

Preparation and characterization of polysulfone membrane coated with poly(ether block amide) for oxygen enrichment process

Kok Chung Chong ^{a,*}, Yin Yin Chan ^a, Woei Jye Lau ^b, Soon Onn Lai ^a, Ahmad Fauzi Ismail ^b, Hui Shan Thiam ^a

^a Universiti Tunku Abdul Rahman, Lee Kong Chian Faculty of Engineering and Science, Jalan Sungai Long, Bandar Sungai Long, 43300 Kajang, Malaysia

^b Advanced Membrane Technology Research Centre (AMTEC), Universiti Teknologi Malaysia, 81310 Skudai, Johor, Malaysia

* Corresponding author: chongkc@utar.edu.my

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Abstract

Oxygen enriched air (OEA) is widely applied in various areas such as chemical and medical applications. Currently, cryogenic distillation and pressure swing adsorption are the two common technologies that being commercially used for the production of OEA. However, these two techniques are not economically favorable due to required intensive energy and large built-up area. With the advancement of membrane technology in separation process, it garners the interest from both industrial and academic to explore the feasibility of membrane in gas separation. In this study, polysulfone (PSF) hollow fiber membranes with poly(ether block amide) (PEBAX) coating were used for the separation of O₂/N₂ gas. The hollow fiber membranes used in this work were fabricated by phase inversion spinning process using PSF pellet, along with N,N-dimethylacetamide (DMAc) and ethanol (EtOH) as solvent and co-solvent, whereas tetrahydrofuran (THF) as additive. The fabricated membrane exhibited dense structure in the inner layer whereas finger like layer at the outer surface. The formation of this structure was attributed by rapid phase inversion of the solution arose from strong solvent used. The EDX surface mapping analysis confirmed the formation of PEBAX coating on the membrane surface. Gas permeation study in this work illustrated that the pristine PSF membrane exhibited better gas separation performance relative to the PEBAX coated membrane with 20% higher in terms of permeance. The results obtained from this work suggested that the PEBAX coating enhanced the membrane surface but not certain to improve the gas separation performance. Further study on the PEBAX materials for the membrane coating is essential to polish its potential in gas separation.

Keywords: Polysulfone, PEBAX, oxygen, nitrogen, gas separation

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INTRODUCTION

Oxygen enriched air (OEA) is widely used in various fields such as commercial, chemical and medical fields. OEA can be used for sewage treatment, enhancement combustion process and relatively novel application on hybrid system to capture carbon (Belaissaoui *et al.*, 2014). An increase in pure oxygen cost is one of the factors that contributes to the elevated demand on OEA in controlling the emission of combustion or gasification (Chong *et al.*, 2017). Current industrial technologies involved in gas separation are PSA and cryogenic distillation. In the production of OEA, cryogenic distillation can yield the product with above 99% purity of O₂ in large scale production which may exceed 100 tons/day while OEA manufactured by PSA is able to attain oxygen purity at about 95% with medium production capacity ranging from 20–100 tons/day. However, both technologies are not economically sound and consume high amount of energy (Belaissaoui *et al.*, 2014). Besides, amine solvent adsorption which is an alternative commercial technique to cryogenic distillation in carbon capture is costly and requires regular recharging (Anderson, Wang and Lin, 2012).

Attributed to all these drawbacks rooted from the current available technologies, researchers and industrial players pay their attention on the membrane technology. Membrane technology is regarded as a potential technique that can replace current industrial gas separation

techniques. This is due to the fact that membrane system is a relatively smaller in size as compared to traditional systems, mechanically simple and highly flexible enabling it to be scaled up easily (Bernardo and Clarizia, 2013). In addition, polymeric membrane is of interest as it is low cost and ease to process (Kim and Lee, 2013).

Despite of the membrane technology possesses advantages as stated above; its application in the industry of gas separation is still narrow as membrane technology is only able to cater for small scale production (25 tons/day) and to produce lower gas purity as reported by Belaissaoui *et al.* (2014). Hence, this research was mainly focused on the evaluation of performance of polymeric membrane in gas separation by using different amounts of dope solution and also characterization of membrane.

In order to be competitive with other traditional gas separation techniques, a polymeric membrane should possess three main properties which are the permeability of gases involved in gas separation, selectivity towards desired product and good mechanical and thermal characteristics (Powell and Qiao, 2006). The specific membrane material is the factor that affects the flux and selectivity of gases moving across the membrane since different polymers may consist of distinct cross-linkage, morphology and porosity (Smith and Klosek, 2001).

In this study, the coating of PEBAX material on the surface of self-fabricated membrane was evaluated in terms of its effect on the O₂/N₂ gas separation performance. Few membrane characteristics studies were performed to study the properties of the membrane in term of morphology, cross section and the quality of coating on the surface. The gas separation performance of O₂/N₂ separation was evaluated by gas permeation study.

EXPERIMENTAL

Materials

The polymer used in this study was polysulfone, which obtained in pellet form from Amoco Chemicals. The solvent used to prepare dope solution was N,N-dimethylacetamide (DMAc) with purity more than 99.5% while ethanol (EtOH) with purity more than 99% was used as co-solvent, both were obtained from Merck. Tetrahydrofuran (THF) with more than 99.5% purity from QReC was added into the dope solution as an additive for solubility enhancement of PSF in the solvent. In this study, polyether-block-amide (PEBAX) obtained from Arkema was employed as membrane coating material. The purpose of PEBAX coating on the membrane surface was to improve the performance of membrane in gas separation by forming a better selective layer on the membrane outer surface.

Preparation of dope solution

The membrane dope solution composition used in this study was detailed and presented in Table 1. The PSF pellets were placed in the vacuum oven for 24 h at 70°C to remove its moisture content before usage. The dried PSF pellets were mixed with DMAc, EtOH and THF at the predefined ratio to obtain the dope solution. The mixture was stirred for 24 h with continuous stirring and heating at 60°C until a homogeneous dope solution was attained. The dope solution was then undergone a sonication process in an ultrasonic bath for degassing to completely remove the bubbles trapped in the solution.

Table 1 Composition of dope solution.

Materials	Weight percentage (wt%)
PSF	30
DMAc	30
EtOH	10
THF	30

Fabrication of hollow fiber membrane

Phase inversion process was employed to prepare the polysulfone hollow fiber membrane in this work. The dope solution was poured into a spin reservoir and then conveyed to a spinneret with the aid of gear pump. The outer and inner diameters of spinneret were 0.6 mm and 0.3 mm, respectively. The dope solution and bore liquid were flowed out from the spinneret through the annular space and center, respectively as illustrated in Fig. 1. The bore liquid flow rate was adjusted at 1.0 mL/min.

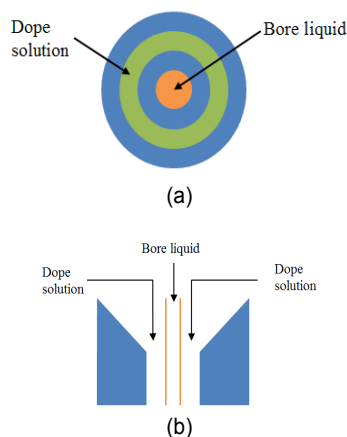


Fig. 1 The view of spinneret in term of (a) the top (b) cross-section.

The hollow fiber membranes were then delivered to a coagulation bath that filled with tap water at 25 °C. The immersion of membrane in water was carried out for one to two days prior to the drying process in a vacuum oven in order to remove the solvent residues. The hollow fiber membranes were then dried in the vacuum oven at 70°C for 24 hours to completely remove the moisture content before usage in the experimental studies. The information of spinning conditions in the phase inversion process was tabulated in Table 2.

Table 2 Spinning conditions of hollow fiber membrane by phase inversion process.

Condition	Value
Bore fluid type	Distilled water
Bore fluid temperature (°C)	25
Bore fluid flow rate (mL/min)	1.0
External coagulant type	Tap water
External coagulant temperature (°C)	25
Spinneret dimension OD/ID (mm/mm)	0.6/0.3
Air gap distance (cm)	30
Room relative humidity (%)	55 ± 5

Preparation of hollow fiber membrane coating

PEBAX coating solutions were consisted of three different PEBAX concentrations at 1 wt%, 3 wt% and 5 wt%, which then were labelled as PSF-1, PSF-3 and PSF-5, respectively. The hollow fiber membranes utilized in this study were tabulated in Table 3 for ease of reference. The coating solution was prepared by dissolving PEBAX pellet in a mixture of solvent that consisted of ethanol/water at a ratio of 70/30 with the aid of hot plate stirrer at the temperature of 75°C. The mixture was stirred until a homogenous solution was is obtained. Dip-coating method was employed to coat the fabricated hollow fiber membranes with PEBAX solution. During the coating process, 100 mL PEBAX solution was first transferred to a measuring cylinder and then the membranes prepared were dipped in the solution for 10 min. Later, the hollow fiber membranes were cured in the vacuum oven at 50°C for 6 hours. The coating procedure was repeated five times in order to obtain a perfect coating.

Table 3 Hollow fiber membrane engaged in gas permeation study.

Membrane	Description
PSF	Pristine membrane
PSF-1	Membrane with 1 wt% PEBAX coating
PSF-3	Membrane with 3 wt% PEBAX coating
PSF-5	Membrane with 5 wt% PEBAX coating

Membrane characterization study

In this study, scanning electron microscopic (SEM, Hitachi, S3400N) was used to examine the morphology of fabricated hollow fiber membranes. In order to obtain clear SEM images with a perfect-cut structure, pre-treatment was carried where the membranes were immersed in liquid nitrogen and broken cryogenically. Later, the membranes were coated with a layer of gold onto the surface by using sputter coating machine (Emitech, SC7620). Additionally, the coating quality of PEBAX on membrane surface was examined by energy dispersive X-ray spectrometer (EDX) under an acceleration voltage of 20 kV.

Gas permeation study

Gas permeation study of the PSF hollow fiber membranes was schematically illustrated in Fig. 2 using pure N₂ and O₂ gas with purity more than 99.99%. Five hollow fiber membranes were arranged together as a bundle with length of 25 cm, before being inserted into a membrane module and sealed firmly at one end by epoxy resin adhesive. The soap-bubble flow meter consisted of two inlets in which one inlet was connected to the module whereas the other inlet was linked to soap container. During the gas permeation study, pure gas was flowed into the module through the shell side driven by the gas

pressure at 5 bar. The gas was diffused across the membranes and flowed out from the lumen side of the module where the flow rate was recorded by the soap-bubble flow meter.

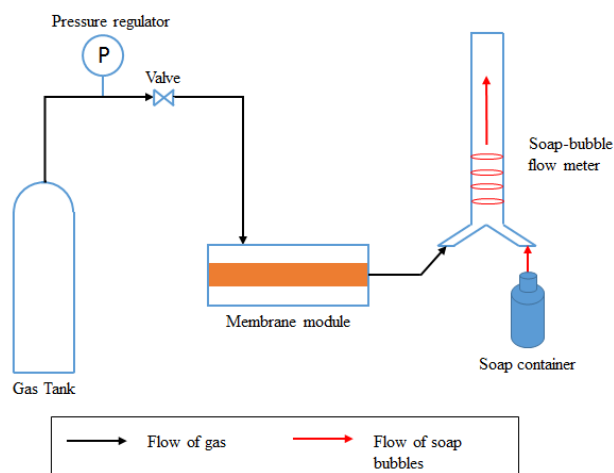


Fig. 2 Schematic diagram of gas permeation study.

Membrane permeability can be defined as the diffusion of gas across the membrane induced by the transmembrane pressure difference. It can be calculated by the following equation,

$$P_A = \frac{Ql}{A\Delta P} \frac{273.15 \times 10^6}{T} \quad (1)$$

where P_A is the membrane permeability, Q is the volumetric flowrate of gas ($\text{cm}^3 \text{ s}^{-1}$, STP), A is the effective membrane area in the unit of cm^2 , ΔP is the pressure difference across the membrane in the unit of cmHg , T is the operating temperature that is kept constant at 300 K. The unit of the permeability is usually represented as Barrer ($1 \text{ Barrer} = 10^{-10} \text{ cm}^3 (\text{STP}) \text{ cm/cm}^2 \text{ s cm Hg} = 3.35 \times 10^{-16} \text{ mol m} / \text{m}^2 \text{ s Pa}$).

Membrane permeance (GPU or $10^{-6} \text{ cm}^3 (\text{STP})/\text{cm}^2 \text{ cm Hg}$) is defined as the permeability (P_A) per unit length (l) of membrane which can be determined using Eq. 2 :

$$\frac{P_A}{l} = \frac{Q}{A\Delta P} \frac{273.15 \times 10^6}{T} \quad (2)$$

Other than permeance and permeability, selectivity is another important parameter that evaluates the performance of membrane. The selectivity of binary gas separation ($\alpha_{A/B}$) is represented by the ratio of the permeability of gas A (P_A) to the permeability of gas B (P_B) can be determined by Eq. 3:

$$\alpha_{A/B} = \frac{P_A}{P_B} \quad (3)$$

RESULTS AND DISCUSSION

Membrane morphology

The membrane morphology for the hollow fiber membrane coated with PEBAX (PSF-1, PSF-3 and PSF-5) was shown in Fig. 3. All the fabricated hollow fiber membranes illustrated dense structure at the inner layer with the finger-like layer at the outer surface. This structure was attributed by the rapid phase inversion of the solution arose from strong solvent used in this study (Wu *et al.*, 2006). The diameter of the membrane fabricated from this study was consistent which was in the range of 310 to 370 μm (Table 4). The characteristics study showed that the concentration of PEBAX was not obviously affect the wall thickness of PSF membrane where the wall thickness of the membrane was in the range of 50 to 54 μm . The EDX results on the quality of the PEBAX coating on the surface were tabulated in Table 5. Nitrogen was found to be presented only in the

PSF-1, PSF-3 and PSF-5 as nitrogen is the was contained in the amide group in PEBAX. It was interesting to note that the content of nitrogen was increased as the coating concentration of the PEBAX on hollow fiber membrane was increased, showing that the PEBAX was successfully coated on the surface of the membrane.

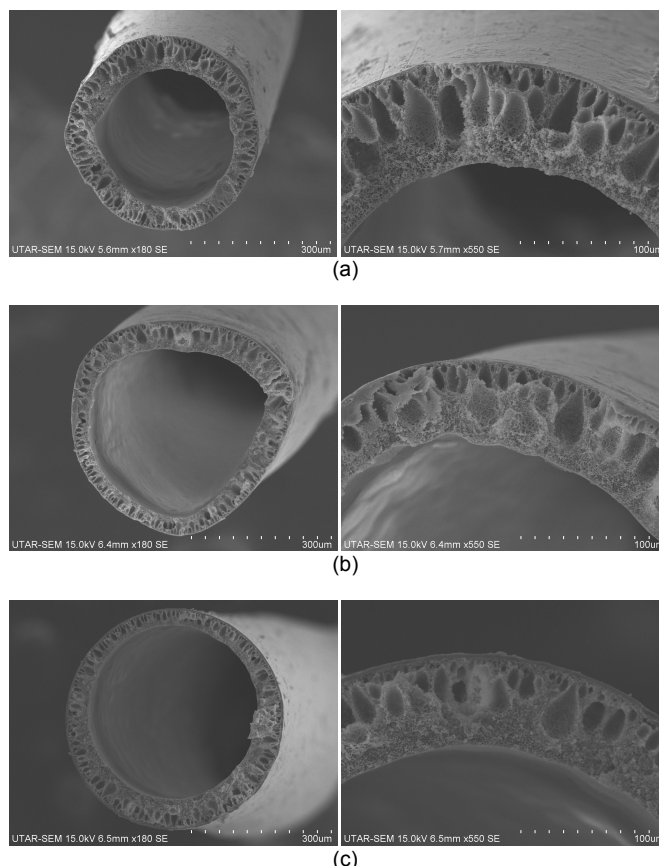


Fig. 3 Membrane cross sectional morphology of PEBAX coated PSF membrane (a) PSF-1 (b) PSF-2 (c) PSF-3.

Table 4 Membrane characteristics.

Membrane	Internal diameter (μm)	Wall Thickness (μm)
PSF	370 \pm 10	50 \pm 4
PSF-1	310 \pm 10	54 \pm 8
PSF-3	350 \pm 5	51 \pm 7
PSF-5	360 \pm 5	50 \pm 7

Table 5 EDX analysis for the membrane.

Membrane	Carbon	Oxygen	Sulphur	Nitrogen
PSF	82.61	14.19	3.20	Not detected
PSF-1	76.00	17.82	3.08	3.10
PSF-3	75.61	16.70	3.01	4.68
PSF-5	74.44	17.54	3.09	4.93

Membrane Performance in OEA Production

The gas permeation results in O_2/N_2 separation performance by PSF membranes were recorded in Table 6. It was found that both permeability of oxygen and O_2/N_2 selectivity were decreased when the concentration of PEBAX on the membrane was increased. The coating of PEBAX solution on the membrane surface was able to improve the smoothness of the membrane surface, nevertheless, it still possessed little advantage in the separation performance. The results obtained in the gas permeation study were opposed to the literature studies whereby the addition of PEBAX coating on the membrane surface was able to improve the permeability while

maintaining the selectivity of the gas separation (Wang et al. 2014). However, it was noteworthy to mention that the works done by previous literature reviews were carried out using flat sheet membrane coated with PEBAX solution (Ren et al., 2012, Ahmadpour et al., 2014 and Wang et al. 2014). The lower gas permeation results obtained from this study might attribute to the PEBAX coating as some of the hollow fiber membranes were tended to stick to each other during the curing process. Hence, the additional optimization was required in our future work in order to release the potential of PEBAX coating on the hollow fiber membrane. On the other hand, the performance of pristine PSF membrane was attributed to its superior gas separation performance which eventually led to higher yield of OEA production.

Table 6 Performance of PSF membrane on O₂/N₂ separation.

Membrane	Permeance (GPU)		Permeability (Barrer)		O ₂ /N ₂ selectivity
	O ₂	N ₂	O ₂	N ₂	
PSF	51.00	13.00	12.75	3.25	3.92
PSF-1	39.81	10.73	9.95	2.68	3.71
PSF-3	36.15	11.52	9.04	2.88	3.14
PSF-5	33.28	10.14	8.32	2.54	3.28

CONCLUSION

In this study, PSF hollow fiber membranes were successfully fabricated and coated with PEBAX materials under different weight concentrations. Membrane characterization and gas permeation studies were conducted to understand the properties of the membranes and evaluate the membrane performance in the O₂/N₂ gas separation for OEA production. The EDX results indicated that PEBAX was successfully coated on the surface of the membrane with the presence of nitrogen from amide group. Results from gas permeation study suggested that the PEBAX coating exhibited modest advantages relative to the pristine membrane in O₂/N₂ gas separation but it was able to enhance the surface of the hollow fiber membrane. The result from this study proposed that PEBAX materials were able to act as membrane surface enhancement and further study on the PEBAX materials on the membrane coating was essential to release its potential in gas separation.

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