



Cite this: DOI: 10.1039/d4ta08839e

Unravelling the origin of enhanced CO₂ selectivity in amine-PIM-1 during mixed gas permeation†Carmen Rizzuto,^{†a} Francesca Nardelli,^{†§b} Marcello Monteleone,^{†a} Lucia Calucci,^{†*bc} C. Grazia Bezzu,^{†d} Mariolino Carta,^{†d} Elena Tocci,^{†*a} Elisa Esposito,^{†a} Giorgio De Luca,^{†a} Bibiana Comesaña-Gándara,^{†e} Neil B. McKeown,^{†f} Bekir Sayginer,^{†g} Peter M. Budd,^{†h} Johannes C. Jansen^{†a} and Alessio Fuoco^{†*a}

Previously, it has been reported that amine-PIM-1, a polymer of intrinsic microporosity obtained by reduction of nitrile groups of PIM-1 to primary amine groups, shows enhanced CO₂ selectivity during mixed gas permeation studies with respect to single gas measurements for gas pairs involving CO₂. This distinct and potentially useful behaviour was ascribed to the affinity of CO₂ for the polymer amine groups. Here, we demonstrate that enhanced selectivity originates from both CO₂ physisorption and chemisorption. A combination of ¹³C and ¹⁵N solid-state NMR spectroscopic analyses of a CO₂-loaded amine-PIM-1 membrane allowed the identification and quantitative determination of both chemisorbed and physisorbed species and the characterization of polymer-CO₂ interactions. Experiments with ¹³C isotopically enriched CO₂ unequivocally demonstrated the conversion of 20% of the NH₂ groups into carbamic acids at 298 K and a CO₂ pressure of 1 bar. Chemisorption was supported by the strong heat of CO₂ adsorption for amine-PIM-1 that was estimated as 50 kJ mol⁻¹. Molecular dynamics simulations with models based on the experimentally determined polymer structure gave a detailed description of intra- and interchain hydrogen bond interactions in amine-PIM-1 after chemisorption, as well as of the effect of chemisorption on polymer porosity and physisorption.

Received 12th December 2024
Accepted 23rd April 2025

DOI: 10.1039/d4ta08839e

rsc.li/materials-a

1. Introduction

Global warming poses an existential threat to humanity and Earth's ecosystem. This crisis demands urgent and transformative action to reduce greenhouse gas emissions and

mitigate their catastrophic impact.¹ Carbon capture and storage (CCS) is recognised as one of the most promising technologies capable of delivering net emissions reductions at the scale required to limit global temperature rise to below 2 °C.² The intergovernmental panel on climate change estimates that CCS will need to contribute about one-sixth of the required CO₂ emission reductions by 2050.³ CCS is particularly important for reducing emissions from industrial processes such as steel, cement, and chemicals production, which cannot be easily replaced by renewable technologies. These industries account for around 45% of the CO₂ captured between 2015 and 2050 in a 2 °C scenario.³ The success of carbon capture depends on developing and optimising materials capable of selectively capturing, storing, and eventually releasing CO₂. These materials must demonstrate high selectivity, capacity, and stability across various conditions.⁴ Key materials studied for carbon capture include adsorbents (*e.g.*, metal-organic frameworks (MOFs) and activated carbon), which possess high surface area and porosity, enabling them to “physically” trap large quantities of CO₂.^{5,6} Adsorbents are also widely used for carbon capture. Typically, they are composed of amine-based solutions that can chemically react with CO₂ to form a compound that can be later decomposed. Both these classes of materials are often functionalized to enhance their affinity for CO₂.⁷

^aInstitute on Membrane Technology (CNR-ITM), National Research Council of Italy, via P. Bucci 17/C, Rende, CS, 87036, Italy. E-mail: alessio.fuoco@cnr.it; e.tocci@itm.cnr.it

^bInstitute of Chemistry of Organometallic Compounds (ICCOM-CNR), via G. Moruzzi 1, Pisa, 56124, Italy. E-mail: lucia.calucci@pi.iccom.cnr.it

^cCenter for Instrument Sharing, University of Pisa (CISUP), Lungarno Pacinotti 43/44, Pisa, 56126, Italy

^dDepartment of Chemistry, Faculty of Science and Engineering, Swansea University, Swansea, SA2 8PP, UK

^eIU CINQUIMA, University of Valladolid, Paseo Belén 5, Valladolid, 47011, Spain

^fEaStCHEM, School of Chemistry, University of Edinburgh, Joseph Black Building, David Brewster Road, Edinburgh, Scotland, EH9 3JF, UK

^gDepartment of Chemistry, Faculty of Arts and Sciences, Kırsehir Ahi Evran University, Kırsehir, 40100, Türkiye

^hDepartment of Chemistry, School of Natural Sciences, The University of Manchester, Manchester, M13 9PL, UK

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4ta08839e>

‡ These two authors contributed equally.

§ Present address: Department of Chemistry and Industrial Chemistry, University of Pisa, via G. Moruzzi 13, 56124 Pisa, Italy.

Membranes for gas separation provide an alternative and potentially more energy-efficient technology for CCS.^{8,9} Permeable and selective materials can separate CO₂ from gas mixtures through a combination of molecular sieving, realized through tailored pore sizes, and affinity for CO₂, often achieved by incorporating carefully selected functional groups.^{10–12} However, developing membrane materials with enhanced sieving capabilities and affinity for CO₂ is required to fulfil the promise of more efficient and cost-effective CO₂ capture.^{11,13}

Polymers of intrinsic microporosity (PIMs) are a class of amorphous glassy polymers characterised by a rigid and contorted molecular structure that inhibits efficient chain packing in the solid state. This typically results in the formation of high free-volume elements.¹⁴ The unique combination of microporosity and processability makes PIMs excellent candidate materials for energy storage, catalysis, and environmental applications.^{15–17} In particular, PIMs show potential as membranes for gas separation, with important applications such as carbon dioxide removal from natural gas (CO₂/CH₄) and flue gas (CO₂/N₂).^{18–20} Their molecular structures, combined with the potential to incorporate various functional groups into the polymeric backbone, enable PIMs to achieve a remarkable trade-off between permeability and selectivity that allows them to define and even surpass the Robeson upper bounds for various gas pairs.²¹ The continuous evolution and development of synthetic strategies has recently culminated in the publication of a series of PIMs used to establish the new upper bounds for CO₂-based gas separations.²² PIMs performance can be further enhanced through strategies like chemical modification,²³ crosslinking,²⁴ and incorporation of MOFs and COFs for the efficient formation of mixed matrix membranes.^{25,26}

PIM-1, the archetypal polymer of intrinsic microporosity and the best-studied,²⁷ possesses two nitrile groups per repeat unit, which can be reduced to primary amine groups that have been shown to improve selectivity for Lewis acidic gases such as CO₂.²⁸ In our previous work, we found that amine-PIM-1 exhibits remarkably low ideal gas selectivity for the CO₂/N₂ and CO₂/CH₄ gas pairs.^{28,29} We attributed this behaviour to specific interactions between the functionalized polymer and CO₂ on the basis of the unusual trend of the gas diffusion coefficients as a function of the squared effective diameter.³⁰ In a more recent paper, the group led by Smith at MIT published intriguing research on functionalised PIM-1.³¹ They studied six different versions of PIM-1, including amine-PIM-1, where nitrile groups were converted into various functional groups to tune the backbone's affinity towards CO₂. The permselectivity

data were assessed using both single and mixed gases (with increasing CO₂ concentration from 10% to 90% for CO₂/CH₄). It was found that, in some cases, the performance with mixed gases improved compared to that predicted by the single gases, especially after physical aging. This improvement was attributed to a mechanism known as “competitive gas sorption”, which allows the more permeable gas (CO₂) to pass more favourably through the membrane, while the less permeable gas (either CH₄ or N₂) is blocked. An enhanced solubility-selectivity arises from the high affinity of the polymeric backbone for CO₂, leading to an overall improvement in CO₂/CH₄ and CO₂/N₂ mixed-gas permselectivity. Especially important proved to be the conversion of the nitrile to amine groups to obtain amine-PIM-1 (or PIM-NH₂), which showed an exceptional improvement of the selectivity with mixed gases up to 140% for CO₂/CH₄ and 250% for CO₂/N₂.

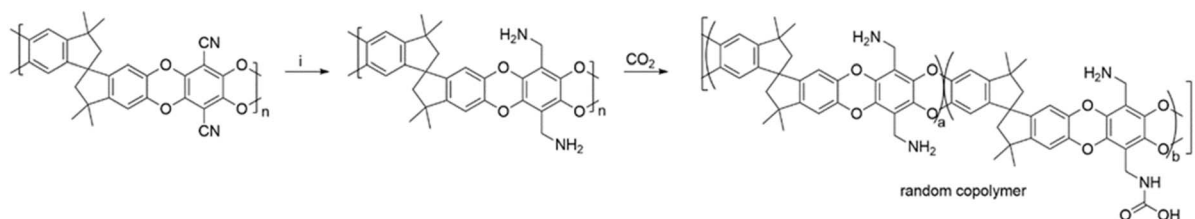
Here, we confirm the enhanced CO₂ selectivity of amine-PIM-1 during mixed gas permeation, and we provide an in-depth experimental and computational study that explains the enhanced affinity of this polymer for CO₂. By exploiting solid-state nuclear magnetic resonance (SSNMR), a powerful technique for identifying and quantifying physisorbed and chemisorbed CO₂ species in amine-functionalized solid sorbents,^{32–37} we demonstrate that CO₂ is partially chemisorbed on amine-PIM-1, forming carbamic acids by reaction with primary amine groups on the membrane surface (Scheme 1). Partial CO₂ chemisorption is also supported by isosteric heat of adsorption. Molecular dynamics (MD) simulations on chemisorbed amine-PIM-1 models, prepared with a number of amine groups converted into carbamic acid groups as determined by SSNMR measurements, emphasize the complexity and critical role of hydrogen bonding in influencing the material's properties and interactions.

2. Experimental

2.1 Materials

¹³C labelled (99.0 atom %) carbon dioxide (¹³CO₂) was purchased from Sigma Aldrich.

Single gases for permeation measurements were supplied by Pirossigeno at a minimum purity of 99.9995%. Certified gas mixtures for permeation measurements were supplied by Sapio at a purity of ±0.01% from the certified concentration (CO₂/CH₄ mixture with 52.11 mol% CO₂ and 47.89 mol% CH₄, and N₂/O₂ mixture with 81.45 mol% N₂ and 18.55 mol% O₂).



Scheme 1 Post-modification of PIM-1 to give amine-PIM-1 and formation of chemisorbed-amine-PIM-1 following CO₂ loading. (i) borane dimethyl sulphide 5.0 M in diethyl ether.



2.2 Membrane preparation

The membranes for the permeation measurements were prepared as reported previously,²⁹ while the powder for BET analysis and the membranes for SSNMR were prepared by a slightly different procedure to guarantee a conversion of nitrile to amine groups more than 95%, herein reported. Reduction of nitrile groups of PIM-1 membranes was carried out using borane-dimethyl sulphide complex in diethyl ether, a non-solvent for the PIM-1 precursor that enables the preparation of amine-PIM-1 membrane samples. Thus, PIM-1 membranes were placed in Petri dishes and then covered with 5.0 M borane-dimethyl sulphide complex in diethyl ether. Subsequently, the Petri dishes were lidded and, in turn, stowed inside a closed plastic container at room temperature to induce and maintain a saturated atmosphere. After one day of reaction time, membranes were collected and placed in Petri dishes of ethanol to remove the excess of borane-dimethyl sulphide complex. Subsequently, membranes were firstly soaked in Petri dishes of 1.0 M methanolic HCl overnight and then in Petri dishes of 5% aqueous NaOH solution for 3 hours. Finally, membranes were rinsed repeatedly in plenty of water and soaked in methanol for 24 h before being dried at 323 K under vacuum. The molecular structure and physical properties of the synthesized materials were confirmed by TGA, XRD and IR measurements (ESI Fig. S1–S3†).

2.3 Gas adsorption measurements

N₂ (77 K and 298 K) and CO₂ (273 K, 298 K, and 308 K) adsorption/desorption measurements of polymer powders were made using an Anton Paar Nova 600. Samples were degassed over 8 h at 80 °C under high vacuum prior to analysis. The gases were supplied by BOC (N₂ purity > 99.999%, CO₂ purity > 99.995%). The data were analysed using the software Anton Paar Kaomi for NOVA, which is provided with the instrument. The Brunauer–Emmett–Teller (BET) surface area was calculated from N₂ adsorption isotherms at a relative pressure $P/P_0 < 0.1$. Non-local density functional theory (NLDFT) analysis on CO₂ adsorption isotherms was performed to calculate the pore size distribution and volume, considering a carbon equilibrium transition kernel at 273 K based on a slit-pore model; the kernel is based on a common, one centre, Lennard-Jones model. Heats of adsorption were calculated from the CO₂ curves measured at 273 K, 298 K, and 308 K. The data were analysed with the Anton Paar Kaomi software and fitted with the Langmuir–Freundlich equation and heat was calculated *via* the Clausius–Clapeyron equation.

2.4 Gas permeation measurements

Pure and mixed gas permeation tests were carried out from 1 to 6 bar of feed pressure using a custom-made constant pressure/variable volume instrument, equipped with a quadrupole mass filter (HPR-20 QIC, Hiden Analytical). The experimental set-up and the procedures are described elsewhere.^{38,39} For mixed gas permeation tests, certified mixtures of CO₂/CH₄ (52.11/47.89 vol%), and N₂/O₂ (81.45/18.55 vol%) were used. The

measurements were performed on an amine-PIM-1 membrane with 85% conversion and aged for 750 days to guarantee the membrane stability during the measurements. Aging of the membranes took place during their natural storage under atmospheric conditions between the individual measurements.

2.5 SSNMR measurements

SSNMR experiments were carried out on a Bruker Avance NEO 500 spectrometer working at the Larmor frequency of 500.13, 125.77, and 50.68 MHz, for ¹H, ¹³C, and ¹⁵N, respectively, equipped with a 4 mm double-channel (H/F-X) CP/MAS probe. ¹³C chemical shifts were referenced to the signal of adamantane at 38.48 ppm and calculated for the other nuclei using the unified scale recommended by IUPAC.⁴⁰ All measurements were conducted at 298 K and under magic angle spinning (MAS) conditions at a frequency of 15 kHz using air as spinning gas, unless otherwise specified.

¹³C quantitative direct excitation (DE) MAS spectra were acquired by applying high-power proton decoupling with a recycle delay of 100 s and 16 scans for the sample loaded with ¹³CO₂ and a recycle delay of 60 s, using a flip angle of 60° and accumulating about 1600 scans, for the sample loaded with CO₂.

¹H–¹³C cross-polarization (CP) MAS experiments were carried out using a recycle delay of 2 s and accumulating 128–288 scans for the sample loaded with ¹³CO₂ and 1000 scans for the unloaded sample and the sample loaded with CO₂. Contact time values ranging from 0.5 to 8 ms were employed. ¹H–¹³C CP MAS experiments were performed at different temperatures ranging from 263 to 303 K.

¹H–¹⁵N CP MAS experiments were recorded using a recycle delay of 2 s and a contact time of 4 ms, accumulating 1600 scans; for these experiments the MAS frequency was 8 kHz.

For the sample loaded with ¹³CO₂, the following experiments were also recorded. ¹H–¹³C HETCOR experiments with FSLG homonuclear decoupling⁴¹ were acquired using a recycle delay of 2 s and contact time values from 0.5 to 3 ms, accumulating 16 scans for each of the 80 increments in the indirect dimension. ¹³C DE spectra were recorded with a MAS frequency of 2.5 kHz without applying proton decoupling, using a recycle delay of 1 s, and accumulating 6400 scans.

The analysis of the spinning side bands pattern of ¹³C DE MAS spectra recorded with the MAS frequency of 2.5 kHz was performed using the solid line shape analysis module (SOLA) included in the software Topspin by Bruker.

Samples for SSNMR measurements were prepared using a home-made cell provided with a mechanical lever operated from outside enabling the capping of the rotor without disturbing the cell atmosphere. In particular, the activated sample was prepared by heating overnight under vacuum (0.1 mbar) at the temperature of 423 K. The membrane was packed into the NMR rotor (4 mm external diameter) and the rotor was capped under N₂ atmosphere. For the CO₂-loaded samples, the activated membrane was loaded with either CO₂ or ¹³CO₂ (*i.e.* ¹³C isotopically enriched CO₂) at 1 bar pressure and the rotor was capped under the gas atmosphere after the equilibrium was reached.



2.6 Molecular modelling

Molecular dynamics simulations were performed using the BIOVIA software package.⁴² The COMPASS (condensed-phase optimized molecular potentials for atomistic simulations studies) force field^{43,44} was used, according to previous studies on PIM-1-like materials.^{29,45–48}

Molecular dynamics simulations were performed on amine-PIM-1 and on chemisorbed-amine-PIM-1.

The amine-PIM-1 polymer chain for the initial packing with the amorphous cell module consisted of 30 monomers (1860 atoms). Each chain contained 60 amines (see Scheme 1). Every packing model contained three polymer chains. A total of 5580 atoms were grown in a 3D model under periodic boundary conditions. Details on the preparation of the two simulation boxes can be found in our previous paper.²⁹

For the simulations of the chemisorbed-amine-PIM-1, a chain of 30 monomer units is constructed with 12 carbamic acid moieties in accordance with the SSNMR measurements. The chain was grown with a torsion angle of 180° in an amorphous cell module. The simulation boxes contain three chains of amorphous polymer (5790 atoms), 300 argon atoms, and the number of CO₂ corresponding to the adsorbate molecules that have not reacted with the amine groups of the polymer at the pressure of 1 bar. Two simulation boxes were selected and equilibrated following a procedure based on three steps: (1) deleting 100 Ar atoms for each step, (2) anneal dynamics run over the 300–700 K temperature range, (3) NPT-MD simulations (constant number of particles (*N*), pressure (*P*), and temperature (*T*)) at 1 bar and 298 K for 100 ps. The electrostatic and van der Waals interactions were studied using the group-based method⁴⁹ with a cut-off distance of 12.5 Å. The unit cell parameters of the 3D-triclinic lattice were $a = b = c = 41.78$ Å, $\alpha = \beta = \gamma = 90^\circ$, cell volume = 72 962.9 Å³. NPT-MD simulations were performed at 298 K and 1 bar for a simulation time of 20 ns with a timestep of 1 fs. The initial velocities were randomly evaluated. Temperature and pressure were controlled using the Berendsen method.⁵⁰ We have inserted 120 carbon dioxide (CO₂) molecules into the simulation cells for the physisorbed system, corresponding to the amount adsorbed at 1 bar, as determined by experimental data. For the chemisorbed system, 36 CO₂ molecules are incorporated into the polymer chains, while the remaining 90 molecules are free within the simulation box.

The theoretical adsorption isotherms were evaluated by the grand canonical Monte-Carlo (GCMC) configurational bias method,⁵¹ implemented in the NVT ensemble (constant number of particles (*N*), volume (*V*), and temperature (*T*)) within the 0–5 bar fugacity range at 298 K.⁴²

The fractional free volume (FFV) was calculated according to the Bondi equation (eqn (1)).⁵²

$$\text{FFV} = \frac{V_{\text{tot}} - 1.3V_{\text{vdW}}}{V_{\text{tot}}} \quad (1)$$

where V_{tot} is the volume of the simulation box, V_{vdW} is the van der Waals volume calculated with the group contribution method, and the 1.3 factor is the packing coefficient used to convert the van der Waals volume to the occupied volume.

2.7 Radial distribution function

The radial distribution function, $g_{AB}(r)$, between particles of *A* and *B* is defined as follows (eqn (2)):

$$g_{AB}(r) = \frac{\langle g_B(r) \rangle}{\langle g_B \rangle_{\text{local}}} = \frac{1}{\langle g_B \rangle_{\text{local}}} \frac{1}{N_A} \sum_{i \in A} \sum_{j \in B} \frac{\delta(r_{ij} - r)}{4\pi r^2} \quad (2)$$

where $g_B(r)$ is the particle density of type *B* at a distance *r* around particles *A*, and $g_{B \text{ local}}$ is the particle density of type *B* averaged over all spheres around particles *A* having radius r_{max} . The radial distribution function is employed to evaluate the specific interactions between the CO₂ molecules and the polymer groups.

2.8 Calculated BET surface area

In the computational BET surface determination, the gas surface accessible area was determined using N₂ as probe having Teplyakov-Mearns' radius (1.52 Å).⁵³ The procedure calculates the accessible area described by the probe centre as it rolls over a scaled van der Waals surface. The gas surface accessible areas were also calculated using the Teplyakov-Mearns' radii for O₂ (1.44 Å), CO₂ (1.51 Å), and CH₄ (1.59 Å).

2.9 Pore size distribution analysis

The pore size distributions were determined using a geometry-based analysis on porous materials with a Voronoi decomposition of the space and lattice precision of 0.1 Å implemented in Zeo++ code.^{54,55} The pore size distributions were calculated by exploring the available void spaces with a probe of zero radius.

3. Results and discussion

For amine-PIM-1, the pure and mixed gas permeabilities for the O₂/N₂ gas pair are closely correlated, which is consistent with the relatively low and similar solubility of both gases in the polymer. In both cases, a weak but significant decrease in permeability as a function of pressure is observed (ESI Fig. S4†), typical of samples exhibiting dual mode behaviour. The selectivity is slightly higher and the nitrogen permeability slightly lower in the gas mixture with respect to the pure gases, probably because of a small degree of competitive permeation. In contrast, as shown in Fig. 1a and b, the pressure dependence of CO₂ and CH₄ permeation properties in amine-PIM-1 exhibits a dramatic difference between the pure and the mixed gas measurements. This difference is much higher with respect to the O₂/N₂ gas pair for the same polymer (ESI Fig. S4†), as well as to the CO₂/CH₄ gas pair in the parent polymer PIM-1 (ESI Fig. S5†). In the literature, this behaviour is known as sorption-enhanced mixed gas transport³¹ and can be ascribed to specific CO₂ affinity with the amine functional groups.

Thus, isothermal gas adsorption studies were conducted on PIM-1 and amine-PIM-1 to evaluate the difference in their CO₂ uptake and to better understand the increased affinity of this gas for the aminated PIM. Previous studies^{28,29} showed that converting the nitrile groups of PIM-1 into primary amine groups leads to stronger inter-chain hydrogen bonding and



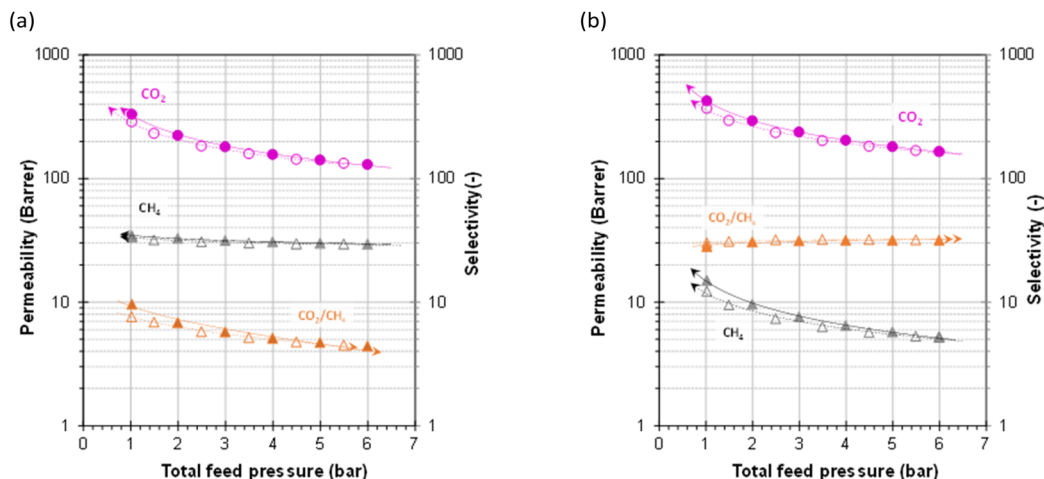


Fig. 1 Pressure dependence of permeability and selectivity in amine-PIM-1 for (a) pure CO₂ and CH₄, and (b) for a CO₂/CH₄ mixture (52.11/47.89 vol%). Filled symbols represent the pressure-increase steps and open symbols represent the subsequent pressure-decrease steps.

a drastic reduction in free volume, slowing down N₂ adsorption kinetics at 77 K, typically used to assess the porosity and surface area of porous materials.⁵⁶ However, our repeated studies show that amine-PIM-1 exhibits substantial nitrogen adsorption at 77 K, from which an apparent BET surface area (S_{BET}) of 645 m² g⁻¹ can be calculated. Although lower than that of PIM-1 (S_{BET} ~750 m² g⁻¹), this value indicates that there is significant intrinsic microporosity available for the physisorption of CO₂ (ESI Fig. S6†).

CO₂ adsorption studies further revealed that, despite its lower porosity, amine-PIM-1 has significantly higher CO₂ uptake than PIM-1 (ESI Fig. S6b†), confirming that converting nitriles into amines does indeed improve CO₂ affinity. The pore size distribution (PSD) analysis, calculated *via* NLDFT from CO₂ adsorption at 273 K, shows values for both polymers (peaks at 3.5–8.5 Å) that are typical for a PIM. Notably, the peak centred around 3.5 Å, indicating greater apparent ultra-microporosity, is much more intense for amine-PIM-1 than for PIM-1 (ESI Fig. S6c†), as expected for the greater CO₂ adsorption at low pressure. CO₂ uptake at different temperatures (273, 298, and 308 K) (ESI Fig. S7†) allowed the calculation of the isosteric heat of adsorption (Q_{st}), which is crucial for understanding CO₂ affinity and assessing adsorption mechanism in PIM-1 and amine-PIM-1 (ESI Fig. S6d†). Interestingly, the Q_{st} for amine-PIM-1 (~50 kJ mol⁻¹) is almost twice that of PIM-1 (~27 kJ mol⁻¹). Physisorption in microporous materials is generally governed by van der Waals interactions with enthalpy values typically ranging between 10 and 50 kJ mol⁻¹,⁵⁷ while chemisorption involves stronger, often covalent or ionic interactions, leading to enthalpy values above this range,⁵⁶ and thus a Q_{st} value of 50 kJ mol⁻¹ marks an estimated boundary between physisorption and chemisorption.⁵⁸ Thus, the herein calculated value for amine-PIM-1 indicates that CO₂ chemisorption occurs to some extent, similarly to what previously reported for an amine functionalized adsorbent⁵⁹ and a MOF.⁶⁰

This is further confirmed by the presence of weak hysteresis in the adsorption/desorption isotherms at all measured

temperatures (ESI Fig. S8†). Thus, while physisorption may still be accounted as the primary adsorption mechanism, a non-negligible amount of CO₂ is chemisorbed in the polymer matrix. To identify, quantify, and characterise physisorbed and chemisorbed CO₂ species, various 1D and 2D SSNMR experiments were recorded on amine-PIM-1 membrane before and after its loading with CO₂ or ¹³CO₂ (1 bar at RT). The comparison of ¹³C magic angle spinning (MAS) spectra shown in Fig. 2 highlights the presence of physisorbed CO₂ (peak at 125 ppm) and two different chemisorbed species (peaks at 158 and 161 ppm). Signals for these species are indeed present only in the spectra of the membrane loaded with CO₂ or ¹³CO₂ (Fig. 2b–d), together with the signals arising from the different membrane carbons,^{28,29} also observed in the spectrum of the membrane not exposed to CO₂ (Fig. 2a). The signals with isotropic chemical

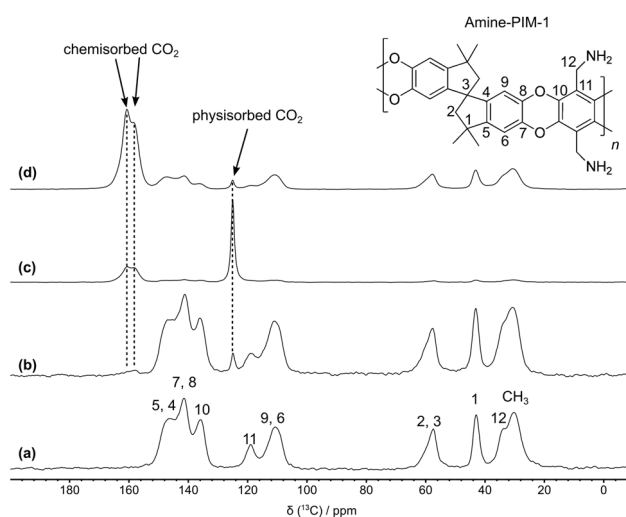


Fig. 2 Quantitative ¹³C DE MAS spectra of: (a) amine-PIM-1; (b) amine-PIM-1 loaded with CO₂; (c) amine-PIM-1 loaded with ¹³CO₂. (d) ¹H-¹³C CP MAS spectrum of amine-PIM-1 loaded with ¹³CO₂ recorded with a contact time of 2 ms. All the spectra were acquired at 298 K at a spinning frequency of 15 kHz.



shift (δ_{iso}) values of 158 and 161 ppm, typical of carbamic acid species differing in hydrogen bonding environment,⁶¹ clearly indicate that, at ambient temperature and atmospheric pressure, CO₂ has access to and reacts with amine groups of amine-PIM-1. The integral areas of the membrane and CO₂ carbon's peaks in the quantitative ¹³C direct excitation (DE) spectra of the CO₂- or ¹³CO₂-loaded membrane (Fig. 2b and c) allowed an estimate of the physisorbed and chemisorbed species: overall these species are present in a 1:1 molar ratio with the polymeric unit of amine-PIM-1, which is in agreement with the finding from CO₂ uptake at 298 K (2 mmol g⁻¹), while the molar ratio between physisorbed and chemisorbed species is 1:0.7 (ESI Fig. S9†). These results indicate that almost one out of five amine groups of amine-PIM-1 reacts with CO₂ to form chemisorbed species, as sketched in Scheme 1.

As far as the identification of the chemisorbed species is concerned, the chemical shift values measured in the ¹³C spectra point to carbamic acid species differing in hydrogen bonding environment,⁶¹ although they are not decisive for a sound assignment of the signals, since δ_{iso} values between 153 and 168 ppm have been reported in the literature for both carbamic acid and ammonium carbamate carbons,^{33,62} two species that can form in amine-PIM-1 if CO₂ reacts with one or two amine groups, respectively.⁶³ Therefore, to undoubtedly identify the formed species, a combination of different experiments was adopted, including the acquisition of ¹H-¹⁵N cross polarization (CP) MAS spectra and 2D ¹H-¹³C HETCOR spectra, as well as the determination of the chemical shift tensor components for the two carbons resonating at 158 and 161 ppm.

Despite the low signal-to-noise ratio arising from the low natural abundance of ¹⁵N (0.36%), the ¹H-¹⁵N CP MAS spectrum of CO₂-loaded amine-PIM-1 clearly shows a signal at 83 ppm ascribable to the nitrogen of the amidic NH group of carbamic acids or carbamates, and a signal at 24 ppm, also observed in the spectrum of the pristine amine-PIM-1 membrane, associated to unreacted amine groups (Fig. 3).^{33,61,64,65} However, no signal is observed at 32–36 ppm,

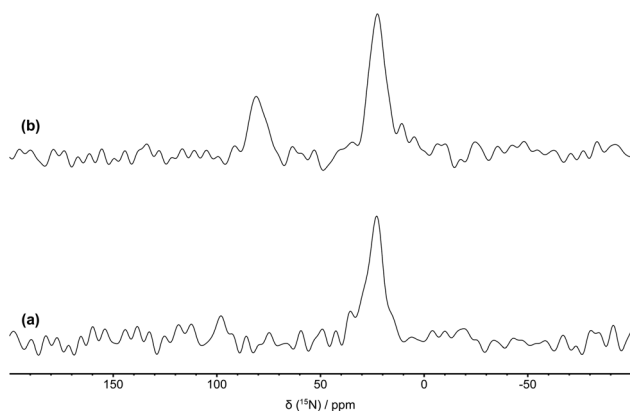


Fig. 3 ¹H-¹⁵N CP MAS spectra recorded with a contact time of 4 ms on: (a) pristine amine-PIM-1 membrane and (b) amine-PIM-1 membrane loaded with CO₂ (i.e., chemisorbed-amine-PIM-1). The spectra were acquired at 298 K at a spinning frequency of 8 kHz.

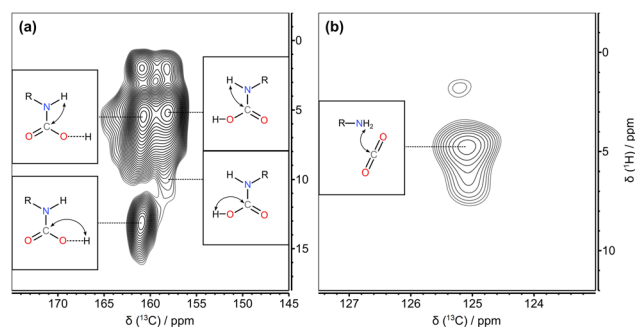


Fig. 4 Expansion of the ¹H-¹³C HETCOR MAS spectra of chemisorbed-amine-PIM-1 loaded with ¹³CO₂ recorded with a contact time of (a) 0.5 ms and (b) 1.5 ms.

where the nitrogen of ammonium groups is expected to resonate,^{33,61,64,65} suggesting that only carbamic acid species are formed.

The 2D ¹H-¹³C HETCOR spectrum recorded on the ¹³CO₂-loaded amine-PIM-1 membrane with a short contact time of 0.5 ms (Fig. 4a and ESI Fig. S12a†) shows cross peaks between the ¹³C nucleus resonating at 158 ppm and ¹H nuclei resonating at 5.2 (NH) and 10.4 (COOH) ppm, and between the ¹³C with δ_{iso} = 161 ppm and ¹H nuclei resonating at 5.4 (NH) and 13.2 (COOH) ppm. The observed differences, especially that concerning the COOH ¹H chemical shift, can be accounted for by a different arrangement of two forms of carbamic acid, the higher chemical shift value being associated to a form with the COOH group involved in a strong H-bond.

The attribution of the two chemisorbed species to carbamic acids involved or not/weakly involved in H-bonding was corroborated by the symmetry of the chemical shift tensors of the COOH carbons determined through the analysis of MAS spectra recorded at slow spinning frequency. According to Čendak *et al.*,⁶⁶ an axial tensor is expected for protonated carbamic acid not/weakly involved in H-bonds, while a change towards an orthorhombic tensor is observed for carbamic acids involved in strong H-bonds and for the deprotonated carbamic acid (carbamate).⁶⁷ Here, the principal components of the chemical shift tensors of the carbons resonating at 158 and 161 ppm were reconstructed from the analysis of the side band manifolds in the ¹³C MAS spectrum recorded at a MAS frequency of 2.5 kHz (Fig. 5). The chemical shift tensor of the species with δ_{iso} = 158 ppm has a nearly axial symmetry with principal components δ_{11} = 220 ppm, δ_{22} = 129 ppm, and δ_{33} = 125 ppm. On the other hand, the chemical shift tensor of the species with δ_{iso} = 161 ppm has an orthorhombic tensor with principal components δ_{11} = 212 ppm, δ_{22} = 157 ppm, and δ_{33} = 114 ppm; the tensor components were expressed using the Mehring notation, where $\delta_{11} \geq \delta_{22} \geq \delta_{33}$. From the tensor's principal components, values of A ($A = \delta_{11} + \delta_{33} - \delta_{22}$) and d (with $d = \delta_{11} - \delta_{\text{iso}}$ when $|\delta_{11} - \delta_{\text{iso}}| > |\delta_{33} - \delta_{\text{iso}}|$ and $d = \delta_{33} - \delta_{\text{iso}}$ when $|\delta_{11} - \delta_{\text{iso}}| < |\delta_{33} - \delta_{\text{iso}}|$) were also determined. A is 216 and 169 ppm, and d is 62 and 51 ppm for carbons with δ_{iso} of 158 and 161 ppm, respectively. According to findings by Gu and McDermott on amino acids⁶⁷ and by Čendak *et al.* on carbamic



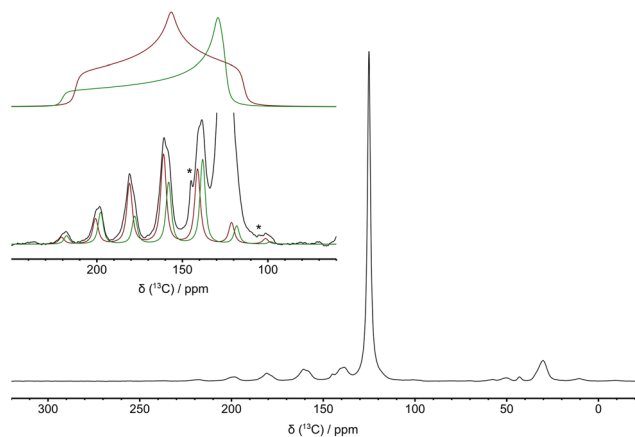


Fig. 5 ^{13}C DE MAS spectrum of amine-PIM-1 loaded with $^{13}\text{CO}_2$ recorded using a MAS frequency of 2.5 kHz. In the inset, simulations of the side band manifolds of signals of carbamic acids with δ_{iso} of 158 and 161 ppm (bottom) and the corresponding reconstructed chemical shift tensors (top) are shown in green and red, respectively. Asterisks indicate spinning side bands of physisorbed CO_2 undergoing anisotropic motion.

acids/carbamates,⁶⁶ these parameters are associated to two protonated forms of carbamic acid, one not (or weakly) involved in H-bonds ($\delta_{\text{iso}} = 158$ ppm) and the other involved in strong H-bonds ($\delta_{\text{iso}} = 161$ ppm). Overall, the SSNMR data indicate that two types of carbamic acids form in the reaction of CO_2 with amine groups of amine-PIM-1 in dry conditions, one strongly involved and one not (or weakly) involved in hydrogen bonding with a hydrogen bond acceptor.

Further insights into hydrogen bonds between polymer moieties were achieved through MD calculations. Simulations on chemisorbed-amine-PIM-1 revealed complex inter- and intramolecular hydrogen bonding networks within and between the polymer chains, between amine/amine, amine/dioxin, and carbamic acid/dioxin groups (Fig. 6a–d). This confirms that the carbamic acids formed in the amine-PIM-1 are involved in different kinds of hydrogen bonds.

The intra- and inter-chain interactions play a pivotal role in the membrane's architecture, its free volume, and structural stabilization, and they commonly decrease the permeability and enhance the membranes' selectivity. Such packing effects were previously noted in PIM-2 (ref. 68) and various post-synthetically modified PIMs.^{69–75} However, to the best of our knowledge, this is the first time that this packing effect is deeply studied in the absence or presence of a molecule that changes the molecular structure of the polymer. The chemisorption of CO_2 into amine-PIM-1, in the form of newly incorporated carbamic acid moieties, affects its morphology, leading to decreased spacing between chains in comparison to “pristine” amine-PIM-1. This results in a reduction of the BET surface area and the fractional free volume (FFV).

In fact, simulations on amine-PIM-1 molecular boxes calculate a BET surface area of $732.6 \pm 0.8 \text{ m}^2 \text{ g}^{-1}$, while a much lower BET surface area, *i.e.* $602.6 \pm 0.2 \text{ m}^2 \text{ g}^{-1}$, is calculated using the chemisorbed-amine-PIM-1 molecular model, showing

that the presence of the carbamic acid has a strong influence on the polymer network, and specifically leads to a reduction of the FFV. The herein simulated FFV of amine-PIM-1 is 0.28 ± 0.02 , which is larger with respect to that calculated for the chemisorbed-amine-PIM-1 (0.25 ± 0.01), again showing that the extra formed hydrogen bonds are leading to a lower FFV. This is further confirmed by the analysis of the accessible surface area of N_2 , O_2 , CO_2 , and CH_4 (Table S1†), which shows that all the gases experience a lower accessible surface area in chemisorbed-amine-PIM-1 with respect to amine-PIM-1.

MD simulations reveal that amine-PIM-1 exhibits a broad range of pore sizes, from 1.1 to 14.1 Å, with predominant diameters between 6.0 and 6.1 Å (Fig. 6e). In contrast, chemisorbed-amine-PIM-1 shows a narrower distribution, ranging from 0.6 to 9.7 Å, with the most prevalent pore sizes around 4.3–4.4 Å (Fig. 6f), indicating a notable presence of smaller voids within the polymer structure. Considering that the simulated distributions capture a wider range of pore sizes due to the use of a zero-sized probe, results of both amine-PIM-1 and chemisorbed-amine-PIM-1 are consistent with the experimental pore size distribution analysis, calculated *via* NLDFT from CO_2 adsorption at 273 K, which highlights peaks between 3.5 and 8.5 Å (ESI Fig. S6c†). The chemisorbed-amine-PIM-1 model, with its predominant pore sizes around 4.3–4.4 Å, aligns more closely with the experimental values than the amine-PIM-1 model, confirming that CO_2 chemisorption and the formation of carbamic acid significantly alter the polymer's structure. The free volume and slice visualizations of chemisorbed-amine-PIM-1 and amine-PIM-1 are reported in ESI Fig. S10 and S11.†

^1H - ^{13}C CP and HETCOR MAS spectra were also recorded on the $^{13}\text{CO}_2$ -loaded amine-PIM-1 membrane to highlight interactions between physisorbed CO_2 and membrane moieties. In the ^1H - ^{13}C CP MAS spectra (Fig. 2 and ESI Fig. S13†) the signal of physisorbed CO_2 carbon at 125 ppm is observed together with signals from the membrane and carbamic acids' carbons at all contact time values employed (from 0.5 to 8 ms), indicating that adsorbed CO_2 establishes interactions with the membrane hydrogens. In particular, the signal of physisorbed CO_2 steadily increases with the contact time up to the longest interval explored (8 ms), suggesting that the magnetization transfer from ^1H to ^{13}C nuclei progressively occurs in time from hydrogens in molecular fragments increasingly distant from CO_2 . Correspondingly, a correlation peak is observed between the signal of physisorbed CO_2 and the ^1H signal of interacting amine groups at 4.7 ppm in the ^1H - ^{13}C HETCOR MAS spectrum recorded with a contact time of 1.5 ms, while correlation peaks are observed with all membrane protons for a longer contact time of 3 ms (ESI Fig. 12b and c†). This evidence indicates that stronger dipolar interactions are established between CO_2 ^{13}C and $-\text{NH}_2$ ^1H nuclei, which are ascribable to a closer proximity of CO_2 to amine groups in the membrane due to Lewis's acid–base interactions.

However, a comparison of the relative intensities of the signals from chemisorbed and physisorbed species in the quantitative ^{13}C DE MAS spectra and in the CP MAS spectra (Fig. 2 and ESI Fig. S13†) suggests that only one part of



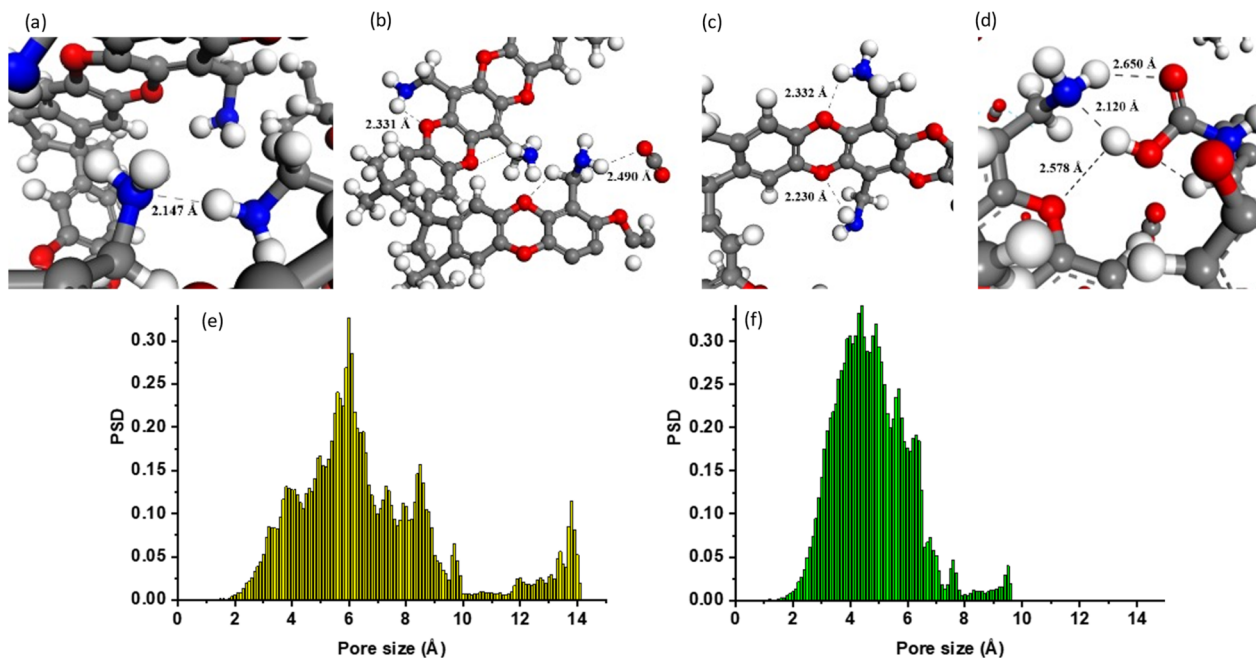


Fig. 6 Detailed snapshots of the interactions between (a) amine/amine, (b) dioxin/ $\text{CH}_2\text{-NH}_2$, (c) amine/dioxin, and (d) amine/carbamic acid groups observed after the simulation time of 20 ns. In chemisorbed-amine-PIM-1, the H atom is white, the O atom is red, the N atom is blue, and the C atom is grey (settings: maximum hydrogen-acceptor distance 2.5 Å, minimum donor-hydrogen-acceptor angle 90°). Average pore size distribution of (e) amine-PIM-1 and (f) chemisorbed-amine-PIM-1, both calculated by exploring the available void spaces with a probe of zero radius.

physisorbed CO_2 can get in close proximity to the membrane and establish dipolar interactions strong enough to allow magnetization transfer. The remaining physisorbed CO_2 is located further from the membrane surface and/or undergoes fast dynamics. This can also be seen in the DE MAS spectrum recorded at the spinning frequency of 2.5 kHz shown in Fig. 5 where both features of CO_2 undergoing isotropic (sharp peak at 125 ppm) and restricted anisotropic motions (side bands marked with asterisks) inside the membrane are observed at RT. However, a detailed analysis of the anisotropic CO_2 signal is hampered by the very low intensity of the sidebands. The observed behavior is different from that reported for $^{13}\text{CO}_2$ -loaded PIM-1, tetrazole- (TZ-PIM), and methyl tetrazole-functionalized (MTZ-PIM) PIM-1, for which only an isotropic line is observed for CO_2 in ^{13}C static spectra at RT.⁷⁶ Indeed, for PIM-1 an isotropic line shape resulting from the averaging of the CO_2 ^{13}C CSA by translational hopping of CO_2 among randomly oriented sites was already observed at 100 K, while it occurred at slightly higher temperatures for TZ-PIM (125 K) and MTZ-PIM (150 K). Our findings indicate stronger interactions of a part of physisorbed CO_2 with amine-PIM-1 adsorption sites with respect to PIM-1 and tetrazole-functionalized PIM-1 membranes, which hamper fast isotropic motion in the membrane pores even at room temperature.

^{13}C CP MAS spectra recorded at different temperatures between 263 and 303 K show a progressive decrease of the chemisorbed CO_2 signal intensities (ESI Fig. S14†). Although these spectra are not quantitative, this observation could be ascribed to the decrease of chemisorbed species by increasing

the temperature. These trends can be related to the decrease in CO_2 -polymer affinity and Langmuir site sorption capacity observed in a previous work,²⁹ as well as to CO_2 adsorption isotherms measured at different temperatures in the present work (ESI Fig. S8†).

Furthermore, the MD simulations indicated that physisorbed CO_2 molecules establish hydrogen bonds with both the unreacted amine groups and carbamic acid groups of chemisorbed-amine-PIM-1. These interactions are identified using radial distribution functions, $g(r)$, between the oxygen atoms of CO_2 and selected atoms of the polymer (Fig. 7a). The first peak of the green curve obtained for the unreacted amine group, centered at 1.67 Å, indicates a strong direct hydrogen bond between the hydrogen of the unreacted amine group and the closest oxygen of CO_2 , while the second broad peak centered at 3.97 Å (Fig. 7a), and displayed at 3.81 Å in a snapshot (Fig. 7c), indicates the interaction between the second oxygen of the same CO_2 molecule and the hydrogen of the amine group. The narrow first peaks of the curves related to -NH (blue) and -OH (red) of the carbamic acid are found at approximately 1.85–2.02 Å, while two broad and more intense peaks are observed, centered at 3.97 Å for the -NH and 5.07 Å for the -OH groups. The -NH of the unreacted amine groups (green curve) is in full overlap with the -NH of the carbamic acid, showing comparable patterns, while the -OH is at a close distance, helping to form a hydrogen bond network around CO_2 . In fact, all together the observed $g(r)$ peaks correspond to weak hydrogen bonds between CO_2 and different chains and/or different sections of the same chain in the chemisorbed-amine-PIM-1, indicating an



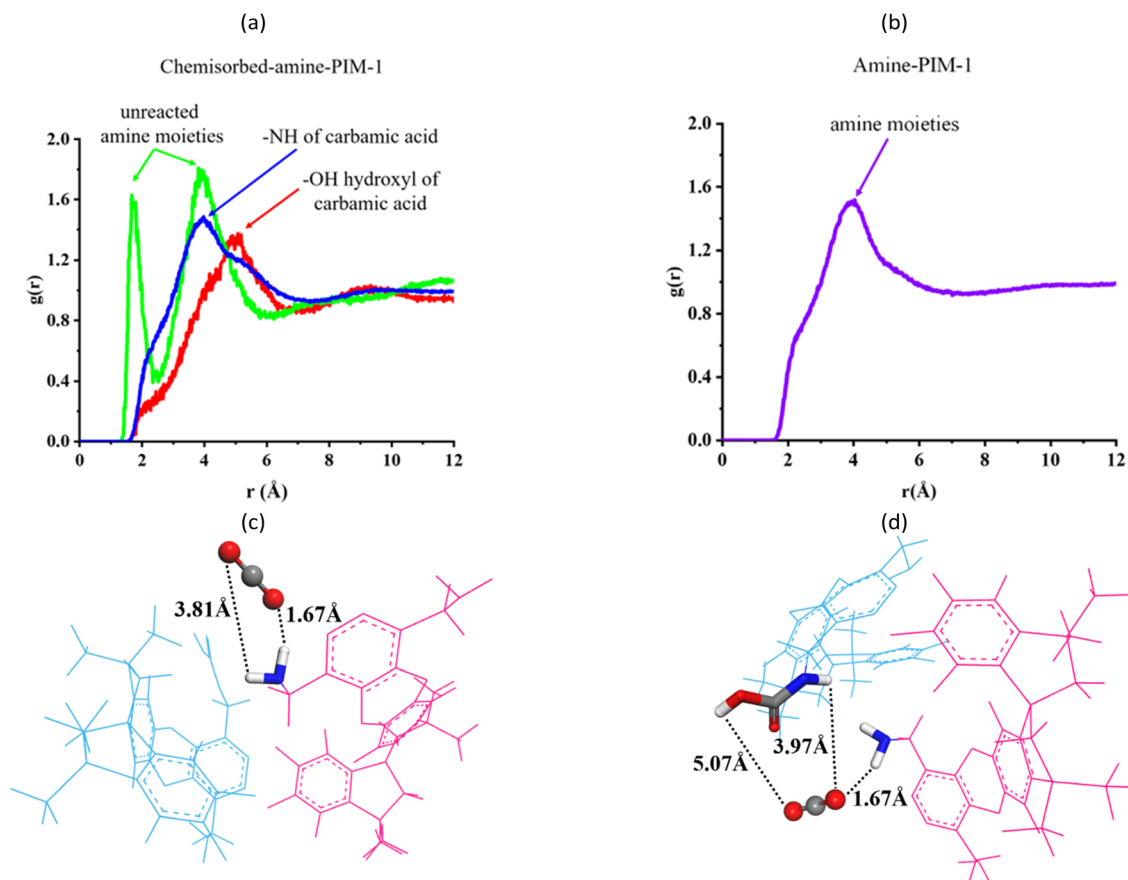


Fig. 7 (a) Radial distribution function between the O atoms of CO₂ and: the H atoms of the unreacted amine groups (green curve), the H atom of -NH- group of carbamic acid (blue curve), and the H atom of -OH group of carbamic acid (red curve) in chemisorbed-amine-PIM-1. (b) Radial distribution function between the O atoms of CO₂ and the H atoms of the amine groups of amine-PIM-1 (violet curve). (c) Indicative hydrogen bonds between the O atoms of CO₂ and the H atoms of unreacted amine groups corresponding to the two peaks of the green curve in a. (d) Examples of hydrogen bonds between the O atoms of CO₂ and one H atom of unreacted amine (1.67 Å), the H atom of -NH- group (blue curve in a) of carbamic acid (3.97 Å), and the H atom of the -OH group (red curve in a) of carbamic acid (5.07 Å). The different polymeric chains are indicated in light blue and pink. Colour code: the C atom is grey, the O atom is red, the H atom is white, and the N atom is blue.

organized structure hosting CO₂ resembling a “pseudo-pocket”. Despite being located approximately 4–5 Å away from the sites, in accordance with the ¹H–¹³C HETCOR MAS spectra (Fig. 4 and ESI Fig. 12†), CO₂ coordinates simultaneously with both carbamic acid and unreacted amine groups. These results suggest that, on average, CO₂ moves with restricted mobility when coordinated to capture sites, while uncoordinated CO₂ has greater degree of freedom. This is confirmed by SSNMR studies, which reveal that CO₂ undergoes both anisotropic motions, reflecting restricted mobility, and isotropic motions, indicating greater freedom.

A different situation is observed in models of amine-PIM-1 with physisorbed CO₂, where only unreacted amine groups are present in the simulation box. In the $g(r)$ function (Fig. 7b), a subtle first peak appears at 2.03 Å, indicating a specific interaction distance between the amine groups and CO₂ molecules. A broader and more intense peak at 3.95 Å suggests a secondary interaction distance. Unlike the chemisorbed case, the first peak is not clearly distinguishable. Detailed analysis of our simulations shows that the number of CO₂ molecules corresponds to an experimental pressure of 1 bar. At this

pressure, the total number of CO₂ molecules is 120, significantly higher than the 90 molecules used in the chemisorbed system (the remaining molecules being chemisorbed).

Due to the large number of CO₂ molecules in the polymeric model, the interaction weakens with the formation of multiple shells, moving further from the membrane surface and undergoing faster dynamics, as confirmed by ¹³C DE MAS and CP MAS spectra. The strong hydrogen bonding with unreacted amine hydrogens is not visually apparent but rather emerges as a statistical effect. Recent analyses of physisorbed systems at different pressures and temperatures have revealed a strong relationship between the amine groups and the C=O bonds of CO₂ molecules, particularly at pressures below 1 bar.²⁹ As pressure and temperature increase, the $g(r)$ distribution decreases, indicating a weaker interaction. At higher pressures, the interaction diminishes due to the large number of CO₂ molecules in the polymeric model, causing the amine groups to “lose sight” of them due to faster CO₂ dynamics. Similarly, at higher temperatures, the interaction between the amine groups and CO₂ weakens as vibrational modes intensify.



4. Conclusions

Our study confirmed the enhanced CO₂ selectivity of amine-PIM-1 in mixed gas with respect to pure gas permeation for gas pairs involving CO₂ and provided sound evidence of the strong CO₂ affinity for this polymer through a combination of experimental and computational data. Adsorption isotherms revealed that amine-PIM-1 is a microporous polymer with higher CO₂ adsorption capacity with respect to the archetypal PIM-1, and a considerably higher isosteric heat of adsorption (about 50 kJ mol⁻¹ for amine-PIM-1 vs. 27 kJ mol⁻¹ for PIM-1), which suggests the presence of chemisorption. Indeed, chemisorbed species were detected and quantified by combining different SSNMR experiments. Two carbamic acid species were found, characterized by different strength of hydrogen bond interactions with the polymer moieties, accounting for about 40% of the detected CO₂ at 1 bar of pressure at RT, and resulting from the reaction of CO₂ with about 20% of amine groups in dry conditions. The number of chemisorbed species was found to decrease by increasing the temperature. Remarkably, our study reveals that chemisorption occurs also in the absence of water, while the latter is fundamental to enhance the CO₂ permselectivity in most facilitated transport membranes. Physisorbed CO₂ was also revealed and characterized by SSNMR spectroscopy. Populations of CO₂ in more and less restricted environments were found, undergoing anisotropic and isotropic dynamics, respectively. Molecular dynamics simulations supported both isothermal adsorption and SSNMR findings. Through the analysis of models for amine-PIM-1 before and after CO₂ chemisorption (chemisorbed-amine-PIM-1), a detailed description of the interactions between polymer moieties was provided. A complex network of hydrogen bonds was found involving amine/amine, amine/dioxin, and amine/carbamic acid groups, which results in a decreased surface area and a narrower pore size distribution for chemisorbed-amine-PIM-1. Moreover, different states corresponding to different interactions with amine and carbamic acid moieties were identified for physisorbed CO₂. Coordination of CO₂ in a pseudo-pocket environment was found in chemisorbed-amine-PIM-1 where CO₂ strongly interacts with unreacted amine groups, indicating that chemisorption promotes physisorption with stronger interactions. In contrast to more common facilitated transport membranes that need water to favour CO₂ transport by carbamate formation, the formation of carbamic acid under dry conditions in the voids of microporous polymers may be an interesting strategy to develop novel, highly selective membranes for CO₂ capture. To summarize, while both sorption-driven affinity (physisorption/chemisorption) and pore-size modifications contribute to CO₂ selectivity, *Q_{st}* data and mixed-gas permeation trends highlight a synergistic relationship, where sorption-driven affinity is the predominant factor in CO₂ uptake, while pore-size modifications *via* hydrogen bonding enhance overall selectivity by restricting CH₄ and N₂ permeability. This structural refinement reinforces CO₂ permselectivity rather than directly increasing CO₂ uptake. The combined influence of enhanced CO₂ affinity and tailored

polymer microstructure leads to the observed improvement in CO₂-based gas separations. The elucidation of the transport properties in amine-PIM-1 may be relevant to the development of next generation membrane materials with enhanced CO₂-philic behaviour.

Data availability

The data supporting this article have been included as part of the ESI† and are available at Zenodo at <https://zenodo.org/communities/dam4co2/>.

Author contributions

CR: conceptualization, data curation, formal analysis, investigation, methodology, resources, visualization, writing – original draft, writing – review & editing; FN: conceptualization, data curation, formal analysis, investigation methodology, resources, visualization, writing – original draft, writing – review & editing; MM: formal analysis, investigation, validation, visualization, writing – original draft; LC: conceptualization, formal analysis, methodology, project administration, supervision, writing – original draft, writing – review & editing; CGB: formal analysis, investigation, methodology, resources, writing – original draft, writing – review & editing; MC: conceptualization, formal analysis, funding acquisition, resources, supervision, writing – original draft, writing – review & editing; ET: conceptualization, data curation, formal analysis, methodology, project administration, supervision, writing – review & editing; EE: investigation, validation, writing – review & editing; GDL: validation, writing – review & editing; BCG: investigation, validation, writing – review & editing; NBM: funding acquisition, resources, supervision, writing – review & editing; BS: formal analysis, investigation, resources, writing – review & editing; PMB: funding acquisition, methodology, supervision, writing – review & editing; JCJ: formal analysis, methodology, project administration, supervision, writing – review & editing; AF: conceptualization, data curation, formal analysis, funding acquisition, methodology, project administration, resources, supervision, writing – original draft, writing – review & editing.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Acknowledgements

The authors are gratefully acknowledging the Ministry for Universities and Research (MUR) of Italy for financial support under the program PRIN 2020 under the project “doMino” (2020P9KBKZ) and the Grants TED2021-131170A-I00 and CNS2022-135430 funded by MICIU/AEI/10.13039/501100011033 and by the European Union NextGenerationEU/PRTR. The research leading to these results has received funding from the European Union's Horizon Europe research and innovation



programme under grant agreement No 101115488, project DAM4CO₂ and by UK Research and Innovation (UKRI) under the UK government's Horizon Europe funding guarantee, grant numbers 10083164 and 10091537.

References

- 1 T. M. Letcher, *The Root Causes of Global Warming and the New Normal*, Elsevier, 2024.
- 2 W. F. Lamb, T. Gasser, R. M. Roman-Cuesta, G. Grassi, M. J. Gidden, C. M. Powis, O. Geden, G. Nemet, Y. Pratama, K. Riahi, S. M. Smith, J. Steinhauser, N. E. Vaughan, H. B. Smith and J. C. Minx, *Nat. Clim. Change*, 2024, **14**, 644–651.
- 3 A. C. Ruane, <https://www.nationalgrid.com/stories/energy-explained/what-is-ccs-how-does-it-work>, accessed June 2024.
- 4 R. L. Siegelman, E. J. Kim and J. R. Long, *Nat. Mater.*, 2021, **20**, 1060–1072.
- 5 B. Chen, D. Fan, R. V. Pinto, I. Dovgaliuk, S. Nandi, D. Chakraborty, N. García-Moncada, A. Vimont, C. J. McMonagle, M. Bordonhos, A. Al Mohtar, I. Cornu, P. Florian, N. Heymans, M. Daturi, G. De Weireld, M. Pinto, F. Nouar, G. Maurin, G. Mouchaham and C. Serre, *Adv. Sci.*, 2024, **11**, 1–15.
- 6 A. Koli, A. K. Battu, R. K. Motkuri and S. Sabale, *Biomass Convers. Biorefin.*, 2024, **14**, 10177–10188.
- 7 X. Y. D. Soo, J. J. C. Lee, W. Y. Wu, L. Tao, C. Wang, Q. Zhu and J. Bu, *J. CO₂ Util.*, 2024, **81**, 1–33.
- 8 E. Favre, R. Bounaceur and D. Roizard, *J. Membr. Sci.*, 2009, **328**, 11–14.
- 9 K. Xie, Q. Fu, G. G. Qiao and P. A. Webley, *J. Membr. Sci.*, 2019, **572**, 38–60.
- 10 Z. Dai and L. Deng, *Sep. Purif. Technol.*, 2024, **335**, 1–27.
- 11 A. Junaidi, U. Zulfiani, S. Khomariyah, T. Gunawan, N. Widiastuti, N. Sazali and W. N. W. Salleh, *RSC Adv.*, 2024, **14**, 2311–2319.
- 12 P. Sherugar, A. M. Antony, N. A. H. M. Nordin, S. A. Patil and M. Padaki, *Fuel*, 2024, **361**, 1–13.
- 13 D. B. Gutierrez, E. B. Caldon, R. D. Espiritu and R. C. Advincula, *MRS Commun.*, 2021, **11**, 391–401.
- 14 P. M. Budd, K. J. Msayib, C. E. Tattershall, B. S. Ghanem, K. J. Reynolds, N. B. McKeown and D. Fritsch, *J. Membr. Sci.*, 2005, **251**, 263–269.
- 15 A. R. Antonangelo, N. Hawkins, E. Tocci, C. Muzzi, A. Fuoco and M. Carta, *J. A. Chem. Soc.*, 2022, **144**, 15581–15594.
- 16 M. O. Amin, E. Al-Hetlani, A. R. Antonangelo, H. Zhou and M. Carta, *Appl. Water Sci.*, 2023, **13**, 1–11.
- 17 F. Topuz, M. H. Abdellah, P. M. Budd and M. A. Abdulhamid, *Polym. Rev.*, 2024, **64**, 251–305.
- 18 X. Feng, J. Zhu, J. Jin, Y. Wang, Y. Zhang and B. Van der Bruggen, *Prog. Mater. Sci.*, 2024, **144**, 1–56.
- 19 H. Zhou, C. Rayer, A. R. Antonangelo, N. Hawkins and M. Carta, *ACS Appl. Mater. Interfaces*, 2022, **14**, 20997–21006.
- 20 Y. Li, J. Brückel, M. Jereb, A. Zupanc, S. P. Hirvonen, S. Hietala, M. Kemell, Y. Wu, O. Fuhr, R. D. Jansen-van Vuuren, M. Carta and S. Bräse, *Adv. Funct. Mater.*, 2024, **2401957**, 1–8.
- 21 Y. Wang, B. S. Ghanem, Y. Han and I. Pinnau, *Curr. Opin. Chem. Eng.*, 2022, **35**, 1–9.
- 22 B. Comesaña-Gándara, J. Chen, C. G. Bezzu, M. Carta, I. Rose, M. C. Ferrari, E. Esposito, A. Fuoco, J. C. Jansen and N. B. McKeown, *Energy Environ. Sci.*, 2019, **12**, 2733–2740.
- 23 M. Tian, S. Rochat, H. Fawcett, A. D. Burrows, C. R. Bowen and T. J. Mays, *Adsorption*, 2020, **26**, 1083–1091.
- 24 T. Joo, T. H. Lee, S. J. Kaser, W. N. Wu, S. Wi, J. Y. Yeo and Z. P. Smith, *Chem. Mater.*, 2024, **36**, 4275–4290.
- 25 E. Esposito, M. Carta, A. Fuoco, M. Monteleone, B. Comesaña-Gándara, E. Gkaniatsou, C. Sicard, S. Wang, C. Serre, N. B. McKeown and J. C. Jansen, *J. Membr. Sci.*, 2024, **697**, 1–14.
- 26 F. Emamverdi, J. Huang, N. M. Razavi, M. J. Bojdys, A. B. Foster, P. M. Budd, M. Böhning and A. Schönhals, *Macromolecules*, 2024, **57**, 1829–1845.
- 27 P. M. Budd, B. S. Ghanem, S. Makhseed, N. B. McKeown, K. J. Msayib and C. E. Tattershall, *Chem. Commun.*, 2004, **4**, 230–231.
- 28 C. R. Mason, L. Maynard-Atem, K. W. J. Heard, B. Satilmis, P. M. Budd, K. Friess, M. Lanci, P. Bernardo, G. Clarizia and J. C. Jansen, *Macromolecules*, 2014, **47**, 1021–1029.
- 29 B. Satilmis, M. Lanč, A. Fuoco, C. Rizzuto, E. Tocci, P. Bernardo, G. Clarizia, E. Esposito, M. Monteleone, M. Dendisová, K. Friess, P. M. Budd and J. C. Jansen, *J. Membr. Sci.*, 2018, **555**, 483–496.
- 30 A. Fuoco, C. Rizzuto, E. Tocci, M. Monteleone, E. Esposito, P. M. Budd, M. Carta, B. Comesaña-Gándara, N. B. McKeown and J. C. Jansen, *J. Mater. Chem. A*, 2019, **7**, 20121–20126.
- 31 K. Mizrahi Rodriguez, F. M. Benedetti, N. Roy, A. X. Wu and Z. P. Smith, *J. Mater. Chem. A*, 2021, **9**, 23631–23642.
- 32 L. Mafra, T. Čendak, S. Schneider, P. V. Wiper, J. Pires, J. R. B. Gomes and M. L. Pinto, *J. Am. Chem. Soc.*, 2017, **139**, 389–408.
- 33 R. Afonso, M. Sardo, L. Mafra and J. R. B. Gomes, *Environ. Sci. Technol.*, 2019, **53**, 2758–2767.
- 34 D. Pereira, R. Fonseca, I. Marin-Montesinos, M. Sardo and L. Mafra, *Curr. Opin. Colloid Interface Sci.*, 2023, **64**, 1–19.
- 35 S. M. Pugh and A. C. Forse, *J. Magn. Reson.*, 2023, **346**, 1–11.
- 36 M. Ilkaeva, R. Vieira, J. M. P. Pereira, M. Sardo, I. Marin-Montesinos and L. Mafra, *J. Am. Chem. Soc.*, 2023, **145**, 8764–8769.
- 37 M. Sardo, T. Morais, M. Soares, R. Vieira, M. Ilkaeva, M. A. O. Lourenço, I. Marín-Montesinos and L. Mafra, *Chem. Commun.*, 2024, **60**, 4015–4035.
- 38 M. Monteleone, E. Esposito, A. Fuoco, M. Lanč, K. Pilnáček, K. Friess, C. Bezzu, M. Carta, N. McKeown and J. C. Jansen, *Membranes*, 2018, **8**, 73.
- 39 S. C. Fraga, M. Monteleone, M. Lanč, E. Esposito, A. Fuoco, L. Giorno, K. Pilnáček, K. Friess, M. Carta, N. B. McKeown, P. Izák, Z. Petrusová, J. G. Crespo, C. Brazinha and J. C. Jansen, *J. Membr. Sci.*, 2018, **561**, 39–58.
- 40 R. K. Harris, E. D. Becker, S. M. C. De Menezes, R. Goodfellow and P. Granger, *Solid State Nucl. Magn. Reson.*, 2002, **22**, 458–483.



- 41 B.-J. Van Rossum, H. Förster, F. Förster and H. J. M. De Groot, *J. Magn. Reson.*, 1997, **124**, 516–519.
- 42 Accelrys Software Inc., *P. U. G. P. S, BIOVIA Package; Ex Material Studio 7.0; Classical Simulation Theory Section; Sorption User Guide*, Accelrys Software Inc., San Diego, CA, USA, 2013.
- 43 H. Sun, *J. Phys. Chem. B*, 1998, **102**, 7338–7364.
- 44 H. Sun, P. Ren and J. R. Fried, *Comput. Theor. Chem.*, 1998, **8**, 229–246.
- 45 K.-S. Chang, K.-L. Tung, Y.-F. Lin and H.-Y. Lin, *RSC Adv.*, 2013, **3**, 10403.
- 46 L. Zhao, D. Zhai, B. Liu, Z. Liu, C. Xu, W. Wei, Y. Chen and J. Gao, *Chem. Eng. Sci.*, 2011, **68**, 101–107.
- 47 L. Zhang, W. Fang and J. Jiang, *J. Phys. Chem. C*, 2011, **115**, 11233–11239.
- 48 O. Hölck, M. Böhning, M. Heuchel, M. R. Siegert and D. Hofmann, *J. Membr. Sci.*, 2013, **428**, 523–532.
- 49 T. A. Halgren, *J. Am. Chem. Soc.*, 1992, **114**, 7827–7843.
- 50 H. J. C. Berendsen, J. P. M. Postma, W. F. Van Gunsteren, A. Dinola and J. R. Haak, *J. Chem. Phys.*, 1984, **81**, 3684–3690.
- 51 R. L. C. Akkermans, N. A. Spenley and S. H. Robertson, *Mol. Simul.*, 2013, **39**, 1153–1164.
- 52 A. Bondi, *J. Chem. Phys.*, 1964, **68**, 442–451.
- 53 V. Teplyakov and P. Meares, *Sep. Purif. Technol.*, 1990, **4**, 66–74.
- 54 T. F. Willems, C. H. Rycroft, M. Kazi, J. C. Meza and M. Haranczyk, *Microporous Mesoporous Mater.*, 2012, **149**, 134–141.
- 55 M. Pinheiro, R. L. Martin, C. H. Rycroft, A. Jones, E. Iglesia and M. Haranczyk, *J. Mol. Graph. Model.*, 2013, **44**, 208–219.
- 56 M. Thommes, K. Kaneko, A. V. Neimark, J. P. Olivier, F. Rodriguez-Reinoso, J. Rouquerol and K. S. W. Sing, *Pure Appl. Chem.*, 2015, **87**, 1051–1069.
- 57 M. M. Dubinin, *Chem. Rev.*, 1960, **2**, 235–241.
- 58 L. W. Bruch, *Surf. Sci.*, 1983, **125**, 194–217.
- 59 W. Si, S. Ye, D. Zhang, B. Yang, Y. Hou, Z. Li, X. Zhang, J. Zhu and L. Lei, *Can. J. Chem. Eng.*, 2019, **97**, 697–701.
- 60 D. D. Zhou, X. W. Zhang, Z. W. Mo, Y. Z. Xu, X. Y. Tian, Y. Li, X. M. Chen and J. P. Zhang, *Energy Chem.*, 2019, **1**, 1–35.
- 61 R. W. Flaig, T. M. Osborn Popp, A. M. Fracaroli, E. A. Kapustin, M. J. Kalmutzki, R. M. Altamimi, F. Fathieh, J. A. Reimer and O. M. Yaghi, *J. Am. Chem. Soc.*, 2017, **139**, 12125–12128.
- 62 M. L. Pinto, L. Mafra, J. M. Guil, J. Pires and J. Rocha, *Chem. Mater.*, 2011, **23**, 1387–1395.
- 63 K. Basaran, B. U. Topcubasi and T. Davran-Candan, *J. CO₂ Util.*, 2021, **47**, 1–16.
- 64 C. H. Chen, D. Shimon, J. J. Lee, S. A. Didas, A. K. Mehta, C. Sievers, C. W. Jones and S. E. Hayes, *Environ. Sci. Technol.*, 2017, **51**, 6553–6559.
- 65 D. Shimon, C. H. Chen, J. J. Lee, S. A. Didas, C. Sievers, C. W. Jones and S. E. Hayes, *Environ. Sci. Technol.*, 2018, **52**, 1488–1495.
- 66 T. Čendak, L. Sequeira, M. Sardo, A. Valente, M. L. Pinto and L. Mafra, *Chem.-Eur. J.*, 2018, **24**, 10136–10145.
- 67 Z. Gu and A. McDermott, *J. Am. Chem. Soc.*, 1993, **115**, 4282–4285.
- 68 A. Fuoco, B. Satilmis, T. Uyar, M. Monteleone, E. Esposito, C. Muzzi, E. Tocci, M. Longo, M. P. De Santo, M. Lanč, K. Friess, O. Vopička, P. Izák and J. C. Jansen, *J. Membr. Sci.*, 2020, **594**, 1–12.
- 69 K. Mizrahi Rodriguez, A. X. Wu, Q. Qian, G. Han, S. Lin, F. M. Benedetti, H. Lee, W. S. Chi, C. M. Doherty and Z. P. Smith, *Macromolecules*, 2020, **53**, 6220–6234.
- 70 R. Swaidan, B. S. Ghanem, E. Litwiller and I. Pinnau, *J. Membr. Sci.*, 2014, **457**, 95–102.
- 71 J. W. Jeon, D. G. Kim, E. H. Sohn, Y. Yoo, Y. S. Kim, B. G. Kim and J. C. Lee, *Macromolecules*, 2017, **50**, 8019–8027.
- 72 N. Du, H. B. Park, G. P. Robertson, M. M. Dal-Cin, T. Visser, L. Scoles and M. D. Guiver, *Nat. Mater.*, 2011, **10**, 372–375.
- 73 J. Weber, N. Du and M. D. Guiver, *Macromolecules*, 2011, **44**, 1763–1767.
- 74 C. R. Mason, L. Maynard-Atem, N. M. Al-Harbi, P. M. Budd, P. Bernardo, F. Bazzarelli, G. Clarizia and J. C. Jansen, *Macromolecules*, 2011, **44**, 6471–6479.
- 75 N. Du, G. P. Robertson, J. Song, I. Pinnau and M. D. Guiver, *Macromolecules*, 2009, **42**, 6038–6043.
- 76 J. K. Moore, R. M. Marti, M. D. Guiver, N. Du, M. S. Conradi and S. E. Hayes, *J. Phys. Chem. C*, 2018, **122**, 4403–4408.

