Epitaxial heterostructures consisting of an yttrium doped barium cerate (BCY) layer sandwiched between two yttrium doped barium zirconate (BZY) thin layers have been deposited on insulating (001) MgO substrates by pulsed laser deposition. The first BZY layer was aimed at improving the lattice match with the MgO substrate, the second at protecting the BCY layer. Ionic conductivity has been studied in the 300 – 600°C temperature range as a function of the BCY thickness. Due to the absence of blocking grain boundaries, heterostructures showed a conductivity larger than that of BCY pellets sintered under optimized conditions.

Introduction

Thin films have attracted considerable interest as electrolyte materials for micro solid oxide fuel cells (µ-SOFCs) operating at intermediate temperatures (600°–800°C) (1, 2). The decrease of the operating temperature has been pursued via the development of high temperature proton conducting electrolytes (HTPCs) (3-5). Several perovskite oxides, such as doped BaCeO₃, SrCeO₃, BaZrO₃, and SrZrO₃ show proton conductivity in hydrogen and water vapour containing atmosphere (6-8). Among these materials, BCY has the largest proton conductivity in the intermediate temperature range (about 10⁻²S cm⁻¹ at 600°C) (9). However, these compounds suffer from poor chemical stability due to degradation in the presence of acid gases (e.g., CO₂ and steam (10, 11). On the other hand, the yttrium doped BaZrO₃(BZY) family exhibits a good chemical stability, but their proton conductivity is relatively low (about 10⁻³S cm⁻¹ at 600°C) (12-15). Therefore, a variety of attempts have been addressed to find the best compromise between high proton conductivity and high chemical stability, such as, for instance, modifying processing parameters or coupling cerates and zirconates by solid solutions of the two compounds (12, 13, 16). These studies suggested that the average grain dimensions decrease with an increase of Zr content in the barium cerate oxides. As a consequence of the higher grain boundary density, a lower conductivity, relative to the pure barium cerate compound, has been observed.

Pulsed Laser Deposition (PLD) is a flexible technique suitable for depositing thin oxide films and complex heterostructures. It is gaining increasing attention for the
fabrication of SOFC materials, due to its capability of depositing high quality films of complex oxides (17,18). Recently, a number of studies have been carried out on textured BZY thin films grown by PLD (3,5,19). The excellent crystalline quality of BZY films identified them as being characterized by the largest conductivity measured for such material, because of the absence of blocking grain boundaries (4).

BCY is a distorted perovskite (i.e. orthorhombic structure with \(a=0.622\) nm, \(b=0.624\) nm and \(c=8.767\) nm, pseudo-cubic lattice parameter \(a=0.44\) nm) (20). Generally, in order to obtain epitaxial films, a good crystallographic match between film and substrate is required, both for the crystallographic structures and for the lattice parameters of the unit cells. BCY films have been grown on monocristalline (001) MgO and porous Al\(_2\)O\(_3\) (21). Such a substrate choice was mostly dictated by the need of having a negligible substrate contribution to the overall conductivity (both ionic and electronic). However, while in the case of Al\(_2\)O\(_3\) there is no match at all between film and substrate, the lattice match for MgO is also very poor. Namely, MgO has a rock salt cubic structure with a lattice parameter of 0.421 nm to be compared with the pseudo-perovskite structure of BCY with a lattice parameter of 0.44 nm. As a consequence of the large crystallographic structural differences between BCY and MgO, the conductivity values of the BCY films, directly deposited onto bare MgO, resulted to be two orders of magnitude lower with respect to the bulk values observed in previous studies (21, 22).

Aiming at improving both the structural quality (and thus the proton conductivity) and the chemical stability of BCY films, we deposited, by PLD, sandwich heterostructures consisting of a BaCe\(_{0.8}\)Y\(_{0.2}\)O\(_{3-x}\) layer, about 160, 480 and 800 nm thick, comprised between two BaZr\(_{0.8}\)Y\(_{0.2}\)O\(_{3-x}\) thin layers about 30 nm thick. BZY has the same crystallographic structure as BCY but a smaller pseudocubic lattice parameter (\(a=0.421\) nm) that matches the MgO lattice parameter very well. The first BZY layer was aimed at releasing the structural lattice mismatch between MgO and BCY, the top most BZY layer at protecting the BCY film from the atmosphere. The overall structural and electrochemical performance of MgO/BZY/BCY/BZY heterostructures were investigated in this paper.

**Experimental**

BCY and BZY targets for the PLD growth were prepared starting from Ba(NO\(_3\))\(_2\) (3% H\(_2\)O, Aldrich, 99.999%), Ce(NO\(_3\))\(_3\)-6H\(_2\)O (Aldrich, 99.9%), (CH\(_3\)CO\(_2\))\(_x\)Zr(OH)\(_y\) (Aldrich) and Y(NO\(_3\))\(_3\)-6H\(_2\)O (Aldrich, 99.9%). BCY and BZY dense targets were finally sintered in air at 1500°C for 5h and 1600°C for 8h, respectively.

The thin films of BCY with a 30 nm buffer layer and 30 nm cap layer of BZY were deposited by PLD on single crystalline MgO (001) substrates. The KrF excimer pulsed laser source (\(\lambda=248\) nm) was operated at 10 Hz, with an energy density of 3 J/cm\(^2\), the substrate temperature was about 700 °C and the O\(_2\) pressure used during the deposition was about 0.1 Pa. The PLD system was equipped with a multtarget carousel and a Reflection High Energy Electron Diffraction (RHEED) system for the in situ diagnostic of the growth. The BCY and BZY targets were mounted on the carousel and the whole heterostructure was grown in a single run without exposing individual layers to contamination from the atmosphere. In our experimental setup a deposition rate of about
0.22 Å per laser shot was obtained for both BCY and BZY. X-ray diffraction analysis (XRD) was used to determine the targets composition and the films lattice parameters. Cu Kα radiation was used. Rocking curves were measured in order to estimate the film’s mosaic spread. The surface morphology and the cross section of the deposited films were investigated by field-emission scanning electron microscopy (FE-SEM, LEO Supra 35).

Impedance measurements were carried out in wet 5% H2 in Ar (~3% vol. H2O) by using a frequency response analyzer, coupled with a dielectric interface. The temperature ranged between 300 and 630 °C and the frequency range was 5 MHz-0.01 Hz, with a 500 mV alternate applied voltage. An in-plane electrode configuration was used for the impedance characterization. Au paste electrodes with dimensions of 2 mm × 5 mm were painted onto the film surface and then fired at 600 °C.

To check the chemical stability of the BCY heterostructures, electrochemical measurements were repeated every day for 14 days.

**Results and Discussion**

The heterostructure growth mechanism was monitored by in-situ RHEED. Fig.1 (a) shows (from bottom to top) the RHEED patterns of the sample obtained depositing on an MgO substrate a BZY buffer layer and then a BCY layer. For the MgO substrate (500 µm thick), in agreement with the rock salt structure factor, only the (00), (02) and (02) diffraction streaks are visible. The BZY buffer layer RHEED pattern showed the typical feature of a perovskite structure, where all (0l) streaks are allowed. The presence of diffraction streaks in the RHEED pattern of both the BZY and BCY layers demonstrates that the growth occurs cube-on-cube in a quasi 2D mode. Dashed vertical lines are a guide for the eye and show that the position of (02) and (02) streaks for MgO and BZY coincide, as expected in view of the fact that the cubic lattice parameters of these compounds are very similar. In the case of the BCY pattern, it is possible to notice that the distance between (02) and (02) streaks is smaller than for BZY, as expected because of the larger BCY lattice constant. Fig.1 (b) shows a schematic sketch of the epitaxial relationship among the crystallographic cells of BCY, BZY, and MgO. The role of the BZY buffer layer is to accommodate the structural misfit (perovskite vs. rock salt) with MgO without need for any lattice misfit accommodation. The lattice misfit is exclusively accommodated by the BZY/BCY, purely perovskitic, interface. Such a two-step process led to an improved quality of the BCY layer, in terms of structural properties, relative to the direct MgO/BCY interface.

The XRD pattern (Figure 2) of the as grown BZY/BCY/BZY (with a BCY layer 800 nm thick nm) sandwich heterostructures on MgO substrates shows very intense (00l) peaks. No peaks from different phases or differently oriented grains can be detected in the spectra. In the inset of Figure 2 is reported a typical BCY rocking-curve around the (002) reflection. The full width at half maximum is about 0.8°. The in-plane lattice parameter was measured from the asymmetric (103) reflection. A value of 0.44nm for the in-plane pseudocubic lattice parameter was found, in agreement with the out-of-plane value. Such a value is in good agreement with the BCY bulk lattice parameter and demonstrates that the epitaxial strain in BCY is fully relaxed.
Figure 1. (a) BCY in situ RHEED pattern of MgO, BZY and BCY, from bottom to top, (b) schematic sketch of the structural relationship between BCY, BZY, and MgO.

Figure 2. Θ-2Θ X ray diffraction spectrum of a BZY/BCY/BZY heterostructure on a MgO (00l) substrate. Beside the intense (002) peak from the substrate only (00l) peaks from BCY and BZY can be observed. The inset shows the rocking curve of the (002) BCY reflection.
Figure 3 shows the FE-SEM cross section micrograph of the BZY/BCY/BZY 800 nm thick heterostructure where sharp interfaces between substrate and all film layers can be observed. The BCY layer fracture surface appears to be dense and uniform as typical epitaxial film growth. No grain boundary defects are detectable as it is expected for a cube-on-cube growth in a quasi 2D mode. The heterostructure surface is, except for some droplets barely visible on the top of the micrograph, quite smooth and homogeneous as typical for PLD deposition.

The ionic conductivity of the BCY sandwich heterostructures was investigated by Electrochemical Impedance Spectroscopy (EIS). Measurements were carried out in wet 5%H₂ in Ar atmosphere. It has to be noted that the MgO substrate resistance (≥ 10¹⁰Ω at 600 °C) is several orders of magnitude larger than the total resistance measured at the same temperature, which guarantees that the measured heterostructure resistance is dominated by the BCY film contribution (23). Figure 4(a) shows the complex impedance plots (Nyquist plots) at different temperatures of the 800 nm thick BCY heterostructure, shown as a representative example. A single semicircle with a negligible depression was observed at high frequencies at all temperatures in agreement with relevant literature on epitaxial thin films (24,2). The signal at low frequencies is attributable to electrode polarization. Fig.4(b) shows the spectroscopic plots of the impedance imaginary part (Bode plot). The bulk and the grain boundary contributions cannot be discriminated, Bode plots show in fact the presence of a single peak at high frequencies suggesting the presence of a single conduction mechanism.

Figure 5 shows the Arrhenius plots for BZY/BCY/BZY sandwich heterostructures of different thickness together with data relative to a BCY sintered pellet reported as reference (13). For all heterostructures, conductivity was slightly higher in the whole range of temperature investigated, reaching a maximum value of 1.1×10⁻² Scm⁻¹ at 630 °C for the heterostructure with the BCY layer 800 nm thick.
At variance with what previously observed for La$_{0.8}$Sr$_{0.2}$Ga$_{0.8}$Mg$_{0.2}$O$_3$ (LSGM) thin films deposited by PLD on the same substrate, conductivity was almost independent from film thickness. LSGM thin films were characterized by a highly textured columnar structure at the nanometer scale (26). Such columnar grains, coalesced during the growth so that the surface region was less defective than the interface region and correspondingly the conductivity increased with film thickness. Otherwise, the conductivity of BZY/BCY/BZY sandwich heterostructures is not affected by the blocking effect of columnar grains in agreement with the SEM observation. The activation energy value for the BCY heterostructures resulted to be very close to the value relative to sintered BCY pellets ($0.58 \pm 0.1\text{eV}$ versus $0.49 \pm 0.1\text{eV}$). The slightly improved electrochemical performance of the films can be attributed to the improved crystallographic quality.

Figure 4. Temperature dependent Nyquist plots (a) and $-\text{Im}(Z)$ versus frequency plots (Bode plot) (b), measured in wet 5%H$_2$ in Ar for a BZY/BCY/BZY heterostructure with a BCY layer 800nm thick.
Figure 5. Electrical conductivity in wet 5% H₂ in Ar atmosphere, as a function of the temperature for three BZY/BCY/BZY heterostructures with the BCY layer 160 nm, 400 nm and 800 nm thick, red stars, green triangles and blue squares respectively. The conductivity for the BCY sintered pellet (black circles) is also reported.

Figure 6. Nyquist plots measured over different days during the thermal stability test in operating conditions. The number next to each plot indicates the day of the test. The inset of the Fig. 6 shows the EIS thermal stability test as a function of time in hours. The black triangles represent the BCY sandwich heterostructure and the red circles represent the BCY sintered pellet.
The chemical stability of the BZY/BCY/BZY sandwich heterostructure (with a BCY 800 nm thick) in wet 5% H2-containing Ar atmosphere at 630 °C was investigated and compared to that of a BCY sintered pellet. Conductivity measurements were performed as a function of time. Figure 6 shows the Nyquist plots of the BZY/BCY/BZY sandwich heterostructure at 630 °C obtained at different days. The number next each plot indicates the day of testing. In the inset of Fig. 6 the conductivities of the heterostructure and of the pellet vs time are reported. The heterostructure conductivity remained almost unchanged for 12 days, only a slight decrease was revealed between the 12th day (R = 1.2 x10² S cm⁻¹) and the 13th day (R = 8.8x10⁻³ S cm⁻¹). For the BCY sintered pellet, a drastic decrease of the conductivity was observed on the 9th day (σ = 8.4×10⁻³ S cm⁻¹ on the 1st day and σ = 7×10⁻⁴ S cm⁻¹ on the 9th day). From the tenth day on it was not possible to measure conductivity because of the complete degradation of the pellet. Such a result showed that the capped BCY heterostructure has an improved chemical stability in a water-containing atmosphere in comparison to the BCY sintered pellet.

To investigate the decrease of the conductivity observed from 13th day of measurement in the BCY sandwich heterostructures, XRD analysis of the: (a) as-grown and (b) aged (after electrical measurements) BCY sandwich heterostructure was carried out and shown in Fig. 7. The as-grown BCY sandwich heterostructure shows (00l) peaks, exclusively while the heterostructure aged for 14 days in wet 5%H₂ in Ar at 630 °C, does not show any extra peaks such as attributable to the presence of either CeO₂ or Ba(OH)₂ phases (30). However, in the XRD pattern shown in Fig. 7 (b), it is possible to notice an intense (110) BCY peak, indicating a much stronger degree of polycrystallinity of the aged heterostructures in comparison to the as grown film. It can thus be concluded that the aging phenomenon of BCY sandwich heterostructure is not driven by chemical reactivity but by a microstructural reconstruction at high temperature (630 °C).
consequent increase in the degree of polycrystallinity negatively affects the transport properties because of the occurrence of the grain boundaries contribution in the polycrystalline microstructure.

Conclusions

Highly textured BZY/BCY/BZY heterostructures were grown by PLD on (001) MgO substrates. SEM and XRD investigations indicate a good crystalline quality as a result of the BZY thin buffer layer. The epitaxial growth of the films resulted in conductivity larger than that reported for BCY sintered pellets as a result of the negligible effect of perpendicular blocking grain boundaries. The heterostructure chemical stability issues have been investigated in a water-containing atmosphere at working temperatures (630 °C). An improved chemical stability of the BCY sandwich heterostructure in comparison to sintered BCY pellets was attributed to the epitaxial BZY cap layer. However, aging in wet 5%H2 in Ar caused a microstructural change from epitaxial to polycrystalline. The morphology variation led to the occurrence of grain boundaries which negatively affect the overall conductance.

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References