Effects of Mg\textsuperscript{2+} addition on structure and electrical properties of gadolinium doped ceria electrolyte ceramics

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Abstract

Series of Gd\textsuperscript{3+} and Mg\textsuperscript{2+} co-doped ceria (Ce\textsubscript{0.8}Gd\textsubscript{0.2-x}Mg\textsubscript{x}O\textsubscript{1.9-δ}) powders were prepared by a sol-gel method and electrolyte ceramics were obtained by sintering at 1300 °C. Thermogravimetric and differential scanning calorimetry, X-ray diffraction, scanning electron microscope and electrochemical impedance spectroscopy were used for structural, morphology and electrical characterization of the prepared samples. Well crystalline cubic fluorite structured composite was confirmed after calcination at 700 °C and the electrolyte ceramics sintered at 1300 °C for 4 h was quite dense with uniform microstructure. The electrochemical analysis results displayed that the highest conductivity has the Ce\textsubscript{0.8}Gd\textsubscript{0.14}Mg\textsubscript{0.06}O\textsubscript{1.87} compound, i.e. 0.0203 S/cm at 800 °C. Therefore, it was concluded that co-doping with Gd\textsuperscript{3+} and Mg\textsuperscript{2+} could enhance the electrical properties of the CeO\textsubscript{2} based solid electrolytes.

Keywords: ceramic electrolytes, co-doping, ceria, electrical properties

I. Introduction

Solid oxide fuel cell (SOFC) has been studied deeply as an advanced future renewable energy device due to its high-efficiency energy conversion and environmental friendliness [1–4]. Electrolyte plays the most important role in SOFC, as it determines the operating temperature to a large extent. CeO\textsubscript{2}-based electrolytes attracted considerable attention as their oxygen ion conductivity in intermediate temperature range (600–800 °C) is higher than that for yttria stabilized zirconia (YSZ). The conductivity can be even increased by doping with trivalent rare earth ions. When ceria is doped with rare earth and alkaline earth cations, oxygen vacancy is formed to compensate the charge balance in the lattice [5]. The defect reaction equation is as follows:

\[ \text{Gd}_2\text{O}_3 + 2\text{CeO}_2 \rightarrow 2\text{Gd}_\text{Ce}' + 3\text{O}_2 + \text{V}_0'' \]  (1)

The rare earth and alkaline earth ions (Gd\textsuperscript{3+}, Sm\textsuperscript{3+}, Y\textsuperscript{3+}, Ca\textsuperscript{2+} and Sr\textsuperscript{2+}) doped ceria at different concentrations have been extensively studied as an upcoming alternate solid electrolyte to YSZ [6–10]. Among them, Gd\textsuperscript{3+} doped ceria (GDC) is considered as promising electrolyte material due to its high oxygen ion conductivity, low polarization resistance and compatibility with electrodes [5,11].

Recently, co-doping with two or more aliovalent cations has been reported as more efficient strategy [12–17]. Muhammed Ali’s finding [18] confirmed that ionic conductivity of multi-doped ceria (Ce\textsubscript{0.8}Sm\textsubscript{0.1}Ba\textsubscript{0.05}Er\textsubscript{0.05}O\textsubscript{2-δ}) is considerably influenced by dopant radius and sintering temperatures with respect to surface morphology, microstructure and oxygen vacancy radius. Co-doped ceria was studied by Arabaci [19], who showed that Ce\textsubscript{0.9-x}Gd\textsubscript{x}Er\textsubscript{0.1}O\textsubscript{1.9-δ} materials can be used as electrolytes for IT-SOFC applications instead of GDC. Maheshwari [7] found that Sr\textsuperscript{2+}-Gd\textsuperscript{3+} co-doped CeO\textsubscript{2} samples show significantly higher total electrical than the electronic conductivities which recommended them as good materials for SOFC electrolytes. The addition of 1 mol% CuO in gadolinium doped ceria showed no detrimental effect on the total electrical conductivity [20].

In view of the above discussions, in the present investigation, gadolinium and magnesium co-doped ceria powders were prepared by a sol-gel route. The effects of Mg addition on the properties of ceria...
based electrolytes, the structure and the conductivity of Ce$_{0.8}$Gd$_{0.2-x}$Mg$_x$O$_{1.9-\delta}$ samples were investigated.

II. Experimental

2.1. Sample preparation

Ultrafine Ce$_{0.8}$Gd$_{0.2-x}$Mg$_x$O$_{1.9-\delta}$ powders (x = 0, 0.02, 0.04 and 0.06) were prepared by a sol-gel method. In this process, analytical pure cerium nitrate (Ce(NO$_3$)$_3$·6H$_2$O, containing 39 wt.% of CeO$_2$), gadolinia (Gd$_2$O$_3$, 99.99 wt.%), magnesium nitrate (Mg(NO$_3$)$_2$·6H$_2$O, analytical reagent) were used as starting materials. All raw materials were purchased from the China National Medicines Corporation Ltd. The experimental flowchart of Ce$_{0.8}$Gd$_{0.2-x}$Mg$_x$O$_{1.9-\delta}$ preparation using the sol-gel method is shown in Fig. 1. The stoichiometric amount of precursors were dissolved in deionized water and mixed together. Citric acid was then added as complexing agents to form a homogeneous solution, the molar ratio of citric acid to metal ions was 1.5 : 1. The obtained mixture was continuously stirred using a magnetic agitator to form a homogeneous solution at 80 °C and then converted to a viscous gel. The synthesized Ce$_{0.8}$Gd$_{0.2-x}$Mg$_x$O$_{1.9-\delta}$ powders were uniaxially pressed into discs of 11 mm in diameter and 0.5 mm in thickness under a pressure of 200 MPa. Finally, the green discs were sintered in a high-temperature furnace at 1300 °C for 4 h.

The xerogels were studied by simultaneous thermogravimetric and differential scanning calorimetry (TG-DSC) (Netzsch STA 409PC, Germany) under a flow of air at a rate of 10 °C/min. The phase of the calcined powders was analysed by X-ray diffraction (XRD, Rigaku, Japan). Scans were taken with a 2θ step of 0.02° (over the range of 20–80°) using Cu Kα radiation (18 kV, λ = 0.154056 nm). The microstructure was analysed with a scanning electron microscope (SEM, Model S4800). By using alternating-current impedance spectroscopy (Model CHI660D, China), the conductivity of the specimens was measured over the temperature range of 500 to 800 °C in air. Before measurements, Ag paste was painted onto either side of the discs and fired at 700 °C for 30 min to erase the solvent. Data were collected in the frequency range 0.1 Hz–100 kHz; registered impedance spectra were analysed using ZSimpWin software.

III. Results and discussion

3.1. TG-DSC analysis

Figure 2 shows TG-DSC plots of the dried Ce$_{0.8}$Gd$_{0.14}$Mg$_{0.06}$O$_{1.87}$ precursor gel measured in temperature range from 50 to 700 °C in air atmosphere. An initial weight loss of 30 wt.% below 300 °C was seen in Fig. 2 which correlated with the evaporation of the adsorbed water and the decomposition of organics such as citric acid. A dramatic weight loss of ∼30 wt.% can be observed in the temperature range 300–380 °C accompanied by an intense exothermic peak in the DSC curve, which can be attributed to the decomposition of organic groups and NO$_3$–. At the temperatures above 380 °C, there is no remarkable changes in TG curve. However, in DSC curve a wide exothermic peak (500–650 °C) can be detected, indicating that the phase transition and crystallization of the sample have happened. Thus, we selected 700 °C as the calcination temperature of the synthesized Ce$_{0.8}$Gd$_{0.2-x}$Mg$_x$O$_{1.9-\delta}$ powders, as it is sufficient to obtain the desired crystalline structure.

2.2. Characterization

The xerogels were studied by simultaneous thermogravimetric and differential scanning calorimetry (TG-DSC) (Netzsch STA 409PC, Germany) under a flow of air at a rate of 10 °C/min. The phase of the calcined powders was analysed by X-ray diffraction (XRD, Rigaku, Japan). Scans were taken with a 2θ step of 0.02° (over the range of 20–80°) using Cu Kα radiation (18 kV, λ = 0.154056 nm). The microstructure was analysed with a scanning electron microscope (SEM, Model S4800). By using alternating-current impedance spectroscopy (Model CHI660D, China), the conductivity of the specimens was measured over the temperature range of 500 to 800 °C in air. Before measurements, Ag paste was painted onto either side of the discs and fired at 700 °C for 30 min to erase the solvent. Data were collected in the frequency range 0.1 Hz–100 kHz; registered impedance spectra were analysed using ZSimpWin software.

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3.2. Phase structure

Figure 3 shows the XRD patterns of the Ce$_{0.8}$Gd$_{0.14}$Mg$_{0.06}$O$_{1.87}$ (x = 0.06) powders calcined at 500, 600 and 700 °C for 2 h. The typical diffraction peaks of cubic fluorite structure occurred

![Figure 1. Experimental flowchart of preparation of samples by the sol-gel method](image1)

![Figure 2. TG-DSC curves of Ce$_{0.8}$Gd$_{0.14}$Mg$_{0.06}$O$_{1.87}$ xerogel](image2)
at 500 °C, but the intensity of the diffraction peaks was weak, and some characteristic peaks were not observed. This indicates that the sample was not well crystallized at this temperature. When the calcination temperature increased to 700 °C, the appearance of the typical diffraction peaks indicated that the powders crystallized well and formed cubic fluorite structure with space group $Fm\overline{3}m$ of CeO$_2$ (PDF No. 34-0394). Thus, fine Ce$_{0.8}$Gd$_{0.14}$Mg$_{0.06}$O$_{1.87}$ powders with cubic fluorite-type structure were successfully prepared after calcined at 700 °C for 2 h, which was consistent with the TG-DSC results.

XRD patterns for the samples with different Mg-dopant concentrations are shown in Fig. 4. All powders calcined at 700 °C have a single-phase fluorite structure. It can be seen that no secondary phase could be identified. According to previous studies, doping with ions having different valences and different ionic radii will change the lattice parameter of ceria and induce corresponding oxygen vacancies generate the internal stress in the ceria matrix [21–23]. The lattice parameters of the prepared Ce$_{0.8}$Gd$_{0.2-x}$Mg$_x$O$_{1.9-\delta}$ samples are basically not changed considerably with the Mg-doping concentration, and remained $a = b = c = 0.5418$ nm. This observation could indicate that solubility of Mg$^{2+}$ in the GDC lattice is very low. The reason could be the difference in ionic radii of Mg$^{2+}$ (0.089 nm) and Ce$^{4+}$ (0.097 nm). Thus, the addition of Mg might cause a distortion of the lattice as well as the increase of lattice energy.

3.3. Microstructure analysis

Micrographs of the Ce$_{0.8}$Gd$_{0.14}$Mg$_{0.06}$O$_{1.87}$ ceramics sintered at 1300 °C for 4 h are shown in Fig. 5. As can be seen, the density of the sample sintered at 1300 °C is high and the grains are connected to each other with only few pores visible. As the addition of Mg$^{2+}$ increases, the change in density is not significant, the average grain size was in the range of 1–2 $\mu$m.

It was reported that ceria electrolytes with density above 95% of theoretical density were produced by sintering at 1200 or 1250 °C [20,24,25]. However, for densification of the Mg-doped GDC the sintering temperature of 1300 °C was necessary. The reason could be that Mg$^{2+}$ is not dissolved into GDC completely, but enriched partially the grain boundaries of the samples. The high melting point of the MgO phase ($\sim$ 2852 °C) and the low solubility of Mg$^{2+}$ [17], could give some support to this idea. However, phase structure analysis indicates that there is no formation of any new phase.

![Figure 3. XRD patterns of the Ce$_{0.8}$Gd$_{0.14}$Mg$_{0.06}$O$_{1.87}$ powders calcined at: a) 500, b) 600 and c) 700 °C](image-url)

![Figure 4. XRD patterns of Ce$_{0.8}$Gd$_{0.2-x}$Mg$_x$O$_{1.9-\delta}$ with different dopant concentrations](image-url)

![Figure 5. SEM images for Ce$_{0.8}$Gd$_{0.14}$Mg$_{0.06}$O$_{1.87}$ sintered at 1300 °C for 4 h in air: a) surface and b) fresh fracture](image-url)
This demonstrates that the content of undissolved MgO at the grain boundaries, if exists, is lower than the detection limits of X-ray diffraction.

3.4. Electrochemical performance

Electrochemical impedance spectroscopy (EIS) has been used to determine electrochemical properties of the solid electrolyte. From the impedance spectroscopy, it is possible to obtain the information on the contribution of grains, grain boundaries and electrode polarization to the total conductivity [25–27]. Impedance spectra for the Ce$_{0.8}$Gd$_{0.2-x}$Mg$_x$O$_{1.9-\delta}$ ceramics measured at different temperatures under air are presented in Figs. 6 and 7. The conductivities can be calculated from the resis-
ent temperatures can be obtained using ZSimpWin software. Then the conductivities at different temperatures obtained by fitting the impedance spectra using the equation:

\[ \sigma = \frac{L}{RS} \]  

(2)

where \( L \) is the sample thickness and \( S \) is the electrode area of the sample.

Conductivity and resistance values of the samples at different temperatures are listed in Table 1. A minor decrease in conductivity was observed in the samples with low Mg-concentration (x = 0.02 and 0.04). However, the increase in conductivity was observed when the dopant content increases to \( x = 0.06 \) reaching 0.0203 S/cm at 800 °C. The reason may be that with the increasing of Mg content, the oxygen vacancies are more mobile which results in the increased conductivity [28]. Incorporation of Mg into GDC lattice enhanced the oxygen ionic conductivity also through the increasing of the activation energy of Mg-doped GDC, This indicated the high-ness of Mg as a co-dopant for the optimization of electrical properties of Ce-based electrolytes. This indicated that \( \text{Ce}_{0.8} \text{Gd}_{0.2} \text{Mg}_{0.06} \text{O}_{1.9-\delta} \) is a promising electrolyte material for intermediate-temperature SOFC.

The total electrical conductivity of \( \text{Ce}_{0.8} \text{Gd}_{0.2} \text{Mg}_{x} \text{O}_{1.9-\delta} \) can be described in terms of the relationship of \( \ln(\sigma \cdot \cdot T) \) versus 1000/T, as plotted in Fig. 8 over the temperature range 500–800°C. As can be seen from Fig. 8, the relationship between conductivity and temperature of \( \text{Ce}_{0.8} \text{Gd}_{0.2} \text{Mg}_{x} \text{O}_{1.9-\delta} \) obeyed the Arrhenius relation:

\[ \sigma = A \exp \left( \frac{-E_a}{K \cdot T} \right) \]  

(3)

where \( \sigma \) is the conductivity of the sample, \( E_a \) is the conduction activation energy, \( K \) is Boltzmann constant and \( T \) is the absolute temperature. The conductivity activation energy of \( \text{Ce}_{0.8} \text{Gd}_{0.14} \text{Mg}_{0.06} \text{O}_{1.87} \) is calculated to be 0.853 eV.

Overall, analysis results suggest that Mg may be used as a co-dopant for the optimization of electrical properties of \( \text{Ce}_{0.8} \text{Gd}_{0.2} \text{O}_{1.9-\delta} \) based electrolytes. This indicated that \( \text{Ce}_{0.8} \text{Gd}_{0.2} \text{Mg}_{x} \text{O}_{1.9-\delta} \) is a possible electrolyte material for IT-SOFC.

IV. Conclusions

\( \text{Ce}_{0.8} \text{Gd}_{0.2} \text{Mg}_{x} \text{O}_{1.9-\delta} \) powders (x = 0, 0.02, 0.04 and 0.06) were synthesized by the sol-gel method and sin-

<table>
<thead>
<tr>
<th>Composition</th>
<th>Reference</th>
<th>Total ionic conductivity ([\text{S/cm}])</th>
<th>Conductivity activation energy ([\text{eV}])</th>
</tr>
</thead>
<tbody>
<tr>
<td>( \text{Ce}<em>{0.8} \text{Gd}</em>{0.18} \text{Mg}<em>{0.02} \text{O}</em>{1.9-\delta} )</td>
<td>Present work</td>
<td>0.0061</td>
<td>0.972</td>
</tr>
<tr>
<td>( \text{Ce}<em>{0.8} \text{Gd}</em>{0.14} \text{Mg}<em>{0.06} \text{O}</em>{1.9-\delta} )</td>
<td>Present work</td>
<td>0.0203</td>
<td>0.853</td>
</tr>
<tr>
<td>( \text{Ce}<em>{0.82} \text{Gd}</em>{0.14} \text{Er}<em>{0.08} \text{O}</em>{1.86} )</td>
<td>[8]</td>
<td>0.042</td>
<td>0.664</td>
</tr>
<tr>
<td>( \text{Ce}<em>{0.85} \text{Sr}</em>{0.05} \text{Gd}<em>{0.1} \text{O}</em>{2-\delta} )</td>
<td>[7]</td>
<td>0.0635</td>
<td>1.13</td>
</tr>
<tr>
<td>( \text{Ce}<em>{0.85} \text{Sm}</em>{0.15} \text{Sc}<em>{0.05} \text{O}</em>{2-\delta} )</td>
<td>[20]</td>
<td>0.0192</td>
<td>1.106</td>
</tr>
</tbody>
</table>

Table 1. Conductivity and correlative parameters of the samples at different temperatures

<table>
<thead>
<tr>
<th>( x )</th>
<th>( L [\text{cm}] )</th>
<th>( S [\text{cm}^2] )</th>
<th>( R_{tot} [\Omega] )</th>
<th>( \sigma [\text{S/cm}] )</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.02</td>
<td>0.118</td>
<td>1.09</td>
<td>188.11</td>
<td>67.43</td>
</tr>
<tr>
<td>0.04</td>
<td>0.11</td>
<td>1.13</td>
<td>261.7</td>
<td>12.78</td>
</tr>
<tr>
<td>0.06</td>
<td>0.114</td>
<td>1.11</td>
<td>86.41</td>
<td>5.05</td>
</tr>
</tbody>
</table>

Table 2. Conductivity and activation energy of co-doped CeO\(_2\) electrolytes at 800 °C
tered at 1300 °C. XRD analysis confirmed the single-phase fluorite structure can be formed at a relatively low calcination temperature. SEM observations confirmed that the electrolytes sintered at 1300 °C for 4 h have dense structure. Based on the EIS data, the co-doped electrolyte with 6 mol% MgO content has the highest conductivity, i.e., 0.0203 S/cm at 800 °C. It is believed that the oxygen vacancies are activated when GDC is doped with Mg$^{2+}$, causing the increase of conductivities of the Ce$_{0.85}$Gd$_{0.15}$Mg$_{0.05}$O$_{1.87}$ electrolyte in comparison to the Gd singly doped CeO$_2$ electrolytes. Thus, co-doping with Mg led to an improvement in conductivity confirming that Ce$_{0.98}$Gd$_{0.02}$Mg$_x$O$_{1.96}$ is a possible electrolyte material for IT-SOFC.

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