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Amides Do Not Always Work: Observation of Guest Binding in an Amide-Functionalised Porous Metal-Organic Framework

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ABSTRACT: An amide-functionalised metal organic framework (MOF) material, MFM-136, shows a high CO₂ uptake of 12.6 mmol g⁻¹ at 20 bar and 298 K. MFM-136 is the first example of acylamide pyrimidyl isophthalate MOF without open metal sites, and thus provides a unique platform to study guest binding, particularly the role of free amides. Neutron diffraction reveals that, surprisingly, there is no direct binding between the adsorbed CO₂/CH₄ molecules and the pendant amide group in the pore. This observation has been confirmed unambiguously by inelastic neutron spectroscopy. This suggests that introduction of functional groups solely may not necessarily induce specific guest-host binding in porous materials, but it is a combination of pore size, geometry, and functional group that leads to enhanced gas adsorption properties.

Recent developments in materials chemistry and crystal engineering have shown that metal-organic frameworks (MOFs) have promising properties that complement or compete favourably with zeolites and activated carbons in various applications.¹ MOFs are crystalline porous coordination polymers consisting of polyatomic organic ligands linked to metal ions/clusters by covalent bonds.² MOFs have shown great promise for gas adsorption and storage owing to their high porosity and internal surface area, and tuneable functionality on the pore surface for selective gas binding. Generation of open metal sites³ and incorporation of pendant functional groups⁴ at the pore surface are two dominant methods of functionalising MOF cavities. For example, MOFs with open Cu(II) sites can show strong adsorption affinity to molecular H₂.⁵ Recently, the detailed binding mechanisms to saturated and unsaturated light hydrocarbons has been rationalised in a hydroxyl-functionalised MOF.⁶ Within the field of carbon capture, materials functionalised with amines (-NH₂), imines (-NH), and amides (-CONH) dominate, largely because of their potential to form specific interactions with CO₂, leading to highly selective CO₂ uptakes. Although high CO₂ adsorption has been observed in a number of amine-, imine-, and amide-

functionalised MOFs,⁷ molecular insight into the direct binding between adsorbed CO₂ molecules and porous host (especially towards these functional groups) is largely lacking. Recently, direct H₂N(δ⁻)... (δ⁺)CO₂ binding has been observed in a Zn(II) MOF incorporating amine groups that protrude into the pore, providing structural insight into the observed high CO₂ adsorption in this material.⁴

The incorporation of pendent amide (-CONH-) and/or amine groups into MOFs is thus regarded generally as a promising approach to enhance CO₂ uptake due to the formation of hydrogen bonds with amides serving as both hydrogen bond acceptors (*via* C=O) and donors (*via* N-H). A series of amide-functionalised MOFs have been synthesised and shown to exhibit high CO₂ uptakes and selectivities.^{7b,8} Likewise, computational studies attribute this to the specific binding and formation of hydrogen bonds between adsorbed CO₂ molecules and free amide or amine groups thus enhancing adsorption affinity and selectivity for CO₂.^{7c,8c,9} However, to date there are few physical investigations on the precise role of amides in CO₂ binding in MOFs. The challenge of such investigations is further increased in MOF systems containing open metal sites owing to the inevitable competition for guest binding between the open metal sites and the organic functional group(s) in the pore. Here, we report the synthesis, structure and gas adsorption properties of an amide-functionalised pyrimidyl Cu(II)-carboxylate MOF, MFM-136, which shows a high CO₂ adsorption capacity (12.6 mmol g⁻¹ at 20 bar and 298 K). In MFM-136, all Cu(II) sites are fully coordinated to carboxylate and pyrimidyl groups, affording a pore environment without open metal sites. This gives an ideal environment for studying the binding interaction between amides and adsorbed CO₂ molecules since it eliminates the competitive binding of CO₂ on the open Cu(II) sites. In this situation, we can clearly evaluate the precise role of free amides in guest binding in the pore. Combined neutron diffraction and inelastic neutron spectroscopy have revealed the preferred binding sites for CO₂ in the pore and the corresponding host-guest binding dynamics. Surprisingly, there is no direct binding between adsorbed CO₂/CD₄ and free

amides in this case, representing the first example of systematic experimental investigation of guest binding in amide-functionalised MOFs.

Solvothermal reaction of 5-[4-(pyrimidin-5-yl)benzamido]isophthalic acid (H_2L , Figure 1a) with $Cu(NO_3)_2 \cdot 3H_2O$ in DMF at $80^\circ C$ for 16 h yields MFM-136 as green single crystals. MFM-136 crystallises in space group $R\bar{3}2$ and shows a 3D binodal (3, 6)-connected network with a rare eea topology.^{8f,10} In MFM-136, the binuclear $\{Cu_2(RCO_2)_4\}$ paddlewheels coordinate to two pyrimidyl nitrogen atoms from two different ligands at both axial positions, resulting in the absence of open Cu(II) sites in the entire structure (Figure 1). The metal-ligand connectivity in MFM-136 affords two types of cages (**A** and **B**). Cage **A** is surrounded by twelve $\{Cu_2\}$ paddlewheel units and six linkers and has a prolate-ellipsoid shape (length 24.9 Å, width 10.6 Å). Cage **B** is enclosed by six $\{Cu_2\}$ paddlewheel units and six linkers and has a more spherical shape (length 16.2 Å, width 12.5 Å). The overall structure is an alternate packing of these two types of cages to give a highly porous and robust framework material with a void fraction of 54% and BET surface area of $1634 \text{ m}^2 \text{ g}^{-1}$ (Figure S6).

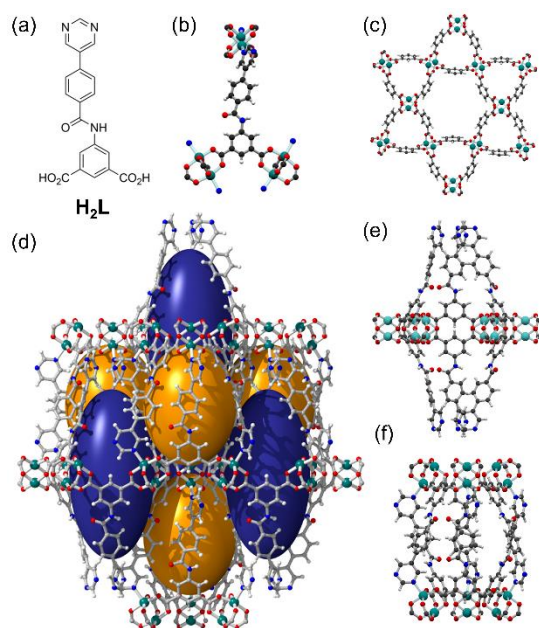


Figure 1. (a) Chemical structure of ligand H_2L . (b) Coordination environment of L^{2-} in MFM-136. (c) View along the c -axis of the *Kagome* lattice in MFM-136. (d) View of the alternate packing of large cages **A** (blue) and small cages **B** (orange) in MFM-136. Views of (e) the large cage **A** and (f) small cage **B** both along the b -axis. Colours: C: black; H: white; O: red; N: blue; Cu: teal.

At 273 K, the CO_2 sorption isotherms of desolvated MFM-136 show an uptake of 7.3 mmol g^{-1} at 1 bar and 14.3 mmol g^{-1} at 20 bar, representing the highest CO_2 uptake in mono-amide-functionalised MOFs reported to date (Table S2). Methane adsorption in MFM-136 gives a lower uptake of 2.9 mmol g^{-1} at 1 bar and 8.3 mmol g^{-1} at 20 bar at 273 K. The experimental CO_2 adsorption isotherms show good agreement with grand canonical Monte Carlo (GCMC) simulations (Figure 2). In contrast, MFM-136 shows negligible N_2 uptake under the same conditions, leading to selectivities for CO_2/N_2 and CO_2/CH_4 of

27:1 and 6.3:1, respectively, at 273 K. The isosteric heats of adsorption for CO_2 and CH_4 in MFM-136 are calculated using the Virial method as 25.6 and 16.0 kJ mol^{-1} , respectively, at low surface coverage. The selective CO_2 uptake in MFM-136 is lower than the leading ultra-microporous MOFs;¹¹ however the high capacity indicates MFM-136 remains a promising candidate in the separation of CO_2 over CH_4 and N_2 .¹² The lack of open Cu(II) sites in the pores of MFM-136 prevents strong binding to water molecules, which often triggers framework collapse or hydrolysis in MOFs containing open metal sites.¹³ Previously reported MOFs containing amides in the absence of open metals sites have exhibited high CO_2 capacities;¹⁴ however the role of the amides in CO_2 binding was not defined structurally.

It is reported that the excellent uptake of CO_2 in amide-functionalised MOFs is a consequence of specific CO_2 -amide interactions based upon hydrogen bond formation between the amide $-NH(\delta^+)$ and the $O(\delta^-)$ of CO_2 .^{7c,8a} To gain experimental insight, preferred binding sites in MFM-136 have been determined by *in situ* neutron powder diffraction (NPD) as a function of gas loading (CO_2 and CD_4). NPD patterns were recorded at 7 K for the desolvated material and at loadings of 1.8 and 2.3 CO_2/Cu and 1.1 CD_4/Cu . Fourier difference map analysis of the NPD patterns revealed positions of the adsorbed CO_2 and CD_4 molecules, which were further developed by Rietveld refinement. All binding sites were checked carefully for their unambiguous presence in the final structural model; *i.e.*, a parallel refinement without each of the binding sites was carried out to confirm the presence of each site by comparing the R factors and the residual peaks.

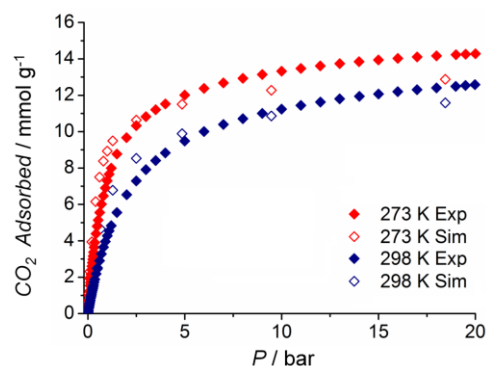


Figure 2. Experimental and simulated adsorption isotherms for CO_2 in desolvated MFM-136 at 273 and 298 K.

The NPD data at a loading of 1.8 CO_2/Cu reveals eight binding sites **A-H** distributed between cages **A** and **B** (Figure 3). The CO_2 molecules are constrained to be linear with equal C-O bond lengths, whilst their crystallographic occupancies and positions (including orientations) have been refined. At low CO_2 loading, adsorbate-adsorbate interactions will be negligible meaning that the site occupancies directly reflect the binding strength between CO_2 and the framework. Three CO_2 sites **A-C** have significantly higher occupancies (0.65, 0.44 and 0.40, respectively) than the remaining five sites **D-H** (ranging within 0.26 - 0.11); surprisingly none of the sites makes an apparent hydrogen bonding interaction with the amide moiety. Site **A** resides on a three-fold symmetry axis in the centre of a triangular pocket formed by a $[(Cu_2)_3(isophalate)_3]$ unit,

where the CO₂ makes three identical long contacts to the phenyl rings [O1A_{CO₂}...ring centroid = 4.21(2) Å]. Site **B** lies close to a pyrimidine ring (O1B_{CO₂}...ring centroid = 3.22(9) Å) and makes a shorter contact with the aromatic C-H group on a phenyl ring (O1B_{CO₂}...C32 = 3.27(7) Å) along with a longer distance to an amide N-H (O1B_{CO₂}...N29 = 3.96(8) Å). Site **C** lies close to an isophthalate phenyl ring (O3C_{CO₂}...ring centroid = 2.99(6) Å). The shortest contact between a guest CO₂ and amide nitrogen atom is observed for the low occupancy site **F** (occupancy = 0.15) where the guest accepts a weak hydrogen bond (N29...O3F_{CO₂} = 3.22(20) Å; <N-H...O = 122(5)°). The absence of CO₂ molecules adjacent to the uncoordinated pyrimidine nitrogen atom is not surprising as the site is sterically hindered by a neighbouring phenyl ring; however, no such impediment exists around the amide N-H site which is fully accessible to the guest molecules. The preference of CO₂ to make multiple long range contacts with phenyl and pyrimidyl rings rather than accepting hydrogen bonds from the amide is striking and contrary to the assumptions which have previously informed the design philosophy of MOF materials for CO₂ capture. Upon increasing the guest loading to 2.3 CO₂/Cu(II), the occupancy of site **A** nears saturation (0.96) and remains distinctly higher than that of remaining sites **B-H** (range 0.50 – 0.18). In the structure of MFM-136 loaded with 1.1 CD₄/Cu(II), an equivalent site to site **A** in centre of the triangular pocket (C1A_{CD₄}...ring centroid = 4.33(2) Å) is observed to have the highest occupancy of 0.36. Additional CD₄ binding sites with lower occupancies were observed without notable interaction to the MOF host (Figure S19). Up to now, crystallographic characterisations of adsorbed gas molecules in MOFs have been mostly limited to one or two binding sites for materials with narrow pores.³⁻⁶ Simultaneous refinement of a large number of sites as reported in this work is made possible by the neutron diffraction data which gives equal prominence to the light guests (particularly for CD₄) and heavy framework.

The absence of adsorbed CO₂ molecules at the pendant amide group could be due to the transition between “dynamic” and “kinetic” products in which the adsorbed CO₂ has great mobility to translate/diffuse along the pore and form interactions with amide groups in a “come and go” fashion. The static crystallographic experiment can only paint a picture averaged over an extended time scale. Hence, only more stable environments of CO₂ can be seen from the diffraction study. Thus to gain a direct insight into the binding dynamics of adsorbed CO₂ molecules and the free amide groups, inelastic neutron spectroscopy (INS) was measured for MFM-136 as a function of CO₂ loading (Figure 4). INS spectra for the bare MOF show multiple features which have been identified *via* DFT calculations (Figure S20). Specifically, the peak at 69 meV corresponds to the out-of-plane wagging modes of the N-H group, and peaks around 110-160 meV originate from the motion of aromatic C-H groups and deformational modes of the phenyl rings. Comparison of the INS spectra for bare and CO₂-loaded MFM-136 shows very small changes to the overall vibrational peaks except for a guest-host stiffening effect as evidenced by a global shift of peaks to slightly higher energy. Indeed, the N-H motion (69 meV) has no detectable changes upon CO₂ loading, while the aromatic C-H groups show small changes as confirmed by the difference spectra (Figure 4b), including a small increase in intensity at 116 meV (assigned as out-of-plane C-H bending on the isophthalate ring) and a decrease at 136 meV (assigned as in-plane C-H bending on all phenyl rings).

This result is in excellent agreement with the NPD study and reaffirms the conclusion made from the diffraction experiment that direct CO₂ binding to the amide groups in the pore is absent.

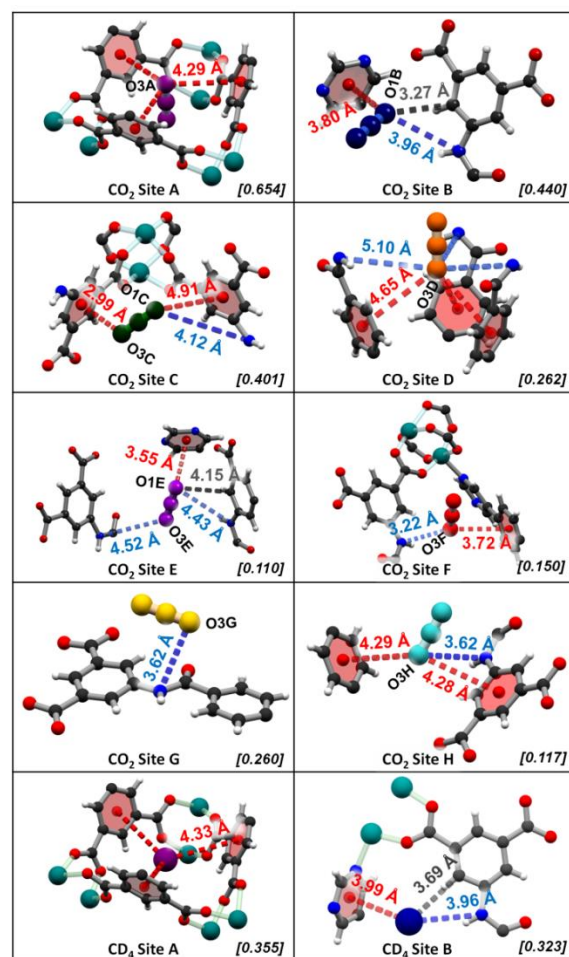


Figure 3. Binding sites of guests in MFM-136 at loadings of 1.8 CO₂/Cu(II) and 1.1 CD₄/Cu(II) elucidated from Rietveld refinement of NPD data. Colours: carbon, black; hydrogen, white; oxygen, red; nitrogen, blue; copper, teal; CO₂/CD₄ guests, purple/dark blue/green for sites **A/B/C**, respectively. Refined chemical occupancies of guest molecules inset.

Analysis of the CO₂-MOF interaction energy landscape determined during the GCMC simulation of the isotherm shows that the strongest predicted guest adsorption locations are in agreement with site **A**, followed by sites around the periphery of cage **A** corresponding to sites **B-E** (Figure S21). As in the NPD and INS studies, no strong adsorption was observed in the regions surrounding the N-H group.

In summary, a (3,6)-connected pyrimidyl carboxylate acylamide decorated MOF with a rare *eea*-topology has been synthesised. The amide-functionalised MOF exhibits high CO₂ uptake capacities and selectivity over CH₄ and N₂. Although

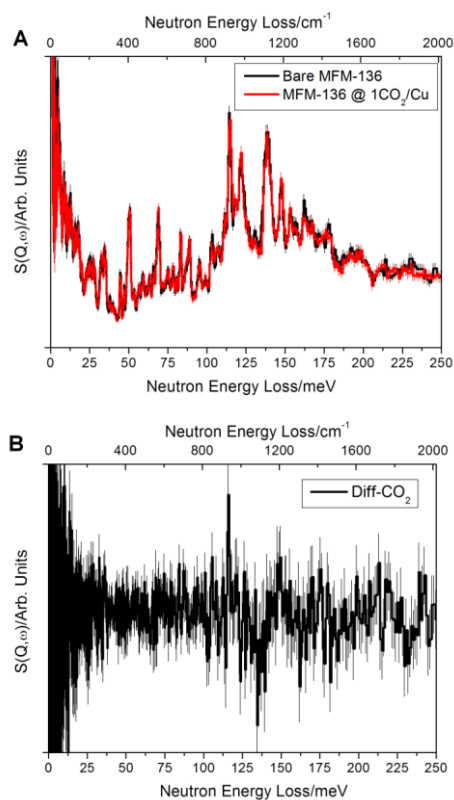


Figure 4. (a) Overlay of the INS spectra for bare and CO₂-loaded MFM-136; (b) difference INS spectrum for the bare and CO₂-loaded MFM-136.

it was anticipated that the amide moieties would actively participate in gas adsorption, the NPD and INS data reveals otherwise. The strongest binding site for both adsorbed CO₂ and CD₄ molecules are at the phenyl-isophthalate rings, and there is an absence of direct binding between adsorbed gas molecules and the pendent amide group in the pore. This has been confirmed by INS which shows retention of vibrational motion of the amide group upon CO₂ binding. This study indicates that introduction of functional groups in MOF structures may not necessarily form strong binding sites for gas molecules. Future investigation of the impact of a combination of functional groups and pore geometry is currently underway.

ASSOCIATED CONTENT

Supporting Information

Synthesis procedures, characterization, and additional analysis of crystal structures. [CCDC-1452775](#), [1481608-1481610](#) and [1504702](#) contain the supplementary crystallographic data for this paper. The Supporting Information is available free of charge on the ACS Publications website.

AUTHOR INFORMATION

Notes

The authors declare no competing financial interests.

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REFERENCES

- (1) Allendorf, M. D.; Stavila, V. *CrystEngComm* **2015**, *17*, 229.
- (2) Guillerm, V.; Kim, D.; Eubank, J. F.; Luebke, R.; Liu, X.; Adil, K.; Lah, M. S.; Eddaoudi, M. *Chem. Soc. Rev.* **2014**, *43*, 6141.
- (3) (a) Bloch, E. D.; Queen, W. L.; Krishna, R.; Zadrozny, J. M.; Brown, C. M.; Long, J. R. *Science* **2012**, *335*, 1606. (b) Xiang, S.; Zhou, W.; Zhang, Z.; Green, M. A.; Liu, Y.; Chen, B. *Angew. Chem. Int. Ed.* **2010**, *49*, 4615. (c) Queen, W. L.; Hudson, M. R.; Bloch, E. D.; Mason, J. A.; Gonzalez, M. I.; Lee, J. S.; Gygi, D.; Howe, J. D.; Lee, K.; Darwish, T. A.; James, M.; Peterson, V. K.; Teat, S. J.; Smit, B.; Neaton, J. B.; Long, J. R.; Brown, C. M. *Chem. Sci.* **2014**, *5*, 4569.
- (4) Vaidhyanathan, R.; Iremonger, S. S.; Shimizu, G. K. H.; Boyd, P. G.; Alavi, S.; Woo, T. K. *Science* **2010**, *330*, 650.
- (5) (a) Lin X.; Telepeni I.; Blake A. J.; Dailly A.; Brown C. M.; Simmons.; J. M.; Zoppi M.; Walker G. S.; Thomas K. M.; Mays T. J.; Hubberstey P.; Champness N. R.; Schröder M. *J. Am. Chem. Soc.* **2009**, *131*, 2159. (b) Yan Y.; Yang S.; Blake A. J.; Schröder M.; *Acc. Chem. Res.* **2014**, *47*, 296.
- (6) Yang, S.; Ramirez-Cuesta, A. J.; Newby, R.; Garcia-Sakai, V.; Manuel, P.; Callear, S. K.; Campbell, S. I.; Tang, C. C.; Schröder, M. *Nat. Chem.* **2015**, *7*, 121.
- (7) (a) Li, B.; Zhang, Z.; Li, Y.; Yao, K.; Zhu, Y.; Deng, Z.; Yang, F.; Zhou, X.; Li, G.; Wu, H.; Nijem, N.; Chabal, Y. J.; Lai, Z.; Han, Y.; Shi, Z.; Feng, S.; Li, J. *Angew. Chem. Int. Ed.* **2012**, *51*, 1412. (b) Du, L.; Yang, S.; Xu, L.; Min, H.; Zheng, B. *CrystEngComm* **2014**, *16*, 5520. (c) Lee, C. H.; Huang, H. Y.; Liu, Y. H.; Luo, T. T.; Lee, G. H.; Peng, S. M.; Jiang, J. C.; Chao, I.; Lu, K. L. *Inorg. Chem.* **2013**, *52*, 3962. (d) Vaidhyanathan, R.; Iremonger, S. S.; Shimizu, G. K. H.; Boyd, P. G.; Alavi, S.; Woo, T. K. *Angew. Chem. Int. Ed.* **2012**, *51*, 1826.
- (8) (a) Alsmail, N. H.; Suyetin, M.; Yan, Y.; Cabot, R.; Krap, C. P.; Lu, J.; Easun, T. L.; Bichoutskaia, E.; Lewis, W.; Blake, A. J.; Schröder, M. *Chem. Eur. J.* **2014**, *20*, 7317. (b) Zheng, B.; Bai, J.; Duan, J.; Wojtas, L.; Zaworotko, M. J. *J. Am. Chem. Soc.* **2011**, *133*, 748. (c) Zheng, B.; Yang, Z.; Bai, J.; Li, Y.; Li, S. *Chem Commun.* **2012**, *48*, 7025. (d) Xiong, S.; He, Y.; Krishna, R.; Chen, B.; Wang, Z. *Cryst. Growth Des.* **2013**, *13*, 2670. (e) Chen, Y.-Q.; Qu, Y.-K.; Li, G.-R.; Zhuang, Z.-Z.; Chang, Z.; Hu, T.-L.; Xu, J.; Bu, X.-H. *Inorg. Chem.* **2014**, *53*, 8842. (f) Chen, Z.; Adil, K.; Weselinski, L. J.; Belmabkhout, Y.; Eddaoudi, M. *J. Mater. Chem. A* **2015**, *3*, 6276. (g) Lu, Z.; Bai, J.; Hang, C.; Meng, F.; Liu, W.; Pan, Y.; You, X. *Chem. Eur. J.* **2016**, *22*, 6277.
- (9) Duan, J.; Yang, Z.; Bai, J.; Zheng, B.; Li, Y.; Li, S. *Chem Commun.* **2012**, *48*, 3058.
- (10) Wei, Y.-S.; Lin, R.-B.; Wang, P.; Liao, P.-Q.; He, C.-T.; Xue, W.; Hou, L.; Zhang, W.-X.; Zhang, J.-P.; Chen, X.-M. *CrystEngComm* **2014**, *16*, 6325.
- (11) (a) Nandi, S.; De Luna, P.; Daff, T. D.; Rother, J.; Liu, M.; Buchanan, W.; Hawari, A. I.; Woo, T. K.; Vaidhyanathan, R. *Sci. Adv.* **2015**, *1*, e1500421. (b) Nugent, P.; Belmabkhout, Y.; Burd, S. D.; Cairns, A. J.; Luebke, R.; Forrest, K.; Pham, T.; Ma, S.; Space, B.; Wojtas, L.; Eddaoudi, M.; Zaworotko, M. J. *Nature* **2013**, *495*, 80.
- (12) Wang, J.; Huang, L.; Yang, R.; Zhang, Z.; Wu, J.; Gao, Y.; Wang, Q.; O'Hare, D.; Zhong, Z. *Energy Environ. Sci.* **2014**, *7*, 3478.
- (13) Singh, M. P.; Dhumal, N. R.; Kim, H. J.; Kiefer, J.; Anderson, J. A. *J. Phys. Chem. C* **2016**, *120*, 17323.
- (14) (a) Keceli, E.; Hemgesberg, M.; Grunker, R.; Bon, V.; Wilhelm, C.; Philippi, T.; Schoch, R.; Sun, Y.; Bauer, M.; Ernst, S.; Kaskel, S.; Thiel, W. R. *Microporous Mesoporous Mater.* **2014**, *194*, 115. (b) Debatin, F.; Thomas, A.; Kelling, A.; Hedin, N.; Bacsik, Z.; Senkovska, I.; Kaskel, S.; Junginger, M.; Mueller, H.; Schilde, U.; Jaeger, C.; Friedrich, A.; Holdt, H.-J. *Angew. Chem., Int. Ed.* **2010**, *49*, 1258.

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