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## Evaluating solvothermal and mechanochemical routes towards the metal–organic framework $Mg_2(m\text{-dobdc})^\ddagger$

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Metal–organic frameworks bearing coordinatively unsaturated Mg(II) sites are promising materials for gas storage, chemical separations, and drug delivery due to their low molecular weights and lack of toxicity. However, there remains a limited number of such MOFs reported in the literature. Herein, we investigate the gas sorption properties of the understudied framework  $Mg_2(m\text{-dobdc})$  ( $dobdc^{4-} = 4,6\text{-dioxido-1,3-benzenedicarboxylate}$ ) synthesized under both solvothermal and mechanochemical conditions. Both materials are found to be permanently porous, as confirmed by 77 K  $N_2$  adsorption measurements. In particular,  $Mg_2(m\text{-dobdc})$  synthesized under mechanochemical conditions using exogenous organic base displays one of the highest capacities reported to date (6.14 mmol g<sup>-1</sup>) for  $CO_2$  capture in a porous solid under simulated coal flue gas conditions (150 mbar, 40 °C). As such, mechanochemically synthesized  $Mg_2(m\text{-dobdc})$  represents a promising new framework for applications requiring high gas adsorption capacities in a porous solid.

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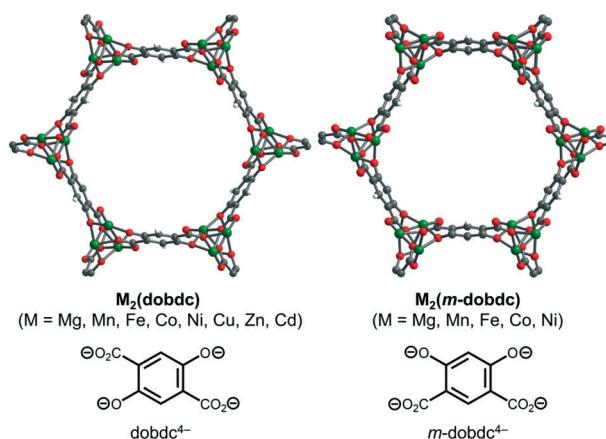
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## Introduction

Metal–organic frameworks (MOFs) are porous, crystalline extended solids that consist of inorganic metal nodes or secondary building units (SBUs) bridged by polytopic organic linkers.<sup>1</sup> Their uniquely modular structures coupled with high internal surface areas have enabled numerous applications in drug delivery, catalysis, chemical separations, and gas storage.<sup>2–5</sup> Frameworks containing coordinatively unsaturated metal centers, also known as open-metal site MOFs, have been extensively studied due to their ability to strongly interact with guest molecules.<sup>6</sup> In particular, the canonical  $M_2(dobdc)$  ( $M = Mg, Mn, Fe, Co, Ni, Cu, Zn, Cd$ ;  $dobdc^{4-} = 2,5\text{-dioxido-1,4-benzenedicarboxylate}$ ), MOF-74, or CPO-27 family of frameworks features hexagonal one-dimensional channels decorated with a high density of exposed M(II) centers (Fig. 1, left).<sup>7</sup> Among the reported isostructural metal variants, the Mg analogue,  $Mg_2(dobdc)$ , is particularly promising due to its low cost, lack of toxicity, and high gravimetric and volumetric adsorption capacities for a range of adsorbates.<sup>8–12</sup> As such, the identification of new porous frameworks bearing accessible Mg(II) sites is highly desirable.

A closely related family of frameworks,  $M_2(m\text{-dobdc})$  ( $M = Mg, Mn, Fe, Co, Ni; m\text{-dobdc}^{4-} = 4,6\text{-dioxido-1,3-benzenedicarboxylate}$ ), have been reported to possess an even higher density of exposed cationic sites than MOF-74 materials due to slight differences in the ligand field around the metal center.<sup>13</sup> However,  $Mg_2(m\text{-dobdc})$  prepared under solvothermal conditions was initially reported to be non-porous due to difficulties associated with removing coordinating solvents such as *N,N*-dimethylformamide (DMF)



**Fig. 1** Structures of  $M_2(dobdc)$  ( $M = Mg, Mn, Fe, Co, Ni, Cu, Zn; dobdc^{4-} = 2,5\text{-dioxido-1,4-benzenedicarboxylate}$ , left) and  $M_2(m\text{-dobdc})$  ( $M = Mg, Mn, Fe, Co, Ni; m\text{-dobdc}^{4-} = 4,6\text{-dioxido-1,3-benzenedicarboxylate}$ , right). Green, gray, white, and red spheres correspond to magnesium, carbon, hydrogen, and oxygen, respectively.

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or methanol (MeOH) from the framework pores.<sup>13</sup> Although  $Mg_2(m\text{-dobdc})$  was later reported to be porous when prepared under mechanochemical conditions,<sup>14</sup> much about its intrinsic gas sorption properties, such as the accessibility of the Mg(II) sites to guest molecules,<sup>8</sup> remains relatively unknown.

Herein, we systematically investigate the synthesis and gas sorption properties of fully desolvated  $Mg_2(m\text{-dobdc})$ . Careful washing and activation of  $Mg_2(m\text{-dobdc})$  prepared under traditional solvothermal (ST) conditions, termed  $Mg_2(m\text{-dobdc})\text{-ST}$ , enables access to a material with a similar 77 K  $N_2$  Brunauer–Emmett–Teller (BET) surface area as closely related  $Mg_2(dobdc)$ . In addition, we report an improved mechanochemical (MC) method to reliably prepare  $Mg_2(m\text{-dobdc})\text{-MC}$  using exogenous organic base.<sup>14</sup> Beyond the inherent advantages in scalability and waste minimization offered by mechanochemical syntheses,<sup>15–17</sup>  $Mg_2(m\text{-dobdc})\text{-MC}$  exhibits higher gas adsorption capacities than  $Mg_2(m\text{-dobdc})\text{-ST}$ . Indeed, at 150 mbar of  $CO_2$  and 40 °C, conditions relevant to  $CO_2$  capture from coal flue gas,<sup>18</sup>  $Mg_2(m\text{-dobdc})\text{-MC}$  exhibits an higher  $CO_2$  uptake (6.14 mmol g<sup>-1</sup>) than even  $Mg_2(dobdc)$  (5.28 mmol g<sup>-1</sup>).<sup>12,19</sup> This represents one of the highest capacities reported to date for  $CO_2$  capture in a porous solid under simulated coal flue gas conditions.<sup>20</sup> Overall, our findings suggest that  $Mg_2(m\text{-dobdc})$  prepared under mechanochemical conditions represents a promising and scalable alternative to ubiquitous  $Mg_2(dobdc)$  for applications in chemical separations, gas storage, and beyond.

## Results and discussion

We commenced our studies into the synthesis and gas sorption properties of  $Mg_2(m\text{-dobdc})$  by optimizing the synthesis of  $H_4m\text{-dobdc}$  (Fig. 2). In our hands, the previously reported synthesis of  $H_4m\text{-dobdc}$  from resorcinol *via* the Kolbe–Schmitt reaction was only modestly reproducible, often yielding the monocarboxylic acid **1** instead of  $H_4m\text{-dobdc}$  (Fig. 2). Two modifications to the standard preparation were identified to make the synthesis of  $H_4m\text{-dobdc}$  more reliable (see ESI† section 2 for details). First, starting from commercially available **1** in place of resorcinol improved the reproducibility of the reaction, likely because only a single carboxylation reaction must take place to yield  $H_4m\text{-dobdc}$ .<sup>21</sup> Second, the reaction temperature (250 °C) was found to be a critical parameter and best monitored using an internal thermocouple placed directly in the solvent-free reaction mixture. With these modifications in place, we were able to reliably synthesize  $H_4m\text{-dobdc}$  on >5 g scale in a single batch.

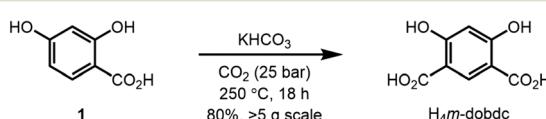


Fig. 2 Synthesis of  $H_4m\text{-dobdc}$  from **1**.

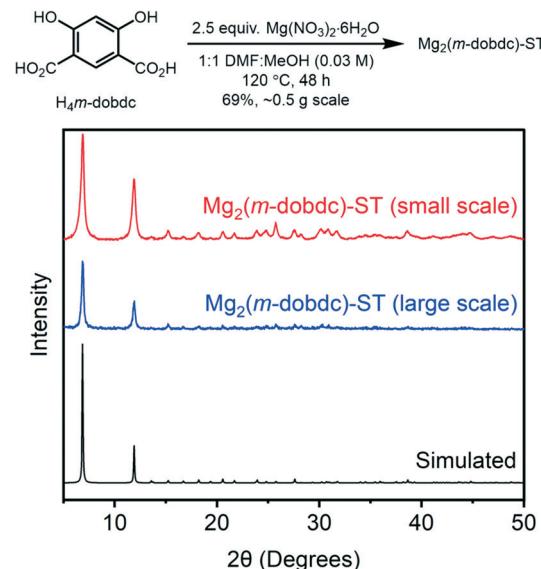


Fig. 3 PXRD ( $\lambda = 1.5406 \text{ \AA}$ ) patterns of  $Mg_2(m\text{-dobdc})\text{-ST}$  synthesized on small and large scale under solvothermal conditions. The simulated pattern based on the previously reported single-crystal X-ray diffraction structure of the isostructural framework  $Co_2(m\text{-dobdc})$  is included for reference.<sup>25</sup>

The previously reported small-scale solvothermal synthesis of  $Mg_2(m\text{-dobdc})$  employed  $Mg(NO_3)_2\cdot 6H_2O$  as the Mg precursor in 2:1 DMF:MeOH at 120 °C.<sup>13</sup> In order to identify the optimal solvothermal conditions for preparing  $Mg_2(m\text{-dobdc})$ , several combinations of amide (DMF or *N,N*-dimethylacetamide) and alcohol (MeOH, ethanol,  $H_2O$ ) solvents were evaluated (ESI† Table S1, see ESI† section 3 for details). Other methods previously reported for the

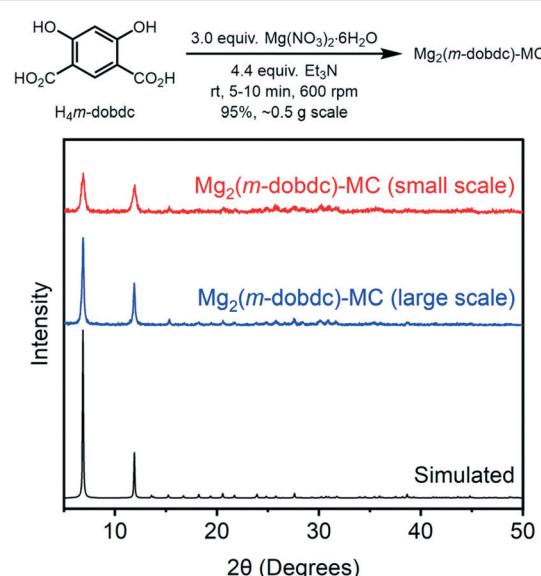


Fig. 4 PXRD ( $\lambda = 1.5406 \text{ \AA}$ ) patterns of  $Mg_2(m\text{-dobdc})\text{-MC}$  synthesized on small and large scale under mechanochemical conditions. The simulated pattern based on the previously reported single-crystal X-ray diffraction structure of the isostructural framework  $Co_2(m\text{-dobdc})$  is included for reference.<sup>25</sup>

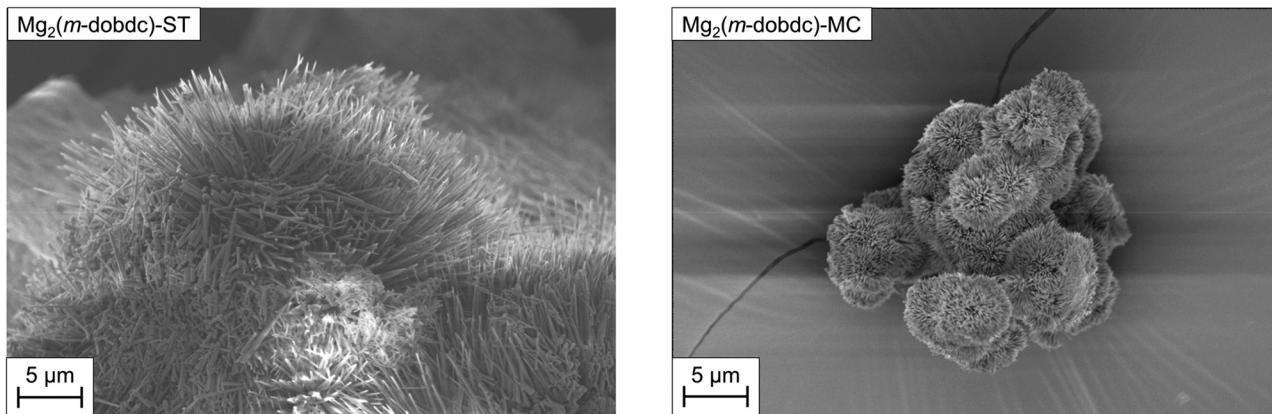


Fig. 5 SEM images of Mg<sub>2</sub>(m-dobdc)-ST (left) and Mg<sub>2</sub>(m-dobdc)-MC (right).

preparation of Mg<sub>2</sub>(dobdc), such as employing Mg(OAc)<sub>2</sub>·4H<sub>2</sub>O as a basic Mg precursor, were tested as well.<sup>22–24</sup> Characterization of the produced solids by powder X-ray diffraction (PXRD)<sup>25</sup> validated that combining H<sub>4</sub>m-dobdc and Mg(NO<sub>3</sub>)<sub>2</sub>·6H<sub>2</sub>O in 1:1 DMF:MeOH (0.03 M) at 120 °C for 48 h was optimal to yield highly crystalline Mg<sub>2</sub>(m-dobdc)-ST (Fig. 3, ESI† Fig. S5). This synthesis can be readily scaled to produce Mg<sub>2</sub>(m-dobdc)-ST on at least 0.5 g scale (Fig. 3, ESI† Fig. S6) and is reproducible as well (ESI† Fig. S17). Soaking the resulting MOF in DMF at 120 °C to remove residual starting materials, then in MeOH at 60 °C to remove DMF, and then in acetone at room temperature to remove MeOH, was sufficient to remove coordinating solvents and soluble impurities from Mg<sub>2</sub>(m-dobdc)-ST, as confirmed by acid digestion and analysis of the resulting solution by <sup>1</sup>H NMR (ESI† Fig. S13). Unfortunately, higher reaction concentrations (>0.1 M) led to impure materials (not shown), limiting the scalability of this solvothermal method.<sup>26</sup>

Having optimized the solvothermal synthesis of Mg<sub>2</sub>(m-dobdc), we set out to improve the mechanochemical synthesis of this material for comparison. Previously, we reported the mechanochemical synthesis of Mg<sub>2</sub>(dobdc) using *N,N*-diisopropylethylamine (Hünig's base) as both the base required to deprotonate the linker precursor and as the liquid to facilitate liquid-assisted grinding in a planetary ball mill.<sup>14</sup> This method could be generalized to the synthesis of porous Mg<sub>2</sub>(m-dobdc) with modest crystallinity. We hypothesized that careful optimization of the Mg precursor (Mg(NO<sub>3</sub>)<sub>2</sub>·6H<sub>2</sub>O, Mg(OAc)<sub>2</sub>·4H<sub>2</sub>O, or MgO), base (Hünig's base, Et<sub>3</sub>N, or 2,6-lutidine), and grinding time (1, 5, or 10 min) would enable the synthesis of Mg<sub>2</sub>(m-dobdc)-MC with

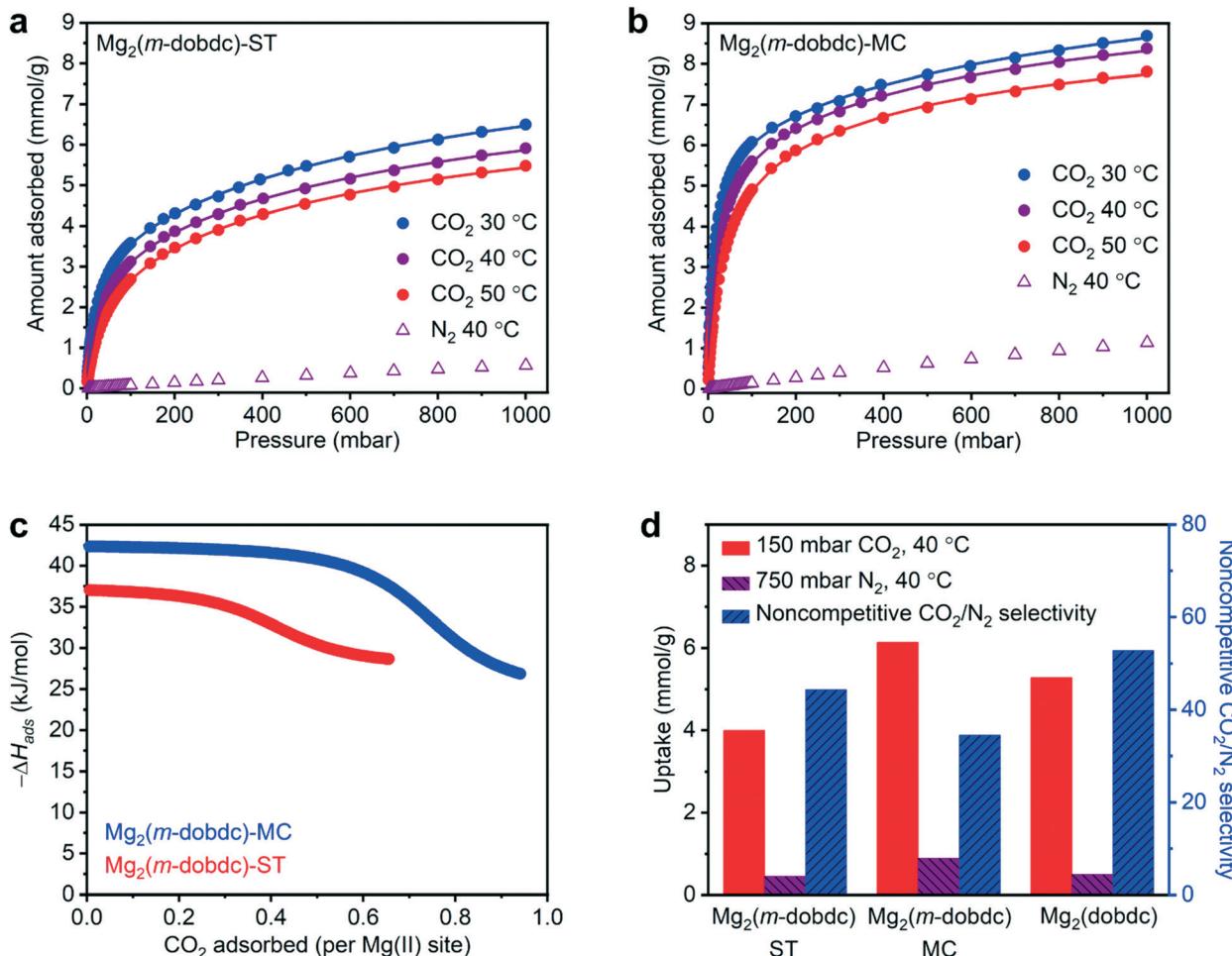
maximum crystallinity and porosity (see ESI† section 4 for details). Indeed, the combination of Mg(NO<sub>3</sub>)<sub>2</sub>·6H<sub>2</sub>O, Et<sub>3</sub>N, and 5–10 min grinding time was optimal to yield crystalline Mg<sub>2</sub>(m-dobdc)-MC (Fig. 4, ESI† Fig. S18 and S19). This method could be reproducibly carried out with 10 min of grinding at 600 rpm to produce Mg<sub>2</sub>(m-dobdc)-MC on 0.5 g scale in excellent yield (Fig. 3, ESI† Fig. S20 and S31). Notably, this mechanochemical synthesis bypasses the use of toxic DMF,<sup>27</sup> representing a green alternative to the solvothermal synthesis of Mg<sub>2</sub>(m-dobdc)-ST. No MOF was obtained with 2,6-lutidine, likely because it is not basic enough to fully deprotonate H<sub>4</sub>(m-dobdc).

With optimized samples of Mg<sub>2</sub>(m-dobdc)-ST and Mg<sub>2</sub>(m-dobdc)-MC in hand, we compared their crystallite morphologies (Fig. 5) and porosities (Table 1) in order to understand how synthesis procedure affects the physical properties of Mg<sub>2</sub>(m-dobdc). Characterization of Mg<sub>2</sub>(m-dobdc)-ST by scanning electron microscopy (SEM) revealed that it is composed of crystalline needles >5 μm in length on average (Fig. 5 left, ESI† Fig. S12). A needle-like morphology for Mg<sub>2</sub>(m-dobdc)-ST is consistent with that previously reported for single crystals of the isostructural framework Co<sub>2</sub>(m-dobdc).<sup>25,28</sup> Similarly, Mg<sub>2</sub>(m-dobdc)-MC is comprised of needles <1 μm in length (Fig. 5, right, ESI† Fig. S26). The smaller crystallites for mechanochemically synthesized Mg<sub>2</sub>(m-dobdc)-MC likely arise due to rapid deprotonation of H<sub>4</sub>m-dobdc by triethylamine during the reaction.<sup>14,29</sup>

Careful activation of Mg<sub>2</sub>(m-dobdc)-ST and Mg<sub>2</sub>(m-dobdc)-MC under high vacuum (<10 μbar) at 180 °C for at least 24 h was sufficient to fully remove solvent molecules from both frameworks. Their porosities were assessed by collecting 77 K N<sub>2</sub> adsorption isotherms (ESI† Fig. S9 and S23). As expected, Mg<sub>2</sub>(m-dobdc)-ST and Mg<sub>2</sub>(m-dobdc)-MC are both microporous, with BET and Langmuir surface areas comparable to those reported for related Mg<sub>2</sub>(dobdc) (Table 1).<sup>10,12</sup> The BET surface area of Mg<sub>2</sub>(m-dobdc)-MC (1653 ± 2 m<sup>2</sup> g<sup>-1</sup>) is somewhat higher than that of Mg<sub>2</sub>(m-dobdc)-ST (1556 ± 2), reflecting a greater degree of accessible pores and/or the presence of insoluble, amorphous impurities in the latter material. We note that heating Mg<sub>2</sub>(m-dobdc) with

Table 1 77 K N<sub>2</sub> BET and Langmuir surface areas of Mg<sub>2</sub>(m-dobdc)-ST and Mg<sub>2</sub>(m-dobdc)-MC. Literature values reported for related Mg<sub>2</sub>(dobdc) are included for comparison

Material	BET surface area (m <sup>2</sup> g <sup>-1</sup> )	Langmuir surface area (m <sup>2</sup> g <sup>-1</sup> )
Mg <sub>2</sub> (m-dobdc)-ST	1556 ± 2	1971 ± 3
Mg <sub>2</sub> (m-dobdc)-MC	1653 ± 2	1964 ± 38
Mg <sub>2</sub> (dobdc)	1495 (ref. 12)	1905–1957 (ref. 10 and 12)



**Fig. 6** 30 °C, 40 °C, and 50 °C CO<sub>2</sub> and 40 °C N<sub>2</sub> adsorption isotherms in a) Mg<sub>2</sub>(*m*-dobdc)-ST and b) Mg<sub>2</sub>(*m*-dobdc)-MC. The lines correspond to individual fits to the dual-site Langmuir model. A data point was considered equilibrated when <0.01% pressure change occurred over a 30 s interval. c) Enthalpy of adsorption ( $-\Delta H_{\text{ads}}$ ) values for Mg<sub>2</sub>(*m*-dobdc)-MC and Mg<sub>2</sub>(*m*-dobdc)-ST determined using simultaneous fits to dual-site Langmuir models. d) Summary of CO<sub>2</sub> and N<sub>2</sub> uptake values and non-competitive CO<sub>2</sub>/N<sub>2</sub> selectivities relevant to CO<sub>2</sub> capture from coal flue gas for Mg<sub>2</sub>(*m*-dobdc)-ST and Mg<sub>2</sub>(*m*-dobdc)-MC. The corresponding values for Mg<sub>2</sub>(dobdc) reported in the literature are included for reference.<sup>19</sup>

ramp rates faster than 1 °C min<sup>-1</sup> or to temperatures greater than 180 °C consistently led to lower surface areas, likely due to partial pore collapse. Nonetheless, these findings confirm that highly porous and crystalline Mg<sub>2</sub>(*m*-dobdc) can be readily prepared under both solvothermal and mechanochemical conditions.

Given the microporosity of both Mg<sub>2</sub>(*m*-dobdc) samples and the high capacities of closely related Mg<sub>2</sub>(dobdc) for a range of adsorbates,<sup>10,12</sup> we assessed the potential suitability of Mg<sub>2</sub>(*m*-dobdc)-ST and Mg<sub>2</sub>(*m*-dobdc)-MC for gas capture applications using CO<sub>2</sub> scrubbing from coal flue gas as a representative separation. As such, CO<sub>2</sub> adsorption and desorption isotherms at 30 °C, 40 °C, and 50 °C and N<sub>2</sub> adsorption and desorption isotherms at 40 °C were collected for Mg<sub>2</sub>(*m*-dobdc)-ST (Fig. 6a, ESI† Fig. S14 and S15) and Mg<sub>2</sub>(*m*-dobdc)-MC (Fig. 6b, ESI† Fig. S28 and S29). In all cases, gas sorption was found to be completely reversible. Both materials exhibit steep uptake at low CO<sub>2</sub> pressures, indicative of strong interaction of CO<sub>2</sub> with exposed Mg(II)

sites.<sup>12</sup> The maximum CO<sub>2</sub> uptakes for both materials at 30 °C and 1 bar of CO<sub>2</sub> are 6.50 mmol g<sup>-1</sup> for Mg<sub>2</sub>(*m*-dobdc)-ST and 8.69 mmol g<sup>-1</sup> for Mg<sub>2</sub>(*m*-dobdc)-MC; the latter value is similar to that predicted for binding one CO<sub>2</sub> per Mg(II) site in this material (8.24 mmol g<sup>-1</sup>). The higher CO<sub>2</sub> capacity of Mg<sub>2</sub>(*m*-dobdc)-MC than Mg<sub>2</sub>(*m*-dobdc)-ST is consistent with its higher 77 K N<sub>2</sub> BET surface area and suggests that the mechanochemically synthesized MOF has more accessible Mg(II) sites.

To confirm that CO<sub>2</sub> binding in Mg<sub>2</sub>(*m*-dobdc) materials likely occurs at coordinatively unsaturated Mg(II) sites, the CO<sub>2</sub> adsorption isotherms were fit using dual-site Langmuir models both independently and simultaneously (ESI† Tables S2 and S4). The independent dual-site Langmuir model fits are included in Fig. 6a and b and represent good fits to the experimental data. Using the Clausius–Clapeyron relationship, the differential enthalpies of adsorption ( $-\Delta H_{\text{ads}}$ ) as a function of CO<sub>2</sub> loading were calculated (Fig. 6c, ESI† Fig. S16 and S30). The  $-\Delta H_{\text{ads}}$  values at low coverage

are comparable for  $\text{Mg}_2(m\text{-dobdc})\text{-MC}$  (42 kJ mol<sup>-1</sup>) and  $\text{Mg}_2(m\text{-dobdc})\text{-ST}$  (37 kJ mol<sup>-1</sup>) and are similar to those previously reported for  $\text{CO}_2$  adsorption in MOFs bearing accessible Mg(II) sites as well (38–43 kJ mol<sup>-1</sup>).<sup>10,30</sup> The  $-\Delta H_{\text{ads}}$  plots support that  $\text{Mg}_2(m\text{-dobdc})\text{-MC}$  contains more accessible Mg(II) sites than  $\text{Mg}_2(m\text{-dobdc})\text{-ST}$ , as the strong binding of  $\text{CO}_2$  drop off at higher loadings in this material (~0.7  $\text{CO}_2$  per Mg vs. ~0.4  $\text{CO}_2$  per Mg in  $\text{Mg}_2(m\text{-dobdc})\text{-ST}$ ).

Among MOFs,  $\text{Mg}_2(\text{dobdc})$  possesses one of the highest reported  $\text{CO}_2$  capacities (5.28 mmol g<sup>-1</sup>) under conditions relevant to  $\text{CO}_2$  capture from coal flue gas (150 mbar, 40 °C) (Fig. 6d).<sup>10,18,19</sup> Coupled with minimal uptake of  $\text{N}_2$  at 750 mbar and 40 °C (0.50 mmol g<sup>-1</sup>),<sup>19</sup> this high  $\text{CO}_2$  uptake at low pressures makes  $\text{Mg}_2(\text{dobdc})$  a promising material for  $\text{CO}_2/\text{N}_2$  separations. While the  $\text{CO}_2$  uptake of  $\text{Mg}_2(m\text{-dobdc})\text{-ST}$  at 150 mbar and 40 °C (3.99 mmol g<sup>-1</sup>) is less than that reported for  $\text{Mg}_2(\text{dobdc})$ , likely due to its dearth of accessible Mg(II) sites,  $\text{Mg}_2(m\text{-dobdc})\text{-MC}$  exhibits a higher capacity for  $\text{CO}_2$  (6.14 mmol g<sup>-1</sup>) than  $\text{Mg}_2(\text{dobdc})$  under these conditions (Fig. 6d). Previous studies have suggested that the metal centers of  $\text{M}_2(m\text{-dobdc})$  MOFs are slightly more Lewis acidic than those of  $\text{M}_2(\text{dobdc})$  MOFs,<sup>13</sup> which may account for the enhanced  $\text{CO}_2$  uptake at low pressures in  $\text{Mg}_2(m\text{-dobdc})\text{-MC}$ . Consistently, the  $\text{N}_2$  capacity of  $\text{Mg}_2(m\text{-dobdc})\text{-MC}$  at 750 mbar and 40 °C (0.89 mmol g<sup>-1</sup>) is higher than that reported for  $\text{Mg}_2(\text{dobdc})$  as well. Given the unclear suitability of calculating selectivities in open metal site MOFs using ideal adsorbed solution theory (IAST),<sup>31</sup> we elected to calculate non-competitive  $\text{CO}_2/\text{N}_2$  selectivities under conditions relevant to coal flue gas capture for  $\text{Mg}_2(m\text{-dobdc})\text{-ST}$ ,  $\text{Mg}_2(m\text{-dobdc})\text{-MC}$ , and  $\text{Mg}_2(\text{dobdc})$  instead (Fig. 6d). The non-competitive  $\text{CO}_2/\text{N}_2$  selectivity calculated for  $\text{Mg}_2(\text{dobdc})$  is the highest (53),<sup>19</sup> followed by  $\text{Mg}_2(m\text{-dobdc})\text{-ST}$  (44), and then  $\text{Mg}_2(m\text{-dobdc})\text{-MC}$  (35). The diminished non-competitive  $\text{CO}_2/\text{N}_2$  selectivity for  $\text{Mg}_2(m\text{-dobdc})\text{-MC}$  is due to the higher uptake of  $\text{N}_2$  in this material. Nonetheless, these findings support that  $\text{Mg}_2(m\text{-dobdc})\text{-MC}$  is competitive with the widely studied MOF  $\text{Mg}_2(\text{dobdc})$  for this representative separation. Further, the superior gas sorption performance of  $\text{Mg}_2(m\text{-dobdc})\text{-MC}$  over  $\text{Mg}_2(m\text{-dobdc})\text{-ST}$  indicates that mechanochemical methods may be preferable for the scalable synthesis of this framework.

## Conclusions

Owing to their low cost and high gravimetric gas storage capacities, MOFs bearing high densities of coordinatively unsaturated Mg(II) centers are highly sought after. We have demonstrated that  $\text{Mg}_2(m\text{-dobdc})$  synthesized under mechanochemical conditions is a promising new Mg-based MOF due to its strong binding and high capacity for  $\text{CO}_2$  at low pressures. Notably, the work presented herein represents a rare example in which a mechanochemically synthesized MOF displays superior gas sorption properties compared to material synthesized under traditional solvothermal conditions.<sup>14–17</sup> Moving forward,

mechanochemical methods will prove to be a valuable alternative to solvothermal syntheses for the preparation of high-quality MOFs for applications in gas storage, chemical separations, and drug delivery.

## Author contributions

P. J. M. and R. M. M. conceived the project. E. Y. C. and R. M. M. carried out all experiments. The manuscript was written through the contributions of all authors, and all authors approved of the final version.

## Conflicts of interest

P. J. M. is listed as an inventor on several patents related to the application of MOFs for gas capture.

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