

CrystEngComm

rsc.li/crystengcomm



ISSN 1466-8033

PAPER

Ishtvan Boldog, Christoph Janiak *et al.*
Metal-organic framework structures of fused hexagonal motifs with cuprophilic interactions of a triangular Cu(I)_3 (pyrazolate-benzoate) metallo-linker



Cite this: *CrystEngComm*, 2022, 24, 3675

Metal–organic framework structures of fused hexagonal motifs with cuprophilic interactions of a triangular Cu(I)₃(pyrazolate-benzoate) metallo-linker†

Saskia Menzel,^{‡a} Tobias Heinen,^{‡a} Ishtvan Boldog,^{‡*a} Thi Hai Yen Beglau,^a Shanghua Xing,^{ab} Alex Spieß,^a Dennis Woschko^{‡a} and Christoph Janiak^{‡*a}

The reaction of the N,O-heteroditopic bifunctional ligand 4-(3,5-dimethyl-1*H*-pyrazol-4-yl)benzoic acid (H₂-mpba) with Cu(NO₃)₂·2.5H₂O and Zn(NO₃)₂·4H₂O or Zn(CH₃COO)₂·2H₂O in *N,N*-dimethylformamide (DMF) results in concomitant formation of three bimetallic metal–organic frameworks (MOFs) with open structures, [Me₂NH₂][Zn₄(Cu₃(mpba)₃)₃(Me₂NH)-(DMF)₂] (**1**), [Zn₆(Cu₃(mpba)₃)₄(DMF)₅] (**2**) and [Zn₃(Cu₃(mpba)₃)₂(DMF)₃-(H₂O)][Zn₄(μ₄-O)(Cu₃(mpba)₃)₂(H₂O)₄] (**3**). Compounds **1** and **3** are isolable in phase pure form under specified conditions. The HSAB hard/soft metal–ligand selectivity (Cu^I-N and Zn–O) allows the structure assembly based on the 3-*c* trigonal planar {Cu₃(pz)₃} moiety (pz = pyrazolate) and various 4-6-*c* {Zn_x(O)_y(COO)_z} (*x* = 2–4, *y* = 0–1, *z* = 4–6) secondary building units to be realized neatly. The *in situ* formed softer Cu^I atoms coordinate with the softer pyrazolate nitrogen donor atoms of the pyrazolate–carboxylate ligand, leaving the carboxylate groups for linking to the Zn atoms. The triangular and trinuclear {Cu₃(mpba)₃}³⁻ metallacycle acts as a tricarboxylate linker between the Zn_{*x*} nodes and can be regarded as an expanded version of the benzene-1,3,5-tricarboxylate linker. In all structures the {Cu₃(pz)₃} moiety is stacked in infinite columns by cuprophilic interactions suggesting the initial formation of the Cu₃(mpba) metallo-ligand. The variable geometry Zn clusters, which are paddle-wheel type, pyramidal and trigonal-prismatic, adapt to the stacked orientation of the carboxylate groups from the {Cu₃(mpba)₃}³⁻ metallacycles. As a result, the formed structures could be viewed as derived from stacked hexagonal honeycomb **hcb** nets, *via* ‘fusion’ of alternating stacked nodes. The resulting nets retain the geometric similarity to the **hcb** net when viewed along the normal direction to the stack, and the individual topologies represent open structures. The minority phase **2** features 2D layers with a ‘dual-strand’ **hcb** net, {4³}{4³·6¹²}. **1** consists of parallel polycatenation, 2D + 2D → 3D, of triple layers with a point symbol of {4·6²}{4²·6⁶·8²}{6³}{6⁵·8}, while **3** presents a pair of interpenetrated 3D + 3D {4·6·8}{4²·6⁵·8⁵·10³} nets as a rare case of hetero-interpenetration. The binodal net in **3** is a rare **sqc**-3,6-Fdd2-2 type **sqc**-subnet and the first example in a MOF. In spite of the interpenetration, **3** retains significant porosity and could be activated to demonstrate permanent porosity, thereby classifiable as a MOF with an Ar-based BET surface area of 762 m² g⁻¹ (87 K) and a CO₂ adsorption capacity at 1 bar of 78 cm³ g⁻¹ (273 K) and 46 cm³ g⁻¹ (293 K). The structure organization similarities of **1**–**3**, stipulated by cuprophilic interactions, could have a general importance for linear heteroditopic ligands analogous to H₂mpba offering a ‘fused-**hcb**’ design approach towards open structures.

Received 25th February 2022,
Accepted 24th April 2022

DOI: 10.1039/d2ce00268j

rsc.li/crystengcomm

^a Institut für Anorganische Chemie und Strukturchemie, Heinrich Heine-Universität Düsseldorf, Universitätsstraße 1, Düsseldorf 40225, Germany.

E-mail: boldogi@hhu.de, janiak@uni-duesseldorf.de

^b Hoffmann Institute of Advanced Materials, Shenzhen Polytechnic, 7098 Liuxian Blvd, Nanshan District, Shenzhen 518055, China

† Electronic supplementary information (ESI) available: Synthesis, crystal photographs, X-ray data and refinement details, additional crystallographic figures, topological analysis, atomic absorption spectroscopy, SEM EDX, TGA, IR, XPS, gas sorption data, and linker flexibility. CCDC 2105164–2105166. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d2ce00268j>

‡ These authors contributed equally.

Introduction

The research area of metal–organic frameworks (MOFs) has reached a high degree of maturity with highly varied topologies^{1,2} and wide-spread properties.^{3–5} MOFs are built from metal clusters, also called secondary building units (SBUs), and bridging organic ligands, so-called linkers.^{6,7} In the rational design of MOFs, the isoreticularity principle foresees the use of linkers with different sizes but the same geometry to obtain MOFs of the same topology. Benzene-



1,3,5-tricarboxylate (btc^{3-} , Scheme 1) is an important linker, used for example in the prototypical MOF Cu-btc (HKUST-1),⁸ Zn-btc^{9–12} or Cd-btc.¹⁰ Extended versions of this linker are 4,4',4''-benzene-1,3,5-triyl-tribenzoate, btb^{3-} , 4,4',4''-(benzene-1,3,5-triyl-tris(benzene-4,1-diyl)tribenzoate, bbc^{3-} = 1,3,5-tris(4'-carboxy[1,1'-biphenyl]-4-yl)benzene, tcbbp^{3-} , 4,4',4''-(benzene-1,3,5-triyl-tris(ethyne-2,1-diyl)tribenzoate, bte^{3-} , 4,4',4''-s-triazine-2,4,6-triyl-tribenzoate, tatb^{3-} (Scheme 1) and 4,4',4''-(triazine-2,4,6-triyl-tris(benzene-4,1-diyl)tribenzoate (tapb^{3-}) as used in the MOFs listed in Table 1.

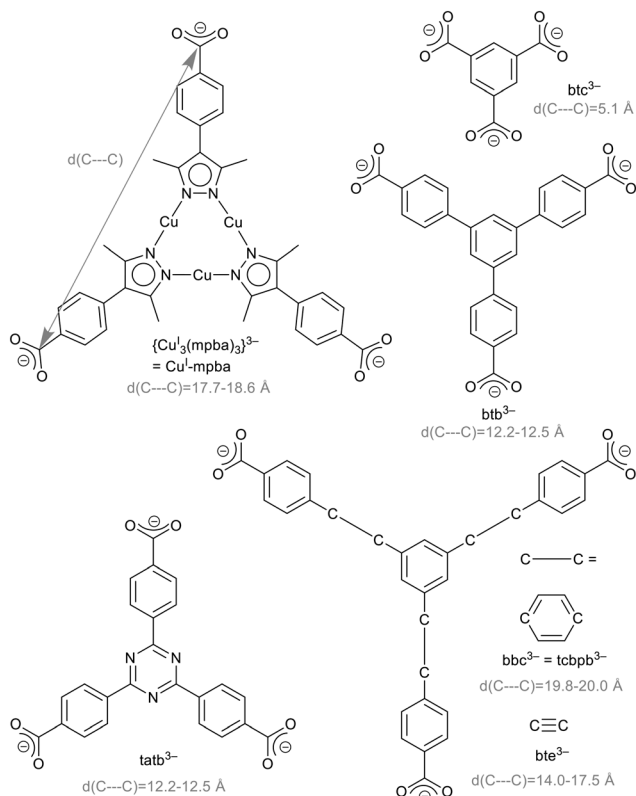
The Zn-MOFs with the trigonal linkers Zn-btc,^{9–12} MOF-177 (btb^{3-}),^{13,14} MOF-180 (btc^{3-})¹³ and MOF-200 (bbc^{3-})¹³ are isorecticular to each other, and so is DUT-40 (bbc^{3-}) to DUT-41 (bbc^{3-}).¹⁵

The limits of isorecticular expansion as well as the influence of the ligand's nature in this regard is an interesting conceptual question. Our interest is focused on the triangular tri-copper metallo-linker, $\{\text{Cu}_3(\text{mpba})_3\}^{3-}$ (in short Cu^{I} -mpba), which can be seen as belonging to the btc-type linker family (Scheme 1). Practically the only MOF-type open framework compound featuring the Cu^{I} -mpba unit is PCN-91, reported by Zhou and co-workers, which is a Cu^{II}

coordination polymer with a Pt_3O_4 underlying net.¹⁶ Cu^{I} -mpba is constructed from three Cu^{I} atoms, which linearly coordinate pyrazolate nitrogen donor atoms of three deprotonated bifunctional 4-(3,5-dimethyl-1H-pyrazol-4-yl)benzoic acid (H_2mpba) ligands. In comparison to other trigonal linkers, the Cu^{I} -mpba metallo-linker is quite large with an edge length $d(\text{C}—\text{C})$ of 17.7–18.6 Å (Scheme 1). Its size is comparable to one of the largest typically used trigonal linkers bbc^{3-} (= tcbbp^{3-}) ($d(\text{C}—\text{C})$ = 19.8–20.0 Å, Scheme 1). Furthermore, Cu^{I} -mpba is not rigid and planar but somewhat flexible and distorted (*vide infra*), similar to bbc^{3-} and bte^{3-} .^{13,30} The arms of the latter two ligands deflect from the plane of the central benzene ring by up to 10° (Fig. S44, ESI†). If the benzoate groups bend all out of the central ring plane to one side, a bowl-shaped linker is formed.^{15,30} The linker bte^{3-} appears to be the most flexible example as the ethynyl-benzoate groups do not only bend out of the central ring plane but also deviate from a 120° angle between them (Fig. S44†).³⁰ The flexibility of the Cu^{I} -mpba ligand stems from the flexibility of the coordination-bonded core, with pyrazolate-phenyl non-planarity due to steric repulsion between the methyl groups of pyrazolate and the phenyl *ortho*-hydrogen atoms, which also endows the ligand branches with a limited rotational degree of freedom.

The trinuclear Cu^{I} -pyrazolate arrangement is well known in $\{\text{Cu}_3(\text{pz})_3\}$ clusters (pz = 3,5-R-pyrazolate, where R is a group with intermediary steric volume, typically Me, sometimes CF_3 or Ph, and only rarely H. In the latter case the $\{\text{Cu}_2(\text{pz})_2\}$ cyclic motif is rather realized). The more widespread, structurally close form of that cluster is $\{\text{Cu}_3(\mu_3\text{-OH})(\text{pz})_3\}^{n-}$, based on $\text{Cu}(\text{I/II})$ or $\text{Cu}(\text{II})$ only (importantly, the pz = 3,5-*H*-pyrazolate ligand is very typical for the $\mu_3\text{-OH}$ -centered clusters, unlike for the $\text{Cu}(\text{I})$ -only cluster).³⁷ As special feature of copper(I), the $\{\text{Cu}_3(\text{pz})_3\}$ clusters are very prone to associate *via* intermolecular cuprophilic interactions, which could affect the course of self-assembly of potential MOFs.^{38,39} There is a fortunate opportunity for the synthesis of potential mixed-metal $\text{M}^{\text{II-IV}}/\text{Cu}^{\text{I}}$ pyrazolate-carboxylate coordination polymers starting from Cu^{II} as a precursor in *N,N*-dimethylformamide (DMF) and similar amide solvents. The latter are favorable for the crystallization of both carboxylate and pyrazolate coordination polymers, while elevated temperatures (80–120 °C) ensure partial or complete reduction of Cu^{II} to Cu^{I} .⁴⁰

The doubly deprotonated bifunctional pyrazolate-carboxylate ligand mpba²⁻, constituting $\{\text{Cu}_3(\text{mpba})_3\}^{3-}$, has already been used in the synthesis of a few reported coordination-bonded frameworks with other coordination bonded clusters, namely Zn^{II} ,^{41–43} Co^{II} ⁴³ or Cu^{II} .^{16,44} It is also worth mentioning the reported, mostly 2D, structures of the partially deprotonated pyrazole-carboxylate ligand Hmpba^- with $\text{Co}(\text{II})$, $\text{Ni}(\text{II})$, $\text{Zn}(\text{II})$, $\text{Cu}(\text{II})$, $\text{Cd}(\text{II})$ ⁴⁵ and the mixed-metal 1D bimetallic coordination polymer $\{[(\text{Ti}_3\text{O})_3(\text{iPro})_8][\text{Hmpba}]_2[\text{Cu}_2\text{I}_2]_{0.5}\}$ (with iPro = isopropoxide).⁴⁶ In the case of the mixed-valent $\text{Cu}^{\text{I/II}}$ framework compound of PCN-91 (ref. 16) and the 1D $\text{Ti}^{\text{IV}}-\text{Cu}^{\text{I}}$ coordination polymer, the harder carboxylate



Scheme 1 Representation of extended versions of the benzene-1,3,5-tricarboxylate, (btc^{3-}) linker: $\{\text{Cu}_3(\text{mpba})_3\}^{3-}$, 4,4',4''-benzene-1,3,5-triyl-tribenzoate (btb^{3-}), 4,4',4''-(benzene-1,3,5-triyl-tris(benzene-4,1-diyl)tribenzoate (bbc^{3-}) = 1,3,5-tris(4'-carboxy[1,1'-biphenyl]-4-yl)benzene (tcbbp^{3-}), 4,4',4''-(benzene-1,3,5-triyl-tris(ethyne-2,1-diyl)tribenzoate (bte^{3-}) and 4,4',4''-s-triazine-2,4,6-triyl-tribenzoate (tatb^{3-}). The separation between the carbon atoms of the carboxyl groups is given as a size indicator in gray.



Table 1 Extended versions of the benzene-1,3,5-tricarboxylate, (btc³⁻) linker and their MOF-examples

Ligand ^a	Example ^c
btc ³⁻	Cu-btc (HKUST-1), ⁸ Zn-btc, ^{9–12} Cd-btc ¹⁰
btb ³⁻	MOF-14(Cu), ¹⁷ MOF-143(Cu), ¹⁸ DUT-9(Ni), ¹⁹ MOF-177(Zn), ^{13,14,20} [Cd ₃ (btb) ₂ (bib)(DMF) ₃], ²¹ {[Ba ₃ (btb) ₂ (NMP) ₂ (H ₂ O) ₂]-2NMP·H ₂ O}, ²² {[Co(Hbtb)(pdp)]·H ₂ O}, ²³ [Cd ₂ (tbib)(btb)(H ₂ O) ₂] _n X, ²⁴ {[Eu(btb)(H ₂ O) ₃]-1.5DMA·5H ₂ O} _n (MHT-1), ²⁵ MUV-12, ²⁶ FJU-120 (ref. 27)
bbc ³⁻ = tcbpb ³⁻	{[Zn ₄ (OH) ₂ (H ₂ O) ₂ (py) ₂ (tcbpb) ₂]-DMF·py·3H ₂ O}, ²⁸ {[Zn ₈ (OH) ₄ (tcbpb) ₄]-2DMF·2EtOH·3H ₂ O}, ²⁸ Al-tcbpb, ²⁹ MOF-399(Cu), ¹⁸ DUT-40(Zn), ¹⁵ DUT-41(Zn), ¹⁵ MOF-200(Zn), ²⁰ MOF-1005(Zr) ³⁰
bte ³⁻	MOF-180(Zn), ²⁰ MOF-1004(Zr) ³⁰
tatb ³⁻	[Ln ₂ (tatb) ₂ (DMSO) ₆] (Ln = La, Ce, Pr, MOF-388Sm, Eu), ³¹ {[Ln ₂ (tatb) ₂ (DMSO) ₅ (CH ₃ OH)]·DMF·DMSO·2H ₂ O} (Ln = Nd, Gd, Tb), ³¹ [Cu ₂ (tatb) ₂ (H ₂ O) ₃], ³² PCN-9(Co), ³³ MIL-142(Sc)-tatb, ³⁴ [Co ₄ (SO ₄) ₃ (F) ₃ (tpt) ₂ (tatb)], ³⁵ [Fe(tatb)(<i>p/m</i> -bib)] ³⁶
tapb ^{3-b}	MOF-388(Cu) ¹⁸

^a See Scheme 1. ^b Not shown in Scheme 1, but tapb is like bbc with a triazine instead of a benzene core. ^c Other abbreviations used: (bib = 1,4-bis(2-methylimidazol-1-yl)butane), NMP = *N*-methyl pyrrolidone, pdp = 2-(4,6-di(pyridine-2-yl)pyridine-2-yl)pyridine, tbib = 1,3,5-tri(1*H*-benzo[*d*]-imidazol-1-yl)benzene, tpt = 2,4,6-tris(4-pyridyl)-1,3,5-triazine, *p/m*-bib = *para/meta*-bis((1*H*-imidazole-1-yl)methyl)benzene.

group combines with the HSAB-harder metal ions Cu^{II} and Ti^{IV} and the HSAB- softer pyrazolate or pyrazole donor is coordinated by Cu^I (according to the HSAB principle).⁴⁷ The hard/soft selectivity of the O,N heteroditopic mbpa²⁻ ligand offers a potentially productive approach towards mixed metal coordination polymers, based on the *in situ* synthesized {Cu₃(mpba)₃}³⁻ cluster.

Carboxylate coordination polymers also featuring the {Cu₃(pz)₃} cluster are rare. According to the CSD search, there are five compounds of this type, with three important MOF representatives, namely the mentioned PCN-91,¹⁶ FDM-4 (ref. 48) and FMD-8.⁴⁹ The two other compounds are the CFA-13 MOF⁵⁰ with an extended {Cu₃(pz)₃} cluster in the form of Cu₃(3,5-CF₃-pyrazolate)₄, and the [Cu^{II}{Cu₃(Hmpba)(mpba)₂}] 1D coordination polymer⁵¹ reported by us. The FDM-4,5,6,7,8 series of Zn/Cu mixed-metal coordination polymers with surface areas up to 3728 m² g⁻¹, based on the fully deprotonated short 4-carboxypyrazole ligand as a sole organic ligand, or together with the 1,4-benzenedicarboxylate or 2,6-naphthalenedicarboxylate ligand, is particularly interesting. The series features reversible redox transformation between the {Cu₃(pz)₃} and {Cu₃^{II}(OH)(pz)₃} cluster without compromising the integrity of the structure (as some of the structures are reported only for the oxidized cluster, the CSD search does not cover the Cu^I-only variants in all cases). As mentioned above, 3,5-*H*-pyrazoles frequently constitute the {Cu₃(OH)(pz)₃} clusters, but they are less frequent in geometrically well-formed {Cu₃(pz)₃} clusters. The FDM-4-8 series demonstrate that in some cases both possibilities could be realized, resulting in highly porous robust structures, which substantiate the interest in mixed-metal and mixed-ligand complexes for longer ligands like mpba²⁻, whose potential is proven by the known PCN-91 material.

The Cu^I-mpba metallo-linker has not been used in the construction of isorecticular Zn-MOF structures (as well as any other bimetallic MOFs) and in the current contribution we explore this possibility. The bifunctional ligand precursor 4-(3,5-dimethyl-1*H*-pyrazol-4-yl)benzoic acid (H₂mpba) was reacted with copper(II) nitrate in DMF in the presence of

zinc(II) salts and, if necessary, a modulator (HBF₄) aiming for Zn frameworks with the metallo-linker, akin to those with the btc, btb, bte or bbc linker (*cf.* Table 1). The importance of the HSAB selectivity regarding the mixed valent Cu^I/Cu^{II} metals and the N,O-heteroditopic ligand is explicitly discussed for PCN-91.¹⁶ The use of Cu^I/Zn^{II} in this work could be viewed as a continuation of that principle aiming for the utilization of zinc (oxo)carboxylate species as secondary building units.

Results and discussion

The solvothermal reaction between copper(II) nitrate, zinc(II) nitrate and H₂mba in *N,N*-dimethylformamide (DMF) at 80–90 °C for several days resulted in the formation of the bimetallic networks [Me₂NH₂][Zn₄{Cu₃(mpba)₃}(Me₂NH)(DMF)₂] (**1**), [Zn₆{Cu₃(mpba)₃}(DMF)₅] (**2**) and [Zn₇{Cu₃(mpba)₃}(O)(H₂O)₅(DMF)₃] (**3**). The synthesis conditions had to be carefully adjusted with regard to time, temperature, Cu : Zn stoichiometry and the amount of HBF₄ modulator, as otherwise mixtures of **1**, **2** and **3** also form (Table S1, Scheme S1, ESI†).

MOF examples in the literature with Cu(I)₃(pyrazolate-carboxylate) motifs include PCN-91,¹⁶ FDM-4-8,^{48,49} CFA-13 (ref. 50) and [Cu^{II}{Cu₃(Hmpba)(mpba)₂}]⁵¹. The syntheses were all performed using Cu(II) as a precursor and in *N,N*-dimethylformamide (DMF) or similar amide solvents. DMF is favorable for the crystallization of both carboxylate and pyrazolate coordination polymers, while elevated temperatures (80–120 °C) in amide solvents ensure partial or complete reduction of Cu(II) to Cu(I).⁵²

The {Cu₃(pz)₃} SBU of PCN-91 forms as a result of the *in situ* reduction of Cu^{II} (Cu(NO₃)₂·2.5H₂O precursor) in DMA and two drops of pyridine at elevated temperature (120 °C) and prolonged time (24 h). In the work of Tu and co-workers, the short 4-pyrazolecarboxylate linker was used to synthesize bimetallic, redox active materials. By varying the Cu : Zn ratio and the solvent mixture (ratio of DMF and NMP or DMF and DMA (only used for FDM-5)) the materials FDM-4 and FDM-5 were synthesized under solvothermal conditions. By adding the organic ligand 1,4-benzene dicarboxylic acid the material

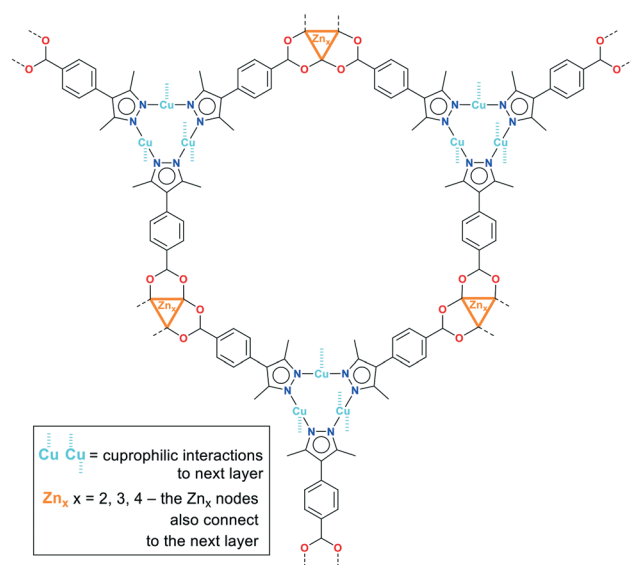


FDM-6 was produced. FDM-8 and the substituted derivatives NH_2 -FDM-8 and C_4H_4 -FDM-8 were synthesized under solvothermal conditions by adding 1,4-benzenedicarboxylate acid, 2,6-naphthalenedicarboxylate acid (for FDM-8) and 2-aminoterephthalic acid or 1,4-naphthalenedicarboxylate acid for NH_2 -FDM-8 and C_4H_4 -FDM-8. In all FDM MOFs, the $\{\text{Cu}_3(\text{pz})_3\}$ moiety was obtained by *in situ* reduction from the Cu(II)-salt ($\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$) in DMF/NMP or DMF/DMA at elevated temperature (85 °C for FDM-4 and FDM-5, 100 °C for FDM-6, FDM-8 and NH_2 -FDM-8 and C_4H_4 -FDM-8) and prolonged time (72 h for FDM-4 and FDM-5 and 16 h for FDM-6, FDM-8 and NH_2 -FDM-8).

Regarding the question of pK_a , the carboxylic acid and pyrazole groups feature very different acidities (~ 5 vs. 14), but this does not preclude the concomitant formation of complexes even for Zn(II). The HSAB hard-hard soft-soft matching between Cu^{I} and pyrazolate as well as Zn^{II} and carboxylate makes the formation of the respective complexes highly predictable.

Compounds **1** and **3** could be obtained reproducibly as yellowish clear crystals (Fig. S1 and S2, ESI †) in phase pure form. Compound **2** was difficult to obtain in pure phase and in single crystal form, and it was only observed as an admixture of **1** and **3**. The identity of the three compounds could be elucidated by single-crystal diffraction (see below). The experimental and simulated powder X-ray diffractograms were matched and the correspondence was checked by Le Bail fitting (Fig. S3, ESI † compound **2**, which could be obtained only in the form of a powder, is the least pure).

A common feature of the structures of **1–3** (Scheme 2) is the structural changeability of the Zn-SBU clusters, featuring



Scheme 2 Common features of the structures of **1–3** with Zn_x nodes of different nuclearities ($x = 2–4$) and the trigonal $\{\text{Cu}_3(\text{mpba})_3\}^{3-}$ nodes giving a hexagonal-ring layer. Both the Zn_x nodes and the cuprophilic interactions from $\{\text{Cu}_3(\text{mpba})_3\}^{3-}$ connect to the next layer.

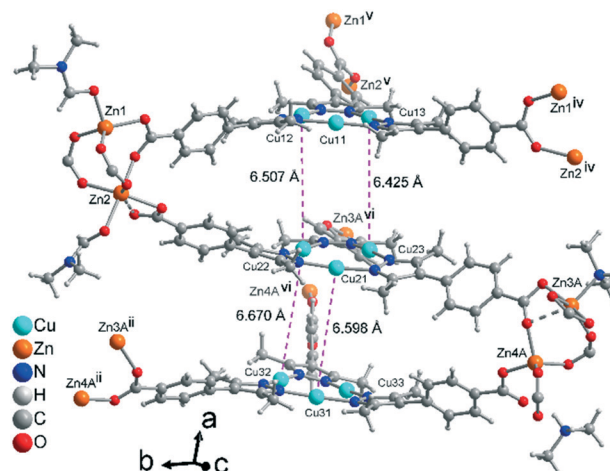


Fig. 1 Extended asymmetric unit of **1** with the two located coordinating DMF solvent molecules, one dimethylamine solvent molecule and one dimethylammonium counter-cation (for further details on these amine/ammonium groups see Fig. S4, ESI †). Each $\{\text{Cu}_3(\text{mpba})_3\}^{3-}$ unit belongs to a different layer. In view of the structure complexity, a ball-and-stick presentation is used and only the major contribution to the disorder of Zn3 and Zn4 is shown. The Zn3/4 disorder is described in more detail in the ESI † . The distances between the Cu atoms (pink dashed lines) correspond to twice the distances of typical cuprophilic $\text{Cu}\cdots\text{Cu}$ contacts, as the interlayer space is filled by a symmetry equivalent Cu^{I} -mpba unit (the stack is shown in Fig. 3 and 4). Symmetry codes: (ii) $x, y + 1, z$; (iv) $x, y - 1, z$; (v) $x, y - 1, z - 1$; (vi) $x, y, z - 1$.

nuclearities of 2–4 (distorted tetrahedral and pentagonal as well as trigonal-prismatic SBU-shapes) and frequent disorder. Such unconventional changeability contrasts the situation in PCN-91 with Cu(II), where the conventional $\{\text{Cu}_2(\text{O}_2\text{C})_4\}$ -paddle-wheel motif is formed.¹⁶ Neither of **1–3** is isostructural to PCN-91, even if a paddle-wheel $\{\text{Zn}_2(\text{O}_2\text{C})_4\}$ -SBU is possible.⁵³ Instead, the structures of **1–3** have a common structure-guiding motif of columnar stacks, composed of Cu^{I} -mpba associated by cuprophilic interactions. The Zn-clusters appear to be flexible and adapt to the requirements imposed by the Cu^{I} -mpba stacks, which results in similarities in structure-organization dominated by the formation of hexagonal coordination-bonded rings within honeycomb **hcb** layers (Scheme 2; the summary of the work on the topological level is given below in Fig. 9, which follows the detailed description of the structures).

Crystal structure of **1**

The formula of compound **1** was established as $[\text{Me}_2\text{NH}_2][\text{Zn}_4\{\text{Cu}_3(\text{mpba})_3\}_3(\text{Me}_2\text{NH})(\text{DMF})_2]$. The formation of dimethylammonium $[\text{Me}_2\text{NH}_2]^+$ from DMF and its presence as a counter-cation has been seen in zinc-MOFs with benzene-1,4-dicarboxylate,⁵⁴ furan-2,5-dicarboxylate⁵⁵ or 1,1'-biphenyl-3,3',4,4'-tetracarboxylate.⁵⁶ The asymmetric unit in **1** consists of three symmetry-independent $\{\text{Cu}_3(\text{mpba})_3\}^{3-}$ units, each belonging to a different layer, which are associated *via* coordination with Zn-atoms (Fig. 1). The three



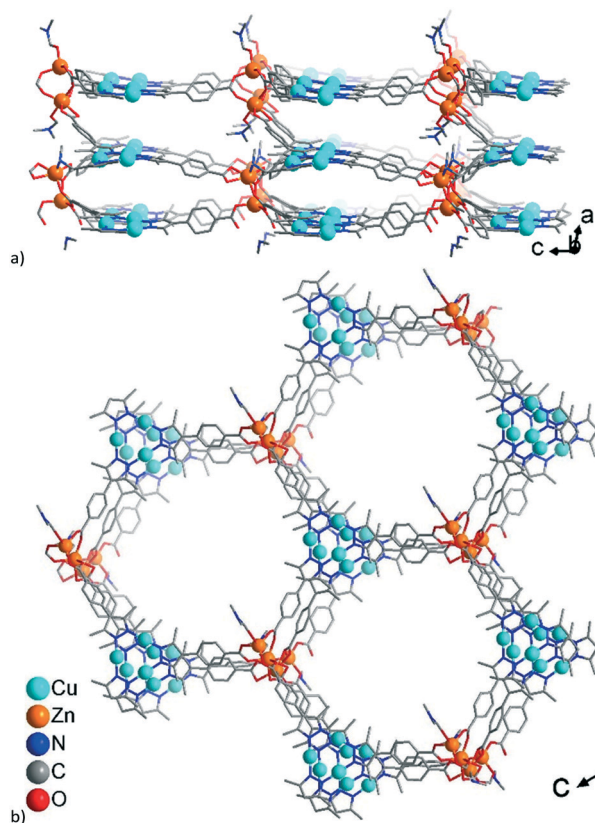


Fig. 2 Wire-frame model of (a) side-view and (b) on-top view (along the *b* and *a* axis, respectively) of a section of a single 3-layer stack in **1**. Only the major contribution to the disorder of Zn3 and Zn4 is depicted. See also the space-filling presentation in Fig. S5.†

layers give three-layer 2D sheets, referred to as ‘stacks’ (Fig. 2).

There are two dinuclear Zn-carboxylate clusters, which could be regarded as disrupted paddle-wheel arrangements. In the first cluster, the Zn1 atom has a tetrahedral $\{ZnO_{4+1}\}$ environment with an additional weaker Zn–O bond, while Zn2 has an octahedral environment, involving a chelating carboxylate, thereby not completing the usual paddle-wheel arrangement (both atoms coordinate one DMF molecule). In the second cluster, the Zn3 and Zn4 atoms are disordered over two different positions, A (81%) and B (19%), assuming alternating roles (Fig. S4, ESI† for the details of the disorder). In part A, the Zn3A atom coordinates a nitrogen atom from a dimethylamine solvent molecule in addition to five carboxylate oxygen atoms, two of which belong to a chelating carboxylate. This results in a strongly distorted octahedron as a coordination environment. The Zn4A atom is tetrahedrally coordinated by four oxygen atoms, two of which belong to monodentate carboxylates. In the absence of the second disordered component (Zn4B), the latter carboxylates are involved in hydrogen bonding with the charge compensating dimethylammonium cation, ensuring the coordination of five carboxylates per Zn_2 -cluster, instead of four.

The connection of the $\{Cu_3(mpba)_3\}^{3-}$ metallo-linkers with the Zn_2 -SBUs leads to hexagonal rings with the Cu_3 -centroid

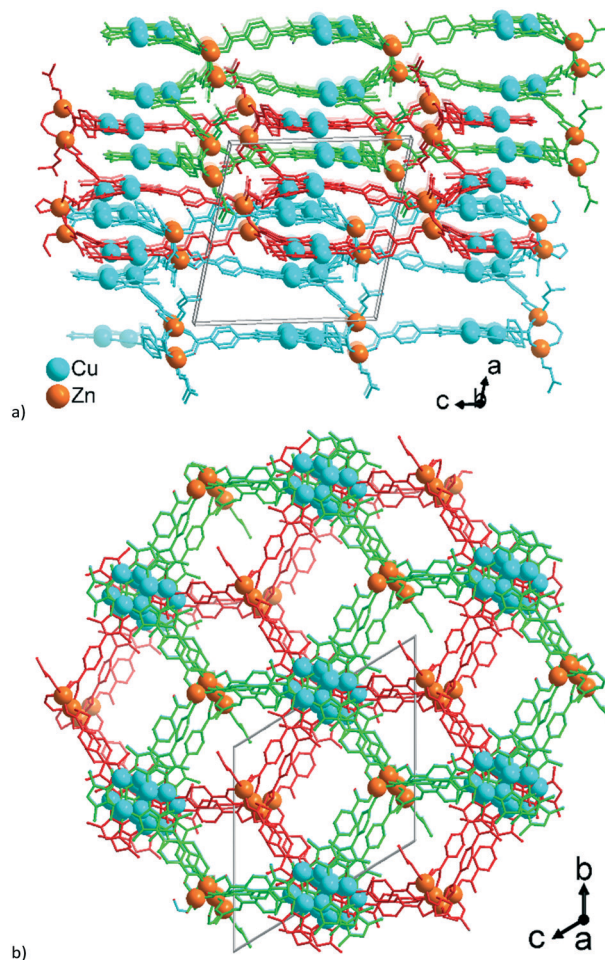


Fig. 3 Wire-frame model of (a) side-view and (b) on-top view (along the *b* and *a* axis, respectively) of a section of three polycatenated 3-layer stacks in **1**. The three stacks are differentiated by their green, red and cyan color. Hydrogen atoms are not shown for clarity. Only the major contribution to the disorder of Zn3 and Zn4 was used in the drawing. See also the space-filling presentation in Fig. S6, ESI.†

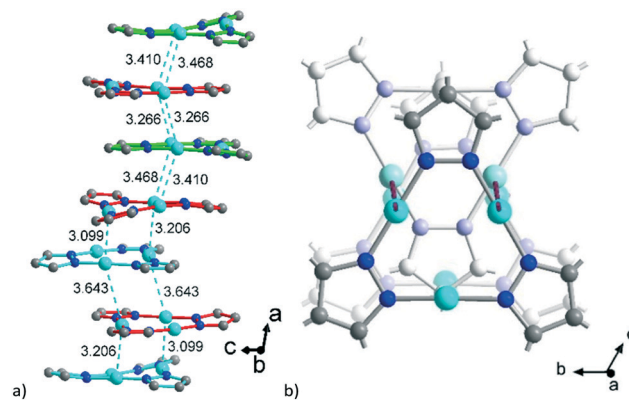


Fig. 4 a) Stacking of the $\{Cu_3(pz)_3\}$ moieties in **1** with $Cu^1 \cdots Cu^1$ contacts (in cyan, in Å) between the polycatenated stacks which are differentiated by the same green, red and cyan color as in Fig. 3; the methyl groups are not shown for clarity. b) Alternation of the axial rotation of the $\{Cu_3(pz)_3\}$ moieties.



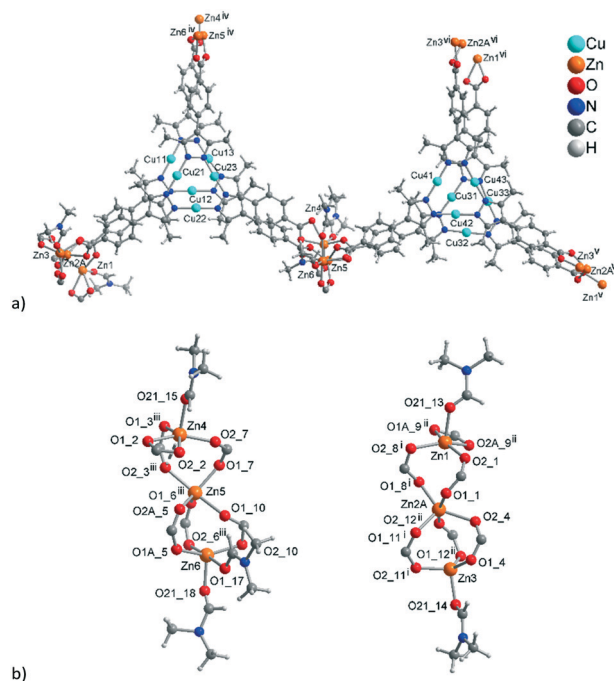


Fig. 5 (a) Extended asymmetric unit of **2**. The emphasis in this image is placed on the slightly shifted eclipsed position of the metallo-linkers. The inter-linker Cu...Cu contacts range from 3.57–3.79 Å (Fig. 6c). (b) The two trinuclear hourglass-like SBUs. Symmetry codes: (i) $x - 1, y - 1, z + 1$; (ii) $x - 1, y, z + 1$; (iii) $x, y + 1, z$; (iv) $x, y - 1, z$; (v) $x + 1, y + 1, z - 1$; (vi) $x + 1, y, z - 1$. Only the major contribution to the disordered carboxylates and Zn2 atoms is shown, and the Zn2 disorder is described in more details in Fig. S9, ESI.†

and Zn₂-centroid as vertices and the mpba-ligands as edges (Fig. 2b). While $\{\text{Cu}_3^{\text{I}}(\text{mpba})_3\}^{3-}$ is a three-connected node, the Zn1/Zn2 group acts as a four-connected node and the Zn3/Zn4 group as a five-connected node. Hence, each Zn1/Zn2 group belongs to the top and middle layer, and each Zn3/Zn4 group to the middle and bottom layer (according to the orientation in Fig. 1 and 2a).

The single 3-layer stack in **1** has a >6.4 Å separation between the metallo-linkers (Fig. 1, 2a, and S5a†) and possesses an open structure with hexagonal pores propagating along the *a*-direction (Fig. 2b and S5b†). A dense packing, however, minimizes the energy of a structure. Therefore, open frameworks often show phenomena of interpenetration, polycatenation or entanglement.^{57–62} In **1** the lower/upper half of a 3-layer stack polycatenates into the upper/lower halves of its neighbors (Fig. 3a and S6a†). At the same time the stacks are packed along the *a*-direction in a staggered fashion in an ABAB sequence, such that the $\{\text{Cu}_3(\text{pyrazolate})_3\}$, *i.e.*, $\{\text{Cu}_3(\text{pz})_3\}$, parts of the metallo-linkers are stacked on top of each other (Fig. 3b and 4) and the Zn-SBUs fill most of the voids of the polycatenated stack (Fig. 3b and S6b, ESI†). Thereby, the available space in a single 3-layer stack is filled (compare Fig. 2b and 3b).

Two adjacent 3-layer stacks polycatenate in a parallel manner in a 2D + 2D motif. The infinite 2D + 2D polycatenation of upper/lower parts of neighboring 3-layer

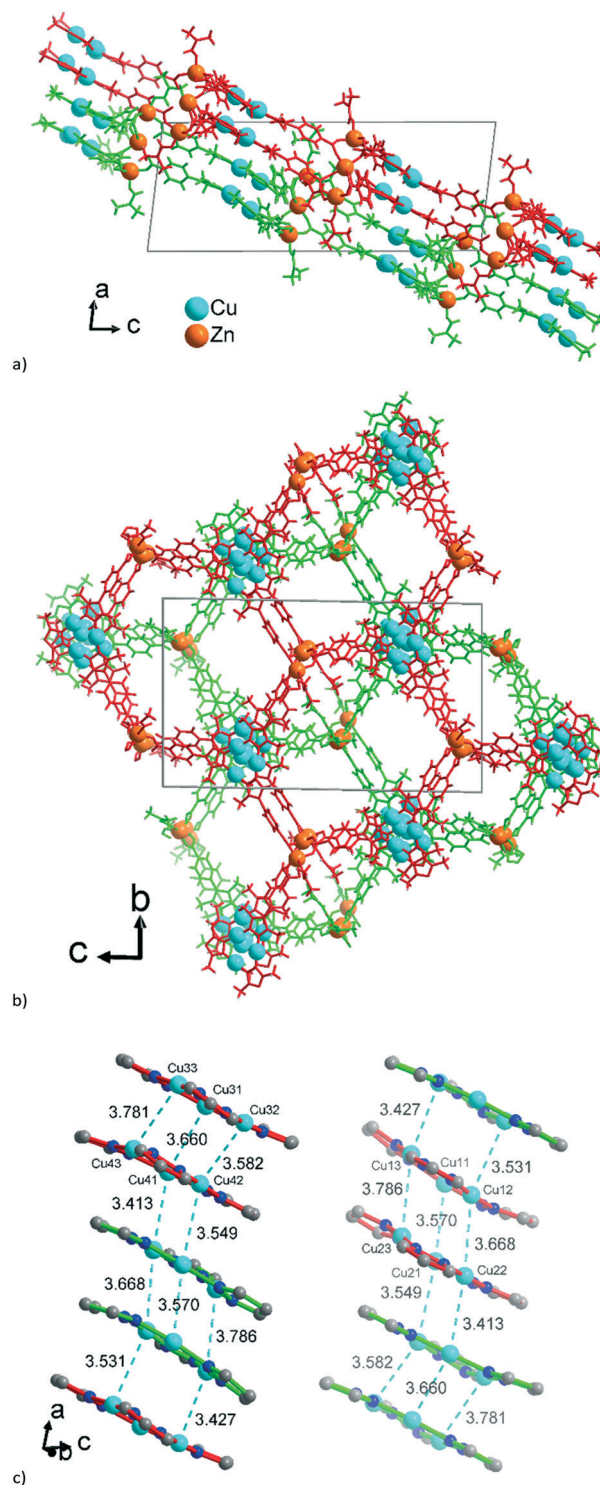


Fig. 6 Wire-frame model of (a) side-view and (b) on-top view (along the *b* and *a* axis, respectively) of two adjacent 2-layer stacks in **2**. The two stacks are differentiated by their green and red color. See also the space-filling presentation in Fig. S10, ESI.† (c) Stacking of the $\{\text{Cu}_3(\text{pz})_3\}$ moieties in **2** with Cu...Cu contacts (in cyan, distances in Å) between the layers. The inversion-symmetry relation of adjacent stacks renders the packing for the two symmetry-independent parts of the asymmetric unit of **2** (*cf.* Fig. 5a) equivalent. Thus, the red $\{\text{Cu}_3(\text{pz})_3\}$ units in the left column correspond to the green $\{\text{Cu}_3(\text{pz})_3\}$ units in the right column in Fig. 6c.



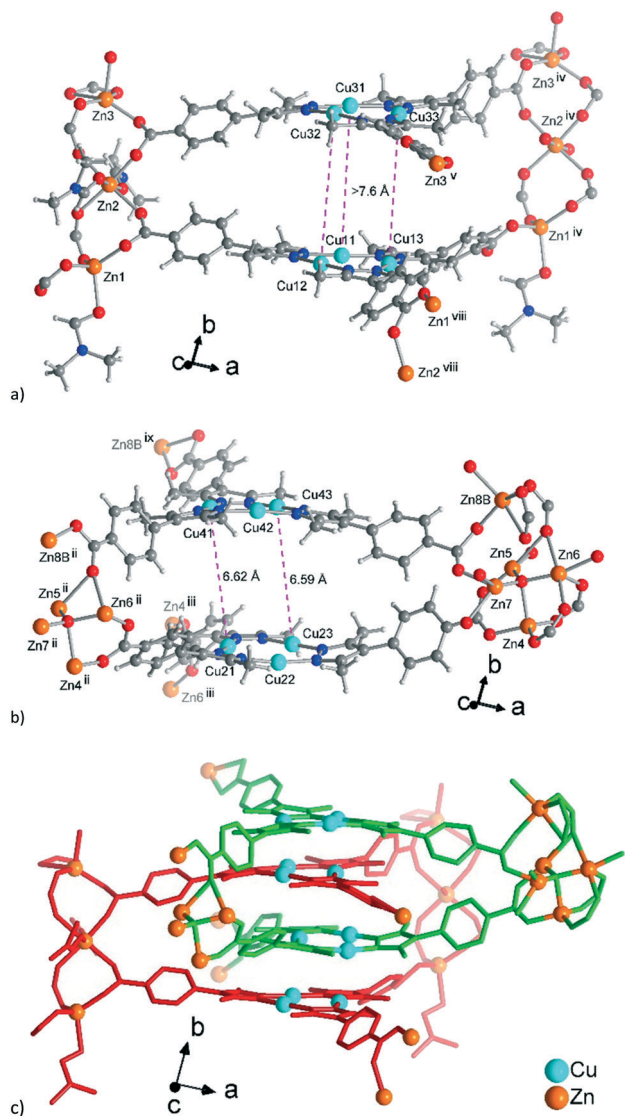


Fig. 7 Extended asymmetric units in **3**. (a) Network A, constituted by the trinuclear Zn-SBUs with branches ending with Zn atoms showing the connections. (b) Similar image for network B constituted by the tetranuclear Zn-SBUs. The Zn atoms in the middle of the images which point to the front (in a) and to the rear (in b) propagate the structure along the *b*-axis. The distances between the Cu atoms (pink dashed lines) correspond to twice the distances of typical cuprophilic Cu \cdots Cu contacts. (c) The mode of interpenetration of network A, shown with red bonds and network B, shown with green bonds. In (b) the major disorder contributions to the Zn-atoms in the structure are shown. From the disorder, the Zn-SBU in (b) looks like a pentanuclear interpretation, yet the 80% occupancy gives a Zn₄-SBU (see Fig. S11, ESI† for details). Symmetry codes: (ii) $x - 1/2, y - 1/2, z$; (iii) $x - 1/2, -y - 3/2, z - 1/2$; (iv) $x + 1/2, y + 1/2, z$; (v) $x + 1/2, -y - 3/2, z + 1/2$; (viii) $x + 1/2, -y - 5/2, z + 1/2$; (ix) $x - 1/2, -y - 1/2, z - 1/2$.

stacks along the perpendicular direction to the stack leads to a 3D network. Interpenetration which results in a higher dimensionality than the discrete network, that is, 2D \rightarrow 3D, is termed polycatenation.⁶⁰ A similar upper/lower part 2D + 2D polycatenation, giving an overall 3D network, is also frequently seen for (4,4)la double-layer **sql** nets.⁶¹

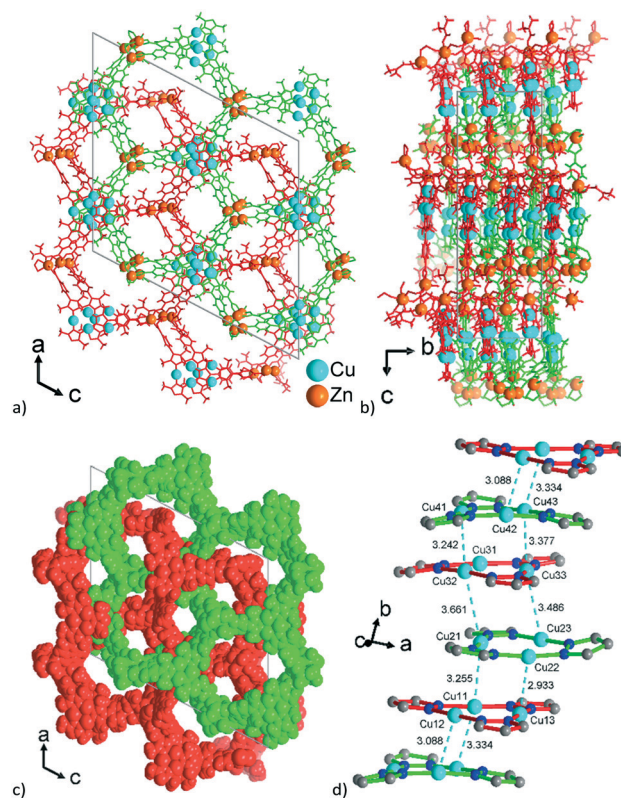


Fig. 8 View along the hexagonal channels of the two interpenetrating 3D-frameworks in **3** as (a and b) wire-frame, and (c) space-filling modes. Network A in red with the Zn₃-SBU and network B in green with the Zn₄-SBU (cf. Fig. 7). (d) Stacking of the {Cu₃(pz)₃} moieties in **3** with Cu¹ \cdots Cu¹ contacts (in cyan, distances in Å) between the layers of the interpenetrating frameworks.

The polycatenation is invoked and stabilized through the inter-stack Cu¹ \cdots Cu¹ contacts, with lengths between 3.099 Å and 3.643 Å (Fig. 4) which thus correspond to typical cuprophilic interactions.^{43,63–65} We note that these inter-stack Cu¹ \cdots Cu¹ separations are in the same range as the Cu¹ \cdots Cu¹ distances within the {Cu₃(pz)₃} clusters, which lie between 3.169 Å and 3.257 Å (Fig. S7, ESI†).

Crystal structure of **2**

The asymmetric unit of **2** with the established formula of [Zn₆{Cu₃(mpba)₃}₄(DMF)₅] (Fig. 5a) consists of four symmetry-independent {Cu₃(mpba)₃}³⁻ metallo-linkers. They are pairwise stacked at short Cu¹ \cdots Cu¹ distances of 3.413 Å and 3.786 Å and connected through two symmetry-independent {Zn₃(O₂C-)₆(DMF)₂₋₃} SBUs (Fig. 5b) to a two-dimensional 2-layer stack. The three Zn atoms in {Zn₃(O₂C-)₆(DMF)₂₋₃} form unsymmetrical hourglass-like trinuclear SBUs where the central Zn atoms (Zn2A, Zn5) rather have a square-pyramidal than bipyramidal environment (geometric index⁶⁶ $\tau = 0.42$ and 0.32) coordinated by five carboxyl-O atoms from five different metallo-linkers (Fig. 5b).

The terminal Zn atoms possess either a tetrahedral (Zn3) or distorted square-pyramidal environment (Zn1, Zn4,



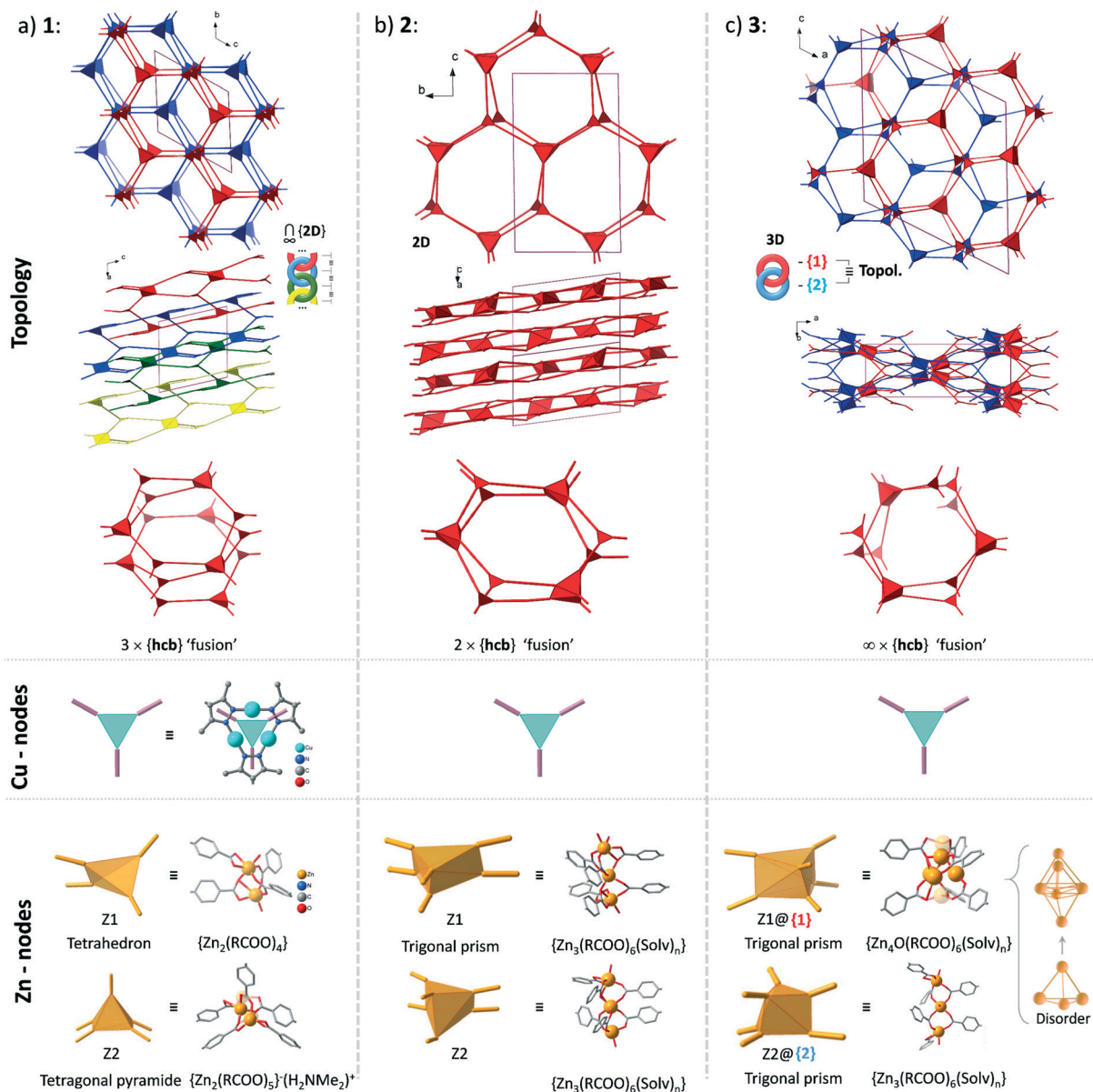


Fig. 9 Topological overview of the structures of 1–3 (the coloration of the separate networks is assigned for convenience: all structures contain only one type of network regarding their topology). The crystallographically distinct Cu-based nodes are structurally- and in most cases topologically equivalent (the exception is the structure of 1, where there are two topologically distinct Cu-nodes among three crystallographically different ones with point symbols of $\{4\cdot6^2\}$ and $\{6^3\}$; see Table S4, ESI†). The Zn-based nodes are distinct in every case, except for the topological equivalence of the two different nodes in the structure of 3. For an illustration of the interpenetration of hexagonal rings in the structures of 1 and 3 see Fig. S15, ESI†.

Zn6 $\tau = 0.23, 0.08, 0.54$), although Zn4 and Zn6 have additional long (>2.5 Å) contacts completing a distorted octahedral environment.⁶⁶ The two Zn_3 -SBUs possess three crystallographically different Zn^{II} ions. The coordination of the terminal Zn atoms Zn1, Zn3 and Zn4 is constructed from three carboxyl groups of metallo-linkers and one DMF molecule. For the five-coordinated atoms Zn1 and Zn4 one of the carboxyl groups is chelating and the DMF ligand is in the apical position. The five-coordinated Zn6 atom is bound to three monodentate carboxylates and two DMF ligands.

The 2D 2-layer stacks in 2 run parallel to the $(1\ 0\ 1)$ plane (Fig. 6a). The combination of the $\{Cu_3^I(mpba)_3\}^{3-}$ metallo-linkers with the Zn_3 -SBUs leads again to a structure based on hexagonal rings, when the Cu_3 -centroids and the midpoints between two Zn atoms in the hourglass SBUs are considered as nodes of the underlying net and the mpba ligands as edges (Fig. 6b). Two adjacent 2-layer stacks, related through a symmetry operation, become staggered along the a -direction, such that the $Cu_3(pyrazolate)_3$, *i.e.*, the $Cu_3(pz)_3$, parts of the metallo-linkers are again stacked on top of each other (Fig. 6c), as was also seen in 1 (*cf.* Fig. 3).



The terminal DMF ligands at the Zn₃ hourglass-SBUs stick out of the stack and protrude into the adjacent stack (Fig. 6a).

Crystal structure of 3

The established formula of **3**, [Zn₃{Cu₃(mpba)₃}₂(DMF)₃(H₂O)]₂[Zn₄(μ₄-O){Cu₃(mpba)₃}₂(H₂O)₄], reflects the presence of two symmetry-independent two-fold interpenetrating neutral networks, based on different Zn-SBUs. Network A = [Zn₃{Cu₃(mpba)₃}₂(DMF)₃(H₂O)] has the trinuclear Zn-SBU {Zn₃(O₂C-)₆(DMF)₃} (Fig. 7a). Network B = [Zn₄(μ₄-O){Cu₃(mpba)₃}₂(H₂O)₄] has the tetranuclear {Zn₄(μ₄-O)(O₂C-)₆(H₂O)₄} Zn-SBU (Fig. 7b). While the trinuclear SBU could be well-refined, the interpretation of the second Zn-SBU is ambiguous. The conventional {Zn₄(μ₄-O)(O₂C-)₆} cluster is octahedral, while the demands imposed by the crystal structure of **3** prescribe a distorted trigonal-prismatic shape. The refined atoms, constituting the cluster, have large thermal displacement factors, allowing multiple interpretations. Based on the Zn : Cu ratio found by analytics and following general chemical logic, a complex, unresolved overlap of nearly regular {Zn₄(μ₄-O)(O₂C-)₆} clusters was assumed (Fig. S12, ESI†). An alternative, more straightforward interpretation is a pentanuclear {Zn₅(μ₄-O)(OH)₂(O₂C-)₆} cluster with a minor disorder. Also, an admixture of Cu(I) such that the disordered {Zn₄(μ₄-O)} cluster might be {Zn₄Cu(μ₄-O)} could not be completely ruled out. Despite the different possible Zn₄-SBU interpretations, the localization of the {Cu₃(mpba)₃} moieties and hence the connectivity in the framework is unambiguous.

Aside from the two different Zn SBUs, the asymmetric unit of each network consists of two symmetry-independent {Cu₃(mpba)₃}³⁻ metallo-linkers, which are stacked in an eclipsed fashion at twice the distance, ~6.5 Å, of optimal cuprophilic interaction between the {Cu₃(pz)₃} moieties, leaving room for an interpenetrating network, as in **1**. The distance between the closest Cu^I atoms is over 6.5 Å (Fig. 7 and 8). Four of the six carboxylate groups of the two eclipsed {Cu₃(mpba)₃}³⁻ metallo-linkers are pairwise 'clamped' by two of the Zn₃ or Zn₄ SBUs, similar to the arrangement in the structure of **2**. The difference is that the remaining two carboxylate branches extend the network to the third dimension, perpendicular to the layers (along the *b* axis, Fig. 7). The connection of the {Cu₃(mpba)₃}³⁻ metallo-linkers with the Zn-SBUs in **3** leads again to a structure dominated by hexagonal rings in both of the interpenetrating frameworks (Fig. 8). The inter-linker cuprophilic contacts^{43,63-65} condition the tight interpenetration with Cu^I...Cu^I distances between 2.933 Å and 3.661 Å (Fig. 8c) as in **1** (Fig. 4) and **2** (Fig. 6c).

Comparison of the structural organization and topology

Structures **1-3** demonstrate a distinct similarity regarding both the formation of hexagonal ring motifs (Fig. 3b, 6b, and

Fig. 8a respectively; Fig. S14a, ESI†) and the columnar association of all present {Cu₃(pz)₃} moieties by Cu^I...Cu^I cuprophilic interactions (the high similarity of the columnar arrangements with staggered neighbor units is shown in Fig. S13†). The hexagonal rings are situated in an eclipsed way within a separate network/topology constituting open channels. The channel openings are partially occupied or blocked by polycatenated (**1**), just neighboring (**2**) or interpenetrated (**3**) networks which are arranged with respect to each other in a slipped or staggered mode similar to the ABAB layer sequence in graphite (Fig. 9).

The Zn-SBU nuclearity increases from Zn₂ in **1** over hourglass-like Zn₃ in **2** to Zn₃ and Zn₄ in **3** (Fig. 9). The connectivity ensured by the Zn-SBU adapts to the demands imposed by the stacked Cu^I-mpba linker arrangement, conditioned by the cuprophilic interactions (Fig. S13, ESI†). The flexibility in the Zn coordination sphere with tetrahedral, trigonal-bipyramidal to square-planar or octahedral coordinated Zn atoms must be seen as instrumental to allow the formation of (energetically) similar structures.

From the topological analysis on the level of the 3-c {Cu₃(pz)₃} node and of the Zn-SBU nodes of higher connectedness, the next point symbols were found: {4²·6²·8²}{6³}{6⁵·8} for **1** (a new 2D net), {4³·6¹²}{4³}₂ for **2** (3,6L66 net), and {4⁶·8²}{4²·6⁵·8⁵·10³} for **3** (**sqc-3,6-Fdd2-2** net) (the topological analysis was performed by ToposPro,^{67,68} see Fig. 9 for the node representations and Table S4, ESI† for node stoichiometry, vertex symbols and TD10).

The structures of **1-3** could be conceptualized through their relation with the hexagonal net, **hcb**. The latter is an expected outcome, when the trigonal {Cu₃(pz)₃} SBUs are combined in a coplanar fashion. The cuprophilic interactions, perpendicular to the nets, would favor a stacked arrangement (the few reported structures of this kind are based on short non-planar bifunctional 3,5-R₂-pyrazolates of copper(I),^{69,70} possessing 3D structures). The structures of **1-3** could be viewed as derivatives of the prototypal stacked **hcb** structure. Some of the alternating Cu-nodes are removed and the respective connectivity is added to the node in the layer above and/or below. The nodes with the increased connectivity are represented by a Zn-cluster and such redistribution of connectivity leads to 'fusing' of the separate **hcb**-nets. The Zn-clusters act as tetrahedral 4-c and tetragonal-pyramidal 5-c in **1** and trigonal-prismatic 6-c in **2** and **3** (Fig. 9). The Zn-clusters adopt different compositions and geometries, invariably ensuring the near coplanarity of the {Cu₃(pz)₃} SBUs within the columnar stacks, associated by cuprophilic interactions, that run uninterrupted through the whole structure (Fig. 4, 6c, 8c and S13, ESI†).

The simplest structural organization of 'fused' **hcb** nets is observed in **2** (Fig. 5, 6 and 9). The trigonal-prismatic Zn clusters correspond to two triangular nodes joined together in parallel. The resultant planar 2D net could be viewed as a "double-strand" ("dual-layer") **hcb** net with two stacked nets fused together at alternating nodes. The Cu-nodes are associated by cuprophilic interactions both within the



formed 2D network and between them forming a stack of networks, whose projection along the *a*-axis is nearly identical to a geometrically regular **hcb** net (Fig. 6 and 9). The structure features two crystallographically distinct, but chemically and topologically equivalent trinuclear Zn-clusters, $\{\text{Zn}_3(\text{COO})_6\}$. Three chelating carboxylates adjoin the neighboring Zn atoms disposed in row of three; two sets of such triples are in an approximately staggered disposition, ensuring the trigonal-prismatic shape of the node (Fig. 5b and 9). Similar structural organization is observed for a number of other tritopic carboxylates featuring an $\{\text{M}_3(\text{COO})_6\}$ SBU/cluster or functionally very similar unit ($\text{M} = \text{Zn}, \text{Co}, \text{Cd}$) (among 32 entries with the 3,6L66 net topology found in the Topcryst database,⁷¹ nearly all are coordination polymers and 23 follow this organization; see also the footnote to Table S4†). We regard the 2-layer stack in **2** as the kinetic product from which the 3-layer stack in **1** and the infinite-layer stack in **3** develop.

The structure of **1** consists of 2D nets, which could be represented as a ‘fusion’ of three stacked hexagonal nets (Fig. 2 and 9). Half of the alternating nodes in the top and bottom layers are represented by 4-c and 5-c Zn-nodes (with an excess connectivity over 3-c of one and two, respectively), which distribute the connectivity of the imaginary triangular node in the middle layer. Each pair of the neighboring triple-layer stacks is polycatenated in a parallel fashion, with the Cu-nodes belonging to the neighboring topologies interdigitating to form a tight stack (Fig. 3 and 4). The triple-layered topologies are polycatenated *via* pairwise interpenetration and topologically constitute a $2\text{D} + 2\text{D} \rightarrow 3\text{D}$ structure. The dinuclear Zn-nodes, $\{\text{Zn}_2(\text{RCOO})_4\}$ and $\{\text{Zn}_2(\text{RCOO})_5\}$ are close to a paddlewheel geometry with an added carboxylate in the second case, which is charge compensated by a nearby $[\text{Me}_2\text{NH}_2]^+$ cation.

The structure of **3** consists of two interpenetrated 3D nets, which are crystallographically and chemically distinct, but topologically equivalent (Fig. 7). Thereby, it represents formally a case of hetero-interpenetration, even if the topologies are very similar. The ‘fusion’ of the imaginary stacked **hcb** nets takes place not on the same level, but on alternating levels. Each of the two different six connected Zn-clusters acts as a trigonal node within its layer (Fig. 9) and the remaining excess connectivity ($2 + 1$) is redistributed above and below, respectively, the layer at the expense of the imaginary Cu-nodes of the prototypal stacked **hcb** net structure. It is interesting to note that each of the two distinct 6-c connected trigonal-prismatic Zn-nodes, $\{\text{Zn}_4\text{O}(\text{RCOO})_6\} = \text{Z1}$ and $\{\text{Zn}_3(\text{RCOO})_6\} = \text{Z2}$ (*cf.* Fig. 9), belongs only to one of the two interpenetrating 3D nets. The Z2 node is similar to the one in **2**, while Z1 is formally close to the expected zinc-oxidocarboxylate cluster (octahedral SBU at the highest symmetry) but with strong geometric distortions accompanied with a complex, not completely resolved crystallographic disorder (Fig. S11, ESI†). The binodal net observed in **3** is also known as the **sqc-3,6-Fdd2-2** type **sqc**-subnet (for net relations, see ref. 72). It is relatively

rare, with only five representatives known in the Topcryst database.⁷¹ All of them are dense coordination polymers based on short bi-azole azole-carboxylate ligands and bear no direct similarity in structural organization with **3**. Hence the hetero-interpenetrated **3** is topologically unique as a MOF.

The general structure-formation logic for **1–3** explains the formation of a class of ‘fused’ **hcb**-nets rather than a singular compound. This class could be compared with the class of pillared **hcb** nets, which is also highly variable when the pillaring could occur both on one and both sides of the planes (connection isomerism). For the case of the ‘fused’ nets, which could be viewed as a zero-length pillaring, the variety stems from variable number of connections for a fused node (4–6 in this work).

The summary on topological characteristics of the nets in structures **1–3** is given in Table S4, ESI†. Except for the simple ‘dual-strand’ **hcb** net in **2**, the other cases are highly special. It is instructive to observe the dominance of 4- and 6-rings (and their exclusivity as smaller cycles). This is the reflection of the ‘fused’ **hcb** structure-organization. When two stacked **hcb** nets are fused *via* alternating vertices a 4-ring is formed (Fig. S16, ESI†). It is a sole outcome for the case of **2** with only one pair of fused-**hcb** nets, but larger rings appear when the fusing of the **hcb**-layers occurs on different levels as in **1** and **3**.

An aspect which should not be overlooked is the flexibility of the $\{\text{Cu}_3(\text{mpba})_3\}^{3-}$ metallo-linker, which surpasses the bbc^{3-} and bte^{3-} flexibilities (*vide supra*, Fig. S44, ESI†). This flexibility can be characterized in terms of separate contributors, as a system of rigid elements, connected by hinges.⁷³ The flexibility factors regarding the $\{\text{Cu}_3(\text{mpba})_3\}^{3-}$ metallo-linker are based on (i) the $\{\text{Cu}_3(\text{pz})_3\}$ cluster level off-plane deformations, (ii) an equivalent flexibility of the phenyl connectors to the carboxylate group, and (iii) the flexibility due to a rotational degree of freedom between the pyrazolate and phenyl moieties. In the example structure of **2**, with its four crystallographically different $\{\text{Cu}_3(\text{mpba})_3\}^{3-}$ metallo-linkers, the benzoate groups bend out of the plane through the three Cu atoms by 0.7° to 14.5° , most of them between 6° and 8° (Fig. S45, ESI†).

Cu/Zn analysis

In view of the disorder in the Zn-SBUs of all three compounds (*cf.* Fig. 1, 3b and 7b, see also Fig. S4, S9 and S11, ESI†) the elemental analysis of the Cu:Zn ratio and the confirmation of the sole presence of the Cu(I) oxidation state (exclusion of Cu(II)) became important. It seemed necessary to ensure that the Zn-SBUs were indeed Zn-only SBUs with no admixture of Cu(II).

The X-ray photoelectron survey spectra (Fig. S23, ESI†) confirm the expected presence of Cu, Zn, O, N and C in the materials. The Cu 2p XP spectra for all three compounds display the core band at around 933 eV for Cu 2p_{3/2} which corresponds to the typical position of Cu(I) (Fig. S24a–S26a, ESI†).⁷⁴ The near perfect single-peak fit indicates the



presence of only one type of Cu(I) species. Cu(II) species are not present in any of the compounds, according to the absence of the characteristic band at ~ 934 eV in the Cu 2p XP spectra.⁷⁴ The band at ~ 1022 eV in the 2p XPS spectra corresponds to Zn(II) and is observed for all three materials (Fig. S24b–S26b, ESI†).⁷⁵

The shape of the Cu LMM peak from the Auger electrons can also be useful in determining the chemical state of Cu. It is particularly indicative in the determination of Cu metal *versus* Cu(I) in the absence of Cu(II) species.⁷⁶ The Cu-LMM peak is observed at a kinetic energy of 916.5 eV in the Auger spectrum, which is a part of the XP spectrum (Fig. S27, ESI†). The modified Auger parameter α' was calculated by adding the binding energy of the Cu 2p_{3/2} photoelectron peak and the kinetic energy of the Cu L3M45M45 Auger peak.⁷⁶ The value of the modified Auger- α' parameter for Cu LMM and the Cu 2p_{3/2} electron was about (933 + 916.5 =) 1849 eV, which confirmed the +1 oxidation state of Cu.^{77,78} For Cu(II) the Auger kinetic energy should lie at 918 eV and the modified Auger- α' parameter should be about 1851 eV.^{77,78} The value of the modified Auger- α' parameter for Zn LMM (988 eV, Fig. S28, ESI†) and the Zn 2p_{3/2} electrons (1022 eV) was approximately 2010 eV, which also confirms the +2 oxidation state of Zn.^{75,79}

The fitting of the C 1s spectra (Fig. S29, ESI†) yielded three bands centered at 285, 286 and 289 eV which can be assigned to C–H/C–C/C=C, C–N, and –COO, respectively.⁸⁰ The O 1s band observed at 531 eV (Fig. S29, ESI†) is due to Zn-bound oxygen, whereas the band at around 533 eV represents organic C–O.^{81,82} The fitting of the N 1s curves shows two types of component peaks (Fig. S29, ESI†), with the major contribution corresponding to Cu–N (399 eV) and the minor one to amide N–(C=O) (400 eV) from coordinated and residual DMF solvent molecules.⁸³

From atomic absorption spectroscopy (AAS) (Table S5, ESI†) and scanning electron microscopy energy-dispersive X-ray spectroscopy (SEM-EDX) (Section S17–S20, ESI† Tables S6 and S7 ESI†), the molar Cu:Zn ratio for **1** was 1:0.44 (AAS) and 1:0.44 (EDX, Table S4, ESI†), which is in good agreement with the expectation from the X-ray refinement (Cu:Zn = 9:4 = 1:0.44). For **3** the analyses gave 1:0.53 (AAS) and 1:0.53 (EDX, Table S3, ESI†), slightly below the value from the X-ray refinement (Cu:Zn = 12:7 = 1:0.58).

Thermogravimetric and gas sorption studies

Thermogravimetric analysis (TGA) of the as-synthesized material **1** (Fig. S21a, ESI†) showed a mass loss of 18 wt% up to ~ 300 °C, which can be attributed to the loss of 9 DMF molecules (calc. 17.8% for 9 DMF). This is in good agreement with the refined coordinated (2 DMF) and SQUEEZE-removed DMF molecules (~ 7.5 DMF, Table S2, ESI†) per formula unit. For compound **2**, TGA gave a mass loss of 26 wt% up to ~ 270 °C, which can be ascribed to the loss of 18 DMF molecules (calc. 26.1% for 18 DMF) (Fig. S21b, ESI†). This agrees well with the found (5 DMF) and SQUEEZE-removed DMF

molecules (~ 14 DMF, Table S2, ESI†) per formula unit. The TGA of the as-synthesized material **3** (Fig. S21b, ESI†) exhibited a mass loss of 20% up to ~ 300 °C, which can be attributed to the loss of 14 DMF molecules (calc. 20.0% for 13.6 DMF). This result is in acceptable agreement with the refined (3 DMF) and squeezed (~ 12 DMF, Table S2, ESI†) DMF molecules per formula unit. The TGA of the activated compound **3-act.** (Fig. S21b, ESI†) did not exhibit any mass loss up to ~ 300 °C. After 300 °C decomposition of all three materials occurred with a mass loss up to 300 °C of 25–30 wt% (Fig. S21, ESI†).

In view of the structural similarity of **1–3** regarding the formation of hexagonal nets in staggered stacking mode (Fig. 3b for **1**, Fig. 6b for **2** and Fig. 8a for **3**; Fig. S13 for comparison) it is no surprise that all three compounds yield a similar specific surface area from a crystal void calculation with CrystalExplorer⁸⁴ and a similar specific pore volume from CrystalExplorer, Mercury^{85,86} and PLATON⁸⁷ (Table S11, ESI†).

Various attempts to activate **1** and to prove the presence of a porous material were not successful. Either direct activation in a vacuum under heating or milder approaches *via* preceding solvent exchange or supercritical drying yielded amorphous materials (**1-act.** Fig. S3, ESI†) which did not demonstrate appreciable Ar (87 K) or CO₂ (195 K) adsorption.

The activation attempts of **3** by heating under vacuum were more successful, and the resultant **3-act.** material demonstrated significant crystallinity, even if inferior to **3** as judged by the peak broadening in the PXRD pattern (Fig. S3, ESI†). The latter also witnesses a structural transformation caused by the loss of solvent molecules (Fig. S21b, ESI†). The two-fold interpenetrated 3D framework in **3** expectedly provides higher structural resilience and prevents a complete structural collapse compared to the polycatenated 2D framework in **1**. But even in **3** there is no prerequisite for a highly rigid structure because of the ligand flexibility (*vide supra*). Notwithstanding the crystallinity, **3-act.** demonstrated only a small N₂ uptake at 77 K with a hysteresis spanning the whole measurement range, indicating kinetic hindrances (Fig. S30, ESI†). At 87 K (liquid argon temperature) argon fills narrow micropores at significantly higher relative pressures in comparison with nitrogen at 77 K.^{88,89} Accordingly, Ar with also a smaller kinetic diameter (3.4 *vs.* 3.64 Å for N₂) and weaker interaction with the adsorbent implying faster equilibration rates compared to N₂ was selected for further studies. The adsorption of Ar at 87 K by **3-act.** was indeed unhindered, followed a type I isotherm (Fig. 10) and allowed for BET surface analysis. The material possesses a total pore volume of 0.30 cm³ g⁻¹ at $P/P_0 = 0.95$, a micro pore volume of 0.25 cm³ g⁻¹ at $P/P_0 = 0.1$, and a BET surface area of 762 m² g⁻¹ (Fig. S31b, ESI†). The desorption shows a small hysteresis loop (type H4) which is typical for micro-mesoporous materials.⁸⁹

CO₂ (and also CH₄) gas sorption studies were performed for the **3-act.** material at 273 K and 293 K. Due to the higher



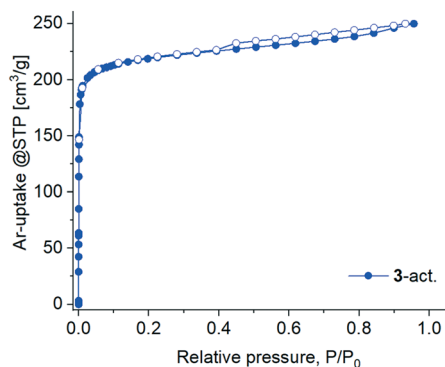


Fig. 10 Ar sorption isotherm at 87 K for 3-act.

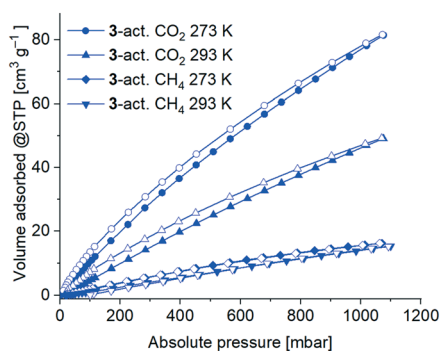


Fig. 11 CO₂ and CH₄ sorption isotherms of 3-act. at 293 K and 273 K with adsorption as filled symbols and desorption as open symbols.

temperatures only minor kinetic hindrances were observed. The adsorption isotherms are of type Ia (Fig. 11), albeit not reaching saturation for CO₂ at 1 bar, which again underlines the microporous character of the material.

As expected, the uptake of the slightly larger non-polar gas CH₄ was significantly lower with 15 cm³ g⁻¹ (273 K) and 14 cm³ g⁻¹ (293 K) than for CO₂ 78 cm³ g⁻¹ (273 K) and 46 cm³ g⁻¹ (293 K). Consequently, the determined CO₂:CH₄ selectivity by the Henry plots of the adsorption isotherms results in a higher selectivity for CO₂ (4.8:1 at 273 K, 3.5:1 at 293 K; Fig. S32, Table S10, ESI†).

The isosteric enthalpy of adsorption at zero coverage, ΔH_{ads}^0 for CO₂ and CH₄ from adsorption isotherms at 273–293 K (Fig. 11) was calculated with the virial fit method (Fig. S33 and S34, ESI†).⁹⁰ As expected, the values of the isosteric enthalpy of adsorption ΔH_{ads} are significantly higher (*i.e.* more negative) for CO₂ than for CH₄ at the same gas uptake. At zero coverage ΔH_{ads}^0 (CO₂) is -29 kJ mol⁻¹ and ΔH_{ads}^0 (CH₄) is -15 kJ mol⁻¹. Compared to other MOFs, these values are at the lower end for ΔH_{ads}^0 .^{91–93}

Conclusions

The N,O-heteroditopic semi-flexible mbpa³⁻ ligand with carboxylate and dimethylpyrazolate tails was used for the exploration of mixed-metal Cu(I)/Zn(II) coordination networks, aiming to combine the probable {Cu₃(3,5-dimethylpyrazolate)₃}

and Zn₄O(CO₂)₆ SBUs. An alternative viewpoint on the synthesis in polar media is the *in situ* formation of the {Cu₃-(mpba)₃}³⁻ metallo-ligand, as an intermediate product, which is engaged in minimal ligand-exchange concurrence in a hard-ligand environment. Accordingly, {Cu₃(mpba)₃}³⁻ is viewed as a large triangular ligand, interesting in the context of reticular chemistry, where 1,3,5-benzenetricarboxylate and expanded analogues play an important role.

Three coordination networks, 1–3, were obtained in a course of low-temperature solvothermal synthesis from *N,N*-dimethylformamide and, remarkably, there is a condition-range when all the three compounds formed simultaneously. In all compounds columnar stacks of Cu₃(3,5-dimethylpyrazolate)₃ clusters associated by cuprophilic interactions are the dominant structural feature. Parallel orientation of the stacks is the most general way to achieve an efficient packing; the various 2-4(5)-nuclear Zn-SBUs are rather adapting to this requirement. The resulting networks bear similarities to stacked **hcb** nets, which would be realized for trigonal SBU-only assembly. The additional connectivities of the Zn-SBU clusters ‘fuse’ the **hcb** nets together, yielding layered 2D, layered polycatenated 2D → 3D or 3D frameworks. Despite the interpenetration, the permanent porosity for 3 with a 3D porous structure was demonstrated by Ar, CO₂ and CH₄ sorption.

Thus, the structures 1–3 demonstrate similar underlying network-formation principles resulting in what could be named a ‘fused’ **hcb** structure class. The structure prediction could be only fuzzy or vague here, and yet such limited design accuracy could be productive. The {Cu₃(mpba)₃}³⁻ metallo-linker has structural degrees of freedom which allow for certain self-adjustments regarding the relative angular positions of the carboxylate groups and this appears favorable for an explorative search approach not based on an exact prediction of a particular structural type. Further elongation of the ligand might yield similar structures due to the good stabilization granted by the cuprophilic stacking. Interesting possibilities of defective structures could be foreseen by using co-ligands with terminated connectivity, like [Cu₃(3,5-dimethylpyrazolate)₃], which could be incorporated in the stacks and decrease the general connectivity.

It is worth stressing the interest in mixed-metal MOFs based on heteroditopic ligands. In the case of mbpa, the Cu^I/Al^{III} combination seems to be particularly interesting due to the multitude of possible aluminum oxidocarboxylate MOFs. Another interesting aspect is the use of the limited-prediction approach when a dominant self-assembly factor is present and the semi-flexible ligands/clusters allow for some self-adjustment (ligands with a high level of flexibility are generally not compatible with permanent porosity). For example, for the octahedral {Zn₄O(COO)₆} and the triangular planar {Cu₃(pz)₃} SBUs a multitude of combination possibilities exist, provided that certain self-adjustment is possible. Among the suitable alternating two vertex 3,6-c nets, the rutile **rtl** net possesses the highest symmetry and suitable



node geometries, however its demands an adjusting turn of the neighboring SBUs by $\sim 45^\circ$, which could be supported by the mpba moiety of the $\{\text{Cu}_3(\text{mpba})_3\}^{3-}$ linker. Despite a multitude of possibilities, all three obtained compounds 1–3 follow the same structure-formation logic: the cuprophilic interactions between the $\{\text{Cu}_3(\text{pz})_3\}$ SBUs, frequently occurring for such structures, evidently play an important pre-organizing role and dictate the course of the self-assembly, which is allowed by the geometric self-adjustment provided by the ligand and concomitant flexible cluster actualization.

Experimental section

Chemicals

Reagents and solvents were obtained from commercial sources and used as received without further purification. $\text{Cu}(\text{NO}_3)_2 \cdot 2.5\text{H}_2\text{O}$ (98%) was purchased from Alfa Aesar, $\text{Zn}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$ (98%) from ALFA Chemistry and $\text{Zn}(\text{CH}_3\text{COO})_2 \cdot 2\text{H}_2\text{O}$ (98%) from VWR. *N,N*-Dimethylformamide (DMF) p.a. was obtained from Riedel-de Haën, tetrafluoroboric acid (HBF_4 , 48 wt% in H_2O) from Sigma-Aldrich and conc. nitric acid from Chemsolute. The synthesis of 4-(3,5-dimethyl-1*H*-pyrazol-4-yl)benzoic acid ($\text{H}_2\text{-mpba}$) was carried out according to the literature procedure.⁴⁵

Instrumentation

Elemental analysis measurements of carbon, hydrogen, nitrogen and sulfur (CHNS) were done with an Elementar Vario MICRO cube. IR-spectra were recorded on a Bruker FT-IR Tensor 37 spectrometer in the 4000–500 cm^{-1} region with 2 cm^{-1} resolution as KBr disks. Powder X-ray diffraction (PXRD) measurements were carried out with a Rigaku MiniFlex600 powder diffractometer using a flat silicon, low background sample holder at 40 kV, 60 mA (600 W) for $\text{Cu-K}\alpha$ radiation ($\lambda = 1.54184 \text{ \AA}$). In all diffractograms, the most intense reflection was normalized to 1. Simulated powder X-ray diffractograms were obtained from single-crystal data using the MERCURY 3.5.1 software program.⁹⁴ Thermogravimetric analysis (TGA) was performed using a Netzsch TG 209 F3 Tarsus from 20 to 600 $^\circ\text{C}$ with a heating range of 5 K min^{-1} under nitrogen atmosphere. Atomic absorption spectroscopy (AAS) was carried out on a PinAAcle 900T from Perkin-Elmer with a copper or zinc hollow cathode in acetylene/air-flame at 2300 $^\circ\text{C}$. Scanning electron microscopy (SEM) images were obtained using a Jeol JSM-6510LV QSEM advanced electron microscope equipped with a LaB_6 cathode at 5–20 kV. The microscope was equipped with a Bruker Xflash 410 silicon drift detector for energy-dispersive X-ray (EDX) spectroscopy. The samples were coated with gold (Au) using a Jeol JFC 1200 finecoater (20 mA for 25 s) before the measurements. X-ray photoelectron spectroscopy (XPS) measurements were made on an ULVAC-PHI VersaProbe II microfocus X-ray photoelectron spectrometer. Experimental XP spectra were fitted by the

CasaXPS program, version 2.3.19PR1.0, copyright 1993–2018 Casa Software Ltd. Volumetric gas sorption measurements up to one bar were performed on a Quantachrome Autosorb iQ MP on sample amounts of at least 30 mg, which were degassed/activated under vacuum (10^{-3} mbar) at 200 $^\circ\text{C}$, for 3 h.

Single crystal X-ray structure determination

Single crystal X-ray structures. Suitable single crystals of 1–3 were carefully selected under a polarized-light microscope and mounted in a perfluorinated oil drop. For 1, the single-crystal diffraction data was collected using a Bruker D8 VENTURE diffractometer with a CCD area detector (PHOTON III C14) and an INCOATEC microfocus sealed tube 3.0 for $\text{Mo-K}\alpha$ radiation ($\lambda = 0.71073 \text{ \AA}$) with a multilayer mirror monochromator. The data were collected under a cold nitrogen gas-stream at 100(2) K using ω - and ϕ -scans. Data reduction was performed by APEX 3 (ref. 95) (integration) and TWINABS-2012/1 (ref. 96) was used for absorption correction.

For 2 and 3, the single-crystal diffraction data was collected using a Rigaku XtaLAB Synergy S four-circle diffractometer with a Hybrid Pixel Array Detector and a PhotonJet X-ray source for $\text{Cu-K}\alpha$ radiation ($\lambda = 1.54184 \text{ \AA}$) with a multilayer mirror monochromator. The data were collected under a cold nitrogen gas-stream at 100.0(1) K using ω -scans. Data reduction and absorption correction were performed by CrysAlisPro 1.171.41.61a [Rigaku Oxford Diffraction, 2020 Yarnton, UK].

All three structures were solved by direct methods using SHELXT-2015.⁹⁷ Full-matrix least squares refinements on F^2 were carried out with SHELXL-2015. Crystal data and details on the structure refinement are given in Table S2, ESI.† Graphics were drawn with the program Diamond.⁹⁸ The structure of 3 was refined as a 2-component inversion twin.

In 1–3 in favour of clarity, the mpba²⁻ units and DMF molecules were organized in residues (RESI). All hydrogen atoms were positioned geometrically and refined using riding models with $U_{\text{iso}}(\text{H}) = 1.2 \cdot U_{\text{eq}}(\text{CH}_{\text{arom.}})/U_{\text{eq}}(\text{NH})$ or $1.5 \cdot U_{\text{eq}}(\text{CH}_3)$.

In 1–3 highly disordered solvent molecules were removed *via* the SQUEEZE (Platon) procedure (structures 1 and 3)⁸⁷ or by solvent masking as implemented in OLEX 2 (structure 2). In 1 the resulting solvent accessible volume of 2751 \AA^3 per unit cell contained 604 electrons, which may correspond to 15 DMF molecules (40 e^- each) as the solvent of crystallization per unit cell or 7.5 DMF molecules per given asymmetric formula unit ($Z = 2$). In 2 the resulting solvent accessible volume of 4630 \AA^3 per unit cell contained 1131 electrons, which may correspond to 28 DMF molecules as the solvent of crystallization per unit cell or 14 DMF molecules per given asymmetric formula unit ($Z = 2$). In 3 the resulting solvent accessible volume of 7369 \AA^3 per unit cell contained 1894 electrons, which may



correspond to 47 DMF molecules as the solvent of crystallization per unit cell or ~12 DMF molecules per given asymmetric formula unit ($Z = 4$).

In **1** the Zn-cluster which contains the dimethylamine and dimethylammonium species which are disordered in an 81:19 ratio, affecting both Zn atoms and the proton of the counterion.

In **2** the phenyl rings of RESI 5, 9 and 12 are disordered over two positions due to rotation along the 1,4-axis which is a well-known phenomenon. Furthermore, one DMF molecule coordinating to Zn4 is disordered over two positions in an 86:14 ratio.

In **3** the zinc oxidocarboxylate coordination-bonded cluster ($\{Zn_4O\}$) involving the Zn4 to Zn8a,b atoms features a complex disorder, which could be described in the first approximation as three overlapping $\{Zn_4O\}$ arrangements. The disorder is driven by the demand imposed by the crystal structure/topology: six carboxylates, which are located within convenient reach, are not arranged suitably to sustain the regular octahedral $\{Zn_4O(RCOO)_6\}$ cluster. The 'three overlapping pyramids' model with a nearly common base (Zn5, Zn6, Zn7) assumes a much lower occupancy of the 'capping/axial' Zn-atoms (Zn4, Zn8a,b), however the refined equivalent U_{iso} values do not indicate dramatic differences. While the relatively large temperature factors could mask additional non-evident disorder possibilities, there is another possible reason for the higher occupancy of the 'capping' atoms: at least for Zn8 there is a weak residual electronic density in the vicinity. Instead of speculating the possibility of an increased number of Zn atoms in the cluster, the disorder was modelled as even *via* ascribing 0.8 occupancy for each of the five Zn atoms (the Zn8 atom is additionally split into two components, Zn8a and Zn8b, with refined contributions). This concise approach reflects both the experimental observations and the limited knowledge about disorder, whose occurrence is, however, not unusual in the given structural setting. Accordingly, the cluster composition, involving the Zn4 to Zn8a,b atoms, is assumed to be $\{Zn_4O(H_2O/DMF)_4(RCOO)_6\}$. The coordinated water molecules (O5S to O8S atoms) are a simplified description of site-sharing H₂O/DMF ligands; the too weak residual electron density made the refinement of the DMF constituents not practical.

The structural data have been deposited with the Cambridge Crystallographic Data Center under deposition numbers CCDC 2105164–2105166.†

Synthetic procedures

$[Me_2NH_2][Zn_4\{Cu_3^I(mpba)_3\}_3(Me_2NH)(DMF)_2]$ (**1**). 20.0 mg (0.092 mmol) of H₂mpba, 21.5 mg (0.092 mmol) of Cu(NO₃)₂·2.5H₂O and 16.9 mg (0.065 mmol) of Zn(NO₃)₂·4H₂O were each dissolved in 1 of mL *N,N*-dimethylformamide (DMF). The three solutions were combined in a glass vial equipped with a screw cap,

sealed and transferred in a preheated oven at 90 °C for six days. After cooling to room temperature, the formed clear yellow parallelepiped-shaped crystals (Fig. S1†) were filtered and washed three times with 3 mL of DMF each (yield 25 mg, 56%, based on zinc salt). Calc. for C₁₀₈H₉₀Cu₉N₁₈O₁₈Zn₄ (2761.42 g mol⁻¹) C 46.97, H 3.28, N 9.13; found C 47.20, H 3.31, N 9.15%. IR: (KBr, Fig. S22, ESI†): $\tilde{\nu} = 3435$ (br), 3067 (br), 2567 (br), 2501 (br), 2360 (br), 2341 (br), 2273 (br), 1661 (s), 1608 (s), 1542 (s), 1403 (br), 1284 (w), 1253 (w), 1179 (s), 1145 (w), 1093 (w), 1062 (w), 1034 (m), 1014 (w), 865 (w), 847 (w), 812 (w), 782 (s), 714 (s), 692 (m), 660 (w), 634 (m), 610 (w), 565 (w), 517 (s), 442 (w) cm⁻¹.

$[Zn_6\{Cu_3^I(mpba)_3\}_4(DMF)_5]$ (**2**). 20.0 mg (0.092 mmol) of H₂mpba, 21.5 mg (0.092 mmol) of Cu(NO₃)₂·2.5H₂O and 15.2 mg (0.070 mmol) of Zn(CH₃COO)₂·2H₂O were each dissolved in 1 mL of DMF. The three solutions were combined in a glass vial with a screw cap. After a few minutes, a greenish precipitate formed. This was dissolved again by adding 30 μL of aqueous tetrafluoroboric acid (HBF₄, 48% min w/w). The glass vial was then sealed and transferred in a preheated oven at 90 °C for nine days. After this time, an off-white precipitate was observed. After cooling to room temperature, the product was filtered and washed three times with 3 mL DMF (yield 15 mg, 35%, based on zinc salt). Calc. for C₁₄₄H₁₂₀Cu₁₂N₂₄O₂₄Zn₆ (3725.60 g mol⁻¹) C 46.42, H 3.25, N 9.02; found C 46.68, H 3.60, N 9.18%. IR: (KBr, Fig. S22, ESI†): $\tilde{\nu} = 3429$ (br), 3184 (br), 3070 (br), 3042 (br), 2920 (m), 2854 (br), 1934 (br), 1811 (br), 1677 (s), 1610 (s), 1586 (w), 1537 (w), 1493 (w), 1405 (br), 1281 (br), 1257 (br), 1178 (w), 1145 (w), 1090 (w), 1013 (m), 860 (w), 805 (w), 777 (w), 712 (w), 660 (w), 609 (w), 559 (w), 516 (br), 442 (w) cm⁻¹.

$[Zn_3\{Cu_3^I(mpba)_3\}_2(DMF)_3(H_2O)] [Zn_4(\mu_4-O)\{Cu_3^I(mpba)_3\}_2(H_2O)_4]$ (**3**). 20.0 mg (0.092 mmol) of H₂mpba, 21.5 mg (0.092 mmol) of Cu(NO₃)₂·2.5H₂O and 15.2 mg (0.070 mmol) of Zn(CH₃COO)₂·2H₂O were each dissolved in 1 mL of DMF. The three solutions were combined in a glass vial equipped with a screw cap. After a few minutes, a greenish precipitate formed. It was dissolved again by adding 10 μL of aqueous tetrafluoroboric acid (HBF₄, 48% min w/w). The glass vial was then sealed and transferred in a preheated oven at 90 °C for six days. The formed clear yellow trapezoidal prismatic crystals (Fig. S2a and b, ESI†) were filtered and washed three times with 3 mL DMF to obtain 28 mg (72%, based on the zinc salt). Calc. for C₁₄₄H₁₀₅Cu₁₂N₂₄O_{23.67}Zn₇ (3791.00 g mol⁻¹) C 45.62, H 3.19, N 8.87; found C 45.60, H 3.12, N 8.72%. IR: (KBr, Fig. S22, ESI†): $\tilde{\nu} = 3435$ (br), 3067 (w), 3042 (w), 2953 (w), 2919 (w), 2854 (w), 2807 (w), 2568 (w), 2501 (w), 2361 (br), 2342 (br), 2273 (br), 1937 (w), 1813 (w), 1668 (s), 1608 (s), 1543 (s), 1492 (s), 1386 (br), 1284 (w), 1255 (w), 1179 (s), 1143 (w), 1094 (w), 1062 (w), 1034 (m), 1014 (w), 905 (w), 866 (w), 847 (w), 812 (w), 782 (s), 714 (s), 694 (w), 661 (w), 634 (m), 610 (w), 565 (w), 517 (s), 438 (w), 406 (w) cm⁻¹.



Author contributions

This manuscript was written through contributions of all the authors. All the authors have given approval to the final version of this manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors thank Mrs Birgit Tommes for carrying out the FT-IR experiments and Annette Ricken for the AAS measurements. Dr. Christian Schürmann (Rigaku) and Dr. Tobias Stürzer (Bruker AXS) are thanked for their crucial crystal-screening and data-collection efforts. The Rigaku X-ray diffractometer was funded by the Deutsche Forschungsgemeinschaft (DFG, German Research Foundation) through project number 440366605. We thank the CeMSA@HHU (Center for Molecular and Structural Analytics@Heinrich Heine University) for recording the mass-spectrometric and the NMR-spectroscopic data.

Notes and references

- O. K. Farha and J. T. Hupp, *Acc. Chem. Res.*, 2010, **43**, 1166–1175.
- M. Taddei, *Coord. Chem. Rev.*, 2017, **343**, 1–24.
- J. D. Evans, B. Garai, H. Reinsch, W. Li, S. Dissegna, V. Bon, I. Senkovska, R. A. Fischer, S. Kaskel, C. Janiak, N. Stock and D. Volkmer, *Coord. Chem. Rev.*, 2019, **380**, 378–418.
- J. R. Long and O. M. Yaghi, *Chem. Soc. Rev.*, 2009, **38**, 1213–1214.
- X. Han, S. Yang and M. Schröder, *Nat. Rev. Chem.*, 2019, **3**, 108–118.
- S. R. Batten, N. R. Champness, X.-M. Chen, J. Garcia-Martinez, S. Kitagawa, L. Öhrström, M. O’Keeffe, M. Paik Suh and J. Reedijk, *Pure Appl. Chem.*, 2013, **85**, 1715–1724.
- H.-C. J. Zhou and S. Kitagawa, *Chem. Soc. Rev.*, 2014, **43**, 5415–5418.
- S. S. Y. Chui, S. M. F. Lo, J. P. H. Charmant, A. G. Orpen and I. D. Williams, *Science*, 1999, **283**, 1148–1150.
- L. Xie, S. Liu, B. Gao, C. Zhang, C. Sun, D. Li and Z. Su, *Chem. Commun.*, 2005, 2402–2404.
- L. H. Xie, J. B. Lin, X. M. Liu, W. Xue, W. X. Zhang, S. X. Liu, J. P. Zhang and X. M. Chen, *Sci. China: Chem.*, 2010, **53**, 2144–2151.
- C. Ordonez, M. Fonari, J. Lindline, Q. Wei and T. Timofeeva, *Cryst. Growth Des.*, 2014, **14**(11), 5452–5465.
- W. W. Lestari, M. Arwinawati, R. Martien and T. Kusumaningsih, *Mater. Chem. Phys.*, 2018, **204**, 141–146.
- H. K. Chae, D. Y. Siberio-Pérez, J. Kim, Y. B. Go, M. Eddaoudi, A. J. Matzger, M. O’Keeffe and O. M. Yaghi, *Nature*, 2004, **427**, 523–527.
- H. Furukawa, M. A. Miller and O. M. Yaghi, *J. Mater. Chem.*, 2007, **17**, 3197–3204.
- I. M. Hauptvogel, V. Bon, R. Grünker, I. A. Baburin, I. Senkovska, U. Mueller and S. Kaskel, *Dalton Trans.*, 2021, **41**, 4172–4179.
- Z. W. Wei, D. Q. Yuan, X. L. Zhao, D. F. Sun and H.-C. Zhou, *Sci. China: Chem.*, 2013, **56**, 418–422.
- B. Chen, M. Eddaoudi, S. T. Hyde, M. O’Keeffe and O. M. Yaghi, *Science*, 2001, **291**, 1021–1023.
- H. Furukawa, Y. B. Go, N. Ko, Y. K. Park, F. J. Uribe-Romo, J. Kim, M. O’Keeffe and O. M. Yaghi, *Inorg. Chem.*, 2011, **50**, 9147–9152.
- K. Gedrich, I. Senkovska, N. Klein, U. Stoeck, A. Henschel, M. R. Lohe, I. A. Baburin, U. Mueller and S. Kaskel, *Angew. Chem., Int. Ed.*, 2010, **49**, 8489–8492.
- H. Furukawa, N. Ko, Y. B. Go, N. Aratani, S. B. Choi, E. Choi, A. O. Yazaydin, R. Q. Snurr, M. O’Keeffe, J. Kim and O. M. Yaghi, *Science*, 2010, **329**, 424–428.
- J. Wang, C. Rao, L. Lu, S. Zhang, M. Muddassir and J. Liu, *CrystEngComm*, 2021, **23**, 741–747.
- M. Wang, J. Liu, J. Jin, D. Wu, G. Yang, W.-Y. Zhang and Y.-Y. Wang, *CrystEngComm*, 2021, **23**, 663–670.
- W. Li and J.-S. Tan, *J. Mol. Struct.*, 2021, **1237**, 130359.
- R. A. Agarwal, *Cryst. Growth Des.*, 2021, **21**, 2046–2055.
- M.-H. Tang, Y. Shi, X.-L. Jiang, H. Xu, Y. Ma and B. Zhao, *J. Mater. Chem. C*, 2021, **9**, 9643–9649.
- E. Lopez-Maya, N. M. Padial, J. Castells-Gil, C. R. Ganivet, A. Rubio-Gaspar, F. G. Cirujano, N. Almora-Barrios, S. Tatay, S. Navalon and C. Marti-Gastaldo, *Angew. Chem., Int. Ed.*, 2021, **60**, 11868–11873.
- J. Chen, T. Chen, S. Xiang, J. Zhang and Z. Zhang, *Z. Anorg. Allg. Chem.*, 2021, **647**, 1301–1304.
- D. Sun, Y. Ke, T. M. Mattox, S. Parkin and H.-C. Zhou, *Inorg. Chem.*, 2006, **45**, 7566–7568.
- D. Saha, R. Zacharia, L. Lafi, D. Cossement and R. Chahine, *Int. J. Hydrogen Energy*, 2012, **37**, 5100–5107.
- S. Lee, H.-B. Bürgi, S. A. Alshmiri and O. M. Yaghi, *J. Am. Chem. Soc.*, 2018, **140**, 8958–8964.
- J. Rong, W. Zhang and J. Bai, *CrystEngComm*, 2016, **18**, 7728–7736.
- D. Sun, S. Ma, Y. Ke, D. J. Collins and H.-C. Zhou, *J. Am. Chem. Soc.*, 2006, **128**, 3896–3897.
- S. Ma and H.-C. Zhou, *J. Am. Chem. Soc.*, 2006, **128**, 11734–11735.
- R. R. R. Prasad, C. Pleass, A. L. Rigg, D. B. Cordes, M. M. Lozinska, V. M. Georgieva, F. Hoffmann, A. M. Z. Slawin and P. A. Wright, *CrystEngComm*, 2021, **23**, 804–812.
- T. Liu, X. Shen, X. Shen, C. He, J. Liu and J.-J. Liu, *CrystEngComm*, 2021, **23**, 4667–4673.
- K. Wang, S. Wang, J. Liu, Y. Guo, F. Mao, H. Wu and Q. Zhang, *ACS Appl. Mater. Interfaces*, 2021, **13**, 15315–15323.
- A. M. Toader, M. C. Buta, F. Cimpoesu, A.-I. Toma, C. M. Zalaru, L. O. Cinteza and M. Ferbinteanu, *Chemistry*, 2021, **3**, 411–439.
- N. V. S. Harisomayajula, S. Makovetskyi and Y.-C. Tsai, *Chem. – Eur. J.*, 2019, **25**, 8936–8954.
- K. Singh, J. R. Long and P. Stavropoulos, *J. Am. Chem. Soc.*, 1997, **119**, 2942–2943.



- 40 (a) S. M.-F. Lo, S. S.-Y. Chui, L.-Y. Shek, Z. Y. Lin, X. X. Zhang, G.-H. Wen and I. D. Williams, *J. Am. Chem. Soc.*, 2000, **122**, 6293–6294; (b) X.-C. Huang, J.-P. Zhang, Y.-Y. Lin, X.-L. Yua and X. M. Chen, *Chem. Commun.*, 2004, 1100–1101; (c) K. Tomar, A. Verma and P. K. Bharadwaj, *Cryst. Growth Des.*, 2018, **18**, 2397–2404; (d) N. Grosjean, C. Descorme and M. Besson, *Appl. Catal., B*, 2010, **97**, 276–283.
- 41 C.-T. He, P.-Q. Liao, D.-D. Zhou, B.-Y. Wang, W.-X. Zhang, J.-P. Zhang and X.-M. Chen, *Chem. Sci.*, 2014, **5**, 4755–4762.
- 42 Z.-M. Ye, C.-T. He, Y.-T. Xu, R. Krishna, Y. Xie, D.-D. Zhou, H.-L. Zhou, J.-P. Zhang and X.-M. Chen, *Cryst. Growth Des.*, 2017, **17**, 2166–2171.
- 43 C. Heering, I. Boldog, V. Vasylyeva, J. Sanchiz and C. Janiak, *CrystEngComm*, 2013, **15**, 9757–9768.
- 44 S. Menzel, S. Millan, S.-P. Höfert, A. Nuhnen, S. Gökpınar, A. Schmitz and C. Janiak, *Dalton Trans.*, 2020, **49**, 12854–12864.
- 45 M. R. Bryant, A. D. Burrows, C. M. Fitchett, C. S. Hawes, S. O. Hunter, L. L. Keenan, D. J. Kelly, P. E. Kruger, M. F. Mahon and C. Richardson, *Dalton Trans.*, 2015, **44**, 9269–9280.
- 46 C. Wang, C. Liu, H.-R. Tian, L.-J. Li and Z.-M. Sun, *Chem. – Eur. J.*, 2018, **24**, 2952–2961.
- 47 (a) A. E. Martell and R. D. Hancock, in *Metal Complexes in Aqueous Solutions*, ed. J. P. Fackler, Plenum Press, 1996; (b) R. B. Martin, in *Encyclopedia of Inorganic Chemistry*, ed. R. B. King, Wiley, 1994, pp. 2185–2189; (c) R. H. Crabtree, in *Encyclopedia of Inorganic Chemistry*, ed. R. B. King, Wiley, 1994, p. 809.
- 48 B. Tu, Q. Pang, H. Xu, X. Li, Y. Wang, Z. Ma, L. Wenig and Q. Li, *J. Am. Chem. Soc.*, 2017, **139**, 7998–8007.
- 49 B. Tu, L. Diestel, Z.-L. Shi, W. R. L. N. Bandara, Y. Chen, W. Lin, Y.-B. Zhang, S. G. Telfer and Q. Li, *Angew. Chem., Int. Ed.*, 2019, **58**, 5348–5353.
- 50 J. Fritzsche, D. Denysenko, M. Grzywa and D. Volkmer, *Dalton Trans.*, 2017, **46**, 14907–14915.
- 51 S. Menzel, S.-P. Höfert, S. Öztürk, A. Schmitz and C. Janiak, *Z. Anorg. Allg. Chem.*, 2021, **647**, 803–808.
- 52 (a) S. M.-F. Lo, S. S.-Y. Chui, L.-Y. Shek, Z. Y. Lin, X. X. Zhang, G.-H. Wen and I. D. Williams, *J. Am. Chem. Soc.*, 2000, **122**, 6293–6294; (b) X.-C. Huang, J.-P. Zhang, Y.-Y. Lin, X.-L. Yua and X. M. Chen, *Chem. Commun.*, 2004, 1100–1101; (c) K. Tomar, A. Verma and P. K. Bharadwaj, *Cryst. Growth Des.*, 2018, **18**, 2397–2404; (d) N. Grosjean, C. Descorme and M. Besson, *Appl. Catal., B*, 2010, **97**, 276–283.
- 53 H. Li, M. Eddaoudi, T. L. Groy and O. M. Yaghi, *J. Am. Chem. Soc.*, 1998, **120**, 8571–8572.
- 54 J. Xiao, H. Zhou and A.-H. Yuan, *Acta Crystallogr., Sect. E: Struct. Rep. Online*, 2009, **65**, m1071.
- 55 Y. Mao and P. Y. Zavalij, *Acta Crystallogr., Sect. C: Struct. Chem.*, 2018, **74**, 1719–1724.
- 56 L. Zou, S. Yao, J. Zhao, D.-S. Li, G. Li, Q. Huo and Y. Liu, *Cryst. Growth Des.*, 2017, **17**, 3556–3561.
- 57 I. A. Baburin, V. A. Blatov, L. Carlucci, G. Ciani and D. M. Proserpio, *J. Solid State Chem.*, 2005, **178**, 2452–2474.
- 58 I. A. Baburin, V. A. Blatov, L. Carlucci, G. Ciani and D. M. Proserpio, *CrystEngComm*, 2008, **10**, 1822–1838.
- 59 V. A. Blatov, L. Carlucci, G. Ciani and D. M. Proserpio, *CrystEngComm*, 2004, **6**, 377–395.
- 60 L. Carlucci, G. Ciani and D. M. Proserpio, *Coord. Chem. Rev.*, 2003, **246**, 247–289.
- 61 L. Carlucci, G. Ciani, D. M. Proserpio, T. G. Mitina and V. A. Blatov, *Chem. Rev.*, 2014, **114**, 7557–7580.
- 62 E. Y. Alexandrov, V. A. Blatov and D. M. Proserpio, *CrystEngComm*, 2017, **19**, 1993–2006.
- 63 B. Liu, Y.-C. Qiu, G. Peng and H. Deng, *CrystEngComm*, 2010, **12**, 270–276.
- 64 R. Clerac, F. A. Cotton, L. M. Daniels, J. Gu, C. A. Murillo and H. C. Zhou, *Inorg. Chem.*, 2000, **39**, 4488–4493.
- 65 K. Singh, J. R. Long and P. Stavropoulos, *J. Am. Chem. Soc.*, 1997, **119**, 2942–2943.
- 66 A. W. Addison, T. N. Rao, J. Reedijk, J. van Rijn and G. C. Verschoor, *J. Chem. Soc., Dalton Trans.*, 1984, 1349–1356.
- 67 V. A. Blatov, A. P. Shevchenko and D. M. Proserpio, *Cryst. Growth Des.*, 2014, **14**, 3576–3586.
- 68 E. V. Alexandrov, V. A. Blatov, A. V. Kochetkov and D. M. Proserpio, *CrystEngComm*, 2011, **13**, 3947–3958.
- 69 J.-P. Zhang and S. Kitagawa, *J. Am. Chem. Soc.*, 2008, **130**, 907–917.
- 70 J. He, Y.-G. Yin, T. Wu, D. Li and X.-C. Huang, *Chem. Commun.*, 2006, 2845–2847.
- 71 E. V. Alexandrov, A. P. Shevchenko and V. A. Blatov, *Cryst. Growth Des.*, 2019, **19**, 2604–2614.
- 72 V. A. Blatov and D. M. Proserpio, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2009, **65**, 202–212.
- 73 L. Sarkisov, R. L. Martin, M. Haranczyk and B. Smit, *J. Am. Chem. Soc.*, 2014, **136**, 2228–2231.
- 74 (a) S. Zhang and L. Zhao, *Nat. Commun.*, 2019, **10**, 4848; (b) A. Ahmed, C. M. Robertson, A. Steiner, T. Whittles, A. Ho, V. Dhanak and H. Zhang, *RSC Adv.*, 2016, **6**, 8902–8905; (c) H. Rupp and U. Weser, *Bioinorg. Chem.*, 1976, **6**, 45–59.
- 75 (a) S. Bera, S. Dhara, S. Velmurugan and A. K. Tyagi, *Int. J. Spectrosc.*, 2011, **2012**, 1–4; (b) X-ray photoelectron spectroscopy (XPS) reference page 27.08.2021, Link: <http://www.xpsfitting.com/search/label/Zinc>.
- 76 X-ray photoelectron spectroscopy (XPS) reference page 30.01.2022, Link: <http://www.xpsfitting.com/2012/01/copper.html>.
- 77 M. M. Sun and Y. Kim, *Bull. Korean Chem. Soc.*, 2001, **22**, 748–752.
- 78 M. C. Biesiner, *Surf. Interface Anal.*, 2017, **49**, 1325–1334.
- 79 L. Yin, T. Tsang, I. Adler and E. Yellin, *J. Appl. Phys.*, 1972, **43**, 3464–3467.
- 80 (a) D. Qu, M. Zheng, J. Li, Z. Xie and Z. Sun, *Light: Sci. Appl.*, 2015, **4**(e364), 1–8; (b) F. Arcudi, L. Dordevic and M. Prato, *Angew. Chem., Int. Ed.*, 2016, **55**, 2107–2112.
- 81 Thermo scientific XPS simplified, 18.08.2021, Link: <https://xpssimplified.com/elements/oxygen.php>.
- 82 S. U. Awan, S. K. Hasanain, J. Rashid, S. Hussain, S. A. Shah, M. Z. Hussain, M. Rafique, M. Aftab and R. Khan, *Mater. Chem. Phys.*, 2018, **211**, 510–521.
- 83 (a) W. He, R. Ifraemov, A. Raslin and I. Ho, *Adv. Funct. Mater.*, 2018, **28**, 1707244; (b) Z. Zude, Z. Xiong, Z. Tao, Y. Humaing and L. Qingliang, *J. Mol. Struct.*, 1999, **478**, 23–27.



- 84 (a) P. R. Spackman, M. J. Turner, J. J. McKinnon, S. K. Wolff, D. J. Grimwood, D. Jayatilaka and M. A. Spackman, *J. Appl. Crystallogr.*, 2021, **54**, 1006–1011; (b) CrystalExplorer21.5, © 2005–2021, University of Western Australia, 2021, <https://crystalexplorer.net/>.
- 85 Mercury 2021.2.0, Program for Crystal Structure Visualisation, Exploration and Analysis from the Cambridge Crystallographic Data Center, Copyright CCDC 2001–2021, <http://www.ccdc.cam.ac.uk/mercury/>.
- 86 Mercury 4.0: from visualization to analysis, design and prediction, C. F. Macrae, I. Sovago, S. J. Cottrell, P. T. A. Galek, P. McCabe, E. Pidcock, M. Platings, G. P. Shields, J. S. Stevens, M. Towler and P. A. Wood, *J. Appl. Crystallogr.*, 2020, **53**, 226–235.
- 87 A. L. Spek, Structure Validation in Chemical Crystallography, *Acta Crystallogr., Sect. D: Biol. Crystallogr.*, 2009, **65**, 148–155; A. L. Spek, PLATON– A Multipurpose Crystallographic Tool, Utrecht University, Utrecht, The Netherlands, 2008; L. J. Farrugia, PLATON for Windows implementation, Version 280317, University of Glasgow, Scotland, 2020.
- 88 J. Rouquerol, F. Rouquerol, K. S. W. Sing, P. Llewellyn and G. Maurin, *Adsorption by Powders and Porous Solids: Principles, Methodology and Applications*, Academic Press, 2014.
- 89 M. Thommes, K. Kaneko, A. V. Neimark, J. P. Olivier, F. Rodriguez-Reinoso, J. Rouquerol and K. S. W. Sing, *Pure Appl. Chem.*, 2015, **87**, 1051–1069.
- 90 A. Nuhnen and C. Janiak, *Dalton Trans.*, 2020, **49**, 10295–10307.
- 91 K. Sumida, D. L. Rogow, J. A. Mason, T. M. McDonald, E. D. Bloch, Z. R. Herm, T.-H. Bae and J. R. Long, *Chem. Rev.*, 2012, **112**, 724–781.
- 92 A. Das and D. M. D'Alessandro, *CrystEngComm*, 2015, **17**, 706–718.
- 93 D. M. D'Alessandro, B. Schmit and J. R. Long, *Angew. Chem., Int. Ed.*, 2010, **49**, 6058–6082.
- 94 C. F. Macrae, I. J. Bruno, J. A. Chisholm, P. R. Edgington, P. McCabe, E. Pidcock, L. Rodriguez-Monge, R. Taylor, J. van de Streek and P. A. Wood, *J. Appl. Crystallogr.*, 2008, **41**, 466–470.
- 95 APEX3, SAINT, SADABS and XP, Bruker AXS Inc., Madison, Wisconsin, USA, 2019.
- 96 Bruker, TWINABS-2012/1, Bruker AXS Inc., Madison, Wisconsin, USA, 2012.
- 97 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Adv.*, 2015, **71**, 3–8.
- 98 K. Brandenburg, *Diamond Crystal and Molecular Structure Visualization, Version 4.6.6*, Crystal Impact GbR, Bonn, Germany, 1997–2021.

