



Cite this: RSC Adv., 2017, 7, 56426

Received 30th September 2017
Accepted 8th December 2017DOI: 10.1039/c7ra10831a
rsc.li/rsc-advances

Zwitterionic polymers showing upper critical solution temperature behavior as draw solutes for forward osmosis

Changha Ju and Hyo Kang  *

We synthesized the zwitterionic homopolymer polysulfobetaine (PBET), which shows upper critical solution temperature (UCST) characteristics, *via* free radical polymerization in order to investigate its suitability as a draw solute for the forward osmosis (FO) process. The FO and recovery properties of PBET were systematically investigated. The UCST characteristics of the monomer sulfobetaine (BET) were not confirmed, while the UCST of PBET was confirmed to be approximately 40 °C. This result suggests that PBET can be easily recovered from aqueous solutions by cooling them to below the UCST. In an active layer facing feed solution (AL-FS) system containing 20 wt% PBET at a temperature of 50 °C, the water flux and reverse solute flux of PBET were found to be approximately 3.22 LMH and 0.36 gMH, respectively. Thus, we investigated the suitability of homopolymers having UCST characteristics as draw solutes for the FO process for the first time.

1 Introduction

Membrane separation has been widely used as a water treatment technology in order to provide fresh water to humankind. One of the membrane separation processes, forward osmosis (FO), shows superior energy efficiency to other methods, and FO has therefore received some attention as a next-generation water treatment technology.^{1,2} However, improving the energy efficiency of the recovery of draw solutes from water is important for the commercialization of FO technology.³ An economic method of recovering draw solutes has been actively investigated as interest in energy efficiency has increased, with attention being paid to the recovery characteristics of draw solutes.⁴

Draw solutes are largely divided into inorganic-based, organic-based, and inorganic/organic hybrid draw solutes. Materials that have been identified as inorganic-based draw solutes include conventional inorganic salts.⁵ For example, Elimelech *et al.* have investigated ammonium bicarbonate (NH_4HCO_3) that was derived from two substances, ammonia (NH_3) and carbon dioxide (CO_2), as a draw solute. However, the decomposition of NH_4HCO_3 into NH_3 and CO_2 in the separation process requires a high temperature (>60 °C), and it is known that it is difficult to completely remove decomposed NH_3 from solutions.⁶ As an alternative to the approaches used previously, one study has reported the direct use of an inorganic draw

solution that had undergone FO as a fertilizer, without a prior recovery process.⁷

Organic-based draw solutes include organic molecules, polyelectrolytes, hydrogels, and responsive polymers.^{8–13} The molecular sizes of such organic molecules are large compared to inorganic molecules, and their reverse solute flux is low compared to inorganic draw solutes.¹⁰ Studies have reported that pure water can be obtained through distillation after FO filtration using volatile organic molecules such as dimethyl ether as draw solutes.^{5,14,15} Polyelectrolytes^{16,17} have the advantage of negligibly small reverse solute fluxes in the FO process when compared to the organic molecules previously described.¹⁸ As another example, hydrogels typically form composites with FO membranes that capture water in a cross-linked hydrogel network during the FO process. The hydrogel can then be dehydrated by various external stimuli to remove this water, resulting in the volume of the hydrogel contracting and pure water being obtained.^{19–21} However, hydrogels have a disadvantage in that the difference between the osmotic pressure of the draw solution and the feed solution is reduced due to the Gibbs–Donnan effect. Responsive polymers can be separated from water by controlling their solubility in response to external stimuli such as CO_2 and temperature.^{22–27} For example, amine-type polymers are quaternized in response to external CO_2 . The polarity of these quaternized polymers, which have ionic charges, induces the movement of water. On the other hand, the dequaternized polymer loses its ionic charge and become an insoluble salt, allowing it to be separated from water.²² Thermo-responsive polymers are draw solutes that utilize lower critical solution temperature (LCST) and upper critical solution temperature (UCST) characteristics, and they

Department of Chemical Engineering, Dong-A University, 37 Nakdong-Daero 50beon-gil, Saha-gu, Busan 604-714, Korea. E-mail: hkang@dau.ac.kr; Fax: +82 200 7728; Tel: +82 51 200 7720



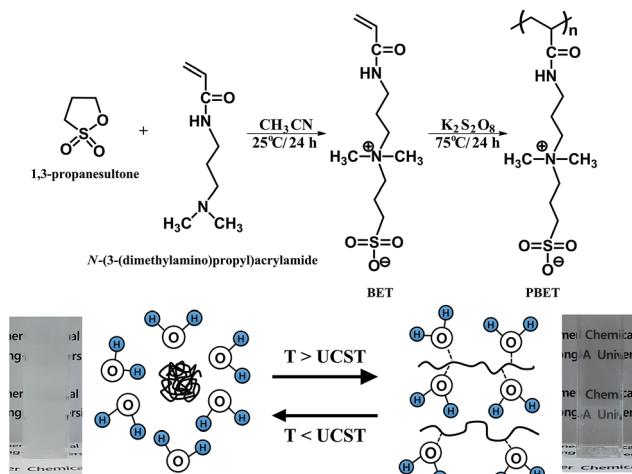


Fig. 1 Synthetic route to zwitterionic homopolymer polysulfobetaine (PBET) and schematic diagram of upper critical solution temperature (UCST) behavior.

can be simply separated from water by controlling their solubility through temperature changes.^{23–28} A previous study reported that a thermo-responsive copolymer contains monomers that are responsible for the osmotic pressure and the recovery properties, respectively, and that the copolymer can have a trade-off relationship between the osmotic pressure and the recovery depending on the molar ratio of the monomers that induce each property.²⁴ Although research on thermo-responsive draw solutes with LCST characteristics is active, few studies have reported the development of draw solutes using UCST characteristics.

Inorganic/organic hybrid draw solutes include hydroacid complexes and functionalized MNPs (magnetic nanoparticles). Hydroacid complexes are coordination complexes of metals with organic molecules that induce high osmotic pressure and have low reverse solute fluxes.^{29–31} Functionalized MNPs are prepared by the ligand exchange of organic molecules that induce osmotic pressure on nano-sized metals with magnetic properties, allowing them to be separated from water using a magnetic field after FO.^{32,33} Some studies have investigated multi-functional MNPs by attaching organic molecules responsive to temperature, light, and CO₂ to MNPs.^{34–36}

In this study, we synthesized a zwitterionic homopolymer of sulfobetaine that is known to have UCST properties that are biocompatible³⁷ and responsive to temperature changes³⁸ as a draw solute candidate for the FO process (Fig. 1). The feasibility of a homopolymer with UCST characteristics as a next-generation draw solute for FO was investigated systematically, including a study of the FO characteristics and recovery method of this draw solute.

2 Experimental

2.1. Reagents and instrumentations

1,3-Propanesultone (PS) and *N*-(3-(dimethylamine)propyl)acrylamide were purchased from Tokyo Chemical Industry Co., Ltd.,

and acetonitrile was purchased from Sigma Aldrich Co., LLC. Potassium persulfate was purchased from Fluka. All reagents were used without further purification. Fourier transform-infrared (FT-IR) spectrometry (Thermo Fisher Scientific, NICOLET 380) and Fourier transform nuclear magnetic resonance (NMR) spectrometry (Agilent, MR400 DD2) were used to identify the synthesized structure. Infrared spectrometry was performed on the attenuated total reflection (ATR) mode between 4000 and 670 cm^{−1}. The number average molecular weight (M_n) and polydispersity index were investigated by gel permeation chromatography (GPC). Relative molecular weight measurement was performed using a Shodex GPC-101 with columns SB-806 M HQ in series with a refractive index (RI) detector. Poly(ethylene oxide) and poly(ethylene glycol) were used as standard materials for calibration. The thermal properties of the samples were analysed by measuring weight changes according to temperature using a thermogravimetric analyser (SETARAM Instrumentation, SETSYS Evolution). The rheological properties of the samples were investigated by measuring the shear rate changes using a cone and plate rheometer with a diameter of 40 mm and an angle of 2° (TA Instrument, AR G2, stress control type). The osmotic pressure was measured using an osmometer (KNAUER, SEMI-MICRO OSMOMETER K-7400) by the freezing point depression method. The UCST was measured by observing the turbidity of the draw solution according to the temperature change using a thermocouple (WOOJIN Instrument, WJ-3500). In addition, UCST was undoubtedly determined by observing the transmittance at a wavelength of 550 nm using UV-Vis spectrophotometer (V-1100 DIGITAL SPECTROPHOTOMETER, EMCLAB instruments) coupled with temperature controller (TC-200P, Misung Scientific. Co. Ltd.) by increasing the temperature from 25 to 50 °C. The water flux was determined by measuring the height difference between the solutions on both sides of a home-made U-shaped tube which face two L-shaped glass tubes using clamp. The reverse solute flux was measured by using a conductivity meter (METTLER TOLEDO, Seven2Go pro) to determine the conductivity difference before and after the experiment.

2.2. Synthesis

We synthesized the zwitterionic homopolymer polysulfobetaine (PBET) using free radical polymerization in order to confirm its usability as a draw solute for FO. The synthetic route used was as follows: 30.61 g (0.251 mol) of PS and 39.15 g of *N*-(3-(dimethylamino)propyl)acrylamide (0.251 mol, 100 mol% PS) were dissolved in acetonitrile (41.15 g, 1.003 mol, 400 mol% PS). The temperature was maintained at 25 °C, and the mixed solution was reacted for 24 h under a nitrogen atmosphere. The resulting solution was precipitated with ethanol and separated via filtration. The solvent was then dried to obtain monomer *N,N'*-dimethyl(acrylamidopropyl)ammonium propane sulfonate (BET). Nuclear magnetic resonance (NMR) spectrometry is used to determine and confirm BET composition (Fig. 2(a)).

¹H-NMR of BET [400 MHz, D₂O, δ/ppm]: 1.97–2.04 (m, 2H, (−CH₂−CH₂−CH₂−)), 2.12–2.20 (m, 2H, (−CH₂−CH₂−CH₂−)), 2.91–2.95 (t, 2H (−CH₂−CH₂−SO₃[−])), 3.06 (s, 6H, (CH₃−N⁺−CH₃)), 3.30–



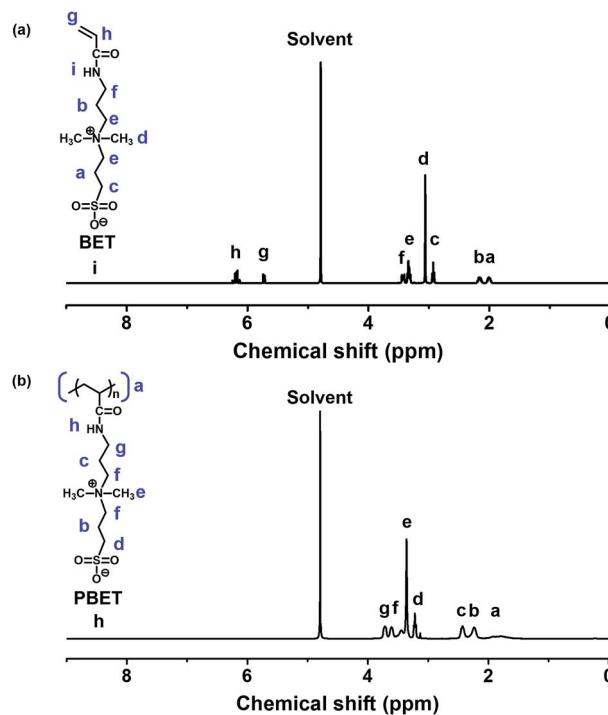


Fig. 2 Nuclear magnetic resonance (NMR) spectra: (a) monomer *N,N'*-dimethyl(acrylamidopropyl)ammonium propane sulfonate (BET), (b) polysulfobetaine (PBET).

3.36 (t, 4H, ($-\text{CH}_2\text{N}^+-\text{CH}_2-$)), 3.41–3.45 (t, 2H, ($-\text{NH}-\text{CH}_2-\text{CH}_2-$)), 5.71–5.74 (d, 2H, ($\text{H}_2\text{C}=\text{CH}-\text{CO}-$)), 6.13–6.25 (t, 1H, ($\text{H}_2\text{C}=\text{CH}-\text{CO}-$)).

BET (20 g) and potassium persulfate (1 g, 5 wt% of BET) were mixed into 80 mL of distilled water and held at a temperature of 75 °C for 24 h under a nitrogen atmosphere. The solution was then added to a dialysis tube (Membrane Filtration Products, Inc., Cellu Sep T1 (molecular weight cut off: 3500 Da)) for 24 h to remove the impurities by dialysis. The solution was then heated to 110 °C to remove the remaining water, allowing pure PBET ($M_n = 18\,000$ and polydispersity index = 2.30) to be obtained. Nuclear magnetic resonance (NMR) spectrometry is used to determine and confirm PBET composition (Fig. 2(b)).

$^1\text{H-NMR}$ of PBET [400 MHz, D_2O , δ/ppm]: 1.64–1.97 (m, 3H, ($-\text{CH}_2-\text{CH}-$)), 2.18–2.33 (m, 2H, ($-\text{CH}_2-\text{CH}_2-\text{CH}_2-$)), 2.38–2.47 (t, 2H, ($-\text{CH}_2-\text{CH}_2-\text{CH}_2-$)), 3.20–3.24 (t, 2H, ($-\text{CH}_2-\text{CH}_2-\text{SO}_3^-$)), 3.36 (s, 6H, ($\text{CH}_3\text{N}^+-\text{CH}_3$))), 3.42–3.46 (t, 2H, ($-\text{CH}_2-\text{CH}_2-\text{N}^+-$)), 3.57–3.65 (t, 2H, ($-\text{N}^+-\text{CH}_2-\text{CH}_2-$)), 3.68–3.75 (t, 2H, ($-\text{NH}-\text{CH}_2-\text{CH}_2-$)).

2.3. FO performance

Water flux, which is an important property for the FO process, was investigated. The water flux was measured using a small-scale FO system connected to a home-made, cap-fixed L-shaped glass tubes.^{24,27} A membrane (Hydration Technologies Inc. (HTI), thin film composite FO membrane) was placed in a channel (diameter: 2.3 cm) between two glass tubes. The PBET draw solution was added into one tube, and distilled water was added into the other. The glass tubes were placed into a water

bath held at a constant temperature. Measurement of water flux was performed in the active layer facing feed solution method (AL-FS). The temperature was maintained at 50 ± 1 °C during the FO process using a temperature control device (BROOKFIELD, TC-500), and the solution was stirred using a magnetic bar and a solenoid (AS ONE, OCTOPUS CS-4). The water flux was determined by measuring the height difference of solution before and after the FO process. The water flux (J_w , ($\text{L m}^{-2} \text{h}^{-1}$, LMH)) was calculated from the volume change of the draw solution using eqn (1).

$$J_w = \frac{\Delta V}{A\Delta t} \quad (1)$$

where ΔV is the volume change of the draw solution over time Δt and A is the effective membrane area ($4.15 \times 10^{-4} \text{ m}^2$).

The reverse solute flux (J_s) was determined by analysing the quantity of solute that diffused from the draw solution into the feed solution during the FO process, and the total dissolved solid (TDS) of the feed solution was also measured. The reverse solute flux (J_s , $\text{g m}^{-2} \text{h}^{-1}$, gMH) was calculated from the difference between the conductivities of the feed solution before and after the FO process using eqn (2).

$$J_s = \frac{\Delta(CV)}{A\Delta t} \quad (2)$$

where ΔC (mol L^{-1}) is the concentration change of the feed solution after time Δt and ΔV (L) is the volume change after time Δt .

3 Results and discussion

3.1. Synthesis and characterization of zwitterionic homopolymer polysulfobetaine (PBET)

The synthetic route to zwitterionic homopolymer polysulfobetaine (PBET) is shown in Fig. 1, where it can be seen that PBET is a zwitterionic homopolymer of sulfobetaine. The chemical structures of the synthesized monomer and homopolymer were identified by $^1\text{H-NMR}$ spectrometry, respectively. We synthesized the zwitterionic monomer *N,N'*-dimethyl(acrylamidopropyl)ammonium propane sulfonate (BET). The $^1\text{H-NMR}$ spectrum and assignments of the respective peaks of monomer BET was indicated in Fig. 2(a). $^1\text{H-NMR}$ spectrum of BET indicates the presence of protons from the vinyl group of BET ($\delta = 5.71\text{--}5.74$ ppm (peak h), 6.13–6.25 (peak h)). The proton peaks from the quaternized ammonia cation ($\delta = 3.06$ ppm (peak d)) show the inclusion of the cation moiety in the zwitterionic BET. The anion moiety in the zwitterionic BET is not observed that is not containing hydrogen because of structure of sulfur trioxide. Based on the $^1\text{H-NMR}$ analysis, we identified that BET was successfully synthesized, and then synthesized polysulfobetaine (PBET) via free radical polymerization. The disappearance of the vinyl double bond peaks at 5.71–5.74 ($\text{H}_2\text{C}=\text{CH}-\text{CO}-$) and 6.13–6.25 ($\text{H}_2\text{C}=\text{CH}-\text{CO}-$) ppm and inclusion of BET due to the presences of the proton peaks of amide and quaternized ammonium after the free radical polymerization demonstrate the successful synthesis of PBET as shown in $^1\text{H-NMR}$ spectrum of PBET (Fig. 2(b)). We synthesized



the zwitterionic homopolymer PBET *via* free radical polymerization using BET as monomer. The synthesized materials were analysed using FT-IR spectroscopy to confirm that PBET had been synthesized. As shown in Fig. 3, the PBET infrared spectrum confirmed the presence of the amide group ($-\text{CO}-\text{NH}-$) through the $\text{C}=\text{O}$ stretching peak at 1625 cm^{-1} and the $\text{N}-\text{H}$ stretching peak at 1521 cm^{-1} . The peak at 1037 cm^{-1} relates to the stretching of $\text{O}=\text{S}=\text{O}$, and the peak at 944 cm^{-1} indicates the stretching of a quaternary ammonium cation. The characteristic IR peaks of the functional groups were identical to those seen in the FT-IR spectra of other polysulfobetaines, confirming that polysulfobetaine was synthesized.^{39,40}

3.2. Thermogravimetry

To analyse the thermal properties of PBET, its pyrolysis behavior was measured up to $800\text{ }^\circ\text{C}$ at a heating rate of $10\text{ }^\circ\text{C min}^{-1}$ under an argon atmosphere. The thermogravimetric curve is shown in Fig. 4. The first weight loss, which began at approximately $100\text{ }^\circ\text{C}$, was due to the loss of the water absorbed in the PBET.⁴⁰ According to the thermogram, the total water content of the polymer was approximately 5 wt%, and thermal decomposition of PBET did not occur in this temperature range. The second transition was confirmed to be caused by the thermal decomposition of the polymer between $254\text{ }^\circ\text{C}$ and $483\text{ }^\circ\text{C}$, with the weight decreasing by approximately 62% in this

range. At this range, the polymer backbone and branched groups such as quaternized ammonium, and sulfonate groups were fragmented, whereupon hydrocarbons were formed. Finally, generated hydrocarbons remained about 15 wt% char at $800\text{ }^\circ\text{C}$.⁴¹ This pyrolysis behavior is similar to that of polysulfobetaine, as reported previously by other research groups.^{41,42} Therefore, this thermal analysis indicates that the homopolymer PBET having thermal stability at operating temperature higher than UCST is available for draw solute in the forward osmosis process, of which UCST characteristics will be mentioned later.

3.3. Viscosity

Viscosity is an important factor that affects the FO behavior of a draw solution, and has a substantial effect on the efficiency of the FO process. Typically, a solution with a high viscosity can induce ICP (internal concentration polarization) and ECP (external concentration polarization) on the membrane, which can reduce the water flux of the draw solution.^{18,43} The viscosity of PBET was measured as $50\text{ }^\circ\text{C}$, which is higher than the phase separation temperature of PBET, as will be discussed later. For comparison purposes, the viscosity of the monomer BET was measured at room temperature and at $50\text{ }^\circ\text{C}$ by varying the shear rate from 0.1 to $100\text{ [s}^{-1}\text{]}$. The results are shown in Fig. 5. Fig. 5(a) and (b) show that the viscosity of BET was observed in the range between 1 cP and 1.47 cP at room temperature and $50\text{ }^\circ\text{C}$, with no significant changes in viscosity according to concentration being indicated. As shown in Fig. 5(c), the viscosity of PBET was approximately 1.62, 3.65, 7.36, and 17.69 cP according to concentration of 5, 10, 15, and 20 wt% at $50\text{ }^\circ\text{C}$, respectively. These results are similar to those of other draw solutes, where increases in viscosity with increasing concentration were also observed.^{16,44} This confirms that the polymer PBET had a relatively higher viscosity than its monomer, BET.

3.4. Osmotic pressure

The osmotic pressure of a draw solution is closely related to its concentration, suggesting that osmotic pressure is an important factor when determining FO performance. We used the freezing point depression method to measure the osmotic pressure in order to investigate the usability of PBET as a draw solute. The osmotic pressure of the monomer BET was measured to be approximately 194, 379, 630, and $898\text{ mOsmol kg}^{-1}$ at concentrations of 5, 10, 15, and 20 wt%, respectively (Fig. 6). These results show that the osmotic pressure increased with increasing concentration, as has been observed in other electrolyte systems that can induce osmotic pressure.^{16,45} However, since PBET is insoluble at room temperature, it was impossible to measure its osmotic pressure using the freezing point depression method.

3.5. Recovery properties

The separation of draw solutes from water in the FO process enables the sustainable use of draw solutes. In order to overcome the trade-off between the osmotic pressure and recovery temperature that occurs in thermo-responsive copolymer-based

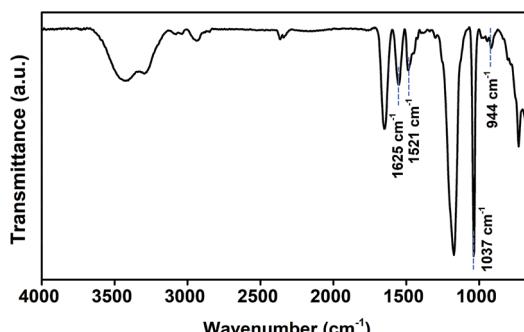


Fig. 3 Fourier transform-infrared (FT-IR) spectrum of PBET.

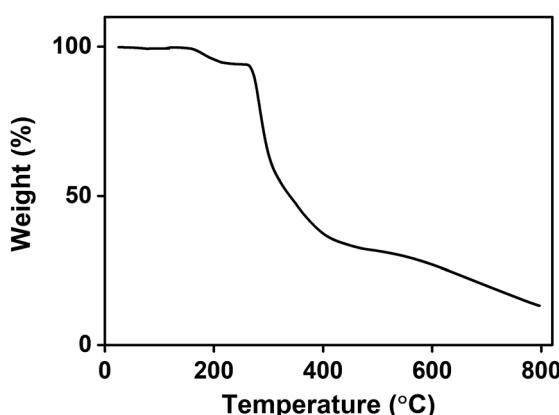


Fig. 4 Thermogravimetric curve of PBET.



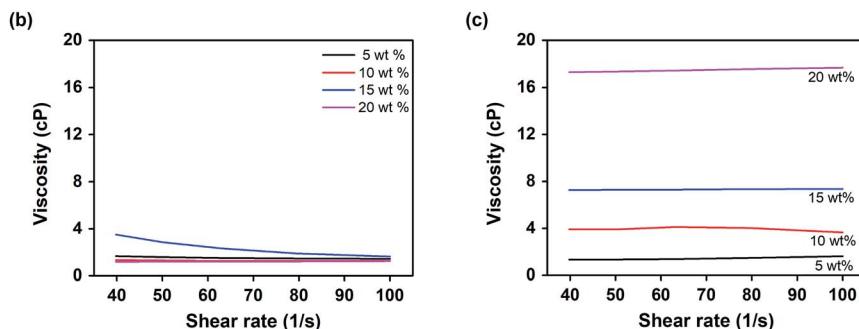


Fig. 5 Viscosity according to the shear rate: (a) BET at room temperature, (b) BET at 50 °C, (c) PBET at 50 °C.

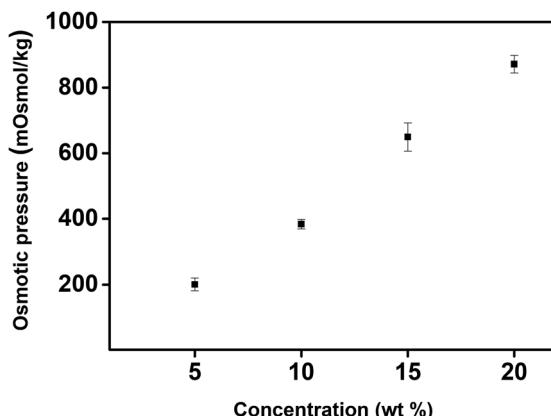


Fig. 6 Osmotic pressure according to the concentration of *N,N'*-dimethyl(acrylamidopropyl)ammonium propane sulfonate (BET) measured by freezing point depression method.

draw solutes, a homopolymer-type draw solute was designed and used in this study. The UCST is defined as the critical temperature at which the draw solute and water in a draw solution change from a homogeneous phase to a heterogeneous phase. To confirm that the draw solute and water were separated into heterogeneous phases, phase separation temperature should be determined transmittance curve of BET and PBET in aqueous solutions according to the temperature change using UV-Vis spectrophotometer. The monomer BET did not show any changes in transmittance curve according to the temperature change from 0 °C to 100 °C (figure not shown), which suggests that BET and water cannot be separated from each other by changing the temperature. Fig. 7 shows transmittance curve of PBET aqueous solution according to the temperature change. The UCST was closely examined by observing the transmittances using UV-Vis spectrophotometer at 550 nm. The transmittance curve exhibits drastic change at approximately 40 °C, which can signify the critical temperature. When the temperature of the PBET solution was higher than the critical temperature, the transmittance of the PBET aqueous solution was approximately 100%. This result indicates that PBET and water were in homogeneous phase. However, when the solution temperature was lower than the critical temperature, the transmittance of the PBET aqueous solution was approximately 0%, which PBET and water were separated into each other becoming

heterogeneous phase. As shown in Fig. 1, the solubility of PBET in an aqueous solution depends on the temperature. When the solution temperature is lower than the critical temperature, the ion-ion interactions between the sulfobetaine moieties are stronger than the sulfobetaine-water interactions causing the polymer chains to aggregate and resulting in PBET forming a heterogeneous phase with water. On the other hand, when the solution temperature is higher than the critical temperature, the sulfobetaine-water interaction becomes stronger than the ion-ion interactions between the sulfobetaine moieties, and PBET and water form a homogeneous phase. The critical temperature for the phase separation of PBET and water was found to be approximately 40 °C. We believe that the UCST of PBET was about 40 °C as a result of the transmittance according to the temperature. Therefore, PBET can be simply separated from water by controlling the solubility through temperature changes, and the energy required for this process can be obtained using geothermal heat or the waste heat of power plants.

3.6. Water flux and reverse solute flux

The water flux and reverse solute flux were measured, giving an indication of the FO performance of recoverable PBET (Fig. 8). Measurements of water flux and reverse solute flux were performed in the AL-FS method using home-made glass tubes. The

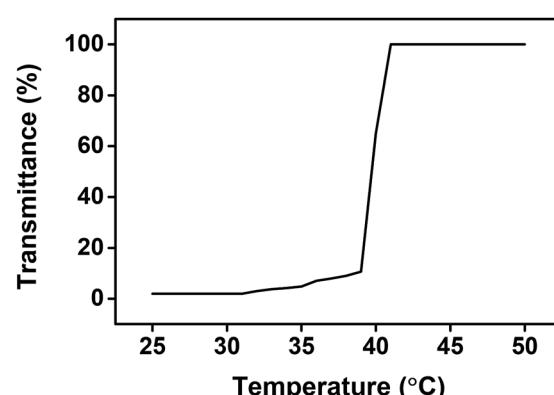


Fig. 7 Transmittance curve of PBET aqueous solution according to the temperature change using UV-Vis spectrophotometer.



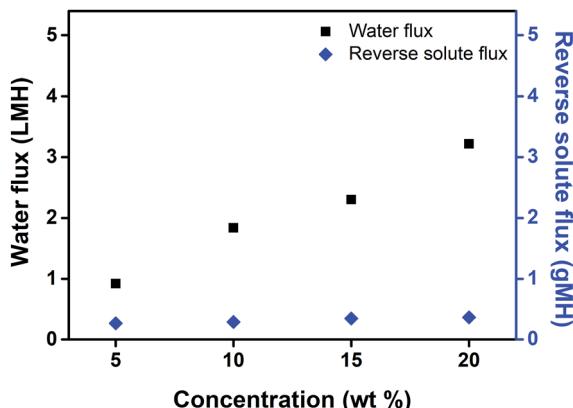


Fig. 8 Water flux and reverse solute flux of PBET according to the concentration at 50 °C during forward osmosis process.

measurements were performed at a temperature of 50 °C, which was above the UCST. Typically, the concentration of the draw solution has an important effect on the FO performance because an increase in the concentration of the draw solution results in an increase in the osmotic pressure, and an increase in osmotic pressure results in an increase in the water flux.⁶ As expected, the water flux increased to 0.92, 1.39, 2.30, and 3.22 LMH as the concentration of the draw solutes was increased to 5, 10, 15, and 20 wt%, respectively. The measurement of the reverse solute flux was performed simultaneously under the same conditions at which the water flux measurements were conducted, and the amount of reversely diffused draw solutes was determined by comparing the conductivity of the feed solution before and after the FO process. Previous studies have reported that the average pore size of the membrane is small compared to the size of draw solute in an aqueous solution, leading to imperfect prevention of back diffusion to the feed solution.⁴⁶ The reverse solute flux is generated due to the difference between the concentrations of the draw solution and the feed solution, and the concentration gradient generated on the active layer of the membrane increases according to the difference between the concentrations of the solutions. Thus, the reverse solute flux increases as the concentration of the draw solution increases.⁴⁷ The reverse solute flux was 0.27, 0.29, 0.35, and 0.36 gMH at draw solute concentrations of 5, 10, 15, and 20 wt%, respectively.

4 Conclusions

We synthesized polysulfobetaine (PBET), a type of zwitterionic homopolymer, by using free radical polymerization in order to confirm its usability as draw solute in forward osmosis. In aqueous solutions, PBET shows UCST characteristics, which plays an important role in imparting recoverability to PBET. The UCST of PBET was found to be approximately 40 °C. The water flux of PBET was identified to be approximately 3.22 LMH at 20 wt% and at 50 °C in active layer facing feed solution (AL-FS) system, and the reverse solute flux was identified to be approximately 0.36 gMH under the same conditions. Therefore,

a thermo-responsive homopolymer containing zwitterionic moieties that shows an UCST has been shown to be a new draw solute that may surpass the limitations of existing technologies, such as the trade-off between osmotic pressure and recovery temperature in copolymer systems. As a result, this research provides a deep understanding of new ways of developing draw solutes, and could provide inspiration for the design and synthesis of thermo-responsive organic materials.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This research was supported by the Basic Science Research Program through the National Research Foundation of Korea (NRF) funded by the Ministry of Education, Science and Technology (NRF-2015R1C1A1A01051654). Financial supports by the Dong-A University Research Fund are gratefully acknowledged.

Notes and references

- 1 P. H. Gleick, in *Water in crisis, Pacific Institute for Studies in Dev., Environment & Security. Stockholm Env. Institute*, Oxford Univ. Press, 1993, p. 473.
- 2 L. Chekli, S. Phuntsho, H. K. Shon, S. Vigneswaran, J. Kandasamy and A. Chanan, *Desalination Water Treat.*, 2012, **43**, 167–184.
- 3 S. Zou, H. Yuan, A. Childress and Z. He, *Environ. Sci. Technol.*, 2016, **50**, 6827–6829.
- 4 D. L. Shaffer, J. R. Werber, H. Jaramillo, S. Lin and M. Elimelech, *Desalination*, 2015, **356**, 271–284.
- 5 J. R. McCutcheon, R. L. McGinnis and M. Elimelech, *Desalination*, 2005, **174**, 1–11.
- 6 A. Achilli, T. Y. Cath and A. E. Childress, *J. Membr. Sci.*, 2010, **364**, 233–241.
- 7 S. Phuntsho, H. K. Shon, S. Hong, S. Lee and S. Vigneswaran, *J. Membr. Sci.*, 2011, **375**, 172–181.
- 8 E. M. Garcia-Castello, J. R. McCutcheon and M. Elimelech, *J. Membr. Sci.*, 2009, **338**, 61–66.
- 9 R. E. Kravath and J. A. Davis, *Desalination*, 1975, **16**, 151–155.
- 10 K. Lutchmiah, L. Lauber, K. Roest, D. J. Harmsen, J. W. Post, L. C. Rietveld, J. B. van Lier and E. R. Cornelissen, *J. Membr. Sci.*, 2014, **460**, 82–90.
- 11 H. Y. Ng, W. Tang and W. S. Wong, *Environ. Sci. Technol.*, 2006, **40**, 2408–2413.
- 12 H. T. Nguyen, S.-S. Chen, N. C. Nguyen, H. H. Ngo, W. Guo and C.-W. Li, *J. Membr. Sci.*, 2015, **489**, 212–219.
- 13 Y. Cai, W. Shen, J. Wei, T. H. Chong, R. Wang, W. B. Krantz, A. G. Fane and X. Hu, *Environ. Sci.: Water Res. Technol.*, 2015, **1**, 341–347.
- 14 G. Batchelder, *US Pat. 3 171 799*, 1962.
- 15 N. Sato, Y. Sato and S. Yanase, *Desalination*, 2014, **349**, 102–105.
- 16 Q. Ge, J. Su, G. L. Amy and T.-S. Chung, *Water Res.*, 2012, **46**, 1318–1326.



17 E. Tian, C. Hu, Y. Qin, Y. Ren, X. Wang, X. Wang, P. Xiao and X. Yang, *Desalination*, 2015, **360**, 130–137.

18 Q. Ge, P. Wang, C. Wan and T.-S. Chung, *Environ. Sci. Technol.*, 2012, **46**, 6236–6243.

19 D. Li, X. Zhang, J. Yao, G. P. Simon and H. Wang, *Chem. Commun.*, 2011, **47**, 1710–1712.

20 Y. Cai, W. Shen, S. L. Loo, W. B. Krantz, R. Wang, A. G. Fane and X. Hu, *Water Res.*, 2013, **47**, 3773–3781.

21 J. Wei, Z.-X. Low, R. Ou, G. P. Simon and H. Wang, *Water Res.*, 2016, **96**, 292–298.

22 Y. Cai, W. Shen, R. Wang, W. B. Krantz, A. G. Fane and X. Hu, *Chem. Commun.*, 2013, **49**, 8377–8379.

23 M. Noh, Y. Mok, S. Lee, H. Kim, S. H. Lee, G.-w. Jin, J.-H. Seo, H. Koo, T. H. Park and Y. Lee, *Chem. Commun.*, 2012, **48**, 3845–3847.

24 J.-j. Kim, J.-S. Chung, H. Kang, Y. A. Yu, W. J. Choi, H. J. Kim and J.-C. Lee, *Macromol. Res.*, 2014, **22**, 963–970.

25 Y. Kohno, S. Saita, Y. Men, J. Yuan and H. Ohno, *Polym. Chem.*, 2015, **6**, 2163–2178.

26 Y. Zhong, X. Feng, W. Chen, X. Wang, K.-W. Huang, Y. Gnanou and Z. Lai, *Environ. Sci. Technol.*, 2015, **50**, 1039–1045.

27 J.-j. Kim, H. Kang, Y.-S. Choi, Y. A. Yu and J.-C. Lee, *Desalination*, 2016, **381**, 84–94.

28 W. Li, J. Li, J. Gao, B. Li, Y. Xia, Y. Meng, Y. Yu, H. Chen, J. Dai, H. Wang and Y. Guo, *Biomaterials*, 2011, **32**, 3832–3844.

29 Q. Ge and T.-S. Chung, *Chem. Commun.*, 2013, **49**, 8471–8473.

30 Q. Ge, F. Fu and T.-S. Chung, *Water Res.*, 2014, **58**, 230–238.

31 N. T. Hau, S.-S. Chen, N. C. Nguyen, K. Z. Huang, H. H. Ngo and W. Guo, *J. Membr. Sci.*, 2014, **455**, 305–311.

32 M. M. Ling and T.-S. Chung, *Desalination*, 2011, **278**, 194–202.

33 S. Y. Park, H.-W. Ahn, J. W. Chung and S.-Y. Kwak, *Desalination*, 2016, **397**, 22–29.

34 Q. Zhao, N. Chen, D. Zhao and X. Lu, *ACS Appl. Mater. Interfaces*, 2013, **5**, 11453–11461.

35 S. Chen, C. X. Guo, Q. Zhao and X. Lu, *Chem.-Eur. J.*, 2014, **20**, 14057–14062.

36 H. Han, J. Y. Lee and X. Lu, *Chem. Commun.*, 2013, **49**, 6122–6124.

37 S. Jiang and Z. Cao, *Adv. Mater.*, 2010, **22**, 920–932.

38 O. Azzaroni, A. A. Brown and W. T. Huck, *Angew. Chem.*, 2006, **118**, 1802–1806.

39 Y.-F. Mi, Q. Zhao, Y.-L. Ji, Q.-F. An and C.-J. Gao, *J. Membr. Sci.*, 2015, **490**, 311–320.

40 E. O. Ningrum, A. Purwanto, H. Ni'mah, S. Sumarno, D. I. Dewitasari and E. O. Mulyadi, *AIP Conf. Proc.*, 2017, **1840**, 090007.

41 R. Lalani and L. Liu, *Polymer*, 2011, **52**, 5344–5354.

42 Z. Dong, J. Mao, M. Yang, D. Wang, S. Bo and X. Ji, *Langmuir*, 2011, **27**, 15282–15291.

43 P. H. Duong, S. Chisca, P.-Y. Hong, H. Cheng, S. P. Nunes and T.-S. Chung, *ACS Appl. Mater. Interfaces*, 2015, **7**, 3960–3973.

44 M. L. Stone, A. D. Wilson, M. K. Harrup and F. F. Stewart, *Desalination*, 2013, **312**, 130–136.

45 J. R. McCutcheon, R. L. McGinnis and M. Elimelech, *J. Membr. Sci.*, 2006, **278**, 114–123.

46 W. A. Phillip, J. S. Yong and M. Elimelech, *Environ. Sci. Technol.*, 2010, **44**, 5170–5176.

47 C. Y. Tang, Q. She, W. C. Lay, R. Wang and A. G. Fane, *J. Membr. Sci.*, 2010, **354**, 123–133.

