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# Facile synthesis of MOFs with uncoordinated carboxyl groups for selective CO<sub>2</sub> capture *via* postsynthetic covalent modification

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A postsynthetic covalent strategy involving dual-acyl chloride has been developed to introduce uncoordinated carboxyl groups into amine containing metal–organic frameworks (MOFs). The carboxyl group functionalized MOFs have been characterized by various techniques, including X-ray diffraction patterning, scanning electron microscopy, Fourier transform infrared spectroscopy, nuclear magnetic resonance, thermal gravimetric analysis, and gas adsorption. Results clearly indicated uncoordinated carboxyl groups were successfully grafted to the MIL-101(Cr)–NH<sub>2</sub> framework. In addition, most of the amine groups (>80%) were grafted with carboxyl groups, which indicates this method is very effective. The thermal stability and adsorption selectivity of CO<sub>2</sub>/N<sub>2</sub> were substantially enhanced, albeit the BET surface areas and total pore volumes were reduced. These observations could be explained by the effect of elimination of macropores in the framework due to the projecting of new functional groups in pore apertures. Here the successful fabrication of a MOF with uncoordinated carboxyl groups provides the possibility of efficiently modifying other MOFs.

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## Introduction

Metal–organic frameworks (MOFs), a new class of porous materials, are made by linking tunable inorganic and organic units.<sup>1</sup> In the last two decades, a particularly large number of MOFs have been successfully synthesized,<sup>2–7</sup> and the versatile characteristics of their chemical moieties have endowed MOFs with many special properties relevant to fuel cell,<sup>8</sup> catalysis<sup>9</sup> and biomedical applications.<sup>10</sup> Because of their ultrahigh porosity and the possibility of functionality, they play an important role in gas adsorption and separation.<sup>11–15</sup> For instance, MOFs are becoming one of the most promising candidates to reduce carbon dioxide (CO<sub>2</sub>) emissions, which is useful to resolve the increasingly severe global warming problem.<sup>13,16,17</sup> However, very limited MOFs show satisfactory CO<sub>2</sub> capture performance at low pressure and ambient temperature due to the weak interaction between CO<sub>2</sub> and the frameworks. In order to improve the adsorption performance, tremendous efforts have been devoted to functionalization of MOFs. To design MOFs with desired properties, the introduction of various functional groups into MOFs has attracted considerable interests. Postsynthetic covalent modification provides a variety of means to chemically modify the structures of MOFs,<sup>18–21</sup> among which the

direct grafting of functional groups on the organic linkers is effective, and has been widely applied.<sup>22–24</sup>

Interestingly, considerable reports on postsynthetic covalent modification involve the use of amine-contained MOFs, which is beneficial for enhancing the CO<sub>2</sub> capture performance. This is partly because of their convenient property to incorporate the 2-amino-1,4-benzenedicarboxylate into a number of BDC-derived MOFs. Amine groups in MOFs not only exhibit affinities with acidic gases molecules such as CO<sub>2</sub>, but also offer binding sites to accept functional groups. Previous studies have shown that amine groups in framework could be covalently modified by reagents such as peptide coupling,<sup>25</sup> acetic anhydride<sup>26</sup> and aldehyde.<sup>27,28</sup> Indeed, a wide range of chemical moieties has been successfully incorporated into MOFs *via* amine groups such as photo-switchable functional groups,<sup>29</sup> halo- and azo dye-functional groups<sup>22</sup> *et al.* Recently, Bai and co-workers have inserted –CONH– groups into rht-type MOFs through acylamide-bridging linkers by *in situ* solvothermal reaction.<sup>30,31</sup> Especially, uncoordinated carboxyl group functionalized MOFs exhibit promise for application in separation of transition metal ions,<sup>32</sup> catalysis<sup>33</sup> and sensing of ions.<sup>34</sup>

In a previous study, we obtained a type of amine-functionalized MIL-101(Cr) nanoparticles *via* a direct synthetic method.<sup>35</sup> The resulting MIL-101(Cr)–NH<sub>2</sub> possesses high surface area, good thermal stability, and selective adsorption of CO<sub>2</sub>.<sup>13,36</sup> In addition, it contains mesoporous cavities, which are likely to preserve sufficient remaining guest accessibility after functionalization with a bulky group. Inspired by the aforementioned advantages of MIL-101(Cr)–NH<sub>2</sub> and carboxyl group

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functionalized MOFs, it was selected in our project to explore the possibility of grafting uncoordinated carboxyl group *via* postsynthetic covalent modification in the present study. In this article, MIL-101(Cr)-NH<sub>2</sub> was covalently modified by two kinds of dual-acyl chlorides for the first time, under a mild condition and simple process. Results show that uncoordinated carboxyl groups are successfully introduced to MIL-101(Cr)-NH<sub>2</sub> framework. In addition, functionalized MIL-101(Cr)-NH<sub>2</sub> exhibited enhanced thermal stabilities, and higher CO<sub>2</sub>/N<sub>2</sub> ideal selectivities than the pristine MIL-101(Cr)-NH<sub>2</sub>. We anticipate that the proposed strategy is also applicable for the functionalization of other MOFs.

## Experimental

### Materials

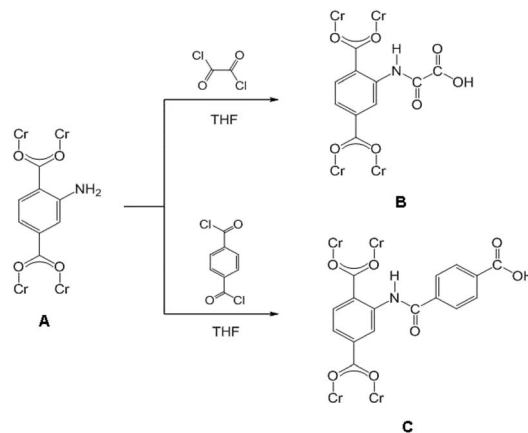
Chromium nitrate [Cr(NO<sub>3</sub>)<sub>3</sub>·9H<sub>2</sub>O, TCI], 2-aminoterephthalic acid (H<sub>2</sub>N-H<sub>2</sub>BDC, Sigma-Aldrich), sodium hydroxide (NaOH, Aladdin), *N,N*-dimethylformamide (DMF, TCI), tetrahydrofuran (THF, TCI), oxalyl chloride [(COCl)<sub>2</sub>, TCI], triethylamine (TEA, TCI), and terephthaloyl dichloride (C<sub>8</sub>H<sub>4</sub>Cl<sub>2</sub>O<sub>2</sub>, TCI) were used as received from vendors without further purification. Deionized water and THF were used as solvents.

### Synthesis of MIL-101(Cr)-NH<sub>2</sub>

MIL-101(Cr)-NH<sub>2</sub> was synthesized mainly according to the procedure reported in our previous work,<sup>35</sup> with an optimized amount of sodium hydroxide. Typically, 2-aminoterephthalic acid (H<sub>2</sub>N-H<sub>2</sub>BDC) (800 mg, 2 mmol), chromic nitrate (360 mg, 2 mmol) and sodium hydroxide (4 mmol) were dispersed in deionized water (15 mL) and ultrasonically mixed for 30 min. Subsequently, the solution was heated in a 50 mL Teflon-lined autoclave at 150 °C for 12 h under autogenous pressure, and then left to cool down to ambient temperature. The green products were collected by centrifugation and washed by DMF first to remove the most unreacted 2-aminoterephthalic acid on the surface, and then the obtained powder was cleaned by hot alcohol at 90 °C for 6 h twice in an autoclave to remove unreacted 2-aminoterephthalic acid in the pores. Finally, the products were dried at 90 °C under vacuum.

### Postsynthetic covalent modification

Instruments in the reactions were thoroughly dried prior to use. Firstly, MIL-101(Cr)-NH<sub>2</sub> (150 mg) powder was placed in a quartz tube and activated at 160 °C and 10<sup>-2</sup> kPa for 10 h to remove the remained guest molecules. Then the MIL-101(Cr)-NH<sub>2</sub> powder was gradually dispersed into the solution, containing dry tetrahydrofuran (THF) (8 mL) and oxalyl chloride (156 mg, 1.23 mmol) or terephthaloyl dichloride (250 mg, 1.23 mmol), under continuous fierce stirring. A certain amount of triethylamine (TEA) was dropwisely added into the mixture after 2 h, and further stirred overnight at ambient temperature. The precipitates were collected by centrifugation and then thoroughly washed by THF and water before drying at 100 °C under vacuum. The obtained samples were denoted as MIL-101(Cr)-amide-1 and MIL-101(Cr)-amide-2 to represent the treatment by



**Scheme 1** Schematic illustration of postsynthetic covalent modification of MIL-101(Cr)-NH<sub>2</sub> (A) using dual-acyl chloride. The resulting products: MIL-101(Cr)-amide-1 (B) and MIL-101(Cr)-amide-2 (C).

oxalyl chloride and terephthaloyl dichloride, respectively. All processes were carried out under anhydrous and argon atmosphere. The reaction process was illustrated as Scheme 1.

### Characterization

Powder X-ray diffraction (PXRD) patterns were obtained on a Bruker AXS D8 Advance diffractometer using Cu K $\alpha$  radiation at room temperature at a voltage of 40 kV and 40 mA. The morphologies of the as-prepared samples were examined using a field emission scanning electron microscope (FESEM, Hitachi, S-4800). Nuclear magnetic resonance (NMR) measurements were performed using a Bruker FT-NMR-spectrometer, Avance III (<sup>1</sup>H: 400 MHz). Thermal gravimetric analysis (TGA) was conducted in air from room temperature to 800 °C at a heating rate of 10 °C min<sup>-1</sup> using a system provided by Mettler Toledo (model TGA/DSC1) apparatus. Fourier transform infrared spectroscopy (FTIR) of the samples was recorded on KBr/sample pellets in a TENSOR 27 apparatus to determine the functional groups. The BET surface area was calculated over the range of relative pressures between 0.05 and 0.10 bar with a N<sub>2</sub> adsorption isotherm at 77.3 K (ASAP 2020, Micromeritics). Adsorption of CO<sub>2</sub> and N<sub>2</sub> at pressures up to 0.1 MPa were measured at 298 K by a standard volumetric analyzer (ASAP 2020, Micromeritics). Before each measurement, the samples were evacuated at 150 °C for 6 h to remove the guest molecules. All the gases used in the study had a minimum purity of 99.99%.

### Adsorption

The as-prepared functionalized MIL-101(Cr) samples were immediately measured after moving from the vacuum. Before each measurement, the sample was evacuated at 160 °C for 12 h. In the adsorption measurement, the apparatus first gave a desired pressure of the gas to the sample holder, and the pressure values were recorded in the sample holder to calculate the adsorption amount.

The adsorption isotherms of CO<sub>2</sub> and N<sub>2</sub> were measured using a volumetric apparatus (ASAP 2020, Micromeritics). The



relationship of adsorption amount *versus* gas bulk pressure was determined using the Langmuir–Freundlich<sup>37,38</sup> fit for the isotherms:

$$\frac{Q}{Q_m} = \frac{B \times P^{(1/t)}}{1 + B \times P^{(1/t)}}$$

here,  $Q$  represents the uptake amount at a certain equilibrium pressure ( $P$ ), with  $Q_m$  as the theoretical maximum. Both  $B$  and  $t$  are the equation constants. To obtain the accurate pressure  $P$ , corresponding to the same adsorption amount, the above equation can be rearranged to:

$$P = \left( \frac{Q/Q_m}{B - B \times Q/Q_m} \right)^t$$

The adsorption selectivities of  $\text{CO}_2/\text{N}_2$  ( $\alpha$ ) for all the samples were estimated by the following equation,<sup>39,40</sup> where  $Q_i$  is the

$$\alpha = \frac{Q_{\text{CO}_2}/P_{\text{CO}_2}}{Q_{\text{N}_2}/P_{\text{N}_2}}$$

adsorption amount of component  $i$ , at a partial pressure of component  $i$ ,  $P_i$ . The adsorption amount of the components is

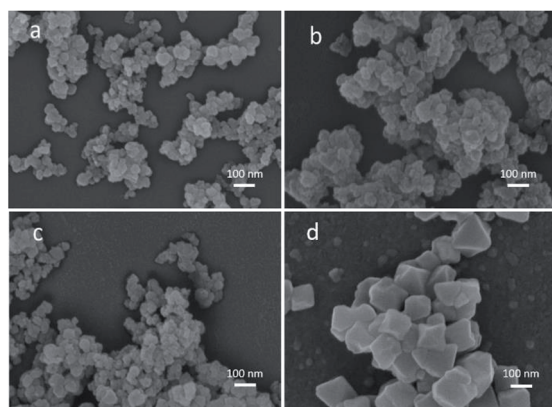
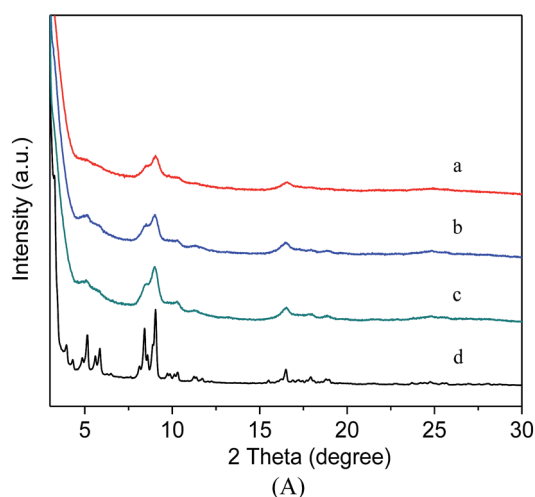


Fig. 1 Powder XRD patterns (A) and SEM images (B) of MIL-101(Cr)–NH<sub>2</sub> (a), MIL-101(Cr)–amide-1 (b), MIL-101(Cr)–amide-2 (c), MIL-101(Cr) (d).

the molar excess adsorption determined without converting to the absolute value.

## Results and discussion

Fig. 1A describes the PXRD results for the as-synthesized MIL-101(Cr), MIL-101(Cr)–NH<sub>2</sub> and the two samples prepared by postsynthetic covalent modification. It is evident that the XRD patterns of the three samples are similar to the characteristic peaks for the MIL-101(Cr), indicating high purity of our products,<sup>31</sup> thus the pore structures are preserved during the covalent modification process. Here the three modified samples also show broader and weaker Bragg reflections of the XRD patterns than the MIL-101(Cr), because the corresponding particle sizes are smaller than the latter.<sup>41</sup> These observations are comparatively verified in the SEM images. As shown in Fig. 1B, the morphologies of MIL-101(Cr)–NH<sub>2</sub> remain nearly intact after the functional incorporation, while the crystal size is very small, much smaller than that of MIL-101(Cr).

To further ensure the successful incorporation of the MIL-101(Cr)–NH<sub>2</sub> framework, IR spectra was collected on the as-synthesized MIL-101(Cr)–NH<sub>2</sub> and products by postsynthetic covalent modification. For MIL-101(Cr)–NH<sub>2</sub>, the double peaks at 3490 cm<sup>−1</sup> and 3363 cm<sup>−1</sup> are clearly observed (Fig. 2), which are assigned to the asymmetrical and symmetrical stretching vibration of the amine groups. In the lower frequency region, the two peaks around 1593 cm<sup>−1</sup> correspond to the N–H bending vibration and the C–N stretching vibrations (1340 cm<sup>−1</sup> and 1259 cm<sup>−1</sup>) of aromatic amines, respectively.<sup>29</sup> Besides, the incorporation of amino-dicarboxylate within the structure is confirmed by the characteristic –(O–C–O)– stretching vibrations at 1500 cm<sup>−1</sup> and 1430 cm<sup>−1</sup>.<sup>42</sup> As for MIL-101(Cr)–amide-1 and MIL-101(Cr)–amide-2, three additional signals including N–H stretching vibration (3310 cm<sup>−1</sup> and 3313 cm<sup>−1</sup>) of amide groups, C=O stretching vibration (1700 cm<sup>−1</sup> and 1710 cm<sup>−1</sup>) and C–N stretching vibration (1271 cm<sup>−1</sup> and 1281 cm<sup>−1</sup>) are directly observed. Here the IR spectra of the samples have validated the

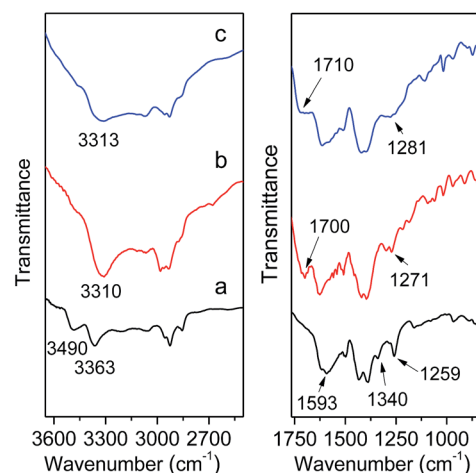


Fig. 2 IR spectra of MIL-101(Cr)–NH<sub>2</sub> (a), MIL-101(Cr)–amide-1 (b), MIL-101(Cr)–amide-2 (c).



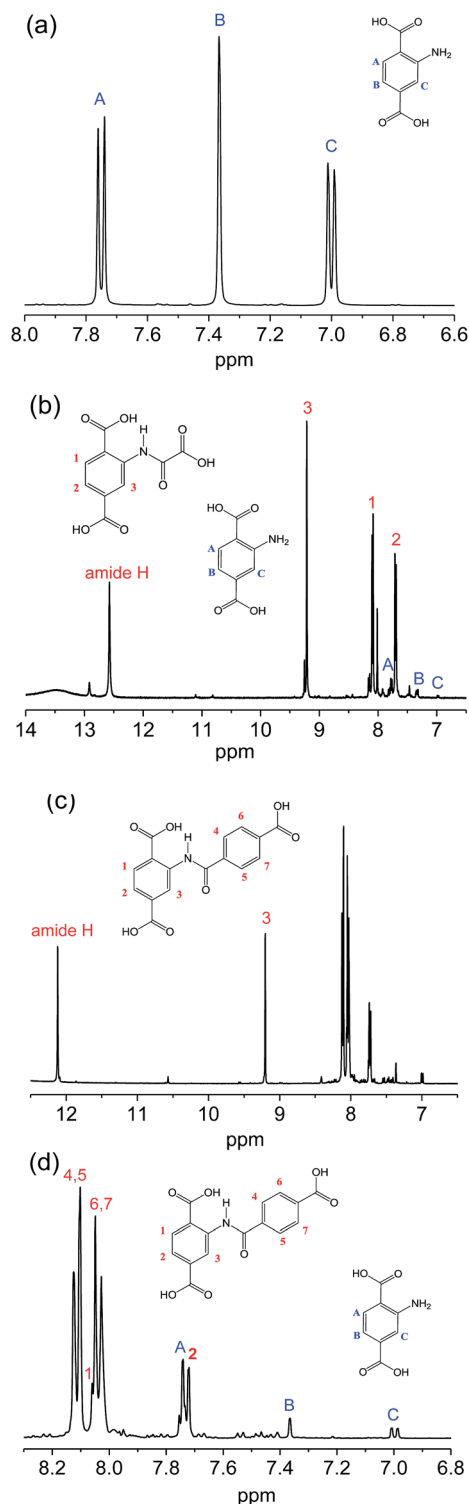


Fig. 3  $^1\text{H}$ -NMR spectra of 2-aminoterephthalic acid (a), the digested products from MIL-101(Cr)-amide-1 (b), and MIL-101(Cr)-amide-2 (c and d).

presence of amide and carboxyl groups in the MIL-101(Cr)-amide-1 and MIL-101(Cr)-amide-2 samples, respectively.

Meanwhile, NMR-measurements were also conducted to inspect the linker structures of the original and the modified

samples. Fig. 3 illustrates the  $^1\text{H}$ -NMR spectra of 2-aminoterephthalic acid ( $\text{H}_2\text{N-H}_2\text{BDC}$ ) and the digested products from the two samples. For  $\text{H}_2\text{N-H}_2\text{BDC}$  in Fig. 3a, signals at 7.75, 7.37 and 7.00 ppm are assigned to protons A, B and C on the benzene. For  $\text{H}_2\text{BDC-amide-1}$  in Fig. 3b, the signals are mainly lumped together, and the characteristic peaks for  $\text{H}_2\text{N-H}_2\text{BDC}$  is hardly seen. The peak at 12.57 ppm is assigned to the amide proton, with the signals at 9.23, 8.09, 7.70 ppm for the proton 3, 1 and 2 (numbering in the figure), respectively. Similarly, Fig. 3c and d describe the majority of characteristic information for  $\text{H}_2\text{BDC-amide-2}$ , despite having few weak signals due to the presence of unreacted  $\text{H}_2\text{N-H}_2\text{BDC}$ . The signal at 12.12 ppm and 9.20 ppm are produced by the amide proton and the proton 3 (numbering in the figure), respectively, and the other protons are marked in the region of 8.2 ppm to 7.6 ppm. The conversion rates of amine groups calculate from the integral data are above 97% and nearly 87% for MIL-101(Cr)-amide-1 and MIL-101(Cr)-amide-2, respectively, implying the high conversion rate of the amino groups in our work. On the contrary, the conversion rates of amine in IRMOF-3 and MIL-68(In)- $\text{NH}_2$  are less than 13% and 10% according to the research, respectively.<sup>27,43</sup> Cohen and co-workers have reported that the rate for the covalent modification of IRMOF-3, UMCM-1- $\text{NH}_2$  and DMOF-1- $\text{NH}_2$  are around 32–76%.<sup>44</sup> Therefore, dual-acyl chloride is very suitable for the amine functionalization of MOFs modified by postsynthetic covalent. In conclusion, the IR and  $^1\text{H}$ -NMR data have successfully verified the accomplishment of covalent modification of MIL-101(Cr)- $\text{NH}_2$  by reacting with dual-acyl chloride.

Moreover, the pore networks of the samples are explored to elucidate a better understanding of their adsorption mechanism.

Fig. 4 plot the adsorption-desorption loops of the three samples by  $\text{N}_2$ , and a type I adsorption behavior occurred, with small hysteresises due to the interstitial voids formed by the small crystal particles. Comparing with MIL-101(Cr)- $\text{NH}_2$ , BET surface areas and total pore volumes are substantially reduced in both MIL-101(Cr)-amide-1 and MIL-101(Cr)-amide-2, due to the presence of carboxyl groups projecting into the pores. The

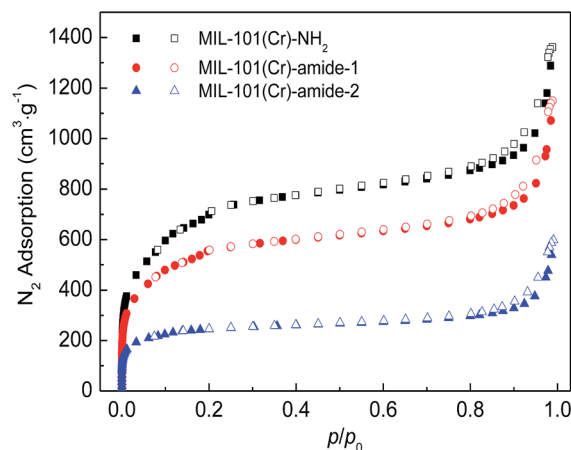


Fig. 4  $\text{N}_2$  adsorption-desorption isotherms on MIL-101(Cr)- $\text{NH}_2$ , MIL-101(Cr)-amide-1 and MIL-101(Cr)-amide-2 samples at 77.3 K.





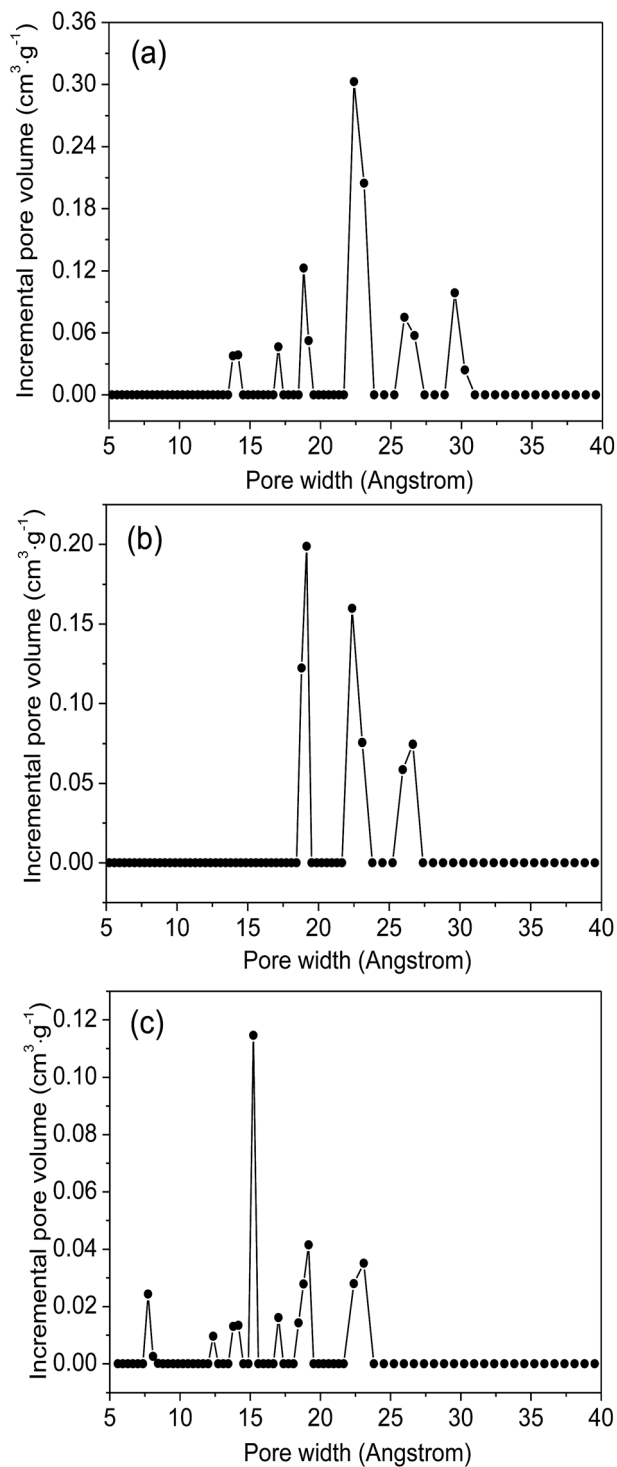


Fig. 5 NLDFT pore size distribution of MIL-101(Cr)-NH<sub>2</sub> (a), MIL-101(Cr)-amide-1 (b) and MIL-101(Cr)-amide-2 (c).

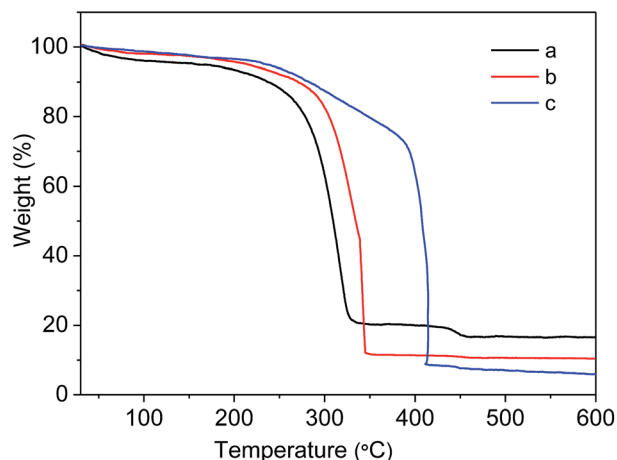


Fig. 6 TGA curves: (a) MIL-101(Cr)-NH<sub>2</sub>, (b) MIL-101(Cr)-amide-1 and (c) MIL-101(Cr)-amide-2.

observed decrements of surface areas have been also found in many other studies for various MOFs modified by postsynthetic covalent.<sup>45</sup> Fig. 5 provides the pore size distributions of the samples based on the interpretation of nonlinear density functional theory (NLDFT), and the results implies that the average pore sizes of MIL-101(Cr)-amide-1 and MIL-101(Cr)-amide-2 become smaller than that of MIL-101(Cr)-NH<sub>2</sub>. It is evident that such a result is caused by the elimination of the larger pore in MIL-101(Cr)-amide-1 and MIL-101(Cr)-amide-2. On the other hand, despite two smallest pores are also eliminated in MIL-101(Cr)-amide-1, some pores are newly-created in MIL-101(Cr)-amide-2, locating at 7.7 Å, 12.4 Å and 15.2 Å, and this is due to the partial occupation of larger groups. The results of the pore parameters are outlined in Table 1. It have found that the surface area and pore volume of MIL-101(Cr) with functional groups decreased with the increases in group sizes. Further, the proportion of micropores agree with the size variation of the functional groups, and this is important to enhance the CO<sub>2</sub>/N<sub>2</sub> selectivity for its smaller kinetic size than N<sub>2</sub>.

To evaluate the stability of as-prepared MIL-101(Cr)-NH<sub>2</sub> and the samples under postsynthetic covalent modification, the thermal gravimetric analysis was carried out. As shown in Fig. 6, MIL-101(Cr)-NH<sub>2</sub> starts to decompose at 250 °C in the air, similar to the discovery reported in the literature.<sup>35</sup> In contrast, the decomposing temperatures of MIL-101(Cr)-amide-1 and MIL-101(Cr)-amide-2 increase up to 280 °C and 380 °C, respectively, which is slightly lower than that of the value for ZIF-8 (*ca.* 500 °C), but higher than most of MOF materials.<sup>46</sup> It have reported that the thermal stability of MOFs is improved by reducing the primary amine to a secondary one,<sup>28</sup> thus the

Table 1 Pore parameters of MIL-101(Cr)-NH<sub>2</sub>, MIL-101(Cr)-amide-1 and MIL-101(Cr)-amide-2 samples

Sample	$S_{\text{BET}}$ (m <sup>2</sup> g <sup>-1</sup> )	$V_{\text{total}}$ (cm <sup>3</sup> g <sup>-1</sup> )	$V_{\text{micro}}$ (cm <sup>3</sup> g <sup>-1</sup> )	$V_{\text{micro}}/V_{\text{total}}$
MIL-101(Cr)-NH <sub>2</sub>	2558	2.11	0.18	8.53%
MIL-101(Cr)-amide-1	2005	1.78	0.16	8.99%
MIL-101(Cr)-amide-2	861	0.93	0.20	21.51%



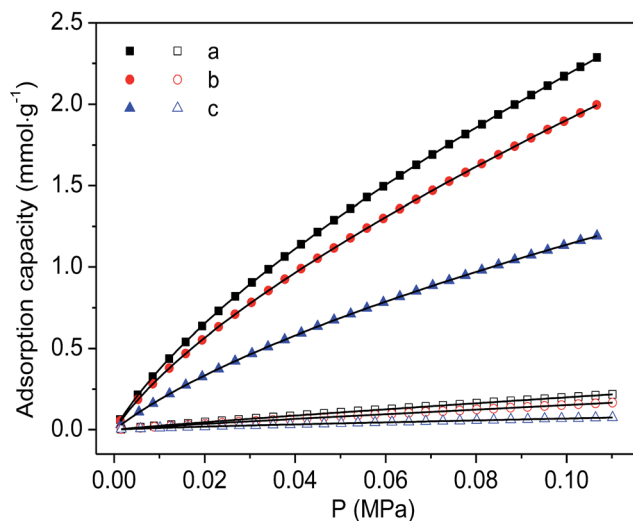


Fig. 7 Adsorption isotherms of CO<sub>2</sub> (solid) and N<sub>2</sub> (hollow) on the samples at 298 K and the fitted isotherms by DSL model: (a) MIL-101(Cr)-NH<sub>2</sub>, (b) MIL-101(Cr)-amide-1 and (c) MIL-101(Cr)-amide-2.

incorporation of larger groups in the framework favors the improvement of the thermal stability.

Finally, the CO<sub>2</sub> and N<sub>2</sub> adsorption isotherms were conducted at a temperature of 298 K to evaluate the adsorption capability of the as-prepared functionalized MIL-101(Cr) materials. As shown in Fig. 7, although the CO<sub>2</sub> adsorption amounts of the two modified samples are consistently smaller than the corresponding adsorption of MIL-101(Cr)-NH<sub>2</sub> within the entire pressure range, relatively higher ideal selectivities of CO<sub>2</sub> over N<sub>2</sub> are produced by the incorporation of amide and carboxyl groups (Fig. 8). Specifically, the ideal selectivities at 0.1 MPa are up to 10.93, 12.49 and 16.44 for MIL-101(Cr)-NH<sub>2</sub>, MIL-101(Cr)-amide-1 and MIL-101(Cr)-amide-2, respectively. Amide groups have been previously reported to enhance CO<sub>2</sub> adsorption.<sup>31</sup>

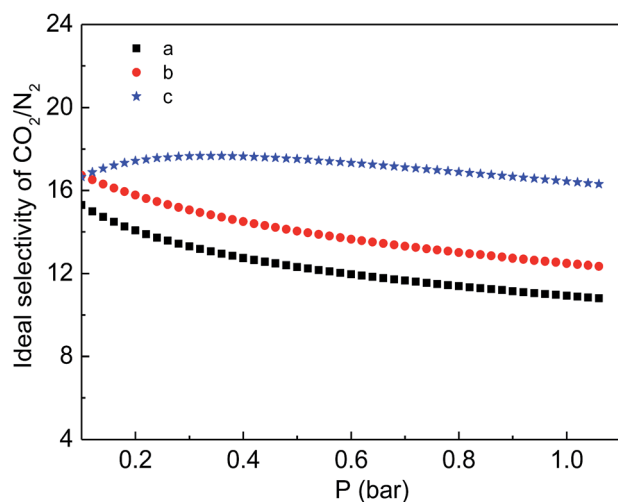


Fig. 8 Ideal separation factor of CO<sub>2</sub>/N<sub>2</sub> for the samples at 298 K: (a) MIL-101(Cr)-NH<sub>2</sub>, (b) MIL-101(Cr)-amide-1 and (c) MIL-101(Cr)-amide-2.

Further, the incorporation of carboxyl groups into the MOFs is also beneficial to the CO<sub>2</sub> capture, and the performance is even better than the results achieved by amine groups,<sup>47</sup> due to a higher heat of adsorption for carboxyl groups, thus a higher efficiency of adsorption behavior at low pressure occurred. Herein the high selectivities in MIL-101(Cr)-amide-1 and MIL-101(Cr)-amide-2 are physically caused by the strong interactions between the modified groups and CO<sub>2</sub> molecules. Moreover, since the quadrupole moment of N<sub>2</sub> ( $-4.6 \times 10^{-40} \text{ C m}^2$ ) is much lower than the value of CO<sub>2</sub> ( $-14 \times 10^{-40} \text{ C m}^2$ ),<sup>48</sup> the enhanced polarizability (electrostatic interaction) of the framework caused by the incorporated groups is also an important factor for the selectivity enhancement.<sup>49</sup> Thus, all of these factors contributes to a higher uptake of CO<sub>2</sub> than of N<sub>2</sub>.

## Conclusions

In summary, uncoordinated carboxyl groups were successfully grafted on MIL-101(Cr)-NH<sub>2</sub> framework, and a general and facile method was developed to modify MOFs by dual-acyl chloride at mild conditions. In addition, most of the amine groups of MIL-101(Cr)-NH<sub>2</sub> were successfully grafted with carboxyl groups in the present work, indicating the proposed strategy is very efficient. The carboxyl groups functionalized samples exhibited enhanced selectivities of CO<sub>2</sub>/N<sub>2</sub> in comparison with the original MIL-101(Cr)-NH<sub>2</sub> at low pressure and ambient temperature. Besides, the thermal stabilities of the functionalized samples were also enhanced. This could be caused by the positive evolution of pore due to the introducing of carboxyl groups into the pores apertures. The uncoordinated carboxyl groups functionalized samples will be used for fabrication of MOF-based mixed matrix membrane and separation of transition metal ions, which are currently underway in our laboratory. Overall, the present work provides a high efficient strategy to functionalize amine-containing MOF with uncoordinated carboxyl groups.

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