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# Enhancing lithium-ion conductivity: impact of hausmannite nanofiller on PVDF–HFP/PEG blend nanocomposite polymer electrolytes

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A new series of PVDF–HFP/PEG-based nanocomposite polymer electrolytes (NCPEs) have been fabricated using hausmannite ( $\text{Mn}_3\text{O}_4$ ) nanoparticles as the nanofiller and  $\text{LiClO}_4$  as the lithium-ion source via the solvent casting method. A pristine PVDF–HFP NCPE sample with 2 wt% nanofiller was also prepared for comparison. The  $\text{Mn}_3\text{O}_4$  nanoparticles were synthesized by the precipitation method using CTAB as a templating agent and  $\text{MnCl}_2 \cdot 4\text{H}_2\text{O}$  as the precursor. FTIR spectroscopy showed that while pristine PVDF–HFP forms a nonpolar  $\alpha$ -phase, the incorporation of salt and nanofiller induced a mixed  $\beta$  and  $\gamma$  crystal phase, indicating interaction between the matrix and additives. Surface morphology studies showed that the NCPEs had a denser surface than pristine PVDF–HFP, with no PEG spherulite formation detected in polarized optical micrographs. Electrochemical impedance spectroscopy revealed that the 2% blend NCPE exhibited the highest ion conductivity of  $3.1 \times 10^{-4} \text{ S cm}^{-1}$  at 80 °C, an order of magnitude higher than the pristine NCPE ( $5.1 \times 10^{-5} \text{ S cm}^{-1}$ ). Temperature-dependent ion conductivity followed Arrhenius behavior, indicating a thermally activated ion hopping mechanism. The dielectric relaxation peak shifted to higher frequency with increasing temperature, suggesting faster ion dynamics and improved conductivity.

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## Introduction

Lithium-ion batteries (LIBs) are widely used as a reliable power source in many portable electronic devices,<sup>1</sup> electric vehicles,<sup>2</sup> and in space applications,<sup>3</sup> due to their lightweight, low cost, high-energy density and specific capacity.<sup>4</sup> The commercial electrolytes employed in conventional LIBs are organic solvents that are highly volatile, flammable, and also associated with leakage issues.<sup>5</sup> Moreover, lithium dendrites extending between the electrodes, could lead to short-circuiting.<sup>6</sup> Polymers, as solid polymer electrolytes (SPEs), have been extensively investigated for the last several decades as safer alternatives to hazardous liquid electrolytes for LIBs.<sup>7</sup> A number of polymers, including poly(ethylene oxide) (PEO),<sup>8,9</sup> poly(vinylidene fluoride-co-hexafluoropropylene) (PVDF–HFP),<sup>10,11</sup> poly(acrylonitrile) (PAN),<sup>12</sup> poly(methyl methacrylate) *etc.*<sup>13</sup> and various polymer blends, such as PVDF/PMMA,<sup>14</sup> PEO/PEG-*b*-P(MA-POSS),<sup>9</sup> PVDF/PEO-*b*-PMMA,<sup>15</sup> and PEO/PVDF–

HFP,<sup>16</sup> *etc.* have been extensively explored as hosts in the fabrication of flexible SPEs.

To enhance the film quality, salt dissociation, and ion conductivity, non-aqueous high dielectric constant liquids are sometimes incorporated as plasticizers into the SPE compositions.<sup>17,18</sup> Other strategies that could be employed for improving ion conductivity and other electrochemical characteristics of the SPEs, include copolymerization,<sup>19,20</sup> blending,<sup>21,22</sup> network formation,<sup>23</sup> use of branched copolymers<sup>24,25</sup> and the addition of metal oxide nanoparticles, such as  $\text{Al}_2\text{O}_3$ ,<sup>26</sup>  $\text{SiO}_2$ ,<sup>27</sup> and  $\text{ZnO}$ <sup>28</sup> *etc.* to the polymer matrix.<sup>13,29,30</sup>

PVDF–HFP has been more commonly investigated as a separator in LIBs. However, a few rare studies have also focused on PVDF–HFP as the host in SPEs and nanocomposite polymer electrolytes (NCPEs) with nanofiller dispersed in the PVDF–HFP matrix.<sup>31,32</sup> As an example, Arwish *et al.*<sup>11</sup> investigated the effect of graphene oxide (GO) nanofiller on structural, electrochemical and mechanical properties of the PVDF–HFP/Pluronic based NCPEs. The electrochemical and physical properties significantly improved with the dispersion of GO nanofiller to the PVDF/Pluronic matrix and ionic conductivity reaches  $1.2 \times 10^{-6} \text{ S cm}^{-1}$  with 0.4 wt% GO addition (pristine PVDF/Pluronic SPE has an ionic conductivity of  $\approx 0.73 \times 10^{-6} \text{ S cm}^{-1}$  at room temperature with 60 wt% Pluronic) with improved mechanical properties. Prabu *et al.*<sup>33</sup> fabricated PVDF–HFP/CeO<sub>2</sub>/SiC (SiC – silicon carbide) NCPEs. The ac

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conductivity of the pristine PVDF–HFP ( $1.67 \times 10^{-4} \text{ S cm}^{-1}$ ) increased to  $1.44 \times 10^{-3} \text{ S cm}^{-1}$  for 15% CeO<sub>2</sub> and SiC filler.

Hausmannite (Mn<sub>3</sub>O<sub>4</sub>), a manganese oxide mineral (Mn<sup>2+</sup>Mn<sup>3+</sup><sub>2</sub>O<sub>4</sub>) is found as brownish black crystals or granular masses in contact metamorphic zones and high temperature hydrothermal veins.<sup>34</sup> Mn<sub>3</sub>O<sub>4</sub>, with a spinel structured unit cell, has 24 cations of both trivalent and divalent forms of Mn and 32 oxygen atoms. The following structural characteristics are responsible for their intriguing physicochemical properties: (i) oxide ions are closely packed in cubic form, (ii) Mn<sup>2+</sup> occupies the tetrahedral site, (iii) Mn<sup>3+</sup> occupies the octahedral site, and (iv) the d<sup>4</sup> state of the Mn(III) atoms in high spin configuration generates Jahn-Teller distortion.<sup>35</sup> The metal oxide interacts with the polymer chains, provides a conduction pathway along the polymer matrix, decreases the crystallinity and improves the mechanical properties of the electrolyte membranes. Hausmannite nanofiller (Mn<sub>3</sub>O<sub>4</sub>) has wide ranging applications in energy storage devices, catalysis, electronics, and sensors.<sup>36</sup>

Here, we employ hausmannite (Mn<sub>3</sub>O<sub>4</sub>) NPs as the nanofiller in PVDF–HFP/PEG blend based NCPEs with LiClO<sub>4</sub> as the source of lithium ions and with a composition of [85 wt% PVDF–HFP/15 wt% PEG/10 wt% LiClO<sub>4</sub>/*x* wt% Mn<sub>3</sub>O<sub>4</sub>], where *x* represents the weight percent of hausmannite NPs. For comparison purposes, a PVDF–HFP based pristine NCPE sample with 2 wt% nanofiller was also prepared and characterized. LiClO<sub>4</sub> was chosen because it is highly soluble in various solvents and possesses high thermal stability and relatively low hydrophobicity. The large size of ClO<sub>4</sub><sup>−</sup> anion helps in salt dissociation and reduces the crystallinity of the polymer host to facilitate lithium-ion transport.<sup>37</sup> The main aim is to evaluate the effect of hausmannite nanofiller on the ion conductivity of PVDF–HFP/PEG blend based NCPEs. A detailed account of the structural, thermal, and electrical characterization has been presented.

## Experimental

### Materials

Poly(vinylidene fluoride-co-hexafluoropropylene) (PVDF–HFP, *M<sub>w</sub>* = 400 000 g mol<sup>−1</sup>, 99.99%), and poly(ethylene glycol)

(PEG, *M<sub>w</sub>* = 8000 g mol<sup>−1</sup>, 99%) were purchased from Sigma-Aldrich. Lithium perchlorate (LiClO<sub>4</sub>, 99.98%) was obtained from Analar. Manganese chloride (MnCl<sub>2</sub>·4H<sub>2</sub>O, 97%), tetrahydro furan (THF, 99%) and cetyltrimethylammonium bromide (CTAB, 99%) were obtained from Sigma-Aldrich. PVDF–HFP, PEG and LiClO<sub>4</sub> were vacuum dried for 24 h at 50 °C before use.

### Preparation of hausmannite (Mn<sub>3</sub>O<sub>4</sub>) nanoparticles

Hausmannite nanoparticles (Mn<sub>3</sub>O<sub>4</sub>) were prepared at ambient temperature by employing the reported method.<sup>38</sup> Typically, 0.23 M MnCl<sub>2</sub>·4H<sub>2</sub>O (4.551 g, 100 mL) was introduced to the 0.45 M (100 mL) sodium hydroxide (NaOH) aqueous solution. Then, CTAB (as the templating agent) (1.38 mM) was dispersed *via* moderate magnetic stirring for 24 h. The reaction mixture was filtered and the precipitate was washed several times with distilled water. The obtained product (brownish color) was dried at 80 °C for 24 h.

### Preparation of nanocomposite polymer electrolyte films

The preparation of PVDF–HFP/PEG blend film involves dissolving 0.425 mg of PVDF–HFP in 8 mL THF followed by the addition of 0.075 mg PEG and stirring for 24 h at 45 °C. The homogeneous solution was then cast on a Teflon Petri dish and dried at ambient temperature, followed by vacuum drying for 2–3 days at 45 °C. For the preparation of nanocomposite polymer electrolytes, additional steps of adding known amounts of salt (dissolved in THF) and nanofiller (dispersed in THF and sonicated) to the polymer solution were also involved. PVDF–HFP based pristine NCPE with 2 wt% nanofiller and PVDF–HFP/PEG based blend NCPEs of various compositions (with 2, 4, and 6 wt% Mn<sub>3</sub>O<sub>4</sub> NPs) were prepared. The sample acronyms and their explanations are given in Table 1.

### Characterizations

FTIR spectra were recorded using a Bruker Tensor VERTEX 70 by KBr pellets at 4 cm<sup>−1</sup> resolution and in the 400–4000 cm<sup>−1</sup> range. X-ray diffraction (XRD) analysis was carried out on an X-ray Bruker diffractometer (Germany) in the range of 3 to 80° by diffraction of Cu K $\alpha$  radiations at a scan rate of 5° minute<sup>−1</sup>. The average crystallite size was calculated using the Debye–Scherrer formula (eqn (1)).<sup>39,40</sup>

$$D = \frac{K\lambda}{(\beta \cos \theta)} \quad (1)$$

where *D* represents the average crystallite size, *K* is a shape factor (typically around 0.9),  $\lambda$  is the X-ray wavelength (1.5406 Å for Cu K $\alpha$  radiation),  $\beta$  is the full width half maximum (FWHM) of the diffraction peak, and  $\theta$  is the Bragg diffraction angle.<sup>39,40</sup> DSC analysis was performed on a Netzsch DSC 204 F1 Phoenix (Germany) by following a heating–cooling–reheating cycle at a scan rate 10 °C min<sup>−1</sup> from −50 °C to 220 °C. TGA analysis was conducted on a Shimadzu TGA 50 analyzer with a Pt cell with a scan rate of 10 °C min<sup>−1</sup> and 20–800 °C temperature range. A scanning electron microscope (SEM), Thermo Fischer NOVA NanoSEM 450-FEISEM, and energy dispersive spectrometer (Oxford Instruments, UK) were employed to analyze the surface

Table 1 Sample acronyms and their meanings

Sample ID	Composition <sup>a</sup>
PVDF–HFP	100 wt% PVDF–HFP
PVDF–HFP/PEG	85 wt% PVDF–HFP/15 wt% PEG
2% Pristine NCPE	85 wt% PVDF–HFP/10 wt% LiClO <sub>4</sub> /2 wt% Mn <sub>3</sub> O <sub>4</sub>
2% Blend NCPE	85 wt% PVDF–HFP/15 wt% PEG/10 wt% LiClO <sub>4</sub> /2 wt% Mn <sub>3</sub> O <sub>4</sub>
4% Blend NCPE	85 wt% PVDF–HFP/15 wt% PEG/10 wt% LiClO <sub>4</sub> /4 wt% Mn <sub>3</sub> O <sub>4</sub>
6% Blend NCPE	85 wt% PVDF–HFP/15 wt% PEG/10 wt% LiClO <sub>4</sub> /6 wt% Mn <sub>3</sub> O <sub>4</sub>

<sup>a</sup> The salt and filler were added with respect to the total weight of PVDF–HFP and PEG matrix. For example, in 2% blend NCPE, PVDF–HFP and PEG had an 85 and 15 wt% ratio, respectively. 10 wt% salt was added with respect to the total weight of the blend matrix (PVDF–HFP and PEG) and 2 wt% filler was added with respect to the total weight of the blend matrix (PVDF–HFP and PEG).



structure and composition of the prepared electrolytes. All images were obtained at an accelerating voltage of 10 kV. A polarized optical microscope (POM) furnished with a Zeiss AxioCam MRc camera and linked to AxioVision Control software (Zeiss, Oberkochen, Germany) was used for surface morphology analysis. X-ray photoelectron spectroscopy (XPS) analysis was carried out with a Nexsa G2 from Thermo Scientific, UK, using a flood gun for static charge compensation and monochromatized Al-K $\alpha$  radiation (1486.6 eV) with a 400  $\mu$ m spot size. At an ultra-high vacuum of 10–8 mbar, the pass energy was set to 200 eV for the survey spectra and 50 eV for the high-resolution scans. Using Avantage V6.4.1, data collection and peak assessment were carried out. A universal testing machine (UTM- Z010, Zwick/Roell) was used to conduct the tensile test at room temperature. A testing speed of 27 mm min<sup>-1</sup> with a pre-load of 0.1 N was employed. For the measurement, a sample with a total length of 20 mm, width of grip section of 6 mm, a width of 2.02 mm, and a gauge length of 12 mm was used.

Electrochemical impedance spectroscopy (EIS) was conducted by using a dielectric spectrometer (Broadband, Germany) employing a novo-control GmbH. A length gauge (MT25B) was used for measuring the film thickness. For EIS the electrolyte film was sandwiched between the gold-plated electrodes of diameter 20 mm and 100  $\mu$ m separation distance. A frequency range of 10<sup>-2</sup>–10<sup>7</sup> Hz and 1 V potential was employed to monitor the current response. For the data acquisition, the sample was equilibrated for 20 min at 25 °C, followed by sample heating up to 80 °C under a nitrogen atmosphere in steps of 15–20 K difference.

## Results and discussion

### Hausmannite (Mn<sub>3</sub>O<sub>4</sub>) nanoparticles

Hausmannite (Mn<sub>3</sub>O<sub>4</sub>) nanoparticles were prepared following the reported literature, employing CTAB as the templating agent and MnCl<sub>2</sub>·4H<sub>2</sub>O as the precursor.<sup>38</sup> The XRD profile depicted in Fig. 1(a) shows that the prepared Mn<sub>3</sub>O<sub>4</sub> is highly crystalline, with characteristic diffraction peaks, which agrees well with JCPDS Card No. 024-0734 with a tetragonal (Hausmannite) crystal structure.<sup>41</sup> The average crystallite size of ~30 nm was calculated from the diffraction peaks using the Debye–Scherrer

approximation.<sup>42,43</sup> Fig. 1(b) depicts the SEM image of the prepared Mn<sub>3</sub>O<sub>4</sub> NPs that reveals the cuboidal shape of the particles with some degree of agglomeration.<sup>44</sup>

### Water content of the prepared nanocomposite polymer electrolytes

PVDF–HFP-based pristine and PVDF–HFP/PEG blend based NCPEs with various filler content have been fabricated and characterized for their structural, electrochemical, and dielectric behaviors. However, before carrying out any structural or electrochemical studies, we quantified the water content of the prepared polymer electrolytes. Due to the hygroscopic nature of the lithium salts, there is always a possibility of water absorption by the electrolyte films. Water traces in polymer electrolytes cause various effects, e.g., the interaction between polymer matrix and water may solvate the polymer, which results in free lithium ions and likely contributes to increased ion mobility in the hydrated system.<sup>45</sup> Additionally, water also acts as a plasticizer for polymers, thus decreasing the glass transition temperature ( $T_g$ ) of the amorphous phase, reducing crystallinity, and enhancing the side chain mobility.<sup>46,47</sup> Therefore, it is recommended to report the water content of the fabricated polymer electrolytes. As an example, Forsyth *et al.*<sup>48</sup> calculated the water content in the PAN (poly(acrylonitrile)) based electrolyte and found that the electrolyte film with 35 wt% LiTf salt had a water content of 3%, whereas the film containing 65 wt% LiTf had a water content of 6%. They suggested that water entered the electrolyte film *via* the complexation of lithium salt with the polymer matrix. Karl Fischer titration was used in the current study to estimate the water content of the fabricated NCPE films. Karl Fischer titration is a fast method for reliably quantifying both free and bound water in the sample.<sup>49</sup> The calculated water content of the prepared electrolyte films is in the range of 1% as given in Table 2.

### PVDF crystal phase and the effect of additives

PVDF is a polymorph that, based on chain conformation, self-assembles into five different crystal phases, namely,  $\alpha$ ,  $\beta$ ,  $\gamma$ ,  $\delta$ , and  $\epsilon$ . The most commonly observed  $\alpha$ -phase is electro inactive with no net dipole moment in the unit cell due to antiparallel

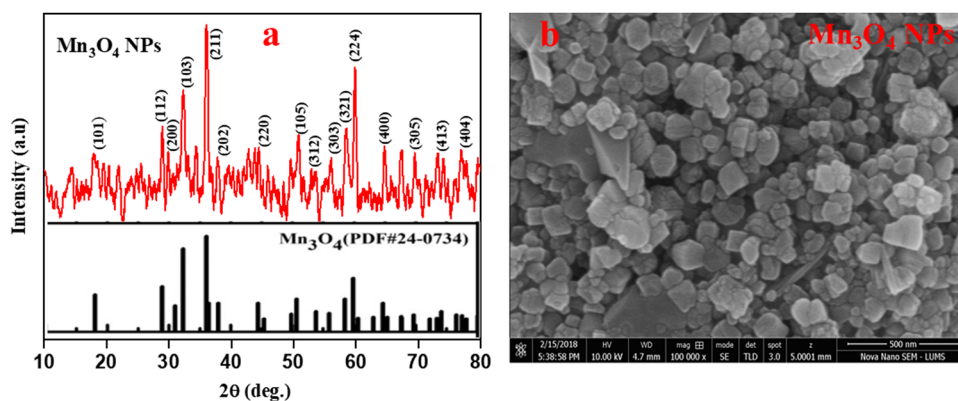


Fig. 1 (a) XRD spectra, and (b) SEM micrograph of the as-prepared Mn<sub>3</sub>O<sub>4</sub> NPs.



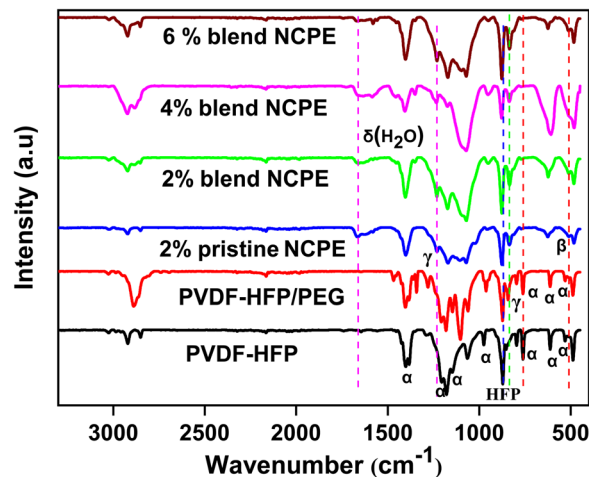
**Table 2** Water content determined by the Karl Fischer method in the fabricated films

Sample	Mass (mg)	Water content (%)
PVDF-HFP	8.1	0.23
PVDF-HFP/PEG	13.3	—
2% Pristine NCPE	11.4	0.42
2% Blend NCPE	25.0	1.00
4% Blend NCPE	30.0	1.20
6% Blend NCPE	30.0	0.38

packing of dipoles, while the  $\beta$  and  $\gamma$  phases are electroactive.<sup>50</sup> The higher content of the  $\beta$  and  $\gamma$  phase in the material is the subject of high interest for potential applications in various fields, including, sensing, electro optic devices, data storage, actuation, and energy harvesting.<sup>51</sup> PVDF is a partially crystalline linear polymer with  $[-CH_2-CF_2-]_n$  as the backbone. The structural repeat unit possess two dipoles, *i.e.*, one is due to the  $CH_2$  while the other is due to the  $CF_2$  moiety.<sup>52</sup> The PVDF crystal phase formation depends on the process employed in the sample preparation. PVDF chains generally self-assemble into a helical  $\alpha$ -phase with a chain conformation of (TGTG) (T and G, respectively, stand for *trans* and *gauche* conformations) where chains pack into an antipolar unit cell.<sup>53</sup> As depicted in Fig. 2, the FTIR spectrum of the prepared pristine PVDF-HFP film reveals the characteristic peaks of the  $\alpha$ -phase at 531, 613, 761, 796, 853, 974, 1206, and 1383  $cm^{-1}$ .<sup>50,54</sup> Furthermore, the peak at 872  $cm^{-1}$  is due to the amorphous phase of PVDF due to the incorporation of HFP.<sup>55</sup> The bands at 1383, 2923, 760, and 1401  $cm^{-1}$  correspond to symmetric, asymmetric stretching, rocking, and deformation vibration of the  $CH_2$  group, respectively.<sup>56</sup> The peaks at 1179 and 1147  $cm^{-1}$  are assigned to symmetric and antisymmetric stretching of the  $CF_2$  group, respectively, while the peak at 613  $cm^{-1}$  is due to  $CF_2$  bending and 794  $cm^{-1}$  is due to  $CF_3$  stretching.<sup>57</sup> Blending PVDF-HFP with PEG shifted some of the electro inactive non-polar  $\alpha$ -phase peaks to a more polar electroactive phase ( $\gamma$ -phase) along with the emergence of some new peaks due to PEG. Thus, the peak at 853  $cm^{-1}$  shifted to 843  $cm^{-1}$  due to a change in TGTG conformation to  $T_3GT_3G$  for the  $\gamma$ -phase.<sup>57</sup> The intense peak observed at 1103  $cm^{-1}$  in the PVDF-HFP/PEG blend is due to C-O-C bending.<sup>58</sup> Thus, the FTIR spectrum confirms the formation of a mixed  $\alpha$  and  $\gamma$  phase in the blend matrix.

The FTIR spectra of the pristine and blend NCPEs reveal the formation of mixed electroactive  $\beta$  and  $\gamma$  phases with no traces of the  $\alpha$ -phase peaks. Thus, the peak appearing at 510  $cm^{-1}$  is due to the  $\beta$  phase while the peaks at 831 and 1231  $cm^{-1}$  are due to the formation of the  $\gamma$  phase of PVDF-HFP. This could be attributed to the ion-dipole and dipole-dipole interactions in the fabricated composite solid electrolytes<sup>50,59</sup> Similar observation of phase shifting from the  $\alpha$  to  $\beta$  and  $\gamma$  phase was also reported by Mallikarjun *et al.*<sup>60</sup> and stated that mixed states reveal the fulfillment of the amorphous nature of the SPE.

The ion-dipole interaction of the C-O-C moiety of the PEG chain with salt in blend NCPEs is evident from the peak shift from  $\approx 1103$   $cm^{-1}$  in the PVDF-HFP/PEG blend to  $\approx 1070$   $cm^{-1}$  in blend NCPEs. Thus, in addition to acting as a plasticizer,



**Fig. 2** FTIR spectra of pristine PVDF-HFP, PVDF-HFP/PEG blend, pristine NCPE, and blend NCPEs with different concentrations of  $Mn_3O_4$  NPs.

PEG also interacts with  $Li^+$  cation, thus, assisting in salt dissociation and increasing the charge carrier concentration in the blend NCPEs. The peak at 624  $cm^{-1}$  is due to free  $ClO_4^-$  anions, indicating the dissolution of lithium salt in the polymer matrix.<sup>61</sup> The peak observed at 1660  $cm^{-1}$  is due to the bending mode of the adsorbed water molecule, due to the hygroscopic nature of  $LiClO_4$  salt.<sup>62</sup>

### X-ray diffraction analysis

The structural characteristics of the fabricated films were investigated by X-ray diffraction (XRD). Fig. 3 depicts the XRD pattern of pristine PVDF-HFP, PVDF-HFP/PEG blend, and the respective NCPEs. For pristine PVDF-HFP, the XRD pattern reveals relative sharp crystalline peaks at  $\sim 18.6^\circ$ ,  $\sim 20.3^\circ$ ,  $\sim 26.9^\circ$ , and  $\sim 38.9^\circ$  along with a halo, which demonstrates the semicrystalline nature of PVDF-HFP.<sup>63,64</sup> The sharp peaks correspond to the crystalline phase of PVDF-HFP due to VDF units while the halo is associated with the amorphous phase due to HFP moieties.<sup>65</sup> The XRD patterns of  $\alpha$ ,  $\beta$ , and  $\gamma$ -phases exhibit an intense peak in the region  $\sim 20^\circ$  but only  $\alpha$  and  $\gamma$ -phases display additional peaks close to  $\sim 18^\circ$ . Also, the XRD pattern of the  $\alpha$ -phase also displays a relatively strong peak around  $26^\circ$ .<sup>54</sup> However, due to the lower crystallinity of the copolymer and poorly resolved crystalline peaks, it is not usually possible to detect the correct crystal phase of the PVDF in PVDF-HFP from the XRD pattern, nevertheless, by taking clues from the FTIR spectrum of the pristine PVDF-HFP, we can safely suggest the  $\alpha$ -phase for the pristine PVDF-HFP film.

Blending PEG with PVDF-HFP results in the broadening of the crystalline peaks; an indication of the decreased crystallinity of PVDF-HFP. The formation of the  $\beta$ -phase is evident from the appearance of a prominent peak at  $\sim 20.3^\circ$  in blend NCPEs, however, the presence of other small peaks does not rule out the existence of all three phases ( $\alpha$ ,  $\beta$ , and  $\gamma$ ) in blend NCPEs. As discussed earlier, FTIR spectra also revealed the formation of mixed crystal phases for the respective samples. Additionally, no visible/distinguishable peak due to PEG could



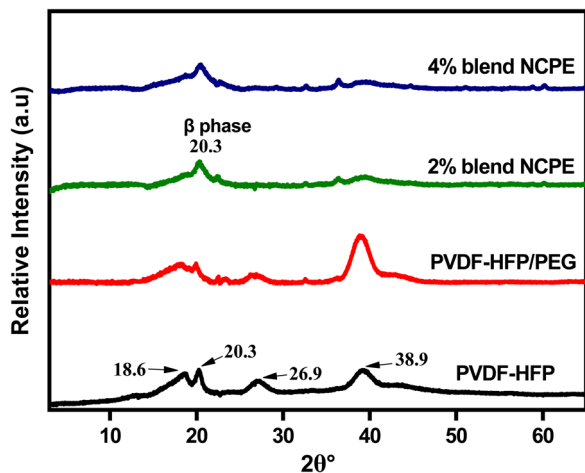


Fig. 3 XRD patterns of pristine PVDF-HFP, PVDF-HFP/PEG blend and blend NCPEs with various concentrations of  $\text{Mn}_3\text{O}_4$  NPs.

be detected in the XRD pattern of the PVDF-HFP/PEG blend and blend NCPEs that could be attributed to the uniform distribution of the PEG in the blend matrix. The absence of peaks due to undissociated  $\text{LiClO}_4$  in the XRD patterns of the NCPEs suggests the homogeneous dispersion of salt in the polymer matrix.<sup>66</sup>

#### DSC analysis

Pristine PVDF-HFP exhibits (Fig. 4) a strong endothermic peak at  $\sim 142.1$  °C, which is attributed to the melting temperature of the crystalline phase of PVDF-HFP. The incorporation of PEG in the matrix has only slightly affected (broadened and the peak maximum shifted to  $\sim 140.7$  °C) the melting peak of PVDF-HFP. The rather smaller effect could be attributed to phase segregation as is evident from the melting endotherm of the PEG at  $\sim 60$  °C in the DSC thermogram of the blend.<sup>67</sup> For blend NCPEs, no PEG melting peak could be detected, which

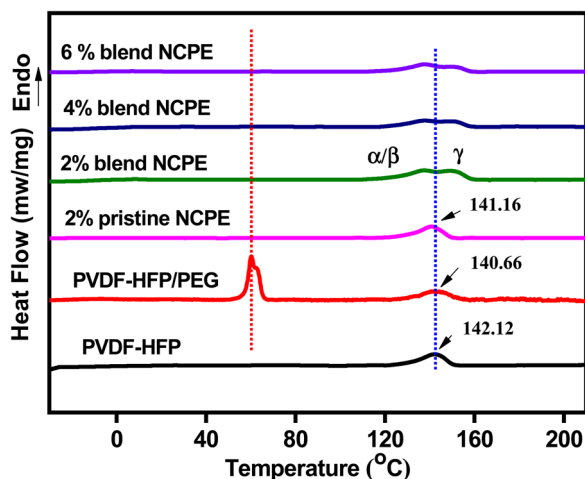


Fig. 4 DSC thermograms of PVDF-HFP, PVDF-HFP/PEG blend, pristine NCPE and blend NCPEs with various concentrations of  $\text{Mn}_3\text{O}_4$  NPs (2nd heating cycle).

Table 3 Thermal transitions and melting enthalpies obtained from the DSC thermograms of the fabricated films

Sample	$T_m$ (°C) ( $\alpha$ , $\beta$ )	$T_m$ (°C) ( $\gamma$ )	$T_c$ (°C)	$\Delta H_m$ (mJ $\text{mg}^{-1}$ )
PVDF-HFP	142.1	—	113.2	71
PVDF-HFP/PEG	140.7	—	110.8	48
2% Pristine NCPE	141.2	—	115.0	64
2% Blend NCPE	137.5	151.2	109.7	15
4% Blend NCPE	137.8	151.3	108.0	30
6% Blend NCPE	138.4	151.6	111.0	25

suggests a homogeneous dispersion of PEG in the blend NCPE matrix. There is a slight variation in melting temperature of the PVDF upon blending or in pristine NCPE, however, a subtle change in melting behavior could be seen for blend NCPEs. Here, the melting peak of the PVDF-HFP splits into two peaks. We can argue that the higher melting peak ( $T_m \sim 151$  °C) represents melting of the  $\gamma$ -phase crystals while the lower temperature melting peak ( $T_m \sim 138$  °C) could be associated with the melting of the  $\alpha/\beta$ -crystalline phases. This further reinforces the FTIR and XRD data discussed above, which suggest the formation of mixed crystal phases in blend NCPEs. It is to be noted that  $\gamma$ -phase crystals melt at higher temperature as compared with  $\alpha$  or  $\beta$ -crystalline phases ( $\alpha$  and  $\beta$  crystalline phases have a similar melting temperature range).<sup>50</sup> To calculate the % crystallinity of a semicrystalline polymer one needs the melting enthalpy of the respective polymer in 100% crystalline state.

The PVDF in the PVDF-HFP copolymer can never be achieved in 100% crystalline state, therefore, % crystallinity cannot be calculated from the DSC data. However, we have qualitatively assessed the crystallinity of the PVDF-HFP in the fabricated samples by comparing the corresponding melting enthalpies  $\Delta H_m$  (mJ  $\text{mg}^{-1}$ ) of the PVDF-HFP calculated from the respective melting peak areas as given in Table 3. There is a significant depression in melting enthalpy and hence crystallinity of the blend NCPEs as compared with the pristine PVDF-HFP or pristine NCPE. This could be attributed to the complex interaction pattern between PEG, additives:  $\text{Mn}_3\text{O}_4$  NPs, and  $\text{LiClO}_4$ , and PVDF-HFP that may offer a strong hindrance to the crystallization of PVDF-HFP chains.<sup>68</sup>

The minimum melting enthalpy and hence crystallinity was calculated for 2% blend NCPE. Further increase in the filler content seems to have the opposite effect on crystallinity. Increasing the filler content above the optimum level leads to their self-aggregation in the matrix, thus leaving the polymer chains unaffected and they can self-organize more efficiently and hence increase crystallinity.

#### Thermal gravimetric analysis

Thermal gravimetric analysis (TGA) was carried out under  $\text{N}_2$  atmosphere to assess the thermal stability of the prepared NCPEs, and the data are depicted in Fig. 5. Pristine PVDF-HFP initiates degradation at  $\sim 480$  °C and follows a single step degradation process. PVDF-HFP decomposes due to chain scission reactions, and inter-chain and intra-chain molecular transfer reactions.<sup>69</sup> The PVDF-HFP/PEG blend degrades in two steps, *i.e.*, the first degradation step at 350 °C is due to the



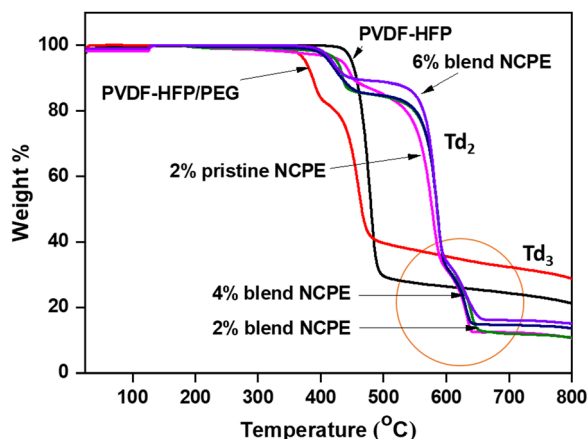


Fig. 5 TGA thermograms of PVDF–HFP, PVDF–HFP/PEG blend, 2% pristine NCPE and blend NCPEs with various concentrations of  $\text{Mn}_3\text{O}_4$  NPs.

decomposition of PEG,<sup>70</sup> with almost 21% mass loss while the second step is due to the degradation of the PVDF–HFP at 410 °C. Three steps of degradation could be seen for the pristine and blend NCPEs. The 1st step of degradation occurs at ~400 °C which is due to PEG degradation while the 2nd ( $T_{d2}$ ) and 3rd ( $T_{d3}$ ) decomposition steps observed at ~550 °C and ~590 °C are due to the PVDF–HFP chains and PVDF–HFP–salt complex, respectively. It is to be noted that both the pristine and blend NCPEs reveal weight loss at ~400 °C, thus, one could argue that the weight loss at ~400 °C could be associated with the degradation of both the  $\text{LiClO}_4$ <sup>71</sup> and PEG. It is observed that PVDF–HFP in both the pristine and blend NCPEs exhibits higher thermal stability as compared to the pristine PVDF–HFP and PVDF–HFP/PEG blend.

This suggests that the inert filler ( $\text{Mn}_3\text{O}_4$ ) enhances the thermal stability of the fabricated NCPEs. A minor, sudden weight increase (<1%) is observed in the TGA curves (Fig. 5) for certain samples within the temperature range of ~130 °C to 140 °C. Normally, the weight of a sample decreases as the temperature rises, due to the degradation of the material into gaseous products. However, in some cases, a weight gain can occur during TGA analysis. This phenomenon can have several potential causes, such as oxidation (reaction with oxygen), gas adsorption by porous materials (e.g., activated carbon), or instrumental artifacts.<sup>72–74</sup> In our case, oxidation can be ruled out, as the TGA analysis was conducted under a nitrogen atmosphere. Additionally, nitrogen adsorption is unlikely since the sample is not porous. Therefore, the small and sudden weight increase observed in the TGA curves for some samples is most likely due to an instrumental artifact. Importantly, this effect is minimal and does not impact the overall thermal behavior or interpretation of the results. It is to be noted that for all the fabricated NCPEs no significant weight loss could be observed up to 400 °C, which ensures their safe practical application under thermally challenging conditions.

### Surface morphology and composition

To find the effect of hausmannite NPs on surface morphology and distribution in the prepared electrolytes, FESEM and EDS

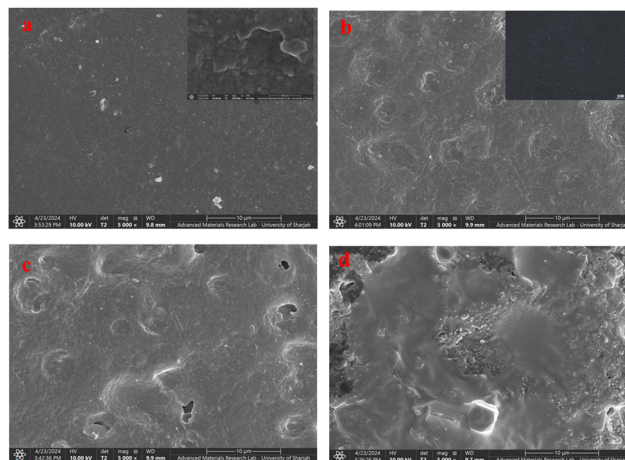


Fig. 6 FESEM images of the (a) pristine PVDF–HFP, (b) PVDF–HFP/PEG blend (inset shows the POM micrograph of the PVDF–HFP/PEG blend), (c) 2% pristine NCPE and (d) 2% blend NCPE.

analyses were performed. The SEM images shown in Fig. 6, reveal a more compact and smoother surface for pristine PVDF–HFP film as compared with the PVDF–HFP/PEG blend or the NCPEs. The polarized optical micrograph (inset Fig. 6(b)) of the PVDF–HFP/PEG blend film did not reveal any spherulite formation by the PEG, suggesting that no PEG surface segregation takes place in the blend film.

Fig. 7(a) depicts the EDS analysis of the distribution of various elements for the 2% blend NCPE exhibiting a uniform distribution of Mn (from nanofiller), Cl (from salt), O (from PEG and salt), and F (from PVDF–HFP) in the polymer matrix.

To further validate the composition and the presence of filler in the composite samples, XPS analysis was carried out. As shown in Fig. 7(b), the XPS survey of 2% pristine NCPE shows their composition, validating the presence of Mn filler. The core shell spectrum of C(1s) peak deconvolution shows two dominant peaks observed at 284.2 eV and 292.1 eV for C–C and C–F respectively, related to the PVDF–HFP polymer matrix.<sup>75</sup> The XPS analysis of F(1s) shows two peaks occurring at 653 eV and 642 eV, related to the F–C and F–Li.<sup>76</sup> This shows the interaction of  $\text{Li}^+$  with the polymer matrix. The binding energy of Li(1s) is 56.1 eV whereas Cl(2p) is 208.6 eV, both related to  $\text{LiClO}_4$  salt. The core shell deconvoluted spectrum for Mn(2p) is also shown in Fig. 7(b)(c), with two distinct peaks observed at 641 eV and 653 eV for  $\text{Mn}(2p)_{3/2}$  and  $\text{Mn}(2p)_{1/2}$  respectively, related to  $\text{Mn}_3\text{O}_4$  filler.<sup>77</sup> Fig. 7b also displays the XPS survey of 2% blend NCPE, along with the core shell spectrum of C(1s), F(1s) and Mn(2p) (Fig. 7b(e–g)). The binding energy of C(1s) in the 2% blend NCPE is 285.14 eV. The binding energies related to O(1s) and O(2s), respectively, are 532.7 and 25.7 eV in the 2% blend NCPE. The F(1s) and F(2s) have binding energies, respectively, of 688.0 and 27.33 eV. As shown in Fig. 7(b) and (f), the F–Li peak is intense in the 2% pristine NCPE as compared to the 2% blend NCPE. This suggests that  $\text{Li}^+$  in pristine NCPE interacts only with the F atom of the matrix whereas in the blend NCPE, the  $\text{Li}^+$  interaction occurs with both components of the matrix *i.e.* F of the PVDF–HFP and O of the PEG.



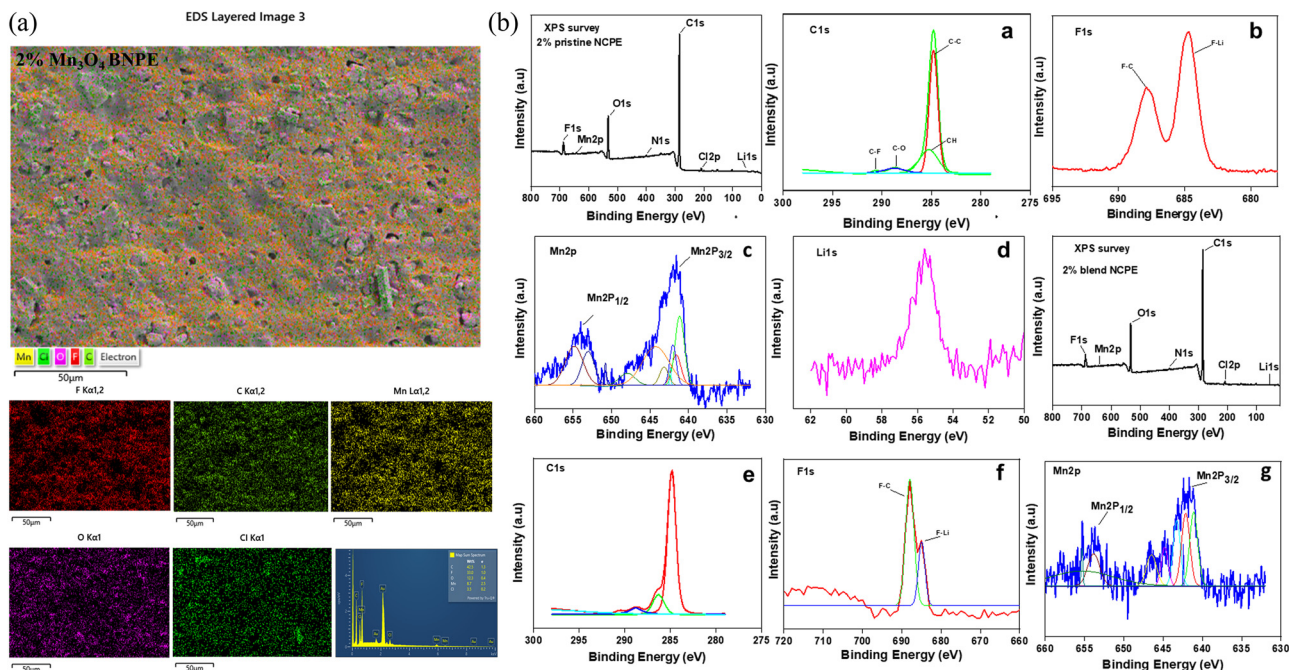


Fig. 7 (a) EDS analysis of the 2% blend NCPE, showing the atom distribution and composition of the film. (b) XPS survey of 2% pristine NCPE, along with the core shell spectrum of various atoms (a)–(d) and XPS survey of the 2% blend NCPE, along with the core shell spectrum of various atoms (e)–(g).

## Tensile properties

The stress–strain curves were recorded to examine the tensile behavior of the fabricated films as shown in Fig. 8. The stress–strain curve displays two distinct regions; an initial linear region depicting the elastic behaviors, followed by the non-linear region representing the nonreversible plastic deformation region.<sup>11,78</sup> As shown in Fig. 8, the stress–strain curve of the pristine PVDF–HFP and PVDF–HFP/PEG blend shows a sharp linear increase in stress (elastic region), reaching a maximum value followed by the irreversible strain deformation and breaking at a much lower strain value. The tensile strength data of the fabricated films are shown in Table 4.

This indicates the brittle nature of the neat PVDF–HFP and PVDF–HFP/PEG blend films. The NCPEs possess more ductile

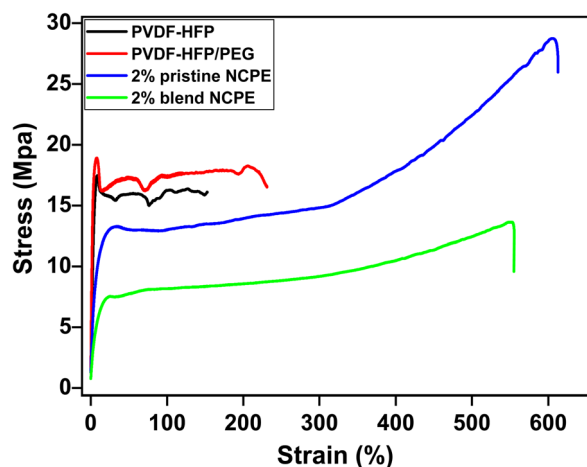


Fig. 8 Typical stress–strain curves of the pristine PVDF–HFP, PVDF–HFP/PEG blend, 2% pristine and 2% blend NCPE.

behavior as evident from the significantly higher elongation at break value as compared to pristine PVDF–HFP and the PVDF–HFP/PEG blend. The ductile behavior and excellent flexibility of the 2% blend NCPE are advantageous for better electrode/electrolyte interfacial content in lithium ion batteries.<sup>79</sup>

## Electrochemical impedance spectroscopy

Electrochemical impedance spectroscopy (EIS) was employed to determine the ion conductivity of the fabricated films in the temperature range  $25\text{ }^{\circ}\text{C} \leq T \leq 80\text{ }^{\circ}\text{C}$ .

Fig. 9(a) shows the log–log plot of  $\sigma'(\omega)$  and frequency for the 2% blend NCPE, measured at various temperatures. As shown in Fig. 9(a), the log–log plot consists of three regions: (i) the low-frequency region below the plateau is due to the electrode polarization, (ii) the higher frequency region which is above the plateau is called the ac conductivity or universal dielectric response, and (iii) the frequency independent plateau region. The dc conductivity ( $\sigma_0$ ) is extracted from the plateau in the log–log plot by applying the so-called Dyre fitting function (eqn (2)).<sup>80</sup>

$$\sigma'(\omega) = \frac{\sigma_0 \omega \tau \tan^{-1}(\omega \tau)}{\frac{1}{4} [\ln\{1 + (\omega \tau)^2\}]^2 + \{\tan^{-1}(\omega \tau)\}^2} \quad (2)$$

where  $\tau$  and  $\sigma_0$  are the dielectric relaxation time and dc

Table 4 The tensile strength of the fabricated films

Sample ID	Tensile strength (MPa)
PVDF–HFP	16.2
PVDF–HFP/PEG	18.9
2% pristine NCPE	28.7
2% blend NCPE	13.6



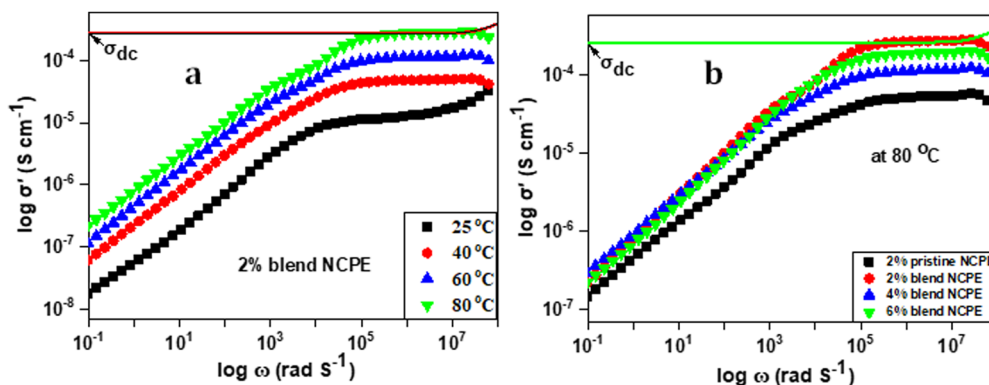


Fig. 9 Log–log plot of (a) the 2% blend NCPE, measured at various temperatures and (b) the 2% pristine NCPE and blend NCPEs measured at 80 °C.

conductivity, respectively, which are the fitting parameters. The  $\tau$  is the inverse of angular frequency ( $\tau = 1/\omega$ ). Fig. 9(b) depicts the log–log plot for 2% pristine NCPE and blend NCPEs with various concentrations of  $\text{Mn}_3\text{O}_4$  NPs. The calculated parameters are tabulated in Table 5.

It is evident from Fig. 9(a) that the dc conductivity increases with increasing temperature as manifested by the shifting of the plateau along the y-axis with increasing temperature. The maximum ion conductivity was calculated to be  $3.1 \times 10^{-4} \text{ S cm}^{-1}$  at 80 °C. This could be attributed to faster segmental mobility and faster ion dynamics at higher temperatures. Fig. 9(b) and Table 5 reveal that 2% blend NCPE displays the maximum ion conductivity which decreases with further increase in filler content. For NCPEs, there is always an optimum filler concentration above which the ion conductivity deteriorates and this trend is attributed to the filler aggregation above the optimum concentration.<sup>8,11</sup> The aggregates block the conduction pathways of the ions that result in lowering ion conductivity. As discussed above for 4% and 6% blend NCPEs the DSC data revealed slightly higher crystallinity as compared with the 2% blend NCPE and that was attributed to the filler aggregation in the matrix.

The data also reveal a significantly higher ion conductivity for blend NCPE as compared with the pristine NCPE with the same composition (2% nanofiller). For example, ion conductivity was measured to be  $5.1 \times 10^{-5} \text{ S cm}^{-1}$  for the 2% pristine NCPE, while for the 2% blend NCPE the value was found to be  $3.1 \times 10^{-4} \text{ S cm}^{-1}$ , which is an order of magnitude higher than that for the 2% pristine NCPE. As discussed above PEG has a dual role in blend NCPEs. It acts as a plasticizer that reduces the PVDF–HFP crystallinity and by interacting with lithium salt

it facilitates its dissociation that in turn results in a higher charge carrier concentration. Furthermore, it is a better conductor of lithium ions as compared with the pristine PVDF–HFP. Thus, we can conclude that the blend PVDF–HFP/PEG matrix offers a more robust ion conducting environment as compared with the pristine PVDF–HFP based NCPE. There are many reports on the incorporation of metal oxides as a nanofiller in PVDF–HFP for improving the ion conductivity.<sup>81–83</sup> Subramania *et al.*<sup>84</sup> incorporated a  $\text{ZrO}_2$  nanofiller into PVDF–HFP based polymer electrolytes and electrochemical properties were analyzed at the optimum nanofiller content (8%  $\text{ZrO}_2$ ). Nidhi *et al.*<sup>85</sup> fabricated PVDF–HFP based NCPEs by incorporating various ceramic oxide nanofillers, such as  $\text{MgO}$ ,  $\text{ZnO}$ ,  $\text{TiO}_2$ , and  $\text{Al}_2\text{O}_3$ . The maximum ion conductivity of  $1.25 \times 10^{-5} \text{ S cm}^{-1}$  was achieved for  $\text{ZnO}$ -based NCPEs at room temperature and the conductivity increased to  $1.24 \times 10^{-4} \text{ S cm}^{-1}$  at 110 °C, which is relatively lower in comparison to blend NCPEs in the current work.

The Arrhenius equation (eqn (3)) was applied to find the temperature-dependent ion conductivity behavior of the fabricated pristine and blend NCPEs as depicted in Fig. 10.

$$\sigma_{\text{dc}} = \sigma_{\infty}^{\text{Arr}} \exp \left( \frac{-E_a}{K_B T} \right) \quad (3)$$

where  $K_B$  is the Boltzmann constant,  $E_a$  represents the activation energy and  $\sigma_{\infty}^{\text{Arr}}$  is the infinitely high temperature ion conductivity. The data shown in Fig. 10 suggest that all the fabricated NCPEs follow temperature dependent Arrhenius behavior. Thus, the ion conductivity is a thermally activated process following the ion hopping mechanism with random jumps of lithium ion along the VDF and PEG units.<sup>86,87</sup>

The activation energy calculated from the respective slopes of the Arrhenius plots (Fig. 10) revealed the lowest value (0.16 eV) (Table 4) for the 2% blend NCPE among the investigated pristine and blend NCPEs, suggesting that the 2% blend NCPE matrix offers the smallest energy barrier to the ion mobility. This trend is in accordance with the ion conductivity behavior of this sample (Table 5). The exact mechanism of the ion diffusion in the blend matrix is still unclear, however, the interaction of lithium ion with PEG, and PVDF–HFP of the matrix is evident from the FTIR, XRD, and DSC data presented above. Furthermore, the temperature dependent ion conductivity follows the Arrhenius

Table 5 The activation energy ( $E_a$ ), dc conductivity ( $\sigma_{\text{dc}}$ ), and the dielectric relaxation time ( $\tau$ ) of the fabricated films

Sample ID	$E_a$ (eV)	$\sigma_{\text{dc}}$ ( $\text{S cm}^{-1}$ , 80 °C)	$\tau$ (s)
PVDF–HFP	—	—	—
PVDF–HFP/PEG	—	—	—
2% Pristine NCPE	0.23	$5.1 \times 10^{-5}$	$5.93 \times 10^{-8}$
2% Blend NCPE	0.16	$3.1 \times 10^{-4}$	$5.24 \times 10^{-8}$
4% Blend NCPE	0.24	$1.12 \times 10^{-4}$	$6.32 \times 10^{-8}$
6% Blend NCPE	0.27	$1.85 \times 10^{-4}$	$6.55 \times 10^{-8}$





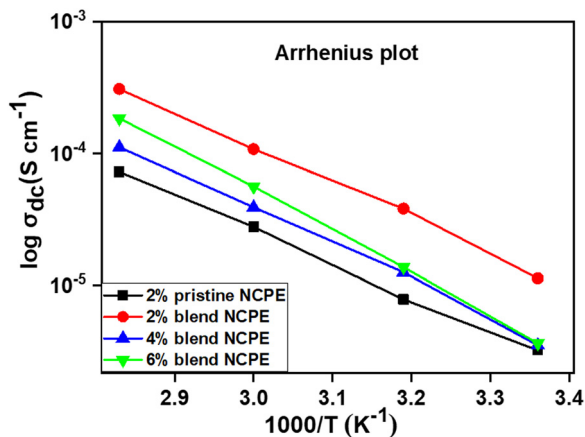


Fig. 10 Temperature dependent Arrhenius plots of pristine NCPE and blend NCPEs with various concentrations of  $\text{Mn}_3\text{O}_4$  NPs.

behavior. Thus, one can conclude the ion diffusion is a thermally activated process and follows the ion hopping mechanism with jumps from one coordinated site to another when voltage is applied. The presence of PEG in the matrix and the Lewis acid–base interaction between the  $\text{Li}^+$  ether oxygen of the PEG backbone offers additional ion conducting pathways in the blend matrix as compared with the pristine PVDF–HFP matrix.

Table 6 reveals the comparison of ion conductivity of some of the reported PVDF–HFP-based composite electrolytes with various nanofillers.

### Dielectric properties

The frequency and temperature-dependent dielectric behavior of the fabricated electrolytes were investigated to obtain insight into the ion transport phenomenon. Fig. 11(a) and (b) represent the dielectric constant ( $\epsilon'$ ) and dielectric loss ( $\epsilon''$ ) as a function of frequency for the 2% blend NCPE measured at various temperatures. Both the  $\epsilon'$  and  $\epsilon''$  sharply rise in the lower frequency regime followed with a flat tail in the higher frequency region. The lower frequency sharp increase is due to the space charge polarization phenomenon, which results from the accumulation of charge carriers at the electrode/electrolyte interface and indicates the non-Debye relaxation process.<sup>92,93</sup> As shown in Fig. 11, both  $\epsilon'$  and  $\epsilon''$  increase with an increase in temperature at a given frequency.<sup>94</sup> At higher frequencies, the ions do not have enough time to align according to the direction of the electric field, hence no long range ion diffusion and accumulation takes place that results in lowering the value of both the dielectric constant and dielectric loss at higher frequencies.<sup>92,95</sup>

Table 6 Comparison of various reported PVDF–HFP and various nanofiller based composite electrolytes

Polymer matrix	Filler	Conductivity ( $\text{S cm}^{-1}$ )	Ref.
PVDF–HFP	ZnO	$1.25 \times 10^{-5}$ (room temp)	85
PVDF–HFP/Pluronic	GO	$1.2 \times 10^{-6}$ (room temp)	88
PVDF–HFP/PEO	M–TiO <sub>2</sub>	$7.21 \times 10^{-4}$ (room temp)	89
PVDF–HFP	SiO <sub>2</sub>	$7.1 \times 10^{-4}$ (room temp)	90
PVDF–HFP	GO	$4.23 \times 10^{-4}$ (room temp)	91
PVDF–HFP/PEG	$\text{Mn}_3\text{O}_4$	$3.1 \times 10^{-4}$ (80 °C)	This work

The loss tangent ( $\tan \delta = \epsilon''/\epsilon'$ ) vs. frequency plot is also depicted in Fig. 11(c) for the 2% blend NCPE measured at various temperatures. The plots show a single dielectric relaxation peak. The relaxation peak shifts to higher frequency as the temperature increases. This finding could be attributed to the comparatively fast ion dynamics and chain segmental motion with increasing temperature that improves the ion conductivity upon heating.<sup>9</sup>

### Electric modulus

To further understand the ion dynamics process as a function of frequency and temperature, the dielectric data were transformed into an electric modulus formalism. The dielectric data in modulus formalism suppress the electrode polarization effect observed in the low-frequency region, therefore, any lower frequency relaxation phenomenon overlooked by the dielectric formalism can be evaluated.<sup>96,97</sup> The complex modulus and permittivity are related by eqn (4):

$$M^* = \frac{1}{\epsilon^*(\omega)} = M' + iM'' = \frac{\epsilon'}{\epsilon'^2 + \epsilon''^2} + i\frac{\epsilon''}{\epsilon'^2 + \epsilon''^2} \quad (4)$$

where  $M'$  is the real part and  $M''$  is the imaginary part of the electric modulus. In the lower frequency region,  $M''$  offers information about the long-range motion of charged species, *i.e.*, related to the ion conductivity. For a pure conduction process, a relaxation peak is present in the spectrum of the imaginary component of the electric modulus, with no relaxation peaks in the dielectric loss spectrum.<sup>98</sup> For the dielectric relaxation process, however, the relaxation peak may be seen in both the  $M''$  and  $\epsilon''$  spectra.<sup>99,100</sup> A relaxation peak in the  $\epsilon''$  versus frequency spectrum indicates that dipoles in the polymer chain have reoriented, which causes dielectric relaxation.

Therefore, it is useful to identify the long-range conduction processes and separate them from the localized dielectric relaxation processes by comparing the experimental data in the  $M^*$  and  $\epsilon^*$  formalisms.

Fig. 12 represents the  $M'$  and  $M''$  vs. frequency for the 2% blend NCPE measured at various temperatures. Both  $M'$  and  $M''$  values reach zero at lower frequencies and sharply increase at higher frequencies.<sup>101</sup> At lower frequencies, the formation of a long tail is assigned to electrode polarization that generates a large capacitance associated with the electrode/electrolyte interface and a high dielectric constant.<sup>102</sup> At higher frequencies, the increased electric modulus value is attributed to the electrolyte bulk properties.<sup>102,103</sup> The increase in  $M''$  in the higher frequency region is an indication of the conductivity relaxation process, however, the relaxation peak could not be detected as it seems to be beyond the frequency limits of the instrument.<sup>104,105</sup>

### Role of PEG on the structural and electrochemical properties of the electrolyte membrane

PEG is a semicrystalline, thermoplastic polymer that can act as a plasticizer matrix when a lower molecular weight  $<10\,000 \text{ g mol}^{-1}$  is used.<sup>106</sup> A plasticizer reduces the intra and inter molecular interaction between the polymer chains, thus reducing the glass transition temperature and crystallinity of the polymer chains and hence the salt dissociation of the



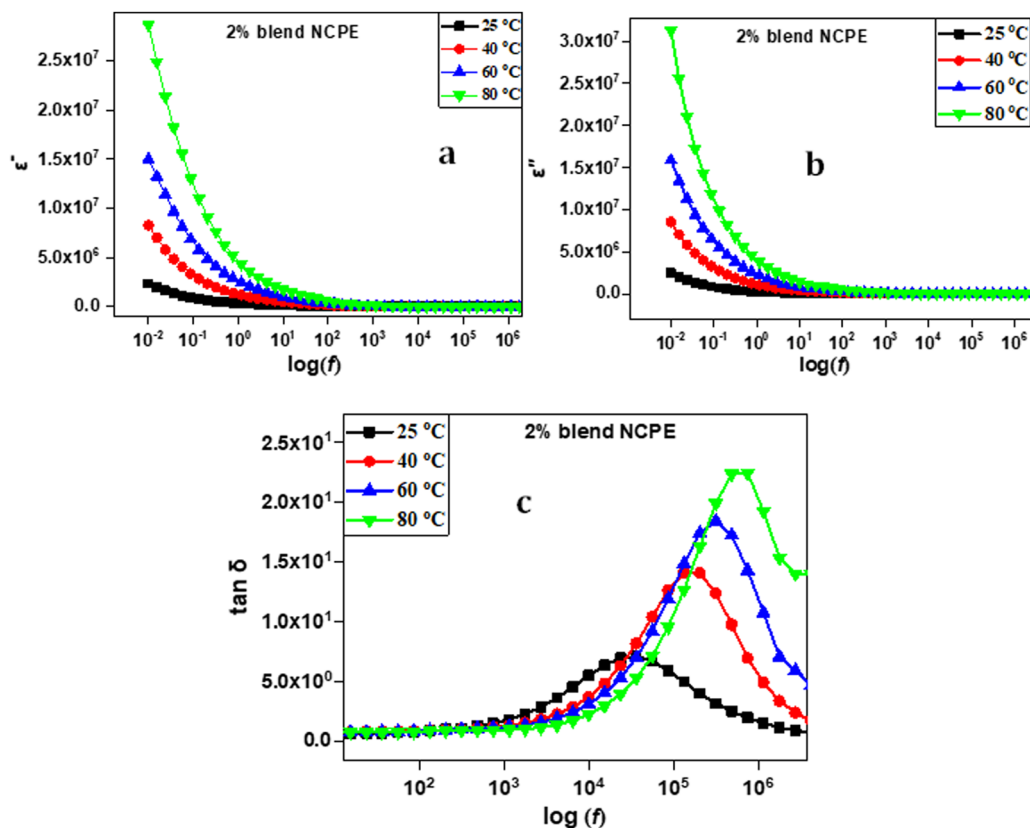


Fig. 11 Dielectric data for the 2% blend NCPE representing frequency dependent (a) dielectric constant, (b) dielectric loss and (c) loss tangent at various temperatures.

polymer matrix. Hence, PEG is regarded as the most efficient plasticizer for PVDF–HFP as it has good stability, dispersibility, and high mobility with large exclusion volumes and non-toxicity.<sup>107</sup> In the present study, low molecular weight PEG was introduced to the PVDF–HFP matrix and various structural, morphological and electrochemical changes were observed. As shown in the FTIR data, adding PEG to the pristine PVDF–HFP matrix shifted some of the electro inactive nonpolar  $\alpha$ -phase peaks to the polar electroactive phase. The higher content of  $\beta$  and  $\gamma$  phases in the material is the subject of high interest for

potential applications in various fields, including sensing, electro optic devices, data storage, actuation, and energy harvesting.<sup>51</sup> As shown from the DSC data, the melting temperature slightly decreased upon the addition of PEG to the PVDF–HFP matrix. In the PVDF–HFP/PEG matrix and blend NCPEs, the melting enthalpy decreased. This indicates an increase in amorphous content and decrease in crystallinity as compared with the pristine PVDF–HFP. It is suggested that PEG chains disrupt the inter PVDF–HFP chain interactions and hinder their crystallization to some extent. The EIS data also

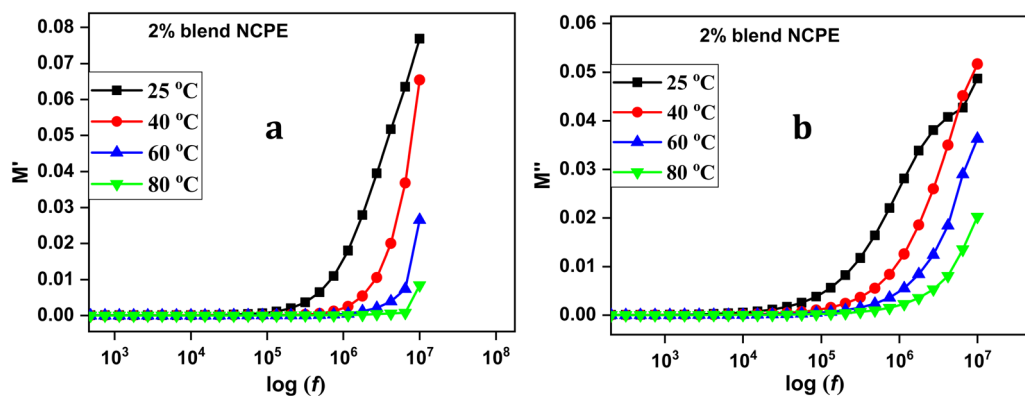


Fig. 12 Electric modulus data (a) real part and (b) imaginary part of electric modulus of the 2% blend NCPE as a function of frequency at various temperatures.



revealed a significantly higher ion conductivity for PEG based blend NCPE as compared with the pristine NCPE with the same composition (2% nanofiller). As discussed above PEG has a dual role in blend NCPEs. It acts as a plasticizer that reduces the PVDF–HFP crystallinity and by interacting with lithium salt it facilitates its dissociation that in turn results in higher charge carrier concentration and so higher ion conductivity. Furthermore, it is a better conductor of lithium ions as compared with the pristine PVDF–HFP. Thus, we can conclude that the blend PVDF–HFP/PEG matrix offers a more robust ion conducting environment as compared with the pristine PVDF–HFP based NCPE.

## Conclusions

The fabrication and characterization of pristine PVDF–HFP and PVDF–HFP/PEG blend-based nanocomposite polymer electrolytes (NCPEs), namely (i) PVDF–HFP/LiClO<sub>4</sub>/2 wt% Mn<sub>3</sub>O<sub>4</sub> and (ii) PVDF–HFP/PEG/LiClO<sub>4</sub>/x wt% Mn<sub>3</sub>O<sub>4</sub>, where x represents the weight percent of Mn<sub>3</sub>O<sub>4</sub> nanofiller (ranging from 2 to 6%) has been carried out. FTIR spectroscopy confirmed the formation of a nonpolar  $\alpha$ -crystal phase by pristine PVDF–HFP, which transformed into a mixed  $\alpha$  and  $\gamma$ -phase in the PVDF–HFP/PEG blend. The  $\alpha$ -crystal phase completely disappeared and was replaced by the  $\beta$  and  $\gamma$  crystal phase in NCPEs. This was attributed to strong interaction between the PVDF and the additives that forces the PVDF chains to pack into polar crystal phases. That was also substantiated by DSC analysis, where the melting peak of PVDF splits into two peaks corresponding to the melting of  $\gamma$ -phase crystals at higher temperature and  $\beta$ -phase crystals at lower temperature. Electrochemical impedance spectroscopy (EIS) revealed a maximum ion conductivity of  $3.1 \times 10^{-4} \text{ S cm}^{-1}$  at 80 °C for the 2% blend NCPE, which is an order of magnitude higher as compared to  $5.1 \times 10^{-5} \text{ S cm}^{-1}$  at 80 °C for the 2% pristine NCPE. The ion conductivity and activation energy data suggest that the PVDF–HFP/PEG blend matrix, due to the presence of PEG, offers more efficient conducting pathways for the transportation of ions as compared with the pristine PVDF–HFP matrix. The dielectric relaxation peak shifted to higher frequency (lower relaxation time) with increasing temperature, which is attributed to faster ion dynamics and conductivity with increasing temperature.

## Author contributions

KHK: conceptualization, investigation, formal analysis, methodology, writing – original draft; AZ: writing, formal analysis; HR: methodology, formal analysis; IA: investigation, methodology, formal analysis; GSK: methodology, investigation, formal analysis, HH: conceptualization, funding acquisition, supervision, writing – original draft.

## Data availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

## Conflicts of interest

The authors declare no competing financial interest.

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