Investigation of Thermal, Structural and Electrical Properties of LaMnO$_3$-Sigma

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Abstract: LaMnO$_3$ is one of the most intensively studied cathode materials for solid oxide fuel cell (SOFC). In the present study, LaMnO$_3$-d$_x$ cathode materials were synthesized by combustion method. The properties of synthesized materials like thermal, structural, electrical and electrochemical were investigated. Thermogravimetric analysis (TGA/DTA) analysis confirms that the calcination temperature (1000°C) is the appropriate temperature for the preparation of the materials using La(NO$_3$)$_3$.6H$_2$O, MnSO$_4$.H$_2$O, Co(NO$_3$)$_2$.6H$_2$O, Fe(NO$_3$)$_3$.9H$_2$O and C$_6$H$_8$O$_7$.H$_2$O precursors. The X-ray powder diffraction (XRD) results of the materials reveal the formation of the hexagonal structure with R3C space group Fd$ar{3}$m. The scanning electron microscope (SEM) characterization shows that the prepared samples have slightly porous structure with agglomerated particles. The energy dispersive spectroscopy (EDS) analysis is also confirms the presence of La, Mn and O elements in all synthesized materials. From the fourier transform infrared spectroscopy (FTIR) analysis, the most significant absorption bands located at 1629.8 cm$^{-1}$ and 589.9 cm$^{-1}$ wave numbers are identified. The room temperature conductivity of the sample is found to be 6.3×10$^{-3}$ for LaMnO$_3$ cathode material. From the dielectric constant $\varepsilon'$ as a function of frequency observed that value of $\varepsilon'$ maximum at lower frequencies and it begins to drop and becomes constant at higher frequencies.

Keywords: Perovskites, SOFC, Cathode Materials

1. Introduction

A major concern of international community is the requirement of energy to an increasing demand under the constraints of a limited fossil fuel supply. The development of energy generation methods that are not limited to fuel supply has, therefore, been widely investigated for many years. The urgency of such research has further increased since the environmental issues associated with fossil fuel combustion (global warming, ocean acidification, smog, low air quality) have become well recognized [1, 2].

One solution to this energy challenge is the hydrogen powered fuel cell. This cell operates on a potentially limitless supply of fuel through different method while producing water and heat as byproduct of the electrochemical reaction that generates electricity as can be seen in equation 1 [3].

$$\frac{1}{2}O_2 + H_2 \rightarrow H_2O + Heat + Electricity$$ (1)

There are different types of fuel cells and they are named according to the material used in the electrolyte. These types are alkaline fuel cell (AFC), phosphoric acid fuel cell (PAFC), polymeric electrolyte membrane fuel cell (PEMFC), molten carbonate fuel cell (MCFC), and solid oxide fuel cell (SOFC). In addition to the above category fuel cell also differ in other important aspects such as efficiency and operating temperature [4].

Solid oxide fuel cell (SOFC) is a type of fuel cell which consists of a perovskite structure. The above material structure is widely applied and studied. Except for sealant, all other components of the SOFC can potentially consist of perovskite ceramics. The most common materials found in the single SOFC configuration are Yittra Stabilized Zirconia (YSZ) in the electrolyte, Ni/YSZ cement in the anode and Strontium doped lanthanum manganite (LSM) in the cathode. The cathode is the most common material in which reduction takes place. According to the outcome of different research the composition of the material,
synthesis method and fabrication have significant impact on the cathode material of SOFC. In most cases, cathode is composed of a perovskite material [4]. Therefore, this study focuses only on cathode perovskite material.

This research work focuses on a mixture of different oxides of manganese which crystallize in perovskite structure called manganites. Manganites have a broad stoichiometric formula of ABO$_{3-\delta}$, where A is lanthanide element and B is manganese. This also includes the ability to generate an exact or no oxygen stoichiometry. One of the main characteristics of manganites perovskites compared with other families of oxides in the wide variety of substitutions in its crystal structure. The lanthanides are among the 25 elements which can occupy the position at A sites. On the other hand, apart from the manganese, almost 50 different elements can occupy the B-site [5].

Manganite compounds crystallize in the structure ABO$_3$ perovskites can have an ideal cubic structure of Pm3m space group, orthorhombic Pbnm space group or rhombohedra R3C space group [6].

The stoichiometry with the corresponding valance states to manganite is A+$x$B$^{2+}$O$_3^{-2}$ with a body centered (BCC) cubic unit cell whose center sits the A$^{3+}$ cation that is usually the largest one; B$^{2+}$ cations occupy the eight apex of the cell and O$^{-2}$ an ions occupy the mid points between cations and in the middle of the edges of the bcc cell [7].

The materials used in SOFC must satisfy many requirements in terms of electrochemical properties. Besides, chemical and physical compatibility must exist between the materials for each component. As SOFC runs in high temperature, a chemical decomposition or a chemical reaction can’t occur during its operation and the chosen materials must have a similar thermal expansion coefficient in order to avoid the formation of cracks during thermal cycling. Perovskite oxides display special attention due to their superior electrical, magnetic, catalytic and superconductive properties that can be an advantage to technological and basic materials science investigations [8].

This work investigates LaMnO$_{3.5}$ perovskite, and the primary objectives are on the synthesis of the advanced ceramic perovskites nano powder using citric acid combustion approach and characterization using different physical and thermal techniques. Among the different synthesis methods, the reason for the selection of this method is that it is characterized by fast heating rates; high temperatures and short reaction times [9]. Thermal decomposition behavior of the compounds were investigated by thermal analysis method. The crystalline phases were developed on heat treatment and the structures of all compounds were studied by XRD and FTIR. Morphology of the powders was characterized by SEM. The electrical and electrochemical properties were conducted using complex impedance spectroscopy (CIS).

2. Method

2.1. Experimental Method of Synthesis

LaMnO$_{3.5}$ perovskites was synthesized by combustion synthesis method using nitrate salts as starting materials and C$_6$H$_{12}$O$_6$H$_2$O as chelating agent. Stoichiometric amounts of La (NO$_3$)$_3$.6H$_2$O and Mn (NO$_3$)$_2$.6H$_2$O was dissolved in some amount of deionized water and mixed together. The mixed metal nitrate solution then added to the citric acid solution in 1:2 molar ratios. Then the solution was placed on magnetic stirrer making the stirrer switch on unlike the heat is switched off. The stirring procedure without applying heat carried on for one hour and fourteen minutes.

After that, aqueous ammonia solution was added drop wise to the mixed solution on heating condition 70°C under constant stirring using magnetic stirrer until the pH of the solution maintains about 7.

The mixed solution with pH7 was constantly stirred by magnetic stirrer on a hot plate at 80°C for about 12 hours in the water bath and stirred continuously to obtain a dark brown gel and combustion takes place. Then after it completely burnt out and form a powder. Further, the obtained Powder was grounded for one hour using get mortar.

The calcinations were carried at a temperature of 1000°C for five hour to obtain composite oxide powder. The various steps involved in combustion method for synthesis of cathode materials are shown in the flow chart of Figure 1 below.

2.2. Pellet Preparation

An ionic, electronic and dielectric property of all synthesized materials was carried out using complex impedance spectroscopy. For this purpose, calcined fine powder was prepared. The preparation process started after combustion. After the final result of combustion the combusted compound placed in a furnace. Applying 1000°C for five hour, the furnace cooled down and using get mortar the result was grounded. Then, polyvinyl alcohol (PVA) was used as a binder and it was released in burning during high temperature sintering to prepare the pellet. After this entire step, the final result was cold pressed in to circular disc shaped pellet at a pressure of 5 tons in a stainless-steel dye set using a hydraulic press. After cold pressed the pellets was sintered at 1000°C for 4h in an alumina crucible in air atmosphere followed by slow cooling process. Subsequently, the pellet was polished by a fine emery paper to make their faces smooth and parallel. The pellets was finally coated with high purity conductive silver paint and dried at 100°C for 30 minutes to remove the moisture.
3. Results and Discussion

3.1. Thermal Analysis

In order to investigate the thermal behavior of the LSM samples, around 15mg of the mixture of La(NO₃)₃·6H₂O, MnSO₄·H₂O and C₆H₈O₇·H₂O precursors were used. The precursors were heated over the temperature range between room temperature and 1100°C under nitrogen atmosphere. The obtained curves are represented in Figures 2. As it can be seen in the figure, thermal decomposition takes place in different stages, depending on the decomposition behavior of

Figure 1. Schematic diagram powder preparation process.
each precursor and the significant weight loss regions as well as the stable phase formation region are also identified.

Figure 2 shows the TGA/DTA behavior of the mixture of La\(\left(\text{NO}_3\right)\)_\text{3},6H\text{2}O, MnSO\text{4},H\text{2}O and C\text{6}H\text{7}O\text{2},H\text{2}O precursors to form LaMnO\text{3} cathode material. It is observed that the first endothermic peak is appeared at 85.2°C with respective weight loss of 3.13% due to the moisture absorbed during storage. Further, the DTA curve observed at 223.6°C which is accompanied by a weight loss of 11.4% refers to the loss of the water used during synthesis procedure. The other DTA peak at 329.7°C intended by a weight loss of 32.7% refers to the decomposition of citric acid and followed by the exothermic DTA peak formed at 583.6°C refers to the combustion and formation of LaMnO\text{3} phase. At higher temperatures, the TGA curve becomes more flattened, indicating the stable phase formation of LaMnO\text{3}. This indicates that 1000°C temperature can be used for preparation of LaMnO\text{3} material using La\(\left(\text{NO}_3\right)\)_\text{3},6H\text{2}O, MnSO\text{4},H\text{2}O and C\text{6}H\text{7}O\text{2},H\text{2}O precursors. Moreover, the total loss of sample weight up to 1100°C is 68.5%.

3.2. Structural Analysis

For further explanation of LaMnO\text{3} in case of structural analysis two materials were synthesized by substitution on the two different sites that is the A-site and B-site. Therefore, the XRD patterns of LaMnO\text{3}, La\(_{0.8}\text{Sr}_{0.1}\text{Co}_{0.1}\text{MnO}_3\) and La\(_{0.8}\text{Sr}_{0.1}\text{Fe}_{0.1}\text{MnO}_3\) cathode materials synthesized by sol-gel combustion method heated at 1000°C for 4 hours in air are shown in Figure 3. It is observed that all the diffraction peaks are strong and sharp, indicating highly crystalline of the cathode materials.

The diffraction peaks which are indexed as (110), (020), (200), (022), (202), (220), (130), (312) and (204) represent of the perovskite phase with hexagonal structure with space group of R-3C [6], which is in line with Joint Committee on Powder Diffraction Standards (JCPDS) card number 00-051-1515. However, additional peaks for Co and Fe substituted samples are appeared, which correspond to the impurity for LaMnO\text{3} compound. However, as compared the XRD of the three samples, the diffraction peaks of La\(_{0.8}\text{Sr}_{0.1}\text{Co}_{0.1}\text{MnO}_3\) and La\(_{0.8}\text{Sr}_{0.1}\text{Fe}_{0.1}\text{MnO}_3\) samples are shifted slightly towards higher angles. This indicates the fact that the substitution of Co\text{2+}, Sr\text{2+} and Fe\text{3+} cations for La\text{3+} cations has led to the decrease in the crystal size of LaMnO\text{3}.

Figure 3. XRD patterns of the synthesized samples.

The lattice parameters, the unit cell volumes and the crystal sizes of each sample were calculated from XRD patterns, and the obtained results are summarized in Table 1. It is found that the lattice parameters, unit cell volumes and crystal sizes of LaMnO\text{3} material are slightly larger than that of La\(_{0.8}\text{Sr}_{0.1}\text{Co}_{0.1}\text{MnO}_3\) and La\(_{0.8}\text{Sr}_{0.1}\text{Fe}_{0.1}\text{MnO}_3\) materials. This is associated with the ionic radius differences of the substituted cations. As compared the substituted materials, Co substituted La\(_{0.8}\text{Sr}_{0.1}\text{Co}_{0.1}\text{MnO}_3\) has larger lattice parameter (a=7.2476Å and c=7.8201Å), unite cell volume (355.7378Å\text{3}) and crystal size (507.63nm) than Fe substituted La\(_{0.8}\text{Sr}_{0.1}\text{Fe}_{0.1}\text{MnO}_3\) material. This behavior can be understood from the ionic radii that is the ionic radius of Co\text{2+} (0.79 Å) [9] is larger than that of Fe\text{3+} (0.64 Å) [10]. The variation of lattice parameters and shift the diffractions line, suggest the incorporation of the dopants in the lattice of both cathodes.
Table 1. Lattice constant, unit cell volume and crystal size of synthesized samples.

<table>
<thead>
<tr>
<th>Samples</th>
<th>Lattice Constants (Å)</th>
<th>UnitCellVolume V (nm³)</th>
<th>CrystalSize from (200) (Å)</th>
</tr>
</thead>
<tbody>
<tr>
<td>LaMnO₃</td>
<td>7.2439 7.8252</td>
<td>355.6097</td>
<td>507.63</td>
</tr>
<tr>
<td>La₀.8Sr₀.1Co₀.1MnO₃</td>
<td>7.2476 7.8201</td>
<td>355.7378</td>
<td>408.16</td>
</tr>
<tr>
<td>La₀.8Sr₀.1Fe₀.1MnO₃</td>
<td>7.2348 7.8305</td>
<td>353.9521</td>
<td>157.63</td>
</tr>
</tbody>
</table>

3.3. SEM and EDS Study

To study the morphology and the grain size distributions of LaMnO₃-δ cathode in the form of powder, the sample is characterized by using scanning electron microscope. Figure 4 shows SEM images of all samples synthesized by sol-gel combustion method at calcination temperature of 1000°C. It is observed that the synthesized samples have fairly uniform grain size distribution, indicating well developed crystallinity. This is consistent with well-defined diffraction peaks observed from XRD patterns. It is also observed that all the prepared samples have porous structure and they have slightly agglomerated particles. The porous structure of the cathode material is important for the insertion and deinsertion of cations during charging and discharging process.

The morphology of LaMnO₃ is more compact as compared to the morphology of the materials found by substitution of elements with ionic radius larger than the elements in the A-site and B-site of the first compound.

The elemental analysis of LaMnO₃ of cathode materials was analyzed by energy dispersive spectroscopy (EDS). As shown in the figure 4, all the synthesized materials show the presence of La, Mn, and O. No other elements are detected in the investigated manganite, which indicates the purity of the synthesized products. This is consistent with the result obtained in the XRD patterns.

3.4. FT-IR Spectroscopy Study

Figure 5 shows the room temperature FT-IR spectra of the synthesized LaMnO₃ and La₀.8Sr₀.1(Co, Fe)₀.1MnO₃ cathode materials in the wavelength range of 350 to 2500 cm⁻¹. As it is observed in the Figure 5, different similar strong absorption bands are identified at different wave number regions. The most significant absorption bands are located between 712.6 and 484.4 cm⁻¹ wave numbers. These bands are assigned to stretching vibrations due to interactions between the cations and oxygen in the synthesized compounds. This indicates the formation of the perovskite-type structures [11].

The band appeared at wavenumber 568.3 cm⁻¹ is responsible for the formation of LaMnO₃ compound, which attributed to stretching vibration of Mn-O band of MnO₆ octahedron. As compared with LaMnO₃, the absorption of Sr, Co and Fe substituted materials are shifted slightly towards to higher wave number regions. This indicates that the substitution of these cations into LaMnO₃ has led to the decrease in octahedral site radius. This result is in good agreement with lower lattice parameters and unit cell volume calculated from the XRD patterns. On the other hand, peaks observed at around 1629.8 and 2365.6 cm⁻¹ are attributed to the deformation mode of absorbed molecular water of the carrier KBr (H₂O)n and CO₂, respectively [12].

Figure 4. SEM and EDS images of LaMnO₃ of cathode materials.

Figure 5. FT-IR spectra of a) LaMnO₃, b) La₀.8Sr₀.1Co₀.1MnO₃. c) La₀.8Sr₀.1Fe₀.1MnO₃ of cathode materials.
La_{0.8}Sr_{0.1}Co_{0.1}MnO_{3} where as the amount of transmittance of La_{0.8}Sr_{0.1}Fe_{0.1}MnO_{3} is shallower than the two compounds

3.5. Electrochemical Impedance Spectroscopy Analysis

Impedance spectra can be plotted in two ways: in the Nyquist representation where the negative imaginary part of the impedance, $Z''$, is plotted versus the real part of the impedance, $Z$ or in the Bode representation where the absolute impedance, $|Z|$, as well as the phase shift is plotted versus the angular frequency. Even if both representations contain, in principle, the same information and can be transformed into each other, for this research work Nyquist plot was used to see the relation between real and imaginary impedance.

Figure 6. Cole-Cole plots for a) LaMnO_{3} b) La_{0.8}Sr_{0.1}Co_{0.1}MnO_{3-δ} c) La_{0.8}Sr_{0.1}Fe_{0.1}MnO_{3-δ} of cathode materials.

In Figure 6 shown above the real versus imaginary impedance for three compounds show single arc, which indicates that the electrical process contribution is from bulk materials (grain interior), which can be modeled as an equivalent circuit comprising of a parallel combination of bulk resistance and bulk capacitance. This can be seen in other work that one arc can be modeled as an equivalent circuit comprising of a parallel combination of bulk resistance and bulk capacitance [13]. The figure 6 starts from zero ohms for both real and imaginary line and goes to 195000 ohm for real and 21000 ohm for imaginary. The figure indicates that there is a direct relationship between real and imaginary for the three compounds. The imaginary value of La_{0.8}Sr_{0.1}Co_{0.1}MnO_{3-δ} greater than LaMnO_{3} and LaMnO_{3} greater than La_{0.8}Sr_{0.1}Fe_{0.1}MnO_{3-δ} on the same value real impedance 6000 ohm. This difference is not the same at higher value of real impedance. Towards the left below 1500 ohm of real impedance a small arc is expected. These arcs are usually related to the interfacial resistance between an electrolyte and a cathode (grain boundary conduction or charge transfer) at an intermediate frequency and the oxygen reduction reaction at the cathode surface (diffusion or concentration polarization) at a low frequency [14].

Figure 7. Real impedance ($Z'$) versus ln(f) for a) LaMnO_{3} b) La_{0.8}Sr_{0.1}Co_{0.1}MnO_{3-δ} c) La_{0.8}Sr_{0.1}Fe_{0.1}MnO_{3-δ} of cathode materials.

According to Figure 7 the real impedance value decreases with increasing the logarithm of frequency. Below 5Hz of ln(f) the real impedance of LaMnO_{3} lies between the other two compounds found by substitution process. The real impedance of LaMnO_{3} decreases as the value of ln(f) increase. The decrease proceeds with no fluctuation like that of La_{0.8}Sr_{0.1}Fe_{0.1}MnO_{3} cathode material. Above 6Hz of ln(f) the compounds found by element substitution on A-site and B-site of LaMnO_{3} show a similar decrease in the amount of real impedance. Above the logarithmic frequency of twelve the real impedance approaches zero for the three synthesized compounds.

Figure 8. Imaginary impedance ($Z''$) versus ln(f) for a) LaMnO_{3} b) La_{0.8}Sr_{0.1}Co_{0.1}MnO_{3-δ} c) La_{0.8}Sr_{0.1}Fe_{0.1}MnO_{3-δ} of cathode materials.

The above Figure 8 shows the comparison of the imaginary impedance with the variation of the frequency. As the frequency increases, the amount of imaginary impedance decreases just like the real impedance. Unlike that of real impedance, the variation of frequency versus imaginary impedance is almost similar from logarithmic frequency of ‘8’ and above for the three compounds. For lower
logarithmic frequency below 6Hz, the imaginary impedance of \( \text{La}_{0.8}\text{Sr}_{0.1}\text{Co}_{0.1}\text{MnO}_{3-\delta} \) greater than \( \text{LaMnO}_3 \), \( \text{La}_{0.8}\text{Sr}_{0.1}\text{Fe}_{0.1}\text{MnO}_{3-\delta} \). The decrease in imaginary impedance of the synthesized material \( \text{LaMnO}_3 \) is nearly similar to the compound \( \text{La}_{0.8}\text{Sr}_{0.1}\text{Fe}_{0.1}\text{MnO}_{3-\delta} \). The variation of dielectric constant \( \varepsilon' \) with frequency was recorded for the compound. Figure 9 shows this variation with frequency on the x-axis and dielectric constant on the y-axis. As the frequency increases the dielectric constant decreases for the compound. Specifically, at around logarithmic frequency of 5Hz \( \text{La}_{0.8}\text{Sr}_{0.1}\text{Co}_{0.1}\text{MnO}_3 \) compound has a larger dielectric constant than \( \text{LaMnO}_3 \) and \( \text{La}_{0.8}\text{Sr}_{0.1}\text{Fe}_{0.1}\text{MnO}_{3-\delta} \) whereas the dielectric constant of \( \text{La}_{0.8}\text{Sr}_{0.1}\text{Fe}_{0.1}\text{MnO}_{3-\delta} \) is the smallest. This implies that \( \text{La}_{0.8}\text{Sr}_{0.1}\text{Co}_{0.1}\text{MnO}_3 \) compound have more available polarization mechanisms as compared to the other compound synthesized for this research work. In general, as frequency increases, the material’s net polarization drops as each polarization mechanism ceases to contribute, and hence its dielectric constant drops. In addition, the room temperature conductivity of all synthesized perovskite materials were calculated using equation and it is found to be \( 6.3 \times 10^{-3} \), \( 8.7 \times 10^{-3} \) and \( 5.8 \times 10^{-3} \) S.cm\(^{-1} \) for \( \text{LaMnO}_3 \), \( \text{La}_{0.8}\text{Sr}_{0.1}\text{Co}_{0.1}\text{MnO}_{3-\delta} \) and \( \text{La}_{0.8}\text{Sr}_{0.1}\text{Fe}_{0.1}\text{MnO}_{3-\delta} \) of cathode materials, respectively.

4. Conclusions

The overall performance of solid oxide fuel cell depends up on its cathode material. The compositions of the material, synthesis method have significant impact on the cathode material of SOFC. The result of DTG shows that the synthesized \( \text{LaMnO}_3 \) materials have both endothermic and exothermic peak with the irrespective temperature. The TGA result of the compound shows the minimum calcination temperature which is 1000°C. The structural property was investigated by using XRD. The result obtained from XRD for the three materials indicates that the lattice constant of \( \text{La}_{0.8}\text{Sr}_{0.1}\text{Fe}_{0.1}\text{MnO}_{3-\delta} \) was larger than the other two but the crystal size was less than the other two materials. The unit cell volume of \( \text{LaMnO}_3 \) material was less than the other two. The SEM was helpful in differentiating the morphology of the synthesized materials. For the material agglomerations can easily be observed in their SEM image. In combination with this the EDS shows the presence of each elements in each compound and the pure formation of the compound. The result from FTIR shows the three most absorption bands. These most significant absorption bands are located between 712.6 and 484.4 cm\(^{-1} \) wave numbers. In addition, the electrical and electrochemical property was revealed by using LCR measurement. Hence, the conductivities of the synthesized material is \( 6.3 \times 10^{-3} \).

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