joy, "Influence of initial particle size on the magnetostriction of sintered cobalt ferrite derived from nanocrystalline powders," *Journal of Magnetism and Magnetic Materials*, vol. 346, pp. 96-102, 2013/11/01/ 2013.
- [24] A. R. West, *Solid state chemistry and its applications*: John Wiley & Sons, 2014.
- [25] S. Castro, M. Gayoso, and C. Rodríguez, "A study of the combustion method to prepare fine ferrite particles," *Journal of solid state chemistry*, vol. 134, pp. 227-231, 1997.
- [26] D. P. Padiyan, A. Marikani, and K. Murali, "Influence of thickness and substrate temperature on electrical and photoelectrical properties of vacuum-deposited CdSe thin films," *Materials Chemistry and Physics*, vol. 78, pp. 51-58, 2003.
- [27] A. K. Zak, W. A. Majid, M. E. Abrishami, and R. Yousefi, "X-ray analysis of ZnO nanoparticles by Williamson-Hall and size-strain plot methods," *Solid State Sciences*, vol. 13, pp. 251-256, 2011.
- [28] B. Cullity, "Elements of X-ray diffraction: a practical approach," ed: Addison-Wesley Publishing Company Inc., California, 1956.
- [29] N. A. Spaldin, *Magnetic materials: fundamentals and applications*: Cambridge University Press, 2010.
- [30] R. Dilip and R. Jayaprakash, "Synthesis and characterization of BaFe_2O_4 nano-ferrites for gas sensor applications," *Energy, Ecology and Environment*, vol. 3, pp. 237-241, 2018.
- [31] H. Saravani, M. R. Esmaeilzadei, and M. T. Ghahfarokhi, "Synthesis and Characterization of Ferromagnetic BaFe_2O_4 Nanocrystals Using Novel Ionic Precursor Complex $[\text{Fe}(\text{opd})_3]_2[\text{Ba}(\text{CN})_8]$," *Journal of*

- Inorganic and Organometallic Polymers and Materials, vol. 26, pp. 353-358, 2016.
- [32] S. Wang, X. Zu, G. Sun, D. Li, C. He, X. Xiang, W. Liu, S. Han, and S. Li, "Highly dispersed spinel (Mg, Ca, Ba)-ferrite nanoparticles: Tuning the particle size and magnetic properties through a modified polyacrylamide gel route," *Ceramics International*, vol. 42, pp. 19133-19140, 2016.
- [33] Y. Yang, Y. Jiang, Y. Wang, Y. Sun, L. Liu, and J. Zhang, "Influences of sintering atmosphere on the formation and photocatalytic property of BaFe₂O₄," *Materials Chemistry and Physics*, vol. 105, pp. 154-156, 2007.
- [34] T. Xie, L. Xu, and C. Liu, "Synthesis and properties of composite magnetic material SrCo_xFe_{12-x}O₁₉ (x=0-0.3)," *Powder Technology*, vol. 232, pp. 87-92, 2012/12/01/ 2012.
- [35] B. P. Barbero, J. A. Gamboa, and L. E. Cadús, "Synthesis and characterisation of La_{1-x}Ca_xFeO₃ perovskite-type oxide catalysts for total oxidation of volatile organic compounds," *Applied Catalysis B: Environmental*, vol. 65, pp. 21-30, 2006/05/15/ 2006.
- [36] A. Gholizadeh, "The effects of A/B-site substitution on structural, redox and catalytic properties of lanthanum ferrite nanoparticles," *Journal of Materials Research and Technology*, 2018.
- [37] M. B. Mohamed, A. M. Wahba, and Z. K. Heiba, "Effect of Zn substitution on structural, magnetic, and electric properties of Ni_{1-x}Zn_xFe_{1.78}Al_{0.2}Gd_{0.02}O₄ nanoparticles," *Journal of Superconductivity and Novel Magnetism*, vol. 28, pp. 3675-3683, 2015.
- [38] L. Kumar and M. Kar, "Influence of Al³⁺ ion concentration on the crystal structure and magnetic anisotropy of nanocrystalline spinel cobalt ferrite," *Journal of Magnetism and Magnetic Materials*, vol. 323, pp. 2042-2048, 2011.
- [39] A. Jaafar, S. Arekat, A. Al-Saie, and M. Bououdina, "Structure and magnetic properties of nanosized BaFe₂O₄ material," *International Journal of Nanoscience*, vol. 9, pp. 575-577, 2010.
- [40] G. B. Teh, S. Nagalingam, and D. A. Jefferson, "Preparation and studies of Co (II) and Co (III)-substituted barium ferrite prepared by sol-gel method," *Materials Chemistry and Physics*, vol. 101, pp. 158-162, 2007.
- [41] A. Thakur, R. Singh, and P. Barman, "Structural and magnetic properties of La³⁺ substituted strontium hexaferrite nanoparticles prepared by citrate precursor method," *Journal of Magnetism and Magnetic Materials*, vol. 326, pp. 35-40, 2013.
- [42] H. Wang, Y. Hai, B. Yao, Y. Xu, L. Shan, L. Xu, J. Tang, and Q. Wang, "Tailoring structure and magnetic characteristics of strontium hexaferrite via Al doping engineering," *Journal of Magnetism and Magnetic Materials*, vol. 422, pp. 204-208, 2017.
- [43] N. A. Merino, B. P. Barbero, P. Ruiz, and L. E. Cadús, "Synthesis, characterisation, catalytic activity and structural stability of LaCo_{1-y}FeyO_{3±λ} perovskite catalysts for combustion of ethanol and propane," *Journal of catalysis*, vol. 240, pp. 245-257, 2006.
- [44] J. Yang, X. Zhou, Z. Chu, W. Hikal, Q. Cai, J. Ho, D. Kundaliya, W. Yelon, W. James, and H. Anderson, "Charge disproportionation and ordering in La_{1/3}Sr_{2/3}FeO_{3.8}," *Journal of Physics: Condensed Matter*, vol. 15, p. 5093, 2003.
- [45] K. Noma, N. Matsushita, S. Nakagawa, and M. Naoe, "Study on S- and R-block construction with additional spinel blocks in sputter-deposited Ba ferrite films," *IEEE Transactions on Magnetics*, vol. 33, pp. 3637-3639, 1997.
- [46] S. Hussain and A. Maqsood, "Structural and electrical properties of Pb-doped Sr-hexa ferrites," *Journal of Alloys and Compounds*, vol. 466, pp. 293-298, 2008.
- [47] S. M. Ghahfarokhi, F. Ranjbar, and M. Z. Shoushtari, "A study of the properties of SrFe_{12-x}Co_xO₁₉ nanoparticles," *Journal of Magnetism and Magnetic Materials*, vol. 349, pp. 80-87, 2014.
- [48] M. George, S. S. Nair, K. Malini, P. Joy, and M. Anantharaman, "Finite size effects on the electrical properties of sol-gel synthesized CoFe₂O₄ powders: deviation from Maxwell-Wagner theory and evidence of surface polarization effects," *Journal of Physics D: Applied Physics*, vol. 40, p. 1593, 2007.
- [49] D. Paramesh, K. V. Kumar, and P. V. Reddy, "Effect of aluminium substitution on the electrical properties of Ni-Zn nanoferrites," *Journal of Magnetism and Magnetic Materials*, vol. 444, pp. 371-377, 2017.
- [50] R. L. Coble, "Sintering crystalline solids. I. Intermediate and final state diffusion models," *Journal of Applied Physics*, vol. 32, pp. 787-792, 1961.
- [51] A. Verma, T. Goel, R. Mendiratta, and P. Kishan, "Magnetic properties of nickel-zinc ferrites prepared by the citrate precursor method," *Journal of Magnetism and Magnetic Materials*, vol. 208, pp. 13-19, 2000.
- [52] M. Bhandare, H. Jamadar, A. Pathan, B. Chougule, and A. Shaikh, "Dielectric properties of Cu substituted Ni_{0.5-x}Zn_{0.3}Mg_{0.2}Fe₂O₄ ferrites," *Journal of Alloys and Compounds*, vol. 509, pp. L113-L118, 2011.
- [53] K. W. Wagner, "Zur theorie der unvollkommenen dielektrika," *Annalen der Physik*, vol. 345, pp. 817-855, 1913.
- [54] C. Koops, "On the dispersion of resistivity and dielectric constant of some semiconductors at audiofrequencies," *Physical Review*, vol. 83, p. 121, 1951.
- [55] N. Sivakumar, A. Narayanasamy, B. Jeyadevan, R. J. Joseyphus, and C. Venkateswaran, "Dielectric relaxation behaviour of nanostructured Mn-Zn ferrite," *Journal of Physics D: Applied Physics*, vol. 41, p. 245001, 2008.
- [56] D. Kothari, S. Phanjoubam, and J. Baijal, "Electrical conduction and dielectric behaviour of the oxidic spinel Li_{0.5+0.5x}Cr_{0.3}Ti_xFe_{2.2-1.5x}O₄," *Journal of Materials Science*, vol. 25, pp. 5142-5146, 1990.
- [57] R. Manjula, V. Murthy, and J. Sobhanadri, "Electrical conductivity and thermoelectric power measurements of some lithium-titanium ferrites," *Journal of Applied Physics*, vol. 59, pp. 2929-2932, 1986.
- [58] M. J. Iqbal and S. Farooq, "Could binary mixture of Nd-Ni ions control the electrical behavior of strontium-barium M-type hexaferrite nanoparticles?," *Materials Research Bulletin*, vol. 46, pp. 662-667, 2011.
- [59] K. M. Batoo, S. Kumar, and C. G. Lee, "Influence of Al doping on electrical properties of Ni-Cd nano ferrites," *Current Applied Physics*, vol. 9, pp. 826-832, 2009.
- [60] S. Joshi, M. Kumar, S. Chhoker, G. Srivastava, M. Jewariya, and V. Singh, "Structural, magnetic, dielectric and optical properties of nickel ferrite nanoparticles synthesized by co-precipitation method," *Journal of Molecular Structure*, vol. 1076, pp. 55-62, 2014.
- [61] I. Afandiyeva, I. Dökme, Ş. Altındal, M. Bülbül, and A. Tataroğlu, "Frequency and voltage effects on the dielectric properties and electrical conductivity of Al-TiW-Pd₂Si/n-Si structures," *Microelectronic Engineering*, vol. 85, pp. 247-252, 2008.
- [62] Z. Haijun, L. Zhichao, M. Chenliang, Y. Xi, Z. Liangying, and W. Mingzhong, "Preparation and microwave properties of Co- and Ti-doped barium ferrite by citrate sol-gel process," *Materials Chemistry and Physics*, vol. 80, pp. 129-134, 2003.