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Using intermolecular interactions to crosslink PIM-1 and modify its gas sorption properties†

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The intermolecular interactions between the “polymer of intrinsic microporosity” PIM-1 and polycyclic aromatic hydrocarbons (PAHs) have been investigated with the aim of modifying the gas sorption and physical properties. Mixing PIM-1 with selected PAHs resulted in rapid precipitation of polymer. Blending PIM-1 with pyrene had a significant effect of the gas sorption properties of the resulting films; dramatically reduced N₂ uptake (77 K), whilst CO₂ uptake at 298 K was only slightly reduced. A gate-opening behaviour was also observed for the N₂ gas sorption (77 K), which was related to the pyrene content of the blend. Using an electron-donating PAH as the additive resulted in a stronger interaction. By exploiting a post-modification strategy after PIM-1 film formation, the absorption of either pyrene or 1-aminopyrene produced films with higher elastic moduli and greatly improved CO₂/N₂ gas sorption selectivities (293 K). Single gas permeability measurements revealed that while the 1-aminopyrene modified film possessed reduced CO₂ permeability, it possessed enhanced CO₂/N₂ selectivity. Importantly, the ageing of the permeability was halted over the 50 days tested, likely due to the physical crosslinking of the polymer chains by 1-aminopyrene.

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Introduction

Polymers of intrinsic microporosity (PIMs) have been the subject of extensive research due to their solution-processability and microporous nature.^{1,2} These polymers possess high free volumes mainly in the microporous range, which is attributed to the contortion of the rigid backbone about a spiro-carbon centre, leading to inefficient packing of the polymer chains.³ A range of PIMs have been prepared utilising an assortment of different monomers¹ but PIM-1 (Fig. 1), the first example of its type, is still regarded as one of the most promising materials for gas separation.⁴ PIM-1 is a linear polymer formed by the

condensation of 5,5,6,6-tetrahydroxy-3,3,3,3-tetramethyl-1,1-spirobisindane with tetrafluoroterephthalonitrile.⁵ This polymer is soluble in a number of common volatile solvents⁶ offering easy solution processing into films and has been the subject of a great deal of studies.^{4,7–17} PIM-1 is interesting for applications in gas separation membranes due to its high permeability and moderate selectivity.¹⁸ Recently, a range of approaches have been published showing a further enhancement of the selectivity of membranes based on PIM-1. These modification approaches generally fall into two categories: modification of the polymer backbone, or crosslinking PIM-1 to form a network. Modifications of the polymer backbone include the hydrolysis of the nitriles to carboxylic acids,⁸ synthesis of analogues of PIM-1 by substituting the nitrile group on the tetrafluoroterephthalonitrile monomer with different sulfone-based functionalities,⁷ converting the nitrile groups of PIM-1 to thioamide groups,¹¹ amidoxime groups¹⁹ or primary amines²⁰ and incorporation of the CO₂-philic tetrazole groups to PIM-1 by

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† Electronic supplementary information (ESI) available: The ESI contains further photographs investigating the precipitation of PIM-1 into solutions of pyrene. Dynamic light scattering analysis of PIM-1 precipitation with pyrene, gas sorption data comparing the effect of length of time used for the methanol soaking of PIM-1 films, a photograph of PIM-1 and PAH blends dried to give films, UV-Vis data for the relationship between pyrene concentration in solution and the amount of pyrene absorbed by the PIM-1 film, visual and SEM comparison of post-modified PIM-1 films, and transmission UV-Vis of PIM-1 films post-modified with pyrene. See DOI: 10.1039/c4ta06070a



Fig. 1 Repeat unit of PIM-1.



Fig. 2 Cartoon representation of physical crosslinking induced by mixing with pyrene additive. (A) Independent rigid PIM-1 polymer chains. (B) Intermolecular and intramolecular physical crosslinking of the PIM-1 chains with pyrene.

reacting the nitrile groups with sodium azide.¹⁷ Crosslinking approaches have included thermal crosslinking of the polymer by the trimerisation of the nitrile groups to triazine,¹³ UV radiation has been used to give an increase in the selectivity, firstly by Chung and co-workers who suggested that the modification in behaviour was due to chain rearrangement²¹ while more recently Sivaniah and co-workers attributed the modification in properties to a photo-oxidative surface modification of the membrane.¹⁶ It has been proposed that crosslinking PIM-1 may have the effect of reducing the pore size of the material thus improving its selectivity. Sivaniah and co-workers have recently demonstrated the use of thermal oxidative crosslinking to tune the properties of PIM-1 by modifying the micropore structure.²²

An interesting method of adjusting the properties of a PIM-1 membrane without modifying the polymer itself is to use additives. Mixed matrix approaches²³ have been shown to be useful in enhancing the gas transport properties of PIM-1 and a range of different materials have been produced by blending PIM-1 with (amongst others) inorganic fillers,²⁴ cage nanocrystals,²⁵ porous nanoparticles,²⁶ carbon nanotubes²⁷ and other polymers.²⁸ Chemical crosslinking with blended polymers has been found to enhance the separation properties.^{29,30} For unrelated polymer systems, supramolecular interactions between polymers have been exploited to provide physical crosslinking used in self-healing polymers.³¹ An example of this approach is the use of charge transfer interactions between electron-rich pyrenyl end-groups and electron-deficient naphthalene diimide units.^{32,33} Here, we propose a novel strategy to modify the properties of PIM-1 by combining the concepts of blending and crosslinking; an additive will be added to the polymer with the aim of facilitating crosslinking *via* physical intermolecular interactions. Given the relatively electron-deficient structure, we hypothesised PIM-1 might interact favourably with electron-rich polycyclic aromatic hydrocarbons (PAHs) (Fig. 2). We have investigated the effect of mixing PIM-1 with a range of PAHs on the gas sorption and mechanical properties of the resulting blends.

Materials and methods

Materials

All materials were used without further purification, 3,3,3',3'-tetramethyl-1-1''-spirobisindane-5,5',6,6'-tetrol was purchased from Apollo (UK), 2,3,5,6-tetrafluorophthalonitrile was purchased from Alfa Aesar (UK) and K₂CO₃ was supplied by Fisher (UK). All other materials were purchased from Sigma-Aldrich (UK).

Synthesis of PIM-1

PIM-1 was synthesised using the procedure described by Budd *et al.*⁵ Briefly, K₂CO₃ (963.2 mmol, 133.12 g) was added to a solution of 3,3,3',3'-tetramethyl-1-1''-spirobisindane-5,5',6,6'-tetrol (120.4 mmol, 40.99 g) and 2,3,5,6-tetrafluorophthalonitrile (120.4 mmol, 24.09 g) in dry DMF (1200 mL). The reaction was stirred at 65 °C for 72 hours. After cooling to room temperature, water (1200 mL) was added. The product was collected by filtration before being reprecipitation from CHCl₃ into MeOH. The resulting yellow solid was dried under vacuum overnight. The yield was 52.42 g (94%). The polymer was analysed by ¹H-NMR, elemental analysis and GPC (Viscotek system employing OmniSEC 4.6 software, TDA Model 302 (right-angle & low-angled light scattering, refractive index, ultra-violet and viscometer detectors coupled to a gpcMAX integrated solvent and sample delivery module). The system was fitted with a Viscogel HHR-H guard column and two Viscogel GMHHR-H columns, at a flow rate of 1 mL min⁻¹, using chloroform). ¹H-NMR (400 MHz, CDCl₃): δ_H 1.31 (6H, s), 1.37 (6H, s), 2.24 (4H, br. d), 6.42 (2H, s), 6.81 (2H, s). Elemental analysis: C₂₉H₂₀N₂O₄. Theory: C 75.64, H 4.38, N 6.08. Found: C 73.26, H 4.27, N 5.96. GPC: *M*_n 70 740 Da, *M*_w 134 641 Da, *M*_w/*M*_n 1.90.

Precipitation experiments

The precipitation of PIM-1 was investigated in a number of PAH solutions. Solutions of PAHs were prepared at 50 mg mL⁻¹ in dichloromethane (DCM), or if this was above the solubility limit of the PAH a saturated solution was prepared and the supernatant was used. A solution of PIM-1 was prepared in DCM at 50 mg mL⁻¹. Three drops (≈0.2 mL) of this solution were added from a 1 mL syringe to each of the PAH solutions. Photographs were taken of samples approximately 10 minutes after PIM-1 addition using a light box (Digital imaging box DIB-1612).

Preparation of PIM-1 and pyrene blends

PIM-1 and pyrene blends were produced as follows. Solutions of PIM-1 or pyrene were prepared at 50 mg mL⁻¹ in DCM. To the appropriate amount of pyrene solution in a 4 mL vial the PIM-1 solution was added, achieving the desired composition with a total volume of 2 mL. The samples were left for 48 hours for the DCM to fully evaporate, before being placed under vacuum at 65 °C for 18 hours. The polymer blends were then immersed in methanol for 3 hours which was then decanted prior to the films being dried under vacuum again at 65 °C for 4 hours. The samples were then broken into smaller pieces and removed



from the vials for gas sorption analysis. The pyrene content of the film was calculated from the initial amount of pyrene solution added.

Post modification of PIM-1 films with pyrene or 1-aminopyrene

Post-modification of PIM-1 films with pyrene or 1-aminopyrene was carried out by the following method. Firstly a PIM-1 film was produced by adding 16–18 g of PIM-1 solution in DCM at 50 mg mL⁻¹ in a 200 mL evaporation dish, which was then left overnight for the DCM to fully evaporate. Pieces of this film PIM-1 were then used to test the effect of different pyrene or 1-aminopyrene content in post-modified films. A control treatment (no modification) was carried out with methanol alone. Four concentrations of pyrene or 1-aminopyrene in methanol were tested up to 7 mg mL⁻¹. In all cases, the films were immersed in these methanolic solutions of pyrene or 1-aminopyrene for 3 hours prior to being dried under vacuum at 65 °C for 4 hours.

Dynamic light scattering analysis

The aggregation of PIM-1 and pyrene was analysed by dynamic light scattering (DLS). PIM-1 was added to a quartz cuvette at 0.39 mg mL⁻¹ in DCM, to which aliquots of pyrene in DCM (at a concentration of 50 mg mL⁻¹) were then added (to achieve a final overall concentration of pyrene of up to 7.6 mg mL⁻¹). The sample was mixed prior to DLS analysis. The scattering of the solutions were measured at a temperature of 25 °C using a Malvern Zetasizer Nano ZS equipped with a 4 mW He-Ne, 633 nm laser. Malvern Zetasizer software version 6.20 was used for data analysis.

UV-Vis characterisation of PIM-1 films post-modified with pyrene or 1-aminopyrene films

The UV-Vis absorption spectra of unmodified and post-modified PIM-1 films were obtained in transmission using a Shimadzu UV-2550 UV-Vis spectrophotometer running the UVProbe software, version 2.34. All spectra were measured from 190–800 nm, with scan speed set to fast and using a slit width of 5 nm.

Characterising pyrene or 1-aminopyrene content in post-modified PIM-1 films

The pyrene or 1-aminopyrene content of the post-modified PIM-1 films was determined by a UV-Vis mass balance approach. A known mass of PIM-1 film was soaked in a known concentration of additive (either pyrene or 1-aminopyrene). After 3 hours, the PIM-1 film was removed and the absorbance of the solution was measured (Shimadzu UV-2550 UV-Vis spectrophotometer) at 332 nm or 282 nm (for pyrene or 1-aminopyrene respectively). The concentration was determined by comparison to calibration from known standards of pyrene or 1-aminopyrene. Any reduction in concentration of the additive was assumed to have been a result of absorption into the PIM-1 film.

Scanning electron microscopy (SEM) of PIM-1 films

Scanning electron microscopy (SEM) analysis was used to analyse the cold-fractured (liquid N₂) edges of PIM-1 unmodified and post-modified films and measure the film thickness. Samples of the films were fixed onto the aluminium stubs with carbon tabs perpendicular to the surface of the stub. The samples were then gold coated for 3 minutes at 20 mA using a sputter-coater (EMITECH K550X) prior to imaging. SEM images were recorded using a Hitachi S-4800 FE-SEM at 3 kV. The mean film thickness was determined by imaging 7 samples of the film, the thickness of the each sample of film was measured in at least 6 places along the film using ImageJ 1.46r. The mean thickness of the film was determined from at total of 137 measurements, providing a value of 49.0 µm with a standard deviation of 5.0 µm.

Nanoindentation analysis

Nanoindentation was carried out using an Agilent nanoindenter G200 (Keysight Technologies, Chandler, AZ, USA) instrument with an ultra-low load dynamic contact module indentation head (DCM-I). The indentations were performed at ambient temperature, aligned normal to the film surface, using a Berkovich tip with a 20 nm radius. The film samples were adhered to aluminium pucks using adhesive pads (STKYDOT, Bruker, Coventry, UK). The continuous stiffness measurement option³⁴ was used at a frequency of 75 Hz for the testing. Each indent was made in the film samples up to a maximum depth of 300 nm of a total film thickness of 49 µm. A Poisson's ratio of 0.3 was assumed in order to calculate the elastic modulus. 36 indentations were made on each sample.

Gas sorption analysis

Initial nitrogen isotherm analyses for the PIM-1 pyrene blends at 77 K were obtained on a Micromeritics ASAP 2420 instrument using an equilibration interval of 30 seconds for relative pressures less than 0.01 and 15 seconds equilibration interval for all other pressures. Prior to analysis, samples were degassed under vacuum for 900 minutes at 80 °C. The free space measurements were carried out after analysis. The cycled isotherm N₂ measurements were obtained using a Micromeritics ASAP 2020 with an equilibration interval of 30 seconds for relative pressures less than 0.15 and 20 seconds equilibration interval for all other pressures. An *in situ* degas was carried out for 4 hours at 80 °C before analysis. Free space measurements were carried out after analysis. The N₂ and CO₂ gas sorption measurements at 298 K were obtained on a Micromeritics ASAP 2020 equipped with a HAAKE Phoenix II circulator. The equilibration intervals were as follows >1.33 mbar = 45 seconds, 1.33–13.33 mbar = 30 seconds, 13.33–133.3 mbar = 25 seconds, 133.3–1100 mbar = 20 seconds. The free space measurements were carried out before analysis as is usual. Gas sorption analysis of the post-modified PIM-1 films was carried out on Micromeritics 3Flex Surface Characterisation Analyser with 3 samples ports each with 100, 10 and 0.1 Torr pressure transducers. Temperature control was achieved using Micromeritics supplied ColdEdge



Technologies Cryocooler model CH-104 with a sample temperature range of 50 K to 310 K with <0.01 K drift. For the N₂ at 77 K analysis an *in situ* degas was carried out for 180 minutes at 80 °C prior, and the equilibration intervals for the following relative pressures were used >0.001 = 45 seconds, 0.001–0.01 = 30 seconds, 0.01–0.1 = 25 seconds, 0.1–1 = 20 seconds. The free space measurements were carried out after analysis. The N₂ and CO₂ gas sorption measurements at 293 K were obtained using the 3Flex using the same settings as used on the ASAP 2020 at 298 K. The CO₂/N₂ gas sorption selectivity was calculated using the respective uptakes at 1000 mbar. All gases used in the gas sorption experiments were at least 99.9995% pure.

Single gas permeability measurements

The pure gas permeabilities of the PIM-1/PAH films were characterised using a home-made setup outlined in Lau *et al.*'s recent work.²⁶ The pure N₂, and CO₂ permeabilities were determined using a constant volume and variable pressure method.³⁵ Gas permeabilities at 2 atm were determined from the rate of downstream pressure build-up rate (dp/dt) obtained when permeation reached a steady state according to the following equation:

$$P = D \times S = \frac{273 \times 10^{10}}{760} \frac{VL}{AT \left[\frac{P_2 \times 76}{14.7} \right]} \left(\frac{dp}{dt} \right)$$

P refers to the permeability of a membrane to a gas and its unit is in Barrer ($1 \text{ Barrer} = 1 \times 10^{-10} \text{ cm}^3 \text{ (STP)-cm cm}^{-2} \text{ s}^{-1} \text{ cmHg}^{-1}$),⁷ D is the average effective diffusivity ($\text{cm}^2 \text{ s}^{-1}$), S is the apparent sorption coefficient/solubility ($\text{cm}^3 \text{ (STP) per cm}^3 \text{ polymer cmHg}$), V is the volume of the downstream chamber

(cm³), L is the film thickness (cm). A refers to the effective area of the membrane (cm²), T is the experimental temperature (K) and the pressure of the feed gas in the upstream chamber is given by p_2 (psia).

Results and discussion

Charge-transfer interactions are intermolecular interactions between π -electron rich (donor) molecules and π -electron-deficient (acceptor) molecules. We hypothesised that the electron-deficient PIM-1 molecule might undergo charge-transfer with PAHs and that these interactions might lead to modification of the physical properties of the material. Addition of a PIM-1 solution to PAH solutions showed immediate precipitation of the polymer in the cases of pyrene, triphenylene, benzo[*a*]pyrene or coronene (Fig. 3). In the case of pyrene and benzo[*a*]pyrene, a slight darkening of the solutions was also observed. No precipitation was observed for any of the other PAHs tested here. There therefore seems to be a requirement for an extended planar conjugated system in order to induce sufficient attraction to the PIM-1 to cause precipitation. Pyrene has previously been shown to form electron donor-acceptor complexes with an electron acceptor molecule.^{36,37} We believe that simple π - π stacking can be ruled out here as the interaction mechanism because precipitation was not seen for all PAHs. Additionally precipitation was also observed when pyrene was dissolved in benzene, where one would expect significant competition if π - π stacking were the driving force for interaction (Fig. S1, ESI†). Mixing the PIM-1 and pyrene solutions resulted in visible precipitation at concentrations of pyrene ≥ 6.25 mg mL⁻¹ (Fig. S2, ESI†). Further investigation of the precipitation process

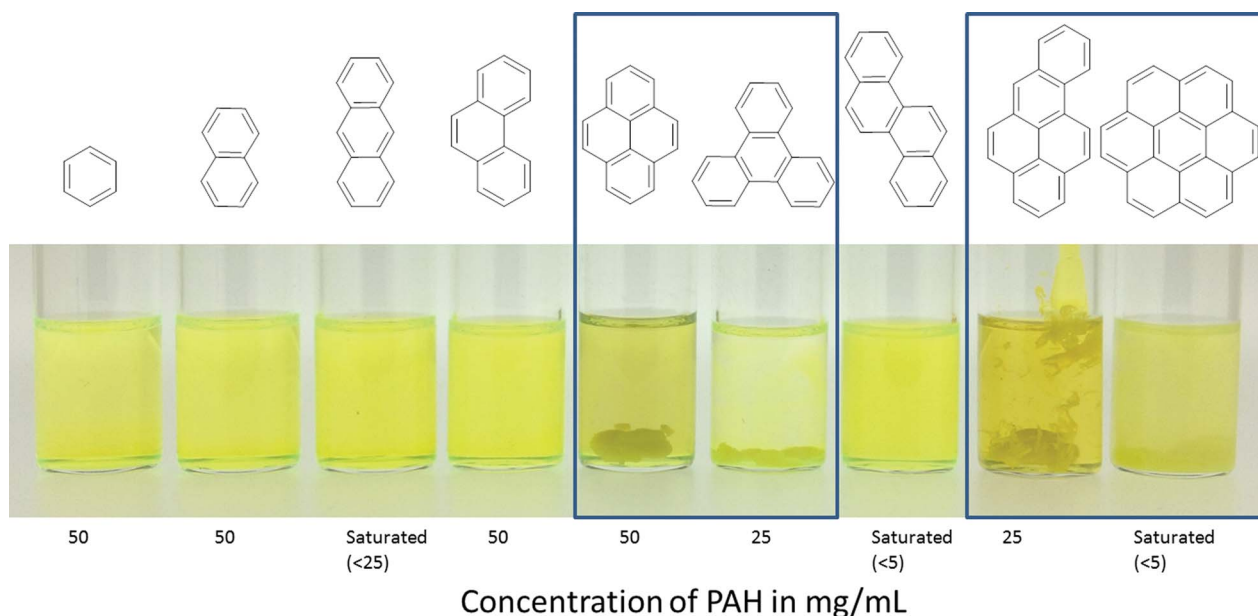


Fig. 3 Effect of the structure of (poly)cyclic aromatic hydrocarbons on the interaction with PIM-1 solution. Three drops of PIM-1 solution (50 mg mL⁻¹ in DCM) were added to PAH solutions (in at the stated concentration DCM). The cyclic aromatic hydrocarbons and PAHs tested were (from left to right): benzene, naphthalene, anthracene, phenanthrene, pyrene, triphenylene, chrysene, benzo[a]pyrene and coronene. Blue boxes indicate when precipitation was visually observed.



Fig. 4 Gas sorption properties for PIM-1 pyrene blends, nitrogen uptake at 1 bar (obtained from nitrogen isotherms at 77 K) and CO₂ uptake at 1 bar (at 298 K) of PIM-1 and pyrene blends as a function of the composition of the film (trend lines are to guide the eye).

by dynamic light scattering (DLS) showed that aggregation of the PIM-1 polymer chains occurred at much lower concentrations (PIM-1 at 0.39 mg mL⁻¹ with pyrene at concentrations above 0.2 mg mL⁻¹, see Fig. S3, ESI†). For further experiments, pyrene was selected as the preferred PAH since this showed a strong interaction with PIM-1, whilst also being inexpensive and abundant. These findings suggest that attractive intermolecular interactions between PIM-1 and certain PAHs results in a physical crosslinking of the polymer causing formation of aggregates and at higher concentrations, precipitation of the polymer.

We were interested in modifying the gas sorption and permeation properties of PIM-1 membranes. We hypothesised that the supramolecularly crosslinking PIM-1 with pyrene would allow these properties to be tuned. Therefore, solutions of PIM-1 and pyrene in DCM were mixed at varying ratios (v/v) and then evaporated to dryness to form films blends. Methanol⁴ or ethanol¹¹ soaking has been shown to be important in enhancing the uptake of PIM-1 films. Therefore, all PIM-1 and pyrene blends were soaked in methanol for 3 hours (longer soaking times had no effect on the N₂ isotherm (see Fig. S4, ESI†)). The 100% PIM-1 film displayed an isotherm with a Brunauer–Emmett–Teller (BET) surface area³⁸ of 686 m² g⁻¹, in agreement with the literature value of approximately 720 m² g⁻¹.⁴ The effect of blending pyrene with PIM-1 was dramatic; at pyrene compositions ≥5% (w/w), essentially no nitrogen uptake was observed (Fig. 4). At lower pyrene contents in the blend, gate-opening behaviour was seen in the isotherm (see Fig. S5, ESI†). This unusual behaviour is currently not well understood and has been reported for PIM-1 in other work.^{22,39} These steps occurred between relative pressures of 0.25–0.5 and appear to be related to the amount of pyrene included in the blend; as the pyrene content increased, the pressure at which pore-opening occurred was also increased. While the isotherms indicated that at lower pressures the equilibration interval was not long enough for complete equilibration, the overall data was of sufficient quality to allow direct comparison between gas

uptakes of the different blends. The N₂ uptakes of the polymer blends showed a dramatic approximately linear decrease with increasing pyrene content from 15.6 mmol g⁻¹ for 100% PIM-1 to 0.7 mmol g⁻¹ for a blend of PIM-1 containing 5% pyrene (Fig. 4). Whilst there was a marked reduction in the nitrogen uptakes of the PIM-1 and pyrene blends, the effect of incorporating pyrene into the film on the CO₂ uptake at 298 K was much less pronounced; CO₂ uptake gradually reduced with increasing pyrene content (Fig. 4).

The dramatic reduction in nitrogen uptake coupled with only a slight loss in CO₂ uptake with 5% pyrene in the film suggests that these films may offer higher selectivity for CO₂ over N₂ and as such may possess superior gas separation characteristics. The gas sorption results of PIM-1 and pyrene blends suggests that the addition of pyrene reduces the pore size of the material. Given the larger size of the N₂ molecule (3.6 Å) than CO₂ (3.3 Å) it is likely that increased crosslinking would narrow the interconnected cavities present in PIM-1²² making the diffusion N₂ into the materials more difficult.

In order to further investigate the observed pore-opening behaviour observed in the N₂ isotherms, cycles of adsorption and desorption were carried out. A PIM-1 (99%) and pyrene (1%) blend was analysed for 3 cycles (Fig. 5A). In these experiments

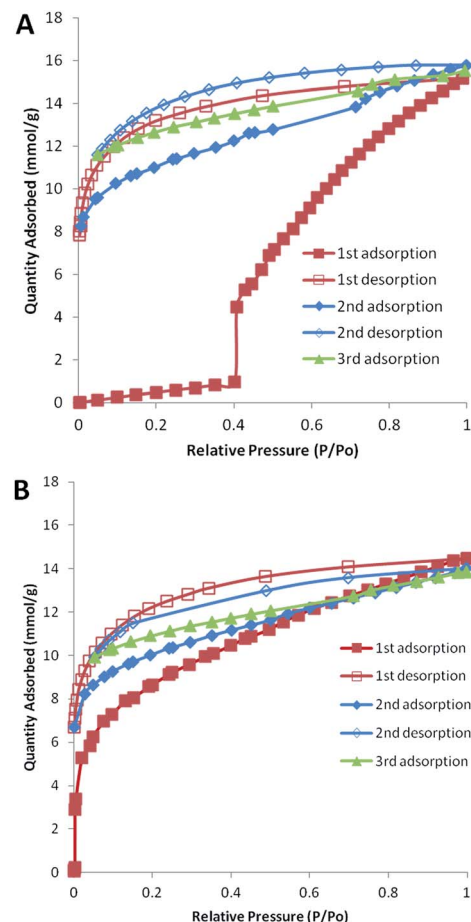


Fig. 5 Cycled adsorption and desorption nitrogen isotherms (77 K). (A) PIM-1 (99%) and pyrene (1%) blend film. (B) PIM-1 film (100%).



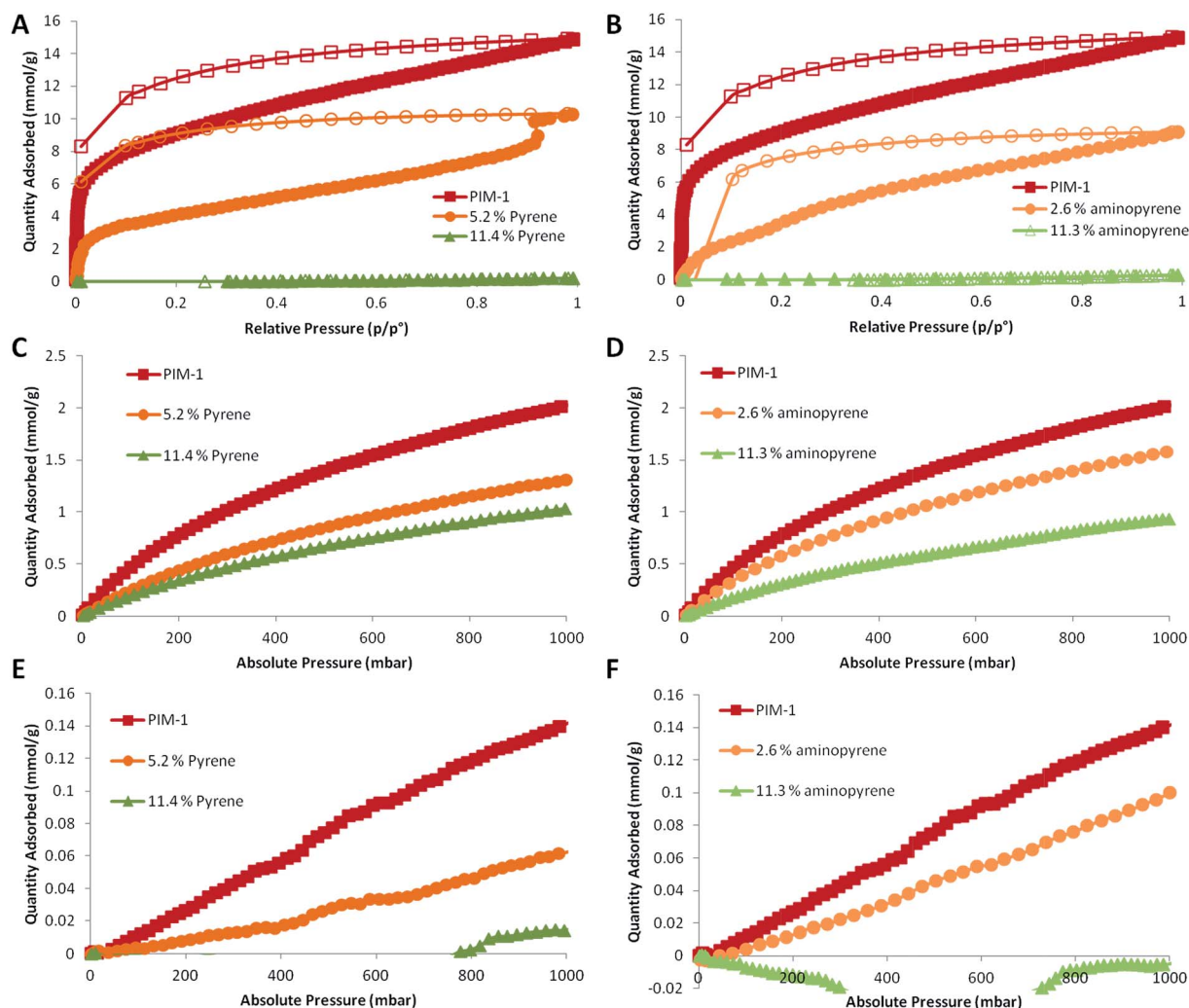


Fig. 9 Gas sorption properties of PIM-1 films post-treated with methanolic pyrene solutions or methanolic aminopyrene solutions. (A) & (B) Nitrogen isotherms (77 K) (legend details the additive content) for pyrene and aminopyrene. (C) & (D) CO₂ isotherms (293 K) (legend details the additive content) for pyrene and aminopyrene. (E) & (F) Nitrogen isotherms (293 K) (legend details the additive content) for pyrene and aminopyrene.

samples with no apparent difference between the different pyrene contents (Fig. S8, ESI†). SEM analysis of a cold-fractured edge of both unmodified PIM-1 film and a film post-modified with pyrene did not reveal any obvious differences (Fig. S9, ESI†). Charge-transfer interactions typically display changes in the photophysical properties of the materials with the emergence of a charge-transfer band (400–650 nm).³⁷ In the case of the PIM-1 film post-modified with pyrene we observed no apparent charge-transfer band (Fig. S10, ESI†). We note that other studies have shown that charge-transfer absorption bands are not always present³⁷ or are very weak.³⁶ However, the UV-Vis absorption of the PIM-1 films post-modified with 1-aminopyrene displayed a more prominent charge-transfer band as a broadening from 500–550 nm in the UV-Vis spectra of the film (Fig. 7). As the amount of 1-aminopyrene incorporated into the film increased, a broader charge transfer band was found.

To probe the mechanical properties of post-modified films we used nanoindentation. This technique has been used

elsewhere to investigate crosslinking in PIM-1 films, with higher crosslinking densities leading to higher elastic moduli.^{16,22} These studies found the elastic modulus of PIM-1 (at a displacement of 100 nm) to be between 2.6 and 4.1 GPa (ref. 16 and 22) in agreement with our value of 2.8 GPa for unmodified PIM-1. We found that post-modification of PIM-1 with both pyrene-based additives produced films with higher elastic moduli, 3.5 GPa and 4.0 GPa for pyrene or 1-aminopyrene modification respectively (Fig. 8). Interestingly, the post-modification of PIM-1 with 1-aminopyrene led to a greater increase in the modulus than pyrene. These nanoindentation results present further evidence for stronger interactions between the PIM-1 polymer chains and pyrene-based additives provides a crosslinking effect, resulting in stiffer films.

The PIM-1 films post-modified with varying amounts of pyrene or 1-aminopyrene additives were then analysed by N₂ and CO₂ gas sorption. The post-modification of PIM-1 films with pyrene additives had a similar effect on the gas sorption



^a A negative uptake was measured due to pressure transducer inaccuracy at very low gas uptake.

The gas permeabilities of PIM-1 are known to decrease over time as a function of polymer ageing.^{4,45,46} Polymer ageing

Figure 1 is a log-log plot showing the relationship between CO₂ permeability and CO₂/N₂ selectivity for various polymers. The x-axis represents Permeability CO₂ (Barrer) on a logarithmic scale from 10 to 10,000. The y-axis represents Selectivity CO₂/N₂ on a logarithmic scale from 10 to 100. A solid black line represents the Robeson upper bound. Data points are shown for PIM-1 (red squares), PIM-1 10% pyrene (blue diamonds), and PIM-1 10% aminopyrene (green triangles). Arrows indicate the direction of improvement for each material. PIM-1 10% aminopyrene shows the highest selectivity for a given permeability, followed by PIM-1 10% pyrene, and then PIM-1.

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occurs as polymer chains attempt to relax towards each other to achieve a thermodynamic equilibrium. We observe the same as trend commonly reported in literature, with the PIM-1 film displaying a decrease in permeability with increased ageing. When a PIM-1 film was post-modified with pyrene, we observed a slight decrease in the rate of ageing. Interestingly, in the case of the PIM-1 film post-modified with 1-aminopyrene, the ageing effect on the permeability was completely arrested and a slight increase in CO₂ permeability was observed over time (Fig. 10). With physical aging, the pore sizes could decrease over time to an extent that the pore size favours CO₂ sorption, resulting in increased enthalpy of CO₂ adsorption. Other researchers have also shown that decreasing the pore volume of other porous materials such as metal-organic frameworks and polymers can lead to increased enthalpy of gas adsorption.^{47,48} Our permeability experiments suggest that the stronger charge-transfer interactions between PIM-1 and 1-aminopyrene (compared to pyrene) may be sufficient to physically crosslink the polymer locking the polymer chains in position, and this rigidification of the PIM-1 chains is the key to stop polymer ageing.

Conclusions

We have demonstrated an extremely facile and inexpensive method for modifying the gas sorption and physical properties of PIM-1. PIM-1 has been shown to interact with a number of PAHs, an extended planar conjugated system appeared to be necessary in order to induce sufficient attraction to the PIM-1 to cause precipitation. We demonstrate that blending pyrene as an additive with PIM-1 led to a dramatic reduction in nitrogen uptake (at 77 K) while only leading to a minor loss of CO₂ uptake (at 298 K). In order to maintain a fully solution processable material, a post-modification strategy with pyrene and 1-aminopyrene was developed. Nanoindentation analysis of the PIM-1 films post-modified with pyrene or 1-aminopyrene revealed that these samples had higher elastic moduli than unmodified PIM-1 films, evidence of the physical crosslinking of the PIM-1 chains. Gas sorption analysis of PIM-1 films post-modified with pyrene and 1-aminopyrene demonstrated that increasing the additive content had a greater effect on reducing the N₂ uptake at 77 K than on lowering the CO₂ uptake at 293 K. The effect on the uptake of N₂ at 293 K was less pronounced, although PIM-1 films post-modified with 11% pyrene or 1-aminopyrene displayed essentially no N₂ uptake, these films displayed vastly improved gas sorption selectivities for CO₂ over N₂ compared to unmodified PIM-1 film.

Analysis of the CO₂ and N₂ gas permeabilities for the post-modified PIM-1 films suggested that while the use of pyrene based additives reduced the overall permeability of the film, the physical crosslinking effect produced by modification of PIM-1 film with 10% 1-aminopyrene was sufficient to stop the usual loss of permeability observed for unmodified PIM-1 with ageing over the 50 days tested.

The effect of blending electron-rich additives into PIM films was highly pronounced. Even only a few percent of inexpensive, commercially available additive (pyrene) was sufficient to significantly modify both the gas sorption properties, and

physical characteristics of the film. Considering the widespread current interest in PIMs for practical applications, this opens up a new strategy for modifying PIM materials. There are now several PIM type polymers reported in the literature which combine porosity, solubility, and an electron-deficient repeating unit. However, the last few years have seen the reports of new porous polymer systems, based on repeating units of pyrene and other related PAHs,^{50–52} which display similar porosity and solubility, but with electron-rich repeating units. For commercial non-porous polymers it is commonplace to blend a polymer with a small amount of co-polymer, or molecular additive, in order to tune physical properties such as glass transition temperature or stiffness. Our results show that a similar methodology could potentially be achieved in PIMs, fully realising the potential for processing that solubility gives these polymers in comparison to other porous materials.

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