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Robust pyridylbenzoate metal–organic frameworks as sorbents for volatile solvents and gases†

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Two activated isostructural porous compounds, [Co(34pba)(44pba)]_n (**1d**) and [Zn(34pba)(44pba)]_n (**2d**) (where 34pba is 3-(pyridin-4-yl)benzoate, 44pba is 4-(pyridin-4-yl)benzoate, and **d** indicates the activated phase) were used for the adsorption of halogenated volatile organic compounds (VOCs), iodine, carbon dioxide (CO₂), and hydrogen. Elucidation of single-crystal structures found that the desolvated phases **1d** and **2d** show rotational disorder of one ring on each linker. The crystal structure of [(Co(34pba)(44pba))·1.6 I₂]_n (**1dI₂**) was obtained by vapour sorption of iodine into **1d**, and this could be desolvated to a phase isostructural with **1d**. Thus the MOF is robust to cycling through sorption/desorption processes with an accompanying order/disorder phase transition. Activation energies for the desorption of dichloromethane, dibromomethane, and iodine were measured as 70, 60, and 77 kJ mol⁻¹ respectively. Both activated compounds were tested as gas sorbents, with **1d** showing higher adsorption capacity than **2d** for both carbon dioxide and hydrogen.

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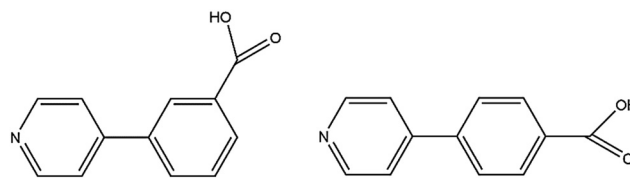
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1. Introduction

Porous metal–organic frameworks (MOFs) have been widely studied in applications such as gas sorption and storage, catalysis, luminescence, magnetism, and sensing.^{1–8} Features such as surface area, solvent-accessible void volume, or pore size play an important role in the applications of a given MOF. Sorption properties are governed by supramolecular interactions between the framework (host) and the adsorbed species (guest).^{9,10} The presence of a specific substituent in a guest molecule allows a given set of specific host–guest supramolecular interactions which may include hydrogen or halogen bonding.^{11,12} These interactions often involve structural changes which can reversibly induce pore opening and closing.^{13–15} Such reversibility in sorption/desorption in response to chemical and physical stimuli are important in the recovery of adsorbents. Thus, host–guest chemistry can be used to design adsorbents for specific purposes. One such application is the removal of harmful volatile organic compounds (VOCs) in the environment.^{16,17}

We previously reported isomorphous MOFs in which the channel double-walls are formed by the related linkers 3-(pyridin-4-yl)benzoic acid (H34pba) and 4-(pyridin-4-yl)benzoic acid (H44pba), Scheme 1. These MOFs showed selective adsorption for the chlorinated VOCs dichloromethane, chloroform, and chlorobenzene.¹⁸ Guest molecules could be removed with the retention of the framework. To further understand the robustness of this framework through the sorption of additional halogenated compounds, we now consider the use of [Co(34pba)(44pba)]_n (**1d**) and [Zn(34pba)(44pba)]_n (**2d**) (where 34pba is 3-(pyridin-4-yl)benzoate, 44pba is 4-(pyridin-4-yl)benzoate, and **d** indicates the activated phase) as adsorbents for the bromo- and iodo-derivatives of the chloro-compounds studied previously. Furthermore, the sorption of iodine vapour, and of carbon dioxide and hydrogen gases by the same adsorbents is also presented here.



Scheme 1 3-(Pyridin-4-yl)benzoic acid (H34pba) and 4-(pyridin-4-yl)benzoic acid (H44pba).

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2. Materials and methods

2.1 Preparation and activation of MOFs and sorption experiments

The preparation of the precursors of the activated adsorbents, $\{[Co(34pba)(44pba)]_n \cdot DMF\}$ (**1**) and $\{[Zn(34pba)(44pba)]_n \cdot DMF\}$ (**2**) were reported previously.¹⁸ To obtain the activated phases **1d** and **2d**, they were heated at 210 °C under vacuum for six hours. This protocol was selected to ensure complete desolvation but not decomposition of the MOFs. The corresponding crystallographic data of **1d** and **2d** are presented here.

Powdered activated samples of mass 5–7 mg were placed in narrow glass vials. The latter were placed into larger vials containing the relevant VOC and then sealed to allow vapour sorption at room temperature (Fig. S1†). The VOCs selected for the study were dibromomethane (CH₂Br₂), bromoform (CHBr₃), bromobenzene (BrBenz), diiodomethane (CH₂I₂), iodoform (CHI₃), iodobenzene (IBenz). Code names for the corresponding phases from **1d** and **2d** are provided in Table 1. Samples were analysed after exposure for between one and fourteen days depending on the vapour pressure of the solvent.

The sorption for iodine in **1d** was carried out in a similar way, using the sublimation of solid iodine to expose single crystals to iodine vapour, with samples taken after two hours, two days, eight days, and finally two weeks. Each sample was analyzed by thermogravimetry (TGA) to determine the mass loss owing to desorption of guests and by powder X-ray diffraction (PXRD) to identify changes in phase. A single crystal was selected for analysis and the structure reported as **1dI₂**. The desorption of iodine from **1dI₂** was performed by soaking crystals of **1dI₂** in methanol for three days, to recover the activated form (**1dI₂d**). As this retained its crystallinity, the single crystal structure of **1dI₂d** is also reported here.

Gas sorption capacity of the **1d** and **2d** adsorbents was investigated for carbon dioxide (CO₂) and hydrogen (H₂) gases using a Micromeritics 3Flex Surface Area Analyzer. After grinding the sample, masses between 130–140 mg were prepared using a Micromeritics Flowprep using a flow of nitrogen over the samples for 2 h with continuous heating at 60 °C. Thereafter, samples were heated at 150 °C under vacuum for 2 h prior to the sorption analysis. The sorption for CO₂ was carried out at various temperatures in order to determine the heat of adsorption (*Q_{st}*), while the sorption capacity for H₂ was only carried out at 77 K. Loading of gas

into the samples was characterized by a pressure change from 0 mmHg to the maximum pressure equilibrium (between 600–1000 mmHg). The complete sorption corresponded to the equilibrium pressure which was followed by the desorption process.

2.2 Thermogravimetric analysis (TGA)

Mass losses attributable to desolvation of guest molecules were determined by thermogravimetric analysis (TGA) using a TA instrument TA-Q500 on 1–2 mg samples in open platinum pans under nitrogen gas flow (50 mL min⁻¹) at a heating rate of 10 °C min⁻¹.

2.3 Powder X-ray diffraction (PXRD)

Powder X-ray diffraction (PXRD) patterns were measured on a Bruker D8 Advance X-ray diffractometer operating in a DaVinci geometry equipped with a Lynxeye detector using Cu-K α -radiation ($\lambda = 1.5406 \text{ \AA}$). X-rays were generated at 30 kV and 40 mA. Samples were placed on a zero-background sample holder and scanned over a range of 4–40° in 2θ with 0.016° step size per second.

2.4 Crystal structure determination

Single crystals were selected using optical microscopy under plane-polarized light. Intensity data were recorded based on a Bruker KAPPA APEX II DUO diffractometer using graphite monochromated Mo-K α radiation ($\lambda = 0.71073 \text{ \AA}$). Data were corrected for Lorentz-polarization effects and for absorption (SADABS¹⁹). The structures were solved by direct methods in SHELXS and refined by full-matrix least-squares on F^2 using SHELXL²⁰ within the XSEED interface.²¹ Non-hydrogen atoms were located in difference electron density maps and were refined anisotropically while hydrogen atoms were placed in calculated positions and refined isotropically. Details of the crystal structures are given in Tables 2 and S1.†

3. Results and discussion

3.1 Crystal structures of **1d** and **2d**

The metal–organic frameworks $\{[Co(34pba)(44pba)]_n \cdot DMF\}$ (**1**) and $\{[Zn(34pba)(44pba)]_n \cdot DMF\}$ (**2**) were reported previously.¹⁸ These form frameworks where each side is comprised of a 34pba and a 44pba linker, giving a double-walled MOF of *bcu* topology. Hour-glass channels (*ca.* 8 Å at the widest point and *ca.* 3.5 Å at the narrowest) run parallel to [100] (Fig. 1). At that time, we were unable to obtain a single crystal of these MOFs after desorption of the guest species, although PXRD showed minimal changes in phase on desorption. Now, after optimizing the desorption process, we are able to report the single crystal structures of both Co(II) (**1d**) and Zn(II) (**2d**) MOFs. As anticipated the frameworks do not change significantly, except that in both **1d** and **2d**, one ring on each of the 34pba and 44pba linkers shows rotational disorder. This is more pronounced for the 44pba moiety, in which the benzyl ring is disordered over two positions with almost

Table 1 Code names of the phases obtained from sorption experiments

Halogenated compound	Sorbent 1d	Sorbent 2d
Dibromomethane, CH ₂ Br ₂	1dCH ₂ Br ₂	2dCH ₂ Br ₂
Bromoform, CHBr ₃	1dCHBr ₃	2dCHBr ₃
Bromobenzene, BrBenz	1dBrBenz	2dBrBenz
Diiodomethane, CH ₂ I ₂	1dCH ₂ I ₂	2dCH ₂ I ₂
Iodoform, CHI ₃	1dCHI ₃	2dCHI ₃
Iodobenzene, IBenz	1dIBenz	2dIBenz
Iodine, I ₂	1dI ₂	—



Table 2 Crystallographic data

Compound	1d	2d ^a	1d ₁	1d ₁ d
Formula	C ₂₄ H ₁₆ N ₂ O ₄ Co	C ₂₄ H ₁₆ N ₂ O ₄ Zn·0.25H ₂ O	C ₂₄ H ₁₆ N ₂ O ₄ Co·1.6I ₂	C ₂₄ H ₁₆ N ₂ O ₄ Co
Formula mass (g mol ⁻¹)	455.34	466.26	658.36	455.34
Crystal size (mm ³)	0.080 × 0.090 × 0.14	0.030 × 0.060 × 0.090	0.080 × 0.12 × 0.18	0.080 × 0.12 × 0.18
Crystal system	Monoclinic	Monoclinic	Monoclinic	Monoclinic
Space group	<i>P</i> 2 ₁ / <i>c</i>	<i>P</i> 2 ₁ / <i>c</i>	<i>P</i> 2 ₁ / <i>c</i>	<i>P</i> 2 ₁ / <i>c</i>
<i>a</i> (Å)	10.3931(14)	10.2197(5)	10.114(3)	10.4014(11)
<i>b</i> (Å)	16.027(2)	16.1263(8)	16.501(4)	16.1330(17)
<i>c</i> (Å)	14.996(2)	14.8826(7)	14.700(4)	14.6878(15)
β (°)	98.243(2)	98.059(2)	97.159(4)	98.482(2)
<i>V</i> (Å ³)	2472.2(6)	2428.5(2)	2434.3(11)	2437.7(4)
<i>T</i> (K)	100(2)	100(2)	293(2)	293(2)
<i>Z</i>	4	4	4	4
<i>D</i> _c (g cm ⁻³)	1.223	1.275	1.796	1.241
μ (Mo-K α) (mm ⁻¹)	0.722	1.042	2.766	0.733
<i>F</i> (000)	932	954	1271	932
Range scanned, θ (°)	1.870–27.963	2.526–28.318	2.029–26.425	1.886–25.043
No. reflections collected	22 903	139 665	18 540	13 599
No. unique reflections	5889	6052	4970	4301
No. reflections with $I \geq 2\sigma(I)$	3893	4793	3691	3067
Parameters/restraints	335/0	354/0	334/3	358/0
Goodness of fit, <i>S</i>	1.042	1.031	1.110	1.031
Final <i>R</i> indices ($I \geq 2\sigma(I)$)	<i>R</i> ₁ = 0.0655 <i>wR</i> ₂ = 0.1677	<i>R</i> ₁ = 0.0518 <i>wR</i> ₂ = 0.1329	<i>R</i> ₁ = 0.0687 <i>wR</i> ₂ = 0.1838	<i>R</i> ₁ = 0.0430 <i>wR</i> ₂ = 0.0959
Final <i>R</i> indices (all data)	<i>R</i> ₁ = 0.1085 <i>wR</i> ₂ = 0.1878	<i>R</i> ₁ = 0.0715 <i>wR</i> ₂ = 0.1459	<i>R</i> ₁ = 0.0934 <i>wR</i> ₂ = 0.1956	<i>R</i> ₁ = 0.0749 <i>wR</i> ₂ = 0.1071
Min, max e ⁻ density/(e Å ⁻³)	0.697, -0.585	0.608, -0.546	2.179, -0.643	0.488, -0.361

^a 2d was found to contain ca. 0.25 water molecules per Zn, which are disordered over five sites.

equal site occupancies, while in the 34pba moiety the pyridyl ring is disordered over two positions with the major position having approximately 80% occupancy. This disorder is illustrated in Fig. 2.

3.2 Sorption of halogenated VOCs by 1d and 2d

In this study, we have extended the range of halogenated volatile organic compounds which are included by 1d and 2d. In doing so, we expected to see differences in loading

capacity of the frameworks based on differences in halogen substituent, owing to their differences in polarity, vapour pressure and ability to participate in non-covalent interactions such as halogen bonding.

The volumes of the solvent-accessible voids in 1d and 2d were estimated using Mercury²² with a probe radius of 1.2 Å

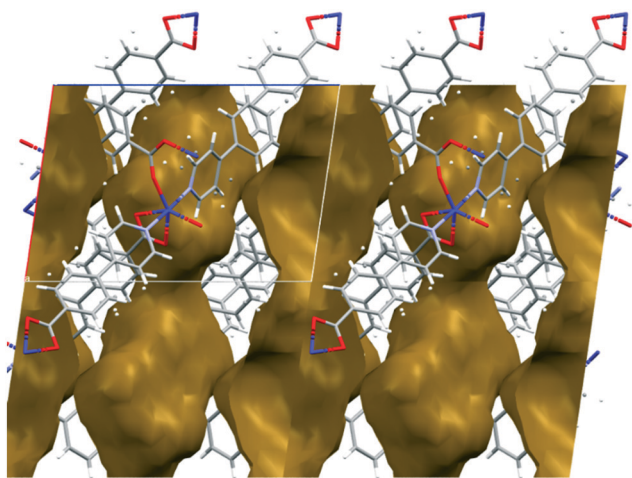


Fig. 1 Channels in the structure of 1d run parallel to [100] and have an hour-glass shape (widest point ca. 8 Å, narrowest point ca. 3.5 Å).

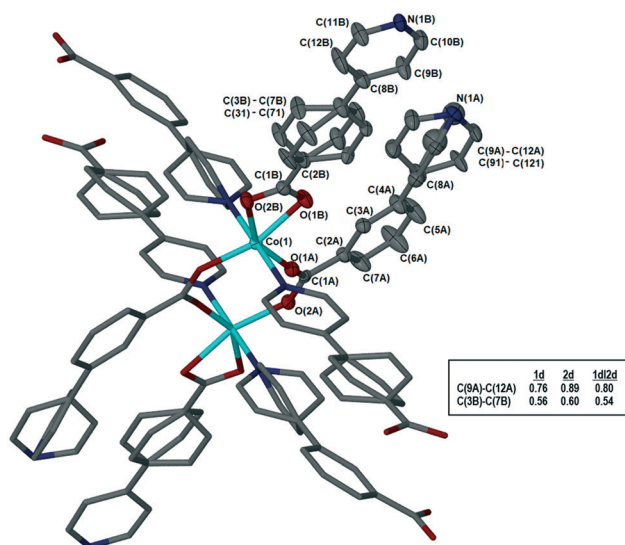


Fig. 2 Coordination geometry and atom labelling, illustrated for 1d. The asymmetric unit is shown with thermal ellipsoids at 50% probability; hydrogen atoms omitted for clarity. Inset table shows site occupancies of the major components for the disordered rings in 2d, and 1d₁d.



and a grid spacing of 0.2 Å. These were estimated as 481 Å³ (20%) per unit cell for **1d** and 571 Å³ (24%) per unit cell for **2d**. Results of the sorption experiments are presented in Table 3. The loading capacity values (L_c) were calculated from TGA analysis (Fig. S2†) using the crystallographically derived void volume and the liquid density of the corresponding solvent. We approximated the maximum loading capacity (ML_c) for the empty frameworks using eqn (1):

$$ML_c = (\text{solvent accessible void volume}) / (Z \times \text{molecular volume}) \quad (1)$$

The molecular volumes of the VOCs were calculated using their liquid density according to eqn (2):

$$\text{Molecular volume of solvent} = \frac{\text{Molecular mass}}{N_A \times \text{density}} \quad (2)$$

The loading capacity (L_c) in the proposed formula $\{[M(34pba)(44pba)]_x \text{ solvent}\}_n$ for both systems is lower than the maximum loading capacity. The dihalomethanes have lower molecular volumes and higher vapour pressures than their corresponding haloforms, which may account for their higher loading capacities. The aromatic halogenated VOCs have lower vapour pressures, but their loading capacity was less affected. In part, this is probably due to the additional interactions possible for these compounds (see next paragraph). In general, higher sorption (% loading capacity) was observed in **1d** (over **2d**) for each individual VOC.

Single crystal data was obtainable for the inclusion of the chlorinated VOCs in Table 3 in the framework of **1d**. These structures were previously reported¹⁸ and showed that the guest VOCs are included in the MOF channels and held in place by Cl $\cdots\pi$ and C–H $\cdots\pi$ interactions and by $\pi\cdots\pi$

interactions between chlorobenzene and the MOF walls (Fig. 3). The crystallographic information for these structures is reproduced in the ESI,† Table S1.

Unfortunately, single crystals could not be obtained for the remaining sorption products, but PXRD analysis confirms that only subtle changes in the frameworks are evident in all cases (Fig. S3†).

3.3 Sorption of iodine by **1d**: structure and solvatochromism

Using a similar process to that described above, we followed the sorption of iodine into **1d** over a period of two weeks using TGA and found that the maximum mass loss of 31% was reached after 8 days (Fig. 4). This corresponds to 1.6 iodine molecules per $[Co(34pba)(44pba)]$ unit.

The sorption process was characterized using PXRD where gradual changes of patterns up to eight days were observed, Fig. 5(top). Most notable is a gradual decrease in peak intensities at 8° and 15° while there was a gradual increase in the peak intensities at 9 and 22.5°. New peaks appear at 10.5 and 25°. These changes can be attributed to the interaction of iodine molecule and the channels of the **1d** framework (see crystal structure of **1dI₂** which is described below). The uptake of iodine could be followed visually (Fig. S4†).

The desorption of iodine from **1dI₂** into methanol could be monitored by following the colour changes that occur at a constant temperature of 22 °C (Fig. 6). The PXRD (Fig. 5 bottom) indicates the robustness of the phase changes during the recovery of iodine into methanol.

The sorption of iodine by **1d** was sufficiently slow that the crystal used retained its single crystallinity and allowed us to obtain a fully elucidated structure, **1dI₂** (Table 2 and Fig. 7).

Table 3 Uptake of selected solvents by the activated phases **1d** and **2d**

VOC	Experimental mass loss, TGA (%)	Temperature range of mass loss (°C)	Loading capacity, L_c (x in formula): $\{[M(34pba)(44pba)]_x \text{ solvent}\}_n$	ML_c	% loading capacity
1d					
CH ₂ Cl ₂ ^a	14.0	60–154	0.9	1.3	69
CH ₂ Br ₂	23.1	70–200	0.8	1.2	67
CH ₂ I ₂	33.3	70–260	0.8	1.0	80
CHCl ₃ ^a	17.1	118–285	0.8	1.0	80
CHBr ₃	19.2	91–236	0.4	1.0	40
CHI ₃	26.0	125–278	0.4	0.8	50
ClBenz ^a	13.0	87–264	0.6	0.8	75
BrBenz	19.2	38–235	0.7	0.8	88
IBenz	19.1	40–277	0.5	0.7	71
2d					
CH ₂ Cl ₂ ^a	11.0	88–220	0.7	1.4	50
CH ₂ Br ₂	23.0	78–266	0.8	1.2	67
CH ₂ I ₂	10.0	87–290	0.2	1.1	18
CHCl ₃ ^a	13.3	110–232	0.6	1.1	55
CHBr ₃	12.7	109–233	0.3	1.0	30
CHI ₃	10.0	105–248	0.1	0.9	11
ClBenz ^a	11.0	61–252	0.5	0.9	55
BrBenz	14.0	58–278	0.5	0.8	63
IBenz	10.6	58–270	0.3	0.8	38

^a Reported in ref. 18.



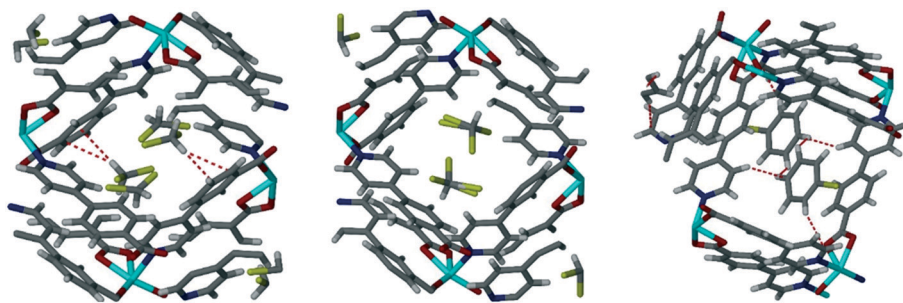


Fig. 3 Single crystal structures of chlorinated VOCs in **1d**, from left to right **1dCH₂Cl₂**, **1dCHCl₃**, **1dClBenz**. Non-covalent interactions are shown with dotted lines.¹⁸

In common with all previously reported structures in which the MOF contains guest molecules, the framework is perfectly ordered, and is isostructural with the structures in which the MOF contains dimethylformamide, dichloromethane, chloroform, or chlorobenzene. To quantify the amount of iodine present in **1dI₂**, we used the SQUEEZE routine in Platon²³ to estimate the void size and number of electrons present. For **1dI₂** this indicated solvent-accessible voids of 644 Å³ and 418 e⁻/unit cell, which corresponds to *ca.* 2 iodine molecules per [Co(34pba)(44pba)] unit (in good agreement with the TGA data). We were able to locate several disordered iodine molecules occupying the same cavity; these were modelled over three sites with total occupancy adding to 0.8 iodine molecules per [Co(34pba)(44pba)] unit. We note that residual electron density in the same area indicates the possibility of further iodine being present but we were unable to model this in a chemically meaningful way. For the iodine molecules we could model we can identify a number of non-covalent interactions of the form I...π and I...I between the iodine molecules and the channel walls and between iodine molecules themselves. These are shown in Fig. 7 and detailed in Table 4.

After the desorption in methanol, a single crystal was analyzed, to give the crystal structure **1dI₂d** (Table 2). **1dI₂d**

reverts to the same structure as **1d**, with identical disorder re-appearing in the pyridyl rings of the 34pba linker and in the benzoic acid rings of 44pba. Thus this material can be taken through successive desolvation, sorption, and desolvation cycles without significant changes.

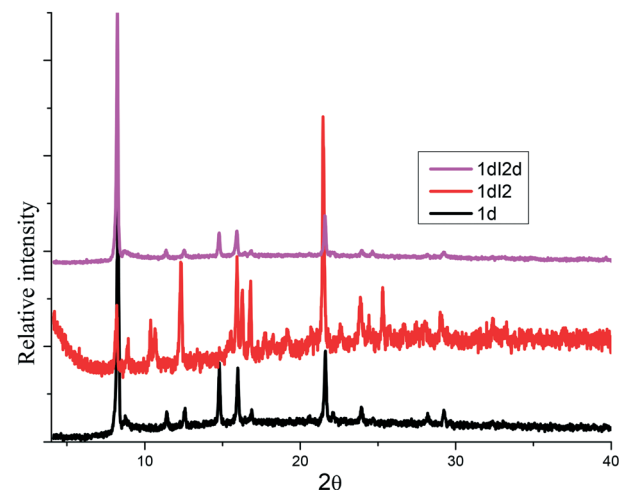
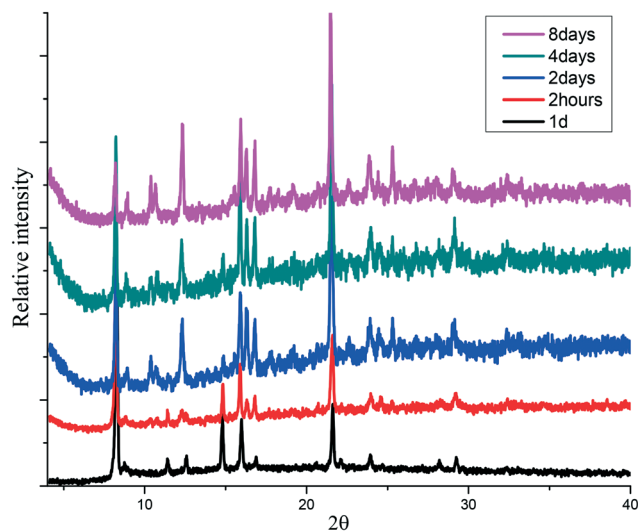


Fig. 5 (top) Gradual phase changes related to the sorption of iodine in **1d** to form **1dI₂**, (bottom) PXRD for iodine desorption from **1dI₂** into methanol to form **1dI₂d**.

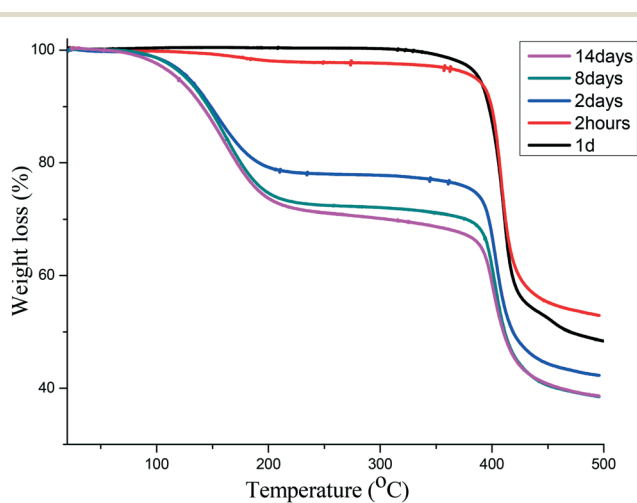


Fig. 4 TGA of iodine sorption by **1d** at different time intervals. The maximum is reached after 8 days.



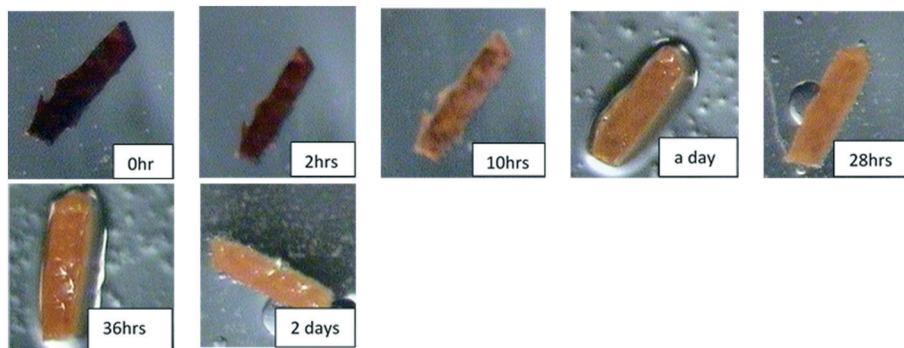


Fig. 6 Photographs showing colour change on the release of iodine from **1dI₂** into methanol.

We are now able to compare the similar structures obtained for the [Co(34pba)(44pba)] MOF **1**, which include six inclusion compounds and two desolvated compounds. As already reported, the compounds are isostructural, with no significant trends emerging in looking at unit cell parameters, though the *b*-axis shows the greatest variation, from 15.3 Å (in **1d**) to 17.8 Å (in **1DMF**). Conformational analysis shows that the rings of both linkers are close to coplanar for inclusion compounds where guests have larger molecular volumes and are more twisted when smaller guests (*e.g.* CH₂Cl₂) are included. When no guest is included (in **1d** or **1dI₂d**) the rings have even greater flexibility. This is particularly pronounced for the pyridyl ring of 34pba in which the major component of the disordered ring is rotated by *ca.* 70°. We noted that the other disordered ring (on the 44pba) aligns with its major component in the same orientation as in **1dCH₂Cl₂** and its minor component aligning with the other solvated compounds. These features are illustrated in Fig. 8. The flexibility of the MOF is confirmed by an analysis of the void space available in each instance

(Table S5[†]), which increases with solvent molecular volume. To confirm the robustness of the MOF an individual crystal was taken through several cycles of desorption and sorption, which showed consistent changes in unit cell parameters (Table S6[†]).

3.4 Kinetics of desorption for **1dCH₂Cl₂** and **1dCH₂Br₂** and **1dI₂**

While **1d** and **2d** show similar sorption trends for halogenated VOCs in terms of adsorption capacity and desorption temperature, (Fig. S2[†]), we noted that the TGA traces for halogenated VOCs of aliphatic compounds show a single desorption step. On the other hand, halogenated VOCs containing aromatic ring showed a two-step desorption process. This may be attributed to the aromatic stacking interactions as seen the **1dClBenz** structure (Fig. 3) which cause the guests to be more strongly retained and result in a larger temperature range for desorption.

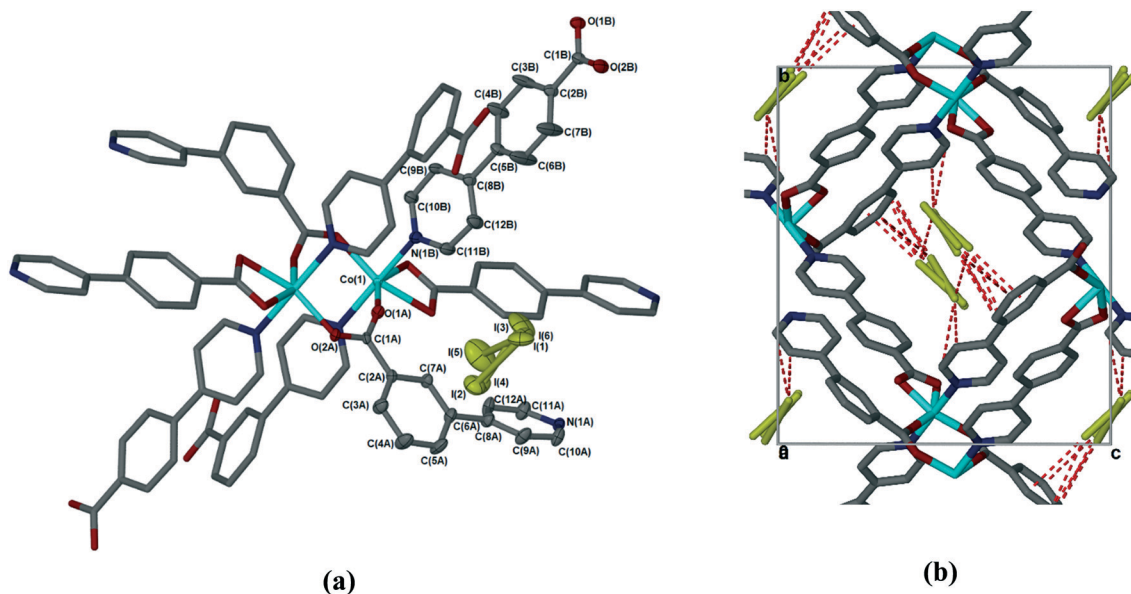


Fig. 7 (a) Coordination geometry and SBU in **1dI₂**. The asymmetric unit is shown with thermal ellipsoids at the 50% probability level. (b) Weak interactions (dotted lines) between the iodine guest molecule and MOF in **1dI₂**. Hydrogen atoms have been omitted for clarity.



Table 4 Non-covalent interactions in **1dI₂**

Atoms	Interatomic distance (Å)
I2–C4A	3.566
I2–C5A	3.628
I3–C9A	3.604
I3–C10A	3.636
I4–C4A	3.533
I4–C5A	3.460
I4–C6A	3.662
I5–C5A	3.820
I5–C6A	3.625
I3–I5	3.875

TGA can also be used to determine the activation energy (E_a) of guest desorption processes, and we have applied this here to the desorption of the related guests dichloromethane and dibromomethane from **1dCH₂X₂**, as well as to the desorption of iodine from **1dI₂**. A series of isothermal TG curves were obtained for each compound at several temperatures just below the desorption temperature observed in Fig. S2.† Each desorption curve was then fitted against a range of plausible desorption mechanisms,²⁴ and a rate constant (k) was determined at each temperature (Fig. S5†). An Arrhenius plot then gave the value of the activation energy for each desorption process. The conversion vs. time curves are shown in Fig. S6,† and the activation energies determined are given in Table 5. The kinetics of desorption for CH₂Cl₂, CH₂Br₂, and I₂ all showed a best fit to the 3D diffusion model, which indicates that the guest molecules interact with the wall of the adsorbent in a spherical zone. The E_a of desorption of CH₂Cl₂ and CH₂Br₂ are approximately equal while that for iodine is higher, probably owing to the increased non-covalent interactions which iodine is capable of making with the channel walls.

3.5 Sorption of carbon dioxide and hydrogen gases in **1d** and **2d**

Activated **1d** and **2d** were investigated for the sorption of CO₂ and H₂ at different temperatures. The heat of adsorption of CO₂ was also determined. Fig. 9 shows the adsorption traces for selected runs using both **1d** and **2d**. At the temperatures

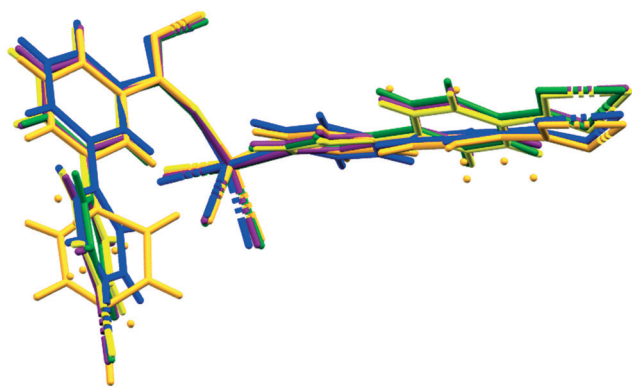


Fig. 8 Molecular overlays of the asymmetric unit for **1d** (orange), **1dCH₂Cl₂** (blue), **1dCHCl₃** (green), **1dClBenz** (yellow), **1dI₂** (purple).

Table 5 Activation energy for the isothermal desorption of CH₂Cl₂, I₂, and CH₂Br₂ from **1d**

Compound	E_a (kJ mol ⁻¹)	R^2 Coeff
1dCH₂Cl₂	70	0.96
1dCH₂Br₂	60	0.97
1dI₂	77	0.90

studied, the sorption behaviour was a type-Ib isotherm. In this model of sorption, the adsorbent is characterized by small external surfaces having a pore size with distributions over a broader range and the presence of wider mesopores with possibility of narrow micropores (≤ 2.5 nm).²⁵ The steep uptake observed at low p/p_0 is due to enhanced adsorbent–gas interactions in narrow micropores. Table 6 records the higher adsorption capacity at 195 K and 676 mmHg for both samples. **1d** showed more than a double 114 cm³ (STP) g⁻¹ (2.35 molecules per ASU) compared to that of **2d** with 52 cm³ (STP) g⁻¹ (1.11 molecules per ASU). Little hysteresis was observed in both isostructural frameworks at the lower temperature (195 K) owing to the lower mobility of gas molecules. Stage A (Fig. 9 top) may indicate the presence of some mesopores in **1d** which are absent in **2d**. Furthermore, this stage can also explain the flexibility of the ligand to open the pores and allow additional gas uptake.²⁶ The highest adsorption recorded at 298 K and 910 mmHg for **1d** and **2d** was 41.0 cm³ (STP) g⁻¹ (0.84 molecules per ASU) and 17.7 cm³ (STP) g⁻¹ (0.37 molecules per ASU), respectively. Sample

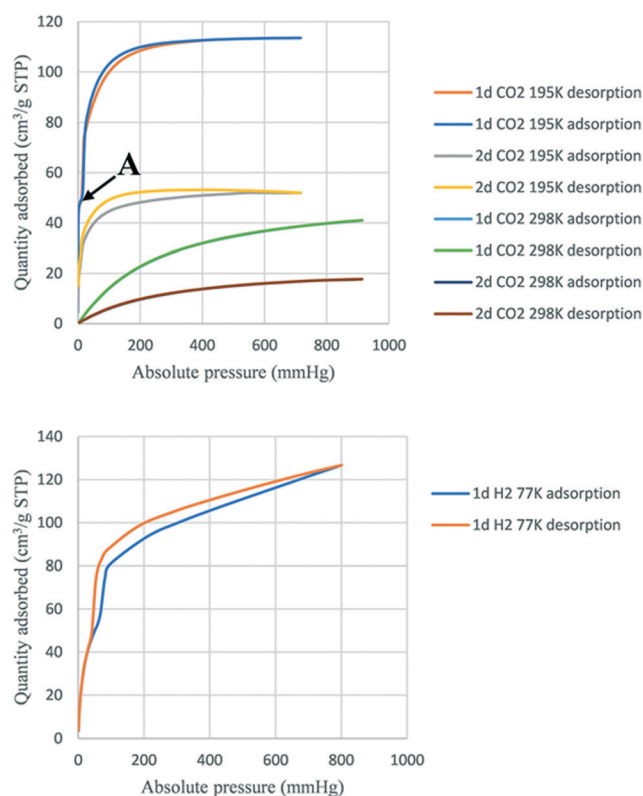


Fig. 9 Sorption of CO₂ (top) and H₂ (bottom).



Table 6 Sorption capacity for CO₂

No	Sample	Volume adsorbed (cm ³ g ⁻¹)	Pressure (mmHg)	Temperature (K)	Corresponding to (mmol g ⁻¹)
1	2d	52.0	676	195	2.40
	1d	114			5.06
2	2d	22.4	784	273	1.00
	1d	47.6			2.10
3	2d	21.7	803	278	1.00
	1d	47.0			2.10
4	2d	19.7	910	288	0.90
	1d	44.6			2.00
5	2d	18.8	900	293	0.84
	1d	43.1			1.92
6	2d	17.7	910	298	0.80
	1d	41.0			1.80

1d showed higher adsorption capacity than **2d** at a similar temperature and pressure.

The trend of higher adsorption capacity in **1d** compared to **2d** was observed for solvent vapours including the halogenated VOCs reported here and the volatile amines reported in ref. 18, and may be the result of a slightly smaller void space in **1d** than **2d**. However, this attribution of adsorption capacity related to the size of pores has been controversial in other reported structures.²⁶ An alternative explanation is that the smaller size of a solvent-accessible void volume causes stronger interactions between the guest and the framework, depending on their respective structures and functionalization. The determination of isosteric heat of adsorption (Q_{st}) of CO₂ showed that the sample **1d** adsorbed between 0.8 mmol and 1.38 mmol with heats of adsorption (Q_{st}) between 29.8 kJ and 30.3 kJ, while sample **2d** adsorbed between 0.1 mmol and 0.4 mmol CO₂ (Q_{st} 28.5 kJ and 28.9 kJ). These results suggest that there is a higher interaction in **1d** than in **2d**.²⁷ The uptake for hydrogen gas in **1d** was 120 cm³ (STP) g⁻¹ at 800 mmHg at 77 K corresponding to 2.44 molecules of H₂ per ASU. In contrast, **2d** did not adsorb hydrogen under the same conditions.

4. Conclusion

The double-walled [Co(34pba)(44pba)]_n-MOF (**1d**) was studied for its ability to take up vapours and gases from the desolvated, activated phase. This MOF has proved to be robust to several cycles of sorption and desorption, as evidenced by both single-crystal and powder diffraction. This is facilitated by the flexibility of the pyridylbenzoate linkers, in particular the ability of the rings to rotate and thus occupy additional space when the compound is desolvated. A similar effect was observed for the Zn(II) analogue (**2d**).

Inclusion of halogenated volatile organic compounds found that, for each series of VOCs studied, the uptake was generally higher for chlorinated over brominated and iodinated compounds. While only the chlorinated species CH₂Cl₂, CHCl₃, and ClBenz afforded single crystal structures, TGA and PXRD analysis indicates that the type of interaction present in the Br-

and I-derivatives are likely to be similar. On the other hand, the sorption of iodine into **1d** could be followed in the powder form, and also afforded single crystals which are stabilized by I...π and I...I interactions. The desorption of iodine from **1d**₂ gave **1dI₂d** whose structure was also fully elucidated, confirming that the desorption–sorption–desorption cycle proceeded without significant loss of crystallinity.

The isothermal kinetic desorption of I₂, CH₂Cl₂, and CH₂Br₂ from **1d** indicates that iodine requires more energy for the desorption in the range of the reported ones. The two adsorbents **1d** and **2d** were both capable of adsorption of carbon dioxide gas while only **1d** adsorbed hydrogen. In general, across all types of vapour and gas sorption tested, the trend was for **1d** to show stronger interactions and higher adsorption capacity. Further studies to determine the reasons for this observation are underway.

Author contributions

Conceptualization S. A. B.; data acquisition and analysis C. A. N.; assistance with crystallography and analysis S. A. B., C. L. O.; assistance with gas sorption N. C.; supervision S. A. B., C. L. O.; writing – original draft C. A. N., S. A. B.; all authors assisted with writing – review and editing.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 B. Wang, P. Wang, L.-H. Xie, R.-B. Lin, J. Lv, J.-R. Li and B. Chen, *Nat. Commun.*, 2019, **10**, 3861.
- 2 N. Gcwensa, N. Chatterjee and C. L. Oliver, *Inorg. Chem.*, 2019, **58**, 2080.
- 3 S. Huh, *Catalysts*, 2019, **9**, 34.
- 4 X.-D. Yang, C. Chen, Y.-Z. Zhang, L.-X. Cai, B. Tan and J. Zhang, *Dalton Trans.*, 2016, **45**, 4522.
- 5 D. N. Jiang, C. Huang, J. Zhu, P. Wang, Z. M. Liu and D. Fang, *Coord. Chem. Rev.*, 2021, **444**, 214064.
- 6 K. Hyojin and C. S. Hong, *CrystEngComm*, 2021, **23**, 1377.
- 7 X. Yang and D. Yan, *Chem. Sci.*, 2016, **7**, 4519.
- 8 X. Yang and D. Yan, *Chem. Commun.*, 2017, **53**, 1801.
- 9 M. Kobalz, J. Lincke, K. Kobalz, O. Erhard, J. Bergmann, D. Lässig, M. Lange, J. Möllmer, R. Gläser, R. Staudt and H. Krautscheid, *Inorg. Chem.*, 2016, **55**, 3030.
- 10 G. Mehlana, S. A. Bourne, G. Ramon and L. Öhrström, *Cryst. Growth Des.*, 2013, **13**, 633.



- 11 L. Mei, C. Wang, L. Wang, Y. Zhao, Z. Chai and W. Shi, *Cryst. Growth Des.*, 2015, **15**, 1395.
- 12 L. Wang, Y. Li, F. Yang, Q. Liu, J.-P. Ma and Y.-B. Dong, *Inorg. Chem.*, 2014, **53**, 9087.
- 13 Q.-Y. Yang, P. Lama, S. Sen, M. Lusi, K.-J. Chen, W.-Y. Gao, M. Shivanna, T. Pham, N. Hosono, S. Kusaka, J. J. Perry, S. Ma, B. Space, L. J. Barbour, S. Kitagawa and M. J. Zaworotko, *Angew. Chem., Int. Ed.*, 2018, **57**, 5684.
- 14 K. Davies, S. A. Bourne, L. Öhrström and C. L. Oliver, *Dalton Trans.*, 2010, **39**, 2869.
- 15 G. Mehlana, S. A. Bourne and G. Ramon, *CrystEngComm*, 2014, **16**, 8160.
- 16 N. A. Khan, Z. Hasan and S. H. Jhung, *J. Hazard. Mater.*, 2013, **244–245**, 444.
- 17 A. Bacchi, S. A. Bourne, G. Cantoni, S. A. M. Cavallone, S. Mazza, G. Mehlana, P. Pelagatti and L. Rhighi, *Cryst. Growth Des.*, 2015, **15**, 1876.
- 18 C. A. Ndamyabera, S. C. Zacharias, C. L. Oliver and S. A. Bourne, *Chemistry*, 2019, **1**, 111.
- 19 G. M. Sheldrick, *SADABS, Version 2.05*, University of Göttingen, Germany, 2007.
- 20 G. M. Sheldrick, *Acta Crystallogr., Sect. C: Struct. Chem.*, 2015, **71**, 3.
- 21 L. J. Barbour, *J. Appl. Crystallogr.*, 2020, **53**, 1141.
- 22 C. F. Macrae, I. J. Bruno, J. A. Chisholm, P. R. Edgington, P. McCabe, E. Pidcock, L. Rodriguez-Monge, R. Taylor, J. van de Streek and P. A. Wood, *J. Appl. Crystallogr.*, 2008, **41**, 466.
- 23 A. L. Spek, *Acta Crystallogr., Sect. C: Struct. Chem.*, 2015, **71**, 9.
- 24 A. Khawam and D. R. Flanagan, *J. Phys. Chem. B*, 2006, **110**, 17315.
- 25 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.
- 26 T. K. Trung, N. A. Ramsahye, P. Trens and N. Tanchoux, *Microporous Mesoporous Mater.*, 2010, **134**, 134.
- 27 N. Chatterjee and C. L. Oliver, *Cryst. Growth Des.*, 2018, **18**, 7570.

