Experimental and Theoretical Study of the Structural, Magnetic and Electronic Properties of the Ba₅GdSbO₉ Perovskite

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Abstract: In this work the procedure to the synthesis of Ba₅GdSbO₉ complex perovskite by the solid-state reaction method is reported. Theoretically a study of the crystalline and electronic structure was performed into the framework of the Density Functional Theory (DFT). The most stable structure is obtained to be a rhombohedral perovskite with a lattice constant a = 6.0840 Å. Due the occurrence of a mean energy gap of 2.84 eV close to the Fermi level for both up and down spin polarizations this material is classified as insulator. The effective magnetic moment of material obtained from the calculations was 7.0 μB. The crystalline structure was analyzed through the X-ray diffraction technique and Rietveld refinement of the experimental data. Results are strongly in agreement with those theoretically predicted. Magnetic response was studied from measurements of magnetic susceptibility as a function of temperature. Results reveal the paramagnetic feature of this material in the temperature regime from 50 K up to 300 K. From the fitting with the Curie law the effective magnetic moment was obtained to be 8.1 μB, which is slightly higher than the theoretical value for the Gd³⁺ isolated cation predicted by the theory of paramagnetism. The energy gap obtained from experiments of diffuse reflectance is relatively in agreement with the theoretical predictions. The dielectric constant as a function of applied frequencies at room temperature was measured. Results reveal a decreasing behavior with a high value of dielectric constant at low applied frequencies.

Keywords: Perovskite Material, Structure, Magnetic Feature, Electronic Properties

1. INTRODUCTION

Recently, there has been a great interest in the electronic oxide ceramics oxides with ABO₃ perovskite structure due to the chemical versatility to evaluate different combinations of transition metal cation in the B site and obtain a variety of magnetic and electric properties [1, 2]. Ideally, a simple ABO₃ perovskite possesses a 3-dimensional cubic network of BO₆ octahedra with the cation A into the void formed by eight octahedral with AO₁₂ coordination [3]. If the A and B sites contain a mixture of cations, distorted complex perovskites structures with lower symmetries are obtained. Distortions are mainly related with octahedral tilting due to a small cation driving a rotating of octahedral respect the crystallographic axes. Other kind of distortion is the cation displacement caused through first and second-order Jahn-Teller effects. Specially, the 1:1 B-site based class with chemical formula A₅BB'O₉ (or double perovskites) is the most frequently complex perovskite encountered [4]. From the distortions and composition of the double perovskites, new properties and applications are found in ferroelectric devices with large dielectric constants and frequency dispersion [5], magnetoresistance [6], solar energy conversion [7], catalysis [8], solid-oxide fuel cells [9] and microwave resonator [10]. In this paper we propose the synthesis and characterization of Ba₅GdSbO₉ ceramic material. We describe the crystalline structure of this double perovskite and perform morphological and compositional analyses. Furthermore, we present results of measurements of the magnetic response as a function of temperature. Moreover, considering that in recent years the density functional theory (DFT) has constituted in a strong tool to study electronic properties in
perovskite-like material [11], we carried out a study of the electronic properties of these materials, in order to establish the type of hybridization between the orbitals of GdO₆ and SbO₆ octahedra present in the structure.

2. EXPERIMENTAL

Samples were synthesized through the standard solid-state reaction recipe. Precursor powders of Gd₂O₃, Sb₂O₃, and Ba₃CO₃ (Aldrich 99.9%) were stoichiometrically mixed according to the chemical formula Ba₃GdSbO₈. Mixture was ground to form a pellet and annealed at 1000 °C for 30 h. The samples were then reground, repelletized and sintered at 1100 °C for 40 h and 1200 °C for 40 h. X-ray diffraction (XRD) experiments were performed by means of a PW1710 diffractometer with λCuKα=1.54064 Å. Rietveld refinement of the diffraction patterns was carried out by the GSAS program [12]. Morphological studies were performed by means scanning electron microscopy (SEM) experiments through the utilization of VEGA 3 equipment, and chemical composition of samples was analyzed by energy dispersive X-ray (EDX) technique. Field cooling measurements of the magnetic susceptibility as a function of temperature were carried out by using an MPMS Quantum Design SQUID. Diffuse reflectance experiments were performed by using a VARIAN Cary 5000 UV-Vis-NIR spectrophotometer, which has an integration sphere with a PMT/Pbs detector. The relative dielectric constant was measured by using an Agilent HP4194A-350 frequency analyzer.

3. CALCULATION METHOD

In order to determine the electronic and band structures we applied the Full-Potential Linear Augmented Plane Wave method (FP-LAPW) within the framework of the Kohn-Sham Density Functional Theory (DFT) [13], and adopted the Generalized Gradient (GGA) approximation for the exchange-correlation energy due to Perdew, Burke and Ernzerhof [14]. The self-consistent process is developed by the numeric package Wien2k [13]. Taking the experimental unit cell data as input, the structure studied in this work were fully relaxed with respect to their lattice parameters and the internal degrees of freedom compatible with the space group symmetry of the crystal structure. The resulting energies versus volume functions have been fitted to the equation of state due to Murnaghan [15] in order to obtain de minimum energy value, the bulk modulus, its pressure derivative and the equilibrium lattice parameters and associated volume. The muffin-tin radiiuses used for Ba₃GdSbO₈ were 2.20; 2.50; 2.12 and 1.82 Bohr for Ba, Gd, Sb and O respectively, angular momentum up to l = 10 inside the muffin-tin sphere, a maximum vector in the reciprocal space of Gmax=12.0; RMT*Kmax = 7.0 and a mesh of 1000 points in the first Brillouin zone (equivalent to a maximum of 250 k points in the irreducible Brillouin zone). Finally, the convergence criterion for the self-consistent calculation was 0.0001 Ry for the total energies, 0.0001 Bohr in the charge and 1.0 mRy/μ.a in the internal forces. Spin polarization was included in the calculations.

4. RESULTS AND DISCUSSION

Figure 1 shows the XRD patterns obtained for Ba₃GdSbO₈. In the picture black symbols represent the experimental data and line corresponds to simulated pattern by means of GSAS code. Base line is the difference between theoretical and experimental results. Refinement parameters of figure 1 were χ²=5.236, R(F2)=5.84%. From the Rietveld refinement it was established that these materials crystallize in a rhombohedral perovskite structure, space group R-3 ($\bar{3}$148). The structural parameter obtained from the refinement was a=5.9882(1) Å. These results are 99.3% in agreement with the theoretical values obtained from the Structure Prediction Diagnostic Software SPhDS [16], which predicts a=6.0293 Å and α=57.8634°. In these ceramic materials the explanation of distortion from the ideal cubic perovskite structure is clear because the double perovskite have the generic formula $A_2BB'O_6$, and for this type of material the tolerance factor $\tau$, is
calculated by the ratio $\tau = \frac{r_A + r_o}{\sqrt{3(r_B + r'_B + r_o)}}$, where $r_A$, $r_B$, $r'_B$, and $r_o$ are the ionic radii of the A, B, B', and O ions, respectively.

If $\tau$ is equal to unity, there is ideal cubic perovskite structure, and if $\tau < 1$ the structures is distorted from the cubic symmetry. The value of tolerance factor obtained for the complex perovskite was 0.9081. The Wyckoff positions obtained from the refinement analysis are presented in Table 1.

From the structural analysis above, the expected structure of the $\text{Ba}_2\text{GdSbO}_6$ complex perovskite is that presented in Figure 2. Clearly, a tolerance factor as far from the ideal value is related to distortions from the cubic cell. As showed in Figures 2a and 2b, there is a difference between directions of the $\text{GdO}_6$ and $\text{SbO}_6$ octahedra. The most important has to do with the nature of the rhombohedral structure, since the trigonal angle $\alpha = 60,0290^\circ$ is quite far from 90° degrees corresponding to the ideal cubic cell. Furthermore, the tilt of the $\text{GdO}_6$ and $\text{SbO}_6$ octahedra occurs out of phase for an angle $\beta = 16,4650^\circ$, corresponding to the Glazer notation a−a−a−, whose symmetry involves reflections, signaling odd-odd-odd reflections, which is characterized by cationic ordering and simple

<table>
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<tr>
<th>Atom</th>
<th>Wyckoff Site</th>
<th>$x$</th>
<th>$y$</th>
<th>$z$</th>
</tr>
</thead>
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<tr>
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<td>2c</td>
<td>0,2500</td>
<td>0,2500</td>
<td>0,2500</td>
</tr>
<tr>
<td>Gd</td>
<td>1a</td>
<td>0,0000</td>
<td>0,0000</td>
<td>0,0000</td>
</tr>
<tr>
<td>Sb</td>
<td>1a</td>
<td>0,5000</td>
<td>0,5000</td>
<td>0,5000</td>
</tr>
<tr>
<td>O1</td>
<td>6f</td>
<td>-0,1771</td>
<td>-0,3613</td>
<td>0,2705</td>
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</table>
octahedral tilting.

SEM images of the material shown in figure 3 reveal a qualitative approximation to the surface microstructure. In picture (a) evidences the formation of clusters of polyhedral grains and interstitial particulate grains. From picture (b) it is clear that the grains could end up having a few tens of nm, forming groups, which have appearance of clusters of several μm sizes.

By applying the EDX microprobe of the VEGA3 TESCAN microscope, the percentages of the constituent elements of the material were obtained, as reported in table 2. These values confirm that the material does not contain other elements than those corresponding to the expected stoichiometry Ba₂GdSbO₆. However, the experimental result reveals an apparent low content of the oxygen anion against an increase in the percentage of Ba, Gd and Sb cations, which may be because the resolution of the device is slightly affected because oxygen is a very light element compared to the other constituents of the compound.

Measurements of magnetic susceptibility as a
Table 2. Compositional percentages of elements for Ba2GdSbO6 obtained from EDX.

<table>
<thead>
<tr>
<th>Element</th>
<th>Theoretical %</th>
<th>Experimental %</th>
</tr>
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<tbody>
<tr>
<td>Ba</td>
<td>42.28</td>
<td>43.33</td>
</tr>
<tr>
<td>Gd</td>
<td>24.20</td>
<td>24.81</td>
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<tr>
<td>Sb</td>
<td>18.74</td>
<td>19.10</td>
</tr>
<tr>
<td>O</td>
<td>14.78</td>
<td>12.76</td>
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Fig. 3. Granular topology of the surface obtained from SEM images for the Ba$_2$GdSbO$_6$ perovskite-like material.

A function of temperature reveal the paramagnetic character of the Ba$_2$GdSbO$_6$ complex perovskite. Figure 4 shows the Curie fitting performed following the equation $X = X_0 + (C/T)$. In the Curie equation, $C = N\mu_{eff}^2/3K_B$ is the Curie constant, $N$ is Avogadro’s number, $\mu_{eff}$ is the effective magnetic moment ($\mu_{eff} = P_{eff} \mu_B$), $P_{eff}$ represent the effective Bohr magneton number, $\mu_B$ is the Bohr magneton, $K_B$ is the Boltzmann constant and $X_0 = 0.00139$ emu/mol is the temperature independent susceptibility term. From the Curie constant $C$ the effective magnetic moments for Ba$_2$GdSbO$_6$ material was calculated to be 8.1 $\mu_B$. This value is 98% in agreement with the theoretical expected moments obtained from the Hund’s rules [19]. The difference is attributed to other magnetic effects due to the hybridization of

Fig. 4. Magnetic behaviour of Ba$_2$GdSbO$_6$ material obtained from measurements of susceptibility as a function of temperature. Red line corresponds to the fit with the Curie law.
the d-Sb and f-Gd orbital and non-preferential spin orientations in the GdO₃ and SbO₃ octahedra due structural distortions.

In order to obtain the most accurate result, the optimal lattice parameter corresponding to the minimal energy value was calculated as the total energy versus volume. The total energy was calculated by fittings with the Murnaghan’s state equation [15]. From total energy as a function of volume the ideal lattice parameter was determined to be a=6.0840 Å. Figure 5 shows the band structure for both up and down spin polarizations. In the picture, E=0 eV corresponds to the Fermi level. It is observed that these materials evidence weak semiconductor behavior with gap energy through the Fermi level from -0.05 eV up to 2.81 eV (width 2.86 eV) for the spin up polarization and between -0.05 eV and 2.78 eV (width 2.83 eV) for the spin down orientation.

The Partial Densities of States (DOS) for the Ba₂GdSbO₆ complex perovskite are exemplified in figure 6. As in the figure 5, the energy zero corresponds to the Fermi level reference. It is important to elucidate that close the Fermi level there are contributions due to the O-2p hybridizations but Gd-4f spin up orbitals are responsible for priority contributions. Very incipient contributions of Sb-3d states are observed. This behavior can be clearly seen in Figures 5 and 6 in the energy range between -2.87 and -0.05 eV. Localized states attributed to Sb-3d, Ba-6s and O-2p appear far the Fermi level, below -3.00 eV. On the other hand, in the conduction band, it is observed in figures 5 and 6 that available states are majority due to the Gd-4f spin down orbitals with very small contributions of the Sb-3d and O-2p spin up and down levels.

The magnetic moment of mixed charge density was calculated to be 7.0 μₜ for Ba₂GdSbO₆. This value is relatively different from the value obtained experimentally. However, an important element to be taken into account has to do with the clutter of Gd and Sb cations in the structure of the double perovskite, which can not only distorted octahedra but also modify the magnetic response of the material.

Results of diffuse reflectance experiments are shown in figure 7. Reflectance values were acquired at 1 nm intervals over the 300–2500 nm,
as shown in figure 7a. The absorption of UV-Vis-NIR radiation reveals that the molecules of the Ba$_2$GdSbO$_6$ material cause the excitation of electrons from the ground state to exited state, which produces electronic jumps between quantum levels. The quantum characteristics energy, which depends from the electronic configuration, is calculated from the radiated energy. By applying the Kubelka-Munk model, currently used to the measurement of the band gap in powder samples, the respective band gap for the Ba$_2$GdSbO$_6$ perovskite is obtained as showed in figure 7b. Structural defects or vacancies can produce spontaneous absorption that appears for several wavelengths, as observed in figure 7a.

From the adjustment to the Kubelka-Munk equation it was determined that the energy gap of the Ba$_2$GdSbO$_6$ is 3.61 eV at room temperature, which is 33% different from the theoretic prediction given by the DOS calculations for T=0 K. This difference occurs because in the DFT calculations, the exchange and correlation potential by Perdew, Burke and Ernzerhof [14] gives a very good approximation for the valence and conduction density of states, but is not very accurate for determining the energy gap in
semiconductors due to self-interaction errors [18-19], for which it would be more interesting to use a potential specifically designed for this purpose, that uses local functional without Hartree-Fock exchange [20-21].

The dielectric characteristics of this compound were obtained from measurements of the dielectric constant as a function of frequency performed. The corresponding results are shown in figure 8. As shown in the picture, at low frequencies (ν=100 Hz) the value of the dielectric constant εr=5400, which is a tendency of the giant-dielectric materials [22]. The value of the dielectric constant decreases dramatically with increasing frequency between 100 Hz and 1000 Hz. Above ν=1000 Hz, the dielectric constant decreases smoothly to the value εr=40 at ν=10 kHz. There are reports according to which it is expected that most giant dielectric constants are due interconnected effects of polarization [23]. In semiconductors, for example, the formation of Schottky barriers at the contacts could lead to depletion regimes. Another example has to do with the formation of internal barrier layer capacitors through grain boundaries, which produce relaxations Maxwell-Wagner type, resulting in very high dielectric constant [24]. It is not excluded that the behavior of the dielectric constant depending on the frequency has to do with the occurrence of relaxation processes attributed to hopping of oxygen O²⁻ ions and (O2)²⁻ peroxide between multiple off-center sites of the crystal [22].

5. CONCLUSIONS

The Ba₂GdSbO₆ complex has been synthesized by the solid-state reaction procedure. A study of the crystalline and electronic structure was performed through the Linearized Augmented Plane Wave method (FP-LAPW) into the framework of the Density Functional Theory (DFT). The results reveal that a rhombohedral perovskite is the more stable structure with a lattice parameter a=6.0840 Å. Close to the Fermi level a mean energy gap of 2.84 eV was observed for both up and down spin polarizations, which permitted to classify this material as insulator. The largest contribution to the effective magnetic moment, due to the Gd⁺⁺ cations, evidenced a value of 7.0 μB. The crystalline structure was experimentally determined by means measurements in a Panalytical XPert Pro X-ray diffractometer with a Rietveld refinement of the diffraction pattern through de GSAS code. The experimental lattice parameter a=5.9882(1) Å is 98.4% in agreement with those theoretically predicted. Measurements
of magnetic susceptibility as a function of temperature reveal the paramagnetic feature of this material. From the fitting with the Curie law the effective magnetic moment was obtained to be 8.1μ₀, which is slightly higher that the theoretical value for the Gd³⁺ isolated cation predicted by μ_{eff} = g\sqrt{J(J+1)}\mu_B, where g represents the Landé’s factor and J is the quantic number related to the 4f electronic orbital. Experiments of diffuse reflectance and fittings with the Kubelka-Munk equation permitted to obtain an energy gap of 3.61 eV, which differs 33% when compared with the theoretical value predicted by DFT. This difference is attributed to the temperature effect.

ACKNOWLEDGEMENTS

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