

Cite this: *Mater. Adv.*, 2021,  
2, 426

# Adsorption properties of acetylene, ethylene and ethane in UiO-66 with linker defects and NO<sub>2</sub> functionalization†

H. Pandey,<sup>ab</sup> T. Barrett,<sup>bc</sup> M. D. Gross<sup>bd</sup> and T. Thonhauser<sup>id\*ab</sup>

Separation of small hydrocarbon gases is crucial for many industrial processes, but the high energy demand of current separation techniques encourages exploration of alternative sorption-based approaches. Here, we investigate the metal organic framework UiO-66(Zr) as possible solid adsorbent with a focus on the fundamental sorption mechanisms of C<sub>2</sub> hydrocarbons in the presence of linker defects and NO<sub>2</sub> functionalization. To this end, we directly measure heats of adsorption with calorimetry and correlate our results with *ab initio* calculations. In particular, we study adsorption of C<sub>2</sub>H<sub>2</sub>, C<sub>2</sub>H<sub>4</sub> and C<sub>2</sub>H<sub>6</sub> at 293 K and 195 K. At 293 K, the measured low-coverage enthalpy is found to be an average enthalpy of multiple adsorption sites. However, at 195 K, the low-coverage enthalpy corresponded to the strongest binding site, *i.e.* the linker defect site. We find that C<sub>2</sub>H<sub>2</sub> exhibits the highest adsorption enthalpy with a significant gap of ~6 kJ mol<sup>-1</sup> to C<sub>2</sub>H<sub>4</sub> and ~9 kJ mol<sup>-1</sup> to C<sub>2</sub>H<sub>6</sub>. Our *ab initio* calculations show excellent agreement with our experiments and are crucial to uncover the mechanisms that lead to the observed differences in binding affinity as a function of coverage. We also find that binding enthalpies increase for all three gases by adding -NO<sub>2</sub> functional groups, which even further increase the gap to ~9 kJ mol<sup>-1</sup> and ~10 kJ mol<sup>-1</sup> for C<sub>2</sub>H<sub>4</sub> and C<sub>2</sub>H<sub>6</sub>, respectively. Our findings provide mechanistic insight that suggests that linker defects in UiO-66(Zr) have promise to improve its effectiveness as a solid adsorbent in C<sub>2</sub>H<sub>2</sub>, C<sub>2</sub>H<sub>4</sub> and C<sub>2</sub>H<sub>6</sub> applications and that functionalization may be used to increase specificity.

Received 21st September 2020,  
Accepted 5th December 2020

DOI: 10.1039/d0ma00722f

rsc.li/materials-advances

## 1 Introduction

C<sub>2</sub>H<sub>2</sub>, C<sub>2</sub>H<sub>4</sub> and C<sub>2</sub>H<sub>6</sub> are amongst the most widely used hydrocarbon gases. They form vital components for various industrial processes such as the chemical conversion to other hydrocarbons,<sup>1,2</sup> formation of aromatic compounds,<sup>3</sup> synthesis of commercially useful products (*e.g.* ethylene glycol),<sup>4</sup> production of various polymers including polyethylene, polyvinylchloride and polyester<sup>5</sup> and as fuel for power generation.<sup>6–8</sup> During natural gas extraction a mixture of C<sub>2</sub>H<sub>2</sub>, C<sub>2</sub>H<sub>4</sub> and C<sub>2</sub>H<sub>6</sub> is obtained that has to be separated and purified before use. C<sub>2</sub>H<sub>2</sub>/C<sub>2</sub>H<sub>4</sub> and C<sub>2</sub>H<sub>2</sub>/C<sub>2</sub>H<sub>6</sub> separation is of particularly interest due to their extensive use in many feedstock industrial

processes that require them to be in highly pure form.‡ Hence, purification and separation has become a critical step for the natural gas and petrochemical industries. Unfortunately, the most common separation method for light hydrocarbons, *i.e.* cryogenic separation,<sup>9–11</sup> requires large amounts of energy; 45–55% of industrial energy consumption in the US is taken up by gas separation processes, making up 10–15% of the total US energy consumption.<sup>12</sup> Replacing cryogenic separation processes with less energy intense methods, such as solid adsorbents and membrane-based separation, have thus seen a surge of interest.<sup>13–18</sup>

Metal organic frameworks (MOFs) have emerged not only as alternative solid adsorbent for gas separation, purification and capture,<sup>19–27</sup> but they are also interesting for catalysis,<sup>28–31</sup> chemical sensing/detection,<sup>32,33</sup> drug delivery,<sup>34–36</sup> multiferroics<sup>37–39</sup> and electrochemical energy storage.<sup>40,41</sup> This vast versatility results from their nanoporous and modular building-block nature, which can be tailored chemically and structurally. In addition, MOFs are relatively easy to synthesize with high yield, versatility and reproducibility.<sup>42</sup> Over the last decade, their use for industrial separation of hydrocarbons has made significant progress<sup>18,43–46</sup> and separation of

<sup>a</sup> Department of Physics, Wake Forest University, Winston-Salem NC 27109, USA. E-mail: thonhauser@wfu.edu

<sup>b</sup> Center for Functional Materials, Wake Forest University, Winston-Salem NC 27109, USA

<sup>c</sup> Department of Chemistry, Wake Forest University, Winston-Salem, NC 27109, USA

<sup>d</sup> Department of Engineering, Wake Forest University, Winston-Salem, NC 27109, USA

† Electronic supplementary information (ESI) available. See DOI: 10.1039/d0ma00722f

‡ *E.g.* acetylene is explosive and can react with the impurities present resulting in massive explosions.





differential energy plus RT.<sup>65</sup> Differential heats of adsorption and isotherms were measured at 195 K and 293 K for C<sub>2</sub>H<sub>2</sub>, C<sub>2</sub>H<sub>4</sub>, and C<sub>2</sub>H<sub>6</sub> (Fig. S4, ESI†).

### 2.3 Computational details

Our *ab initio* calculations were performed at the density functional theory (DFT) level, using VASP (Vienna *ab Initio* Simulation Package)<sup>66,67</sup> in conjunction with the vdW-DF functional.<sup>68–71</sup> The SCF convergence condition was set to 10<sup>−3</sup> meV and the structure was optimized until all forces between atoms were smaller than 1 meV Å<sup>−1</sup>. The plane-wave energy cut-off was set at 600 eV and, due to the size of the unit cell, only the  $\Gamma$ -point was used. Without guest molecule, the unit cell of UiO-66(Zr) contains 108 atoms (Fig. 1) and that of UiO-66(Zr)-NO<sub>2</sub> contains 126. Since the Zr atoms contain unpaired electrons, initial simulations were performed spin-polarized. However, spin effects cause differences in binding energies of only <0.1% so that we used non-spin polarized calculations henceforth. Guest molecules were introduced in the MOF at different sites to find the favorable binding sites and allowed to fully relax. The unit cell shape and volume was initially allowed to change during relaxation, but our calculations showed that this has only a minor effect on the binding energies (<1.5%), so the unit cell dimensions were kept fixed for all subsequent relaxations. Binding energies were calculated as energy differences of suitable fragments. Note that enthalpy contributions to the binding energy for C<sub>2</sub>H<sub>2</sub>, C<sub>2</sub>H<sub>4</sub>, and C<sub>2</sub>H<sub>6</sub> tend to be small at finite temperature, *i.e.* on the order of 2–3 kJ mol<sup>−1</sup> (see *e.g.* ESI of 23).

## 3 Results and discussion

### 3.1 MOF structure

Dehydroxylated UiO-66(Zr) is a zirconium-based MOF,<sup>72</sup> made of [Zr<sub>6</sub>(μ<sub>3</sub>-O)<sub>6</sub>] clusters connected to each other by

benzenedicarboxylate (BDC) linkers [OOC-C<sub>6</sub>H<sub>4</sub>-OOC]<sup>−2</sup>. Each cluster has 12 BDC linkers binding it to 12 other clusters thus giving a charge neutral structure. A unit cell of UiO-66(Zr) used for our calculations consisted of 6 Zr atoms interconnected with 6 BDC linkers. An extended version of the unit cell having 12 complete linkers is shown in Fig. 1. UiO-66 belongs to a special family of porous materials with 2 types of pores: an octahedral pore (11 Å diameter) and a tetrahedral pore (8 Å diameter). The structure of UiO-66(Zr) is well-known to contain missing linker defects and the presence of water molecules bound to the metal centers at the missing linker sites has been identified.<sup>64,73</sup> Following our specific synthesis procedure in Section 2.1, our MOF exhibits linker defects and our thermogravimetric analysis (TGA) in Fig. S3 (ESI†) indicates that UiO-66(Zr) has 1.35 missing linkers and UiO-66(Zr)-NO<sub>2</sub> has 0.60 missing linkers per unit cell. For our theoretical calculations – to avoid large super cells – we thus use the UiO-66(Zr) model depicted in Fig. 2 with one missing linker per unit cell. Nonetheless, in our partition-function occupancy-probability calculations we do use the exact numbers of missing linkers and thus available primary binding sites from our TGA results.

The symmetry of our MOF structure can be exploited to reduce the number of possible binding sites that have to be considered. One end of a BDC linker connects to one Zr atom to form an arm that repeats itself over the whole UiO-66 structure. Along this BDC-Zr-arm we can identify five potential binding sites: Site A1: Zr metal site with a linker missing, Site A2: Zr metal site with no missing linker, Site B: oxygen atom attached to the Zr metal, Site C: carbon atom between the oxygen and the benzene ring and Site D: benzene ring in the BDC linker. Finally, we find another binding site called Site E, where – depending on the situation – a varying number of guest

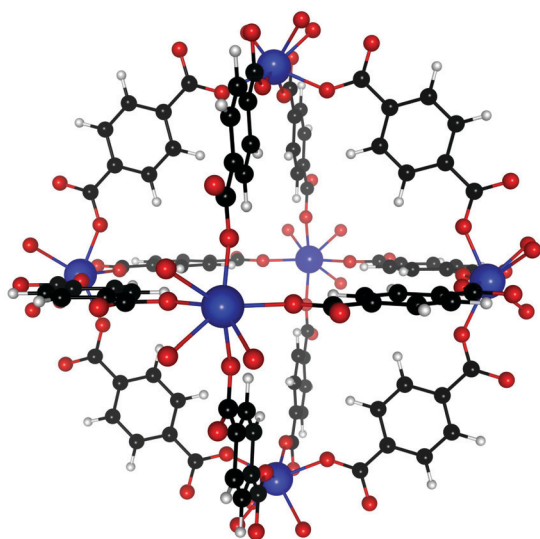


Fig. 1 Structure of UiO-66(Zr), showing the Zr atoms bound to each other by 12 BDC linkers. The pristine unit cell of UiO-66(Zr) consists of 6 Zr atoms and 6 BDC linkers. Color coding: Zr (blue), oxygen (red), carbon (black), and hydrogen (white).

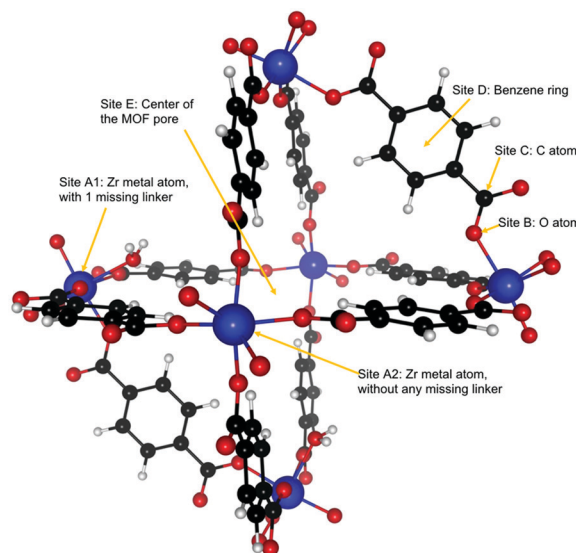


Fig. 2 UiO-66(Zr) structure used in this study with linker defects, *i.e.* two out of every twelve BDC linkers missing, which corresponds to one missing linker per unit cell. The arrows show the six possible binding sites for guest molecules. Note that, due to the symmetry of this structure, each binding site has a particular number of equivalent sites in each unit cell.



molecules can be adsorbed in the center of the MOF pore. These binding sites are depicted in Fig. 2. Although there are potentially six different types of active sites inside the MOF, our calculations show that sites B and C are not preferred by our molecules of interest, *i.e.* when placing  $C_2H_2$ ,  $C_2H_4$  or  $C_2H_6$  near those sites, they eventually drift closer to another nearby stable site like the Zr metal center or the benzene ring.

### 3.2 Adsorption of $C_2H_2$ , $C_2H_4$ and $C_2H_6$ in UiO-66(Zr)

Calorimetry results in the form of isosteric heats of adsorption for  $C_2H_2$ ,  $C_2H_4$  and  $C_2H_6$  in UiO-66(Zr) at 195 K are depicted in Fig. 3 as a function of guest molecule coverage; results at 293 K are given in Fig. S5 (ESI<sup>†</sup>). Heat of adsorption measurements were done for 10 sets of data and the uncertainty in measurement was found to be <2%. The results show that at 293 K the low-coverage heat of adsorption for  $C_2H_4$  and  $C_2H_6$  are nearly identical and 3 kJ mol<sup>-1</sup> lower than  $C_2H_2$ . However, when going to a lower temperature of 195 K, the heats of adsorption are significantly more separated and also shift to noticeably higher values, which is beneficial for separation purposes – we thus focus on the 195 K results.

Combining our calorimetry data and DFT calculations we can qualitatively and quantitatively explain the adsorption behavior in Fig. 3. Our main theoretical results are collected in Table 1, showing binding energies for all three gases corresponding to their primary, secondary, and tertiary binding site. For comparison, we also list the experimental heats of adsorption at 195 K and 293 K for the lowest measured coverage – as can be seen, we find excellent agreement between experiment and our calculated primary binding energies. It is also interesting to see that as primary sites all three molecules prefer Site A1, *i.e.* the Zr metal site with a

missing linker. However, the secondary and tertiary binding sites differ from molecule to molecule.

Looking at the curve for the  $C_2H_2$  heat of adsorption in Fig. 3, we measure at 195 K for the low coverage of 0.1 mmol g<sup>-1</sup> a value of 39.75 kJ mol<sup>-1</sup>, in excellent agreement with our calculated value of 39.52 kJ mol<sup>-1</sup> for the primary binding site. As we increase the coverage to *e.g.* 1 molecule per unit cell (0.66 mmol g<sup>-1</sup>) on average we see that the heat drops to a value below the primary binding site, 37.1 kJ mol<sup>-1</sup>. The heat continues to drop with increasing coverage until we reach approximately 3 guest molecules per unit cell (2 mmol g<sup>-1</sup>) and thereafter stays constant at approximately 33 kJ mol<sup>-1</sup>. The initial drop and then the plateau in the heat of adsorption of  $C_2H_2$  can be perfectly explained by partition-function occupancy-probability calculations based on our theoretical binding energies and the number of available binding sites in Table 1. Note that one missing linker leaves two open metal Sites A1, but  $C_2H_2$  binds in such a way – taking up the space between both Sites A1 – that there is practically only one binding site available (which is not the case for  $C_2H_4$  and  $C_2H_6$ ). For the secondary binding site of  $C_2H_2$  (Site D), the molecule sits in a triangular space surrounded by three linkers, which – according to our binding calculations – can hold up to two guest molecules with 34.51 kJ mol<sup>-1</sup> each. For our cell in Fig. 2 there are four such triangular spaces available, providing a maximum of 8 secondary sites. Our calculated heats of adsorption (blue dashed line in Fig. 3) are in excellent agreement with experiment and fully captures and explains the overall trend. Note that our calculated values are well within chemical accuracy (1 kcal mol<sup>-1</sup> = 4.2 kJ mol<sup>-1</sup>) over the entire range of coverages. The drop and the plateau is thus a result of the small number of available primary sites and the large number of secondary sites. We expect a further drop in the heat of adsorption down to ~29 kJ mol<sup>-1</sup> for coverages above 9 molecules per unit cell.

The case of  $C_2H_4$  is similar to the case of  $C_2H_2$ . But, there are two main differences: (i) the difference in binding energies between primary, secondary, and tertiary sites is much less compared to  $C_2H_2$ , and (ii) there are only four secondary sites available. Note that Site E in this case can hold up to four guest molecules. As a result, primary, secondary, and even tertiary sites get filled up more linearly with coverage and we thus observe a consistent decrease in heat as a function of coverage for  $C_2H_4$  in Fig. 3. Our calculated binding energy at the primary site of 32.49 kJ mol<sup>-1</sup> is in very good agreement to the measured low-coverage value of 33.39 kJ mol<sup>-1</sup> and our partition-function occupancy-probability calculations capture the linear decrease with coverage very well, albeit with a slightly too shallow slope (green line in Fig. 3).

Finally, for  $C_2H_6$  molecules we find a similar behavior, but the number of available binding sites are slightly different from the other cases. Here, due to the larger size of  $C_2H_6$ , Site E can only hold three guest molecules. Overall, our calculated primary site binding energy of 32.10 kJ mol<sup>-1</sup> is once more in excellent agreement with the low-coverage measured value of 31.12 kJ mol<sup>-1</sup> and our calculated heats capture the overall trend extremely well over the entire range of coverage (red line in Fig. 3). We note a small increase in the experimental heat of adsorption for  $C_2H_6$  at low concentrations – this slight increase

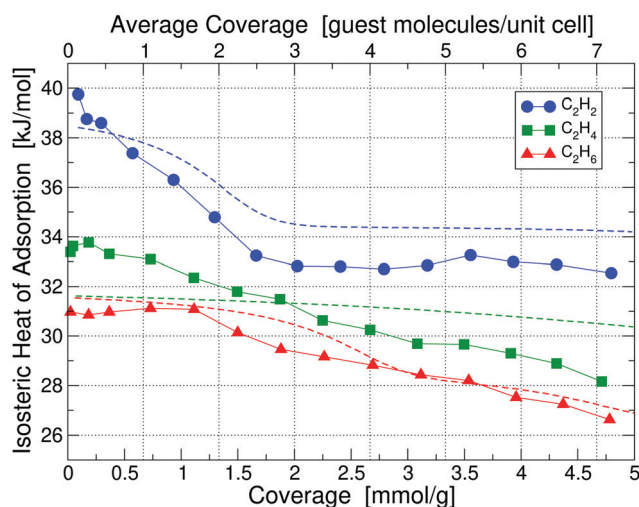


Fig. 3 Isosteric heats of adsorption of  $C_2H_2$ ,  $C_2H_4$  and  $C_2H_6$  in UiO-66(Zr) at 195 K, obtained by calorimetry measurements; results for 293 K are shown in Fig. S5 (ESI<sup>†</sup>). The x-axis is given in the chemically more useful units of [mmol g<sup>-1</sup>] at the bottom, whereas we have converted it to units of [guest molecule per unit cell] on the top, which is more useful for theoretical calculations. Dashed lines correspond to our partition-function occupancy-probability calculations based on our theoretical binding energies and availability of binding sites.





**Table 1** Calculated binding energies of C<sub>2</sub>H<sub>2</sub>, C<sub>2</sub>H<sub>4</sub> and C<sub>2</sub>H<sub>6</sub> in UiO-66(Zr) [kJ mol<sup>-1</sup>]. Experimental results for the lowest measured coverage at 195 K and 293 K are given for comparison. Sites are labeled according to Fig. 2. The number of equivalent sites per cell is also given. A1\* indicates that the guest molecule is bound close to site A1, but in a different orientation

Molecule	Exp. low coverage		Primary			Secondary			Tertiary		
	195 K	293 K	Energy	Site	Equiv.	Energy	Site	Equiv.	Energy	Site	Equiv.
C <sub>2</sub> H <sub>2</sub>	39.75	32.08	39.52	A1	1	34.51	D	8	29.02	A2	4
C <sub>2</sub> H <sub>4</sub>	33.39	29.11	32.49	A1	2	30.94	A2	4	29.40	E	4
C <sub>2</sub> H <sub>6</sub>	31.12	28.99	32.10	A1	2	28.95	A1*	2	26.70	E	3

is on the order of 0.2–0.3 kJ mol<sup>-1</sup> and falls within the uncertainty of our measurements.

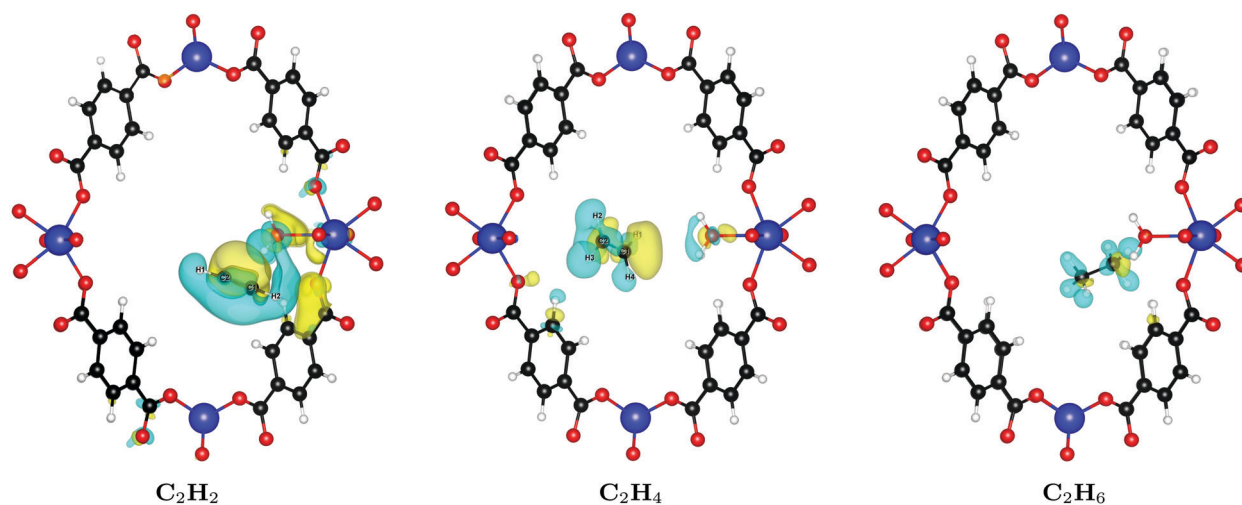
In order to understand the large difference of 28% in binding affinity between C<sub>2</sub>H<sub>2</sub> and C<sub>2</sub>H<sub>6</sub>, we calculate the induced charge density upon binding of both molecules, *i.e.* the charge density rearrangement upon forming the bond; results are depicted in Fig. 4. As can be seen, the interaction of C<sub>2</sub>H<sub>2</sub> with the framework is noticeably stronger compared to C<sub>2</sub>H<sub>4</sub> or C<sub>2</sub>H<sub>6</sub>. The reason for this behavior is that – because of its linear shape and thus smaller cross-section – it can sit in the groove formed between two water molecules at the linker defect site and interact with both, resulting in the increased binding energy. The smallest distance between C<sub>2</sub>H<sub>2</sub> and the water molecules is only 2.59 Å. Because of its shape, C<sub>2</sub>H<sub>4</sub> cannot fit in the same way and gets pushed further away into the pore, with a smallest distance of 3.34 Å. The same is true for the even larger C<sub>2</sub>H<sub>6</sub>, which shows even less charge rearrangement and interaction (smallest distance of 4.09 Å).

### 3.3 Adsorption of C<sub>2</sub>H<sub>2</sub>, C<sub>2</sub>H<sub>4</sub>, and C<sub>2</sub>H<sub>6</sub> in NO<sub>2</sub> functionalized UiO-66(Zr)

We also studied the effect of functionalizing UiO-66(Zr) with NO<sub>2</sub> by adding one NO<sub>2</sub> functional group on each BDC linker. The resulting new structure after relaxation is shown in Fig. 5 and the experimental calorimetry data is depicted in Fig. 6. We observe an overall rise in the adsorption enthalpy for all three gases as

compared to the unfunctionalized case in Fig. 3. For the energy at low coverage the values are increased by a magnitude of 3.5 kJ mol<sup>-1</sup> for C<sub>2</sub>H<sub>2</sub> and only 1 kJ mol<sup>-1</sup> for C<sub>2</sub>H<sub>4</sub> and 2 kJ mol<sup>-1</sup> for C<sub>2</sub>H<sub>6</sub>. This further increases the separation between C<sub>2</sub>H<sub>2</sub> and C<sub>2</sub>H<sub>4</sub> from 6.4 kJ mol<sup>-1</sup> to 8.9 kJ mol<sup>-1</sup> and from 8.6 kJ mol<sup>-1</sup> to 10 kJ mol<sup>-1</sup> for C<sub>2</sub>H<sub>6</sub>, with important implications for the separation of these molecules.

Calculated binding energies are shown in Table 2. The measured heats of adsorption for C<sub>2</sub>H<sub>2</sub> of 43.27 kJ mol<sup>-1</sup> at 0.04 mmol g<sup>-1</sup> coverage is in good agreement with our calculated primary binding energy of 45.12 kJ mol<sup>-1</sup>; similarly for the cases of C<sub>2</sub>H<sub>4</sub> and C<sub>2</sub>H<sub>6</sub>. Due to the presence of NO<sub>2</sub> on each linker, the available pore size for binding is reduced, which generally leads to a lower number of available binding sites for guest molecules in Table 2. As in the unfunctionalized case, primary binding always occurs at Site A1. The behavior of the isosteric heats of adsorption in Fig. 6 follows the same overall trend as compared to the unfunctionalized case and can be explained on the same basis of binding energies and site occupancy probabilities (see the discussion in Section 3.2 why sometimes there is only one Site A1 available while in other cases there are two). One noticeable qualitative difference is that the heat for C<sub>2</sub>H<sub>2</sub> keeps decreasing with coverage, whereas for the unfunctionalized case it plateaus up to the maximum coverage measured. This behavior can be explained by the



**Fig. 4** Induced charge density upon formation of the bond between UiO-66(Zr) and C<sub>2</sub>H<sub>2</sub> (left), C<sub>2</sub>H<sub>4</sub> (middle), and C<sub>2</sub>H<sub>6</sub> (right). Yellow lobes show charge accumulation and blue ones charge depletion. All isolevels are set at 0.0002 e Å<sup>-3</sup>. The interaction of C<sub>2</sub>H<sub>2</sub> with the framework is noticeably stronger compared to C<sub>2</sub>H<sub>4</sub> or C<sub>2</sub>H<sub>6</sub>. Although the perspective is deceiving, the distance of the guest molecule to the water bound to Site A1 increases from the left to the right.



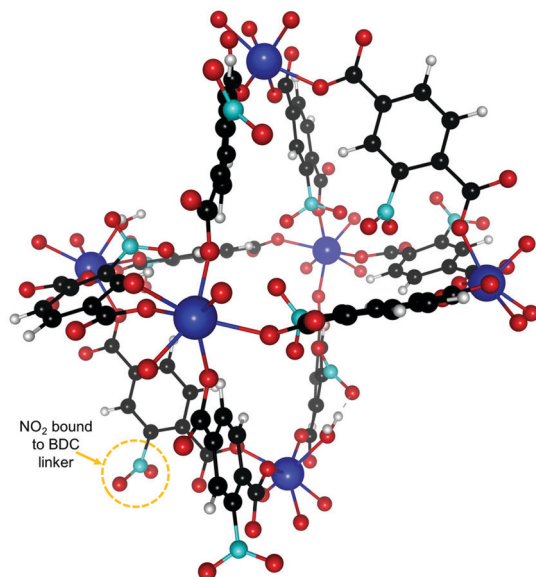


Fig. 5 NO<sub>2</sub> functionalized UiO-66(Zr) unit cell. One NO<sub>2</sub> was attached to every BDC linker. The binding sites are similar to the unfunctionalized case in Fig. 2 and we are thus using the same labeling.

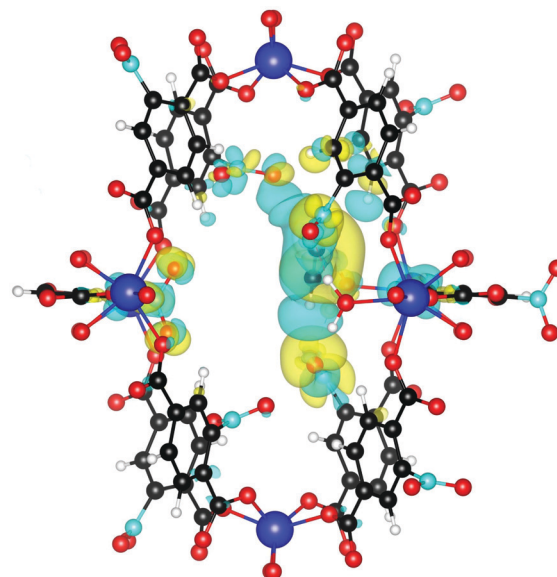


Fig. 7 Induced charge density upon formation of the bond between NO<sub>2</sub> functionalized UiO-66(Zr) and C<sub>2</sub>H<sub>2</sub>. Yellow lobes show charge accumulation and blue ones charge depletion. All isolevels are set at 0.0002 e Å<sup>-3</sup>. Compared to Fig. 4 we see that the guest molecule now also shows additional interactions with three NO<sub>2</sub> functional group, leading to the increased binding energy.

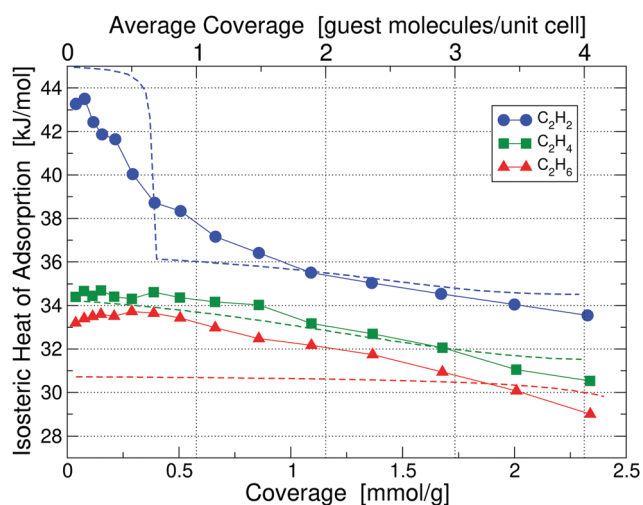


Fig. 6 Isosteric heats of adsorption of C<sub>2</sub>H<sub>2</sub>, C<sub>2</sub>H<sub>4</sub> and C<sub>2</sub>H<sub>6</sub> in UiO-66(Zr)-NO<sub>2</sub> at 195 K, obtained by calorimetry measurements; results for 293 K are shown in Fig. S6 (ESI†). Dashed lines correspond to our partition-function occupancy-probability calculations based on our theoretical binding energies and availability of binding sites.

much smaller number of available secondary binding sites in the functionalized case, which increases the probability of guest molecules occupying tertiary sites earlier.

Our partition-function occupancy-probability calculations again capture the overall trends well and are within chemical accuracy of the experimental results for all coverages. The noticeable kink for C<sub>2</sub>H<sub>2</sub> in our theoretical results, which is not present in the experimental results, is an artefact of our slightly too high calculated primary site energy and probably a slightly underestimated secondary site value. If both values would move 1 kJ mol<sup>-1</sup> in the right direction, the kink would disappear. The slope for C<sub>2</sub>H<sub>6</sub> is also underestimated due to our slightly underestimated primary site value.

Finally, we investigate why the functionalized MOF can bind the guest molecules stronger than the unfunctionalized case. To this end, we plot the induced charge density of C<sub>2</sub>H<sub>2</sub> in Fig. 7. It can be seen that the guest molecule now also shows an additional interaction with three NO<sub>2</sub> functional groups, leading to the increased binding energy. C<sub>2</sub>H<sub>2</sub> is at a distance of only 2.18 Å from the closest NO<sub>2</sub> and at 2.66 Å and 3.02 Å from

**Table 2** Calculated binding energies of C<sub>2</sub>H<sub>2</sub>, C<sub>2</sub>H<sub>4</sub> and C<sub>2</sub>H<sub>6</sub> in NO<sub>2</sub> functionalized UiO-66(Zr) [kJ mol<sup>-1</sup>]. Experimental results for the lowest measured coverage at 195 K and 293 K are given for comparison. Sites are labeled according to Fig. 2. The number of equivalent sites per unit cell is also given. Asterisks for binding sites indicate that the guest molecule is bound close to that site, but in a different orientation

Molecule	Exp. low coverage		Primary			Secondary			Tertiary		
	195 K	293 K	Energy	Site	Equiv.	Energy	Site	Equiv.	Energy	Site	Equiv.
C <sub>2</sub> H <sub>2</sub>	43.27	37.21	45.12	A1	1	37.12	A1*	1	34.51	E	3
C <sub>2</sub> H <sub>4</sub>	34.40	32.41	35.57	A1	1	33.96	A1*	2	31.49	E	2
C <sub>2</sub> H <sub>6</sub>	33.20	33.86	31.15	A1	2	30.75	E	1	26.43	E*	1





