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Design and synthesis of a single ion conducting block copolymer electrolyte with multifunctionality for lithium ion batteries

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Abstract

A novel single ion conducting block copolymer electrolyte (SI-co-PE) for applications in lithium-ion batteries is presented. The block copolymer is made of alternative ethylene oxide (EO) and aromatic segments to tune the glass transition temperature (T_g), the mechanical strength and the porosity of the material for achieving high electrochemical performance of lithium-ion batteries. Upon blending with a PVDF-HFP binder via a solution cast method, a gel SI-co-PE membrane with high ionic conductivity, a wide electrochemical window and a high lithium transference number was obtained. Excellent electrochemical stability and battery performance at both room temperature and 80 °C with various charge/discharge rates were demonstrated. The study underscores the fundamental importance of polymer electrolyte microstructures in battery performance enhancement and suggests ways to

improve the design of new electrolyte materials for development of better battery devices with a long cycle life.

Keywords: lithium ion battery, gel single ion conducting polymer electrolyte, block copolymer, battery performance.

1. Introduction

Gel polymer electrolytes (GPEs) containing liquid dual-ion inorganic lithium salts in low T_g flexible polymer matrices have made remarkable advances in the last decade and have been widely used in a variety of commercial applications, most notably in portable electronic devices.¹⁻⁸ GPEs exhibit many advantages over solid polymer electrolytes (SPEs) with better compatibility with electrodes in the charge-discharge cycling processes and substantially higher ion conductivity because of the introduction of organic solvents into polymer matrices, which reduce interfacial resistance and facilitate ion transport. Compared with liquid electrolytes, GPEs usually display a wider electrochemical window and high thermal stability.⁹ Nevertheless, there still remain several key issues that need to be resolved for the GPE materials to compete with the conventional liquid electrolytes to achieve even broader applicability while maintaining substantially improved safety.¹⁰ First, a high lithium ion transfer number close to unity would significantly raise the coulomb efficiency of battery devices.^{11,12} Second, solution retention capacity must be enhanced for stable device performance. In principle, a high ionic conductivity requires a high carrier concentration.¹³ For a GPE with a dual-ion lithium salt in a polymer matrix to achieve high ion conductivity, appropriate porosity including the

pore size, the volume fraction and the interconnectivity of pore domains in the polymer membrane is required to accommodate a large number of charge carriers.^{14,15} In practice, a porous membrane with an average pore size of more than 1 μm usually causes an electrolyte leakage, leading to a deterioration of solution retention capacity, a degradation of ionic conductivity and a compromise of device safety.¹⁶ Indeed, it has been a formidable challenge to obtain a GPE with both a high ionic conductivity and a strong mechanical strength because a membrane with high porosity weakens its mechanical strength.¹⁷ Furthermore, a strong concentration gradient inherent to dual-ion based electrolytes during battery operation leads to dendrite growth and a low lithium ion transference number (<0.3), which ultimately limits power delivery.^{18,19}

Gel single ion conducting polymer electrolytes (Gel-SIPEs) have been found to offer an effective solution to the concentration gradient problem with a high lithium ion transference number close to unity.²⁰⁻²² Several studies have demonstrated that lithium ion batteries assembled with gel-SIPE membranes display excellent electrochemical stability and device performance.²³⁻²⁸ In this paper, we report design and synthesis of a gel single ion conducting block copolymer electrolyte (Gel-SI-co-PE) with a structure shown in Figure 1. The Gel-SI-co-PE membranes were fabricated using PVDF-HFP as a binder. All the membranes display high ion transference numbers close to unity. Three essential properties, i.e. ion conductivity, solution retention capacity and mechanical strength, were optimized simultaneously through tuning the ratio of the aliphatic and the aromatic segments of the polymers to

achieve high battery performance.²⁹ The two chemically dissimilar polymers, labelled as PEEIA and LiPEEPSI in Figure 1, are polymerized to form a block copolymer covalently bonded end-to-end. This architecture gives rise to local segregation or “microphase separation”, resulting in formation of microporous domains to absorb a sufficient amount of solvent and to effectively prevent solution leakage from occurring with a pore diameter of less than 1 μm ³⁰ as well as construction of continuous ion conducting channels upon formation of micropores.³¹ Finally, the charge delocalized diphenylsulfonimide (DPSI) anions³² were impregnated to the block copolymer backbone to minimize the concentration polarization effect upon charge/discharge usually observed in dual-ion based electrolytes. Lithium ions are thus attached to the immobilized polyanions through a weak electrostatic interaction with high mobility.³³⁻³⁹ The membrane of the SI-co-PE complex was fabricated with a PVDF-HFP binder via a solution cast method and the thickness and flexibility of the resulting film were entirely tuneable. The membrane displays a low glass transition temperature as well as a high melting temperature. A macroporous morphology with an average pore size of ca. 200 nm and excellent mechanical strength of 27 MPa tensile stress was obtained. In addition, the membrane exhibits high electrochemical stability up to 4.3 V of oxidation voltage, high ionic conductivity of $3.9 \times 10^{-4} \text{ S cm}^{-1}$ at 25 °C and $1.2 \times 10^{-3} \text{ S cm}^{-1}$ at 80 °C and a high lithium ion transference number close to unity. Finally, excellent cyclic performance of the lithium battery assembled with the SI-co-PE based membrane at both an ambient temperature and an elevated temperature was demonstrated.

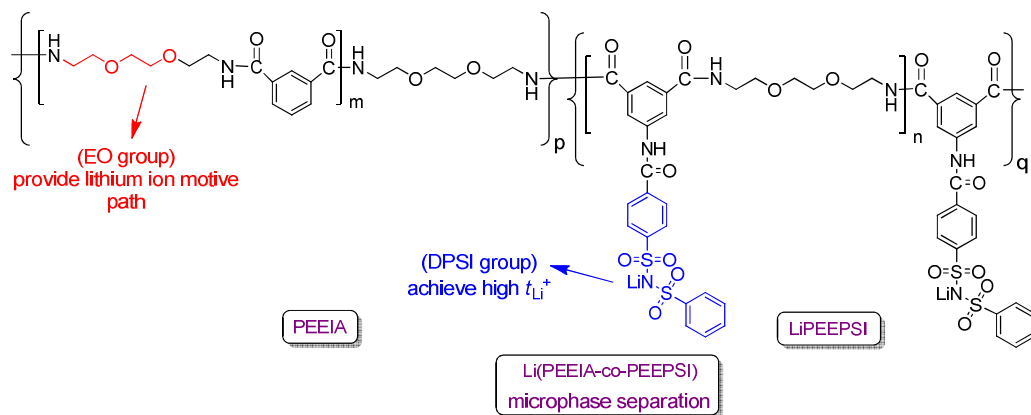


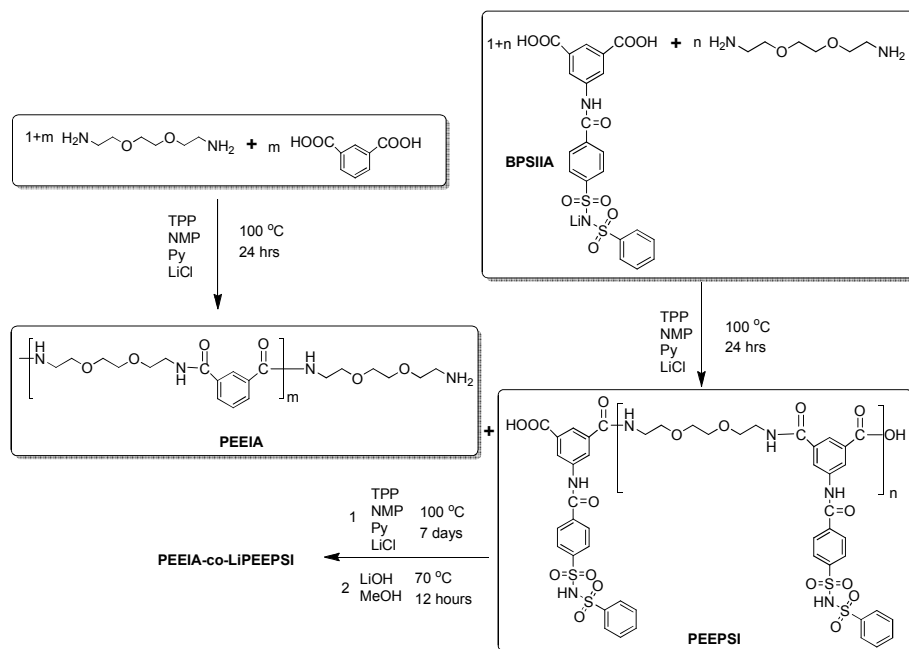
Figure 1. Chemical structure of the single ion conducting block copolymer electrolyte PEEIA-co-LiPEEPSI.

2. Experimental

2.1 Materials

P-toluenesulfonamide ($\geq 98\%$) (Sigma-Aldrich), benzenesulfonchloride (99%) (Sigma-Aldrich), $KMnO_4$ (GCE), lithium hydroxide monohydrate (Sigma-Aldrich), 2,2'-(ethylenedioxy)bis(ethylamine) (98%) (Sigma-Aldrich) and PVDF-HFP (Sigma-Aldrich) were used as received. Dimethyl-5-aminoisophthalate (Sigma-Aldrich), calcium chloride (GCE) and isophthalic acid (99%) (Sigma-Aldrich) were dried overnight in a vacuum oven at $80\text{ }^\circ\text{C}$ before use. Pyridine (Alfa-Aesar) was dried over KOH, N-methyl-2-pyrrolidone (NMP) (VWR) was dried over phosphorus (V) oxide. Triphenylphosphine (TPP) (Alfa-Aesar), analytical grade acetone, ethylene carbonate (EC) (Sigma-Aldrich), methanol, tetrahydrofuran (THF) (RCI labscan), and propylene carbonate (PC) (Sigma-Aldrich) were used as received without further purification. Cambridge Isotope Laboratories deuterated dimethyl sulfoxide (DMSO- d_6) solvent was used for the NMR tests.

2.2 Synthesis of the PEEIA-co-LiPEEPSI



Scheme 1: Polymerization scheme of PEEIA, PEEPSI to form PEEIA-co-LiPEEPSI via a polycondensation reaction.

Synthesis of the homopolymers. The precursor, bis(phenylsulfonyl imide isophthalate amide (BPSIIA), was synthesized following the procedure described in our previous work (Supporting Information).⁴⁰ The homopolymers, poly((2,2'-ethylenedioxy) bis(ethylamine) isophthalic acid amide) (PEEIA) and lithium poly((2,2'-ethylenedioxy)bis(ethylamine) bis(phenylsulfonyl imide) isophthalate amide) (PEEPSI), were separately synthesized using the procedure described in Ref. 26. Isophthalic acid (1.1255 g, 6.667 mmol) and 2,2'-(ethylenedioxy) bis(ethylamine) (1.0374 g, 7 mmol) were added into a mixture of 1.60 g of lithium chloride, 5 ml of TPP, 20 mL of NMP and 15 ml of pyridine for the synthesis of PEEIA. BPSIIA (1.1514 g, 3 mmol) and 2,2'-(ethylenedioxy) bis(ethylamine) (0.4234

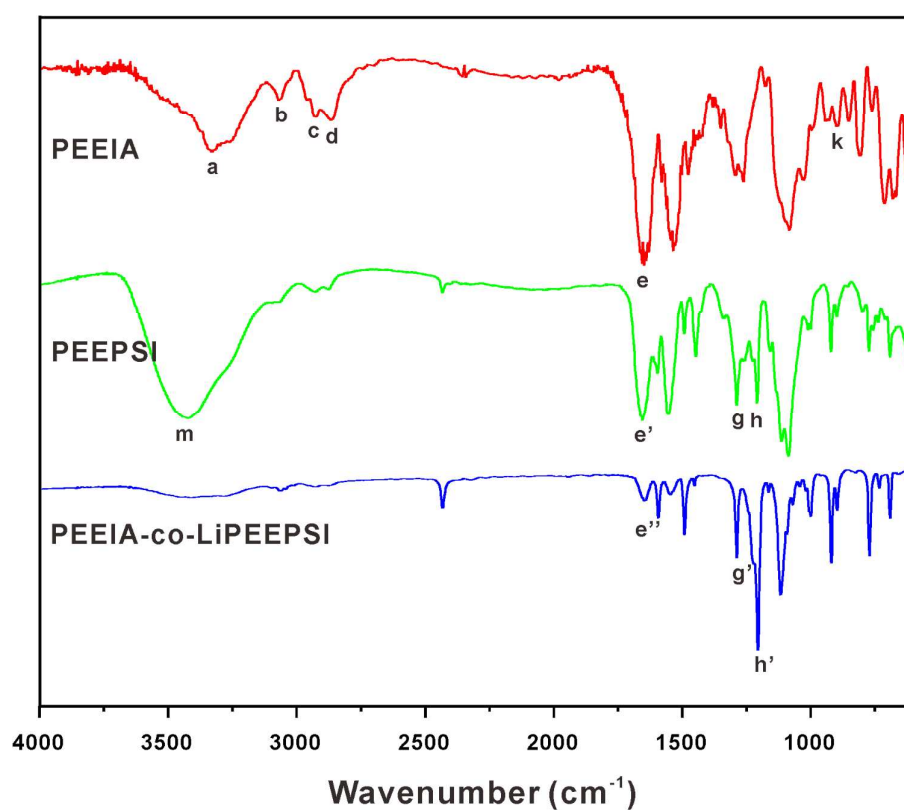
g, 2.857 mmol) were added into a mixture of 1.60 g of lithium chloride, 5 ml of TPP, 20 ml of NMP and 15 ml of pyridine for the synthesis of PEEPSI. The reactions were first kept at 100 °C for 12 hours under argon atmosphere overnight. After that, an excess of additional 10 % mole of 2,2'-(ethylenedioxy)bis(ethylamine) and BPSIIA were added into the reaction mixtures, respectively. These moieties were left to reflux for another 5 hours. The hot mixtures were separately poured into 100 ml of acetone with stirring and washed repeatedly with acetone until fine powders were obtained. The lithiation of PEEPSI was carried out by an acid-base neutralization reaction with LiOH·H₂O in methanol at 70 °C for 12 hours. Upon solvent removal, LiPEEPSI was dried at 80 °C for 24 hours.

Synthesis of the block copolymer. PEEIA (0.1380 g, 0.0105 mmol) and LiPEEPSI (0.5000 g, 0.0105 mmol) were added into a mixture of 0.16 g of lithium chloride, 0.7 ml of TPP, 2 ml of NMP and 1.5 ml of pyridine in a glove box. The reaction was conducted at 100 °C under argon atmosphere for a week. After that, the mixture was poured into 60 ml of acetone to produce a pale pink precipitate. The polymer was washed continuously with acetone. The block copolymer (PEEIA-co-LiPEEPSI) was collected via vacuum filtration and dried in a vacuum oven at 80 °C for 24 hours. The molecular weights of the homopolymers and block copolymer were characterized by gel permeation chromatography (GPC) (Table 1).

Table 1. Molecular weights and PDI of PEEIA, PEEPSI and PEEIA-co-PEEPSI.

	M_n	M_w	PDI
PEEIA	13,600	31,416	2.31
PEEPSI	49,271	87,512	1.77
PEEIA-co-PEEPSI	57,284	111,224	1.94

2.3 Characterizations

**Figure 2.** The FTIR spectra of the PEEIA, PEEPSI and PEEIA-co-LiPEEPSI.

The FTIR spectra of PEEIA, PEEPSI and PEEIA-co-PEEPSI are shown in Figure 2. The characteristic bands of PEEIA at 3300 cm^{-1} (a) and 893 cm^{-1} (k) indicate a presence of primary amine at the terminals of PEEIA. The broad peak of PEEPSI at around 3475 cm^{-1} (m) is assigned to the O-H bond stretch arising from the

COOH groups at the terminals of PEEPSI. The peaks at 1649 cm^{-1} (e), 1654 cm^{-1} (e') and 1649 cm^{-1} (e'') are assigned to amide C=O stretching of PEEIA, PEEPSI and PEEIA-co-PEEPSI, respectively. The peaks at 1339 cm^{-1} (g) and 1158 cm^{-1} (h) of PEEPSI and 1325 cm^{-1} (g') and 1165 cm^{-1} (h') of PEEIA-co-PEEPSI reflect the O=S=O asymmetric stretching and symmetric stretching, respectively.

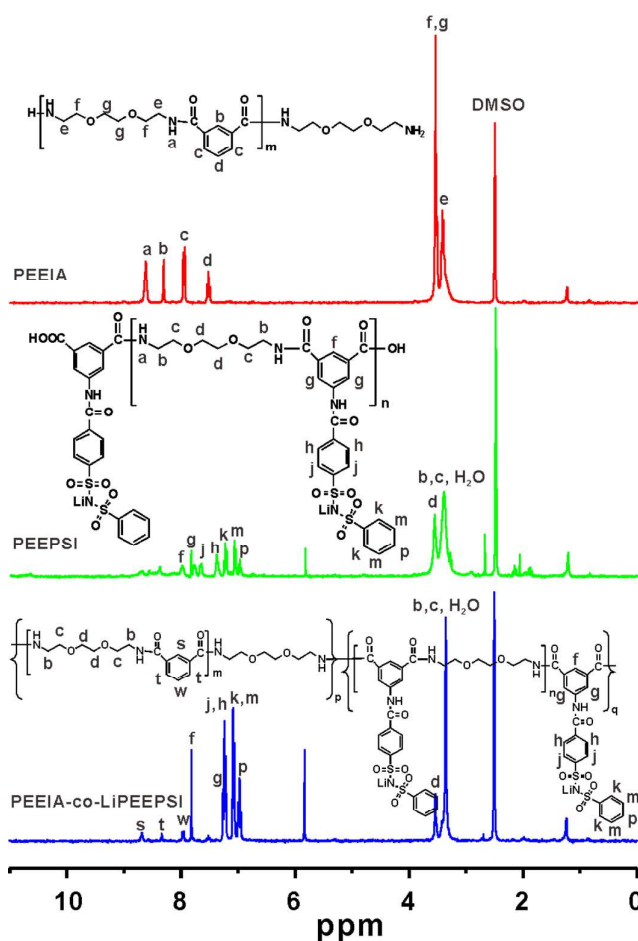


Figure 3. The ^1H NMR spectra of PEEIA PEEPSI and PEEIA-co-LiPEEPSI.

The ^1H NMR spectra of PEEIA, PEEPSI and PEEIA-co-LiPEEPSI are shown in Figure 3. The spectrum of PEEIA shows clearly defined peaks at 7.52 ppm, 7.96 ppm 8.31 ppm and a broader signal at 8.62 ppm whose integration ratio corresponds

well with the number of protons in the aromatic ring and the amide bond. The signals at 3.4 - 3.5 ppm correspond to the protons in 2,2'-(ethylenedioxy)bis(ethylamine). For the spectrum of PEEIA, the signals for hydrogen (H_{f-p}) on the BPSIIA segments are clearly identified and the signals at 3.4 - 3.5 ppm, corresponding to the protons in 2,2'-(ethylenedioxy)bis(ethylamine), are readily visible. All the proton signals of PEEIA and PEEPSI are observed in the spectrum of the PEEIA-co-LiPEEPSI block copolymer. Comparing the 1H NMR spectra of PEEIA-co-PEEPSI and the homopolymers, the proton signals of the aromatic segments on PEEIA (H_{b-d}) and on the aromatic ring in PEEPSI (H_{f-p}) shift slightly to the lower frequencies upon copolymerization to form PEEIA-co-PEEPSI.

2.4 Preparation of single-ion conducting polymer electrolyte membranes (SIPEMs)

The SIPE membranes were fabricated using PVDF-HFP as a binder via a solution cast method without using a pore inducing agent. PVDF-HFP and PEEIA-co-LiPEEPSI with an equivalent mass ratio were dissolved in DMF at 80 °C to form a homogenous solution. The composite polymer was then casted onto a glass petri dish, which was kept at 80 °C for overnight for solvent evaporation. After that, the membrane was peeled off from the petri dish and transferred into a vacuum oven for overnight at 80 °C to remove a trace of the solvent. The dried membrane was subsequently transferred into a glove box and soaked in an EC/PC (v:v, 1:1) solvent for overnight before performing electrochemical characterization and battery performance test. The choice of organic solvent greatly influences the properties of

the electrolyte. For example, high dielectric constant of organic solvent can prompt the delocalization of the Li-ion while low viscosity of organic solvent enhances Li-ion mobility in the electrolyte.⁴¹⁻⁴⁵ Here the objective of the present study is to demonstrate the concept of single ion conducting block copolymer electrolytes, rather than to select an optimal solvent, for applications in lithium ion batteries. Therefore, EC/PC solution, which has been extensively used in literature,^{46,47} was selected in the present work. For comparison, a blend membrane of PVDF-HFP and LiPEEPSI with an equivalent mass ratio was also prepared using the same procedure. The thickness of the membrane is in the range of 20 - 30 μm . The methods to test the properties and performance of the membranes are described in details in Supporting Information.

2.5 Battery test and fabrication of cathode film.

Battery performance was studied using a multichannel battery testing instrument Arbin BT-2000 in which the cell was charged and discharged at different C rates over the voltage range 3.8 - 2.5 V. The coin cell (CR2025 type) was made by sandwiching SIPE membrane between a lithium metal foil and a LiFePO_4 electrode. The cathode was prepared by casting a homogenous mixture of 75 % LiFePO_4 , 10 % carbon black, 10 % PVDF, and 5% bis(4-carboxyl benzene benzene sulfonyl) imide (CBBSI) as a supporting electrolyte in NMP solvent onto an aluminium foil. The cathode was subsequently dried in a vacuum oven at 80 $^\circ\text{C}$ overnight.

3. Results and Discussion

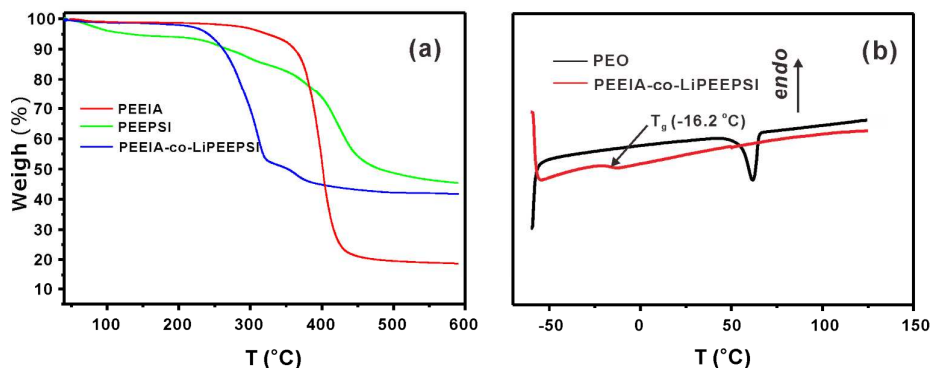


Figure 4. (a) The TG curve and (b) DSC curves of the PEEIA-co-LiPEEPSI SIPE.

The thermal stability of the PEEIA-co-LiPEEPSI membrane was measured with TGA and DSC techniques (Figure 4). The membrane remains stable until 220 °C, at which degradation occurs. The degradation temperature is substantially higher than the conventional operation temperature range of lithium ion batteries (Figure 4a). The DSC curve shown in Figure 4b indicates that PEEIA-co-LiPEEPSI exhibits a low glass transition temperature (T_g) of -16.2 °C, which is attributed to the increase of the randomness of the copolymer due to the introduction of the flexible EO leads into the PEEIA-co-LiPEEPSI copolymer.⁴⁸ The T_g is higher than that of PEO (-59 °C), likely resulting from the hydrogen bonding interaction between amide groups. The lithium ion mobility can still be comparable to that of PEO based polymer membranes at room temperature due to the low T_g of PEEIA-co-LiPEEPSI.

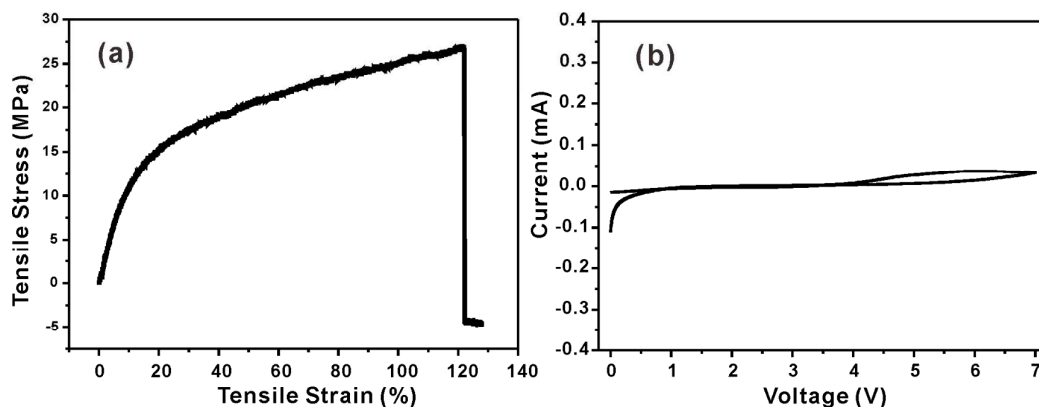


Figure 5. (a) The mechanical stability and (b) electrochemical stability of the PVDF-HFP/PEEIA-co-LiPEEPSI membrane.

Figure 5 shows the mechanical and electrochemical stabilities of the PVDF-HFP/PEEIA-co-LiPEEPSI membrane. The PVDF-HFP/PEEIA-co-LiPEEPSI membrane displays an excellent mechanical strength of 27 MPa (Figure 5a), substantially higher than the corresponding values (ca. 20 MPa) of most PVDF-HFP blended SIPE membranes.³⁸ This result indicates that the PVDF-HFP/PEEIA-co-LiPEEPSI membrane is both mechanically and dimensionally stable, which is highly beneficial for prolonging the cyclic life of batteries. The electrochemical stability of the PVDF-HFP/PEEIA-co-LiPEEPSI membrane was determined via measurement of cyclic voltammetry (CV) on a Li|PVDF-HFP/PEEIA-co-LiPEEPSI|Stainless steel (SS) cell with the CV curve shown in Figure 5b.⁴⁹ The high electrochemical stability up to 4.3 V suggests that the PVDF-HFP/PEEIA-co-LiPEEPSI membrane might be useful for batteries that need a wider electrochemical window than what can be offered by conventional liquid electrolytes. We further measured the lithium ion transference number of the

PVDF-HFP/PEEIA-co-LiPEEPSI membrane (t_{Li^+}) in a Li|PVDF-HFP/PEEIA-co-LiPEEPSI|Li symmetric cell at room temperature using the steady-state current method (Figure S3). The measured values required to derive t_{Li^+} are summarized in Table S1. The value of t_{Li^+} of the PVDF-HFP/PEEIA-co-LiPEEPSI membrane was found to be 0.91, substantially higher than that of conventional liquid electrolytes (< 0.3) and similar to those of the reported SIPEs.²⁰

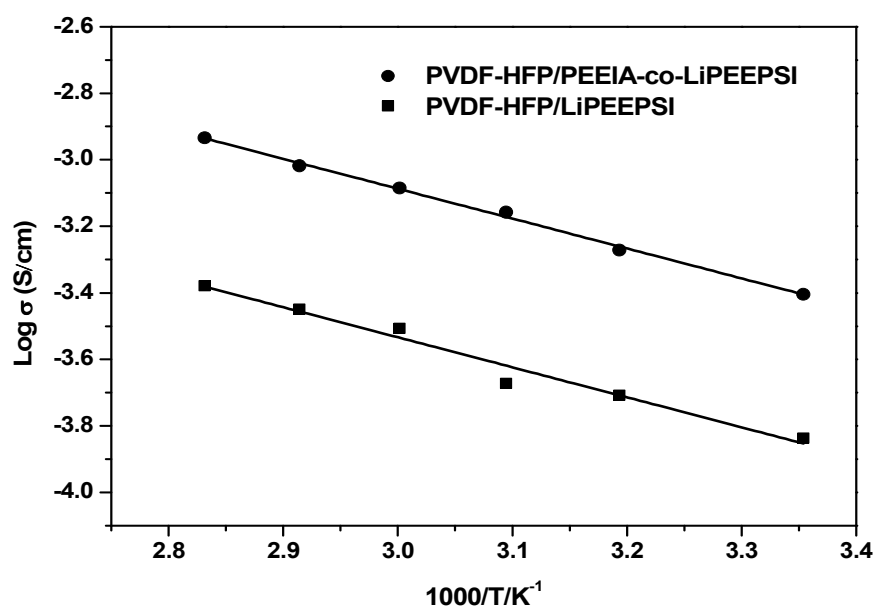


Figure 6. The ionic conductivities of the PVDF-HFP/PEEIA-co-LiPEEPSI membranes and the PVDF-HFP/ LiPEEPSI membranes vs. the inverse of temperature.

The temperature dependence of the ionic conductivity of the PVDF-HFP/PEEIA-co-LiPEEPSI membrane is depicted in Figure 6. For comparison, the conductivity of the PVDF-HFP/LiPEEPSI membrane is also shown here. The ionic conductivity of the PVDF-HFP/PEEIA-co-LiPEEPSI membrane is 3.39×10^{-4} S

cm^{-1} at room temperature and reaches to $1.16 \times 10^{-3} \text{ S cm}^{-1}$ at $80 \text{ }^\circ\text{C}$. In contrast, the ionic conductivity of the PVDF-HFP/LiPEEPSI membrane is $1.46 \times 10^{-4} \text{ S cm}^{-1}$ at room temperature and $4.18 \times 10^{-4} \text{ S cm}^{-1}$ at $80 \text{ }^\circ\text{C}$, about three times lower than the value of the PVDF-HFP/PEEIA-co-LiPEEPSI membrane. The excellent ionic conductivity of PEEIA-co-LiPEEPSI is chiefly attributed to the contribution of the EO segments in the polymer backbone with an additional contribution from the porous morphology of the material, which enables a high solvent uptake to enhance ion mobility without compromising dimensional stability of the membrane.

It should be noted here that blending the nonconductive PVDF-HFP with PEEIA-co-LiPEEPSI would inevitably affect the ion conductivity of the membrane. Rani et al. showed that for PVDF-HFP blended GPE membranes, ion conductivity increases as PVDF-HFP content decreases,⁵⁰ which is not surprising since the number of charge carriers in a membrane with a low content of PVDF-HFP is higher. We observed a similar phenomenon in our experiments and made significant effort to use only a minimum amount of PVDF-HFP in the membrane to enhance ionic conductivity while maintaining sufficient mechanical strength. The optimal ratio of PVDF-HFP to PEEIA-co-LiPEEPSI was found to be 1:1.

The high solvent uptake (141 wt. %) is evidenced from the surface SEM images of the PVDF-HFP/PEEIA-co-LiPEEPSI membrane shown in Figure 7 (a and b). Importantly, the continuous ion conducting channels are well established upon microphase separation, which is imperative for Li-ion batteries. In contrast, the

PVDF-HFP/LiPEEPSI membrane exhibits essentially no porosity as clearly shown in Figure 7c.

In general, the morphology of the blend membranes is affected by the compatibility of the polymers.²⁹ For example, the aliphatic PVDF-HFP is compatible with the aliphatic segments but not compatible with the aromatic segments in the polymer electrolyte. As a consequence, the morphology of the PVDF-HFP/LiPEEPSI membrane is not porous (Figure 7c) while the PVDF-HFP/PEEIA-co-LiPEEPSI membrane exhibits pores (Figure 7b).

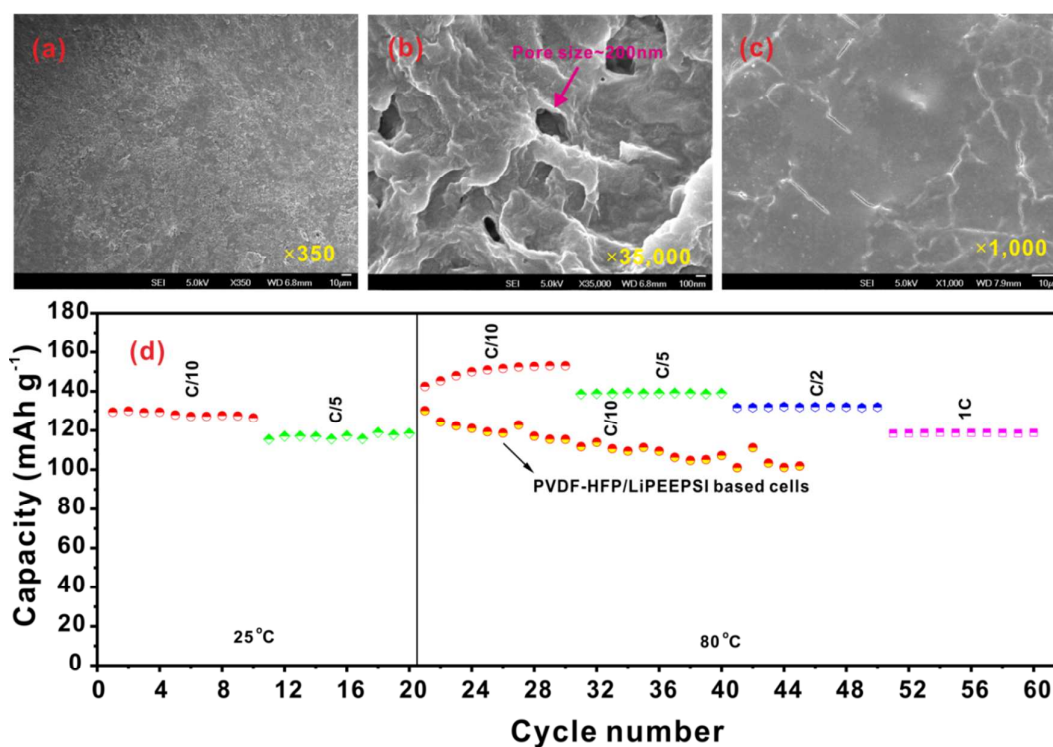


Figure 7. The SEM images of the PVDF-HFP/PEEIA-co-LiPEEPSI membrane: (a) $\times 350$, (b) $\times 35,000$; the SEM image of the surface morphology of the PVDF-HFP/LiPEEPSI membrane (c) $\times 1,000$. The cycle performance of the prototype Li|PVDF-HFP/SIPE membranes|LiFePO₄ cells (d).

The battery performance of the prepared SIPE membranes in Li-ion batteries was investigated at various C-rates at 25 °C and 80 °C, respectively. Figure 7d shows the discharge capacity vs. the cycle numbers of the battery cells. Unlike most of the SIPE based batteries,^{21,27,38} the battery of the Li|PVDF-HFP/PEEIA-co-LiPEEPSI|LiFePO₄ is operative at room temperature and the discharge capacity was found to be 130 mAh g⁻¹ at 0.1 C and 118 mAh g⁻¹ at 0.2 C. At 80 °C, the performance was significantly improved both from discharge capacity and C-rate and the discharge capacity becomes 150 mAh g⁻¹ at 0.1 C and 140 mAh g⁻¹ at 0.2 C, close to the theoretical capacity. More remarkably, the battery maintains a high discharge capacity of 130 mAh g⁻¹ at 0.5 C and 120 mAh g⁻¹ at 1 C, respectively, without obvious capacity fading after 60 cycles. In comparison, a battery equipped with the PVDF-HFP/LiPEEPSI membrane is only operative at an elevated temperature and a low C-rate of 0.1 C with a much lower average discharge capacity of 110 mAh g⁻¹ (Figure 7). In addition, the battery displays unstable cycle performance. The performance of both batteries suggests that the microstructure of SIPE membranes can profoundly influence operability of battery cells. Indeed, the superior performance of the PVDF-HFP/PEEIA-co-LiPEEPSI membrane is largely attributed to the nicely formed continuous ion transport channels arising from its macroporous morphology. In contrast, the PVDF-HFP/LiPEEPSI membrane without proper porosity prevents establishment of continuous ion transport channels and thus leads to inferior performance.

4. Conclusions

A single ion block copolymer (PEEIA-co-PEEPSI) was successfully prepared via copolymerization of the two carefully designed homopolymers, PEEIA and PEEPSI upon a lithiation reaction with $\text{LiOH}\cdot\text{H}_2\text{O}$. The homopolymers and the copolymer were characterized using FTIR, ^1H NMR, GPC, TGA and DSC spectroscopic techniques. Subsequently, a PEEIA-co-LiPEEPSI based SIPE membrane was fabricated by blending with a PVDF-HFP binder using a solution cast method. The physicochemical properties of the membrane were then systematically tested. The PEEIA-co-LiPEEPSI membrane displays good thermal stability up to 220 °C, a low T_g of -16.9 °C and a melting point higher than 125 °C. Importantly, the membrane exhibits a macroporous structure with a high solvent retention capacity up to 141 wt.%, leading to high ionic conductivity of $3.39 \times 10^{-4} \text{ S cm}^{-1}$ at room temperature and $1.16 \times 10^{-3} \text{ S cm}^{-1}$ at 80 °C. In addition, the material is electrochemically stable with a wide electrochemical window up to 4.3 V. The measured ion transference number is 0.91, substantially higher than the value found in dual-ion based electrolytes. The membrane displays an excellent mechanical strength up to 27 MPa. More remarkably, a battery assembled with the PVDF-HFP/PEEIA-co-LiPEEPSI membrane displays outstanding performance both at room temperature and 80 °C with a highly stable discharge capacity at various C-rates. Even at 1 C, the average discharge capacity still maintains at 120 mAh g^{-1} without performance fading after 60 cycles. In comparison, a battery equipped with the PVDF-HFP/LiPEEPSI membrane with essentially no proper porosity and ion transport channels in the polymer matrix displays only a modest discharge capacity of

110 mAh g⁻¹ at 80 °C and 0.1 C. The results underscore the fundamental importance of the microstructure of polymer electrolytes in battery performance enhancement and suggest ways to improve the design of new electrolyte materials for development of better battery devices with high longevity and high discharge capacity.

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