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# Experimental and theoretical studies on the formation of pure $\beta$ -phase polymorphs during fabrication of polyvinylidene fluoride membranes by cyclic carbonate solvents†

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The use of highly toxic solvents presents significant risks to both the environment and human health. Therefore, the adoption of green solvents will be crucial for achieving sustainable membrane production. This work reports the use of inexpensive environmentally friendly biobased and biodegradable cyclic carbonate solvents, namely ethylene carbonate (EC), propylene carbonate (PC), and butylene carbonate (BC), to fabricate polyvinylidene fluoride (PVDF) membranes. The solvent dependence of the phase inversion mechanisms, morphology, crystalline structures, and polymorphism of the prepared PVDF membranes were investigated. Polymorph analysis revealed that membrane fabrication in EC or PC yielded exclusively the  $\beta$ -phase product, whereas PVDF membrane fabrication in BC yielded a mixture of  $\alpha$  and  $\beta$  phase material. The mechanism of  $\beta$ -phase formation was investigated using molecular dynamics simulation and shown to depend on the extent of hydrogen bonding at the polymer–solvent interface. The PVDF membrane formed in EC exhibited the highest porosity and pure water permeability, and was therefore tested in direct contact membrane distillation (DCMD), exhibiting promising results in terms of permeate flux and salt rejection. These results suggest that large-scale production of piezoelectric PVDF membranes using green solvents should be practically feasible.

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## 1. Introduction

The performance of the low-energy intensive separation units such as membrane separation technologies depends strongly on the properties of the separatory material interacting with the chemical constituents of an industrial stream. One group of interesting materials considered for membrane preparation is fluoropolymers among which poly(vinylidene fluoride)

(PVDF) is the most popular due to its attractive properties and good processability.

Homopolymeric PVDF contains 59.4 wt% fluorine and 3 wt% hydrogen.<sup>1</sup> The crystalline phase of PVDF has three different molecular conformations (TGTG', TTT, and TTTGTTG') and five crystal polymorphs: phase I ( $\beta$ -phase), phase II ( $\alpha$ -phase), phase III ( $\gamma$ -phase),  $\delta$ , and  $\epsilon$ . Phases I, II, and III (see Fig. 1) are the most frequent crystalline polymorphs of PVDF. The  $\alpha$ -phase is kinetically favoured, but the  $\beta$ -phase is the most thermodynamically stable. Detailed information on each crystalline polymorph is presented in Table 1.

$\beta$ -Phase PVDF has useful piezo-, pyro-, and ferroelectric properties. Piezoelectric PVDF membranes vibrate when an AC current is passed through them. This property has recently attracted much interests because it could potentially be exploited to control membrane fouling. For example, Darestani *et al.*<sup>2</sup> showed that fouling of piezoelectric PVDF membranes can be controlled by a synergistic combination of piezoelectric vibration and high crossflow velocity. Similar results were reported by Zhang *et al.*,<sup>3</sup> whose studies on PVDF/acetyl tributyl citrate/ionic liquid systems revealed that applying an AC current to PVDF membranes increased the water permeate flux

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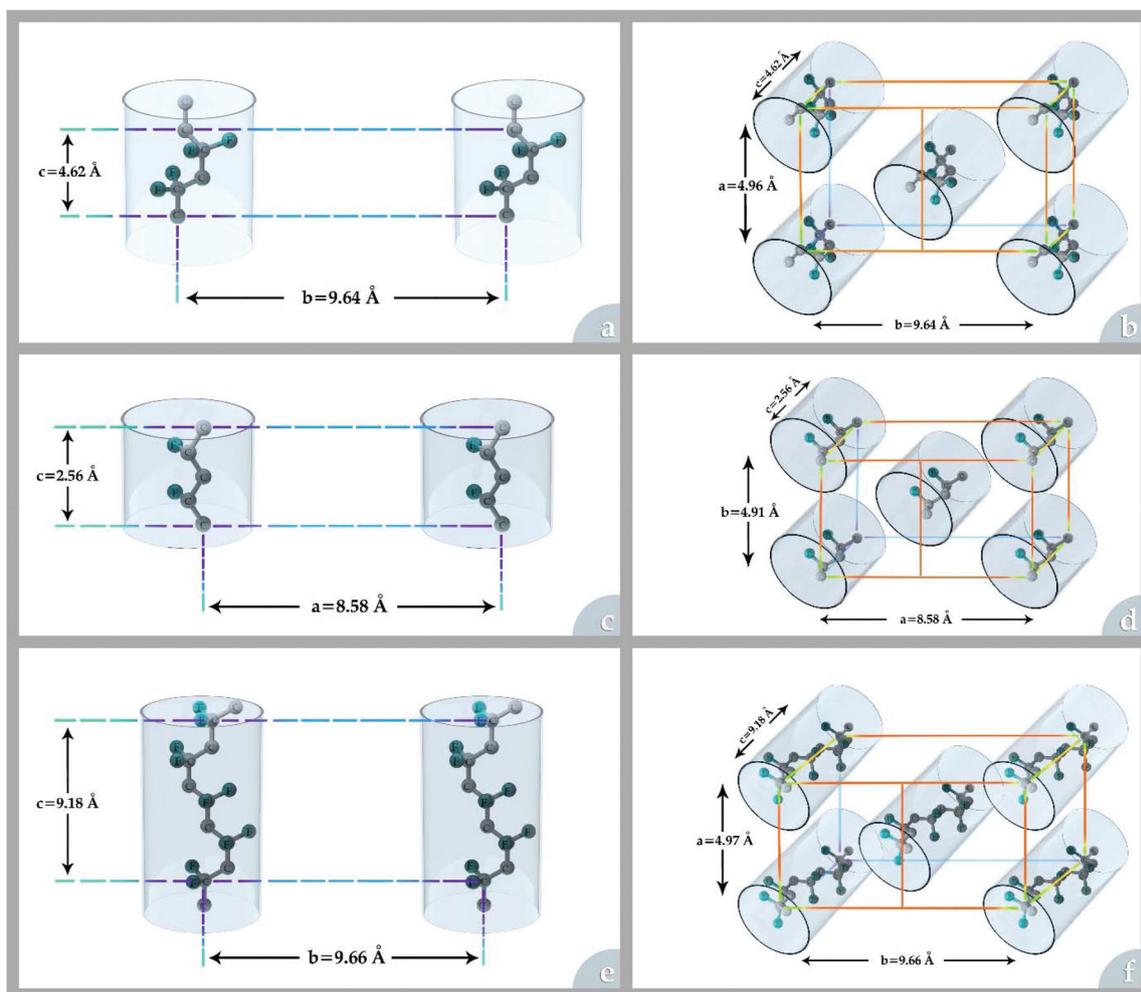


Fig. 1 The conformations and crystal cells for most frequent crystalline polymorphs of PVDF: (a) and (b)  $\alpha$ -phase (c) and (d)  $\beta$ -phase and (e) and (f)  $\gamma$ -phase.

Table 1 Properties of PVDF crystals with different polymorphs<sup>6–8</sup>

Polymorph phase	Phase I ( $\beta$ -phase)	Phase II ( $\alpha$ -phase)	Phase III ( $\gamma$ -phase)
Melting point ( $^{\circ}\text{C}$ )	167–172	167–172	179–180
Crystal system	Orthorhombic	Monoclinic	Monoclinic
Space group	$Cm2m(C(14/2v))$	$P21/c(C(5/2h))$	$C121(C(3/2))$
Polarity	Strong	None	Intermediate
Lattice constants	$a = 8.58 \text{ \AA}$ $b = 4.91 \text{ \AA}$ $c(\text{f.a.}) = 2.56 \text{ \AA}$	$a = 4.96 \text{ \AA}$ $b = 9.64 \text{ \AA}$ $c(\text{f.a.}) = 4.62 \text{ \AA}$ $\beta = 90^{\circ}$	$a = 8.66 \text{ \AA}$ $b = 4.93 \text{ \AA}$ $c(\text{f.a.}) = 2.58 \text{ \AA}$ $\beta = 97^{\circ}$
Electroactivity	High piezo-, pyro-, and ferro-electric	None	Intermediate
Number of chains per lattice	2	2	2
Density ( $\text{g cm}^{-3}$ )	1.81	1.77	1.80
FTIR peak ( $\text{cm}^{-1}$ )	445, 470, 511, 600, 840, 1270	408, 532, 612, 766, 795, 855, 976, 1182, 1400	431, 512, 776, 795, 812, 833, 840, 1233
X-ray diffraction peak ( $^{\circ}$ )	20.26	17.66, 18.30, 19.90, 26.56	18.5, 19.2, 20.4
Molecular conformation	TTT	TGTG'	TTTGTG'
Molecular structure			



by around 90.6% for a CaCO<sub>3</sub> aqueous solution and 53.8% for a Bovine Serum Albumin (BSA) aqueous solution. These results indicate that vibration reduces the fouling potential of piezoelectric PVDF membranes during filtration, regardless of the nature of the fouling materials, but it is most efficient for systems with larger particle sizes (e.g. CaCO<sub>3</sub>). In another work using the same polymeric system, Cao *et al.*<sup>4</sup> observed that applying a mild AC voltage to  $\beta$ -phase PVDF membranes reduced contaminants deposition on the membrane surface and concluded that this was due to current-induced *in situ* vibrations that increased water permeate flux and reduced the tendency for membrane fouling. Chen *et al.*<sup>5</sup> suggested that the vibrations of piezoelectric PVDF membranes might also generate waves in the surrounding medium (e.g. water or air), and developed self-cleaning membranes based on this concept. Particularly when the velocity of vibration is high, such vibrations generate lift forces due to shearing, along with an inertial force acting on the membrane surface and pores. These forces drive the membrane's main cleaning mechanism.

As for other polymeric membranes, the phase inversion method is the most common technique employed for PVDF membrane preparation in both industry and academia. In this method, the desired amount of polymer is dissolved in an appropriate solvent, after which phase separation is induced by an external factor. Commonly used phase inversion methods include nonsolvent induced phase separation (NIPs), temperature induced phase separation (TIPs), vapor induced phase separation (VIPs) and evaporation induced phase separation (EIPs).

The choice of solvent used for the dissolution of PVDF has become a critical issue in recent years. An ideal solvent should have good compatibility with the polymer, a high boiling point, a low molecular weight (to facilitate extraction of residual solvent from membrane structure), while also being inexpensive and overall environmentally friendly. The later property is critical for sustainable membrane production because the most popular solvents for fabricating PVDF membranes are toxic and harmful to the environment. Xie *et al.*<sup>9</sup> evaluated the sustainability of membrane technology and analyzed the parameters that negatively affected it. The three solvents most commonly used in industrial membrane production at present are *N*-methyl-2-pyrrolidone (NMP), *N,N*-dimethylformamide (DMF), and *N,N*-dimethylacetamide (DMAC). According to the International Agency for Research on Cancer (IARC),<sup>10</sup> all three are potential teratogenic compounds that may cause congenital fetal malformation.<sup>11</sup> Other commonly used solvents include diphenyl ketone (DPK), which tends to form carcinogenic compounds under UV radiation,<sup>12</sup> and phthalate-based solvents, most of which accumulate in biological tissues.<sup>13,14</sup> The need to move away from these harmful solvents is amplified by the large volume of wastewaters generated during industrial membrane fabrication, which exceeds 50 billion liters annually.<sup>15</sup>

During the past decade, several green solvents have been proposed for PVDF membrane fabrication, including glycerol triacetate (triacetin),<sup>16,17</sup> maleic acid dibutyl ester,<sup>18</sup> triethylene glycol diacetate,<sup>19</sup> acetyl tributyl citrate,<sup>11</sup> tributyl

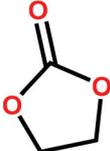
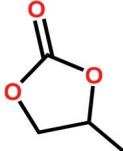
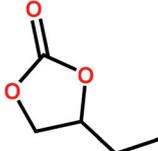
citrate,<sup>20</sup> methyl-5-(dimethylamino)-2-methyl-5-oxopentanoate (Polarclean®),<sup>21,22</sup> Cyrene®,<sup>23</sup> dimethyl isosorbide,<sup>24</sup> and ionic liquids (ILs).<sup>25</sup> Wu *et al.*<sup>26</sup> adopted the strategy of mixing the toxic solvent with a green solvent to decrease the negative environmental impact of the preparing ultrafiltration membranes. However, two factors must be considered when evaluating these reports and the suitability of green solvents for PVDF membrane fabrication. First, many of these studies considered only membrane fabrication without addressing the overall solvent life cycle. Second, many of these solvents are only available in limited quantities and are costlier than toxic solvents.<sup>27</sup> Therefore, they are often only suitable for smaller industrial applications involving high value products. For example, ILs offer tunable properties based on the wide range of cations and anions from which they can be prepared. Although all ILs have negligible vapor pressure and are non-flammable, their toxicity is poorly characterized, which is a significant problem for their potential use as green solvents.<sup>28</sup> Furthermore, their biodegradability depends strongly on the properties of their constituent cations and anions.<sup>29</sup> Another barrier to large-scale industrial uses of ILs is the difficulty and cost of their synthesis.

Organic carbonates, which have been known since the 1950s, are an interesting alternative to the above mentioned solvents. They are available in large quantities at low prices (the total worldwide production of organic carbonates in 2007 was around 1 megaton). They are the esters of carbonic acid that exhibit low (eco)toxicity and complete biodegradability.<sup>30,31</sup> Unfortunately, their uses in membrane preparation are underexplored. To date, studies regarding organic carbonates are limited to propylene carbonate (PC). For instance, Su *et al.*<sup>32</sup> and Yang *et al.*<sup>33</sup> studied the PVDF membrane formation by using several solvents, including PC. Cui *et al.*<sup>34</sup> studied the effect of diluents mixture (PC/dioctyl terephthalate) on PVDF hollow fiber membrane characteristics. Fang *et al.*<sup>35</sup> used a mixture of PC and diphenyl carbonate for polymer dope solution preparation and PC solution as an extruding solvent for the outer layer to prepare dual-layer PVDF hollow fiber membrane. Recently, Rasool *et al.*<sup>36</sup> studied the potential of organic carbonates for membrane preparation *via* NIPs method using various polymers (not including PVDF) with organic carbonates and mixtures of organic carbonates with the toxic solvent NMP. However, PVDF was not used in their study although it is one of the popular polymers industrially employed for the fabrication of membranes. Meanwhile, the study by Yadav *et al.*<sup>27</sup> has confirmed that membrane fabricated with ethylene carbonates exhibits the lowest environmental impact and cost as compared with PVDF with toxic solvents (NMP, DMAC and DMF). Moreover, in PVDF membrane literature, there is still a lack of information about the formation of PVDF membranes, particularly  $\beta$ -phase PVDF, in solvents such as cyclic organic carbonates, and the effect of the solvent's chemical structure on PVDF polymorphism.

In this study, we propose the preparation of PVDF membranes using three green, cheap, and biodegradable cyclic carbonate solvents: ethylene carbonate (EC), PC, and butylene car-



Table 2 Cyclic carbonate solvents used to dissolve PVDF and their properties

			
	Ethylene carbonate	Propylene carbonate	1,2-butylene carbonate
Density at 25 °C (g cm <sup>-3</sup> )	1.3312	1.1981	1.14
Melting point, °C	36.3	-55.0	-45.0
Boiling point, °C	248.2	241.7	238.7
Solubility in water at 25 °C	Soluble	Soluble	Low solubility
Dispersion Force ( $\delta_D$ , MPa <sup>0.5</sup> )	19.4 <sup>a</sup>	20 <sup>a</sup>	16.9 <sup>b</sup>
Polar Force ( $\delta_P$ , MPa <sup>0.5</sup> )	21.7 <sup>a</sup>	18.0 <sup>a</sup>	18.9 <sup>b</sup>
Hydrogen bond force ( $\delta_H$ , MPa <sup>0.5</sup> )	5.1 <sup>a</sup>	4.1 <sup>a</sup>	9.1 <sup>b</sup>
R <sub>a</sub>	11.40	9.94	6.52

<sup>a</sup> Data collected from Hansen Solubility Parameters: A User's Handbook.<sup>37</sup> <sup>b</sup> Data collected from Chernyak.<sup>38</sup>

bonate (BC) by a combined NIPs and TIPs method. The chemical structures and properties of these solvents are presented in Table 2. Particular emphasis is placed on identifying a scalable approach for producing pure  $\beta$ -phase PVDF membranes. The mechanism of  $\beta$ -phase formation in carbonate solvents is investigated experimentally and confirmed by molecular dynamics simulation. The physicochemical properties of the as-prepared membranes are studied and the most promising membrane application for distilled water production by direct contact membrane distillation (DCMD) is investigated and compared with previous publications.

## 2. Materials and methods

### 2.1. Materials

Polyvinylidene fluoride (PVDF) polymer (Solef 1015, MW = 534 000 g mol<sup>-1</sup>) was kindly supplied by Solvay Specialty Polymers. The cyclic organic carbonate solvents EC, PC, and BC were purchased from Tokyo Chemical Industry Co. Ltd and used without purification. Kerosene was purchased from Sigma Aldrich Inc. and used as a wetting agent for porosity measurement.

### 2.2. Membrane preparation

Polymer solutions were prepared by dissolving the appropriate amount of PVDF polymer (15, 17.5 and 20 wt%) in a cyclic organic carbonate solvent (EC, PC, or BC). The solution was then stirred at 130 °C until it became clear and homogeneous, after which it was cast on a glass plate kept at 50 °C using an automatic film applicator (200  $\mu$ m thickness) and immersed in a coagulation bath held at 20 °C. After coagulation, the produced membranes were rinsed with distilled water and dried for 24 h.

### 2.3. Membrane characterization

A Field Emission Scanning Electron Microscope (FE-SEM) from Carl Zeiss was used to obtain membrane surface and cross-section images. The membrane samples were freeze-

cracked in liquid nitrogen and mounted on conductive carbon tape. To reduce sample charging under the electron beam, samples were coated with a 5 nm thick palladium layer using a Quorum Q150T-ES Sputter Coater. The SEM images were analyzed by using ImageJ software in order to determine the diameter of the spherulites. At least three SEM images for each membrane were analyzed, and the diameters of a total number of 100 spherulites were measured. Statistical analysis was applied in order to determine the spherulites diameter by Normal (Gaussian) Distribution (*i.e.* the histogram of the spherulites diameter).

Pure water permeability (PWP) tests were conducted at room temperature using a dead-end Amicon stirred cell. The water fluxes through the membranes were measured at a trans-membrane pressure of 1 bar. The effective diameter of the filtration area was 40 mm. Due to the strong hydrophobicity of PVDF, the investigated membranes were first immersed in an ethanol solution and then washed with water prior to the PWP experiment. Initially, the membrane was pre-compacted at a pressure above 1 bar for 10 min to establish a stable permeate flux before testing. The average water permeate flux was obtained using three membrane samples prepared under the same conditions, and each membrane was tested at least three times. The PWP (L m<sup>-2</sup> h<sup>-1</sup> bar<sup>-1</sup>), was determined using the following equation:

$$\text{PWP} = \frac{Q}{A \cdot t \cdot \Delta P} \quad (1)$$

where  $Q$  is the volume of water permeated through the membrane (in liters),  $A$  is the effective filtration area of the membrane (m<sup>2</sup>),  $t$  is the time corresponding to the volume of collected water (in hour) and  $\Delta P$  is the operating transmembrane pressure fixed at 1 bar for all measurements.

After measuring the dry weight of the membranes, they were wetted by immersion in kerosene for 24 h, after which their wet weight was determined using the following equation:

$$\varepsilon (\%) = \frac{\frac{(m_w - m_d)}{\rho_k}}{\frac{(m_w - m_d)}{\rho_k} + \frac{m_d}{\rho_p}} \times 100\% \quad (2)$$



Here,  $m_w$  and  $m_d$  are the weights of the wet and dry membranes, respectively.  $\rho_k$  and  $\rho_p$  are the density of kerosene ( $0.81 \text{ g cm}^{-3}$ ) and polymer ( $1.78 \text{ g cm}^{-3}$ ), respectively. For each membrane, the porosity was determined from the average of five measurements.

X-Ray Diffraction (XRD) was performed using a PANalytical X'Pert PRO X-ray diffractometer with Cu-K $\alpha$  radiation (wavelength:  $0.154 \text{ nm}$ ) operated at  $40 \text{ kV}$  and  $40 \text{ mA}$ . The samples were scanned in the  $2\theta$  range of  $10^\circ$  to  $40^\circ$  at a rate of  $0.5^\circ \text{ min}^{-1}$  using a silicon-based specimen holder. The  $1/2^\circ$  divergence slit size was fixed for all measurements.

Fourier transform infrared spectroscopy of the membranes was carried out using a Bruker (Vertex 80) spectrometer in the range of  $600\text{--}1600 \text{ cm}^{-1}$  with a resolution of  $2 \text{ cm}^{-1}$ , using the ATR (attenuated total reflection) transmission mode.

A differential scanning calorimeter (DSC, Q-20, USA) was used to measure the crystallization temperature ( $T_c$ ) of the PVDF/solution samples. To this end, PVDF polymer was first dissolved in the appropriate cyclic carbonate solvent (EC, PC, or BC) at the desired concentration (15, 17.5, or 20 wt%) at  $150 \text{ }^\circ\text{C}$  until a homogenous solution formed. The sample was placed in an enclosed specific experimental pan and heated to  $180 \text{ }^\circ\text{C}$  for  $10.0 \text{ min}$ , then cooled to  $20 \text{ }^\circ\text{C}$  at  $10 \text{ }^\circ\text{C min}^{-1}$ . The  $T_c$  value was then determined by software analysis.

The Liquid Entry Pressure (LEP) is the minimum pressure that must be applied over a liquid before it penetrates into non-wetted (dry) membrane pores. LEP measurements for distilled water and  $30 \text{ g L}^{-1}$  NaCl aqueous solution were performed using a dead-end filtration module. The applied pressure was gradually increased by  $5 \text{ kPa}$  every  $4 \text{ min}$  using nitrogen gas ( $\text{N}_2$ ). Mean values and standard deviations were obtained by testing three different samples taken from different parts of the same membrane.

The "bubble-point" test was used to determine the maximum pore size of a porous membrane using the standard test method described elsewhere.<sup>39</sup> The gas pressure needed for a liquid to penetrate a pore is inversely proportional to the pore size. Thus, for a porous membrane, liquid penetration occurs first through the largest pores. As the gas pressure rises, water is gradually pushed into the largest pore of the membrane. Once the gas pressure becomes high enough to overcome the capillary force of the liquid inside the pores, the liquid appears on the other side of the membrane. In this case, the Young-Laplace equation is used to calculate the maximum pore diameter  $d_{p, \text{max}}$  ( $\mu\text{m}$ ):

$$d_{p, \text{max}} = 4\gamma \cos \theta / P \quad (3)$$

where  $\gamma$  ( $\text{N m}^{-1}$ ) is the liquid's coefficient of surface tension,  $\theta$  ( $^\circ$ ) is the contact angle of the liquid on the pore wall, and  $P$  (Pa) is the liquid entry pressure. The pore structure is usually irregularly shaped, so the size of the pores is defined as the diameter of a circle whose area is equal to the pores' cross-

section. When the wetting effect of the liquid is perfect, the angle  $\theta$  is assumed to be  $\sim 0$ , so eqn (5) is simplified to:

$$d_{p, \text{max}} = \frac{4\gamma}{P}. \quad (4)$$

## 2.4 Molecular dynamics simulation

Molecular dynamics (MD) simulations were performed to investigate the polymorphism of PVDF polymers in cyclic carbonate solvents (EC, PC, and BC). In the first step, the solubility parameter and density of the pure solvents were estimated. The formation of  $\beta$ -phase and  $\alpha$ -phase PVDF in the three solvents was then investigated, and the simulations' results were analyzed by considering the Radial Distribution Function (RDF), the relative concentration, and hydrogen bonding.

The simulations were performed with Materials Studio package (BIOVIA Accelrys, version 4.3) using the COMPASS force field (Condensed phase Optimized Molecular Potentials for Atomistic Simulation Studies).<sup>40</sup> Previous studies have shown that this force field gives acceptable results for polymers and their solvents.<sup>41–44</sup>

For each solvent, a simulation box containing 100 solvent molecules with an initial density of  $0.4 \text{ g cm}^{-3}$  was constructed and equilibrated for  $1 \text{ ns}$  to estimate the density and solubility parameters of the pure solvents in the NPT ensemble (*i.e.* with a constant number of molecules, temperature, and pressure) at  $1 \text{ bar}$  (Berendsen barostat<sup>45</sup>) and  $300 \text{ K}$  (Nose thermostat<sup>46</sup>). Periodic boundary conditions, the Verlet algorithm,<sup>47</sup> and a time step of  $1 \text{ fs}$  were used in all MD simulations.

The simulation boxes containing PVDF surfaces in contact with the solvent were constructed using the equilibrium density of the solvent to investigate the interactions between the solvent and the PVDF surface. For this purpose, supercells for different crystalline polymorphs (*i.e.*,  $\alpha$ -PVDF and  $\beta$ -PVDF) were created using the appropriate crystalline coordinates (see Table 1), repeating periodically along the  $x$ ,  $y$  and  $z$ -axes. The corresponding crystal sizes are listed in Table S.1,<sup>†</sup> and details of the simulation box setup are presented in Table S.2.<sup>†</sup> As an example, the simulation boxes for the  $\alpha$  and  $\beta$  PVDF phases in contact with the EC are shown in Fig. 2.

The simulation boxes were equilibrated using the NVT ensemble (*i.e.* a constant number of molecules, volume, and pressure) for  $300 \text{ ps}$ , after which the MD simulation was run for  $10 \text{ ns}$  using the NPT ensemble at  $300 \text{ K}$  and  $1 \text{ bar}$  to obtain the final equilibrium PVDF surface and solvent molecule distribution. Structures were sampled at  $10 \text{ ps}$  intervals along the MD trajectory. Finally, all possible hydrogen bonds between the solvent molecules and the PVDF surface were calculated.<sup>48</sup>

The interactions of EC, PC, and BC with the PVDF surface were evaluated by computing the RDF at  $300 \text{ K}$ . The RDF represents the probability density of finding atoms  $j$  at a distance  $r$  from atom  $i$ , averaged over the equilibrium density, and can be expressed as follows:

$$g_{ij}(r) = \frac{d(N_{ij}(r))}{dV(r)} \quad (5)$$



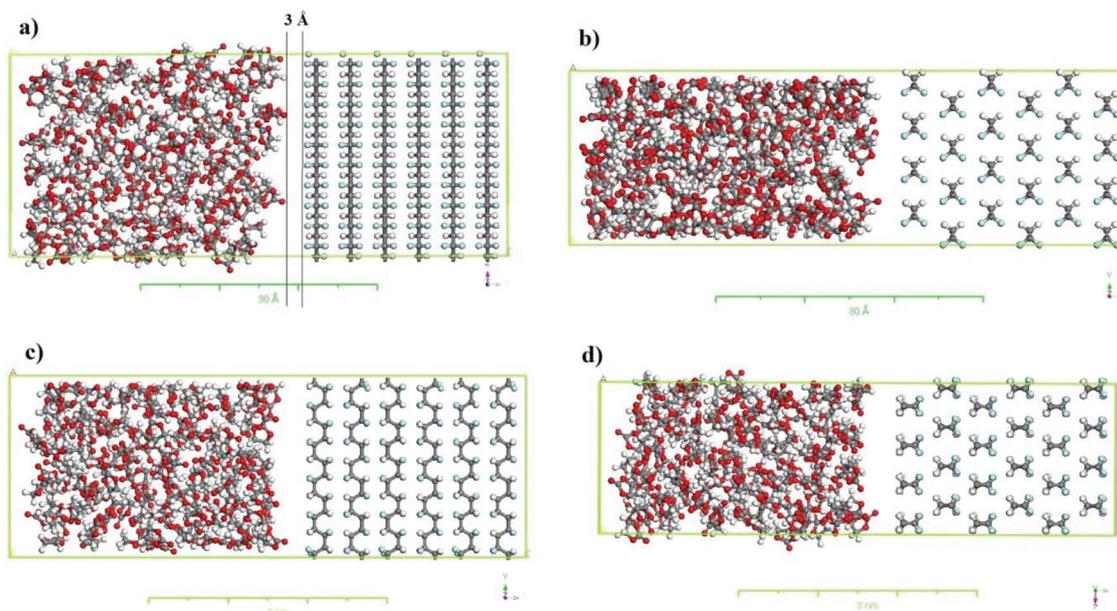


Fig. 2 Simulation box for  $\beta$ -phase PVDF in EC (a) front view and (b) top view; and  $\alpha$ -phase PVDF in EC (c) front view and (d) top view. Carbon, fluorine, oxygen, and hydrogen are shown in grey, cyan, red, and white, respectively.

where  $d(N_{ij}(r))$  is the average number of times the  $j$  atoms are found in a spherical shell of thickness  $dr$  at a distance  $r$  from atoms  $i$ .

### 2.5. Direct contact membrane distillation (DCMD)

The membrane with the highest porosity and pure water permeability was used for desalination by DCMD using a lab-scale set-up (Fig. S1†). The membrane was placed between two chambers of a stainless steel flat sheet module (model CPR920, Convergence Industry B.V; Netherlands). DCMD was initially performed using distilled water with an electrical conductivity below  $4.5 \mu\text{S cm}^{-1}$  to detect potential leakage in the set-up. The permeate flux ( $J_w$ ) was obtained from eqn (6) by monitoring the mass of permeate over time using a precision balance (PCE Iberica S.L. Instruments, model PCE-BSH 10000) connected to a computer:

$$J_w = \frac{Q}{A \cdot t} \quad (6)$$

where  $Q$  is the volume of water permeated through the membrane (in kg),  $A$  is the effective membrane superficial area ( $0.006 \text{ m}^2$ ), and  $t$  is the time corresponding to the volume of collected water (in hour).

The feed and permeate temperatures were controlled by an electromantle thermostat (model EMX5000/SCE, Electrothermal; UK) and a recirculation chiller (model CFT-75, VESLAB instruments. Int.; USA), respectively. The feed and distillate were circulated co-currently through their respective channels using Magnetic Coupling Water Pumps (Xylem Flojet, HPR6/8, Totton Pumps Ltd, UK). TMU thermometers (Papouch store s.r.o.; Czech Republic) with USB interfaces were installed at both the inlets and outlets of the membrane

module and were connected to a computer, in which a software automatically recorded the inlet and outlet temperatures every 30 s. The permeate inlet temperature ( $T_{p,\text{in}}$ ) was maintained at  $20 \text{ }^\circ\text{C}$ , while the feed inlet temperature ( $T_{f,\text{in}}$ ) was varied from  $40$  to  $80 \text{ }^\circ\text{C}$ . Distilled water and an aqueous solution of  $30 \text{ g L}^{-1}$  NaCl were used as feed. Both the feed and permeate flow rates were kept constant at  $\sim 0.51 \text{ L min}^{-1}$ . The electrical conductivity of both the feed and permeate solutions was measured with a conductivity meter (edge, model HI2003, Hanna instruments Inc., USA) and used to determine the corresponding salt concentrations and salt separation factor ( $\beta$ ) based on following equation:

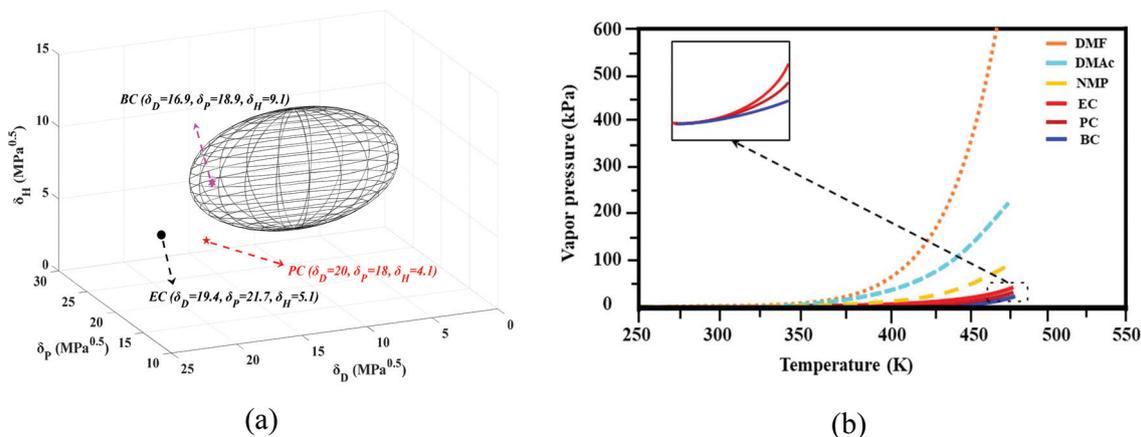
$$\beta(\%) = \left( 1 - \frac{\frac{C_{p,\text{initial}} + C_{p,\text{final}}}{2}}{\frac{C_{f,\text{initial}} + C_{f,\text{final}}}{2}} \right) \cdot 100 \quad (7)$$

where  $C_{p,\text{initial}}$ ,  $C_{p,\text{final}}$ ,  $C_{f,\text{initial}}$ , and  $C_{f,\text{final}}$  are the initial and final NaCl concentrations in the permeate and the feed solutions, respectively.

## 3. Results and discussion

The group contribution method was used to determine the solubility parameters of the cyclic organic carbonates (EC, PC, and BC) for PVDF. Based on these parameters, it is expected that PVDF, which has a solubility parameter of  $\delta = 23.2 \text{ MPa}^{0.5}$  ( $\delta_D = 17.1 \text{ MPa}^{0.5}$ ,  $\delta_P = 12.6 \text{ MPa}^{0.5}$ ,  $\delta_H = 10.6 \text{ MPa}^{0.5}$ ,  $R_0 = 5$ )<sup>49–52</sup> should be soluble in BC but not in PC and EC (see Fig. 3a). However, it is worth considering the impact of the temperature on solvation and the accuracy of the Hansen solubility parameter. As noted by other researchers, calculated





**Fig. 3** (a) Solubility region of PVDF in the Hansen space (hydrogen bonds ( $\delta_H$ ), intermolecular force ( $\delta_P$ ) and dispersion force ( $\delta_D$ )) and the positions of ethylene carbonate (EC), propylene carbonate (PC), and butylene carbonate (BC) and (b) vapor pressure of some popular polar aprotic solvents for membrane preparation and cyclic carbonates examined in this study.

solubility parameters represent only rough approximations of a system's actual behavior.<sup>36</sup> In this study, increasing the temperature enhanced the solvation power of the studied cyclic carbonates. The three solvents (EC, PC, and BC) dissolved PVDF at elevated temperatures (130 °C), allowing membrane formation to be achieved by the TIPs method. These solvents are water-soluble, allowing phase inversion to be induced by solvent/nonsolvent exchange (NIPs). Consequently, both the TIPs and NIPs mechanisms contributed to the studied membrane formation process.

A key advantage of cyclic carbonates in membrane preparation is their low vapor pressure. Fig. 3b compares the vapor pressure of some polar aprotic solvents that are widely used in membrane preparation to that of cyclic carbonates (the thermodynamic correlations used to estimate the vapor pressure are presented in the ESI together with Table S.3†). The vapor pressures of EC, PC, and BC are much lower than those of the polar aprotic solvents, indicating that cyclic carbonates are preferable from both safety and sustainability perspectives.

### 3.1 Membrane characteristics

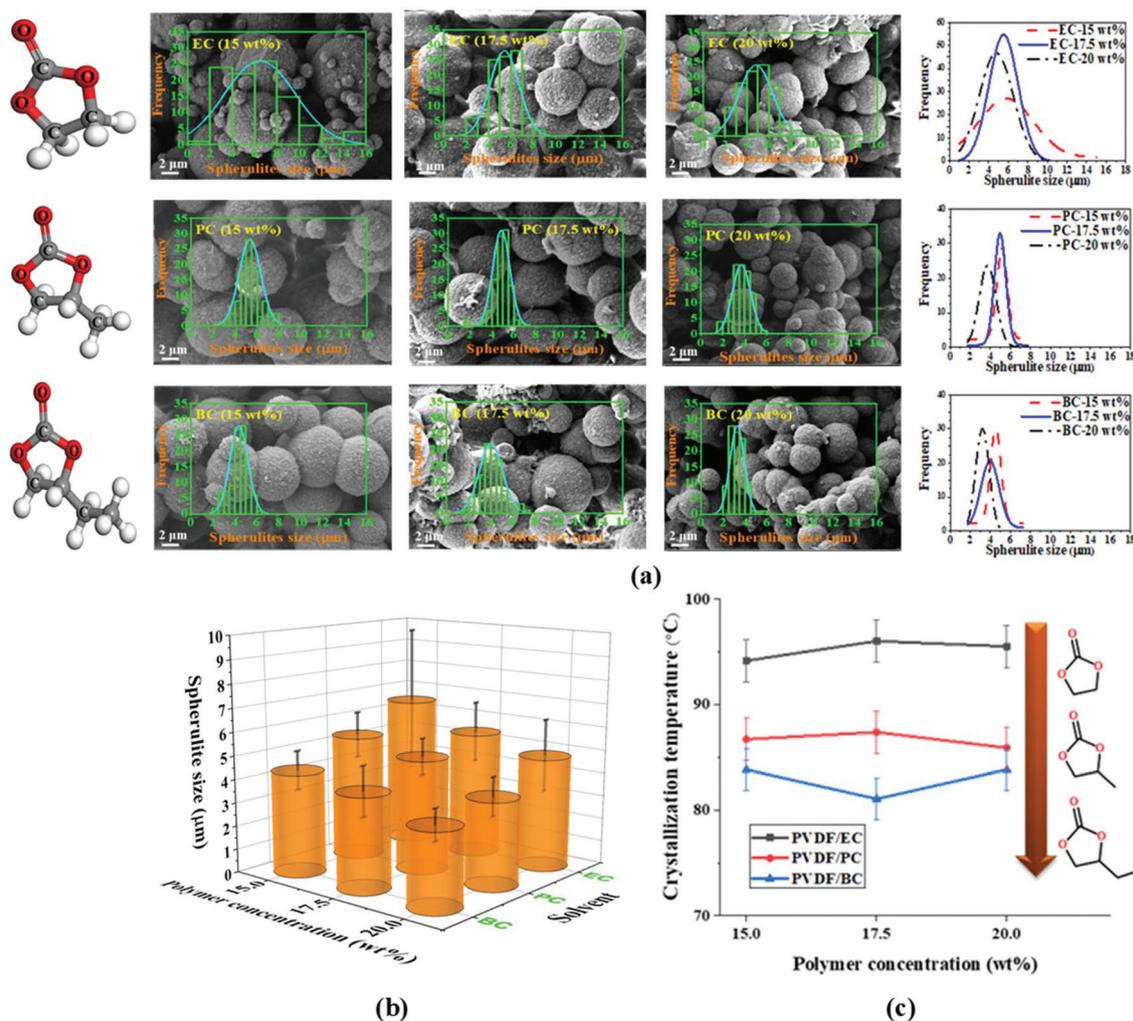
The TIPs process is a non-equilibrium process, so its analysis requires consideration of temperature effects. In particular, the cooling conditions are a key kinetic parameter that determines the spherulitic structure of the formed PVDF membrane. In this study, we kept the polymer solution and quenching bath temperature constant, at room temperature  $20 \pm 2$  °C, for two reasons. First, other research studies have already investigated the effects of varying these parameters,<sup>11,19,53,54</sup> and second the change of the quenching bath temperature affects energy consumption and environmental impact of membrane fabrication, both of which are important for sustainable membrane fabrication.

All prepared membranes in this study exhibit a spherulitic structure. Fig. 4a and b shows the morphology, spherulitic size, and spherulitic distribution of the PVDF membranes pre-

pared with cyclic carbonate solvents. This can be attributed to the phase inversion mechanism (*i.e.*, solid-liquid phase inversion) of semi-crystalline PVDF in cyclic carbonate solvents. Due to the favorable interaction between PVDF and the solvents (EC, PC, and BC), PVDF crystallization occurs before liquid-liquid phase separation, resulting in a spherulitic structure. Fig. 4c shows the crystallization temperature of PVDF in EC, PC, and BC at different concentrations (15, 17.5, and 20 wt%). A decrease of the crystallization temperature can be seen with the increase of the alkyl chain length of the cyclic organic carbonate solvents but it maintains almost the same with the increase of the PVDF concentration.

The nucleation density is an important parameter in solid-liquid phase inversion because spherulites form by growing from a nucleus into lamellae that in turn gives rise to a sheaf-like structure known as axialites, which finally develop into spherulites.<sup>55</sup> For the same solvent, the nucleation density is directly proportional to the polymer content of the solution. The effect of the polymer concentration (15, 17.5, and 20 wt%) on the morphological structure of the PVDF membrane is presented in Fig. 4a and b. As can be seen, the nucleation density clearly increases with the increase of the polymer concentration, and the final spherulite size decreases with the increase of the nucleation density. The reduction in spherulite size and the volume of the inter-spherulite spaces is confirmed by a concomitant decrease in the overall membrane porosity, as shown in Fig. 5a. Similar trends were observed in other studies for PVDF membranes prepared with toxic solvents.<sup>56</sup> Additionally, spherulites formed at higher polymer concentrations exhibited more uniform structures. The polymer's crystallization depends on the magnitude of the crystallization driving force (*i.e.* the difference in the polymer's chemical potential in the crystalline and solution phases). At low polymer concentrations, the driving force is not strong enough to induce the growth of all formed nuclei, resulting in the formation of some fully developed spherulitic with some smaller spherulitic structures (Fig. 4a and b).





**Fig. 4** (a) Morphologies together with histograms and comparison of normal Gaussian distribution of the spherulites diameter (Left) when using ethylene carbonate (EC), propylene carbonate (PC) and butylene carbonate (BC); (b) 3D graph comparing the spherulitic size and (c) crystallization temperatures of PVDF membranes prepared with different polymer concentrations and solvent alkyl chain lengths, and (c) crystallization temperature of PVDF membranes prepared with different polymer concentrations and solvent alkyl chain lengths.

Changing the solvent's chemical structure affected the final morphological structure of the membrane as shown in Fig. 4a. Specifically, increasing the solvent's alkyl chain length increased the solvent's compatibility with PVDF (Fig. 4a). This in turn decreased the crystallization temperature (Fig. 4c) and strengthened the thermodynamic driving force of nucleation, leading to an increased nucleation density (Fig. 4a). This phenomenon results in a decrease of both the spherulitic size and the inter-spherulitic space. As stated before, these changes in membrane morphology affect the overall porosity. Lengthening the alkyl side chain of the solvent structure reduced the volume of the inter-spherulitic spaces and a decrease of the overall porosity (see Fig. 5a).

The PWP of the prepared PVDF membranes with different solvents and different polymer concentrations is shown in Fig. 5b. As reported previously, longer alkyl side chain of the solvent molecule reduced the spherulitic size, the inter-

spherulitic space, and the overall porosity. This results in a decline of the PWP. A similar trend was observed upon increasing the polymer concentration. Higher polymer concentrations tended to increase the nucleation density, reducing the volume of the inter-spherulitic spaces, overall porosity, and PWP. However, this trend was not followed in the case of EC prepared with 20 wt% PVDF. In this case, the PWP was actually greater than that of the membrane prepared with 17.5 wt% PVDF. This unexpected result can be attributed to a significant reduction of the thickness of the membrane's skin layer at the higher polymer concentration (*i.e.* 20 wt%), which is readily apparent in the SEM images presented in Fig. S.2a.†

### 3.2 Polymorphism experimental observation

The XRD and FTIR spectra of the PVDF membranes prepared with different solvents and polymer concentrations are pre-



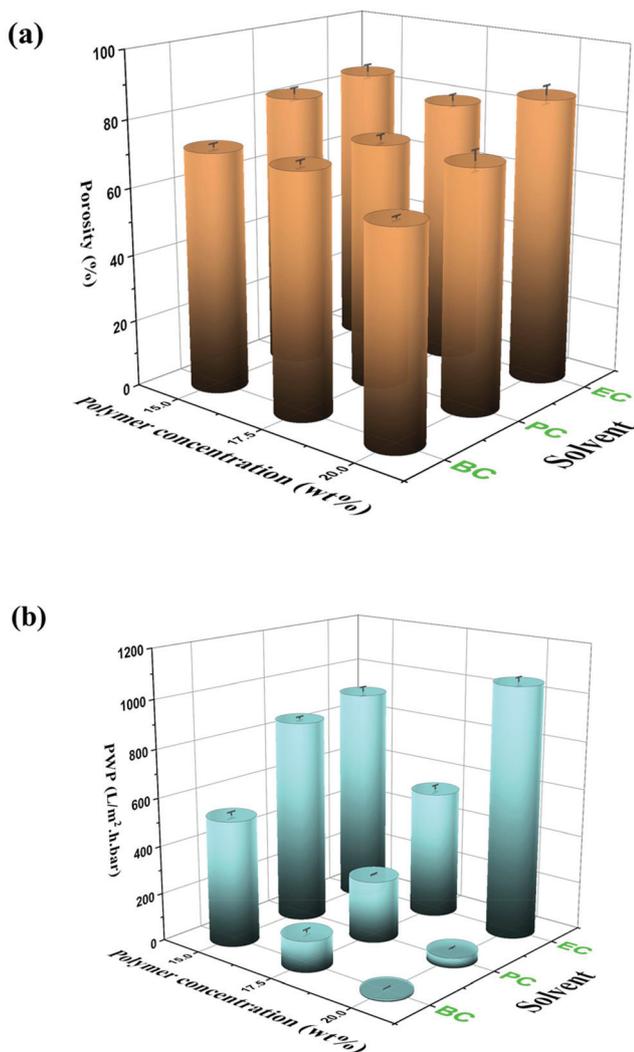


Fig. 5 (a) Porosity and (b) PWP of the PVDF/EC, PVDF/PC and PVDF/BC membranes prepared with different polymer concentrations.

sented in Fig. 6. The  $\beta$ -phase predominates in the membranes prepared with EC and PC solvents because the only visible peak for these membranes is that at  $20.26^\circ$ . The FTIR spectra of these two membranes prepared with EC and PC supported the finding that the  $\beta$ -phase predominated in these cases because their transmittance peaks were registered at  $840 \text{ cm}^{-1}$  and  $1275 \text{ cm}^{-1}$ ,<sup>57,58</sup> indicating  $\beta$ -phase formation. Meanwhile, no peaks were detected at  $488, 532, 615, 766, 795, 854, 976, 1383,$  and  $1423 \text{ cm}^{-1}$ ,<sup>59,60</sup> indicating the absence of the  $\alpha$ -phase. Similar results were obtained for the membranes prepared with different polymer concentrations, indicating that the polymer concentration did significantly affect the polymorph distribution.

The membranes prepared with BC exhibited different trends. The XRD spectra indicated the decrease of the  $\beta$ -phase formation as the polymer concentration was increased from 15 to 20 wt%. There were no clear  $\alpha$  peaks in the FTIR spectra of the membrane formed with 15 wt% polymer concentration,

but the FTIR data clearly indicated the coexistence of the  $\alpha$  and  $\beta$ -phases for the membranes prepared with higher polymer concentrations 17.5 and 20 wt%. Additionally, the relative abundance of the  $\alpha$ -phase increased with the polymer concentration. It may be explained by the relatively long alkyl side chain of BC that weakens the forces favoring oriented dipole packing, preventing adoption of the TTT conformation and leading to a preference for the thermodynamically more stable  $\alpha$ -phase (TGTG) conformation.

The formation of the  $\beta$ -phase can be attributed to the strong electrostatic interaction of the PVDF chain with EC and PC solvents. The polarity of the solvent, the dipole-dipole interactions, and the hydrogen bonding at the interface between PVDF nuclei and the solvent molecules will all favor the conformation that optimizes the packing of the  $\text{CH}_2\text{-CF}_2$  dipoles (*i.e.* the TTT conformation).<sup>61</sup> The role of the solvent polarity in  $\beta$ -phase formation was highlighted by Gregorio *et al.*<sup>62</sup> and Hassankiadeh *et al.*<sup>21</sup> As shown in Fig. 7, cyclic carbonates have greater dipole moments than other solvents commonly used in membrane fabrication. This favors the formation of the all-*trans* (TTT) conformation and subsequently  $\beta$ -phase predomination. Large solvent dipoles will be stabilized by the net dipole of  $\beta$ -PVDF more strongly than by the non-polar  $\alpha$ -phase. In the  $\beta$ -phase, PVDF has a permanent dipole moment of 2.1 Debye per monomer unit, whereas the dipole moment per monomer unit of the  $\alpha$ -phase is only 1.3 Debye.<sup>63</sup> When a solvent with a high dipole moment interacts with PVDF, strong electrostatic interactions will favor the formation of locally extended chain coils. Therefore, the all-*trans* conformation (*i.e.* the  $\beta$ -phase) is readily formed in solvents with high dipole moments. Conversely, when PVDF is dissolved in solvents with a lower dipole moment, the solvent interacts less strongly with PVDF and results in the formation of unexpanded or shrunken globules favoring therefore the twisted  $\alpha$ -phase conformation.<sup>64</sup>

It must be mentioned that hydrogen bonding at the interface of PVDF and solvent is another key factor governing  $\beta$ -phase formation. Interfacial hydrogen bonds are formed between hydrogen or fluorine atoms in PVDF and the solvent. To clarify the contribution of such interfacial hydrogen bonding in the formation of  $\beta$ -phase PVDF in carbonate solvents, MD simulations were performed as stated previously.

### 3.3 Polymorphism simulation observation

The density and the solubility parameter of the pure cyclic organic carbonate solvents were calculated from the results of MD simulations with the COMPASS force field and are shown in Table 3. The simulations' results of the solubility parameter were underestimated by 11.3 to 16.9%; however the density simulations' results agree well with the experimental indicating that MD described the solvents' behaviors reasonably well. The interactions between the solvents (EC, PC, and BC) and PVDF surfaces ( $\alpha$  and  $\beta$ -phase) were then investigated by MD. The relative concentrations of EC and  $\beta$ -PVDF in contact with each other in the simulation box before and after the simulation are shown in Fig. 8a. Before the simulation, the relative



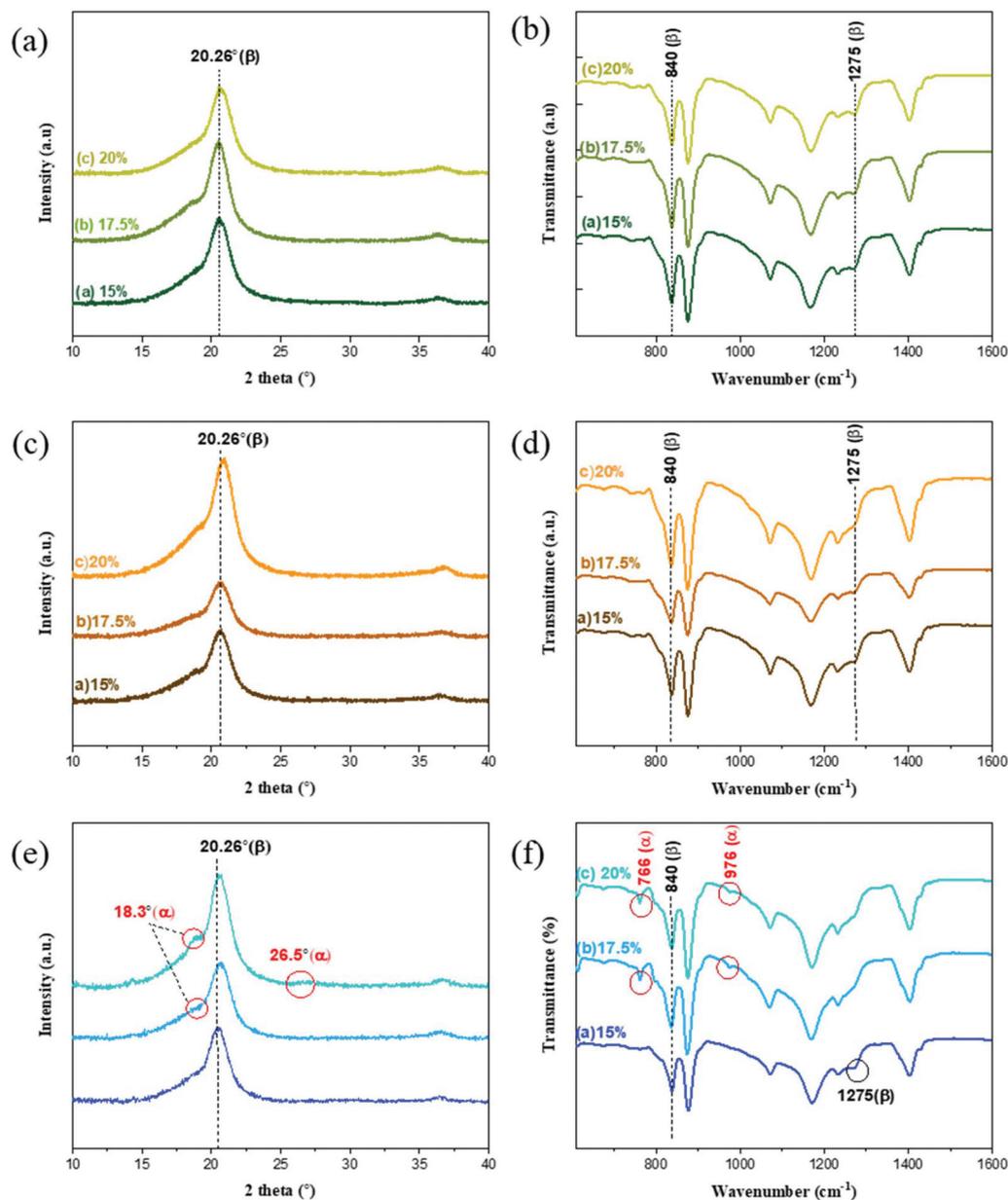


Fig. 6 XRD (left) and FTIR (right) data of the PVDF membranes prepared with different polymer concentrations with (a and b) EC, (c and d) PC and (e and f) BC solvents.

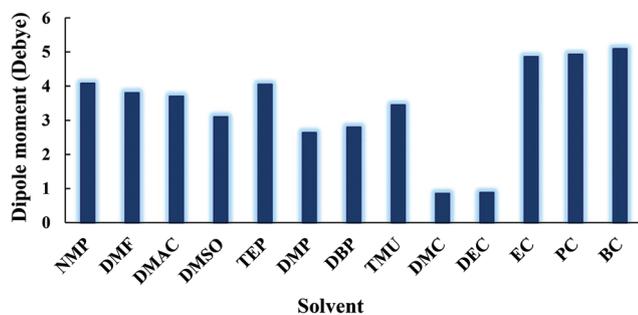


Fig. 7 Dipole moments of various solvents and cyclic organic carbonates.

Table 3 Simulation-based and experimental density and solubility parameter of the three cyclic organic carbonate solvents

Solvent	Density ( $\text{g cm}^{-3}$ )		Solubility parameter ( $(\text{J cm}^{-3})^{0.5}$ )	
	Simulation	Exp.	Simulation	Exp.
EC	$1.27 \pm 0.02$	1.33	$26.22 \pm 0.80$	29.55
PC	$1.17 \pm 0.02$	1.19	$23.63 \pm 0.60$	27.21
BC	$1.11 \pm 0.02$	1.14	$22.38 \pm 0.40$	26.93



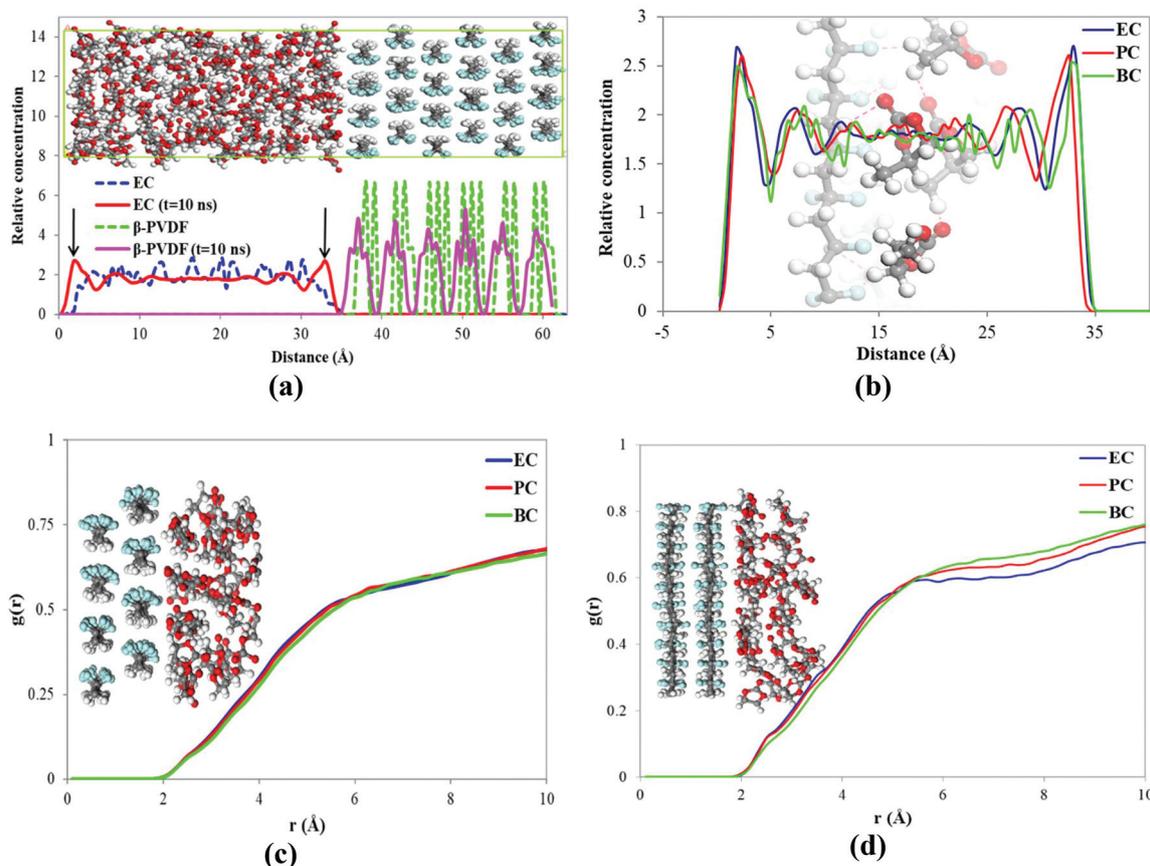
concentration of EC in the simulation box indicates that the solvent molecules were evenly dispersed; their minimum distance from the PVDF surface is 3 Å (see Fig. 2) and there is no sharp peak in their relative concentration diagram (Fig. 8a). However, at the end of simulation, two peaks of 1.79 and 32.99 Å (because of the periodic boundary conditions used in the MD simulation) appeared in the relative concentration of EC (red line), confirming that the solvent molecules oriented themselves close to the  $\beta$  surface. Additionally, there are regular sharp peaks in the relative concentration of solvent molecules in contact with PVDF before the simulation and at its end, confirming that the crystal structure of  $\beta$ -phase PVDF is stable during the MD simulation. A snapshot of the simulation box at the end of simulation (see Fig. 8a) also indicated that the crystal structure of  $\beta$ -PVDF was stable.

Fig. 8b shows the relative concentrations of three different solvents (EC, PC and BC) in contact with the  $\beta$ -phase PVDF surface in the simulation box at the end of the simulation. There are two sharp peaks at 1.79 and 32.99 Å, indicating that solvent molecules accumulated near the  $\beta$ -PVDF surface. This can be attributed to an attraction between the solvent molecules and the  $\beta$ -PVDF surface that is similar in magnitude for all three solvents. The degree of solvent accumu-

lation at the surface decreased in the order EC > PC > BC. This indicates that EC exhibited a greater number of interactions with the  $\beta$ -phase than PC and BC at shorter distances.

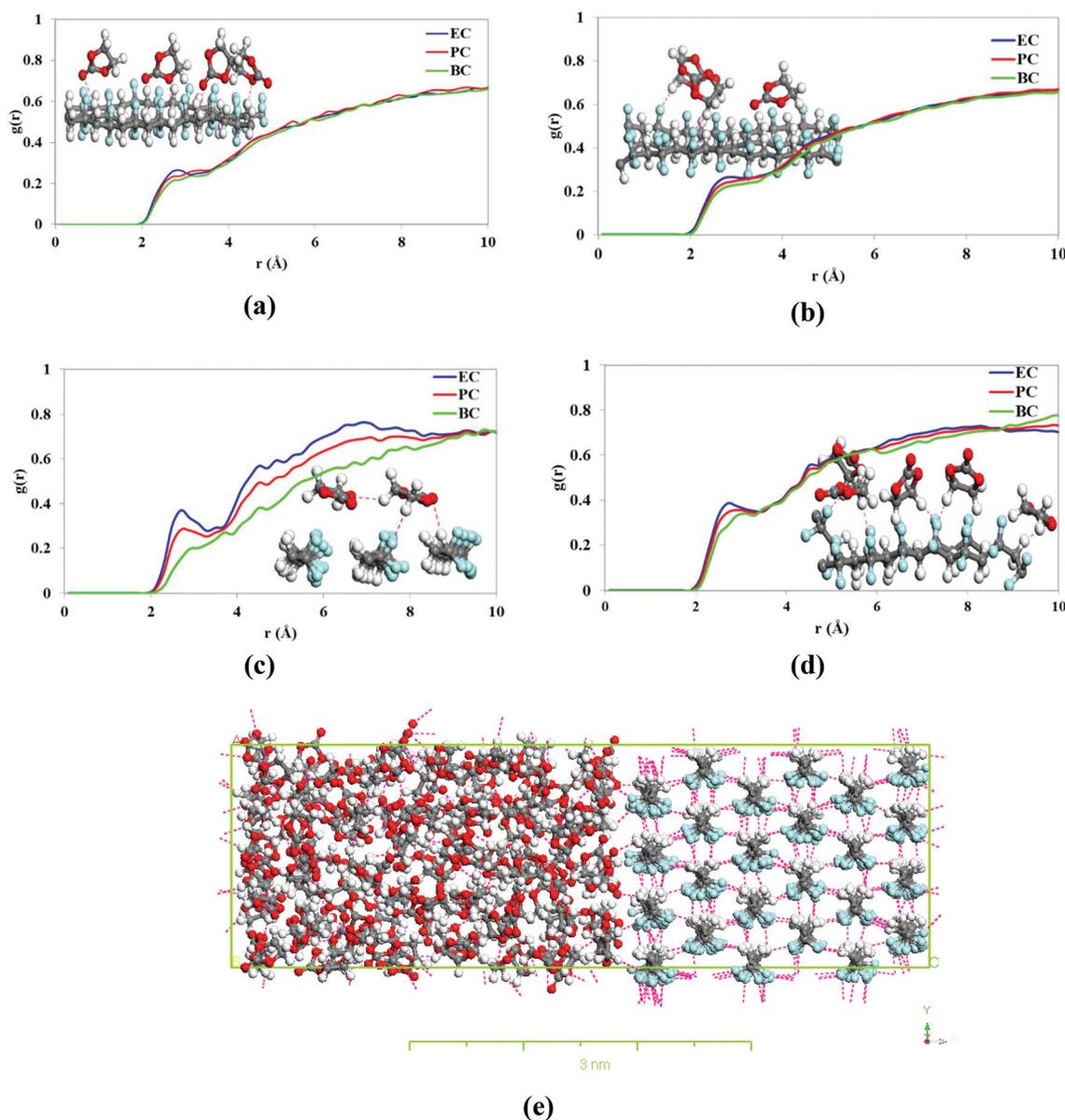
The RDFs of solvent molecules with PVDF surfaces are shown in Fig. 8c and d. There is no sharp peak in the RDF of solvent molecules with the PVDF surface because not surface. However, the PVDF the to access have all solvent molecules could result confirm that a number of solvent molecules are located 1.9 Å away from the PVDF surface, which can be explained by interfacial hydrogen bonding. The tendency of solvent molecules to be found at the PVDF–solvent interface again decreases in the order EC > PC > BC.

The RDFs of solvent atoms with H and F atoms of PVDF chains are shown in Fig. 9. The RDF plot for O atoms in solvent molecules and H atoms of  $\beta$ -PVDF at the interface (Fig. 9a) shows a relatively small peak at 2.7 Å. The largest and smallest peaks at this point were observed for EC and BC, respectively. Similarly, the results plotted in Fig. 9b indicate a relatively small peak at 2.7 Å in the RDFs for solvent H atoms and  $\beta$ -PVDF F atoms and this was larger for EC compared to PC and BC peaks. The RDFs of solvent O and H atoms with the H and F atoms of the  $\alpha$ -PVDF surface displayed in Fig. 9c



**Fig. 8** (a) Relative concentrations of EC in contact with  $\beta$ -phase at the beginning and end of the simulation; (b) the relative concentrations of solvents in contact with the  $\beta$ -PVDF surface along the simulation box at the end of simulation; RDF of solvent molecules with (c)  $\beta$ -PVDF and (d)  $\alpha$ -PVDF surfaces.





**Fig. 9** RDFs of (a) solvent O atoms with H atoms of  $\beta$ -PVDF, (b) solvent H atoms and  $\beta$ -PVDF F atoms, (c) solvent O atoms with H atoms of  $\alpha$ -PVDF, (d) solvent H atoms with  $\alpha$ -PVDF F atoms, and (e) snapshot of the simulation box showing the hydrogen bonding in EC in contact with the  $\beta$ -PVDF surface.

and d also show a relatively short peak at 2.7 Å being the longest and shortest peaks at this distance for EC and BC, respectively. These results confirm that the number of hydrogen bonds between the solvent and the  $\alpha$ -PVDF surface is maximized in EC and minimized in BC. In other words, as the solvent's carbon chain length increases, the accessibility of the surface to the solvent molecules (and thus the number of interfacial hydrogen bonds) decreases. This was also observed for the  $\beta$ -PVDF surface. Therefore, the results in Fig. 9 confirm that the number of hydrogen bonds between the solvent and the PVDF surface is maximized in EC and minimized in BC, for both  $\alpha$ - and  $\beta$ -PVDF. This is consistent with our experimental results and expectations; BC has a longer alkyl side

chain that disrupts interfacial interactions and therefore cannot interact strongly with the PVDF chain and promotes the  $\beta$ -phase orientation.

The simulation box observed for EC in contact with the  $\beta$ -PVDF surface at the end of the simulation shown in Fig. 9e clearly demonstrates extensive hydrogen bonding between solvent molecules and also between solvent molecules at the surface and the polymer chain (pink lines). Previous studies<sup>65</sup> have shown that C–O and C–F moieties are hydrogen bond acceptors that form C–H...F–C and C–O...H–C interactions. The hydrogen bonds between the solvent molecules and the polymer surface are thus formed between O and H atoms of solvent and H and F atoms of  $\beta$ -PVDF.



To evaluate the stability of the PVDF structures in contact with various solvents, the RDFs of the H and F atoms of PVDF chains were calculated and the results are shown in Fig. 10. The RDFs of the  $\beta$ -phase surface in contact with various solvents (see Fig. 10a) were identical, indicating that  $\beta$ -PVDF has a stable structure in all three solvents. Five peaks are observed at the same distances ( $r$ ) for the three solvents, confirming the presence of the  $\beta$ -phase polymorph. However, the intensity of the peaks indicates that the relative abundance of the  $\beta$ -phase content decreases in the order EC > PC > BC. In other words, increasing the solvent's alkyl side chain lengths reduces the intensity of the RDF peak, indicating a reduced relative abundance of  $\beta$ -PVDF.

At the end of the simulation, the RDFs for EC with the two phases ( $\alpha$  and  $\beta$ ) of the PVDF surface were very similar, indicating that over the course of the simulation, the  $\alpha$ -phase was partially transformed to the  $\beta$ -phase when in contact with EC. One would therefore expect PVDF polymer to adopt  $\beta$ -phase crystal conformations in the presence of EC, and that the relative abundance of  $\beta$ -PVDF structure would depend on the solvent's alkyl side chain length.

The number of hydrogen bonds at the surface-solvent interface was estimated based on the number of contacts within 3 Å for each solvent and PVDF polymorph. There was little variation in the numbers of hydrogen bonds formed with EC for the two surfaces due to their similar contact surfaces (Table 4). However, it was clear that the number of hydrogen bonds decreased as the length of the solvent alkyl side chain increased. This can be attributed to the fact that longer alkyl side chains interrupt the interactions of solvent O atoms with PVDF H atoms. Therefore, EC interacts strongly with PVDF chains and promotes their orientation in the  $\beta$ -phase while BC is less effective in this role.

The simulation boxes for the three solvents in contact with the  $\alpha$ -PVDF surface are shown in Fig. S.3.† It is clear that the  $\alpha$ -phase of PVDF is converted into the  $\beta$ -phase and it is stable in that state in the solvent EC, whereas there are depressions and changes in the structure of the  $\alpha$ -phase PVDF surface in PC and BC. In addition, the results presented in Fig. S.3.† also

**Table 4** Numbers of hydrogen bonds at the surface-solvent interface for different PVDF polymorphs and solvents

Solvent	Number of interfacial hydrogen bonds	
	$\alpha$ -PVDF	$\beta$ -PVDF
EC	75	76
PC	72	74
BC	71	69

confirm that the PVDF polymer chain in contact with EC always adopts the  $\beta$ -phase structure, and that the polymer's  $\beta$ -phase ratio decreases as the solvent alkyl side chain length increases.

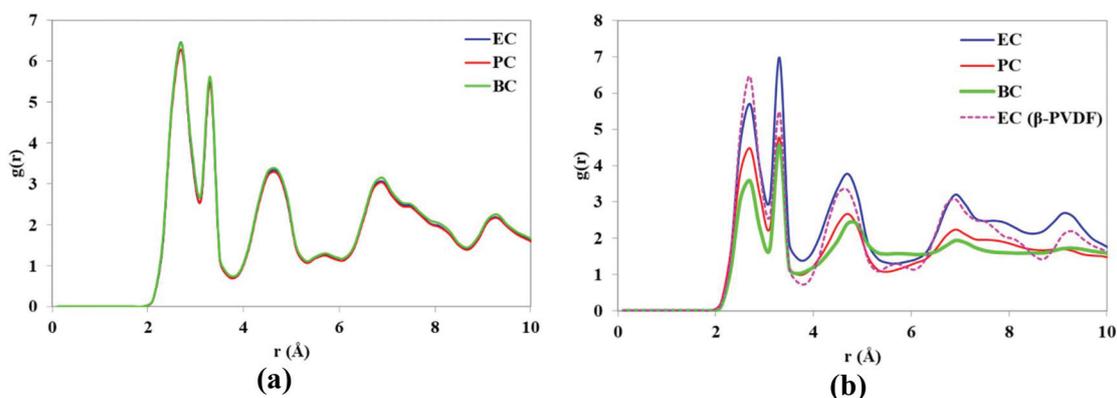
### 3.4 DCMD experiments

Among all prepared PVDF membranes with different cyclic carbonate solvents, the one prepared with EC and a polymer concentration of 20 wt% (PVDF/EC-20) exhibited the highest porosity and PWP. Its DCMD performance was therefore investigated. Table 5 shows the measured LEP values of this PVDF/EC-20 membrane for distilled water and 35 g L<sup>-1</sup> NaCl aqueous solution. These data are comparable to those of commercial membranes commonly used in membrane distillation confirming the suitability of the membrane PVDF/EC-20 in this technology. For instance, TF1000 (Gelman) and FALP (Millipore) have LEP values of 48 and 48.3 kPa, respectively.<sup>66</sup>

Fig. 11(a) shows the effects of the inlet feed temperature on the DCMD permeate flux ( $J_w$ ) and the salt rejection factor ( $\beta$ ) when using as feed distilled water and NaCl (35 g L<sup>-1</sup>) aqueous

**Table 5** Characteristics of the PVDF membrane prepared with the cyclic organic solvent EC and 20 wt% PVDF polymer (PVDF/EC-20): total thickness ( $\delta$ ), liquid entry pressure (LEP), and maximum pore size  $d_{p, \max}$

$\delta$ ( $\mu\text{m}$ )	LEP <sub>H<sub>2</sub>O</sub> (kPa)	LEP <sub>35 g L<sup>-1</sup> (NaCl)</sub> (kPa)	$d_{p, \max}$ ( $\mu\text{m}$ )
165.0 $\pm$ 7.1	52.5 $\pm$ 3.5	70.0 $\pm$ 7.1	1.932 $\pm$ 0.26



**Fig. 10** RDFs of H and F atoms of (a)  $\beta$ -PVDF and (b)  $\alpha$ -PVDF (the RDF of  $\beta$ -PVDF in EC is included for comparative purposes).



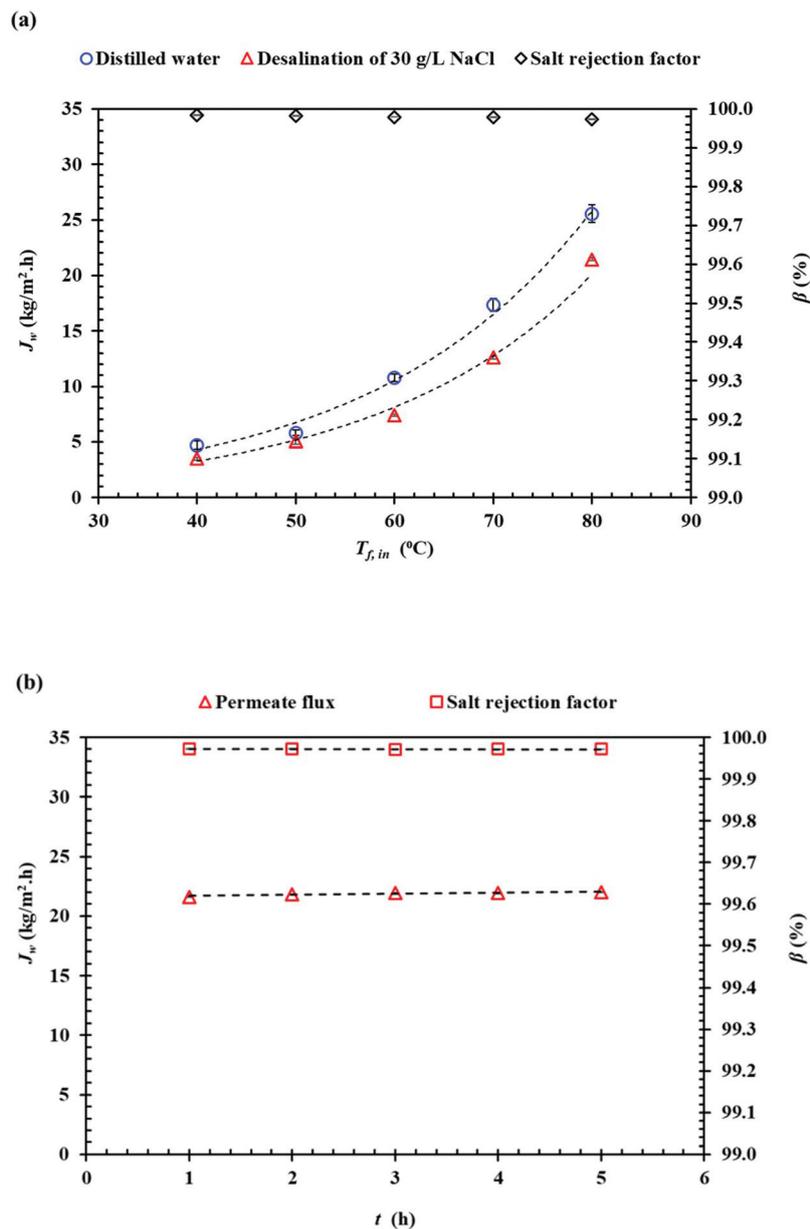


Fig. 11 (a) Permeate flux,  $J_w$  and salt rejection factor ( $\beta$ ) of the PVDF/EC-20 flat-sheet membrane as a function of the inlet feed temperature ( $T_{f,in}$ ) when using as feed distilled water and NaCl (30 g L<sup>-1</sup>) aqueous solution ( $T_{p,in} = 20$  °C) and (b) as a function of time for  $T_{f,in} = 80$  °C,  $T_{p,in} = 20$  °C.

solutions. Exponential (Arrhenius-type) relationships between the DCMD permeate flux and the feed temperature were observed. As a result, the permeate flux increased by 81.56% for distilled water and 83.34% for 30 g L<sup>-1</sup> (NaCl) aqueous solution when the feed temperature increased from 50 to 80 °C while maintaining the permeate temperature at 20 °C. It is well known that in membrane distillation process, the operating feed temperature is the predominant parameter affecting the permeate flux due to the exponential increase of the vapor pressure with temperature. This trend has been observed in various studies.<sup>67–76</sup> In addition, for the saline solution, the permeate flux was decreased by 25.5% and 16.3% at 40 °C and 80 °C, respectively, compared to the corresponding permeate

flux obtained for distilled water. This reduction of the permeate flux upon the adding of NaCl to the aqueous feed solution is attributed to the reduced water vapor pressure of the feed solution and to the concentration polarization effect.<sup>68</sup> As can be seen in Fig. 11(b) both the permeate flux and the salt rejection factor were maintained stable with time. In general, the salt rejection factors were above 99.97% and the electrical conductivity ( $\Omega_{p,f}$ ) of the permeate was below 23.8  $\mu\text{S cm}^{-1}$ . Compared to other PVDF membranes prepared with other solvents for membrane distillation summarized in Table 6, the PVDF/EC-20 membrane prepared with the green cyclic carbonate solvent EC exhibit a competitive DCMD desalination performance.



Table 6 DCMD permeate flux ( $J_w$ ) of the PVDF/EC-20 membrane and other commercial PVDF membranes proposed for membrane distillation reported in the literature

Polymer	Solvent/non-solvent & additive <sup>a</sup>	Membrane distillation configuration <sup>a</sup>	$J_w$ <sup>b</sup> (kg m <sup>-2</sup> h <sup>-1</sup> ); $\alpha$ <sup>b</sup> (%)	Operating conditions <sup>a</sup>	Ref.
PVDF (Kynar 761)	DMF/Ethanol & GO-ODA	AGMD	16.7; ( $\alpha$ = 98.3%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 80 °C; $T_c$ = 15 °C	77
PVDF (534 kDa)	DMF/H <sub>2</sub> O & GO-APTS	AGMD	6.2; ( $\alpha$ > 99.9%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 80 °C; $T_c$ = 20 °C	78
PVDF (Kynar 761)	DMF/H <sub>2</sub> O & c(Graphene + LiCl)	AGMD	20.5; ( $\alpha$ = 99.9%)	RO brine as a feed; $T_f$ = 60 °C; $T_c$ = 20 °C	79
PVDF (534 kDa)	DMF/H <sub>2</sub> O & rGO	AGMD	7.0; ( $\alpha$ > 99.99%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 80 °C; $T_c$ = 20 °C	80
PVDF (Kynar 740)	NMP/(isopropanol + H <sub>2</sub> O) & SiO <sub>2</sub>	VMD	10.0; ( $\alpha$ > 99.99%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 27 °C; $P_v$ = 94.8 kPa	81
PVDF (Kynar 740)	DMAC/H <sub>2</sub> O & SiO <sub>2</sub>	VMD	3.0; ( $\alpha$ > 99.9%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 35 °C; $P_v$ = 1.5 kPa	82
PVDF (Kynar 740)	DMAC/H <sub>2</sub> O	VMD	16.06; ( $\alpha$ = 7.89)	10 g L <sup>-1</sup> of CHCl <sub>3</sub> in H <sub>2</sub> O $T_f$ = 25 °C; $P_v$ = 1.666 kPa	83
PVDF (Kynar 740)	DMAC/H <sub>2</sub> O & CuO	VMD	1.25 ( $\alpha$ > 99.9%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 27.5 °C; $P_v$ = 1.2 kPa	84
PVDF (Kynar 740)	DMAC/MWCNTs	VMD	5.64; ( $\alpha$ > 99.8%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 50 °C; $P_v$ = 2.306 kPa	85
PVDF (275 kDa)	NMP/graphene & H <sub>2</sub> O	DCMD	59.5; ( $\alpha$ > 99.9%)	~30 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 72.5 °C; $T_p$ = 22.5 °C	86
PVDF (HSV900)	DMAC/H <sub>2</sub> O	DCMD	11.0; ( $\alpha$ = 99.99%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 70 °C; $T_p$ = 25 °C	87
PVDF (FR-904)	DMF/(acetone & H <sub>3</sub> PO <sub>4</sub> )	DCMD	12.5; ( $\alpha$ > 99%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 50 °C; $T_p$ = 20 °C	53
PVDF (275 kDa)	TEP/(2-propanol & H <sub>2</sub> O)	DCMD	40.0; (salt passage ~0 mmol m <sup>-2</sup> h <sup>-2</sup> )	58 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 60 °C; $T_p$ = 20 °C	88
PVDF (Kynar 760)	DMAC/ <i>n</i> -propanol & H <sub>2</sub> O	DCMD	9.3; ( $\alpha$ = 99.98%)	58 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 50 °C; $T_p$ = 20 °C	89
PVDF (FR-904)	DMAC/acetone & H <sub>3</sub> PO <sub>4</sub>	DCMD	47.6; ( $\alpha$ > 99.9%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 53 °C; $T_p$ = 20 °C	90
PVDF (Dp, 1000)	DMAC/H <sub>2</sub> O & LiCl	DCMD	9.71; ( $\alpha$ > 98%)	10–20 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 80 °C; $T_p$ = 20 °C	91
PVDF (Solef® 6010)	DMAC/PG & H-SiO <sub>2</sub>	DCMD	27.4; ( $\alpha$ > 99.98%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 70 °C; $T_p$ = 15 °C	92
PVDF (400 kDa)	DMAC/H <sub>2</sub> O & LiCl	VMD	22.4; ( $\alpha$ = 99.9%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 73 °C; $P_v$ = 31 kPa	93
PVDF (Solef® 1015)	EC/H <sub>2</sub> O	DCMD	25.6	Distilled water as feed; $T_f$ = 80 °C; $T_p$ = 20 °C	This work
			21.9; ( $\alpha$ > 99.9%)	35 g L <sup>-1</sup> NaCl as a feed; $T_f$ = 80 °C; $T_p$ = 20 °C	

<sup>a</sup> PVDF, polyvinylidene fluoride; EC, ethylene carbonate; DCMD, direct contact membrane distillation; H<sub>3</sub>PO<sub>4</sub>, phosphoric acid; SiO<sub>2</sub>, superhydrophobic nanoparticles silicon dioxide;  $T_f$ , feed temperature;  $T_p$ , permeate temperature;  $T_c$ , coolant inlet temperature;  $P_v$ , apply vacuum pressure; TEP, triethyl phosphate; LiCl, lithium chloride; NaCl, sodium chloride; RO, reverse osmosis; rGO, reduced graphene oxide; NMP, *N*-methyl-2-pyrrolidone; SiO<sub>2</sub>, hydrophilic nanoparticles; f-SiO<sub>2</sub>, octadecyltrimethoxysilane (OTMS)-modified silica nanoparticles; CuO, hydrophilic spherical nanoparticles;  $D_p$ , degree of polymerization; PG, 1,2-propylene glycol; H-SiO<sub>2</sub>, hydroxyl-rich silica nanoparticle; GO-ODA, octadecylamine functionalized graphene oxide; GO-APTS, 3-(aminopropyl) triethoxysilane functionalized graphene oxide; MWCNTs; multi-walled carbon nanotubes; CHCl<sub>3</sub>, chloroform. <sup>b</sup> Permeate flow data reported from literature with unit conversion if necessary.  $\alpha$ , rejection factor.



## 4. Conclusions

PVDF membranes with differing morphologies and properties were prepared using inexpensive, green, biodegradable cyclic carbonate solvents (EC, PC, and BC). The effect of the polymer concentration in the dope solution was investigated observing the formation of spherulitic structures. The spherulite size decreased as the polymer concentration and alkyl chain length of the solvent was increased (EC < PC < BC). The porosity and pure water permeability of the prepared membranes decreased as the spherulite size was decreased. When using the solvent EC or PC, PVDF crystals were formed solely in the  $\beta$  phase, independently of the PVDF concentration. However, with the solvent BC, these were formed as mixtures of  $\alpha$  and  $\beta$  phase crystals. Molecular dynamics simulation yielded results consistent with the experimental data. The PVDF membrane prepared with the solvent EC and 20 wt% PVDF (PVDF/EC-20) exhibiting the highest pure water permeability and porosity was further tested in DCMD experiments. Stable and competitive DCMD desalination performance (a permeate flux up to 25.6 kg m<sup>-2</sup> h<sup>-1</sup> with a salt rejection factor below 23.8  $\mu$ S cm<sup>-1</sup>) was obtained for this membrane. The achieved results in this study indicate that cyclic organic carbonates are attractive green solvents for PVDF membrane fabrication for different applications (membrane distillation, microfiltration, etc.).

## Nomenclature

$A$	Effective filtration area of membrane (m <sup>2</sup> )
$C_{f,final}$	Final (NaCl) concentration in the feed solution (g L <sup>-1</sup> )
$C_{f,initial}$	Initial (NaCl) concentration in the permeate solution (g L <sup>-1</sup> )
$C_{p,final}$	Final (NaCl) concentration in the permeate solution (g L <sup>-1</sup> )
$C_{p,initial}$	Initial (NaCl) concentration in the permeate solution (g L <sup>-1</sup> )
$d(N_{ij}(r))$	Average number of times the $j$ atoms are comprised in a spherical shell of the thickness $dr$ at a distance $r$ from atoms $i$
$d_{p,max}$	Maximum pore diameter ( $\mu$ m)
$g_{ij}(r)$	Probability density of finding atoms $j$ at a distance $r$ from atom $i$
$J_w$	Water flux (kg m <sup>-2</sup> h <sup>-1</sup> )
$Q$	Volume of water permeated through the membrane (kg)
$m_d$	Weights of the dry membranes (g)
$m_w$	Weights of the wet membranes (g)
$P$	Liquid entry pressure (kPa)
$T_C$	Crystallization temperature (°C)
$T_{f,in}$	Feed inlet temperature (°C)
$T_m^0$	Melting temperature (°C)
$T_{p,in}$	Permeate inlet temperature (°C)
$t$	Time corresponding to a certain volume of collected water (h)

$\beta$	Salt rejection factor (%)
$\delta$	Solubility parameter (MPa <sup>0.5</sup> )
$\delta_D$	Dispersion Force (MPa <sup>0.5</sup> )
$\delta_P$	Polar Force (MPa <sup>0.5</sup> )
$\delta_H$	Hydrogen bond force (MPa <sup>0.5</sup> )
$\rho_p$	Density of polymer (g cm <sup>-3</sup> )
$\rho_k$	Density of kerosene (g cm <sup>-3</sup> )
$\Delta T$	Temperature difference (°C)
$\theta$	Water contact angle (°)
$\Omega_{p,f}$	Final electrical conductivity ( $\mu$ S cm <sup>-1</sup> )
$\gamma$	Coefficient of surface tension of the liquid (N m <sup>-1</sup> )

## Conflicts of interest

There are no conflicts to declare.

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## References

- B. Ameduri, *Chem. Rev.*, 2009, **109**, 6632–6686.
- M. T. Darestani, H. G. L. Coster and T. C. Chilcott, *J. Membr. Sci.*, 2013, **435**, 226–232.
- J. Zhang, P. Cao, Z. Cui, Q. Wang, F. Fan, M. Qiu, X. Wang, Z. Wang and Y. Wang, *AIP Adv.*, 2019, **9**, 115219.
- P. Cao, J. Shi, J. Zhang, X. Wang, J. T. Jung, Z. Wang, Z. Cui and Y. M. Lee, *J. Membr. Sci.*, 2020, **603**, 118037.
- D. Chen and C. Pomalaza-Ráez, *Sep. Purif. Technol.*, 2019, **215**, 612–618.
- M. Li, H. J. Wondergem, M.-J. Spijkman, K. Asadi, I. Katsouras, P. W. M. Blom and D. M. de Leeuw, *Nat. Mater.*, 2013, **12**, 433–438.
- Z. Cui, N. T. Hassankiadeh, Y. Zhuang, E. Drioli and Y. M. Lee, *Prog. Polym. Sci.*, 2015, **51**, 94–126.
- S. Barrau, A. Ferri, A. Da Costa, J. Defebvin, S. Leroy, R. Desfeux and J.-M. Lefebvre, *ACS Appl. Mater. Interfaces*, 2018, **10**, 13092–13099.
- W. Xie, T. Li, A. Tiraferri, E. Drioli, A. Figoli, J. C. Crittenden and B. Liu, *ACS Sustainable Chem. Eng.*, 2021, **9**, 50–75.
- W. H. Organization, *IARC Monographs on the Evaluation of Carcinogenic Risks to Humans*, Lyon, France, 2012.
- Z. Cui, N. T. Hassankiadeh, S. Y. Lee, J. M. Lee, K. T. Woo, A. Sanguineti, V. Arcella, Y. M. Lee and E. Drioli, *J. Membr. Sci.*, 2013, **444**, 223–236.
- M. C. Rhodes, J. R. Bucher, J. C. Peckham, G. E. Kissling, M. R. Hejtmancik and R. S. Chhabra, *Food Chem. Toxicol.*, 2007, **45**, 843–851.



- 13 U. Heudorf, V. Mersch-Sundermann and J. Angerer, *Int. J. Hyg. Environ. Health*, 2007, **210**, 623–634.
- 14 M. Becker, S. Edwards and R. I. Massey, *Environ. Sci. Technol.*, 2010, **44**, 7986–7991.
- 15 M. Razali, J. F. Kim, M. Attfield, P. M. Budd, E. Drioli, Y. M. Lee and G. Szekely, *Green Chem.*, 2015, **17**, 5196–5205.
- 16 S. Rajabzadeh, M. Teramoto, M. H. Al-Marzouqi, E. Kamio, Y. Ohmukai, T. Maruyama and H. Matsuyama, *J. Membr. Sci.*, 2010, **346**, 86–97.
- 17 N. Ghasem, M. Al-Marzouqi and N. Abdul Rahim, *Sep. Purif. Technol.*, 2012, **99**, 91–103.
- 18 Z. Cui, Y. Cheng, K. Xu, J. Yue, Y. Zhou, X. Li, Q. Wang, S.-P. Sun, Y. Wang, X. Wang and Z. Wang, *Polymer*, 2018, **141**, 46–53.
- 19 Z. Cui, N. T. Hassankiadeh, S. Y. Lee, K. T. Woo, J. M. Lee, A. Sanguineti, V. Arcella, Y. M. Lee and E. Drioli, *J. Membr. Sci.*, 2015, **473**, 128–136.
- 20 M. Liu, Z.-l. Xu, D.-g. Chen and Y.-m. Wei, *Desalin. Water Treat.*, 2010, **17**, 183–192.
- 21 N. T. Hassankiadeh, Z. Cui, J. H. Kim, D. W. Shin, S. Y. Lee, A. Sanguineti, V. Arcella, Y. M. Lee and E. Drioli, *J. Membr. Sci.*, 2015, **479**, 204–212.
- 22 W. Xie, A. Tiraferri, B. Liu, P. Tang, F. Wang, S. Chen, A. Figoli and L.-Y. Chu, *ACS Sustainable Chem. Eng.*, 2020, **8**, 91–101.
- 23 T. Marino, F. Galiano, A. Molino and A. Figoli, *J. Membr. Sci.*, 2019, **580**, 224–234.
- 24 F. Russo, F. Galiano, F. Pedace, F. Aricò and A. Figoli, *ACS Sustainable Chem. Eng.*, 2020, **8**, 659–668.
- 25 D. Y. Xing, W. Y. Dong and T.-S. Chung, *Ind. Eng. Chem. Res.*, 2016, **55**, 7505–7513.
- 26 Q. Wu, A. Tiraferri, H. Wu, W. Xie and B. Liu, *ACS Omega*, 2019, **4**, 19799–19807.
- 27 P. Yadav, N. Ismail, M. Essalhi, M. Tysklind, D. Athanassiadis and N. Tavajohi, *J. Membr. Sci.*, 2021, **622**, 118987.
- 28 J. Ranke, S. Stolte, R. Störmann, J. Arning and B. Jastorff, *Chem. Rev.*, 2007, **107**, 2183–2206.
- 29 J. R. Harjani, J. Farrell, M. T. Garcia, R. D. Singer and P. J. Scammells, *Green Chem.*, 2009, **11**, 821–829.
- 30 T. Sakakura and K. Kohno, *Chem. Commun.*, 2009, 1312–1330, DOI: 10.1039/B819997C.
- 31 B. Schöffner, F. Schöffner, S. P. Verevkin and A. Börner, *Chem. Rev.*, 2010, **110**, 4554–4581.
- 32 Y. Su, C. Chen, Y. Li and J. Li, *J. Macromol. Sci., Part A: Pure Appl. Chem.*, 2007, **44**, 99–104.
- 33 H.-C. Yang, Q.-Y. Wu, H.-Q. Liang, L.-S. Wan and Z.-K. Xu, *J. Polym. Sci., Part B: Polym. Phys.*, 2013, **51**, 1438–1447.
- 34 Z. Cui, S. Xu, J. Ding, J. Zhang, B. He, H. Wang and J. Li, *Polymers*, 2018, **10**, 719.
- 35 C. Fang, S. Rajabzadeh, W. Liu, H.-C. Wu, N. Kato, Y. Sun, S. Jeon and H. Matsuyama, *J. Membr. Sci.*, 2020, **596**, 117715.
- 36 M. A. Rasool, P. P. Pescarmona and I. F. J. Vankelecom, *ACS Sustainable Chem. Eng.*, 2019, **7**, 13774–13785.
- 37 C. M. Hansen, *Hansen Solubility Parameters: A User's Handbook, Second Edition*, CRC Press, 2007.
- 38 Y. Chernyak, *J. Chem. Eng. Data*, 2006, **51**, 416–418.
- 39 ASTM F316-03, *Standard Test Methods for Pore Size Characteristics of Membrane Filters by Bubble Point and Mean Flow Pore Test*, ASTM International, West Conshohocken, PA, 2003, DOI: 10.1520/F0316-03.
- 40 H. Sun, *J. Phys. Chem. B*, 1998, **102**, 7338–7364.
- 41 E. Tocci, C. Rizzuto, F. Macedonio and E. Drioli, *Ind. Eng. Chem. Res.*, 2020, **59**, 5267–5275.
- 42 M. Rahmati, M. Jangali and H. Rezaei, *J. Mol. Liq.*, 2019, **296**, 111781.
- 43 N. Ghahramani and M. Rahmati, *Int. J. Heat Mass Transfer*, 2020, **154**, 119487.
- 44 H. Iesavand, M. Rahmati, D. Afzali and S. Modiri, *Mater. Sci. Eng., C*, 2019, **105**, 110010.
- 45 H. J. C. Berendsen, J. P. M. Postma, W. F. v. Gunsteren, A. DiNola and J. R. Haak, *J. Chem. Phys.*, 1984, **81**, 3684–3690.
- 46 S. Nosé, *J. Chem. Phys.*, 1984, **81**, 511–519.
- 47 L. Verlet, *Phys. Rev.*, 1967, **159**, 98–103.
- 48 T. Marino, N. Russo, E. Tocci and M. Toscano, *Theor. Chem. Acc.*, 2001, **107**, 8–14.
- 49 P. Sukitpaneinit and T.-S. Chung, *J. Membr. Sci.*, 2009, **340**, 192–205.
- 50 J. T. Jung, J. F. Kim, H. H. Wang, E. Di Nicolo, E. Drioli and Y. M. Lee, *J. Membr. Sci.*, 2016, **514**, 250–263.
- 51 A. Bottino, G. Capannelli, S. Munari and A. Turturro, *J. Polym. Sci., Part B: Polym. Phys.*, 1988, **26**, 785–794.
- 52 Y. K. Ong, N. Widjojo and T.-S. Chung, *J. Membr. Sci.*, 2011, **378**, 149–162.
- 53 D. Hou, H. Fan, Q. Jiang, J. Wang and X. Zhang, *Sep. Purif. Technol.*, 2014, **135**, 211–222.
- 54 F. Shi, J. Ma, P. Wang and Y. Ma, *J. Taiwan Inst. Chem. Eng.*, 2012, **43**, 980–988.
- 55 L. Li, C.-M. Chan, K. L. Yeung, J.-X. Li, K.-M. Ng and Y. Lei, *Macromolecules*, 2001, **34**, 316–325.
- 56 G.-L. Ji, C.-H. Du, B.-K. Zhu and Y.-Y. Xu, *J. Appl. Polym. Sci.*, 2007, **105**, 1496–1502.
- 57 A. Salimi and A. A. Yousefi, *Polym. Test.*, 2003, **22**, 699–704.
- 58 X. Cai, T. Lei, D. Sun and L. Lin, *RSC Adv.*, 2017, **7**, 15382–15389.
- 59 V. V. Kochervinskii, *Russ. Chem. Rev.*, 1996, **65**, 865–913.
- 60 D. Yang, S. Tornga, B. Orlor and C. Welch, *J. Membr. Sci.*, 2012, **409–410**, 302–317.
- 61 X. He and K. Yao, *Appl. Phys. Lett.*, 2006, **89**, 112909.
- 62 J. R. Gregorio and M. Cestari, *J. Polym. Sci., Part B: Polym. Phys.*, 1994, **32**, 859–870.
- 63 C.-g. Duan, W. N. Mei, W.-G. Yin, J. Liu, J. R. Hardy, S. Ducharme and P. A. Dowben, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2004, **69**, 235106.
- 64 T. Nishiyama, T. Sumihara, E. Sato and H. Horibe, *Polym. J.*, 2017, **49**, 319–325.
- 65 B. K. Saha, A. Saha, D. Sharada and S. A. Rather, *Cryst. Growth Des.*, 2018, **18**, 1–6.
- 66 M. Khayet and T. Matsuura, *Membrane distillation: principles and applications*, Elsevier, 2011.



- 67 J. Mengual, M. Khayet and M. Godino, *Int. J. Heat Mass Transfer*, 2004, **47**, 865–875.
- 68 M. Khayet, *Adv. Colloid Interface Sci.*, 2011, **164**, 56–88.
- 69 F. A. Banat and J. Simandl, *Desalination*, 1994, **95**, 39–52.
- 70 A. Alkhubiri, N. Darwish and N. Hilal, *Desalination*, 2012, **287**, 2–18.
- 71 M. Essalhi and M. Khayet, *J. Membr. Sci.*, 2013, **433**, 180–191.
- 72 M. Khayet, M. P. Godino and J. I. Mengual, *Sep. Sci. Technol.*, 2005, **39**, 125–147.
- 73 M. Khayet and J. I. Mengual, *Desalination*, 2004, **168**, 373–381.
- 74 M. Wang, G. Liu, H. Yu, S.-H. Lee, L. Wang, J. Zheng, T. Wang, Y. Yun and J. K. Lee, *ACS Appl. Mater. Interfaces*, 2018, **10**, 13452–13461.
- 75 F. Guo, A. Servi, A. Liu, K. K. Gleason and G. C. Rutledge, *ACS Appl. Mater. Interfaces*, 2015, **7**, 8225–8232.
- 76 X. Li, W. Qing, Y. Wu, S. Shao, L. E. Peng, Y. Yang, P. Wang, F. Liu and C. Y. Tang, *ACS Appl. Mater. Interfaces*, 2019, **11**, 47963–47971.
- 77 J. Zahirifar, J. Karimi-Sabet, S. M. A. Moosavian, A. Hadi and P. Khadiv-Parsi, *Desalination*, 2018, **428**, 227–239.
- 78 S. Leaper, A. Abdel-Karim, B. Faki, J. M. Luque-Alled, M. Alberto, A. Vijayaraghavan, S. M. Holmes, G. Szekely, M. I. Badawy, N. Shokri and P. Gorgojo, *J. Membr. Sci.*, 2018, **554**, 309–323.
- 79 Y. C. Woo, Y. Kim, W.-G. Shim, L. D. Tijing, M. Yao, L. D. Nghiem, J.-S. Choi, S.-H. Kim and H. K. Shon, *J. Membr. Sci.*, 2016, **513**, 74–84.
- 80 A. Abdel-Karim, J. M. Luque-Alled, S. Leaper, M. Alberto, X. Fan, A. Vijayaraghavan, T. A. Gad-Allah, A. S. El-Kalliny, G. Szekely, S. I. A. Ahmed, S. M. Holmes and P. Gorgojo, *Desalination*, 2019, **452**, 196–207.
- 81 M. H. Emamirad and S. Javadpour, *J. Environ. Health Sci. Eng.*, 2020, **18**, 495–504.
- 82 J. E. Efome, M. Baghbanzadeh, D. Rana, T. Matsuura and C. Q. Lan, *Desalination*, 2015, **373**, 47–57.
- 83 M. Khayet and T. Matsuura, *Ind. Eng. Chem. Res.*, 2001, **40**, 5710–5718.
- 84 M. Baghbanzadeh, D. Rana, T. Matsuura and C. Q. Lan, *Desalination*, 2015, **369**, 75–84.
- 85 R. Zhou, D. Rana, T. Matsuura and C. Q. Lan, *Sep. Purif. Technol.*, 2019, **217**, 154–163.
- 86 C. Athanasekou, A. Sapalidis, I. Katris, E. Savopoulou, K. Beltsios, T. Tsoufis, A. Kaltzoglou, P. Falaras, G. Bounos and M. Antoniou, *Polym. Eng. Sci.*, 2019, **59**, E262–E278.
- 87 F. Abdulla AlMarzooqi, M. Roil Bilad and H. Ali Arafat, *Appl. Sci.*, 2017, **7**, 181.
- 88 S. Nejati, C. Boo, C. O. Osuji and M. Elimelech, *J. Membr. Sci.*, 2015, **492**, 355–363.
- 89 C.-Y. Kuo, H.-N. Lin, H.-A. Tsai, D.-M. Wang and J.-Y. Lai, *Desalination*, 2008, **233**, 40–47.
- 90 D. Hou, G. Dai, J. Wang, H. Fan, Z. Luan and C. Fu, *Desalination*, 2013, **326**, 115–124.
- 91 M. Tomaszewska, *Desalination*, 1996, **104**, 1–11.
- 92 C. Wu, W. Tang, J. Zhang, S. Liu, Z. Wang, X. Wang and X. Lu, *J. Membr. Sci.*, 2017, **543**, 288–300.
- 93 H. Fan, Y. Peng, Z. Li, P. Chen, Q. Jiang and S. Wang, *J. Polym. Res.*, 2013, **20**, 134.

