Enhanced Electrochemical Properties of Non-stoichiometric Layered Perovskites, $\text{Sm}_{1-x}\text{BaCo}_2\text{O}_5+d$, for IT-SOFC Cathodes

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In this study, electrochemical properties of layered perovskites having non-stoichiometric compositions ($\text{Sm}_{1-x}\text{BaCo}_2\text{O}_{5+d}$, $x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, \text{and } 0.15$) were analyzed for the direct application of cathode materials for Intermediate Temperature-operating Solid Oxide Fuel Cells (IT-SOFC). From the $\text{Sm}_{1-x}\text{BaCo}_2\text{O}_{5+d}$ oxide systems calcined at $1,100^\circ\text{C}$ for $8\text{~h}$, single phase ($\text{SmBaCo}_2\text{O}_{5+d}$, SBCO_1) was maintained only in the case of the $x = 0$ composition. In the compositions of $x = 0.05-0.10$, BaCo$_{2.6}$ was mixed with the pattern of SBCO. In addition, in the composition of $x = 0.15$, it was confirmed that BaCo$_{2.6}$ and CoO phases coexisted with SBCO. In the compositions of $\text{Sm}_{1-x}\text{BaCo}_2\text{O}_{5+d}$, the overall Area Specific Resistance (ASR) values decreased as the removal amount of Sm increased from $x = 0.05-0.10$; then, the values increased for compositions from $x = 0.15$. For example, the ASRs of SBCO_1, $\text{Sm}_{0.95}\text{BaCo}_2\text{O}_{5+d}$ (SBCO_0.95), $\text{Sm}_{0.90}\text{BaCo}_2\text{O}_{5+d}$ (SBCO_0.90), and $\text{Sm}_{0.85}\text{BaCo}_2\text{O}_{5+d}$ (SBCO_0.85) measured at $600^\circ\text{C}$ were 0.301, 0.147, 0.119, and 0.179 $\Omega\text{~cm}^2$, respectively. In particular, SBCO_0.90 was found to have an excellent ASR property of about $0.035\Omega\text{~cm}^2$ at $700^\circ\text{C}$. Typical properties of the metal–insulator transition (MIT) electrical conductivity were shown in all measured compositions. The temperature at which MIT occurred increased as the non-stoichiometric composition increased.

Keywords: layered perovskite, cathode, intermediate temperature-operating solid oxide fuel cell, non-stoichiometric composition, area specific resistance, electrical conductivity

HIGHLIGHTS

- $\text{Sm}_{1-x}\text{BaCo}_2\text{O}_{5+d}$ ($x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, \text{and } 0.15$) oxide systems having non-stoichiometric compositions were synthesized by solid state reaction.
- BaCo$_{2.6}$ phase was identified from the composition of $x = 0.05$ or more in $\text{Sm}_{1-x}\text{BaCo}_2\text{O}_{5+d}$; CoO phase was also found to exist for the composition of $x = 0.15$
• All composition showed higher electrical conductivity values than 100 S/cm in the temperature ranges of 500–750 °C.
• Area specific resistance (ASR) of Sm$_{1-x}$BaCo$_{2}$O$_{5+d}$ was 0.15 Ω · cm$^2$ or less at 650°C. SBCO_0.90 showed the lowest value of 0.119 Ω · cm$^2$ at 600 °C.

INTRODUCTION

As environmental problems such as global warming, caused by the indiscriminate use of fossil fuels since the beginning of the industrial revolution in the 19th century have recently emerged, interest in eco-friendly energy has been expanding (Wigley, 1998; Hoard, 2015). Accordingly, many countries around the world have been implementing nature-friendly policies and spurring green energy development. There are various nature-friendly energy generation methods, including wind, hydroelectric, and solar power. However, these power generation systems have disadvantages such as restrictions on installation locations and high initial cost. Therefore, there is need for development of other energy methods that can replace these systems and mitigate their problems (Rodman and Meentemeyer, 2006; Kosnik, 2008; Ingole and Rakhonde, 2015).

Fuel cells or fuel cell systems based on hydrogen energy have been emerging as alternative energy conversion methods to replace the energy generating from fossil fuel. Especially, among various fuel cell types, solid oxide fuel cells (SOFC) are the focus of attention recently. An SOFC is an energy converting device that produces water and generates electricity via the chemical reaction of hydrogen and oxygen (Minh, 2004).

SOFCs are known to show higher values of energy efficiency and power density than those of other fuel cells. SOFCs, which are made of ceramics, are composed of a dense electrolyte and two porous electrodes (cathode and anode). The SOFC structure is simpler than that of other types of fuel cell. In addition, due to their high exchange current density and the kinetics characteristics that occur under relatively high temperature operating conditions, SOFCs do not require any precious or expensive catalysts for operation.

While SOFCs have advantages of high efficiency at high temperature and a stable ceramic structure, their high-temperature operating conditions can rather act as a disadvantage, causing problems such as thermal degradation of ceramic materials, sealing issues, and rapid oxidation of metal materials used as interconnectors. To solve these problems, research on Intermediate Temperature-operating Solid Oxide Fuel Cells (IT-SOFC) that feature lowered operating temperature in the intermediate temperature range (500–750°C) has been actively conducted. However, the main obstacle to this lowered-temperature-range SOFCs is the decreased electrochemical activity of the two electrodes (cathode and anode), as well as lowered ionic conductivity and increased ohmic resistance of the electrolyte. Significantly, the major source of voltage loss in IT-SOFCs is the cathode materials. Therefore, to achieve high power densities at the reduced temperatures, it is necessary to develop cathode materials that show excellent electrochemical reaction and enhanced/advanced low polarization resistance (Minh, 1993; Tu et al., 1997).

Recently, to replace existing IT-SOFC cathode materials, many research groups have investigated new concept-based cathode materials with a chemical composition of layered perovskite, because this structure yields superior oxide-ionic diffusivity, excellent oxygen surface exchange coefficients, high oxygen transport properties, and higher electronic conductivity (Kim et al., 2010; Joung et al., 2013; Zhang et al., 2016). In this new perovskite structure ($A^{'/}\!/B^{'/}_{2}$O$_{5+d}$ chemical composition), various lanthanide atoms such as La, Pr, Nd, and Sm can be substituted into the $A'$-site; Ba can be replaced in the $A'$-/site; and transition metal ions of Co, Mn, and Fe can be located at the B-site.

The first application of a layered perovskite structure as cathode of SOFC (Zhou et al., 1993) showed that a difference in coordination number occurs due to the difference in ion radius between the $A'$ and $A'$-/ions; thus, oxygen vacancies are formed, yielding excellent electrochemical properties (Akahoshi and Ueda, 2001; Frontera et al., 2003; Kim et al., 2007, 2010; Kim and Manthiram, 2015). The layered perovskite structure into which elemental Co was substituted on the B-site is comprised of a stacked structure with [CoO$_2$]-[BaO]-[CoO$_2$]-[LnO$_6$] on the c-axis.

In the [LnO$_6$] layer, there is a large difference between the ionic radius of Ln and the ionic radius of Ba$^{2+}$, resulting in a pyramid structure of CO$_2$ and an octahedral structure of CO$_6$. Finally, oxygen vacancies are generated in the [LnO$_6$] layer due to this structural feature. Accordingly, structure is known to exhibit higher electrochemical properties because it has excellent oxygen mobility properties and at the same time shows excellent surface diffusion properties. For example, according to Ding et al., who investigated the composition of PrBa$_{0.5}$Sr$_{0.5}$Co$_2$O$_{5+d}$, this composition exhibited remarkable properties in the 500–700°C temperature range and showed an area specific resistance (ASR) value of 0.23 Ω · cm$^2$ at 650°C (Ding and Xue, 2010). In addition, the SmBaCo$_2$O$_{5+d}$ layered perovskite had an ASR of 0.13 Ω · cm$^2$ at 650°C (Kim et al., 2009). However, the thermal shock and long-term stability of the Co substituted layered perovskite at high temperatures have been questioned due to the high thermal expansion coefficient (TEC) (Tietz, 1999; Pelosiato et al., 2015).

Strategies to develop not only excellent cathode materials for IT-SOFCs by substituting various materials into layered perovskite, but also non-stoichiometric cathode materials synthesized by removing individual elements of layered perovskite are being pursued (Kostogloudis and Fikos, 1999; Wang et al., 2014; Sun et al., 2016; Yi et al., 2016). Based on the composition PrBaCo$_2$O$_{5+d}$, PrBa$_{0.92}$Co$_2$O$_{5+d}$, in which 0.08 mol% of Ba was removed from the $A'$-/site, showed better electrochemical properties than the composition PrBaCo$_2$O$_{5+d}$. In particular, PrBa$_{0.92}$CoO had an excellent ASR value of 0.166 Ω · cm$^2$ at 600°C (Idrees et al., 2019). It can be seen from this result that the electrochemical properties were enhanced in the non-stoichiometric composition in which the $A'$-/site is removed stepwise from the composition of layered perovskite LnBaCo$_2$O$_{5+d}$. 

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**Electrochemical Properties of Sm$_{1-x}$BaCo$_{2}$O$_{5+d}$**

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**Area specific resistance (ASR)** of Sm$_{1-x}$BaCo$_{2}$O$_{5+d}$ was 0.15 Ω · cm$^2$ or less at 650°C. SBCO_0.90 showed the lowest value of 0.119 Ω · cm$^2$ at 600 °C.
In the case of the composition of PrBaCo$_2$O$_{5+d}$ with partially removed A$^\prime$-site, not A$^\prime$/-site, the concentration of oxygen vacancies inside the crystal structure increased as the Pr-deficiency level increased. Especially, the Pr$_{0.95}$BaCo$_2$O$_{5+d}$ composition showed ASR values of 0.113, 0.054, and 0.028 $\Omega$·cm$^{-2}$ at 600, 650, and 700$^\circ$C (Jiang et al., 2014). As another example, the oxygen content of PrBaCo$_2$O$_{5+d}$ was 5.74 and the value of Pr$_{0.92}$BaCo$_2$O$_{5+d}$ was reduced to 5.62 when the amount of Pr removed increased. In addition, Pr$_{0.92}$BaCo$_2$O$_{5+d}$ was found to have excellent ASR values of 0.081 and 0.041 $\Omega$·cm$^{-2}$ at 700 and 750$^\circ$C (Zhang et al., 2016). Therefore, in the case of the non-stoichiometric composition of the layered perovskite, excellent electrochemical properties can be obtained when the lanthanide element is controlled. In summary, oxygen vacancies in the lattice of layered perovskite can be activated through non-stoichiometry of the A$^\prime$-site, thereby promoting the ORR reaction.

The goal of this research is to investigate the phase synthesis characteristics of Sm$_{1-x}$BaCo$_2$O$_{5+d}$ (x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, and 0.15) showing non-stoichiometric compositions in which samarium (Sm) elements were stepwise removed at the A$^\prime$-site of a SmBaCo$_2$O$_{5+d}$ layered perovskite by using solid state reaction. Significantly, the electrochemical properties of Sm$_{1-x}$BaCo$_2$O$_{5+d}$ oxide systems were also investigated to apply this material as cathode material for direct application to IT-SOFCs.

**EXPERIMENTAL SECTION**

**Synthesis**

Sm$_{1-x}$BaCo$_2$O$_{5+d}$ (x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, and 0.15) oxide systems having layered perovskite structure were synthesized using Solid State Reaction (SSR). The starting materials were Sm$_2$O$_3$ (samarium oxide, Alfa Aesar, 99.9%), BaCO$_3$ (barium carbonate, Samchun, 99.0%), and Co$_3$O$_4$ (cobalt oxide, High Purity Chemicals, 99.9%); an electronic balance (WBA-320) was used for accurate weighing to the third decimal place.

In this study, two calcinations were performed for more accurate synthesis of single phase. The weighed materials were mixed and calcined for 6 h at 1,000$^\circ$C in air. After the first calcination into uniform and small particles, wet ball milling was performed at room temperature for about 1 day to make the composite powder. The powder after ball milling was again calcined at a rate of 5$^\circ$C/min in air, and secondary calcination was performed at 1,100$^\circ$C for 8 h. Resulting materials after the two calcinations were pulverized to form fine powders. The compositions and abbreviations are summarized in Table 1.

**Sample Preparation**

Ce$_{0.9}$Gd$_{0.1}$O$_{2-d}$ (CGO91, Solvay) powders were used as the electrolyte of symmetrical half cells (Sm$_{1-x}$BaCo$_2$O$_{5+d}$/CGO91/Sm$_{1-x}$BaCo$_2$O$_{5+d}$). After weighing 2.5 g of CGO91, it was put into a metal mold and compression molding was performed by applying a force of 2 × 10$^3$ kg/m$^2$. After that, sintering was performed at 1,450$^\circ$C for 6 h in air. Cathode ink made using a-terpieal(Kanto Chemical), Butvar(Sigma Aldrich), and acetone with cathode powder; composite ink was also prepared by mixing electrolyte powder and cathode material at a weight ratio of 50:50. Both sides of the sintered electrolyte support were printed with the prepared cathode ink, and the printed cell was sintered at 1,000$^\circ$C for 1 h in the air condition. After the sintering, surface area of both sides of cathode was measured and found to be 0.785 cm$^2$. To confirm the electrical conductivity characteristics of the composites as identified through XRD analysis, a sample was prepared and the electrical conductivity was measured. After making a bar-type sample with dimensions of ~2.8 × 5.5 × 23 mm through compression molding, sample was sintered at 1,100$^\circ$C for 3 h in air.

**Characterizations**

To understand the crystal structure of the cathode powder synthesized by SSR method, X-ray diffraction analysis was used. The experimental equipment was an X-ray Diffractometer (Model: SmartLab/Rigaku), operated using a Cu K$_\alpha$ filter under conditions of 9 kW, 45 kV, and 200 mA. A range of 10–140$^\circ$ (2$\theta$) was employed with steps of 0.02$^\circ$. Microstructural properties of Sm$_{1-x}$BaCo$_2$O$_{5+d}$ (x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, and 0.15) oxide systems were examined using a field-emission scanning electron microscope (FESEM/Hitachi SU5000). To measure the electrical conductivity, a bar-type sample was prepared and the electrical conductivity was measured through the DC 4-probe method. A Keithley 2400 Source meter was used as measuring device; after connecting the sample with Pt-wire, the electrical conductivity was measured by increasing the temperature in steps of 50$^\circ$C in a temperature range of 50–900$^\circ$C. Additionally, the current range was measured from 0 to 1A through steps of 0.05A, and the limit of voltage was set to 20V. Impedance measurement was performed to confirm the electrochemical properties of the synthesized Sm$_{1-x}$BaCo$_2$O$_{5+d}$ (x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, and 0.15) oxide systems. According to existing literature, SmBaCo$_2$O$_{5+d}$ does not react with Ce$_{0.9}$Gd$_{0.1}$O$_{2-d}$ (CGO91), so there is no problem to use it with CGO91 (Kim et al., 2009). Therefore, the symmetrical half cell (SBCO$_X$/CGO91/SBCO$_X$) after the sintering was connected using Pt mesh, and the impedance was measured in a temperature range of 500–750$^\circ$C in an air atmosphere. Frequency range was 0.05 Hz to 2.5 MHz.

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**Table 1**

| Composition | Abbreviation |
|-------------|--------------|
| SmBaCo$_2$O$_{5+d}$ | SBCO$_1$ |
| Sm$_{0.99}$BaCo$_2$O$_{5+d}$ | SBCO$_{0.99}$ |
| Sm$_{0.98}$BaCo$_2$O$_{5+d}$ | SBCO$_{0.98}$ |
| Sm$_{0.97}$BaCo$_2$O$_{5+d}$ | SBCO$_{0.97}$ |
| Sm$_{0.96}$BaCo$_2$O$_{5+d}$ | SBCO$_{0.96}$ |
| Sm$_{0.95}$BaCo$_2$O$_{5+d}$ | SBCO$_{0.95}$ |
| Sm$_{0.90}$BaCo$_2$O$_{5+d}$ | SBCO$_{0.90}$ |
| Sm$_{0.85}$BaCo$_2$O$_{5+d}$ | SBCO$_{0.85}$ |
The area specific resistance (ASR) of the synthesized composition was calculated using the impedance results obtained through electrochemical analysis. Using the impedance results with ohmic resistance removed, ASR was calculated according to the difference between the low intercept and high intercept of the real X-axis. Impedance was measured using a multi-channel electrochemical analyzer (Model nStat, HS Technologies).

RESULTS AND DISCUSSION

Figure 1 summarizes the XRD results of Sm$_{1-x}$BaCoO$_{5+d}$ ($x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, \text{and} 0.15$) calcined at 1,100°C for 8 h to investigate the synthesis characteristics of these oxide systems with respect to the non-stoichiometric compositions. In addition, indicators for BaCoO$_{2.6}$ phase (▼) and CoO (▼) were added to all XRD results. The characteristic XRD patterns discovered in the vicinity of 23.4°, 33.0°, 40.4°, 46.7°, 59.1°, 69.2°, 78.1°) indicate that the Sm$_{1-x}$BaCoO$_{5+d}$ oxide systems were synthesized as a single phase from the stoichiometric SmBaCoO$_{5+d}$ (SBCO) composition in which Sm was not removed and the non-stoichiometric composition in which 0.01, 0.02, 0.03, and 0.04 mol% of Sm (SBCO$_{0.99}$, SBCO$_{0.98}$, SBCO$_{0.97}$, and SBCO$_{0.96}$) were partially removed. However, when Sm was removed at 0.05 mol% or more, BaCoO$_{2.6}$ phase appeared (SBCO$_{0.95}$, SBCO$_{0.90}$). Especially, when Sm was removed 0.15 mol%, CoO phase was additionally identified (SBCO$_{0.85}$).

Supplementary Figure 1 summarizes the SEM images of the Sm$_{1-x}$BaCoO$_{5+d}$ ($x = 0, 0.01, 0.05,$ and 0.10) oxide systems. Overall particle size decreases from (A) to (C), which results in porous microstructures as the non-stoichiometry composition increases. Especially, the particular microstructural characteristics can be seen in Supplementary Figures 1C,D. For example, in Supplementary Figure 1D, in which a part of SBCO$_{0.95}$ is magnified, white crystals can be seen protruding like arrowheads on the cathode surface. Particle size of arrowheads is very small, so it was impossible to carry out EDS (Energy-Dispersive x-ray Spectroscopy) analysis. However, as mentioned in the XRD results, it is known that the white arrowheads are formed by crystal structure of BaCoO$_{2.6}$. In other words, it can be thought that a small amount of secondary phase particles was generated on the cathode surface. For the SBCO$_{0.90}$ composition, which further increased the non-stoichiometry, as can be seen in Supplementary Figures 1E,F, particles like pebbles can be seen attached to the cathode surface. It is found that the particle size increased as the concentration of BaCoO$_{2.6}$ increased via the increase of the non-stoichiometry.

Figures 2A,B show that the overall ASR value decreases as the tendency toward non-stoichiometric composition increases (= the removal amount of Sm increases). For example, the ASRs of SBCO$_{1}$, SBCO$_{0.95}$, and SBCO$_{0.90}$ were about 0.301, 0.147, and 0.119 Ω · cm$^2$ at 600°C. These ASR values are closely related to the composition because the lowest ASR value was observed for the SBCO$_{0.90}$ composition. However, in the composition of SBCO$_{0.85}$ with an increased non-stoichiometric condition, the ASR value measured at 600°C was 0.179 Ω · cm$^2$, which is relatively higher than the ASR values of SBCO$_{0.95}$ and SBCO$_{0.90}$ measured at 600°C. That is, from the non-stoichiometric compositions of Sm$_{1-x}$BaCoO$_{5+d}$, the characteristics of the percolation composition showing the lowest ASR value in the chemical composition of Sm$_{0.98}$BaCoO$_{5+d}$ can be found. This tendency can also be confirmed in the layered perovskite composition of LnBaCoO$_{5+d}$ in which Pr was substituted into the A$^1$-site of the layered perovskite (Zhang et al., 2016; Idrees et al., 2019).

For example, according to the literature on ASR characteristics caused by non-stoichiometric composition of Pr$_{1-x}$BaCoO$_{5+d}$ ($x = 0–0.1$), PrBaCoO$_{5+d}$ (PBCO) and P$_{0.92}$BaCoO$_{5+d}$ (PBCO$_{0.92}$) exhibit ASR values of about 0.097 and 0.081 Ω · cm$^2$ at 700°C (Zhang et al., 2016). In the case of electrochemical properties related to the non-stoichiometric composition that reduce the amount of Pr in Pr$_{1-x}$BaCoO$_{5+d}$ oxide systems, when the degree of non-stoichiometry of the A$^1$-site increased, the ASR tended to decrease; at the same time, the percolation composition was also exhibited.

As the degree of non-stoichiometry of the A$^1$-site increases in the Sm$_{1-x}$BaCoO$_{5+d}$ oxide systems, the value of the activation energy tends to decrease. This trend can also be confirmed in works in the literature that have studied the electrochemical properties of similar layered perovskite structure composition in which the A$^1$-site is partially removed (Jiang et al., 2014; Yi et al., 2016). As a result of calculating the activation energy values, as shown in Figure 2A, Sm$_{0.98}$BaCoO$_{5+d}$ and Sm$_{0.96}$BaCoO$_{5+d}$ compositions have slightly increased activation energy values, but overall activation energy showed a tendency to decrease. For example, the activation energy of Sm$_{0.98}$BaCoO$_{5+d}$ was 1.094 eV. As degree of non-stoichiometry of A$^1$-site increases, the activation energy decreases. The values of Sm$_{0.92}$BaCoO$_{5+d}$ and Sm$_{0.90}$BaCoO$_{5+d}$ were calculated and found to be 0.970 and 0.948 eV. In addition, the value (0.987 eV) of Sm$_{0.88}$BaCoO$_{5+d}$ increased slightly compared to that of Sm$_{0.90}$BaCoO$_{5+d}$. When comparing the activation values of the Sm$_{1-x}$BaCoO$_{5+d}$ oxide systems synthesized in this experiment with those of recently reported cathode materials showing excellent electrochemical reaction, it can be determined that Sm$_{1-x}$BaCoO$_{5+d}$ oxide systems have relatively lower activation energy (Clematis et al., 2019; Anbo et al., 2020). In general, the activation energy indicates the strength of the energy barrier that must be overcome when oxygen ions move through the oxygen vacancies (Kaur and Singh, 2020).

For example, as mentioned above, Jiang et al. reported on the compositions of PBCO and P$_{0.95}$BCO; these compositions showed activation energy values of 1.19 and 0.98 eV (Jiang et al., 2014). This activation energy phenomenon was caused by the increase of concentration of pathways through which oxygen ions could move because the deficiency of Pr in A$^1$-sites resulted in many more oxygen vacancies in the PrO layer. The activation energy value was 0.948 eV, excellent even when compared with those of other compositions of layered perovskite (Jiang et al., 2014; Yi et al., 2016). In addition, as shown in Figure 2B, Sm$_{0.90}$BaCoO$_{5+d}$ had the lowest ASR values in this experiment, showing values of 0.119, 0.061, and 0.035 Ω · cm$^2$ at 600, 650, and 750°C, respectively.
FIGURE 1 | X-ray diffraction (XRD) patterns of Sm$_{1-x}$BaCo$_2$O$_{5+d}$ (x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, and 0.15) oxide systems calcined at 1,100°C for 8 h.

The results of an equivalent circuit of SBCO-0.90 are given in Table 2 and Figure 3 to confirm the rate determining step (RDS). In Table 2, the impedance results measured at 600°C using the Sm$_{1-x}$BaCo$_2$O$_{5+d}$ oxide systems were divided into R$_1$, R$_2$, and R$_3$.

In general, the impedance results of SOFC cathode materials can be classified by the frequency range (Murray et al., 1998). In equivalent circuits, the resistance from the high frequency (HF, 10$^3$ Hz–) is directly related with the charge transfer (R$_1$) when oxygen ions move across the interface between the cathode and the electrolyte. The resistance from the middle frequency (MF, 10$^2$–1 Hz) range is attributed to the resistance (R$_2$) that occurs when oxygen ions move inside the bulk of the cathode. The lower frequency (LF, below 1 Hz) arc is ascribed to oxygen dissociation and bulk or surface oxygen diffusion process (R$_3$). Therefore, ASR results were analyzed by subdividing them into R$_1$, R$_2$, and R$_3$.

In Figures 3A–C, the results for an equivalent circuit consisting of R$_1$, R$_2$, and R$_3$ and the impedance results measured directly for the SBCO-0.90 composition at 550, 600, and 650°C are summarized. Figure 3D presents the impedance results of SBCO-0.90 measured in the temperature range of 500–750°C, as well as the values R$_1$, R$_2$, and R$_3$ separating these results, obtained using an equivalent circuit.

As can be seen from Figures 3A–C, R$_2$ takes up the largest proportion of the total resistance. These results are also summarized in Table 2. In addition, all the compositions of Sm$_{1-x}$BaCo$_2$O$_{5+d}$ (x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, and 0.15) applied in this experiment showed the same results. For example, R$_2$ of SBCO-1 was calculated and found to be 0.2162 Ω · cm$^2$, which is about 72% of the total ASR; however, R$_1$ (0.074 Ω · cm$^2$) and R$_3$ (0.0104 Ω · cm$^2$) occupied only 28% of the total resistance. This trend does not change with increasing degree of non-stoichiometry of compositions and the R$_2$ values of SBCO-0.95 and SBCO-0.90 are 0.1112 and 0.0847 Ω · cm$^2$, with proportions of about 75 and 71% of the total resistance. Therefore, it can be
FIGURE 2 | Area specific resistance (ASR) results of (A) Sm$_{1-x}$BaCo$_2$O$_{5+d}$ oxide systems with respect to temperature, and (B) ASR dependence of cathodic polarization with respect to compositions at 600, 650, and 700°C.
determined that the RDS of Sm$_{1-x}$BaCo$_2$O$_{5+d}$ ($x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10$, and 0.15) oxide systems tested is $R_3$, which is the resistance when oxygen ions move in the cathode bulk. In addition, in Table 2, all $R_1$, $R_2$, and $R_3$ ASR values decreased as the non-stoichiometry compositions increased. Especially for the SBCO$_{0.95}$ composition, ASR decreases rapidly. The reason for the rapid decrease in ASR results for SBCO$_{0.95}$ is the effect of BaCoO$_2$ phase, as described in the above XRD and SEM results. In the case of the SBCO$_{0.90}$ composition, which showed increased concentration of BaCoO$_2$, the ASR value was lower than that of SBCO$_{0.95}$.

According to Qi et al., who studied the composition of the perovskite structure BaCo$_{0.6}$Zr$_{0.4}$O$_{3-δ}$ (BZC-BC) in nanocomposite form, advanced electrochemical properties were reported when a secondary phase of BaCo$_{0.96}$Zr$_{0.04}$O$_{3-δ}$ transformed into BaZr$_{0.82}$Co$_{0.18}$O$_{3-δ}$ main phase. For example, single-phase BaZr$_{0.80}$Co$_{0.20}$O$_{3-δ}$ (BZC2) showed a power density of 33 mW·cm$^{-2}$ at 750°C. However, the BCZ-BC oxide system containing the secondary phase (BaCo$_{0.96}$Zr$_{0.04}$O$_{3-δ}$) showed a value of about 1,530 mW·cm$^{-2}$, an increase of 46 times compared to the case of using single phase at the same temperature; this implies that the composition BZC-BC including secondary phase exhibited excellent properties. In addition, in the case of multi-phase cathode materials mixed with various compositions containing Ba and Co based oxide, it was reported that BaCoO$_{3-δ}$ generates a large number of oxygen vacancies that promote the oxygen reduction reaction (ORR) by assisting the movement of reduced oxygen ions (Qi et al., 2020). Therefore, the composition of SBCO$_{0.90}$ applied in this experiment, which has the lowest ASR characteristics, is affected by activated ORR caused by BaCoO$_2$.

**Figure 4** summarizes the impedance results of the Sm$_{1-x}$BaCo$_2$O$_{5+d}$ ($x = 0, 0.05, 0.10$, and 0.15) oxide systems measured at 600°C. ASR values decrease as the effect of degree of non-stoichiometry composition increased. However, in the SBCO$_{0.85}$ composition, the $R_3$ value increased sharply and the total ASR increased. From these results, it can be determined that the CoO phase found in the composition of SBCO$_{0.85}$, shown in **Figure 1**, somewhat limits the movement of oxygen ions in the bulk state of the cathode.

Additionally, electrochemical analysis was carried out using composite cathodes prepared by mixing CGO91 powder and SBCO$_{0.90}$. For the fabrication of the composite cathodes, the synthesized SBCO$_{0.90}$ was mixed with CGO91 at a weight ratio of 1:1. The summarized ASR results of the composite cathodes are given in **Figure 5** and Table 3. As can be seen in **Figure 5**, the two cathode materials of single phase SBCO$_{0.90}$ and composite phase exhibited similar ASR values over the entire measured temperature range. For example, the composite cathode showed ASR values of 0.031 and 0.024 Ω·cm$^{-2}$ at 700 and 750°C; it showed a slight difference of only 0.004 Ω·cm$^{-2}$ compared to the ASR value of SBCO$_{0.90}$. In the general case of composite cathodes, ASRs measured from the composite cathode tend to decrease. However, a large amount of CGO91 could rather interfere with the movement of oxygen ions in the bulk and on the surface of the cathode, and so composite did not significantly affect the reduction of resistance of the composite cathode applied in this experiment.

From these results, it can be confirmed that the effect of the composite cathode is not significant for the non-stoichiometric composition state in which the quantitative composition of the A-site of the layered perovskite changes. However, the thermal expansion problem of the SOFC cathode operating at high temperature range can be solved when the composite cathode concept is applied. In other words, thermal stability can be secured by lowering the thermal expansion coefficient (TEC). Co-based materials have high TEC of $20.2 \times 10^{-6}$K$^{-1}$ when SOFC is operated in high temperature operating temperature range; however, CGO91 has a relatively low TEC of $12.1 \times 10^{-6}$K$^{-1}$ (Kim et al., 2009). Therefore, the high TEC value generated from the cathode material substituted with Co can lower the coefficient of thermal expansion through composite cathodes.

Kim et al., who synthesized the composition of SmBaCo$_2$O$_{5+d}$ through in a composite with CGO91 at an ~50:50 ratio, reported that the TEC of $20.2 \times 10^{-6}$K$^{-1}$ decreased by about 66% to $13.3 \times 10^{-6}$K$^{-1}$ (Kim et al., 2009). By mixing the electrolyte materials and cathode materials, the thermal stability increases while the TECs rapidly decrease. Therefore, it is suitable to use a composite cathode with excellent electrochemical properties and relatively lower TEC when applying single phase SBCO$_{0.90}$ at a relatively high temperature ($650°C$). However, it is desirable to use a single phase cathode when using at a relatively lower temperature range.

The ASRs of Sm$_{1-x}$BaCo$_2$O$_{5+d}$ ($x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10$, and 0.15) applied in this study were 0.083, 0.095, 0.088, 0.074, 0.075, 0.041, 0.035, and 0.045 Ω·cm$^{-2}$, respectively, at 700°C. All compositions can be sufficiently used as cathode materials at 700°C because these ASR results are all 0.15 Ω·cm$^{-2}$ or less (Steele, 1996). The ASRs of SBCO$_{0.97}$, SBCO$_{0.96}$, SBCO$_{0.95}$, and SBCO$_{0.85}$ are 0.130, 0.129, 0.072, and 0.085 Ω·cm$^{-2}$ at 650°C. Especially, the SBCO$_{0.90}$ composition has an ASR of 0.130 Ω·cm$^{-2}$, and can thus be used as a cathode material at a relatively low temperature of 600°C.

The electrical conductivity results of the Sm$_{1-x}$BaCo$_2$O$_{5+d}$ ($x = 0, 0.01, 0.02, 0.03, 0.04, and 0.05$) oxide systems are

| Composition  | $R_x$ (2.5 MHz–100 Hz) Ω·cm$^2$ | $R_2$ (100 Hz–1 Hz) Ω·cm$^2$ | $R_3$ (1 Hz–0.05 Hz) Ω·cm$^2$ |
|--------------|-------------------------------|-------------------------------|-------------------------------|
| SBCO$_{1}$   | 0.0741                        | 0.2162                        | 0.0104                        |
| SBCO$_{0.99}$| 0.0650                        | 0.2405                        | 0.0167                        |
| SBCO$_{0.98}$| 0.0658                        | 0.2628                        | 0.0152                        |
| SBCO$_{0.97}$| 0.0650                        | 0.1826                        | 0.0118                        |
| SBCO$_{0.96}$| 0.0569                        | 0.1702                        | 0.0135                        |
| SBCO$_{0.95}$| 0.0279                        | 0.1112                        | 0.0082                        |
| SBCO$_{0.90}$| 0.0250                        | 0.0847                        | 0.0091                        |
| SBCO$_{0.85}$| 0.0317                        | 0.1355                        | 0.0115                        |
summarized in Figure 6A. According to the literature related to the electrical conductivity behavior of SmBaCo$_2$O$_{5+d}$ (SBCO, in this study SBCO$_1$), SBCO showed a maximum electrical conductivity value (570 S/cm) in the range of 200–250°C; then, the electrical conductivity started to decrease from 250°C, indicating Metal Insulator Transition (MIT) behavior (Kim et al., 2009). The electrical conductivity tendency of MIT in the LnBaCo$_2$O$_{5+d}$ system is influenced by various factors, such as small degree of polaron hopping and oxygen content (Maignan et al., 1999; Moon et al., 2001).

However, in Figures 6A,B, it can be seen that the temperature at which the change in electrical conductivity occurs is directly related to the non-stoichiometric composition. For the compositions of SBCO$_1$, SBCO$_{0.99}$, SBCO$_{0.98}$, and SBCO$_{0.97}$, which are relatively less influenced by the non-stoichiometric composition, the electrical conductivity value increases from 50°C and then begins to decrease at 200°C. For the SBCO$_{0.96}$ and SBCO$_{0.95}$ compositions, which show increased influence of the non-stoichiometric composition in the A-site, the electrical conductivity began to decrease at 250°C, as shown in Figure 6A. On the other hand, the electrical conductivity of SBCO$_{0.90}$ decreased from 650°C and the electrical conductivity values of SBCO$_{0.85}$ began to decrease at 450°C, as shown in Figure 6B.

From these results, it can be concluded that when the influence of the non-stoichiometric composition increases, temperature of the MIT behavior increases. The electrical conductivities of materials having non-stoichiometric compositions, namely the Sm$_{1-x}$BaCo$_2$O$_{5+d}$ ($x = 0$, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, and 0.15) oxide systems, are summarized in Figure 6C for the intermediate temperature range (500–750°C). The values of electrical conductivity increase as the effect of non-stoichiometric composition increases ($x \leq 0.05$). That is, the electrical conductivity increases when a relatively small amount of Sm is removed in the range of Sm$_{1-x}$BaCo$_2$O$_{5+d}$ ($x = 0$, 0.01, 0.02, 0.03, 0.04, and 0.05). Especially the SBCO$_{0.95}$ composition...
exhibited electrical conductivity values of about 917 and 732 S/cm at 500 and 600°C and showed the highest electrical conductivity values among the measured compositions. However, when the non-stoichiometric composition increased from \( x = 0.05 \) in the Sm\(_{1-x}\)BaCo\(_2\)O\(_{5+d}\) \( (x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, \) and \( 0.15 \)) oxide systems, the electrical conductivity value decreased. It can be concluded that this is because a large amount of BaCoO\(_{2.6}\) phase activates the conductivity of ions but inhibits the transfer of charge carriers (Chen et al., 2011; Liu et al., 2011).

Therefore, it can be seen that a percolation phenomenon that affects the electrical conductivity according to the non-stoichiometry is found in the Sm\(_{1-x}\)BaCo\(_2\)O\(_{5+d}\) \( (x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, \) and \( 0.15 \)) oxide systems. In addition, the conductivity characteristics found in the non-stoichiometric compositions exceeding \( x = 0.04 \) can be determined by the influence of the secondary phase. However, the detailed influence of the BaCoO\(_{2.6}\) composition for electrical conductivity in SmBaCo\(_2\)O\(_{5+d}\) oxide systems is not discussed now and will be reported later.

Therefore, it can be determined that as the deficiency level of A'-site increases, the electrical conductivity behavior conditions in this experiment changed. In addition, the lowest conductivity value in the IT range was 169 S/cm, a value exceeding 100 S/cm. This is a value that exceeds the standard that must be satisfied for material to be used as cathode for SOFC; therefore, all compositions can be suitable for use as cathode material for IT-SOFC (Tu et al., 1997). In addition, SBCO_0.95, having the highest electrical conductivity values in the IT ranges, also showed electrical conductivity values higher than those of NdBaCo\(_2\)O\(_{6-d}\) and the compositions of LnBaCo\(_2\)O\(_{5+d}\) (Ln = La, Pr, Nd, Sm, Gd, Y) (Kim and Manthiram, 2015; Yi et al., 2016). Especially SBCO_0.90, which shows excellent electrochemical properties, also showed a high electrical conductivity value of 500 S/cm or more in the IT range.
The electrochemical properties of materials having non-stoichiometric compositions in which Sm of the $A^\prime$-site was partially removed in SmBaCo$_2$O$_{5+d}$ oxide systems were studied for direct application of these materials as cathodes for IT-SOFC. In the case of the Sm$_{1-x}$BaCo$_2$O$_{5+d}$ ($x = 0, 0.01, 0.02, 0.03, 0.04, 0.05, 0.10, \text{and} 0.15$) oxide systems synthesized by SSR, single phase composition was confirmed for Sm$_{0.96}$BaCo$_2$O$_{5+d}$ (SBCO$_{0.96}$). However, secondary phase of BaCoO$_2$ was found for compositions of Sm$_{0.95}$BaCo$_2$O$_{5+d}$ (SBCO$_{0.95}$) in Sm$_{1-x}$BaCo$_2$O$_{5+d}$ oxide systems. Additional phase of CoO appeared for the SBCO$_{0.85}$ composition.

All of the synthesized compositions showed excellent electrochemical properties, with ASR values $< 0.15 \ \Omega \cdot \text{cm}^2$ at 650°C. The Sm$_{0.96}$BaCo$_2$O$_{5+d}$ (SBCO$_{0.90}$) showed the lowest ASR characteristics, with values of 0.119, 0.061, and 0.035 $\Omega \cdot \text{cm}^2$ at 600, 650, and 700°C. Additionally, the ASRs of the composite cathodes prepared by mixing with an electrolyte using CGO91 and SBCO$_{0.90}$ were 0.139 and 0.056 $\Omega \cdot \text{cm}^2$ at 600 and 650°C. All compositions exhibit electrical conductivity values of more than 100 S/cm over the temperature range (500–750°C) for IT-SOFC. It was also confirmed that the behavior conditions of MIT change as the non-stoichiometry composition of the $A^\prime$-site increases. SBCO$_{0.90}$ composition, which had the best electrochemical properties, showed the maximum (568 S/cm)
and minimum (387 S/cm) electrical conductivity values in the temperature range of 500–900°C.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author/s.

**AUTHOR CONTRIBUTIONS**

CK was responsible for conducting all experiments, analyzing experimental data, and writing papers. SW teaches experimental methods and informs questions about results. KS, S-WB, HK, and WC helped to write the manuscript. KS provided additional help in thesis publication. Finally, JK coordinated the overall experimental design, direction, and schedule. In addition, he reviewed the paper and gave direction. All authors contributed to the article and approved the submitted version.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fchem.2021.633868/full#supplementary-material
