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GRAPHICAL ABSTRACT

Facile Synthesis of Triazine-Triphenylamine Based Microporous Covalent Polymer Adsorbent for Flue Gas CO₂ Capture

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A triazine-triphenylamine based microporous covalent organic polymer was prepared through Friedel-Craft reaction under mild conditions. The Framework Lewis basicity favors the interaction with CO₂, which results in enhanced CO₂ capacity and high CO₂/N₂ selectivity, promoted the material as a potential adsorbent for flue gas CO₂ capture.

Facile synthesis of triazine-triphenylamine-based microporous covalent polymer adsorbent for flue gas CO₂ capture

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Abstract

The sustainable capture and sequestration of CO₂ from flue gas emission is an important and unavoidable challenge to control greenhouse gas release and climate change. In this report, we describe a triazine-triphenylamine-based microporous covalent organic polymer under mild synthetic conditions. ¹³C and ¹⁵N solid-state NMR and FTIR analyses confirm the linkage of the triazine and triphenylamine components in the porous polymer skeleton. The material is composed of spherical particles 1.0 to 2.0 μm in size and possesses a high surface area (1104 m²/g). The material exhibits superb chemical robustness under acidic and basic conditions and high thermal stability. Single-component gas adsorption exhibits an enhanced CO₂ uptake of 3.12 mmol/g coupled with high sorption selectivity for CO₂/N₂ of 64 at 273 K and 1 bar, whereas the binary gas mixture breakthrough study using a model flue gas composition at 298 K shows a high CO₂/N₂ selectivity of 58. The enhanced performance is attributed to the high Lewis basicity of the framework, as it favors the interaction with CO₂.

Keywords: covalent porous polymer, CO₂ capture, microporous, flue gas, gas adsorption

1. Introduction

Over the past several decades, significant research effort has been devoted to carbon capture and sequestration (CCS) due to widespread concerns of the rapid increase in the CO₂ level in the atmosphere,

which creates adverse effects such as global climate change and ocean acidification [1-5]. Aqueous amine scrubbing is a commercialized CO₂ capture process, but it is corrosive and energy intensive and thus economically unfeasible for large-scale CCS applications [6, 7]. To date, various CCS technologies have been developed, such as cryogenic distillation, membrane separation and adsorption [4]. Adsorption by porous solid adsorbents (PSAs) has been proposed as one of the most feasible alternatives [7-9]. The process, in principle, incurs a lower energy penalty than amine scrubbing, but the synthesis of high-performance PSAs that meet all practical requirements remains a large challenge [8, 10]. A number of porous materials are under intensive investigation, including activated carbons [11, 12], functionalized silicas [13, 14], metal organic frameworks (MOFs) [15, 16], covalent organic frameworks (COFs) [17, 18], and molecular organic solids [19, 20]. A special group of triazine-based microporous covalent organic polymer (TMCOP) materials developed recently have attracted much interest because these materials have shown promising gas adsorption performances and superb chemical robustness and thermal stability [21-28]. However, most TMCOPs are synthesized under harsh reaction conditions such as high temperature above 673 K and high pressure [21-24]. Herein, we report the synthesis of a novel TMCOP from cyanuric chloride and triphenylamine in the presence of the Lewis acid AlCl₃. A similar TMCOP made from the same starting materials has been reported by Xiong and coworkers [29]. However, their synthesis required a strong acid, CH₃SO₃H, in large excess (14 times excess), which requires careful handling. Our synthetic conditions are much milder, the synthetic procedure is facile and easy to scale-up, and more importantly, our product has a larger surface area, enhanced gas selectivity and superb chemical and thermal robustness.

2. Experimental Section

2.1. Materials

Cyanuric chloride (99%), triphenylamine (Reagent Plus, 99%), *o*-dichlorobenzene (DCB; anhydrous, 99%), anhydrous aluminum chloride (AlCl₃) and hydrochloric acid (HCl) were all purchased from Sigma-

Aldrich. Solvents such as tetrahydrofuran (THF), 1,4-dioxane, chloroform (CHCl_3), ethanol (EtOH), methanol (MeOH) and acetone were also purchased from Sigma-Aldrich. All of the chemicals were used as received without further purification.

2.2. Solvothermal synthesis

In a typical synthetic procedure, anhydrous aluminum chloride (480 mg, 3.0 mmol) was first dissolved in 10 mL of DCB in a flame-dried Schleck tube under sonication. Next, cyanuric chloride (184 mg, 1.0 mmol) was dispersed in 5 mL of degassed DCB through sonication and then added to the first solution. Triphenylamine (245 mg, 1.0 mmol) was quickly dissolved in 5 mL of degassed DCB and added to the first solution in a dropwise fashion using a syringe under vigorous stirring. The mixture was allowed to stir at room temperature for 6 h. The whole process was carried out under an argon environment. The solution was then refluxed at 463 K for 36 h under an argon environment. After cooling to room temperature, the reddish precipitate was collected by filtration, washed sequentially with dry DCB, methanol, ethanol, acetone, THF and dichloromethane, and finally dried in a drying oven. The sample was purified by Soxhlet extraction using methanol and then treated with 1 M HCl-ethanol solution.

2.3. Characterization techniques

Transmission electron microscopy (TEM) images were taken on an FEI Tecnai twin microscope operated at 120 kV. The samples were first ground into a fine powder and dispersed in ethanol. The solvent was evaporated, leaving the sample deposited on a 400-mesh copper TEM grid. Scanning electron microscopy (SEM) images were obtained using an FEI Nova Nano630 SEM and an FEI Helios NanoLab™ 600 DualBeam operated at 10 kV. The Brunauer-Emmett-Teller (BET) specific surface area was measured on an ASAP 2420 system (Micromeritics) using nitrogen at 77 K. Prior to all gas adsorption measurements, the samples were degassed at 453 K for 24 h under high vacuum. The pore size distribution was derived from the adsorption branch using nonlinear density functional theory (NLDFT).

The total pore volume was estimated from the amount of nitrogen adsorbed at a relative pressure of 0.99. Single-component CO₂ and N₂ adsorption isotherms were measured on an ASAP 2050 sorption analyzer (Micromeritics). ¹³C and ¹⁵N NMR measurements were performed on a Bruker 400 M MAS system, and adamantane was used as the internal reference. The rotation frequency was set to 5 kHz. Fourier transform infrared spectroscopy (FTIR) was performed on a Nicolet iS10 smart FTIR spectrometer (Thermo Scientific, USA) equipped with a smart OMNI transmission over a range of 4000 cm⁻¹ to 400 cm⁻¹. Thermogravimetric analysis (TGA) and differential thermal analysis (DTA) were carried out on a thermal analyzer TG 209 (Netzsch) under N₂ flow (20 mL/min). The binary gas mixture breakthrough experiments were carried out at 298 K. Detailed information on the breakthrough setup, as well as descriptions of the calculation of the Henry selectivity, IAST selectivity and isosteric heat of adsorption (Q_{st}), are all available in our previous publications [30, 31].

3. Results and Discussion

Scheme 1 illustrates the formation of the TMCOP structure through a possible Friedel-Craft reaction mechanism in presence of a Lewis acid catalyst, AlCl₃ [32, 33]. Briefly, AlCl₃ first interacts with cyanuric chloride to generate a carbocation complex. Then, the triphenylamine ligand attacks the carbocation. The product becomes more nucleophilic and hence leads to a subsequent polymerization process to generate the porous polymeric network. The fast reaction rate and non-rigid network allows the system to be controlled kinetically instead of thermodynamically, which favors a highly porous but amorphous structure [21, 34]. The covalent bonding and the triazine groups offer high thermal and chemical robustness to the network, while the high density of nitrogen Lewis base favors the adsorption of CO₂, which provides high CO₂ uptake, as well as high selectivity.

Scheme 1: Schematic representation of the redox mechanism for the formation of TMCOP through a polymerization reaction in the presence of a Lewis acid catalyst, AlCl_3 .

The morphology and microstructure of the TMCOP powder are shown in the SEM and TEM images in Fig. 1. Fig. 1a and 1b shows a spherical morphology with relatively uniform sizes of approximately 1.0-2.0 μm . In the high-resolution TEM images (Fig. 1c), low electron density spots (pores) were observed throughout the specimen with an average size of approximately < 1.0 nm, which was further validated by the NLDFT pore size distribution obtained from the nitrogen sorption analysis (inset of Fig.

2A). The wide-angle powder X-ray diffraction confirmed the typical amorphous nature of the TMCOP material, as shown in Fig. 1f [30].

Figure 1: FE-SEM images of the TMCOP material (a); TEM image at low magnification (b) and HRTEM image (c) at higher magnification of the TMCOP material; FE-SEM images of the TMCOP material after immersion in water for a week (d) and in 0.1 (M) hydrochloric acid in ethanol (e); wide-angle powder X-ray diffraction of the TMCOP material (f).

The ^{13}C NMR spectrum of the TMCOP material is shown in Fig. 2. The ^{13}C NMR spectral band detected at 164.882 ppm indicates the presence of the aromatic carbons in the triazine moiety of the cyanuric chloride component. All three carbons of the triazine moiety are equivalent in the TMCOP material [7]. The other broad peaks at 130-160 ppm correspond to the aromatic carbons in the benzene rings in the triphenylamine moiety [32]. The broad spectra depicted the overlap of all four types of carbon atoms (all peaks are expected to be very close) shown in the proposed scheme. In addition, the ^{15}N CP-MAS spectrum showed two resonance peaks at -72.50 and -357.657 ppm, indicating that the two types of nitrogen functionalities are present in the TMCOP material. The signal at -72.50 ppm corresponds to the nitrogen in the triazine component, whereas the other signal at -357.657 ppm signifies the presence of the tertiary nitrogen atom in the triphenylamine moiety.

Figure 2: Schematic representation of the structural bonding in the TMCOP network (a); ^{13}C (b) and ^{15}N (c) solid-state MAS NMR spectra of the porous TMCOP material.

Fig. 3 shows the FTIR spectra of TMCOP, cyanuric chloride and triphenylamine. Several strong vibrational bands were detected in the 1200 to 1550 cm^{-1} region, representing typical stretching modes of CN heterocycles [7, 35]. The nodal lines resulting from standing waves were used to detect the quadrant and semicircle vibrations of the heteroaromatic rings. The distinct bands related to the quadrant (1590 cm^{-1}) and semicircle stretching (1490 cm^{-1}) of the triazine ring and the characteristic breathing mode band of the triazine unit at 813 cm^{-1} were observed in the TMCOP spectrum, as denoted by the

green dotted line in Fig. 3 [36]. The band at 760 cm^{-1} corresponding to the bending vibration of C-H in the present 1,4-disubstitued benzene is seen in the spectrum of the presented network [29]. The stretching vibrational bands near 850 cm^{-1} were observed in the cyanuric chloride spectrum as a result of the C-Cl bond, whereas this vibrational band disappeared in the porous TMCOP network, which implies complete substitution of the three chloride atoms in the cyanuric chloride molecule, as marked by the magenta dotted line in Fig. 3 [7, 25].

Figure 3: FTIR spectroscopy of (a) TMCOP, (b) cyanuric chloride and (c) triphenylamine.

To investigate the thermal stability of the TMCOP material, thermogravimetric analysis of the TMCOP material was carried out up to 1173 K at a ramping rate of 10 K min^{-1} under a continuous N_2 atmosphere, as shown in Fig. 4a. Two sharp drops were observed in the TGA curve. One is before 360 K, which is due to the loss of adsorbed moisture and trapped solvent from the surface and pore interior. The other one is in between 800 and 900 K and is due to the structural decomposition. The experiment demonstrated the high thermal stability of the TMCOP structure, even though it is entirely composed of organic constituents.

The TMCOP material is insoluble in water, and almost all common organic solvents. We further investigated its stability towards humidity, air and/or acid or alkaline media. For example, the material was immersed in water for a week and in 0.1 M HCl or 0.1 M NaOH in ethanol for two days. Then, we investigated the materials by SEM image analyses, and it was observed that the materials retain their spherical microstructure morphology, as shown in Fig. 1d and Fig. 1e.

The nitrogen physisorption isotherm and corresponding pore size distribution of the TMCOP material are shown in Fig. 4b. The obtained isotherm shows a typical type I curve as defined by IUPAC. The observed steep increase in the gas uptake isotherm at low relative pressure (P/P_0) up to 0.09 indicates the presence of a permanent microporosity [37]. The calculated Brunauer-Emmett-Teller (BET) surface area of $1104 \text{ m}^2\text{g}^{-1}$ is somewhat lower compared to the Langmuir surface area of $1302 \text{ m}^2\text{g}^{-1}$. The NLDFT method was used to calculate the pore size distribution, and the majority of the observed pores are micropores approximately 0.65 nm in size. Another peak at 1.19 nm in the supermicropore region was also observed, and this is possibly due to the spatial linkage of the monomers in the polymeric network. No hysteresis loop was observed in the P/P_0 region of 0.40 to 0.80, confirming the absence of large mesopores. The N_2 sorption isotherm showed that desorption branch does not coincide with the adsorption branch. This may be due to the existence of the narrow micropores that kinetically restrict the exit of adsorbed nitrogen from the pore interior during pressure releasing desorption process [29]. The isotherm is not completely closed at $P/P_0 = 0.20$, which is typical for microporous materials [10, 38]. Referring to the results reported by Xiong et al. [29], the reported BET surface is in the range of 428-894 m^2g^{-1} , and the pore size distribution widely spreads from 5 Å to 30 Å. Thus, our sample has a much higher BET surface area and a more uniform pore size distribution.

Figure 4: (a) TGA analysis of the porous TMCOP material and (b) N₂ sorption isotherms of TMCOP measured at 77 K (adsorption: filled circles and desorption: empty circles). The corresponding pore size distribution determined using NLDFT is shown in the inset.

The single-component adsorption isotherms for CO₂ and N₂ at different temperatures and 1 bar pressure and the corresponding selectivities are shown in Fig. 5a-b. In the studied pressure range, the CO₂ uptake increases with increased pressure and does not reach equilibrium or a saturated state [4, 7]. The CO₂ uptake at 1 bar is approximately 3.12 and 1.54 mmol/g at 273 K and 298 K, respectively, whereas the N₂ uptake at 1 bar is approximately 0.20 and 0.12 mmol/g at 273 K and 298 K, respectively. The CO₂/N₂ selectivity in the low-pressure range (< 0.1 bar) was estimated using Henry's law [30, 39]. The calculated Henry selectivity at 273 K is 64, whereas at 298 K, it is 48, estimated from the linear plot of CO₂ and N₂ uptake at 273 K and 298 K, respectively. Compared to the results reported by Xiong et al., where the CO₂ uptake is around 1.31-2.50 mmol/g at 273 K and 0.78-1.40 mmol/g at 298 K and the CO₂/N₂ selectivity is approximately 27.1-33.7 at 273 K and 25.6-38.7 at 298 K [29], both the CO₂ uptake and CO₂/N₂ selectivity are significantly improved by our new synthetic method. The IAST method was further used to calculate the selectivity of CO₂ over N₂ at 10:90 and 15:85 feed molar compositions (model flue gas composition) [7, 40]. The obtained results are given in Figure 5b, which shows that the

selectivity of TMCOP increases with the total feed pressure at a given temperature, while it decreases with increasing temperature at a given feed pressure [41]. For example, at temperature of 273 K and CO₂ to N₂ molar composition of 15:85, the IAST selectivity of CO₂ to N₂ increases from 34 to 46 when total pressure increases from 0.2 to 1 bar, while if the total pressure is fixed at 1 bar, the estimated IAST selectivity (CO₂: N₂ 15:85) decreases from 46 to 25 when temperature increases from 273 K to 298 K. The reason for this trend is because that at low pressure most adsorption sites are unsaturated, so the IAST selectivity is close to the single component Henry selectivity; with rising feed pressure and loading, the competition for adsorption sites between CO₂ and N₂ intensifies; therefore the IAST adsorption selectivity rises due to the much higher affinity of CO₂ to the network. Similar observation was also reported in the literature [30, 41]. When temperature increases (such as 298 K), the interaction of CO₂ to the network becomes weaker and therefore the selectivity will decrease at a given feed pressure compare to the 273 K, as shown in Fig. 5b. The isosteric heat of adsorption Q_{st} for CO₂ is shown in Fig. 5c. The Q_{st} value calculated in the low uptake range (i.e., low-pressure range) represents the interaction strength between the gas and the adsorbent, i.e., the TMCOP network, whereas at higher loadings, it represents the interactions between the gas molecules. At low loading, the estimated Q_{st} value for CO₂ is 26.7 kJ mol⁻¹. The estimated Q_{st} value for CO₂ of the studied TMCOP material was compared with those of interesting materials reported in the literature; the reported materials listed in Table 1 include PPF-3 (19.4 kJ mol⁻¹) [42], PECONF-1 to PECONF-3 (22.2-24.9 kJ mol⁻¹) [10], BILP-2 (18.4 kJ mol⁻¹), CTPP (34 kJ mol⁻¹) [7], Azo-POF-1 to Azo-POF-2 (26.6-27.5 kJ mol⁻¹) [39], Ene-POF-1 to Ene-POF-2 (24.5-24.6 kJ mol⁻¹) [39] and NOP-19 to NOP-21 (28-37 kJ mol⁻¹). Recently, by incorporating different functional groups (such as EtNH₂ or alkynes), modified COF adsorbents showed Q_{st} values ranging from 15.3 to 20.9 kJ mol⁻¹ [43], and for other classes of functional materials, CMP-1-COOH = 33 kJ mol⁻¹ and BILPs = 26.7-28.8 kJ mol⁻¹. The triazine-based frameworks with enriched Lewis base functionality intrinsically favor interactions with CO₂ over those with N₂. In fact, both CO₂ and N₂ are apolar molecules with similar kinetic diameters, typically 0.34 nm and 0.364 nm, respectively, [16, 44, 45]. Moreover, the quadrupole moment of CO₂ is

2.85 times larger, and its polarizability is 1.5 times more than that of N_2 [37, 45]. All of these differences lead to CO_2 interacting more favorably than N_2 with the Lewis base-enriched TMCOP [46-48], consequently leading to high CO_2 uptake through the dipole and quadrupole interactions of CO_2 molecules.

Figure 5: (a) CO_2 and N_2 sorption isotherms for TMCOP at 273 K, 285 K and 298 K at 1 bar; (b) CO_2/N_2 selectivity of TMCOP using the IAST method with a model flue gas in different composition of $CO_2:N_2$; 10:90 (S1), 15:85 (S2) at 273 K, and 10:90 (S3), 15:85 (S4) 298 K at 1 bar; (c) isosteric heat of adsorption of TMCOP at different CO_2 uptake [N.B: The isosteric heat of adsorption is negative.]; (d) column breakthrough experimental results for the TMCOP material using a CO_2 and N_2 gas mixture in a

model flue gas feed composition of CO₂:N₂ (15:85 v/v) measured at 298 K and 1 bar pressure after activation by continuous He flow at 473 K for 12 h.

To demonstrate the effectiveness of the TMCOP material in real applications, we performed breakthrough experiments at ambient pressure and temperature (1 bar; 298 K) under kinetic flow conditions with a model flue gas mixture (15:85 v/v), as shown in Fig. 5d [7, 40]. The obtained breakthrough curves can be split into three phases based on the adsorption characteristics [49]. In phase I, the unsaturated TMCOP adsorbent simultaneously captured both CO₂ and N₂ from the mixed gas feed stream. In phase II, the TMCOP continuously uptakes CO₂; however, no additional N₂ was captured, as the adsorbent was saturated with N₂ in phase I. Moreover, in this phase, the TMCOP adsorbent may competitively and/or preferentially uptake CO₂ while a portion of the N₂ (that was adsorbed in the previous phase I) desorbs into the release gas flow. This is the reason why extra N₂ molecules are present in the release stream in phase II. This observation suggested that the TMCOP material can selectively capture CO₂ over N₂ from a mixed gas feed stream. In phase III, the flow rate and composition of the release stream are the same as that of the feed stream because TMCOP was already saturated with both gases from the prior phases. The estimated adsorption capacities for CO₂ and N₂ are 0.70 mmol/g and 0.068 mmol/g, respectively, which translates to a high CO₂/N₂ selectivity of 58.2. The calculated CO₂/N₂ selectivity is very close to the IAST selectivity. The regeneration characteristics of the TMCOP material were also tested in breakthrough experiments by regenerating the material under relatively mild conditions with helium purging (5 mL/min, 353 K for 0.5 h). The regenerated material exhibits adsorption capacities similar to the CO₂/N₂ selectivity observed in the repeated breakthrough experiments.

Table 1: Comparison of CO₂ and N₂ uptake and CO₂/N₂ separation selectivity of reported porous covalent triazine-based polymers with the synthesized TMCOP material.

Material	CO ₂ uptake (mmol/g)	N ₂ uptake (mmol/g)	CO ₂ /N ₂ selectivity	Q _{st} of CO ₂ (kJ mol ⁻¹)	Reference
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	273 K	298 K	273 K	298 K	273 K	298K		
CTF-1	2.47	1.41	-	-	20	-	27.5	[24]
MCTP-1	4.64	3.08	-	-	-	15	32	[50]
CTP-4	-	2.27	-	-	-	20	33	[32]
NOP-1	2.50	1.40	0.09	-	25.6	-	33.8	[51]
PPF-3	2.09	-	-	-	20	-	21.8	[42]
PAF-26-COOH	2.32	1.43	0.18	0.14	-	20	28.1	[52]
NPOF-1	2.84	1.52	0.29	0.22	-	8	21.7	[53]
NPOF-1-NO ₂	3.97	2.52	0.31	0.23	-	20	29.2	[53]
PAF-5	-	0.27	-	-	-	9.3	26	[54]
TMCOP	3.12	1.54	0.20	0.12	45.6	24.5	26.7	This work

The comparison of the performance of the studied TMCOP material in the breakthrough selectivity of CO₂/N₂ with the different reported interesting materials is addressed here. Well-known amorphous materials, such as activated carbon and metal-anchored activated carbon (KNC-A-K), resulted in CO₂/N₂ selectivities of 4.5 and 44, respectively, [55], as calculated from breakthrough experiments with a 10:90 v/v feed composition of CO₂/N₂ at 298 K and 1 bar. The porous covalent triazine-based materials CTF-1 and fluorinated FCTF-1 exhibited breakthrough selectivities of 18 and 77, respectively, [47], and MOFs, such as SIFSIX-2-Cu-I, exhibited a breakthrough selectivity of 72 [56]. Functional metal-doped materials such as Ni/DOBDC (Ni-MOF-74 or CPO-27-Ni) showed flue gas breakthrough selectivities ranging from 22 to 38 [57]. In view of the above comparisons and progress, the flue gas capture and separation performances of the similar types of triazine-based materials reported in the literature, as given in the comparison in Table 1, demonstrated that the studied TMCOP material has comparable or better performance in some cases.

4. Conclusions

The Lewis base-enriched and electron-rich microporous covalent organic polymer adsorbent demonstrated enhanced CO₂ uptake and high CO₂/N₂ separation, as revealed by both single-component gas adsorption studies and column breakthrough experiments, giving the material potential as an adsorbent for flue gas CO₂ capture. The observed enhanced CO₂ uptake and CO₂/N₂ selectivity is due to quadrupole-dipole interactions between the CO₂ molecules and the Lewis base sites (lone pair of electrons

on the electronegative nitrogen functionality). Furthermore, the material offers permanent porosity with high surface area and exhibits superior robustness, chemical durability and thermal durability. Thus, the unique nanostructure and enhanced properties (stability, enhanced flue gas capture and separation) of the TMCOP material highlight it as a promising adsorbent material for environmental remediation.

List of Parameters/Symbols with unit

BET = Brunauer-Emmett-Teller (m^2g^{-1}).

(P/P_0) = Relative pressures.

Q_{st} = Isostatic heat of adsorption (kJ mol^{-1}).

F_i = Influent flow rate of gas (mL/m).

F_e = Effluent flow rate of gas (mL/min).

q = Absolute adsorption capacity of gas (mmol/g).

α = breakthrough selectivity of gases.

y = Molar fraction of gas in the gas mixture.

R = Universal gas constant (8.315 J/mol K).

IAST = Ideal adsorption solution theory.

q_{sat} = Saturation adsorption capacity (mmol/g).

q_T = Total amount of adsorption (mmol/g).

M = Molarity (mole/liter).

Kinetic Diameter ($\text{\AA}$).

Dipole moment (esu cm).

Quadrupole moment (esu cm^2).

Polarizability (cm^3).

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Highlights

- (1) A triazine-based microporous covalent polymer was prepared under mild conditions.
- (2) High surface area (1104 m²/g), CO₂ uptake (3.12 mmol/g) and Henry selectivity (64).
- (3) High CO₂/N₂ selectivity of 58 in mixture breakthrough study.