

ARTICLE

Received 13 Aug 2012 | Accepted 5 Dec 2012 | Published 15 Jan 2013

DOI: 10.1038/ncomms2359

Unprecedented high-temperature CO₂ selectivity in N₂-phobic nanoporous covalent organic polymers

Hasmukh A. Patel¹, Sang Hyun Je¹, Joonho Park¹, Dennis P. Chen¹, Yousung Jung¹, Cafer T. Yavuz¹ & Ali Coskun^{1,2}

Post-combustion CO₂ capture and air separation are integral parts of the energy industry, although the available technologies remain inefficient, resulting in costly energy penalties. Here we report azo-bridged, nitrogen-rich, aromatic, water stable, nanoporous covalent organic polymers, which can be synthesized by catalyst-free direct coupling of aromatic nitro and amine moieties under basic conditions. Unlike other porous materials, azo-covalent organic polymers exhibit an unprecedented increase in CO₂/N₂ selectivity with increasing temperature, reaching the highest value (288 at 323 K) reported to date. Here we observe that azo groups reject N₂, thus making the framework N₂-phobic. Monte Carlo simulations suggest that the origin of the N₂ phobicity of the azo-group is the entropic loss of N₂ gas molecules upon binding, although the adsorption is enthalpically favourable. Any gas separations that require the efficient exclusion of N₂ gas would do well to employ azo units in the sorbent chemistry.

¹Graduate School of Energy, Environment, Water and Sustainability (EEWS), Korea Advanced Institute of Science and Technology (KAIST), Daejeon 305-701, Republic of Korea. ²Department of Chemistry, Korea Advanced Institute of Science and Technology (KAIST), Daejeon 305-701, Republic of Korea. Correspondence and requests for materials should be addressed to C.T.Y. (email: yavuz@kaist.ac.kr) or to A.C. (email: coskun@kaist.ac.kr).

Nanoporous polymers^{1–6} (pore width 1–100 nm) show considerable promise in gas capture⁷, storage⁸ and separation technologies⁹, and as nano-filtration membranes³, catalysts¹⁰, drug delivery vehicles¹¹ and charge carriers^{12,13}. These mostly organic polymers can easily be designed and constructed *via* facile synthetic protocols. To date, several crystalline and amorphous nanoporous organic materials with tunable functionality have been developed—namely, COF¹⁴, PIM¹⁵, HCP and CMP^{16,17}, CTF¹⁸, PAF¹⁹, PPN²⁰, POF²¹, BILP²², EOF²³, PECONF²⁴ and COP²⁵.

CO₂ discharge into the atmosphere has risen with the increasing consumption of fossil fuels³, adversely affecting the fight against global warming²⁶. CO₂ scrubbing by ethanolic aqueous amines²⁷ is yet to be challenged as a practical solution, despite the claims that numerous porous solids have better physicochemical properties and lower energy penalties^{2,28}. In order to avoid energy-intensive CO₂ capture, we have to resort to physisorptive processes (enthalpy of adsorption, $Q < 40 \text{ kJ mol}^{-1}$) rather than chemisorptive ones ($Q > 40 \text{ kJ mol}^{-1}$) (ref. 29). Physisorption, however, brings new challenges. First, there is the lack of CO₂ selectivity as there is no chemical bond being formed. Second, when the functional groups are tuned in order to enhance selectivity, CO₂ uptakes are hindered³⁰. Third, no structure has been identified that rejects N₂, a crucial but missing link in boosting CO₂ selectivity in post-combustion conditions.

In this work, we introduce the concept of N₂ phobicity in order to achieve high CO₂/N₂ selectivities without compromising CO₂ uptake. Our findings show an unprecedented increase in selectivity with increasing temperature, a previously unknown behaviour. To achieve this objective, we have synthesized a whole new class of nitrogen-rich nanoporous polymers, the azo-linked covalent organic polymers (azo-COPs) and found that their azo (–N=N–) groups reject N₂ gas selectively. Azo-COPs, with Brunauer–Emmett–Teller (BET) surface areas up to $729 \text{ m}^2 \text{ g}^{-1}$, can store significant amount of CO₂ (up to 151.3 mg g^{-1}) and show a very high CO₂/N₂ selectivity (288) at 50 °C, an industrially relevant temperature (>40 °C). Azo-COPs are found to be extremely stable up to 350 °C in air as well as in boiling water for a week. A larger entropy loss associated with N₂ adsorption onto the azo groups of the framework as compared with CO₂ explains the high selectivity as well as its temperature dependence. This unique property of azo-COPs, along with their thermal and chemical stability, makes them ideal candidates for post-combustion CO₂ separations.

Results

Synthesis. Azo compounds can be prepared by simple synthetic protocols³¹, where aromatic amines are oxidized using stoichiometric amounts of metal catalysts, resulting in the formation of desired products. Although effective, these methods are not sustainable and environmentally benign because of the need for rare or transition metals³². Here, we have followed a metal catalyst-free synthesis employing the direct coupling of a nitro aromatic compound, tetrakis(4-nitrophenyl)methane, **A**, and amines, 4,4',4'',4'''-methanetetrayltetraaniline, **B1**, *p*-phenylenediamine, **B2**, and benzidine, **B3**, under basic conditions for the formation of interwoven, highly porous and amorphous aromatic azo-COPs (Fig. 1).

Physicochemical characterizations. The formation of azo-COPs was revealed by Fourier transform infrared spectroscopy (FTIR), elemental analysis, cross-polarization magic-angle spinning (CP/MAS) ¹³C NMR and ultraviolet–visible spectroscopies (Supplementary Figs S1–S3). The characteristic stretching band

for –N=N– functionality in the FTIR spectrum is clearly visible at 1,447 and $1,403 \text{ cm}^{-1}$ along with the respective bands for aromatic rings³³. The FTIR bands located at 1,520 and $1,340 \text{ cm}^{-1}$ correspond to N–O stretching mode, suggesting the presence of unreacted terminal nitro groups. The presence of terminal nitro groups is also evidenced by elemental analysis, which indicates the presence of oxygen in azo-COPs (Supplementary methods section). In addition, the presence of bands in the range of $3,600\text{--}3,000 \text{ cm}^{-1}$ indicates that there are terminal amino groups present in the structure. The presence of chemical shifts in CP/MAS ¹³C NMR spectra of azo-COPs, namely, azo-COP-1, azo-COP-2 and azo-COP-3 (also named as COP-68, -69, -70), (www.porouspolymers.com) located at 150.2, 144.7, 129.5, 123.4 and 54.9 ppm confirmed the formation of the azo-linked aromatic polymers^{19,34}. Azobenzene moieties have a characteristic ultraviolet–visible spectra where an intense absorption band at 360 nm indicates the presence of *trans*-azobenzene unit³⁵. The solid-state ultraviolet–visible analysis revealed³⁴ that the as-synthesized azo-COPs have predominantly *trans* configuration and feature low band gaps (Supplementary Figs S3, S4).

Porous polymers are known³⁶ to be amorphous solids. Broad powder X-ray diffraction patterns of azo-COPs (Supplementary Fig. S5) refer to their amorphous nature. Small-angle X-ray scattering analysis of azo-COPs, however, exhibited (Supplementary Fig. S6) mesostructured character for azo-COP-1 and -2, whereas azo-COP-3 has a completely amorphous framework. The thermogravimetric analysis of azo-COPs (Supplementary Fig. S7) indicate that they are stable up to 350 °C in air and 300 °C under N₂ environment. The mass losses can be assigned to desorption of trapped solvent molecules and the decomposition of the framework. It is well-known²⁵ that CO₂ scrubbing operations require high chemical stability as harsh conditions such as steam are employed frequently. In order to investigate the water stability of azo-COPs, we kept the polymers in boiling water for a week and measured their surface areas. Azo-COPs did not show any noticeable loss of surface area or CO₂ capacity (Supplementary Fig. S8), indicating that network structures are highly robust. The bulk-scale morphology of azo-COPs was examined (Supplementary Figs S9, S10) with field-emission scanning electron microscope and transmission electron microscope. Irregular-shaped particles of similar sizes are arranged to give a macroporous surface. The reasonably monodisperse grains are consistent with a homogeneous reaction, an essential feature for a large-scale industrial production. Azo-COP-2 has the smallest particles of all three, hence its enhanced gas sorption. Azo compounds feature low band gaps (for example, 2.22–2.35 eV for azo-COPs), evaluated from ultraviolet–visible diffuse reflectance spectroscopy (Supplementary Fig. S4). Azo-bridged polymeric materials are also widely studied³⁷ for their optical storage applications. As optically stored information is only stable below glass transition temperature (T_g), a high T_g means more potential in optical data storage. All three azo-COPs show T_g values of 158.6 °C (Supplementary Fig. S11) in a differential scanning calorimetry (DSC) investigation, suggesting their potential for incorporation into optical storage devices.

Argon physisorption measurements at 87 K were performed³⁸ (Fig. 2) in order to evaluate the pore structure of the azo-COPs. We have shown that azo-COP-1, -2 and -3 are microporous and exhibit typical type-I reversible sorption profiles. The BET surface areas were found to be 635, 729 and $493 \text{ m}^2 \text{ g}^{-1}$ (BET linear plot, Supplementary Fig. S12 and Supplementary Table S1), and total pore volumes of 0.39, 0.44 and $0.32 \text{ cm}^3 \text{ g}^{-1}$ for azo-COP-1, azo-COP-2 and azo-COP-3, respectively. The pore width maxima of azo-COP-1, azo-COP-2 and azo-COP-3 centred at 0.7, 0.48 and

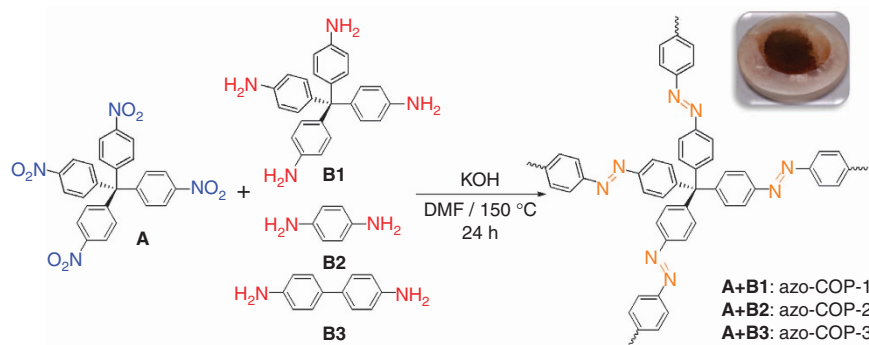


Figure 1 | Syntheses of a series of azo-COPs. Orange powders of azo-COPs were obtained by reacting tetrakis(4-nitrophenyl)methane, **A**, with three aromatic amines, namely 4,4',4'',4'''-methanetetrayltetraaniline (**B1**, azo-COP-1), *p*-phenylenediamine (**B2**, azo-COP-2) and benzidine (**B3**, azo-COP-3), in DMF at 150 °C in the presence of potassium hydroxide under an atmosphere of N₂.

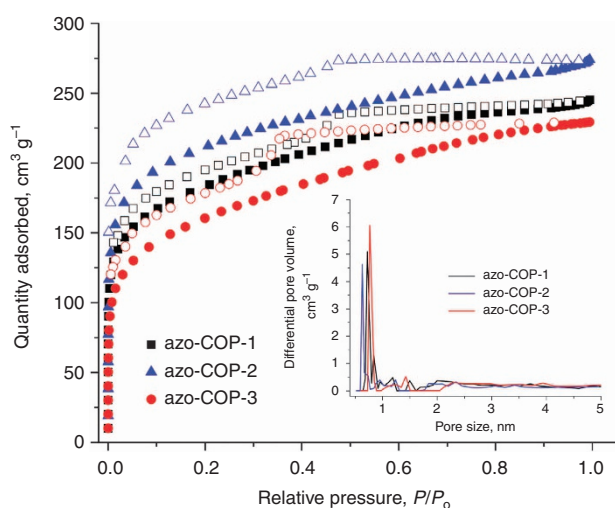


Figure 2 | Argon adsorption-desorption isotherms of azo-COPs at 87 K.

Filled and empty symbols represent adsorption and desorption, respectively. For nitrogen isotherms at 77 K and BET linear plots of azo-COPs see Supplementary Fig. S12. Inset: pore volume versus pore size. All the experiments were carried out using 1 g of each azo-COP.

0.8 nm, respectively. The hysteresis at low pressures may be attributed to the pore networks effect³⁶ and also to the irreversible binding of gas molecules to the micropore surface. The calculated BET surface areas obtained from N₂ sorption isotherms of azo-COPs at 77 K (Supplementary Figs S13, S14) commensurate with the ones from Argon physisorption measurements. The BET surface areas of azo-COPs were also measured using CO₂ as an adsorbate at 273 K (ref. 39), demonstrating surface areas of 303, 334 and 206 m² g⁻¹ for azo-COP-1, azo-COP-2 and azo-COP-3, respectively (Supplementary Fig. S15).

Selective gas sorption. CO₂ and N₂ are two apolar molecules with similar and small kinetic diameters, namely 3.30 and 3.64 Å (ref. 40), respectively. Their quadrupole moments (2.85 to 1) and polarizabilities (1.5 to 1), however, differ enough to design CO₂-philic solids, which would be neutral towards N₂. Owing to the presence of an electron-deficient carbon atom, CO₂ is expected to interact with protic electronegative functionalities (for example, amines) leading to a strong chemisorption, while N₂ remains impartial. So far, this simple interaction has led to a plethora of nanoporous polymers where the nitrogen-rich

functionalities, such as triazine^{1,18}, tetrazole³, imidazole⁴¹, phosphazene²⁴, imide⁴² and amines^{25,43}, are built in. In all these and many other situations², CO₂-philicity is the main driving force for the CO₂/N₂ selectivity, without taking into account any chemical architecture that would repel N₂ molecules selectively.

The significance of the microporosity and the nitrogen-rich framework of azo-COPs have been investigated for their uptake of CO₂ and N₂ gas molecules. The CO₂ and N₂ adsorptions of the azo-COPs were measured up to 1 bar at 263, 273, 298 and 323 K in order to mimic post-combustion CO₂ capture conditions (Fig. 3, Table 1). Before the measurements, the azo-COPs were degassed at 150 °C for 5 h (see Methods) for both activation purposes and also to avoid any contribution from *cis*-azobenzene units. All the measurements were carried out using 1 g of azo-COPs to minimize errors associated with the mass of the adsorbents. The CO₂ isotherms are completely reversible and show an adsorption capacity of 145.4 mg g⁻¹ for azo-COP-1, 151.3 mg g⁻¹ for azo-COP-2 and 103.3 mg g⁻¹ for azo-COP-3 at 263 K. This high affinity is a consequence of the favourable interactions of the polarizable CO₂ molecules through dipole-quadrupole interactions with the framework, and also the inherent microporosity of azo-COPs. At 273 K, CO₂ uptake is 107.6 mg g⁻¹ for azo-COP-1, 112.4 mg g⁻¹ for azo-COP-2 and 85.1 mg g⁻¹ for azo-COP-3. As expected, the CO₂ uptake of azo-COPs decreases at 298 K, to 65.3 mg g⁻¹ for azo-COP-1, 67.3 mg g⁻¹ for azo-COP-2 and 53.6 mg g⁻¹ for azo-COP-3. These values are still comparable to the very high surface area porous polymers²⁵, such as PPN-4 (ref. 20) (65 mg g⁻¹) and PAF-1 (ref. 19) (45 mg g⁻¹). A smooth adsorption-desorption cycle implies that the interaction between CO₂ and the networks are weak enough to regenerate these azo-COPs without too much energy. Similar behaviour has been reported where adsorption on basic sites do not require heating to regenerate porous polymers^{41,44}. The CO₂ isosteric heat of adsorption (Q_{st}) is calculated from the adsorption data at 263, 273, 298 and 323 K⁴⁵. The Q_{st} for azo-COP-1, azo-COP-2 and azo-COP-3 are in the range of 24.8–32.1 kJ mol⁻¹, values which are comparable to those obtained with the previously reported⁴⁵ porous polymers. Slightly higher heats of adsorption are ascribed to the presence of -N=N- moieties in the framework of the azo-COPs.

For a post-combustion CO₂ adsorption, CO₂ should be adsorbed preferentially over N₂ at high temperatures (> 40 °C). To the best of our knowledge, however, all the porous polymers reported so far^{2,41,44} exhibit significant loss in CO₂/N₂ selectivity on going from 273 to 298 K (Supplementary Table S2). Contrary to the common trend, azo-COPs show (Table 1) higher CO₂/N₂ selectivities with increasing temperatures. The selectivities were

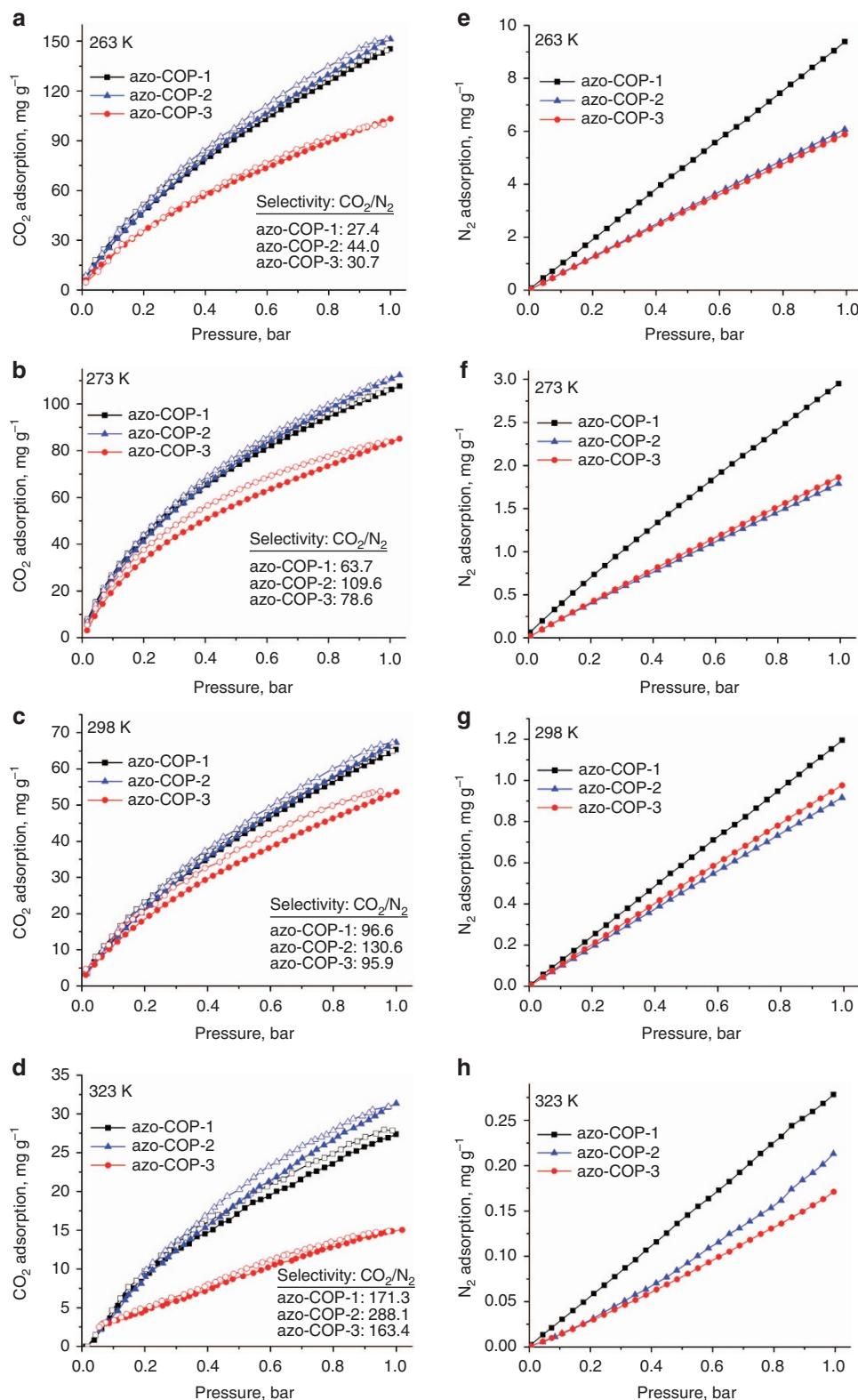


Figure 3 | CO₂ and N₂ adsorption-desorption isotherms of azo-COPs at different temperatures. CO₂ (a–d) and N₂ (e–h) gas isotherms measured up to 1 bar at 263, 273, 298, 323 K. Filled and empty symbols represent adsorption and desorption, respectively. Inset: respective CO₂/N₂ selectivity. Increase in the CO₂/N₂ selectivity with increasing temperature proves the N₂ phobicity of the framework while retaining its CO₂-philic character.

calculated using the Ideal Adsorbed Solution Theory (IAST) for CO₂:N₂ mixture in the ratio of 0.15:0.85 (Table 1, Supplementary Figs S16–S20) and Henry's Law constants from the isotherms in the linear low pressure (<0.1 bar) range

(Supplementary Table S2, Supplementary Figs S21–S23). The IAST CO₂/N₂ selectivities of the azo-COPs range from 27.4 to 44.0 at 263 K, 63.7 to 109.6 at 273 K, 95.9 to 130.6 at 298 K and become 163.4 to 288.1 at 323 K, which is being the highest

Table 1 | CO₂ and N₂ uptakes, selectivities and isosteric heat of adsorptions of azo-COPs.

T (K) =	CO ₂ adsorption (mg g ⁻¹)				N ₂ adsorption (mg g ⁻¹)				Selectivity* CO ₂ /N ₂				Q _{st} † (kJ mol ⁻¹)	
	263	273	298	323	263	273	298	323	263	273	298	323	CO ₂	N ₂
azo-COP-1	145.4	107.6	65.3	27.4	9.39	2.95	1.19	0.28	27.4	63.7	96.6	171.3	29.3	24.1
azo-COP-2	151.3	112.4	67.3	31.4	6.07	1.79	0.92	0.21	44.0	109.6	130.6	288.1	24.8	19.5
azo-COP-3	103.3	85.1	53.6	15.0	5.89	1.86	0.98	0.17	30.7	78.6	95.9	163.4	32.1	21.0

*From IAST at CO₂:N₂ ratio of 0.15:0.85 at 1 bar.†Q_{st}: Isosteric heat of adsorption.

reported CO₂/N₂ selectivity at 323 K to date. N₂ uptake drops ~70% at a temperature increase from 273 to 298 K, compared with a ~40% drop for CO₂. This unprecedented behaviour cannot be explained only by conventional CO₂ affinities. A new concept, namely N₂ phobicity, is the result.

When the CO₂/N₂ selectivities found here are compared with those for other nitrogen-rich porous polymers (Supplementary Table S2), azo-COPs show a lot less affinity towards N₂. For example, covalent triazine-based frameworks (CTFs) reveal a mere CO₂/N₂ selectivity of 31.2 at 273 K. Our earlier work on triazines²⁵ leads to the understanding⁴⁶ that an imidic bond (–C=N–) is not N₂-phobic. Benzimidazole-linked polymers⁴¹ show selectivities up to 113 at 273 K, being reduced to 71 at 298 K. Although nitrogen-rich organic cage frameworks^{30,47} show high CO₂/N₂ selectivities up to 213 at 298 K, their surface areas (<10 m² g⁻¹) and CO₂ capture capacities (up to 7.15 mg g⁻¹) are very low. Other structures with protic nitrogens also behave in a similar fashion^{43,44,48}.

Computational studies. As the aromatic rings present in the azo-COPs cannot be the reason for N₂ phobicity^{19,20}, we studied the effects of the –N=N– moieties in the framework on the near rejection of N₂ at higher temperatures, 323 K, as well as the temperature dependence of the increased selectivity. Incidentally, similar observations have been noted in a very recent report⁴⁹, where azo-bridged linkers are compared with ethylenic ones in crystalline metal-organic frameworks (MOFs) for CO₂ uptake. The authors, however, ignored the obvious N₂-phobic behaviour of azo-MOFs when N₂ uptakes are compared. Under the same conditions (195 K), argon uptake follows a nonspecific interaction, thus serving as a control for the N₂-phobic behaviour of the framework incorporating azo-bridged linkers. Our findings are commensurate with their data, in the sense that taken together, they prove the existence of N₂ phobicity.

We first compared the relative binding affinity of the –N=N– group towards CO₂ versus N₂ using *trans*-azobenzene as a model system (Fig. 4a). The lowest energy structure of the CO₂/*trans*-azobenzene complex indicates that CO₂ preferentially binds with the azo-group of *trans*-azobenzene with binding energy (BE) of 17.0 kJ mol⁻¹ while N₂ preferentially binds to the phenyl-group with BE = 11 kJ mol⁻¹, rather than to azo (BE = 9 kJ mol⁻¹). CO₂ also has a favourable interaction with aromatic ring involving large stabilization energy, 14.4 kJ mol⁻¹. As the experimental heat of adsorption for the present COPs observed for CO₂ is 25–32 kJ mol⁻¹ (Table 1), it appears that CO₂ interacts with more than one azo- or aromatic functionalities in COPs owing to the amorphous structural nature. Owing to a difficulty in structure prediction for the amorphous polymers³⁶, we instead considered the coordination pillared-layer network (CPL-4)⁵⁰, a closest crystalline MOF with azo moiety. Incidentally, our computed binding energies of the azo-group with CO₂ and N₂ are similar to the observed CO₂ and N₂ heats of adsorption⁵⁰ for CPL-4, -20 and -12 kJ mol⁻¹, respectively,

where the multiple interactions with more than one functional group is not feasible owing to the structural rigidity of the MOFs. Of the two interaction moieties in the experimental azo-COPs, azo and aromatic rings, the azo-group is clearly the deciding factor for N₂ phobicity considering a large difference in azo-affinity for CO₂ versus N₂ (17.0 versus 9.0 kJ mol⁻¹), while the phenyl ring has a relatively comparable binding affinity towards both CO₂ and N₂, 14.4 and 11.0 kJ mol⁻¹, respectively. These results support the experimental trend that more azo-substitution increases the selectivity. The *ab initio* relative binding affinity for CO₂ versus N₂ (3–8 kJ mol⁻¹) is in reasonable agreement with the experimental relative heat of adsorption data for various azo-COPs (5–9 kJ mol⁻¹). Energy decomposition analysis⁵¹ suggests that the frozen density repulsion as well as the lack of sufficient dispersion interactions between N₂ gas molecules and the azo groups are responsible for the moderate binding strength of N₂/azo interaction compared with the stronger CO₂/azo binding (Supplementary Fig. S24). Despite the favourable heat of adsorption (–19.5 kJ mol⁻¹), the fact that N₂ is nearly rejected, we can conclude that the rejection must have its origin in entropy. From this viewpoint, an upper bound of the adsorption entropy for N₂ might be estimated approximately as:

$$\Delta S(\text{azo-COP-2: N}_2) < \frac{\Delta H(\text{N}_2)}{T} = \frac{-19.5 \text{ kJ mol}^{-1}}{323 \text{ K}} = -60.4 \text{ JK}^{-1} \text{ mol}^{-1} \quad (1)$$

In order to understand the observed temperature dependence of the selectivity, we recall that the selectivity is related to

$$\Delta \Delta G_{\text{ads}} = \Delta G_{\text{ads}}(\text{CO}_2) - \Delta G_{\text{ads}}(\text{N}_2) \quad (2)$$

Where ΔG_{ads} is the free energy of adsorption and its temperature derivative is the entropy of adsorption (assuming constant heat capacity or enthalpy)

$$\Delta S_{\text{ads}} = -\frac{\partial \Delta G_{\text{ads}}}{\partial T} \quad (3)$$

In other words, the rate of increase in adsorption free energy with temperature is determined by the entropy change upon gas adsorption. As

$$\Delta G_{\text{ads}} = \Delta H_{\text{ads}} - T\Delta S_{\text{ads}} \quad (4)$$

in usual enthalpy–entropy compensation, the higher BE (large negative ΔH_{ads}) would cause a tighter association with a corresponding loss in entropy (large negative ΔS_{ads}). Therefore, the conventional enthalpy–entropy compensation would predict $\Delta \Delta G_{\text{ads}}$ to increase (or selectivity to decrease) with temperature owing to a negative

$$\Delta \Delta S_{\text{ads}} = \Delta S_{\text{ads}}(\text{CO}_2) - \Delta S_{\text{ads}}(\text{N}_2), \quad (5)$$

that is, $\partial \Delta \Delta G_{\text{ads}} / \partial T = -\Delta \Delta S_{\text{ads}} > 0$, as in most previous temperature-dependent selectivity measurements^{2,41,44}. However, the present azo-COPs show the opposite temperature dependence of the CO₂/N₂ selectivity, suggesting positive $\Delta \Delta S_{\text{ads}}$, that is, the entropy loss in the event of N₂ adsorption should

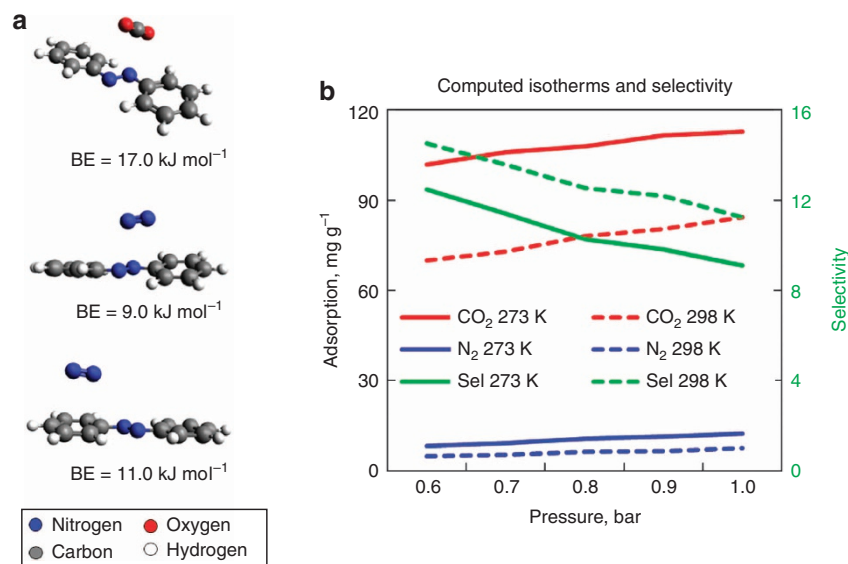


Figure 4 | Computational analysis for the gas affinity of azo linkages. (a) The RIMP2 minimum energy structures of CO₂/*trans*-azobenzene and N₂/*trans*-azobenzene complexes with the associated binding energies (BEs). (b) The computed isotherms for CO₂ (red) and N₂ (blue) obtained through GCMC simulation and the CO₂/N₂ selectivity (green) at 273 K (solid line) and 298 K (dashed line).

be larger than that associated with the CO₂ adsorption (Supplementary Fig. S25).

In order to support this entropic origin qualitatively, we performed the grand canonical Monte Carlo (GCMC) simulations and analysed the trajectories on CPL-4. Figure 4b shows that the isotherms of CO₂ (red) and N₂ (blue), and the selectivity (green) at 273 (solid line) and 298 K (dashed line) within the pressure range between 0.6 and 1.0 bar. For convenience, we use the computed selectivity simply defined as the uptake ratios of different gases. As temperature rises, the gas uptakes decrease as expected; however, the selectivity increases (from solid to dashed lines) as in experiments.

The interaction energy distribution between the gas and framework and the distance distribution between the centre of mass of the gas molecule and the centre of the azo-group are simulated at 263 K and 1.0 bar (Supplementary Fig. S26). For both distributions in energy (0.6 versus 4.1 kJ mol⁻¹ for N₂ versus CO₂) and configuration (0.48 versus 0.70 Å) space, the widths of distributions corresponding to the CO₂ binding with azo are much broader than those for N₂. A similarly narrower distance distribution between N₂ and the aromatic ring (width being 0.50 Å) was also observed. No significant change in the energy distribution was observed with pressure up to 1 bar (Supplementary Fig. S27), indicating that the binding sites are roughly the same regardless of the loading. The narrower distance distributions obtained for N₂ binding qualitatively suggest that azo–N₂ interactions have configuration space that is much less accessible for binding than the case of CO₂, which makes the adsorbed N₂ retain lower (more unfavourable) entropy than the adsorbed CO₂. The net effect is then positive $\Delta\Delta S_{\text{ads}}$, leading to negative temperature dependence in $\partial\Delta\Delta G_{\text{ads}}/\partial T = -\Delta\Delta S_{\text{ads}} < 0$, or increased selectivity with temperature. We note, however, that our interpretation assumes constant (or weak) temperature dependence for enthalpy of adsorption for CO₂ and N₂, which needs to be further investigated.

Discussion

We have evolved a catalyst-free synthesis of robust, N₂-phobic, nanoporous azo-COPs with surface areas up to 729 m² g⁻¹.

These azo-COPs reveal carbon dioxide uptake twice as much as the industrial standard, monoethanolamine, while the selectivity for carbon dioxide over nitrogen is 288 at 50 °C. The increased selectivity for carbon dioxide over nitrogen, as well as the thermal and water stability of the azo-COPs at industrially relevant temperatures, suggests potential applications for these materials in post-combustion carbon dioxide capture and in air separations. It is not unlikely that these N₂-phobic azo-COPs will provide design principles for the development of next-generation porous polymers for highly selective gas separations, such as natural gas purification.

Looking into the broader picture, CO₂-related gas separations require the following four fundamental guidelines for the proposed adsorbent materials: (1) water stability²⁹, (2) selectivity³, (3) thermal stability¹ and (4) cost²⁵. Any newly discovered sorbent candidate has to be evaluated for all these prospects. Not to mention, studying temperatures < 40 °C usually mislead in identifying the best sorbent as both pre-combustion and post-combustion processes run at warmer temperatures.

Methods

Synthesis of azo-COPs-1,2,3. All reagents and solvents employed in the synthesis were commercially available and used as received without further purification. In a typical synthesis, orange powders of azo-COPs were obtained in 40–57% yields by reacting tetrakis(4-nitrophenyl)methane with aromatic amines—for example, benzidine, *p*-phenylenediamine, 4,4',4'',4'''-methanetetrayltetraaniline—in DMF in the presence of KOH under an inert atmosphere of N₂, where the reaction temperature was raised to 150 °C and stirred for 24 h. Once completed, the reaction was cooled to room temperature and stirred in an excess of distilled water for 1 h. The precipitate was filtered off and washed with warm distilled water (× 4), followed by washing with Me₂CO (× 3) and THF (× 3). Subsequently, an orange precipitate was dried at 110 °C under vacuum for 4 h. All the azo-COPs were reproducible on multigram scales. The synthesis of monomers and detailed synthetic routes for azo-COPs are provided in the Supplementary methods section.

Characterization methods. FTIR spectra were recorded on KBr pellets using a Perkin-Elmer FTIR spectrometer. Solid-state CP/MAS ¹³C NMR spectra of azo-COPs were collected by a Bruker Avance III 400 WB NMR spectrometer. Liquid ¹H and ¹³C NMR spectra were taken using DMSO-*d*₆ as the solvent in a 400 MHz Bruker NMR spectrometer. Thermogravimetric analysis was performed on a NETZSCH-TG 209 F3 instrument by heating the samples to 800 °C at a rate of 10 °C min⁻¹ under air and N₂ atmospheres. DSC of azo-COPs was measured by NETZSCH DSC 204 F1 instrument (Germany) at a scan rate of 10 °C min⁻¹ in a nitrogen flow in the range of –60 to +280 °C. X-ray diffraction patterns of the

samples were acquired from 2 to 70° by a Rigaku D/MAX-2500 (18 kW) micro-area X-ray diffractometer. Small-angle X-ray scattering data for powdered azo-COPs were obtained on a Rigaku D/MAX-2500 diffractometer. The morphology of azo-COPs was studied by field emission scanning electron microscopy on a Philips XL30SFEF instrument. TEM images were collected by field emission transmission electron microscope (200 KV, Tecnai F20). Samples for TEM investigations were prepared by putting an aliquot of alcoholic dispersion of azo-COPs onto an amorphous carbon substrate supported on a copper grid. The excess liquid was then removed, and the grid was allowed to dry at room temperature. The absorbance spectra of azo-COPs were collected on a ultraviolet–visible spectrophotometer where the powder samples were sandwiched between two quartz glasses and irradiated by a 365-nm ultraviolet lamp (8 W; 230 V; ~50/60 Hz from Vilber Lourmat, France). In order to evaluate the porosity of azo-COPs, Ar and N₂ adsorption isotherms were obtained with a Micromeritics ASAP 2020-accelerated surface area and porosimetry analyser at 87 and 77 K, respectively, after the samples had been degassed at 150 °C for 5 h under vacuum. The adsorption–desorption isotherms (Ar at 87 K and N₂ at 77 K) were obtained to give the pore parameters, including BET ($P/P_0 = 0.01–0.25$) and Langmuir ($P/P_0 = 0.1–0.35$) surface area, pore size and pore volume. The pore size distribution of azo-COPs was calculated from the Ar and N₂ adsorption isotherms by the Non-Local Density Functional Theory method using a cylindrical pore model. IAST data were calculated using OriginPro v8.5 (Supplementary methods section). Pore volume of azo-COPs was calculated at $P/P_0 = 0.95$. The low-pressure CO₂ and N₂ adsorption–desorption isotherms for azo-COPs were measured at 263, 273, 298 and 323 K using a static volumetric system (ASAP 2020, Micromeritics Inc., USA). The temperature during adsorption and desorption was kept constant using a circulator. Before the adsorption measurements, the samples were dried at 110 °C for 24 h. The samples were further activated *in situ* by increasing the temperature at a heating rate of 1 K min⁻¹ up to 423 K under vacuum (5×10^{-3} mm Hg) and the temperature and vacuum was maintained for 5 h before taking the sorption measurements. All the adsorption–desorption experiments were carried out twice to ensure the reproducibility. There were no noticeable differences in the isotherm points obtained from both experiments. Heats of adsorption were determined from CO₂ and N₂ adsorption isotherms at 263, 273, 298 and 323 K using a Micromeritics ASAP 2020 instrument and the standard calculation routines in the Datamaster offline data reduction software (Micromeritics).

Computational details. Quantum chemical calculations for the binding energies of CO₂ and N₂–*trans*-azobenzene complex and their energy decomposition analysis were performed using Q-CHEM⁵². To describe nonbonding interactions between the framework and gas molecules correctly, we used the RIMP2/VDZ(d) geometry optimizations, and the final binding energies were further refined with the complete basis set limit using the RIMP2/aug-cc-pV(DT)Z extrapolation. The isotherms and equilibrium ensembles for the CO₂ and N₂ uptakes in the CPL-4 frame containing the azo-group were obtained using GCMC simulations with the Cerius2 v4.0 software. The force field parameters and charges for CO₂ were adopted from Harris *et al.*⁵³—Dreiding 2.21 and equilibrium charge method.

References

- Thomas, A. Functional materials: from hard to soft porous frameworks. *Angew. Chem. Int. Ed.* **49**, 8328–8344 (2010).
- Dawson, R., Cooper, A. I. & Adams, D. J. Nanoporous organic polymer networks. *Prog. Polym. Sci.* **37**, 530–563 (2012).
- Du, N. Y. *et al.* Polymer nanosieve membranes for CO₂-capture applications. *Nat. Mater.* **10**, 372–375 (2011).
- Kitagawa, S., Kitaura, R. & Noro, S. Functional porous coordination polymers. *Angew. Chem. Int. Ed.* **43**, 2334–2375 (2004).
- Férey, G. Hybrid porous solids: past, present, future. *Chem. Soc. Rev.* **37**, 191–214 (2008).
- Yaghi, O. M. *et al.* Reticular synthesis and the design of new materials. *Nature* **423**, 705–714 (2003).
- Xiang, S. *et al.* Microporous metal-organic framework with potential for carbon dioxide capture at ambient conditions. *Nat. Commun.* **3**, 1–9 (2012).
- Cote, A. P. *et al.* Porous, crystalline, covalent organic frameworks. *Science* **310**, 1166–1170 (2005).
- Budd, P. M. *et al.* Polymers of intrinsic microporosity (PIMs): robust, solution-processable, organic nanoporous materials. *Chem. Commun.* 230–231 (2004).
- Kaur, P., Hupp, J. T. & Nguyen, S. T. Porous organic polymers in catalysis: opportunities and challenges. *ACS Catal.* **1**, 819–835 (2011).
- Zhao, H. Y. *et al.* Targeted synthesis of a 2D ordered porous organic framework for drug release. *Chem. Commun.* **47**, 6389–6391 (2011).
- Wan, S. *et al.* Covalent organic frameworks with high charge carrier mobility. *Chem. Mater.* **23**, 4094–4097 (2011).
- Jiang, J. X., Trewin, A., Adams, D. J. & Cooper, A. I. Band gap engineering in fluorescent conjugated microporous polymers. *Chem. Sci.* **2**, 1777–1781 (2011).
- El-Kaderi, H. M. *et al.* Designed synthesis of 3D covalent organic frameworks. *Science* **316**, 268–272 (2007).
- McKeown, N. B. & Budd, P. M. Exploitation of intrinsic microporosity in polymer-based materials. *Macromolecules* **43**, 5163–5176 (2010).
- Wood, C. D. *et al.* Hydrogen storage in microporous hypercrosslinked organic polymer networks. *Chem. Mater.* **19**, 2034–2048 (2007).
- Jiang, J. X. *et al.* Conjugated microporous poly(aryleneethynylene) networks. *Angew. Chem. Int. Ed.* **46**, 8574–8578 (2007).
- Schwab, M. G. *et al.* Catalyst-free preparation of melamine-based microporous polymer networks through schiff base chemistry. *J. Am. Chem. Soc.* **131**, 7216–7217 (2009).
- Ben, T. *et al.* Targeted synthesis of a porous aromatic framework with high stability and exceptionally high surface area. *Angew. Chem. Int. Ed.* **48**, 9457–9460 (2009).
- Yuan, D. Q., Lu, W. G., Zhao, D. & Zhou, H. C. Highly stable porous polymer networks with exceptionally high gas-uptake capacities. *Adv. Mater.* **23**, 3723–3725 (2011).
- Katsoulidis, A. P. & Kanatzidis, M. G. Phloroglucinol based microporous polymeric organic frameworks with -OH functional groups and high CO₂ capture capacity. *Chem. Mater.* **23**, 1818–1824 (2011).
- Rabbani, M. G., Reich, T. E., Kassab, R. M., Jackson, K. T. & El-Kaderi, H. M. High CO₂ uptake and selectivity by triptycene-derived benzimidazole-linked polymers. *Chem. Commun.* **48**, 1141–1143 (2012).
- Rose, M., Bohlmann, W., Sabo, M. & Kaskel, S. Element-organic frameworks with high permanent porosity. *Chem. Commun.* 2462–2464 (2008).
- Mohanty, P., Kull, L. D. & Landskron, K. Porous covalent electron-rich organonitridic frameworks as highly selective sorbents for methane and carbon dioxide. *Nat. Commun.* **2**, 401–406 (2011).
- Patel, H. A. *et al.* High capacity carbon dioxide adsorption by inexpensive covalent organic polymers. *J. Mater. Chem.* **22**, 8431–8437 (2012).
- Yavuz, C. T. *et al.* Markedly improved CO₂ capture efficiency and stability of gallium substituted hydroxalcalites at elevated temperatures. *Chem. Mater.* **21**, 3473–3475 (2009).
- Rochelle, G. T. Amine scrubbing for CO₂ capture. *Science* **325**, 1652–1654 (2009).
- Zulficar, S. *et al.* Amidoximes: promising candidates for CO₂ capture. *Energy Environ. Sci.* **4**, 4528–4531 (2011).
- Keskin, S., van Heest, T. M. & Sholl, D. S. Can metal-organic framework materials play a useful role in large-scale carbon dioxide separations? *ChemSuschem* **3**, 879–891 (2010).
- Jin, Y. H. *et al.* Highly CO₂-selective organic molecular cages: what determines the CO₂ selectivity. *J. Am. Chem. Soc.* **133**, 6650–6658 (2011).
- Grirrane, A., Corma, A. & Garcia, H. Preparation of symmetric and asymmetric aromatic azo compounds from aromatic amines or nitro compounds using supported gold catalysts. *Nat. Protoc.* **5**, 429–438 (2010).
- Zhao, R. *et al.* One step synthesis of azo compounds from nitroaromatics and anilines. *Tetrahedron Lett.* **52**, 3805–3809 (2011).
- Saiz, L. M., Oyanguren, P. A. & Galante, M. J. Polyurethane-epoxy based azopolymers: influence of chemical structure over photoinduced birefringence. *React. Funct. Polym.* **72**, 478–485 (2012).
- Xie, S. A., Natansohn, A. & Rochon, P. Compatibility studies of some azo polymer blends. *Macromolecules* **27**, 1489–1492 (1994).
- Bandara, H. M. D. & Burdette, S. C. Photoisomerization in different classes of azobenzene. *Chem. Soc. Rev.* **41**, 1809–1825 (2012).
- McKeown, N. B. & Budd, P. M. Polymers of intrinsic microporosity (PIMs): organic materials for membrane separations, heterogeneous catalysis and hydrogen storage. *Chem. Soc. Rev.* **35**, 675–683 (2006).
- Viswanathan, N. K. *et al.* Surface relief structures on azo polymer films. *J. Mater. Chem.* **9**, 1941–1955 (1999).
- Uribe-Romo, F. J. *et al.* A crystalline imine-linked 3-D porous covalent organic framework. *J. Am. Chem. Soc.* **131**, 4570–4571 (2009).
- Tian, J., Thallapally, P. K., Dalgarno, S. J., McGrail, P. B. & Atwood, J. L. Amorphous molecular organic solids for gas adsorption. *Angew. Chem. Int. Ed.* **48**, 5492–5495 (2009).
- D'Alessandro, D. M., Smit, B. & Long, J. R. Carbon dioxide capture: prospects for new materials. *Angew. Chem. Int. Ed.* **49**, 6058–6082 (2010).
- Rabbani, M. G. & El-Kaderi, H. M. Template-free synthesis of a highly porous benzimidazole-linked polymer for CO₂ capture and H₂ storage. *Chem. Mater.* **23**, 1650–1653 (2011).
- Farha, O. K. *et al.* Synthesis, properties, and gas separation studies of a robust diimide-based microporous organic polymer. *Chem. Mater.* **21**, 3033–3035 (2009).
- Dawson, R., Stockel, E., Holst, J. R., Adams, D. J. & Cooper, A. I. Microporous organic polymers for carbon dioxide capture. *Energy Environ. Sci.* **4**, 4239–4245 (2011).
- Laybourn, A. *et al.* Branching out with amins: microporous organic polymers from difunctional monomers. *Polym. Chem.* **3**, 533–537 (2012).
- Dawson, R., Adams, D. J. & Cooper, A. I. Chemical tuning of CO₂ sorption in robust nanoporous organic polymers. *Chem. Sci.* **2**, 1173–1177 (2011).
- Ren, S. J. *et al.* Porous, fluorescent, covalent triazine-based frameworks via room-temperature and microwave-assisted synthesis. *Adv. Mater.* **24**, 2357–2361 (2012).

47. Jin, Y. H. *et al.* Microwave-assisted syntheses of highly CO₂-selective organic cage frameworks (OCFs). *Chem. Sci.* **3**, 874–877 (2012).
48. Drage, T. C. *et al.* Materials challenges for the development of solid sorbents for post-combustion carbon capture. *J. Mater. Chem.* **22**, 2815–2823 (2012).
49. Nagaraja, C. M., Haldar, R., Maji, T. K. & Rao, C. N. R. Chiral porous metal organic frameworks of Co(II) and Ni(II): synthesis, structure, magnetic properties, and CO₂ uptake. *Cryst. Growth Des.* **12**, 975–981 (2012).
50. Garcia-Ricard, O. J., Silva-Martinez, J. C. & Hernandez-Maldonado, A. J. Systematic evaluation of textural properties, activation temperature and gas uptake of Cu₂(pzdc)₂L [L = dipyriddy-based ligands] porous coordination pillared-layer networks. *Dalton Trans.* **41**, 8922–8930 (2012).
51. Khaliullin, R. Z., Cobar, E. A., Lochan, R. C., Bell, A. T. & Head-Gordon, M. Unravelling the origin of intermolecular interactions using absolutely localized molecular orbitals. *J. Phys. Chem. A* **111**, 8753–8765 (2007).
52. Shao, Y. *et al.* Advances in methods and algorithms in a modern quantum chemistry program package. *Phys. Chem. Chem. Phys.* **8**, 3172–3191 (2006).
53. Harris, J. G. & Yung, K. H. Carbon dioxide's liquid-vapor coexistence curve and critical properties as predicted by a simple molecular model. *J. Phys. Chem.* **99**, 12021–12024 (1995).

Acknowledgements

This work was made possible by NPRP grants # 08-670-1-124 and # 5-499-1-088 from the Qatar National Research Fund (a member of Qatar Foundation). The statements made herein are solely the responsibility of the authors. C.T.Y. and Y.J. acknowledge the financial support by grants from Korea CCS R&D Centre, IWT (NRF-2012-C1AAA001-

M1A2A2026588), and WCU programme (R31-2008-000-10055-0) funded by the Ministry of Education, Science and Technology of Korean government, and KAIST EEWs Initiative. This research was also supported by the KUSTAR-KAIST Institute, Korea, under the R&D programme supervised by the KAIST.

Author contributions

H.A.P., C.T.Y. and A.C. designed the study. H.A.P., J.P., Y.J., C.T.Y. and A.C. analysed the data and wrote the paper. D.P.C and S.H.J. prepared monomers. H.A.P. and S.H.J. synthesized and analysed porous polymers. J.P. and Y.J. performed the theoretical analyses and interpreted the anomalous temperature-dependent selectivity increase. All authors discussed the results and commented on the manuscript.

Additional information

Supplementary Information accompanies this paper at <http://www.nature.com/naturecommunications>

Competing financial interests: The authors declare no competing financial interests.

Reprints and permission information is available online at <http://npg.nature.com/reprintsandpermissions/>

How to cite this article: Patel, H.A. *et al.* Unprecedented high-temperature CO₂ selectivity in N₂-phobic nanoporous covalent organic polymers. *Nat. Commun.* **4**:1357 doi: 10.1038/ncomms2359 (2013).