IONIC CONDUCTIVITY AND ELECTRICAL PROPERTIES OF CARBOXYMETHYL CELLULOSE - NH₄Cl SOLID POLYMER ELECTROLYTES

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Abstract
In this present work, carboxymethyl cellulose (CMC) – ammonium chloride (NH₄Cl) solid polymer electrolyte (SPE) films were prepared by solution casting method. The ionic conductivity and electrical properties of SPE films were investigated using Electrical Impedance Spectroscopy. SPE film containing 16 wt. % NH₄Cl exhibited the highest ionic conductivity of 1.43 x 10⁻³ S/cm at ambient temperature, 303K. The temperature dependence SPE films showed an Arrhenius-type relation where the regression values obtained from the log conductivity versus reciprocal temperature is close to unity (R² = 1). The electrical properties have been measured as a function of frequency of εᵣ, εᵢ, Mᵣ, Mᵢ, shown a non-Debye type behavior. 

Keywords: Solid polymer electrolyte, carboxymethyl cellulose, ammonium chloride, ionic conductivity, electrical properties

1. Introduction
Solid polymer electrolytes (SPEs) have received widespread attention due to their technological applications in rechargeable batteries, super capacitors, fuel cells, gas sensors and electrochromic power sources [1]. SPEs have several been reported to exhibit several advantages including favorable electrical, optical and mechanical properties, ease of fabrication in the thin film form and ability to form effective electrode–electrolyte contacts [2].
PVA, PVC and PEO are some examples of biodegradable polymers used to prepare SPE films [2-7]. One of the promising materials which are biodegradable is carboxymethyl cellulose (CMC).

CMC deserves a special attention due to its superior mechanical, good electrical properties and easy processability. CMC contains a hydrophobic polysaccharide backbone and it shows CMC one of hydrophilic carboxyl groups. Hence, it is showing water-soluble features [8].

This research focuses to employ CMC as a polymer host due to numerous advantages as previously reported [8]. CMC was doped with NH$_4$Cl as ammonium ions, NH$_4^+$ were believed to be responsible for the ionic conduction in SPE [9]. SPE films were characterized using electrical impedance spectroscopy (EIS) to investigate the ionic conductivity and electrical behavior in a temperature range of 303 K – 353 K.

2. Materials and method

2.0 g of CMC (Acros Organic Co.) was dissolved in 100 ml distilled water at room temperature. A varied amount of NH$_4$Cl (Sigma Aldrich) in weight percentage (0 – 20 wt. %) was added into the CMC solution. The mixed solution was stirred until complete dissolution was achieved and cast into petri dishes before left to dry at room temperature for film to form.

Electrical impedance spectroscopy (EIS) was utilized to determine the ionic conductivity of SPE films in the frequency range of 50 Hz to 1MHz. The range of the frequency has been used because of the bulk resistance, $R_b$ exists at that range. Electrical measurements were performed on a HIOKI 3532-50 LCR Hi-Tester by using temperatures between 303K and 353K. SPE film was placed between the blocking stainless steel electrodes of a sample cell holder connected EIS. The bulk resistance ($R_b$) of SPE film was obtained from the plot of imaginary part (-$Z_i$) versus real part ($Z_r$) of impedance. The ionic conductivity ($\sigma$) was calculated from the equation,

$$\sigma = \frac{t}{R_b A}$$

(1)
where, $t$ is the thickness and $A$ (cm$^2$) is the electrode-electrolyte contact area of SPE films.

3. Results and Discussion

3.1. Salt dependence of ionic conductivity

Figure 1 illustrates the ionic conductivity plot of CMC – NH$_4$Cl SPE films. It is shown that the decreases ionic conductivity starts when NH$_4$Cl concentration is beyond 16 wt. %. Therefore, the highest conductivity measured is $1.43 \times 10^{-3}$ S/cm at 16 wt. % of NH$_4$Cl. The increase of ionic conductivity with the addition of NH$_4$Cl concentration can be attributed to ion dissociation between host polymer and ionic dopant. As the ionic dopant content increases, more protons are supplied due to the dissociation of the ionic dopant into the polymer matrix

$$
\sigma_a \exp \left( \frac{-E_a}{kT} \right)
$$

(2)

Above 16 wt. % NH$_4$Cl the conductivity decreases and can be attributed to the reassociation of the ions into neutral aggregates [11, 12].

Figure 2 shows the Arrhenius plot for CMC – NH$_4$Cl solid polymer electrolyte films. Refer to Figure 2, log conductivity of SPE films was observed to obey Arrhenius behavior when $R^2 \approx 1$. Regression value, $R^2$ is shown in Table 1.

Arrhenius plot describes the thermally activated process for SPEs. The relationship between conductivity and temperatures of SPE illustrated by slopes of log $\sigma$ versus 1000/T. According to [6], the ionic conductivity increases with increasing of temperatures due to mobility of carrier of ion concentration.

From the slope of Arrhenius plot, the activation energy, $E_a$, can be calculated [13],
where \( \sigma_o \) is the exponential factor, \( k \) is the Boltzmann constant and \( T \) is the absolute temperature.

Refer to Table 1 tabulates the regression value and activation energy. From calculation of activation energy, the results were found to increase and then decrease at 12 wt. % in \( \text{NH}_4\text{Cl} \) concentration. Comparing with the values of ionic conductivity, it is inversely proportional with activation energy. According to [7, 12], the value of ionic conductivity values do not show any sudden jump indicating the fact that the polymer electrolyte exhibit a completely amorphous structure. The activation energy is a combination of energy of charge carrier creation (defect formation) and the energy of ion migration [14]. Therefore, it suggests that the value of \( E_a \) is due to the energy that is required to provide a conductive condition for the migration of ions [3].

![Arrhenius plot for CMC - \( \text{NH}_4\text{Cl} \) solid polymer electrolyte films](image)

**Table 1. The regression values and activation energy of CMC – \( \text{NH}_4\text{Cl} \) solid polymer electrolyte films.**

<table>
<thead>
<tr>
<th>Sample</th>
<th>Regression value, ( R^2 )</th>
<th>Activation energy, ( E_a \times 10^{-3} ) (eV)</th>
</tr>
</thead>
<tbody>
<tr>
<td>( \text{NH}_4\text{Cl} – 0 )</td>
<td>0.99</td>
<td>3.74</td>
</tr>
<tr>
<td>( \text{NH}_4\text{Cl} – 2 )</td>
<td>0.98</td>
<td>3.37</td>
</tr>
<tr>
<td>( \text{NH}_4\text{Cl} – 6 )</td>
<td>0.93</td>
<td>3.28</td>
</tr>
<tr>
<td>( \text{NH}_4\text{Cl} – 8 )</td>
<td>0.98</td>
<td>3.17</td>
</tr>
<tr>
<td>( \text{NH}_4\text{Cl} – 12 )</td>
<td>0.98</td>
<td>2.00</td>
</tr>
<tr>
<td>( \text{NH}_4\text{Cl} – 14 )</td>
<td>0.99</td>
<td>1.82</td>
</tr>
</tbody>
</table>
### 3.2. Dielectric studies

The dielectric study was performed on the highest ionic conductivity SPE film (16 wt. % NH₄Cl). Figure 3 shows the plot of frequency dependence of dielectric constant, $\varepsilon_r$, while figure 4 shows the plot of frequency dependence of dielectric loss, $\varepsilon_i$, at selected temperature. Dielectric constant, $\varepsilon_r$, is known as stored charge in a material while dielectric loss, $\varepsilon_i$, is a measure of energy losses to move ions when the polarity of electric field turns rapidly.

The $\varepsilon_r$ and $\varepsilon_i$ can be defined as [8, 15-17],

$$
\varepsilon_r(\omega) = \frac{Z_i}{\omega C_o (Z_r^2 + Z_i^2)} \quad (3)
$$

$$
\varepsilon_i(\omega) = \frac{Z_r}{\omega C_o (Z_r^2 + Z_i^2)} \quad (4)
$$

where $C_o = \frac{\varepsilon_o A}{t}$, $\varepsilon_o$ is the permittivity of free space. $\omega = 2\pi f$ and $f$ is frequency.

From Figure 3, it can be observed that there were no appreciable relaxation peaks observed in the frequency range employed in this study. No relaxation peaks are observed could be used as the indicator to show that the increasing conductivity is mainly attributed to the increasing free mobile ions [18]. Both $\varepsilon_r$ and $\varepsilon_i$ rise sharply at low frequencies indicating that electrode polarization and space charge effects have occurred confirming non-Debye dependence [19]. Otherwise, at high frequencies, the occurrence of periodic reversal of the electric field so fast that there is no excess ion diffusion in the direction of the field. Polarization due to charge accumulation decreases, leading to the observed decrease in dielectric constant and dielectric loss [4, 20].

### 3.3. Modulus studies

Modulus studies highlight the bulk dielectric behavior and suppress the effects of electrode polarization. The real electric modulus, $M_r$, in Figure 5 and imaginary modulus, $M_i$, in Figure 6 can be analyzed using equations,

$$
M_r = \frac{\varepsilon_r}{(\varepsilon_r^2 + \varepsilon_i^2)} \quad (5)
$$

$$
M_i = \frac{\varepsilon_i}{(\varepsilon_r^2 + \varepsilon_i^2)} \quad (6)
$$

The real part of electrical modulus, $M_r$, versus frequency, $\omega$, at selected temperatures is shown in Figure 5. The long tail in the plot appeared at lower frequencies indicating that the SPE film is capacitive in nature conductors [13].

| NH₄Cl – 16 | 0.98 | 1.39 |
| NH₄Cl – 18 | 0.99 | 1.71 |
| NH₄Cl – 20 | 0.99 | 2.75 |
The slightly peaking curve at higher frequencies ($>10^5$ Hz) might have been caused by the bulk effect [21].

The imaginary part of electrical modulus, $M_i$, increases with the increasing frequency, however decreases with the temperature. This indicates to the relaxation time for the proton at high temperature is shorter than at lower temperature.

Fig. 3. $\varepsilon_r$ versus $\omega$ for CMC – 16 wt. % NH$_4$Cl at selected temperatures.

Fig. 4. $\varepsilon_i$ versus $\omega$ for CMC – 16 wt. % NH$_4$Cl at selected temperatures.
4. Conclusions

The CMC - NH₄Cl solid polymer electrolytes exhibited highest ionic conductivity of $1.43 \times 10^{-3}$ (16 wt. % NH₄Cl) at ambient temperature. The conductivity arises with temperature following the Arrhenius behavior. The electrical behavior of the SPE films showed a strong dependence on frequencies and temperatures. The frequency dependence of $\varepsilon_r$, $\varepsilon_i$, $M_r$, $M_i$ were observed to be non-Debye type.

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