Yttria and Ceria Doped Zirconia Thin Films Grown by Pulsed Laser Deposition

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The Yttria stabilized Zirconia (YSZ) is a standard electrolyte for solid oxide fuel cells (SOFCs), which are potential candidates for next generation portable and mobile power sources. YSZ electrolyte thin films having a cubic single phase allow reducing the SOFC operating temperature without diminishing the electrochemical power density. Films of 8 mol% Yttria stabilized Zirconia (8YSZ) and films with addition of 4 weight% Ceria (8YSZ + 4CeO2) were grown by pulsed laser deposition (PLD) technique using 8YSZ and 8YSZ + 4CeO2 targets and a Nd-YAG laser (355 nm). Films have been deposited on Soda-Calcia-Silica glass and Si(100) substrates at room temperature. The morphology and structural characteristics of the samples have been studied by means of X-ray diffraction and scanning electron microscopy. Films of a cubic-YSZ single phase with thickness in the range of 1-3 µm were grown on different substrates.

Keywords: solid oxide fuel cells (SOFCs), Yttria-stabilized Zirconia (YSZ), pulse laser deposition (PLD), thin films

1. Introduction

Solid oxide fuel cells (SOFCs) have been studied intensively because they exhibit a clean and efficient power generation and do not require high purity fuel. SOFCs are typically composed of two porous electrodes with a dense electrolyte of a solid oxide ceramic material in between. The anode and cathode must be porous in order to allow gas transportation, whereas the electrolyte must avoid electronic charge and gas transportation across the cell. Other desired characteristics for the cell elements are chemical stability under operational conditions, high electronic conductivity for the electrodes and electrical interconnections, high ionic conductivity and almost zero electronic conductivity for the electrolyte, chemical and mechanical compatibility for all the cell components and gases, as well as low cost. Tetragonal Zirconia exhibits high toughness but its ionic conductivity is much lower than that of cubic Zirconia. Besides, it shows a degradation process in the presence of hydroxyl ions (OH–) at low temperature (lower than 400 °C), which is inconvenient in a fuel cell. The cubic Zirconia can be fully stabilized by the addition of soluble aliovalent oxides such as CaO, MgO or Y2O3. The doping introduces oxygen vacancies in the lattice, improving the ionic conductivity at elevated temperatures. Yttria-stabilized Zirconia (YSZ) in particular exhibits a remarkable thermal stability and degradation resistance. The ionic conductivity in a bulk material of the Yttria doped Zirconia (ZrO2;Y2O3) reaches a maximum for an Yttria content in the solid solution between 8 and 12 mol% (8YSZ-12YSZ)

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The YSZ is a standard electrolyte for SOFCs, which are potential candidates for next generation portable and mobile power sources. Standard YSZ electrolytes operate at high temperatures (800-1000 °C) due to their ionic conductivity properties which become efficient enough only at this temperature range. However, at such high temperatures thermal and mechanical degradation of the cell materials drastically limit its lifetime. For most applications it is necessary to decrease the operation temperature without decreasing its working performance.

Decreasing the operational temperature decreases the YSZ ionic conductivity, but this can be compensated by reducing the thickness of the electrolyte, and thus reducing ohmic losses. Thin and dense YSZ electrolytes can be fabricated by different techniques such as physical or chemical vapor deposition (PVD, CVD), plasma enhanced CVD (PECVD), sol-gel method, electrochemical vapor deposition, suspension or atmospheric plasma spraying, pulsed laser deposition (PLD), etc. PLD allows producing films with thickness from 1 nm up to one micron scale, whereas other techniques allow producing thicker films in the range of 100 nm to 20 µm. PLD is commonly used to produce thin films of a great variety of materials because the stoichiometry obtained is transferred from the target to the film. Moreover, this technique produces films with good adherence and allows growing crystalline thin films. The thickness and structure of the film can be modified changing the deposition parameters, such as: laser wavelength, laser fluence, deposition time, substrate material, substrate temperature, etc.
It has been observed that Ce-based electrolytes have high conductivity at temperatures lower than 800 °C\(^2\). It is of interest to investigate the effect of CeO\(_2\) addition to the 8YSZ, produced by PLD. In this work, thin films of cubic single phase of 8YSZ + 4 wt.% CeO\(_2\) grown on Si(100) substrate by means of PLD are presented for the first time, as far as the authors know. Also thin films of cubic single phase of 8YSZ have been produced by the same technique. The morphology, structure, thickness and porosity of the films were studied using X-ray diffraction (XRD) and scanning electron microscopy (SEM).

### 2. Experimental

Film samples of 8YSZ and 8YSZ + 4 wt.% CeO\(_2\) were produced by PLD on Soda-Calcia-Silica glass and polished Si(100) substrates with different deposition times using a pulsed Nd-YAG laser. The substrates were situated at 25 mm in front of the target. Table 1 summarizes the samples’ codes, compositions, substrates and processing conditions of each studied samples. The targets were prepared with the same composition of the films to be deposited. They were prepared from powders by ball milling, powder dry pressing and air sintering disk shapes of 25 mm in diameter and 1.5 mm thickness between 1450 °C and 1560 °C. The Nd-YAG laser used in this work (Spectra-Physics, Model Lab-190, USA) can be operated at 266 nm, 355 nm, 532 nm, and 1064 nm wavelengths. In this work pulses were produced at 355 nm wavelength with 8 ~ 12 ns width, an energy density of 3 J.cm\(^{-2}\) per pulse and a 30 Hz repetition rate using the Q-switching operating mode. The laser beam is reflected and guided by a set of mirrors and focusing lenses yielding a maximum laser energy density to ablate the target. The beam was oriented at 15 degrees with respect to the surface of the target. The films were grown in a chamber with air atmosphere with a dynamic vacuum of 133 mbar. At a higher pressure, no films grew, only powder deposited on the substrates was observed. The chamber is mounted on a Zaber’s T-series high precision PC controlled X-Y table that allows ablating a selected area on the target. The target was moved in X and Y directions having an irradiation time of 30 seconds for each position.

The morphology, microstructure and the thickness of the deposited films were analysed by secondary electron (SE) images of the surfaces and cross-sections of each sample using a Jeol 6510 LV SEM and a Philips XL30 SEM equipped with an energy dispersive X-ray (EDX) detector. The chemical compositions were checked by EDX analysis using accelerating voltages of 15 to 25 kV. Phase purity and crystallinity were studied by X-ray diffraction (XRD) in Bragg-Brentano geometry using a BRUKER D8 advanced diffractometer with monochromatic CuK\(_{\alpha}\) radiation (\(\lambda = 0.154060\) nm) and Lynx Eye one dimension detector.

### 3. Results and Discussion

Figure 1 shows the XRD results of 8YSZ (M1 and M2 samples) and 8YSZ + 4CeO\(_2\) (M3 and M4 samples) films deposited on glass and Si(100) substrates. These results show that cubic-YSZ polycrystalline single phase films were grown on both types of substrates. Diffractograms of the films and the targets are similar having both materials the same phase. The inset in the Figure 1 shows a detail of the peaks corresponding to the planes (531) and (600) from both 8YSZ + 4CeO\(_2\) targets. It is observed that the addition of CeO\(_2\) to 8YSZ produces peak broadening and shifts toward lower angles (see Figure 1 and inset).

The unit cell parameter determined for the 8YSZ cubic-phase grown on both types of substrates (samples M1 and M2) is 0.5319 nm, which is in agreement with the unit cell value given in the standards of the International database ICDD card Ner. 00-030-1468. No contribution from tetragonal-YSZ phase was identified neither in the film (see Figure 1) nor in the target (see inset of Figure 1). No significant texture in the films were observed, which means that no preferential epitaxial growth was induced on the Si(100) substrate.

The addition of 4 wt.% CeO\(_2\) to the 8YSZ produces peak broadening and shifts toward lower angles in the X-ray diffractogram of the samples M3 and M4 with respect to the X-ray diffractogram samples M1 and M2 in agreement with the observation in the targets (see inset Figure 1). These shifts can be explained by a Vegard law\(^{26,27}\) taking into account that the substitution of Zr by Ce increases the parameters of the unit cell due to the difference in the cationic radii (\(r_{\text{Ce}^{4+}} = 0.97\) Å and \(r_{\text{Zr}^{4+}} = 0.84\) Å in 8-fold coordination\(^{28}\)).

The classical Williams-Hall representation was used to separate the broadening of the XRD peaks induced by the crystallite size from that arising from the strains distribution. It was determined that all the films have very similar crystallite size of ~100 nm. Therefore, the largest peak broadening for the Ce doped samples respect to the undoped Ce samples is due to the larger strains distribution (7 times larger) related to the local strain generated by the difference in the cation size between Ce\(^{4+}\) and Zr\(^{4+}\).

Figure 2 shows EDX spectrum of the sample M2, 8YSZ film grown on Si (100). No contaminant elements were found in any of the samples, as shown in the EDX spectra. Additional peaks from the substrates were also observed.

Figure 3a, b shows SE images from cross-section of samples M3 and M4 respectively. Different layers that form the deposited films can be seen (Figure 3a). Porosity is also observed in some cross sections (see arrows in Figure 3a, b).

<table>
<thead>
<tr>
<th>Sample</th>
<th>Material</th>
<th>Substrate</th>
<th>Pressure (mbar)</th>
<th>Deposition time (minutes)</th>
</tr>
</thead>
<tbody>
<tr>
<td>M1</td>
<td>8YSZ</td>
<td>Soda-Calcia-Silica glass</td>
<td>133</td>
<td>383</td>
</tr>
<tr>
<td>M2</td>
<td>8YSZ</td>
<td>Polished Si(100)</td>
<td>133</td>
<td>121</td>
</tr>
<tr>
<td>M3</td>
<td>8YSZ + 4CeO(_2)</td>
<td>Polished Si(100)</td>
<td>133</td>
<td>275</td>
</tr>
<tr>
<td>M4</td>
<td>8YSZ + 4CeO(_2)</td>
<td>Polished Si(100)</td>
<td>133</td>
<td>150</td>
</tr>
</tbody>
</table>
Figure 1. X-ray diffractogram of 8YSZ and 8YSZ + 4CeO₂ films deposited on glass and Si(100) substrates, inset X-ray diffractograms of 8YSZ and 8YSZ + 4CeO₂ targets. In particular no peaks corresponding to tetragonal phase were found in the 2θ range at 43°, 52° and 53° in the films and from 123° to 132° in the targets (see inset). C: cubic-YSZ phase.

Figure 2. EDX spectrum of the sample M2. Accelerating voltage used was 15 kV.

Figure 3. SE images from cross-section of the samples M3 (a) and M4 (b). The arrows show the porosity in the samples.
Figure 4 shows the thickness of the film grown on Si(100) substrate at room temperature as a function of deposition time. The average value of the thickness of each film was measured on the cross section SE images, as it is shown in Figure 3b. The particulates on the film surface were not taken into account in the measurements. The error bars represent the standard deviation of thickness measured at different point along the cross section. Despite the fact that the ablation points were changed by moving the target with an interval of 30 seconds, the plume deflection effect due to the surface roughness produced by the ablation process led to slightly non-uniform thickness of the films. Thicknesses obtained were between 1 to 3 µm and are correlated with deposition time. The relationship between thickness and deposition time is slightly smaller than linear regression. This phenomenon has been observed previously and it is suggested to be correlated with a reduction in the amount of target material being vaporized by the laser beam. Krajnovich and Vazquez have observed that the thickness-increasing trend stops with prolonged deposition time when periodic structures such as ripples, ridges, and cones are formed on the target material.

Figure 5 shows SE images of the top view on the films surface of 8YSZ an 8YSZ + 4CeO$_2$ samples. Some micro-sized fragments or particulates as 1-5 µm spherical or quasi-spherical clusters are clearly visible on the surface of all the samples. The presence of the particulates is a characteristic of the PLD films. The sample M1 has irregular particulates, which are different to the ones deposited in the other samples. The gas pressure in the chamber limits the plasma...
plume expansion, which increases the probability of collisions between the ablated particles and thus leads to the nucleation and growth of clusters (particulates) that are deposited on the substrate during the film growth. Porosity is also observed in some of the films (M2, M3) produced on Si(100) substrate. It is expected a more compact film of Yttria and Ceria doped Zirconia cubic phase can be obtained at a lower pressure than 100 mbar.

4. Conclusion

Films of cubic phase of Yttria and Ceria doped Zirconia, 8YSZ and 8YSZ + 4 wt.% CeO₂, have been obtained by means of PLD technique using an Nd-YAG laser with 355 nm wavelength. The ablation process was produced on sintered targets at low pressure air atmosphere. The films grew without any texture and with the same cubic structure present in the targets. All the samples show crystallites of the cubic Zirconia phase with size of ~100 nm.

The cubic phase in the 8YSZ films has a unit cell parameter of 0.5319 nm. Whereas, the cubic phase in the 8YSZ + 4 wt.% CeO₂ films has larger unit cell and local strain distribution generated by the larger cation size of Ce⁺⁺ respect to the Zr⁺⁺.

The growth rate is smaller than a linear correlation between thickness and deposition time due to a decrease in the ablated material from the target during the ablation process. All films grown on Si(100) substrates show similar surface morphology with similar particulates size.

Yttria and Ceria doped Zirconia films can be produced with cubic phase without any texture on crystalline and amorphous substrates by means of PLD technique. Further work on the films properties are required in order to evaluate the performance to the Yttria and Ceria doped Zirconia cubic phase to be used as electrolyte in solid oxide fuel cells.

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