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Pore volume regulated CO₂ adsorption in C–C bonded porous organic frameworks†

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Three C–C bonded porous organic polymers are assembled by reacting rigid monomers (triformylphenol, triformylphloroglucinol, triformylresorcinol) with pyrrole in a catalyst-free solvothermal reaction. The resulting black-colored amorphous polymers yield moderate Brunauer–Emmett–Teller surface areas (213 m² g^{−1} to 277 m² g^{−1}). All the polymers exhibit the same pore size but different pore volumes. The impact of pore volume on the adsorption capacity has been investigated. The POFs show excellent CO₂ adsorption properties with moderate heat of adsorption values (31.9–37.6 kJ mol^{−1}) and good IAST selectivity for CO₂/N₂ and CO₂/CH₄ separation. The working capacity of the POFs determined using mixed gas IAST calculations shows that these POFs can preferentially adsorb CO₂ from the CO₂/N₂ gas mixture. POFs show facile adsorption kinetics with a self-diffusion coefficient on the order of 10^{−9} m² sec^{−1}. The POFs offer excellent stability under acidic conditions and retain their working capacity after long humidity exposure. The stability of POFs under harsh acidic conditions has also been established by 273 K CO₂ adsorption, which shows a negligible fall in the CO₂ saturation capacity.

Introduction

The increasing atmospheric carbon dioxide concentrations due to anthropogenic emissions is a major environmental concern.¹ The current level of CO₂ has increased to 420 ppm and is responsible for global warming.² Hence strategies to mitigate CO₂ emissions are urgently needed. A desirable way to reduce carbon dioxide emissions into the environment is by capturing and storing carbon dioxide (CCS) from coal and gas-burning power stations.³ However, the large energy penalty in the capture process has limited the growth of CCS. The energy input necessary for the regeneration of the material is the primary cause of the high cost.³ For instance, the traditional CO₂ capture method based on alkanolamine solutions produces an energy cost of about 30% of the power plant's output.⁴ To tackle this, creating new generation CO₂ capture materials by fine-tuning the thermodynamics of the interaction between CO₂ and the adsorbent is a worthy pursuit.⁴

In this regard, porous materials have drawn interest from academia and industry across many scientific fields and have shown great promise. Porous organic frameworks (POFs) are an important class of porous materials constructed from lightweight

elements linked by strong covalent bonds that show great potential in a wide range of applications, such as gas storage and separation,^{1,5–15} catalysis,^{16,17} chemical sensing,^{18,19} optoelectronics,^{20,21} and energy storage.^{22,23} They have shown great promise due to their low density, high porosity, exceptional thermal and chemical stability, tunable pore size, control of the number of functional groups, *etc.* Along with high porosity, the presence of heteroatoms plays a crucial role in high and selective CO₂ capture; the heteroatoms, especially nitrogen, are known to interact with CO₂ *via* dipole–quadrupole interactions, which is why a lot of work has been carried out on amine grafting and amine impregnation to boost the CO₂ capture performance.^{24,25}

High CO₂ uptake, working capacity and selectivity over other gases (particularly CO₂/N₂ selectivity) under humid conditions, fast adsorption kinetics, cheap production cost, and low energy consumption during adsorption and regeneration phases are crucial requirements that must be met by an effective adsorbent.^{26,27} While constructing a sorbent for CO₂ separation, microstructural qualities such the surface area, pore volume, and pore size must be carefully taken into account in addition to surface functionalization and properties of the gas.²⁸ Materials with pore sizes close to the kinetic diameter of the gas (3.3 Å CO₂, 3.8 Å of CH₄ and 3.64 Å of N₂) would be more suitable due to the possibility of multiple gas–surface interactions.²⁹ However, the correlation between pore volume and CO₂ capacity is less explored and is relevant to porous polymers.³⁰

In 2008, Thomas *et al.* reported the trimerization of aromatic nitriles (Covalent Triazine Frameworks). They gained much

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Fig. 2 (A) Solid state NMR of POFs showing the characteristic peaks. (B) IR-spectra of the POFs showing the peaks corresponding to the different stretching frequencies. (C) Thermo-gravimetric analysis of the POFs showing the high thermal stability. (D)–(F) FE-SEM morphology of 1, 2 and 3 respectively. (SEM images with the scale bar are shown in the ESI†).

of the polymers (Fig. 2B). These results are in agreement with the solid-state magic angle spinning NMR. The thermo-gravimetric analysis proves the excellent thermal stability ($> 350\text{ }^{\circ}\text{C}/623\text{ K}$) for all three polymers; around 20% of the total mass is lost up to $350\text{ }^{\circ}\text{C}$ temperature, and this can be attributed to the solvents residing in the ultra micropores of the polymers (Fig. 2C). Powder X-ray diffraction does not show any sharp peaks confirming the amorphous nature of the POFs (Fig. S1, ESI†). The FE-SEM images show a very homogeneous and similar morphology for all three polymers; they consist of spherical balls inter-grown into clusters (Fig. 2D–F and Fig. S2–S4, ESI†). TEM images also showed similar morphology with uniform micropores on the surface (Fig. S5, ESI†).

The permanent porosity is established by 273 K CO_2 adsorption isotherms; all three POFs show a Type I isotherm. 1, 2 and 3 show CO_2 uptakes of 3, 2.6 and 2 mmol g^{-1} at 273 K, respectively (Fig. 3A–C). The BET fit to the isotherms yields a BET area of 277, 235 and $213\text{ m}^2\text{ g}^{-1}$ and a Langmuir surface area of 345, 289, and $260\text{ m}^2\text{ g}^{-1}$ for 1, 2, and 3, respectively (Fig. S6–S11, ESI†). There is a consistent decrease in the CO_2 uptake capacity with the increase in the $-\text{OH}$ groups in the polymers. The non-local density functional theory (NL-DFT) fit



Fig. 3 (A) Equilibrium CO_2 adsorption isotherms on 1 at different temperatures. (B) Equilibrium CO_2 adsorption isotherms on 2 at different temperatures. (C) Equilibrium CO_2 adsorption isotherms on 3 at different temperatures. (D) Pore size and pore volume distribution of POFs obtained by NLDFT fit to the 273 K CO_2 isotherm. (E) Heat of adsorption plot for POFs calculated using the Virial method. (F) IAST selectivity of POFs for CO_2/N_2 and CO_2/CH_4 gas mixtures.

to the adsorption branch of the 273 K CO_2 isotherm yields a pore size of 0.55 nm in all three cases. However, the pore volume reduced with the increase in the number of hydroxyl groups in the framework (Fig. 3D). Since the POFs are isostructural with the same pore size, a direct correlation between the CO_2 uptake and the pore volume can be made. Hence to design better materials, it is not sufficient to have only ultra-micropores, but maintaining the pore volume on the higher side of the scale is also crucial. Based on this observation, an ideal case would be to have a material with small apertures ($< 0.6\text{ nm}$), which allows only the selective gas molecules to enter through the molecular sieving effect, with a cage-like cavity of larger volumes that can collect large amounts of gas resulting in high working capacity and selectivity. The CO_2 isotherms were also performed at 298 K and 313 K for all the POFs. The CO_2 heat of adsorption was calculated by fitting the pure component isotherms to the virial model and nominal heat of adsorption values were obtained; 37.6, 35.7, and 31.9 kJ mol^{-1} for 1, 2 and 3, respectively, at zero coverage. They decrease linearly with an increase in the loading of gas molecules, signifying the pore-filling mode of adsorption and the absence of any specific strong binding site or the cooperativity among the guest molecules (Fig. 3E and Fig. S12–S14 and



Table S1, ESI†). In order to estimate the ability of our POFs towards selective CO₂ capture, we measured the pure component N₂ and CH₄ isotherms at ambient conditions and 0.11, 0.09 and 0.07 mmol g⁻¹ of nitrogen and 0.55, 0.4 and 0.28 mmol g⁻¹ of methane were adsorbed by **1**, **2** and **3**, respectively. The pure component adsorption isotherms were fitted using the Dual Site Langmuir (DSL) model, and IAST was employed to calculate the selectivity at different concentrations (Fig. S15–S23 and Table S2, ESI†). The CO₂/N₂ IAST selectivity for the flue gas composition was estimated to be 98, 222, and 171 for **1**, **2**, and **3**, respectively. In the case of CO₂/CH₄, the selectivity for a 50 : 50 composition is 17, 45, and 28 for **1**, **2**, and **3**, respectively (Fig. 3F). These numbers are relatively lower compared to the CO₂/N₂ selectivity because of high methane uptakes as compared to nitrogen. To determine the CO₂ working capacity of the polymers under the mixed gas stream (15CO₂:85N₂), ideal adsorbed solution theory (IAST) was employed. The mixed gas isotherms were simulated using the IAST ++ software,³⁸ which reveals that the saturation capacity of CO₂ for a 15CO₂:85N₂ gas mixture is 0.87, 0.89 and 0.65 mmol g⁻¹ for **1**, **2** and **3**, respectively (Fig. 4A).

Practical CO₂ separation is carried out under dynamic conditions; the adsorbent should have facile diffusion within the polymer. To determine the adsorption kinetics, the CO₂ self-diffusion coefficient was calculated as a function of CO₂ loading and it comes out to be $5 \times 10^{-9} \text{ m}^2 \text{ s}^{-1}$ (Fig. 4B), which is much superior to the traditional sorbents such as zeolites. Stability is another key aspect that determines the quality of the adsorbent. The flue gas from which we aim to capture CO₂ consists of water vapors, and acidic gases such as SO_x and NO_x in addition to CO₂ and N₂.³⁹ Since in the flue gas, CO₂ and N₂ are mixed with acidic gases, demonstrating the stability of adsorbents under such

harsh acidic conditions would be meaningful. This is a key property of all-organic C–C bonded porous polymers which could give them an edge over MOFs or other inorganic sorbents. To verify this, we carried out 273 K CO₂ adsorption on the POF samples, which were soaked in 2 M HCl for 24 hours. And no loss in the saturation CO₂ capacity at 273 K was observed (Fig. 4C). Additionally, the POF samples exposed to 99% RH at 50 °C for 6 hours did not show any fall in the CO₂ uptake compared to the pristine POFs (Fig. 4C), revealing their hydrolytic stability. The cycling stability of the best-performer, **1**, was established by the TGA cycling experiment, which shows that the CO₂ working capacity does not fall even after 12 consecutive adsorption-desorption cycles (Fig. 4D).

Water vapor in the flue gas can decompose the adsorbent if it is unstable in humid conditions.⁴⁰ Since water is a polar molecule, it can interact more strongly with the binding sites present in the adsorbent, and hence the positions which are supposed to be occupied by the CO₂ molecules can be taken up by the water molecules, resulting in the loss of working capacity.^{41,42} To avoid this, the humid flue gas stream is pre-treated to adsorb water before it is fed to the CO₂-capturing column, but this adds a parasitic load on the CCS process.⁴³ Hence if the adsorbent has water-repelling properties, it would be an additional advantage.^{37,44} To decipher this, water sorption isotherms were carried out at two different temperatures for all the POFs, and a linear isotherm with a low uptake for **1** was observed. However, the uptake increased going from **1** to **3**; also, the shape of the isotherm changed from near-linear to Langmuir, depicting the stronger interaction of water molecules with the framework (Fig. 5A). Notably, **1**, having the least number of OH functionalities, exhibits the most favorable CO₂ adsorption characteristics and has the least interaction with water. In order to check the surface hydrophobicity, we performed contact angle measurements. We observed that **1** is the most hydrophobic and **3** is the least, and **2** lies in between, all with a contact angle of > 90 (Fig. 5B–D). This confirms that **1**



Fig. 4 (A) Binary CO₂ and N₂ adsorption isotherms simulated using IAST for 15CO₂:85N₂. (B) High resolution 273 K adsorption isotherm of **1** used in this diffusion modelling. Inset shows the CO₂ self-diffusion coefficients from kinetics measurement. (C) 273 K CO₂ adsorption isotherms performed on samples exposed to different conditions. (D) Gravimetric CO₂–N₂ cycling performed on **1**.

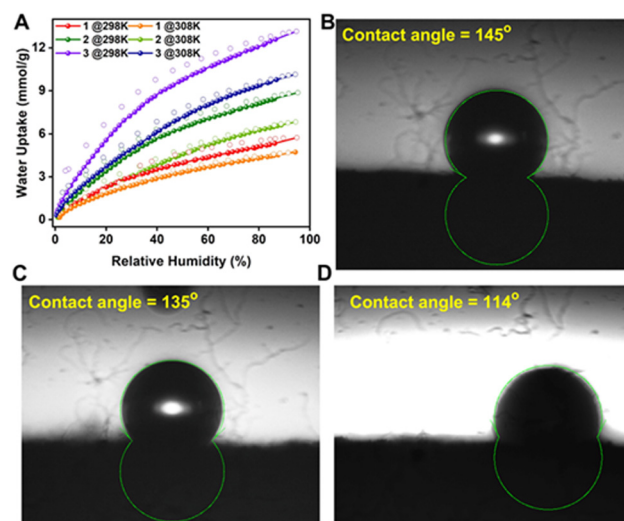


Fig. 5 (A) Water sorption isotherms for **1**, **2**, and **3**. (B)–(D) Water contact angle of **1**, **2**, and **3**.



has the most affinity for water, whereas 3 has the least. The plausible reason is that the number of framework OH groups capable of hydrogen bonding with the water increases from 1 to 3. Hence an increased affinity towards the water in 3 is observed. This could also mean that the hydrophobicity and hydrophilicity can be tuned in these POFs by fixing the number of water-attracting/repelling groups.

Conclusions

In conclusion, we have synthesized three ultramicroporous POFs with similar pore sizes and different pore volumes in a catalyst-free manner. The difference in the pore volumes leads to various working capacities for CO₂. They possess moderate HOA and good CO₂/N₂ and CH₄ IAST selectivity. Water sorption isotherms and contact angle measurements on the three polymers revealed the hydrophilic/hydrophobic character of the materials, which is correlated with the number of phenolic –OH groups present in the material. The C–C bonded framework makes the polymers stable to harsh acidic conditions as well as under a high relative humidity of 99%, which is relevant to the humid flue gas capture. In the future, we aim to synthesize polymers with optimal pore sizes and the pore volume identified in this work, to access smooth adsorption–desorption kinetics. Also, making a comparable polymer with no –OH functionality would give more insights into the role of the phenolic groups in providing CO₂ selectivity over water. Work is in progress.

Experimental

Synthetic procedures

Synthesis of IISERP-POF15 (1). A solvothermal reaction between 2,4,6-triformylphenol (1 mmol) and pyrrole (2 mmol) in a solution containing 5 ml 1,4-dioxane was carried out at 493 K for 48 h. A black colored powder was isolated by filtration and was washed with THF. The air dried sample gave a yield of ~88%.

Synthesis of IISERP-POF16 (2). A solvothermal reaction between 2,4,6-triformylresorcinol (1 mmol) and pyrrole (2 mmol) in a solution containing 5 ml of 1,4-dioxane was carried out at 493 K for 48 h. A black colored powder was isolated by filtration and was washed with THF. The air dried sample gave a yield of ~88%.

Synthesis of IISERP-POF17 (3). A solvothermal reaction between 2,4,6-triformylphloroglucinol (1 mmol) and pyrrole (2 mmol) in a solution containing 5 ml of 1,4-dioxane was carried out at 493 K for 48 h. A black colored powder was isolated by filtration and was washed with THF. The air dried sample gave a yield of ~88%.

Conflicts of interest

There are no conflicts to declare.

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