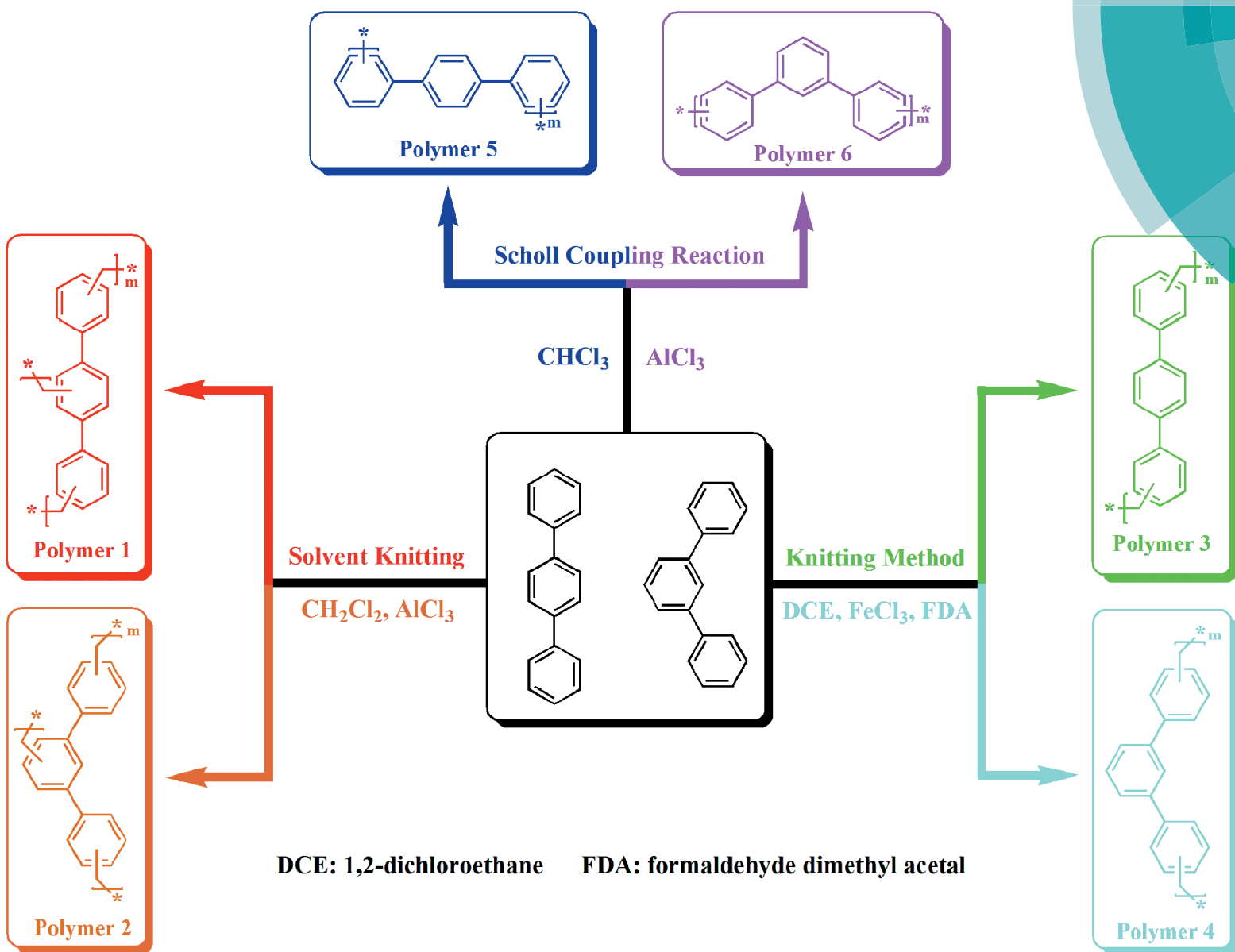


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Effects of synthesis methodology on microporous organic hyper-cross-linked polymers with respect to structural porosity, gas uptake performance and fluorescence properties



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Effects of synthesis methodology on microporous organic hyper-cross-linked polymers with respect to structural porosity, gas uptake performance and fluorescence properties†

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The structural characteristics of hyper-cross-linked polymers (HCPs) make them interesting for a wide variety of applications. To form knitted HCPs various synthesis approaches including solvent knitting, a knitting method using formaldehyde dimethyl acetal (FDA) as an external cross-linker and the Scholl coupling reaction have been used. However, a systematic study of these strategies has not been explored. Here HCPs have been made using facile synthesis methodologies and characterized to reveal resulting Brunauer–Emmett–Teller (BET) surface areas up to 2278 m² g^{−1} and 2064 m² g^{−1} with CO₂ adsorption capacities of 20.89 wt% and 18.95 wt% (at 273.15 K/1.00 bar), respectively. Detailed information is also given with respect to distinctions in textural properties, gas uptake performances and optical properties. More importantly, our comparative studies have revealed the significance and efficiency of solvent knitting when used to fabricate HCP materials with good porosities and gas uptake abilities. Among these approaches, the knitting method with FDA as an external cross-linker is the most time-efficient technique, whereas the Scholl coupling reaction has advantages for the production of fluorescent polymers. Based on the proposed mechanisms, changes in structural features can also be rationally explained.

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Introduction

Porous hyper-cross-linked polymers (HCPs) that can be prepared in a facile manner using the Friedel–Crafts reaction under mild conditions are attracting increasing interest.¹ Due to extensive cross-linking, the resultant porous polymer's polymer chains will not collapse into a dense and nonporous state, while additionally their highly cross-linked nature confers an unusually high thermal stability that is not commonly found in organic polymers.² With the advantages of a large surface area, low cost, relatively low heats of adsorption, excellent chemical robustness (*e.g.* resistance to strong acids and bases),³ they have found increasingly wide applications in gas capture and separation,^{4–7} energy storage,^{7–9} volatile organic compound adsorption,^{10,11} uptake of chemical warfare agents,¹² catalysis,^{13–15} and drug release.¹⁶ These properties

have sparked in recent years an interest in their rational design and synthesis.¹⁷

The development of HCPs dates back to the introduction of hyper-cross-linked polystyrene in the early 1970s.² Since then, much effort has been devoted to its development. For example, Cooper and co-workers showed the carbonization of HCPs with KOH activation can be used to generate highly porous structures with a BET surface area of 4334 m² g^{−1} adsorbing 3.5 mmol g^{−1} CO₂ at 298 K/1.00 bar and 3.6 wt% H₂ at 77.3 K/1.00 bar.¹⁸ Jiang *et al.*, presented a novel and facile strategy for making high surface area porous carbon nanotubes (PCNTs) with outstanding porosities and an excellent electrochemical performance by direct carbonization of 1D hyper-cross-linked polymer nanotubes.¹⁹ Our group has previously developed a cobalt coordinated polymer with a BET surface area of 1360 m² g^{−1}, which efficiently catalyzed the coupling of CO₂ with various substituted epoxides at room temperature and atmospheric pressure.²⁰ Currently many interesting strategies are being pursued to obtain high surface area polymers with tailored porosities. Building blocks, such as aromatic hydrocarbons,^{21–23} polycyclic aromatic hydrocarbons,^{19,21,24} and heterocyclic compounds,^{25,26} are used to synthesize HCPs. Among these materials a microporous organic polymer derived from *N*4,*N*4'-di-1-naphthalenyl-*N*4,*N*4'-di-2-naphthalenyl-[1,1'-biphenyl]-4,4'-diamine has a BET surface area of 1717 m² g^{−1}

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and a CO₂ adsorption of 18.85 wt% at 273.15 K/1.00 bar.²⁷ Further attempts to optimize structural properties, solvents,^{21–23,28,29} external cross-linkers,^{22,28} and catalysts,^{21,23,24,28,29} have been made, *e.g.* a fluoranthene-based polymer synthesized from anhydrous AlCl₃ in dichloromethane showed a CO₂ adsorption of 24.79 wt% at 273.15 K/1.00 bar under optimized capture conditions.²⁴

To make HCPs with high surface areas and controlled porosities, many synthesis strategies such as the 'knitting method' with formaldehyde dimethyl acetal (FDA) as an external cross-linker,²¹ the Scholl coupling reaction²² and the solvent knitting method²³ have been used. In 2011, our group put forward the knitting method, using FDA as an external cross-linker and a Lewis acid (anhydrous FeCl₃) catalyst in 1,2-dichloroethane.²¹ The resulting HCP materials had well formed microporous structures with various enclosed functional groups. This strategy allows for the use of a variety of 'building blocks' under facile reaction conditions and inexpensive reagents.²¹ Consequently, this strategy has been used to synthesize diverse polymer networks. In 2014, our group incorporated functional groups in microporous organic polymers (MOPs), in a protocol based on the Scholl coupling reaction catalyzed by anhydrous AlCl₃ in the presence of chloroform.²² Recently in 2017, our group proposed the solvent knitting method to make 2D layered HCPs by ultrasonic-assisted solvent exfoliation.²³ These HCPs are usually made using Friedel–Crafts alkylation with the use of dichloroalkane as an economical solvent, stable electrophilic reagent, and external cross-linker. Such materials' porous structures and surface areas can be controlled by altering factors such as the molar ratio of the catalyst (AlCl₃), the sizes of the building blocks, and the solvent's chain length. Because this method is simple, one-step and cost-effective, it offers opportunities for knitting polymers with high surface areas and gas adsorption capacities. This knitting strategy is considered a low-cost, scalable and efficient polymerization technique for creating HCP materials for real applications.²³

These distinct approaches possess the following distinguishing features. The knitting strategy with FDA as an external cross-linker is conducted in atmospheric air. When catalyzed by FeCl₃ and traces of atmospheric water a greater driving force for the reaction is created. Based on the Friedel–Crafts reaction, this method with FDA cross-linking and building block molecules usually gives a moderate reaction rate.²¹ The Scholl coupling reaction and the solvent knitting method are performed under a nitrogen atmosphere. In the absence of water AlCl₃ will not be hydrolysed, which will favor the reactions. Scholl coupling, with the elimination of a C–C bond, has the fastest reaction rate without any cross-linker.²² In contrast, the solvent knitting method with an improved Friedel–Crafts alkylation and a low activity cross-linker (CH₂Cl₂), has the slowest reaction rate, which is helpful to enhance the free-packing of building blocks and knitting polymers with large surface areas and abundant micropores.²³

The existing methodologies result in HCPs with abundant micropores and high surface areas. More significantly, great

interest has been aroused to explore these strategies to gain insights into the design and development of HCP materials. This work focuses on the generation of microporous networks using the previously mentioned strategies that employ aromatic hydrocarbons such as 1,4-diphenylbenzene and 1,3-diphenylbenzene to fabricate novel microporous HCPs. It is worth noting that the prepared polymers have BET surface areas up to 2278 m² g^{−1} and 2064 m² g^{−1} and CO₂ adsorption capacities of 20.89 wt% and 18.95 wt% at 273.15 K/1.00 bar, which have rarely been reported. Furthermore, with respect to porous structures, gas uptakes and the fluorescence of the resultant polymers, this work has demonstrated that: (I) the solvent knitting method has a higher efficiency for knitting polymers giving better porous structures and gas uptake performances; (II) the knitting method with FDA as an external cross-linker is the most time efficient; (III) the Scholl coupling reaction can be used to prepare fluorescent polymers. We finally offer a detailed insight into the reaction mechanisms that lead to these distinctions.

Experimental section

Materials

1,4-Diphenylbenzene (1,4-DPB), 1,3-diphenylbenzene (1,3-DPB) and formaldehyde dimethyl acetal (FDA) were purchased from Aladdin Chemical Reagent Corp. (Shanghai, China) and used as received. Dichloromethane (CH₂Cl₂), 1,2-dichloroethane (DCE), chloroform (CHCl₃), hydrochloride (HCl), methanol, ethyl acetate, dimethyl sulfoxide (DMSO), *N,N*-dimethylformamide (DMF), *N,N*-diethylformamide (DEF), ethanol, anhydrous ferric chloride (FeCl₃) and anhydrous aluminum chloride (AlCl₃) were acquired from the Sinopharm Chemical Reagent Ltd Co. (Shanghai, China) and used as received. Unless stated otherwise, all the solvents and chemicals were of analytical grade and obtained from local suppliers and used without further purification.

Preparation of polymers

Synthesis of polymer **1** by the solvent knitting method. Under a nitrogen atmosphere, 1,4-DPB (2 mmol, 0.460 g) was dispersed in CH₂Cl₂ (8 mL) for 30 minutes, anhydrous AlCl₃ (24 mmol, 3.204 g) was immediately added to the solution which was then allowed to react at 20 °C for 4 h, 30 °C for 8 h, 40 °C for 12 h, 60 °C for 12 h, and 80 °C for 24 h under vigorous stirring. After cooling to room temperature, the solid product was quenched using 20 mL of HCl–H₂O (*v/v* = 2 : 1), washed several times with deionized water and ethanol, further purified by Soxhlet extraction with ethanol for 48 h, and finally dried in a vacuum oven at 70 °C for 24 h. The polymer was obtained as a black solid. Yield: 140%.

Polymer **2** was prepared in a manner similar to polymer **1** by treating 1,3-DPB (2 mmol, 0.460 g) with anhydrous AlCl₃ (24 mmol, 3.204 g) in CH₂Cl₂ (8 mL). The polymer material was again obtained as a black solid. Yield: 135%.

Polymer 3 was synthesized by the knitting method with FDA as an external cross-linker. Under an air atmosphere, anhydrous FeCl_3 (18 mmol, 2.925 g) was added to a solution of 1,4-DPB (2 mmol, 0.460 g) and FDA (18 mmol, 1.370 g) in DCE (8 mL). The mixture was heated at 45 °C for 5 h and then 80 °C for 19 h with vigorous stirring. After cooling to room temperature, the solid product was washed with methanol several times, further purified by Soxhlet extraction with methanol for 48 h, and finally dried in a vacuum oven at 70 °C for 24 h. The polymer material was obtained as a brown solid. Yield: 137%.

Polymer 4 was also synthesized by the knitting method as described above for polymer 3, using 1,3-DPB (2 mmol, 0.460 g), anhydrous FeCl_3 (12 mmol, 1.950 g) and FDA (12 mmol, 0.913 g) in DCE (8 mL). The polymer was again obtained as a brown solid. Yield: 140%.

Polymer 5 was synthesized by the Scholl coupling reaction method. Under a nitrogen atmosphere, anhydrous AlCl_3 (24 mmol, 3.204 g) was added to a solution of 1,4-DPB (2 mmol, 0.460 g) in CHCl_3 (8 mL) with vigorous stirring at 58 °C, the reaction proceeded for 48 h. The solid product was washed once with ethanol, twice with $\text{HCl-H}_2\text{O}$ (v/v = 2 : 1), and then three times with ethanol; followed by ethanolic Soxhlet extraction with ethanol for 48 h, prior to drying in a vacuum oven at 70 °C for 24 h. The polymer was obtained as a black solid. Yield: 97%.

Polymer 6 was likewise prepared by the Scholl coupling reaction as in the above procedure for polymer 5 using 1,3-DPB (2 mmol, 0.460 g) and anhydrous AlCl_3 (24 mmol, 3.204 g) in CHCl_3 (8 mL). The polymer was obtained as a brown solid. Yield: 98%.

Characterization

Fourier transform infrared (FT-IR) spectra were recorded using a Bruker VERTEX 70 FT-IR spectrometer (wavenumber range 4000–500 cm^{-1}) at room temperature and atmospheric pressure. Solid state ^{13}C cross-polarization/magic-angle spinning nuclear magnetic resonance (^{13}C CP/MAS NMR) spectra were recorded on a WB 400 MHz Bruker Avance II spectrometer and collected with a spinning rate of 20 kHz by using a 2.5 mm double-resonance MAS probe. The field-emission scanning electron microscopy (FE-SEM) images were recorded on a FEI Sirion 200 field-emission scanning electron microscope operating at 10 kV. All the samples were dried in a vacuum oven at 70 °C for 24 h and then sputter-coated with platinum before measurement. The transmission electron microscopy (TEM) images were recorded on a Tecnai G2 F30 microscope (FEI Corp. Holland) operating at 200 kV. Thermogravimetric analysis (TGA) was performed from room temperature to 850 °C, using a PerkinElmer Instrument Pyris1 TGA with a heating rate of 10 °C min^{-1} under a nitrogen atmosphere. The solid state fluorescence spectra were recorded using an FP-6500 fluorescence spectrometer (Jasco, JPN). The absolute solid state fluorescence quantum yield of polymers was tested by using an Absolute PL Quantum Yield Spectrometer F4600 (Hitachi, JPN) at room temperature and atmospheric pressure. All the samples were degassed at 120 °C

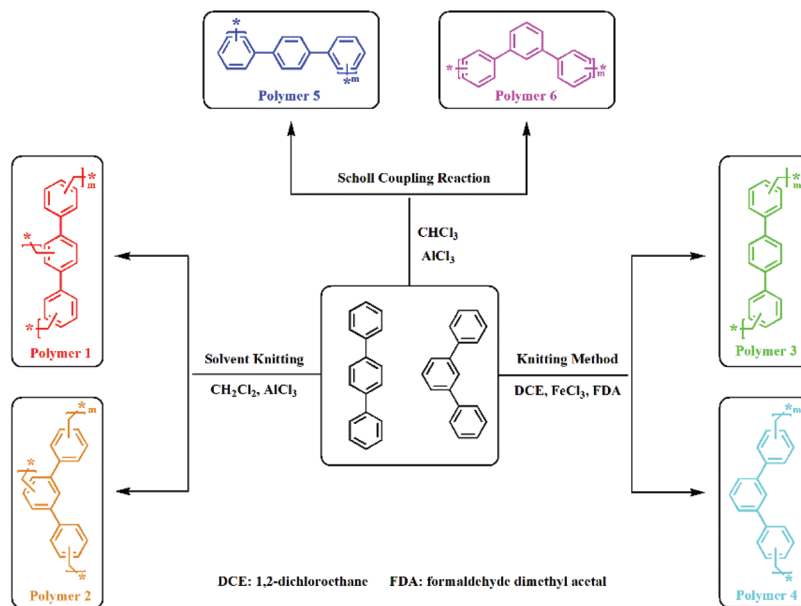
for a minimum of 8 h under vacuum of 10^{-5} bar prior to being measured. N_2 isotherms were collected on a Micromeritics ASAP 2020 M analyzer at 77.3 K. Gas (CO_2 , N_2 and CH_4) adsorption and desorption isotherms were measured on a Micromeritics ASAP 2020 M analyzer at 273.15 and 298.15 K. Pore size distribution was calculated from N_2 adsorption isotherms using a Tarazona nonlocal density functional theory (NLDFT) model assuming a slit pore geometry. Total pore volumes were calculated from nitrogen sorption isotherms at relative pressure $P/P_0 = 0.995$.

Results and discussion

In this work, HCP materials were prepared by three synthesis methods to gain better insights into the distinctions in structural porosities, gas uptakes and fluorescence properties of microporous organic HCPs. Of note, the Scholl coupling polymerization reaction proceeds²² without the addition of an external cross-linker. Therefore, the monomers react directly with a fast reaction rate. While the knitting method, based on the Friedel–Crafts reaction, has FDA as an external cross-linker,²¹ and in contrast with the Scholl coupling reaction, it shows a slower reaction rate. The solvent knitting method employs an improved Friedel–Crafts alkylation,²³ in which dichloromethane plays the role of both an external cross-linker and solvent. Due to the lower reaction activity of dichloromethane compared to FDA, it has the slowest reaction rate, which is advantageous to form networks with layered structures. Since the reaction mechanism is closely associated with the materials' structure, the mechanisms associated with these methodologies vary, which may explain the diversity of the resultant polymer networks.

Scheme 1 shows the 'building blocks' and proposed pathways leading to the formation of HCPs by the solvent knitting method, the knitting method with FDA as an external cross-linker and the Scholl coupling reaction. The building blocks are soluble in organic solvents such as ethyl acetate, dimethyl sulfoxide, *N,N*-dimethylformamide and *N,N*-diethylformamide, while the obtained polymers are insoluble in these solvents. It has been demonstrated that the polymerization normally leads to a decrease in polymer solubility.

FT-IR spectra were recorded to confirm the starting materials and polymers' structural features. Taking 1,3-DPB-based polymers as an example. The broad bands at 3600–3100 cm^{-1} in polymers 2 and 4 are largely attributed to adsorbed water in the networks created. The bands at 3100–3000 cm^{-1} are the C–H stretching vibrations of the benzene ring, while the bands at ~ 1508 cm^{-1} , 1477 cm^{-1} and 1437 cm^{-1} are assigned to C=C stretching vibrations of the benzene ring, which is found in both polymeric networks 2, 4, 6 and their 'building blocks' (Fig. 1b). The absorption bands at 2910 cm^{-1} and 2829 cm^{-1} are the C–H stretching vibrations of the methylene bridges, seen in polymers 2 and 4 (Fig. 1b), and the FT-IR spectra of 1,4-DPB and 1,4-DPB-based polymer materials are shown in Fig. 1a.



Scheme 1 Proposed synthesis pathways to microporous organic HCP networks, together with the building blocks' chemical structures.

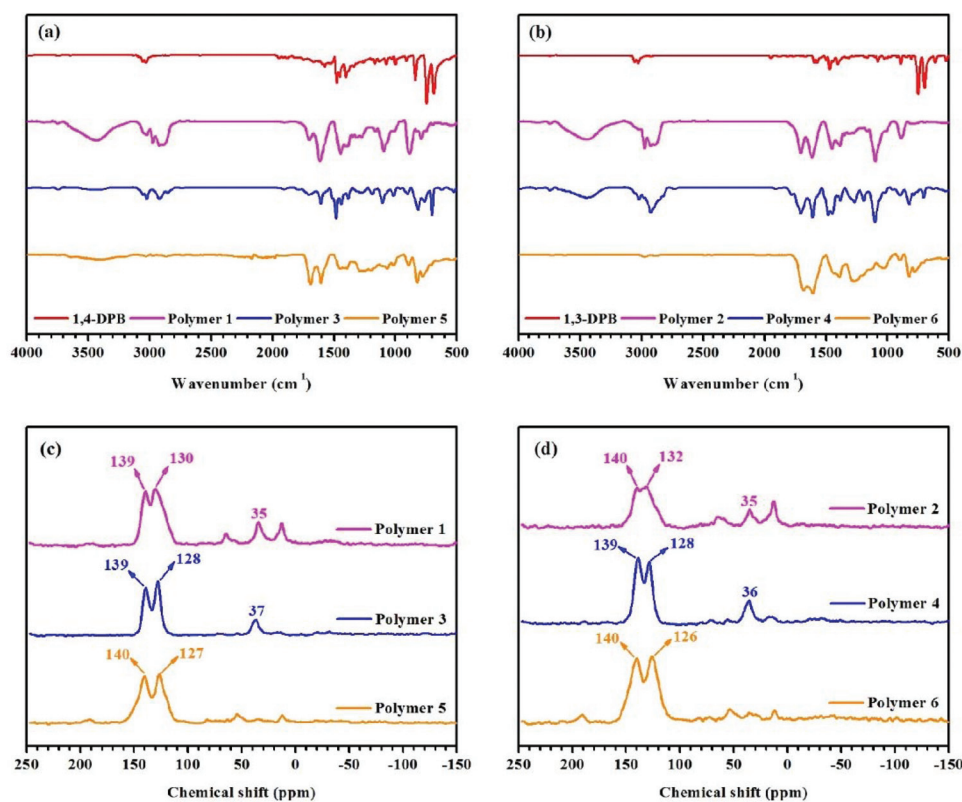


Fig. 1 FT-IR spectrum of (a) 1,4-DPB and 1,4-DPB-based polymer materials, and (b) 1,3-DPB and 1,3-DPB-based polymer materials; solid state cross-polarization (CP) ^{13}C MAS natural abundance NMR spectrum of (c) 1,4-DPB-based polymer materials, and (d) 1,3-DPB-based polymer materials.

Following the FT-IR study, solid state ^{13}C CP/MAS NMR spectroscopy was used to investigate the structural characteristics of the polymers. Well-resolved signals are seen in

Fig. 1d. For example, the singlets at ~ 137 ppm belong to the substituted aromatic carbon and the singlets near to 130 ppm are attributed to the unsubstituted aromatic carbon,¹⁵ while

the resonance at ~ 35 ppm in polymer 2 is ascribed to the methylene linkers – derived from CH_2Cl_2 (solvent and an external cross-linker).^{23,27} Meanwhile, there are also doublets in the range 70–10 ppm in polymer 2, attributable to methylene bridges (resulting from hyper-cross-linking) and residual dichloromethane. As previously stated, dichloromethane acts as an external cross-linker and solvent in the solvent knitting strategy. Similarly, in polymer 4 the signal at ~ 36 ppm is assigned to the methylene linkers, derived from $\text{CH}_3\text{OCH}_2\text{OCH}_3$ (an external cross-linker).²⁵ Moreover, the resonances at ~ 126 and 140 ppm in polymer 6 are respectively due to the non-coupled and coupled aromatic carbons.²² A detailed graphical explanation of the polymers derived from 1,4-DPB is given in Fig. 1c.

Taking together the FT-IR and solid state ^{13}C CP/MAS NMR spectra we have shown the successful generation of HCPs by precursor cross-linking using three different techniques.

The surface morphologies and porous nature of the polymers were studied by FE-SEM and TEM. Fig. S2–S4 (ESI†) show FE-SEM images of these networks, which have the irregular blocks aggregated together. Fig. S1 (ESI†) shows the morphologies of their parent monomers, which are amorphous in nature. By comparing the morphologies of polymers and their building blocks we can deduce that these synthesis methodologies did not distinctly influence the morphology of polymer materials. Fig. S5–S7 (ESI†) show TEM images of the polymers showing the amorphous and porous structures of the HCPs obtained by using different strategies.

TGA analysis (Fig. 2) was used to record the thermal stability of all the HCPs. As can be seen, polymer materials from polymer 1 to polymer 4 are stable up to approximately 400°C under a nitrogen atmosphere, which is a little higher than those of polymers 5 and 6. Thus it is evident that, polymer networks from 1 to 4 prepared using the knitting strategy tend to be more stable than polymers 5 and 6 synthesized by the Scholl coupling reaction. This is primarily attributed to the different degrees of polymer cross-linking resulting from the use of differing synthesis methodologies. Furthermore, their obvious thermal stability (with weight loss only occurring with temperatures above 400°C) is ascribed to polymeric network degradation.

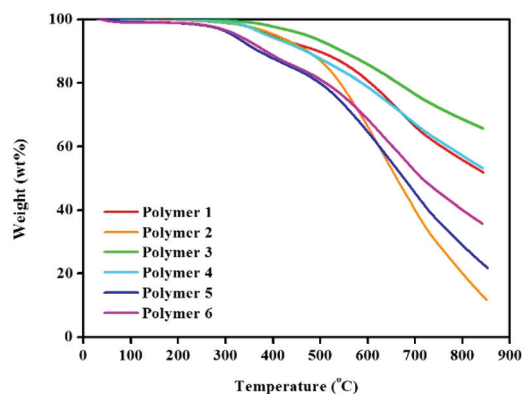


Fig. 2 TGA analysis of materials from polymer 1 to polymer 6.

The porous nature of these hyper-cross-linked polymers was shown by nitrogen adsorption and desorption analysis at 77.3 K and 1.00 bar. Fig. 3a shows the high nitrogen gas uptake at a low relative pressure ($P/P_0 < 0.001$) thereby highlighting the microporous structure of the resultant polymers. The observed hysteresis loops in the middle of all isotherms imply the mesopores. No obvious rise in the nitrogen adsorption isotherms at high relative pressures ($P/P_0 > 0.9$) can be seen, which suggests that macropores are nearly absent in these polymers. The pore size distribution of polymers was estimated using N_2 adsorption isotherms employing the NLDFT model. As seen from Fig. 3b, these polymers with different textures have hierarchical pore size distributions. Besides the peaks (2–10 nm) in the mesoporous region, the pore size is mainly located within 2 nm in the microporous section, including the ultramicroporous region, whose diameter is no more than 0.7 nm. Table 1 lists the porosity parameters of all HCP samples. For clarity, 1,3-DPB-based polymers are discussed in detail below. The BET surface areas of the polymeric networks varied from $1092\text{ m}^2\text{ g}^{-1}$ to $2278\text{ m}^2\text{ g}^{-1}$, the Langmuir surface areas ranged from $1335\text{ m}^2\text{ g}^{-1}$ to $2797\text{ m}^2\text{ g}^{-1}$, the micropore areas were from $692\text{ m}^2\text{ g}^{-1}$ to $1443\text{ m}^2\text{ g}^{-1}$, the total pore volumes ranged between $0.58\text{ cm}^3\text{ g}^{-1}$ and $1.08\text{ cm}^3\text{ g}^{-1}$, with micropore volumes between $0.28\text{ cm}^3\text{ g}^{-1}$ to $0.57\text{ cm}^3\text{ g}^{-1}$. It was found that when compared to polymers produced by the knitting method, with FDA as an external cross-linker and the Scholl coupling reaction, polymers prepared by the solvent knitting method have the highest BET surface areas, Langmuir surface areas, micropore areas, pore volumes and micropore volumes. Similarly, 1,4-DPB-based polymer networks including polymers 1, 3, and 5 also follow the same trend. This is probably due to the polymerization strategies. As already mentioned, the solvent knitting method proceeds without the addition of an external cross-linker. In this strategy, dichloromethane plays the role of both the solvent and cross-linking agent. The activity of dichloromethane is very low, thus it has the slowest reaction rate, thereby enhancing the cross-linking degree, while improving the free-packing of building blocks with the resulting polymers tending to produce abundant micropores and a much higher surface area (Table 1).

Porous materials with ultramicroporous structures are often used for gas sorption and storage *e.g.* in CO_2 sorption. Microporous organic HCPs were examined in this study for their potential for CO_2 capture at 273.15 and 298.15 K up to a pressure of 1.13 bar (Fig. 4 and Table 1). For each sample, it can be seen, the CO_2 adsorption capacity continuously increased with the feed pressure, however saturation is not seen in experimental pressures ranging up to 1.13 bar, clearly indicating that a larger CO_2 uptake can be expected with additional elevations in CO_2 pressure. As a result, it was demonstrated that polymer 2 with a BET surface area of $2278\text{ m}^2\text{ g}^{-1}$ and a Langmuir surface area of $2797\text{ m}^2\text{ g}^{-1}$ had a CO_2 adsorption of 20.89 wt% at 273.15 K/1.00 bar. This is comparable to many reported materials such as the carbazolic porous organic framework Cz-POF-3 (4.77 mmol g^{-1} , $S_{\text{BET}} =$

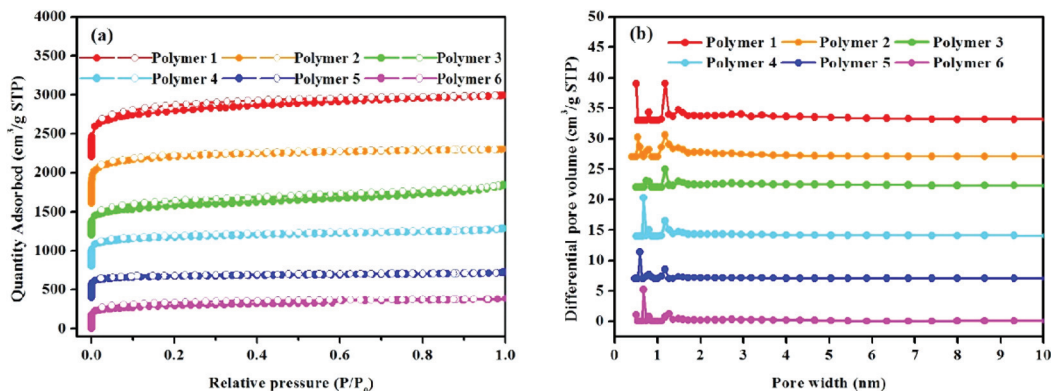


Fig. 3 (a) Nitrogen adsorption and desorption isotherms at 77.3 K and 1.00 bar, and (b) pore distribution curves of the pore size calculated using nonlocal density functional theory (slit pore models, differential pore volumes, pore width). Filled symbols represent adsorption and open symbols represent desorption.

Table 1 Porous properties, CO₂ uptake capacities and isosteric heat of adsorption values of polymer samples

Samples	S_{BET}^a (m ² g ⁻¹)	S_{L}^b (m ² g ⁻¹)	MA ^c (m ² g ⁻¹)	PV ^d (cm ³ g ⁻¹)	MPV ^e (cm ³ g ⁻¹)	CO ₂ uptake ^f (wt%)	CO ₂ uptake ^g (wt%)	Q_{st}^h (kJ mol ⁻¹)
Polymer 1	2064	2780	921	1.23	0.41	18.95	9.54	26.30–26.04
Polymer 2	2278	2797	1443	1.08	0.57	20.89	11.83	21.97–21.60
Polymer 3	1315	1771	533	0.99	0.24	13.70	7.87	27.22–24.90
Polymer 4	1421	1726	935	0.75	0.37	16.47	9.26	26.20–24.95
Polymer 5	1040	1227	778	0.49	0.30	15.04	8.82	27.34–25.97
Polymer 6	1092	1335	692	0.58	0.28	15.11	9.00	28.01–26.68

^a Apparent surface areas calculated from nitrogen adsorption isotherms at 77.3 K using the BET equation. ^b Surface areas calculated from nitrogen adsorption isotherms at 77.3 K using the Langmuir equation. ^c *t*-Plot micropore areas. ^d Pore volumes calculated from the nitrogen isotherm at $P/P_0 = 0.995$ and 77.3 K. ^e *t*-Plot micropore volumes calculated from the nitrogen isotherm at $P/P_0 = 0.050$. ^f CO₂ uptake determined volumetrically using a Micromeritics ASAP 2020 M analyzer at 1.00 bar and 273.15 K. ^g CO₂ uptake determined volumetrically using a Micromeritics ASAP 2020 M analyzer at 1.00 bar and 298.15 K. ^h Isosteric heat of adsorption of polymer materials determined volumetrically using a Micromeritics ASAP 2020 M analyzer at 273.15 and 298.15 K.

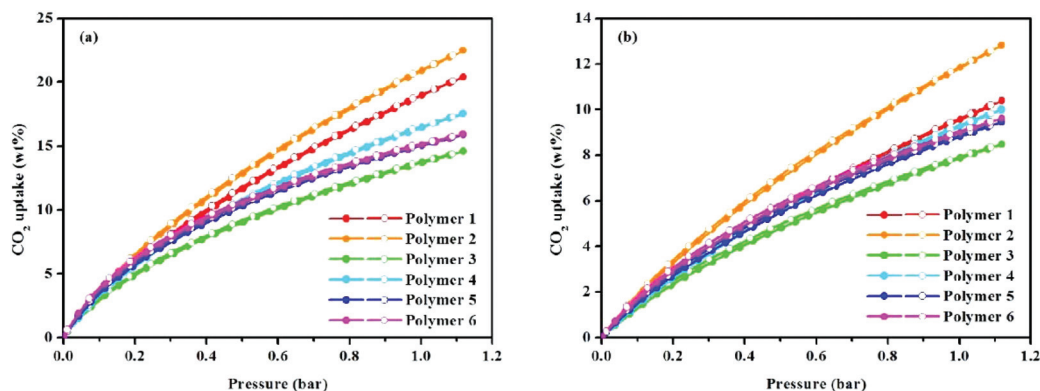


Fig. 4 (a) Volumetric CO₂ adsorption isotherms and desorption isotherms up to 1.13 bar at 273.15 K, and (b) volumetric CO₂ adsorption isotherms and desorption isotherms up to 1.13 bar at 298.15 K.

1927 m² g⁻¹ at 273.15 K/1.00 bar),³⁰ CP-CMP5 (20.11 wt%, $S_{\text{BET}} = 2241$ m² g⁻¹ at 273.15 K/1.13 bar),³¹ CP-CMP6 (19.14 wt%, $S_{\text{BET}} = 1764$ m² g⁻¹ at 273.15 K/1.13 bar),³¹ CPOP-1 (21.2 wt%, $S_{\text{BET}} = 2220$ m² g⁻¹ at 273.15 K/1.00 bar),³² and PPN-CH₂DETA (15.8 wt%, $S_{\text{BET}} = 555$ m² g⁻¹ at 298.15 K/1.00

bar).³³ In addition, the CO₂ uptake is much higher than some types of porous materials with ultrahigh surface areas or high total pore volumes under similar conditions, e.g. COF-102 (6.8 wt%, $S_{\text{BET}} = 3622$ m² g⁻¹),³⁴ PAF-1 (9.1 wt%, $S_{\text{BET}} = 4077$ m² g⁻¹),³⁵ CMPN-3 (2.36 cm³ g⁻¹, 3.82 wt%),³⁶ and

COP-10 (about 9.1 wt%, $S_{\text{BET}} = 3337 \text{ m}^2 \text{ g}^{-1}$ at 298.15 K/1.00 bar).³⁷ The CO_2 adsorption capacity of polymer 2 is much higher than many HCP materials such as 1,3,5-triphenylbenzene-based HCP (15.9 wt%, $S_{\text{BET}} = 1059 \text{ m}^2 \text{ g}^{-1}$),²¹ tetraphenylmethane-based HCP (1.66 mmol g^{-1} , $S_{\text{BET}} = 1679 \text{ m}^2 \text{ g}^{-1}$)³⁸ and binaphthol-based HCP (17.4 wt%, $S_{\text{BET}} = 1015 \text{ m}^2 \text{ g}^{-1}$).¹⁹ Moreover, the CO_2 uptake ability of polymer 2 also outperforms commercially available BPL carbon (2.09 mmol g^{-1} , at 273 K/1 bar),²⁶ POSS-based HPPs (0.62–1.04 mmol g^{-1}),^{39,40} and organic–inorganic hybrid porous Fc-HPP (4.87 wt%, $S_{\text{BET}} = 1015 \text{ m}^2 \text{ g}^{-1}$, at 273 K/101 kPa),⁴¹ thus highlighting a comparative CO_2 uptake capacity. From the gas sorption analysis, it is evident that, the solvent knitting method enhances CO_2 capture to a greater extent than does either the knitting method with FDA (an external cross-linker) or the Scholl coupling reaction. Also of note is that the CO_2 uptake of polymer 3 is the lowest among all the samples, even though its surface area is higher than that of either polymer 4 or 5. This is primarily due to the polymer's pore size distribution. Because a previous study has shown that, in addition to the surface area, the pore size and pore size distribution are also closely related to the CO_2 adsorption capacity of microporous organic polymers.⁴² From Fig. 3b, it can be seen, polymer 3 with a pore size $>0.7 \text{ nm}$ lacks ultramicropores, while polymers with a pore size $<0.7 \text{ nm}$ possess ultramicropores. Since a pore size $<1 \text{ nm}$ preferentially adsorbs CO_2 molecules,⁴³ especially the ultramicropore, whose diameter is comparable to the kinetic diameter of CO_2 and is therefore prone to increasing the interactions between the CO_2 molecules and the pore walls,⁴⁴ it is not hard to understand, that although the surface area of polymer 3 is moderate its CO_2 uptake performance is the lowest. To better understand the binding affinity of polymers towards CO_2 , the isosteric heat (Q_{st}) of polymers was calculated by fitting CO_2 adsorption isotherms obtained at 273.15 and 298.15 K using the Clausius–Clapeyron equation,⁴⁵ which was found to be in the ranges of 26.30–26.04, 21.97–21.60, 27.22–24.90, 26.20–24.95, 27.34–25.97, and 28.01–26.68 kJ mol^{-1} for polymer 1 to polymer 6 (Fig. S8,† Table 1). These results make it clear that there is a normal physical adsorption interaction between the CO_2 molecules and polymer networks. While in comparison with polymers having a higher isosteric heat of adsorption such as Py-1 (36 kJ mol^{-1}),²⁶ PI-1 (34 kJ mol^{-1}),⁴⁶ and PECONF-4 (34 kJ mol^{-1}),⁴⁷ the moderate Q_{st} values of the prepared polymers are probably due to the lack of heteroatoms *e.g.* nitrogen atoms, which are known to improve the binding affinity towards CO_2 molecules by promoting dipole–quadrupole interactions.²⁶

In addition to the studies on CO_2 uptake capacity and isosteric heat of adsorption, gas adsorption selectivity including CO_2/N_2 and CO_2/CH_4 is another significant parameter relevant to polymer performance. Single-component physisorption isotherms for CO_2 , N_2 and CH_4 were measured by volumetric methods at 273.15 and 298.15 K, see Fig. 5. Gas adsorption values are listed in Table 2. The gas adsorption selectivity of the polymers towards CO_2 over N_2 and CH_4 was calculated using the initial slope ratios from the Henry's law constants of

the pure gas adsorption isotherms at the same temperature and a low pressure coverage of less than 0.3 bar. From Fig. S9–S14 (ESI†) and Table 2, the CO_2/N_2 selectivities at 273.15 K are ~ 16.7 , 16.9, 26.1, 17.2, 21.6, and 19.9 for polymers 1 to 6, respectively. The experimental results also prove that polymer 3 has the highest CO_2/N_2 selectivity *i.e.* 26.1 at 273.15 K. As stated, the incorporation of heteroatoms into the polymer's framework is an effective way to enhance the gas sorption selectivity of the polymer networks. The fabricated polymers mainly comprise carbon, hydrogen and a lack of heteroatoms. Though much lower than KFUPM-1 (141),⁴⁸ PVAm/PEI-*g*-ZIF-8/mPSf (79.9),⁴⁹ TrzPOP-1 (61),⁵⁰ bare Pebax membranes (57),⁵¹ and PMOP (47.1),⁵² the CO_2/N_2 value of polymer 3 at 273.15 K is comparable to those of heteroatom-containing scaffolds such as CPOP-1 (25),³² pTOC (27),⁵³ Network-2 (26),²⁹ Network-4 (27),²⁹ NOP-3 (27.1),⁵⁴ PI-ADNT (25),⁵⁵ APOP-1-OH (26.0),⁵⁶ and NPTN-3 (25)⁵⁷ at 273 K, 1,1-bi-2-naphthol (26),⁵⁸ Glc-3 (27),⁵⁹ and Ara-1 (26)⁵⁹ at 298 K, and better than the porous polymer frameworks (PPFs) (14.5–20.4),⁶⁰ NPTN-2 (22),⁵⁷ JUC-141 (21.62),⁶¹ and CTFs (16.1–24.1)⁶² at 273 K, hydroxyl group functionalized polymers (16),⁵⁸ Gal-1 (21),⁵⁹ Glc-2 (18),⁵⁹ Glc-1 (18),⁵⁹ and CTF-20-400 (19)⁶³ at 298 K. As shown in Fig. S9–S14 (ESI†) and Table 2, the CO_2/CH_4 selectivities at 273.15 K are ~ 4.2 , 4.1, 6.3, 4.0, 4.8, and 4.5 for materials from polymers 1 to 6 respectively. It is apparent that polymer 3 has the best CO_2/CH_4 selectivity at 273.15 K. Although lower than those of materials such as IITKGP-8 (17.7),⁶⁴ MaSOPd₅₀ (15),⁶⁵ 6FDA-durene (12.45 ± 0.29),⁶⁶ and $[\text{Co}_2(5,4\text{-PMIA})_2(\text{TPOM})_{0.5}]\cdot\text{x solvent}$ (9),⁶⁷ the CO_2/CH_4 selectivity value of polymer 3 at 273.15 K is comparable to APOP-1-F (6.4),⁵⁶ ALP-1 (6.0),⁶⁸ CP-CMP-7 (6.0),³¹ and APOP-1-ONa (6.7)⁵⁶ at 273 K, APOP-1-ONa (6.6),⁵⁶ APOP-1-F (6.3),⁵⁶ and ALP-3 (6.0)⁶⁸ at 298 K, and exceeds TBMIDM (4.5),⁶⁹ Azo-CMP2 (5.3),⁷⁰ P-2 (3.0),⁷¹ FCDTPA-K-900 (3.4),²⁸ TBMIDE (4.3),⁶⁹ BO-CMP-2 (4.9),⁷² and OBO-CMP-1 (4.7)⁷² at 273 K, P-1 (4.0),⁷¹ and APOP-4 (4.7),⁵⁶ P-2 (3.0),⁷¹ and TPI-3@IC (2.0)⁷³ at 298 K. To understand the relationship between the structure and performance, following the gas sorption selectivity of polymers at 273.15 K, the CO_2/N_2 and CO_2/CH_4 selectivities of all polymers at 298.15 K were assessed. As seen in Fig. S15–S20 (ESI†) and Table 2, the CO_2/N_2 and CO_2/CH_4 selectivity values are in the ranges of 12.0–22.1 and 3.2–5.6 from polymers 1 to 6. It is apparent that polymer 3 has a CO_2/N_2 selectivity of 22.1 and a CO_2/CH_4 selectivity of 5.6, values that are comparable to other MOPs.

From the above gas sorption polymer data, the selectivity of CO_2/N_2 is much higher than that of CO_2/CH_4 at both 273.15 and 298.15 K. There are many influencing factors such as the pore size and physicochemical nature of polymers, the kinetic diameter of the gas molecules, and differences in thermodynamic behaviour of the pairs of gases.⁷⁴ To have a convincing explanation and better understanding the selective adsorption behavior and the critical temperature (T_c), which is a thermodynamic factor, should be taken into consideration. Specifically, the critical temperatures of common gas molecules like CO_2 , N_2 and CH_4 are 304.2, 190.6 and 126.2 K, which

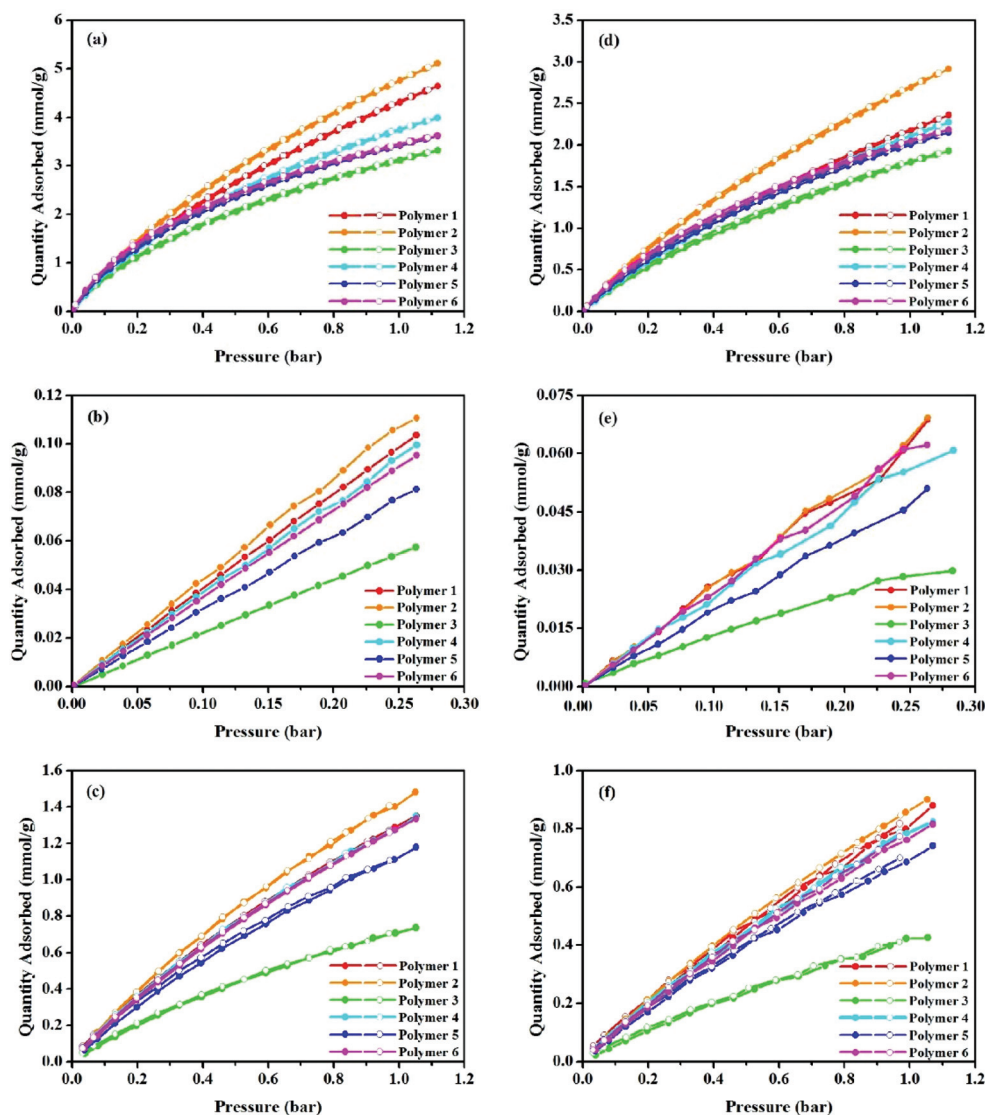


Fig. 5 Adsorption isotherms of CO₂ (a), N₂ (b), CH₄ (c) gases at 273.15 K, and CO₂ (d), N₂ (e), CH₄ (f) gases at 298.15 K for polymer 1 to polymer 6. Filled symbols represent adsorption and open symbols represent desorption.

Table 2 N₂ and CH₄ uptakes, and gas adsorption selectivity of polymer samples

Samples	N ₂ uptake ^a (wt%)	N ₂ uptake ^b (wt%)	CH ₄ uptake ^c (wt%)	CH ₄ uptake ^d (wt%)	CO ₂ /N ₂ ^e	CO ₂ /N ₂ ^f	CO ₂ /CH ₄ ^g	CO ₂ /CH ₄ ^h
Polymer 1	0.28	0.18	2.08	1.30	16.7	12.0	4.2	3.2
Polymer 2	0.30	0.18	2.27	1.38	16.9	15.2	4.1	3.7
Polymer 3	0.15	0.08	1.14	0.68	26.1	22.1	6.3	5.6
Polymer 4	0.27	0.16	2.06	1.26	17.2	12.8	4.0	3.4
Polymer 5	0.22	0.13	1.80	1.11	21.6	16.7	4.8	3.9
Polymer 6	0.25	0.17	2.05	1.23	19.9	13.7	4.5	3.9

^a N₂ uptake determined volumetrically using a Micromeritics ASAP 2020 M analyzer at 273.15 K and 0.25 bar. ^b N₂ uptake determined volumetrically using a Micromeritics ASAP 2020 M analyzer at 298.15 K and 0.25 bar. ^c CH₄ uptake determined volumetrically using a Micromeritics ASAP 2020 M analyzer at 273.15 K and 1.00 bar. ^d CH₄ uptake determined volumetrically using a Micromeritics ASAP 2020 M analyzer at 298.15 K and 1.00 bar. ^e Adsorption selectivity of CO₂ over N₂ calculated by employing the initial slope ratio from the Henry's law constants at 273.15 K and a low pressure coverage of less than 0.3 bar. ^f Adsorption selectivity of CO₂ over N₂ calculated by employing the initial slope ratio from the Henry's law constants at 298.15 K and a low pressure coverage of less than 0.3 bar. ^g Adsorption selectivity of CO₂ over CH₄ calculated by employing the initial slope ratio from the Henry's law constants at 273.15 K and a low pressure coverage of less than 0.3 bar. ^h Adsorption selectivity of CO₂ over CH₄ calculated by employing the initial slope ratio from the Henry's law constants at 298.15 K and a low pressure coverage of less than 0.3 bar.

can be ranked in the order $T_c(\text{CO}_2) > T_c(\text{CH}_4) > T_c(\text{N}_2)$.⁷⁵ It has been documented that the gas solubility coefficient in a polymer is positively correlated with its critical temperature.⁷⁶ Due to having the lowest T_c value, the N_2 molecule has the smallest uptake performance. Due to having the highest T_c value, which makes it easy to condense CO_2 within the small pores, the CO_2 molecule has the highest adsorption capacity,⁷⁷ making it largely responsible for the higher selectivity of CO_2/N_2 when compared to its corresponding CO_2/CH_4 gas pair.

Studies have shown that light-emitting MOPs are promising candidates for applications in optoelectronics,^{78,79} chemosensing,^{80,81} light-harvesting antennas,⁸² molecular sensing devices,⁸³ and ion conductivity.⁸⁴ Due to their extensive conjugation we decided to explore the polymers' fluorescence properties. After studying the solid state fluorescence emission spectra (Fig. 6) it was found that, polymers 5 and 6 were fluorescent with $\lambda_{\text{em}} = 280 \text{ nm}$ and $\lambda_{\text{ex}} = 466 \text{ nm}$, while almost no fluorescence was observed for polymers 1 to 4. Meanwhile, the absolute solid state fluorescence quantum yield of polymers 5 and 6 was found to be 0.23 and 0.18 respectively, which demonstrates that polymers knitted by the Scholl coupling reaction may also have potential in sensing, electric conduction, and other applications.

Based on our findings, these HCPs prepared by using different strategies have distinctly different structural features and gas uptake abilities that result from their building blocks' chemical and aggregation structures⁴² and their consequent influence on the commonly employed Friedel–Crafts alkylation

reaction. During the solvent knitting reaction method, CH_2Cl_2 , which acts as both the solvent and an external cross-linker, firstly reacts with AlCl_3 to produce complex $\text{CH}_2\text{Cl}_2 \cdot \text{AlCl}_3$. Since the interaction force between the chlorine atom and the methylene has been weakened, the $\text{CH}_2\text{Cl}_2 \cdot \text{AlCl}_3$ will form CH_2^{2+} and AlCl_4^- , and then the CH_2^{2+} will react with the building block (R). By substituting hydrogen atoms from R, the methylene groups will cross-link R and form polymer networks, with hydrogen chloride as the only byproduct. Due to the methylene groups, the π -conjugated systems and consequently the fluorescence of the resultant polymers will be destroyed. For the knitting method with FDA as an external cross-linker, it is a simple one-step (two stage 45°C for 5 h and then 80°C for 19 h) Friedel–Crafts reaction catalyzed by anhydrous FeCl_3 in 1,2-dichloroethane. In this reaction, the external cross-linker $\text{CH}_3\text{OCH}_2\text{OCH}_3$ activated by FeCl_3 will turn into $\text{CH}_3\text{O}^+\text{CH}_2^+\text{OCH}_3$, by reacting with R, CH_3O^+ will change into the only byproduct CH_3OH by hydrolysis in the presence of trace water, leaving the building blocks bridged by methylene groups. Because of the methylene groups, the π -conjugated systems in polymer materials are interrupted. The Scholl coupling is typically performed at 58°C for 48 h in which R will react with AlCl_3 in $\text{AlCl}_3[\text{H}^+]$ and chloroform, normally resulting R^+ and AlCl_3H . R^+ then engages in a nucleophile substitution reaction with another R, resulting in the elimination of two aryl-bound hydrogen atoms to form a new aryl–aryl bond, leading to novel polymer networks. Since the building blocks are cross-linked by a carbon–carbon single

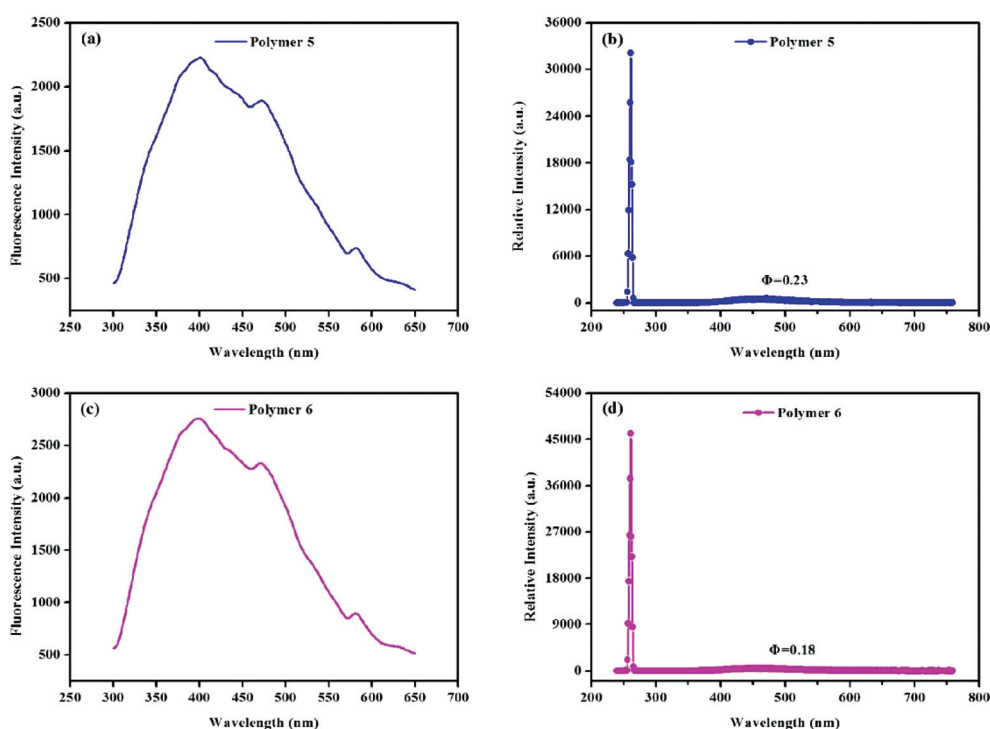


Fig. 6 Solid state fluorescence excitation spectrum of polymer 5 (a, $\lambda_{\text{em}} = 280 \text{ nm}$, $\lambda_{\text{ex}} = 466 \text{ nm}$), the absolute solid state fluorescence quantum yield of polymer 5 (b, $\Phi = 0.23$), the solid state fluorescence excitation spectrum of polymer 6 (c, $\lambda_{\text{em}} = 280 \text{ nm}$, $\lambda_{\text{ex}} = 466 \text{ nm}$), and the absolute solid state fluorescence quantum yield of polymer 6 (d, $\Phi = 0.18$).

bond, the polymers can have extended π -conjugated systems and three-dimensional networks of π - π stacked aryl rings, which are conducive to maintaining fluorescence. It is worth mentioning that, due to the bridging by methylene groups, polymers obtained by the solvent knitting method and the knitting method with FDA as an external cross-linker tend to have similar pore size distributions (Fig. 3b). Due to the connection type (carbon-carbon single bonds), polymers based on the Scholl coupling reaction tend to have narrow pore size distributions (Fig. 3b). Additionally, the solvent knitting method requires 60 h, the knitting method with FDA as an external cross-linker needs 24 h and the Scholl coupling reaction requires 48 h. Therefore, the solvent knitting method is the most time-consuming while the knitting method with FDA as an external cross-linker is the most time-efficient.

Conclusions

A series of hyper-cross-linked polymers based on two aromatic hydrocarbon building blocks and three versatile methodologies *i.e.* the solvent knitting method, the knitting method with FDA as an external cross-linker and the Scholl coupling reaction were made. For the resulting polymers the BET surface area can be as high as $2064 \text{ m}^2 \text{ g}^{-1}$ and $2278 \text{ m}^2 \text{ g}^{-1}$, the CO_2 uptake capacity can be up to 18.95 wt% and 20.89 wt% at 273.15 K/1.00 bar. The systematic and comprehensive study which focused on the porous texture, gas adsorption capacity and fluorescence properties of the prepared materials shows that the solvent knitting method generally produces polymers with better porosities and gas uptake performances. The knitting method with FDA as an external cross-linker is the fastest to generate polymer networks. The Scholl coupling reaction facilitates the formation of knitting fluorescent polymers. It is important to note that these distinctions among the generated microporous HCPs may be largely due to the underlying organic reaction mechanisms. Our study is an attempt to provide original insights into the rational design and synthesis of microporous organic HCPs and offer a route to novel future polymers.

Conflicts of interest

There are no conflicts to declare.

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