Dielectric characteristics of fast Li ion conducting garnet-type Li\textsubscript{5+2x}La\textsubscript{3}Nb\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12} (x = 0.25, 0.5 and 0.75)

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Here, we report the dielectric characteristics of Li-stuffed garnet-type Li\textsubscript{5+2x}La\textsubscript{3}Nb\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12} (x = 0.25, 0.5 and 0.75) in the temperature range about −53 to 50 °C using AC impedance spectroscopy. All the investigated Li-stuffed garnet compounds were prepared, under the same condition, using conventional solid-state reaction at elevated temperature in air. The Nyquist plots show mainly bulk contribution to the total Li\textsuperscript{+} ion conductivity for Li\textsubscript{5}La\textsubscript{3}Nb\textsubscript{1.75}Y\textsubscript{0.25}O\textsubscript{12} (Li\textsubscript{5+}–Nb) and Li\textsubscript{5}La\textsubscript{3}Nb\textsubscript{1.5}Y\textsubscript{0.5}O\textsubscript{12} (Li\textsubscript{5+}–Nb), while both bulk and grain-boundary effects are visible in the case of Li\textsubscript{5}La\textsubscript{3}Nb\textsubscript{1.25}Y\textsubscript{0.75}O\textsubscript{12} (Li\textsubscript{5+}–Nb) phase at ∼−22 °C. Non-Debye relaxation process was observed in the modulus AC impedance plots. The dielectric loss tangent of Li\textsubscript{5+2x}La\textsubscript{3}Nb\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12} are compared with that of the corresponding Ta analogue, Li\textsubscript{5+2x}La\textsubscript{3}Ta\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12} and showed a decrease in peak intensity for the Nb-based garnet samples which may be attributed to a slight increase in their Li\textsuperscript{+} ion conductivity. The relative dielectric constant values were also found to be higher for the Ta member (>60 for Li\textsubscript{5+2x}La\textsubscript{3}Ta\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12}) than that of the corresponding Nb analogue (<50 for Li\textsubscript{5+2x}La\textsubscript{3}Nb\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12}) at below room temperature. A long-range order Li\textsuperscript{+} ion migration pathway with relaxation time (τ\textsubscript{0}) 10\textsuperscript{−18}–10\textsuperscript{−15} s and an activation energy of 0.59–0.40 eV was observed for the investigated Li\textsubscript{5+2x}La\textsubscript{3}Nb\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12} garnets and is comparable to that of the corresponding Ta-based Li\textsubscript{5+2x}La\textsubscript{3}Ta\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12} garnets.

Introduction

Currently, lithium ion batteries (LIBs) attain much attention because of their high volumetric and gravimetric energy densities, which make them suitable for possible applications from portable devices to electric vehicles and grid scale to support the peak power demand. However, there is a long-term remaining safety concern related to the presently used organic polymer-based Li\textsuperscript{+} ion electrolytes in LIBs and they must be replaced with thermally stable fast Li\textsuperscript{+} ion conductors. To this end, several families of inorganic metal oxides and non-oxides have been investigated, of which recently discovered, Li-stuffed garnet compounds were prepared, under the same condition, using conventional solid-state reaction at elevated temperature in air. The Nyquist plots show mainly bulk contribution to the total Li\textsuperscript{+} ion conductivity for Li\textsubscript{5}La\textsubscript{3}Nb\textsubscript{1.75}Y\textsubscript{0.25}O\textsubscript{12} (Li\textsubscript{5+}–Nb) and Li\textsubscript{5}La\textsubscript{3}Nb\textsubscript{1.5}Y\textsubscript{0.5}O\textsubscript{12} (Li\textsubscript{5+}–Nb), while both bulk and grain-boundary effects are visible in the case of Li\textsubscript{5}La\textsubscript{3}Nb\textsubscript{1.25}Y\textsubscript{0.75}O\textsubscript{12} (Li\textsubscript{5+}–Nb) phase at ∼−22 °C. Non-Debye relaxation process was observed in the modulus AC impedance plots. The dielectric loss tangent of Li\textsubscript{5+2x}La\textsubscript{3}Nb\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12} are compared with that of the corresponding Ta analogue, Li\textsubscript{5+2x}La\textsubscript{3}Ta\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12} and showed a decrease in peak intensity for the Nb-based garnet samples which may be attributed to a slight increase in their Li\textsuperscript{+} ion conductivity. The relative dielectric constant values were also found to be higher for the Ta member (>60 for Li\textsubscript{5+2x}La\textsubscript{3}Ta\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12}) than that of the corresponding Nb analogue (<50 for Li\textsubscript{5+2x}La\textsubscript{3}Nb\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12}) at below room temperature. A long-range order Li\textsuperscript{+} ion migration pathway with relaxation time (τ\textsubscript{0}) 10\textsuperscript{−18}–10\textsuperscript{−15} s and an activation energy of 0.59–0.40 eV was observed for the investigated Li\textsubscript{5+2x}La\textsubscript{3}Nb\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12} garnets and is comparable to that of the corresponding Ta-based Li\textsubscript{5+2x}La\textsubscript{3}Ta\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12} garnets.

Dielectric characteristics of fast Li ion conducting garnet-type Li\textsubscript{5+2x}La\textsubscript{3}Nb\textsubscript{2−x}Y\textsubscript{x}O\textsubscript{12} (x = 0.25, 0.5 and 0.75)

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conduction mechanism in the garnet structure.\textsuperscript{18,19} Furthermore, several studies have been carried out to understand the chemical composition and Li\(^+\) ion dynamics relationship in these Li-stuffed garnets using neutron diffraction, NMR, electrochemical AC impedance spectroscopy and DFT calculations.\textsuperscript{8,10,19–31} The present study is focused on analyzing the dielectric properties of Li\(_{5+2x}\)La\(_{3}Nb\(_{2-x}\)Y\(_{x}\)O\(_{12}\) (x = 0.25, 0.5 and 0.75) using electrochemical AC impedance spectroscopy to better understand the Li\(^+\) ion conduction mechanism in the Li-stuffed garnets.

### Experimental

Single phase cubic structured garnet-type Li\(_{5+2x}\)La\(_{3}Nb\(_{2-x}\)Y\(_{x}\)O\(_{12}\) (x = 0.25, 0.5 and 0.75 which are represented as Li\(_{5.5}\)-Nb, Li\(_{6}\)-Nb and Li\(_{6.5}\)-Nb, respectively) were prepared by solid-state reaction at elevated temperature.\textsuperscript{32} The stoichiometric amount of the precursors of high purity LiNO\(_{3}\) (99%, Alfa Aesar), La\(_{2}\)O\(_{3}\) (99.99%, Alfa Aesar, preheated at 900 °C for 24 h), Nb\(_{2}\)O\(_{5}\) (99.5%, Alfa Aesar), and Y(NO\(_{3}\))\(_{3}\) (99.9%, Alfa Aesar) were ball milled to ensure homogeneity. 10 wt% excess LiNO\(_{3}\) was added to compensate the loss of lithium oxide while sintering. Heating at different stages were carried out at 700 °C/6 h, 900 °C/24 h and 1100 °C/6 h. The dielectric analyses were carried out using the AC impedance spectroscopy employing a Solartron 1260 impedance/gain-phase analyzer (0.01 Hz to 1 MHz) in the temperature range of -53 to 50 °C in air. The temperature was controlled below room temperature using dry ice.\textsuperscript{33} The pellet garnet samples for AC impedance measurements were painted with Au paste on both sides and were cured for 1 h at 600 °C.

### Results and discussion

#### AC impedance spectroscopy

Typical AC impedance spectra of cubic structured garnet-type Li\(_{5+2x}\)La\(_{3}Nb\(_{2-x}\)Y\(_{x}\)O\(_{12}\) (x = 0.25, 0.5 and 0.75 which are represented as Li\(_{5.5}\)-Nb, Li\(_{6}\)-Nb and Li\(_{6.5}\)-Nb, respectively) at -22 °C are shown in Fig. 1. The Li\(_{5.5}\)-Nb and Li\(_{6}\)-Nb phases of Li\(_{5+2x}\)La\(_{3}Nb\(_{2-x}\)Y\(_{x}\)O\(_{12}\) show a single semicircle, which corresponds to bulk (b) contribution at higher frequency range and a spike at the low-frequency range due to polarization resistance. The line spacing through the data points is fitted using an equivalent circuit consisting of (R\(_{b}\)CPE\(_{b}\))(CPE\(_{e}\)) for Li\(_{5.5}\)-Nb and Li\(_{6}\)-Nb and (R\(_{b}\)CPE\(_{b}\))(R\(_{gb}\)CPE\(_{gb}\))(CPE\(_{e}\)) for Li\(_{6.5}\)-Nb.

![Fig. 1 Typical complex impedance spectra obtained using Li\(^+\) ion blocking Au electrodes for Li\(_{5+2x}\)La\(_{3}Nb\(_{2-x}\)Y\(_{x}\)O\(_{12}\) (Li\(_{5.5}\)-Nb), Li\(_{6}\)-Nb, and total (bulk + grain-boundary) Li\(^+\) ion conductivity of Li\(_{6.5}\)-Nb. The line passing through the data points is fitted using an equivalent circuit consisting of (R\(_{b}\)CPE\(_{b}\))(CPE\(_{e}\)) for Li\(_{5.5}\)-Nb and Li\(_{6}\)-Nb and (R\(_{b}\)CPE\(_{b}\))(R\(_{gb}\)CPE\(_{gb}\))(CPE\(_{e}\)) for Li\(_{6.5}\)-Nb.](image)

\[\sigma = \frac{1}{R} \left( \frac{l}{a} \right) \]  \quad (1)

\[\log_{10} 10^{\alpha} (\text{S/m}) = \left( -\frac{E}{10^{\frac{E}{4.5}}} \right) \]  \quad (2)

Li\(_{5.5}\)-Li\(_{6}\)-Li\(_{6.5}\) phase. These factors might be contributing to the presence of grain-boundary impedance in Li\(_{5.5}\)-Nb compared to Li\(_{5.5}\)-Nb and Li\(_{6}\)-Nb. The Li\(^+\) ion conductivity (σ) from the low-frequency intercept point of Z′ (imaginary part of impedance) to the X-axis (Z′, real part of the impedance) (Fig. 1). The Li\(^+\) ion conductivity was calculated using the eqn (1):

\[\sigma = \frac{1}{R} \left( \frac{l}{a} \right) \]  \quad (1)

\[\log_{10} 10^{\alpha} (\text{S/m}) = \left( -\frac{E}{10^{\frac{E}{4.5}}} \right) \]  \quad (2)

Fig. 2 Arrhenius plots of bulk Li ion conductivity of Li\(_{5+2x}\)La\(_{3}Nb\(_{2-x}\)Y\(_{x}\)O\(_{12}\) (Li\(_{5.5}\)-Nb), Li\(_{6}\)-Nb, and total (bulk + grain-boundary) Li\(^+\) ion conductivity of Li\(_{6+2x}\)La\(_{3}Nb\(_{2-x}\)Y\(_{x}\)O\(_{12}\) (Li\(_{6.5}\)-Nb). The line passing through the data points is the fitted line using eqn (2).
where $l$ is the sample thickness and $a$ is the area of current collector. The Li$^+$ ion conductivity dependence on temperature was represented as the Arrhenius plots (Fig. 2) and they can be described using the eqn (2):

$$
\sigma T = A \exp \left( \frac{E_a}{kT} \right)
$$

where $T$ is the temperature, $A$ is the pre-exponential factor, $E_a$ is the activation energy, and $k$ is the Boltzmann’s constant ($1.38 \times 10^{-23}$ J K$^{-1}$). The Li$^+$ ion conductivity of samples increases from Li$_{5.5}$La$_3$Nb$_{1.75}$Y$_{0.25}$O$_{12}$ (Li$_{5.5}$–Nb) to Li$_{6.5}$La$_3$Nb$_{1.25}$Y$_{0.75}$O$_{12}$ (Li$_{6.5}$–Nb), which is consistent with the increase in cell parameter ($a$) calculated from powder X-ray diffraction (PXRD). The cell constant was found to be 12.8582(5) Å, 12.9136(4) Å and 12.9488(11) Å for Li$_{5.5}$La$_3$Nb$_{1.75}$Y$_{0.25}$O$_{12}$ (Li$_{5.5}$–Nb), Li$_6$La$_3$Nb$_{1.5}$Y$_{0.5}$O$_{12}$ (Li$_6$–Nb), and Li$_{6.5}$La$_3$Nb$_{1.25}$Y$_{0.75}$O$_{12}$ (Li$_{6.5}$–Nb), respectively. The activation energy calculated for Li$_{5.5}$–Nb, Li$_6$–Nb and Li$_{6.5}$–Nb phases are 0.59, 0.58 and 0.42 eV, respectively, in the temperature range of $-53$ to $50$ °C (Table 1) which is in agreement with the Li$^+$ ion conductivity trend.

Table 1 The bulk (except bulk + grain-boundary for Li$_{6.5}$–Nb) conductivity of Li$_{5.5}$La$_3$Nb$_{2-x}$Y$_x$O$_{12}$ at $-22$ and $25$ °C and the activation energy calculated from the Arrhenius plot (Fig. 2) at temperature range $-53$ to $50$ °C

| Sample | $\sigma_{-22}$ (S cm$^{-1}$) | $\sigma_{25}$ (S cm$^{-1}$) | $E_a$ (eV) |
|--------|-----------------------------|-----------------------------|------------|
| Li$_{5.5}$La$_3$Nb$_{1.75}$Y$_{0.25}$O$_{12}$ (Li$_{5.5}$–Nb) | 1.50 $\times$ 10$^{-6}$ | 7.18 $\times$ 10$^{-3}$ | 0.59 |
| Li$_{6}$La$_3$Nb$_{1.5}$Y$_{0.5}$O$_{12}$ (Li$_6$–Nb) | 4.20 $\times$ 10$^{-6}$ | 1.87 $\times$ 10$^{-4}$ | 0.58 |
| Li$_{6.5}$La$_3$Nb$_{1.25}$Y$_{0.75}$O$_{12}$ (Li$_{6.5}$–Nb) | 1.27 $\times$ 10$^{-5}$ | 2.99 $\times$ 10$^{-4}$ | 0.42 |

The relationship of conductivity ($\sigma$) and frequency is expressed by the Jonscher universal power law:

$$
\sigma_\omega = \sigma_{dc} + A\omega^n
$$

where $\omega$ is the angular frequency, $\sigma_{dc}$ is the dc conductivity, $A$ is the proportionality constant, and $n$ is the power factor which varies from 0 to 1. Shown in Fig. 3 is the Li$^+$ ion conductivity as a function of frequency for Li$_{5.5-2x}$La$_3$Nb$_{2-x}$Y$_x$O$_{12}$ in the temperature range from $-53$ to $50$ °C. The dispersion region seen at lower frequency range is due to the polarization due to Li$^+$ ion-blocking electrode. The frequency independent dc plateau region is seen at intermediate frequency range followed by a dispersion region. This could be explained by jump relaxation model which indicates that at intermediate frequencies, the ions hop from one site to the neighboring vacancy site contributing to the dc conductivity, and at higher frequencies the correlated alternative

![Fig. 3](image-url) The electrical conductivity of (a) Li$_{5.5}$La$_3$Nb$_{1.75}$Y$_{0.25}$O$_{12}$ (Li$_{5.5}$–Nb), (b) Li$_6$La$_3$Nb$_{1.5}$Y$_{0.5}$O$_{12}$ (Li$_6$–Nb) and (c) Li$_{6.5}$La$_3$Nb$_{1.25}$Y$_{0.75}$O$_{12}$ (Li$_{6.5}$–Nb) as a function of frequency at different temperatures obtained using AC impedance spectroscopy with Li$^+$ ion blocking Au electrodes.
hopping along with the relaxation of ions contribute to the conductivity relaxation at the dispersion region.\textsuperscript{35,36} For garnet-type oxide, the jumping of Li\textsuperscript{+} ions from one octahedral site to another octahedral site (for \textit{e.g.}, 48g-site to the 96h-site) followed by the readjustment of surrounding ions causes the conductivity relaxation.\textsuperscript{25}

The real ($\varepsilon'$) and imaginary ($\varepsilon''$) part of permittivity ($\varepsilon$) was computed using the eqn (4) and (5) as shown below:\textsuperscript{37}

$$\varepsilon' = \frac{Z''}{\omega C_0 (Z'^2 + Z''^2)}$$  \hspace{1cm} (4)

$$\varepsilon'' = -\frac{Z'}{\omega C_0 (Z'^2 + Z''^2)}$$  \hspace{1cm} (5)

where $C_0$ is the vacuum capacitance of the cell which is represented as $C_0 = \varepsilon_0 (a/l)$ (where $\varepsilon_0$ is the permittivity of free space $= 8.854 \times 10^{-14}$ F cm$^{-1}$). Fig. 4 shows the relation between the real part of permittivity ($\varepsilon'$) and frequency at different temperatures. It is noticeable that the dielectric constant decreases with frequency showing a constant minimum value. The upturn at the lower frequency region may be due to the blocking electrode-electrolyte polarization at the interface. When the frequency is increased, the periodical reversal of the field is increased and as a result the oscillating charges do not contribute to the dielectric constant. This causes a decrease in dielectric constant with increase in frequency. Dipolar polarization leads to formation of a plateau at the low-frequency region at higher temperature, which is clearly visible in the case of Li$_{6.5}$–Nb phase and also in the Ta series, Li$_{5+x}$La$_3$Ta$_{2-x}$Y$_x$O$_{12}$ ($x = 0.25$, 0.5 and 0.75).\textsuperscript{25} Shifting of the plateau towards high frequencies with increasing temperature indicates the increase in frequency of dipolar rotations, as Li\textsuperscript{+} ion mobility is enhanced due to the thermal activation.\textsuperscript{38}

The plot of imaginary part of the dielectric permittivity ($\varepsilon''$) against frequency is illustrated in Fig. 5a–c. $\varepsilon''$ is the dielectric loss factor which could be the combination of dipolar loss and conduction loss. The conduction loss is predominant at lower temperatures and appears as a straight line and shows inverse relationship to the angular frequency ($\omega$) especially for Li$_{5.5}$–Nb and Li$_6$–Nb phases (Fig. 5a and b). In all the cases,
the peak/plateau at lower frequencies due to dipolar polarization was observed (Fig. 4).

The dielectric loss tangent as a function of frequency of $Li_{5+2x}La_{3}Nb_{2}xY_{x}O_{12}$ is shown in Fig. 6. The peak observed in $Li_{5.5-Nb}$ and $Li_{6-Nb}$ phases can be attributed to the dielectric loss due to Li–Li dipolar rotations under the applied electrical field. There seem to be two relaxation peaks for $Li_{6.5-Nb}$ phases at intermediate frequencies, suggesting that an additional polarization loss occurs when compared to the $Li_{5.5-Nb}$ and $Li_{6-Nb}$ phases. The higher frequency peak may be attributed to the dielectric loss due to Li–Li dipolar rotation in the bulk. The lower frequency peak in Fig. 6c seems to be due to the dielectric loss, as a result of grain-boundary polarization. This is consistent with the appearance of a grain-boundary arc in the AC impedance spectrum of $Li_{6.5-Nb}$ phase (Fig. 1). The shift in loss tangent peaks towards higher frequencies with increase in temperature, similar to the real part of permittivity ($\varepsilon'$) as a function of temperature, indicates that both grain-boundary space charge polarization and Li–Li dipolar interaction in bulk are thermally activated processes.

In Fig. 7, the loss tangent of both Nb with Ta members of $Li_{5+2x}La_{3}M_{2-x}Y_{x}O_{12}$ ($M = Nb, Ta$) at different temperatures is compared. A similar plot is observed, except for $Li_{6.5-Nb}$ phase of Nb series $Li_{5+2x}La_{3}Nb_{3}xY_{x}O_{12}$, corresponding to bulk and grain-boundary contribution. However, very small values of peak intensity of $Li_{6.5-Nb}$ phases of Nb series $Li_{5+2x}La_{3}Nb_{3}xY_{x}O_{12}$ compared to that of the Ta series $Li_{5+2x}La_{3}Ta_{3}xY_{x}O_{12}$ indicated that either there are fewer dipoles in the Nb garnets than in the Ta garnets or the mobility of the ions is higher in the presently investigated Nb garnets leading to a quick response of Li–Li dipoles to external electric field and resulting in less dielectric loss. Overall, the dielectric relaxation can be considered due to the main-body interactions as is common for the ionic conductor. The relative permittivity/dielectric constant ($\varepsilon_r$) of $Li_{5+2x}La_{3}M_{2-x}Y_{x}O_{12}$ ($M = Nb, Ta$) were calculated using the capacitance ($C$), as given in eqn (6) and the results are summarized in Fig. 8.

$$C = \frac{\varepsilon_0 a}{\varepsilon_r}$$  

The dielectric constant slightly increases with an increase in the Li content in the Ta series materials, although there is no trend in variation of $\varepsilon_r$ with the Li content in the Nb series. With the increase in temperature, the value of the dielectric constant...
is increased in both Nb and Ta based garnets. The dielectric constant of the Nb garnets is found to be relatively higher than that of the Ta garnets. Generally, the higher the dielectric constant, the greater is the polarizability of the medium. Thus, the higher dielectric constant (Fig. 8) and lower dielectric loss (Fig. 7) in the Nb series materials seems to indicate that their Li$^+$ ion mobility is higher compared to that of the Ta garnets since both Nb and Ta oxides exhibit similar ionic radius and electronic structure. A comparison of Li$^+$ ionic conductivity of both Nb and Ta phases of Li$^{5+2}x$La$_3M_2$/Co$_x$Y$_x$O$_{12}$ at specific temperatures is shown in Table 2. Using modulus analysis, it is possible to understand the bulk relaxation properties in detail by suppressing the lower frequency phenomena, especially the electrode effect. The complex dielectric modulus is expressed as eqn (7):

$$M = j\omega C_0[Z' + Z'']$$

where the real and imaginary parts can be separated as $M' = \omega C_0Z'$ and $M'' = -\omega C_0Z''$. The electric modulus accounts for the electric field relaxation in the material at a constant electric displacement. The modulus spectra of Li$_{5.5}$–Nb, Li$_{6}$–Nb and Li$_{6.5}$–Nb phases of Li$_{5+2}x$La$_3$M$_2$/Co$_x$Y$_x$O$_{12}$ samples, measured at a temperature range of $-53$ to $-21$ °C as a function of frequency is shown in Fig. 9. The electric modulus ($M''$) shows a frequency independent behavior at the lower frequency region for all the samples. This indicates that the electrode polarization does not contribute to the electric modulus and the long-range migration of Li$^+$ ion by hopping from one site to the neighboring site. The presence of a peak in the modulus spectra has been ascribed to the relaxation associated with Li$^+$ ion mobility. The appearance of a high frequency side peak at lower temperatures could be due to the Li$^+$ ion re-orientation relaxation as Li$^+$ ion moves from one octahedron to another octahedron around the immobile tetrahedral site. The shift in peak position towards higher frequencies with increase in temperature indicates that Li$^+$ ion relaxation re-orientation is a thermally activated phenomenon. Also, it is noticeable that the peak tend to disappear in the Li$_{6.5}$–Nb phase which can be explained in terms of long-range order migration of the Li$^+$ ion instead of local migration as seen in other members. The relaxation time, $\tau$, is related to the frequency maximum in the modulus plots, $f_{M''}$ as shown in eqn (8):

$$\tau = \frac{1}{2\pi f_{M''}}$$
which can actually reveal the short range and/or the long range migration of Li$^+$ in the crystal structure. Fig. 10 shows the Arrhenius behavior of the relaxation time, $t$, for the local motion of Li$^+$ according to the eqn (9):

$$
t = t_0 \exp \left( \frac{-E_m}{k_B T} \right) \tag{9}
$$

where $t_0$ is the relaxation time at infinite temperature and $E_m$ is the migration energy. The migration energy calculated for Li$_{5.5}$–Nb, and Li$_6$–Nb and Li$_{6.5}$–Nb phases are 0.59, and 0.52, and 0.40 eV, respectively which is close to the theoretically calculated
The calculated relaxation time, $\tau$, was in the range of $10^{-15}$ to $10^{-18}$ s which decreases with Li content and is comparable with that of Ta family. Li$_{5+2x}$La$_3$Ta$_2$Y$_x$O$_{12.25}$ The AC impedance or NMR analysis of some of the ionic conductors is mentioned here to get an idea of how the relaxation time varies in different electrolytes. In the polymer nano-composites of polyethylene oxide and lithium perchlorate, Li$^+$ ions show relaxation times of micro seconds ($10^{-6}$ s). The fastest Na ion conducting, Na$^+$-alumina, has exhibited a relaxation time $t_0$ of $10^{-12}$ s. Solid-state $^7$Li NMR analysis of Li$_{1.3}$Al$_{0.15}$Y$_{0.15}$Ti$_{1.7}$(PO$_4$)$_3$ showed a relaxation time in the order of $10^{-13}$ s. For (BiI$_3$)$_{0.4}$–(Ag$_3$PO$_4$)$_{0.6}$ electrolyte, the mobile Ag$^+$ ions showed a relaxation time of $10^{-7}$ to $10^{-11}$ s.

**Conclusions**

The transport mechanism of Li$_{5+2x}$La$_3$Nb$_2$Y$_x$O$_{12}$ ($x = 0.25$, 0.5 and 0.75) was studied in terms of their dielectric behavior at below room temperature. The bulk Li$^+$ ion conductivity of the samples increases with an increase in the Li content and follows the trend, Li$_{5.5}$La$_3$Nb$_1.75$Y$_{0.25}$O$_{12}$ (Li$_{5.5}$–Nb) < Li$_6$La$_3$Nb$_1.5$Y$_{0.5}$O$_{12}$ (Li$_6$–Nb) < Li$_{6.5}$La$_3$Nb$_1.25$Y$_{0.75}$O$_{12}$ (Li$_{6.5}$–Nb) and also follows the
cell constant trend. The Li$^+$ ion conductivity as a function of frequency was found to follow the Jonscher universal power law. The modulus plots indicate a non-Debye behavior for Li$^+$ ion relaxation. Relatively higher dielectric constants for the Nb members compared to that of the Ta members of Li$_{5+2x}$La$_x$Nb$_{2-x}$Y$_2$O$_{12}$ (M = Nb, Ta) could be explained in terms of the higher dipole moments and greater Li$^+$ ion mobility. The calculated relaxation time, $\tau$ was $10^{-18}$, $10^{-17}$ and $10^{-15}$ s and the activation energy observed was 0.59, 0.52 and 0.40 eV for Li$_{5.5}$-Nb, Li$_6$-Nb and Li$_{6.5}$-Nb, respectively, of Li$_{5+2x}$La$_x$Nb$_{2-x}$Y$_2$O$_{12}$ from the relaxation profile.

Acknowledgements

The funding support from the Natural Sciences and Engineering Research Council of Canada (NSERC) through Discovery Grants (DG) program is acknowledged for this work.

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