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# Influence of microstructural variations on morphology and separation properties of polybutadiene-based polyurethanes†

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Polybutadiene-based polyurethanes with different *cis/trans*/1,2-vinyl microstructure contents are synthesized. The phase morphology and physical properties of the polymers are investigated using spectroscopic analysis (FTIR and Raman), differential scanning calorimetry (DSC), X-ray scattering (WAXD and SAXS) and atomic force microscopy (AFM). In addition, their gas transport properties are determined for different gases at 4 bar and 25 °C. Thermodynamic incompatibility and steric hindrance of pendant groups are the dominant factors affecting the morphology and properties of the PUs. FTIR spectra, DSC, and SAXS analysis reveal a higher extent of phase mixing in high vinyl-content PUs. Moreover, the SAXS analysis and AFM phase images indicate smaller microdomains by increasing the vinyl content. Smaller permeable soft domains as well as the lower phase separation of the PUs with higher vinyl content create more tortuous pathways for gas molecules and deteriorate the gas permeability of the membranes.

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## Introduction

Knowledge about the structure–property relationships of block copolymers is necessary for selecting proper materials for any specific application. Most properties of block copolymers, such as polyurethanes (PUs), are directly related to their multiphase morphology.<sup>1–3</sup> From the theoretical perspective, the difference between the polarity of the urethane hard segment and polyol soft segment (*i.e.*, thermodynamic incompatibility) leads to a phase-separated structure.<sup>4</sup> Besides, higher chain mobility (*i.e.*, kinetic factor) of either soft or hard segments can improve phase separation.<sup>5,6</sup> Nevertheless, complete phase segregation rarely takes place in PU systems. The intra- and inter-segmental hydrogen bonding in the hard and soft domains determine the block structures. Besides, the crystallinity and glass transition temperature of each block determine the morphology.<sup>7</sup>

PU morphology can be manipulated by the chemistry of its components (*e.g.*, polyol and diisocyanates), thermal annealing,<sup>8,9</sup> and synthesis methods.<sup>10</sup> The extent of phase segregation was found to have a major impact on the mechanical and thermal properties of PUs.<sup>10</sup> The dissolution of soft segments in the hard domains disrupts hydrogen bonding and the molecular order in urethane chains that could lead to lower mechanical properties.<sup>11–13</sup> It was shown that using symmetrical monomers in PU synthesis can considerably improve phase separation and mechanical properties.<sup>14–16</sup> Moreover, gas permeation in PU membranes is significantly influenced by the degree of phase separation between the hard and soft segments. Since the soft domains are the main diffusion pathway for gas molecules in PU membranes, phase intermixing can adversely affect gas transport properties. A wide range of gas permeability has been found for different PUs, ranging from 10 to 1000 barrer (1 barrer = 10<sup>-10</sup> cm<sup>3</sup> (STP) cm cm<sup>-2</sup> s<sup>-1</sup> cmHg<sup>-1</sup>), which discloses the impact of morphology on the separation properties of PU membranes.<sup>17,18</sup>

The degree of phase separation can be altered by the extent of hydrogen bonding, steric hindrance and incompatibility between the two blocks. The urethane-urea bonds or cross-linking the hard segments favor phase segregation.<sup>19,20</sup> Moreover, moderate phase mixing was observed in polyether-based PUs because of the association of oxygen groups in polyether with urethane –NH groups in the hard segments. Polyethylene glycol (PEG) and polypropylene glycol (PPG) are among the most common polyethers in the PU industry. Compared to PPG, PEG has a higher tendency to mix with the urethane hard segments

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due to its lower hydrophobicity. Therefore, the gas permeability of PEG-based PU membranes is inferior due to their lower phase separation and semi-crystalline structure as compared to the PPG-based membranes with amorphous soft domains.<sup>21</sup> In addition to thermodynamic parameters, it was claimed that steric hindrance and chain conformation play an important role in the disruption of hydrogen bonding between the hard and soft segments.<sup>22–24</sup> In our previous publication, a hydroxyl-terminated PEG-*b*-PPG block copolymer was used for PU synthesis.<sup>21</sup> The methyl side group in PPG spatially hindered the interaction between the PEG and the hard segment. The membranes showed a *ca.* 400% improvement in CO<sub>2</sub> permeability compared to the PEG-based PU membrane. Although polyols with methyl and/or fluorine pendant groups would enhance phase segregation,<sup>25,26</sup> the presence of bulky side groups in the chain extender has the opposite effect.<sup>10</sup> Moreover, tuning the morphology of block copolymer membranes for a better gas separation performance was accomplished by the incorporation of nanoparticles in the polymer matrix.<sup>27–31</sup> Interaction of fillers with the functional groups of either soft or hard blocks reduces phase-intermixing in the PU matrix.

The development of PU systems with low miscibility between hard and soft segments is desirable in gas separation membranes. Polydimethylsiloxane (PDMS) and hydroxyl-terminated polybutadiene (HTPB) are well-known polyols in this regard. Their lower solubility parameter compared to other polyols results in a well-phase-separated PU structure, the properties of which are entirely independent of the hard segment chemistry.<sup>32</sup> Additionally, the PUs based on PDMS and HTPB soft segments exhibit high gas permeability, excellent hydrophobicity and good thermal and oxidation stability.<sup>17,33</sup> The large polarity difference between the urethane and PBD creates isolated hard domains with a nearly sharp interface boundary.

The morphology of polybutadiene-based PUs has been studied over the past years.<sup>34–36</sup> Nevertheless, the impact of structural configuration (*i.e.*, 1,2-vinyl, *cis*, *trans*) on morphology and properties of PUs has not yet been investigated. The pendant 1,2-vinyl moieties in PBD can inhibit the interactions between the two segments and also can suppress the PU crystallinity. In contrast, the viscosity and  $T_g$  of the polyol, as two critical parameters in the determination of PU morphology, increase with vinyl content. Also, the polarity and chain mobility of PUs can be influenced by the *cis/trans* ratio in the polyol.<sup>37</sup> In this study, hydroxyl-terminated PBDs with different vinyl contents (7%, 20%, 65%, and 90%) were utilized for PU synthesis. Subsequently, the effect of *cis* and *trans* microstructures on the properties of PUs was evaluated by X-ray scattering methods and thermo-mechanical analysis. The gas transport properties of the synthesized membranes were also explored for a variety of gases.

## Experimental

### Materials

Four types of hydroxyl-terminated polybutadiene with different vinyl contents (7%, 20%, 65%, and 90%) were kindly provided

by Nippon Soda Co., Japan, and Cray Valley Co., USA. All polyols were dried under vacuum at 80 °C for 48 h to remove residual water before synthesis. The chemical structure and properties of HTPBDs were shown in Table 1. Isophorone diisocyanate (IPDI), super dehydrated dimethylacetamide (DMA), super dehydrated tetrahydrofuran (THF), and dimethylformamide (DMF) were obtained from Wako Pure Chemical Industries, Ltd., Japan and used as received. Octanediamine (ODA, Tokyo Chemical Industry) and dibutyltin dilaurate (DBTDL, Wako Pure Chemical Industries) were used as chain extender and catalyst, respectively, after being dried over 4 Å molecular sieve (Nacalai Tesque, Japan).

### Polymer synthesis

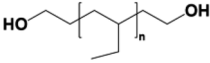
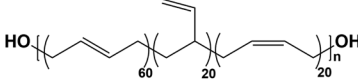
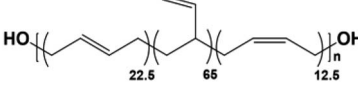
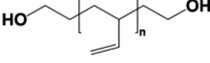
The chemical reaction was shown in Fig. S1.† An exact amount of IPDI (10 mmol) was added dropwise into the stirred HTPBD solution (5 mmol) in 20 mL DMAc/THF (1/1 vol%). 0.1 mL DBTDL was added to the reactor, and the mixture was stirred under Ar atmosphere at 70 °C for 2 h to obtain the NCO-terminated prepolymer. The polymerization was proceeded by the addition of 5 mmol ODA (dissolved in 20 mL THF) into the cooled-room temperature reaction mixture and stirred few minutes. After the completion of the reaction, an exact amount of the mixture was cast into Teflon Petri-dishes, and the membrane films were formed under controlled evaporation at 50 °C for 24 h. The prepared membranes with the thickness of 60–70 μm were further dried under vacuum at 80 °C for 24 h before testing. The synthesized polymers were denoted PU(*t,c,v*), wherein *t*, *c*, and *v* respectively stand for *trans*, *cis* and vinyl isomer contents in HTPBD.

### Characterization

The chemical structure of the synthesized PUs was determined by Raman (Horiba XploRA, Japan) and FTIR (Shimadzu IR Tracer-100 spectrometer, Japan) spectroscopies. Raman spectra were collected in the range of 3100–400 cm<sup>−1</sup> with a resolution of 1 cm<sup>−1</sup> and a laser excitation wavelength of 532 nm. ATR-FTIR spectra were obtained in the range of 4000–600 cm<sup>−1</sup> with a resolution of 4 cm<sup>−1</sup> and 128 scans. The morphology of the samples was determined by wide-angle X-ray diffraction (WAXD, Rigaku RINT XRD, Japan) using CuKα radiation ( $\lambda = 1.54 \text{ \AA}$ ) and the scattering range of  $2\theta = 5\text{--}40^\circ$ . Small-angle X-ray scattering (SAXS, NANO-Viewer, Rigaku, Co, Ltd., Japan) was performed using a 1.2 kW rotating-anode X-ray generator (40 kV, 30 mA). The X-ray wavelength and sample to detector distance were 1.54 Å and 1300 mm, respectively. Thermal transition temperatures of the PUs were detected by differential scanning calorimetry (DSC, Bruker DSC 3100SA, Germany) under N<sub>2</sub> atmosphere in the range of −100 °C to 200 °C and at a scanning rate of 10 °C min<sup>−1</sup>. Thermogravimetric analysis (TGA, Rigaku Thermo Plus EVO 2, Japan) was performed under an air atmosphere at the heating rate of 10 °C min<sup>−1</sup> up to 800 °C. Phase morphological analysis was carried out by atomic force microscopy (AFM, Nano wizard 2, JPK Instruments, Japan) under AC mode with a 6 N m<sup>−1</sup> cantilever.



Table 1 Chemical structure and physical properties of polyols

Sample	Polyol	Chemical structure	$M_n$ (g mol <sup>-1</sup> )	1-2, Vinyl%	Trans%	Cis%	$T_g$ of polyol (°C)
PU(0,0,7)	GI-3000®		3100	7	—	—	-38.0
PU(60,20,20)	Poly bd®R-45HTL		2800	20	60	20	-80.0
PU(22.5,12.5,65)	Krasol®LBH-P3000		3200	65	22.5	12.5	-48.0
PU(0,0,90)	G-3000®		3000	90	—	—	-17.0

### Gas permeation test

Pure gas permeability of the PU membranes to H<sub>2</sub>, O<sub>2</sub>, N<sub>2</sub>, CH<sub>4</sub>, and CO<sub>2</sub> was measured using the constant volume-variable pressure method at 4 bar and 25 °C. The permeate pressure and temperature were monitored by a sensor (Keller PAA 33X) connected to Labview software. The permeability of each membrane was measured three times to assess the reproducibility of the data. Gas transport across polymeric membranes is described by the solution-diffusion mechanism, where the permeability is the product of diffusivity coefficient ( $D$ ) and solubility coefficient ( $S$ ):  $P = D \times S$ . The gas permeability is given by the following equation:

$$P = J_i \frac{l}{\Delta p} = 10^{10} \frac{273.15}{76} \frac{V}{AT} \left( \frac{dp}{dt} \right) \frac{l}{\Delta p} \quad (1)$$

where  $P$ ,  $l$  and  $\Delta p$  are the permeability coefficient (barrer), membrane thickness (cm) and the pressure difference across the membrane (bar), respectively.  $V$ ,  $A$  and  $T$  are the volume in the permeate side, the active area of the membrane and measurement temperature, respectively. The ideal selectivity ( $\alpha_{A/B}$ ) of the membranes was calculated from pure gas permeation data:

$$\alpha_{ij} = P_i/P_j \quad (2)$$

where  $P_i$  and  $P_j$  are the permeability coefficients of the penetrants  $i$  and  $j$ , respectively.

## Results and discussion

### Spectroscopic analysis

Fig. 1a represents evidence for the existence of unsaturated C=C bonds and carbonyl groups in the synthesized PUs. The disappearance of the NCO peak in FTIR spectra at 2300 cm<sup>-1</sup> indicates the completion of the reaction. The bands appearing at 907 cm<sup>-1</sup>, 995 cm<sup>-1</sup>, and 3070 cm<sup>-1</sup> are assigned to the C=C bending vibrations and stretching mode of 1,2-vinyl substitutes,<sup>38</sup> which are absent in the PU(0,0,7) sample. Besides, the

band at 965 cm<sup>-1</sup> refers to the bending C=C vibrations of the *trans*-1,4-PBD microstructure,<sup>38,39</sup> where the PU(60,20,20) with 60% *trans* isomer exhibits the most prominent peak than other samples. The obvious peaks at 1640 cm<sup>-1</sup> and 1710 cm<sup>-1</sup> correspond to the bonded and free carbonyl groups, respectively, which are helpful in comprehending the phase separation of PUs.<sup>40</sup> The presence of pendant functional groups impedes the formation of hydrogen bonding between the hard and soft segments. Nevertheless, the thermodynamic parameter is the dominant factor for the lower phase separation of PU(22.5,12.5,65) compared to that of PU(60,20,20). The R45-HT polyol is less polar than LBHP-3000 due to its higher density of *trans*-isomer. Therefore, R45-HT has a lower tendency to mix with the polar urethane segments. The steric effect of the pendant functional groups is significant in similar PU(0,0,90) and PU(0,0,7) structures. The higher phase separation of PU(0,0,7) is associated with a higher content of bulky ethyl groups, which effectively disrupt the hydrogen bonding between the urethane -NH groups and the polyol. The A value which is a well-known parameter for the evaluation of the bulkiness of functional groups is 1.75 kcal mol<sup>-1</sup> and 1.35 kcal mol<sup>-1</sup> for ethyl and 1,2-vinyl substitutes, respectively.<sup>41,42</sup>

The degree of phase separation can be evaluated by hydrogen bonding index (HBI), calculated from eqn (3):

$$\text{HBI} = \frac{I_{\text{C=O, bonded}}}{I_{\text{C=O, free}}} \quad (3)$$

where  $I_{\text{C=O, bonded}}$  and  $I_{\text{C=O, free}}$  correspond to the absorbance intensity of bonded and free carbonyl peaks. The HBI value of the synthesized PUs decreased as follows: PU(0,0,7)(5.6) > PU(60,20,20)(3.0) > PU(22.5,12.5,65)(2.3) > PU(0,0,90)(1.4), which is in agreement with the above discussion. The disruption of hydrogen bonding due to the steric effect of bulky functional groups gives rise to a larger HBI value.

Understanding the PU characteristics depends on a comprehensive study of the polybutadiene structure; thus, it helps analyze the different isomers in PBD quantitatively. Raman spectroscopy detects the C=C stretching vibrations in the *trans*



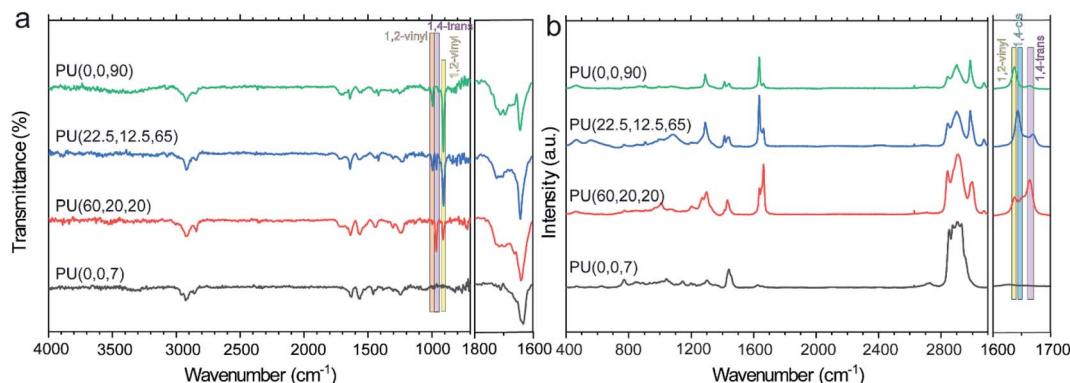


Fig. 1 FTIR (a) and Raman (b) spectra of the synthesized polyurethanes.

unit since they are infrared inactive.<sup>43</sup> As shown in Fig. 1b, the peaks at  $1631\text{ cm}^{-1}$ ,  $1651\text{ cm}^{-1}$  and  $1663\text{ cm}^{-1}$  are ascribed to the C=C stretching modes of 1,2-vinyl, *cis* and *trans* in the polybutadiene-based PUs.<sup>44,45</sup> The molar content of each configuration can be evaluated by the integration of the corresponding peak. For example, the vinyl content of PU(0,0,90), PU(22.5,12.5,65), PU(60,20,20) and PU(0,0,7) was calculated to be around 80%, 61%, 25%, and 0%, respectively, which are in a good agreement with the pure polyol specifications reported in Table 1.

### Thermomechanical properties

Thermal transitions in block copolymers provide useful information about their morphology. DSC thermograms were collected for each sample in the second heating cycle to erase

the thermal history (Fig. 2a). Glass transition temperature ( $T_g$ ) of the soft segment is an indication of the phase separation; however, the synthesized polymers display neither melting peaks nor  $T_g$  for the hard segments because of the asymmetric structure of IPDI.<sup>19</sup>  $T_g$  of the PUs was interpreted in comparison with its value for the pure polyols in Table S1.† The highest  $T_g$  was observed at  $-2\text{ }^\circ\text{C}$  for the PU(0,0,90), which also has a significant deviation from the  $T_g$  of GI-3000 ( $\Delta T_g = 15\text{ }^\circ\text{C}$ , Table S1†). In contrast,  $T_g$ s of PU(0,0,7) and PU(60,20,20) are about the same as their pure polyols, GI-3000 and R45-HT, indicating high purity of the soft domains (*i.e.*, greater phase separation). Besides, the slope of the  $T_g$ s of PU(0,0,7) and PU(60,20,20) is sharper than that of the other samples. This can be attributed to the lower phase mixing which can lead to a lower restriction on chain mobility. The trend of phase

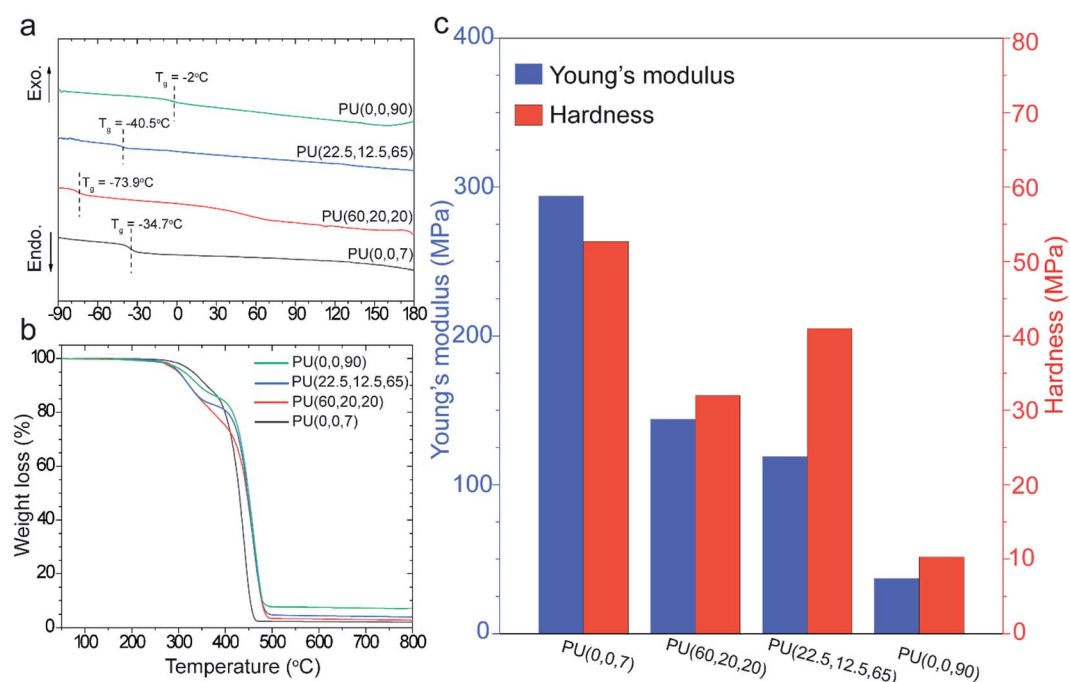


Fig. 2 DSC thermograms of the synthesized PUs (a); thermogravimetical analysis of the synthesized PUs under air atmosphere (b); mechanical properties of the prepared membranes using nanoindentation method (c).



segregation observed in DSC was supported by FTIR, where PU(0,0,90) and PU(0,0,7) have the highest and lowest phase mixing, respectively.

Thermal stability of the synthesized PUs was studied by TGA in Fig. 2b. Typically, PUs undergo a two-step degradation process; the decomposition commences by cleavage of urethane bonds in the hard segments at around 300 °C. Although the polymers consist of the same hard segment chemistry, the temperature and decomposition rate are different in each sample. For example,  $T_{10\%}$  (the temperature at which 10% weight loss occurs) of PU(0,0,90) at 330 °C improves to 362 °C for PU(0,0,7) (Table S1†). It is supposed that PU(0,0,7) benefits from stronger hydrogen bonds in the hard domains since it has lower interferences with the polyols. Therefore, the energy required for decomposition is larger than other samples with lower phase separation. This behavior was also reported in previous works.<sup>20,46</sup> Although degradation of the hard segments starts earlier for the PUs with unsaturated 1,2-vinyl groups, the decomposition of these samples, *i.e.*, PU(60,20,20), PU(22.5,12.5,65) and PU(0,0,90), at higher temperature (above 400 °C) were improved compared to PU(0,0,7). The crosslinking reactions proceeded *via* 1,2-vinyl groups at high temperatures delayed the degradation. The higher residual char yield of the unsaturated PUs confirms this hypothesis (Table S1†).

Type and content of microstructures (vinyl, *cis*, *trans*) in polybutadienes determine the mechanical properties of the synthesized PUs. Fig. 2c compares elastic modulus ( $E$ ) and hardness ( $H$ ) of the samples, measured by the nanoindentation method. An overall decay in the mechanical properties was observed by increasing the vinyl content. For example, the elastic modulus of saturated PU(0,0,7) decreased from 293.9 MPa to 36.2 MPa for PU(0,0,90), a significant loss of about 85%. It is believed that the lower phase separation, higher  $T_g$  and the distorted structure of the PUs (refer to XRD section) in the presence of vinyl pendant groups are responsible for this decline.<sup>13</sup> In addition, the higher modulus of PU(60,20,20) ( $E = 151$  MPa) than PU(22.5,12.5,65) ( $E = 102$  MPa) can be explained by higher *trans*-microstructure content in the R-45HT butadiene backbone.<sup>47</sup> Nevertheless, we are uncertain about the influence of 1,2-vinyl groups on the hardness of the PUs. It seems a lower hardness of PU(60,20,20) and PU(22.5,12.5,65) compared to other samples is related to the higher flexibility of these polymers due to the presence of *trans* isomers.

### Morphology and microstructure

AFM phase images of the PUs display remarkable changes in the phase separation. The difference in modulus of the hard and soft domains manifests itself in the color profile of the captured images, where darker and lighter areas refer to the hard-rich and soft-rich domains, respectively. Fig. 3a shows smaller and more phase-mixed structures for the samples with higher 1,2-vinyl contents. The average size of the soft domain for PU(0,0,90) was 10–30 nm, while it increased to 50–70 nm in PU(0,0,7). The larger domain size of the PU(0,0,7) compared to other samples was also confirmed by SAXS data (ESI†). It seems the self-assembly of the soft segments to form soft domains is

restricted more by increasing the vinyl content. The vinyl groups can act as anchors in the polymer backbone, which decreases the chain mobility (*i.e.*, higher viscosity).

The influence of vinyl groups on the microstructure of the synthesized PUs was investigated by WAXD, as shown in Fig. 3b. A broad amorphous peak was observed at  $2\theta = 15\text{--}20^\circ$  for all samples. As observed in Fig. 3b, the peak shifts to lower frequencies by increasing the vinyl content. This increase in chain spacing by vinyl content can be explained by the repulsion effect due to the dipole–dipole interactions between the vinyl and urethane C=O groups.<sup>48,49</sup>

The average size of the microdomains can be evaluated from the SAXS patterns (shown in Fig. 3c) using the peak analysis of the Lorentz-corrected curves (Fig. S2 and Table S2†). The estimated volume fraction of the microdomains obtained from the analysis of the one-dimensional correlation function.<sup>50–52</sup> (described in ESI†) is higher than the nominal volume fraction of hard or soft segments derived from the molar volumes of each monomer (Table S2†), which reveals the occurrence of phase mixing in the hard microdomains of the synthesized polymers.<sup>53</sup>

### Gas permeation

Table 2 demonstrates the separation performance of the PU membranes for CO<sub>2</sub>, N<sub>2</sub>, CH<sub>4</sub>, H<sub>2</sub>, and O<sub>2</sub> at 4 bar at 25 °C. CO<sub>2</sub> permeability of the membranes is higher than the permeability of H<sub>2</sub> despite its larger gas molecular size. The same result was also seen for CH<sub>4</sub> permeation with a bigger dynamic diameter ( $d_k = 3.8$  Å) than N<sub>2</sub> ( $d_k = 3.64$  Å). The order of gas permeability through the PU membranes is  $P_{(\text{CO}_2)} > P_{(\text{H}_2)} > P_{(\text{O}_2)} > P_{(\text{CH}_4)} > P_{(\text{N}_2)}$ , which is a common trend in rubbery materials such as PDMS and Pebax.<sup>54,55</sup> In general, the gas transport properties in PU membranes strongly rely on the gas solubility, which is controlled by the critical temperature ( $T_c$ ) of the gas: CO<sub>2</sub> (304.2 K) > CH<sub>4</sub> (190.9 K) > O<sub>2</sub> (154.6 K) > N<sub>2</sub> (126.3 K) > H<sub>2</sub> (33.2 K).<sup>56</sup>

The differences in morphology of the hard and soft microdomains across the chemically-similar samples can lead to different gas permeability. As can be observed from Table 2, the gas permeability of the samples with lower vinyl content is generally higher than their high-vinyl-content counterparts. This can be due to the higher chain flexibility of samples with lower vinyl content, which leads to a higher fractional free volume in the soft domains. It is commonly observed in rubbery polymers that a decrease in  $T_g$  leads to a higher gas diffusivity.<sup>57</sup> In addition, volume fraction, size and purity of the microdomains can affect gas permeability of the samples with different vinyl contents.

In comparing PU(0,0,7) and PU(0,0,90), the reduction in the gas permeability with vinyl content can also be attributed to a lower purity of the soft microdomains in PU(0,0,90) (*i.e.*, lower phase separation), as confirmed by DSC and FTIR results. Besides, the size of soft microdomains (Fig. 3a and Table S2†) in PU(0,0,90) is smaller than that in PU(0,0,7). Therefore, gas molecules face higher tortuosity in the permeable soft phase, leading to a lower gas permeability of PU(0,0,90).



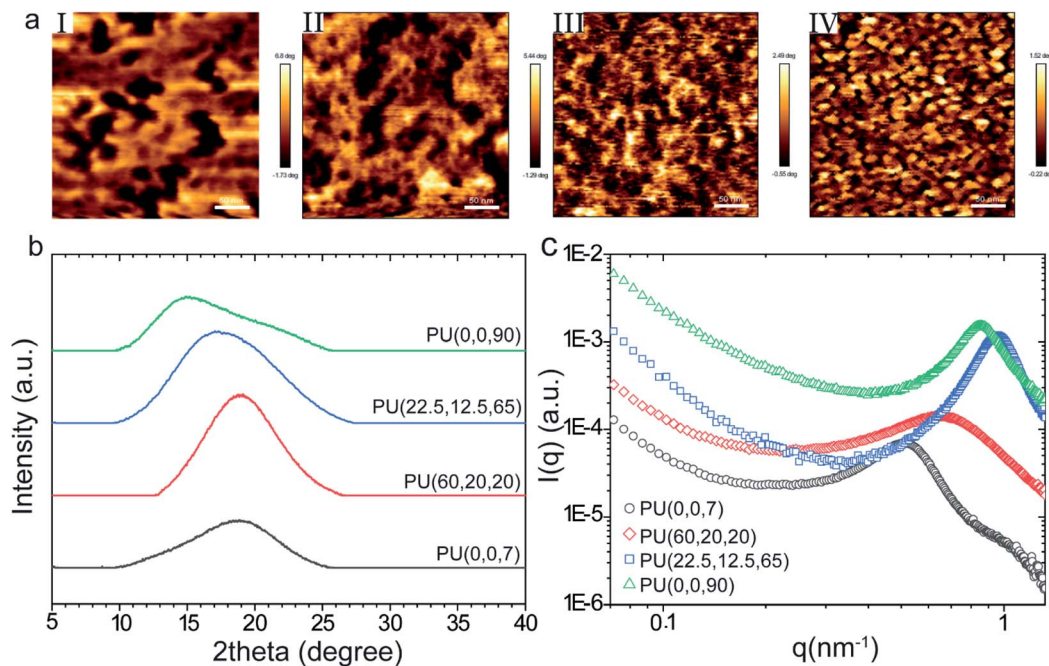


Fig. 3 AFM phase images (a) for the prepared membranes: (I) PU(0,0,7), (II) PU(60,20,20), (III) PU(22.5,12.5,65), (IV) PU(0,0,90), scale bar is 50 nm; WAXD (b) and SAXS (c) diffraction patterns of the synthesized PUs.

Regarding the morphology of PU(60,20,20) and PU(22.5,12.5,65), similar to above, the size of soft microdomains (Fig. 3a and Table S2†) decreased with an increase in vinyl content. Additionally, based on DSC and FTIR results, the purity of the soft domains is only slightly affected in these samples by increasing the vinyl content. Consequently, the higher gas permeability of PU(60,20,20) can also be attributed to the larger microdomains and the lower  $T_g$  of the soft segments in this sample.

As can be observed in Table 2, the  $\text{CO}_2/\text{N}_2$  and  $\text{CO}_2/\text{CH}_4$  selectivities of PU(0,0,90) are larger than those of PU(0,0,7), which can be attributed to a higher phase mixing and a more tortuous pathway for gas species in the former sample compared to the latter, as discussed earlier. Conversely, the  $\text{CO}_2/\text{N}_2$  and  $\text{CO}_2/\text{CH}_4$  selectivities are almost unchanged among PU(60,20,20) and PU(22.5,12.5,65). This result can be attributed to the similarity in the extent of phase mixing in the soft microdomains of these samples. It can also be observed in Table 2 that  $\text{O}_2/\text{N}_2$  selectivity is almost the same across all samples, being slightly higher in PU(0,0,90). The higher  $\text{O}_2/\text{N}_2$

selectivity of PU(0,0,90) can be associated with the higher sieving ability of the membrane due to its smaller microdomains and more rigidified structure compared to other samples (higher soft domain  $T_g \sim -2^\circ\text{C}$ ).

## Conclusion

A series of polybutadiene-based polyurethane structures with different amounts of *trans*, *cis* and vinyl structural configurations were synthesized. The effect of microstructural variations on the phase morphology, thermo-mechanical properties and gas transport behavior of the PU films were studied. The thermal stability of the PUs was slightly improved (up to  $25^\circ\text{C}$ ) by increasing the vinyl contents. However, the elastic modulus and hardness deteriorated more than 80% for the PUs with a large amount of vinyl groups. Higher *trans* content in PU(60,20,20) improved its chain mobility ( $T_g = -73.9^\circ\text{C}$ ) and phase separation compared to other samples. The  $\text{CO}_2$  permeability of PU(60,20,20)  $\sim 74$  barrer was more than twice the permeability of the PUs without *trans*-microstructure, PU(0,0,7)

Table 2 Gas permeability and ideal selectivity of the PU membranes at 4 bar and  $25^\circ\text{C}$

Membrane	Permeability (barrer)					Selectivity		
	$\text{H}_2$	$\text{CO}_2$	$\text{O}_2$	$\text{N}_2$	$\text{CH}_4$	$\text{CO}_2/\text{N}_2$	$\text{CO}_2/\text{CH}_4$	$\text{O}_2/\text{N}_2$
PU(0,0,7)	$28.8 \pm 0.8$	$34.7 \pm 1.8$	$8.4 \pm 0.3$	$2.7 \pm 0.1$	$4.7 \pm 0.3$	$12.9 \pm 0.8$	$7.4 \pm 0.6$	$3.1 \pm 0.2$
PU(60,20,20)	$23.4 \pm 0.6$	$79.8 \pm 3.0$	$10.8 \pm 0.5$	$3.5 \pm 0.1$	$11.6 \pm 0.7$	$22.8 \pm 1.1$	$6.9 \pm 0.5$	$3.1 \pm 0.2$
PU(22.5,12.5,65)	$29.2 \pm 0.5$	$65.2 \pm 3.2$	$9.5 \pm 0.3$	$3.0 \pm 0.2$	$9.3 \pm 0.5$	$21.7 \pm 1.8$	$7.0 \pm 0.5$	$3.2 \pm 0.2$
PU(0,0,90)	$19.4 \pm 0.4$	$28.1 \pm 0.8$	$4.1 \pm 0.2$	$1.1 \pm 0.1$	$2.7 \pm 0.1$	$25.5 \pm 2.4$	$10.4 \pm 0.5$	$3.7 \pm 0.4$



and PU(0,0,90). SAXS results and AFM phase images revealed smaller microdomains for the samples with higher vinyl content, which created a more tortuous diffusion pathway for gas molecules within the membranes. Besides, the intermixing of hard and soft segments increased by increasing the vinyl contents (*i.e.*, lower phase separation).

## Author contributions

Ali Pournaghshband: conceptualization, methodology, investigation, data analysis, writing-review & editing. Mahdi Shahrooz: data analysis, investigation, software, writing-review & editing. Takuma Yamamoto: experiments & investigation, data analysis. Ansori Muchtar: experiments & investigation, data analysis. Masateru M. Ito: investigation, data analysis. Daisuke Yamaguchi: data analysis, software, validation. Mikihito Takenaka: data analysis, validation. Easan Sivaniah: supervision; conceptualization; validation; writing-review & editing. Behnam Ghalei: supervision; conceptualization; validation; writing-review & editing.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 A. Fakhar, M. Sadeghi, M. Dinari, M. Zarabadipoor and R. Lammertink, *Eur. Polym. J.*, 2020, **122**, 109346.
- 2 M. Radjabian and V. Abetz, *Prog. Polym. Sci.*, 2020, 101219.
- 3 Z. Wang, A. P. Isfahani, K. Wakimoto, B. B. Shrestha, D. Yamaguchi, B. Ghalei and E. Sivaniah, *ChemSusChem*, 2018, **11**, 2744–2751.
- 4 A. Fakhar, M. Sadeghi, M. Dinari and R. Lammertink, *J. Membr. Sci.*, 2019, **574**, 136–146.
- 5 M. A. Hood, B. Wang, J. M. Sands, J. J. La Scala, F. L. Beyer and C. Y. Li, *Polymer*, 2010, **51**, 2191–2198.
- 6 Y. Li, T. Gao and B. Chu, *Macromolecules*, 1992, **25**, 1737–1742.
- 7 Y. Li, T. Gao, J. Liu, K. Linliu, C. R. Desper and B. Chu, *Macromolecules*, 1992, **25**, 7365–7372.
- 8 J. T. Koberstein and A. F. Galambos, *Macromolecules*, 1992, **25**, 5618–5624.
- 9 N. Hossieny, V. Shaayegan, A. Ameli, M. Saniei and C. Park, *Polymer*, 2017, **112**, 208–218.
- 10 P. Krol, *Prog. Mater. Sci.*, 2007, **52**, 915–1015.
- 11 Q. Tian, G. Yan, L. Bai, X. Li, L. Zou, L. Rosta, A. Wacha, Q. Li, I. Krakovský and M. Yan, *Polymer*, 2018, **147**, 1–7.
- 12 J. Koberstein, A. Galambos and L. Leung, *Macromolecules*, 1992, **25**, 6195–6204.
- 13 L. M. Leung and J. T. Koberstein, *Macromolecules*, 1986, **19**, 706–713.
- 14 C. Prisacariu, *Polyurethane elastomers: from morphology to mechanical aspects*, Springer Science & Business Media, 2011.
- 15 M. Sadeghi, M. A. Semsarzadeh, M. Barikani and B. Ghalei, *J. Membr. Sci.*, 2011, **385**, 76–85.
- 16 S. Sami, E. Yildirim, M. Yurtsever, E. Yurtsever, E. Yilgor, I. Yilgor and G. L. Wilkes, *Polymer*, 2014, **55**, 4563–4576.
- 17 H. B. Park, C. K. Kim and Y. M. Lee, *J. Membr. Sci.*, 2002, **204**, 257–269.
- 18 A. Pournaghshband Isfahani, M. Sadeghi, K. Wakimoto, B. B. Shrestha, R. Bagheri, E. Sivaniah and B. Ghalei, *ACS Appl. Mater. Interfaces*, 2018, **10**, 17366–17374.
- 19 A. P. Isfahani, B. Ghalei, R. Bagheri, Y. Kinoshita, H. Kitagawa, E. Sivaniah and M. Sadeghi, *J. Membr. Sci.*, 2016, **513**, 58–66.
- 20 A. P. Isfahani, B. Ghalei, K. Wakimoto, R. Bagheri, E. Sivaniah and M. Sadeghi, *J. Mater. Chem. A*, 2016, **4**, 17431–17439.
- 21 A. P. Isfahani, M. Sadeghi, K. Wakimoto, A. H. Gibbons, R. Bagheri, E. Sivaniah and B. Ghalei, *J. Membr. Sci.*, 2017, **542**, 143–149.
- 22 C. W. Meuse, X. Yang, D. Yang and S. L. Hsu, *Macromolecules*, 1992, **25**, 925–932.
- 23 J.-L. Hong, C. P. Lillya and J. C. Chien, *Polymer*, 1992, **33**, 4347–4351.
- 24 M. M. Coleman, G. J. Pehlert and P. C. Painter, *Macromolecules*, 1996, **29**, 6820–6831.
- 25 P. J. Flory and M. Volkenstein, *Biopolymers*, 1969, **8**, 699–700.
- 26 W. P. Chen, D. J. Kenney, K. C. Frisch, S. W. Wong and R. Moore, *J. Polym. Sci., Part B: Polym. Phys.*, 1991, **29**, 1513–1524.
- 27 A. A. Shamsabadi, A. P. Isfahani, S. K. Salestan, A. Rahimpour, B. Ghalei, E. Sivaniah and M. Soroush, *ACS Appl. Mater. Interfaces*, 2020, **12**, 3984–3992.
- 28 G. Huang, A. P. Isfahani, A. Muchtar, K. Sakurai, B. B. Shrestha, D. Qin, D. Yamaguchi, E. Sivaniah and B. Ghalei, *J. Membr. Sci.*, 2018, **565**, 370–379.
- 29 A. P. Isfahani, M. Sadeghi, S. Nilouyal, A. Muchtar, G. Huang, M. Ito, D. Yamaguchi, E. Sivaniah and B. Ghalei, *J. Mater. Chem. A*, 2020, **8**, 9382–9391.
- 30 M. Sadeghi, A. P. Isfahani, A. A. Shamsabadi, S. Favakeh and M. Soroush, *J. Appl. Polym. Sci.*, 2020, **137**, 48704.
- 31 S. H. Yuan, A. P. Isfahani, T. Yamamoto, A. Muchtar, C. Y. Wu, G. Huang, Y. C. You, E. Sivaniah, B. K. Chang and B. Ghalei, *Small Methods*, 2020, 2000021.
- 32 B. Bengtson, C. Feger, W. MacKnight and N. Schneider, *Polymer*, 1985, **26**, 895–900.
- 33 S.-L. Huang and J.-Y. Lai, *J. Membr. Sci.*, 1995, **105**, 137–145.
- 34 C. Li, S. L. Goodman, R. M. Albrecht and S. L. Cooper, *Macromolecules*, 1988, **21**, 2367–2375.
- 35 I. Krakovský, H. Urakawa and K. Kajiwara, *Polymer*, 1997, **38**, 3645–3653.
- 36 I. Fink, B. Eling, E. Pösel and G. A. Luinstra, *J. Polym. Sci., Part A: Polym. Chem.*, 2018, **56**, 1162–1172.



- 37 Z. Cao, Q. Zhou, S. Jie and B.-G. Li, *Ind. Eng. Chem. Res.*, 2016, **55**, 1582–1589.
- 38 Y.-J. Li, N. Nakamura, Y.-F. Wang, M. Kodama and T. Nakaya, *Chem. Mater.*, 1997, **9**, 1570–1577.
- 39 N. Akram, K. M. Zia, R. Sattar, S. Tabassum and M. Saeed, *J. Appl. Polym. Sci.*, 2019, **136**, 47289.
- 40 M. Sadeghi, A. A. Shamsabadi, A. Ronasi, A. P. Isfahani, M. Dinari and M. Soroush, *Chem. Eng. Sci.*, 2018, **192**, 688–698.
- 41 D. P. White, J. C. Anthony and A. O. Oyefeso, *J. Org. Chem.*, 1999, **64**, 7707–7716.
- 42 J. A. Hirsch, *Top. Stereochem.*, 1967, **1**, 199.
- 43 S. Cornell and J. Koenig, *Macromolecules*, 1969, **2**, 540–545.
- 44 S. Poshyachinda, H. G. Edwards and A. F. Johnson, *Polymer*, 1991, **32**, 338–342.
- 45 D. J. Nagle, M. Celina, L. Rintoul and P. M. Fredericks, *Polym. Degrad. Stab.*, 2007, **92**, 1446–1454.
- 46 S.-L. Huang, P.-H. Chang, M.-H. Tsai and H.-C. Chang, *Sep. Purif. Technol.*, 2007, **56**, 63–70.
- 47 R. M. Shankar, T. K. Roy and T. Jana, *J. Appl. Polym. Sci.*, 2009, **114**, 732–741.
- 48 J. A. Brydson, *Plastics materials*, Elsevier, 1999.
- 49 P. Liu, P. McCarren, P. H.-Y. Cheong, T. F. Jamison and K. Houk, *J. Am. Chem. Soc.*, 2010, **132**, 2050–2057.
- 50 O. Glatter and O. Kratky, *Small angle X-ray scattering*, Academic press, 1982.
- 51 M. Shahrooz, M. Sadeghi, R. Bagheri and M. Laghaei, *Macromolecules*, 2016, **49**, 4220–4228.
- 52 G. Strobl and M. Schneider, *J. Polym. Sci., Polym. Phys. Ed.*, 1980, **18**, 1343–1359.
- 53 J. Sewell, *J. Appl. Polym. Sci.*, 1973, **17**, 1741–1747.
- 54 S. Matteucci, Y. Yampolskii, B. D. Freeman and I. Pinnau, *Mater. Sci. Membr. Gas Vap. Sep.*, 2006, **1**, 1–2.
- 55 A. A. Shamsabadi, F. Seidi, E. Salehi, M. Nozari, A. Rahimpour and M. Soroush, *J. Mater. Chem. A*, 2017, **5**, 4011–4025.
- 56 Q. Song, S. Nataraj, M. V. Roussanova, J. C. Tan, D. J. Hughes, W. Li, P. Bourgoïn, M. A. Alam, A. K. Cheetham and S. A. Al-Muhtaseb, *Energy Environ. Sci.*, 2012, **5**, 8359–8369.
- 57 B. Freeman, Y. Yampolskii and I. Pinnau, *Materials science of membranes for gas and vapor separation*, John Wiley & Sons, 2006.

