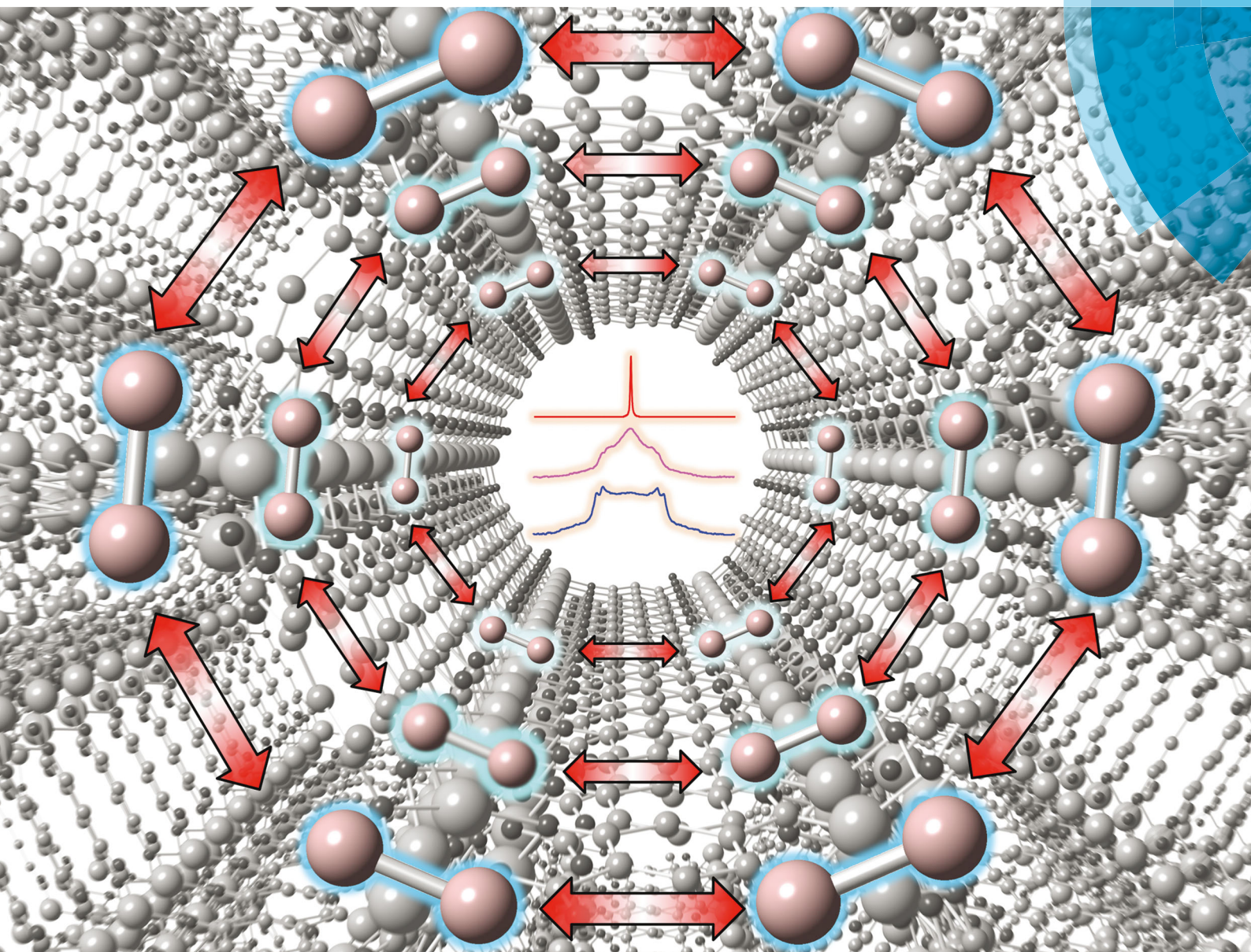


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# Grasping hydrogen adsorption and dynamics in metal–organic frameworks using $^2\text{H}$ solid-state NMR†

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Record greenhouse gas emissions have spurred the search for clean energy sources such as hydrogen ( $\text{H}_2$ ) fuel cells. Metal–organic frameworks (MOFs) are promising  $\text{H}_2$  adsorption and storage media, but knowledge of  $\text{H}_2$  dynamics and adsorption strengths in these materials is lacking. Variable-temperature (VT)  $^2\text{H}$  solid-state NMR (SSNMR) experiments targeting  $^2\text{H}_2$  gas (i.e.,  $\text{D}_2$ ) shed light on  $\text{D}_2$  adsorption and dynamics within six representative MOFs: UiO-66, M-MOF-74 ( $\text{M} = \text{Zn}, \text{Mg}, \text{Ni}$ ), and  $\alpha\text{-M}_3(\text{COOH})_6$  ( $\text{M} = \text{Mg}, \text{Zn}$ ).  $\text{D}_2$  binding is relatively strong in Mg-MOF-74, Ni-MOF-74,  $\alpha\text{-Mg}_3(\text{COOH})_6$ , and  $\alpha\text{-Zn}_3(\text{COOH})_6$ , giving rise to broad  $^2\text{H}$  SSNMR powder patterns. In contrast,  $\text{D}_2$  adsorption is weaker in UiO-66 and Zn-MOF-74, as evidenced by the narrow  $^2\text{H}$  resonances that correspond to rapid reorientation of the  $\text{D}_2$  molecules. Employing  $^2\text{H}$  SSNMR experiments in this fashion holds great promise for the correlation of MOF structural features and functional groups/metal centers to  $\text{H}_2$  dynamics and host–guest interactions.

Carbon dioxide ( $\text{CO}_2$ ) is associated with the greenhouse effect and global warming.  $\text{H}_2$  fuel cells and other “greener” solutions are the future energy sources for automobiles, however, many proposals for  $\text{H}_2$  storage involve tanks for compressed  $\text{H}_2$ ,<sup>1</sup> which is flammable and explosive. Safer alternatives for  $\text{H}_2$  storage, such as crystalline materials,<sup>1,2</sup> are desired. Metal–organic frameworks (MOFs) are ordered three-dimensional structures consisting of metal centers or metal–inorganic units joined by organic linkers. By varying the MOF topology, metal center, and linkers,<sup>3</sup> tailored large surface areas and guest binding strengths are possible. Several MOFs have shown  $\text{H}_2$  adsorption and storage capabilities,<sup>4</sup> including UiO-66,<sup>5,6</sup> MOF-74,<sup>7</sup>  $\alpha\text{-Mg}_3(\text{COOH})_6$ ,<sup>8</sup> and  $\alpha\text{-Zn}_3(\text{COOH})_6$ .<sup>9</sup>

X-ray diffraction is widely used to investigate gas adsorption in MOFs but cannot reliably locate  $\text{H}_2$ . Neutron diffraction can

find  $\text{H}_2$  but little motional details can be obtained. Adsorption isotherms provide  $\text{H}_2$  capacity yet yield little positional or dynamic data. IR spectroscopy may indicate  $\text{H}_2$  adsorption and binding sites, but  $\text{H}_2$  motion remains unknown. Computational methods can estimate  $\text{H}_2$  location and dynamics, yet demand experimental verification. In order to (i) move MOFs toward practical incorporation as  $\text{H}_2$  storage media, and (ii) enhance  $\text{H}_2$  capacity in future MOFs, knowledge of the dynamic behavior of  $\text{H}_2$  in today's MOFs is critical. SSNMR is a sensitive probe of the local nuclear electronic and magnetic environment, and provides rich information on MOFs from the perspective of the metals,<sup>10</sup> organic linkers,<sup>11</sup> guest molecules,<sup>12</sup> and dynamic components.<sup>13,14</sup> NMR interactions are generally anisotropic (directionally-dependent) with respect to the magnetic field and are influenced by dynamics in a predictable manner; rich motional information can be extracted from spectral simulations.<sup>15</sup>

When dynamics are of interest in NMR, the  $^2\text{H}$  isotope is preferred over  $^1\text{H}$ .  $^2\text{H}$  has a spin of 1 and is subject to the anisotropic quadrupolar interaction (QI) between the nuclear quadrupole moment and surrounding electric field gradients (EFGs). Any motion that reorients the  $^2\text{H}$  EFG tensor influences the QI; since the  $^2\text{H}$  SSNMR spectrum is dominated by the QI,  $^2\text{H}$  SSNMR is a powerful probe of guest dynamics.  $^1\text{H}$  is a spin-1/2 nucleus that suffers from very strong  $^1\text{H}$ – $^1\text{H}$  homonuclear dipolar coupling in solids, resulting in broad  $^1\text{H}$  SSNMR spectra that rarely yield useful information.  $^2\text{H}$  SSNMR has proven to be an effective tool for studying the binding in metal–dihydrogen complexes,<sup>16–18</sup> and has been successfully used to probe  $\text{D}_2$  mobility within a Ru-modified MOF.<sup>19</sup> Wright *et al.* have shown that  $^2\text{H}$  SSNMR provides rich information on the dynamics of deuterated linkers and guests in MOFs and microporous materials.<sup>20–24</sup> Herein, we use VT  $^2\text{H}$  SSNMR to probe  $\text{H}_2$  adsorption in a series of different MOFs, studying the significant differences in  $\text{D}_2$  adsorption behavior and dynamics within UiO-66, M-MOF-74 ( $\text{M} = \text{Zn}, \text{Mg}, \text{Ni}$ ), and  $\alpha\text{-M}_3(\text{COOH})_6$  ( $\text{M} = \text{Mg}, \text{Zn}$ ).

UiO-66<sup>6</sup> is a three-dimensional MOF composed of  $\text{Zr}_6\text{O}_4(\text{OH})_4$  units and 1,4-benzenedicarboxylate (BDC) linkers, with large octahedral and tetrahedral cages with pore sizes *ca.* 11 Å and 8 Å,

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† Electronic supplementary information (ESI) available: Full details of MOF synthesis,  $\text{D}_2$  loading, and SSNMR experiments. Illustrations of the MOFs,  $\text{H}_2/\text{D}_2$  adsorption sites in MOF-74, additional  $^2\text{H}$  SSNMR spectra and simulations, tables, and powder XRD patterns are also included. See DOI: 10.1039/c6cc03205b

‡ These authors contributed equally to this work.

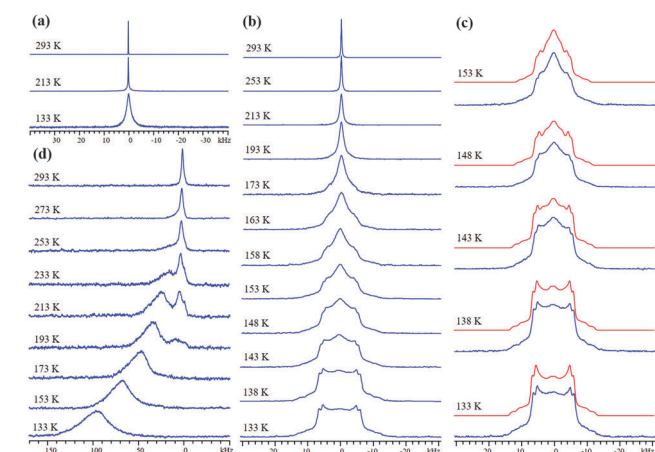




respectively (Fig. S1, ESI†). VT  $^2\text{H}$  SSNMR spectra of  $\text{D}_2$  in UiO-66 (Fig. S2, ESI†) feature a narrow resonance from 293 K to 133 K with a full width at half height (FWHH) of 45 Hz throughout. Although  $\text{D}_2$  is adsorbed in UiO-66, the large pores permit rapid diffusion of  $\text{D}_2$  guests through the MOF void space,<sup>5</sup> eliminating the spectral broadening effects of the QI and giving rise to a sharp, motionally-averaged  $^2\text{H}$  resonance. The narrow lineshape indicates that any localized MOF- $\text{D}_2$  interactions are relatively weak and easily overcome by diffusion. The tetrahedral and octahedral pore geometries may also contribute to the observed narrow resonance.

M-MOF-74 (M = metal) is composed of metal centers connected by 2,5-dioxido-1,4-benzenedicarboxylate (dobdc) linkers that form honeycomb-shaped channels *ca.* 11 Å wide with metal centers at each vertex (Fig. S3, ESI†). Each metal center is connected to five oxygen atoms from four linkers in the as-made MOF, along with a sixth oxygen from a water molecule that can be removed *via* activation. The resulting coordinatively-unsaturated open metal site (OMS) strongly encourages adsorption of  $\text{H}_2$ .<sup>7,25</sup> The nature of the metal center influences  $\text{H}_2$  binding affinity, which increases in the order  $\text{Zn} < \text{Mg} < \text{Ni}$ .<sup>7</sup> The  $^2\text{H}$  SSNMR spectrum of  $\text{D}_2$  in Zn-MOF-74 features a narrow resonance of 80 Hz FWHH at 293 K (Fig. 1(a)), implying that  $\text{D}_2$  undergoes rapid isotropic reorientation in the MOF. The FWHH increases to 120 Hz at 213 K and 1920 Hz at 133 K. At lower temperatures,  $\text{D}_2$  motion is reduced, leading to a large increase in FWHH. The  $^2\text{H}$  resonance widths in Zn-MOF-74 are much larger than in UiO-66 at all temperatures, indicating that  $\text{D}_2$  is considerably less mobile in Zn-MOF-74 due to metal- $\text{H}_2$  interactions; however, the lack of spectral features at 133 K confirms that  $\text{D}_2$  dynamics remain rapid, and efficient  $\text{H}_2$  diffusion pathways in the MOF-74 family are known to exist.<sup>26</sup>

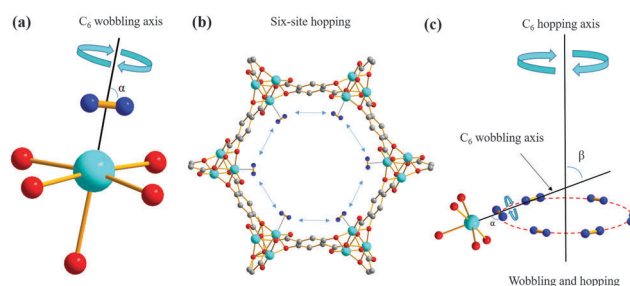
Mg-MOF-74 has a higher affinity for  $\text{H}_2/\text{D}_2$  and yields the VT SSNMR spectra in Fig. 1(b). The narrow resonance at 293 K corresponds to highly mobile  $\text{D}_2$  gas, while emerging broad spectral features at 173 K indicate the onset of significant  $\text{D}_2$  adsorption.



**Fig. 1** The experimental static VT  $^2\text{H}$  SSNMR spectra of adsorbed  $\text{D}_2$  within (a) Zn-MOF-74, (b and c) Mg-MOF-74, and (d) Ni-MOF-74 are depicted. Experimental (blue) and simulated (red)  $^2\text{H}$  SSNMR spectra in Mg-MOF-74 at low temperatures are shown in (c). The sharp resonance at *ca.* 0 kHz in all spectra at high temperatures corresponds to mobile or very weakly adsorbed  $\text{D}_2$ ; this resonance is broadened in (d) due to the influence of the paramagnetic Ni metal center.

At 153 K, adsorbed  $\text{D}_2$  gives rise to two broad  $^2\text{H}$  powder patterns along with a narrow central resonance from free  $\text{D}_2$  in the center of the pores. The wide  $^2\text{H}$  lineshapes at 133 K are intense, well-defined, and associated with most of the  $\text{D}_2$  in Mg-MOF-74 (Fig. 1(c)). By comparing the  $^2\text{H}$  quadrupolar coupling constant ( $C_Q$ ) of gaseous  $\text{D}_2$  (225 kHz<sup>27</sup>) to the apparent  $C_{QS}$  of  $\text{D}_2$  in Mg-MOF-74 (Table S1, ESI†),  $\text{D}_2$  dynamic information can be extracted.<sup>18</sup> Observed  $C_Q(^2\text{H})$  values in Mg-MOF-74 are far less than 225 kHz, and are also smaller than the  $C_Q(^2\text{H})$  range of *ca.* 30–120 kHz associated with  $\eta^2$  bonding between a metal and  $\text{D}_2$  in metal-dihydrogen complexes,<sup>18,28–30</sup> implying that fast  $\text{D}_2$  motion exists here and that the metal- $\text{D}_2$  interactions in Mg-MOF-74 are weaker than the bonding in metal-dihydrogen complexes. It should be noted that at the experimental temperatures and magnitudes of quadrupolar coupling in this study, any spectral effects arising from  $\text{D}_2$  rotational tunneling are expected to be negligible.<sup>31,32</sup>

The same two powder patterns and NMR parameters persist, along with the sharp component, when the loading level is halved to 0.1  $\text{D}_2/\text{Mg}$  (Fig. S4 and S5, Table S2, ESI†). The two broad  $^2\text{H}$  powder patterns of  $\text{D}_2$  in Mg-MOF-74 reveal that there are two similar, but nonequivalent  $\text{D}_2$  molecules ( $\text{D}_2(1)$  and  $\text{D}_2(2)$ ) adsorbed on two different OMSs, as indicated by their QI parameters and  $\text{D}_2$  motions (see Appendix A in the ESI†), while the narrow third resonance indicates that highly mobile  $\text{D}_2$  is also present. Previous studies have suggested that there may be two very similar  $\text{H}_2$  adsorption sites of nearly identical adsorption enthalpy localized on the OMS near our experimental loading levels with slightly different interaction geometries.<sup>33,34</sup> There are 6 OMSs located near the same cross-sectional plane per channel (Fig. 2(b)). At a low loading level of 0.1  $\text{D}_2/\text{metal}$ , on average, there is less than one  $\text{D}_2$  guest per channel cross-section. Thus, it is reasonable to assume that no two  $\text{D}_2$  molecules are adsorbed simultaneously on the same OMS at any given time. It is likely that each channel cross-section contains only 1  $\text{D}_2$  molecule that is in rapid exchange among the 6 sites (*vide infra*). Based on the intensity ratios in Table S1 (ESI†), *ca.* 59% of the channel cross-sections



**Fig. 2** The localized  $C_6$  rotation, or wobbling, of  $\text{D}_2$  molecules adsorbed on the OMS is shown in (a). The black line represents the metal- $\text{D}_2$  vector; the cone of  $\text{D}_2$  wobbling movement traces out an angle  $\alpha$  about the wobbling axis. A schematic of  $\text{D}_2$  hopping in MOF-74 is shown in (b). The combination of wobbling ( $C_6$  rotation,  $\alpha$ ) and six-site hopping ( $C_6$  rotation,  $\beta$ ) of  $\text{D}_2$  molecules in MOF-74 is shown in (c), where the wobbling axis makes an angle  $\beta$  with respect to the hopping axis. The colors red, grey, blue and cyan correspond to oxygen, carbon, deuterium and the metal center, respectively. Our data indicate that there are two very similar  $\text{D}_2$  adsorption sites on the OMS, giving rise to two separate  $^2\text{H}$  SSNMR powder patterns.



are populated by D<sub>2</sub>(1) and 41% of the channel cross-sections are occupied by D<sub>2</sub>(2).

Using the known <sup>2</sup>H QI parameters of D<sub>2</sub> gas ( $C_Q(^2\text{H}) = 225 \text{ kHz}$ ,  $\eta_Q = 0$ ),<sup>27</sup> simulations<sup>15</sup> of motionally-averaged <sup>2</sup>H spectra (Fig. S6, ESI†) reveal that common D<sub>2</sub> dynamics exist at the two similar adsorption sites on the OMS in Mg-MOF-74 (Fig. 2 and Table S3, ESI†). D<sub>2</sub> undergoes a local rotation or “wobbling” modeled by a sixfold rotation about the minimum energy configuration with respect to the OMS (Fig. 2(a)), as well as a non-localized six-site hopping along the pore edge (Fig. 2(b)). The combined motional model is shown in Fig. 2(c). The wobbling describes a rotation of the D–D bond about the axis passing through the OMS (Fig. 2(a)), as defined by angle  $\alpha$ . It is likely that the wobbling rotation represents a model for some kind of rotational diffusion on the OMS. The hopping occurs between six nearly coplanar OMSs, where three reside in one plane, and the other three OMSs are in a very proximate plane slightly offset along the longitudinal direction of the channel. It should be noted that since the wobbling and hopping motional rates are in the fast exchange limit, there are an infinite number of  $C_n$  ( $n \geq 3$ ) jumping motions that could lead to the observed powder patterns. The dynamic behavior of D<sub>2</sub> at both adsorption sites is strikingly similar, supporting the notion of two H<sub>2</sub> adsorption sites of nearly identical interaction geometry and enthalpy on the OMS: each has D<sub>2</sub> wobbling angles increasing from *ca.* 74° at 293 K to 82° at 133 K, along with constant hopping angles of *ca.* 60° (Table S3, ESI†). The larger wobbling angle at low temperatures enhances the interaction of D<sub>2</sub> with the Mg site. It should also be noted that simulation of fast-exchange SSNMR spectra can sometimes lead to more than one motional model. However, we were unable to simulate the split “horns” or characteristic “shoulders” of the low-temperature spectra using a single adsorption site with any type of possible motions (Appendix A). These results also suggest that there does not seem to be significant exchange of D<sub>2</sub> between the two adsorption sites; increasing the possibility that the sites are somehow segregated or correspond to slightly different types of OMSs in separate MOF channels. The calculated H<sub>2</sub> position in Co-MOF-74<sup>35</sup> strongly agrees with the orientations in our motional model, while the types of D<sub>2</sub> dynamics resemble those of CO<sub>2</sub> and CO in MOF-74.<sup>36–38</sup> To confirm at low temperatures that the D<sub>2</sub> motional rate remains in the fast (*i.e.*,  $\geq 10^7 \text{ Hz}$ ) and not in the intermediate regime, quadrupolar echo experiments with different inter-pulse delays were performed, resulting in unchanged spectra (Fig. S7 and S8, Table S4, ESI†).

Ni-MOF-74 has a stronger H<sub>2</sub>/D<sub>2</sub> binding affinity.<sup>7</sup> Unlike diamagnetic Zn<sup>2+</sup> and Mg<sup>2+</sup>, Ni<sup>2+</sup> is paramagnetic in this MOF. The <sup>2</sup>H magnetic dipole couples with those of proximate unpaired electrons, resulting in spectral broadening and unusual chemical shifts when D<sub>2</sub> is near Ni<sup>2+</sup>. VT <sup>2</sup>H SSNMR spectra of D<sub>2</sub> in Ni-MOF-74 (Fig. 2(d)) at room temperature reveal that D<sub>2</sub> is mobile and/or rapidly exchanges with adsorbed D<sub>2</sub>. The broad 1.2 kHz FWHH confirms the proximity of D<sub>2</sub> to Ni<sup>2+</sup>. At 273 K, a second, broader resonance emerges at 140 ppm, corresponding to D<sub>2</sub> adsorbed on the OMS. At 213 K, the broad resonance is at 375 ppm and is the dominant feature. The broad resonance increases in frequency,

width, and relative intensity as temperature is reduced; the narrower resonance at lower frequency vanishes at 173 K, implying that most D<sub>2</sub> is adsorbed. At 133 K a broad, featureless <sup>2</sup>H resonance of FWHH 30 kHz centered at about 1550 ppm is evident, confirming that a majority of D<sub>2</sub> is adsorbed onto the OMS. The featureless nature of these resonances prohibits detailed analysis.<sup>39</sup>

$\alpha\text{-Mg}_3(\text{COOH})_6$  and  $\alpha\text{-Zn}_3(\text{COOH})_6$  are porous MOFs<sup>8,9,40</sup> with one-dimensional zig-zag channels of small *ca.* 4–5 Å diameter (Fig. S9, ESI†). The <sup>2</sup>H spectrum of D<sub>2</sub> in  $\alpha\text{-Mg}_3(\text{COOH})_6$  at 293 K features a narrow resonance of 375 Hz FWHH (Fig. S10(a), ESI†) arising from mobile D<sub>2</sub>, flanked by a less intense powder pattern 40 kHz broad at 193 K from adsorbed D<sub>2</sub>, in agreement with adsorption isotherms.<sup>8,40</sup> At 133 K the broad powder patterns are of higher S/N, reflecting increased D<sub>2</sub> adsorption in  $\alpha\text{-Mg}_3(\text{COOH})_6$ , and are convoluted with a broad distribution of intensity possibly from disordered D<sub>2</sub>; other small guests such as ethanol and methanol are also disordered in this MOF.<sup>40</sup> The <sup>2</sup>H SSNMR spectra of D<sub>2</sub> in  $\alpha\text{-Zn}_3(\text{COOH})_6$  are of similar lineshape and breadth (Fig. S10(b), ESI†), implying that both MOFs have similar H<sub>2</sub> adsorption strengths. The intense narrow resonance persists at lower temperatures in  $\alpha\text{-Zn}_3(\text{COOH})_6$ , indicating that there is more mobile H<sub>2</sub> in  $\alpha\text{-Zn}_3(\text{COOH})_6$  at all temperatures. The complicated powder patterns are clear evidence of multiple D<sub>2</sub> adsorption sites within both  $\alpha\text{-Mg}_3(\text{COOH})_6$  and  $\alpha\text{-Zn}_3(\text{COOH})_6$ . At 0.1 D<sub>2</sub>/metal loading, in addition to free gas, <sup>2</sup>H spectra may be simulated using three and two adsorption sites in  $\alpha\text{-Mg}_3(\text{COOH})_6$  and  $\alpha\text{-Zn}_3(\text{COOH})_6$ , respectively. In strong agreement with our experimental observations is a very recent study that has located three main H<sub>2</sub> adsorption sites in  $\alpha\text{-Mg}_3(\text{COOH})_6$ .<sup>41</sup>

The <sup>2</sup>H spectrum of  $\alpha\text{-Zn}_3(\text{COOH})_6$  is less ambiguous, of higher resolution, and is more straightforward to simulate (Fig. S10(c), Tables S3 and S5, ESI†). The well-defined <sup>2</sup>H SSNMR spectra of D<sub>2</sub> in  $\alpha\text{-Zn}_3(\text{COOH})_6$  yield relatively high  $\eta_Q$  values along with  $C_Q$  values *ca.* 5–10 kHz greater than those of D<sub>2</sub> in Mg-MOF-74 at all equivalent temperatures, reflecting reduced D<sub>2</sub> mobility and perhaps a stronger D<sub>2</sub>-MOF interaction in  $\alpha\text{-Zn}_3(\text{COOH})_6$ . In contrast, the very complicated <sup>2</sup>H spectra of  $\alpha\text{-Mg}_3(\text{COOH})_6$  permit only a preliminary motional simulation at this time (Fig. S11, ESI†). A combined wobbling and hopping of D<sub>2</sub> occurs at all adsorption sites in both systems. D<sub>2</sub> hops between two adsorption sites in  $\alpha\text{-Zn}_3(\text{COOH})_6$ , reminiscent of the twofold hopping by CO<sub>2</sub> guests in the  $\alpha\text{-M}_3(\text{COOH})_6$  MOF.<sup>42</sup> It is notable that the changes in D<sub>2</sub> motional angles within Zn are not very pronounced across the experimental temperature range (Table S3, ESI†). At this point, the nature of the adsorption sites is unclear from SSNMR experiments. We are currently performing more detailed studies in order to extract detailed D<sub>2</sub> motional and adsorption information from both  $\alpha\text{-M}_3(\text{COOH})_6$  MOFs.

<sup>2</sup>H SSNMR spectroscopy has revealed unique insights into H<sub>2</sub> dynamics and adsorption locations within these six MOFs; this information is unavailable or very difficult to obtain using traditional MOF characterization methods such as adsorption isotherms and X-ray diffraction. Complementary methods such as neutron diffraction, inelastic neutron scattering, and VT infrared spectroscopy are necessary to locate adsorption locations, but yield limited motional data. Variation of the MOF topology,



metal center, and linker gives rise to very different D<sub>2</sub> dynamic behaviors and adsorption strengths. Further studies employing lower temperatures (*i.e.*, <100 K) are necessary to reduce H<sub>2</sub> mobility and obtain richer dynamic knowledge in these systems, however, these experiments are not yet possible with the equipment available to us. We have used relatively low levels of D<sub>2</sub> loading to simplify these systems for spectral simulations; the next step is to explore higher D<sub>2</sub> loading levels in order to understand the implications for H<sub>2</sub> motion in a practical hydrogen storage setting. The comprehensive molecular-level knowledge of H<sub>2</sub> dynamics and adsorption in MOFs available from <sup>2</sup>H SSNMR spectroscopy will undoubtedly assist in establishing clear links between H<sub>2</sub> dynamics and high-capacity H<sub>2</sub> storage in porous materials, with clear applications in green energy solutions.

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