Original Article

Fabrication of Ba Doped LaMnO₃ Nanomaterials by Microwave Combustion Method

Tran Thi Ha¹, Pham Nguyen Hai², Tran Thi Uyen², Nguyen Ngoc Dinh², Nguyen Thi Ha Thu³, Ho Khac Hieu⁴,⁵, Nguyen Viet Tuyen²,*

¹Faculty of Basic Sciences, Hanoi University of Mining and Geology, 18 Vien Duc Thang, Hanoi, Vietnam
²Faculty of Physics, VNU University of Science, 334 Nguyen Trai, Thanh Xuan, Hanoi, Vietnam
³Air Defence - Air Force Academy, Kim Son, Son Tay, Hanoi, Vietnam
⁴Institute of Research and Development, Duy Tan University, 03 Quang Trung, Da Nang, Vietnam
⁵Faculty of Natural Sciences, Duy Tan University, 03 Quang Trung, Hai Chau, Da Nang, Vietnam

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Abstract: The major energy sources that human are relying on are fossil fuel and it is exhausted day by day. Furthermore, exploitation and usage of fossil fuel also bring in many negative impacts on environmental pollution. Solid oxide fuel cell (SOFC) has been considered as a potential solution for such environmental and energy problems. Seeking a facile, cost and time saving process to synthesize Ba doped LaMnO₃ is very important for development of SOFC applications because it helps to reduce the cost of commercial SOFC. LaMnO₃ doped with Ba is more cost effective than Sr doped LaMnO₃ because less rare earth elements are used. At the same time, its conductivity is still good enough with an appropriate thermal expansion matched with those of other parts of SOFC based on yttrium stabilized zirconia (YSZ).

In this paper, Ba doped LaMnO₃ nanoparticles (LBMO), a promising material for making cathode of SOFC, were prepared by microwave combustion method. This material has many advantages. Effect of the amount fuel in the combustion reaction on the products was studied by various methods such as: X-ray diffraction measurement, scanning electron microscopy, energy dispersive X-ray spectroscopy. The results showed that each doping rate requires an appropriate amount of fuel to obtain pure and crystalline product. The obtained LBMO nanoparticles are crystalline in hexagonal phase at doping ratio of 0.2 and orthorhombic phase at doping ratio of equal or larger than 0.3.

Keywords: Perovskite, microwave combustion, Ba doped LaMnO₃ nano particle.

*Corresponding author.
Email address: nguyenviettuyen@hus.edu.vn
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1. Introduction

In the next several decades, minimizing emission gases due to burning fossil fuels is extremely important to reduce the impact on environment pollution. A promising solution is SOFC because it can transform energy from chemical bonding of fuel (bio-gas, natural gas, hydrogen, etc.) directly into electricity via electrochemical reactions at high yield while reduce much toxic emission gases [1–4]. The most commonly used materials in cathode of SOFC are ABO$_3$ perovskites, where A is a rare earth metal (La, Sr, etc.) and B is a transition metals such as: Mn, Fe, Co, Ni or the combination of these metals [2, 5–7]. The catalytic ability of oxidation reduction process is provided by cation B. With an appropriate selection of A and B, a number of oxygen vacancies are generated during working process and thus support for the transport of a large amount of oxygen to the electrolyte layer [8–12]. Lanthanum manganite is a perovskite with p type conduction, fulfills the requirements for cathodes of SOFC, which are high conductivity, suitable thermal expansion, stability in both oxidized and reduced ambient at high temperature [8, 9]. Recently, Ba was reported as a promising dopant, which can help to increase the conductivity and magnetic property of LaMnO$_3$ [15–18].

In this research, Ba which is an earth alkali metal of abundant amount and low cost, was a doped into LaMnO$_3$ with the hope that partial replacement of rare earth elements with low cost one can help to gain better price and electrical properties of cathode of SOFC.

2. Experiment

![Diagram of synthetic process of LBMO by microwave combustion method.](image)

The synthetic process is summarized in the diagram shown in Figure 1. Precursor materials are La$_2$O$_3$ (99.99%), Mn(NO$_3$)$_2$$\cdot$4H$_2$O (99%) and glycine NH$_2$CH$_2$COOH (99%). La$_2$O$_3$ was dissolved in HNO$_3$ acid (65%) to obtain La(NO$_3$)$_3$ solution. Then, La(NO$_3$)$_3$, Ba(NO$_3$)$_2$$\cdot$4H$_2$O and Mn(NO$_3$)$_2$$\cdot$4H$_2$O were mixed together to form a solution with La : Ba : Mn molar ratio of (1-x) : x : 1 where x = 0.2; 0.3; 0.5; 0.6; 0.7 is the doping rate. In the next step, glycine of appropriate amount was added to the solution. A set of La$_{1-x}$Ba$_x$MnO$_3$ samples was prepared and denoted as BxGy where, x is the doping
rate and y is defined as molar ratio between glycine and Mn. The obtained solution would be heated by a magnetic stirrer at 100 °C. After 40 min the color of solution would turn into milky gel. The gel was transferred into a microwave oven and heated in 2 min at power of 800 W. The microwave supported for the combustion reaction and produced black powder.

3. Results and Discussion

Glycine plays an important role in the formation of metal complexes, which guarantees a homogenous reaction at atomic level. Moreover, glycine supports for the reduction of nitrate group in combustion process and liberates a huge amount of heat for the formation of pervoskite crystal. Hence, investigating the effect of fuel amount on structure and morphology of nanoproduct prepared by microwave combustion is very important. A small amount of fuel which is below threshold will not provide enough energy for the reaction to occur. On the other hand, excess amount of fuel will also degrade the quality of the product because generated gas in the reaction will dissipate the useful heat for the reaction. Simultaneously, doping concentration also changes kinetics of reaction. The XRD results (Figure 2) show that when same amount fuel is utilized (y=5), the quality of the product changes a lot depending on the doping rate (x = 0.3 or 0.4). XRD diffraction patterns show that La$_{0.6}$Ba$_{0.4}$MnO$_3$ sample was in crystalline form, implied by characteristic peaks of LaMnO$_3$, without dopant or impurity. Meanwhile, sample La$_{0.7}$Ba$_{0.3}$MnO$_3$ was amorphous even though the same ratio of fuel to precursors was applied. Thus, La$_{0.7}$Ba$_{0.3}$MnO$_3$ nanomaterial was prepared with three different y values: 4.5; 4.75 and 5. It should be noted that y of less than 4.5 resulted in non crystalline product as demonstrated by XRD patterns in Figure 3.

![Figure 2. XRD patterns of LBMO power samples with doping rate x = 0.3 and 0.4.](image-url)
Figure 3. XRD patterns of $\text{La}_{0.7}\text{Ba}_{0.3}\text{MnO}_3$ powder prepared with different value of $y$: 4.5; 4.75 and 5.

The above results suggest that each doping rate $x$ will require a corresponding appropriate fuel to precursors ratio.

Figure 4. XRD patterns of LBMO samples of different doping rate $x = 0.2; 0.3; 0.4; 0.5; 0.6; 0.7$.

The detailed investigation showed that fuel ratio of $y = 4$ offers $\text{La}_{0.8}\text{Ba}_{0.2}\text{MnO}_3$ of hexagonal phase. For samples with doping rate $x = 0.3; 0.4; 0.5$, the fuel to precursor ratio $y$ equals 4.75; 5; 5.25.
respectively will produce single phase nanopowder of orthorhombic structure. Samples with doping rate \(x\) greater than 0.6 cannot be obtained in crystalline form at any value of fuel ratio \(y\). These results are in agreement with theoretical calculation by Ahmed [19], where the authors showed that the greater the doping concentration is, the harder for the product can be formed due to the difference in ion radius of Ba and La. Ahmed et al. calculated the dependence of formation energy of LBMO materials on doping rate. The results are summarized in Table 1.

Table 1. Forming energy of LBMO product of different doping rate [19].

<table>
<thead>
<tr>
<th>X</th>
<th>0.125</th>
<th>0.25</th>
<th>0.375</th>
<th>0.5</th>
<th>0.625</th>
<th>0.75</th>
<th>0.875</th>
</tr>
</thead>
<tbody>
<tr>
<td>(E_{\text{form}}) (meV/atom)</td>
<td>438.2</td>
<td>518.6</td>
<td>588.0</td>
<td>676.1</td>
<td>773.5</td>
<td>883.9</td>
<td>984.6</td>
</tr>
</tbody>
</table>

A high formation energy implies difficulties of doping process. As doping rate increases, the formation energy grows correspondingly. Thus, a high doping rate will require more heat provided by fuel. However, the gas product dissipates some portion of heat to environment, which is not useful for the combustion reaction. This explains for the optimum value of fuel ratio for each doping concentration as investigated above.

Figure 5. XRD patterns of LBMO samples with doing rate \(x\) = 0.2; 0.3; 0.4; 0.5 in a 2\(\theta\) range of 32 and 33\(^\circ\).

As can be seen from Figure 5, the strongest peak of LBMO shows a clear shift toward higher angle as the doping concentration increases. This peak shift is an indication of replacement of La in the lattice by Ba ion. The decrease in distance between two neighbor planes with increasing of doping concentration can be explained by the smaller radius of Ba\(^{2+}\) ion compared with that of La. The lattice parameters of LBMO samples were estimated by using the following formula for distances between neighbor planes of orthorhombic and hexagonal structures, while the crystal size was
estimated by Sherrer formula. The estimated data (shown in Table 2) are in consistent with the values reported in literature for LaMnO$_3$ materials [3, 11].

Table 2. Lattice parameters and crystal size of LBMO samples of different doping concentration $x = 0.2; 0.3; 0.4; 0.5$.

<table>
<thead>
<tr>
<th>Sample</th>
<th>$a$</th>
<th>$b$</th>
<th>$c$</th>
<th>$D$</th>
<th>Phase</th>
</tr>
</thead>
<tbody>
<tr>
<td>$La_{0.8}Ba_{0.2}MnO_3$</td>
<td>4.147Å</td>
<td>4.147Å</td>
<td>13.463Å</td>
<td>19.60nm</td>
<td>Hexagonal</td>
</tr>
<tr>
<td>$La_{0.7}Ba_{0.3}MnO_3$</td>
<td>5.523Å</td>
<td>5.698Å</td>
<td>7.803Å</td>
<td>21.61nm</td>
<td>Orthorhombic</td>
</tr>
<tr>
<td>$La_{0.6}Ba_{0.4}MnO_3$</td>
<td>5.523Å</td>
<td>5.680Å</td>
<td>7.790Å</td>
<td>24.43nm</td>
<td>Orthorhombic</td>
</tr>
<tr>
<td>$La_{0.5}Ba_{0.5}MnO_3$</td>
<td>5.482Å</td>
<td>5.678Å</td>
<td>7.736Å</td>
<td>26.44nm</td>
<td>Orthorhombic</td>
</tr>
</tbody>
</table>

Elemental percentage of the nanoproducts determined by EDS spectra agrees well with the nominal content of the starting materials. For example, in the obtained $La_{0.8}Ba_{0.2}MnO_3$ sample (Figure 6), the atomic element ratio of La, Ba and Mn is $8:2:9.3$. This is close to the stoichiometric ratio $8:2:10$.

![Figure 6. EDS spectrum of LBMO sample with $x=0.2$.](image)

When doping Ba into LaMnO$_3$ lattice, Ba can either replace for La or go into interstition position, stay at the grain boundary, or dissociate to form another phase of Ba for instance BaCO$_3$. However, the combination of XRD and EDS data suggest that the second case can be neglected because the amount of Ba in the product consists with the nominal doping concentration while the product are single phase even at high doping concentration.

Microstructure of the product can affect greatly to the catalyst yield of ceramic materials, thus, the micromorphology of the samples was investigated by scanning electron microscope. The SEM images in Figure 7 show that the obtained products are nanoparticles in range of 100–300 nm. The uniform particle size demonstrates the potential of using the nanoproduct as a precursor materials for fabrication of thin films of perovskite for cathode of SOFC.
4. Conclusion

Ba doped LaMnO$_3$ nanopowder was prepared by microwave combustion method with many advantages such as: simple setup, low cost, time saving. The study showed that glycine as the fuel of the combustion reaction plays a critical role on the formation of perovskite nanomaterials. A fine control of fuel amount is necessary for the production of single phase product. The preparation of Ba
doped LaMnO$_3$ nanoparticles of single phase, high quality opens the possibility to fabricate nanocrystalline thin films of this material by electrophoresis for SOFC applications.

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