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## Improving MOF stability: approaches and applications

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Metal–organic frameworks (MOFs) have been recognized as one of the most important classes of porous materials due to their unique attributes and chemical versatility. Unfortunately, some MOFs suffer from the drawback of relatively poor stability, which would limit their practical applications. In the recent past, great efforts have been invested in developing strategies to improve the stability of MOFs. In general, stable MOFs possess potential toward a broader range of applications. In this review, we summarize recent advances in the design and synthesis of stable MOFs and MOF-based materials *via de novo* synthesis and/or post-synthetic structural processing. Also, the relationships between the stability and functional applications of MOFs are highlighted, and finally, the subsisting challenges and the directions that future research in this field may take have been indicated.

### 1. Introduction

Over the past two decades, the emergence of metal–organic frameworks (MOFs) which are constructed by metal ions/clusters and multidentate organic linkers *via* coordination bonds (reticular synthesis) has significantly enriched the domain of

porous materials.<sup>1</sup> Based on the flexibility of the constituents' geometry, size, and functionality, MOFs with a crystalline nature generally possess extremely high surface areas (typically ranging from 1000 to 10 000 m<sup>2</sup> g<sup>−1</sup>) and tunable pore sizes/characteristics.<sup>1</sup> These aspects endow MOFs with fantastic functionalities/properties for a variety of applications, for example, gas adsorption and separation, catalysis, sensors, drug delivery, proton conduction, *etc.*<sup>1–18</sup> For various applications of MOFs, stability, mainly including chemical, thermal and mechanical stability, is one of the major concerns. The chemical stability of MOFs, specifically, refers to their resistance to the effects of exposure to various chemicals in their environment, *e.g.* moisture, solvents, acids, bases, and aqueous

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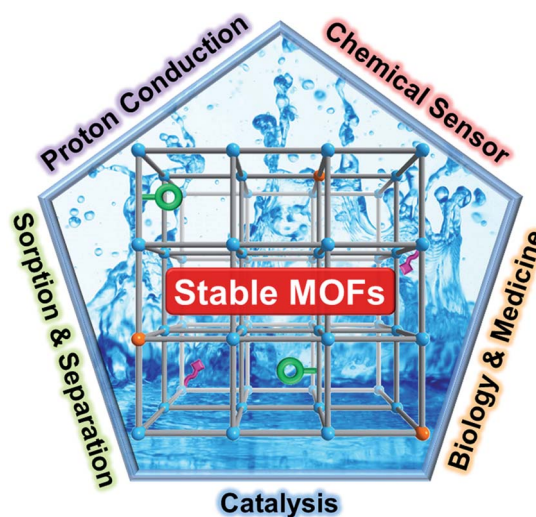


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Over the past two decades, various promising methods have been developed to tune MOF stability, especially chemical stability, which might be the most important and significant prerequisite of MOFs for their diverse applications.<sup>19,37–39</sup> Therefore, we believe that it is a good time for us to provide a systematic review of recent advances in the preparation of stable MOFs and their applications (Scheme 1). Our discussions begin with a summary of strategies that have been developed for improving the chemical stability of MOFs while emphasizing the key factors in the synthesis of stable MOFs and MOF-based materials. Next, the unique attributes of stable MOFs and their extensive applications, including adsorption and separation, heterogeneous catalysis, fluorescence sensing, biological and medical application, and proton conductivity, are introduced. At the end of the review, a summary and proposed future developments in this research field will be presented. Hopefully, this review will inspire the enthusiasm of scientists in chemistry and materials science and interdisciplinary researchers toward this field.

Although many MOFs are vulnerable to structural destruction even in an ambient atmospheric environment due to the lability of coordination bonds between metal ions and ligands, a growing number of MOFs with exceptional chemical stability have been reported in recent years. In general, two main types of approaches have been adopted to improve the chemical stability of MOFs. One is the *de novo* synthesis of stable unknown MOFs and the other is to improve the stability of existing MOFs. The chemical stabilities of some representative MOFs are summarized in Table 1.



The development of stable MOFs can be divided into several important stages and that began with the advent of the first porous MOF (MOF-5).<sup>27</sup> In the early stage, the study of MOF stability was mainly focused on their thermal and mechanical stability.<sup>21,28–35</sup> With the in-depth understanding of the framework collapse of MOFs in air,<sup>36</sup> the development of stable MOFs entered the second stage. Numerous water-stable MOFs were reported based on their structural designability and chemical tunability. In the third stage, different techniques of crystal growth and crystal structure determination gave rise to a large number of extraordinarily stable MOFs, which effectively

MOFs	Metal	Ligands	Testing condition <sup>a</sup>	Time	Ref.
MAF-X27-Cl	Co(II)	bbta	0.001 M HCl, 1.0 M KOH	1 week	88
H <sub>3</sub> [(Cu <sub>4</sub> Cl) <sub>3</sub> -(BTtri) <sub>8</sub> ]	Cu(II)	BTtri	Boiling water	3 days	87
FJI-H14		BTta	pH = 3	1 day	
BUT-155		tdhb	pH = 2–12 (373 K)	24 h	95
USTC-6		4,4'-(Hexafluoroisopropylidene) diphthalicanhydride	pH = 4–10, boiling water	24 h	97
Ni <sub>3</sub> (BTP) <sub>2</sub>	Ni(II)	BTP	pH = 2–10	7 days	109
PCN-601		TPP	Water	1 month	
PCN-602(Ni)		TPPP	Boiling solutions of pH = 2–14	14 days	83
ZIF-8	Zn(II)	MeIM	Saturated NaOH (100 °C)	24 h	25
USTC-7		TZBPDC	0.1 mM HCl (RT)		
oCB-MOF-1			1 M Na <sub>2</sub> CO <sub>3</sub> , K <sub>3</sub> PO <sub>4</sub> , KF, pH = 4–14	24 h	84
MIL-53(Al)			Boiling benzene, methanol, and water	7 days	79
NH <sub>2</sub> -MIL-53(Al)			8 M NaOH (100 °C)	24 h	
Al-TCPP			Boiling organic solvents, pH = 2–12 (RT)	12 h	94
PCN-333(Al)			Water (RT)	4 months	
467-MOF			pH = 2–12 (RT)	15 h	108
			pH = 4–12	3 days	44
				2 months	
			pH = 5–8 (RT)	1 week	49
			pH = 3–9	Overnight	50
			pH = 0–13 (RT)	36 h	74
			Water (RT)	30 days	
			Boiling water	Over 1 week	
AITCS-1			pH = 1–11, aqua regia solution	24 h	75
MIL-53(Cr)	Cr(III)	BDC	Solution of 7.0 × 10 <sup>-2</sup> M NaOH	6 h	21
			Solution of 7.0 × 10 <sup>-2</sup> M HCl	48 h	
MIL-101(Cr)			pH = 0–12	2 months	44
MIL-100(Cr)			Water	12 months	64
PCN-333(Fe)	Fe(III)	BTC	pH = 3–9	Overnight	50
PCN-600(Fe)		TATB	pH = 2–11	24 h	61
MIL-100(Fe)		TCPP-Fe	pH = 2–11	24 h	61
PCN-250(Fe <sub>2</sub> Co)	Fe(II,III), Co(II)	BTC	pH = 1–10	7 days	65
		BTB, 4,4'-(diazene-1,2-diyl) dibenzoate	Water	6 months	60
			pH = 1–12	24 h	
			pH = 3.5–10 (373 K)	24 h	106
[(CH <sub>3</sub> ) <sub>2</sub> NH <sub>2</sub> ] <sub>2</sub> [Eu <sub>6</sub> (μ <sub>3</sub> -OH) <sub>8</sub> (NDC) <sub>6</sub> (H <sub>2</sub> O) <sub>6</sub> ]	Eu(III)	NDC			
[La(TTCA) (H <sub>2</sub> O)]·DMF·H <sub>2</sub> O	La(III)	TTCA	Boiling water, pH = 1–13	7 days	52
[La(BTB)H <sub>2</sub> O]		BTB	pH = 2–14 (100 °C)	3 days	54
USTC-8(In)	In(III)	TCPP	pH = 0–11	12 h	76
MIL-177-LT	Ti(IV)	mdip	Water (RT)	30 days	66
			Boiling water	24 h	
			Conc. HCl, HNO <sub>3</sub> , H <sub>2</sub> SO <sub>4</sub> and HPO <sub>4</sub> , aqua regia (RT)	7 days	
UiO-66	Zr(IV)	BDC	pH = 0–12	2 months	44
PCN-777		TATB	pH = 3–11	12 h	51
DUT-98		CPCDC	Boiling water or solutions of 2–6 M HCl (298 K)	24 h	53
PCN-222(Fe)		TCPP-Fe	Boiling water, conc. HCl	24 h	55
PCN-223		TCPP	pH = 0–10	24 h	56
PCN-224(Ni)		TCPP-Ni	pH = 0–11	24 h	57
PCN-225		TCPP	pH = 1–11	12 h	58
PCN-228		TCP-1	pH = 0–12	24 h	59
PCN-229		TCP-2			
PCN-230		TCP-3			
BUT-12		CTTA	Conc. HCl, pH = 10	24 h	63
BUT-13		TTNA			
NUPF-1		4,4',4'',4'''-((4,4',4'',4'''-(Porphyrin-5,10,15,20-tetrayl) tetrakis (benzoyl))tetrakis(azanediyl)) tetrabenzoic acid	12 M HCl, pH = 9	3 days	67
MOF-808		BTC	Boiling water, conc. HCl, pH = 10	24 h	68



Table 1 (Contd.)

MOFs	Metal	Ligands	Testing condition <sup>a</sup>	Time	Ref.
MIP-202(Zr)		Aspartic acid	Boiling water, pH = 0–12	—	73
Zr(H <sub>4</sub> L <sup>1</sup> )		Tetraphenylsilane tetrakis-4-phosphonic acid (H <sub>8</sub> L <sup>1</sup> )	pH = 12 (100 °C), conc. HNO <sub>3</sub> (100 °C)	7 days	77
ZrPP-1		THPP	0.1 M HCl to 20 M NaOH	7 days	78
UiO-67-o-2CF <sub>3</sub>		BPDC-o-2CF <sub>3</sub>	Boiling water, 8 M HCl, 50 ppm NaF aqueous solution	24 h	100

<sup>a</sup> RT: room temperature. RH: relative humidity. Conc.: concentrated.

## 2.1 Governing MOFs towards high stability *via de novo* synthesis

It is widely believed that the degradation of MOFs generally has two aspects, namely, the metal–ligand bond breaking, and the formation of products that are more stable as compared with pristine MOFs.<sup>40,41</sup> Hence, the chemical stability of MOFs depends strongly on the intrinsic structures of MOFs (internal factors), including the charge density of metal ions, connection numbers of metal ions/clusters, basicity and configuration as well as hydrophobicity of ligands, *etc.* Following these considerations, judicious selection of metal nodes and design of linkers enable the construction of MOFs with sound stability.<sup>42</sup>

**2.1.1 High-valent metal-containing MOFs.** Based on the HSAB principle, with the same coordination environments, high-valent metals with high charge density (hard acids), including Zr<sup>4+</sup>, Cr<sup>3+</sup>, Al<sup>3+</sup>, Fe<sup>3+</sup>, *etc.*, tend to coordinate with O donor ligands (hard bases) (Fig. 1) to form MOFs with strong coordination bonding, thus usually presenting good chemical stability. Despite the relatively low pK<sub>a</sub> of carboxylate linkers, these high-valent metal ions often require more linkers to balance the charges. This can lead to a high connection number

of metal clusters and further enhance the stability of the resultant MOFs. In addition, the strong coordination bonds and the low pK<sub>a</sub> of carboxylate linkers endow the high-valent metal ion-containing MOFs with a decent level of acidic stability.

As an early example, Férey and co-workers fabricated a Cr(III)-based MOF, MIL-101(Cr), with very large pore sizes (2.9 and 3.4 nm) and Brunauer–Emmett–Teller (BET) surface area (~4000 m<sup>2</sup> g<sup>−1</sup>) *via* the combination of targeted chemistry and computational design.<sup>43</sup> By virtue of the inertness of Cr–carboxylate oxygen bonds, MIL-101(Cr) is highly resistant to the attack of acid and alkali, making it stable in aqueous solutions of pH = 0–12 for 2 months.<sup>44</sup> Using a similar strategy, a variety of high-valent metal-containing MOFs with high chemical stability have been reported, including MIL-100(Cr), MIL-53(Cr), UiO series, MIL-125, Al-TCPP, PCN series, *etc.*<sup>22,45–68</sup> However, the strong M–O bonds often lead to the poor crystallinity and the small sizes of resultant MOF particles. This increases the difficulty of structural characterization with single-crystal X-ray diffraction. For this reason, most of the crystal structures of the above-mentioned MOFs are determined by using powder X-ray diffraction.



Fig. 1 Some representative O donor ligands, with their pK<sub>a</sub>s referred in this text.





To address this issue, Behrens's group developed a modulated synthesis strategy to regulate the size of Zr-MOFs from nanosized crystallites to large single crystals, and reported the first Zr-MOF single crystal that was sufficiently large for single-crystal X-ray diffraction.<sup>69</sup> By adjusting the concentration of the modulator (*i.e.* monocarboxylic acid) in the synthetic system, the nucleation rate of MOFs can be controlled, which ultimately leads to the formation of highly crystalline products or even large single crystals.<sup>69,70</sup> With this in mind, Zhou's group employed Fe-tetrakis(4-carboxyphenyl)porphyrin,  $\text{Zr}^{4+}$  and benzoic acid separately as the ligand, metal ion and modulator to yield needle-shaped single crystals, PCN-222(Fe), also called MOF-545 and MPPF-6.<sup>55,71,72</sup> The stability experiments showed that this MOF was able to survive even in concentrated aqueous HCl solution for 24 h due to the strong Zr–O bonds in the 8-connected Zr-clusters and the chelating effect between Fe(III) and porphyrin. A similar phenomenon was observed in other PCN-series reported by the same team.<sup>50,56–62</sup> In fact, monocarboxylic acids can also work as solvents in the preparation of stable Zr(IV)-based MOFs. The common solvent in MOF synthesis, dimethylformamide (DMF), was replaced by a mixture of monocarboxylic acid (formic acid) and acetic anhydride to afford a large-pore MOF, MIP-200, based on  $\text{Zr}^{4+}$  and 3,3',5,5'-tetracarboxydiphenylmethane ( $\text{H}_4\text{mdip}$ ).<sup>73</sup> Remarkably, MIP-200 exhibited excellent stability under extremely harsh conditions, including concentrated strong acids (HCl and  $\text{HNO}_3$ ), highly concentrated  $\text{H}_3\text{PO}_4$  and  $\text{H}_2\text{SO}_4$ , and  $\text{NH}_4\text{OH}$  vapor. The modulated synthesis strategy applies equally to trivalent metal-based MOFs. Using formic acid as the modulator, Du's group successfully fabricated an Al-based MOF (467-MOF) based on a flexible ligand (4,4',4''-[benzene-1,3,5-triyl-tris(oxy)]tribenzoic acid,  $\text{H}_3\text{BTB}$ ).<sup>74</sup> Chemical stability investigation for the 467-MOF indicated that it remained stable in aqueous solutions of HCl or NaOH of a broad pH range from 1 to 11. Similarly, with the aid of hydrofluoric acid, large and good-quality single crystals of Al-MOF, ALTCS-1, were obtained based on tetrakis(4-oxycarbonylphenyl)silane.<sup>75</sup> Strikingly, ALTCS-1 could be stable in both aqueous solutions of pH 1 to 11 and aqua regia solution for at least 24 h. Other  $\text{M}^{3+}$  ions like  $\text{In}^{3+}$  can also form stable MOFs with O donor linkers. Recently, Jiang's group synthesized an unusual out-of-plane (OOP) porphyrin-based MOF, USTC-8(In), by employing  $\text{In}(\text{OH})_3$  and  $\text{HNO}_3$  separately as the metal precursor and modulator.<sup>76</sup> Due to the strong coordination bonds between  $\text{In}^{3+}$  ions and porphyrin carboxylate ligands, USTC-8(In) could remain intact in aqueous solutions with pH values ranging from 2 to 11 for 12 h.

Note that high-valent metal-containing MOFs with remarkable chemical stability include not only carboxylate-based MOFs, but also those with phosphonate and phenolate-based frameworks. Sun's group selected tetraphenylsilane tetrakis-4-phosphonic acid as the ligand and constructed a zirconium-phosphonate network that displayed distinctive tolerance to aqueous solutions with a wide range of pH values (1–12) and concentrated acids.<sup>77</sup> As for phenolate-based MOFs, the relatively high  $\text{p}K_{\text{a}}$  of the phenolate ligands can afford stronger M–O bonds with  $\text{Zr}^{4+}$  ions in comparison to carboxylate linkers. Lin's

group demonstrated the fabrication of a series of zirconium polyphenolate-(metallo)porphyrin MOFs, in which the highly stable 5-membered rings generated by the chelating effect of polyphenols greatly improved the stability of resultant MOFs (Fig. 2a).<sup>78</sup> Remarkably, the prototypic ZrPP-1 not only remained intact in strong acid (HCl, pH = 1) but also presented high robustness in saturated NaOH solution for 168 h (Fig. 2b), offering an ideal platform for diverse applications.

To summarize, the basic principle for the fabrication of stable high-valent metal-containing MOFs is the strong coordination bond, high coordination number of metal ions, inertness of metal and others, which enhance the MOF tolerance of hydrolysis before framework collapse occurs.

**2.1.2 Low-valent metal containing MOFs.** Apart from high-valent metal ions, low-valent metal ions, including  $\text{Zn}^{2+}$ ,  $\text{Co}^{2+}$ ,  $\text{Ni}^{2+}$ ,  $\text{Fe}^{2+}$ , and  $\text{Ag}^+$ , can be considered as soft acids to construct highly stable MOFs with suitable N-containing linkers (soft bases) (Fig. 3). The high  $\text{p}K_{\text{a}}$  of azoles and the strong coordination bonds usually endow these MOFs with remarkable stability in basic solutions. In general, the higher the  $\text{p}K_{\text{a}}$  for the N donor ligand, the more stable the resultant MOFs under humid conditions.

As a representative example, zeolitic imidazolate framework (ZIF)-8 (also called MAF-4) which possesses a zeolite-type topology composed of four-coordinated  $\text{Zn}^{2+}$  ions linked by imidazolate linkers is very stable in an aqueous environment.<sup>79,80</sup> In particular, the structure of ZIF-8 can be well maintained even in 8 M aqueous NaOH for 24 h at 100 °C, indicating the exceptional stability of this framework under alkaline conditions. A similar stability can be observed also in other Zn-based ZIF series like ZIF-68, -69, and -70.<sup>81</sup> Not limited to imidazoles, reacting the pyrazole-derived ligand with suitable low-valent metal salts can also lead to the formation of MOFs with high chemical stability. Volkmer's group deliberately selected 1,4-bis[(3,5-dimethyl)-pyrazol-4-yl]benzene ( $\text{H}_2\text{bdpb}$ ) and Co(II) salts to fabricate a hydrolytically stable MOF (MFU-1), which is isostructural with MOF-5.<sup>82</sup> The relatively strong bonds in the  $\{\text{CoON}_3\}$  coordination units of MFU-1 give rise to the decent thermodynamical stability of the framework under

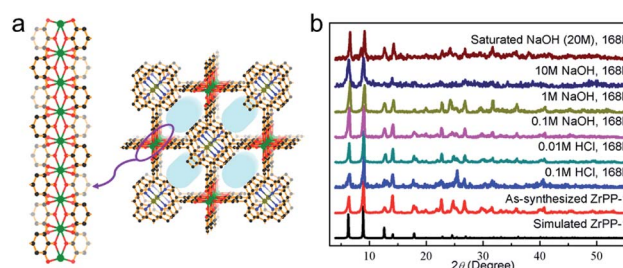


Fig. 2 (a) The structure of ZrPP-1 constructed from  $\text{Zr}^{4+}$  and 5,10,15,20-tetrakis(3,4,5-trihydroxyphenyl)porphyrin with the view of the  $\text{Zr}^{\text{IV}}$ -pyrogallate (=1,2,3-trioxobenzene) coordination chain. Color code: C, black; O, red; N, blue; Zr, green; and metal (in the porphyrin core), olive green. H atoms are omitted for clarity. (b) Powder X-ray diffraction (PXRD) patterns of ZrPP-1 under different conditions. Adapted from ref. 78 with permission from Wiley-VCH, copyright 2018.





Fig. 3 Some representative N donor ligands, with their  $pK_a$ s referred in this text.

ambient conditions for over 6 months. Similarly, using the reaction between the pyrazol-type ligand (1,3,5-tris(1*H*-pyrazol-4-yl)benzene,  $H_3BTP$ ) and  $Ni^{2+}$  ions, an ultrastable framework,  $Ni_3(BTP)_2$ , was obtained and the chemical resistance experiments indicated that it was stable in boiling aqueous solutions of pH 2 to 14 for 2 weeks.<sup>83</sup> Recently, Zhou, Li and coworkers utilized a top-down strategy to rationally synthesize an alkali stable MOF, PCN-601, based on a pyrazolate-based porphyrinic ligand (Fig. 4a).<sup>25</sup> All the strong coordination bonds, relatively

short ligand length, and high connectivities of the porphyrinic ligand and  $[Ni_8]$  cluster ( $[Ni_8(OH)_4(H_2O)_2Pz_{12}]$ , Pz = pyrazolate) guaranteed the high chemical resistance of PCN-601 to saturated NaOH solution ( $\sim 20$  M) at  $100^\circ\text{C}$ . In view of the small window size of PCN-601 ( $\sim 2.1 \times 8.0$  Å), the groups then fabricated PCN-602(Ni) which is isostructural with PCN-601 using the reticular synthesis strategy (Fig. 4a).<sup>84</sup> Apart from the high resistance to alkaline solution (pH = 14), PCN-602(Ni) also displayed reasonable tolerance to aqueous solutions with coordinating anions (e.g.  $CO_3^{2-}$ ,  $PO_4^{3-}$ ,  $F^-$ ) (Fig. 4b and c), which was very rare in other high-valent metal-based MOFs like PCN-224(Co), PCN-222(Fe), and PCN-600(Mn). Actually, the investigation of  $[Ni_8]$  cluster-based MOFs can be traced back to the year 2010.<sup>85,86</sup> However, unlike carboxylate-based MOFs and the above-mentioned PCN-601 and PCN-602(Ni),  $[Ni_8]$  cluster-based MOFs fabricated with 4,4'-bis(1*H*-pyrazol-4-yl) biphenyl and 2,6-bis(1*H*-pyrazol-4-yl)pyrrolo[3,4-*f*]isoindole-1,3,5,7(2*H*,6*H*)-tetrone are polycrystalline products.<sup>85</sup>

The self-assembly from low-valent metal ions and triazole-based linkers can also afford highly stable MOFs. Long's group reported a triazolate-bridged framework, Cu-BTTr, which displayed high tolerance even after soaking in boiling water and in a solution of HCl (pH = 3) for 3 days and 1 day, respectively.<sup>87</sup> Lu *et al.* discovered that MAF-X27-Cl as a triazole-based framework was able to retain its structural integrity not only in acidic solution (0.001 M HCl) but also in strong alkaline solution (1.0 M KOH) for over 1 week.<sup>88</sup> It should be noted that, when tetrazole-based linkers are utilized as ligands, the resultant low-valent metal containing MOFs often present weak stability on account of their relatively low basicity and weak coordination ability.<sup>19,89</sup>

In short, low-valent metal ions afford hydrostable skeletons with N donor linkers on account of the relatively high basicity (high  $pK_a$ ) of these linkers. However, these resultant frameworks usually show poor resistance to acidic solutions due to the fairly strong affinity between protons and azolate groups.

**2.1.3 Mixed-metal MOFs.** Given the important roles of metal ions in the fabrication of stable MOFs, the strategy of introducing two or more types of metal ions into the metal clusters of MOFs has been developed. Some mixed-metal MOFs have been found to display enhanced stability as compared to single metal MOFs. There may be several reasons for this: (1) the formation of stronger coordination bonds compared with pristine bonds; (2) the enhancement of the inertness of metal clusters; (3) the improvement of surface hydrophobicity.<sup>90,91</sup> Hence, the design principle toward the fabrication of stable mixed-metal MOFs is the replacement of existing metal ions with more inert species.

For instance, to improve the hydrostability of MOF-5,  $Zn^{2+}$  ions were partially replaced with  $Ni^{2+}$  ions to give the Ni-doped MOF-5.<sup>90</sup> In particular, when the Ni content reached 22%, the resultant Ni-doped MOF-5, due to the stable  $Ni_x(Zn)_{4-x}O^{6+}$  cluster secondary building units (SBUs), was able to retain its integrity when exposed to static air conditions ( $25^\circ\text{C}$ , 30–37% RH) for one week. Furthermore, Ni-doping increased the Langmuir specific surface area and pore size of the resultant MOF on account of the prevention of framework



Fig. 4 (a) Reticular design and construction of PCN-601 and PCN-602(Ni) from TPP<sup>4-</sup> and TPPP<sup>4-</sup> ligands, respectively. Color code: C, gray; O, red; N, blue; and Ni, cyan. H atoms are omitted for clarity. (b) PXRD patterns of PCN-602(Ni) under different conditions. (c) N<sub>2</sub> sorption isotherms of the pristine PCN-602(Ni) sample and of those after soaking in different aqueous solutions (inset: photographs of PCN-602(Ni), PCN-224(Co), PCN-222(Fe), and PCN-600(Mn) soaked in different solutions for 24 h: deionized water, 1 M KF, 1 M Na<sub>2</sub>CO<sub>3</sub>, and 1 M K<sub>3</sub>PO<sub>4</sub> aqueous solutions, from left to right). Adapted from ref. 84 with permission from the American Chemical Society, copyright 2017.

interpenetration. Employing a similar strategy, a few metal ions, including  $\text{Cu}^{2+}$ ,  $\text{Cd}^{2+}$ , and  $\text{Fe}^{2+}$ , were separately doped into a gyroidal MOF, STU-1.<sup>91</sup> Unlike the poor hydrostability of pure STU-1, it was found that all metal doped STU-1s retained their morphology and crystallinity even after soaking in boiling water for 7 days. Beyond that, after doping with metal ions, the significantly increased surface hydrophobicity of STU-1 was proved by water vapor adsorption isotherm curves. The authors speculated that the enhanced hydrophobicity could be ascribed to the perturbation on the MOF surface by doping with metal ions, which impeded the formation of water clusters on the pore surface and improved the water stability of the resultant MOFs.

Indeed, this strategy powerfully enhances the chemical stability of MOFs. However, this advantage is offset by the fact that many metal ions are difficult to dope into the framework by this process. Additionally, the sites of doped metal ions are usually indistinct even when observed by means of the single-crystal X-ray diffraction technology. These factors pose challenges for the mechanism study.

**2.1.4 MOFs based on the azole-containing carboxylic acid ligand.** As mentioned previously, the strong coordination interactions between high-valent metal ions and carboxylic acid linkers or between low-valent metal ions and azole linkers are prone to generate microcrystals or amorphous solids, which pose difficulty for single-crystal X-ray structure determination. In contrast, the weak coordination interaction between low-valent transition metal ions and carboxylates is beneficial to the growth of good-quality MOF crystals, which are usually less stable though large enough for structural characterization. In this case, employing linkers with both azole and carboxylic acid groups may be the route to reconcile the above contradiction and generate MOFs with high stability and large single crystals suitable for structure determination.

By substituting the carboxylate ligand (BDC) with an azole-containing carboxylic acid ligand (3,5-dimethyl-4-carboxypyrazole,  $\text{H}_2\text{dmcapz}$ ), Montoro *et al.* successfully obtained a MOF-5 type framework with  $\text{Zn}^{2+}$  ions.<sup>92</sup> Because of more robust Zn–N bonds, this MOF was highly stable in water and boiling organic solvents for 24 h as compared with the moisture-sensitive MOF-5. In addition, the methyl groups of the linker improved the hydrophobicity of the MOF, which provided protection to the weak Zn–O bonds against hydrolysis. Even when 4-(3,5-dimethyl-1H-pyrazol-4-yl)benzoic acid ( $\text{HMe}_2\text{pzC}_6\text{H}_4\text{CO}_2\text{H}$ ) was employed as a linker, the resultant MOF-5 analogs could withstand prolonged contact with the water/DMF mixture.<sup>93</sup> Using a  $\pi$ -conjugated ligand, 4'-(1H-tetrazol-5-yl)-[1,10-biphenyl]-3,5-dicarboxylic acid ( $\text{H}_3\text{TZBPDC}$ ), involving both soft- and hard-base coordination sites, a zinc-based MOF, USTC-7, with large single-crystal sizes, was successfully synthesized by Jiang's group.<sup>94</sup> As expected, USTC-7 demonstrated its outstanding chemical stability by preserving its framework integrity not only in various boiling solvents but also in pH = 2–12 aqueous solutions for 12 h. Besides zinc azolate-carboxylate frameworks, recently, Liang *et al.* reported a highly stable MOF, FJI-H14, synthesized by a self-assembly process from  $\text{Cu}^{2+}$  ions and 2,5-di(1H-1,2,4-triazol-1-yl)terephthalic acid ( $\text{H}_2\text{BTTA}$ ).<sup>95</sup> Both the penta-coordinated Cu(II) ion subunit formed by Cu–N bonds

and Cu–O bonds, and the abundant free N atoms effectively improved the stability of FJI-H14 under acidic and alkaline conditions.

**2.1.5 Hydrophobic ligand.** To reduce the affinity of MOFs to moisture/water, hydrophobic properties of MOFs are enhanced by binding hydrophobic groups to the linkers or by creating them around the metal nodes. The water repellent functional groups around the metal clusters can effectively protect the weak coordination bonds from the attack by water molecules and enhance the MOF stability in water.

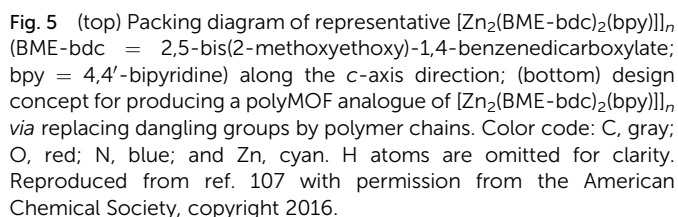
To illustrate the effectiveness of hydrophobic ligands, a methyl group modified MOF-5 was synthesized.<sup>96</sup> The  $\text{H}_2$  adsorption isotherm revealed that the  $\text{H}_2$  uptake capacity of 2,5-dimethyl-modified MOF-5 was preserved even after a four-day exposure to ambient air. Chen *et al.* designed and synthesized an octatopic carboxylic ligand, 3,3',5,5'-tetrakis(3,5-dicarboxyphenyl)-2,2',4,4',6,6'-hexamethylbiphenyl, to prepare a copper(II)-paddlewheel MOF (BUT-155).<sup>97</sup> Compared with the common copper(II)-paddlewheel MOFs, the high connectivity and abundance of hydrophobic methyl groups endowed BUT-155 with exceptional chemical stability in aqueous solutions of a wide pH range (4–10) and boiling water for 24 h.

To investigate the influence of side-chain length, a series of isostructural NbO-type MOFs were tested.<sup>98</sup> Upon extending the length of dialkoxy-substituents (from  $\text{C}_1$  to  $\text{C}_6$ ) on the organic linker, the moisture stability of the resultant Cu(II)-MOF improved noticeably while the thermal stability decreased. Furthermore, an inevitable problem was the gradual drop in porosity of the resultant MOFs after the incorporation of these hydrophobic functional groups. In addition to the length of the side chain, the position of the hydrophobic groups relative to metal clusters also plays a critical role in improving MOF chemical stability.<sup>99,100</sup> A convincing comparison was made between MOF-508 and its two analogs (SCUTC-18 and SCUTC-19).<sup>99</sup> Compared with bipyridine of MOF-508, there were two methyl-substituents, respectively, located at the *ortho*-positions and *meta*-positions of the coordinating N atoms of bipyridine in SCUTC-18 and SCUTC-19. Upon exposure to humid air for 30 days, only the porous structure of SCUTC-18 was maintained due to the introduction of methyl-substituents at sites more adjacent to the metal clusters.

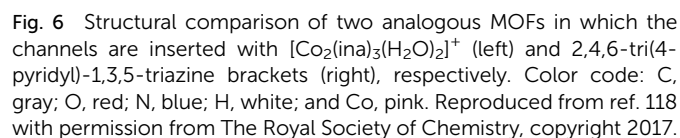
Apart from alkyl chains,<sup>96–101</sup> introduction of fluorinated groups (*e.g.*  $-\text{F}$ ,  $-\text{CF}_3$ , *etc.*) or other hydrophobic substituent groups into organic linkers is also an efficient means for enhancing MOFs' moisture/water resistance.<sup>102–114</sup> Jiang's group reported a copper-based MOF, USTC-6, based on 4,4'-(perfluoropropane-2,2-diyl)diphthalic acid.<sup>109</sup> Although most Cu–O coordination bonds in MOFs are vulnerable to hydrolysis, USTC-6 exhibits exceptional tolerance to water and aqueous solutions in the pH range 2 to 10 on account of the hydrophobicity of the corrugated  $-\text{CF}_3$  surface. Cohen's group developed a bridging co-ligand strategy to construct a series of polymer-metal-organic frameworks (polyMOFs) (Fig. 5).<sup>107</sup> Remarkably, all polyMOFs inherited the hydrophobicity from the substituted  $\text{H}_2\text{BDC}$  derivatives with water contact angles (WCA) of 110–120°. As expected, several polyMOFs displayed outstanding stability even after exposure to 90% RH at 25 °C for 7 days.







In the case of MIL-88 type MOFs, organic linkers with  $C_3$  symmetry are the ideal size-matching MBBs. For example, both  $[\text{Co}_2(\text{ina})_3(\text{H}_2\text{O})_2]^+$  (ina = isonicotinate) and 2,4,6-tri(4-pyridyl)-1,3,5-triazine (tpt) were inserted into the open channels during the MOF formation (Fig. 6).<sup>118</sup> The grand canonical Monte Carlo (GCMC) simulations and density functional theory (DFT) calculations revealed that the framework energy was drastically



For example, the stabilities of two isostructural pillared MOFs, MOF-508 and DMOF, upon exposure to water were compared by Walton's group.<sup>121</sup> Upon exposure to RH = 90%, the surface area of non-interpenetrated DMOF dramatically declined (from 1980 to 7 m<sup>2</sup> g<sup>-1</sup>). In contrast, the two-fold interpenetrated MOF-508 showed no significant change in the surface area after the same treatment. This clearly demonstrated the enhanced framework stability resulting from catenation. A similar phenomenon was also observed in two-fold catenated Co-BTTB-DMBPY and Zn-BTTB-DMBPY.<sup>122</sup> Actually, the influence of the interpenetration direction on framework stability is also very apparent. Using the solvothermal reaction of Zn(NO<sub>3</sub>)<sub>2</sub> and 4-(3,5-dimethyl-1*H*-pyrazol-4-yl)benzoic acid (H<sub>2</sub>mpba), three isomeric MOFs with different polar nets were reported.<sup>123</sup> Interestingly, although all these three MOFs present



4-fold interpenetrating structures, the different interpenetration directions endow them with entirely different levels of porosity and stability. Molecular mechanics (MM) calculations revealed that the framework energy of the nonporous isomer was the lowest. As expected, after being immersed in water, the other two nanoporous isomers transformed into the most stable framework (nonporous isomer).

Recently, Bu's group demonstrated that a self-penetrated network was more stable than an interpenetrated framework (Fig. 7).<sup>124</sup> By employing 5,5'-((thiophene-2,5-dicarbonyl)bis(azanediyl))diisophthalic acid and 5,5'-([2,2'-bipyridine]-5,5'-dicarbonyl)bis(azanediyl)diisophthalic acid separately as the organic linkers, NKU-112 with a two-fold interpenetrated framework and NKU-113 with a self-interpenetrated framework were prepared. In NKU-113, the coordinated DMF molecules of metal clusters were substituted by chelating the bipyridine moiety of 5,5'-([2,2'-bipyridine]-5,5'-dicarbonyl)bis(azanediyl)diisophthalic acid. This greatly increased the framework rigidity of NKU-113 as compared to NKU-112. After solvent exchange and supercritical drying, NKU-113 retained its framework and displayed a moderate BET surface area of 1486 m<sup>2</sup> g<sup>-1</sup>, while only a negligible amount of N<sub>2</sub> was adsorbed in NKU-112 due to its collapsed structure.

## 2.2 Improving the chemical stability of existing MOFs

Given that the rational strategy for preparing stable MOFs *via de novo* syntheses remains limited, a large number of strategies for enhancing the stability of existing MOFs have been developed. In this section, four major approaches, including post-synthetic exchange, post-synthetic modification, hydrophobic surface treatment, and composite fabrication, will be discussed.

**2.2.1 Post-synthetic exchange.** As one of the important post-functionalization methods, post-synthetic exchange (PSE)

can offer a promising tool to improve the physical and chemical attributes of MOFs *via* metal-ion metathesis, linker exchange or the replacement of the counterions of ionic MOFs. Along this line, the PSE process might increase the bond strength of coordination bonds in the unstable SBUs of MOFs or tune the water-resistance of MOFs without disturbing the framework structure.

As regards metal-ion metathesis, an early example was found in SDU-1.<sup>125</sup> By soaking into a Cu<sup>2+</sup> containing solution, the Zn<sup>2+</sup> ions of SDU-1 were exchanged by Cu<sup>2+</sup> ions in a single-crystal-to-single-crystal (SC-SC) fashion, giving a stable Cu-SDU-1. After the removal of the guest solvent, the N<sub>2</sub> adsorption capacity of Cu-SDU-1 was ~3 times higher than that of SDU-1, demonstrating the increase of framework stability upon PSE. In 2014, Zhou's group developed a postsynthetic metathesis and oxidation strategy to transform the labile PCN-426-Mg into two robust MOFs, PCN-426-Fe(III) and PCN-426-Cr(III).<sup>126</sup> Although PCN-426-Mg was labile in water, PCN-426-Fe(III) could remain intact even after immersion in water for one day. When it came to PCN-426-Cr(III), essentially unchanged crystallinity was found in solutions in the range from pH = 12 to extremely acidic conditions for over 12 h. Inspired by the high stability of Cr(III)-based MOFs, PCN-333(Cr) was obtained by the reduction of PCN-333(Fe) with Cr(II) ions and the subsequent reductive labilization process.<sup>127</sup> PCN-333(Fe) could only maintain its structural integrity in aqueous solutions with a pH range from 3 to 9, while PCN-333(Cr) was stable at pH ranging from 0 to 11. Wang *et al.* demonstrated that exchange solvent played a vital role in the PSE process of MOFs.<sup>128</sup> Acetone was proven to be a promising solvent for a postsynthetic metal exchange due to the direct participation of the carbonyl group *via* coordination interaction with metal ions. With the aid of acetone, several Fe/Cr-exchanged MOFs were obtained from the Fe(III)-based counterparts under very mild conditions and all these modified MOFs presented enhanced chemical stability.

Compared with investigations into postsynthetic metal exchange, there are fewer investigations into postsynthetic linker exchange and counterion exchange for improving the stability of MOFs. Utilizing DUT-67 as the template, the polarity of the inner surface was tuned through the postsynthetic exchange of the modulator by fluorinated monocarboxylates, such as trifluoroacetic acid, pentafluorobenzoic acid, perfluorooctanoic acid, *etc.*<sup>129</sup> The incorporation of fluorinated linkers rendered the resultant MOF hydrophobic, greatly increasing their tolerance to water. Upon water adsorption at 298 K, only 0.3% porosity loss was observed in the perfluorooctanoic acid-modified DUT-67, while the same treatment caused a porosity loss of 34% in the parent DUT-67. As for postsynthetic counterion exchange, a series of alkylammonium cations were employed to adjust the surface wettability of bio-MOF-1.<sup>130</sup> Water stability tests demonstrated that the large-sized tetraethylammonium (TEA) and tetrabutylammonium (TBA) cations with hydrophobic nature effectively protected bio-MOF-1 from the attack of water molecules. In contrast, significant deformation took place in the parent bio-MOF-1 and the small-sized tetramethylammonium (TMA) or hydrophilic



Fig. 7 Schematic diagram showing the framework and the modification strategy. The blue-colored balls, silver-colored sticks, yellow-colored sticks, and red-colored cylinders represent metal SBUs, ligands, coordinate bonds and pores, respectively. Reproduced from ref. 124 with permission from The Royal Society of Chemistry, copyright 2018.



triethylhydroxylmethylammonium cation modified frameworks after exposure to water vapor.

**2.2.2 Post-synthetic modification.** The organic linkers and metal clusters in MOFs provide great opportunities to a vast range of chemical transformations, which means that the introduction of particular functional groups *via* the post-synthetic modification (PSM) approach can alter the pore environments of MOFs.<sup>131</sup> In this section, the improved resistance of MOFs towards moisture and different solutions is discussed by incorporating appropriate functional groups based on this strategy.

As a very early example, IRMOF-3 was modified with various alkyl anhydrides to give moisture-resistant IRMOF-3-AMX ( $X = 1-6, 15, iPr, tBu$ ).<sup>132</sup> The longer the chain of alkyl anhydride was, the more hydrophobic the resultant modified MOF became. In sharp contrast to the significantly reduced crystallinity of IRMOF-3 (WCA:  $\sim 0^\circ$ ), no evident change was found in the crystallinity of IRMOF-3-AM15 (WCA:  $123 \pm 5^\circ$ ) upon exposure to ambient air for 4 days, revealing the protective effects from the modified alkyl chains on MOFs. The same approach was also applied to  $NH_2$ -MIL-53(Al) and  $Cu_3(NH_2BTC)_2$ , greatly increasing their stability.<sup>132,133</sup> Using a diazotization reaction between amidogen and 1-methylindole,  $NH_2$ -UiO-66 was successfully transformed into  $UiO-66-N=N-ind$ .<sup>134</sup> By comparison, the resistance to the acidic and alkaline environment of  $UiO-N=N-ind$  was expanded from the pH range of 1–9 ( $NH_2$ -UiO-66) to 1–12. However, the action was not limited to the aforementioned functional groups. Hydrophobic phenylisocyanate could also be modified on the framework to protect the MOF from the attack of water molecules.<sup>135</sup>

As for the post-synthetic modification of metal clusters, hydrophobic groups were grafted at the  $Zr_6$ -oxo nodes of NU-1000 *via* the solvent-assisted ligand incorporation (SALI) process.<sup>136</sup> The dangling organic carboxylates could effectively increase the tolerance of  $Zr_6$ -oxo nodes to water, albeit with the reduction of pore volume. Specifically, perfluorodecanoic-acid-functionalized NU-1000 could maintain its crystallinity and porosity even after 20 cycles of water vapor adsorption and desorption, proving its strong hydrostability. Moreover, when monotopic carboxylate was replaced with the phosphonate ligand, the resultant MOF exhibited higher resistance to hydroxide as a result of stronger bonds between the phosphonate ligand and  $Zr_6$ -oxo nodes.<sup>137</sup> The same effect was also found in the PCN-222 system by employing the SALI strategy with diphenylphosphinic acid,<sup>138</sup> demonstrating the universal applicability of this method. Recently, Kim's group obtained a super-hydrophobic MOF (WCA:  $161^\circ$ ),  $NH_2$ -UiO-66-shp, by modifying the  $Zr_6$ -oxo clusters of  $NH_2$ -UiO-66 (WCA:  $0^\circ$ ) with phenylsilane *via* pore-surface engineering (Fig. 8).<sup>139</sup> The stability tests demonstrated that  $NH_2$ -UiO-66-shp not only possessed the same acidic stability as  $NH_2$ -UiO-66 but also showed excellent tolerance to 0.1 M NaOH solution for 5 h, surpassing the alkali labile  $NH_2$ -UiO-66.

**2.2.3 Hydrophobic surface treatment.** Although the chemical stability of MOFs can be significantly improved by the above methods, the porosities and adsorption properties are often drastically reduced by the occupation of the channels by

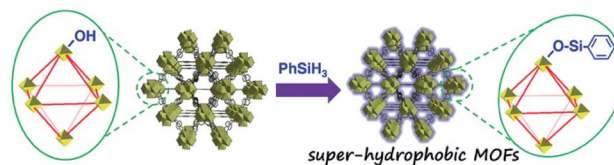


Fig. 8 Preparation diagram of super-hydrophobic  $NH_2$ -UiO-66-shp. Reproduced from ref. 139 with permission from Wiley-VCH, copyright 2019.

dangling functional groups. To avoid this, hydrophobic surface treatment has been developed to be applied to the exterior surface of MOFs to protect the MOFs from water while preserving their porosity to a large degree. In general, hydrophobic surface treatment mainly includes surface coating, post-synthetic thermal annealing, and post-synthetic surface modification.

Surface hydrophobic coating is a very simple and direct yet effective approach to improve MOF stability with retained porosity. In 2014, Zhang *et al.* developed a facile and general polydimethylsiloxane (PDMS)-coating strategy to enhance the stability of MOFs in the presence of moisture or water (Fig. 9a).<sup>140</sup> The hydrophobic PDMS coating layer yielded impressive results. The hydrostability of three randomly selected and structurally different MOFs, namely, MOF-5, HKUST-1, and ZnBT, was significantly enhanced. The coating layer prevented the attack from water. It is worth noting that the BET surface area and gas uptake performance of the PDMS-

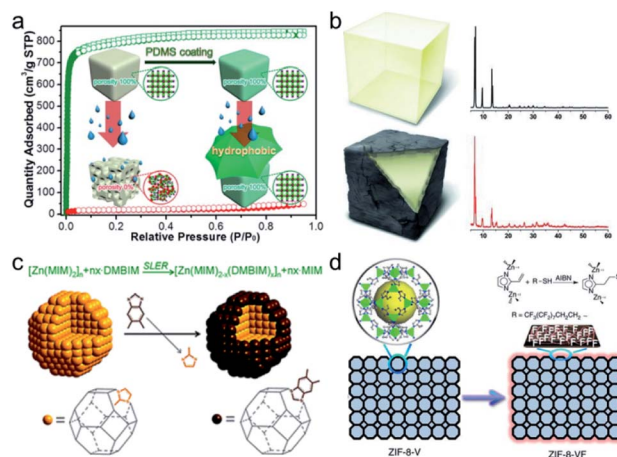


Fig. 9 Schematic representation of (a) PDMS-coating on the MOF surface for improved moisture/water resistance, (b) showing MOF-5 (top) and MOF-5 after thermal modification to produce amorphous carbon-coated MOFs (bottom) and related powder XRD patterns of corresponding samples (right), (c) shell-ligand-exchange-reaction process of ZIF-8, and (d) imparting amphiphobicity on ZIF-8-V *via* grafting perfluoroalkyl groups on the exterior surface. (a) Reproduced from ref. 140 with permission from the American Chemical Society, copyright 2014. (b) Reproduced from ref. 145 with permission from Wiley-VCH, copyright 2012. (c) Reproduced from ref. 147 with permission from The Royal Society of Chemistry, copyright 2013. (d) Reproduced from ref. 148 with permission from Nature Publishing Group, copyright 2016.

coated MOFs are almost maintained, demonstrating the good permeability of the coating layer. Furthermore, the PDMS coating layer on the surface of MOFs did not block the catalytically active sites from coming into contact with the substrates during the catalytic reaction. When the PDMS-coated MOF and the pristine MOF are subjected to the same water vapor treatment, the PDMS-coated MOF is observed to retain its catalytic activity better than the pristine MOF. Since this work, a number of similar approaches have been developed and applied to improve MOF stability.<sup>141–144</sup> Qian *et al.* successfully deposited a hydrophobic organosilicone layer on the exterior surface of MOFs *via* a facile solution-immersion process without any heat treatment.<sup>141</sup> Upon exposure to water for 5 days, it was found that the porosity of the MOFs HKUST-1, MIL-125, and ZIF-67 was retained to a great extent due to the enhanced hydrophobicity. Hou *et al.* developed a post-synthetic polymerization method to form polymer brushes on the external surfaces of MOFs.<sup>142</sup> Interestingly, upon irradiation with UV-light, polymerization could be initialized by active hydrogen species (surface radicals) on the MOFs to generate the polymer brushes on the MOFs to obtain MOF-based materials with enhanced chemical stabilities. Recently, Queen's group successfully developed a two-step post-synthetic polymerization method to improve the stability of seven different types of MOFs.<sup>143</sup> First, polydopamine (PDA) as an adhesive was firstly coated on the surface of these MOFs under an oxygen atmosphere. Then, based on a Michael addition, the hydrophobic 1*H*,1*H*,2*H*,2*H*-perfluorodecanethiol was covalently grafted on the surface of the MOF@PDA that resulted from the previous step. As expected, all the highly hydrophobic MOF composites displayed superior chemical stability under acidic or alkaline conditions compared with the pristine MOFs. Thermal polymerization of protective layers on the MOF surfaces can also strengthen their chemical stability. Using direct polymerization of fluorinated 4-undecylcatechol (fdcat), HKUST-1 was coated with hydrophobic layers under mild reaction conditions.<sup>144</sup> Compared with unmodified HKUST-1, after soaking in water for a week, the permeable coatings protected HKUST-1 crystals against water molecules while retaining 92% of the porosity.

Inspired by the high hydrostability of "carbonaceous grease", Park's group prepared highly moisture-resistant black-colored MOF-5 through a simple heat treatment at a specific temperature followed by immediate cooling to room temperature (Fig. 9b).<sup>145</sup> When the pyrolysis temperature was raised to 530 °C, the BET surface area of the resultant material (1740 m<sup>2</sup> g<sup>-1</sup>) was half of that (3450 m<sup>2</sup> g<sup>-1</sup>) of pristine MOF-5. Nevertheless, thermally modified MOF-5 retained its crystallinity and porosity to a great extent upon exposure to ambient air for at least two weeks. This indicated its high hydrostability. Gadipelli *et al.* investigated the framework decomposition of MOF-5 crystals by varying the annealing temperature.<sup>146</sup> After annealing MOF-5 at 400 °C (that is, below the onset decomposition temperature of 500 °C) for 3 h, the resultant sample could retain certain porosity even after exposure to air for 15 days.

As an example of post-synthetic surface modification, 2-methylimidazolate located on the exterior surface of ZIF-8

particles was substituted with 5,6-dimethylbenzimidazole (DMBIM) *via* a shell-ligand-exchange-reaction (SLER) to improve the hydrothermal stability (Fig. 9c).<sup>147</sup> After four cycles of isobutanol sorption, the uptake amount of ZIF-8 fell to ~0 caused by the collapse of the structure. In contrast, no obvious change in adsorption capacity was found in the DMBIM modified ZIF-8 thanks to the maintenance of the framework. By deliberately utilizing 2-vinyl-imidazole as the organic linker, Sun *et al.* successfully fabricated ZIF-8-V, a ZIF-8 type framework and hydrophobically treated its external surface with per-fluoroalkyl groups based on the thiol-ene click reaction (Fig. 9d).<sup>148</sup> The resultant ZIF-8-VF was an amphiphobic material and was able to remain intact upon exposure to 100% RH under a CO<sub>2</sub> atmosphere at 45 °C for over 720 h, while almost all ZIF-8 structures degraded under the above conditions in 240 h. The method was also applied to MOF-5, producing a similar effect. Analogously, alkylphosphonic acids with long alkyl chains were modified on the Zr<sub>6</sub>-nodes on the exterior surface of MOFs, namely, UiO-66, UiO-66-SO<sub>3</sub>H, and PCN-222, without any disturbance to the internal pores.<sup>149</sup> By virtue of the external superhydrophobic modification, the tolerance of the resultant MOFs toward different harsh environments was appreciably enhanced.

**2.2.4 Composite.** The high porosity and compatibility of MOFs allow the hybridization of MOFs with other materials such as polymers, graphite oxide, carbon nanotubes, *etc.*<sup>150–164</sup> By growing MOFs on the surface or inside the pores of other substrates, or incorporating other materials into the cavities of MOFs, the resultant MOF composites can be imparted with the combined characteristics of the two components or acquire even new functions such as hydrophobicity, hydrostability, mechanical properties, and so on.

In 2009, Park's group successfully incorporated acid-treated multiwalled carbon nanotubes (MWCNTs) into MOF-5 to give a hybrid composite (MOFMC).<sup>150</sup> This composite growth helped with removing the guest molecules during activation which enhanced the Langmuir specific surface area. Moreover, the rigid MWCNTs could protect the moisture-sensitive MOF-5, making MOFMC stable even after one week in air. The same effects can also be achieved when MWCNTs are replaced by graphite oxide or carboxyl-functionalized attapulgite.<sup>151,152</sup> Matzger's group produced MOF-5-polystyrene composites using the polymerization of styrene in the presence of MOF-5.<sup>153</sup> Polystyrene was uniformly distributed throughout the MOF-5 particles. The resultant composites were endowed with much better moisture stability at RH of 53% as compared with the pristine MOF-5. Apart from polystyrene, polynaphthylene (PN) could also be formed in the channels of MOF-5 as partitions *via* a radical reaction, affording a hydrophobic PN@MOF-5 composite with significantly improved hydrostability.<sup>154</sup> In fact, so long as the hydrophobic molecules are introduced into pores, MOFs would display improved water stability. Decoste's group prepared hydrophobic HKUST-1 by treating the parent MOF with plasma-enhanced chemical vapor deposition (PECVD) of perfluorohexane (PFH).<sup>155</sup> With the protection from adsorbed PFH, hydrophobic HKUST-1 could remain resistant to the attack by water for at least 24 h.



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toxic and corrosive ammonia ( $\text{NH}_3$ ), the Morris group synthesized a Cu(II)-based MOF, STAM-17-OEt, based on the 5-ethoxy isophthalate linker.<sup>172</sup> Interestingly, upon exposure to a humid environment, the weak Cu–O bonds located on the paddlewheel cluster of the dehydrated MOF would transform into its original hydrated form. This conversion prevents the breakup of the Cu–O bonds in the paddlewheel clusters which hold the framework together, which improves the stability of the framework. STAM-17-OEt was then employed as an adsorbent for  $\text{NH}_3$  and infrared spectroscopy demonstrated that the MOF structure was retained even after 5 days of the ammonia breakthrough experiment (90% RH). DeCoste *et al.* prepared a set of chemically stable mixed-matrix membranes (MMMs) based on HKUST-1 and polyvinylidene difluoride (PVDF).<sup>161</sup> By virtue of the enhanced hydrophobicity and unsaturated  $\text{Cu}^{2+}$  sites, the HKUST-1 MMMs exhibited outstanding  $\text{NH}_3$  uptake performance and a large proportion of  $\text{NH}_3$  adsorption capacity survived at 90% RH at 25 °C for 28 days. Given the ultra-efficient metal ion permeation, excellent stability and processability of the MOF membrane, Liu *et al.* incorporated these two MOFs into various polymer matrixes to create a sequence of UiO-66 and  $\text{NH}_2$ -UiO-66 membranes. They employed the membranes as adsorbents in the recovery of palladium Pd(II) and platinum Pt(IV) ions from strongly acidic solutions (pH = 1.0).<sup>176</sup> By virtue of the high affinity to Pt/Pd and the striking stability caused by the strong  $\pi$ – $\pi$  interactions between the polymer and the MOFs, the resultant membranes showed amazing adsorption capacities for the above ions in strongly acidic solutions, and also excellent recycle performance. Recently, a hydrophobic porous coordination polymer, USTC-6, was uniformly grown throughout a graphene oxide-modified sponge to yield a macroscopic USTC-6@GO@sponge adsorbent with excellent chemical stability for highly efficient oil/water separation.<sup>109</sup> Remarkably, USTC-6@GO@sponge could be combined with tubes and a self-priming pump to fabricate a model apparatus for continuous oil recovery from water, showing a promising future for stable MOFs for oil/water separation.

Undoubtedly, highly stable MOFs have exhibited their unique advantages in sorption and separation. The immediate challenges lie in the design of MOF-based adsorbents such as MOF membranes with low (production and regeneration) cost, desirable working capacity and selectivity, and long-term stability according to the demands of practical use. Beyond these, more efforts are needed in mechanistic studies of the co-adsorption behaviour of some adsorbates in MOFs.

### 3.2 Heterogeneous catalysis

The Lewis acid/basic sites, Brønsted acid sites, and redox active sites on the metal clusters and/or the organic linkers make stable MOFs efficient heterogeneous catalysts for various reactions.<sup>12,15,17,177</sup> Since many of these reactions take place in water or even in acidic or basic solutions, MOF-based catalysts should be stable to tolerate such conditions.

By employing MIL-101- $\text{SO}_3\text{H}$  as the acid catalyst, reasonable hydrolysis efficiency of cellulose was achieved in aqueous solution systems.<sup>178</sup> To beneficially utilise the outstanding

chemical stability of PCN-602(Ni), Lv *et al.* employed PCN-602 with  $\text{Mn}^{3+}$ -porphyrin centers (denoted as PCN-602(Mn)) for the C–H bond halogenation under harsh conditions (in aqueous solutions and dichloromethane mixtures of 0.165 M NaClO).<sup>84</sup> Thanks to the presence of the Mn-porphyrin centre and rigid framework structure, a high yield of chlorocyclohexane (92%) was achieved on PCN-602(Mn) after 5 h reaction, much higher than that (8%) achieved by the homogeneous catalyst, Mn(TPP) Cl. Apart from organocatalysis, many electrocatalytic and photocatalytic properties of MOFs have also been studied in aqueous or even in acidic or basic solutions. For instance, the stable MIL-101(Fe) was chosen as an effective catalyst for visible-light-driven water oxidation.<sup>179</sup> With the aid of  $[\text{Ru}(\text{bpy})_3]^{2+}$ , MIL-101(Fe) presented a decent initial turnover frequency ( $0.10 \text{ s}^{-1}$ ) with a high oxygen yield of 36.5%. Furthermore, the recycling tests and X-ray photoelectron spectra (XPS) results showed that MIL-101(Fe) was stable after water oxidation. Lu *et al.* obtained an alkaline-stable MOF, MAFX27-OH, through the post-synthetic ion exchange of MAF-X27-Cl during a linear sweep voltammetric procedure.<sup>88</sup> In 1.0 M KOH aqueous solution, MAFX27-OH with both OMSs and hydroxides offered an overpotential of 292 mV at  $10.0 \text{ mA cm}^{-2}$  with remarkable durability in 20 h of electrocatalytic oxygen evolution reaction (OER) processes. Shen *et al.* developed a modular synthesis strategy to stabilize the  $\text{Co}_2(\text{RCOO})_4(\text{H}_2\text{O})_2$  (R = substituent group) cluster into MCF-37 by exchanging the original paddle-wheel type Fe-carboxylate core, affording a modified alkaline-stable MOF, MCF-49 (Fig. 11).<sup>180</sup> By virtue of the high catalytic activity of  $\text{Co}_2(\text{RCOO})_2(\text{L}^T)_2$  ( $\text{L}^T$  = terminal ligand) towards electrocatalytic OER and the high alkaline stability of the framework, MCF-49 showed a very low overpotential of 225 mV at  $10 \text{ mA cm}^{-2}$  with nearly 100% Faraday efficiency in aqueous solution (pH = 13). In terms of hydrogen evolution reaction (HER), Qin *et al.* rationally designed two highly stable polyoxometalate (POM)-based MOFs, NENU-500 and NENU-501, and employed them as HER electrocatalysts in 0.5 M  $\text{H}_2\text{SO}_4$  aqueous solution.<sup>181</sup> NENU-500 gave a noticeably low onset overpotential of 180 mV with a Tafel slope of  $96 \text{ mV dec}^{-1}$ . In addition, the two MOFs demonstrated their ultrahigh stability by continuing their activities even after 2000 cycles. Based on the outstanding chemical stability and unique photocatalytic properties of porphyrins, Al-TCPP was utilized as a photocatalyst for hydrogen generation from water.<sup>49</sup> With the aid of methyl viologen and ethylenediaminetetraacetic acid, Al-TCPP gave an

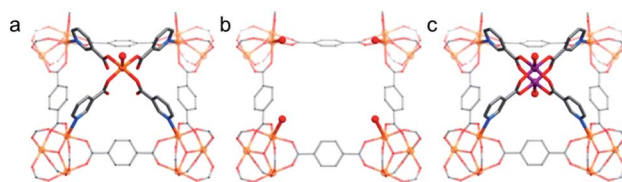


Fig. 11 Key local structures of (a) MCF-37, (b) the  $[\text{Fe}_3(\mu_3\text{-O})(\text{BDC})_3(\text{L}^1)_3]$  scaffold, (c) MCF-49. Color code: C, gray; O, red; N, blue; Fe, orange; and Co, purple. H atoms are omitted for clarity. Adapted from ref. 180 with permission from the American Chemical Society, copyright 2017.



As stated above, enhanced stability has widened the scope for the use of MOFs and MOF-based materials as catalysts. In future, it can be expected that the use of operando spectroscopy for gaining a better understanding of the catalytic mechanism, precise control of active sites, and multifunction synergy will be of great importance for heterogeneous catalysis using stable MOFs. In addition, to address the challenges involved in MOF catalysis such as mass transfer, conductivity, and high cost, is also the important consideration.

In conclusion, stable MOFs have shown great potential for sensing various analytes. However, there are many improvements necessary and several issues to be solved before practical implementation. First, more effort should be devoted to improve sensitivity, selectivity, response time, detection limit and long-term stability of fluorescent MOF materials based on the requirements of practical use. Second, further investigation



into MOF-based turn-on sensors and their turn-on mechanism will prove valuable. Last but not least, the integration of MOFs into sensing devices is still at a very early stage. Hence, more endeavour is necessary in this direction.

### 3.4 Biological and medical applications

For their tunable pore sizes, high surface area, and versatile physical and chemical properties, MOFs are extensively used in biological and medical applications such as biosensing, drug release, biomimetic catalysis, *etc.*<sup>5,195–201</sup> For these applications, MOFs with excellent chemical stability are a must. They should possess strong resistance to hydrolysis/collapse in physiological environments in which they would be expected to function, *e.g.* stomach acidity, intestinal alkalinity and the peristalsis in the esophagus, stomach, and intestines.

As for biosensing, a representative example is Cu-TCA constructed with  $\text{Cu}_2(\text{O}_2\text{CR})_4$  paddlewheel units and tricarboxytriphenyl amine (TCA) linkers.<sup>202</sup> Upon the addition of NO, the coordination interaction between NO and Cu(II) complexes reduced  $\text{Cu}^{2+}$  to  $\text{Cu}^+$  ions and caused the recovery of luminescence from Cu-TCA in aqueous solutions. The brightness of emission of Cu-TCA made it an ideal candidate for biological imaging of NO in living cells. Wu *et al.* utilized mixed ligands to fabricate a lanthanide-based MOF (Ln-MOF) that shows high 48 h stability in water.<sup>203</sup> The competition for the absorption of irradiated light between biomarkers serotonin (5-hydroxytryptamine, HT) and 5-hydroxyindole-3-acetic acid (HIAA) and the organic linkers, and the dynamic quenching process caused fluorescence quenching, thus endowing the Ln-MOF with remarkable sensitivity for selective detection of the two biomarkers in aqueous solution. On the basis of the intrinsic peroxidase-like catalytic activity, two Fe-based MOFs, MIL-68 and MIL-100, were found to behave as colorimetric biosensors to detect  $\text{H}_2\text{O}_2$ .<sup>204</sup> The high chemical stability of MIL-68 and MIL-100(Fe) allowed them to be used to catalyze the oxidation of different peroxidase substrates with  $\text{H}_2\text{O}_2$  in acetate buffer with  $\text{pH} = 4.0$ . Moorthy's group obtained a luminescent and stable homochiral MOF, Zn-PLA, based on a concave shaped pyrene-tetralactic acid.<sup>205</sup> Interestingly, histidine was found to cause more significant fluorescence quenching in the aqueous dispersion of Zn-PLA than many other amino acids. This led to the use of Zn-PLA, an enantiodifferentiating sensor of histidine. The authors speculated that this fluorescence quenching was due to the exchange of dimethylammonium cations in Zn-PLA by deprotonated histidine in water.

As an early attempt at the use of MOFs for drug release, two flexible frameworks, MIL-53(Cr) and MIL-53(Fe), were selected as carrier systems for delivering ibuprofen.<sup>206</sup> Due to the unique flexibility, a very slow and complete release of ibuprofen over a period of three weeks was accomplished with both MIL-53(Cr) and MIL-53(Fe). Deng's group revealed that the guest release rate was strongly correlated with the type and proportion of functional groups in MOFs.<sup>207</sup> By constructing different multivariate MOFs from MIL-101(Fe), the maximum release amount of doxorubicin (DOX) could be shifted from the 17th to the 29th day within a 40 day release period. In order to improve the

biocompatibility of carriers, adenine was employed as a biomolecular ligand to fabricate an ionic framework, bio-MOF-1, which could maintain the structural integrity in biological buffers for weeks.<sup>208</sup> Unlike common MOFs, the release of procainamide from bio-MOF-1 could be triggered by the ionic interactions between the drug and framework. Recently, Farha's group demonstrated the immobilization of insulin in an acid-stable MOF, NU-1000.<sup>209</sup> By virtue of the protective effect from NU-1000, insulin was stable even upon exposure to stomach acid ( $\text{pH} = 1.5\text{--}3.5$ ) and pepsin (Fig. 12). Moreover,  $\sim 40$  wt% of insulin could be released from the host framework under simulated physiological conditions ( $\text{pH} = 7.0$ ) (Fig. 12).

As for the biocatalysis, Reynolds's group utilized Cu-BTTRI with high stability in fresh citrated whole-blood (30 min,  $\text{pH} 7.4$ ,  $37^\circ\text{C}$ ) to produce polymeric medical devices with biomedical grade polyurethane.<sup>210</sup> Fortunately, the catalytic activity from the exposed  $\text{Cu}^{2+}$  active sites of Cu-BTTRI can be retained, and the medical devices realized the surface-localized generation of NO from endogenous sources even in fresh citrated whole blood. Inspired by the bimetallic Zn-OH-Zn active site in phosphotriesterase enzymes, UiO-66 with similar Zr-OH-Zr bonds in  $[\text{Zr}_6\text{O}_4(\text{OH})_4]$  clusters was utilized as a biomimetic catalyst for the hydrolysis of methyl paraoxon in an aqueous solution containing 0.45 M *N*-ethylmorpholine ( $\text{pH} = 10$ ).<sup>211</sup> Besides directly behaving as biocatalysts, stable MOFs can also act as catalyst supports for immobilizing enzymes for biocatalysis.<sup>50,212</sup> Three enzymes of different sizes, namely, horseradish peroxidase (HRP), cytochrome *c* (Cyt *c*) and microperoxidase-11 (MP-11), were separately encapsulated in PCN-333(Al).<sup>50</sup> Compared with the same enzymes immobilized in SBA-15, the enzymes encapsulated in PCN-333(Al) presented much superior catalytic stability through five cycles of *o*-phenylenediamine oxidation due to the outstanding chemical stability of PCN-333(Al) and strong interaction between the



Fig. 12 Schematic illustration of (a) encapsulation of insulin in the mesopores (32 Å) of NU-1000 and exclusion of pepsin from the MOF framework and (b) exposure of insulin and insulin@NU-1000 to stomach acid. Free insulin denatures in stomach acid and is digested by pepsin. The release of insulin from NU-1000 happens when insulin@NU-1000 is exposed to a PBS solution. Reproduced from ref. 209 with permission from the American Chemical Society, copyright 2018.

enzymes and frameworks. Using PCN-888 with three different sized cavities as the carrier, Lian *et al.* fabricated a tandem nanoreactor *via* a unique stepwise encapsulation of two different enzymes (glucose oxidase (GOx) and horseradish peroxidase (HRP)).<sup>212</sup> It was remarkable that strong interactions between immobilized moieties and cages of PCN-888 not only prevented the trypsin digestion of encapsulated enzymes but also ensured the maintenance of catalytic activity through four catalytic cycles.

Nevertheless, the study of MOFs and MOF-based materials for biological and medical applications is still in its infancy. One remaining critical issue is the fabrication of nontoxic MOFs with outstanding chemical stability, excellent biocompatibility, and appropriate pore size and pore volume. In addition, control of MOF particle size is also very important in view of the endocytosis by the living cells and systemic circulation in blood. As for the drug delivery, the degradation mechanism of MOFs should be studied closely before the *in vivo* investigations and clinical applications.

### 3.5 Proton conductivity

Over the past decades, MOFs have attracted rapidly increased attention in the field of proton conductivity on account of their tailorable structures.<sup>213–217</sup> Different from other porous materials, proton conduction of MOFs can be easily controlled by tuning the hydrophilicity and acidity of the surface structure of MOFs. To date, many proton conduction studies on MOFs have been reported, though only a limited number of MOFs displayed both high conductivity and good stability.<sup>218</sup> In principle, the stability of MOFs as proton conductors mainly refers to long-term stability under different temperature and/or humidity conditions. It is important to note here that proton transport relies mainly on the generation of hydrogen-bonded water networks in the cavities of MOFs according to the Grotthuss or proton-hopping mechanism.<sup>213,219,220</sup> Therefore, design, synthesis and modification of MOFs with high chemical stability will provide an excellent platform for investigating their proton-conducting behaviors.

In light of the easy proton release of strong acids such as H<sub>2</sub>SO<sub>4</sub>, UiO-66(SH)<sub>2</sub> was oxidized by H<sub>2</sub>O<sub>2</sub> to afford UiO-66(SO<sub>3</sub>H)<sub>2</sub> with SO<sub>3</sub>H groups covalently linked to the framework, which greatly favored the proton conductivity.<sup>221</sup> At 80 °C and 90% RH, UiO-66(SO<sub>3</sub>H)<sub>2</sub> displayed an extraordinary protonic conductivity of  $8.4 \times 10^{-2}$  S cm<sup>-1</sup> with long-term stability over 96 h, demonstrating its high stability. Ponomareva *et al.* developed a facile method of incorporating nonvolatile H<sub>2</sub>SO<sub>4</sub> and H<sub>3</sub>PO<sub>4</sub> within the pores of MIL-101(Cr) to give H<sub>2</sub>SO<sub>4</sub>@MIL-101 and H<sub>3</sub>PO<sub>4</sub>@MIL-101 for proton conductivity.<sup>222</sup> Specifically, upon confining ~70% H<sub>2</sub>SO<sub>4</sub> (~10 M) or ~80% H<sub>3</sub>PO<sub>4</sub> (~14 M) in MIL-101, the proton conductivity of the resultant composites reached up to  $4.0 \times 10^{-2}$  and  $2.5 \times 10^{-2}$  S cm<sup>-1</sup>, respectively, at low temperature and 20% RH. Not limited to acids, NH<sub>3</sub> and amino were also modified within MOFs *via* the *de novo* process or post-synthetic strategy, improving their proton-conducting ability.<sup>223,224</sup> Wang *et al.* reported a highly stable MOF, MIP-202(Zr), assembled by L-aspartate and Zr<sup>4+</sup> and



Fig. 13 Illustration of the proposed self-adaption mechanism in the flexible MOF (top) and rigid MOF (bottom) with a high density of sulfonic acid sites for proton conduction. Reproduced from ref. 226 with permission from Nature Publishing Group, copyright 2017.

investigated its proton-conduction properties at 90 °C and 95% RH.<sup>224</sup> The interaction between NH<sub>3</sub><sup>+</sup> groups (proton source) and water molecules in the cavities generated an extended hydrogen-bonded network in MIP-202(Zr), endowing it with a desirable and steady proton conductivity of 0.011 S cm<sup>-1</sup>. Hong and co-workers utilized a microwave-assisted solvothermal reaction to obtain Ni-MOF-74, which remained stable even at a pH value as low as 1.8.<sup>225</sup> When Ni-MOF-74 was immersed in sulfuric acid solutions with different pH values, H<sup>+</sup>@Ni-MOF-74 was generated with different proton-conducting properties. Particularly, at pH 1.8, the resultant H<sup>+</sup>@Ni-MOF-74 presented a remarkable proton conductivity of  $2.2 \times 10^{-2}$  S cm<sup>-1</sup> under the conditions of 95% RH and 80 °C. In order to prepare a stable MOF-based electrolyte material with good conductivity at low RH, a highly stable and flexible MOF, BUT-8(Cr)A, was constructed from naphthalene-2,6-dicarboxylate, 4,8-disulfonaphthalene-2,6-dicarboxylate and Cr<sub>3</sub>O(OH)(CO<sub>2</sub>)<sub>6</sub> SBUs.<sup>226</sup> In contrast to other rigid MOFs, BUT-8(Cr)A presented a decent proton conductivity of  $1.27 \times 10^{-1}$  S cm<sup>-1</sup> at 80 °C and 100% RH due to the chemically stable structure with high-density -SO<sub>3</sub>H sites. Its high conductivity could be maintained across a wide range of RH. The authors speculated that this unique feature was ascribable to the water-content-dependent structural transformation in BUT-8(Cr)A (*i.e.* framework 'self-adaption') (Fig. 13).

Although the proton conductivity of certain MOFs has exceeded that of the commercially available Nafion, there still exist many challenging problems to be solved. The key task is to gain a deep understanding of the proton conduction mechanism of MOF materials. Besides, from a practical perspective, more attention should be paid to the commercial development of MOF-based electrochemical devices with desirable mechanical stability.

## 4. Conclusions and perspectives

Over the past two decades, much effort has been dedicated to the *de novo* synthesis of novel MOFs with high stability *via* different strategies, including preparation of high-valent metal-carboxylate or low-valent metal-azolate MOFs, use of azole-containing carboxylate linkers, mixed metal ions, and



hydrophobic ligands, insertion of building blocks, framework interpenetration, *etc.* Beyond these, post-synthetic structural processing and composite material engineering have also been employed to improve the stability of existing MOFs. Amongst MOF chemical stability investigations, water or moisture stability is the most preliminary and important, as handling MOF samples in air (containing moisture) is hardly avoidable for almost all MOF applications. Fortunately, a very large number of MOF-based materials that are stable in air and water have been developed. Moreover, more and more MOFs resistant to harsh chemical environments (acidic/alkaline media) have also been reported. As extreme examples, AITCS-1 shows strong chemical resistance in aqua regia solution over 24 h;<sup>75</sup> the structure of PCN-601 is not disturbed during alkaline treatment in saturated NaOH solution at 100 °C;<sup>25</sup> *etc.* These important breakthroughs together with many other advances give the MOF community great encouragement and confidence. The stability (especially chemical stability) of MOFs, which was once regarded as a critical bottleneck for their real-world applications, may no longer be a severe difficulty and should be addressed in the near future with the continuous efforts of scientists. The preparation of stable MOFs and the strategy of improving MOF stability would become more rational along with powerful prediction of computational chemistry. Accordingly, the applications of MOFs might become widespread and more practicable. As a step further, on the basis of the intriguing properties of stable MOFs, the scope of MOF applications has significantly expanded. In this review, only a minority of MOF applications have been covered. Undoubtedly, the development of MOF stability and its interaction with other research fields will bring more promising and innovative applications.

Although great progress has been made in this field, many challenges remain. Due to the particular requirements of each strategy, it is rather difficult to find a general approach that is applicable to different MOF systems. Also, a number of pre- or post-synthetic modification methods usually cause the change of pore features, which poses challenges for their subsequent applications. To address these issues, some targeted approaches such as hydrophobic surface treatment might be worthy of further investigation. Furthermore, experimental trials can be rationally combined with computational design to afford new framework materials with high stability and particular properties. In addition, so far, there appear to be no systematic investigations on the mechanical stability of MOFs, which is a crucial factor for their industrialization. Maybe, certain specific techniques such as single crystal X-ray diffraction analysis and theoretical calculation will prove helpful in the characterisation of mechanical stability of MOFs. Moreover, it is necessary to establish a unified evaluation index system for assessing the mechanical properties of MOFs. With regard to the experimental stability testing, the combination of multiple means of detection is more reliable to qualify the maintenance of intrinsic properties of MOFs.<sup>227</sup> Particularly, compared with static stability analysis, the investigation of the dynamic change course of MOFs under harsh conditions may be more meaningful.

In the future, the fabrication of very stable MOFs is likely to remain a dream and that would pave the way to their applications to diverse ends. For example, stable and well-designed multivariate MOFs or MOF-based materials, which possess the features of each individual part, can provide extraordinary performance for targeted application. In addition, the precise location control of customisable functional groups within stable MOFs through stepwise synthetic techniques will produce great potential or optimal performance for diverse applications. Moving forward, mechanism analysis related to applications and device-engineering are also the attractive goals for stable MOFs. We also hope that such a Minireview can stimulate the interest of scientists who engage in the interdisciplinary area to explore the value of MOFs from an academic and/or industrial perspective.

In conclusion, MOFs have entered a new period of rapid development and are no longer a category of moisture-sensitive frameworks. The endeavor and progress on enhanced MOF (chemical) stability do greatly promote the development and applications of MOFs. We believe in a bright future for MOF chemistry.

## Conflicts of interest

There are no conflicts to declare.

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