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Self-adjusting binding pockets enhance H₂ and CH₄ adsorption in a uranium-based metal–organic framework†

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A new, air-stable, permanently porous uranium(IV) metal–organic framework U(bdc)₂ (**1**, bdc²⁻ = 1,4-benzenedicarboxylate) was synthesized and its H₂ and CH₄ adsorption properties were investigated. Low temperature adsorption isotherms confirm strong adsorption of both gases in the framework at low pressures. *In situ* gas-dosed neutron diffraction experiments with different D₂ loadings revealed a rare example of cooperative framework contraction ($\Delta V = -7.8\%$), triggered by D₂ adsorption at low pressures. This deformation creates two optimized binding pockets for hydrogen ($Q_{st} = -8.6$ kJ mol⁻¹) per pore, in agreement with H₂ adsorption data. Analogous experiments with CD₄ ($Q_{st} = -24.8$ kJ mol⁻¹) and *N,N*-dimethylformamide as guests revealed that the binding pockets in **1** adjust by selective framework contractions that are unique for each adsorbent, augmenting individual host–guest interactions. Our results suggest that the strategic combination of binding pockets and structural flexibility in metal–organic frameworks holds great potential for the development of new adsorbents with an enhanced substrate affinity.

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Introduction

Metal–organic frameworks are a class of chemically-robust, porous, and often rigid materials, composed of metal ions or clusters connected by bridging organic linkers.^{1–4} The physical and chemical properties of these materials are highly tunable based on choice of metal and linker, and thus metal–organic frameworks have been proposed for a wealth of applications,^{5–9} including catalysis,^{10–15} sensing,^{16–18} carbon capture,^{19–23} gas separations,^{24–26} and gas storage.^{27–31} Metal–organic frameworks have attracted particular interest as candidate gas storage

materials for H₂ and CH₄ that could enable more efficient use of these energy carriers as cleaner fuel alternatives.^{32–38} However, significant advances are still needed to develop frameworks capable of maintaining interactions with these guests at ambient temperatures.^{39–41}

Two main strategies have been developed to achieve strong binding of H₂ and CH₄ in metal–organic frameworks. The first approach utilizes materials with coordinatively-unsaturated metal sites, which can polarize and strongly bind various guests.^{42,43} Representative of this materials class is the framework Ni₂(*m*-dobdc) (*m*-dobdc⁴⁻ = 4,6-dioxido-1,3-benzenedicarboxylate), which is currently the top performing material for ambient temperature, physisorptive H₂ storage.^{33,34} The other strategy exploits tight binding pockets in small-pore frameworks, which can engage in multiple, weak interactions with guest molecules to achieve strong overall guest binding, analogous to shape-selective molecular recognition in enzymes.⁴⁴ An example of how such cumulative dispersion forces can outperform strong interactions at open metal sites is the adsorption of CH₄ in Cu₂(btc)₃ (HKUST-1, btc³⁻ = 1,3,5-benzenetricarboxylate).⁴⁵ This material exhibits open metal sites and binding pockets in direct competition for CH₄ adsorption. Structural characterization of Cu₂(btc)₃ dosed with low pressures of CD₄ confirmed that methane preferably adsorbs at the binding pockets inside small octahedral cages of the framework, rather than through direct interactions at the copper(II) open metal sites. The reason for this behavior is that the multiple interactions inside the pore give rise to a higher

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overall binding energy than that achieved with a CH₄ molecule adsorbed at a single copper(II) center (-21.8 versus -9.4 kJ mol⁻¹, respectively).⁴⁵

Cumulative dispersion interactions between guest molecules and framework pockets decrease exponentially with the adsorbate–framework distances ($F \propto 1/r^6$), and therefore require a precise geometric fit between guest and binding pocket.⁴⁶ For example, as a result of its smaller kinetic diameter relative to CH₄,⁴⁷ H₂ preferentially binds at the open metal sites of Cu₂(btc)₃, rather than in the hexagonal pockets.⁴⁸ The development of new frameworks with efficient binding pockets therefore requires precise optimization for each adsorbate of interest, although achieving this goal by structural design remains a significant challenge.

An alternative approach to circumvent the synthetic intricacy of developing materials with optimized guest-specific binding pockets, are materials that combine small binding pockets with moderate framework flexibility.⁴⁹ Synthetic tuning can thus be used to design crude binding pockets, which are capable of self-adjusting in response to guest adsorption. Together, these design features could enable access to optimal binding pocket geometries for a variety of guests within the same material. Such molecular recognition often relies on initially weak host-guest interactions, highlighting the importance to precisely adjust the energy required for the deformation of a flexible framework and the energy released by guest adsorption.^{50–52}

Flexibility is typically introduced into metal–organic frameworks by utilizing organic linkers with non-rigid stems, by interconnecting metals with non-chelating linkers, or by cross-linking two-dimensional frameworks with additional ditopic but weakly binding linkers.^{53–55} Prominent examples are M(OH)(bdc) (MIL-53; bdc²⁻ = 1,4-benzenedicarboxylate; M = Fe, Cr, Sc, Al, or Ga)^{56–60} and M₃(O)(OH)(H₂O)₂(bdc)₃ (MIL-88; M = Fe, Cr).⁶¹ These frameworks undergo drastic geometric distortions upon guest adsorption, often referred to as framework swelling, which can induce a substantial unit cell volume increase of up to 74%, as shown for example by CO₂ adsorption in Fe(OH)(bdc).⁶² Such large structural changes are too extreme to drive the subtle binding pocket adjustments sought here. One could instead envision limiting the flexibility of non-chelating bdc²⁻ linkers by substantially increasing the number of metal–ligand bonds per metal node. A higher coordination number should limit structural rearrangements by causing steric encumbrance around the metal nodes and increase rigidity by further crosslinking the resulting material. Additionally, a higher ligand-to-metal ratio could result in smaller pore sizes and better binding pockets.

With their tendency to adopt high coordination numbers, actinides are well suited as metal nodes for the development of such materials.⁶³ We chose depleted uranium to test our hypothesis, as it is only mildly radioactive and because a limited but growing number of uranium-based frameworks have already been reported and could guide the synthesis.⁶⁴ Inspired by previous work on the synthesis of porous metal–organic frameworks from uranium(IV) and bdc²⁻ linkers,⁶⁵ we synthesized a new, three-dimensional U(bdc)₂ phase (**1**) with permanent porosity and a moderate level of structural flexibility. Using

a combination of gas adsorption studies and *in situ* powder neutron diffraction experiments, we demonstrate that this framework undergoes an adjustable contraction of its pores to accommodate and strongly bind H₂ and CH₄, with different levels of contraction and host-guest interactions for each molecule.

Results and discussion

The compound U(bdc)₂·4H₂O (**1**-H₂O) was synthesized through the reaction of U₄(1,4-dioxane)₂ with H₂bdc in *N,N*-dimethylformamide (DMF, <0.15% water content as received) at 140 °C under argon inside a Parr autoclave. After three days, the material was isolated in 79% yield as air-stable, thin emerald green needle-shaped crystals. Single crystal X-ray diffraction analysis was used to determine the structure of **1**-H₂O (Fig. 1), and selected bond distances and angles are given in Table S5 of the ESI.† We note that powder X-ray diffraction patterns collected for bulk samples of **1**-H₂O match the simulated pattern determined from single-crystal data, confirming the bulk purity of the crystalline material (see ESI, Fig. S7†).

Compound **1**-H₂O crystallizes in the space group *C2/c* and features eight-coordinate uranium centers in a distorted square-antiprismatic environment. Each uranium(IV) is



Fig. 1 (a) Single crystal X-ray diffraction structure of **1**-H₂O viewed along the *c*-axis, showing the parallelepipedal pores. (b) Truncated structure showing one of the pores of **1**-H₂O along the crystallographic *c*-axis, with the two identical binding pockets of the pore depicted as blue spheres. (c) The same view as in (b), rotated by 90° to visualize the bowl-shaped arrangement of three bdc²⁻ linkers that form the cap of each binding pocket. Orange, red, and grey spheres represent U, O, and C atoms, respectively; solvent and H atoms are omitted for clarity.



coordinated to one oxygen atom of eight different bdc^{2-} linkers, and all linkers are coordinated to four different uranium ions in a bridging fashion. This motif results in an overall framework structure consisting of distorted parallelepipedal pores (Fig. 1a) formed by chains of uranium(IV) centers that propagate along the crystallographic c -axis (see ESI, Fig. S17†) and are bridged by bdc^{2-} linkers bent in a concave and convex fashion.

Each pore is formed by two opposing, inwardly bent bdc^{2-} linkers at the sides and is capped at the top and bottom by a bowl-shaped arrangement of three additional bdc^{2-} linkers (Fig. 1b and c). The resulting geometry yields two identical binding pockets per pore that are ~ 5 Å in diameter and related by an inversion center.

In the as-synthesized framework, each binding pocket is occupied by two disordered water molecules, yielding the composition $\text{U}(\text{bdc})_2 \cdot 4\text{H}_2\text{O}$, which was also confirmed by thermogravimetric analysis (see ESI, Fig. S6†). While disorder precluded modeling of any specific interactions, the guest water molecules are likely to engage in hydrogen bonding with each other and with the highly polarized U–O bonds. We note that the structure of the pores is such that guests could engage in a variety of additional interactions, including with the linker π -systems and arene C–H bonds.

Activated $\text{U}(\text{bdc})_2$ (**1**) was obtained by heating **1**– H_2O at 260 °C for 10 h under dynamic vacuum. Combustion analysis confirmed the empirical formula for **1** and the removal of guest water molecules. Nitrogen adsorption isotherms obtained at 77 K for four different samples revealed the activated material is permanently porous, with an average Langmuir surface area of $497 \pm 6 \text{ m}^2 \text{ g}^{-1}$ (see ESI, Fig. S1†). Powder X-ray diffraction analysis confirmed that **1** remains crystalline with a slightly different structure from that of its solvated analogue (see ESI, Fig. S7†). Interestingly, while many flexible frameworks contract or even fully collapse to a nonporous structure upon solvent removal,^{37,66–68} activation of **1**– H_2O to give **1** results in an expansion of the framework along the crystallographic b -axis, from 12.598(1) to 12.812(1) Å. This change results in an increase in the unit cell volume from 1887.4(3) to 1902.1(2) Å³, while retaining the $C2/c$ space group. This behavior upon guest removal indicates that the framework binding pockets are indeed able to contract to improve interactions with adsorbates.

We sought to study the flexibility of **1** in more detail using H_2 and CH_4 (with kinetic diameters of 2.9 and 3.8 Å, respectively)¹⁸ as probe molecules of interest for potential gas storage applications. The low-pressure H_2 adsorption isotherm for $\text{U}(\text{bdc})_2$ at 77 K exhibits an initial steep rise to $\sim 3.5 \text{ mmol g}^{-1}$ at 115 mbar, which is indicative of the presence of strong adsorption sites (Fig. 2). We note that this loading corresponds to the theoretical capacity expected for adsorption of one H_2 molecule per adsorption pocket (two per pore). With further increasing pressure, the quantity of adsorbed H_2 increases very gradually to an apparent saturation value of $\sim 4.9 \text{ mmol g}^{-1}$ at 1.2 bar. A dual site Langmuir model was used to fit independently H_2 adsorption data collected at 77 and 87 K (see Section 3 of the ESI, Fig. S2 and Table S1†), and the Clausius–Clapeyron equation was then employed to calculate the isosteric heat (Q_{st}) of H_2 adsorption as a function of loading (Fig. 2, inset). For loadings



Fig. 2 Hydrogen adsorption isotherm for **1**, measured at 77 K. Inset: loading-dependent isosteric heat of adsorption (Q_{st}) for H_2 in **1**.

up to 2.5 mmol g^{-1} , H_2 adsorbs exclusively at primary binding sites in the framework pockets (see below) to give a Q_{st} of -8.6 kJ mol^{-1} . Notably, this value is larger in magnitude than the H_2 isosteric heat of adsorption in activated carbon materials (-5.0 to -6.4 kJ mol^{-1})⁶⁹ and the majority of frameworks with coordinatively-saturated metal sites (-4.1 to -8.8 kJ mol^{-1}).⁷⁰

The CH_4 adsorption isotherm for **1** obtained at 195 K exhibits a steep uptake similar to that characterized for H_2 at low pressures, again indicative of strong interactions between CH_4 and the binding pockets of the framework (Fig. 3). A dual site Langmuir model was used to simultaneously fit isotherm data collected at 195, 273, 298, and 308 K, and the Clausius–Clapeyron equation was then employed to calculate a value of $Q_{\text{st}} = -24.8 \text{ kJ mol}^{-1}$ at low loadings (see ESI, Fig. S5†). Notably, this value surpasses that determined for $\text{Zn}_4\text{O}(\text{bdc})_3$ (MOF-5; $Q_{\text{st}} = -12.3 \text{ kJ mol}^{-1}$) and even values for frameworks with snugly fitting pore window, such as $\text{Cu}_2(\text{btc})_3$ ($Q_{\text{st}} = -17.1 \text{ kJ mol}^{-1}$), or strongly polarizing open metal cation sites, as in $\text{Ni}_2(\text{dobdc})$ ($\text{dobdc}^{4-} = 2,5\text{-dioxido-1,4-benzenedicarboxylate}$; $Q_{\text{st}} = -20.6 \text{ kJ mol}^{-1}$).^{41,45} It is clear that the binding pockets in



Fig. 3 Methane adsorption isotherms for **1**, measured at the indicated temperatures.



U(bdc)₂ can strongly interact with both H₂ and CH₄, despite their different sizes, which suggests that the framework may distort or flex to optimize interactions with different guest molecules. In order to study the framework-guest interactions in more detail, we turned to *in situ* gas-dosing powder neutron diffraction. The powder neutron diffraction pattern of activated **1** at 9 K was first collected as a reference for gas dosing experiments (see ESI, Fig. S9†). Dosing with 0.3 equiv. of D₂ per pore results in clear changes in the powder pattern, particularly visible at low values of scattering angle 2θ (Fig. 4 and S10†). Specifically, the reflections of **1** decrease in intensity while a second set of peaks arises, ascribed to a new crystalline phase **1-D₂**. Upon increasing the loading to 0.7 equiv. of D₂, both phases are still present, although the peaks of **1** diminish further and the peaks of newly formed **1-D₂** gain in intensity (Fig. 4 and S11†). The two phases coexist up to a loading of at least 1.5 equiv. D₂ per pore (see ESI, Fig. S12†), and their interconversion is best followed by evaluating the high intensity, low angle peaks at $2\theta \approx 11.85^\circ$ for **1** and at $2\theta \approx 12.63^\circ$ for **1-D₂** in Fig. 4. These data strongly suggest that cooperative effects drive an adsorbate-induced framework distortion from **1** to **1-D₂**. Such a mechanism is in contrast to a gradual and homogeneous uptake of D₂, or a gradually changing degree of distortion depending on the D₂-loading. As a result, before achieving saturation loading, some individual crystallites of the sample will be distorted, such that both binding pockets per pore are occupied with a D₂ molecule, while others will remain in the activated structure of **1**. At the highest D₂ dosing level of 2.5 equiv. per pore, the diffraction pattern of the sample contains only reflections associated with **1-D₂** (see ESI, Fig. S13†).

In situ powder neutron diffraction experiments were also carried out by dosing **1** with 0.7 and 1.5 equiv. of CD₄ (Fig. S14 and S15†). The data from these experiments suggest a similar cooperative transformation from **1** to an adsorbed phase **1-CD₄**.



Fig. 4 Rietveld refinement fits of powder neutron diffraction patterns ($\lambda = 2.0772 \text{ \AA}$, $T = 9 \text{ K}$) collected for activated **1** and activated **1** dosed with 0.3, 0.7, and 1.5 equiv. of D₂ per pore. Black (lower) and blue (upper) tick marks indicate calculated Bragg peak positions for **1** and the D₂ adsorbed phase, **1-D₂**, respectively. Arrows indicate peaks that best illustrate the conversion of **1** to **1-D₂** with increasing D₂ loading.

Rietveld refinements were applied to all powder neutron diffraction data (see Section 6 in the ESI†) in order to elucidate adsorbate-induced structural deformations and characterize specific adsorption sites for D₂ and CD₄. Selected unit cell parameters determined for the different structures are summarized in Table 1. Based on unit cell volume, **1** contracts to a greater extent to accommodate D₂ than it does in the presence of CH₄.

Surprisingly, a structural comparison of **1** with **1-D₂** and **1-CD₄** reveals an almost constant coordination environment around the uranium nodes in each phase. The framework flexibility instead relies on a tilting of the parallelepipedal pores, which is clearly seen by comparing the pore geometries of **1** and **1-D₂** dosed with 2.5 equiv. D₂, as shown in Fig. 5a. Here, each structure is overlaid with an idealized parallelogram with corners defined by the uranium ions. Dosing with 2.5 equiv. of D₂ results in a change of the idealized parallelogram angles, φ_1 and φ_2 , from $77.1(1)^\circ$ and $102.902(3)^\circ$ in **1** to $65.03(6)^\circ$ and $114.973(1)^\circ$ in **1-D₂**. In order to accommodate this rearrangement, the two unique dihedral angles, ω_1 and ω_2 , between the bdc^{2-} O-C-O planes and the neighboring O-U...U-O planes adjust from $151(1)^\circ$ and $168.4(9)^\circ$ in **1** to $138.5(4)^\circ$ and $174.9(5)^\circ$ in **1-D₂** (Fig. 5b).

The resulting hinge-type bending between the UO₈ nodes and linkers is analogous to the change that occurs in the structure of the flexible framework Cr(OH)(bdc) upon water adsorption.⁵³ In particular, water adsorption is accompanied by a unit cell volume decrease from 1486.1 to 1012.6 Å³, as well as a decrease of the symmetrical dihedral angles, ω , from 179.8° to 162.3° . The analogous idealized Cr...Cr...Cr angles φ_1 and φ_2 in



Fig. 5 (a) Illustration of pores in **1** and **1-D₂** (dosed with 2.5 equiv. D₂ per pore). The pore contraction upon D₂ dosing is highlighted by the colored blue and purple areas within the crystallographic *ab*-plane. Tilting of the pores to facilitate contraction is illustrated by idealized parallelograms (dashed lines) with corners defined by uranium atoms. Adsorbed D₂ molecules inside the pores of **1-D₂** are represented as aquamarine spheres for adsorption site I and as brown spheres for adsorption site II. Each D₂ super atom at site II is shared between two neighboring pores, accordingly the two depicted site II super atoms together account for an occupancy of one D₂ molecule per pore. (b) Comparison of the dihedral angles ω_1 as described in the text (ω_2 is not shown, but is the corresponding angle at the other end of the linker), representing the main structural distortion undergone by **1** upon adsorption of D₂. Orange, red, and grey spheres represent U, O, and C atoms, respectively.



Table 1 Unit cell parameters (space group $C2/c$) for activated **1** and **1** dosed with different adsorbates, along with selected angles describing the pore deformation upon guest adsorption. Angles φ_1 and φ_2 describe the $U\cdots U\cdots U$ angles of the idealized parallelogram spanned by the four corner uranium atoms within the crystallographic ab -plane of each pore, as illustrated in Fig. 5a. Angles ω_1 and ω_2 describe the dihedral angle between $O-U\cdots U-O$ planes and $O-C-O$ planes at each end of a bdc^{2-} linker, as illustrated in Fig. 5b for **1** and **1-D₂**. One standard deviation for all values is given in parentheses

Sample	a (Å)	b (Å)	c (Å)	V (Å ³)	φ_1 (°)	φ_2 (°)	ω_1 (°)	ω_2 (°)
1	17.587(1)	12.812(1)	9.2999(5)	1902.1(2)	77.1(1)	102.902(3)	151(1)	168.4(9)
1-D₂ (0.3 equiv.)	18.361(7)	11.35(1)	9.335(2)	1782(2)	67.51(1)	112.491(9)	148.7(3)	159.6(3)
1-D₂ (0.7 equiv.)	18.402(3)	11.324(5)	9.333(1)	1781.4(8)	67.4(2)	112.576(1)	130(2)	175(3)
1-D₂ (1.5 equiv.)	18.456(1)	11.233(1)	9.3508(4)	1775.8(2)	66.86(6)	113.143(1)	141.4(8)	170(1)
1-D₂ (2.5 equiv.)	18.665(1)	10.9486(8)	9.3838(5)	1754.5(2)	65.03(6)	114.973(1)	138.5(4)	174.9(5)
1-CD₄ (1.5 equiv.)	18.031(1)	11.9665(7)	9.3206(4)	1839.3(2)	71.52(5)	108.483(1)	142(3)	171.1(9)
1-DMF	18.2658(7)	12.0252(5)	9.3579(3)	1865.6(1)	71.63(4)	108.373(1)	166(11)	164(4)

Cr(OH)(bdc) change drastically from 75.9° and 104.2° (activated) to 44.8° and 135.2° (hydrated).⁵⁶ Interestingly, in distinct contrast to **1**, Cr(OH)(bdc) distorts very little upon interaction with D₂ (<4 equiv. per pore), adopting a symmetric dihedral angle $\omega = 178.4^\circ$ and Cr \cdots Cr \cdots Cr angles of 80.6° and 99.4°, concomitant with a very small unit cell volume increase to 1534.5 Å³.^{58,71} The change in the structure of **1** upon dosing with 2.5 equiv. of D₂ results in a much more drastic change in unit cell volume, from 1902.1(2) to 1754.5(2) Å³. We rationalize that the greater deformation of **1** arises as a result of its better ability to enshroud H₂ within its pores, which leads to a greater adsorption enthalpy ($Q_{st} = -8.6$ kJ mol⁻¹ vs. -6.9 kJ mol⁻¹ for Cr(OH)(bdc)),⁷² and a larger driving force for structural rearrangement. Thus, the smaller pores within the framework of **1** are able to optimize binding through multiple stabilizing interactions, whereas the comparably large pores of Cr(OH)(bdc) provide fewer contacts.

In order to elucidate the hydrogen binding sites in **1**, we treated the D₂ molecules as “super atoms” in our analysis of the diffraction data (see ESI, Section 6†).^{73,74} We first analyzed the structure of **1** loaded with 1.5 equiv. of D₂ per pore (corresponding to less than one D₂ per pocket) to enable an accurate structure determination in the absence of adsorbate–adsorbate interactions.^{75–77} As expected, the D₂ super atoms were located in both binding pockets of each pore, with an occupancy of 75% per site. Each D₂ super atom is situated within van der Waals contact distance of three H atoms of bdc^{2-} linkers, the π -system of the outer pocket-capping bdc^{2-} linker, and two oxygen atoms of the UO₈ coordination polyhedron (Fig. 6a). The D₂ \cdots H contact distances of 2.96(1), 2.98(1), and 3.15(2) Å indicate moderately strong van der Waals interactions.^{74,78,79} The distance from D₂ to the centroid of the nearest benzene ring is 3.45(1) Å, which is indicative of a modest D₂ \cdots π interaction,⁸⁰ while the closest D₂ \cdots O contact is 3.59(1) Å. In the structure of **1** dosed with 2.5 equiv. of D₂, an additional D₂ molecule was located in the center of the pore (site II, see Fig. 5a). The D₂ molecules at site II are stabilized by four symmetry equivalent D₂ \cdots D₂ interactions at a distance of 3.10(1) Å, as well as by weak C–H \cdots D₂ contacts with the linkers (3.31(3) and 3.55(1) Å). As discussed above, in order to accommodate these interactions, the pores of **1** contract significantly around the D₂ molecules, decreasing the unit cell volume by as much as 7.8% and

shrinking the binding pocket diameter from 5.0 to 3.6 Å (see ESI, Section 6†). The crucial role of this structural distortion is further exemplified by considering a hypothetical D₂ super atom at the fractional coordinates of site I in fully activated **1**. In this environment, the D₂ \cdots framework interactions are elongated beyond meaningful van der Waals contact distances (see ESI, Section 6†).

Rietveld refinement of *in situ* powder neutron diffraction data collected for **1** dosed with 1.5 equiv. of CD₄ confirmed that the molecule occupies the same adsorption pocket as D₂ (site I). Due to the larger size of CD₄ relative to D₂, a less pronounced contraction of the framework is sufficient to enable similar host–guest contacts. The unit cell volume of **1-CD₄** contracts by 3.4% to 1839.3(2) Å³, resulting in a binding pocket diameter of 4.1 Å. Notably, the adsorbed CD₄ molecules are well-ordered as a result of a large number of specific host–guest interactions (Fig. 6b). The nearest D \cdots arene distances are 2.740(9) Å (side wall of the pore) and 3.407(8) Å (outer linker of bowl-shaped cap). These relatively short distances are consistent with those determined previously from studies of methane adsorption on benzene (2.1–3.8 Å) and support the characterized orientation of CD₄ inside the pore of U(bdc)₂.^{45,81–83} Additional D \cdots H van der Waals contacts at 2.57(1), 2.69(2), and 2.81(2) Å stabilize and orient the adsorbed CD₄ molecules within the binding pocket. Adsorbed CD₄ further interacts with three O atoms of two independent UO₈ nodes at distances of 3.10(2), 3.25(1), and 3.50(1) Å. Lastly, adsorbate–adsorbate interactions based on D \cdots D contacts at 2.84(1), 3.10(1), and 3.20(1) Å stabilize CD₄ inside each pore.^{84,85} Together, the wealth of stabilizing contacts explains the competitively high heat of adsorption ($Q_{st} = -24.8$ kJ mol⁻¹) for methane in U(bdc)₂. It is worth noting that methane is also expected to occupy a second adsorption site at higher loadings, as seen for D₂. This observation is supported by the CH₄ adsorption data shown in Fig. 3. Here, initial steep uptake is associated with saturation of the binding pockets of site I until a loading of 3.5 mmol g⁻¹. The onset of far more gradual CH₄ uptake to 4.1 mmol g⁻¹ at 1.2 bar suggests additional methane adsorption at a second, weaker binding site.

Finally, we sought to study the distortion of **1** in the presence of an even larger guest molecule and prepared crystals of **1-DMF** by soaking the framework in dry DMF (kinetic diameter of 5.5 Å).⁸⁶ The structure of **1-DMF** was determined from Rietveld



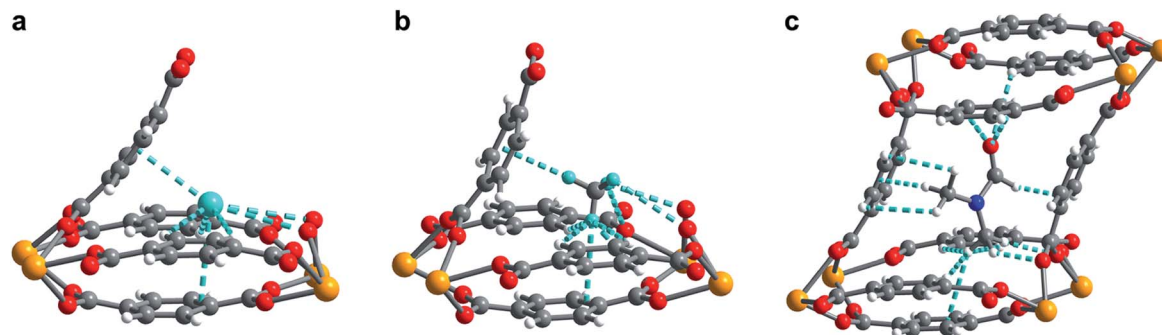


Fig. 6 (a) Neutron powder diffraction structure of D_2 adsorbed at site I in $1-D_2$ after dosing **1** with 2.5 equiv. D_2 per pore. (b) Neutron powder diffraction structure of CD_4 adsorbed at site I in $1-CD_4$ after dosing **1** with 1.5 equiv. CD_4 per pore. (c) Powder X-ray diffraction structure of DMF adsorbed inside the pore of $U(bdc)_2$. For clarity, only selected interactions of DMF with the framework are depicted. All framework–guest interactions are depicted as aquamarine colored dashed lines. For clarity in (a) and (b), the pore is truncated and cut in half diagonally, showing only one of the two adsorption pockets of the pore. Orange, red, grey, blue, aquamarine, and white spheres represent U, O, C, N, D, and H atoms, respectively.

refinement of powder X-ray diffraction data obtained at 298 K (see ESI, Fig. S8†). The framework indeed contracts to optimize interactions with DMF, but the unit cell volume decreases by only 1.9% (compared to 7.8% and 3.4% in the cases of 2.5 equiv. D_2 and 1.5 equiv. CD_4 , respectively) and the pocket diameter only decreases to 4.2 Å. While two molecules of the smaller guests D_2 and CD_4 can simultaneously occupy the two binding pockets in each pore, the larger DMF molecule occupies the whole pore space, bridging both pockets (Fig. 6c). As a result, DMF is stabilized by seven of the eight bdc^{2-} linkers that form the surrounding pore. While precise contact distances are obscured by disorder of DMF over two positions in the structure, the general identity of host–guest interactions in $1-DMF$ is clear. In particular, DMF binds through eight $H\cdots H$ contacts, three $C-H\cdots\pi$ -interactions, three $O\cdots H$ contacts involving the carbonyl and arene $C-H$ moieties, and interactions of two DMF $C-H$ groups with O atoms of two UO_8 nodes (all distances between 2 and 4 Å).

Conclusions

Porous adsorbents with small binding pockets can engage in strong, selective host–guest interactions, and are therefore of interest for applications including gas capture and storage. However, it remains a significant challenge to tune a binding pocket structure to optimize interactions with a specific target molecule. In this work, we show that combining moderate flexibility with small pores in the new flexible metal–organic framework $U(bdc)_2$ (**1**) eliminates the need for precise tuning of pore geometry. Indeed, this material is capable of uniquely adjusting its pore and binding pocket geometry to optimize host–guest interactions in the presence of even very weakly adsorbing molecules, such as H_2 and CH_4 . Temperature-dependent H_2 and CH_4 adsorption isotherms yielded isosteric heats of adsorption of -8.6 and -24.8 kJ mol^{-1} for H_2 and CH_4 , respectively, confirming comparatively strong interactions of **1** with both gases, despite their different sizes. *In situ* powder neutron diffraction experiments with D_2 and CD_4 revealed that

cooperative effects drive a spontaneous adjustment of the binding pockets in **1** to generate multiple stabilizing interactions between each adsorbate and the framework, which are not achieved without pore contraction. Altogether, our results demonstrate the utility of frameworks featuring self-adjusting binding pockets that can flex in response to different guest molecules. The design principles applied for our model material **1** are transferable to non-radioactive frameworks and suggest further exploration of such materials will be advantageous in the search for adsorbents with improved gas storage properties.

Conflicts of interest

There are no conflicts to declare.

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References

- H. Furukawa, K. E. Cordova, M. O'Keeffe and O. M. Yaghi, *Science*, 2013, **341**, 1230444.
- N. Stock and S. Biswas, *Chem. Rev.*, 2012, **112**, 933.
- M. Eddaoudi, J. Kim, N. Rosi, D. Vodak, J. Wachter, M. O'Keeffe and O. M. Yaghi, *Science*, 2002, **295**, 469.
- S. Yuan, L. Feng, K. Wang, J. Pang, M. Bosch, C. Lollar, Y. Sun, J. Qin, X. Yang, P. Zhang, Q. Wang, L. Zou, Y. Zhang, L. Zhang, Y. Fang, J. Li and H.-C. Zhou, *Adv. Mater.*, 2018, **30**, 1704303.
- T. Islamoglu, S. Goswami, Z. Li, A. J. Howarth, O. K. Farha and J. T. Hupp, *Acc. Chem. Res.*, 2017, **50**, 805.
- W. Lu, Z. Wei, Z.-Y. Gu, T.-F. Liu, J. Park, J. Park, J. Tian, M. Zhang, Q. Zhang, T. Gentle, M. Bosch and H.-C. Zhou, *Chem. Soc. Rev.*, 2014, **43**, 5561.
- A. U. Czaja, N. Trukhan and U. Müller, *Chem. Soc. Rev.*, 2009, **38**, 1284.
- C. H. Hendon, A. J. Rieth, M. D. Korzyński and M. Dincă, *ACS Cent. Sci.*, 2017, **3**, 554.
- A. Bétard and R. A. Fischer, *Chem. Rev.*, 2012, **112**, 1055.
- Q. Wang and D. Astruc, *Chem. Rev.*, 2020, **120**, 1438.
- D. J. Xiao, E. D. Bloch, J. A. Mason, W. L. Queen, M. R. Hudson, N. Planas, J. Borycz, A. L. Dzubak, P. Verma, K. Lee, F. Bonino, V. Crocell, J. Yano, S. Bordiga, D. G. Truhlar, L. Gagliardi, C. M. Brown and J. R. Long, *Nat. Chem.*, 2014, **6**, 590.
- S. A. Burgess, A. Kassie, S. A. Baranowski, K. J. Fritzscheing, K. Schmidt-Rohr, C. M. Brown and C. R. Wade, *J. Am. Chem. Soc.*, 2016, **138**, 1780.
- S. M. Cohen, Z. Zhang and J. A. Boissonnault, *Inorg. Chem.*, 2016, **55**, 7281.
- Y.-B. Huang, J. Liang, X.-S. Wang and R. Cao, *Chem. Soc. Rev.*, 2017, **46**, 126.
- J. Lee, O. K. Farha, J. Roberts, K. A. Scheidt, S. T. Nguyen and J. T. Hupp, *Chem. Soc. Rev.*, 2009, **38**, 1450.
- L. E. Kreno, K. Leong, O. K. Farha, M. Allendorf, R. P. van Duyne and J. T. Hupp, *Chem. Rev.*, 2012, **112**, 1105.
- M. G. Campbell, D. Sheberla, S. F. Liu, T. M. Swager and M. Dincă, *Angew. Chem., Int. Ed.*, 2015, **54**, 4349.
- I. Stassen, N. Burch, A. Talin, P. Falcaro, M. Allendorf and R. Ameloot, *Chem. Soc. Rev.*, 2017, **46**, 3185.
- K. Sumida, D. L. Rogow, J. A. Mason, T. M. McDonald, E. D. Bloch, Z. R. Herm, T.-H. Bae and J. R. Long, *Chem. Rev.*, 2012, **112**, 724.
- T. M. McDonald, J. A. Mason, X. Kong, E. D. Bloch, D. Gygi, A. Dani, V. Crocellà, F. Giordanino, S. O. Odoh, W. S. Drisdell, B. Vlasisavljevich, A. L. Dzubak, R. Poloni, S. K. Schnell, N. Planas, K. Lee, T. Pascal, L. F. Wan, D. Prendergast, J. B. Neaton, B. Smit, J. B. Kortright, L. Gagliardi, S. Bordiga, J. A. Reimer and J. R. Long, *Nature*, 2015, **519**, 303.
- R. L. Siegelman, P. J. Milner, A. C. Forse, J.-H. Lee, K. A. Colwell, J. B. Neaton, J. A. Reimer, S. C. Weston and J. R. Long, *J. Am. Chem. Soc.*, 2019, **141**, 13171.
- M. Ding, R. W. Flaig, H.-L. Jiang and O. M. Yaghi, *Chem. Soc. Rev.*, 2019, **48**, 2783.
- D. J. Babu, G. He, J. Hao, M. T. Vahdat, P. A. Schouwink, M. Mensi and K. V. Agrawal, *Adv. Mater.*, 2019, **31**, 1900855.
- B. R. Barnett, M. I. Gonzalez and J. R. Long, *Trends Chem.*, 2019, **1**, 159.
- H. Li, K. Wang, Y. Sun, C. T. Lollar, J. Li and H.-C. Zhou, *Mater. Today*, 2018, **21**, 108.
- D. E. Jaramillo, D. A. Reed, H. Z. H. Jiang, J. Oktawiec, M. W. Mara, A. C. Forse, D. J. Lussier, R. A. Murphy, M. Cunningham, V. Colombo, D. K. Shuh, J. A. Reimer and J. R. Long, *Nat. Mater.*, 2020, **19**, 517.
- B. Li, H.-M. Wen, W. Zhou and B. Chen, *J. Phys. Chem. Lett.*, 2014, **5**, 3468.
- R. E. Morris and P. S. Wheatley, *Angew. Chem., Int. Ed.*, 2008, **47**, 4966.
- D. A. Reed, D. J. Xiao, H. Z. H. Jiang, K. Chakarawet, J. Oktawiec and J. R. Long, *Chem. Sci.*, 2020, **11**, 1698.
- S. Ma and H.-C. Zhou, *Chem. Commun.*, 2010, **46**, 44.
- I. Senkowska, E. Barea, J. A. R. Navarro and S. Kaskel, *Microporous Mesoporous Mater.*, 2012, **156**, 115.
- P. García-Holley, B. Schweitzer, T. Islamoglu, Y. Liu, L. Lin, S. Rodriguez, M. H. Weston, J. T. Hupp, D. A. Gómez-Gualdrón, T. Yildirim and O. K. Farha, *ACS Energy Lett.*, 2018, **3**, 748.
- M. T. Kapelewski, S. J. Geier, M. R. Hudson, D. Stück, J. A. Mason, J. N. Nelson, D. J. Xiao, Z. Hulvey, E. Gilmour, S. A. FitzGerald, M. Head-Gordon, C. M. Brown and J. R. Long, *J. Am. Chem. Soc.*, 2014, **136**, 12119.
- M. T. Kapelewski, T. Runčevski, J. D. Tarver, H. Z. H. Jiang, K. E. Hurst, P. A. Parilla, A. Ayala, T. Gennett, S. A. FitzGerald, C. M. Brown and J. R. Long, *Chem. Mater.*, 2018, **30**, 8179.
- A. Ahmed, S. Seth, J. Purewal, A. G. Wong-Foy, M. Veenstra, A. J. Matzger and D. J. Siegel, *Nat. Commun.*, 2019, **10**, 1568.
- Y. He, W. Zhou, G. Qian and B. Chen, *Chem. Soc. Rev.*, 2014, **43**, 5657.
- J. A. Mason, J. Oktawiec, M. K. Taylor, M. R. Hudson, J. Rodriguez, J. E. Bachman, M. I. Gonzalez, A. Cervellino, A. Guagliardi, C. M. Brown, P. L. Llewellyn, N. Masciocchi and J. R. Long, *Nature*, 2015, **527**, 357.
- K. V. Kumar, K. Preuss, M.-M. Titirici and F. Rodríguez-Reinoso, *Chem. Rev.*, 2017, **117**, 1796.
- M. D. Allendorf, Z. Hulvey, T. Gennett, A. Ahmed, T. Autrey, J. Camp, E. Seon Cho, H. Furukawa, M. Haranczyk, M. Head-Gordon, S. Jeong, A. Karkamkar, D.-J. Liu, J. R. Long, K. R. Meihaus, I. H. Nayyar, R. Nazarov, D. J. Siegel, V. Stavila, J. J. Urban, S. P. Veccham and B. C. Wood, *Energy Environ. Sci.*, 2018, **11**, 2784.
- M. P. Suh, H. J. Park, T. K. Prasad and D.-W. Lim, *Chem. Rev.*, 2012, **112**, 782.



- 41 J. A. Mason, M. Veenstra and J. R. Long, *Chem. Sci.*, 2014, **5**, 32.
- 42 M. I. Gonzalez, J. A. Mason, E. D. Bloch, S. J. Teat, K. J. Gagnon, G. Y. Morrison, W. L. Queen and J. R. Long, *Chem. Sci.*, 2017, **8**, 4387.
- 43 D. Denysenko, M. Grzywa, J. Jelic, K. Reuter and D. Volkmer, *Angew. Chem., Int. Ed.*, 2014, **53**, 5832.
- 44 P. G. Boyd, A. Chidambaram, E. García-Díez, C. P. Ireland, T. D. Daff, R. Bounds, A. Gładysiak, P. Schouwink, S. M. Moosavi, M. M. Maroto-Valer, J. A. Reimer, J. A. R. Navarro, T. K. Woo, S. Garcia, K. C. Stylianou and B. Smit, *Nature*, 2019, **576**, 253.
- 45 Z. Hulvey, B. Vlasisavljevich, J. A. Mason, E. Tsivion, T. P. Dougherty, E. D. Bloch, M. Head-Gordon, B. Smit, J. R. Long and C. M. Brown, *J. Am. Chem. Soc.*, 2015, **137**, 10816.
- 46 D. G. Truhlar, *J. Chem. Educ.*, 2019, **96**, 1671.
- 47 N. Mehio, S. Dai and D.-e. Jiang, *J. Phys. Chem. A*, 2014, **118**, 1150.
- 48 Y. Liu, C. M. Brown, D. A. Neumann, V. K. Peterson and C. J. Kepert, *J. Alloys Compd.*, 2007, **446–447**, 385.
- 49 A. P. Katsoulidis, D. Antypov, G. F. S. Whitehead, E. J. Carrington, D. J. Adams, N. G. Berry, G. R. Darling, M. S. Dyer and M. J. Rosseinsky, *Nature*, 2019, **565**, 213.
- 50 Y. Gu, J.-J. Zheng, K.-I. Otake, K. Sugimoto, N. Hosono, S. Sakaki, F. Li and S. Kitagawa, *Angew. Chem., Int. Ed.*, 2020, DOI: 10.1002/anie.202003186.
- 51 F.-X. Coudert, M. Jeffroy, A. H. Fuchs, A. Boutin and C. Mellot-Draznieks, *J. Am. Chem. Soc.*, 2008, **130**, 14294.
- 52 B. Chen, S. Xiang and G. Qian, *Acc. Chem. Res.*, 2010, **43**, 1115.
- 53 A. Schneemann, V. Bon, I. Schwedler, I. Senkovska, S. Kaskel and R. A. Fischer, *Chem. Soc. Rev.*, 2014, **43**, 6062.
- 54 L. Sarkisov, R. L. Martin, M. Haranczyk and B. Smit, *J. Am. Chem. Soc.*, 2014, **136**, 2228.
- 55 J.-P. Zhang, H.-L. Zhou, D.-D. Zhou, P.-Q. Liao and X.-M. Chen, *Natl. Sci. Rev.*, 2018, **5**, 907.
- 56 C. Serre, F. Millange, C. Thouvenot, M. Noguès, G. Marsolier, D. Louër and G. Férey, *J. Am. Chem. Soc.*, 2002, **124**, 13519.
- 57 M. Vougo-Zanda, J. Huang, E. Anokhina, X. Wang and A. J. Jacobson, *Inorg. Chem.*, 2008, **47**, 11535.
- 58 G. Férey, M. Latroche, C. Serre, F. Millange, T. Loiseau and A. Percheron-Guégan, *Chem. Commun.*, 2003, 2976.
- 59 F. Millange, N. Guillou, M. E. Medina, G. Férey, A. Carlin-Sinclair, K. M. Golden and R. I. Walton, *Chem. Mater.*, 2010, **22**, 4237.
- 60 J. P. S. Mowat, S. R. Miller, A. M. Z. Slawin, V. R. Seymour, S. E. Ashbrook and P. A. Wright, *Microporous Mesoporous Mater.*, 2011, **142**, 322.
- 61 S. Surblé, C. Serre, C. Mellot-Draznieks, F. Millange and G. Férey, *Chem. Commun.*, 2006, 284.
- 62 N. Guillou, S. Bourrelly, P. L. Llewellyn, R. I. Walton and F. Millange, *CrystEngComm*, 2015, **17**, 422.
- 63 S. R. Daly, P. M. B. Piccoli, A. J. Schultz, T. K. Todorova, L. Gagliardi and G. S. Girolami, *Angew. Chem., Int. Ed.*, 2010, **49**, 3379.
- 64 E. A. Dolgoplova, A. M. Rice and N. B. Shustova, *Chem. Commun.*, 2018, **54**, 6472.
- 65 C. Falaise, A. Assen, I. Mihalcea, C. Volkringer, A. Mesbah, N. Dacheux and T. Loiseau, *Dalton Trans.*, 2015, **44**, 2639.
- 66 V. Bon, I. Senkovska, D. Wallacher, D. M. Töbrens, I. Zizak, R. Feyerherm, U. Mueller and S. Kaskel, *Inorg. Chem.*, 2014, **53**, 1513.
- 67 C. Mellot-Draznieks, C. Serre, S. Surblé, N. Audebrand and G. Férey, *J. Am. Chem. Soc.*, 2005, **127**, 16273.
- 68 M. K. Taylor, T. Runčevski, J. Oktawiec, M. I. Gonzalez, R. L. Siegelman, J. A. Mason, J. Ye, C. M. Brown and J. R. Long, *J. Am. Chem. Soc.*, 2016, **138**, 15019.
- 69 P. Bénard and R. Chahine, *Langmuir*, 2001, **17**, 1950.
- 70 L. J. Murray, M. Dincă and J. R. Long, *Chem. Soc. Rev.*, 2009, **38**, 1294.
- 71 F. M. Mulder, B. Assfour, J. Huot, T. J. Dingemans, M. Wagemaker and A. J. Ramirez-Cuesta, *J. Phys. Chem. C*, 2010, **114**, 10648.
- 72 F. Salles, D. I. Kolokolov, H. Jovic, G. Maurin, P. L. Llewellyn, T. Devic, C. Serre and G. Férey, *J. Phys. Chem. C*, 2009, **113**, 7802.
- 73 R. A. Pollock, J.-H. Her, C. M. Brown, Y. Liu and A. Dailly, *J. Phys. Chem. C*, 2014, **118**, 18197.
- 74 H. Wu, W. Zhou and T. Yildirim, *J. Am. Chem. Soc.*, 2007, **129**, 5314.
- 75 D. Gygi, E. D. Bloch, J. A. Mason, M. R. Hudson, M. I. Gonzalez, R. L. Siegelman, T. A. Darwish, W. L. Queen, C. M. Brown and J. R. Long, *Chem. Mater.*, 2016, **28**, 1128.
- 76 K. Sumida, J.-H. Her, M. Dincă, L. J. Murray, J. M. Schloss, C. J. Pierce, B. A. Thompson, S. A. FitzGerald, C. M. Brown and J. R. Long, *J. Phys. Chem. C*, 2011, **115**, 8414.
- 77 T. Runčevski, M. T. Kapelewski, R. M. Torres-Gavosto, J. D. Tarver, C. M. Brown and J. R. Long, *Chem. Commun.*, 2016, **52**, 8251.
- 78 J. Luo, H. Xu, Y. Liu, Y. Zhao, L. L. Daemen, C. Brown, T. V. Timofeeva, S. Ma and H.-C. Zhou, *J. Am. Chem. Soc.*, 2008, **130**, 9626.
- 79 J. van Kranendonk and H. P. Gush, *Phys. Lett.*, 1962, **1**, 22.
- 80 V. K. Peterson, Y. Liu, C. M. Brown and C. J. Kepert, *J. Am. Chem. Soc.*, 2006, **128**, 15578.
- 81 S. Tsuzuki, K. Honda, T. Uchamaru, M. Mikami and K. Tanabe, *J. Am. Chem. Soc.*, 2000, **122**, 3746.
- 82 A. L. Ringer, M. S. Figgs, M. O. Sinnokrot and C. D. Sherrill, *J. Phys. Chem. A*, 2006, **110**, 10822.
- 83 H. Wu, W. Zhou and T. Yildirim, *J. Phys. Chem. C*, 2009, **113**, 3029.
- 84 E. Tsivion and M. Head-Gordon, *J. Phys. Chem. C*, 2017, **121**, 12091.
- 85 S. Tsuzuki, T. Uchamaru and K. Tanabe, *Chem. Phys. Lett.*, 1998, **287**, 202.
- 86 J. E. ten Elshof, C. R. Abadal, J. Sekulić, S. R. Chowdhury and D. H. A. Blank, *Microporous Mesoporous Mater.*, 2003, **65**, 197.

