Design, Synthesis, Structure and Properties of Ba-Doped Derivatives of SrCo$_{0.95}$Ru$_{0.05}$O$_{3−δ}$ Perovskite as Cathode Materials for SOFCs

Sabina Sydyknazar $^1$, Vanessa Cascos $^{1,\dagger}$, Loreto Troncoso $^{1,2,*}$, Ana Laura Larralde $^3$, María Teresa Fernández-Díaz $^4$ and José Antonio Alonso $^1$

$^1$ Instituto de Ciencia de Materiales de Madrid, C.S.I.C., Cantoblanco, E-28049 Madrid, Spain; sabishas.es@gmail.com (S.S.); vcascos@icmm.csic.es (V.C.); ja.alonso@icmm.csic.es (J.A.A.)

$^2$ Instituto de Materiales y Procesos Termomecánicos, Universidad Austral de Chile, General Lagos, 2086, 5111187 Valdivia, Chile

$^3$ INQUIMAE, Departamento de Química Inorgánica, Analítica y Química Física, Facultad de Ciencias Exactas y Naturales, Universidad de Buenos Aires, Buenos Aires C1428EGA, Argentina; alarralde@qi.fcen.uba.ar

$^4$ Institut Laue Langevin, BP 156X, F-38042 Grenoble, France; ferndiaz@ill.eu

* Correspondence: loreto.troncoso@uach.cl

Received: 29 May 2019; Accepted: 13 June 2019; Published: 18 June 2019

Abstract: We have designed and prepared a novel cathode material for solid oxide fuel cell (SOFC) based on SrCo$_{0.95}$Ru$_{0.05}$O$_{3−δ}$ perovskite. We have partially replaced Sr by Ba in Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3−δ}$ (SBCRO) in order to expand the unit-cell size, thereby improving the ionic diffusion of O$^{2−}$ through the crystal lattice. The characterization of this new oxide has been studied at room temperature by X-ray diffraction (XRD) and neutron powder diffraction (NPD) experiments. At room temperature, SBCRO perovskite crystallizes in the P4/mmm tetragonal space group, as observed from NPD data. The maximum conductivity value of 18.6 S cm$^{-1}$ is observed at 850 $^\circ$C. Polarization resistance measurements on LSGM electrolyte demonstrate an improvement in conductivity with respect to the parent Sr-only perovskite cathode. A good chemical compatibility and an adequate thermal expansion coefficient make this oxide auspicious for using it as a cathode in SOFC.

Keywords: IT-SOFC; cathode; perovskite; neutron diffraction; SrCoO$_3$
compatibility issues. Most of the known MIEC materials belong to the ABO$_3$ perovskite family, where A and B can be partially or totally substituted leading to a wide range of catalytic activities, oxygen stoichiometries and interesting transport properties [5,6].

The high-temperature cubic phase of SrCoO$_{3-\delta}$ is a promising material due to its high electrical conductivity and oxygen permeation flux [7,8]. Depending on the synthesis conditions and the oxygen content, we can obtain different structural distortions such as orthorhombic, tetragonal, and cubic unit cells. Whereas the cubic phase exhibits an excellent ionic and electronic conductivity, the hexagonal phase has very poor properties and, unfortunately, it is thermodynamically stable, which limits the use of this pure material as a cathode in SOFC. As a strategy to prevent this cubic-to-hexagonal phase transition, it was found that replacing few percentages of Co by aliovalent elements like Re, Nb, Sb or Ti was effective to stabilize the cubic phase in a wide temperature range [9–13]. Recently, we observed that replacing 5% of Co by Ru prevents this unwanted transformation and enhances the catalytic oxygen reduction reaction [14]. In present research, we worked on the parent SrCo$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$ with an additional Ba doping at the Sr positions, in order to expand the unit-cell volume, so as to enhance the ion diffusion of O$^{2-}$ ions through the crystal lattice in this MIEC oxide. Our goal was to improve the electrochemical, electrical and catalytic performance of this compound in order to use it as a cathode in SOFC.

This work describes the full structural characterization of the novel material, including neutron powder diffraction data, in complement with transport, dilatometric, and polarization resistance measurements that confirm the excellent performance of this material as cathode in SOFCs.

2. Experimental

2.1. Synthesis

Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$ perovskite was obtained as a polycrystalline powder via soft chemistry, using the citrate-nitrate route. Stoichiometric amounts of Sr(NO$_3$)$_2$ (Strem Chemicals, Newburyport, MA, USA, 99%), Ba(NO$_3$)$_2$ (Merck, Kenilworth, NJ, USA, 99%) and Co(NO$_3$)$_2$·6H$_2$O (Alfa Aesar, Haverhill, MA, USA, 99%) were dissolved in a 10% saturated solution of citric acid. Afterwards, RuO$_2$ (Alfa Aesar, 99.9%) was added as a powder and kept in suspension until the solution was slowly evaporated and an organic resin was formed. The latter was dried at 120 °C and decomposed at 600 °C for 12 h. Finally, the solid was fired in air at 1000 °C for 24 h and used without further treatment.

2.2. Structural Characterization

The product was characterized by X-ray diffraction (XRD) for phase identification and to assess phase purity. The measurement was performed using a Bruker-AXS D8 diffractometer (Bruker, Billerica, MA, USA, 40 kV, 30 mA) in Bragg-Brentano reflection geometry with CuK$_{\alpha}$ radiation ($\lambda = 1.5418$ Å), in the 10–64° 2θ range. A nickel filter allowed the elimination of Cu K$_{\beta}$ radiation.

In order to characterize the oxygen vacancy distribution, a neutron powder diffraction (NPD) study was carried out. A high-resolution pattern was collected in the D2B diffractometer at the Laue-Langevin Institute (Grenoble, France) with a neutron wavelength $\lambda = 1.594$ Å within the angular range of 20 from 10 to 160°. About 2 g of the sample was contained in a cylindrical vanadium holder. A time of 2 h was required to collect a full diffraction pattern at 25 °C. The diffraction data were analyzed by the Rietveld method through the FULLPROF program (version 3.00) [15]. A pseudo-Voigt function was chosen to generate the line shape of the diffraction peaks. The scale factor, background points, zero shift, half-width, pseudo-Voigt corrected for asymmetry parameters, unit-cell parameters, positional coordinates and isotropic displacement factors for metals and anisotropic for oxygen atoms were refined in the final run. Occupancy factors for oxygen atoms were also refined from the collected NPD data.
2.3. Thermal Expansion Measurements and Chemical Compatibility

The mechanical compatibility between this material with the other cell components was determined by thermal expansion measurements in a cylindrical dense pellet (5 mm diameter × 2 mm thickness) obtained by pressing the prepared powder under 200 MPa of uniaxial pressure and sintered at 1050 °C for 12 h in air. The thermal expansion of the sintered sample was studied using a LINSEIS L75/155 °C dilatometer (Linseis Messgeraete, Selb, Germany) between 400 and 900 °C in air.

2.4. dc Measurements

The electrical conductivity was measured in the temperature range between 20 and 850 °C, using the dc four-probe technique in bar-shaped pellets under dc currents between 0.01 and 0.5 A. The currents were applied and collected using a Potentiostat-Galvanostat AUTOLAB PGSTAT 302 from ECO CHEMIE (NorECs Norwegian Electro Ceramics AS, Oslo, Norway).

2.5. ac Measurements

Electrochemical impedance spectroscopy (EIS) measurements were performed in air under open-circuit potential (OCP) conditions in a symmetrical configuration to extract the corresponding values of the electrolyte and electrode contributions. A dense electrolyte pellet of LSGM (La0.83Sr0.17Ga0.8Mg0.2O3−δ) was used and was prepared in our laboratory. An ink of the cathode material was created with the Sr0.9Ba0.1Co0.95Ru0.05O3−δ powders. First, the material was ball-milled in ethanol to break the agglomerates and then the dried powders were mixed with a binder (V-006 from Heraeus, Madrid, Spain). The ink was symmetrically painted over the electrolyte onto both surfaces in Sr0.9Ba0.1Co0.95Ru0.05O3−δ/LSGM/Sr0.9Ba0.1Co0.95Ru0.05O3−δ configuration. The cells were calcined at 1000 °C for 2 h to obtain a good adherence between the cathodes and the electrolytes according to a procedure previously optimized in our laboratories for doped-SrCoO3 cathode powders. Subsequently, two Pt electrodes were painted onto the cathode surfaces and calcined at 800 °C for 1 h to ensure equipotential conditions. EIS was then performed in potentiostatic mode, decreasing the temperature from 900 to 600 °C, with an excitation voltage of 50 mV in the range of 1 kHz to 100 MHz. All the cell impedances were normalized by the superficial area so that they have the units of Ω cm². The electrode endurance was evaluated under a constant cathodic current density of 200 mA cm⁻² at 850 and 800 °C for 20 h. The current passage was interrupted to collect the electrochemical impedance spectra in open circuit. The data obtained were analyzed with the Zview software (version 3.5) (Scribner Associates). The EIS spectra were collected using an AUTOLAB PGSTAT 302 (NorECs Norwegian Electro Ceramics AS, Oslo, Norway) from ECO CHEMIE.

3. Results and Discussion

3.1. Crystallographic Characterization

Sr0.9Ba0.1Co0.95Ru0.05O3−δ perovskite was obtained as a polycrystalline powder. The XRD diffractogram (Figure 1) collected with Cu Kα radiation evidenced the presence of the perovskite cubic phase. In a first approach, the crystal structure was refined on the basis of a simple cubic aristotype defined in the standard perovskite model at Pm-3m space group, with unit-cell parameter of a = 3.9159(5) Å. This is considerably expanded with respect to that corresponding to the parent compound of 3.858(1) Å [14], as expected for the much bigger size of Ba²⁺ (1.61 Å) with respect to Sr²⁺ (1.44 Å) ions in twelvefold coordination [16].
Figure 1. Observed (red crosses) and calculated profiles (full black line) for Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$ perovskite after Rietveld refinement of the crystal structure in a standard perovskite model in the Pm-3m space group, with unit-cell parameter of $a = 3.9159(5)$ Å. The difference plot (full blue line) and allowed Bragg reflections (green vertical lines) are also shown.

A room temperature study of Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$ by neutron powder diffraction (NPD) was very useful to investigate fine structural details. Although the XRD pattern can be interpreted in a cubic model, the NPD data suggest the presence of a tetragonal superstructure with a doubled $c$-axis, as $a = b \approx a_0$, $c \approx 2a_0$ and it was correctly refined in the P4/mmm space group. In this perovskite, the Sr and Ba atoms are distributed at random at $2h$ ($\frac{1}{2}$, $\frac{1}{2}$, $z$) sites, Co1 at $1a$ (0, 0, 0), Co2 and Ru at $1b$ (0, 0, $\frac{1}{2}$), and the three types of oxygen atoms O1 at $2f$ ($\frac{1}{2}$, 0, 0), O2 at $2g$ (0, 0, $z$) and O3 at $2e$ ($\frac{1}{2}$, 0, $\frac{1}{2}$) sites. The occupancy factors of the three types of oxygen atoms were refined. While O2 and O3 displayed conspicuous deficiency, O1 converged to values close to the stoichiometric value, so the occupancy factor was set to unity. Ru was unambiguously located at Co2 sites, thanks to the contrast between scattering lengths for Co (2.49 fm) vs Ru (7.03 fm). Figure 2 illustrates the goodness of the fit for the NPD pattern for the Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$ compound. Table 1 shows the final structural parameters, the displacement factors and the agreement factors after the Rietveld refinement from NPD data at room temperature.
Figure 2. Observed (crosses), calculated (full line) and difference (at the bottom) neutron powder diffraction (NPD) profile for $\text{Sr}_{0.9}\text{Ba}_{0.1}\text{Co}_{0.95}\text{Ru}_{0.05}\text{O}_{3-\delta}$ phase at 25 °C, refined in the tetragonal $\text{P4}/\text{mmm}$ space group. The vertical markers correspond to the allowed Bragg reflections.

Table 1. Structural parameters for $\text{SrCo}_{0.95}\text{Ru}_{0.05}\text{O}_{3-\delta}$ refined from neutron powder diffraction (NPD) data at 25 °C.

| Crystal Data | Refinement |
|--------------|------------|
| $a = 3.87158 (8) \text{ Å}$ | $R_p = 3.07\%$ |
| $c = 7.7928 (2) \text{ Å}$ | $R_{wp} = 3.98\%$ |
| $V = 116.81 (1) \text{ Å}^3$ | $R_{exp} = 2.59\%$ |
| $Z = 2$ | $R_{Bragg} = 8.04\%$ |
| $\chi^2 = 2.41$ | 3199 data points |

| Atoms | x | y | z | $U_{iso}/U_{eq}$ | Occupation (<1) |
|-------|---|---|---|-----------------|-----------------|
| $\text{Sr}$ | 0.50000 | 0.50000 | 0.2572 (4) | 0.0103 (9)* | 0.90000 |
| $\text{Ba}$ | 0.50000 | 0.50000 | 0.2572 (4) | 0.0103 (9)* | 0.10000 |
| $\text{Co}_1$ | 0.00000 | 0.00000 | 0.00000 | 0.005 (2)* | |
| $\text{Co}_2$ | 0.00000 | 0.00000 | 0.50000 | 0.020 (4)* | 0.95000 |
| $\text{Ru}_2$ | 0.00000 | 0.00000 | 0.50000 | 0.020 (4)* | 0.05000 |
| $\text{O}_1$ | 0.50000 | 0.00000 | 0.00000 | 0.0112 (15)* | |
| $\text{O}_2$ | 0.00000 | 0.00000 | 0.7696 (7) | 0.027 (3) | 0.921 (4) |
| $\text{O}_3$ | 0.50000 | 0.00000 | 0.50000 | 0.044 (4) | 0.824 (2) |

Anisotropic Displacement Parameters (Å²)

| Atoms | $U^{11}$ | $U^{22}$ | $U^{33}$ | $U^{12}$ | $U^{13}$ | $U^{23}$ |
|-------|---------|---------|---------|---------|---------|---------|
| $\text{O}_2$ | 0.038 (3) | 0.038 (3) | 0.007 (3) | 0.00000 | 0.00000 | 0.00000 |
| $\text{O}_3$ | 0.024 (3) | 0.073 (6) | 0.036 (4) | 0.00000 | 0.00000 | 0.00000 |

The tetragonal crystal structure (Figure 3) can be defined as a superstructure of perovskite with a doubled $a_0$ axis along the $c$ direction. This superstructure is due to the long-range ordering of the oxygen vacancies, quantified in the neutron refinement, and concentrated at the $\text{O}_2$ and $\text{O}_3$ positions. For this reason, the superstructure could not be detected from XRD data, given the weak scattering factor.
for oxygen atoms. The \( z(\text{O}_2) \) coordinate (Table 1) is slightly shifted from \( \frac{3}{4} \), determining alternating elongated \((\text{Co},\text{Ru})_2\text{O}_6\) octahedral layers, containing \( \text{Co}_2 \) atoms, and flattened \( \text{Co}_1\text{O}_6 \) octahedral layers containing \( \text{Co}_1 \) atoms. The \((\text{Co},\text{Ru})_2\text{O}_6\) octahedra share \( \text{O}_3 \) atoms, while \( \text{Co}_1\text{O}_6 \) octahedra share \( \text{O}_1 \) atoms. Finally, both octahedra layers share \( \text{O}_2 \) atoms. These differences in the octahedral size suggest that \( \text{Co}_1 \) cations might present a higher oxidation state, closer to \( 4^+ \), whereas \( \text{Co}_2 \) is close to \( 3^+ \), assuming a full charge disproportionation. In fact, the presence of \( \text{Co}^{3+} \) with an intermediate-spin state in \( \text{Co}_2 \) sites could cause the Jahn-Teller distortion of the Co–O bond lengths, and \( \text{O}_2^- \) ions would approach \( \text{Co}^{4+} \) cations, moving from \( \text{Co}_2 \) to \( \text{Co}_1 \), justifying the observed shift of \( \text{O}_2 \) atoms.

\[ \text{Figure 3. Tetragonal crystal structure observed for Sr}_{0.9}\text{Ba}_{0.1}\text{Co}_{0.95}\text{Ru}_{0.05}\text{O}_{3-\delta}. \]

3.2. Thermal Expansion Measurements

The thermal expansion coefficient (TEC) was measured by dilatometric analysis, in order to determine the mechanical compatibility of the material with the components of the cell. The analysis was carried out between 400 and 900 °C for several cycles and the data were collected only during the heating runs. Figure 4 shows the thermal expansion of \( \text{Sr}_{0.9}\text{Ba}_{0.1}\text{Co}_{0.95}\text{Ru}_{0.05}\text{O}_{3-\delta} \), which is approximately linear in the considered temperature range, and does not display discontinuities that could suggest decompositions. The TEC parameter, measured in air between 400 and 850 °C, is also shown. The determined value of TEC for the prepared oxide is substantially higher than those that normally have the components of the cell, but it is typical for cobaltites, which have been successfully used as cathodes in SOFC. In fact, the porous nature of the electrode once it is deposited on the electrolyte is able to accommodate the excessive dilation upon heating, avoiding cracking problems or delamination from the electrolyte.
3.3. Chemical Compatibility

The chemical compatibility of the Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3−δ}$ sample with the La$_{0.83}$Sr$_{0.17}$Ga$_{0.8}$Mg$_{0.2}$O$_{3−δ}$ (LSGM) electrolyte has been studied by mixing and grinding both powdered materials in the same quantities and heating the mixture at 1100 °C in air for 12 h. Figure 5 shows a Rietveld analysis of the product, consisting of an unaltered mixture of the starting Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3−δ}$ and LSGM oxides. Both are cubic perovskites with slightly different unit-cell parameters. No reaction between the materials was observed; tiny peaks at 2θ = 15°, 28° and 44° were attributed to a minor impurity phase below the 0.5% level.

Figure 4. Thermal expansion determined by dilatometry of the perovskite Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3−δ}$ collected in two consecutive heating runs.

Figure 5. Rietveld refinement from XRD data of a powdered mixture of the La$_{0.83}$Sr$_{0.17}$Ga$_{0.8}$Mg$_{0.2}$O$_{3−δ}$ (LSGM) electrolyte and the Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3−δ}$ electrode after a thermal treatment in air at 1100 °C, showing both phases unchanged.
3.4. Electrical Conductivity Measurements

3.4.1. dc Measurements

Figure 6 shows the thermal variation of the electrical conductivity of the Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$ perovskite, measured on a sintered bar (10 mm × 3 mm × 3 mm) in air by the four-probe technique. It is compared with that of the parent compound, SrCo$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$. In the low-temperature region, it shows semiconductor behavior with an anomaly at 450 °C, suggesting an insulator-to-metal transition. Above this temperature, a negative slope is observed until 720 °C. The maximum conductivity value at 450 °C is 28.5 S cm$^{-1}$; the conductivity at 850 °C is 18.6 S cm$^{-1}$. Although the electrical conductivity is lower than those presented by other Co-perovskites, with values between 30 and 50 S cm$^{-1}$, it should be sufficient for this material to be used satisfactorily as cathode material. The anomaly at 450 °C is also present in the parent compound (Figure 6, upper curve, taken from [14]), where it was related to a structural rearrangement from the room-temperature tetragonal superstructure to a truly simple cubic structure, where all the oxygen atoms become equivalent [14]. It is worth noting that the incorporation of Ba$^{2+}$ ions to the perovskite structure, expanding the unit-cell size, separate apart the Co–O atoms decreasing the orbital overlap, and for that reason the dc conductivity values decrease with respect to the value observed for the parent compound at 850 °C of 110 S cm$^{-1}$.

![Figure 6](image_url)

**Figure 6.** Thermal evolution of the dc electrical conductivity of Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$ compared with the parent compound Sr$_{0.9}$Co$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$.

3.4.2. ac Measurements

Electrochemical impedance spectroscopy (EIS) determinations seek to separate and identify the global electrode reactions into different contributions evidenced by distinct impedance arcs. The reaction steps are time-dependent and are represented in terms of frequency (s$^{-1}$). Each charge-transfer limiting process (i.e., electrolyte, electrode process, etc.) occurs at a distinct characteristic frequency, resulting in a separate arc of the impedance diagram [15,16]. The time-response of each individual process for the oxygen reduction reaction (ORR) is often modeled in terms of an equivalent circuit. The high frequency (HF) contribution is attributed to the oxygen-ion transfer process from the electrode to the electrolyte. A medium frequency (MF) signal is assigned to oxygen ion diffusion to the perovskite, measured on a sintered bar (10 mm × 3 mm × 3 mm) in air by the four-probe technique. It is compared with that of the parent compound, SrCo$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$. The polarization resistance was investigated by ac impedance spectroscopy in the temperature range of 600 to 900 °C. The Pt/Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$/LSGM/Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$/Pt symmetrical cell
was previously sintered at 1000 °C for 2 h. The data, collected in air, were normalized for the geometric area of the electrodes in the symmetrical cell and halved due to the symmetry of the cell.

The electrochemical characterization shows the same behavior in all the temperature range: (1) an inductive process, product of the electrochemical equipment and Pt wires, (2) an ohmic contribution, i.e., the intersection of the spectrum with the x-axis due to the internal resistance of the LSGM electrolyte, and (3) two semicircles, representing the polarization resistance of the cathode. These last features suggest the presence of two internal processes occurring at the triple phase boundary (TPB): the first one, at high frequencies, corresponds to the electrode/electrolyte interfacial charge transfer resistance, and the second one, at lower frequencies, associated with the oxygen reduction reaction (ORR).

Figure 7 displays the impedance diagrams of the cathode in the temperature range between 750 and 900 °C, showing that the resistance decreases as the temperature increases, as expected. All the spectra were fitted with the equivalent circuit indicated in the inset, using the non-linear least squares fitting of the Z-view program. In the scheme, L represents the inductive process, Re the electrolyte resistance, R1 the charge transfer resistance and R2 the ORR resistance. As mentioned, the sum of R1 and R2 corresponds to the polarization resistance Rp. The fact that the electrode can be polarized by the charge transfer and reduction reactions, implies the appearance of capacitive processes (“double layer” effect), which occurs in parallel to the resistive processes. This is represented in the circuit as a constant phase element (CPE), which represents a time-dependent capacitive element (pseudo-capacitance). The obtained Rp values are presented in Table 2. For comparative reasons, reported Rp values (studied previously by our group) for several SrCoO$_{3-\delta}$ perovskites are also included. They were all studied following the same guidelines in terms of preparation and sintering processes.

### Table 2. Polarization resistances and power densities (with H$_2$ as a fuel) of different cathodes. For the sake of comparison, values of undoped SrCoO$_{3-\delta}$ (2H phase) are also given. For the Power Densities, LSGM and SrMo$_{0.8}$Fe$_{0.2}$O$_{3-\delta}$ perovskites were used as electrolyte and anode material, respectively. For the 2H phase, no power density magnitude was available.

| Cathode (850 °C) | Rp (Ω cm$^2$) | Power Densities (mW cm$^{-2}$) |
|------------------|---------------|-----------------------------|
| Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_3$ | 0.048 | this work |
| SrCo$_{0.9}$Ru$_{0.1}$O$_{3-\delta}$ | 0.110 | 652 [14] |
| SrCo$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$ | 0.070 | 1100 [14] |
| SrCo$_{0.9}$Re$_{0.1}$O$_{3-\delta}$ | 0.066 | 660 [10] |
| SrCo$_{0.95}$Re$_{0.05}$O$_{3-\delta}$ | 0.080 | 570 [10] |
| SrCo$_{0.95}$Ti$_{0.05}$O$_{3-\delta}$ | 0.116 | 824 [13] |
| SrCo$_{0.97}$V$_{0.03}$O$_{3-\delta}$ | 0.025 | 550 [9] |
| SrCoO$_{3-\delta}$ (2H-phase) | 0.362 | - [8] |

Rp values obtained for Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$ are comparable to those previously reported. From the data in Table 2, we can see that not necessarily only a lower polarization resistance will imply a higher power density in a single test cell. In the case of the V$^{5+}$ doped sample, the Rp is only 0.025 Ω cm$^2$, and its power density is 550 mW cm$^{-2}$; instead, for the 10% Ru$^{4+}$ doped sample, the Rp is 0.110 Ω cm$^2$ and its power density is 652 mW cm$^{-2}$, both at 850 °C. This suggests that the doping cation must also have a functional catalytic activity for the ORR reaction. Comparing the results gathered in Table 2, it can be clearly seen that the one that outperforms under these conditions is the Ru$^{4+}$ cation with 5% of doping. The value of 0.048 Ω cm$^2$ determined for the present Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$ sample (vs. 0.070 Ω cm$^2$ for the parent compound) brings us to the conclusion that the introduction of Ba$^{2+}$ into the Sr site effectively decreases the polarization resistance, probably as an effect of the oxide-ion diffusion across the crystal lattice. This is auspicious regarding the performance of this material as cathode for SOFC in the intermediate temperature range.
Figure 7. Impedance spectra obtained at different temperatures in symmetrical cells of $\text{Sr}_{0.9}\text{Ba}_{0.1}\text{Co}_{0.95}\text{Ru}_{0.05}\text{O}_{3-x}$ onto LSGM electrolyte.

The activation energy ($E_a$) for $\text{Sr}_{0.9}\text{Ba}_{0.1}\text{Co}_{0.95}\text{Ru}_{0.05}\text{O}_{3-x}$ can be obtained from the Arrhenius plots of the resistance as a function of temperature (Figure 8). This value of activation energy, $E_a = 1.50 \text{ eV}$, is associated with the process of oxygen reduction reaction and is comparable to that observed for electrodes of the $\text{SrCo}_{1-x}\text{M}_x\text{O}_{3-\delta}$ family, also included in Figure 8. It is also noticeable that the presented Ba-doped electrode presents an improvement in the total conductivity with respect to the parent Ru-containing compound.

Figure 8. Arrhenius plot for $\text{Sr}_{0.9}\text{Ba}_{0.1}\text{Co}_{0.95}\text{Ru}_{0.05}\text{O}_{3}$ compound compared with similar cathodes.

The electrode endurance was studied in open circuit conditions and $dc$ current polarization. Figure 9 shows the symmetric cell impedance under OCV at 0 h compared with a constant current density of 200 mA cm$^{-2}$ after 20 h and 40 h at 850 °C and 800 °C. It is possible to observe that under the electrical flow of direct current, the $dc$ current flux produces an alteration in the electrode polarization.
resistance, increasing the electrolyte resistance and the polarization resistance, probably indicating an incipient reaction in the interface.

![Figure 9](image_url)

**Figure 9.** Impedance spectra of symmetrical cells under open circuit voltage (OCV) (0 h) and constant current density of 200 mA cm$^{-2}$ (20 h) at (a) 850 °C and (b) 800 °C.

3.5. Scanning Electron Microscopy

Figure 10 shows a SEM post-mortem study of the pellet utilized in the EIS tests. The micrograph shows the interphase of the Sr$_{0.9}$Ba$_{0.1}$Co$_{0.95}$Ru$_{0.05}$O$_3$ cathode and the electrolyte (LSGM).

![Figure 10](image_url)

**Figure 10.** SEM micrograph showing the porous cathode layer of Sr$_{0.9}$Ba$_{0.1}$Co$_{0.90}$Ru$_{0.10}$O$_3$ tightly adhered to the dense LSGM electrolyte. The thickness of the cathode is about 25 μm.

An important requirement for the most favorable cathode materials is the porosity. Figure 10 shows a good porosity in the surface of the cathode that favors the diffusion and reduction process of the oxygen throughout the bulk of the cathode. Also, the dense electrolyte layer without cracks or fractures is displayed.
4. Conclusions

In the present work, we have designed a novel cathode material by partially replacing Sr by Ba in SrCo$_{0.95}$Ru$_{0.05}$O$_{3-\delta}$ perovskite. We have shown with an NPD study that this compound is stabilized in a tetragonal perovskite structure at RT, defined in the P4/mmm group, with an expanded unit-cell volume with respect to the parent compound. The tetragonal superstructure is due to the long-range ordering of oxygen vacancies along the c-axis. The dc conductivity seems to be sufficient for a good performance as cathode material in SOFC. A good chemical compatibility was observed with the LSGM electrolyte for 12 h. The TEC of this material is higher than those usually presented in the other cell components, but it is characteristic of perovskites that contain Co. Interestingly, an EIS investigation unveils a lower polarization resistance than the parent compound, together with a reasonable total conductivity. All these properties make this material a good cathode candidate for solid oxide fuel cells performing at intermediate temperatures (IT-SOFC).

Author Contributions: Conceptualization, J.A.A.; Funding acquisition, J.A.A.; Investigation, J.A.A., S.S., V.C., L.T., A.L.L., M.T.F.-D.; Methodology, J.A.A., S.S., V.C., L.T., A.L.L., M.T.F.-D.; Project administration, J.A.A.; Resources, J.A.A.; Writing—original draft, J.A.A., S.S., V.C., L.T., A.L.L., M.T.F.-D.; Writing—review & editing, J.A.A., S.S., V.C., L.T., A.L.L., M.T.F.-D.

Funding: This research was funded by Spanish Ministry of Industry and Competitiveness, project number MAT2017-84496-R, Spain.

Acknowledgments: We thank the financial support of the Spanish Ministry of Industry and Competitiveness to the project MAT2017-84496-R. We thank the Institut Laue-Langevin (ILL) for making all facilities available.

Conflicts of Interest: The authors declare no conflict of interest.

References

1. Acres, G.J.K. Recent advances in fuel cell technology and its applications. J. Power Sources 2001, 100, 60–66. [CrossRef]
2. Shao, Z.; Zhou, W.; Zhu, Z. Advanced synthesis of materials for intermediate-temperature solid oxide fuel cells. Prog. Mater. Sci. 2012, 57, 804–874. [CrossRef]
3. Nicholas, J.D. Highlights from the 2013 National Science Foundation Solid Oxide Fuel Cell Promise, Progress, and Priorities (SOFC-PPP) Workshop. Interface Mag. 2016, 22, 49–54. [CrossRef]
4. Aguadero, A.; Fawcett, L.; Taub, S.; Woolley, R.; Wu, K.T.; Xu, N.; Kilner, J.A.; Skinner, S.J. Materials development for intermediate-temperature solid oxide electrochemical devices. J. Mater. Sci. 2012, 47, 3925–3948. [CrossRef]
5. Chen, D.; Chen, C.; Zhang, Z.; Baiyee, Z.M.; Ciucci, F.; Shao, Z. Compositional engineering of perovskite oxides for highly efficient oxygen reduction reactions. ACS Appl. Mater. Interfaces 2015, 7, 8562–8571. [CrossRef] [PubMed]
6. Zhou, W.; Jin, W.; Zhu, Z. Structural, electrical and electrochemical characterizations of SrNb$_{0.1}$Co$_{0.9}$O$_{3-\delta}$ as a cathode of solid oxide fuel cells operating below 600 °C. Int. J. Hydrogen Energy 2010, 35, 1356–1366. [CrossRef]
7. Deng, Z.Q.; Yang, W.S.; Liu, W.; Chen, C.S. Relationship between transport properties and phase transformations in mixed-conducting oxides. J. Solid State Chem. 2006, 179, 362–369. [CrossRef]
8. De la Calle, C.; Aguadero, A.; Alonso, J.A.; Fernández-Díaz, M.T. Correlation between reconstructive phase transitions and transport properties from SrCo$_{2.5}$ brownmillerite: A neutron diffraction study. Solid State Sci. 2008, 10, 1924–1935. [CrossRef]
9. Cascos, V.; Troncoso, L.; Alonso, J.A. New families of M$^{3+}$-doped SrCo$_{1-x}$M$_{x}$O$_{3-\delta}$ perovskites performing as cathodes in solid-oxide fuel cells. Int. J. Hydrogen Energy 2015, 40, 11333–11341. [CrossRef]
10. Troncoso, L.; Gardey, M.C.; Fernández-Díaz, M.T.; Alonso, J.A. New rhenium-doped SrCo$_{1-x}$Re$_{x}$O$_{3-\delta}$ Perovskites Performing as Cathodes in Solid Oxide Fuel Cells. Materials 2016, 9, 717. [CrossRef] [PubMed]
11. Cascos, V.; Alonso, J.A.; Fernández-Díaz, M.T. Nb$^{5+}$-doped SrCo$_{3-\delta}$ perovskites as potential cathodes for solid-oxide fuel cells. Materials 2016, 9, 579. [CrossRef] [PubMed]
12. Aguadero, A.; Alonso, J.A.; Pérez-Coll, D.; De La Calle, C.; Fernández-Díaz, M.T.; Goodenough, J.B. SrCo$_{0.95}$Sb$_{0.05}$O$_{3–δ}$ as cathode material for high power density solid oxide fuel cells. *Chem. Mater.* 2010, 22, 789–798. [CrossRef]

13. Cascos, V.; Fernández-Díaz, M.T.; Alonso, J.A. Structural and electrical characterization of the novel SrCo$_{1−x}$Ti$_x$O$_{3–δ}$ (x = 0.05, 0.1 and 0.15) perovskites: Evaluation as cathode materials in solid oxide fuel cells. *Renew. Energy* 2019, 133, 205–215. [CrossRef]

14. Cascos, V.; Troncoso, L.; Fernández-Díaz, M.T.; Alonso, J.A. SrCo$_{1−x}$Ru$_x$O$_{3–δ}$ (x = 0.05, 0.1, and 0.15) Perovskites As Outperforming Cathode Material in Intermediate-Temperature Solid Oxide Fuel Cells. *ACS Appl. Energy Mater.* 2018, 1, 4505–4513. [CrossRef]

15. Rodríguez-Carvajal, J. Recent advances in magnetic structure determination by neutron powder diffraction. *Phys. B Condens. Matter* 1993, 192, 55–69. [CrossRef]

16. Shannon, R.D. Revised Effective Ionic Radii and Systematic Studies of Interatomic Distances in Halides and Chalcogenides. *Acta. Cryst.* 1976, 32, 751–767. [CrossRef]

17. Adler, S.B. Factors Governing Oxygen Reduction in Solid Oxide Fuel Cell Cathodes. *Chem. Rev.* 2004, 104, 4791–4844. [CrossRef] [PubMed]