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Two-dimensional conjugated metal–organic frameworks (2D c-MOFs): chemistry and function for MOFtronics

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The 21st century has seen a reinvention of how modern electronics impact our daily lives; silicon-electronics and organic electronics are currently at the core of modern electronics. Recent advances have demonstrated that conductive metal–organic frameworks (MOFs), as another unique class of electronic materials, are emerging to provide additional possibility for multifunctional electronic devices that brings us “MOFtronics”. Typically, two-dimensional conjugated MOFs (2D c-MOFs) are a novel class of layer-stacked MOFs with in-plane extended π -conjugation that exhibit unique properties such as intrinsic porosity, crystallinity, stability, and electrical conductivity as well as tailorable band gaps. Benefiting from their unique features and high conductivity, 2D c-MOFs have displayed great potential for multiple high-performance (opto)electronic, magnetic, and energy devices. In this review article, we introduce the chemical and synthetic methodologies of 2D c-MOFs, intrinsic influences on their electronic structures and charge transport properties, as well as multifunctional applications of this class of materials for MOFtronics and potential power sources for MOFtronics. We highlight the benefits and limitations of thus-far developed 2D c-MOFs from synthesis to function and offer our perspectives in regard to the challenges to be addressed.

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1. Introduction

Metal–organic frameworks (MOFs) represent crystalline coordination polymers,^{1–4} which are a class of porous materials with high surface area that were developed three decades ago.

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Chemical tunability in the porous structures^{5–7} allows various applications in gas storage,^{8,9} separation,^{10,11} catalysis,^{12,13} proton conduction,¹⁴ sensor,^{15,16} etc. The majority of traditional MOFs are electrical insulators which limit their utilization in multifunctional electronic devices.^{17,18} Two-dimensional conjugated MOFs (2D *c*-MOFs) refer to layer-stacked MOFs comprising *ortho*-substituted conjugated building blocks (e.g. benzene, triphenylene) and square-planar linkages with high in-plane conjugation and weak out-of-plane van der Waals interactions. The synthesis of 2D *c*-MOFs has been reported accompanied by intrinsic charge transport since 2012.¹⁹ The extended π -conjugation in the 2D plane is capable of facilitating the delocalization of charge carriers within the network that is beneficial to high mobility and conductivity.²⁰ Currently, over ten types of 2D *c*-MOFs have been developed based on conjugated ligands and linkages.^{21,22} Fig. 1 shows the timeline of the contribution to the synthesis of 2D *c*-MOFs and the discovery of key properties from the community. Besides the preserved characteristics of traditional MOFs, such as tunable porosity, versatile structures and well-defined active sites, 2D *c*-MOFs have exhibited unique chemical and physical features such as high stability, electrochemical activity, photoactivity, tailorable band gaps, superior electrical conductivity, ferromagnetic ordering and topological state,^{23–27} thereby enabling 2D *c*-MOFs for broad applications in MOFtronics, such as (opto)electronics^{28–31} and spintronics,^{32–34} as well as the potential power sources for MOFtronics including battery^{35,36} and supercapacitor devices.^{37,38}

This review article covers the chemistry and charge transport of 2D *c*-MOFs by consideration of structural diversity and structure–electronic property relationship based on the influence of molecular design (type and geometry of ligand, metal) and material structure (redox-activity in linkage, layer stacking and arrangement) on intrinsic charge transport properties.



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well as 2D carbon-rich conjugated polymers for optoelectronic applications, energy storage and conversion, and new energy devices and technologies. He has published more than 480 research articles which have attracted more than 52 000 citations with an H-index of 109.

An overview of the synthetic methodologies including hydro-/solvothermal methods and wet-interface-assisted strategies is further offered. Next, device fabrications and the applications of 2D *c*-MOFs for MOFtronics are presented covering (opto)electronics (e.g. field-effect transistors, superconductors, chemiresistors, photodetectors), spintronics, thermoelectrics, and energy storage devices with the discussion of their achievements and existing challenges. Critical directions for future research are proposed towards MOFtronics within the community.

2. Chemical methodologies of 2D *c*-MOFs

After the first 2D *c*-MOF reported by Yaghi group in 2012,¹⁹ various *ortho*-substituted benzene- and triphenylene (TP)-based building blocks have been developed/employed for the formation of 2D *c*-MOFs, such as 1,2,3,4,5,6-hexahydroxybenzene (HHB),³⁹ 1,2,3,4,5,6-hexaaminobenzene (HAB),⁴⁰ 1,2,3,4,5,6-benzenehexathiol (BHT, also denoted asHTB),⁴¹ 1,2,3,4,5,6-benzenehexaselenol (HSeB),⁴² 2,3,6,7,10,11-hexahydroxytriphenylene (HHTP),¹⁹ 2,3,6,7,10,11-hexaaminotriphenylene (HATP),^{43,44} 2,3,6,7,10,11-triphenylenehexathiol (THT, also denoted as HTTP),⁴⁵ 2,3,6,7,10,11-triphenylenehexaselenol (HSeTP).⁴⁶ Larger conjugated monomers have been developed and explored for the synthesis of 2D *c*-MOFs as well (Fig. 2), e.g. 2,3,8,9,14,15-hexahydroxytrinaphthylene (HHTN),⁴⁷ 2,3,7,8,12,13-hexahydroxytruxene (HHTX),⁴⁸ metal(II) 2,3,8,9,14,16-hexaaminohexaazatrinaphthylene (HAHATNM'₃, M' = Cu, Ni, Co),⁴⁹ 1,2,3,4,5,6,7,8,9,10,11,12-perthiolated coronene (PTC),³² metal(II) 2,3,9,10,16,17,23,24-octahydroxyphthalocyanine (M'Pc(OH)₈, M' = H₂, Cu, Fe, Ni, Zn),^{33,50–53} metal(II) 2,3,9,10,16,17,23,24-octaaminophthalocyanine (M'Pc(NH₂)₈, M' = Cu, Ni),^{54,55} metal(II) 3,4,12,13,21,22,30,31-octahydroxynaphthalocyanine (M'NPc(OH)₈, M' = Ni).⁵¹ Besides, a non-planar conjugated ligand, 2,3,6,7,10,11,14,15-octahydroxydibenzo[*g,p*]chrysene (DBC(OH)₈),⁵⁶ was included into the 2D *c*-MOF family recently.

In fact, the ligands in most of the thus-far constructed 2D *c*-MOFs are partially oxidized to form charge-neutral frameworks, instead of the generally displayed structures which correspond to negatively charged linkages (e.g. Cu₃(HHTP)₂, Fig. 10 in Section 4.1).^{19,23,25,40,55–58} Meanwhile, negatively charged 2D *c*-MOFs are also present in this emerging field, such as Cu₃(HHB)₂ (Section 3.4).^{39,59} Among the reported 2D *c*-MOFs, three types of linkages containing chelating functional groups were usually adopted: metal-bis(dioxolene) (abbreviated as MO₄), metal-bis(diimine) (MN₄, also termed as metal-bis(iminobenzosemiquinoid)), and metal-bis(dithiolene) (MS₄) complexes (M = Fe, Co, Ni, Cu, Zn, Pd, or Pt). Due to the presence of both neutral and charged MO₄-linked networks, the term of dioxolene instead of the normally applied semiquinoid is likely more reasonable.^{25,60} In addition, metal-diimine-dithiolene (MN₂S₂) mixed linkage and metal-bis(diselenolene) (MSe₄)



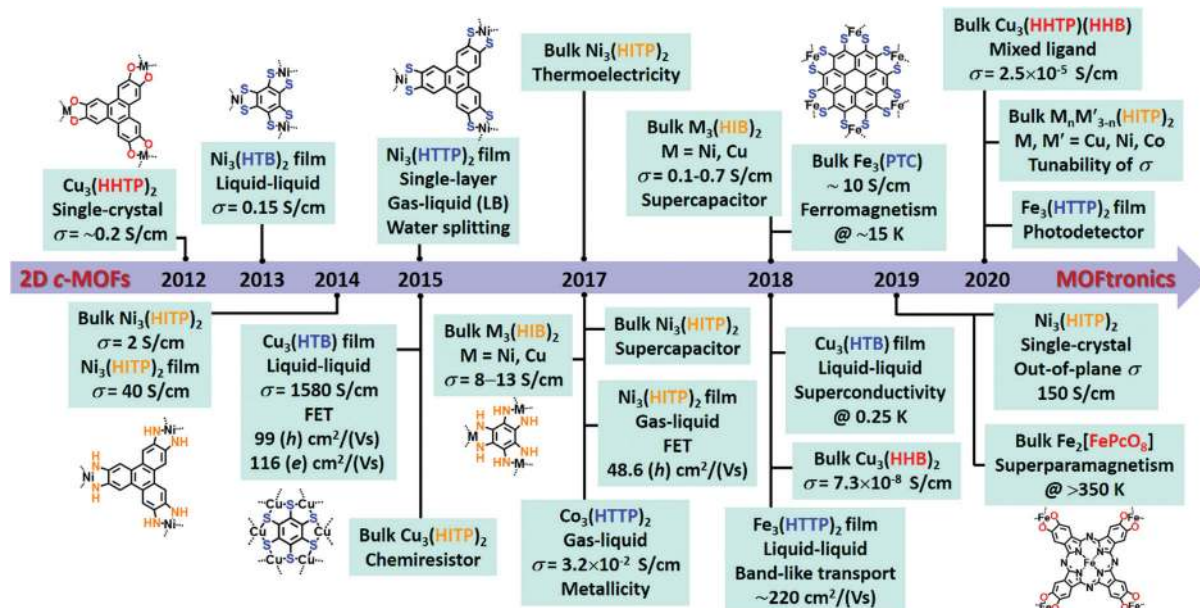


Fig. 1 Timeline in the development of 2D c-MOFs from synthesis to MOFtronics. The ligands containing $-\text{OH}$, $-\text{NH}_2$, and $-\text{SH}$ are marked in red, orange, and blue colour, respectively. The $\text{Cu}_3(\text{HHTP})(\text{HHB})$ 2D c-MOF is also denoted as $\text{Cu}_3(\text{HHTP})(\text{THQ})$ (THQ = tetraoxyquinone in 2D c-MOFs). For 2D c-MOFs, HHTP represents 2,3,6,7,10,11-hexaaoxytriphenylene instead of 2,3,6,7,10,11-hexahydroxytriphenylene for the monomer.

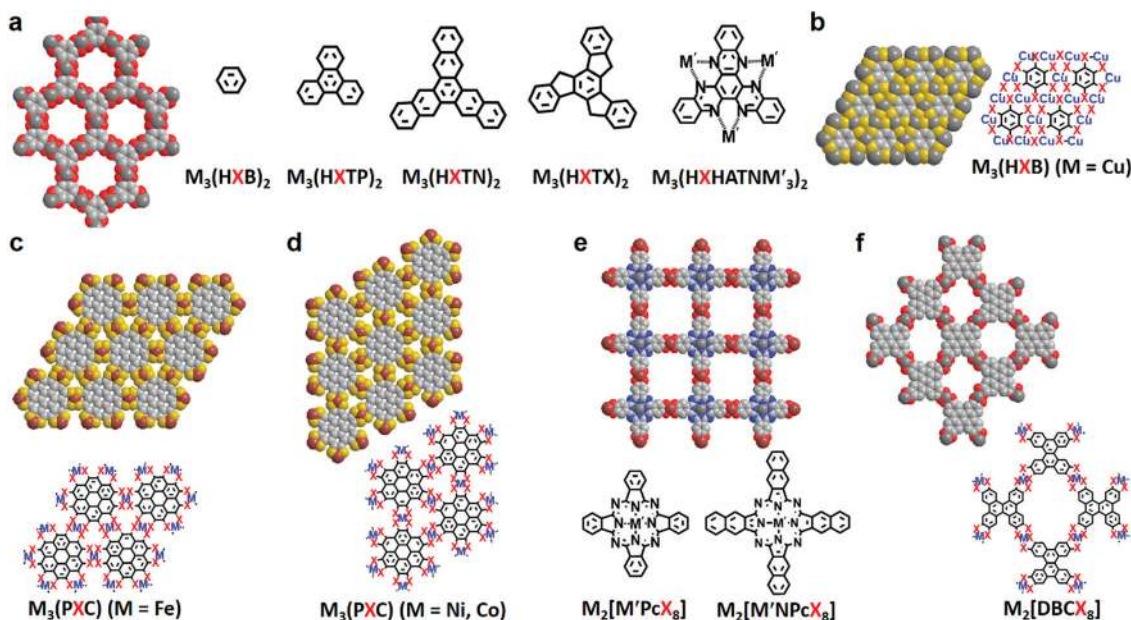


Fig. 2 Structure of various 2D c-MOFs. (a) 2D c-MOFs with honeycomb lattice. $\text{M}_3(\text{HXB})_2$, $\text{M}_3(\text{HXTP})_2$, $\text{M}_3(\text{HXTN})_2$, $\text{M}_3(\text{HXTX})_2$, $\text{M}_3(\text{HXHATNM}'_3)_2$. (b–d) Non-porous 2D c-MOFs. $\text{M}_3(\text{HXB})$ ($\text{M} = \text{Cu}$; $\text{X} = \text{S, Se}$) and $\text{M}_3(\text{PXC})$ ($\text{M} = \text{Fe, Ni, Cu}$; $\text{X} = \text{S}$). (e) 2D c-MOFs with square lattice. $\text{M}_2[\text{M}'\text{PcX}_8]$ ($\text{M} = \text{Fe, Co, Ni, Cu, Zn}$; $\text{M}' = \text{Fe, Ni, Cu, Zn}$; $\text{X} = \text{O, NH}$) and $\text{M}_2[\text{M}'\text{NPcX}_8]$ ($\text{M} = \text{Ni, Cu}$; $\text{M}' = \text{Ni}$; $\text{X} = \text{O}$). (f) Non-planar 2D c-MOF. $\text{M}_2[\text{DBCX}_8]$ ($\text{M} = \text{Cu}$; $\text{X} = \text{O}$).

linkage were also investigated in few reports.^{42,46,61–63} Among these various linkages, Cu and Ni represented the majority of metal nodes, while Fe and Co were widely utilized for the construction of MS_4 . By contrast, the utilization of Zn, Pd, and Pt remains very limited.^{53,64–67}

Due to the increasing ligand library and the diverse compositions, a uniform description shall be helpful for the development of 2D c-MOF families. Based on the mostly accepted style, we label the benzene- and triphenylene-based 2D c-MOFs as $\text{M}_n(\text{HXL})_m$ (Fig. 2a), in which M and L represent the type of



metal node and abbreviation of the building block (B for benzene, TP for triphenylene) respectively for a smallest repeating unit in their chemical structures, while H and X correspond to six substituent X in each ligand (X = O, NH, S, and Se). This can be also applied to trinaphthylene (TN)-, truxene (TX)- and hexaazatrinaphthylene (HATN)-based systems as $M_n(\text{HXTN})_m$, $M_n(\text{HXTX})_m$, and $M_n(\text{HXHATNM}')_{3m}$ (Fig. 2a). Among all these systems, n and m —the number of metal nodes and ligands—are confirmed to be 3 and 2 respectively in general, except for rare instances including $\text{Ni}_3(\text{HHTP})_4$, $\text{Co}_9(\text{HHTP})_4$ (Fig. 10c),¹⁹ $\text{Cu}_3(\text{HTB})$, and $\text{Cu}_3(\text{HSeB})$ ($n = 3$, $m = 1$, Fig. 2b).^{31,42,68} For a non-porous 2D c -MOF based on the fully substituted coronene,³² $\text{M}_3(\text{PXC})$ is adopted (Fig. 2c and d), in which P and C represent full substitution in the ligand, and the abbreviation of coronene, respectively. However, the octa-substituted metallophthalocyanine ($\text{M}'\text{Pc}$) and dibenzo[*g,p*]chrysene (DBC) are complicated systems. Instead of a scientific abbreviation for the ligands, the visualized one ($\text{M}'\text{PcX}_8$ for $\text{M}'\text{Pc}$,^{33,50–52,54,55} DBCX_8 for DBC⁵⁶) is applied, thus leading to $\text{M}_2[\text{M}'\text{PcX}_8]$ (Fig. 2e) and $\text{M}_2[\text{DBCX}_8]$ (Fig. 2f) 2D c -MOFs, respectively. This address is also applicable to the larger metallonaphthalocyanine ($\text{M}'\text{Npc}$) system (Fig. 2e).⁵¹ Notably, three-dimensional (3D) MOFs without layer-stacked structures could be also constructed based on planar ligands (e.g. HHTP, $\text{CoPc}(\text{OH})_8$) through metal-catecholate linkages.^{69–72} Therefore, the structure of 2D c -MOFs highly depends on the employed ligands, metal centers and synthetic conditions.⁷³

3. Chemical structure and electronic property relationship in 2D c -MOFs

Conductivity (σ) represents the transport of carriers of electrons (e) and holes (h), which can be quantified with mobility (μ) and density (n): $\sigma = e(n_e\mu_e + n_h\mu_h)$. Free charge carriers in 2D c -MOFs can be generated through thermal/photo-population or charge carrier injection through redox reactions.^{23,74,75} In this context, both the metal nodes (Frontier orbitals) and the ligands can serve as the sources. The potential charge transport pathways in 2D c -MOFs were ascribed as: through-bond, extended conjugation, through-space, and hopping.²⁵ Particularly, through-bond or extended conjugation pathways in the 2D plane and through-space channel (strong π - π stacking and weak metal-metal interactions) dominate the intrinsic conductivity of 2D c -MOFs, while hopping stems from the charge carrier migration through grain boundaries, defects, *etc.* Many inspiring review articles have provided in-depth understanding on the above transport mechanism as well as the description of current characterization methods toward conductivity measurement, such as two-probe and four-probe (linear four-probe or van der Pauw) configurations.^{17,20,22–25,60,76} Table 1 lists the currently reported 2D c -MOFs and the related studies on charge transport, including the sample form, characterization method, conductivity and mobility. Hereby, we would like to discuss the influence of molecular design (type and geometry of ligand, metal) and material structure (redox-activity in linkage, layer stacking and

arrangement) as well as crystallinity on the charge transport properties for thus-far reported materials.

3.1 Effect of ligand type

The current ligands utilized for the construction of 2D c -MOFs have been mainly based on the planar conjugated building blocks with symmetrical functional groups of $-\text{OH}$, $-\text{NH}_2$, and $-\text{SH}$. Varying the functional groups results in increased bond strength between metals and linkers from MO_4 to MN_4 to MS_4 ,²⁵ thus leading to the apparent enhancement of the in-plane charge transport along the 2D network. Density functional theory (DFT) calculations indicated that the linker substitution in MX_4 (X = O, NH, and S) also affects the dynamic stability of the frameworks; $\text{Zn}_3(\text{HTB})$ is more stable than $\text{Zn}_3(\text{HSeB})$ whilst $\text{Zn}_3(\text{HHB})$ is the most unstable one.⁷⁷ In addition, energy matching of the d orbitals of different transition metals with the π -conjugated ligands generally generates the difference in π -d hybridization, hence tuning the delocalization of carriers in the 2D plane to a different extent (Section 3.3). Among these 2D c -MOFs, MO_4 -linked ones exhibited apparently inferior charge transport to the related 2D c -MOFs linked by the MN_4 coordination complex. For instance, charge-neutral MO_4 -linked $\text{Cu}_3(\text{HHTP})_2$ could display broad conductivity values in the range of 10^{-4} to 1 S cm^{-1} measured by 2-/4-probe measurement in their pellet form at room temperature. By contrast, the pellet samples of MN_4 -linked $\text{Cu}_3(\text{HITP})_2$ could exhibit higher conductivity values in the range of 0.2 – 0.75 S cm^{-1} . Regarding Pc-based $\text{M}_2[\text{M}'\text{PcX}_8]$ 2D c -MOFs, the MN_4 -linked samples also showed superior conductivity than the MO_4 -linked ones. The MS_4 complex presents the highest π -d hybridization among these linkages, and therefore thus-far reported MS_4 -linked 2D c -MOFs have been demonstrated to exhibit excellent conductivity and mobility. For example, the $\text{Ni}_3(\text{HTB})_2$ film showed a conductivity of $\sim 160 \text{ S cm}^{-1}$ and the non-porous $\text{Cu}_3(\text{HTB})$ film could even exhibit a conductivity as high as $\sim 2500 \text{ S cm}^{-1}$ (shown in Table 1). Besides the conductivity, another typical MS_4 -linked sample, $\text{Fe}_3(\text{HTTP})_2$, was observed to present a notable band-like transport with record high charge mobility $> 200 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ amongst the reported 2D c -MOFs due to the high π -d conjugation.⁷⁸ The $\text{Co}_3(\text{HTTP})_2$ 2D c -MOF was recorded to possess a negative temperature-resistivity relationship in the high temperature region and a positive trend at low temperature for semiconducting and metallic behavior, respectively.^{79,80} Recently, MSe_4 -linked bulk 2D c -MOFs have been developed and displayed lower conductivity (e.g. $\text{Cu}_3(\text{HSeB})$: 110 S cm^{-1} , $\text{Co}_3(\text{HSeTP})_2$: $10^{-6} \text{ S cm}^{-1}$) than those of MS_4 -linked bulk 2D c -MOFs (Table 1).^{42,46}

Varying the functional groups will also vary the coordination bond length between metal nodes and linkers in the MX_4 -linkages due to the difference of the atom radius of the O, N, S, and Se (Se > S > O > N), which plays also an important role in the charge transport and other physical properties. For instance, for some selected square-planar Ni-coordinated complexes (Fig. 3A–C), the bond lengths are 1.866, 1.822–1.826, and 2.132–2.170 Å for Ni–O, Ni–N, and Ni–S bonds, respectively.^{81–83}



Table 1 Summary of the literature reported electrical conductivities and mobilities for 2D *c*-MOFs

2D <i>c</i> -MOF	Building block	Type	Method	σ^a (S cm ⁻¹)	μ (cm ² V ⁻¹ s ⁻¹)	Ref.
Metal-bis(dioxolene) linked 2D <i>c</i> -MOFs						
Cu ₃ (HHB) ₂	B	Pellet	vdP ^b	7.8 × 10 ⁻⁸	2.4 (Hall, <i>h</i> ^d)	39
		Pellet	—	2.2 × 10 ⁻⁹		87
Cu ₃ (HHTP) ₂	TP	Pellet ^c	vdP	1.5 × 10 ⁻⁷		88
		Pellet	2-Probe	1.0 × 10 ⁻²		89
		Pellet	2-Probe	3.0 × 10 ⁻³		90
		Pellet	2-Probe	2.0 × 10 ⁻³		91
		Pellet	4-Probe	2.0 × 10 ⁻²		92
		Pellet	vdP	4.5 × 10 ⁻²		93
		Pellet	vdP	2.7 × 10 ⁻²		94
		Pellet	—	0.09-0.1		95
		Film	2-Probe	2.0 × 10 ⁻²	96	
		Film	4-Probe	0.29	97	
Ni ₉ (HHTP) ₄	TP	Film	—	10 ⁻⁴	98	
		Nanorod ^e	4-Probe	0.21	19	
		Nanorod ^e	4-Probe	1.5	95	
		Nanoflake ^e	2-Probe	0.5	95	
		Pellet	2-Probe	6.0 × 10 ⁻³	99	
		Pellet	4-Probe	1.0 × 10 ⁻²	100	
		Pellet	4-Probe	0.1	92	
		Pellet	4-Probe	0.26	101	
		Pellet	vdP	6.8 × 10 ⁻³	94	
		Film	vdP	1.1 × 10 ⁻³	94	
Co ₉ (HHTP) ₄	TP	Hybrid	2-Probe	1.6 × 10 ⁻⁴	102	
		Pellet	2-Probe	2.0 × 10 ⁻³	99	
		Pellet	4-Probe	2.7 × 10 ⁻⁶	92	
		Pellet	vdP	3.2 × 10 ⁻²	94	
		Film	vdP	3.3 × 10 ⁻³	94	
Cu ₃ (HHTP)(HHB)	TP/B	Pellet	2-Probe	2.5 × 10 ⁻⁵	59	
Cu ₃ (HHTN) ₂	TN	Pellet	2-Probe	9.6 × 10 ⁻¹⁰	47	
Cu ₃ (HHTX) ₂	TX	Pellet	2-Probe	8.3 × 10 ⁻⁴	48	
Cu ₂ [DBCO ₈]	DBC	Pellet	—	1 × 10 ⁻²	56	
Cu ₂ [CuPcO ₈]	Pc	Pellet	2-Probe	9.4 × 10 ⁻⁸	50	
Cu ₂ [NiPcO ₈]	Pc	Pellet	2-Probe	1.6 × 10 ⁻⁶	50	
Cu ₂ [NiPcO ₈]	Pc	Pellet	4-Probe	1.4 × 10 ⁻²	51	
Ni ₂ [NiPcO ₈]	Pc	Pellet	4-Probe	7.2 × 10 ⁻⁴	51	
Fe ₂ [FePcO ₈]	Pc	Pellet	vdP	~1 × 10 ⁻⁵	~0.1 (Hall, <i>h</i>)	33
		Pellet	—	—	15 (TRTS)	33
Cu ₂ [NiNPcO ₈]	NPc	Pellet	4-Probe	3.1 × 10 ⁻²		51
Ni ₂ [NiNPcO ₈]	NPc	Pellet	4-Probe	1.8 × 10 ⁻²		51
Metal-bis(diimine) linked 2D <i>c</i> -MOFs						
Cu ₃ (HIB) ₂	B	Pellet	4-Probe	0.1	48.6 (<i>h</i>) 45.4 (<i>h</i>)	38
		Pellet	vdP	13		40
Ni ₃ (HIB) ₂	B	Pellet	4-Probe	0.7		38
		Pellet	vdP	8		40
Co ₃ (HIB) ₂	B	Pellet	4-Probe	0.1–1.57		35
Cu ₃ (HITP) ₂	TP	Pellet	2-Probe	0.2		28
		Pellet	4-Probe	0.75		58
Ni ₃ (HITP) ₂	TP	Pellet	2-Probe	2		57
		Pellet	4-Probe	39		101
		Pellet	4-Probe	1.896		103
		Pellet	—	0.5		104
		Pellet	vdP	50		37
		Pellet	4-Probe	55.4		58
		Pellet	vdP	58.8		105
		Film	—	37.2		104
		Film	2-Probe	8.5 × 10 ⁻³		106
		Film	2-Probe	0.23	106	
		Film	vdP	40	57	
		Co ₃ (HITP) ₂	TP	Hybrid	2-Probe	2.6 × 10 ⁻³
Nanorod ^e	4-Probe			150	95	
Film	FET ^g			—	29	
Film	FET ^g			—	107	
Pellet	4-Probe			8 × 10 ⁻⁴	103	
Co ₃ (HITP) ₂	TP	Pellet	4-Probe	0.024	58	
Co ₃ Cu ₆ (HITP) ₂ ^f	TP	Pellet	4-Probe	5.8 × 10 ⁻³	58	
Ni ₃ (HXHATNNi ₃) ₂	HATN	Pellet	4-Probe	2		
Ni ₂ [NiPc(NH) ₈]	Pc	Film	4-Probe	0.2	54	
Ni ₂ [CuPc(NH) ₈]	Pc	Pellet ^c	vdP	~1 × 10 ⁻⁴	1.6 (Hall, <i>h</i>)	55



Table 1 (continued)

2D <i>c</i> -MOF	Building block	Type	Method	σ^a (S cm ⁻¹)	μ (cm ² V ⁻¹ s ⁻¹)	Ref.
Metal-bis(dithiolene) linked 2D <i>c</i> -MOFs						
Cu ₃ (HTB)	B	Pellet	4-Probe	48–280	116 (e), 99 (h)	108
		Film	4-Probe	1580		31
		Film	FET ^g	—		31
		Film	4-Probe	2500		68
Ni ₃ (HTB) _h	B	Pellet	2-Probe	0.15	116 (e), 99 (h)	74
		Pellet	2-Probe	6.7 × 10 ⁻³		74
		Pellet	2-Probe	2.8		75
		Pellet	2-Probe	160		75
Pd ₃ (HTB) ₂	B	Film	4-Probe	2.8 × 10 ⁻²	116 (e), 99 (h)	65
		Pellet	—	2.4 × 10 ⁻⁸		109
Cu ₃ (HTTP) ₂	TP	Pellet	—	3.6 × 10 ⁻⁴		109
Ni ₃ (HTTP) ₂	TP	Pellet	—	2.4 × 10 ⁻⁹		109
Co ₃ (HTTP) ₂	TP	Pellet	vdP	1.4 × 10 ⁻³	229 (Hall, h) 211 (TRTS)	79
		Film	vdP	3.2 × 10 ⁻²		79
		Film	vdP	3.4 × 10 ⁻²		78
		Film	—	—		78
Fe ₃ (HTTP) ₂	TP	Film	vdP	0.2	229 (Hall, h) 211 (TRTS)	80
		Film	2-Probe	3.9 × 10 ⁻⁶		64
		Pellet	vdP	10		32
		Pellet	4-Probe	9		110
Co ₃ (PTC)	C	Pellet	4-Probe	45		111
Metal-diimine-dithiolene linked 2D <i>c</i> -MOFs						
Ni ₃ (TITTB) ₂ _h	B	Pellet	vdP	0.1	116 (e), 99 (h)	63
		Pellet	vdP	3 × 10 ⁻⁶		62
Metal-bis(diselenolene) linked 2D <i>c</i> -MOFs						
Cu ₃ (HSeB)	B	Pellet	4-Probe	110		42
Co ₃ (HSeTP) ₂	TP	Pellet	2-Probe	10 ⁻⁶		46

^a Electrical conductivity at room temperature. ^b Van der Pauw. ^c Nanosheets. ^d Hall effect. ^e Device based on single-crystal. ^f $a = 2.47$, $b = 0.53$. ^g Field-effect transistor. ^h Reduced 2D *c*-MOF with charge state of -1 . ⁱ Oxidized 2D *c*-MOF with charge state of 0.



Fig. 3 The bond length of metal-linkers (O, NH, S) in the MX₄ linkage. Selected structures of A,⁷⁵ B,⁷⁴ and C⁷⁶ refer to reported metal-complex small molecules.

While for 2D *c*-MOFs, Ni₃(HITP)₂ and Ni₃(HTTP)₂, powder X-ray diffraction (PXRD) indicated prominent (100) peak (Cu-K α radiation) with $2\theta = 4.7^\circ$ (1.88 nm) and 4.5° (1.96 nm) respectively, indicative of larger lattice constant of Ni₃(HTTP)₂ than Ni₃(HITP)₂ due to the presence of its longer Ni–S bond.^{29,45,57} A similar phenomenon was revealed by the DFT calculations: ~ 0.1 nm smaller lattice constant of Mn₃(HIB)₂ than Mn₃(HTB)₂.^{84,85} In a monolayer of Mn₃(HIB)₂, the shorter distance between Mn²⁺ ions is expected to enhance the magnetic coupling; compared with MS₄, N atoms in the MN₄ linkage are more effective at mediating the magnetic couplings. As a result, the monolayer of Mn₃(HIB)₂ is predicted to possess a transition temperature for ferromagnetic coupling at ~ 450 K, which considerably surpasses that of 212 K for a single-layer of Mn₃(HTB)₂.⁸⁴ Nevertheless, it remains not

sufficient to quantitatively compare the influence of the MX₄-linkages on conductivity, due to the few examples and the heterogeneous sample morphologies, potential doping, as well as the lack of unique features of chemical composition and structural geometry for individual samples.

3.2 Effect of ligand geometry

Ligand geometry and symmetry are of key significance for the construction of regular frameworks of 2D *c*-MOFs. Currently, three types of networks have been developed for 2D *c*-MOFs, including hexagonal, square, and honeycomb lattices with polygons of triangles, squares, and hexagons, respectively (Fig. 4). For the construction of hexagonal, square, or honeycomb frameworks, symmetric ligands with C₆ (e.g. PTC), C₄ (e.g. M'PcX₈), or C₃ (e.g. HXTP) geometry respectively are required. Importantly, the resulting framework geometry is relevant to the electronic structure of the achieved 2D *c*-MOF. For example, the hexagonal and square geometries correlate with a structure, where only one type of vertex exists in each unit cell with six neighboring triangles or four neighboring squares (Fig. 4a and b, unit cells are marked red in the middle figures). DFT calculations indicated that only one type of band is presented in the band structures for both geometries.⁸⁶ On the other hand, each unit cell contains two vertices in different environments in a honeycomb network (Fig. 4c). As a result,



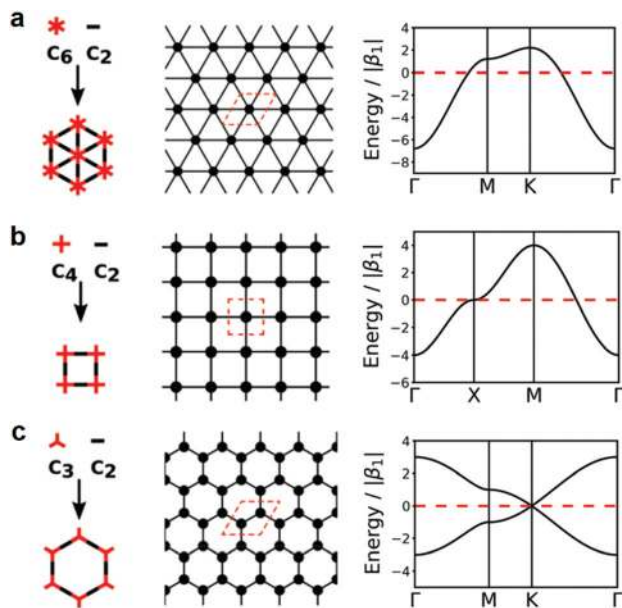


Fig. 4 Topological diagrams of the networks and 2D representations with different geometries (unit cells are marked in red) as well as corresponding band structures. (a) Hexagonal. (b) Square. (c) Honeycomb. C2 monomer refers to metal ions here ref. 86. Copyright 2020, the Royal Society of Chemistry.

two bands appear in the energy band diagram forming the Dirac point.

Though a dual-porous kagome network (Fig. 5a, left) is not presented among thus-far reported 2D *c*-MOFs, the metal atoms in the honeycomb frameworks (e.g. $M_3(\text{HXB})_2$, $M_3(\text{HXTP})_2$) are arranged in the kagome lattice (Fig. 5a, middle), which exhibits a band structure including a Dirac point and a flat band above the Dirac bands (Fig. 5a, right).⁸⁶ The framework with extended π -conjugation in a kagome lattice is predicted to exhibit non-trivial physical properties. For instance, first-principles calculations indicated that a single-layer of $\text{Ni}_3(\text{HTB})_2$ possesses the typical kagome bands: a flat band above two Dirac bands (Fig. 5b and c). The local density of state of a semi-infinite $\text{Ni}_3(\text{HTB})_2$ framework for spin-up and spin-down components indicated nontrivial topological edge states (Fig. 5d and e), indicative of the possible existence of an organic topological insulator.¹¹² Besides, a monolayer of $\text{Ni}_3(\text{HITP})_2$ (Fig. 8b) or $\text{Cu}_3(\text{HITP})_2$ also exhibited Dirac bands near the Fermi level.¹¹³

In addition to the type and geometry of the ligand, the ligand size (e.g. benzene, triphenylene, trinaphthylene), which correlates with diverse densities of building blocks and linkages as well as the π -conjugation degree in the network, is also an important parameter for the electronic structures. However, accompanied by other factors, such as the lattice, layer distance, porous density, etc., the influences of ligand size remain elusive. Besides, the control of π -conjugation degree in the building blocks (e.g. benzene, triphenylene, trinaphthylene, coronene, phthalocyanine) is expected to efficiently vary the conductivity and mobility of 2D *c*-MOFs in both in-plane and cross-plane directions. Nonetheless, because of the differences in the geometry and size of the ligand, a proper relationship between π -conjugation and conductivity has been so far elusive.

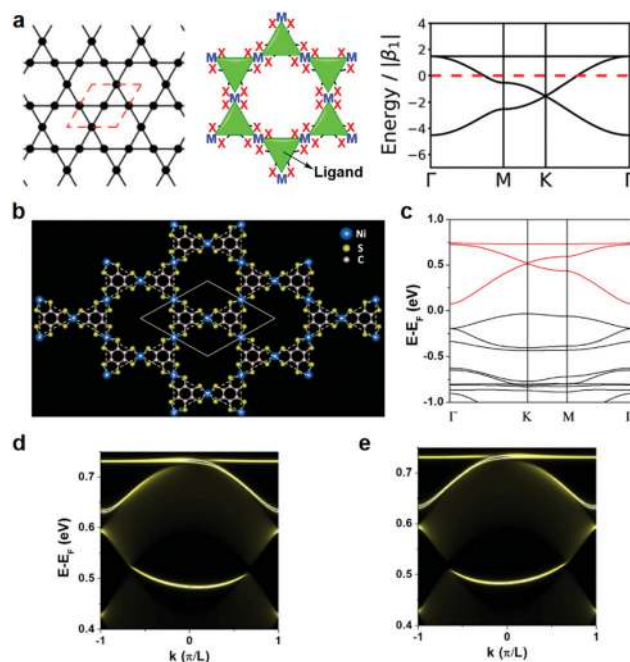


Fig. 5 Kagome lattice in 2D *c*-MOFs. (a) Topological diagram of the kagome network (left), the kagome lattice of metal atoms in a honeycomb network (middle), and the band structure for a kagome network (right).⁸⁶ Copyright 2020, the Royal Society of Chemistry. (b) Kagome lattice of the $\text{Ni}_3(\text{HTB})_2$ monolayer. The dashed lines outline the kagome lattice. (c) The band structure of the $\text{Ni}_3(\text{HTB})_2$ monolayer. (d and e) The semi-infinite edge states of the spin-up and spin-down components for the $\text{Ni}_3(\text{HTB})_2$ monolayer, respectively. Reproduced with permission ref. 112. Copyright 2013, American Chemical Society.

3.3 Effect of metals

The metal centers have been demonstrated to play an important role in the electronic properties of the 2D *c*-MOFs. It is already known that the conductive 3D MOFs with mixed valence Fe atoms ($\text{Fe}^{2/3+}$) show superior electrical conductivity to the pure Fe^{2+} -bridged ones.^{114–117} For instance, fractional reduction of Fe^{3+} atoms in $\text{Fe}_2(\text{BDP})_3$ (BDP = 1,4-benzenedipyrazolate) to access mixed valence state of $\text{Fe}^{2/3+}$ in the framework resulted in a nearly four orders of magnitude higher conductivity along a single crystallographic axis.¹¹⁵ For 2D *c*-MOF systems, many examples possess mixed valence as well, such as $\text{Fe}^{2/3+}$, $\text{Co}^{2/3+}$, and $\text{Cu}^{1/2+}$.^{33,39,56,80,118} Nevertheless, the valence influence on the electronic properties has not been systematically investigated due to the lack of control on the metal valence. For the moment, we would assume a similar conclusion in 2D *c*-MOFs compared with conductive 3D MOFs.

On the other hand, the metal type also plays an important role in the charge transport performance of 2D *c*-MOFs.^{58,119} For instance, first-principles calculations indicated that the band gaps of $M_3(\text{HTB})$ ($M = \text{Mg}, \text{Ca}, \text{Zn}, \text{Cd}, \text{Ge}, \text{and Sn}$) ranged from 1.7 to 3.2 eV.¹²⁰ Besides, the π -d hybridization between d (metal) and π (HTB) bands can considerably delocalize the wave function of band edge states and is of importance in determining their hole effective masses. More specifically, as the π -d coupling increases from the elements of group IIA (Mg, Ca) to



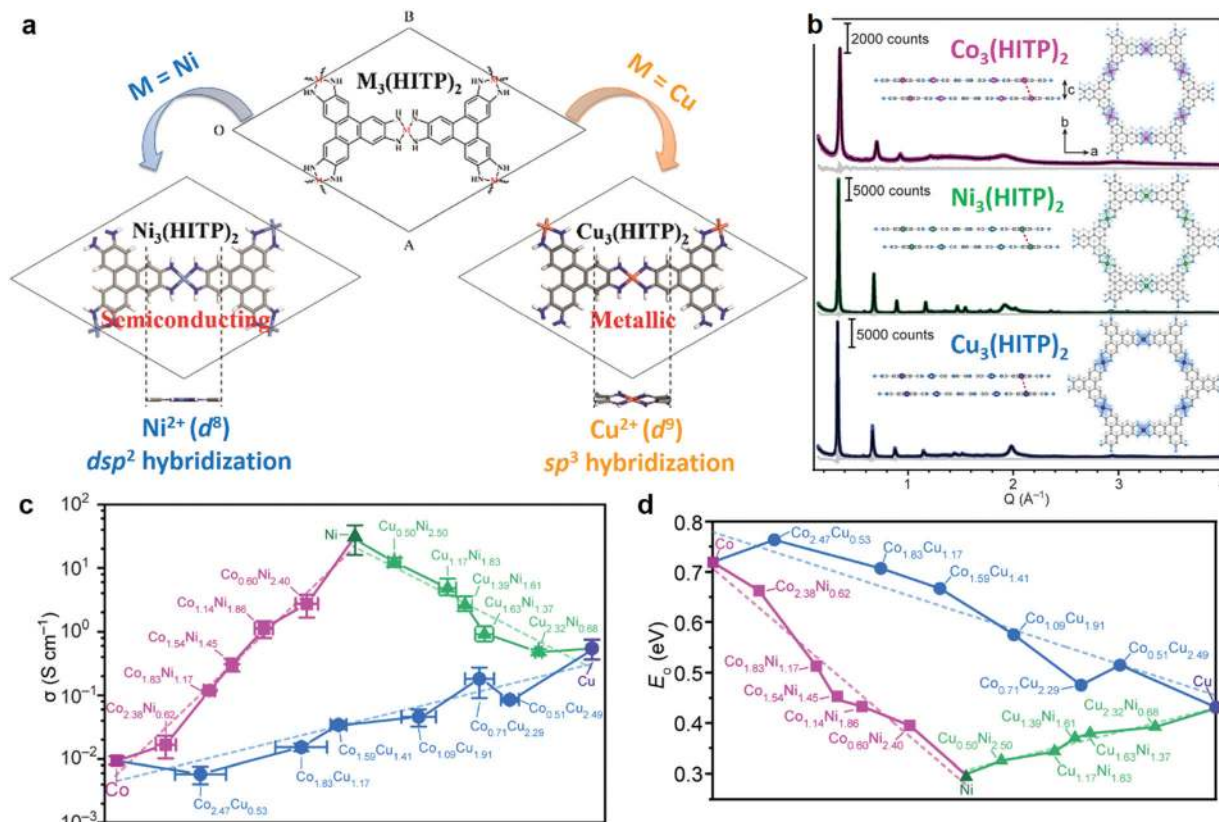


Fig. 6 Metal substitution in 2D *c*-MOFs. (a) Illustration of the geometry of $M_3(\text{HITP})_2$ ($M = \text{Ni}, \text{Cu}$) by substituting Ni (left) with Cu (right).¹¹³ Copyright 2015, the PCCP Owner Societies. (b) Synchrotron X-ray diffraction patterns and Pawley refinements for $M_3(\text{HITP})_2$. The insets show the bilayer of $M_3(\text{HITP})_2$ with reference to the interlayer displacement. (c and d) Electrical conductivity and optical bandgap of $M_nM'_{3-n}(\text{HITP})_2$, respectively. Dashed lines refer to linear fitting. Reproduced with permission ref. 58. Copyright 2020, American Chemical Society.

IVA (Ge, Sn) to IIB (Zn, Cd), the hole effective masses decrease significantly. While for electron transport, from group IIB (Zn, Cd) to IIA (Mg, Ca) to IVA (Ge, Sn), the electron effective masses increase.¹²⁰ Nonetheless, these 2D *c*-MOFs have so far not been constructed in the laboratory. In addition, great efforts from DFT calculations have been also devoted to the well-known $M_3(\text{HITP})_2$ ($M = \text{Ni}, \text{Cu}$) 2D *c*-MOFs regarding the metal substitution. The substitution from Ni to Cu in this system will change both the geometries and the electronic band structures (Fig. 6a). The Ni atoms in $\text{Ni}_3(\text{HITP})_2$ adopt the dsp^2 hybridization in a square-planar geometry, thus forming a planar 2D framework.¹¹³ Nevertheless, for $\text{Cu}_3(\text{HITP})_2$, the Cu atoms adopt the sp^3 hybridization leading to a distorted network in a specific square-grid coordination geometry. In addition, a single-layer of $\text{Cu}_3(\text{HITP})_2$ is metallic whilst the monolayer of $\text{Ni}_3(\text{HITP})_2$ is an indirect semiconductor.¹¹³

Experimentally, varying the metal center in the charge-neutral $M_3(\text{HITP})_2$ 2D *c*-MOF system could readily change the layer distance, which was determined as 3.30, 3.16, and 3.29 Å for $\text{Ni}_3(\text{HITP})_2$, $\text{Cu}_3(\text{HITP})_2$ and $\text{Co}_3(\text{HITP})_2$, respectively (Fig. 6b).⁵⁸ The layer distance refers to the strength of π - π stacking interactions, thereby enabling to define the electronic structure in the cross-plane direction (Section 3.5). According to the PXRD analysis, the substitution of the metal center is also

relevant to the interlayer displacement, which was recorded as 1.56, 0.86, and 1.39 Å respectively for the above 2D *c*-MOFs. Unsurprisingly, conductivities of these 2D *c*-MOF samples differed greatly, which were determined as 55.4, 0.75, and 0.024 S cm^{-1} , respectively. In this work, Dincă and co-workers further demonstrated a series of binary alloys of $M_nM'_{3-n}(\text{HITP})_2$ consisting of two metal species (Co/Cu, Co/Ni, or Cu/Ni) within one network.⁵⁸ Continuous shifts of the interlayer distance and displacement as well as conductivity were determined with an increasing amount of one metal. For instance, increasing the amount of Ni in the $\text{Co}_n\text{Ni}_{3-n}(\text{HITP})_2$ system shifts the values of layer distance and displacement in such a way as to enhance the π - π interactions, which results in a continuous increment of conductivity over 3 orders of magnitude from 0.024 to 55.4 S cm^{-1} (Fig. 6c) and a continuously tuned bandgap over 0.4 eV in fine-scale (Fig. 6d). While for the fully substituted coronene (PTC), the metal substitution from Fe to Ni/Cu results in not only changes in the conductivity but also in the variation of the chemical structures of 2D *c*-MOFs (Fig. 2c and d).^{32,110,111} In addition, the metal atoms play also an important role in the magnetic properties of 2D *c*-MOFs. In particular, due to the unique spin properties of Fe and the strong π - d hybridization between the d/p orbitals in the FeX_4 complex, the resultant 2D *c*-MOFs exhibit nontrivial ferromagnetism (Section 6.3).^{32,33}



3.4 Effect of redox-activity in linkage

Here, the redox activity majorly refers to the charge state of the linkages. Each linkage is able to undergo a two-electron transfer through redox reactions (Fig. 7), which corresponds to diverse charge state of the linkage as 0 (existence of mono-radical), -1 (presence of counter cation), and 1 (presence of counter anion). Among the reported 2D *c*-MOFs, the ligands are generally partially oxidized providing free radicals as charge carriers (charge-neutral framework with a charge state of 0), which correlates with the density (n) of charge carriers in the frameworks.^{19,36,51,56,65,73–75} For example, $\text{Cu}_3(\text{HHB})_2$ possesses a network with denser distributed ligands and metals as well as smaller layer distance ($\sim 3 \text{ \AA}$)³⁹ compared with $\text{Cu}_3(\text{HHTP})_2$ ($\sim 3.3 \text{ \AA}$);⁹⁵ the former one was expected to display higher conductivity due to the stronger cross-plane π - π interactions (Section 3.5). However, conductivity experiments indicated inferior values in the range of $(0.22\text{--}15) \times 10^{-8} \text{ S cm}^{-1}$ for $\text{Cu}_3(\text{HHB})_2$, making it like an insulator. The huge difference would be attributed to the localized charge carriers due to the absence of free radicals with respect to a charge state of -1 . While for $\text{Ni}_3(\text{HTB})_2$, X-ray photoelectron spectroscopy (XPS) analysis revealed mixed charge states of 0 and -1 (ratio of 26 : 74, corresponding to an average charge state of $-3/4$) in the nickel-bis(dithiolene) linkages for the as-synthesized sample (conductivity in the range of $0.15\text{--}2.8 \text{ S cm}^{-1}$), indicative of the existence of few free radicals. It was demonstrated that additionally generated radicals upon oxidation offer abundant free carriers, thereby leading to enhanced conductivity values.^{74,75} For instance, the amount of radicals in $\text{Ni}_3(\text{HTB})_2$ was further increased through oxidation reactions (charge state of 0), leading to significantly advanced conductivity up to 160 S cm^{-1} . By comparison, the reduction reaction on the as-synthesized $\text{Ni}_3(\text{HTB})_2$ led to a decreased conductivity of $6.7 \times 10^{-3} \text{ S cm}^{-1}$, which can be ascribed to the disappearance of those few radicals after chemical reduction (charge state of -1).

3.5 Effect of layer stacking and arrangement

As is seen in 2D *c*-MOFs, linking the *ortho*-substituted conjugated ligands by square-planar metal coordination complexes yields an extended π -conjugated network, thus enhancing the delocalization of charge carriers within the 2D plane. The in-plane charge carrier migration was considered as the dominant pathway of the intrinsic charge transport in 2D *c*-MOFs. Nevertheless, in recent studies, DFT calculations and charge transport characterization based on single-crystals revealed the significance of the through-space pathway in the electronic properties. As mentioned in Section 3.3, a single-layer of $\text{Ni}_3(\text{HITP})_2$ is predicted to be a semiconductor (bandgap of 0.13 eV) (Fig. 8a and b). The increased dimensionality from a

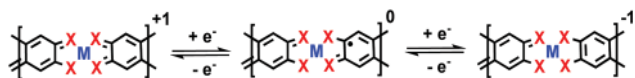


Fig. 7 Redox activity of the linkage of 2D *c*-MOFs through electron transfer.

monolayer to layer-stacked $\text{Ni}_3(\text{HITP})_2$ leads to a significant change from semiconducting to metallic in the electronic band structure (Fig. 8c), which can be attributed to the strong π - π ($d = 3.3 \text{ \AA}$) and weak Ni-Ni ($d = 3.809 \text{ \AA}$) interactions in the direction perpendicular to the $\text{Ni}_3(\text{HITP})_2$ layers.¹¹³ Very recently, the Dincă group characterized the out-of-plane conductivity of $\text{Ni}_3(\text{HITP})_2$ up to 150 S cm^{-1} and demonstrated that interlayer π - π interactions indeed considerably contribute to the conductivity in layer-stacked 2D *c*-MOFs based on single-crystalline samples and related devices.⁹⁵ Thus, it raised up an open question whether high-performance electrically conductive MOFs necessarily require the incorporation of in-plane conjugated structures. Another work from the same group reported layered lanthanide-HHTP MOFs with metal centers located in between the linker layers.¹²¹ The lanthanide-HHTP MOFs do not possess in-plane π - d conjugation but nevertheless display high electrical conductivity (up to 0.05 S cm^{-1}).

Rational tuning the interlayer spacing and modification of layer arrangement can induce significant changes in the electronic band structures as well as conductivities. Great efforts from DFT calculations have been devoted to the calculation of band structures and electrical properties for 2D *c*-MOFs.^{31–33,40,78–80,112,113,122–127} Combining experimental and computational analysis, the reported 2D *c*-MOFs preferred to stack in an AA-eclipsed (AAAA) symmetry (Fig. 8d), rather than the staggered AB (ABAB) mode.^{46,48,50,51,54} Taking carefully in the AA symmetry, the AA-serrated (AA'AA'') stacking (Fig. 8e), so called “slipped-parallel AB stacking” with slightly slipped arrangement in every two layers,^{19,28,32,33,36,39,40,42,52,53,57,59,68,96,97,99} and AA-inclined (AA'A''A'''), that is, the layers are continuously shifted in one direction) stacking (Fig. 8f)⁷⁸ were suggested to be energetically more favored than the AA-eclipsed one in these highly conjugated systems. The stacking—refers to the arrangement of layers—affects the pore geometries¹²⁸ and the electronic properties¹²⁹ of 2D framework materials. Taking a representative 2D covalent organic framework (COF), COF-5,¹³⁰ as an example, the band structure of a single-layer shows flat conduction and valence bands as well as typical signature of the kagome lattice^{68,112,113} along the Γ - K - M - Γ path (in-plane), while stacking the layers with different arrangements varies not only the out-of-plane charge transport, but also the in-plane one (Fig. 8d–g, bottom).¹²⁹ The AA-eclipsed stacked COF-5 retains almost unchanged band structures with preserved signature of the kagome lattice in the in-plane direction (for details see ref. 129), while interlayer interactions induce strong cross-plane band dispersion hence narrowing the band gap from ~ 2.5 (monolayer) to 1.3 eV . However, the AA-serrated and AA-inclined stacking modes strongly alter the in-plane electronic structure, leading to the absence of the kagome characteristic (for details see ref. 121). Compared with the eclipsed layer arrangement, both the serrated and inclined cases exhibit smaller out-of-plane dispersion due to the weaker π - π interactions, which corresponds to a larger band gap of 2.2 eV for both cases. In another instance, $\text{Fe}_3(\text{HITP})_2$ 2D *c*-MOF, the band gap of a monolayer was calculated as 0.3 eV ; it opens up to 0.35 and 0.65 eV for the AA-inclined and AA-eclipsed arrangements, respectively.⁷⁸ Furthermore, the obtained effective



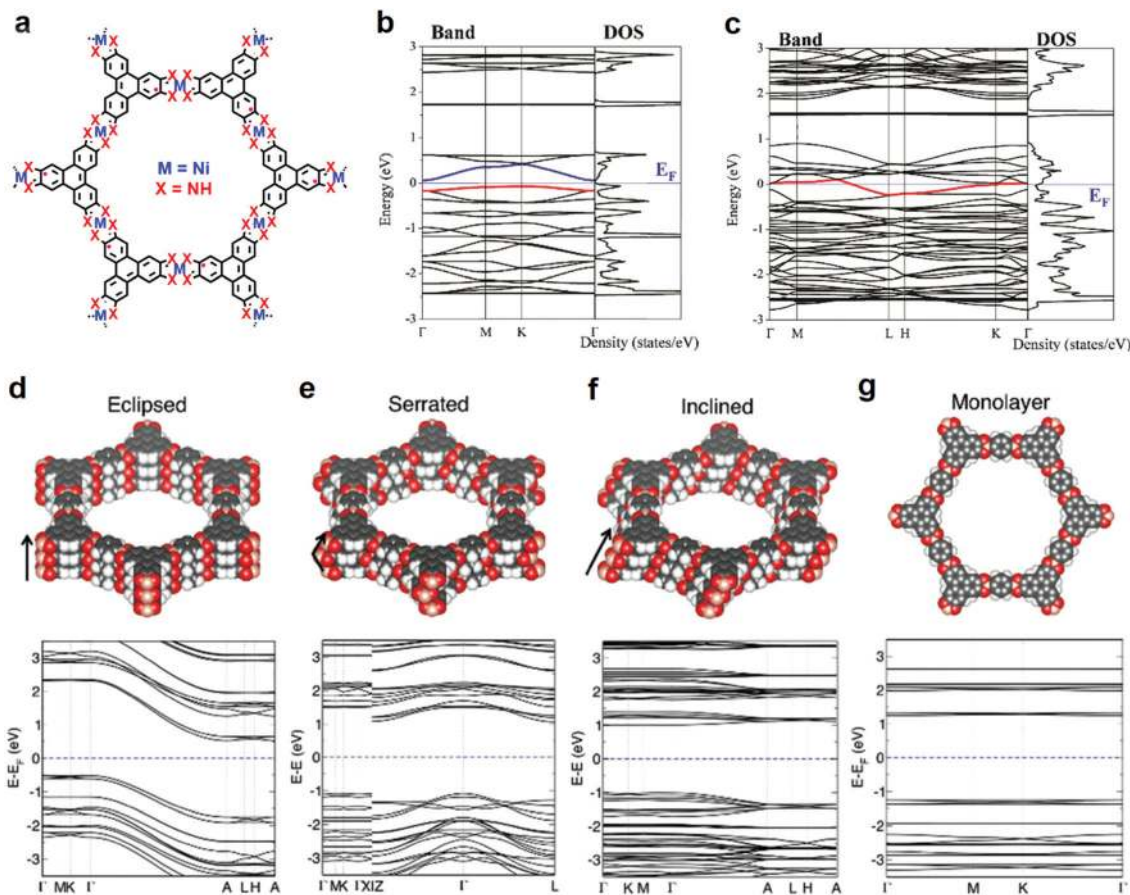


Fig. 8 Electronic band structure with respect to layer stacking and arrangement. (a–c) Band structure for monolayer (b) and layer-stacked (c) $\text{Ni}_3(\text{HITP})_2$.¹¹³ Copyright 2015, the PCCP Owner Societies. (d–f) AA-eclipsed, AA-serrated, and AA-inclined stacking of COF-5 as well as related band structures shown at the bottom. (g) Monolayer and related band structures of COF-5.¹²⁹ Copyright 2020, Wiley-VCH.

mass (m^*) values for AA-inclined stacking ($m_h^* = 0.78$, $m_e^* = 0.98$) and AA-eclipsed stacking ($m_h^* = 1.07$, $m_e^* = 2.08$) differ significantly from those for a monolayer ($m_h^* = 0.45$, $m_e^* = 1.34$) of $\text{Fe}_3(\text{HTTP})_2$. Therefore, to evaluate the band diagrams in the absence of high structural resolution, several potential stacking sequences including not merely AA-eclipsed and AB arrangements merit thorough consideration.

In addition to the foregoing points, high crystallinity or even single-crystal not only offers a high-quality sample but also allows precise structural analysis,^{71,131–133} thus ensuring the establishment of a reliable structure–property relationship. The polycrystalline $\text{Cu}_3(\text{HITP})_2$ possessed a mixed valence of $\text{Cu}^{1/2+}$ in the as-synthesized sample.²⁸ However, all Cu atoms presented to be Cu^{2+} in the single-crystalline sample hence ruling out the contribution of mixed valences to conductivity.⁵⁸ The mixed valence of $\text{Cu}^{1/2+}$ could be ascribed to the presence of defects or amorphous regions in the less crystalline $\text{Cu}_3(\text{HITP})_2$ sample. Furthermore, the conductivity value of the compressed pellet of $\text{Cu}_3(\text{HITP})_2$ single-crystals increased from 0.2 S cm^{-1} for the polycrystalline $\text{Cu}_3(\text{HITP})_2$ to 0.75 S cm^{-1} . In another example, the highly crystalline $\text{Cu}_3(\text{HTB})$ film exhibits a conductivity reaching 2500 S cm^{-1} ,⁶⁸ while the conductivity of the one with low crystallinity (according to the X-ray diffraction and

transmission electron microscopy analysis) was merely determined to be 1580 S cm^{-1} .^{31,134} In addition, the synthesized $\text{M}_3(\text{HIB})_2$ ($\text{M} = \text{Cu}, \text{Ni}, \text{Co}$) films were reported as an electrical insulator in the early work from Lahiri *et al.*,¹³⁵ which is likely attributed to the low crystallinity in these samples. By comparison, their intrinsic metallicity was confirmed in the highly crystalline bulk samples ($\text{M} = \text{Cu}, \text{Ni}$) by ultraviolet-photoelectron spectroscopy. Conductivity measurements on the pellets revealed characteristic values of 8 and 13 S cm^{-1} for $\text{Ni}_3(\text{HIB})_2$ and $\text{Cu}_3(\text{HIB})_2$, respectively.⁴⁰ The polycrystalline $\text{Ni}_3(\text{HITP})_2$ 2D *c*-MOF displayed temperature-dependence suggesting a semiconducting behavior and a conductivity up to $\sim 60 \text{ S cm}^{-1}$,⁵⁷ while single-crystalline $\text{Ni}_3(\text{HITP})_2$ nanorod sample presented metallicity with a higher conductivity of $\sim 150 \text{ S cm}^{-1}$ through the cross-plane direction.⁹⁵ Thus, single-crystalline devices that can minimize the influence of grain boundary and disorder on the charge transport are urgently required for characterizing 2D *c*-MOFs.^{29,57} Besides conductivity, few reports attempted to probe the type of carriers and charge mobility as well as density for 2D *c*-MOFs in the direct current (DC) limits including field-effect transistors and Hall effect measurements, as well as in the alternating current (AC) limits such as contact-free time-resolved terahertz



spectroscopy (TRTS) and flash photolysis-time-resolved microwave conductivity.³³ An in-depth understanding of the mechanism underlying the conductivity and the establishment of a reliable structure–property relationship through combining multi-scaled physical characterization methods merit further exploration.

4. Synthetic methodologies of 2D *c*-MOFs

Currently, various synthetic methodologies have been applied to develop 2D *c*-MOFs as bulk or film samples. Hydro-/solvo-thermal syntheses represent a representative strategy (Fig. 9) that allows diversity in the reaction conditions including the solvent, temperature, volume and pressure, and offers advantages in scalable and high-yield synthesis. They also provide the possibility to achieve moderate-sized (1–10 μm) single-crystals.^{19,95} Nevertheless, the obtained bulk materials generally comprise nanocrystals smaller than 100 nanometers (nm), associating with difficulties in sample processing, device fabrication, and charge transport characterization. Recently, the wet-interface-assisted synthesis^{21,136–142} shows promise in preparing crystalline, large-area, free-standing 2D *c*-MOF thin films. These films ensure facile device integration (notably for electronic devices) and sufficient charge transport^{29,31,36,41,57,68,74,75,78–80,96,97,143} that holds potential to realize the practical applications of these materials for MOF-ronics. Nonetheless, the universality of this interfacial strategy towards large π-conjugated systems, the further improvement of the crystallinity and the mechanical strength of the film as well as the control of the film thickness are still synthetic challenges. In the following sections, we will introduce these

synthetic methodologies toward 2D *c*-MOFs and discuss the current achievements and challenges.

4.1 Bulk 2D *c*-MOFs through hydro-/solvo-thermal syntheses

In 2012, Hmadeh *et al.* reported a series of TP-based MOF-linked 2D *c*-MOFs by heating an aqueous solution of HHTP and metal(II) acetate (M = Cu, Ni, Co) at 85 °C for 24 hours (Fig. 9a and 10a).¹⁹ Scanning electron microscopy (SEM) revealed uniform morphology of the obtained samples as nanorods with length less than 10 μm and diameter of ~100 nm (Fig. 10e). Their crystal structures were resolved by single-crystal X-ray diffraction, PXRD, and high-resolution transmission electron microscopy (HR-TEM). These 2D *c*-MOFs possess two different structures: (1) Cu₃(HHTP)₂ (Fig. 10b) with a layered honeycomb network and (2) M₉(HHTP)₄ (discounting water molecules, M = Co, Ni) (Fig. 10c) comprising alternatively layered honeycomb frameworks of M₃(HHTP)₂(H₂O)₆ (each M²⁺ is coordinated to the linkers in two adjacent deprotonated HHTP molecules and two additional aqua ligands in an octahedral environment) and trinuclear M₃(HHTP)(H₂O)₁₂ clusters (one M²⁺ is coordinated to the linkers in one HHTP molecules and four additional aqua ligands) (Fig. 10d).⁹⁹ Notably, the Cu²⁺ ions in Cu₃(HHTP)₂ are in a square-planar environment without the presence of water ligands. The Brunauer–Emmett–Teller (BET) measurements

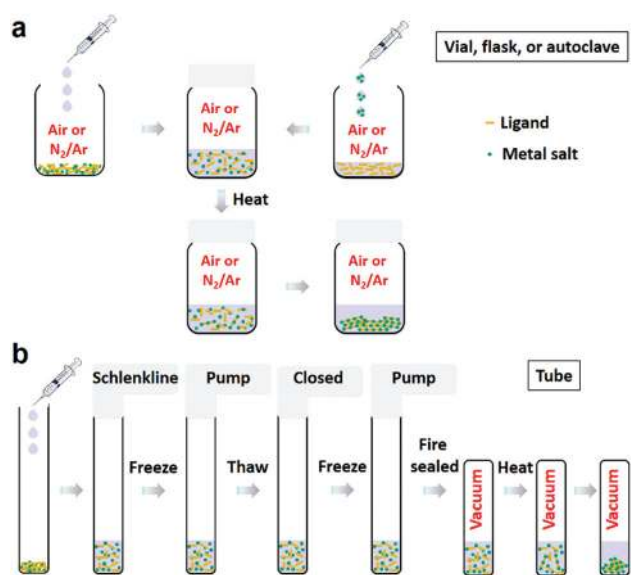


Fig. 9 Schematic illustration of various hydro-/solvo-thermal syntheses methods. (a) Under ambient or inert atmosphere. (b) Under vacuum.

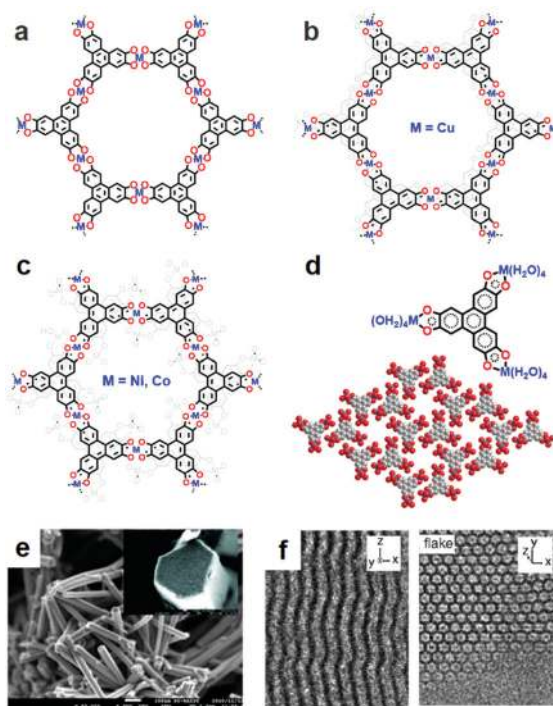


Fig. 10 TP-based 2D *c*-MOFs through hydro-/solvo-thermal syntheses. (a) Generally displayed structure of Cu₃(HHTP)₂ and M₉(HHTP)₄ (M = Ni, Co). (b) Radical structure of bilayer Cu₃(HHTP)₂. (c) Radical structure of layered M₉(HHTP)₄ (M = Ni, Co) without counting the water molecules. (d) Structure of trinuclear M₃(HHTP)(H₂O)₁₂ clusters. (e) SEM image of the nanorod sample. Reproduced with permission ref. 19. Copyright 2012, American Chemical Society. (f) HR-TEM images of Cu₃(HHTP)₂ nanorods (left) and nanoflakes (right). Reproduced with permission ref. 95. Copyright 2019, American Chemical Society.



displayed the specific surface area (SSA) values of these porous materials in the range of 400 to 500 m² g⁻¹. Electron paramagnetic resonance (EPR) suggested a ligand-centered monoradical (Fig. 7, charge state of 0) and confirmed the charge-neutral nature (Fig. 10b and c). In a later report from Day *et al.*, Cu₃(HHTP)₂ layers stacked perpendicular to the axis along each nanorod in a slightly offset fashion with a continuous shift in the *c* direction (Fig. 10f, left).⁹⁵ Combining liquid-phase exfoliation,^{144,145} the authors further synthesized single-crystalline nanoflake-like Cu₃(HHTP)₂ (Fig. 10f, right).

In addition to Cu₃(HHTP)₂, Ni₄-linked Ni₃(HITP)₂ is another representative 2D *c*-MOF sample. In 2014, Sheberla *et al.* reported the hydrothermal synthesis of Ni₃(HITP)₂ based on amine-functionalized TP.⁵⁷ Upon heating an aqueous solution of HATP and NiCl₂ with the existence of NH₃-H₂O in air, Ni₃(HITP)₂ was obtained as bulk nanoparticles with an average crystal size less than 200 nm. Monoanionic *o*-diiminobenzosemiquinonate moieties were confirmed by XPS and elemental analysis, indicative of a charge-neutral network. The further structural analysis revealed the crystal structure of Ni₃(HITP)₂ with unit cell parameters of *a* = *b* = 21.75 Å. In the later studies from the same group, employing NaOAc for the deprotonation, the obtained Ni₃(HITP)₂ showed superior crystallinity in a rod-like morphology with length of up to 2 μm.^{58,95} The HR-TEM image of the individual crystal suggested that the Ni₃(HITP)₂ layers stacked in an almost eclipsed configuration with a periodic interlayer spacing of 3.30 Å. This optimization was also applied to the structurally related charge-neutral Cu₃(HITP)₂ and Co₃(HITP)₂ 2D *c*-MOFs.^{28,58} Meanwhile, HHTP-based MS₄-linked (M = Ni, Co, Fe, Pt) 2D *c*-MOFs were also developed (mainly as films, see Section 4.2),^{45,64,118} and subsequently great efforts have been devoted to the TP-based 2D *c*-MOF family with the aim of exploring their multifunctional properties.^{28,46,59,90,100,106,109,146,147}

Towards frameworks with denser distributed ligands and metals, benzene-based ligands were developed.^{31,39,40,42,74,135} Typically, in 2017, Dou *et al.* synthesized MN₄-linked M₃(HIB)₂ (M = Cu, Ni) 2D *c*-MOFs by mixing HAB, M²⁺, and NH₃-H₂O in DMSO, followed by heating at 60 °C for 2 hours.⁴⁰ The obtained samples were formed by irregularly shaped nanoparticles smaller than 100 nm. The backbones were suggested to be charge-neutral, indicative of the existence of same monoanionic *o*-diiminobenzosemiquinonate moieties as Ni₃(HITP)₂. Unsurprisingly, N₂ adsorption revealed low BET SSA values in the range of 110–150 m² g⁻¹ due to the increment of the density of ligands and metals in the backbone. Using ethylenediamine as the base under an inert atmosphere, Park *et al.* reported CuO₄-linked Cu₃(HHB)₂ 2D *c*-MOF nanocrystals (estimated to be <200 nm).³⁹ The authors showed synthetic diversity through the coordination reaction between Cu²⁺ and either HHB or its derivative tetrahydroquinone (THQ). Elemental analysis suggested the presence of protonated ethylenediamine (NH₃CH₂CH₂NH₃²⁺) as the counter cation to balance the negative charge of the backbones, which corresponds to a charge state of -1 in each linkage. Analogous to M₃(HIB)₂, Cu₃(HHB)₂ exhibited a low BET SSA value of ~143 m² g⁻¹.

A larger π-conjugated coronene-based building block was introduced for the construction of 2D *c*-MOFs by our group in 2016.³² The fully thiol-substituted ligand PTC was incorporated into the 2D *c*-MOF through the coordination with Fe(OAc)₂ in an ammoniacal mixture containing deoxygenated water and dimethylformamide (DMF) at 120 °C for 48 hours. Thus, FeS₄-linked Fe₃(PTC)₂ was achieved as an almost non-porous network with crystalline domains up to a dozen of nm. XPS and Mössbauer spectroscopy suggested the presence of Fe³⁺ species. In addition, we inferred a unit cell with *a* = *b* = ~11.7 Å and an interlayer spacing of ~3.9 Å for this 2D *c*-MOF depending on thorough characterization with the support of PXRD, TEM, XPS, and X-ray absorption spectroscopy. Very recently, Yao *et al.* established a dual-ligand Cu₃(HHTP)(HHB) 2D *c*-MOF with two kinds of ligands of HHTP and THQ in the network.⁵⁹ Definitely, it was a challenge for the controlled synthesis of this dual-ligand 2D *c*-MOF by varying various reaction conditions including the temperature, reaction time, ratio of two ligands, and metal salts, in terms of stronger coordination capability of Cu²⁺ with HHTP than THQ. As a result, the obtained Cu₃(HHTP)(HHB) 2D *c*-MOF presented a morphology of hundreds of nm long nanowires and a unit cell with parameters of *a* = *b* = 17.154 Å and *c* = 6.266 Å for a honeycomb structure. In addition, the charge state of the linkage was -1 with NH₃CH₂CH₂NH₃²⁺ as counter ions. The dual-ligand structure shows the great potential of tunability of the porosity and electronic properties.

Utilizing hydro-/solvothoermal methodologies, many other 2D *c*-MOFs based on various building blocks have been constructed, such as Cu₃(HHTN)₂, Cu₃(HHTX)₂, Ni₃(HHATNM'₃)₂, Cu₂[DBCOS₈], M₂[M'PcX₈], and Ni₂[NiNPcO₈].^{33,47–51,53–56,148} In 2019, we employed a modified solvothoermal protocol applying vacuum for the synthesis of an oxygen-sensitive system (Fig. 9b): Fe₂[FePcO₈] 2D *c*-MOF.³³ This method has also been widely employed for vacuum-promoted synthesis of 2D COFs^{148–150} and 2D non-conjugated MOFs.¹⁵¹ To avoid air-induced oxidation of iron(II) salts, Fe₂[FePcO₈] was synthesized by mixing FePc(OH)₈, Fe(OAc)₂, and KOAc (base) in *N*-methyl-2-pyrrolidone (NMP)/H₂O (v/v = 3/1) in a vacuum-sealed tube after thorough deoxygenation by three cycles of freeze–pump–thaw, followed by heating at 150 °C for 3 days. Thus, the obtained Fe₂[FePcO₈] is comprised of rectangle-shaped crystals in the range of 10–100 nm with a square lattice of *a* = *b* = ~18 Å.

Though hydro-/solvothoermal syntheses have been demonstrated as the most effective and simplest strategy for expanding the 2D *c*-MOF family, this method generally allows barely monitoring or controlling the crystal growth and provides small-sized bulk polycrystals. As a result, integrating the MOF particles into nanodevices and the relevant performance of the thus-far fabricated devices have been largely impeded by the lack of process-capability and accessible active sites as well as sufficient charge carrier transport.

4.2 2D *c*-MOF films through wet-interface-assisted synthesis

Recently, various bottom-up wet interface-assisted synthetic strategies including gas–liquid (GL), liquid–liquid (LL), and



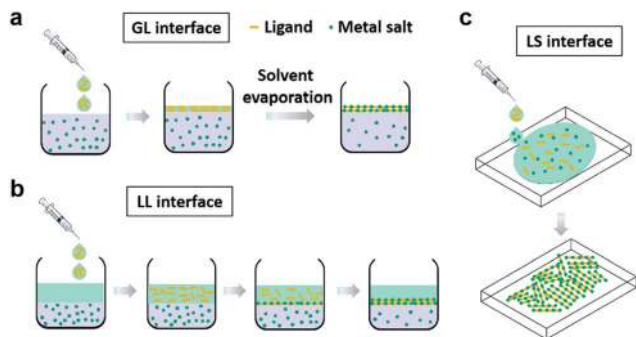


Fig. 11 The interfacial strategies for the synthesis of 2D *c*-MOF films. (a and b) Schematic illustration of gas–liquid and liquid–liquid interfacial methods. (c) Schematic illustration of liquid–solid interfacial synthesis.

liquid–solid (LS) interfacial approaches have been developed to prepare large-area thin 2D *c*-MOF films (Fig. 11).^{21,152–154}

One representative work on the synthesis of 2D *c*-MOF films was reported initially by Kambe *et al.* in 2013 using a LL interfacial strategy (Fig. 12a).^{74,75} An organic phase containing HTB in dichloromethane (DCM) was overlaid with an aqueous solution of Ni(OAc)₂ and NaBr under an argon atmosphere. The water/DCM interface allowed the formation of a 1–2 μm thick Ni₃(HTB)₂ 2D *c*-MOF film with a lateral size of ~100 μm (Fig. 12b). PXRD and TEM analysis (Fig. 12c) indicated a crystal model of Ni₃(HTB)₂ with parameters of *a* = *b* = 14.1 Å and *c* = 7.6 Å. The same group also reported an isostructural Pd₃(HTB)₂ using the above LL interfacial method. Additionally, K₃[Fe(CN)₆] was added as a redox buffer to prevent the reduction

of Pd⁺ to the formation of Pd⁰.⁶⁵ In 2015, Huang *et al.* reported a non-porous 2D *c*-MOF network Cu₃(HTB) (Fig. 12d–f) by coordination of HTB and Cu(NO₃)₂ at the water/DCM interface.³¹ The Cu₃(HTB) 2D *c*-MOF was obtained as a continuous film with a lateral size larger than 1 cm and thickness in the range of 20 to 140 nm. In the Cu₃(HTB) network, each sulfur atom is coordinated to two copper atoms leading to a non-porous structure with denser ligands and metals. SEM and atomic force microscopy (AFM) images showed a continuously smooth upside in the film whilst the downside composed of plate-like nanosheets was fairly rough, especially for the thick films. The authors suggested that the water/DCM interface functioned initially as a platform for the formation of 2D *c*-MOF thin films (the upside in the final film), while the HTB ligand functioned as the transporter of the Cu-ions from the interface to the DCM phase below the interface. Then, the Cu-ions coordinated with the HTB molecules below the interface in a less order due to the weaker interface-confining effect here than that at the interface. Grazing incident X-ray diffraction (GIXRD) revealed a face-on orientation of Cu₃(HTB) in a 60 nm thick film whilst no preferred orientation was observed for the sample thicker than 200 nm, indicative of randomly accumulated nanocrystals on the downside of the film near the water/DCM interface. Lahiri *et al.* also employed GL/LL interfacial methods for the construction of M₃(HIB)₂ (M = Co, Ni, Cu) thin and thick films with thicknesses of 10–13 nm and ~1 μm, respectively.¹³⁵

In 2016, Wu *et al.* reported a modified solution synthesis of large-area Ni₃(HITP)₂ films in a homogeneous aqueous solution of HATP and NiCl₂ using trimethylamine as the base under no perturbation.²⁹ Upon heating the mixture to 60 °C, Ni₃(HITP)₂ nanoparticles self-assembled at the air/water interface and packed further closely to afford a continuous smooth film. The film thickness could reach ~100 nm after 3 min of polymerization. In 2018, our group demonstrated a LL interfacial synthesis of the Fe₃(HTTP)₂ 2D *c*-MOF film (Fig. 12g).⁷⁸ Organic salt of iron(II) acetylacetonate was dissolved in chloroform, which was overlaid by an aqueous phase containing the ligand HTTP and ammonia. Large-area (cm²), free-standing Fe₃(HTTP)₂ films were obtained with a tunable thickness from 20 nm to ~2 μm by increasing the reaction time (Fig. 12h and i). TEM imaging revealed that the samples comprised of crystalline domains of ten to hundreds of nm. A honeycomb structure with a pore size of ~1.9 nm was observed by high-resolution TEM. PXRD displayed (100) and (200) peaks at 2θ = 4.5° and 9.1°, respectively, which demonstrated a honeycomb packing within the *ab* planes. The (001) reflection (peak at 2θ = 27.3°) suggested an ordered stacking along the *c* direction with an interlayer distance of ~0.33 nm. Experimental PXRD patterns agreed well with the simulated inclined AA stacking models. The BET SSA was measured to be ~526 m² g⁻¹. In 2019, the above 2D *c*-MOF film with the thickness ranged from ~84 to ~410 nm was also reported by Clough *et al.* using an ethyl acetate (containing 1% NMP)/water interface.⁸⁰

Apart from GL and LL approaches, LS interface-assisted synthetic methodology (also termed as on-substrate synthesis, Fig. 11c) has recently emerged for preparing 2D *c*-MOF films

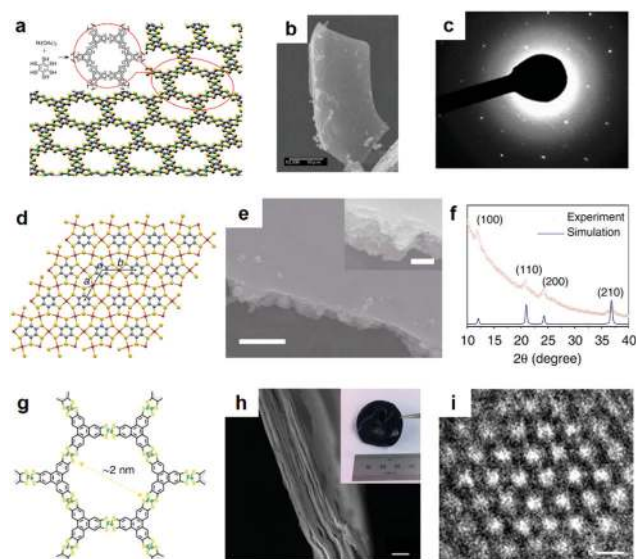


Fig. 12 Selected 2D *c*-MOFs synthesized through the LL interfacial method. (a–c) Ni₃(HTB)₂ 2D *c*-MOF and its SEM and HR-TEM images. Reproduced with permission ref. 74. Copyright 2013, American Chemical Society. (d–f) Lattice structure of Cu₃(HTB)₂ 2D *c*-MOF and its SEM image as well as in-plane GIXRD pattern.³¹ Copyright 2015, Nature Publishing Group. (g–i) Fe₃(HTTP)₂ 2D *c*-MOF and its SEM image from cross-side view and HR-TEM image.⁷⁸ Copyright 2018, Nature Publishing Group.



with a preferred face-on orientation, for example, for HHTP-based systems.^{94,96,155,156} The preliminary on-substrate synthesis was explored by Sheberla *et al.* by placing a quartz substrate in the reaction mixture for the synthesis of bulk $\text{Ni}_3(\text{HHTP})_2$ 2D *c*-MOFs.⁵⁷ $\text{Ni}_3(\text{HHTP})_2$ was obtained as a ~ 500 nm thick crystallite-accumulated macroscopic film on quartz surface. In 2017, Yao *et al.* reported a spray layer-by-layer liquid-phase epitaxial method for preparing face-on orientated $\text{Cu}_3(\text{HHTP})_2$ thin films.⁹⁶ Typically, $-\text{OH}$ functionalized substrates were alternatively exposed to a solution of HHTP and copper(II) acetate in ethanol by a spray method (Fig. 13a and b). The thickness could be controlled from 20 to 100 nm with an average increment of ~ 2 nm after each growing cycle. Top- and side-view SEM images reveal a dense and continuous thin film with a moderately smooth surface on the substrate (Fig. 13c). Later on, Mahringer *et al.* explored a vapor-assisted conversion¹⁰⁶ as an effective way for the synthesis of face-on oriented $\text{Cu}_3(\text{HHTP})_2$ and $\text{M}_9(\text{HHTP})_4$ ($\text{M} = \text{Ni}, \text{Co}$) thin films on various substrates.⁹⁴ The authors utilized an additive of 1-propanol as a modulator for enlarging the crystal size (up to 10 μm for $\text{Cu}_3(\text{HHTP})_2$, 1–2 μm for $\text{Ni}_9(\text{HHTP})_4$, and 3–4 μm for $\text{Co}_9(\text{HHTP})_4$). In a typical synthesis, a water/1-propanol ($v/v = 1/1$) solution of precursor was spread onto an elevated gold substrate in a reaction vessel containing the vapor source of the same water/1-isopropanol mixture, and heated at 85 $^\circ\text{C}$ for 3 h (Fig. 13d). GIXRD patterns in the in-plane and out-of-plane directions indicated a preferential orientation of $\text{Cu}_3(\text{HHTP})_2$ or $\text{M}_9(\text{HHTP})_4$ layers with stacking reflections aligned orthogonally to the surface of the substrates. These samples are free of crack over a large area with thicknesses of ~ 200 nm (Fig. 13e). Cross-section SEM images revealed packed pillared crystallites, corresponding to the observation of GIXRD analysis. The LS interfacial synthesis of high quality thin films allows direct device fabrication by adopting conductive or insulating substrates for electronic devices.^{94,96,97}

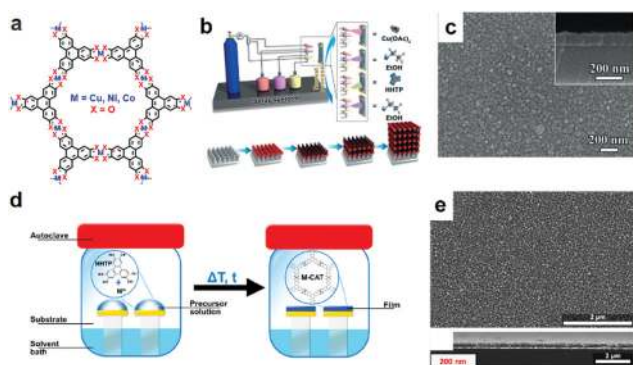


Fig. 13 Oriented HHTP-based 2D *c*-MOF films synthesized through various LS interfacial methods. (a) HHTP-based 2D *c*-MOFs. (b and c) Schematic illustration of the spray layer-by-layer liquid-phase epitaxial method and the SEM image of the obtained $\text{Cu}_3(\text{HHTP})_2$ film. Reproduced with permission ref. 96. Copyright 2017, Wiley-VCH. (d and e) Schematic illustration of the vapor-assisted conversion method and the top-/side-view SEM images. Reproduced with permission ref. 94. Copyright 2019, American Chemical Society.

Apart from 2D *c*-MOF thin/thick films and thin nanosheets, the construction of a single-layer sample with high structural control at the atomic level has remained a synthetic challenge. The single-layer sample will offer distinguished properties due to the 2D material nature,^{157–159} which include, but are not limited to size effects on the intrinsic properties, varied electronic and optical properties due to the absence of interlayer interactions, high exposure of surface active sites, high mechanical flexibility, facile device design and processing, and further construction of multicomponent van der Waals heterostructures.^{45,61,98,160} In 2015, our group reported the preparation of the first example of large-area, free-standing, single-layer 2D *c*-MOFs through Langmuir–Blodgett (LB) assisted interfacial synthesis (Fig. 14),¹⁶¹ by controlling the surface pressure (Fig. 15a and b).⁴⁵

Typically, to produce a single-layer $\text{Ni}_3(\text{HHTP})_2$, a mixed solution of ligand HHTP and DMF/chloroform ($v/v = 1/3$) was spread over the water surface in a LB trough. Subsequently, the HHTP molecules were compressed into a dense self-assembled monolayer through monitoring and increasing the surface pressure to 10 mN m^{-1} , and an aqueous solution of $\text{Ni}(\text{NO}_3)_2$ was injected into the water phase. With the diffusion of the Ni^{2+} ions to the air/water interface, a coordination reaction was triggered between HHTP and Ni^{2+} ions. A free-standing single-layer $\text{Ni}_3(\text{HHTP})_2$ film was obtained and could be directly transferred onto various substrates for further characterization. The thickness of the film was measured to be ~ 0.7 nm by AFM (Fig. 15c and d). As the single-layer sheets showed low stability under electron irradiation, selected area electron diffraction (SAED) was performed by cryogenic TEM at -175 $^\circ\text{C}$, exhibiting a typical hexagonal diffraction pattern. This SAED result at least suggested local crystallinity presenting in the single-layer 2D *c*-MOF with an ordered honeycomb network having a cell size of ~ 2 nm. Such a $\text{Ni}_3(\text{HHTP})_2$ 2D *c*-MOF was transferred onto glassy carbon electrodes for electrocatalytic water splitting. Currently, due to their well-defined structure and abundant active sites as well as intrinsic conductivity, 2D *c*-MOFs have been rising as a promising electrode material for electrochemical energy devices.

These efforts from various interfacial approaches allow the integration of 2D *c*-MOFs into functional devices and thus exploring their multifunctional properties. However, the

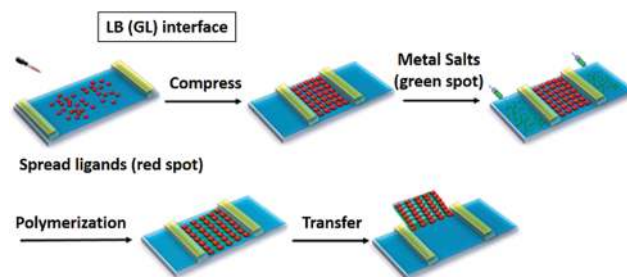


Fig. 14 Schematic illustration of Langmuir–Blodgett air/water (a special GL interface) interface-assisted synthesis. Reproduced with permission ref. 21. Copyright 2018, American Chemical Society.



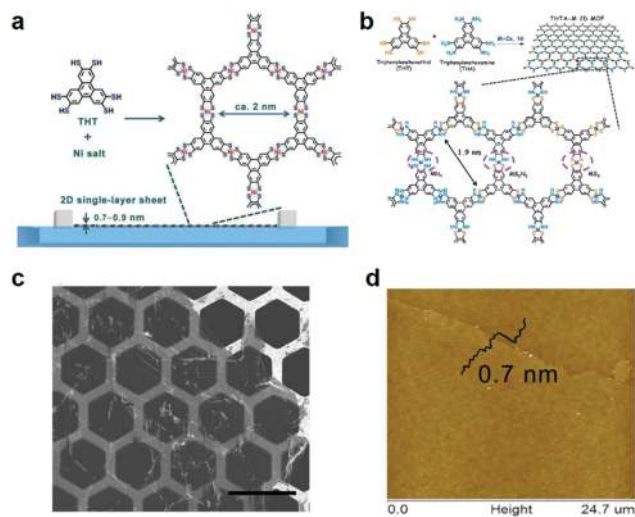


Fig. 15 Single-layer 2D *c*-MOFs synthesized through LB-assisted interfacial synthesis. (a and b) Schematic illustration of the synthesis of $\text{Ni}_3(\text{HTTP})_2$ and a $\text{Co}_3(\text{HITP})(\text{HTTP})$ 2D *c*-MOF comprising mixed ligands of HTTP and HATP. (c and d) SEM and AFM images of single-layer $\text{Ni}_3(\text{HTTP})_2$. Reproduced with permissions.^{45,61} Copyrights 2015 and 2017, Wiley-VCH.

interfacial strategies have been so far rather limited; the successful synthesis strongly relied on the association of high solubility of selective building blocks in water or organic solvents. In addition, accessibility on controlling/tuning orientation remains to be explored. In addition to wet-surface, the on-surface synthesis using Cu(111) or Au(111) substrates under ultrahigh vacuum provides another pathway to develop a single-layer 2D *c*-MOF that guarantees a molecular structural resolution with the support of a scanning tunneling microscope (STM).^{162–165} In 2018, Lischka *et al.* reported a topologically diverse framework upon annealing HATP on Cu(111). A metal-organic trimer composed of three ligand molecules linked by a Cu_3 cluster instead of a single Cu atom was observed, while ring-fused $\text{Ni}_3(\text{HITP})_2$ with a limited number of repeating units appeared in the presence of additional Ni on Cu(111).¹⁶³ Later on, by varying Cu(111) surface to Au(111), Gao *et al.* obtained a monolayer of $\text{Ni}_3(\text{HITP})_2$ with a domain size of ~ 40 nm and resolved its structure at sub-molecular level by STM.¹⁶² Recently, Zhang *et al.* applied the on-surface method for bonding HHB with Cu atoms; however, a single-layer 2D non-conjugated MOF was determined.¹⁶⁵ Despite the preliminary success of the on-surface synthesis, many synthetic challenges remain to be addressed, such as the synthesis of large-area (from μm^2 to cm^2), highly-ordered monolayer networks, the complete lift of the monolayer from the solid surface for device integration, and so on.

5. Device integration based on multi-dispersed 2D *c*-MOFs

As is shown, 2D *c*-MOFs have been synthesized by various synthetic methodologies and have shown diverse morphologies,

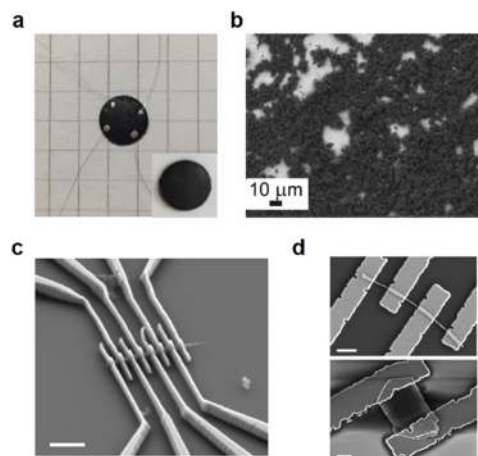


Fig. 16 Device integration of bulk 2D *c*-MOFs. (a) Pellet of 2D *c*-MOFs and device fabrication in the van der Pauw geometry.⁵⁵ Copyright 2020, Wiley-VCH. (b) SEM image of drop-cast $\text{Cu}_3(\text{HTTP})_2$ on an indium-tin-oxide glass slide. Reproduced with permission ref. 28. Copyright 2015, Wiley-VCH. (c) SEM image of the fabricated device based on isolated single-crystalline $\text{Ni}_3(\text{HTTP})_2$ nanorods. (d) SEM images of fabricated devices based on isolated single-crystalline $\text{Cu}_3(\text{HTTP})_2$ nanorods (top) and nanoflakes (down). Reproduced with permission ref. 95. Copyright 2019, American Chemical Society.

such as bulk powder (polycrystalline nanoparticles or single-crystals) and large-area thin films. Regarding the bulk powders, compressed pellets (Fig. 16a) have been widely employed for device integration (other method *e.g.* drop-casting, Fig. 16b) and charge transport characterization. Nonetheless, these pellet samples are generally comprised of nanocrystals with no preferred orientation, co-existing with numerous grain boundaries and defects,¹⁶⁶ which limited the conductivity performance and the unambiguous understanding on the transport mechanism. By comparison, the single-crystalline 2D *c*-MOF samples offer the opportunity for device investigation based on each crystal (nanorods or nanoflakes) and allow precise and reliable analysis of the intrinsic electronic properties (Fig. 16c and d). However, the isolated single-crystalline nanorod or nanoflake sample suffers from synthetic challenges and limitation in practical functional studies.

In regard to addressing the above challenges with respect to application of the bulk crystals (polycrystals and single-crystals) obtained by the hydro-/solvothermal syntheses, top-down exfoliation through methods such as ball-milling^{167,168} and sonication^{144,145,169} (Fig. 17a) is expected to be an efficient approach to achieve thin nanosheets,^{169–172} which can maintain the intrinsic conductivity and porosity, allow high exposure of active sites, as well as facile solution-processible device integration.¹⁷³ Currently, few top-down methods have been applied to the delamination of bulk 2D *c*-MOFs.

Very recently, we employed ball milling exfoliation for the synthesis of $\text{Ni}_2[\text{CuPc}(\text{NH})_8]$ 2D *c*-MOF nanosheets.⁵⁵ Firstly, a highly crystalline bulk $\text{Ni}_2[\text{CuPc}(\text{NH})_8]$ sample was prepared with an average domain size of ~ 200 nm *via* a solvothermal protocol. Then, *via* a NaCl-assisted low energy ball milling strategy (Fig. 18a), bulk $\text{Ni}_2[\text{CuPc}(\text{NH})_8]$ nanocrystals were



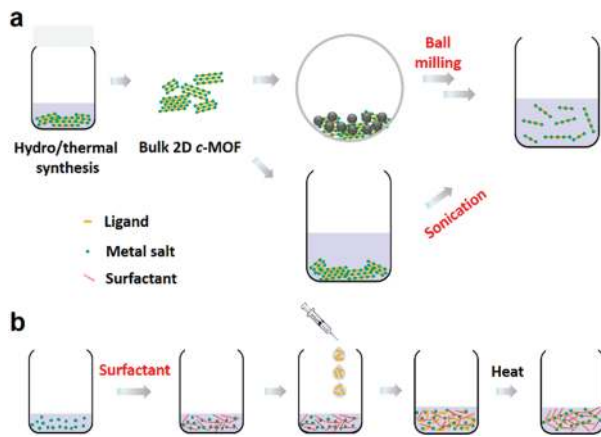


Fig. 17 Schematic illustration of the synthesis of 2D *c*-MOF nanosheets through top-down and bottom-up method. (a) Top-down ball milling and sonication exfoliation. (b) Bottom-up surfactant-assisted synthesis.

mechanically exfoliated into nanosheets with a yield of 40–50%. PXRD patterns revealed a shift of the layer distance from 3.24 to 3.33 Å, indicative of swollen layers after milling. SEM, AFM and HR-TEM together indicated an average lateral size of ~160 nm and a thickness of ~7 nm for these nanosheets (Fig. 18b). The achieved nanosheets could be homogeneously dispersed in DMF for more than six months that allows facile solution-processible micro-supercapacitor device fabrication through filtration (further discussion is in Section 6.5.2).

Even though the top-down exfoliation method offers a possibility in scalable synthesis of nanosheets with accessible active sites and solution-processibility, it remains limited in the control of the morphology, the layer numbers and the lateral domain size. By contrast, bottom-up methods that refer to the direct synthesis of the nanosheets from metal nodes and ligands hold promise in the synthesis of high-quality 2D *c*-MOF nanosheets. Recently, our group introduced a surfactant-mediated synthetic method toward single-crystalline $\text{Cu}_3(\text{HHB})_2$ nanosheets (Fig. 17b).⁸⁸ The sodium dodecyl sulfate

(SDS) anionic surfactant was used as the directing agent for restricting the growth of 2D *c*-MOFs along the *c* direction and weakening the interlayer interactions.^{174,175} In a typical surfactant-mediated hydrothermal synthesis, the precursors of THQ, $\text{Cu}(\text{OAc})_2$, and NaOH (base) in a SDS aqueous solution were ultrasonicated at 50 °C to prepare the nanosheets with an ultrahigh yield of 80–90% (Fig. 18c). As a consequence, the achieved nanosheets presented a thickness of 4.2 ± 1.1 nm and a lateral size of $0.30\text{--}0.65 \mu\text{m}^2$, as well as a much higher BET SSA ($385 \text{ m}^2 \text{ g}^{-1}$) than the contrast powder sample ($119 \text{ m}^2 \text{ g}^{-1}$) obtained in the absence of SDS. HR-TEM and related SAED analysis demonstrated the achievement of single-crystalline nanosheets without noticeable defects (Fig. 18d). The high exposure of active sites in the nanosheet sample allows fast ion/electron diffusion hence ensuring superior performance than the related bulk $\text{Cu}_3(\text{HHB})_2$ as an electrode material (Section 6.5.1).

Synthesis of solution-processible nanosheets will be facile for the fabrication of 2D *c*-MOF based devices *via* spin-coating, filtration, ink-printing, *etc.* In this respect, it requires to address the key questions on the homogeneous dispersion in solution and the control of size, thickness and crystallinity as well as avoiding the re-stacking during solidification. Definitely, another choice is to utilize the free-standing, large-area, highly crystalline 2D *c*-MOF thin films with tunable thickness, which will simplify the device fabrication by direct deposition onto the substrates. To this end, an extraordinary property and a reliable structure–property relationship are expected, thus pushing the functional development for MOFtronics.

6. Functions for MOFtronics

Conductive 2D *c*-MOFs possess a series of unique features, such as large surface to volume ratio, defined active sites, and tailored electronic structure that enables a set of intriguing properties from semiconductor and redox activity to metallicity, ferromagnetism, photoswitching, *etc.* High conductivities in 2D *c*-MOFs have attracted great attention in various MOFtronics including (opto)electronics, spintronics, and thermoelectrics, as well as energy storage devices as potential power sources for MOFtronics.

6.1 Electronics

The emerging 2D *c*-MOFs featuring semiconductor properties, high conductivity, and electrically-transduced sensing have been expected to be outstanding substitutes for potential applications in electronics, such as field-effect transistors, superconductors, chemiresistors, *etc.*

6.1.1 Field-effect transistors (FETs). FETs are a class of three-terminal semiconductor electronic devices with high integration and mass production²⁹ that have played key roles in the current logic and memory devices and sensors. In addition, FETs represent an efficient tool for the study of charge transport properties including the type of charge carries and their mobility values. In 2015, Huang *et al.* fabricated a FET based on the $\text{Cu}_3(\text{HTB})$ film in a bottom-gate bottom-contact

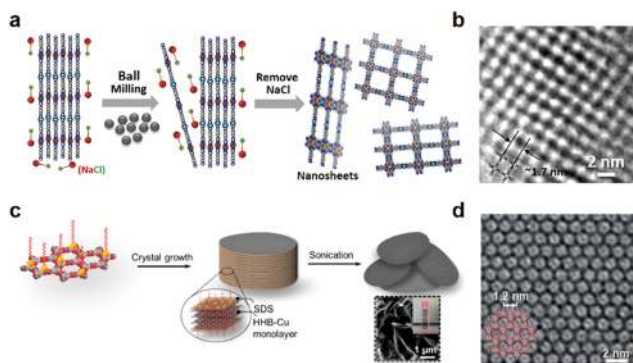


Fig. 18 Synthesis of 2D *c*-MOF nanosheets. (a and b) Schematic illustration of the NaCl-assisted ball-milling exfoliation of $\text{Ni}_2[\text{CuPc}(\text{NH})_8]$ and HR-TEM image of the synthesized nanosheets.⁵⁵ Copyright 2020, Wiley-VCH. (c and d) Schematic illustration of surfactant-assisted synthesis of $\text{Cu}_3(\text{HHB})_2$ 2D *c*-MOF nanosheets and the related HR-TEM image.⁸⁸ Copyright 2020, the Royal Society of Chemistry.



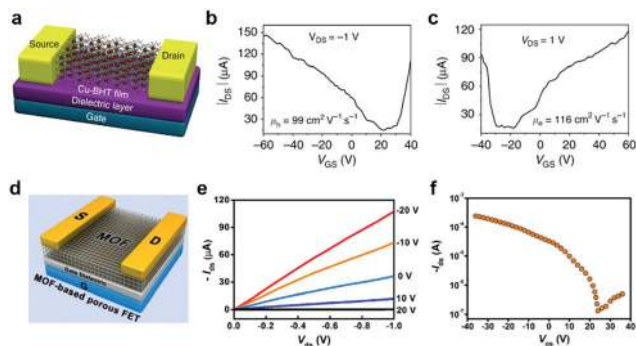


Fig. 19 2D *c*-MOF based FET devices. (a–c) Illustrative schematic of FET based on Cu₃(HTB) and related transfer characteristics.³¹ Copyright 2015, Nature Publishing Group. (d) Ni₃(HITP)₂-based FET. (e and f) Output curves and transfer characteristic plot of Ni₃(HITP)₂, respectively. Reproduced with permission ref. 29. Copyright 2017, American Chemical Society.

geometry (Fig. 19a).³¹ The drain-source current (I_{ds}) versus gate-source voltage (V_{gs}) curves indicated ambipolar behavior with outstanding hole and electron mobility of 99 and 116 cm² V⁻¹ s⁻¹ respectively as well as a current on/off ratio of 10 (Fig. 19b and c). The low on/off ratio correlates with the metallic nature of Cu₃(HTB). Later on, Wu *et al.* applied the Ni₃(HITP)₂ 2D *c*-MOF film in a bottom-gate top-contact geometry (Fig. 19d).²⁹ The curve of I_{ds} as a function of V_{gs} showed an increased channel current with a declined V_{gs} , suggestive of a p-type semiconducting behavior for the as-synthesized Ni₃(HITP)₂ film sample with a mobility up to 48.6 cm² V⁻¹ s⁻¹ (holes as the majority of carriers) and a high current on/off ratio of ~2000 (Fig. 19e and f). Recently, Wang *et al.* integrated the Ni₃(HITP)₂ film synthesized by the LS interfacial method into FETs with bottom-contact device geometry.¹⁰⁷ The FET showed similar performance to the previous report, with a mobility of 45.4 cm² V⁻¹ s⁻¹. Using the FET as a liquid-gated device, this 2D *c*-MOF was further explored in FET-based biosensors.

The porous and tailored structure of 2D *c*-MOFs ensures the diversity of their FET devices. However, many challenges remain to date, such as the formation of the 2D *c*-MOF film with high smoothness and controllable thickness, tunability of band gaps, optimized integration for good contacts, *etc.*

6.1.2 Superconductors. The zero resistance and the Meissner effect characterizing superconductivity has been considered as an important parameter in power transmission, NMR imaging, maglev, and superconducting quantum interference devices (SQUIDs).⁶⁸ Cu₃(HTB) exhibiting a zero band gap and extraordinary conductivity could function as an excellent material candidate for this application.^{31,68,126} Through electrical resistivity, AC magnetic susceptibility, and specific heat measurements, Zhu and co-workers detected a superconducting transition at 0.25 K for Cu₃(HTB) (Fig. 20a), which remains the only example in this field. The transition depended on the magnetic field and disappeared at an applied field of 2500 Oe (Fig. 20b). Moreover, a second superconductivity phase with a transition temperature of ~3 K in the magnetic susceptibility measurement was determined and it is ascribed to the superconductivity of monolayers or few-layers of Cu₃(HTB).

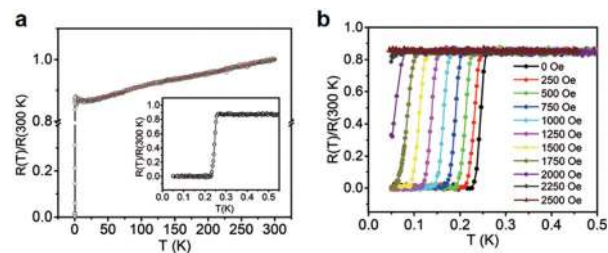


Fig. 20 Superconductivity of Cu₃(HTB). (a) Temperature-dependence of the normalized resistance from 0.05 to 300 K of Cu₃(HTB). (b) Temperature-dependence of the normalized resistance of Cu₃(HTB) in applied magnetic fields up to 2500 Oe. Reproduced with permission ref. 68. Copyright 2018, Wiley-VCH.

This observation is in accordance with the calculated results; the monolayer of Cu₃(HTB) has a predicted transition temperature of 4.43 K, while it decreased to 1.58 K for the bulk one.¹²⁶

6.1.3 Chemiresistors. Gas sensors are desirable for disease diagnosis and prevention as well as environmental protection.²⁴ Benefiting from the large surface to volume ratio, MOFs are able to serve as active materials. Nonetheless, due to the lack of intrinsic electrical conductivity in traditional MOFs, the fabricated sensors have been limited to devices with a complicated transducing mechanism such as luminescence quenching or enhancement.²⁸ Conductive 2D *c*-MOFs hold considerable promise for fast signal transduction in chemiresistive sensors, which represent one of the most intensively studied MOFtronics. The preliminary results reported to date indicate the immense potential of these materials for efficient detection of various gas analytes.

2D *c*-MOF based chemiresistors were demonstrated by Campbell *et al.* in 2015 initially.²⁸ Bulk Cu₃(HITP)₂ 2D *c*-MOF was transferred onto interdigitated gold electrodes by drop-casting an acetone dispersion of Cu₃(HITP)₂ to construct the device. The chemiresistor was capable of detection of NH₃ in parts per million (ppm) concentration level at room temperature, in spite of the existence of 60% relative humidity (RH) (Fig. 21a and b). However, the isostructural Ni₃(HITP)₂ 2D *c*-MOF did not produce any reliable response under identical conditions, indicating the impact of rational modification on metal nodes on the functionality. Later on, Mirica's group reported the utilization of Cu₃(HHTP)₂ and M₉(HHTP)₄ (M = Ni, Co) 2D *c*-MOFs for the detection of NH₃ by preparing the 2D *c*-MOF/graphite hybrids⁹² or assembling the 2D *c*-MOFs on graphitic electrodes¹⁷⁶ through the typical hydrothermal method. In 2018, combining experiment with simulation, Rubio-Gimenez *et al.* ascribed the origin of response of Cu₃(HHTP)₂ toward NH₃ to the coordination of NH₃ to Cu²⁺ in the network.¹²² Kelvin probe measurements revealed the difference of work function after exposing Cu₃(HHTP)₂ to NH₃, and infrared (IR) reflection absorption spectroscopy indicated the appearance of NH₃ stretching bands and the coordination between NH₃ and Cu atom in Cu₃(HHTP)₂. According to the simulation, the coordination leads to slight distortions of the electronic structure and triggers an increase in the band gap from 0.33 to 0.42 eV. Yao *et al.* integrated the Cu₃(HHTP)(HHB)



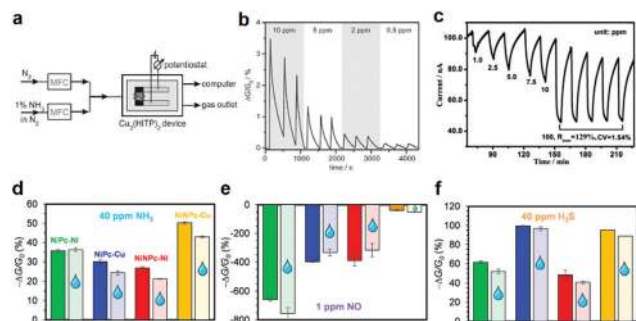


Fig. 21 2D *c*-MOF based chemiresistors for the detection of NH_3 , NO , and H_2S . (a and b) Schematic of the experimental apparatus and relative response curve of $\text{Cu}_3(\text{HHTP})_2$ toward NH_3 at different concentrations. Reproduced with permission ref. 28. Copyright 2015, Wiley-VCH. (c) The response–recovery curve of the $\text{Cu}_3(\text{HHTP})_2$ thin film toward NH_3 at different concentrations. Reproduced with permission ref. 96. Copyright 2017, Wiley-VCH. (d–f) Sensing responses of $\text{M}_2[\text{NiPcO}_8]$ and $\text{M}_2[\text{NiNpCo}_8]$ ($\text{M} = \text{Cu}, \text{Ni}$) upon 30 min exposure to 40 ppm of NH_3 , H_2S , and 1 ppm of NO in dry nitrogen and in the presence of H_2O . Reproduced with permission ref. 51. Copyright 2019, American Chemical Society.

2D *c*-MOF into a gas sensor by the drop-casting method which showed good sensing performance for NH_3 in the concentration range of 1–100 ppm.⁵⁹ IR spectroscopy and PXRD spectroscopy revealed the $\text{NH}_3\text{-Cu}^{+2/2+}$ interaction with an expanded *c* lattice upon exposure to NH_3 . Important advancement was reported by Xu and co-workers using the high quality $\text{Cu}_3(\text{HHTP})_2$ thin film prepared through the LS interfacial approach.⁹⁶ The resulting sensor exhibited a high response of 129% and >10% toward 100 and 1 ppm of NH_3 vapor, respectively (Fig. 21c). The authors calculated the theoretical limit of detection (LOD) to be ~ 0.5 ppm by setting the response as 10%. In addition, the response and recovery were accelerated by 54% and 10% respectively compared with those of the corresponding bulk $\text{Cu}_3(\text{HHTP})_2$ based sensor.

Besides NH_3 , Mirica and co-workers demonstrated $\text{Cu}_3(\text{HHTP})_2$ and $\text{Ni}_9(\text{HHTP})_4$ 2D *c*-MOFs as the active materials for the chemiresistive response to other toxic gases including NO and H_2S .¹⁷⁶ The same group integrated $\text{Ni}_9(\text{HHTP})_4$ or $\text{Ni}_3(\text{HHTP})_2$ into textile-supported gas sensors for the detection of NO and H_2S , which exhibited good mechanical flexibility.¹⁰² The sensors were able to detect NO and H_2S with high performance in the presence of humidity (18% RH) and were fully recoverable after washing the devices with water. Recently, Meng *et al.* reported chemiresistive sensors based on a series of phthalocyanine derivatives-based 2D *c*-MOFs ($\text{M}_2[\text{NiPcO}_8]$ and $\text{M}_2[\text{NiNpCo}_8]$, $\text{M} = \text{Cu}, \text{Ni}$).⁵¹ The resulting devices displayed outstanding humidity-independent ability during the detection and differentiation of NH_3 , H_2S , and NO , and delivered sub-ppm or part-per-billion (ppb) level theoretical LODs at room temperature (Fig. 21d–f).

In 2015, Campbell *et al.* contributed to a cross-reactive sensor array, which was capable of distinguishing five categories of volatile organic compounds (VOCs) (alcohols, amines, ketones/ethers, aromatic and aliphatic hydrocarbons) (Fig. 22a and b).⁹¹ Structurally analogous $\text{Cu}_3(\text{HHTP})_2$ and $\text{M}_3(\text{HHTP})_2$ ($\text{M} = \text{Cu}, \text{Ni}$) 2D *c*-MOFs were applied for the

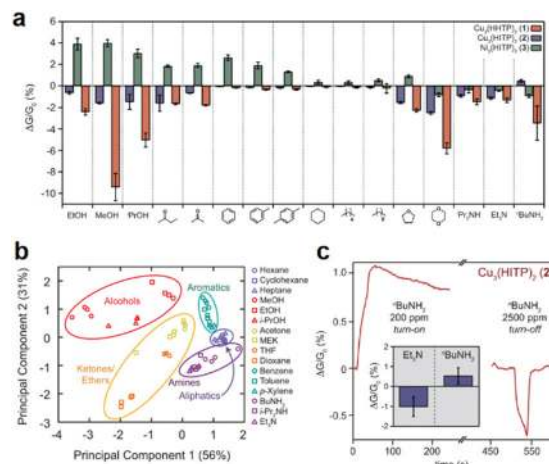


Fig. 22 2D *c*-MOF based chemiresistors for the detection of VOCs. (a and b) Sensing responses and related principal component analysis to various 200 ppm of VOC vapors for the 2D *c*-MOF array: $\text{M}_3(\text{HHTP})_2$ ($\text{M} = \text{Cu}, \text{Ni}$; $\text{X} = \text{O}, \text{NH}$). (c) Sensing response for $\text{Cu}_3(\text{HHTP})_2$ to ${}^n\text{BuNH}_2$. Reproduced with permission ref. 91. Copyright 2015, American Chemical Society.

fabrication of sensors through drop-casting. These 2D *c*-MOF based devices exhibited diverse sensing behaviors toward various VOC vapors at a concentration of 200 ppm. Both varying the metal nodes and ligands led to changes in the response to each gas. Furthermore, all samples delivered a high response to polar VOCs, while no appreciable signals were recorded for the aliphatic hydrocarbons. The authors noted both “turn-on” and “turn-off” responses of $\text{Cu}_3(\text{HHTP})_2$ for ${}^n\text{BuNH}_2$ at 200 and 2500 ppm respectively (Fig. 22c) and speculated a multiple sensing mechanism including charge transfer and hydrogen bonding among these 2D *c*-MOFs. Later on, Hoppe *et al.* prepared a 1–3 μm thick $\text{Cu}_3(\text{HHTP})_2$ film by spray-coating its dispersion on glass slides or flexible polycarbonate foils and fabricated chemiresistive sensors for methanol sensing accordingly.⁹³

Besides conventional analytes, the groups of Mirica and Dincă have applied 2D *c*-MOFs for the detection of ions ($\text{Cu}_3(\text{HHTP})_2$, and $\text{M}_9(\text{HHTP})_4$)¹⁷⁷ and CO_2 ($\text{Cu}_3(\text{HIB})_2$),¹⁷⁸ respectively. Despite numerous successful examples of 2D *c*-MOFs for sensors, the thus-far fabricated gas sensors through the compressing or drop-casting method based on bulk 2D *c*-MOFs obtained by solvothermal synthesis, have brought difficulties in realizing the mass production of devices and exhibited limited performance. Analogous to FETs, the rational tuning of the electronic and chemical structures, improving the quality of thin 2D *c*-MOF films, and further optimizing the integration methods thus leading to high performance remain to be explored. Moreover, fundamental understanding of the sensing mechanisms such as analyte-metal coordination, hydrogen bonding interaction, and guest–host interaction to ascertain neat structure–property relationship requires under investigation.

6.2 Optoelectronics

Conductive 2D *c*-MOFs with tailorable band gaps are appealing materials for optoelectronics, such as photodetectors, organic



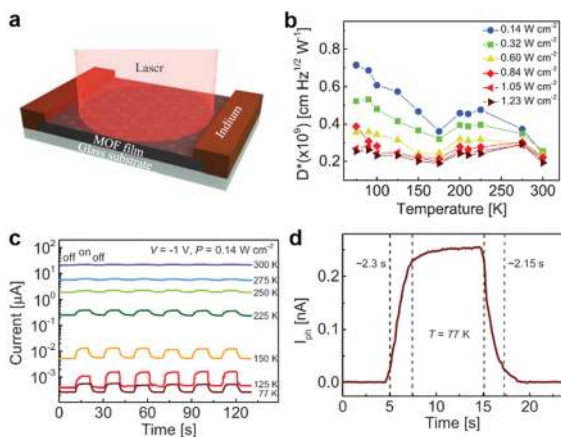


Fig. 23 $\text{Fe}_3(\text{HTTP})_2$ 2D *c*-MOF based photodetector. (a and b) Schematic of a two-terminal photodetector device based on 1.7 μm thick $\text{Fe}_3(\text{HTTP})_2$ and its dependence of detectivity at -1 V bias. (c) Temperature-dependent photoresponse versus time. (d) Zoomed time-resolved response at 77 K showing a response time of ~ 2 s.³⁰ Copyright 2020, Wiley-VCH.

light-emitting devices, solar cells, *etc.*^{30,143,179} Very recently, we and cooperation partner demonstrated a $\text{Fe}_3(\text{HTTP})_2$ based two-terminal photodetector device operated in photoconductive mode (Fig. 23a).³⁰ The record-high mobility and band-like charge transport of $\text{Fe}_3(\text{HTTP})_2$ allow effective transduction of generated signals. The related device was capable of detecting light ranging from UV to NIR range (400–1575 nm). At room temperature, due to thermally activated band-to-band population of charge carriers in this active material, the $\text{Fe}_3(\text{HTTP})_2$ possessing a narrow IR band gap of ~ 0.45 eV displayed limited performance. Cooling the devices to 77 K suppressed this thermal excitation hence leading to a stronger response and higher sensitivity (Fig. 23b–d). Upon 785 nm excitation at 77 K, a lower noise equivalent power was achieved with a specific detectivity of 7×10^8 $\text{cm Hz}^{1/2} \text{W}^{-1}$.

6.3 Spintronics

Merging high conductivity and long range magnetic ordering are highly desirable in spintronics for data storage and processing, such as spin transistors,¹⁸⁰ spin Hall effect transistors,¹⁸¹ spin photovoltaic,¹⁸² spin valves,¹⁸³ *etc.* The traditional active materials such as inorganic compounds or organic/molecular film semiconductors have displayed high performance. Nevertheless, the inorganic compounds and organic/molecular film semiconductors are facing the challenge of lack of tunability in magnetic/conductive properties or adequacy in sufficient mobilities, respectively.³² By contrast, in-plane fully π -conjugated 2D *c*-MOFs with structural diversity and high charge transport enable the interaction between the carriers and localized spins, hence providing promise of tunable magnetic ordering.^{32,67,184}

6.3.1 Ferromagnetic semiconductors based on 2D *c*-MOFs.

Ferromagnetic (FM) semiconductors as emerging materials have attracted considerable attention as FM electrodes in the next generation logic and memory devices.^{151,183,185–189} In 2018, our group reported the FM properties of the 2D *c*-MOF.³² A FM semiconductor based on the non-porous coronene-based

$\text{Fe}_3(\text{PTC})$ 2D *c*-MOF (Fig. 24a) presented a FM ground state at ~ 15 K in the magnetic susceptibility measurements, which is in accordance with the exchange energy ($E_{\text{exchange}} = E_{\text{anti-FM}} - E_{\text{FM}}$) of 1.22 meV and an estimated Curie point of ~ 16 K by DFT simulation (Fig. 24b and c). The distances of neighboring Fe atoms in $\text{Fe}_3(\text{PTC})$ were estimated as 0.54–1.20 and ~ 0.5 nm in the in-plane and out-of-plane direction respectively, which excludes the strong exchange between the d orbitals from Fe atoms directly. Therefore, we proposed an indirect exchange through the network for the observed ferromagnetism and highlighted the impact of high π -d hybridization between the d/p orbitals of the FeS_4 complex and coronene on the indirect exchange interaction. Later on, we developed another FM semiconductor based on the $\text{Fe}_2[\text{FePcO}_8]$ 2D *c*-MOF (Fig. 24d).³³ DFT calculations revealed a higher exchange energy of ~ 300 meV for FM coupling and suggested again an indirect exchange interaction benefiting from the strong π /d hybridization of the orbitals between FeO_4 and iron phthalocyanine. SQUID magnetometry showed a blocked superparamagnetic nature in $\text{Fe}_2[\text{FePcO}_8]$ due to the limited crystal size in the achieved sample and maintained magnetic ordering even at 350 K (Fig. 24e and f).

The contribution of strong π -d hybridization and strong π -conjugation to the charge transport along the entire network remains an open question. Nevertheless, we noted their influence on the extraordinary magnetic ordering among 2D *c*-MOFs. We also consider that both studies utilizing bulk materials did not exclude the impact of the grain boundaries, defects, edges, or the crystallite tilting on the magnetic properties. Improving the crystallinity to achieve single-crystals or single domain films will benefit the understanding of intrinsic magnetic properties and realizing ferromagnetism even at/above room temperature.

6.3.2 Nonmagnetic 2D *c*-MOFs as spacers. In addition to the compelling FM properties, nonmagnetic 2D *c*-MOFs are able to function as channel materials for spacers in spin valves or spin transistors.¹⁸³ In 2019, Song *et al.* applied the $\text{Cu}_3(\text{HHTP})_2$ film to fabricate organic spin valves (OSVs) with vertical architecture (Fig. 24g) and studied its spin injection and transport properties.⁹⁷ The authors selected initially the 100 nm thick $\text{Cu}_3(\text{HHTP})_2$ as the channel material between the top (Co) and bottom ($\text{La}_{0.67}\text{Sr}_{0.33}\text{MnO}_3$, LMSO) FM electrodes to minimize the penetration of Co into the 2D *c*-MOF network in this device configuration and prevent possible short circuit. The fabricated OSV exhibited a large negative magnetoresistance (MR) of 25% at 10 K (Fig. 24h), and this negative MR could still be observed at temperature up to 200 K.

However, the porous structure of $\text{Cu}_3(\text{HHTP})_2$ as a spacer in a vertical spin device may affect its long-term practicability due to the potential inter-diffusion of the upper electrode material of Co through the 1D channel of $\text{Cu}_3(\text{HHTP})_2$ to the LMSO electrode at the bottom.¹⁹⁰ Therefore, non-porous networks for preventing the penetration are more reasonable. Instead, the lateral architecture represents another approach for the integration of porous 2D *c*-MOFs into spintronic devices (Fig. 24i).



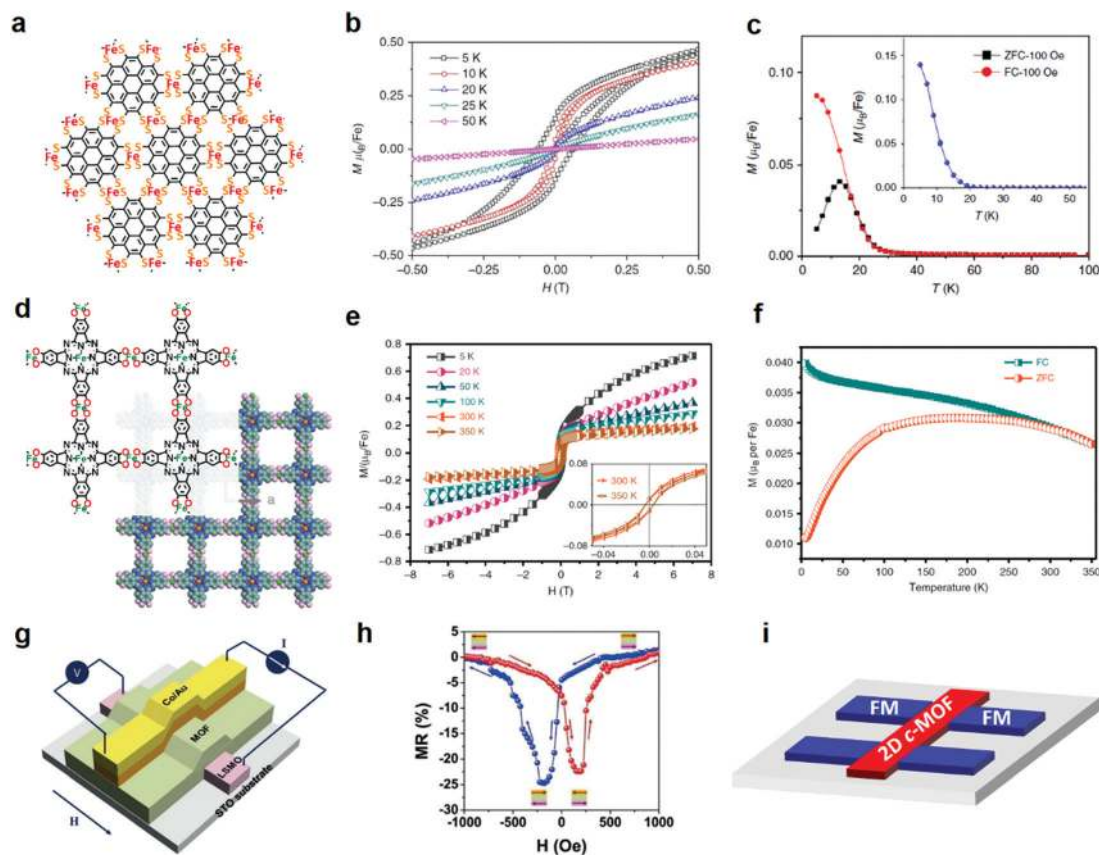


Fig. 24 Spin-properties of 2D *c*-MOFs. (a–c) $\text{Fe}_3(\text{PTC})$ 2D *c*-MOF and its ferromagnetic ground state, magnetic hysteresis loops at different temperatures, as well as field-cooled (FC)/zero-field-cooled (ZFC) magnetization as a function of temperature at an applied field of 100 Oe.³² Copyright 2018, Nature Publishing Group. (d–f) $\text{Fe}_2[\text{FePcO}_8]$ 2D *c*-MOF and its magnetic hysteresis loops at different temperatures, ZFC/FC magnetization at 100 Oe.³³ Copyright 2019, Nature Publishing Group. (g and h) Diagram of the $\text{Cu}_3(\text{HHTP})_2$ based vertical OSV and its magnetoresistance loop for at 10 K. Reproduced with permission ref. 97. Copyright 2020, Wiley-VCH. (i) Schematic illustration of lateral spin valves.

6.4 Thermoelectrics

The porous structure of 2D *c*-MOFs makes them great candidate for thermoelectric applications because the pores are capable of scattering phonons strongly.^{105,110} In 2017, Sun *et al.* applied $\text{Ni}_3(\text{HITP})_2$ to the investigation of its thermoelectric properties through a pellet sample.¹⁰⁵ The device of the pellet showed an ultralow thermal conductivity of $0.21 \text{ W m}^{-1} \text{ K}^{-1}$ and a high thermoelectric figure of merit (ZT) of 1.19×10^{-3} at 25°C . The ZT value increased 17-fold compared with traditional 7,7,8,8-tetracyanoquinodimethane-infiltrated $\text{Cu}_3(\text{benzene-1,3,5-tricarboxylate})_2$ ($\text{Cu}_3(\text{BTC})_2\text{-TCNQ}$) MOF based thermoelectric device. The room temperature conductivity of 58.8 S cm^{-1} correlates to an electronic (κ_e) contributed thermal conductivity of $4.28 \times 10^{-3} \text{ W m}^{-1} \text{ K}^{-1}$, hence suggesting a lattice dominated thermal transport in this 2D *c*-MOF.

Up to here, despite the great progress of 2D *c*-MOFs for (opto)electronics, spintronics, and thermoelectrics, the current studies in these fields rely largely on TP-based 2D *c*-MOFs, due to the contributions of diverse synthetic methodologies for the TP-based systems. The development of MOFtronics towards other 2D *c*-MOF systems is still in its infancy. In-depth understanding on producing/integrating methods and exploring

active materials with appropriate conductivity and band structure merits further exploration.

6.5 Electrochemical energy storage devices

Conductive 2D *c*-MOFs incorporating triphenylene-type monomers and abundant quinone-species, such as *o*-benzosemiquinone and *o*-diiminobenzosemiquinone (Fig. 7),^{36,40,51,56,57,191} have emerged as a novel class of multifunctional electrode materials for electrochemical energy storage. Among plenty of technologies,^{104,192–196} metal-ion batteries and supercapacitors represent two major energy storage devices.¹⁹⁵ Benefiting from the intrinsically conductive and porous properties, 2D *c*-MOFs have shown excellent performance in metal-ion batteries and supercapacitors, which could function as the power sources for the above mentioned MOFtronics.^{197–201}

6.5.1 Metal-ion batteries. Benzene-based 2D *c*-MOFs with dense redox active sites have been widely utilized as active cathode materials for lithium/sodium-ion batteries (Fig. 25a).^{35,36,87,88,202} As a typical example, Wada *et al.* reported the utilization of $\text{Ni}_3(\text{HIB})_2$ 2D *c*-MOFs in Li-ion batteries in 2018.³⁶ As a cathode material, $\text{Ni}_3(\text{HIB})_2$ delivered a high specific capacity of 155 mA h g^{-1} in a 1 M LiPF_6 electrolyte (Fig. 25b). Based on XPS and DFT, the authors proposed a



charge re-arrangement between N and Ni atoms once electron was injected, which shifts $\text{Ni}_3(\text{HIB})_2$ to a more stable structure. The bottom-up synthesized $\text{Cu}_3(\text{HHB})_2$ nanosheets (Section 5) hold promise for lithium-ion batteries owing to the high exposure of active sites.⁸⁸ The discharge capacity was determined as 153 mA h g^{-1} for the nanosheet samples in 1 M LiPF_6 at 0.1 A g^{-1} in $1.3\text{--}2.6 \text{ V}$ vs. Li^+/Li , and it was further pushed to 400 mA h g^{-1} in the potential window of $1.3\text{--}4.0 \text{ V}$ (Fig. 25c). The prepared battery exhibited outstanding cycling stability with a capacity retention of 90% after 1000 cycles. The ultrathin feature from the nanosheets for fast ion diffusion and high utilization of active sites was highlighted, while bulk $\text{Cu}_3(\text{HHB})_2$ powder provided limited performance. Park *et al.* utilized $\text{Co}_3(\text{HIB})_2$ for Na-ion storage in 2018.³⁵ This 2D *c*-MOF was found to be capable of storing three Na^+ per HIB ligand (corresponding to two Na^+ per CoN_4 linkage) with 1 M NaPF_6 electrolyte. As a result, an ultrahigh specific capacity of 291 mA h g^{-1} was recorded. XPS analysis suggested that the reversible redox reactions mainly proceeded on the *o*-diiminobenzosemiquinonate moiety, since no obvious valence change for Co^{2+} was detected. $\text{Ni}_3(\text{HIB})_2$ exhibited a similar capacity of 295 mA h g^{-1} under identical conditions.

Although Li/Na-ion batteries have been extensively studied, the high cost and safety issues impede further development. Instead, less toxic and cheaper aqueous zinc batteries are a good alternative. Recently, Nam *et al.* developed a $\text{Cu}_3(\text{HHTP})_2$ based Zn-ion battery (Fig. 25d) with a high capacity of 228 mA h g^{-1} .⁸⁹ The authors identified the role of H_2O in the performance. In an aqueous electrolyte of a 3 M aqueous solution of $\text{Zn}(\text{CF}_3\text{SO}_3)_2$, the hydrated Zn^{2+} was allowed to be

inserted into the 1D channels in the backbone with a high diffusion rate during discharge. As contrast, significantly higher interfacial resistance and a lower capacity of 144 mA h g^{-1} were recorded for the organic electrolyte.

Apart from serving as active materials, our group has explored 2D *c*-MOFs as the host for the non-conductive cathode material of iodine (I_2) in Na- I_2 batteries.²⁰³ $\text{M}_2[\text{CuPcO}_8]$ ($\text{M} = \text{Fe}, \text{Ni}, \text{Zn}$) were selected for hosting iodine and resolving the poor cycle stability of the I_2 cathode due to the dissolution of polyiodide in the electrolyte. The MO_4 linkages were confirmed to be effective for iodine adsorption; amongst, FeO_4 possessed the highest efficacy. The constructed $\text{Fe}_2[\text{CuPcO}_8]/\text{I}_2$ electrode delivered a high discharge capacity of 208 mA h g^{-1} and outstanding cycling stability with almost no capacity decay after 3200 cycles (Fig. 25e and f).

6.5.2 Supercapacitors. 2D *c*-MOFs have also represented promising electroactive materials for electrical double layer (EDL) capacitors,^{37,101,146,199} which possess a fast charge/discharge process, excellent cyclability, and superior power density with respect to metal-ion batteries.²⁰⁴ The capacitance of EDL capacitors arises from physical ion adsorption and charge separation on the electrode surface. In 2016, Sheberla *et al.* reported the exploration of $\text{Ni}_3(\text{HITP})_2$ for EDL capacitors.³⁷ Pelletized $\text{Ni}_3(\text{HITP})_2$ ($\sigma = 50 \text{ S cm}^{-1}$) with unique 1D channels with diameters of $\sim 1.5 \text{ nm}$ exhibited ideal EDL charge storage with a high gravimetric capacitance of $\sim 110 \text{ F g}^{-1}$ and an areal capacitance of $\sim 18 \text{ } \mu\text{F cm}^{-2}$ at a discharge rate of 0.05 A g^{-1} . The authors highlighted the role of the 1D channel in the diffusion and accommodation of the electrolyte ions to the EDL storage (Fig. 26a). To access insights into the relationship between the pore diameters and the performance, Bi *et al.* performed constant-potential molecular dynamic simulations on three types of 2D *c*-MOFs with room-temperature ionic liquids (RTILs) as the electrolyte.²⁰⁵ The 2D *c*-MOFs, $\text{Ni}_3(\text{HHB})_2$, $\text{Ni}_3(\text{HITP})_2$, and $\text{Ni}_3(\text{HITN})_2$ (HITN = 2,3,8,9,14,15-hexaminoctrinaphthylene), feature different pore sizes in the range of 0.8 to 2.4 nm (Fig. 26b). Among them, $\text{Ni}_3(\text{HITN})_2$ with the largest pore was estimated to exhibit the highest gravimetric capacitance and energy density. However, theory inferred highest values with respect to volumetric capacitance and energy density for $\text{Ni}_3(\text{HHB})_2$ with the narrowest 1D channel. On further reducing the pore size, ions could barely enter the channels leading to smaller or zero capacitance. In addition, the less conductive $\text{Cu}_3(\text{HHTP})_2$ was assembled directly onto a carbon fiber paper (as current collector) to function as electrodes in a EDL capacitor,⁹⁰ which exhibited a gravimetric capacitance of 120 F g^{-1} and an areal capacitance of $\sim 22 \text{ mF cm}^{-2}$ at 0.5 A g^{-1} in polyvinyl alcohol (PVA)/KCl gel electrolyte (Fig. 26c). The authors highlighted the mechanical adhesion between the current collector and the assembled 2D *c*-MOF on top for effective charge transport. Unsurprisingly, the electrode composed of coated bulk $\text{Cu}_3(\text{HHTP})_2$ with a polymer binder on the carbon paper is not that effective.

Apart from the conventional supercapacitors,^{206,207} efforts have been devoted to flexible micro-supercapacitors (MSCs) as well, which are promising micro-power sources for next-generation

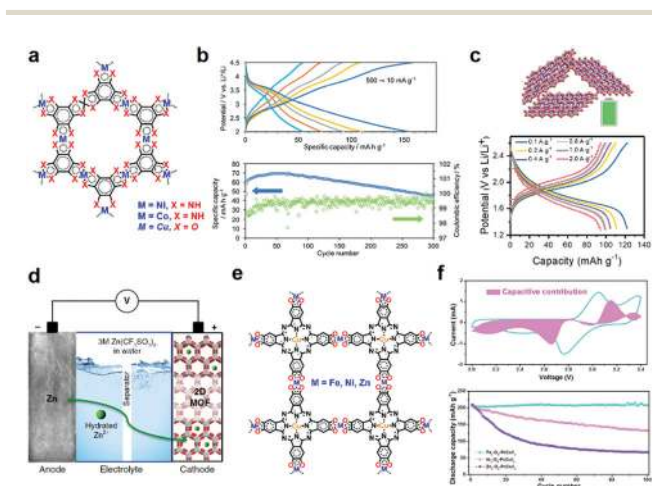


Fig. 25 Selected 2D *c*-MOF based metal-ion batteries. (a) Applied $\text{M}_3(\text{HXB})_2$ 2D *c*-MOFs for metal-ion batteries. When $\text{M} = \text{Cu}$, $\text{X} = \text{O}$, the charge state is -1 . (b) The charge/discharge curves and cycling stability test of $\text{Ni}_3(\text{HIB})_2$ in a Li-ion battery. Reproduced with permission ref. 36. Copyright 2018, Wiley-VCH. (c) Utilization of $\text{Cu}_3(\text{HHB})_2$ thin nanosheets in Li-ion batteries and the charge/discharge curves.⁸⁸ Copyright 2020, the Royal Society of Chemistry. (d) Illustration of $\text{Cu}_3(\text{HHTP})_2$ based aqueous Zn-ion battery.⁸⁹ Copyright 2019, Nature Publishing Group. (e and f) $\text{M}_2[\text{CuPcO}_8]$ as the host in a Na- I_2 battery and its electrochemical performance, the calculated capacitive contribution of the $\text{Fe}_2[\text{CuPcO}_8]/\text{I}_2$ electrode as well as the cycling stability.²⁰³ Copyright 2020, Wiley-VCH.



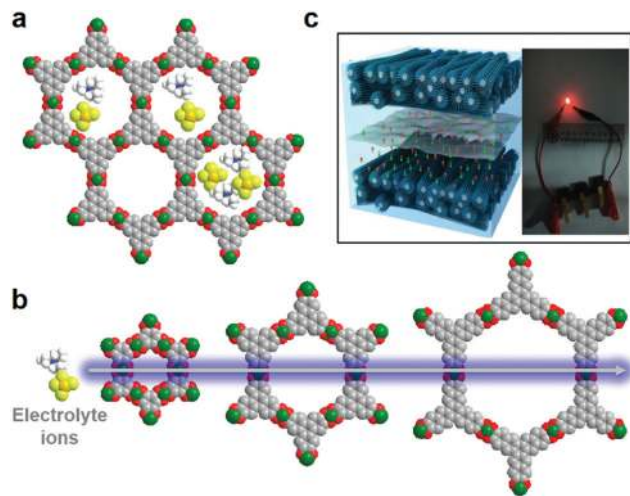


Fig. 26 EDL capacitors based on 2D *c*-MOFs. (a) Schematic of the accommodation of electrolyte ions in 2D *c*-MOF. (b) Schematic of the diffusion of electrolyte ions into various 2D *c*-MOFs with different sized 1D channels. (c) $\text{Cu}_3(\text{HHTP})_2$ on carbon fiber paper for supercapacitors, which powered a red light-emitting-diode. Reproduced with permission ref. 90. Copyright 2017, Wiley-VCH.

portable electronic devices.^{208–211} Very recently, Wu *et al.* applied laser scribing technique assisted solvothermal synthesis to interdigitate $\text{Ni}_9(\text{HHTP})_4$ on 3D porous laser scribed graphene (LSG) on a polyimide (PI) substrate.¹⁰⁰ A CO_2 laser was applied for patterning LSG, followed by O_2 -plasma treatment to serve as a current collector and the platform for the deposition of $\text{Ni}_9(\text{HHTP})_4$ (Fig. 27a and b). The hybrid delivered areal capacitance, energy density, and power density of 15.2 mF cm^{-2} , $4.1 \mu\text{W h cm}^{-2}$, and 7 mW cm^{-2} , respectively. Benefiting from the ultrathin feature of our exfoliated $\text{Ni}_2[\text{CuPc}(\text{NH})_8]$ nanosheets (Section 5) that allow facile solution-processibility through filtration, flexible MSC devices were constructed based on $\text{Ni}_2[\text{CuPc}(\text{NH})_8]$ /graphene hybrids on PI as film electrodes in a PVA/LiCl gel electrolyte (Fig. 27c).⁵⁵ The MSC device exhibited high areal capacitance and power density of 18.9 mF cm^{-2} and 168 mW cm^{-2} respectively as well as excellent cycling stability. Even on alternating the flat and bent states, the device still maintained 86.2% of the initial performance after 3000 cycles (Fig. 27d).

Beside EDL storage, 2D *c*-MOFs have displayed pseudo-capacitive charge storage^{25,26} in aqueous electrolytes, which differs from the EDL mechanism and involves surface reversible faradaic redox reactions.²⁰⁶ The incorporation of the pseudo-capacitive storage with redox reactions generally leads to enhancement in gravimetric and volumetric capacitances. In 2018, Feng *et al.* reported pseudo-capacitors based on $\text{Cu}_3(\text{HIB})_2$ and $\text{Ni}_3(\text{HIB})_2$ in 1 M KOH aqueous electrolyte.³⁸ The reversible pseudo-capacitive behaviors enabled $\text{Cu}_3(\text{HIB})_2$ and $\text{Ni}_3(\text{HIB})_2$ pellets to present a high gravimetric capacitance of 215 and over 400 F g^{-1} , respectively, while the contribution of EDL capacitances was estimated to be less than 20% and 10%, respectively. A $50 \mu\text{m}$ thick $\text{Ni}_3(\text{HIB})_2$ exhibited an excellent volumetric capacitance up to 760 F cm^{-3} . By further

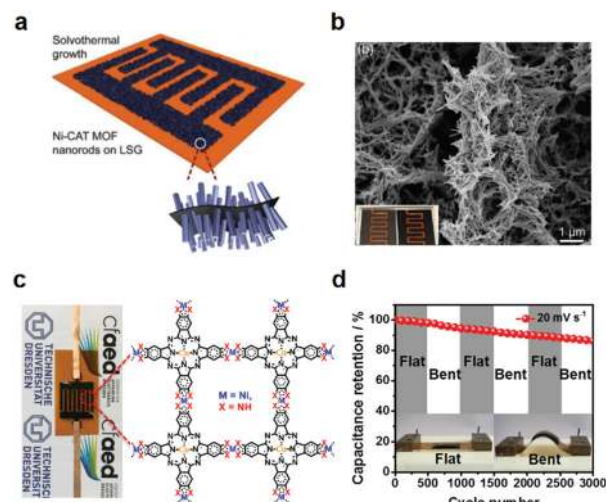


Fig. 27 2D *c*-MOFs based MSCs. (a and b) Illustration of LSG-supported $\text{Ni}_9(\text{HHTP})_4$ for MSC and its SEM image. Reproduced with permission ref. 100. Copyright 2019, Wiley-VCH. (c and d) Flexible MSC device based on $\text{Ni}_2[\text{CuPc}(\text{NH})_8]$ nanosheets.⁵⁵ Copyright 2020, Wiley-VCH.

thickening it to $360 \mu\text{m}$, the volumetric capacitance was revealed to be 460 F cm^{-3} and an areal capacitance was determined to be 17 F cm^{-2} . In addition, similar to those metal-ion batteries,^{35,36} the redox reaction in $\text{Ni}_3(\text{HIB})_2$ was speculated to be ligand-based,³⁷ as evidenced by a set of redox peaks from the ligand and excluded valence change of Ni atoms by XPS. Pseudo-capacitive behavior was further exemplified by the DBC-based $\text{Cu}_2[\text{DBC}_8]$ 2D *c*-MOF.⁵⁶ In a 1 M NaCl aqueous electrolyte, $\text{Cu}_2[\text{DBC}_8]$ displayed a nearly rectangular cyclic voltammetry curve with distinct redox peaks, indicative of both EDL and pseudo-capacitance for this material with estimated contributions of 62 and 38%, respectively. The supercapacitor provided high areal and volumetric capacitances of $\sim 0.9 \text{ F cm}^{-2}$ and 22 F cm^{-3} respectively in solid-state.

As demonstrated, owing to the high intrinsic porosity and abundant redox active linkages, 2D *c*-MOFs have been integrated into multifunctional electrochemical energy storage devices. Hybridization^{212–216} of 2D *c*-MOFs with other materials has been proved to be an approach for advancing the performance. Accordingly, various templates, such as carbon fiber paper,⁹⁰ cellulose nanofibers,¹⁰¹ graphene,¹⁰⁰ copper metal,²¹⁷ and polypyrrole membrane¹⁴⁶ have been utilized as platforms for the deposition of 2D *c*-MOFs. However, these samples are not structurally defined 2D *c*-MOFs anymore and exclude a neat structure–property relationship. Overall, the current studies provide guidelines corresponding to the rational design of chemical structure and modulation on the porosity and pore size.

7. Conclusions and outlook

In this review article, we introduce the diversity of the chemical structures of 2D *c*-MOFs based on various conjugated building blocks including benzene, triphenylene, trinaphthylene, truxene, hexaazatrinaphthylene, coronene, phthalocyanine,



naphthalocyanine, and dibenzo[*g,p*]chrysene with *ortho*-substituted functional groups (–OH, –NH₂, –SH, and –SeH). In addition, the multicomponent 2D *c*-MOFs comprising mixed ligands (e.g. Cu₃(HHTP)(HHB)) or mixed metals (e.g. M_nM'_{3–n}(HITP)₂) show great potential of tunability of the porosity, active sites, and electronic properties. The chemical tunability endows 2D *c*-MOFs with unique chemical and physical features, such as crystallinity, porosity, stability, tailorable band gaps, electrical conductivity, photoactivity, magnetic ordering, and topological state, thereby offering 2D *c*-MOFs great potential for broad applications in multiple MOFtronics. In this context, based on the thus-far reported examples, we provide an overview of the monomer/2D *c*-MOF design, controlled synthesis, device integration, and multifunction of 2D *c*-MOFs.

More specifically, we discuss the structure–electronic property relationship by considering the influence of molecular design (type and geometry of ligand, metal) and material structure (redox-activity in linkage, layer stacking and arrangement) as well as crystallinity. We further summarize the conductivities and mobilities for thus-far reported 2D *c*-MOFs, which reach 2500 S cm^{–1} (Cu₃(HTB), measured by 4-probe configuration) and 220 cm² V^{–1} s^{–1} (Fe₃(HITP)₂, hole mobility measured by Hall effect and TRTS) respectively at room temperature. Theoretically, DFT calculations demonstrated that 2D *c*-MOFs are expected to exhibit some nontrivial physical properties, such as topological state (single-layer of Ni₃(HTB)₂), high-temperature ferromagnetic ordering (Curie temperature of 170, 212, and 450 K for single-layer of Ni₂[MnPc(NH₈)], Mn₃(HTB)₂, and Mn₃(HIB)₂, respectively).^{34,84,85} Experimentally, various synthetic methodologies have been applied to the development of 2D *c*-MOFs as bulk powder (hydro-/solvo-thermal method) or film (wet-interface-assisted method) samples. The hydro-/solvo-thermal method is a straightforward strategy that allows diversity in the reaction conditions and offers large scale and high yield synthesis. Besides the generally obtained morphology of 2D *c*-MOFs as nanocrystals < 100 nm, this method also provides the possibility to synthesize moderate-sized single-crystals (1–10 μm). Among the reported systems, we highlight the significance of single-crystalline nanorods (e.g. Cu₃(HHTP)₂, Ni₃(HITP)₂) and nanoflakes (e.g. Cu₃(HHTP)₂, Cu₃(HHB)₂) on unambiguous understanding of the charge transport properties/mechanisms of 2D *c*-MOFs. By comparison, the wet-interface-assisted strategies including gas–liquid, liquid–liquid, and liquid–solid interfacial approaches show promise in constructing large-area, free-standing 2D *c*-MOF thin films, which ensure sufficient charge transport and facile device integration and hold potential for the realization of practical application of this class of materials for MOFtronics. Particularly, we highlight the construction of single-layer of 2D *c*-MOFs (e.g. Ni₃(HITP)₂, at the LB air/water interface), which is beneficial to the high exposure of active sites and excellent processibility. Next, device fabrications based on the obtained bulk or film samples are presented. To address the processing challenges faced by the bulk crystals, top-down exfoliation and bottom-up surfactant-assisted synthesis achieving homogeneous nanosheets

in solution are considered as efficient strategies to provide accessible active sites and solution-processibility.

Finally, we introduce the broad application of conductive 2D *c*-MOFs for MOFtronics covering (opto)electronics (e.g. FETs, superconductors, chemiresistors, photodetectors), spintronics (e.g. ferromagnetic semiconductors, spacers), and thermoelectrics, as well as the utilization of 2D *c*-MOFs as potential power sources for MOFtronics including battery and supercapacitor devices. Owing to the unique features of 2D *c*-MOFs combining large surface to volume ratio, defined active sites, and tailored electronic structure with high conductivity, photoactivity, or high-temperature ferromagnetic ordering, we especially emphasize the great potential of 2D *c*-MOFs for superconductors, photodetectors, and ferromagnetic semiconductors.

In addition to the foregoing applications, 2D *c*-MOFs are also potential alternatives to traditional electrocatalysts for promoting the reactivity and fundamentally understanding the mechanism^{218–221} in various electrochemical energy conversion reactions.^{222–229} MO₄[–] and MS₄-linkages have been widely demonstrated as the active sites towards the hydrogen evolution reaction^{45,49,61,62,118,230,231} and oxygen evolution reaction.^{54,160,232,233} For instance, the constructed single-layer of Ni₃(HITP)₂ exhibited an overpotential of 333 mV.⁴⁵ Apart from water splitting, 2D *c*-MOFs have emerged as model catalysts to investigate the mechanism during the oxygen reduction reaction.^{52,99,103,234–238} For instance, Co₂[CuPcO₈] delivered excellent activity with a half-wave overpotential of 830 mV, under a four-electron O₂ reduction process.⁵³ In regard to the emerging carbon dioxide reduction reaction (CO₂RR), based on an array of M₂[M'PcO₈] (M = Cu, Zn; M' = Cu, Zn) 2D *c*-MOFs with finely tuned active sites, our group has established a structure–property relationship regarding the selectivity of active sites toward carbon dioxide reduction or the concomitant hydrogen evolution.⁵³ Regardless of the preliminary success, the study towards complicated reactions such as CO₂RR is still at infancy, and in-depth investigation on the mechanisms remains of growing interest. In the past few years, 2D *c*-MOFs stand out for electrochemical molecular capture¹⁰⁹ and detection^{239,240} as well.

The development of 2D *c*-MOFs discussed here leaves no doubt that the extension of this class of materials to MOFtronics is of substantial interest to the next generation of logic and memory devices. In this context, it is important to clarify that the traditional photo-/electroactive materials should not be the metric for 2D *c*-MOFs and MOFtronics. 2D *c*-MOFs may outperform traditional electronic materials, but, this judgement cannot be true without specifying the particular property and device/technique type. For instance, the intrinsic room-temperature mobilities of electrons and holes for crystalline silicon were recorded in the range of 1450–1610 and 300–400 cm² V^{–1} s^{–1}, respectively *via* the Hall effect, which is far beyond the thus-far reported highest values recorded by the same technique for 2D *c*-MOFs (220 cm² V^{–1} s^{–1}). In spite of whether 2D *c*-MOFs become better than silicon in the future, MOFtronics will likely never replace the silicon-electronics. Besides, 2D *c*-MOFs represent layer-stacked framework materials



and the development of single- or few-layer 2D *c*-MOFs is still in its infancy. The lack of control on layer number and the ambiguous contribution of in-plane charge transport lead to difficulties to compare the electronic properties of 2D *c*-MOFs with other 2D materials. Nevertheless, the reported mobilities for 2D *c*-MOFs significantly exceed those of typical crystalline organic semiconductors (mostly $< 20 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$). The band gaps of 2D *c*-MOFs exhibit a range of values comparable to amorphous organic molecules/polymers. In addition, for semiconducting organic molecule/polymer single-crystals, it is quite complicated to thin the crystals into single- or few-layers, whereas free-standing 2D *c*-MOF nanosheets/films represent a kind of advantage in processing and device integration. Therefore, it is clear that 2D *c*-MOFs are a unique class of electronic materials that exhibit exceptional properties, *i.e.*, highly tailorable band structure, high porosity and stability. The immense progress from synthesis to function are highly encouraging and validate their unique properties underlying the applications. Nevertheless, the practical utilization of 2D *c*-MOFs beyond laboratories is yet to be realized. Many challenges remain to be addressed for ascertaining a neat structure–property relationship and promoting the development of this field.

(i) As illustrated, the library of 2D *c*-MOFs have considerably expanded with the development of novel ligands and synthetic methodologies. The known layered 2D *c*-MOFs exhibit anisotropic transport features; the strong π -*d* hybridization and extended π -conjugation together with the interlayer π - π interactions contribute to their charge transport. Nonetheless, the current ligand design and linkage chemistry are quite limited. Regardless of that multicomponent 2D *c*-MOFs allow further expansion of chemical structures and show promise in the tunability of band gaps in fine-scale, related investigation remains rare. Regarding the development of chemical methodologies, rational design of both conjugated ligands and metal nodes is responsible for the high electrical conductivity. More specifically, the molecular design (ligand type/geometry/size, metal valence/type, and π -conjugation degree) and the material structure (redox-activity in linkage, layer stacking and arrangement) are considered to be the major influential factors on the overall high conductivity of 2D *c*-MOFs but remain immature. Concerning the redox-activity in linkage, precise modulation of charge states *via* redox reactions leading to charge carrier (free radicals) injection plays an important role in the charge transport, which is however unexplored for many negatively charged 2D *c*-MOFs (*e.g.* $\text{Cu}_3(\text{HHB})_2$ with a charge state of -1). Meanwhile, after the redox reactions, thorough structural characterization including compositional and crystal analysis of 2D *c*-MOFs thus excluding potential decomposition/doping would be beneficial to ascertain a neat structure–conductivity relationship.

(ii) An important aspect with respect to the recorded charge transport properties (*e.g.* conductivity, metallicity, mobility, *etc.*) is the absence of the contributions of boundaries, defects, and edges in the polycrystalline samples having small domains, which results in an ambiguous charge transport mechanism. Regarding the extraordinary high-temperature ferromagnetic ordering, despite the highlighted significance of strong π -*d*

hybridization and π -conjugation, current investigation did not exclude these foregoing extrinsic factors. Thus, in terms of the fundamental physical property studies, it is clear that their boundaries and effects need to be considered and understood at a deeper level based on reliable methods and devices (*e.g.* single-crystal device). Improving the quality of crystals (crystallinity and domain size) to achieve single-crystals or single-domain films, controlling their morphology, and tuning the orientation of the nanocrystals thus minimizing the extrinsic influence are of key significance before establishing structure–property relationships.

(iii) Proof-of-concept functional investigation of conductive 2D *c*-MOFs have been successfully demonstrated in various MOFtronics. Yet the thus-far fabricated devices through the compressing or drop-casting method based on bulk 2D *c*-MOFs have brought difficulties in realizing the mass production of devices. Unsurprisingly, the reported performances are still far from satisfaction. Furthermore, the current functional studies rely largely on triphenylene-based systems; MOFtronics towards other 2D *c*-MOFs is still in its infancy. In-depth thinking on improving the quality and mechanical strength of 2D *c*-MOF thin films, optimizing the producing/integrating methods leading to high smoothness and good contacts, and exploring active materials with appropriate conductivity and band gap/structure remain to be explored. Moreover, fundamental understanding of the mechanism behind properties thus ascertaining neat structure–property relationship requires investigation. In addition to MOFtronics, the conductive and porous 2D *c*-MOFs also show great promise for developing future energy storage devices. Despite the substantial progress, challenges of probing the accurate charge storage mechanism on redox behavior and ion/electron diffusion remain ahead. In addition, utilization of these bulk porous materials through mechanical compression or deposition with support of binders on conductive platforms correlates with drawbacks including sacrificed pore volume and poor contact hence minimizing the performance in general, which rely on further optimizing the processing methods to overcome.

(iv) Currently, the novel device and new functions for 2D *c*-MOFs are largely limited. In regard to the nontrivial organic topological insulators, a single-crystalline monolayer of the 2D *c*-MOF (*e.g.* $\text{Ni}_3(\text{HTB})_2$) with a large size is still missing. The monolayer of 2D *c*-MOFs offers also other advantages: tunability at the atomic/molecular level and fabrication of 2D *c*-MOF/2D *c*-MOF or 2D *c*-MOF/2D materials (*e.g.* graphene, MoS_2) van der Waals heterostructures. Particularly, van der Waals heterostructures provide the possibility toward atomic/molecular-level structure design and multifunctional properties by selecting appropriate candidates but remained unexplored. Therefore, the major challenges for pushing MOFtronics still rely on the controlled chemical synthesis.

(v) Except for the foregoing points, we face many questions on the synthetic methodologies of 2D *c*-MOFs, which hinder in principle the advancement of 2D *c*-MOFs and MOFtronics with respect to property analysis, device integration, and function. Hydro-/solvothermal syntheses represent the most



representative strategy, but they barely allow control of the crystal growth and the obtained bulk samples are associated with difficulties in sample processing and device integration. Exfoliation of the bulk crystals into nanosheets has been recently explored, but the quality of the obtained nanosheets relying on the bulk samples remains unsatisfied. As mentioned, the development of large-area, free-standing 2D *c*-MOFs thin/thick films and monolayers synthesized *via* wet interface-assisted strategies allow facile device integration, sufficient charge transport, and fast signal transduction, hence offering the possibility toward high performance and long-term stability of MOFtronics. Nevertheless, the interfacial synthetic methods are limited to a certain range of systems (mainly based on benzene or triphenylene building blocks) hence limiting the application of other 2D *c*-MOFs. Towards controlled synthesis of 2D *c*-MOF crystals, rational design of the ligand structure/chemistry, favoring high/moderate solubility in certain solvents and high reversibility during the metal coordination as well as re-balancing the competition between in-plane coordination and out-of-plane van der Waals interactions shall be essential to enlarge the crystal sizes and reduce defects and grain boundaries. In addition, high mechanical strength, controlled orientation, as well as fine-tuned thickness and lateral size of the films are still synthetic challenges. Given all that, the synthesis of 2D *c*-MOFs as single-crystals or single-domain monolayers is urgently required for future advancement.²⁴¹

Over eight years of development, despite many open questions and the huge gap between this stage and the commercialization of MOFtronics, 2D *c*-MOFs have exhibited extraordinary electrical conductivity up to 2500 S cm⁻¹, band-like transport with mobility reaching ~220 cm² V⁻¹ s⁻¹ and tailorable band gaps that particularly encourages further studies. Furthermore, 2D *c*-MOFs feature many other unique properties of tunable porosity, redox and photoactivity, magnetic ordering and high stability, which make them appealing advanced electronic materials for the next-generation of electronic devices. It is expected that scalable production of 2D *c*-MOFs *via* controlled synthesis and facile processing will realize the MOFtronics in the near future.

Conflicts of interest

There are no conflicts of interest to declare.

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References

- H.-C. Zhou, J. R. Long and O. M. Yaghi, *Chem. Rev.*, 2012, **112**, 673.
- J. Liu and C. Wöll, *Chem. Soc. Rev.*, 2017, **46**, 5730.
- S. Krause, V. Bon, I. Senkowska, U. Stoeck, D. Wallacher, D. M. Töbrens, S. Zander, R. S. Pillai, G. Maurin, F.-X. Coudert and S. Kaskel, *Nature*, 2016, **532**, 348.
- S. Yuan, L. Feng, K. Wang, J. Pang, M. Bosch, C. Lollar, Y. Sun, J. Qin, X. Yang, P. Zhang, Q. Wang, L. Zou, Y. Zhang, L. Zhang, Y. Fang, J. Li and H.-C. Zhou, *Adv. Mater.*, 2018, **30**, 1704303.
- L. Liu, L. Li, M. E. Ziebel and T. D. Harris, *J. Am. Chem. Soc.*, 2020, **142**, 4705.
- S. Yuan, L. Huang, Z. Huang, D. Sun, J.-S. Qin, L. Feng, J. Li, X. Zou, T. Cagin and H.-C. Zhou, *J. Am. Chem. Soc.*, 2020, **142**, 4732.
- S. Yuan, J.-S. Qin, J. Li, L. Huang, L. Feng, Y. Fang, C. Lollar, J. Pang, L. Zhang, D. Sun, A. Alsalmé, T. Cagin and H.-C. Zhou, *Nat. Commun.*, 2018, **9**, 808.
- H. Furukawa, K. E. Cordova, M. O'Keeffe and O. M. Yaghi, *Science*, 2013, **341**, 1230444.
- L. J. Murray, M. Dincă and J. R. Long, *Chem. Soc. Rev.*, 2009, **38**, 1294.
- E. J. Kim, R. L. Siegelman, H. Z. H. Jiang, A. C. Forse, J.-H. Lee, J. D. Martell, P. J. Milner, J. M. Falkowski, J. B. Neaton, J. A. Reimer, S. C. Weston and J. R. Long, *Science*, 2020, **369**, 392.
- E. D. Bloch, W. L. Queen, R. Krishna, J. M. Zadrozny, C. M. Brown and J. R. Long, *Science*, 2012, **335**, 1606.
- Q. Yang, Q. Xu and H.-L. Jiang, *Chem. Soc. Rev.*, 2017, **46**, 4774.
- L. Jiao, Y. Wang, H.-L. Jiang and Q. Xu, *Adv. Mater.*, 2018, **30**, 1703663.
- H. Wang, Q.-L. Zhu, R. Zou and Q. Xu, *Chem*, 2017, **2**, 52.
- A. Schneemann, V. Bon, I. Schwedler, I. Senkowska, S. Kaskel and R. A. Fischer, *Chem. Soc. Rev.*, 2014, **43**, 6062.
- L. E. Kreno, K. Leong, O. K. Farha, M. Allendorf, R. P. Van Duyne and J. T. Hupp, *Chem. Rev.*, 2012, **112**, 1105.
- I. Stassen, N. Burtch, A. Talin, P. Falcaro, M. Allendorf and R. Ameloot, *Chem. Soc. Rev.*, 2017, **46**, 3185.
- A. A. Talin, A. Centrone, A. C. Ford, M. E. Foster, V. Stavila, P. Haney, R. A. Kinney, V. Szalai, F. El Gabaly, H. P. Yoon, F. Léonard and M. D. Allendorf, *Science*, 2014, **343**, 66.
- M. Hmadeh, Z. Lu, Z. Liu, F. Gándara, H. Furukawa, S. Wan, V. Augustyn, R. Chang, L. Liao, F. Zhou, E. Perre, V. Ozolins, K. Suenaga, X. Duan, B. Dunn, Y. Yamamoto, O. Terasaki and O. M. Yaghi, *Chem. Mater.*, 2012, **24**, 3511.
- L. Sun, M. G. Campbell and M. Dincă, *Angew. Chem., Int. Ed.*, 2016, **55**, 3566.
- R. Dong, T. Zhang and X. Feng, *Chem. Rev.*, 2018, **118**, 6189.
- W.-H. Li, W.-H. Deng, G.-E. Wang and G. Xu, *EnergyChem*, 2020, **2**, 100029.
- M. Ko, L. Mendecki and K. A. Mirica, *Chem. Commun.*, 2018, **54**, 7873.
- Z. Meng, R. M. Stolz, L. Mendecki and K. A. Mirica, *Chem. Rev.*, 2019, **119**, 478.



- 25 L. S. Xie, G. Skorupskii and M. Dincă, *Chem. Rev.*, 2020, **120**, 8536.
- 26 W.-T. Koo, J.-S. Jang and I.-D. Kim, *Chem*, 2019, **5**, 1938.
- 27 M. G. Campbell and M. Dincă, *Sensors*, 2017, **17**.
- 28 M. G. Campbell, D. Sheberla, S. F. Liu, T. M. Swager and M. Dincă, *Angew. Chem., Int. Ed.*, 2015, **54**, 4349.
- 29 G. Wu, J. Huang, Y. Zang, J. He and G. Xu, *J. Am. Chem. Soc.*, 2017, **139**, 1360.
- 30 H. Arora, R. Dong, T. Venanzi, J. Zscharschuch, H. Schneider, M. Helm, X. Feng, E. Cánovas and A. Erbe, *Adv. Mater.*, 2020, **32**, 1907063.
- 31 X. Huang, P. Sheng, Z. Tu, F. Zhang, J. Wang, H. Geng, Y. Zou, C.-a. Di, Y. Yi, Y. Sun, W. Xu and D. Zhu, *Nat. Commun.*, 2015, **6**, 7408.
- 32 R. Dong, Z. Zhang, D. C. Tranca, S. Zhou, M. Wang, P. Adler, Z. Liao, F. Liu, Y. Sun, W. Shi, Z. Zhang, E. Zschech, S. C. B. Mannsfeld, C. Felser and X. Feng, *Nat. Commun.*, 2018, **9**, 2637.
- 33 C. Yang, R. Dong, M. Wang, P. S. Petkov, Z. Zhang, M. Wang, P. Han, M. Ballabio, S. A. Bräuninger, Z. Liao, J. Zhang, F. Schwotzer, E. Zschech, H.-H. Klaus, E. Cánovas, S. Kaskel, M. Bonn, S. Zhou, T. Heine and X. Feng, *Nat. Commun.*, 2019, **10**, 3260.
- 34 W. Li, L. Sun, J. Qi, P. Jarillo-Herrero, M. Dincă and J. Li, *Chem. Sci.*, 2017, **8**, 2859.
- 35 J. Park, M. Lee, D. Feng, Z. Huang, A. C. Hinckley, A. Yakovenko, X. Zou, Y. Cui and Z. Bao, *J. Am. Chem. Soc.*, 2018, **140**, 10315.
- 36 K. Wada, K. Sakaushi, S. Sasaki and H. Nishihara, *Angew. Chem., Int. Ed.*, 2018, **57**, 8886.
- 37 D. Sheberla, J. C. Bachman, J. S. Elias, C. J. Sun, Y. Shao-Horn and M. Dincă, *Nat. Mater.*, 2017, **16**, 220.
- 38 D. Feng, T. Lei, M. R. Lukatskaya, J. Park, Z. Huang, M. Lee, L. Shaw, S. Chen, A. A. Yakovenko, A. Kulkarni, J. Xiao, K. Fredrickson, J. B. Tok, X. Zou, Y. Cui and Z. Bao, *Nat. Energy*, 2018, **3**, 30.
- 39 J. Park, A. C. Hinckley, Z. Huang, D. Feng, A. A. Yakovenko, M. Lee, S. Chen, X. Zou and Z. Bao, *J. Am. Chem. Soc.*, 2018, **140**, 14533.
- 40 J. H. Dou, L. Sun, Y. Ge, W. Li, C. H. Hendon, J. Li, S. Gul, J. Yano, E. A. Stach and M. Dincă, *J. Am. Chem. Soc.*, 2017, **139**, 13608.
- 41 X. Huang, H. Li, Z. Tu, L. Liu, X. Wu, J. Chen, Y. Liang, Y. Zou, Y. Yi, J. Sun, W. Xu and D. Zhu, *J. Am. Chem. Soc.*, 2018, **140**, 15153.
- 42 Y. Cui, J. Yan, Z. Chen, J. Zhang, Y. Zou, Y. Sun, W. Xu and D. Zhu, *Adv. Sci.*, 2019, **6**, 1802235.
- 43 Y. Jiang, I. Oh, S. H. Joo, O. Buyukcakir, X. Chen, S. H. Lee, M. Huang, W. K. Seong, S. K. Kwak, J.-W. Yoo and R. S. Ruoff, *J. Am. Chem. Soc.*, 2019, **141**, 16884.
- 44 L. Chen, J. Kim, T. Ishizuka, Y. Honsho, A. Saeki, S. Seki, H. Ihee and D. Jiang, *J. Am. Chem. Soc.*, 2009, **131**, 7287.
- 45 R. Dong, M. Pfeiffermann, H. Liang, Z. Zheng, X. Zhu, J. Zhang and X. Feng, *Angew. Chem., Int. Ed.*, 2015, **54**, 12058.
- 46 Y. Cui, J. Yan, Z. Chen, W. Xing, C. Ye, X. Li, Y. Zou, Y. Sun, C. Liu, W. Xu and D. Zhu, *iScience*, 2020, **23**, 100812.
- 47 Z. Meng and K. A. Mirica, *Nano Res.*, 2021, **14**, 369.
- 48 Q. Zhao, S. H. Li, R. L. Chai, X. Ren and C. Zhang, *ACS Appl. Mater. Interfaces*, 2020, **12**, 7504.
- 49 H. Huang, Y. Zhao, Y. Bai, F. Li, Y. Zhang and Y. Chen, *Adv. Sci.*, 2020, **7**, 2000012.
- 50 H. Nagatomi, N. Yanai, T. Yamada, K. Shiraishi and N. Kimizuka, *Chem. – Eur. J.*, 2018, **24**, 1806.
- 51 Z. Meng, A. Aykanat and K. A. Mirica, *J. Am. Chem. Soc.*, 2019, **141**, 2046.
- 52 H. Zhong, K. H. Ly, M. Wang, Y. Krupskaya, X. Han, J. Zhang, J. Zhang, V. Kataev, B. Büchner, I. M. Weidinger, S. Kaskel, P. Liu, M. Chen, R. Dong and X. Feng, *Angew. Chem., Int. Ed.*, 2019, **58**, 10677.
- 53 H. Zhong, M. Ghorbani-Asl, K. H. Ly, J. Zhang, J. Ge, M. Wang, Z. Liao, D. Makarov, E. Zschech, E. Brunner, I. M. Weidinger, J. Zhang, A. V. Krasheninnikov, S. Kaskel, R. Dong and X. Feng, *Nat. Commun.*, 2020, **11**, 1721.
- 54 H. Jia, Y. Yao, J. Zhao, Y. Gao, Z. Luo and P. Du, *J. Mater. Chem. A*, 2018, **6**, 1188.
- 55 M. Wang, H. Shi, P. Zhang, Z. Liao, M. Wang, H. Zhong, F. Schwotzer, A. S. Nia, E. Zschech, S. Zhou, S. Kaskel, R. Dong and X. Feng, *Adv. Funct. Mater.*, 2020, **30**, 2002664.
- 56 J. Liu, Y. Zhou, Z. Xie, Y. Li, Y. Liu, J. Sun, Y. Ma, O. Terasaki and L. Chen, *Angew. Chem., Int. Ed.*, 2020, **59**, 1081.
- 57 D. Sheberla, L. Sun, M. A. Blood-Forsythe, S. Er, C. R. Wade, C. K. Brozek, A. Aspuru-Guzik and M. Dincă, *J. Am. Chem. Soc.*, 2014, **136**, 8859.
- 58 T. Chen, J.-H. Dou, L. Yang, C. Sun, N. J. Libretto, G. Skorupskii, J. T. Miller and M. Dincă, *J. Am. Chem. Soc.*, 2020, **142**, 12367.
- 59 M. S. Yao, J. J. Zheng, A. Q. Wu, G. Xu, S. S. Nagarkar, G. Zhang, M. Tsujimoto, S. Sakaki, S. Horike, K. Otake and S. Kitagawa, *Angew. Chem., Int. Ed.*, 2020, **59**, 172.
- 60 V. Rubio-Giménez, S. Tatay and C. Martí-Gastaldo, *Chem. Soc. Rev.*, 2020, **49**, 5601.
- 61 R. Dong, Z. Zheng, D. C. Tranca, J. Zhang, N. Chandrasekhar, S. Liu, X. Zhuang, G. Seifert and X. Feng, *Chem. – Eur. J.*, 2017, **23**, 2255.
- 62 X. Sun, K.-H. Wu, R. Sakamoto, T. Kusamoto, H. Maeda, X. Ni, W. Jiang, F. Liu, S. Sasaki, H. Masunaga and H. Nishihara, *Chem. Sci.*, 2017, **8**, 8078.
- 63 X. Sun, K.-H. Wu, R. Sakamoto, T. Kusamoto, H. Maeda and H. Nishihara, *Chem. Lett.*, 2017, **46**, 1072.
- 64 J. Cui and Z. Xu, *Chem. Commun.*, 2014, **50**, 3986.
- 65 T. Pal, T. Kambe, T. Kusamoto, M. L. Foo, R. Matsuoka, R. Sakamoto and H. Nishihara, *ChemPlusChem*, 2015, **80**, 1255.
- 66 T. Pal, S. Doi, H. Maeda, K. Wada, C. M. Tan, N. Fukui, R. Sakamoto, S. Tsuneyuki, S. Sasaki and H. Nishihara, *Chem. Sci.*, 2019, **10**, 5218.
- 67 Y. Misumi, A. Yamaguchi, Z. Zhang, T. Matsushita, N. Wada, M. Tsuchiizu and K. Awaga, *J. Am. Chem. Soc.*, 2020, **142**, 16513.
- 68 X. Huang, S. Zhang, L. Liu, L. Yu, G. Chen, W. Xu and D. Zhu, *Angew. Chem., Int. Ed.*, 2018, **57**, 146.



- 69 N. T. T. Nguyen, H. Furukawa, F. Gándara, C. A. Trickett, H. M. Jeong, K. E. Cordova and O. M. Yaghi, *J. Am. Chem. Soc.*, 2015, **137**, 15394.
- 70 J. Huang, Y. He, M.-S. Yao, J. He, G. Xu, M. Zeller and Z. Xu, *J. Mater. Chem. A*, 2017, **5**, 16139.
- 71 R. Matheu, E. Gutierrez-Puebla, M. Á. Monge, C. S. Diercks, J. Kang, M. S. Prévot, X. Pei, N. Hanikel, B. Zhang, P. Yang and O. M. Yaghi, *J. Am. Chem. Soc.*, 2019, **141**, 17081.
- 72 D. L. Turner, T. P. Vaid, P. W. Stephens, K. H. Stone, A. G. DiPasquale and A. L. Rheingold, *J. Am. Chem. Soc.*, 2008, **130**, 14.
- 73 X. Huang, Y. Qiu, Y. Wang, L. Liu, X. Wu, Y. Liang, Y. Cui, Y. Sun, Y. Zou, J. Zhu, W. Fang, J. Sun, W. Xu and D. Zhu, *Angew. Chem., Int. Ed.*, 2020, **59**, 2.
- 74 T. Kambe, R. Sakamoto, K. Hoshiko, K. Takada, M. Miyachi, J. H. Ryu, S. Sasaki, J. Kim, K. Nakazato, M. Takata and H. Nishihara, *J. Am. Chem. Soc.*, 2013, **135**, 2462.
- 75 T. Kambe, R. Sakamoto, T. Kusamoto, T. Pal, N. Fukui, K. Hoshiko, T. Shimojima, Z. Wang, T. Hirahara, K. Ishizaka, S. Hasegawa, F. Liu and H. Nishihara, *J. Am. Chem. Soc.*, 2014, **136**, 14357.
- 76 M. D. Allendorf, R. Dong, X. Feng, S. Kaskel, D. Matoga and V. Stavila, *Chem. Rev.*, 2020, **120**, 8581.
- 77 L.-P. Tang, L.-M. Tang, D. Wang, H.-X. Deng and K.-Q. Chen, *J. Phys.: Condens. Matter*, 2018, **30**, 465301.
- 78 R. Dong, P. Han, H. Arora, M. Ballabio, M. Karakus, Z. Zhang, C. Shekhar, P. Adler, P. S. Petkov, A. Erbe, S. C. B. Mannsfeld, C. Felser, T. Heine, M. Bonn, X. Feng and E. Cánovas, *Nat. Mater.*, 2018, **17**, 1027.
- 79 A. J. Clough, J. M. Skelton, C. A. Downes, A. A. de la Rosa, J. W. Yoo, A. Walsh, B. C. Melot and S. C. Marinescu, *J. Am. Chem. Soc.*, 2017, **139**, 10863.
- 80 A. J. Clough, N. M. Orchanian, J. M. Skelton, A. J. Neer, S. A. Howard, C. A. Downes, L. F. J. Piper, A. Walsh, B. C. Melot and S. C. Marinescu, *J. Am. Chem. Soc.*, 2019, **141**, 16323.
- 81 D. Herebian, E. Bothe, F. Neese, T. Weyhermüller and K. Wieghardt, *J. Am. Chem. Soc.*, 2003, **125**, 9116.
- 82 X. Chen, Y. Hu, D. Wu, L. Weng and B. Kang, *Polyhedron*, 1991, **10**, 2651.
- 83 A. Das, Z. Han, W. W. Brennessel, P. L. Holland and R. Eisenberg, *ACS Catal.*, 2015, **5**, 1397.
- 84 J. Liu and Q. Sun, *ChemPhysChem*, 2015, **16**, 614.
- 85 M. Zhao, A. Wang and X. Zhang, *Nanoscale*, 2013, **5**, 10404.
- 86 M. A. Springer, T.-J. Liu, A. Kuc and T. Heine, *Chem. Soc. Rev.*, 2020, **49**, 2007.
- 87 Q. Jiang, P. Xiong, J. Liu, Z. Xie, Q. Wang, X. Q. Yang, E. Hu, Y. Cao, J. Sun, Y. Xu and L. Chen, *Angew. Chem., Int. Ed.*, 2020, **59**, 5273.
- 88 Z. Wang, G. Wang, H. Qi, M. Wang, M. Wang, S. Park, H. Wang, M. Yu, U. Kaiser, A. Fery, S. Zhou, R. Dong and X. Feng, *Chem. Sci.*, 2020, **11**, 7665.
- 89 K. W. Nam, S. S. Park, R. dos Reis, V. P. Dravid, H. Kim, C. A. Mirkin and J. F. Stoddart, *Nat. Commun.*, 2019, **10**, 4948.
- 90 W.-H. Li, K. Ding, H.-R. Tian, M.-S. Yao, B. Nath, W.-H. Deng, Y. Wang and G. Xu, *Adv. Funct. Mater.*, 2017, **27**, 1702067.
- 91 M. G. Campbell, S. F. Liu, T. M. Swager and M. Dincă, *J. Am. Chem. Soc.*, 2015, **137**, 13780.
- 92 M. Ko, A. Aykanat, M. K. Smith and K. A. Mirica, *Sensors*, 2017, 17.
- 93 B. Hoppe, K. D. J. Hindricks, D. P. Warwas, H. A. Schulze, A. Mohmeyer, T. J. Pinkvos, S. Zailskas, M. R. Krey, C. Belke, S. König, M. Fröba, R. J. Haug and P. Behrens, *CrystEngComm*, 2018, **20**, 6458.
- 94 A. Mähringer, A. C. Jakowetz, J. M. Rotter, B. J. Bohn, J. K. Stolarczyk, J. Feldmann, T. Bein and D. D. Medina, *ACS Nano*, 2019, **13**, 6711.
- 95 R. W. Day, D. K. Bediako, M. Rezaee, L. R. Parent, G. Skorupskii, M. Q. Arguilla, C. H. Hendon, I. Stassen, N. C. Gianneschi, P. Kim and M. Dincă, *ACS Cent. Sci.*, 2019, **5**, 1959.
- 96 M.-S. Yao, X.-J. Lv, Z.-H. Fu, W.-H. Li, W.-H. Deng, G.-D. Wu and G. Xu, *Angew. Chem., Int. Ed.*, 2017, **56**, 16510.
- 97 X. Song, X. Wang, Y. Li, C. Zheng, B. Zhang, C.-a. Di, F. Li, C. Jin, W. Mi, L. Chen and W. Hu, *Angew. Chem., Int. Ed.*, 2020, **59**, 1118.
- 98 V. Rubio-Giménez, M. Galbiati, J. Castells-Gil, N. Almora-Barrios, J. Navarro-Sánchez, G. Escorcía-Ariza, M. Mattera, T. Arnold, J. Rawle, S. Tatay, E. Coronado and C. Martí-Gastaldo, *Adv. Mater.*, 2018, **30**, 1704291.
- 99 E. M. Miner, L. Wang and M. Dincă, *Chem. Sci.*, 2018, **9**, 6286.
- 100 H. Wu, W. Zhang, S. Kandambeth, O. Shekhah, M. Eddaoudi and H. N. Alshareef, *Adv. Energy Mater.*, 2019, **9**, 1900482.
- 101 S. Zhou, X. Kong, B. Zheng, F. Huo, M. Strømme and C. Xu, *ACS Nano*, 2019, **13**, 9578.
- 102 M. K. Smith and K. A. Mirica, *J. Am. Chem. Soc.*, 2017, **139**, 16759.
- 103 Y. Lian, W. Yang, C. Zhang, H. Sun, Z. Deng, W. Xu, L. Song, Z. Ouyang, Z. Wang, J. Guo and Y. Peng, *Angew. Chem., Int. Ed.*, 2020, **59**, 286.
- 104 Y. Zang, F. Pei, J. Huang, Z. Fu, G. Xu and X. Fang, *Adv. Energy Mater.*, 2018, **8**, 1802052.
- 105 L. Sun, B. Liao, D. Sheberla, D. Kraemer, J. Zhou, E. A. Stach, D. Zakharov, V. Stavila, A. A. Talin, Y. Ge, M. D. Allendorf, G. Chen, F. Léonard and M. Dincă, *Joule*, 2017, **1**, 168.
- 106 K. Yuan, T. Song, X. Zhu, B. Li, B. Han, L. Zheng, J. Li, X. Zhang and W. Hu, *Small*, 2019, **15**, 1804845.
- 107 B. Wang, Y. Luo, B. Liu and G. Duan, *ACS Appl. Mater. Interfaces*, 2019, **11**, 35935.
- 108 X. Huang, H. Yao, Y. Cui, W. Hao, J. Zhu, W. Xu and D. Zhu, *ACS Appl. Mater. Interfaces*, 2017, **9**, 40752.
- 109 L. Mendecki, M. Ko, X. Zhang, Z. Meng and K. A. Mirica, *J. Am. Chem. Soc.*, 2017, **139**, 17229.
- 110 Z. Chen, Y. Cui, Y. Jin, L. Liu, J. Yan, Y. Sun, Y. Zou, Y. Sun, W. Xu and D. Zhu, *J. Mater. Chem. C*, 2020, **8**, 8199.
- 111 Z. Chen, Y. Cui, C. Ye, L. Liu, X. Wu, Y. Sun, W. Xu and D. Zhu, *Chem. – Eur. J.*, 2020, **26**, 12868.



- 112 Z. F. Wang, N. Su and F. Liu, *Nano Lett.*, 2013, **13**, 2842.
- 113 S. Chen, J. Dai and X. C. Zeng, *Phys. Chem. Chem. Phys.*, 2015, **17**, 5954.
- 114 L. Sun, C. H. Hendon, S. S. Park, Y. Tulchinsky, R. Wan, F. Wang, A. Walsh and M. Dincă, *Chem. Sci.*, 2017, **8**, 4450.
- 115 M. L. Aubrey, B. M. Wiers, S. C. Andrews, T. Sakurai, S. E. Reyes-Lillo, S. M. Hamed, C.-J. Yu, L. E. Darago, J. A. Mason, J.-O. Baeg, F. Grandjean, G. J. Long, S. Seki, J. B. Neaton, P. Yang and J. R. Long, *Nat. Mater.*, 2018, **17**, 625.
- 116 J. G. Park, M. L. Aubrey, J. Oktawiec, K. Chakarawet, L. E. Darago, F. Grandjean, G. J. Long and J. R. Long, *J. Am. Chem. Soc.*, 2018, **140**, 8526.
- 117 X. Wu, Y. Qiu, Z. Chen, B. Guan, X. Hao, A. I. Rykov, Y. Sun, L. Liu, Y. Zou, J. Sun, W. Xu and D. Zhu, *Angew. Chem., Int. Ed.*, 2020, **59**, 1.
- 118 A. J. Clough, J. W. Yoo, M. H. Mecklenburg and S. C. Marinescu, *J. Am. Chem. Soc.*, 2015, **137**, 118.
- 119 A. C. Hinckley, J. Park, J. Gomes, E. Carlson and Z. Bao, *J. Am. Chem. Soc.*, 2020, **142**, 11123.
- 120 L.-P. Tang, L.-M. Tang, H. Geng, Y.-P. Yi, Z. Wei, K.-Q. Chen and H.-X. Deng, *Appl. Phys. Lett.*, 2018, **112**, 012101.
- 121 G. Skorupskii, B. A. Trumpf, T. W. Kasel, C. M. Brown, C. H. Hendon and M. Dincă, *Nat. Chem.*, 2020, **12**, 131.
- 122 V. Rubio-Gimenez, N. Almora-Barrios, G. Escorcía-Ariza, M. Galbiati, M. Sessolo, S. Tatay and C. Marti-Gastaldo, *Angew. Chem., Int. Ed.*, 2018, **57**, 15086.
- 123 L. Dong, Y. Kim, D. Er, A. M. Rappe and V. B. Shenoy, *Phys. Rev. Lett.*, 2016, **116**, 096601.
- 124 M. E. Foster, K. Sohlberg, C. D. Spataru and M. D. Allendorf, *J. Phys. Chem. C*, 2016, **120**, 15001.
- 125 M. E. Foster, K. Sohlberg, M. D. Allendorf and A. A. Talin, *J. Phys. Chem. Lett.*, 2018, **9**, 481.
- 126 X. Zhang, Y. Zhou, B. Cui, M. Zhao and F. Liu, *Nano Lett.*, 2017, **17**, 6166.
- 127 Y. Jiang, I. Oh, S. H. Joo, Y.-S. Seo, S. H. Lee, W. K. Seong, Y. J. Kim, J. Hwang, S. K. Kwak, J.-W. Yoo and R. S. Ruoff, *J. Am. Chem. Soc.*, 2020, **142**, 18346.
- 128 B. Lukose, A. Kuc and T. Heine, *Chem. – Eur. J.*, 2011, **17**, 2388.
- 129 A. Kuc, M. A. Springer, K. Batra, R. Juarez-Mosqueda, C. Wöll and T. Heine, *Adv. Funct. Mater.*, 2020, **30**, 1908004.
- 130 A. P. Côté, A. I. Benin, N. W. Ockwig, M. O’Keeffe, A. J. Matzger and O. M. Yaghi, *Science*, 2005, **310**, 1166.
- 131 M. J. Kory, M. Wörle, T. Weber, P. Payamyar, S. W. van de Poll, J. Dshemuchadse, N. Trapp and A. D. Schlüter, *Nat. Chem.*, 2014, **6**, 779.
- 132 T. Ma, E. A. Kapustin, S. X. Yin, L. Liang, Z. Zhou, J. Niu, L.-H. Li, Y. Wang, J. Su, J. Li, X. Wang, W. D. Wang, W. Wang, J. Sun and O. M. Yaghi, *Science*, 2018, **361**, 48.
- 133 A. M. Evans, L. R. Parent, N. C. Flanders, R. P. Bisbey, E. Vitaku, M. S. Kirschner, R. D. Schaller, L. X. Chen, N. C. Gianneschi and W. R. Dichtel, *Science*, 2018, **361**, 52.
- 134 J. Ogle, N. Lahiri, C. Jaye, C. J. Tassone, D. A. Fischer, J. Louie and L. Whittaker-Brooks, *Adv. Funct. Mater.*, 2020, 2006920, DOI: 10.1002/adfm.202006920.
- 135 N. Lahiri, N. Lotfizadeh, R. Tsuchikawa, V. V. Deshpande and J. Louie, *J. Am. Chem. Soc.*, 2017, **139**, 19.
- 136 J. Sakamoto, J. van Heijst, O. Lukin and A. D. Schlüter, *Angew. Chem., Int. Ed.*, 2009, **48**, 1030.
- 137 L. Wang, H. Sahabudeen, T. Zhang and R. Dong, *npj 2D Mater. Appl.*, 2018, **2**, 26.
- 138 T. Zhang, H. Qi, Z. Liao, Y. D. Horev, L. A. Panes-Ruiz, P. S. Petkov, Z. Zhang, R. Shivhare, P. Zhang, K. Liu, V. Bezugly, S. Liu, Z. Zheng, S. Mannsfeld, T. Heine, G. Cuniberti, H. Haick, E. Zschech, U. Kaiser, R. Dong and X. Feng, *Nat. Commun.*, 2019, **10**, 4225.
- 139 K. Liu, L. Wang and R. Dong, *J. Mater. Chem. C*, 2020, **8**, 10696.
- 140 K. Liu, H. Qi, R. Dong, R. Shivhare, M. Addicoat, T. Zhang, H. Sahabudeen, T. Heine, S. Mannsfeld, U. Kaiser, Z. Zheng and X. Feng, *Nat. Chem.*, 2019, **11**, 994.
- 141 X. Zhuang, Y. Mai, D. Wu, F. Zhang and X. Feng, *Adv. Mater.*, 2015, **27**, 403.
- 142 X. Feng and A. D. Schlüter, *Angew. Chem., Int. Ed.*, 2018, **57**, 13748.
- 143 S. Liu, Y. C. Wang, C. M. Chang, T. Yasuda, N. Fukui, H. Maeda, P. Long, K. Nakazato, W. B. Jian, W. Xie, K. Tsukagoshi and H. Nishihara, *Nanoscale*, 2020, **12**, 6983.
- 144 A. Ciesielski and P. Samori, *Chem. Soc. Rev.*, 2014, **43**, 381.
- 145 V. Nicolosi, M. Chhowalla, M. G. Kanatzidis, M. S. Strano and J. N. Coleman, *Science*, 2013, **340**, 1226419.
- 146 R. Hou, M. Miao, Q. Wang, T. Yue, H. Liu, H. S. Park, K. Qi and B. Y. Xia, *Adv. Energy Mater.*, 2020, **10**, 1901892.
- 147 X. Du, J. Zhang, H. Wang, Z. Huang, A. Guo, L. Zhao, Y. Niu, X. Li, B. Wu and Y. Liu, *Mater. Chem. Front.*, 2020, **4**, 243.
- 148 M. Wang, M. Ballabio, M. Wang, H.-H. Lin, B. P. Biswal, X. Han, S. Paasch, E. Brunner, P. Liu, M. Chen, M. Bonn, T. Heine, S. Zhou, E. Cánovas, R. Dong and X. Feng, *J. Am. Chem. Soc.*, 2019, **141**, 16810.
- 149 B. P. Biswal, S. Valligatla, M. Wang, T. Banerjee, N. A. Saad, B. M. K. Mariserla, N. Chandrasekhar, D. Becker, M. Addicoat, I. Senkowska, R. Berger, D. N. Rao, S. Kaskel and X. Feng, *Angew. Chem., Int. Ed.*, 2019, **58**, 6896.
- 150 S. Xu, G. Wang, B. P. Biswal, M. Addicoat, S. Paasch, W. Sheng, X. Zhuang, E. Brunner, T. Heine, R. Berger and X. Feng, *Angew. Chem., Int. Ed.*, 2019, **58**, 849.
- 151 L. E. Darago, M. L. Aubrey, C. J. Yu, M. I. Gonzalez and J. R. Long, *J. Am. Chem. Soc.*, 2015, **137**, 15703.
- 152 R. Sakamoto, K. Takada, T. Pal, H. Maeda, T. Kambe and H. Nishihara, *Chem. Commun.*, 2017, **53**, 5781.
- 153 Q. Jiang, C. Zhou, H. Meng, Y. Han, X. Shi, C. Zhan and R. Zhang, *J. Mater. Chem. A*, 2020, **8**, 15271.
- 154 K. Zhao, W. Zhu, S. Liu, X. Wei, G. Ye, Y. Su and Z. He, *Nanoscale Adv.*, 2020, **2**, 536.
- 155 A.-Q. Wu, W.-Q. Wang, H.-B. Zhan, L.-A. Cao, X.-L. Ye, J.-J. Zheng, P. N. Kumar, K. Chiranjeevulu, W.-H. Deng, G.-E. Wang, M.-S. Yao and G. Xu, *Nano Res.*, 2021, **14**, 438.
- 156 L.-A. Cao, M.-S. Yao, H.-J. Jiang, S. Kitagawa, X.-L. Ye, W.-H. Li and G. Xu, *J. Mater. Chem. A*, 2020, **8**, 9085.
- 157 G. Iannaccone, F. Bonaccorso, L. Colombo and G. Fiori, *Nat. Nanotechnol.*, 2018, **13**, 183.



- 158 E. Pomerantseva and Y. Gogotsi, *Nat. Energy*, 2017, **2**, 17089.
- 159 G. Fiori, F. Bonaccorso, G. Iannaccone, T. Palacios, D. Neumaier, A. Seabaugh, S. K. Banerjee and L. Colombo, *Nat. Nanotechnol.*, 2014, **9**, 768.
- 160 M. Zhang, B.-H. Zheng, J. Xu, N. Pan, J. Yu, M. Chen and H. Cao, *Chem. Commun.*, 2018, **54**, 13579.
- 161 R. Makiura, S. Motoyama, Y. Umemura, H. Yamanaka, O. Sakata and H. Kitagawa, *Nat. Mater.*, 2010, **9**, 565.
- 162 Z. A. Gao, C.-H. Hsu, J. Liu, F.-C. Chuang, R. Zhang, B. Xia, H. Xu, L. Huang, Q. Jin, P. N. Liu and N. Lin, *Nanoscale*, 2019, **11**, 878.
- 163 M. Lischka, R. Dong, M. Wang, N. Martsinovich, M. Fritton, L. Grossmann, W. M. Heckl, X. Feng and M. Lackinger, *Chem. – Eur. J.*, 2019, **25**, 1975.
- 164 X. Meng, E. Kolodzeiski, X. Huang, A. Timmer, B. Schulze Lammers, H.-Y. Gao, H. Mönig, L. Liu, W. Xu, S. Amirjalayer, D. Zhu and H. Fuchs, *ChemNanoMat*, 2020, **6**, 1.
- 165 R. Zhang, J. Liu, Y. Gao, M. Hua, B. Xia, P. Knecht, A. C. Papageorgiou, J. Reichert, J. V. Barth, H. Xu, L. Huang and N. Lin, *Angew. Chem., Int. Ed.*, 2020, **59**, 2669.
- 166 H. Qi, H. Sahabudeen, B. Liang, M. Položij, M. A. Addicoat, T. E. Gorelik, M. Hamsch, M. Mundszinger, S. Park, B. V. Lotsch, S. C. B. Mannsfeld, Z. Zheng, R. Dong, T. Heine, X. Feng and U. Kaiser, *Sci. Adv.*, 2020, **6**, eabb5976.
- 167 Y. Peng, Y. Li, Y. Ban, H. Jin, W. Jiao, X. Liu and W. Yang, *Science*, 2014, **346**, 1356.
- 168 I.-Y. Jeon, Y.-R. Shin, G.-J. Sohn, H.-J. Choi, S.-Y. Bae, J. Mahmood, S.-M. Jung, J.-M. Seo, M.-J. Kim, D. Wook Chang, L. Dai and J.-B. Baek, *Proc. Natl. Acad. Sci. U. S. A.*, 2012, **109**, 5588.
- 169 P.-Z. Li, Y. Maeda and Q. Xu, *Chem. Commun.*, 2011, **47**, 8436.
- 170 S. Yang, P. Zhang, F. Wang, A. G. Ricciardulli, M. R. Lohe, P. W. M. Blom and X. Feng, *Angew. Chem., Int. Ed.*, 2018, **57**, 15491.
- 171 Z. Zhang, S. Yang, P. Zhang, J. Zhang, G. Chen and X. Feng, *Nat. Commun.*, 2019, **10**, 2920.
- 172 M. Zhao, Y. Huang, Y. Peng, Z. Huang, Q. Ma and H. Zhang, *Chem. Soc. Rev.*, 2018, **47**, 6267.
- 173 H. Shi, M. Li, A. Shaygan Nia, M. Wang, S. Park, Z. Zhang, M. R. Lohe, S. Yang and X. Feng, *Adv. Mater.*, 2020, **32**, 1907244.
- 174 S. C. Junggeburth, L. Diehl, S. Werner, V. Duppel, W. Sigle and B. V. Lotsch, *J. Am. Chem. Soc.*, 2013, **135**, 6157.
- 175 M. Zhao, Y. Wang, Q. Ma, Y. Huang, X. Zhang, J. Ping, Z. Zhang, Q. Lu, Y. Yu, H. Xu, Y. Zhao and H. Zhang, *Adv. Mater.*, 2015, **27**, 7372.
- 176 M. K. Smith, K. E. Jensen, P. A. Pivak and K. A. Mirica, *Chem. Mater.*, 2016, **28**, 5264.
- 177 L. Mendecki and K. A. Mirica, *ACS Appl. Mater. Interfaces*, 2018, **10**, 19248.
- 178 I. Stassen, J. H. Dou, C. Hendon and M. Dincă, *ACS Cent. Sci.*, 2019, **5**, 1425.
- 179 Z. Jin, J. Yan, X. Huang, W. Xu, S. Yang, D. Zhu and J. Wang, *Nano Energy*, 2017, **40**, 376.
- 180 S. Datta and B. Das, *Appl. Phys. Lett.*, 1990, **56**, 665.
- 181 J. Wunderlich, B.-G. Park, A. C. Irvine, L. P. Zárbo, E. Rozkotová, P. Nemeč, V. Novák, J. Sinova and T. Jungwirth, *Science*, 2010, **330**, 1801.
- 182 X. Sun, S. Vélez, A. Atxabal, A. Bedoya-Pinto, S. Parui, X. Zhu, R. Llopis, F. Casanova and L. E. Hueso, *Science*, 2017, **357**, 677.
- 183 V. A. Dediu, L. E. Hueso, I. Bergenti and C. Taliani, *Nat. Mater.*, 2009, **8**, 707.
- 184 L. Yang, X. He and M. Dincă, *J. Am. Chem. Soc.*, 2019, **141**, 10475.
- 185 I.-R. Jeon, B. Negru, R. P. Van Duyne and T. D. Harris, *J. Am. Chem. Soc.*, 2015, **137**, 15699.
- 186 J. A. DeGayner, I.-R. Jeon, L. Sun, M. Dincă and T. D. Harris, *J. Am. Chem. Soc.*, 2017, **139**, 4175.
- 187 K. S. Pedersen, P. Perlepe, M. L. Aubrey, D. N. Woodruff, S. E. Reyes-Lillo, A. Reinholdt, L. Voigt, Z. Li, K. Borup, M. Rouzières, D. Samohvalov, F. Wilhelm, A. Rogalev, J. B. Neaton, J. R. Long and R. Clérac, *Nat. Chem.*, 2018, **10**, 1056.
- 188 A. E. Thorarinsdottir and T. D. Harris, *Chem. Rev.*, 2020, **120**, 8716.
- 189 X. Li, X. Li and J. Yang, *J. Phys. Chem. Lett.*, 2020, **11**, 4193.
- 190 D. Sun, E. Ehrenfreund and Z. Vally Vardeny, *Chem. Commun.*, 2014, **50**, 1781.
- 191 Y. Shi, M. R. Momeni, Y.-J. Chen, Z. Zhang and F. A. Shakib, *Chem. Mater.*, 2020, **32**, 9664.
- 192 J. Sun, L. Guo, X. Sun, J. Zhang, Y. Liu, L. Hou and C. Yuan, *J. Mater. Chem. A*, 2019, **7**, 24788.
- 193 S. Gu, Z. Bai, S. Majumder, B. Huang and G. Chen, *J. Power Sources*, 2019, **429**, 22.
- 194 W. Zhou, S. Lv, X. Liu, Y. Li and J. Liu, *Chem. Commun.*, 2019, **55**, 11207.
- 195 A. Nazir, H. T. T. Le, C. W. Min, A. Kasbe, J. Kim, C. S. Jin and C. J. Park, *Nanoscale*, 2020, **12**, 1629.
- 196 F. Xu, Y. Zhai, E. Zhang, Q. Liu, G. Jiang, X. Xu, Y. Qiu, X. Liu, H. Wang and S. Kaskel, *Angew. Chem., Int. Ed.*, 2020, **59**, 19460.
- 197 M. Yu, R. Dong and X. Feng, *J. Am. Chem. Soc.*, 2020, **142**, 12903.
- 198 J. Liu, X. Song, T. Zhang, S. Liu, H. Wen and L. Chen, *Angew. Chem., Int. Ed.*, 2020, **59**, 2.
- 199 X. Li, X. Yang, H. Xue, H. Pang and Q. Xu, *EnergyChem*, 2020, **2**, 100027.
- 200 K. Wada, H. Maeda, T. Tsuji, K. Sakaushi, S. Sasaki and H. Nishihara, *Inorg. Chem.*, 2020, **59**, 10604.
- 201 B. Xu, H. Zhang, H. Mei and D. Sun, *Coord. Chem. Rev.*, 2020, **420**, 213438.
- 202 Z. Wu, D. Adekoya, X. Huang, M. J. Kiefel, J. Xie, W. Xu, Q. Zhang, D. Zhu and S. Zhang, *ACS Nano*, 2020, **14**, 12016.
- 203 F. Wang, Z. Liu, C. Yang, H. Zhong, G. Nam, P. Zhang, R. Dong, Y. Wu, J. Cho, J. Zhang and X. Feng, *Adv. Mater.*, 2020, **32**, 1905361.
- 204 P. Zhang, F. Wang, M. Yu, X. Zhuang and X. Feng, *Chem. Soc. Rev.*, 2018, **47**, 7426.



- 205 S. Bi, H. Banda, M. Chen, L. Niu, M. Chen, T. Wu, J. Wang, R. Wang, J. Feng, T. Chen, M. Dincă, A. A. Kornyshev and G. Feng, *Nat. Mater.*, 2020, **19**, 552.
- 206 M. Yu, H. Shao, G. Wang, F. Yang, C. Liang, P. Rozier, C.-Z. Wang, X. Lu, P. Simon and X. Feng, *Nat. Commun.*, 2020, **11**, 1348.
- 207 M. Yu, D. Lin, H. Feng, Y. Zeng, Y. Tong and X. Lu, *Angew. Chem., Int. Ed.*, 2017, **56**, 5454.
- 208 K. Jiang, I. A. Baburin, P. Han, C. Yang, X. Fu, Y. Yao, J. Li, E. Cánovas, G. Seifert, J. Chen, M. Bonn, X. Feng and X. Zhuang, *Adv. Funct. Mater.*, 2020, **30**, 1908243.
- 209 P. Zhang, Y. Li, G. Wang, F. Wang, S. Yang, F. Zhu, X. Zhuang, O. G. Schmidt and X. Feng, *Adv. Mater.*, 2019, **31**, 1806005.
- 210 C. Yang, K. S. Schellhammer, F. Ortmann, S. Sun, R. Dong, M. Karakus, Z. Mics, M. Löffler, F. Zhang, X. Zhuang, E. Cánovas, G. Cuniberti, M. Bonn and X. Feng, *Angew. Chem., Int. Ed.*, 2017, **56**, 3920.
- 211 Y. He, S. Yang, Y. Fu, F. Wang, J. Ma, G. Wang, G. Chen, M. Wang, R. Dong, P. Zhang and X. Feng, *Small Struct.*, 2020, 2000095, DOI: 10.1002/ssstr.202000095.
- 212 Y. Zhong, B. Cheng, C. Park, A. Ray, S. Brown, F. Mujid, J.-U. Lee, H. Zhou, J. Suh, K.-H. Lee, A. J. Mannix, K. Kang, S. J. Sibener, D. A. Muller and J. Park, *Science*, 2019, **366**, 1379.
- 213 F. Bonaccorso, L. Colombo, G. Yu, M. Stoller, V. Tozzini, A. C. Ferrari, R. S. Ruoff and V. Pellegrini, *Science*, 2015, **347**, 1246501.
- 214 G. Wang, J. Zhang, S. Yang, F. Wang, X. Zhuang, K. Müllen and X. Feng, *Adv. Energy Mater.*, 2018, **8**, 1702254.
- 215 H. Peng, J. Raya, F. Richard, W. Baaziz, O. Ersen, A. Ciesielski and P. Samorì, *Angew. Chem., Int. Ed.*, 2020, **59**, 2.
- 216 M. Yu, N. Chandrasekhar, R. K. M. Raghupathy, K. H. Ly, H. Zhang, E. Dmitrieva, C. Liang, X. Lu, T. D. Kühne, H. Mirhosseini, I. M. Weidinger and X. Feng, *J. Am. Chem. Soc.*, 2020, **142**, 19570.
- 217 Q. Ma, P. Yin, M. Zhao, Z. Luo, Y. Huang, Q. He, Y. Yu, Z. Liu, Z. Hu, B. Chen and H. Zhang, *Adv. Mater.*, 2019, **31**, 1808249.
- 218 J. Zhang, Z. Zhao, Z. Xia and L. Dai, *Nat. Nanotechnol.*, 2015, **10**, 444.
- 219 L. Dai, Y. Xue, L. Qu, H.-J. Choi and J.-B. Baek, *Chem. Rev.*, 2015, **115**, 4823.
- 220 G. Chen, J. Zhang, F. Wang, L. Wang, Z. Liao, E. Zschech, K. Müllen and X. Feng, *Chem. – Eur. J.*, 2018, **24**, 18413.
- 221 J. Zhang, G. Chen, K. Müllen and X. Feng, *Adv. Mater.*, 2018, **30**, 1800528.
- 222 G. Chen, T. Wang, J. Zhang, P. Liu, H. Sun, X. Zhuang, M. Chen and X. Feng, *Adv. Mater.*, 2018, **30**, 1706279.
- 223 H. Wang, X.-B. Li, L. Gao, H.-L. Wu, J. Yang, L. Cai, T.-B. Ma, C.-H. Tung, L.-Z. Wu and G. Yu, *Angew. Chem., Int. Ed.*, 2018, **57**, 192.
- 224 M. Yu, Z. Wang, C. Hou, Z. Wang, C. Liang, C. Zhao, Y. Tong, X. Lu and S. Yang, *Adv. Mater.*, 2017, **29**, 1602868.
- 225 H. Zhong, J. Wang, F. Meng and X. Zhang, *Angew. Chem., Int. Ed.*, 2016, **55**, 9937.
- 226 F. Wei, T. Wang, X. Jiang, Y. Ai, A. Cui, J. Cui, J. Fu, J. Cheng, L. Lei, Y. Hou and S. Liu, *Adv. Funct. Mater.*, 2020, **30**, 2002092.
- 227 D. Xing, Y. Wang, P. Zhou, Y. Liu, Z. Wang, P. Wang, Z. Zheng, H. Cheng, Y. Dai and B. Huang, *Appl. Catal., B*, 2020, **278**, 119295.
- 228 W. Xiong, X. Cheng, T. Wang, Y. Luo, J. Feng, S. Lu, A. M. Asiri, W. Li, Z. Jiang and X. Sun, *Nano Res.*, 2020, **13**, 1008.
- 229 W. Sheng, I. Amin, C. Neumann, R. Dong, T. Zhang, E. Wegener, W.-L. Chen, P. Förster, H. Q. Tran, M. Löffler, A. Winter, R. D. Rodriguez, E. Zschech, C. K. Ober, X. Feng, A. Turchanin and R. Jordan, *Small*, 2019, **15**, 1805228.
- 230 C. A. Downes, A. J. Clough, K. Chen, J. W. Yoo and S. C. Marinescu, *ACS Appl. Mater. Interfaces*, 2018, **10**, 1719.
- 231 J. Wang, J. Wang, X. Song, S. Qi and M. Zhao, *Appl. Surf. Sci.*, 2020, **511**, 145393.
- 232 W.-H. Li, J. Lv, Q. Li, J. Xie, N. Ogiwara, Y. Huang, H. Jiang, H. Kitagawa, G. Xu and Y. Wang, *J. Mater. Chem. A*, 2019, **7**, 10431.
- 233 C. Li, L. Shi, L. Zhang, P. Chen, J. Zhu, X. Wang and Y. Fu, *J. Mater. Chem. A*, 2020, **8**, 369.
- 234 E. M. Miner, T. Fukushima, D. Sheberla, L. Sun, Y. Surendranath and M. Dincă, *Nat. Commun.*, 2016, **7**, 10942.
- 235 E. M. Miner, S. Gul, N. D. Ricke, E. Pastor, J. Yano, V. K. Yachandra, T. Van Voorhis and M. Dincă, *ACS Catal.*, 2017, **7**, 7726.
- 236 H. Yoon, S. Lee, S. Oh, H. Park, S. Choi and M. Oh, *Small*, 2019, **15**, 1805232.
- 237 X.-H. Liu, W.-L. Hu, W.-J. Jiang, Y.-W. Yang, S. Niu, B. Sun, J. Wu and J.-S. Hu, *ACS Appl. Mater. Interfaces*, 2017, **9**, 28473.
- 238 S. S. Shinde, C. H. Lee, J.-Y. Jung, N. K. Wagh, S.-H. Kim, D.-H. Kim, C. Lin, S. U. Lee and J.-H. Lee, *Energy Environ. Sci.*, 2019, **12**, 727.
- 239 F. Wu, W. Fang, X. Yang, J. Xu, J. Xia and Z. Wang, *J. Chin. Chem. Soc.*, 2019, **66**, 522.
- 240 M. Ko, L. Mendecki, A. M. Eagleton, C. G. Durbin, R. M. Stolz, Z. Meng and K. A. Mirica, *J. Am. Chem. Soc.*, 2020, **142**, 11717.
- 241 J.-H. Dou, M. Q. Arguilla, Y. Luo, J. Li, W. Zhang, L. Sun, J. L. Mancuso, L. Yang, T. Chen, L. R. Parent, G. Skorupskii, N. J. Libretto, C. Sun, M. C. Yang, P. V. Dip, E. J. Brignole, J. T. Miller, J. Kong, C. H. Hendon, J. Sun and M. Dincă, *Nat. Mater.*, 2020, DOI: 10.1038/s41563-020-00847-7.

