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Authors
Han, KS
Rajput, NN
Wei, X
et al.

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Diffusional motion of redox centers in carbonate electrolytes

Kee Sung Han,1 Nav Nidhi Rajput,2 Xiaoliang Wei,3 Wei Wang,3 Jian Zhi Hu,4 Kristin A. Persson,2 and Karl T. Mueller1,5,a)

1Environmental Molecular Sciences Laboratory, Pacific Northwest National Laboratory, Richland, Washington 99352, USA
2Environmental Energy Technologies Division, Lawrence Berkeley National Laboratory, Berkeley, California 94720, USA
3Energy and Environmental Directorate, Pacific Northwest National Laboratory, Richland, Washington 99352, USA
4Department of Chemistry, Pennsylvania State University, University Park, Pennsylvania 16802, USA
5Fundamental and Computational Sciences Directorate, Pacific Northwest National Laboratory, Richland, Washington 99352, USA

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Ferrocene (Fc) and N-(ferrocenylmethyl)-N,N-dimethyl-N-ethylammonium bistrifluoromethylsulfonimide (Fc1N112-TFSI) were dissolved in carbonate solvents and self-diffusion coefficients (D) of solutes and solvents were measured by 1H and 19F pulsed field gradient nuclear magnetic resonance (NMR) spectroscopy. The organic solvents were propylene carbonate (PC), ethyl methyl carbonate (EMC), and a ternary mixture that also includes ethylene carbonate (EC). Results from NMR studies over the temperature range of 0–50 °C and for various concentrations (0.25–1.7 M) of Fc1N112-TFSI are compared to values of D simulated with classical molecular dynamics (MD). The measured self-diffusion coefficients gradually decreased as the Fc1N112-TFSI concentration increased in all solvents. Since TFSI− has fluoromethyl groups (CF3), D_{TFSI} could be measured separately and the values found are larger than those for D_{Fc1N112} in all samples measured. The EC, PC, and EMC have the same D in the neat solvent mixture and when Fc is dissolved in EC/PC/EMC at a concentration of 0.2 M, probably due to the interactions between common carbonyl structures within EC, PC, and EMC. A difference in D (D_{PC} < D_{EC} < D_{EMC}) and both a higher E for translational motion and higher effective viscosity for PC in the mixture containing Fc1N112-TFSI reflect the interaction between PC and Fc1N112+, which is a relatively stronger interaction than that between Fc1N112+ and other solvent species. In the EC/PC/EMC solution that is saturated with Fc1N112-TFSI, we find that D_{PC} = D_{EC} = D_{EMC} and Fc1N112+ and all components of the EC/PC/EMC solution have the same E for translational motion, while the ratio D_{EC/PC/EMC}/D_{Fc1N112} is approximately 3. These results reflect the lack of available free volume for independent diffusion in the saturated solution. The Fc1N112+ transference numbers lie around 0.4 and increase slightly as the temperature is increased in the PC and EMC solvents. The trends observed for D from simulations are in good agreement with experimental results and provide molecular level understanding of the solvation structure of Fc1N112-TFSI dissolved in EC/PC/EMC. © 2014 AIP Publishing LLC.

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INTRODUCTION

Ferrocene-derived materials have become a focus of interest recently for applications as active materials: they are used as redox centers utilizing the Fc/Fc+ reactions for improving the performance of Li ion batteries.1,2 and for non-aqueous redox flow battery materials.3,4 The flow battery is one of the alternative battery technologies that has more design flexibility than traditional batteries, exhibiting enhanced energy storage capacity, battery response, and cycle/calendar life with easy scale up processes that decrease the cost per unit energy.5 The flow battery uses a flowing catholyte and anolyte, which are stored separately and externally, to generate electricity at the electrodes. Therefore, the concentrations of the active materials in the electrolyte and their diffusivity are key factors for performance. The Fc/Fc+ redox reaction of ferrocene-derived materials has revealed a diffusion-controlled reaction at the electrodes during cell testing.3,4 The contribution of the active materials to the total ionic conductivity of the electrolyte, which is closely related to the overall current of the battery, is related to the individual diffusion coefficient of each component of the mixed electrolyte by the Nernst-Einstein equation.5,7 Therefore, it is valuable to measure the individual diffusion coefficient of each component of the mixed electrolyte as a potential guide to optimize the electrolyte and gain an increased molecular level understanding of battery performance. Pulsed field gradient nuclear magnetic resonance (PFG-NMR) experiments have been used to determine the diffusion coefficient of liquids5,6,8,9 and the contributions of each ion to the total ionic conductivity5,7, and to study the interactions between the solutes and solvents or...
between the diffusant and the matrix resulting in lower diffusion coefficients.9-14 To obtain microscopic understanding of the observed trends in diffusion coefficient found using pulsed field gradient (PFG) NMR experiments, we have performed classical MD simulations.

In this study, the diffusion coefficients of major components of active center/electrolyte mixtures were measured by \(^1\)H and \(^{19}\)F PFG-NMR as a function of concentration in the temperature range of 0–50 °C. The components under study were ferrocene, Fc (Scheme 1); Fc1N112\(^+\) and TFSI\(^-\) in N-(ferrocenylmethyl)-N,N-dimethyl-N-ethylammonium bistri fluoromethylsulfoneimide, Fc1N112-TFSI (Scheme 1); and the solvents EC, PC, and EMC either in the solutions alone (single solvents PC and EMC, as EC is a solid below 40 °C) or within a ternary mixture of EC/PC/EMC (with a weight ratio of 4:1:5, reflecting typical flow battery concentrations). The solvent-solute interactions are interpreted from the measured diffusion coefficients and solvation structure obtained from MD simulations. The results show that the diffusion of the solvent slows with an increase in Fc1N112-TFSI concentration due to the interaction between the Fc1N112\(^+\) and the solvent. However, when Fc is dissolved in EC/PC/EMC at a concentration of 0.2 M, only a subtle change is found in both the diffusion coefficient and the activation energy for translational motion. At concentrations below the saturation point of Fc1N112-TFSI in the ternary solvent mixture, PC molecules in the solution, on average, interact with Fc1N112\(^+\) more strongly than do EC and EMC as shown by the decreased \(D\) of PC relative to that of EC and EMC. The Fc1N112\(^+\) transference number remains constant (∼0.4) over the temperature range in the EC/PC/EMC mixed solvent while it gradually increases with temperature in PC and in EMC.

### EXPERIMENTAL

#### Sample preparations

To improve low ferrocene solubility in the electrolyte (0.2 M in the EC/PC/EMC solvent studied here), a ferrocene-based ionic liquid compound (Fc1N112-TFSI, shown in Scheme 1) was prepared. Starting from (dimethylaminomethyl)ferrocene (FcN11), the Fc1N112-TFSI was synthesized via a nucleophilic substitution reaction utilizing bromoethane to yield the intermediate compound, dimethyl ethyl ferrocenylmethyl ammonium bromide (Fc1N112-Br), followed by anion exchange with TFSI\(^-\) to afford the Fc1N112-TFSI at an overall two-step yield of 91%. The experimental details of the preparation can be found elsewhere.\(^4\)

By virtue of the structural modification, the resulting Fc1N112-TFSI shows a dramatically enhanced solubility (up to 1.7 M) in the EC/PC/EMC solvent system.

#### NMR measurements

Diffusion coefficients \((D)\) of the redox centers and the organic solvent(s) for neat PC and EMC and the ternary mixture EC/PC/EMC (4:1:5 wt.) with and without the Fc or Fc1N112-TFSI redox center were measured by \(^1\)H and \(^{19}\)F PFG simulated echo NMR spectroscopy. Since resonances from EC, PC, EMC, and Fc1N112\(^+\) are separated in one-dimensional \(^1\)H spectra, separate diffusion coefficients could be measured (see Figure S1 in the supplementary material).\(^{29}\) \(^{19}\)F PFG experiments were utilized to follow the dynamics of the TFSI\(^-\) species. Experiments were performed at Larmor frequencies of 599.82 and 564.3 MHz for \(^1\)H and \(^{19}\)F, respectively, over the temperature range of 0 to 50 °C using a 14.1 T (600 MHz \(^1\)H) NMR spectrometer (Agilent) equipped with a 5 mm z-gradient probe (Doty Scientific), which can generate a maximum gradient strength of approximately 31 T/m. The echo heights, \(S(g)\), recorded as a function of gradient strength, \(g\), were fitted with the Stejskal-Tanner equation,

\[
S(g) = S(0)e^{-[D(\gamma g)^2/(\Delta - \delta/3)]},
\]

where \(S(g)\) and \(S(0)\) are the echo heights at the gradient strengths of \(g\) and 0, respectively; \(D\) is the diffusion coefficient; \(\gamma\) is the gyromagnetic ratio of \(^1\)H or \(^{19}\)F; \(\Delta\) is the time interval between the two gradient pulses (also called the diffusion delay); and \(\delta\) is the length of time of the gradient pulse. The gradient strength was varied over 15 equal steps and the maximum gradient strength was chosen according to the echo height at the maximum gradient (to ensure proper sampling of the data for chosen experimental conditions). The sample convection artifact induced by the temperature gradient along the z-direction was found to be negligible in a reference sample of water. However, large convection artifacts were introduced from these organic solvents (see Figure S2 in the supplementary material).\(^{29}\) Therefore, to suppress the convection artifacts a modified bipolar pulsed field gradient stimulated echo pulse sequence (vendor-supplied pulse sequence Dbpsste_cc in VNMRIJ version 3.2 on the Agilent system) was used, which has convection compensation gradients in the center of the two pairs of bipolar gradient pulses.\(^{15}\) Timings for the 90° pulses and the two delays, \(\Delta\) and \(\delta\), were 5.5 \(\mu\)s, 20 ms, and 2 ms, respectively. Below 0 °C, the diffusion coefficient obtained from the neat EC/PC/EMC showed a dependence on the diffusion time \(\Delta\) that indicates restricted diffusion (see Figure S3 in the supplementary material).\(^{29}\) The restricted diffusion is due to the presence of a solid EC/liquid PC/EMC binary phase that exists below approximately 0 °C due to the higher melting point of EC (Table I). This behavior has been previously observed in binary mixed electrolytes\(^{17}\) and predicted in ternary and quaternary mixed electrolytes.\(^{18}\) Therefore, PFG-NMR can be used to determine the solid/liquid
MD simulations

Molecular dynamics simulations of the bulk solutions were performed using the GROMACS MD simulation package version 4.5.3. The force field parameters and other information to run the simulations in GROMACS are available in the supplementary material. The initial configurations were obtained by randomly distributing the molecules in a cubic box, periodic in XYZ directions using PACKMOL. All systems studied here were initially minimized with respect to energy using steepest descent and then using conjugate gradient to relax the strained contacts in the initial configurations. The systems were equilibrated in the isothermal-isobaric ensemble (constant NPT) using the Berendsen barostat to maintain the pressure of 1 bar with a time constant of 2 ps. The production runs of 3 ns were then obtained in the canonical ensemble (NVT) using an improved velocity-rescaling algorithm proposed by Parrinello et al. with a time constant of 0.1 ps over the temperature range of 0 to 50 °C. The Lennard-Jones interactions were truncated at a cut-off of 1.2 nm. The particle-mesh Ewald (PME) method was used to handle long-range electrostatic interactions with a real space cut-off of 1.0 nm, and the Verlet leapfrog integration algorithm was used with a time step of 0.01 fs to solve the equations of motion. Long-range corrections were applied for both energy and pressure. Force fields for all the molecules were nonpolarizable. For Fc1N112+, the force field was developed by using a combination of electronic structure calculations to obtain the partial charges and parameters from the work of Lopes et al. For TFSI−, the force field was taken from Kelkar and Maginn. The Lennard-Jones interactions were truncated at a cut-off of 1.2 nm. The Verlet leapfrog integration algorithm was used with a time step of 0.01 fs to solve the equations of motion. Long-range corrections were applied for both energy and pressure. Force fields for all the molecules were nonpolarizable. For Fc1N112+, the force field was developed by using a combination of electronic structure calculations to obtain the partial charges and parameters from the work of Lopes et al. For TFSI−, the force field was taken from Kelkar and Maginn.

RESULTS AND DISCUSSION

The temperature dependent diffusion coefficients of neat PC and EMC, and the three components EC, PC, and EMC in the ternary mixture of EC/PC/EMC with a weight ratio 4:1:5 are shown in Figure 1. Both PFG and MD simulations indicate that the diffusion coefficients of all components within the neat EC/PC/EMC mixture lie between those of neat PC and EMC over the temperature range utilized in this study. The trends observed from simulations are in agreement with experimental results; however, the simulated dynamics are slower than the experimental results. The slower dynamics from simulations when compared to experiments is a common limitation of non-polarizable force fields, in particular when studying ionic liquids. While the physical properties of EC, PC, and EMC are quite different (Table I), the three components of the mixed organic solvent have the same diffusion coefficients, $D_{EC} = D_{PC} = D_{EMC}$ within experimental error, presumably due to the interactions between EC, PC, and EMC through a common carbonyl structure, O-CO-O, as previously suggested for the binary mixed electrolytes EC/DEC and PC/DEC. Usually, the error in diffusion measurement is less than 5%. A larger error in the measurement is introduced by the background 1H signal from water that was used for gradient coil cooling in this experiment only. The average standard deviation is 6% ± 5% and the maximum error is 11%. For all subsequent measurements, the background 1H signal was removed by using air instead of water for cooling. Then the error reduced to less than 5% and is similar to the size of the symbols (see Figure 2). The diffusion coefficients of the three carbonate components when Fc is dissolved at 0.2 M in EC/PC/EMC are also the same within experimental error (with an average standard deviation of about 4%) and slightly decreased from values found from the neat EC/PC/EMC mixture. The interactions between EC, PC, and EMC and the transalional motion of EC/PC/EMC are unaffected by addition of ferrocene, and 0.2 M Fc is the maximum amount that can be dissolved in this mixture. 

![Figure 1. Diffusion coefficients of neat PC and EMC and of the three components of the ternary mixture of EC/PC/EMC with and without the 0.2 M Fc redox center as a function of inverse temperature from MD simulations and PFG-NMR measurements.](image-url)
ternary solvent. The determination of the Fc diffusion coefficient for 0.2 M Fc EC/PC/EMC is not possible because of the resonance overlap with the solvent peak arising from EMC.

The temperature dependent diffusion coefficients of neat PC, 0.25 M and 1.6 M (sat.) Fc1N112-TFSI dissolved in PC, and of neat EMC, 0.25 M and 1.4 M (sat.) Fc1N112-TFSI dissolved in EMC are shown in Figures 2(a) and 2(b), respectively. The diffusion coefficients of solvent species decreased by addition of Fc1N112-TFSI and that of solvent and solute, i.e., the values for $D_{\text{Fc1N112}^+}$ and $D_{\text{TFSI}^-}$ decreased further by increasing the concentration of Fc1N112-TFSI. The Fc1N112$^+$ cation diffuses slower than the TFSI$^-$ anion basically due to the larger molecular size. The cross sectional lengths of Fc1N112$^+$ and TFSI$^-$ are 1.0 and 0.65 nm, respectively, which were calculated by density functional theory (DFT)-based geometry optimization.\(^4\) The ratio of the diffusion coefficient for the ionic liquid species ($D_{\text{Fc1N112}^+/D_{\text{TFSI}^-}}$) would be 1 if Fc1N112$^+$ and TFSI$^-$ are moving together as an ion pair or estimated to be approximately 0.65 in these solutions when Fc1N112$^+$ and TFSI$^-$ are completely dissociated without any interaction with the solvent molecules based on the Stokes-Einstein theory of diffusion, where $D = k_B T f / 6\pi \eta r_s$, $k_B$ is the Boltzmann constant, $T$ is the absolute temperature, $\eta$ is the viscosity, and $r_s$ is hydrodynamic radius of the diffusing molecule. Therefore, the strength of ion association between Fc1N112$^+$ and TFSI$^-$ can be estimated from the $D_{\text{Fc1N112}^+/D_{\text{TFSI}^-}}$ values.\(^6\) The interaction between TFSI$^-$ and solvent molecules is very weak (to be discussed below). The measured $D_{\text{Fc1N112}^+/D_{\text{TFSI}^-}}$ is 0.59 in the 0.25 M Fc1N112-TFSI dissolved in PC and is $0.67 \sim 0.74$ in other samples (see Figure S4 in supplementary material).\(^29\) These results demonstrate that Fc1N112$^+$ interacts with PC strongly relative to the interactions between Fc1N112$^+$ and other solvent molecules. This result is also confirmed from MD simulations (Table II) where we observed an average higher coordination number of PC around Fc1N112$^+$ in 0.25 M Fc1N112-TFSI dissolved in PC as compared to the coordination number of EMC around the same ion for 0.25 M Fc1N112-TFSI dissolved in EMC. The larger $D_{\text{Fc1N112}^+/D_{\text{TFSI}^-}}$ (~ 0.74) for the component ions in PC and in EMC saturated with Fc1N112-TFSI shows the enhancement of ion association between Fc1N112$^+$ and TFSI$^-$ at the higher Fc1N112-TFSI concentrations. Samples with 0.25, 0.85, and 1.7 M Fc1N112-TFSI dissolved in PC/EMC show $D_{\text{Fc1N112}^+/D_{\text{TFSI}^-}} \approx 0.65$, suggesting that the PC/EMC mixed solvent is better for the solvation of Fc1N112-TFSI.

Diffusion coefficients of the ions and solvent molecules when Fc1N112-TFSI is dissolved in EC/PC/EMC with various concentrations (0.25, 0.85 and 1.7 M) of Fc1N112-TFSI were determined as a function of temperature and are plotted in Figure 3(a). The diffusion coefficients at 323 K are compared as a function of concentration in Figure 3(b). These data indicate that the diffusion coefficients of all components (EC, PC and EMC as well as the Fc1N112$^+$ and TFSI$^-$) gradually decreased as Fc1N112-TFSI concentrations increased. As predicted by the Stokes-Einstein relation and DFT calculation, the measured ratio of $D_{\text{Fc1N112}^+/D_{\text{TFSI}^-}} \approx 0.65$ shows that the Fc1N112$^+$ cation diffuses slower than the TFSI$^-$ anion because Fc1N112$^+$ is larger than TFSI$^-$ while there is a little variation of the $D_{\text{Fc1N112}^+/D_{\text{TFSI}^-}}$ due to the interaction between the molecules in the solutions. All molecules in the

![Graph](image)

**Table II.** Coordination numbers from molecular dynamics simulations of PC and EMC around Fe, N, and the terminal carbon of the ethyl chain of Fc1N112$^+$ in 0.25 M Fc1N112-TFSI dissolved in PC and in 0.25 M Fc1N112-TFSI dissolved in EMC.

<table>
<thead>
<tr>
<th>Sites</th>
<th>PC</th>
<th>EMC</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fe</td>
<td>9.4</td>
<td>7.8</td>
</tr>
<tr>
<td>N</td>
<td>7.4</td>
<td>6.3</td>
</tr>
<tr>
<td>Terminal carbon</td>
<td>5.2</td>
<td>4.4</td>
</tr>
</tbody>
</table>
This effect has been observed previously in similar systems; the diffusion coefficient of EC is smaller than DMC or DEC when LiPF$_6$ is dissolved in EC/DMC/DEC due to relatively the diffusion coefficient of EC is smaller than DMC or DEC. This effect has been observed previously in similar systems; the diffusion coefficient of EC is smaller than DMC or DEC when LiPF$_6$ is dissolved in EC/DMC/DEC due to relatively strong interactions between the solvent molecules and solute.\textsuperscript{6,10,11}

This suggests that the interaction between PC and Fc$_{112}$TFSI is stronger at lower temperatures, similar to the behavior observed due to the interaction between EC and Li$^+$ when LiPF$_6$ is dissolved in a EC/DMC/DEC mixture.\textsuperscript{14}

When the ternary mixture EC/PC/EMC is saturated with Fc$_{112}$TFSI, the diffusion coefficients of the three components were equal ($D_{EC} = D_{PC} = D_{EMC}$). We attribute this to the lack of available free space (volume) for independent diffusion because the diffusion coefficient is determined by the available free space in the highly concentrated samples rather than the interactions between the solutes and the solvent as in the moderately concentrated samples.\textsuperscript{10} The diffusion ratio of Fc$_{112}^{12+}$ to the solvent species, $D_{Fc_{112}/D_{solvent}}$ is approximately 0.5 for the 0.25 and 0.85 M Fc$_{112}$TFSI solutions and approximately 0.3 for the solution saturated with Fc$_{112}$TFSI. The normalized diffusion of Fc$_{112}^{12+}$ with respect to the solvent is relatively faster in the lower concentration solutions compared to the saturated sample. This behavior suggests that more free space for independent diffusion is available in the lower concentration solutions.

When the ternary mixture EC/PC/EMC is saturated with Fc$_{112}$TFSI, the diffusion coefficients of the three components were equal ($D_{EC} = D_{PC} = D_{EMC}$). We attribute this to the lack of available free space (volume) for independent diffusion because the diffusion coefficient is determined by the available free space in the highly concentrated samples rather than the interactions between the solutes and the solvent as in the moderately concentrated samples.\textsuperscript{10} The diffusion ratio of Fc$_{112}^{12+}$ to the solvent species, $D_{Fc_{112}/D_{solvent}}$ is approximately 0.5 for the 0.25 and 0.85 M Fc$_{112}$TFSI solutions and approximately 0.3 for the solution saturated with Fc$_{112}$TFSI. The normalized diffusion of Fc$_{112}^{12+}$ with respect to the solvent is relatively faster in the lower concentration solutions compared to the saturated sample. This behavior suggests that more free space for independent diffusion is available in the lower concentration solutions.
coefficients using the Arrhenius equation, $D = D_0 \exp\left( -\frac{E_a}{RT} \right)$, where $R$ is the gas constant (1.985 cal/mol·K) and $T$ is absolute temperature. The calculated values for $E_a$ gradually increase as the Fc1N112-TFSI concentration increases, which indicates that free space controls the diffusion process in the higher concentration samples. The increase in $E_a$ results from additional thermal energy needed to activate the translational motion in more concentrated samples due to the higher viscosity. Both the activation energies and the diffusion coefficients of a neat and 0.2 M Fc solution in EC/PC/EMC are similar, indicating that the addition of 0.2 M Fc does not affect the translational motion of the solvent and there is no measurable interaction between the Fc and the solvent molecules. The $E_a$ values for translation of Fc1N112$^+$ and PC are the same within experimental error and are larger than those of EC and EMC in 0.25 and 0.85 M Fc1N12-TFSI solutions in mixed EC/PC/EMC solvents because the interaction between the PC and Fc1N112$^+$ is stronger relative to the interaction between Fc1N12$^+$ and either EC or EMC. All components of the solutions in EC/PC/EMC that are saturated with Fc1N112-TFSI have the same $E_a$. These data again suggest that the free volume, which was available for the independent diffusion at lower concentrations, is not available at the higher concentrations of Fc1N112-TFSI.

To investigate the hypothesis that stronger ionic interactions between the PC and Fc1N112-TFSI and the lack of the free space for independent diffusion in the concentrated samples explain the observed diffusion data, the effective viscosity ($\eta^*$) was calculated for each solution at 25°C using measured diffusion coefficients and the viscosity of PC and EMC (Table I). For the calculation it was assumed that these solutions hold to the Stokes-Einstein theory of diffusion. The actual diffusion coefficient varied with the friction factor $f$, which is related to the size of and the interactions that affect the diffusing molecule. Therefore, the friction factor needs to be modified accordingly. In our solutions, there is no justification to apply different hydrodynamic radii to the species when they occur within different samples. So it was assumed that all samples have the same $r_s$ regardless of the Fc1N112-TFSI concentrations. Then the diffusion coefficient of a single solvent, either PC or EMC (labeled $D_s$), and that of the solvent molecules of mixed solvents (labeled $D_{x,\text{solvent}}$) are given by

$$D_x = \frac{k_B T}{6\pi \eta_x r_s^3} \quad \text{and} \quad D_{x,\text{solvent}} = \frac{k_B T}{6\pi \eta_{x,\text{solvent}} r_s^3},$$

where $r_s = r_s^{x,\text{solvent}}$ and $\eta^*$ will be

$$\eta^* = \eta_x \frac{D_x}{D_{x,\text{solvent}}},$$

where $x = \text{PC or EMC}$. The effective viscosities (Figure 5) show that the ternary mixed solvent EC/PC/EMC and the sample with 0.2 M Fc dissolved in EC/PC/EMC hold to the Stokes-Einstein relation. Mixed liquids have unique viscosities which eliminates the need for a different $\eta^*$ for each component. The $\eta^*$ values calculated from PC and EMC are the same in the neat ternary mixture and when the concentration of Fc dissolved in EC/PC/EMC is 0.2 M, and lie between the measured viscosities of neat EMC and neat PC (Table I). The $\eta^*$ of 0.2 M Fc dissolved in EC/PC/EMC increases moderately from that of neat EMC but is smaller than the viscosity of neat PC. The $\eta^*$ is increased with increasing Fc1N112-TFSI concentration. For 0.25 and 0.85 M Fc1N112-TFSI dissolved in EC/PC/EMC, the $\eta^*$ calculated from PC is larger than that from EMC. This observation again presents experimental evidence that the interaction between PC and Fc1N112$^+$ is stronger relative to that between EMC and Fc1N112$^+$. The Fc1N112-TFSI saturated solution in the ternary mixture shows nearly the same $\eta^*$ for PC and EMC. As noted above, this is consistent with a reduction of the available spaces to distinguish the relative strength of the interaction between solvent molecules and Fc1N112$^+$. The variations of $D$, $E_a$, and $\eta^*$ in the EC/PC/EMC solutions due to the presence of Fc1N112-TFSI show that the interaction between Fc1N112$^+$ and solvent molecules...
enhances the solubility of active material further from the EC/PC/EMC solution with Fc, which shows negligible effects.

To analyze the Fc1N112⁺ contribution to the total ionic conductivity of the mixture, the transference numbers of Fc1N112⁺ were calculated using the measured diffusion coefficients,

\[
t_{\text{Fc1N112}} = \frac{n_{\text{Fc1N112}} D_{\text{Fc}}}{n_{\text{Fc1N112}} D_{\text{Fc1N112}} + n_{\text{TFSI}} D_{\text{TFSI}}},
\]

where \(n_{\text{Fc1N112}}\) and \(n_{\text{TFSI}}\) are the number density (molar ratios) of Fc1N112⁺ and TFSI⁻, respectively. The solvent molecules are not included in Eq. (4) due to their negligible charge. The results plotted as a function of Fc1N112-TFSI concentrations in Figure 6 reveal that the Fc1N112⁺ transference number increases slightly with temperature in PC, EMC, and when Fc1N112-TFSI is dissolved at 0.25 M in EC/PC/EMC, while it is nearly the same in 0.85 and 1.7 M (saturated) Fc1N112-TFSI dissolved in EC/PC/EMC. The interaction between Fc1N112⁺ and PC results in a relatively small \(t_{\text{Fc1N112}}\) when Fc1N112-TFSI is dissolved at a concentration of 0.25 M in PC.

Finally, it is insightful to compare the diffusion coefficients of the components within the solution to the values of \(D\) computed with MD simulations. The diffusion coefficients of Fc1N112⁺, TFSI⁻, EC, PC, and EMC in 0.85 M Fc1N112-TFSI dissolved in EC/PC/EMC, computed by MD simulations, are presented in Figure 7 (0.25 and 1.7 M data plotted in Figure S6 in the supplementary material).29 The trends in diffusion coefficient from both experiments and MD simulations indicate that the dynamics of all the species Fc1N112⁺, TFSI⁻, EC, PC, and EMC in the solution decrease with a decrease in temperature (although there is some scatter within experimental error in the simulated data). We observed that \(D_{\text{Fc1N112}} < D_{\text{TFSI}}\) for the temperature range considered in this work which is due to the larger size of Fc1N112⁺ and the stronger interaction of the cation with solvent molecules as compared to TFSI⁻. This trend is consistent at 0.25 M and 1.7 M concentrations as well.

**CONCLUSIONS**

The temperature dependent diffusion coefficients, \(D\) of the neat PC, EMC, and components of the ternary mixture of EC/PC/EMC with and without ferrocene or Fc1N112-TFSI were measured by \(^1\)H and \(^{19}\)F PFG-NMR and computed using classical MD simulations. The data show decreasing \(D\) with the addition of Fc and Fc1N112-TFSI. The diffusion coefficients gradually decreased as the Fc1N112-TFSI concentrations increased. The effective viscosity, which is related to the strength of interactions between the components of the solvent and Fc1N112-TFSI, calculated using the
Stokes-Einstein equation from the $D$ and viscosity of EMC and PC shows the ternary mixture of EC/PC/EMC holds to the Stokes-Einstein relation of diffusion. The higher $E_a$ and $\eta^*$ of PC when FeC1N112-TFSI is dissolved in EC/PC/EMC suggest that the interaction between PC and FeC1N112$^+$ is stronger relative to other components of the solvent. This suggestion is also consistent with the smaller $D_{FeCN112}$ in PC than in EMC from 0.25 M FeC1N112-TFSI dissolved in PC and in EMC. In addition, the large coordination number of PC (from simulations) as compared to EMC confirms the stronger interaction between PC and FeC1N112$^+$. All components of the FeC1N112-TFSI saturated solution of EC/PC/EMC have the same $E_a$ values for translational motion as well as similar $\eta^*$ due to the lack of available free space for independent diffusion. The interaction between FeC1N112$^+$ and solvent molecules enhances the solubility of active material, FeC1N112$, by a factor of 9 from the EC/PC/EMC solution with FeC. The transferance numbers of FeC1N112$^+$ lies around 0.4 and gradually increased with temperature except in the EC/PC/EMC solution with high FeC1N112-TFSI concentrations. This result suggests that the solvent systems that contain less PC are more favorable to the FeC/Fe$^+$ redox reactions in an actual flow battery cell.

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