Highly enhanced electrical properties of lanthanum-silicate-oxide-based SOFC electrolytes with co-doped tin and bismuth in La$_{9.33-x}$Bi$_x$Si$_{6-y}$Sn$_y$O$_{26}$

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Solid oxide fuel cells (SOFCs) are one of the most promising clean energy sources to be developed. However, the operating temperature of SOFCs is currently still very high, ranging between 1073 and 1273 K. Reducing the operating temperature of SOFCs to intermediate temperatures in between 773 and 1073 K without decreasing the conductivity value is a challenging research topic and has received much attention from researchers. The electrolyte is one of the components in SOFCs which has an important role in reducing the operating temperature of the SOFC compared to the other two fuel cell components, namely the anode and cathode. Therefore, an electrolyte that has high conductivity at moderate operating temperature is needed to obtain SOFC with medium operating temperature as well.

La$_{9.33}$Si$_6$O$_{26}$ (LSO) is a potential electrolyte that has high conductivity at moderate operating temperatures when this material is modified by doping with metal ions. Here, we report a modification of the structure of the LSO by partial substitution of La with Bi$^{3+}$ ions and Si with Sn$^{4+}$, which forms La$_{9.33-x}$Bi$_x$Si$_{6-y}$Sn$_y$O$_{26}$ with $x = 0.5, 1.0, 1.5,$ and $y = 0.1, 0.3, 0.5$, in order to obtain an electrolyte of LSO with high conductivity at moderate operating temperatures. The addition of Bi and Sn as dopants has increased the conductivity of the LSO. Our work indicated highly enhanced electrical properties of La$_{7.83}$Bi$_{1.5}$Si$_{5.7}$Sn$_{0.3}$O$_{26}$ at 873 K (1.84 x 10$^{-2}$ S cm$^{-1}$) with considerably low activation energy ($E_a$) of 0.80 eV comparing to pristine La$_{9.33}$Si$_6$O$_{26}$ (0.08 x 10$^{-2}$ S cm$^{-1}$).

Introduction

Solid oxide fuel cells (SOFCs) are energy devices that convert chemical reactions into electricity and produce up to 10 MW of electrical power. An SOFC has three main components: a cathode that reduces fuel, an anode that reduces oxygen gas to become oxygen ions, and a solid electrolyte responsible for delivering oxygen ions, and an anode that reduces oxygen gas to become oxygen ions.

General SOFCs typically operate at high temperatures in the range between 1073 K and 1273 K, which causes accelerated aging and damaging their components. Therefore, lowering the operating temperature is an important issue that must be developed in SOFC. In addition, lowering its working temperature is also necessary to eliminate power loss during operation. However, a decrease in operating temperature usually causes a decrease in SOFC performance. To resolve these problems, the use of high conductivity electrolytes is a very determining factor and must be considered when manufacturing SOFC. High conductivity electrolytes are usually obtained from new types of electrolytes, or modifications to existing electrolytes. However, finding new types of electrolytes is more difficult than modifying existing electrolytes. The usual modification by defect formation in crystals can be done by doping and forming composite. Electrolyte modification usually also correlates with other improvements in physical properties such as reduced brittleness and obtained high-density electrolytes with lower sintering temperature. The electrolyte from apatite type is an alternative electrolyte to overcome this problem because of the large conduction tunnel in apatite lattice structure, facilitating the circulation of free oxide ions which allows the increased conductivity of the electrolyte. LSO with a hexagonal crystal structure with three possible space clusters namely $P3$, $P6_3$, and $P6_3/m$ was first discovered by Nakayama et al. The electrolytes of the LSO generally exhibit insufficient conductivity at lower operating temperatures below 973 K. However, the open structure of the LSO allows modification by the doping technique on the La...
and Si site which makes it possible to obtain sufficient conductivity values at low-temperature operation. Several studies reported that by adding dopants to the apatite lattice structures at the La and Si sites, the conductivity can be increased. Some of the dopants that have been reported to substitute La and Si in LSO are Sr, Ca, Ba, Mg, Cu, Co, W, and Ti. Noviyanti et al. added 0.5 and 1.0 of Bi dopant at the La site, while Xiang et al. reported that 0.5 doping of Sn at the Si site resulted in conductivity values of $2.46 \times 10^{-4}$ S cm$^{-1}$ at 773 K and $5.71 \times 10^{-3}$ S cm$^{-1}$ at 1073 K. Whereas various Bi doping at composition of $0.5 \leq x \leq 2$ on La$_{1-x}$Bi$_x$(SiO$_4$)$_2$O$_3$ resulted in a conductivity of $2.4 \times 10^{-4}$ S cm$^{-1}$ at 973 K. Kebab et al. also reported that doping of Bi in CaBaLa$_x$Bi$_y$(SiO$_4$)$_2$O$_3$ has resulted a conductivity of $2.20 \times 10^{-5}$ S cm$^{-1}$ at 873 K. However, the simultaneous use of Bi and Sn doping at La and Si sites in the LSO-apatite structure has not been reported before. Therefore, this study aims to study the effect of various concentrations of Bi- and Sn-dopant substituted at La and Si sites on the La$_{9.33}$Si$_6$O$_{26}$ structure. To be utilized as an electrolyte of SOFC, the LSO must be synthesized into a solid membrane which is prepared by sintering at high temperatures. It should also be noted that to obtain a high quality and solid membrane, the temperature is an important step to obtain high-quality sintering temperatures of LSO require high temperatures of up to 1073 K. Therefore, the use of dopants that have the dual advantages of increasing conductivity and lowering the sintering temperature is an important step to obtain high-quality membranes. Bi is known to have the ability to increase conductivity as well as reduce sintering temperature. Meanwhile, Sn is known to be suitable support on Si site which can improve the total conductivity. The hydrothermal method was chosen as the synthesis method, in order to obtain a good crystalline powder and high purity. Here, we reported a study of the structure and electrical properties of La$_{9.33}$-Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$ in which the Bi$^{3+}$ and Sn$^{4+}$ were used simultaneously as doping at La and Si sites in various concentration. Our study offers the co-doping strategy of Sn and Bi to enhance the electrical properties several times higher than the pristine LSO and previously reported value.

**Experimental procedure**

The materials of La$_{9.33}$-Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$-apatite type with $x = 0.5$, 1.0, 1.5 and $y = 0.1, 0.3, 0.5$ were prepared from high-purity La$_2$O$_3$ (Sigma Aldrich, 99.999%), Na$_2$SiO$_3$, 5H$_2$O (Sigma Aldrich, 97%), Bi$_2$O$_3$, Sn$_2$O$_3$, and NaOH (Merck, 99%).

The experimental design in the synthesis of La$_{9.33}$-Bi$_x$Si$_6$-Sn$_y$O$_{26}$-apatite type with $x = 0.5$, 1.0, 1.5 and $y = 0.1, 0.3, 0.5$ are listed in Table 1.

Lanthanum oxide was initially calcined for the decarbonation purpose at 1373 K for 10 hours. Then, a certain amount of Na$_2$SiO$_3$ was weighed and dissolved in a 50 mL of a 3 M NaOH solution. A solution of Na$_2$SiO$_3$ was mixed with La$_2$O$_3$, SnO$_2$, and Bi$_2$O$_3$. The mixture was transferred into the autoclave and heated in an oven at 503 K for 3 days. The resulting product was washed with deionized water, then filtered and dried in an oven at 393 K for 24 hours. All samples were pelleted then sintered at 1773 K for 8 hours.

The structures of the La$_{9.33}$-Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$ were determined using a Phillips Analytical X-ray PW3373 diffractometer with a CuK$_\alpha$ radiation source ($\lambda = 0.15$ nm). The diffraction patterns were scanned from 10 to 80°. The crystal lattice parameters and its cell unit volume were analysed by using a General Structure and Analysis Software II (GSAS-II) and refined by using a pattern from JCPDS no. 04-011-7781 and no. 00-049-0443, for La$_{9.33}$-Bi$_x$Si$_6$O$_{26}$ (LSO) and La$_{9.33}$-Bi$_x$Si$_6$-Sn$_y$O$_{26}$, respectively.

The effect of the Bi$^{3+}$ and Sn$^{4+}$ dopant concentrations in the apatite conductivity was tested using Electrochemical Impedance Spectroscopic (EIS, GW Instek 6105G LCR meter). The sintered pellets were coated with silver paste on both sides with the effective area ($A$) of 0.026 cm$^2$, and then heated at 873 K for 10 minutes. The pellets are connected with platinum wire to the electrodes where the silver paste is coated. Impedance spectroscopic measurements were carried out at the range between 20 Hz and 5 MHz, a voltage of 1 V and at a range temperature of 573–1073 K. Based on the EIS measurement, the obtained Nyquist plots were interpreted using EIS analyzer. The conductivity value is calculated using the eqn (1).

\[
\sigma = \frac{L}{AR}\quad(1)
\]

where $\sigma$ = conductivity, $L$ = thickness of pellets (cm), $R$ = resistance (ohms) and $A$ = area of effective electrode (cm).

**Result and discussion**

**Structure**

La$_{9.33}$-Bi$_x$Si$_6$-Sn$_y$O$_{26}$-apatite type electrolytes were successfully synthesized via a hydrothermal method. Fig. 1 depicts

| Target compounds | Sample code | Precursors | Weight (g) | Dopants |
|------------------|-------------|------------|------------|--------|
| La$_{9.33}$Si$_6$O$_{26}$ | I | La$_2$O$_3$ | 1.6165 | — |
| La$_{9.33}$Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$ | II | Na$_2$SiO$_3$, 5H$_2$O | 1.3534 | — |
| La$_{9.33}$Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$ | III | Bi$_2$O$_3$ | 1.4948 | 0.1211 |
| La$_{9.33}$Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$ | IV | Sn$_2$O$_3$ | 1.3003 | 0.0157 |
| La$_{9.33}$Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$ | V | | 1.2445 | 0.1199 |
| La$_{9.33}$Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$ | VI | | 1.1892 | 0.0465 |
| La$_{9.33}$Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$ | VII | | 1.2225 | 0.1188 |
| La$_{9.33}$Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$ | VIII | | 1.2012 | 0.0769 |
| La$_{9.33}$Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$ | IX | | 0.2356 | 0.0457 |
| La$_{9.33}$Bi$_{0.5}$Si$_6$-Sn$_{0.5}$O$_{26}$ | X | | 0.3472 | 0.0449 |

Table 1: The precursors and the target compounds of La$_{9.33}$-Bi$_x$Si$_6$-Sn$_y$O$_{26}$

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The XRD patterns of co-doped La$_{9.33}$Si$_6$O$_{26}$-apatite phases in various value of La with various intensities as shown in Fig. 1. The intensity of the peak at the $2\theta$ of 15° elevated with the increasing content of the Bi-doping. Adding a higher of Bi-doping in the La site of LSO should be eliminating the phase impurities at this $2\theta$ region. However, we found contrastly in this study at which the higher content of Bi with the co-existence of Sn-doping in La$_{9.33-x}$Bi$_x$Si$_{6-y}$Sn$_y$O$_{26}$ may possibly lead to the disintegration of the La ions, resulting in La(OH)$_3$ phase. The Rietveld refinement’s result shows that the entire apatite-type electrolyte has hexagonal-based crystal structure with $P3$ space group. The certain amount of dopant may change the lattice parameter as shown in Table 2. A larger dopant of Bi$^{3+}$, with the radius of 1.17 Å, prefers the La site (La$^{3+}$ radius = 1.15 Å). Meanwhile, a smaller dopant such as Sn$^{4+}$ that exhibits the radius of 0.71 Å prefers the Si site (Si$^{4+}$ radius = 0.41 Å). The greater the Bi concentration, the higher the lattice parameter both on the $a$- and $c$-axis. It is because the size of Bi is larger than that of La, especially seen in samples V and VI with Bi concentrations of 1 and 1.5 respectively, as listed in Table 1 and Fig. 2. The possible reasons for increasing the lattice parameters and volume of the unit cell first and then decreasing gradually are as follow: first, the existence of impurity that disturbing the position of atoms or decreasing the content of several atoms in the crystals structure, second, the simultaneous doping of two atoms causes opposite effects to each other in certain compositions.

The simulations of the positions of La, Si, and O atoms in the LSO structure are shown in Fig. 3. Based on their positions, La, Si, and O atoms are divided into La1, La2, La3, Si, O1, O2, O3, O4, O5, and O6. It is well known that O6 is interstitial oxygen due to a Frenkel defect that is formed due to the influence of Bi and Sn doping. The atomic coordinates for all atoms including the site occupation factor are listed in Table 3.

The bond lengths between the atoms at the LSO for all samples are summarized in Table 4. Changes in bond length of all La–O bonds and Si–O bonds due to the doping effect of Bi and Sn are clearly observed in sample V and sample VI. These changes can also result in the enlargements of the unit cell and create structural imbalances that can affect their physical properties as expected.
Conductivity

All samples were designed with the same void concentration (La = 9.33) and oxygen content (O = 26) so that changes in conductivity could be directly compared and evaluated as the effect of Bi and Sn doping concentrations on doped-LSO. The conductivity of LSO and doped-LSO is shown in Fig. 4 and Table 4. It is found that the doping effect of Bi and Sn appears to significantly affect their conductivity. The Nyquist plot showed decreasing a semi-circle impedance as varied Sn dopants which suggested an increasing of the conductivities. For example, the conductivity of sample V, which $\sigma_{873\,K}$ of $1.37 \times 10^{-2}$ S cm$^{-1}$, increased almost twice comparing to sample III ($\sigma_{873\,K} = 0.74 \times 10^{-2}$ S cm$^{-1}$). Bi and Sn doping also have the effect of reducing the $E_a$ from 0.69 to 0.76 eV, which indicates that the movement of oxide ions is facilitated in the doped-apatite structure. In addition, the doping concentration also significantly affects the increase in conductivity. Increasing the Sn concentration from 5% ($y = 0.3$) to ~8.3% ($y = 0.5$) as in sample IV actually decreased its conductivity. However, on the other hand, a different effect is shown by Bi doping. It was found that increasing the Bi concentration tended to increase its conductivity as well, as shown in samples V and VI. It is found that sample VI of La$_{7.83}$Bi$_{1.5}$Si$_{5.7}$Sn$_{0.3}$O$_{26}$ has the highest value of conductivity as produced by sample IV.

Conductivity of LSO and co-doped LSO-based materials

Table 4

| Bond distance (Å) | Sample code | I  | II  | III | IV  | V   | VI  |
|------------------|-------------|----|-----|-----|-----|-----|-----|
| La1–O1           | 2.576       | 2.577| 2.577| 2.577| 2.581| 2.580|
| La1–O2           | 2.512       | 2.512| 2.513| 2.513| 2.516| 2.516|
| La1–O3           | 2.914       | 2.915| 2.916| 2.916| 2.922| 2.922|
| La2–O1           | 2.430       | 2.430| 2.431| 2.431| 2.435| 2.435|
| La2–O2           | 2.559       | 2.560| 2.561| 2.560| 2.564| 2.564|
| La2–O3           | 2.497       | 2.499| 2.500| 2.499| 2.505| 2.504|
| La3–O1           | 2.716       | 2.718| 2.718| 2.718| 2.724| 2.723|
| La3–O2           | 2.497       | 2.499| 2.500| 2.500| 2.505| 2.504|
| La3–O3           | 2.588       | 2.589| 2.590| 2.589| 2.594| 2.594|
| La3–O4           | 2.435       | 2.435| 2.435| 2.430| 2.437| 2.437|
| La3–O5           | 2.307       | 2.309| 2.310| 2.309| 2.314| 2.313|
| La3–O6           | 2.478       | 2.480| 2.480| 2.480| 2.485| 2.484|
| Si–O1            | 1.606       | 1.607| 1.608| 1.607| 1.611| 1.610|
| Si–O2            | 1.627       | 1.628| 1.629| 1.628| 1.632| 1.631|
| Si–O3            | 1.617       | 1.617| 1.617| 1.617| 1.619| 1.619|
| Si–O4            | 1.626       | 1.626| 1.626| 1.626| 1.628| 1.628|

The significant increase in conductivity values in samples V and VI can be explained by changes in the bond length between La–O and Si–O. High conductivity is found in doped-LSO which have the longest La–O and Si–O bond lengths. The addition of the La–O and Si–O bond lengths caused the channel on the c-axis to expand as well, thereby facilitating the movement of oxide ions. Especially for the effect of adding Sn-dopant to the Si site in samples I to IV, the conductivity value increased with increasing Sn from 0 to 0.3 (sample I to sample III). However, a bit increasing of Sn dopant (0.5) dropped the conductivity almost twice from 0.74 to 0.38 S cm$^{-1}$. The maximum conductivity value was obtained when Sn was 0.3 (sample III). The reason for this data may be explained by looking at the conduction pathway at the position of La at site 6g. An increase in Sn at the Si site in excess of 0.3 can inhibit the ionic conduction pathway by disrupting the position of La at site 6 and pushing it into a narrow conduction pathway, which results in a low conductivity value as produced by sample IV.

The $E_a$ lower than 1.1 eV indicates the ion conduction pathways at which adopting an interstitial ion migration mechanism at the temperature range of 573 to 873 K. The existence of the O6 site obtained a from the simulation results confirmed that the dominant mechanism in the conduction of...
LSO and doped-LSO is the insertion mechanism. The $E_a$ value of each sample together with the value of conductivity is presented in Table 5. Generally, the $E_a$ is lowered after an introduction of Bi dopant with constant Sn concentration. A bit enhancement of Sn dopant to the system drastically declines the $E_a$ along with an increase of $E_a$. With the same Bi content ($x = 0.5$), $E_a$ elevated from 0.69 to 0.87 eV as increasing Sn dopant (from 0.1 to 0.5) which also showed the similar behaviour with the same Sn concentration of 0.3 and increasing of Bi dopant from 0.5 to 1.5 with $E_a$ increased from 0.77 to 0.80 eV.

Conclusions
The materials of co-doping of Bi and Sn in the structure of La$_{9.33-x}$Bi$_x$Si$_6$Sn$_y$O$_{26}$-apatite with $x = 0.5$, 1, and 1.5 and $y = 0.1$, 0.3, and 0.5 were successfully synthesized by using a hydrothermal method. All main peaks of LSO are observed in all samples. The conductivity of the La$_{9.33-x}$Bi$_x$Si$_6$Sn$_y$O$_{26}$-apatite type increased as the Bi-doping content increases. La$_{7.83}$Bi$_{1.5}$Si$_{5.7}$Sn$_{0.3}$O$_{26}$ has the highest value of conductivity as $1.84 \times 10^{-2}$ S cm$^{-1}$ at 873 K with the activation energy ($E_a$) value of 0.80 eV which indicates the adoption of the interstitial ionic conduction migration pathways. Our study can open a new possibility of structure modification of lanthanum-oxide-based SOFC electrolyte to achieve high electrochemical performance for the application in the clean energy generation.

Author contributions
A. R. N.-conceptualisation, writing-original draft, writing-revision, investigation; J. J.-supervision, formal analysis; S. W.-data curation; R. R.-writing-revision, supervision, formal analysis; D. G. S.-resources; Y. T. M.-writing-revision, formal analysis, investigation, data curation; R. S.-writing-revision.

Conflicts of interest
There are no conflicts to declare.

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