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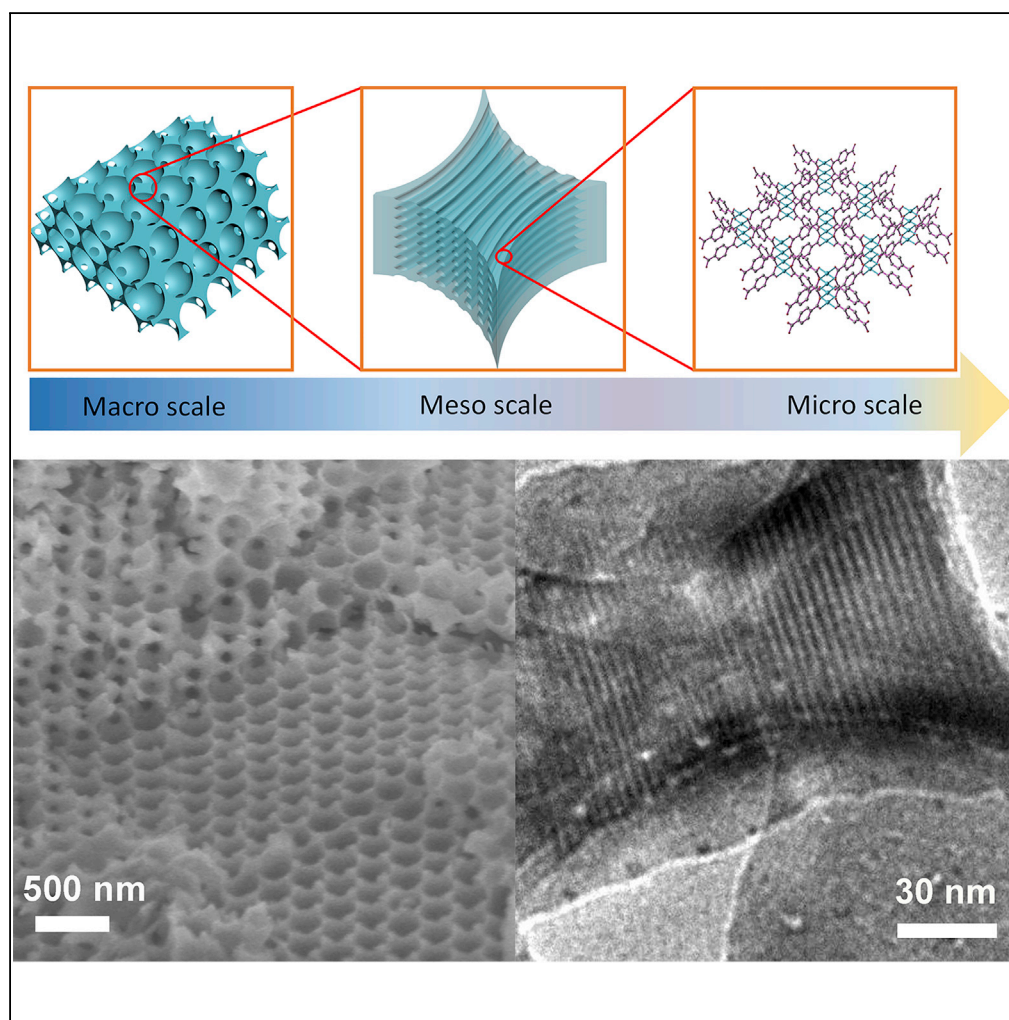
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## Article

# Hierarchical Structure with Highly Ordered Macroporous-Mesoporous Metal-Organic Frameworks as Dual Function for CO<sub>2</sub> Fixation



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**HIGHLIGHTS**

Macro-, meso-, and microporous structures were integrated in one framework

Macroporous-mesoporous MOFs are prepared by co-assembly double template method

Macroporous-mesoporous MOFs possess high activity for chemical fixation of CO<sub>2</sub>

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## Article

# Hierarchical Structure with Highly Ordered Macroporous-Mesoporous Metal-Organic Frameworks as Dual Function for CO<sub>2</sub> Fixation

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## SUMMARY

**As a major greenhouse gas, the continuous increase of carbon dioxide (CO<sub>2</sub>) in the atmosphere has caused serious environmental problems, although CO<sub>2</sub> is also an abundant, inexpensive, and nontoxic carbon source. Here, we use metal-organic framework (MOF) with highly ordered hierarchical structure as adsorbent and catalyst for chemical fixation of CO<sub>2</sub> at atmospheric pressure, and the CO<sub>2</sub> can be converted to the formate in excellent yields. Meanwhile, we have successfully integrated highly ordered macroporous and mesoporous structures into MOFs, and the macro-, meso-, and microporous structures have all been presented in one framework. Based on the unique hierarchical pores, high surface area (592 m<sup>2</sup>/g), and high CO<sub>2</sub> adsorption capacity (49.51 cm<sup>3</sup>/g), the ordered macroporous-mesoporous MOFs possess high activity for chemical fixation of CO<sub>2</sub> (yield of 77%). These results provide a promising route of chemical CO<sub>2</sub> fixation through MOF materials.**

## INTRODUCTION

From natural zeolite to meso- and macroporous materials, porous materials have been extensively applied in ion exchange (Okada et al., 2015; Li et al., 2018; Qi et al., 2015), separation (Gupta et al., 2017; Chu and Pan, 2012; Dumee et al., 2013), catalysis (An et al., 2014; Sun et al., 2016; Qian et al., 2009), drug delivery (Wang et al., 2017; Shin et al., 2017), and other fields (Kitagawa et al., 2004). The skeleton of early porous materials was composed of inorganic compounds. A new type of nanoporous material, metal-organic frameworks (MOFs), which has the properties of both inorganic and organic materials, has drawn great attention in recent years (Yaghi et al., 1995, 2003; Yaghi and Li, 1995; Matsuda et al., 2005; Lin et al., 2016). MOF is a coordination polymer composed of inorganic metal structural units and organic ligands through covalent or ionic covalent bonds. Owing to their unique high specific surface area and adjustable pore structure (Furukawa et al., 2010; Ferey et al., 2005; Aijaz et al., 2014), as well as diverse structure and excellent catalysis performance, MOFs have been widely used as functional materials in the areas of selective catalysis (Liu et al., 2014; Kornienko et al., 2015; Huang et al., 2018; Czaja et al., 2009), gas storage (Ma and Zhou, 2010; Yoon et al., 2013), optoelectronic materials (Chen et al., 2007; Allendorf et al., 2009; Stavila et al., 2014; Choi et al., 2014; Kong et al., 2016; Avci et al., 2018; Guan et al., 2017; Zheng et al., 2017), drug controlled release (Horcajada et al., 2010; Taylor-Pashow et al., 2009; Zheng et al., 2016), and molecular separation (Kang et al., 2014; Zhang et al., 2016; Zheng et al., 2017; Liu et al., 2018). Despite these excellent properties, most of the currently reported MOFs only have microporous regime, which severely obstructs the mass transfer and limits the access of large molecules to the active site (Furukawa et al., 2014; Xuan et al., 2012). Therefore it is significant to develop an efficient way to introduce larger pores into MOFs. Shen et al. (Shen et al., 2018) prepared ordered macroporous-microporous MOF single-crystal materials with the heterogeneous solvent-induced heterogeneous nucleation method using highly ordered polystyrene (PS) microspheres as template. Koo et al. (Koo et al., 2017) synthesized water-stable hierarchical porous MOFs by a selective acid etching process. Ma et al. (Ma et al., 2012) presented mesoporous MOFs by a rational bottom-up method using complicated disulfonate acids as ligands and metal nitrates or chlorides as inorganic precursors at high pressure and high temperature. Zhao et al. (Zhao et al., 2011) proposed the synthesis of MOF nanosphere with well-ordered mesopores in ionic liquid, high pressure, and supercritical CO<sub>2</sub> atmosphere. For surfactant-assisted synthesis, it is difficult to remove the surfactants without compromising the mesoporosity. The larger the pores, the more readily the frameworks collapse.

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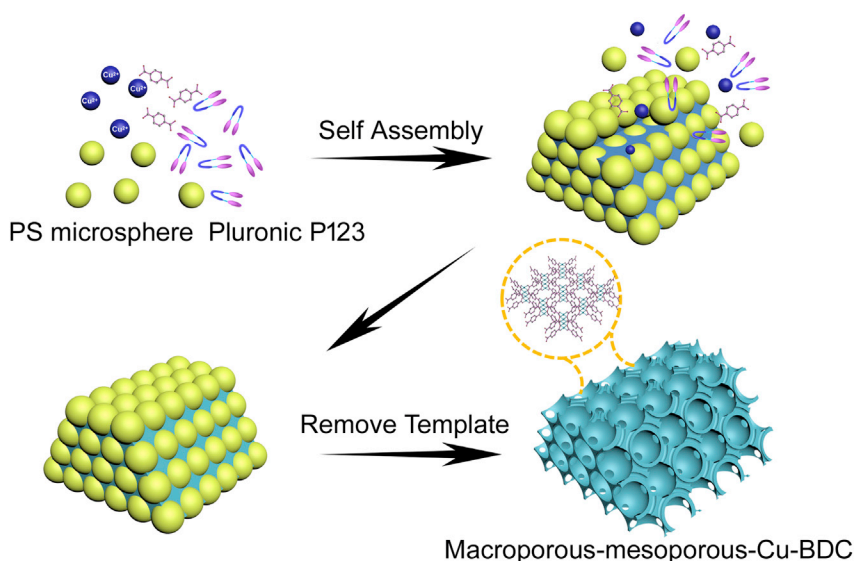
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**Figure 1. Schematic Illustration**

Schematic illustration of the synthesis procedure of the highly ordered macroporous-mesoporous MOFs.

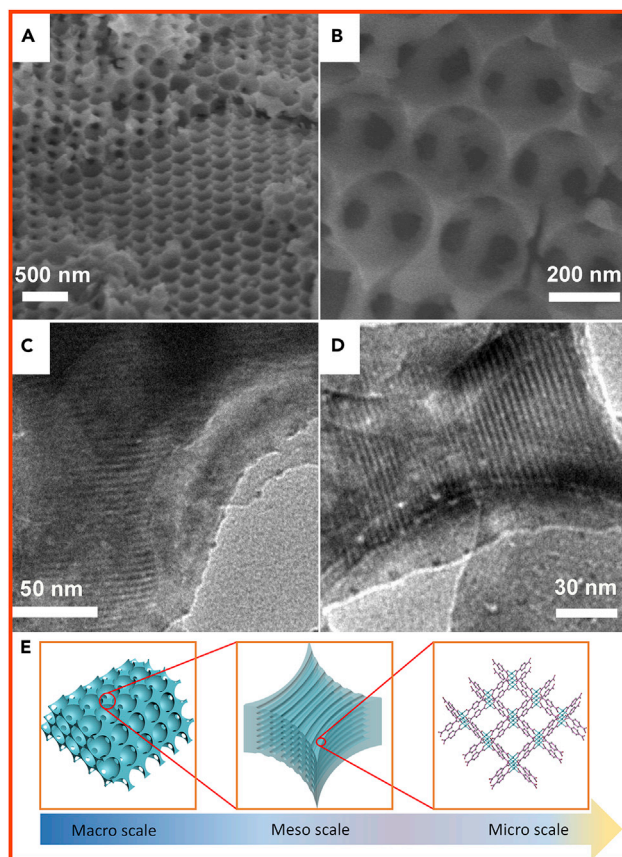
There are so many living organisms with exquisite hierarchical structure and fantastic functions in nature, such as plant stems (Bae et al., 2014), butterfly wings (Peng et al., 2011), lotus leaves (Ensikat et al., 2011), and bone (Meyers et al., 2008), which exhibit highly organized hierarchical structures from the nano- to the micrometer scales, providing us a vivid inspiration and reference. We introduce the bionics ideas into the traditional preparation methods, imitating the hierarchical structure of biological systems to build hierarchical MOF structures and introducing the fine hierarchical structure of natural biomaterials into the MOFs to prepare a new type of hierarchical MOF materials. It is rare to introduce macro- and mesoporous structures into the MOFs at the same time to construct highly organized hierarchical structures, which are similar to those of the living organisms. The new hierarchical MOF materials therefore have highly ordered morphological structures and multilevel distribution in the continuous scale range. The ordered meso- and macrochannels can facilitate accessibility and mass transfer with high efficiency and can be applied in many fields.

As a major greenhouse gas, the continuous increase of carbon dioxide ( $\text{CO}_2$ ) in the atmosphere has caused serious environmental problems (Liao et al., 2015). However,  $\text{CO}_2$  is also an abundant, inexpensive, and nontoxic carbon source. However, due to its thermodynamic and kinetic stability,  $\text{CO}_2$  has been largely limited in chemical synthesis. Herein, we first synthesized the highly ordered macroporous-mesoporous MOFs, novel hierarchical MOF materials with three-scale porous sizes, the macroporous, mesoporous, and microporous structures, in one framework. Furthermore, both the macro- and mesoporous structures are highly ordered. Meanwhile, the ordered macroporous-mesoporous MOFs possess significant capture and selectivity for  $\text{CO}_2$  and can be used to catalyze the reaction of  $\text{CO}_2$  with benzyl halogen for chemical  $\text{CO}_2$  fixation under ambient conditions. These results provide a promising route for chemical  $\text{CO}_2$  fixation through MOF materials.

## RESULTS AND DISCUSSION

We demonstrate a solvent evaporation-induced and co-assembly route to synthesize highly ordered macroporous-mesoporous MOFs with hierarchical structure using PS as the macroporous template and block copolymers as the mesoporous template (Figure 1). Both the macro- and mesoporous templates were assembled into the MOFs at the same time by ethanol evaporation process. After removing the templates, the highly ordered macroporous and mesoporous structures could be successfully introduced into the MOF structure to construct a hierarchical structure by using the one-pot method.

The hard template PS microspheres were uniform with diameters of 455 nm, as can be seen in Figure S2. Figures 2A and 2B show the representative scanning electron microscopic images of the synthesized



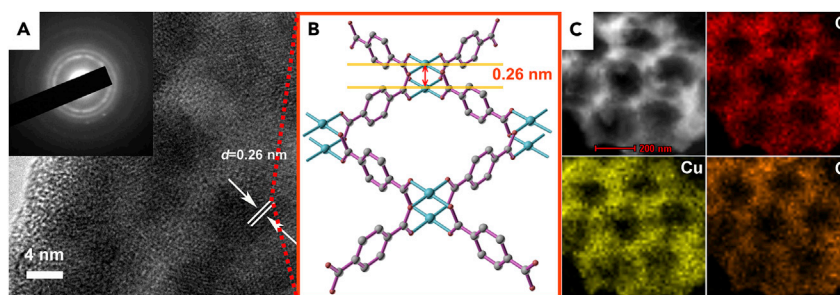
**Figure 2. Pore Structure Characterization of Macro-Meso-Cu-BDC**

(A–D) (A) and (B) Scanning electron microscopic images and (C) and (D) TEM images of macro-meso-Cu-BDC. (E) Schematic illustration of hierarchical structure.

macroporous-mesoporous MOFs (macro-meso-Cu-BDC), wherein the uniform macropores with ordered arrangement can be observed clearly. The well-ordered close-packed structures were organized into a typical face-centered cubic close-packed arrangement ordered over a range of tens of micrometers. The diameter of the macropores was 265 nm, which was smaller than that of the PS microspheres due to shrinkage during dissolution of the PS microspheres in organic solvents. The hexagonal mesoporous structure was confirmed by transmission electron microscopy (TEM). The TEM images (Figures 2C and 2D) show a high degree of periodicity in the mesoporous structure of the macroporous walls, which were in the [110] orientation, and the diameter of the uniform mesopores was 4.1 nm. For clearly conveying the idea here, Figure 2E shows the schematic illustration of the hierarchical structure.

The crystal lattice fringes were observed in high-resolution TEM image (Figure 3A), confirming the high crystallinity of the mesoporous walls. Lattice fringes with interplanar distance of 0.26 nm were observed, corresponding to the (001) plane of Cu-BDC. The crystalline nature of the mesoporous wall of the macro-meso-Cu-BDC was also verified by the observable polycrystalline diffraction rings in the selected area electron diffraction pattern (Figure 3A, inset). The elemental mapping image (Figure 3C) exhibits a homogeneous distribution of all the related elements, including C, O, and Cu, throughout whole macroporous-mesoporous MOFs.

The small-angle X-ray diffraction (XRD) pattern of the synthesized macro-meso-Cu-BDC (Figure 4A) demonstrates a typical hexagonal ( $p6mm$ ) mesophase with a strong diffraction peak at  $2\theta = 2.04^\circ$  and two small peaks at  $2\theta = 2.65^\circ$  and  $2.91^\circ$ , which could be attributed to (100), (110), and (200) reflection, respectively. The corresponding  $d$  spacing is calculated to be 4.2 nm, which agrees well with the TEM result. Figure 4B shows wide-angle XRD patterns of the macro-meso-Cu-BDC. Compared with pure Cu-BDC (Carson et al.,



**Figure 3. High-Resolution TEM, Energy-Dispersive X-Ray Spectroscopic Characterization, and Topology of Macro-Meso-Cu-BDC**

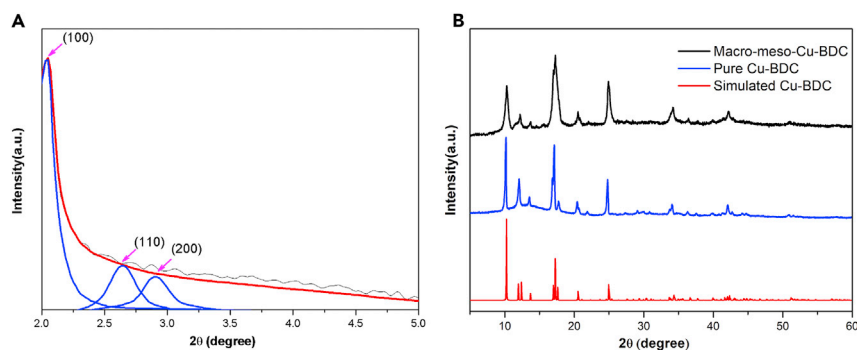
(A) High-resolution TEM image of macro-meso-Cu-BDC (the inset in (A) is the corresponding SAED pattern).  
 (B) Cu-BDC topology.  
 (C) Energy-dispersive X-ray spectroscopic element mappings of macro-meso-Cu-BDC.

2009) (blue line) and simulated Cu-BDC (red line), the reflection peaks of the macro-meso-Cu-BDC fit very well with pure Cu-BDC and simulated Cu-BDC, and no other peaks could be observed in the patterns, which demonstrates the formation of phase-pure Cu-BDC with highly crystalline mesopore walls. Furthermore, the XRD pattern of macro-meso-Cu-BDC shows broader diffraction peaks, suggesting that ordered mesostructure was obtained.

In agreement with XRD, the Fourier transform infrared (FTIR) spectra (Figure S3A) for macro-meso-Cu-BDC and pure Cu-BDC samples also represent the remaining unchanged main building block of the Cu-BDC structure. The appearance of a broad band at  $3,000\text{--}3,700\text{ cm}^{-1}$  indicates the presence of -OH groups and water. The peaks at  $1,576\text{ cm}^{-1}$  and  $1,690\text{ cm}^{-1}$  correspond to the symmetric and asymmetric stretching vibrations of the carboxylate groups in Cu-BDC, respectively (Bhardwaj et al., 2015). In addition, the FTIR spectra for the sample without template removal, the sample with removal of only PS, and the sample with removal of only P123 were measured and demonstrated in Figure S4. The characteristic peak attributable to PS at  $2,800\text{--}3,100\text{ cm}^{-1}$  disappeared, substantiating the fact that the PS template is all removed.

The thermal stabilities of the as-prepared macro-meso-Cu-BDC and pure Cu-BDC were examined by thermogravimetric analysis (TGA) under argon ambience. For the macro-meso-Cu-BDC (Figure S3B), the thermogravimetric profile can be divided into two steps: the vaporization of adsorbed water in the pores occurred at  $130^\circ\text{C}$  and then a sharp weight loss can be observed at temperatures between  $250$  and  $440^\circ\text{C}$ , which reflects the collapse of the framework due to the decomposition of the organic ligand. In the TGA profile of pure Cu-BDC, two obvious weight loss steps in the temperature range of  $190^\circ\text{C}\text{--}255^\circ\text{C}$  and  $325^\circ\text{C}\text{--}460^\circ\text{C}$  corresponding to the removal of the free molecules and the decomposition of the framework, respectively, were respectively found. The weight loss of macro-meso-Cu-BDC at lower temperatures than pure Cu-BDC further confirms the open meso- and macroporous hierarchical structure in the macro-meso-Cu-BDC.

To obtain further structural information, the porosities of pure Cu-BDC and macro-meso-Cu-BDC samples were measured by  $\text{N}_2$  adsorption-desorption isotherms at 77 K, the results of which are shown in Figure 5A. Pure Cu-BDC shows type I isotherms according to the International Union of Pure and Applied Chemistry classification (Sing, 1985), indicating the formation of microporous structures. Compared with pure Cu-BDC, the adsorption isotherms of macro-meso-Cu-BDC exhibited an intermediate mode between type I, characteristic of microporous structure, and type IV, characteristic of mesoporous structure, with a high nitrogen adsorption capacity at very low relative pressures. The hysteresis loop was observed to indicate the existence of mesoporous structure. Furthermore, the corresponding micropore size distribution curves of macro-meso-Cu-BDC and pure Cu-BDC were calculated using nonlocal density functional theory modeling method, and a narrow peak at  $0.67\text{ nm}$  was observed in both the curves. Moreover, as calculated from the Barrett-Joyner-Halenda method (Figure 5B, inset), a uniform mesopore size of  $3.9\text{ nm}$  was obtained in the macro-meso-Cu-BDC, which agrees well with the TEM result. The Brunauer-Emmett-Teller surface area of macro-meso-Cu-BDC was  $592\text{ m}^2/\text{g}$ , which was higher than that of the pure Cu-BDC ( $540\text{ m}^2/\text{g}$ ). It suggests that the macro- and mesoporous structures are in favor of the improvement of the surface areas in the macro-meso-Cu-BDC. The  $\text{CO}_2$  uptake isotherms of the samples measured at



**Figure 4. XRD Characterization of Macro-Meso-Cu-BDC**

(A) Small-angle X-ray diffraction pattern of macro-meso-Cu-BDC.

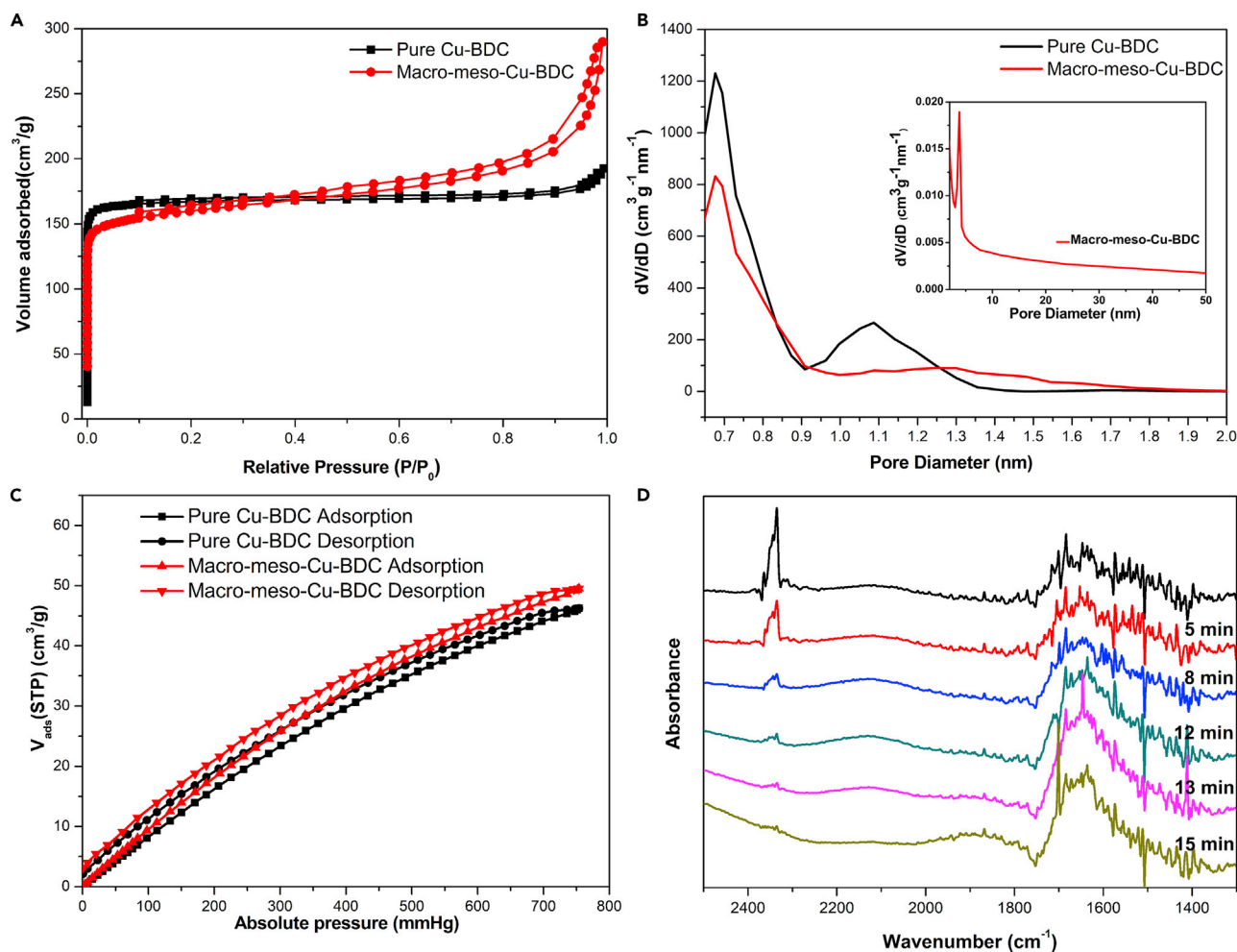
(B) Wide-angle XRD patterns of macro-meso-Cu-BDC, when compared with pure Cu-BDC (blue line) and simulated Cu-BDC (red line).

273 K and up to 1 bar are shown in Figure 5C. The macro-meso-Cu-BDC exhibited a high CO<sub>2</sub> adsorption capacity with 49.51 cm<sup>3</sup>/g, whereas pure Cu-BDC gave a lower CO<sub>2</sub> capacity with only 46.23 cm<sup>3</sup>/g. It suggests that CO<sub>2</sub> adsorption capacity of the macro-meso-Cu-BDC is higher than that of pure Cu-BDC, which is related to the hierarchical structures of macro-meso-Cu-BDC. The large CO<sub>2</sub> adsorption capacity of macro-meso-Cu-BDC may be attributed to its high specific surface area compared with that of pure Cu-BDC.

To further investigate the interaction between macro-meso-Cu-BDC and CO<sub>2</sub>, *in situ* diffuse reflectance infrared Fourier transform (DRIFT) experiments were carried out at room temperature (Figure 5D). During the CO<sub>2</sub> adsorption, the macro-meso-Cu-BDC displayed a distinct and intense band centered at approximately 2,335 cm<sup>-1</sup> corresponding to the ν<sub>3</sub> mode of CO<sub>2</sub>, which showed a significant red shift compared with that of the gas phase (2,349 cm<sup>-1</sup>). It is due to the formation of the H-bonds between CO<sub>2</sub> and the -OH groups of Cu-BDC (Vimont et al., 2007; Lan et al., 2011). There were also several peaks discovered in the range of 1,800 to 1,300 cm<sup>-1</sup>, which originated from the interaction between CO<sub>2</sub> and -OH groups (Ferretto and Glisenti, 2003). After stopping CO<sub>2</sub> supply, the DRIFT spectra were collected under argon purging. It was found that the characteristic band at 2,335 cm<sup>-1</sup> significantly decreased with time and almost disappeared from the spectra after 15-min purging. However, the intensities of peaks at 1,800 to 1,300 cm<sup>-1</sup> remain unchanged, disclosing a strong bonding between CO<sub>2</sub> and the macro-meso-Cu-BDC.

In addition, the basicity of the as-synthesized macro-meso-Cu-BDC and the corresponding pure Cu-BDC were also evaluated by CO<sub>2</sub>-temperature-programmed reduction (TPD) experiments in Figure S5. CO<sub>2</sub> was adsorbed at 25°C and then desorbed in the range of 25°C–700°C. The macro-meso-Cu-BDC and the pure Cu-BDC both showed two obvious CO<sub>2</sub> desorption peaks around 372°C and 424°C and 400°C and 436°C, respectively. It indicated the presence of different surface basic sites. Based on the peak area, higher values of basicity were observed for the macro-meso-Cu-BDC than those for the pure Cu-BDC, indicating a larger CO<sub>2</sub> adsorption capacity and more active sites for CO<sub>2</sub>.

The high CO<sub>2</sub> adsorption capacity and the unique hierarchical multiscale porous framework indicate that the macro-meso-Cu-BDC can be a highly promising heterogeneous adsorbent and catalyst for the chemical fixation of CO<sub>2</sub>. The CO<sub>2</sub> carbonylative coupling reaction with 4-methylbenzyl chloride was first chosen to evaluate the performance of macro-meso-Cu-BDC at relatively mild conditions of 0.1 MPa CO<sub>2</sub> and 100°C; the desired carbonylative coupling product **1** was obtained with yield of 68%, accompanied by the by-product **2** in 3% yield, which reacted with *N,N*-Dimethylformamide (DMF) (Table S1, entry 1). Unlike the macro-meso-Cu-BDC-catalyzed reaction, when pure Cu-BDC was used as the catalyst, the reaction showed a lower yield and selectivity and only a small amount (8%) of **1** was afforded, with **2** in 10% yield (Table S1, entry 2). At the same time, the meso-Cu-BDC has only 13% yield and 6% selectivity (Table S1, entry 3). Comparing various copper salts, the results show that the copper salt does not have any effect on the reaction (Figure S7, Table S1, entries 4–7). Macro-meso-Cu-BDC showed an excellent catalytic performance in the chemical fixation CO<sub>2</sub> reaction, which could significantly enhance the yield of the desired product, because ordered meso- and macrochannels existed in the MOFs, which can facilitate accessibility



**Figure 5. Characterization of Adsorption Capacity of Macro-Meso-Cu-BDC**

(A and B) (A)  $N_2$  adsorption isotherms and (B) pore size distribution analyses based on the nonlocal density functional theory method for macro-meso-Cu-BDC and pure Cu-BDC (the inset in (B) is the corresponding Barrett-Joyner-Halenda pattern).

(C)  $CO_2$  adsorption isotherms of macro-meso-Cu-BDC and pure-Cu-BDC at 273 K.

(D) DRIFT spectrum of  $CO_2$  adsorbed on macro-meso-Cu-BDC at room temperature (black line). Spectra collected with Ar purge for different time lengths: 5, 8, 12, 13, and 15 min.

and mass transfer with high efficiency. Owing to the unique hierarchical pores, high surface area, and high  $CO_2$  adsorption capacity, macro-meso-Cu-BDC exposes more active sites, which is more conducive to adsorbing and activating large amounts of  $CO_2$  to catalyze the reaction at copper active sites.

To demonstrate the scope and applicability of this macro-meso-Cu-BDC for other structurally diverse benzyl halogen, the  $CO_2$  carbonylative coupling reactions with various benzyl halogens are carried out, and the results were summarized in Table 1. The reactions of 4-nitrobenzyl chloride (1a), 4-fluorobenzyl chloride (1b), and 4-methoxybenzyl chloride (1c) that bear strong electron-withdrawing groups on the *para*-position provide high yields for products 2a, 2b, and 2c under the same reaction conditions (entries 1–3, 77%, 71%, and 70%). By only reducing the amount of  $Cs_2CO_3$ , products 2d and 2e were also obtained in good yield from the reactions of benzyl chlorides 1d and 1e bearing an electron-donating group methyl (Me) on the *para*-position (entry 4, 68%) and *meta*-position (entry 5, 66%). Furthermore, 4-*tert*-butylbenzyl chloride (1f) was converted with a yield of 50% (entry 6). Moreover, benzyl chloride (1g) and benzyl bromide (1h) were also utilized in this type of carbonylative coupling reaction. Products 2g and 2h are obtained with yields of 60% and 43%, respectively (entries 7–8). Therefore this catalysis system was quite versatile, as a variety of benzyl halogens could be converted to the corresponding phenyl ester in excellent yields. The



Entry	Benzyl Halogen	Product	Yield (%) <sup>b</sup>
1			77
2			71
3			70
4 <sup>c</sup>			68
5 <sup>c</sup>			66
6 <sup>c</sup>			50
7 <sup>c</sup>			60
8 <sup>c</sup>			43

**Table 1. Summary of the Results of CO<sub>2</sub> Carbonylation Coupling Reactions**

CO<sub>2</sub> fixation with benzyl halogen with different functional groups catalyzed by macro-meso-Cu-BDC.<sup>a</sup>

<sup>a</sup>Reaction conditions: benzyl halogen (2 mmol), NaBH<sub>4</sub> (2 mmol), Cs<sub>2</sub>CO<sub>3</sub> (0.4 mmol), CO<sub>2</sub> (0.1 MPa), catalyst (10 mol %), DMF (8 mL) at 100°C for 8 h.

<sup>b</sup>Isolated yield.

<sup>c</sup>Cs<sub>2</sub>CO<sub>3</sub> (0.2 mmol).

XRD pattern and TEM image of macro-meso-Cu-BDC after catalytic reaction showed that the phase and pore structures of the catalyst remained almost unchanged, which also confirmed the stability of the catalyst (Figure S8).

### Conclusion

In summary, the novel highly ordered hierarchical macro-meso-Cu-BDC with crystalline mesoporous walls was successfully synthesized by using a facile solvent evaporation induced co-assembly route in one step. These hierarchical MOF materials with three-scale porous sizes, which are the macro-, meso-, and microporous structures, have been presented in one framework. The highly ordered meso- and macrochannels can further improve the mass transfer efficiency. Owing to the unique hierarchical pores, high surface area, and high CO<sub>2</sub> adsorption capacity, the macro-meso-Cu-BDC possesses high activity for chemical fixation of CO<sub>2</sub> under mild conditions. These results provide a promising route to chemical CO<sub>2</sub> fixation through MOF materials.

### Limitations of Study

Owing to the poor structural stability of macro-meso-Cu-BDC, we do not know if it can be used for longer period (>50 cycles) catalytic reactions.

### METHODS

All methods can be found in the accompanying [Transparent Methods supplemental file](#).

### SUPPLEMENTAL INFORMATION

Supplemental Information can be found online at <https://doi.org/10.1016/j.isci.2019.05.006>.

### ACKNOWLEDGMENTS

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### AUTHOR CONTRIBUTIONS

Y.Y. and Z.L. conceived the project and designed the experiments. X.X., J.C., and J.X. performed the experiments. D.M. and Z.W. carried out material characterization. R.W. assisted in material characterization. Z.L. and M.L. wrote the manuscript. All authors discussed the results and commented on the manuscript.

### DECLARATION OF INTERESTS

The authors declare no competing interests.

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### REFERENCES

- Aijaz, A., Fujiwara, N., and Xu, Q. (2014). From metal-organic framework to nitrogen-decorated nanoporous carbons: high CO<sub>2</sub> uptake and efficient catalytic oxygen reduction. *J. Am. Chem. Soc.* *136*, 6790–6793.
- Allendorf, M.D., Bauer, C.A., Bhakta, R.K., and Houk, R.J.T. (2009). Luminescent metal-organic frameworks. *Chem. Soc. Rev.* *38*, 1330–1352.
- An, K., Alayoglu, S., Musselwhite, N., Na, K., and Somorjai, G.A. (2014). Designed catalysts from Pt nanoparticles supported on macroporous oxides for selective isomerization of n-Hexane. *J. Am. Chem. Soc.* *136*, 6830–6833.
- Avci, C., Imaz, I., Carné-Sánchez, A., Pariente, J.A., Tasios, N., and Pérez-Carvajal, J. (2018). Self-assembly of polyhedral metal-organic framework particles into three-dimensional ordered superstructures. *Nat. Chem.* *10*, 78–84.
- Bae, W.G., Kim, H.N., Kim, D., Park, S.H., Jeong, H.E., and Suh, K.Y. (2014). 25th Anniversary article: scalable multiscale patterned structures inspired by nature: the role of hierarchy. *Adv. Mater.* *26*, 675–700.
- Bhardwaj, S.K., Bhardwaj, N., Mohanta, G.C., Kumar, P., Sharma, A.L., Kim, K.H., and Deep, A. (2015). Immunosensing of atrazine with antibody-functionalized Cu-MOF conducting thin films. *ACS Appl. Mater. Interfaces* *7*, 26124–26130.
- Carson, C.G., Hardcastle, K., Schwartz, J., Liu, X., Hoffmann, C., and Gerhardt, R.A. (2009).

- Synthesis and structure characterization of copper terephthalate metal-organic frameworks. *Eur. J. Inorg. Chem.* 16, 2338–2343.
- Chen, B., Yang, Y., Zapata, F., Lin, G., Qian, G., and Lobkovsky, E.B. (2007). Luminescent open metal sites within a metal-organic framework for sensing small molecules. *Adv. Mater.* 19, 1693–1696.
- Choi, K.M., Jeong, H.M., Park, J.H., Zhang, Y.B., Kang, J.K., and Yaghi, O.M. (2014). Supercapacitors of nanocrystalline metal-organic frameworks. *ACS Nano* 8, 7451–7457.
- Chu, Y., and Pan, Q.M. (2012). Three-dimensionally macroporous Fe/C nanocomposites as highly selective oil-absorption materials. *ACS Appl. Mater. Interfaces* 4, 2420–2425.
- Czaja, A.U., Trukhan, N., and Mütler, U. (2009). Industrial applications of metal-organic frameworks. *Chem. Soc. Rev.* 38, 1284–1293.
- Dumée, L., He, L., Lin, B., Ailloux, F.M., Lemoine, J.B., Velleman, L., She, F., Duke, M., Orbell, J., and Erskine, G. (2013). The fabrication and surface functionalization of porous metal frameworks—A review. *J. Mater. Chem. A* 1, 15185–15206.
- Ensatat, H.J., Ditsche-Kuru, P., Neinhuis, C., Barthlott, W., and Beilstein, J. (2011). Superhydrophobicity in perfection: the outstanding properties of the lotus leaf. *Beilstein J. Nanotechnol.* 2, 152–161.
- Ferretto, L., and Glisenti, A. (2003). Surface acidity and basicity of a rutile powder. *Chem. Mater.* 15, 1181–1188.
- Ferey, G., Mellot-Draznieks, C., Serre, C., Millange, F., Dutour, J., Surblé, S., and Margiolaki, I. (2005). A chromium terephthalate-based solid with unusually large pore volumes and surface area. *Science* 309, 2040–2042.
- Furukawa, H., Ko, N., Go, Y.B., Aratani, N., Choi, S.B., Choi, E., Yazaydin, A.O., Snurr, R.Q., O’Keeffe, M., Kim, J., and Yaghi, O.M. (2010). Ultrahigh porosity in metal-organic frameworks. *Science* 329, 424–428.
- Furukawa, S., Reboul, J., Diring, S., Sumida, K., and Kitagawa, S. (2014). Structuring of metal-organic frameworks at the mesoscopic/macroscopic scale. *Chem. Soc. Rev.* 43, 5700–5734.
- Guan, B.Y., Yu, X.Y., Wu, H.B., and Lou, X. (2017). Complex nanostructures from materials based on metal-organic frameworks for electrochemical energy storage and conversion. *Adv. Mater.* 29, 1703614.
- Gupta, R.K., Dunderdale, G.J., England, M.W., and Hozumi, A. (2017). Oil/water separation techniques: a review of recent progresses and future directions. *J. Mater. Chem. A* 5, 16025–16058.
- Horcajada, P., Chalati, T., and Serre, C. (2010). Porous metal-organic-framework nanoscale carriers as a potential platform for drug delivery and imaging. *Nat. Mater.* 9, 172–178.
- Huang, N., Drake, H., Li, J., Pang, J., Wang, Y., Yuan, S., Wang, Q., Cai, P., Qin, J., and Zhou, H.C. (2018). Flexible and hierarchical metal-organic framework composites for high-performance catalysis. *Angew. Chem. Int. Ed.* 57, 8916–8920.
- Kang, Z., Xue, M., Fan, L., Huang, L., Guo, L., Wei, G., Chen, B.L., and Qiu, S.L. (2014). Highly selective sieving of small gas molecules by an ultra-microporous metal-organic framework membrane. *Energy Environ. Sci.* 7, 4053–4060.
- Kitagawa, S., Kitaura, R., and Noro, S. (2004). Functional porous coordination polymers. *Angew. Chem. Int. Ed.* 43, 2334–2375.
- Kong, B., Tang, J., Zhang, Y., Jiang, T., Gong, X., Peng, C., Wei, J., Yang, J., Wang, Y., Wang, X., et al. (2016). Incorporation of well-dispersed sub-5-nm graphitic pencil nanodots into ordered mesoporous frameworks. *Nat. Chem.* 8, 171–178.
- Koo, J.Y., Hwang, I.C., Yu, X.J., Saha, S.H., Kim, Y.H., and Kim, K. (2017). Hollowing out MOFs: hierarchical micro- and mesoporous MOFs with tailorable porosity via selective acid etching. *Chem. Sci.* 8, 6799–6803.
- Kornienko, N., Zhao, Y., Kley, C.S., Zhu, C., Kim, D., Lin, S., Chang, C.J., Yaghi, O.M., and Yang, P. (2015). Metal-organic frameworks for electrocatalytic reduction of carbon dioxide. *J. Am. Chem. Soc.* 137, 14129–14135.
- Lan, Y.Q., Jiang, H.L., Li, S.L., and Xu, Q. (2011). Mesoporous metal-organic frameworks with size-tunable cages: selective CO<sub>2</sub> uptake, encapsulation of Ln<sup>3+</sup> cations for luminescence, and column-chromatographic dye separation. *Adv. Mater.* 23, 5015–5020.
- Li, J., Wang, X., Zhao, G., Chen, C., Chai, Z., Alsaedi, A., Hayat, T., and Wang, X. (2018). Metal-organic framework-based materials: superior adsorbents for the capture of toxic and radioactive metal ions. *Chem. Soc. Rev.* 47, 2322–2356.
- Liao, P.Q., Chen, H., Zhou, D.D., Liu, S.Y., He, C.T., Rui, Z., Ji, H.B., Zhang, J.P., and Chen, X.M. (2015). Monodentate hydroxide as a super strong yet reversible active site for CO<sub>2</sub> capture from high humidity flue gas. *Energy Environ. Sci.* 8, 1011–1016.
- Lin, J.M., He, C.T., Liu, Y., Liao, P.Q., Zhou, D.D., Zhang, J.P., and Chen, X.M. (2016). A metal-organic framework with a pore size/shape suitable for strong binding and close packing of methane. *Angew. Chem. Int. Ed.* 55, 4674–4678.
- Liu, J.W., Liu, J., Chen, L., Cui, H., Zhang, J., Zhang, L., and Su, C.Y. (2014). Applications of metal-organic frameworks in heterogeneous supramolecular catalysis. *Chem. Soc. Rev.* 43, 6011–6061.
- Liu, G., Chernikova, V., Liu, Y., Zhang, K., Belmabkhout, Y., Shekhat, O., Zhang, C., Yi, S., Eddaoudi, M., and Koros, W.J. (2018). Mixed matrix formulations with MOF molecular sieving for key energy-intensive separations. *Nat. Mater.* 17, 283–289.
- Ma, S.Q., and Zhou, H.C. (2010). Gas storage in porous metal-organic frameworks for clean energy applications. *Chem. Commun.* 46, 44–53.
- Ma, T.Y., Li, H., Deng, Q.F., Liu, L., Ren, T.Z., and Yuan, Z.Y. (2012). Ordered mesoporous metal-organic frameworks consisting of metal disulfonates. *Chem. Mater.* 24, 2253–2255.
- Matsuda, R., Kitaura, R., Kitagawa, S., Kubota, Y., and Belosludov, R.V. (2005). Highly controlled acetylene accommodation in a metal-organic microporous material. *Nature* 436, 238–241.
- Meyers, M.A., Chen, P.Y., Lin, A.Y.M., and Seki, Y. (2008). Biological materials: structure and mechanical properties. *Prog. Mater. Sci.* 53, 1–206.
- Okada, K., Asakura, G., Tokudome, Y., Nakahira, A., and Takahashi, M. (2015). Macroporous titanate nanotube/TiO<sub>2</sub> monolith for fast and large-capacity cation exchange. *Chem. Mater.* 27, 1885–1891.
- Peng, W., Hu, X., and Zhang, D. (2011). Bioinspired fabrication of magneto-optic hierarchical architecture by hydrothermal process from butterfly wing. *J. Magn. Magn. Mater.* 323, 2064–2069.
- Qi, X.H., Du, K.Z., Feng, M.L., Li, J.R., Du, C.F., Zhang, B., and Huang, X.Y. (2015). A two-dimensionally microporous thiostannate with superior Cs<sup>+</sup> and Sr<sup>2+</sup> ion-exchange property. *J. Mater. Chem. A* 3, 5665–5673.
- Qian, K., Wan, J., Qiao, L., Huang, X., Tang, J., Wang, Y., Kong, J., Yang, P., Yu, C., and Liu, B. (2009). Macroporous materials as novel catalysts for efficient and controllable proteolysis. *Anal. Chem.* 81, 5749–5756.
- Shen, K., Zhang, L., Chen, X., Liu, L., Zhang, D., Han, Y., Chen, J., Long, J., Luque, R., Li, Y., and Chen, B. (2018). Ordered macro-microporous metal-organic framework single crystals. *Science* 359, 206–210.
- Shin, B.Y., Cha, B.G., Jeong, J.H., and Kim, J. (2017). Injectable macroporous ferrogel microbeads with a high structural stability for magnetically actuated drug delivery. *ACS Appl. Mater. Interfaces* 9, 31372–31380.
- Sing, K.S.W. (1985). Reporting physisorption data for gas/solid systems with special reference to the determination of surface area and porosity (recommendations 1984). *Pure Appl. Chem.* 57, 603–619.
- Stavila, V., Talin, A.A., and Allendorf, M.D. (2014). MOF-based electronic and opto-electronic devices. *Chem. Soc. Rev.* 43, 5994–6010.
- Sun, M.H., Huang, S.Z., Chen, L.H., Li, Y., Yang, X.Y., Yuan, Z.Y., and Su, B.L. (2016). Applications of hierarchically structured porous materials from energy storage and conversion, catalysis, photocatalysis, adsorption, separation and sensing to biomedicine. *Chem. Soc. Rev.* 47, 3479–3563.
- Taylor-Pashow, K.M., Rocca, J.D., Xie, Z.G., Tran, S., and Lin, W.B. (2009). Postsynthetic modifications of iron-carboxylate nanoscale metal-organic frameworks for imaging and drug delivery. *J. Am. Chem. Soc.* 131, 14261–14263.
- Vimont, A., Travert, A., Bazin, P., Lavalley, J.C., Daturi, M., Serre, C., Férey, G., Bourrelly, S., and Llewellyn, P.L. (2007). Evidence of CO<sub>2</sub> molecule acting as an electron acceptor on a nanoporous metal-organic-framework MIL-53 or Cr<sup>3+</sup>(OH)

(O<sub>2</sub>C-C<sub>6</sub>H<sub>4</sub>-CO<sub>2</sub>). *Chem. Commun. (Camb.)* 31, 3291–3293.

Wang, B., Prinsen, P., Wang, H., Bai, Z., Wang, H., Luque, R., and Xuan, J. (2017). Macroporous materials: microfluidic fabrication, functionalization and applications. *Chem. Soc. Rev.* 46, 855–914.

Xuan, W., Zhu, C., Liu, Y., and Cui, Y. (2012). Mesoporous metal–organic framework materials. *Chem. Soc. Rev.* 41, 1677–1695.

Yaghi, O.M., and Li, H. (1995). Hydrothermal synthesis of a metal–organic framework containing large rectangular channels. *J. Am. Chem. Soc.* 117, 10401–10402.

Yaghi, O.M., Li, G., and Li, H. (1995). Selective binding and removal of guests in a microporous metal–organic framework. *Nature* 378, 703–706.

Yaghi, O.M., O’Keeffe, M., and Ockwig, N.W. (2003). Reticular synthesis and the design of new materials. *Nature* 423, 705–714.

Yoon, M., Suh, K., Natarajan, S., and Kim, K. (2013). Proton conduction in metal–organic frameworks and related modularly built porous solids. *Angew. Chem. Int. Ed.* 52, 2688–2700.

Zhang, Z., Nguyen, H.T.H., Miller, S.A., Ploskonka, A.M., Decoste, J.B., and Cohen, S.M. (2016). Polymer–metal–organic frameworks (polymofs) as water tolerant materials for selective carbon dioxide separations. *J. Am. Chem. Soc.* 138, 920–925.

Zhao, Y., Zhang, J., Han, B., Song, J., Li, J., and Wang, Q. (2011). Metal–organic framework nanospheres with well-ordered mesopores synthesized in an ionic liquid/CO<sub>2</sub>/surfactant system. *Angew. Chem. Int. Ed.* 50, 636–639.

Zheng, H., Zhang, Y., Liu, L., Wan, W., Guo, P., Nyström, A.M., and Zou, X. (2016). One-pot synthesis of metal–organic frameworks with encapsulated target molecules and their applications for controlled drug delivery. *J. Am. Chem. Soc.* 138, 962–968.

Zheng, S.S., Li, X., Yan, B., Hu, Q., and Xu, Y. (2017). Transition–metal (Fe, Co, Ni) based metal–organic frameworks for electrochemical energy storage. *Adv. Energy Mater.* 7, 1602733.

**ISCI, Volume 15**

**Supplemental Information**

**Hierarchical Structure with Highly**

**Ordered Macroporous-Mesoporous Metal-Organic**

**Frameworks as Dual Function for CO<sub>2</sub> Fixation**

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## Supporting Information

### **Hierarchical Structure with Highly Ordered Macroporous-Mesoporous MOFs as Dual-Function for CO<sub>2</sub> Fixation**

*Zhenxing Li,\* Xiaofei Xing, Dong Meng, Zhengxu Wang, Jingjing Xue, Rui Wang, Junmei Chu, and Mingming Li, and Yang Yang\**

#### **Transparent Methods**

##### *Synthesis of PS microspheres 455*

In a typical synthesis, styrene was washed with NaOH solution and distilled water until the pH value is neutral. 12 ml of styrene and 160 ml of deionized water were added into a 250 ml three-necked round-bottom flask, and nitrogen was bubbled for 15 minutes to remove the oxygen. And then the solution was heated to 70 °C and 10 mL of K<sub>2</sub>S<sub>2</sub>O<sub>8</sub> (0.007 g· mL<sup>-1</sup>) was added. The reaction was continued for 28 hours under nitrogen protection. The obtained mixture was washed with deionized water and ethanol for three times. Finally, white powder PS microspheres were obtained after dried in air at 30 °C.

##### *Synthesis of macro-meso-Cu-BDC*

In a typical synthesis, 1.5 g of P123 was dissolved in 30 ml of ethanol, after P123 was completely dissolved, 1.0 g of PS microspheres (d = 455 nm) were added and continued stirring for another 4 hours to ensure the complete dispersion of the PS microspheres. Then 0.121 g of Cu (NO<sub>3</sub>)<sub>2</sub>•3H<sub>2</sub>O was added to the above solution, and after stirring for 2 hours, 0.083 g of p-phthalic acid was added. After continuing stirring for a further 3 hours at room temperature, the solution was transferred to an oven at 30 °C for 5 days. After removal from the oven, washing several times with DMF (elimination of ligand and PS microspheres). The template P123 was then removed using an acidic ethanol solution (the volume ratio of acetic acid and ethanol=1:50). At last, the above solution was placed in a 60 °C vacuum oven for 10 hours to obtain the blue solid powder.

##### *Synthesis of meso-Cu-BDC*

In a typical synthesis, 1.5 g of P123 was dissolved in 30 ml of ethanol, after P123 was completely dissolved. Then 0.121 g of  $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$  was added to the above solution, and after stirring for 2 hours, 0.083 g of p-phthalic acid was added. After continuing stirring for a further 3 hours at room temperature, the solution was transferred to an oven at 30 °C for 5 days. The template P123 was then removed using an acidic ethanol solution (the volume ratio of acetic acid and ethanol=1:50). At last, the above solution was placed in a 60 °C vacuum oven for 10 hours to obtain the blue solid powder.

#### *Synthesis of pure Cu-BDC.*

0.530 g of  $\text{Cu}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$  was dissolved in 45 ml of DMF, after stirring for 2 hours, 0.362 g of p-phthalic acid was added and continued stirring for another 2 hours. Then the mixture was transferred to a reactor and reacted at 110 °C for 36 hours. After that, the mixture was washed with DMF and ethanol three times and dried at 40 °C for 8 hours in in vacuum to obtain the blue solid powder.

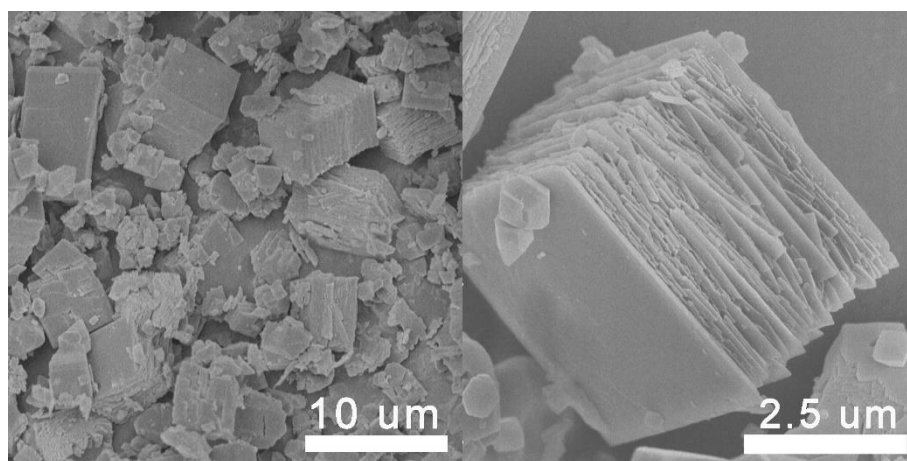
#### *Representative procedure for catalytic reactions*

In a typical experiment, the reaction was performed in a sealed Schlenk tube using benzyl halogen (0.2532 g, 2 mmol),  $\text{NaBH}_4$  (0.0757 g, 1 eq.),  $\text{Cs}_2\text{CO}_3$  (0.0652 g, 0.4 mmol) and macro-meso-Cu-BDC (10 mol %) in anhydrous DMF at 0.1 MPa and 100 °C for 4 h under the  $\text{CO}_2$  atmosphere. The crude product was purified by column chromatography (petroleum ether: ethyl acetate=30:1) to give a white solid. The resulting products were characterized by  $^1\text{H}$ NMR.

#### *Characterization*

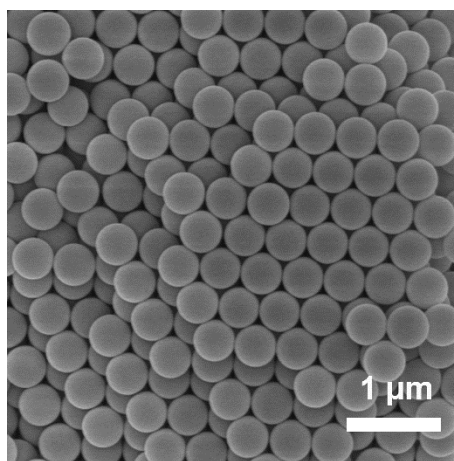
The scanning electron microscopy (SEM) observations were taken by Hitachi SU8010 scanning electron microscopy at an accelerating voltage of 30 kV. Transmission electron microscopy (TEM) and HRTEM images were carried out on a JEM 2100 LaB6 at 200 kV. The energy-dispersive X-ray analysis (EDS) were recorded on a Hitachi SU8010 scanning electron microscopy under a working voltage of 30 kV. The X-ray diffraction (XRD) data were carried out on a Burker D8-advance X-ray power diffractometer with  $\text{Cu-K}\alpha$  radiation ( $\alpha=1.5406 \text{ \AA}$ ). The FT-IR spectra were determined at room temperature on a Perkin Elmer Frontier spectrometer (equipped with a DTGS detector). Thermal gravimetric analyses (TGA) were recorded on Mettler Toledo TGA/DSC 3+ thermogravimetric

analyzer under argon atmosphere at a heating rate of  $10\text{ }^{\circ}\text{C min}^{-1}$  in the temperature range of 100-700  $^{\circ}\text{C}$ .  $\text{N}_2$  adsorption-desorption isotherms were obtained on a Micromeritics ASAP 2460 analyzer (USA) at liquid nitrogen temperature (77 K). The samples were degassed in a vacuum at 80  $^{\circ}\text{C}$  for 6 h prior to the measurement. Pore size distributions were estimated using non-local density functional theory (NLDFT) and BJH (Barret-Joyner-Halenda) model. The surface areas were calculated by the BET (Brunauer-Emmett-Teller) method. The  $\text{CO}_2$  sorption isotherms were measured by an automatic adsorption equipment (Autosorb-iQ-MP). Liquid  $^1\text{H}$ NMR spectra were taken using  $\text{CDCl}_3$  as the solvent at 600 MHz with a JNM-ECA600 spectrometer. The DRIFT spectra of  $\text{CO}_2$  adsorption were performed on a Nicolet iS10 spectrometer with an MCT detector. The samples were activated at 160  $^{\circ}\text{C}$  for 12 h before characterization. After that, the samples purged by argon flow (20 mL/min) for 20 min at room temperature. Then the spectra of  $\text{CO}_2$  adsorption were collected under a  $\text{CO}_2$  flow (3 mL/min) and the weakly adsorbed  $\text{CO}_2$  was removed under an Ar flow (5 mL/min).  $\text{CO}_2$ -TPD experiments were performed on a Quantachrome CHEMBET-3000 TPD instrument equipped with a TCD detector. The samples heated to 120  $^{\circ}\text{C}$  and held at this temperature for 30 min under a He flow (25 mL/min) to eliminate water and adsorbed  $\text{CO}_2$ . Then, the temperature was reduced to 25  $^{\circ}\text{C}$ , and  $\text{CO}_2$  flow (70 mL/min) contacted the samples for 30 min. After that, the samples were submitted to a He flow (25 mL/min) and the  $\text{CO}_2$  desorption was carried out between 25 and 700  $^{\circ}\text{C}$  at a rate of 10  $^{\circ}\text{C}/\text{min}$ .

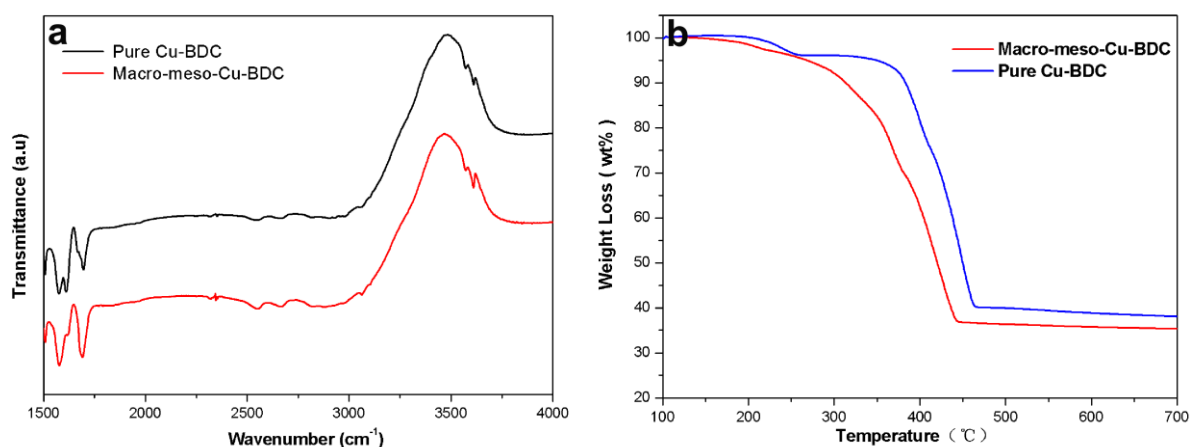


**Figure S1.** SEM image of pure Cu-BDC. (related to Figure 1)

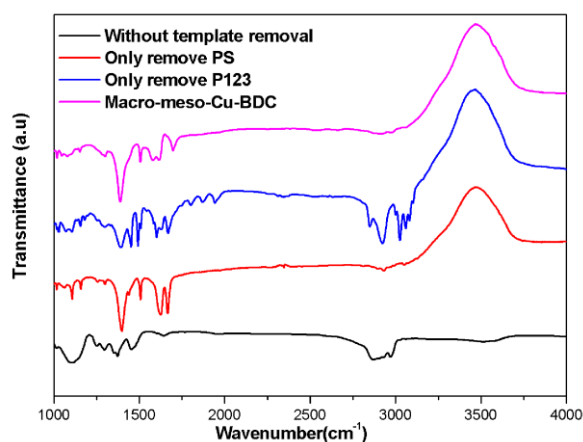




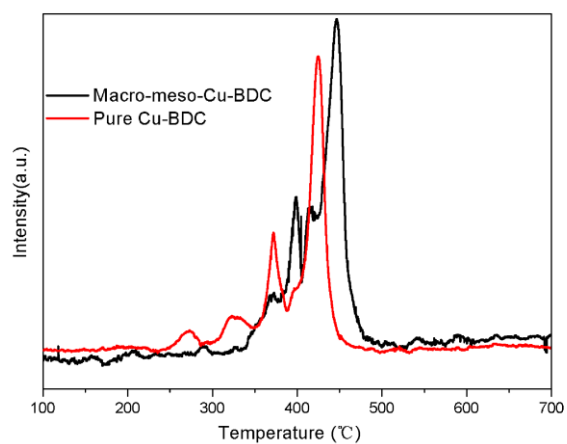
**Figure S2.** SEM image of the prepared PS microspheres with the diameter of 455 nm. (related to Figure 1)



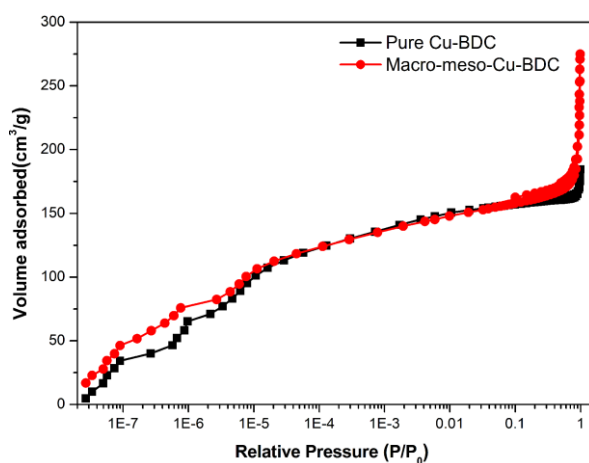
**Figure S3.** (a) FT-IR spectra of macro-meso-Cu-BDC and pure Cu-BDC, respectively. (b) Thermogravimetric analysis (TGA) of macro-meso-Cu-BDC and pure Cu-BDC in Ar atmosphere, respectively, with a heating rate of 5 °C per minute. (related to Figure 3)



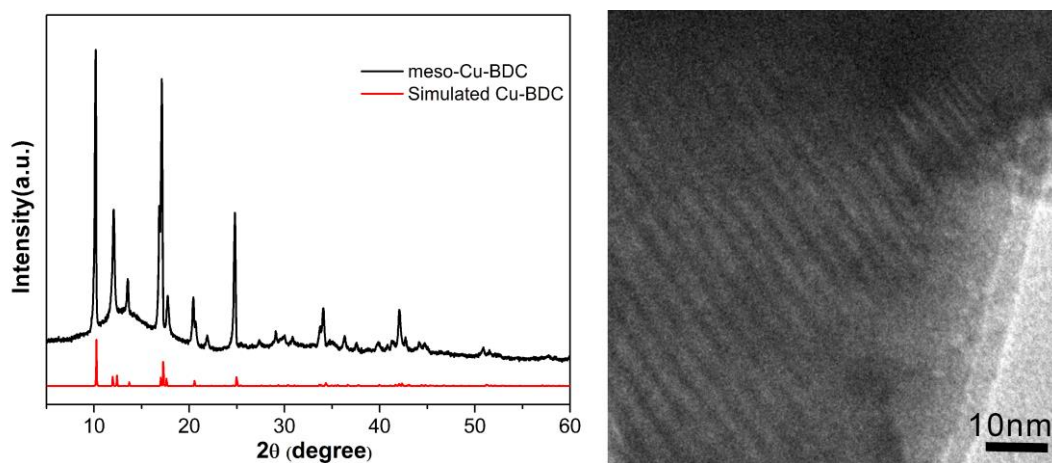
**Figure S4.** FT-IR spectra of macro-meso-Cu-BDC, the sample without template removal, the sample only remove of PS and the sample only remove of P123, respectively. (related to Figure 3)



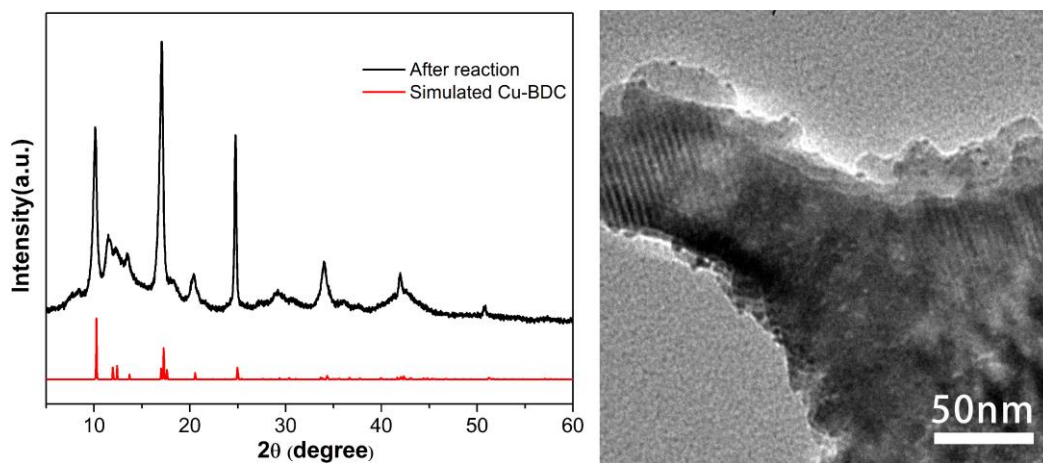
**Figure S5.** CO<sub>2</sub>-TPD spectra of macro-meso-Cu-BDC and pure Cu-BDC. (related to Figure 5)



**Figure S6.** log-scale N<sub>2</sub> adsorption isotherms for macro-meso-Cu-BDC and pure Cu-BDC. (related to Figure 5)



**Figure S7.** XRD pattern and TEM image of meso-Cu-BDC. (related to Table 1)



**Figure S8.** XRD pattern and TEM image of macro-meso-Cu-BDC after catalysis. (related to Table 1)

**Table S1.** CO<sub>2</sub> fixation with 4-methylbenzyl chloride catalyzed by macro-meso-Cu-BDC, pure Cu-BDC and meso-Cu-BDC, respectively.<sup>a)</sup> (related to Table 1)

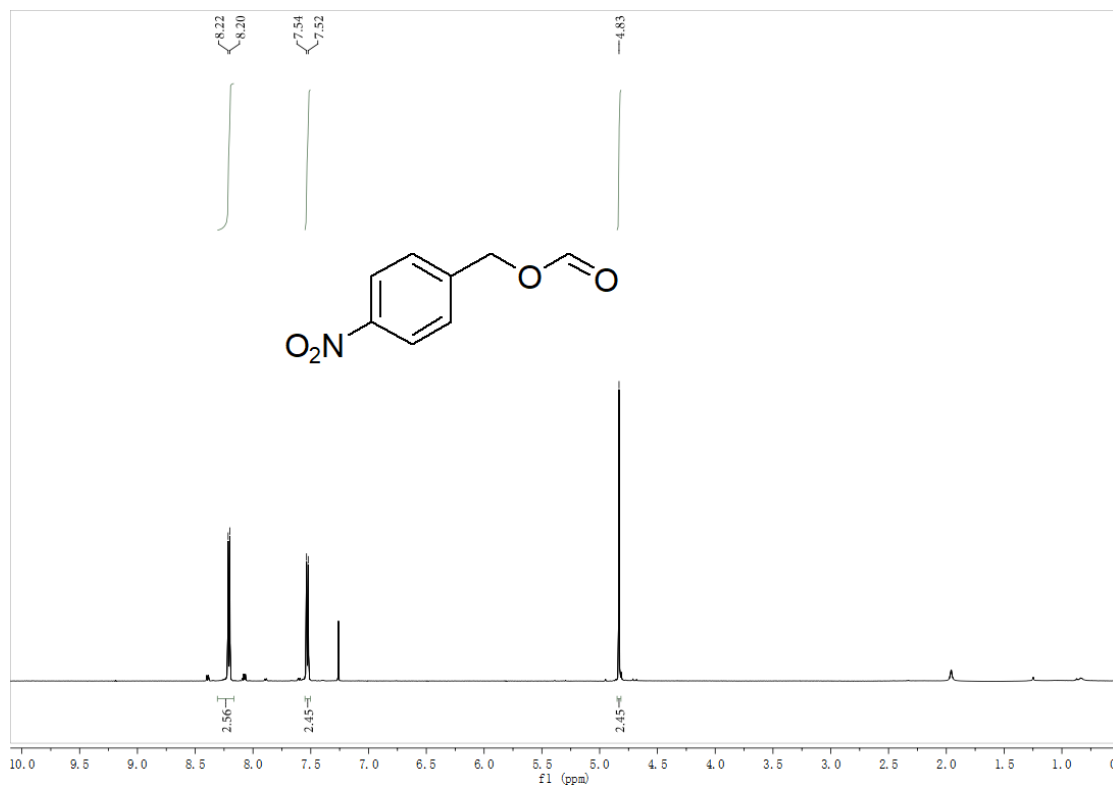
Entry	4-Methylbenzyl chloride	Catalyst	Yield (%) <sup>b)</sup>	
			1	2
1		Macro-meso-Cu-BDC	68	3
2		Pure Cu-BDC	8	10
3		Meso-Cu-BDC	13	6
4		CuCl <sub>2</sub>	Not Detected	Not Detected
5		CuSO <sub>4</sub>	Not Detected	Not Detected
6		Cu(NO <sub>3</sub> ) <sub>2</sub> ·3H <sub>2</sub> O	Not Detected	Not Detected
7		Cu(AC) <sub>2</sub> ·H <sub>2</sub> O	Not Detected	Not Detected
8 <sup>c)</sup>		Macro-meso-Cu-BDC	66	4

a) Reaction conditions: 4-methylbenzyl (2 mmol), NaBH<sub>4</sub> (2 mmol), Cs<sub>2</sub>CO<sub>3</sub>, (0.4 mmol), CO<sub>2</sub> (0.1 MPa), catalyst (10 mol %), DMF (8 mL) at 100 °C for 8 h. b) Isolated yield. c) Reaction conditions: 4-methylbenzyl (2 mmol), NaBH<sub>4</sub> (2 mmol), CsF<sub>3</sub>, (0.4 mmol), CO<sub>2</sub> (0.1 MPa), catalyst (10 mol %), DMF (8 mL) at 100 °C for 8 h.

**<sup>1</sup>HNMR (related to Table 1):**

**Product 2a**

<sup>1</sup>HNMR (600 MHz, CHLOROFORM-D) δ 8.21 (d, J = 8.7 Hz, 3H), 7.53 (d, J = 8.8 Hz, 2H), 4.83 (s, 2H).



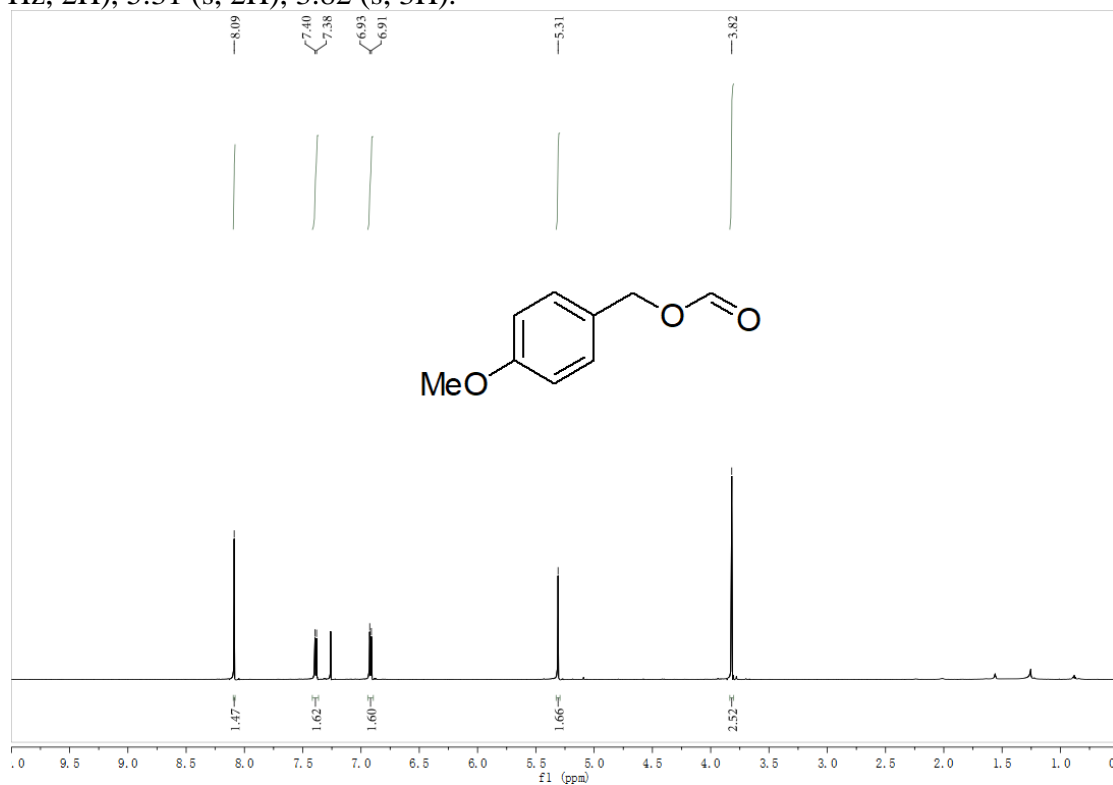
## Product 2b

$^1\text{H}$ NMR (600 MHz, CHLOROFORM-D)  $\delta$  8.11 (s, 1H), 7.43 (d,  $J = 8.5, 5.3$  Hz, 2H), 7.08 (t,  $J = 8.7$  Hz, 2H), 5.34 (s, 2H).



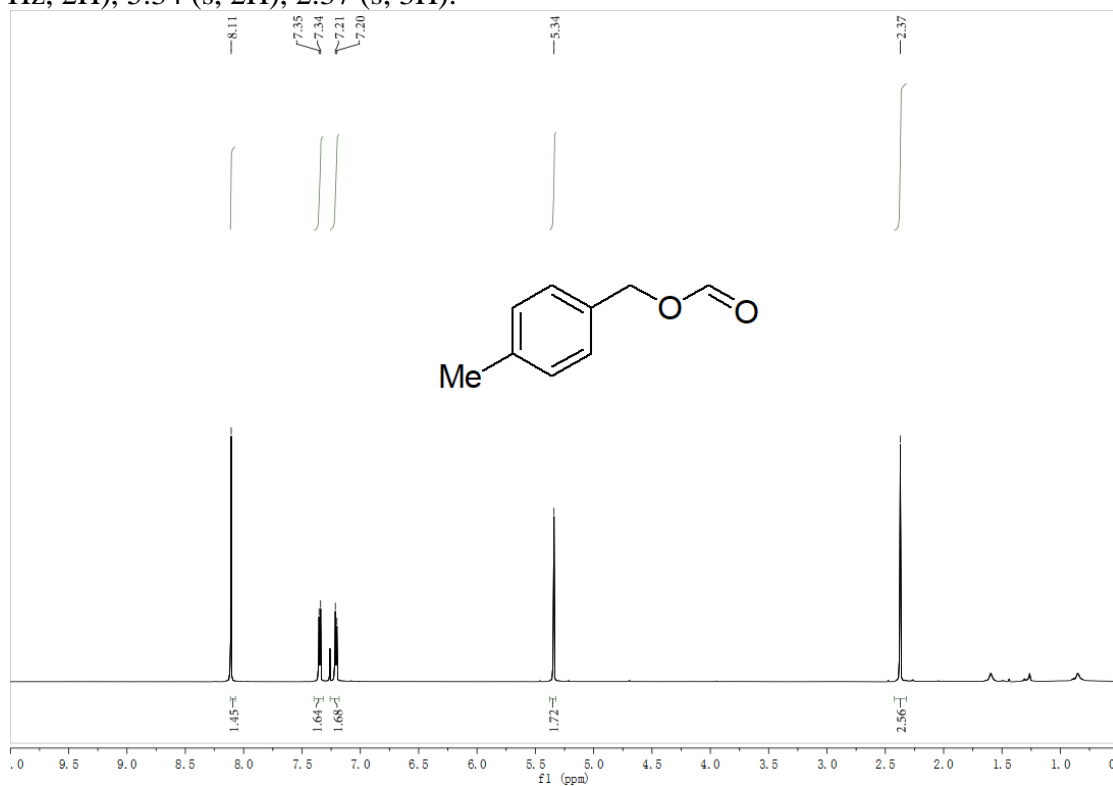
## Product 2c

$^1\text{H}$ NMR (600 MHz, CHLOROFORM-D)  $\delta$  8.09 (s, 1H), 7.39 (d,  $J = 8.4$  Hz, 2H), 6.92 (d,  $J = 8.5$  Hz, 2H), 5.31 (s, 2H), 3.82 (s, 3H).



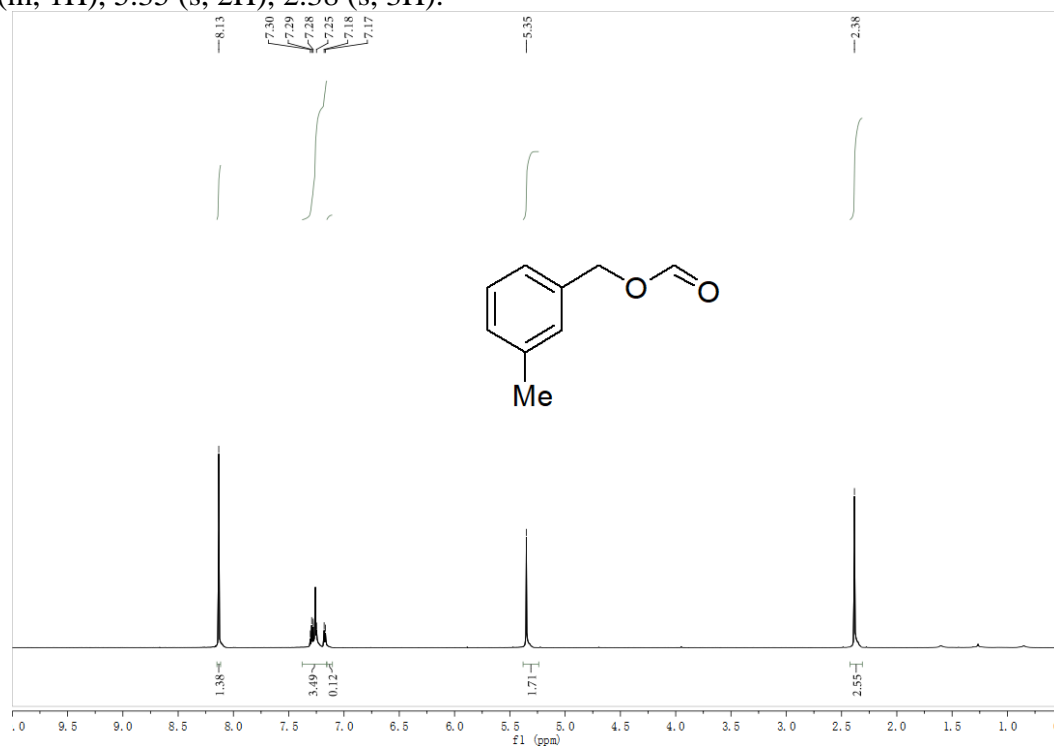
## Product 2d

$^1\text{H}$ NMR (600 MHz, CHLOROFORM-D)  $\delta$  8.11 (s, 1H), 7.35 (d,  $J = 7.9$  Hz, 2H), 7.21 (d,  $J = 7.8$  Hz, 2H), 5.34 (s, 2H), 2.37 (s, 3H).



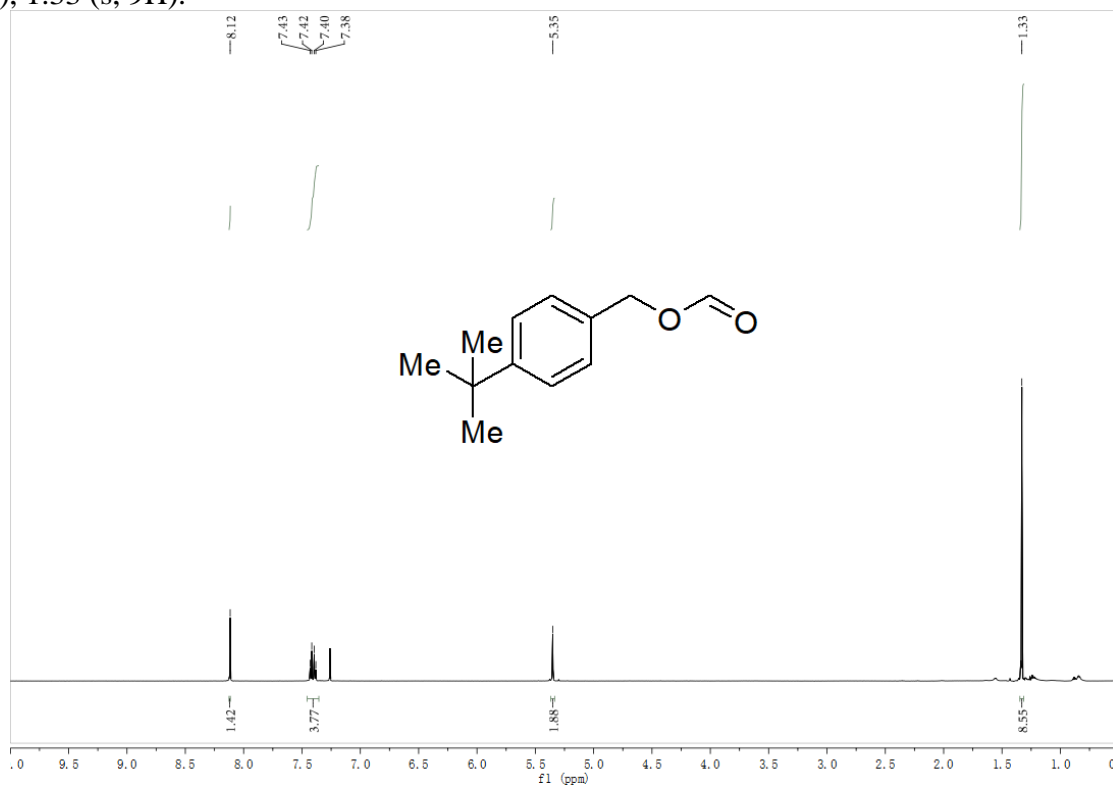
## Product 2e

$^1\text{H}$ NMR (600 MHz, CHLOROFORM-D)  $\delta$  8.13 (s, 1H), 7.25 (d,  $J = 48.7, 7.4$  Hz, 3H), 7.16 – 7.10 (m, 1H), 5.35 (s, 2H), 2.38 (s, 3H).



### Product 2f

$^1\text{H}$ NMR (600 MHz, CHLOROFORM-D)  $\delta$  8.12 (s, 1H), 7.41 (d,  $J = 21.0, 8.4$  Hz, 4H), 5.35 (s, 2H), 1.33 (s, 9H).



### Product 2g

$^1\text{H}$ NMR (600 MHz, CHLOROFORM-D)  $\delta$  8.13 (s, 1H), 7.40 (ddd,  $J = 25.2, 20.8, 7.2$  Hz, 5H), 5.38 (s, 2H).



## Product 2h

$^1\text{H NMR}$  (600 MHz, CHLOROFORM-D)  $\delta$  8.13 (s, 1H), 7.45 (d,  $J = 7.1$  Hz, 2H), 7.38 (t,  $J = 25.5, 7.2$  Hz, 3H), 5.38 (s, 2H).

