

Article

Synthesis, Multinuclear NMR Characterization and Dynamic Property of Organic-Inorganic Hybrid Electrolyte Membrane Based on Alkoxysilane and Poly(oxyalkylene) Diamine

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Abstract: Organic-inorganic hybrid electrolyte membranes based on poly(propylene glycol)-*block*-poly(ethylene glycol)-*block*-poly(propylene glycol) bis(2-aminopropyl ether) complexed with LiClO₄ via the co-condensation of tetraethoxysilane (TEOS) and 3-(triethoxysilyl)propyl isocyanate have been prepared and characterized. A variety of techniques such as differential scanning calorimetry (DSC), Fourier transform infrared (FTIR) spectroscopy, alternating current (AC) impedance and solid-state nuclear magnetic resonance (NMR) spectroscopy are performed to elucidate the relationship between the structural and dynamic properties of the hybrid electrolyte and the ion mobility. A VTF (Vogel-Tamman-Fulcher)-like temperature dependence of ionic conductivity is observed for all the compositions studied, implying that the diffusion of charge carriers is assisted by the segmental motions of the polymer chains. A maximum ionic conductivity value of $5.3 \times 10^{-5} \text{ Scm}^{-1}$ is obtained at 30 °C. Solid-state NMR results provide a microscopic view of the effects of salt concentrations on the dynamic behavior of the polymer chains.

Keywords: organic-inorganic hybrid electrolyte; ionic conductivity; poly(oxyalkylene) diamine; segmental motion

Figure S1. DSC thermograms of TIE(600)-Z hybrid electrolytes with various [O]/[Li] ratios, where Z = (a) ∞ , (b) 32, (c) 24, (d) 16 and (e) 8.

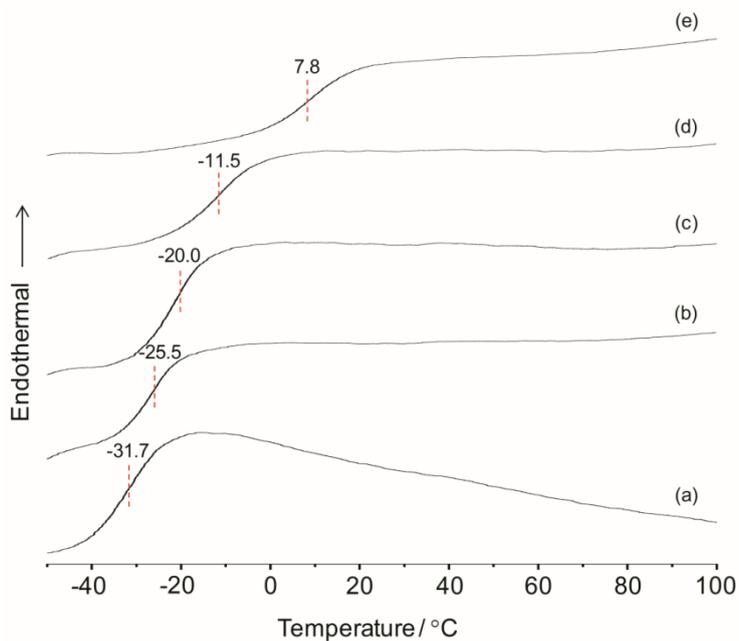


Figure S2. Temperature dependence of ionic conductivity of TIE(600)-Z hybrid electrolytes with Z = (a) 32, (b) 24, (c) 16 and (d) 8.

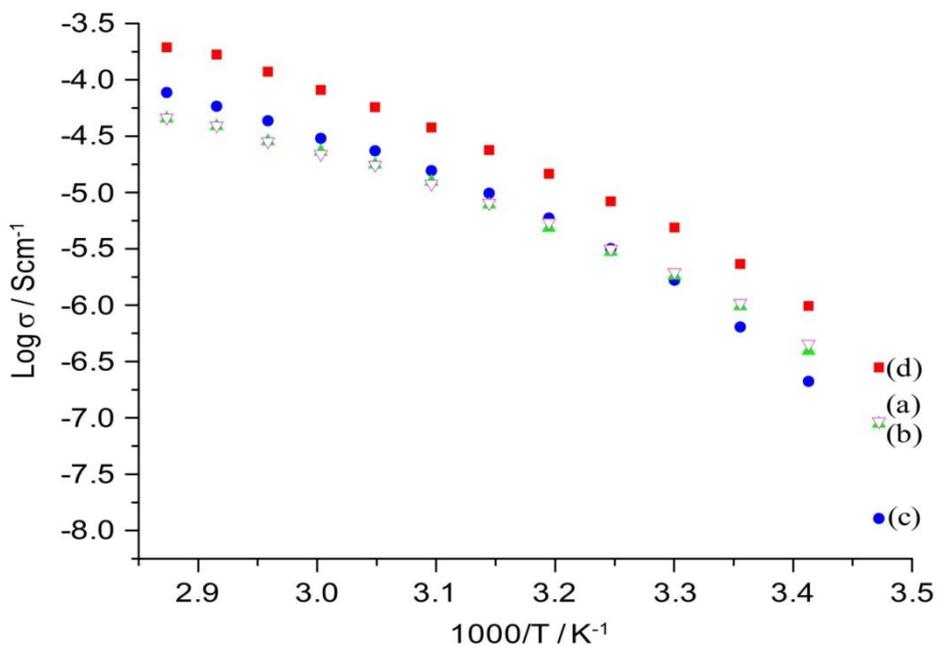
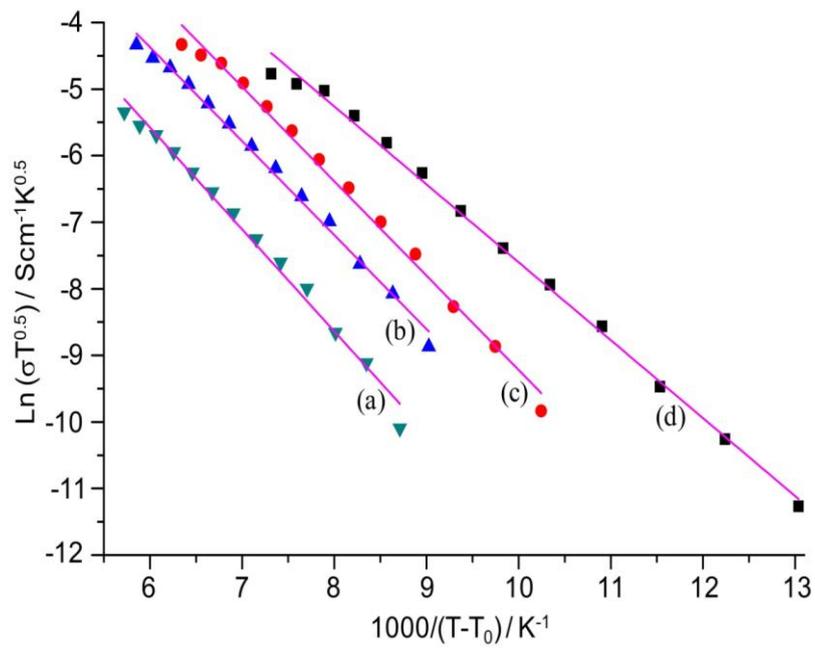


Figure S3. VTF curve fitting results of TIE(2000)-Z hybrid electrolytes with Z = (a) 32, (b) 24, (c) 16 and (d) 8.



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