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 (ΔV = −7.8%), triggered by D₂ adsorption at low pressures. This deformation creates two optimized binding pockets for hydrogen (Q₂ = −8.6 kJ mol⁻¹) per pore, in agreement with H₂ adsorption data. Analogous experiments with CD₄ (Q₂ = −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.

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 Ni₃(m-dobdc) (m-dobdc⁻¹ = 4,6-dioxido-1,3-benzedicarboxylate), which is currently the top-performing material for ambient temperature, physisorptive H₂ storage.44,45 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.46 An example of how such cumulative dispersion forces can outperform strong interactions at open metal sites is the adsorption of CH₄ in Cu₂(btc)₁ (HKUST-1, btc⁻¹ = 1,3,5-benzenetricarboxylate).43 This material exhibits open metal sites and binding pockets in direct competition for CH₄ adsorption. Structural characterization of Cu₂(btc)₁ 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(n) open metal sites. The reason for this behavior is that the multiple interactions inside the pore give rise to a higher...
overall binding energy than that achieved with a CH$_4$ molecule adsorbed at a single copper(ii) center (−21.8 versus −9.4 kJ mol$^{-1}$, respectively).  

Cumulative dispersion interactions between guest molecules and framework pockets decrease exponentially with the adsorbate–framework distances ($F \propto 1/r^9$), and therefore require a precise geometric fit between guest and binding pocket.  

For example, as a result of its smaller kinetic diameter relative to CH$_4$, H$_2$ preferentially binds at the open metal sites of Cu$_2$(btc)$_3$, rather than in the hexagonal pockets.  

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.  

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.

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.  

Prominent examples are M(OH)(btc) (MIL-53; btc$^{2-}$ = 1,4-benzenedicarboxylate; M = Fe, Cr, Sc, Al, or Ga) and M$_2$(O)OH(H$_2$O)$_2$(btc)$_3$ (MIL-88; M = Fe, Cr).  

These frameworks undergo dramatic geometrical 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 CO$_2$ adsorption in Fe(OH)(btc).  

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 btc$^{2-}$ 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.  

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.  

Inspired by previous work on the synthesis of porous metal–organic frameworks from uranium(iv) and btc$^{2-}$ linkers, we synthesized a new, three-dimensional U(btc)$_2$ phase (I) 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 H$_2$ and CH$_4$, with different levels of contraction and host-guest interactions for each molecule.

## Results and discussion

The compound U(btc)$_2$·4H$_2$O (1–H$_2$O) was synthesized through the reaction of U$_4$(1,4-dioxane)$_2$ with H$_2$btc 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–H$_2$O (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–H$_2$O match the simulated pattern determined from single-crystal data, confirming the bulk purity of the crystalline material (see ESI, Fig. S7†).

Compound 1–H$_2$O crystallizes in the space group C2/c and features eight-coordinate uranium centers in a distorted square-antiprismatic environment. Each uranium(iv) is...
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(v) 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 U[bdc]$\cdot$4H$_2$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 U[bdc]$\cdot$ (1) was obtained by heating 1-H$_2$O 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 $497 \pm 6$ m$^2$ 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, the activation pore window, such as Cu$_2$(btc)$_3$ (Q$_{st}$ = 12.3 kJ mol$^{-1}$) and even values for frameworks with snugly fitting pore window, such as Cu$_2$(btc)$_3$ (Q$_{st}$ = 17.1 kJ mol$^{-1}$), or strongly polarizing open metal cation sites, as in Ni$_4$,(dobdc) (dobdc$^{4-}$ = 2,5-dioxido-1,4-benzenedicarboxylate; Q$_{st}$ = 20.6 kJ mol$^{-1}$). It is clear that the binding pockets in the unit cell volume from 1887.4(3) to 1902.1(2) Å$^3$, 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 U[bdc]$\cdot$ at 77 K exhibits an initial steep rise to ~3.5 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 ~4.9 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 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$^{-1}$)$^{96}$ 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$_{st}$ = 24.8 kJ mol$^{-1}$ at low loadings (see ESI, Fig. S5†). Notably, this value surpasses that determined for Zn$_4$O(bdc)$_3$ (MOF-5; Q$_{st}$ = ~12.3 kJ mol$^{-1}$) and even values for frameworks with snugly fitting pore window, such as Cu$_2$(btc)$_3$ (Q$_{st}$ = ~17.1 kJ mol$^{-1}$), or strongly polarizing open metal cation sites, as in Ni$_4$,(dobdc) (dobdc$^{4-}$ = 2,5-dioxido-1,4-benzenedicarboxylate; Q$_{st}$ = ~20.6 kJ mol$^{-1}$). It is clear that the binding pockets in...
U(bdc)$_2$ can strongly interact with both H$_2$ and CH$_4$, 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 D$_2$ per pore results in clear changes in the powder pattern, particularly 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–D$_2$. Upon increasing the loading to 0.7 equiv. of D$_2$, both phases are still present, although the peaks of 1 diminish further and the peaks of newly formed 1–D$_2$ gain in intensity (Fig. 4 and S11†). The two phases coexist up to a loading of at least 1.5 equiv. D$_2$ per pore (see ESI, Fig. S12†), and their interconversion is best followed by evaluating the high intensity, low angle peaks at 2θ = 11.85° for 1 and at 2θ = 12.63° for 1–D$_2$ in Fig. 4. These data strongly suggest that cooperative effects drive an adsorbate-induced framework distortion from 1 to 1–D$_2$. Such a mechanism is in contrast to a gradual and homogeneous uptake of D$_2$, or a gradually changing degree of distortion depending on the D$_2$-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 D$_2$ molecule, while others will remain in the activated structure of 1. At the highest D$_2$ dosing level of 2.5 equiv. per pore, the diffraction pattern of the sample contains only reflections associated with 1–D$_2$ (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 CD$_4$ (Fig. S14 and S15†). The data from these experiments suggest a similar cooperative transformation from 1 to an adsorbed phase 1–CD$_4$.

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 D$_2$ and CD$_4$. 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 D$_2$ than it does in the presence of CH$_4$.

Surprisingly, a structural comparison of 1 with 1–D$_2$ and 1–CD$_4$ reveals an almost constant coordination environment around the uranium nodes in each phase. The framework flexibility instead relies on a tilting of the parallelepipedic pores, which is clearly seen by comparing the pore geometries of 1 and 1–D$_2$ dosed with 2.5 equiv. D$_2$, 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 D$_2$ results in a change of the idealized parallelogram and stretch of the pentagonal bipyramids, which is clearly shown in Fig. 5a.

The resulting hinge-type bending between the UO$_6$ nodes and linkers is analogous to the change that occurs in the structure of the flexible framework Cr(OH)(bdc)$_2$ upon water adsorption. In particular, water adsorption is accompanied by a unit cell volume decrease from 1486.1 to 1012.6 Å$^3$, as well as a decrease of the symmetrical dihedral angles, $\omega_1$ from 179.8° to 162.3°. The analogous idealized Cr⋯Cr⋯Cr angles $\varphi_1$ and $\varphi_2$ in

![Fig. 4](image1.png)

**Fig. 4** Rietveld refinement fits of powder neutron diffraction patterns (λ = 2.0772 Å, T = 9 K) collected for activated 1 and activated 1 dosed with 0.3, 0.7, and 1.5 equiv. of D$_2$ per pore. Black (lower) and blue (upper) tick marks indicate calculated Bragg peak positions for 1 and the D$_2$ adsorbed phase, 1–D$_2$, respectively. Arrows indicate peaks that best illustrate the conversion of 1 to 1–D$_2$ with increasing D$_2$ loading.

![Fig. 5](image2.png)

**Fig. 5** (a) Illustration of pores in 1 and 1–D$_2$ (dosed with 2.5 equiv. D$_2$ per pore). The pore contraction upon D$_2$ 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 D$_2$ molecules inside the pores of 1–D$_2$ are represented as aquamarine spheres for adsorption site I and as brown spheres for adsorption site II. Each D$_2$ 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 D$_2$ molecule per pore. (b) Comparison of the dihedral angles $\omega_1$, as described in the text [i.e., $\omega_2$ 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 D$_2$. Orange, red, and grey spheres represent U, O, and C atoms, respectively.
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⋅⋅π 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)$_2$.54,81–83 Additional D⋅⋅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 UO$_8$ nodes at distances of 3.10(2), 3.25(1), and 3.50(1) Å. Lastly, adsorbate–adsorbate interactions based on D⋅⋅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$_d$ = −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 D$_2$. This observation is supported by the CH$_4$ adsorption data shown in Fig. 3. Here, initial steep uptake is associated with saturation of the binding pockets of site 1 until a loading of 3.5 mmol g$^{-1}$. The onset of far more gradual CH$_4$ 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

**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 $\phi_1$ and $\phi_2$ describe the U⋯U⋯U angles of the idealized parallelogram spanned by the four corner uranium atoms within the crystallographic a-plane of each pore, as illustrated in Fig. 5a. Angles $\omega_3$ and $\omega_2$ describe the dihedral angle between O⋯U⋯O⋯U planes and O⋯C⋯O planes at each end of a bdc$^{2-}$ linker, as illustrated in Fig. 5b for 1 and 1–D$_2$. One standard deviation for all values is given in parentheses.

<table>
<thead>
<tr>
<th>Sample</th>
<th>a (Å)</th>
<th>b (Å)</th>
<th>c (Å)</th>
<th>V (Å$^3$)</th>
<th>$\phi_1$ (°)</th>
<th>$\phi_2$ (°)</th>
<th>$\omega_3$ (°)</th>
<th>$\omega_2$ (°)</th>
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<tr>
<td>1</td>
<td>17.587(1)</td>
<td>12.812(1)</td>
<td>9.2999(5)</td>
<td>1902.1(2)</td>
<td>77.1(1)</td>
<td>112.90(2)</td>
<td>168.4(9)</td>
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<tr>
<td>1–D$_2$ (0.3 equiv.)</td>
<td>18.361(7)</td>
<td>11.351(4)</td>
<td>9.338(2)</td>
<td>1782.2(5)</td>
<td>67.5(1)</td>
<td>112.49(9)</td>
<td>148.7(3)</td>
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<tr>
<td>1–D$_2$ (0.7 equiv.)</td>
<td>18.402(3)</td>
<td>11.324(5)</td>
<td>9.333(1)</td>
<td>1781.4(8)</td>
<td>67.4(2)</td>
<td>112.57(6)</td>
<td>130(2)</td>
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<tr>
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<td>11.233(1)</td>
<td>9.350(8)</td>
<td>1775.8(2)</td>
<td>66.8(6)</td>
<td>113.14(3)</td>
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<td>10.948(6)</td>
<td>9.383(5)</td>
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<td>9.3579(3)</td>
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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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