

Capture of Carbon Dioxide from Air and Flue Gas in the Alkylamine-Appended Metal–Organic Framework $\text{mmen-Mg}_2(\text{dobpdc})$

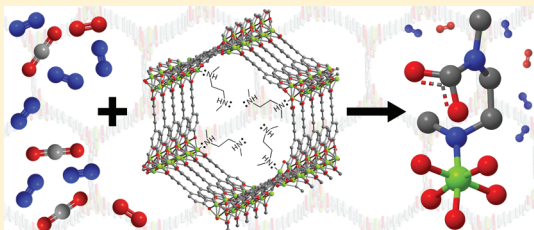
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Supporting Information

ABSTRACT: Two new metal–organic frameworks, $\text{M}_2(\text{dobpdc})$ ($\text{M} = \text{Zn}$ (**1**), Mg (**2**); $\text{dobpdc}^{4-} = 4,4'$ -dioxido-3,3'-biphenyldicarboxylate), adopting an expanded MOF-74 structure type, were synthesized via solvothermal and microwave methods. Coordinatively unsaturated Mg^{2+} cations lining the 18.4-Å-diameter channels of **2** were functionalized with N,N' -dimethylethylenediamine (mmen) to afford $\text{Mg}_2(\text{dobpdc})\text{-}(\text{mmen})_{1.6}(\text{H}_2\text{O})_{0.4}$ ($\text{mmen-Mg}_2(\text{dobpdc})$). This compound displays an exceptional capacity for CO_2 adsorption at low pressures, taking up 2.0 mmol/g (8.1 wt %) at 0.39 mbar and 25 °C, conditions relevant to removal of CO_2 from air, and 3.14 mmol/g (12.1 wt %) at 0.15 bar and 40 °C, conditions relevant to CO_2 capture from flue gas. Dynamic gas adsorption/desorption cycling experiments demonstrate that $\text{mmen-Mg}_2(\text{dobpdc})$ can be regenerated upon repeated exposures to simulated air and flue gas mixtures, with cycling capacities of 1.05 mmol/g (4.4 wt %) after 1 h of exposure to flowing 390 ppm CO_2 in simulated air at 25 °C and 2.52 mmol/g (9.9 wt %) after 15 min of exposure to flowing 15% CO_2 in N_2 at 40 °C. The purity of the CO_2 removed from dry air and flue gas in these processes was estimated to be 96% and 98%, respectively. As a flue gas adsorbent, the regeneration energy was estimated through differential scanning calorimetry experiments to be 2.34 MJ/kg CO_2 adsorbed. Overall, the performance characteristics of $\text{mmen-Mg}_2(\text{dobpdc})$ indicate it to be an exceptional new adsorbent for CO_2 capture, comparing favorably with both amine-grafted silicas and aqueous amine solutions.



1. INTRODUCTION

The concentration of CO_2 in the Earth's atmosphere is presently 390 ppm,¹ an increase of approximately 110 ppm since the start of the Industrial Revolution.² The combustion of fossil fuels is largely responsible for this increase,³ yet fossil fuels will continue to be heavily utilized for energy production during the 21st century. Currently, there is significant interest in the development and implementation of technologies that slow CO_2 emissions and thus forestall the most severe consequences of global warming. For limiting future CO_2 emissions from large, stationary sources like coal-fired power plants, carbon capture and sequestration (CCS) has been proposed.⁴ The CCS process involves the selective removal of CO_2 from gas mixtures, the compression of pure CO_2 to a supercritical fluid, transportation to an injection site, and finally permanent subterranean or submarine storage.⁵ For the retrofit of existing power plants, post-combustion CO_2 capture is a likely configuration. In this design, fuel is burned in air and CO_2 is removed from the effluent. For coal-fired power plants, the largest flue gas components by volume are N_2 (70–75%), CO_2 (15–16%), H_2O (5–7%), and O_2 (3–4%), with total pressures near 1 bar and temperatures between 40 and 60 °C.⁶

Aqueous amine solutions are currently the most viable absorbents for carbon capture under the aforementioned conditions, and they are presently used for the removal of CO_2 from industrial commodities like natural gas.⁷ While a

variety of advanced amines are available, 30% monoethanolamine (MEA) in water is the benchmark solvent against which competing technologies are generally compared. The low solvent cost and proven effectiveness make MEA an attractive absorbent for many applications. However, if MEA were to be utilized for CCS, electricity prices are projected to increase by 86%.⁸ The U.S. Department of Energy has targeted a maximum 35% increase for the cost of electricity produced from a coal power plant that captures 90% of the CO_2 it generates. The diversion of steam from the electricity generation cycle to the solvent regeneration cycle sharply reduces the net electricity output of the plant, drastically increasing electricity costs. Previous work has demonstrated that plant efficiency is highly dependent on the solvent regeneration energy.⁹

Presently, there is significant interest in the development of solid adsorbents that selectively adsorb CO_2 at partial pressures applicable to CCS.¹⁰ Solid adsorbents are promising candidates because the significantly smaller heat capacities of solids may reduce the sensible heat required for regeneration. In addition, solvent loss and corrosion issues resulting from the use of aqueous amines would be minimized if solids adsorbents were instead utilized.⁹

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While CCS is perhaps unlikely to be widely implemented within the next decade,⁵ a number of current industrial processes utilize liquid or solid adsorbents to remove CO₂ from gas mixtures. These processes could benefit greatly from the next generation of adsorbents that are currently being proposed for CCS applications. Currently, aqueous amines are used industrially to separate CO₂ from gas mixtures with high CO₂ partial pressures like natural gas, while solid adsorbents are used to remove CO₂ from mixtures with very low CO₂ partial pressures.

Among the most challenging CO₂ separations is the removal of CO₂ directly from air. Prior to the cryogenic distillation of air for N₂, O₂, and Ar production, CO₂ is removed from the air to minimize solid CO₂ formation on heat exchangers.¹¹ To maximize the capacity of zeolite 13X for CO₂ adsorption, the air stream is dried over alumina, cooled to 5 °C, and pressurized to 5–7 bar. The effective capacity of zeolite 13X, the most widely used adsorbent for this process, under these conditions is ca. 0.35 mmol/g, which corresponds to a crystallographic volumetric capacity of approximately 0.5 mmol/cm³.^{11c,12} In this “pre-purification” step, the CO₂ concentration in the air is reduced to less than 1 ppm. New higher-capacity adsorbents could potentially eliminate the costs associated with pre-cooling feed air and reduce the adsorbent regeneration energy.

Carbon dioxide scrubbers are critical life support systems in confined spaces with limited air exchange, such as spacecraft, submarines, and breathing suits.¹³ Because of the very low capacity of solid adsorbents for 390 ppm CO₂ in unpressurized gas streams, weight and volume limitations prevent the implementation of systems capable of maintaining CO₂ concentrations at atmospheric levels. Thus, the average CO₂ concentration aboard the International Space Station ranges from 3000 to 7000 ppm,¹⁴ which approaches the currently established safe limit for chronic CO₂ exposure.^{14,15} Improved adsorbents could potentially reduce CO₂ concentrations within confined spaces to significantly lower and potentially safer levels, while simultaneously reducing the adsorbent mass and volume.

In addition to current processes that remove CO₂ from air, proposed technologies may benefit greatly from improved air capture adsorbents. For example, alkaline fuel cells (AFCs)¹⁶ and iron-air batteries¹⁷ require CO₂-free O₂ sources to avoid electrolyte side reactions. It has been previously suggested that improved CO₂ adsorbents could solve many issues associated with the operation of AFCs because air purification imposes considerable engineering and financial burdens on the system.¹⁸

Lastly, directly adsorbing CO₂ from the atmosphere combined with geologic sequestration has been proposed as a potential solution for offsetting CO₂ emissions from mobile or diffuse generators.¹⁹ Direct air capture, if widely implemented, could also theoretically reduce atmospheric CO₂ concentrations by capturing historic emissions rather than simply abating future emissions. Significant obstacles remain for direct air capture including its substantially higher cost compared to traditional CCS.²⁰ Yet many estimates rely on the use of traditional inorganic bases, which require the construction of very large physical structures to ensure sufficient surface area for air to contact the adsorbent.²¹ Porous solid adsorbents with high surface areas could potentially drastically reduce the large capital costs associated with air capture.

Recently, amine-functionalized porous solids have been proposed as superior adsorbents for air capture compared to

inorganic bases and zeolites. For example, amine-modified porous polymers have been tested for use aboard spacecraft.²² Silica,^{17,23} alumina,²⁴ and carbon²⁵ adsorbents functionalized with amines have also been analyzed for their efficacy as air capture adsorbents, as have ammonium ion-exchange membranes.²⁶ The primary advantage of amine-functionalized adsorbents is their high capacity for 390 ppm CO₂, in some cases in excess of 2 mmol/g. However, existing materials of this type frequently require hours to reach saturation because of slow adsorption kinetics.

Metal–organic frameworks are a class of porous, crystalline adsorbents that have recently attracted much attention for use in gas separations.^{10,27} The high tunability of their design may enable greater functionality with reduced adsorbent mass and volume compared to traditional solid adsorbents. Among the most interesting features of some metal–organic frameworks is the presence of coordinatively unsaturated metal centers (open metal sites) along the pore surfaces.^{28,29} These five-coordinate metal cations, known to behave as Lewis acids that strongly polarize gas adsorbents, are further amenable to post-synthetic functionalization.³⁰

In chemically robust metal–organic frameworks with well-separated open metal sites, one amine of a diamine molecule can bind to a metal cation as a Lewis base, while the second amine remains available as a chemically reactive adsorption site. The modification of open metal sites within the metal–organic framework Cu-BTTrI with the secondary amine *N,N'*-dimethylethylenediamine (mmen) was recently reported.³¹ Upon grafting mmen onto the exposed Cu²⁺ sites of the framework, a 3.5 times enhancement in CO₂ capacity at 0.15 bar and 25 °C was realized. We believe that the incorporation of very basic alkylamines into framework pores will be a promising strategy for increasing the capacity of metal–organic frameworks for CO₂ uptake at low partial pressures.

The high concentration of open metal sites within the M₂(dobdc) (M = Mg, Mn, Fe, Co, Ni, Zn; dobdc⁴⁻ = 2,5-dioxido-1,4-benzenedicarboxylate; M-MOF-74 or CPO-27-M) series of metal–organic frameworks makes them attractive candidates for diamine functionalization.²⁹ To date, however, we have been unable to synthesize promising amine functionalized derivatives of the M₂(dobdc) series. We hypothesized that the relatively narrow, one-dimensional channels (~11 Å diameter) may be hindering effective diffusion of the diamines into the framework. Thus, we sought to synthesize expanded analogues of the M₂(dobdc) structure via a ligand extension. Larger pores should enable more facile functionalization, enhance gas diffusion, and potentially unlock unrealized functionality within this interesting structural topology replete with open metal sites.

Herein, we report the first expanded analogues of the M₂(dobdc) structure type, featuring 18.4-Å-wide channels and exhibiting exceptional CO₂ adsorption properties upon functionalization with mmen.

2. EXPERIMENTAL SECTION

General. All reagents were obtained from commercial vendors at reagent grade purity or higher and used without further purification.

4,4'-Dihydroxy-(1,1'-biphenyl)-3,3'-dicarboxylic Acid (*H₄dobpdc*). The compounds 4,4'-dihydroxybiphenyl (1.16 g, 6.24 mmol), KHCO₃ (2.00 g, 20.0 mmol), dry ice (4 g), and 1,2,4-trichlorobenzene (3 mL) were added to a PTFE insert within a steel acid digestion bomb (23 mL) and heated at 255 °C for 17 h.³² After cooling to room temperature, the mixture was collected via vacuum filtration and washed with diethyl ether. The solid was suspended in 300 mL of

distilled water and filtered again. To the filtrate, neat HCl was slowly added until a pH between 1 and 2 was reached. The resulting crude product was collected via filtration. Recrystallization using 50 mL of acetone and 50 mL of water per gram of crude material afforded 0.68 g (40%) of pure product as a white powder. Anal. Calcd for $C_{14}H_{10}O_6$: C, 61.32; H, 3.68. Found: C, 61.34; H, 3.60. IR (KBr) 1659 (vs), 1612 (s), 1481 (s), 1450 (vs), 1319 (s), 1290 (s), 1240 (s), 1099 (w), 1047 (w), 894 (w), 825 (m), 792 (m), 723 (w), 686 (m), 573 (w), 530 (w) cm^{-1} . 1H NMR (300 MHz, $dmso-d_6$): δ = 10.75–11.95 (br, 2H), 7.93 (s, 2H), 7.74 (d, 2H, J = 11.2 Hz), 7.00 (d, 2H, J = 8.6 Hz).

$Zn_2(dobpdc)(DEF)_2 \cdot DEF \cdot H_2O$ (DEF-1). To a 2-mL Pyrex tube, $H_4dobpdc$ (4.0 mg, 0.015 mmol), $ZnBr_2 \cdot 2H_2O$ (8.9 mg, 0.034 mmol), and 0.5 mL of mixed solvent (1:1 DEF:EtOH; DEF = N,N -diethylformamide) were added. The tube was sealed and placed in a pre-heated oven at 100 °C. After 72 h, needle-shaped, colorless crystals had formed. The crystals were isolated by filtration and washed with hot DEF to afford 3.7 mg (35%) of product. Anal. Calcd for $C_{29}H_{41}N_3O_{10}Zn_2$: C, 47.84; H, 5.64; N, 5.79. Found: C, 48.21; H, 5.72; N, 5.82. IR (KBr): 1655 (vs), 1610 (s), 1544 (s), 1462 (s), 1412 (vs), 1286 (s), 1234 (s), 1149 (m), 1103 (m), 1047 (w), 881 (m), 825 (m), 760 (w), 690 (m), 586 (m), 505 (w) cm^{-1} .

$Mg_2(dobpdc)(DEF)_2 \cdot DEF_{1.5} \cdot H_2O$ (DEF-2). Into a 10-mL Pyrex cell, $H_4dobpdc$ (24 mg, 0.088 mmol), $MgBr_2 \cdot 6H_2O$ (60 mg, 0.21 mmol), and 3 mL of solvent (1:1 DEF:EtOH) were loaded and sealed with a PTFE cap. The mixture was irradiated in a microwave reactor (CEM Discover) for 30 min at 120 °C. After 30 min, the solution was cooled, and the resulting solid was collected via filtration and washed with hot DEF. The solid was dried under vacuum to yield 57.5 mg (95%) of product as a white powder. Anal. Calcd for $C_{31.5}H_{46.5}Mg_2N_{3.5}O_{10.5}$: C, 54.77; H, 6.78; N, 7.10. Found: C, 54.85; H, 7.07; N, 6.86. IR (KBr): 1661 (vs), 1612 (s), 1570 (s), 1468 (vs), 1419 (s), 1298 (s), 1242 (s), 1149 (m), 1111 (m), 1047 (w), 945 (w), 885 (m), 842 (m), 825 (m), 725 (w), 692 (m), 660 (w), 590 (m), 501 (w), 447 (w) cm^{-1} . Heating at 420 °C for 65 min *in vacuo* yielded the fully activated adsorbent $Mg_2(dobpdc)$ (2).

$Mg_2(dobpdc)(mmen)_{1.6}(H_2O)_{0.4}$ (*mmen-Mg₂(dobpdc)*) or *mmen-2*. A sample of fully activated 2 (77 mg, 0.24 mmol) was immersed in anhydrous hexane, and 20 equiv of N,N' -dimethylethylenediamine (*mmen*, 0.53 mL, 4.8 mmol) was added. The suspension was stirred for one day, filtered, and rinsed copiously with hexanes. The solid was then evacuated of residual solvents at 100 °C for 24 h to afford 87 mg (77%) of product as a gray-white powder. Anal. Calcd for $C_{20.4}H_{26}Mg_2N_{3.2}O_{9.6}$: C, 52.46; H, 5.62; N, 9.60. Found: C, 52.15; H, 5.41; N, 9.52. IR (ATR, neat): 3320 (w), 2952 (w), 2910 (w), 2862 (w), 2806 (w), 1616 (s), 1575 (s), 1538 (w), 1468 (vs), 1421 (vs), 1295 (m), 1244 (s), 1152 (m), 1104 (m), 1053 (m), 1000 (w), 887 (m), 844 (m), 828 (m), 727 (w), 692 (s), 618 (m), 589 (s) cm^{-1} .

X-ray Structure Determination. A crystal of DEF-1 was mounted on a cryoloop under a cooling stream of dinitrogen. Diffraction data were collected with synchrotron radiation using a 6B MX-I ADSC Quantum-210 detector with a silicon (111) double-crystal monochromator at the Pohang Accelerator Laboratory. The ADSC Quantum-210 ADX program (Ver. 1.92) was used for data collection and HKL2000 (Ver. 0.98.699) was used for cell refinement, data reduction, and absorption corrections. The structure was solved by direct methods and refined by full-matrix least-squares analysis using anisotropic thermal parameters for non-hydrogen atoms with the SHELXTL program.³³ The C2 and C5 atoms were isotropically refined due to poor thermal behavior. Guest molecules in the pores were highly disordered and unable to be modeled. To account for this electron density, the program SQUEEZE³⁴ was employed. All hydrogen atoms were calculated at idealized positions and refined using a riding model. Crystal data for DEF-1: empirical formula = $C_{12}H_{14}NO_4Zn$, M_r = 301.61, T = 100(2) K, space group = $P3_221$, a = 21.698(3) Å, c = 6.8690(14) Å, α = 90°, β = 90°, γ = 120°, V = 2800.7(8) Å³, Z = 6, D_{calc} = 1.073 g/cm³, μ = 1.319 mm⁻¹, 13927 reflections collected, 2941 unique (R_{int} = 0.0418), $R1$ = 0.0466, $wR2$ = 0.1204 ($I > 2\sigma(I)$).

Powder X-ray Diffraction. Powder X-ray diffraction data were collected with either a Rigaku Ultima III or a Bruker D8 Advance

diffractometer using Cu $K\alpha$ radiation (λ = 1.5406 Å). The unit cell dimensions of DEF-2 and *mmen-2* were determined by performing a full-pattern decomposition using the Le Bail method, as implemented in TOPAS-Academic.³⁵ Owing to the isomorphism with $Zn_2(dobpdc)$, the trigonal space group $P3_221$ was used for the refinements. Crystal data for DEF-2: a = 21.761(2) Å, c = 6.9721(7) Å, V = 2859.1(5) Å³ (R_{wp} = 0.093, R_p = 0.067). Crystal data for *mmen-2*: a = 21.500(2) Å, c = 6.8275(9) Å, V = 2733.2(6) Å³ (R_{wp} = 0.042, R_p = 0.033).

Gas Adsorption Measurements. Gas adsorption data for pressures in the range 0–1.1 bar were obtained by volumetric methods using a Micromeritics ASAP2020 instrument. All gases were 99.998% purity or higher. Isotherms at 77 K were measured in liquid nitrogen baths. Isotherms at 25, 35, 45, 50, and 75 °C were measured using liquid circulators to maintain a constant temperature. Isotherms at 100 and 120 °C were measured using a heated sand bath controlled by a programmable temperature controller. BET surface areas were calculated from N_2 adsorption at 77 K. DFT pore size distributions and pore sizes were calculated from N_2 adsorption at 77 K with the Micromeritics DFT Plus Models Kit (Ver. 2.02) software suite with cylinder pore geometries for an oxide surface. The compound *mmen-2* was regenerated at 100 °C under dynamic vacuum for 4 h after measurement of each isotherm.

Isosteric Heats of Adsorption Calculations. A dual-site Langmuir–Freundlich equation (eq 1) was employed to model the CO_2 adsorption at 25, 50, and 75 °C for *mmen-2* in the region before the step in the isotherms.

$$q = \frac{q_{sat,A} b_A p^{\alpha_A}}{1 + b_A p^{\alpha_A}} + \frac{q_{sat,B} b_B p^{\alpha_B}}{1 + b_B p^{\alpha_B}} \quad (1)$$

Here, q is the amount of CO_2 adsorbed (mmol/g), p is the pressure (bar), q_{sat} is the saturation capacity (mmol/g), b is the Langmuir–Freundlich parameter ($bar^{-\alpha}$), and α is the Langmuir–Freundlich exponent (dimensionless) for two adsorption sites A and B. In order to model the CO_2 adsorption in the region after the step, a modified Langmuir–Freundlich equation (eq 2) was employed.

$$q = \frac{q_{sat,A} b_A (p - p^*)^{\alpha_A}}{1 + b_A (p - p^*)^{\alpha_A}} + \frac{q_{sat,B} b_B (p - p^*)^{\alpha_B}}{1 + b_B (p - p^*)^{\alpha_B}} + \frac{q_{sat,C} b_C (p - p^*)^{\alpha_C}}{1 + b_C (p - p^*)^{\alpha_C}} \quad (2)$$

Here, adsorption is considered at three sites, A, B, and C, and the extra parameter p^* is used to account for the pressure at which the step in the isotherm occurs and the strongest adsorption sites are first populated. After carefully refining the parameters in eqs 1 and 2, excellent agreement was achieved between the experimental isotherm data and the corresponding isotherm fits (see Figures S10–S13). Using the appropriate isotherm fits, Mathematica software was used to solve for the exact pressures, p , corresponding to constant amounts of CO_2 adsorbed, q , at 25, 50, and 75 °C. The Clausius–Clapeyron equation (eq 3) was then used to calculate the isosteric heats of adsorption, Q_{st} , by determining the slope of the best-fit line for $\ln p$ versus $1/T$ at each loading.

$$(\ln p)_q = \left(\frac{Q_{st}}{R} \right) \left(\frac{1}{T} \right) + C \quad (3)$$

As indicated by the residual sum of square values, R^2 , of close to 1 (see Figure S14), the isotherm data were consistent with the Clausius–Clapeyron equation across the entire loading range considered, even with the changes in the location of the step in the isotherms.

The isosteric heats of adsorption for CO_2 in the unmodified $Mg_2(dobpdc)$ were determined by fitting the adsorption isotherms at 25, 35, and 45 °C with a dual-site Langmuir–Freundlich equation (eq 1). Each temperature was fit independently, and the Clausius–Clapeyron equation was used to determine Q_{st} as a function of loading.

All fit parameters for $Mg_2(dobpdc)$ and *mmen-Mg₂(dobpdc)* are specified in Tables S2–S4.

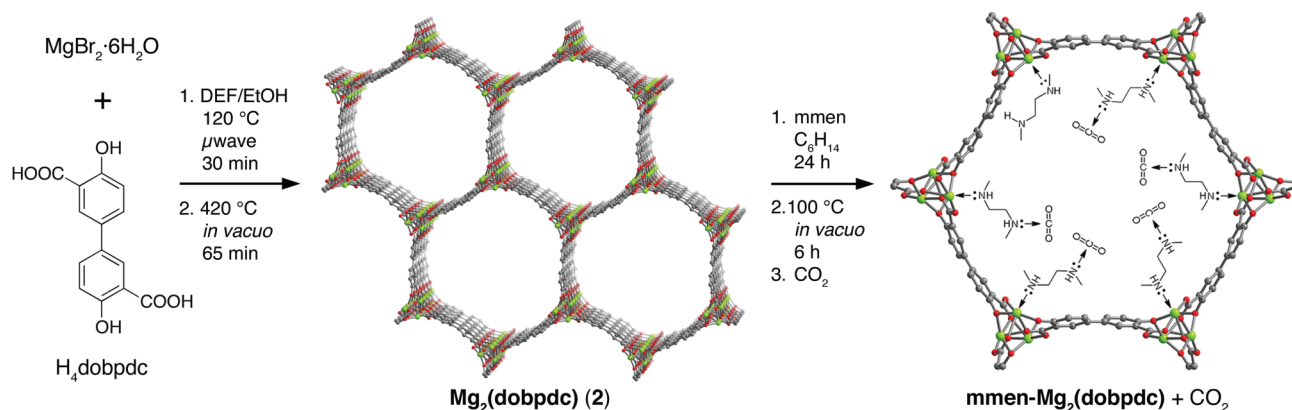


Figure 1. Synthesis of $\text{mmen-Mg}_2(\text{dobpdc})$ ($\text{mmen} = N,N'$ -dimethylethylenediamine; $\text{dobpdc}^{4-} = 4,4'$ -dioxido-3,3'-biphenyldicarboxylate). From the microwave reaction of $\text{MgBr}_2 \cdot 6\text{H}_2\text{O}$ and H_4dobpdc (left), $\text{Mg}_2(\text{dobpdc})$ (**2**) is obtained following evacuation of the as-synthesized solid at high temperatures (middle). The framework structure depicted is that obtained from single crystal X-ray analysis of the isostructural zinc compound DEF-1. Green, red, and gray spheres represent Mg, O, and C atoms respectively; H atoms are omitted for clarity. Addition of an excess of mmen to the evacuated framework yields the amine-appended CO_2 adsorbent $\text{Mg}_2(\text{dobpdc})(\text{mmen})_{1.6}(\text{H}_2\text{O})_{0.4}$ (right).

Model CO_2 Isotherms. Isotherms for CO_2 uptake in mmen-2 at other temperatures were predicted from the experimental isotherm at 75°C via application of a Clausius–Clapeyron relation (eq 4).

$$\ln p_2 = \frac{Q_{\text{st}} + \left(\ln p_1 \left(\frac{RT_1 T_2}{T_2 - T_1} \right) \right)}{\left(\frac{RT_1 T_2}{T_2 - T_1} \right)} \quad (4)$$

Calculated pressures (p_2) at constant loadings (q) were determined assuming an isosteric heat of adsorption (Q_{st}) of 71 kJ/mol for all loadings. Here, p_1 is a pressure experimentally measured at 75°C , R is the universal gas constant, T_1 is 75°C , and T_2 is the temperature for which a model isotherm is desired.

CO_2 Selectivity Calculations. Here, the adsorption capacities of component n (q_n) are defined to be molar excess adsorption capacities determined experimentally without correction for absolute adsorption, and p_n is defined to be the pressure of component n as experimentally measured. Selectivity (S) is defined according to eq 5, while purity is defined according to eq 6.

$$S = \frac{q_{\text{CO}_2}/q_2}{p_{\text{CO}_2}/p_2} \quad (5)$$

$$\text{purity} = \frac{q_{\text{CO}_2}}{q_{\text{CO}_2} + q_2} \times 100\% \quad (6)$$

Thermogravimetric Analysis and Gas Cycling Measurements. Thermogravimetric analyses (TGA) were carried out at ramp rates between 5 and $10^\circ\text{C}/\text{min}$ under a nitrogen flow with a TA Instruments TGA Q5000 (Ver. 3.1 Build 246) or a Scinco TGA N-1000.

Carbon dioxide cycling experiments were performed on the aforementioned TA Instruments analyzer using $15\% \text{CO}_2$ in N_2 (Praxair NI-CD15C-K), 390 ppm CO_2 in air (Praxair AI-CD-390C-K; 390 ppm CO_2 , $21\% \text{O}_2$, balance N_2), CO_2 (Praxair 99.998%), and N_2 (Praxair, 99.9%). A flow rate of $25 \text{ mL}/\text{min}$ was employed for all gases. Prior to cycling, the sample was activated by heating at 150°C for 1 h . For Figure 6, sample mass was normalized to be 0% at the adsorption temperature (25°C for 390 ppm CO_2 ; 40°C for $15\% \text{CO}_2$) under flowing N_2 . For Figure 7, sample mass was normalized to be 0% at 150°C under flowing $100\% \text{CO}_2$. Masses were uncorrected for buoyancy effects. In Figure 6, the difference between the quantity of N_2 adsorbed at the tare temperature and the regeneration temperature likely accounts for the negative apparent mass of mmen-2 at high temperatures.

Differential Scanning Calorimetry. Thermal analysis was performed on a TA Instruments Q200 differential scanning calorimeter equipped with a refrigerated cooling system (RCS40). Through the sample cell, $15\% \text{CO}_2$ in N_2 or N_2 was flowed over T-Zero aluminum pans that were not hermetically sealed. An empty, aluminum T-Zero pan provided the reference sample for thermal analysis. The calorimeter was calibrated with the TA Instruments software package; the melting point of indium (156.60°C) was utilized for the temperature calibration. Sample and pan masses were determined after activation at 150°C for 60 min on a TA Instruments TGA Q5000 via the readout from the internal balance under flowing N_2 . Identical aliquots of mmen-2 were utilized for TGA and DSC measurements to approximate heats of adsorption. Integrated heats were calculated with TA Instruments Universal Analysis software suite.

Other Physical Measurements. Elemental analyses for C, H, and N were performed at the Elemental Analysis Service Center of Sogang University or the Microanalytical Laboratory of the University of California, Berkeley. ^1H spectra were obtained using a 300 MHz Varian instrument. Infrared spectra were obtained from KBr pellets with a Bomen MB-104 spectrometer or on a Perkin-Elmer Spectrum 400 FTIR spectrometer equipped with an attenuated total reflectance (ATR) accessory. For diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) spectra, the Perkin-Elmer spectrometer was equipped a Harrick Praying Mantis Diffuse Reflectance accessory and a temperature-controlled high-pressure gas cell with Swagelok valves connecting $5\% \text{CO}_2$ in He (Praxair certified standard HE CD5C-K) and an oil-free vacuum.

3. RESULTS AND DISCUSSION

Synthesis, Structure, and Activation. Reaction of H_4dobpdc with $\text{ZnBr}_2 \cdot 2\text{H}_2\text{O}$ or $\text{MgBr}_2 \cdot 6\text{H}_2\text{O}$ in $1:1$ DEF:EtOH afforded DEF-1 and DEF-2, respectively (see Figure 1). The coordination environment of the divalent metal cations within **1** and **2** are analogous to those in the $\text{M}_2(\text{dobc})$ series.²⁹ In the crystal structure of DEF-1, four different dobpdc^{4-} ligands and one DEF molecule are bonded to each Zn^{2+} ion in a distorted octahedral geometry. There are three unique O donor types from the dobpdc^{4-} ligand: bridging (μ_2) aryloxy O atoms (O1), bridging (μ_2) carboxylate O atoms (O2), and nonbridging carboxylate O atoms (O3). The equatorial plane of each Zn^{2+} is composed of two *trans*-disposed O1 ligands from different linkers, one O3 donor atom, and one O2 donor atom. An O2 donor atom occupies one axial coordination site, while the other axial site is occupied by an O donor atom from DEF, the reaction solvent. This coordination

mode results in the formation of helical chains of Zn^{2+} atoms running along the c axis of the crystal. The resulting framework consists of a honeycomb lattice of hexagonal, one-dimensional channels approximately 18.4 Å in width. Bound DEF molecules occupy the Zn^{2+} coordination sites along the corners of hexagonal channel walls. As shown in Figure S1, powder X-ray diffraction (PXRD) data indicate DEF-2 to be isostructural with DEF-1.

Heating DEF-2 at 420 °C for 65 min under dynamic vacuum, removed the DEF molecules bound to the metal atoms, completely activating the material and generating open Mg^{2+} coordination sites. Such extreme thermal treatment was necessary because soaking in methanol at 100 °C for 20 h did not lead to exchange of the bound DEF molecules. The porosity of activated **2** was confirmed via N_2 adsorption at 77 K (see Figure S4), resulting in a BET surface area of 3270 m^2/g . Note that, in line with the expanded structure, this is significantly greater than the BET surface area of 1495 m^2/g reported for $\text{Mg}_2(\text{dobdc})$.³⁶

The synthesis and structure of mmen-**2** is depicted schematically at the right of Figure 1. An activated sample of **2** was suspended in hexanes and an excess of mmen was added. As shown by powder X-ray diffraction (see Figure S3), framework crystallinity was not significantly affected by activation or subsequent amine functionalization. A much reduced BET surface area of 70 m^2/g was calculated for mmen-**2**, while DFT pore size distributions indicated a reduction in average pore size (see Figures S5–S7).

Upon exposure of DEF-2 to atmospheric air, the white powder turns blue and a loss of crystallinity occurs, as observed via powder X-ray diffraction measurements. Amine functionalization, however, appears to enhance framework stability, because no similar degradation was observed for mmen-**2** upon exposure to air for one week.

CO₂ Adsorption Isotherms. The CO_2 adsorption data obtained for **2** are depicted in Figure S8. Isothermic heats of adsorption were calculated to approach -44 kJ/mol at low coverage, as shown in Figure S9. This value is similar to those previously reported for the analogous $\text{Mg}_2(\text{dobdc})$ framework.³⁷ The adsorption capacity of **2** at 25 °C is 4.85 mmol/g (17.6 wt %) and 6.42 mmol/g (22.0 wt %) at 0.15 and 1 bar, respectively. The gravimetric capacity of **2** for CO_2 at 0.15 bar exceeds the capacity of all metal–organic frameworks except for $\text{Mg}_2(\text{dobdc})$,^{10c} which adsorbs 6.1 mmol/g (21.2 wt %) at 25 °C and 0.15 bar.³⁶

The alkylamine-functionalized metal–organic framework mmen-**2** displayed an extremely high affinity for CO_2 at extraordinarily low pressures. The CO_2 adsorption isotherms obtained at 25, 50, and 75 °C are presented in Figure 2. At 25 °C and 0.39 mbar, near the current partial pressure of CO_2 in Earth's atmosphere, the compound adsorbed 2.0 mmol/g (8.1 wt %), which is 15 times the capacity of **2**. At the much higher pressure of 5 mbar, the median partial pressure of CO_2 within the International Space Station, the framework adsorbed 2.6 mmol/g (10.3 wt %). For comparison, zeolite 5A, which is currently used aboard the station to adsorb CO_2 , adsorbs 0.85 mmol/g (3.6 wt %, crystallographic volumetric capacity 1.3 mmol/cm^3) at 5 mbar.^{12,38}

At 25 °C, the CO_2 adsorption in mmen-**2** reaches 3.13 mmol/g (12.1 wt %) at 0.15 bar and 3.86 mmol/g (14.5 wt %) at 1 bar. Remarkably, its CO_2 uptake at 1 bar and 25 °C exceeds the amount of N_2 adsorbed at 77 K. Thus, the low surface area measured at 77 K does not appear to accurately reflect the

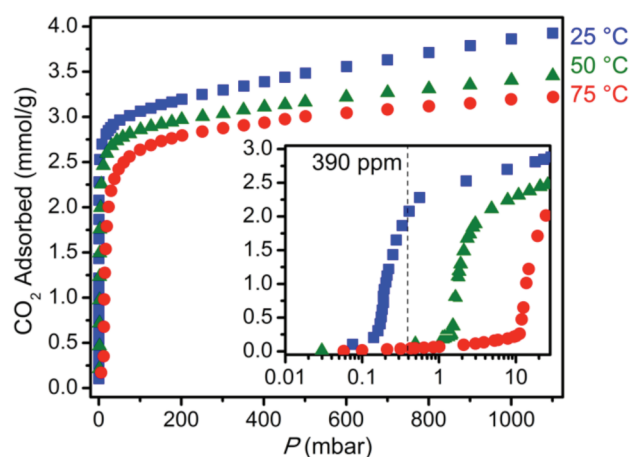


Figure 2. Adsorption of CO_2 in mmen-**2** at 25 °C (blue squares), 50 °C (green triangles), and 75 °C (red circles). Inset: The isotherms at very low pressures exhibit a step that shifts to higher pressures at higher temperatures. The dashed, vertical line marks the current partial pressure of CO_2 in air (390 ppm).

surface area accessible to CO_2 , and the pore size distribution calculated from N_2 adsorption at cryogenic temperatures likely does not accurately represent the true pore size distribution within mmen-**2**. On a per mass basis, the amine-functionalized framework adsorbed less CO_2 than **2** at pressures higher than 20 mbar. The large density difference between the two frameworks is primarily responsible for the lower gravimetric capacity of mmen-**2**. Crystallographic densities of 0.58 and 0.86 g/cm^3 were calculated for **2** and mmen-**2**, respectively. At 0.15 bar, **2** and mmen-**2** adsorb 2.8 and 2.7 mmol/cm^3 , respectively.

Based upon the calculated number of dangling amine groups in mmen-**2**, a loading of 3.4 mmol/g would correspond to one CO_2 per amine, yet uptake of only ca. 2.2 mmol/g was observed. Here, pore blockages, hydrogen bonded amines, or cooperative binding mechanisms between two amines and one CO_2 may be limiting the accessible stoichiometry of mmen-**2**. Thus, significant additional capacity improvements might be realized in the material if conditions can be identified for appending one mmen per magnesium and binding one CO_2 molecule per dangling amine.

Isothermic heat of adsorption calculations were hindered by the presence of a prominent step in the isotherms at low pressures and convex to the pressure axis. Generally, continuous mathematical functions are used to model experimental isotherms, which then become the input parameters for the Clausius–Clapeyron relation. Since we were unable to mathematically model the CO_2 isotherms of mmen-**2** with continuous equations over the entire pressure range, each isotherm was modeled with two Langmuir–Freundlich equations. Data sets corresponding to the adsorption regions before and after the steps were compiled and then modeled individually.

As shown in Figure 3, isosteric heats of adsorption for mmen-**2** were calculated from the 25, 50, and 75 °C isotherm models. At low loadings, heats significantly lower than those expected for chemical adsorption of CO_2 onto an amine were calculated. However, calculated heats quickly approached and maintained a value of -71 kJ/mol, which likely corresponds to the chemical adsorption of CO_2 onto the free amine of mmen. Here, a carbamate with a weak C–N bond is probably formed

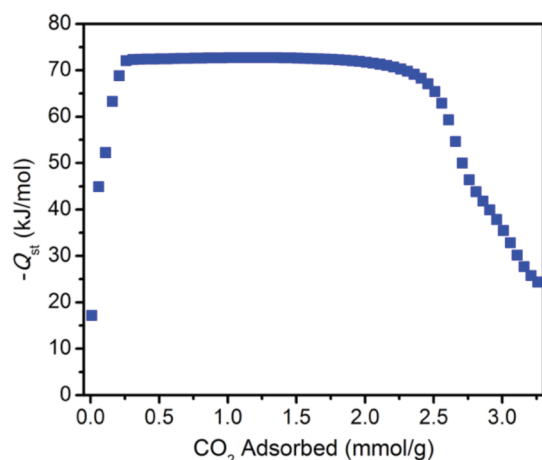


Figure 3. Isosteric heats of CO₂ adsorption onto mmen-2, as calculated using the Clausius–Clapeyron relation. The calculated values indicate that chemical adsorption of CO₂ onto the amines did not occur at very low coverage.

through interaction the lone pair of the free amine of mmen and the electrophilic carbon of CO₂.

In situ diffuse reflectance infrared Fourier transform spectroscopy (DRIFTS) was employed to probe the chemical nature of adsorbed CO₂. At 25 °C, 1 bar of 5% CO₂ in He was introduced into an airtight gas cell containing a sample of activated mmen-2. The difference spectrum between mmen-2 under a 1 bar atmosphere and the activated framework under vacuum is shown in Figure 4. The prominent loss peak at 3316

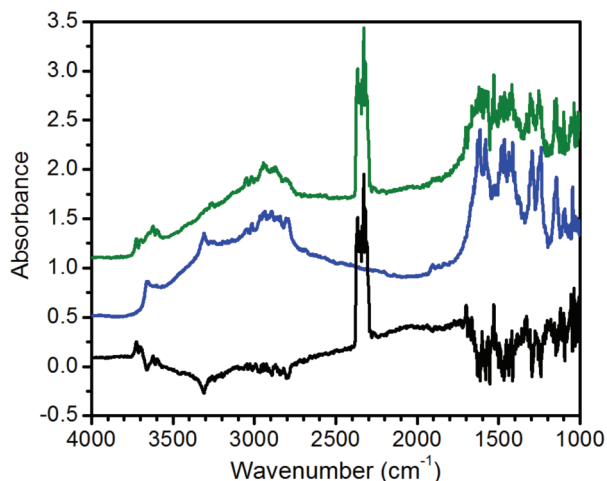


Figure 4. DRIFTS spectra of CO₂ adsorption in mmen-2. The loss band at 3316 cm⁻¹ in the difference spectrum (black line) between evacuated mmen-2 (blue line) and mmen-2 under a 5% CO₂ in He atmosphere (green line) indicates CO₂ adsorption occurs via a chemical adsorption mechanism. The blue line is offset by -0.5 absorbance units for visual clarity.

cm⁻¹, assigned to the NH stretch of free mmen-2, is indicative of chemical adsorption of CO₂ onto amines.

Recent work on alkylamine-grafted silica surfaces have suggested that chemical adsorption of CO₂ onto alkylamines is not possible without neighboring amines or surface hydroxyl groups to stabilize the resulting carbamates; ammonium carbamates or surface-bonded carbamates are formed, respectively.³⁹ The formation of surface-bonded carbamates in

mmen-2 is unlikely due to a lack of surface hydroxyl groups, and the broad NH stretches expected for ammonium cations are not definitively resolvable from the DRIFTS difference spectrum. Furthermore, the slow reversibility of mmen-2 at room temperature (see Figure S15) appears to preclude the formation of ammonium carbamates, which have been reported to desorb CO₂ from primary amines readily at room temperature.³⁹ Additional experiments are necessary to ascertain whether adjacent functional groups are necessary for CO₂ adsorption within such alkylamine-functionalized metal–organic frameworks.

The step in each isotherm marks the pressure at which CO₂ adsorption becomes dominated by chemisorption. Interestingly, the step moves to significantly higher pressures as the adsorption temperature increases. As shown in Figure S16, the location of the step at 75 °C was reproduced four times with little variation. Thus, the step cannot be attributed to slow kinetics, but appears to be an inherent characteristic of mmen-2 CO₂ adsorption isotherms. All batches of mmen-2 have exhibited steps at approximately the same pressures, even when functionalized with different amounts of amines. Figure S17 presents the 75 °C CO₂ adsorption isotherm of Mg₂(dobpdc)(mmen)_{1.75}(H₂O)_{0.25}.

The location of the step is modeled well by a simple Clausius–Clapeyron relation (see eq 4), which predicts how isotherms move as a function of temperature. In Figure 5,

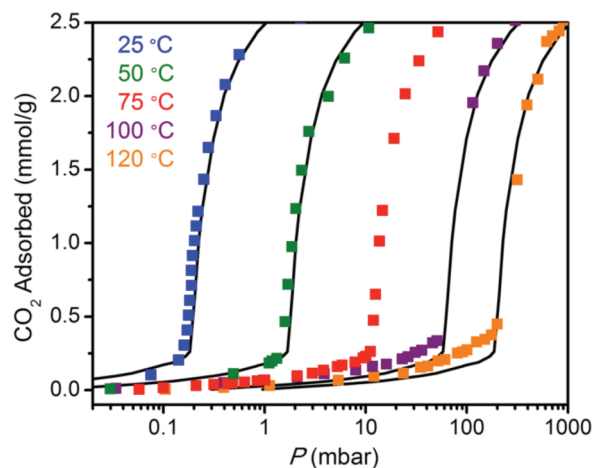


Figure 5. Adsorption of CO₂ in mmen-2 at 25, 50, 75, 100, and 120 °C, represented as blue, green, red, purple, and orange squares, respectively. The locations of the step for the 25, 50, 100, and 120 °C isotherms were modeled with a Clausius–Clapeyron derived equation constructed from the 75 °C data (shown as black lines for each temperature). The heat of adsorption was assumed to be a constant -71 kJ/mol. The onset pressure of each isotherm step was predicted well by the simple model.

experimental isotherms at 25, 50, 75, 100, and 120 °C are shown as blue, green, red, purple, and orange squares, respectively. The black lines through the 25, 50, 100, and 120 °C data are predicted isotherms for those temperatures modeled from the experimental data collected at 75 °C by assuming that the heat of adsorption is -71 kJ/mol for all CO₂ loadings. The black lines are not continuous functions, but connect the calculated points listed in Table S5. Utilizing the approximate heat of adsorption calculated from the 25, 50, and 75 °C isotherms, the model was able to predict the general shapes of the 100 and 120 °C isotherms, as well as the

Table 1. Approximate CO₂ Selectivities of mmen-2 for Selected Gas Mixtures at 25 °C

mixture	comp. 2	pressure (mbar)		uptake (mmol/g)		molar selectivity	est. purity (%)
		CO ₂	comp. 2	CO ₂	comp. 2		
air	N ₂	0.4	800	2.05	0.083	49 000	96
air	O ₂	0.4	200	2.05	0.038	27 000	98
flue gas	N ₂	150	750	3.13	0.079	200	98

pressures at which the steps arise. Thus, it seems that thermodynamics alone account for the movement of the step to higher pressures at higher temperatures.

The existence of the step, however, is unexpected in a strongly adsorbing material with large pores. The amines, as the strongest adsorption sites, should preferentially adsorb CO₂ before other, weaker adsorption sites adsorb significant quantities of CO₂. To explain the nonclassical adsorption behavior of mmen-2, we presently hypothesize that adsorption of CO₂ onto mmen is disfavored at low adsorptive concentrations (the density of gas phase CO₂ in the pores) because of the large positive entropy associated with reorganization of the amines, as required to form a chemical bond with CO₂. Before the chemical potential necessary for amine reorganization afforded to the system by the adsorptive is achieved, CO₂ adsorption is dominated by non-amine or weak amine-CO₂ adsorption sites. The minimum chemical potential required for amine reorganization increases as thermal motion increases, in line with the shift of the step to higher pressures at higher temperatures. Additional experiments and modeling will be required to test this hypothesis and to understand the relationship between pore filling and adsorption in mmen-2. However, the shift of the step to higher pressures at higher temperatures could afford the opportunity for unique regeneration conditions, whereby a weak vacuum could nearly fully regenerate the material at moderate temperatures.

CO₂ Selectivity. Adsorption isotherms for N₂ and O₂ in mmen-2 at 25 °C are shown in Figure S18. The uptake capacities for these gases relative to CO₂ suggest that it would be a highly selective adsorbent. Since we were unable to mathematically model the CO₂ isotherms of mmen-2 with a meaningful, single equation⁴⁰ over the entire pressure range of interest, ideal adsorbed solution theory (IAST)⁴¹ selectivities could not be calculated. Table 1 therefore instead presents the approximate molar selectivities of mmen-2 for relevant gas mixtures, as calculated according to eq 5 above using the excess adsorption capacities. This method for determining selectivities was recently shown to be less accurate than IAST, in part because it generally underestimates the values.³⁷

The selectivity of mmen-2 for CO₂ over N₂ in air is thus estimated to be at least 49 000. However, the purity of gas adsorbed is perhaps a more physically meaningful value than selectivity, and is an important criterion for evaluating adsorbents if captured CO₂ is to be transported or sequestered. The purity of CO₂ adsorbed onto mmen-2, calculated according to eq 6, would likely be more than 96% or 98% for capture from air or flue gas, respectively.

While the purity of gas adsorbed within individual mmen-2 crystallites is very high, as shown in Table 1, the purity of gas adsorbed within a bed of adsorbent will be affected by the composition of gas phase molecules that fill the pore and intercrystalline spaces. For carbon capture applications, the high concentration of N₂ in these pore volumes will negatively affect the selectivity and purity values of real adsorbent beds. However, the approximate selectivity and purity values

calculated in Table S6 indicate that while the purity of CO₂ decreases within increasing void space, any adsorbent bed with less than 80% void space should be capable of achieving CO₂ purity in excess of 90%. From these calculations, we believe mmen-2 to be a promising, high-capacity adsorbent for removing high-purity CO₂ from dry gas mixtures containing N₂ and O₂.

CO₂ Capture from Air. To evaluate the performance of mmen-2 as a regenerable adsorbent, thermogravimetric analysis (TGA) was utilized to monitor sample mass under dynamic environments. The top panel of Figure 6 plots changes in

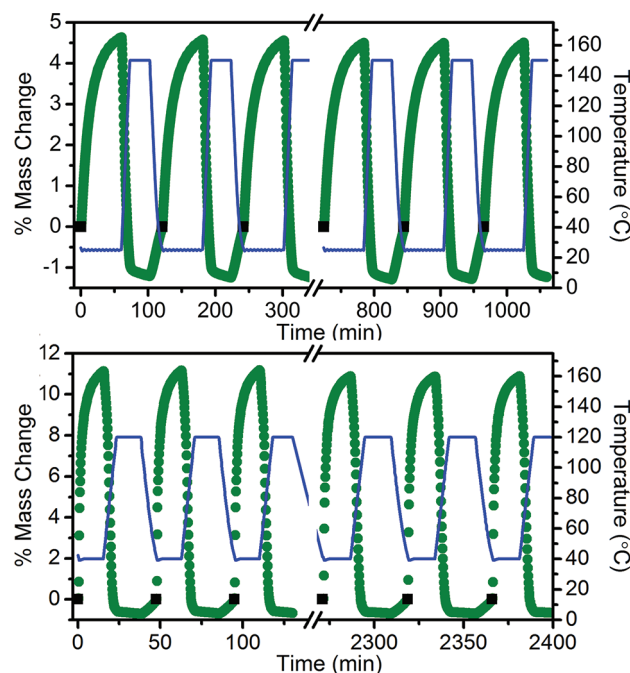


Figure 6. Adsorption–desorption cycling for mmen-2, demonstrating reversible uptake of 1.05 mmol CO₂/g from dry air after 1 h (upper) and 2.52 mmol CO₂/g from dry flue gas after 15 min (lower). Changes in sample mass are plotted as green circles and sample temperatures are plotted as blue lines; CO₂ was introduced into the sample chamber at the points marked by black squares. Samples were regenerated under flowing N₂.

sample mass (normalized to the mass of the framework under N₂ at 25 °C) while simulated air containing 390 ppm CO₂ was flowed over the sample. Despite the very low concentration of CO₂, a 4.6%_{mc} (%_{mc} = percent mass change;⁴² 1.05 mmol/g; 4.4 wt %) was realized after 60 min. The adsorbent was then regenerated under flowing N₂ at 150 °C for 30 min and the cycle repeated 10 times with no apparent loss of capacity.

To the best of our knowledge, no studies evaluating the efficacy of air capture within metal–organic frameworks have yet been reported. The equilibrium capacity (2.0 mmol/g, 1.72 mmol/cm³) of mmen-2 is similar to the capacities of the very best amine-grafted silica and alumina adsorbents reported to

date. However, the kinetics of adsorption appear to be significantly faster in mmen-2 than for amines deposited via evaporation or polymerization methodologies. For example, while the pseudo-equilibrium capacity of an outstanding poly(ethyleneimine) impregnated silica gel was reported to be ca. 2.4 mmol/g, it took nearly 200 min for the silica based adsorbent to realize 4.6%_{mc}, the capacity of mmen-2 for CO₂ after only 60 min.^{23d} The easily accessed amines within mmen-2 appear to enhance adsorption rates greatly, enabling rapid adsorption–desorption cycles to be utilized.

CO₂ Capture from Dry Flue Gas. The capabilities of mmen-2 as an adsorbent for removing CO₂ from the flue gas of coal-fired power stations were also investigated. The bottom panel of Figure 6 presents the dynamic cycling behavior of mmen-2 under the relevant, dry conditions: 15% CO₂ in N₂ flowing over the solid at 40 °C. After adsorbing CO₂ for 15 min, the sample was heated at 120 °C for 15 min under N₂. A capacity of 11.1%_{mc} (2.52 mmol/g, 9.9 wt %) relative to the sample mass of mmen-2 under N₂ at 40 °C was realized. After 50 cycles, only a 0.2%_{mc} capacity loss was observed. Longer adsorption and desorption times did not significantly improve the cycling capacity of the material, nor did higher desorption temperatures. Note that the apparent capacity of mmen-2 greatly exceeds the ca. 2 wt % working capacity⁴³ of aqueous monoethanolamine (MEA) scrubbers, which would likely swing between the same adsorption and desorption temperatures.

If captured CO₂ is to be sequestered, high-purity CO₂ is essential. To desorb the ca. 98% pure CO₂ adsorbed onto mmen-2, a N₂ or air purge cannot be utilized to strip the adsorbent bed. Hence, to approximate the working capacity of mmen-2 using a temperature swing without a N₂ purge, a pure CO₂ purge was utilized instead. A 7.8%_{mc} (1.8 mmol/g, 7.2 wt %) was realized when 15% CO₂ in N₂ at 40 °C was desorbed with 100% CO₂ at 150 °C. In Figure 7, 15% CO₂ was

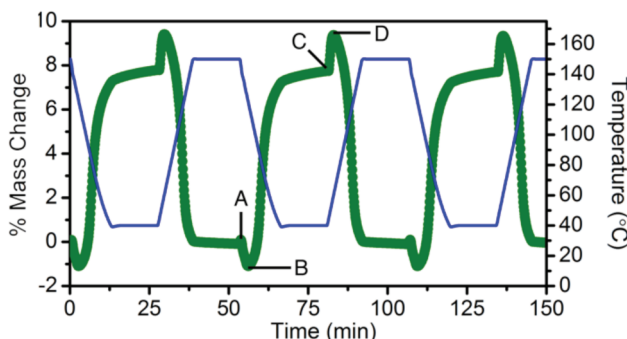


Figure 7. A temperature swing processes, as simulated for CO₂ in mmen-2 using TGA and desorption with a CO₂ purge. A 1.8 mmol/g capacity difference was calculated from the quantity of CO₂ adsorbed at 150 °C under a 100% CO₂ atmosphere (point A) and the quantity of CO₂ adsorbed at 40 °C under flowing 15% CO₂ in N₂ (point C). Changes in sample mass are plotted as green circles and sample temperatures are plotted as blue lines; points B and D arise from introducing 15% CO₂ at 150 °C and 100% CO₂ at 40 °C respectively.

introduced over the sample at 150 °C (point A). The change in atmosphere from 100% to 15% CO₂ partially activated the framework, resulting in a negative effective mass (point B). As the sample cooled to 40 °C under the 15% atmosphere, CO₂ was adsorbed. After 15 min at 40 °C (point C), 100% CO₂ was introduced and the sample was heated at 10 °C/min. The increase in CO₂ concentration at low temperatures temporarily

increased the amount of CO₂ adsorbed (point D). The working capacity of mmen-2 is the difference between the amounts adsorbed at points A and C. The mass of the material at 150 °C under 100% CO₂ was normalized to be 0% mass in Figure 7. The relationship between points A and C and isotherms is illustrated in Figure S19.

Differential Scanning Calorimetry. To approximate the adsorbent regeneration energy, heats of adsorption and desorption were quantified for mmen-2 via differential scanning calorimetry (see Figure 8).⁴⁴ When 15% CO₂ in N₂ at 40 °C

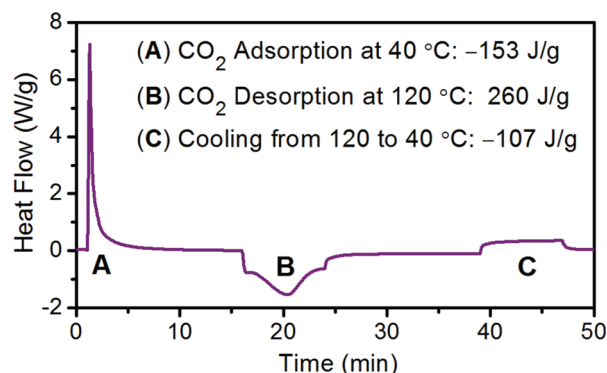


Figure 8. Heat flows from mmen-2, as determined via differential scanning calorimetry. First, 15% CO₂ in N₂ was adsorbed at 40 °C (event A). After 15 min, mmen-2 was heated to 120 °C under N₂ (event B). Lastly, the regenerated material was cooled under N₂ to 40 °C (event C). The integrated heats calculated for each event are given in the legend.

was introduced over a sample of mmen-2 activated at 150 °C under N₂, an exothermic heat flow of −153 J/g was observed (event A). Note that, to enable rapid gas exchange, the pan was not hermetically sealed and the true value of heats evolved are likely slightly larger than those registered by the instrument due to heat loss to the environment. When this heat flow was correlated to the 2.5 mmol/g capacity calculated via TGA adsorption under similar conditions, an approximate, average heat of adsorption of −61 kJ/mol CO₂ was calculated.

Following a 15 min adsorption period, 100% N₂ was introduced into the cell and the sample was heated to 120 °C (event B); 260 J/g were required to heat the sample and maintain the regeneration temperature for 15 min. Lastly, −107 J/g were evolved as the material was cooled under a N₂ atmosphere (event C) from 120 to 40 °C. Based upon this laboratory-scale experiment, approximately 2.34 MJ of energy would be required to regenerate 1 kg of CO₂ adsorbed onto mmen-2. This value compares very favorably with the 3.6–4.5 MJ of energy necessary to release 1 kg of CO₂ from an MEA scrubber.^{5,43}

However, the aforementioned regeneration energies were measured under a N₂ purge, a laboratory-scale analogue for a temperature-vacuum swing regeneration cycle. While a direct measurement of regeneration energy with a 100% CO₂ purge was not possible, calculations shown in the SI indicate that the regeneration energy of mmen-2 under CO₂ will be higher primarily due to the reduced working capacity of the adsorbent. The regeneration energy of mmen-2 under CO₂ is still lower than the regeneration energy of the most highly optimized MEA solutions, and it is likely that optimization of adsorption and desorption temperatures will further lower the heat of regeneration of mmen-2.

Approximate heat capacities (C_p) were calculated for mmen-2. Over the 80 °C temperature range, the average heat required to regenerate CO₂-loaded mmen-2 was 3.3 J/g·K. The regenerated framework released an average of 1.3 J/g·K upon cooling, a reasonable approximation of the heat capacity of mmen-2. For comparison, the heat capacity of a 30 wt % MEA solution without CO₂ is ~3.8 J/g·K over the same 80 °C range.⁴³ The lower regeneration energy of mmen-2 is thus attributable to its lower heat capacity. In addition, the significantly greater working capacity of mmen-2 means that a smaller mass is required to adsorb the same amount of CO₂. The heat required to desorb CO₂ from the amines is similar for mmen-2 and MEA solutions.⁴³

4. CONCLUSIONS AND OUTLOOK

With the new metal–organic frameworks M₂(dobpdc) (M = Zn (1), Mg (2)), an expanded variant of the M₂(dobdc) (MOF-74) structure type has been achieved, leading to enlarged, 18.4-Å-wide channels lined with open metal coordination sites. Functionalization of 2 with mmen, afforded a remarkable new CO₂ adsorbent, mmen-Mg₂(dobpdc). The large capacity, high selectivity, and fast kinetics of this material for adsorbing CO₂ from dry gas mixtures with N₂ and O₂ make it an attractive new adsorbent for applications in which zeolites and inorganic bases are currently used, including the removal of CO₂ from air.

Future work will focus on evaluating the efficacy of alkylamine-appended metal–organic frameworks such as mmen-2 under humid conditions. Whether the presence of water in gas mixtures will significantly affect the stability, capacity, selectivity, or regeneration energy of mmen-2 is presently unknown. In addition, the peculiar CO₂ isotherm shapes warrant further investigation, since a detailed understanding of the adsorption behavior within mmen-2 may enable the synthesis of even better materials for CO₂ capture. Finally, efforts are also underway to improve the separation performance of the material through use of various other polyamines in place of mmen.

■ ASSOCIATED CONTENT

Supporting Information

Powder X-ray diffraction patterns and fits, additional CO₂, N₂, and O₂ isotherms and fits, FTIR spectra, TGA traces, and additional purity and regeneration energy calculations. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

The authors declare no competing financial interest.

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$$\text{wt\%} = \frac{\text{mass of adsorbate}}{\text{mass of adsorbate} + \text{mass of adsorbent}} \times 100\% \quad (8)$$

The mass change registered by the TGA does not account for adsorbate mass in the denominator and thus is not strictly wt %. To clearly differentiate this difference, the abbreviation %_{mc} (percent mass change) is used instead to describe the experimentally observed mass changes. As an additional note, because TGA only records net mass change, the mass of N₂ in the activated framework likely displaced by CO₂ is not registered as part of the %_{mc}.

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