

Interplay between defects, disorder and flexibility in metal-organic frameworks

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Metal-organic frameworks are a novel family of chemically diverse materials, which are of interest across engineering, physics, chemistry, biology and medicine-based disciplines. Since the development of the field in its current form more than two decades ago, priority has been placed on the synthesis of new structures. However, more recently, a clear trend has emerged in shifting the emphasis from material design to exploring the chemical and physical properties of structures already known. In particular, although such nanoporous materials were traditionally seen as rigid crystalline structures, there is growing evidence that large-scale flexibility, the presence of defects and long-range disorder are not the exception in metal-organic frameworks, but the rule. Here we offer some perspective into how these concepts are perhaps inescapably intertwined, highlight recent advances in our understanding and discuss how a consideration of the interfaces between them may lead to enhancements of the materials' functionalities.

As an entirely novel class of nanoporous materials, metal-organic frameworks (MOFs) have attracted a lot of interest and are studied by scientists from a wide variety of backgrounds. This is due to their intrinsic aesthetic appeal and potential industrial applications as selective adsorbents and catalysts, substrates for biosensors and drug delivery, and membranes and films in various nanotechnologies, to list but a few examples. Compared with both dense and nanoporous inorganic materials, MOFs are based on relatively weaker interactions (coordinative bonds, π - π stacking, hydrogen bonds, and so on) and present large numbers of intramolecular degrees of freedom. Although all molecular assemblies and solids show some degree of flexibility (even those considered to be rigid, such as the archetypical MOF-5), there is a propensity among coordination polymers to display large-scale dynamic behaviour, which is typically described by the vague term 'flexibility'¹ (Fig. 1). This appellation contains phenomena that are very diverse both in terms of their microscopic origins and their macroscopic manifestations. The latter include negative thermal expansion, a relatively common trait², as well as anomalous mechanical properties³ such as auxeticity (the presence of a negative Poisson's ratio, where a material becomes thicker when stretched along certain directions) or negative linear compressibility (total volume reduction with expansion along a specific direction under mechanical pressure)⁴. Both phenomena are rare among inorganic and molecular solids and stem from wine-rack, honeycomb and related topologies, which allow the framework to act as a Tinkertoy model with rigid struts and hinges⁵.

Just as there is flexibility in any molecular assembly, there is no such thing as a crystalline solid without defects. The prevalence of defects in nanoporous frameworks was first made evident by the difficulty in comparing fluid adsorption isotherms across experimental measurements performed by different groups, and against computational predictions. The most commonly observed effect was an incomplete activation of framework porosity — resulting in guest molecules blocking the nanopores and negatively impacting surface area and adsorption capacity^{6,7}. It was only after some time that defects intrinsic to the frameworks (for example, linker and inorganic node vacancies, partial metal reduction and

dislocations) started to be characterized and reported⁸. Recently, Li and co-workers described how defects may be introduced into the formation mechanism of some MOFs, leading to mismatched growth and gelation, as opposed to crystallization in long-range ordered structures⁹. Indeed, it is now being recognized that such defects do not necessarily need to have adverse effects but can instead give rise to specific functionalities, such as improving adsorption affinity or catalytic activity¹⁰. Hoskins and Robson's description of MOFs in 1990 as "ordered, truly crystalline" structures is recognized as a milestone in the field¹¹. Subsequently, examples of framework dynamics or rotational disordering of cations within MOFs, along with cases that consider topologically or statically disordered materials (that is, X-ray amorphous species possessing no long-range order), perhaps challenge this view^{12,13}. Comparatively few reports of such systems exist, though it is likely

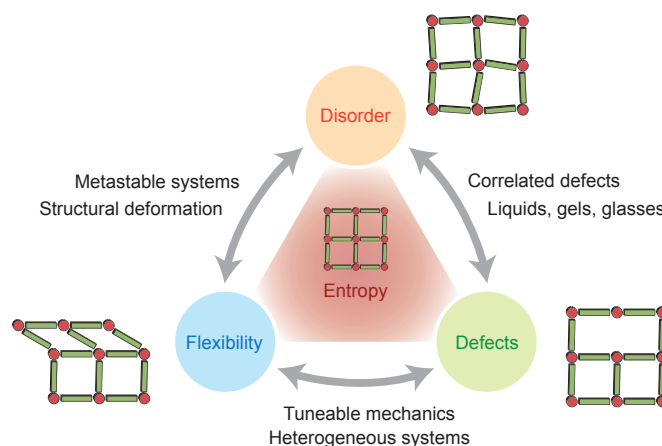


Figure 1 | Flexibility, defects and disorder in metal-organic frameworks.

Illustration of the interplay between the intertwined concepts of flexibility, defects and disorder in metal-organic frameworks, highlighting some of the phenomena (grey arrows) that emerge from their coupling. Entropy plays a central role in all of these systems.

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that many have been synthesized and subsequently discarded due to their lack of crystallinity. Interestingly however, even Robson and Hoskins recognized that their approach to MOFs could yield “amorphous materials ... with many (internal) loose ends”¹¹. Indeed, recent terminology guidelines¹⁴ do not necessarily limit MOF materials to those which exhibit long-range ordering, and disordered MOFs are becoming of increasing interest to the community, driven by an emerging focus on the physical properties of framework structures.

Flexibility, defects and disorder are all related to entropy (Fig. 1). They arise from the high dimensionality of the intramolecular degrees of freedom of the materials and the many corresponding ‘soft’ modes of low energy, as well as the relatively weak nature of the interactions involved in the framework assembly, such as coordinative bonds, π - π stacking and hydrogen bonds. For example, the MIL-53(Al) structure undergoes a thermally induced phase transition in the absence of solvent between 325 and 375 K. The low-temperature narrow-pore form is stabilized by dispersion interactions, though it is the contribution of vibrational entropy that drives the formation of the more porous structure at high temperatures¹⁵. Entropy also plays an important role in the formation of disordered MOF systems upon heating¹⁶, and in some MOFs proves to be a sufficient thermodynamic driving force to overcome unfavourable enthalpies of defect incorporation⁸.

Flexibility and defects

One aspect of the flexibility of MOFs is their general softness (in the terminology of elasticity) or compliance, that is, their high compressibility in comparison to traditional nanoporous materials. This has been widely observed and discussed in the literature¹⁷, based on both experimental measurements and quantum-chemistry predictions, and is linked to the weaker nature of their bonds

(compared to, for example, the very strong Si-O bond in zeolites) and their higher porosities. More dense MOFs (or those with very high connectivities) feature the highest elastic moduli, with bulk moduli in the 25–50 GPa range¹⁸. Flexibility in MOFs is, however, tightly coupled with the potential presence of defects, as can be illustrated in the UiO-66 family of porous isoreticular frameworks.

With a chemical formula based on $[M_6O_4(OH)_4][C_6H_4(COO)_2]_6$ ($M = \text{Ti, Zr, Hf, ...}$) and consisting of twelve connected nodes, these materials display relatively low compressibility. UiO-66 not only presents substantial non-stoichiometry, with up to 10% of terephthalate linkers missing, but the concentration of defects can be controlled in a systematic manner by varying the concentration of a modulator (typically a monocarboxylic acid) during synthesis¹⁹. The capacity of the framework to accommodate a large number of missing linkers has a strong impact on the materials’ properties, both by changing the characteristics of the pore space of the MOF and by introducing sites with higher catalytic activities. In UiO-66(Zr), the inclusion of linker vacancies was used to tune the pore volumes from 0.44 to 1.0 $\text{cm}^3 \text{g}^{-1}$ and the specific surface from 1,000 to 1,600 $\text{m}^2 \text{g}^{-1}$ (ref. 20), with a gas adsorption performance for CO_2 that is higher than the ‘perfect’ defect-free crystal. Vermoortele *et al.* showed that post-synthetic removal of the modulators can lead to large concentrations of coordinatively unsaturated metal sites, and drastically increased activity for several Lewis acid catalysed reactions²¹.

However, the missing linkers also affect the mechanical properties and flexibility of UiO-66 (Fig. 2). Decreasing the number of bidentate linkers from an average of twelve (in the ‘defect-free’ structure) to eight was observed to result in a decrease in bulk, elastic and shear moduli. Anisotropy was also observed to increase upon increasing defect concentrations, whereas, curiously, auxeticity was also introduced at these higher-defect concentrations²². The

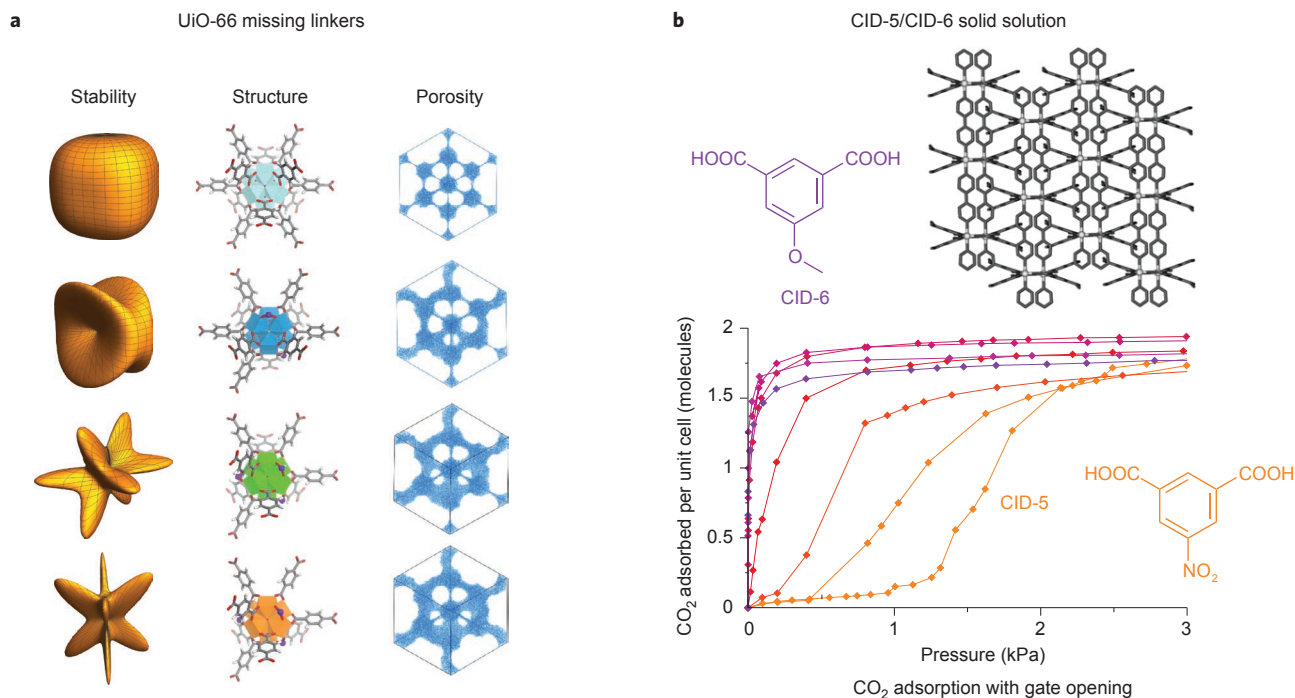


Figure 2 | Interplay between defects and flexibility in metal-organic frameworks. **a**, Defect engineering in the UiO-66(Zr) family of metal-organic frameworks is illustrated by plotting the mechanical stability (represented here by directional Young's modulus; leftmost column, in orange) and accessible porosity (rightmost column, in blue) of several defective structures with an increasing number (from top to bottom) of missing linkers. Figure adapted from ref. 22; RSC. **b**, Tuning of the gate-opening structural transition upon CO_2 adsorption in a CID-5/CID-6 solid solution. The CO_2 adsorption isotherms at 195 K are plotted for CID-5 (orange), CID-6 (purple) and their solid solutions (intermediate colours). The interdigitated structure of the solid solution is shown in the inset (with 48% of CID-5 linker and 52% CID-6 linker). Figure adapted from ref. 24; John Wiley and Sons.

effect on CO₂ uptake of such defects was also studied, with high pressures eliciting increased capacities in the defective structures.

Another example of the intimate interplay between flexibility and defects is based on the use of linker substitution (as opposed to linker removal) to create multivariate MOFs — that is, frameworks encompassing several distinct functionalities by the incorporation of different chemical groups on their organic linkers^{23,24} — through mixing functionalized linkers based on the same backbone^{25,26}. A series of mixed-ligand compounds, starting from the interdigitated layered structures of {Zn(5-NO₂-ip)(bpy)}(0.5DMF·0.5MeOH) and {Zn(5-MeO-ip)(bpy)}(0.5DMF·0.5MeOH) were prepared by Kitagawa *et al.* (ip = isophthalate, bpy = 4,4'-bipyridyl). The flexible gate-opening behaviour (the transition between a closed and open form) of the former compound upon H₂O and CO₂ adsorption was found to be substantially modified through solid solution formation (Fig. 2). Selective adsorption uptakes of CO₂ from a CO₂/CH₄ mixture were investigated, revealing selectivities in the mixed-linker frameworks that overcame the low overall adsorption of the pure 5-NO₂-ip linker system, and the poor separation abilities of the pure 5-MeO-ip linker-containing system.

Post-synthetic modification offers perhaps a more systematic approach to linker substitution, such as the use of solvent-assisted linker exchange (SALE) in Zn- and Cd-based zeolitic imidazolate frameworks (ZIFs)^{27,28}. Here, Hupp and co-workers demonstrated that by exchanging linkers in a relatively simple framework, one can introduce other chemical functions in a manner that is controlled by the exchange kinetics. This process can even be used to synthesize frameworks that were previously unobtainable by direct synthesis methods, because they would crystallize in another (undesirable) topology. Several examples of the use of SALE in flexible, low-density ZIF structures exist²⁹, and the resulting materials show a broad variety of thermal and mechanical behaviour and stability³⁰.

Flexibility and disorder

There are several examples of temperature-induced disorder being used to create flexible materials, showing how entropy intertwines flexibility and disorder in MOFs. Resonant ultrasound spectroscopy was used to probe phase transitions in the formate MOF of composition [NH₄][Zn(HCOO)₃]. It was found that the low-temperature ferroelectric structure, which adopts the polar hexagonal space group P6₃, underwent a transition to a paraelectric phase upon heating over 192 K. The high-temperature framework exists in the non-polar hexagonal space group P6₃/22, and the transition is associated with significant elastic softening due to dynamical disordering of the NH₄⁺ cations within framework channels (Fig. 3)³¹.

A further example of engineering disorder in systems and using it to create flexibility can be found in the UiO framework family. In UiO-67, a planar biphenyl carboxylate ligand connects two Zr₆O₄(OH)₄ nodes, and molecular-dynamics simulations show good agreement between simulated and crystallographic structures³². However, replacement of the linker with azobenzene-4,4'-dicarboxylate, in which the two central nitrogen atoms possess occupational disorder, causes significant dynamic disorder in which the ligands bow in and out of the plane. Consequently, the azobenzene derivative possesses significantly greater flexibility, which is reflected in a significantly lower elastic modulus.

On the other hand, flexibility can also be used to induce disorder. MOFs possess limited mechanical stability, which results in low resistances to structural collapse, or amorphization. Such instabilities have been evidenced under various conditions: hydrostatic compression — with both penetrating and non-penetrating pressure-transmitting fluids, uniaxial compression and ball milling. The microscopic picture of this mechanical instability — established through quantum-chemistry calculations as well as experimental phonon measurements — shows that pressure-induced amorphization is linked to the existence of deformation modes of low elastic

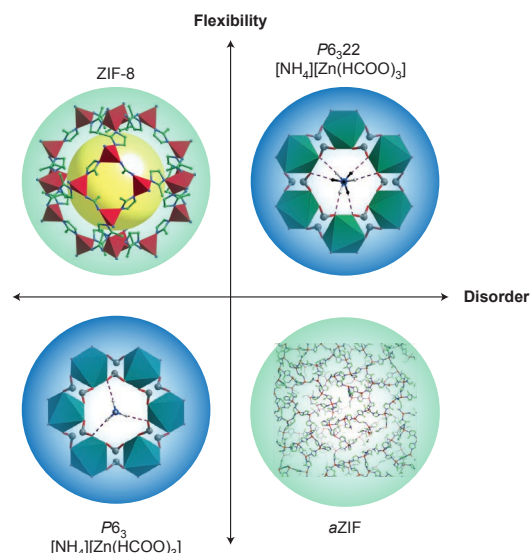


Figure 3 | The relationships between flexibility and disorder in metal-organic frameworks. In green circles: crystalline ZIF-8 (top left), which exhibits local flexibility of its linkers and shear instabilities that lead to amorphization upon application of pressure, forming an amorphous ZIF (aZIF, bottom right), which has a disordered network and no local flexibility. In blue circles: the ordered polar (bottom left) and disordered nonpolar (top right) systems of a zinc formate framework, [NH₄][Zn(HCOO)₃], whose ferroelectric–paraelectric transition is associated with significant elastic softening due to dynamical disordering of the NH₄⁺ cations within framework channels. Figure adapted from ref. 31; RSC.

modulus and their pressure-induced softening (Fig. 3)³⁰. This mechanism has now been confirmed on several MOFs of varying chemical composition and topology. In most cases, the pressure-sensitive soft modes are shear modes, which explain why some MOFs can withstand significant hydrostatic pressure yet are unstable upon solvent evacuation (that is, the first generation MOFs)³³. This key finding was first established experimentally on ZIF-8 [Zn(C₄H₅N₂)₂], whose shear modulus (slightly below 1 GPa) is one order of magnitude lower than the bulk modulus (8 GPa)³⁴, but it is now understood to be a common feature in MOFs.

Defects and disorder

The highly ordered structures of MOFs are often ascribed to their coordination-based self-assembly and highly directional bonding³⁵. For example, Su *et al.* demonstrated that formation of the MIL-53 (Al) framework [Al(OH)(BDC)] proceeded via an initial assembly of reactants into MOF nanoparticles of ~10 nm diameter, before rapid crystallization kinetics led to subsequent aggregation and continued coordination of these clusters into extended MOF particles of ~200 nm in size⁹. Disruption of this secondary long-range aggregate forming step was observed to cause defective coordination in the end product and yielded disordered particle aggregates, or metal–organic gels (Fig. 4). In the example of [Cu₂(ndc)₂(dabco)], defects were introduced in the aggregation stage by using synthetic modulators such as acetic acid to provide competing, connectivity-terminating interactions³⁶. Fe³⁺ and Cr³⁺ dicarboxylate aerogels also formed by introducing defects at a key stage in the formation process, and were described by James *et al.* as a “coherent, spongy network of continuous particles”³⁷. Such work provides pathways to low-density, disordered aggregates of MOF particles and the associated inclusion of hierarchical porosity³⁸.

Instances of defect-induced disorder can also be located within the melting behaviour associated with crystalline MOFs or

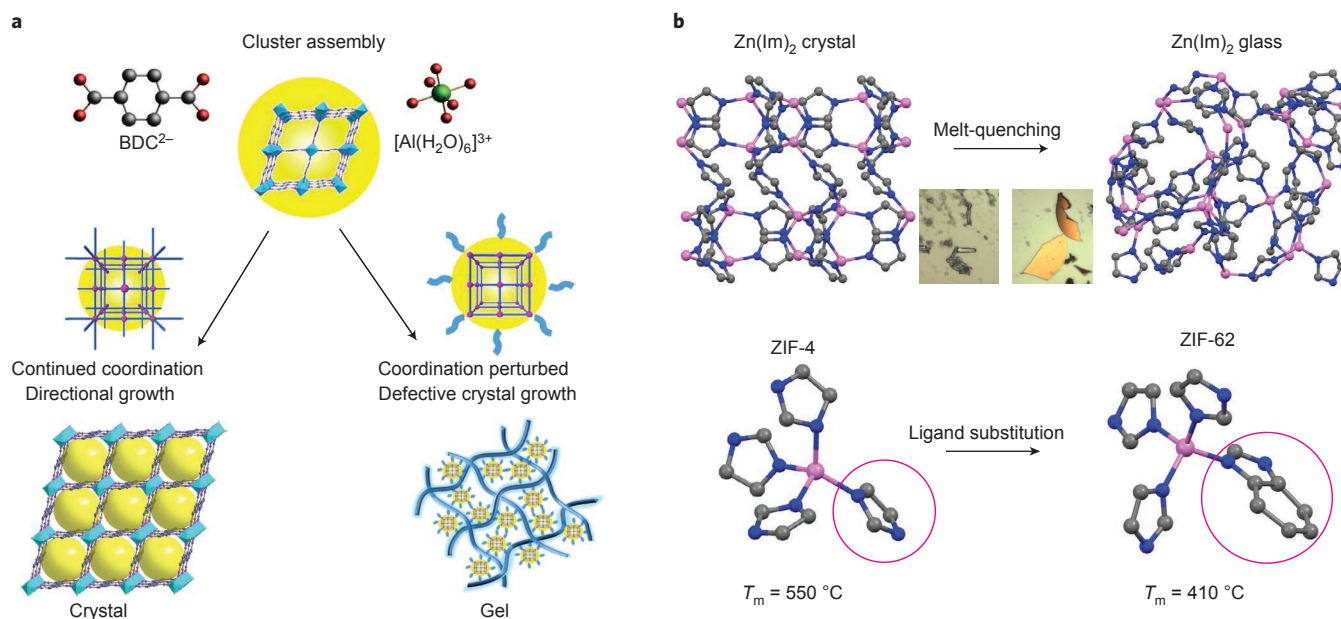


Figure 4 | Coupling between defects and disorder in metal-organic frameworks. **a**, Introduction of mechanistic defects by perturbing coordination during the formation of MIL-53(Al) leads to the formation of a gel based on MOF nanoparticles instead of the crystalline product obtained by unperturbed synthesis. Removal of the solvents via subcritical CO_2 extraction then yields a porous metal-organic aerogel with hierarchical porosity. Adapted from ref. 9; Macmillan. **b**, Defect introduction into ZIFs leads to a lowering in melting point. Partial substitution of the imidazolate linker of ZIF-4 by benzimidazolate (forming ZIF-62) lowers the melting point of the material from 550°C to 410°C , allowing the formation of novel MOF glasses by melt-quenching. Adapted from ref. 16; ACS.

coordination polymers^{12,16,39–41}. In one example, heating a ZIF of composition $[\text{Zn}(\text{C}_3\text{H}_3\text{N}_2)_2]$ to $\sim 550^\circ\text{C}$ resulted in formation of a macroscopic liquid state. Cooling of the resultant liquid resulted in a glass of identical composition, composed of ZnN_4 tetrahedra linked by imidazolate ligands (that is, the same bonding motif as in the starting crystalline state)¹⁶. A mixed-ligand variant was subsequently prepared by addition of dopant quantities of a secondary linker during the initial synthesis, and the resultant, topologically identical material (ZIF-62, $[\text{Zn}(\text{Im})_{1.75}(\text{bIm})_{0.25}]$) was observed to melt at a strikingly lower temperature, of $\sim 400^\circ\text{C}$ (ref. 16).

Linker substitution is not the only example of defects being used to influence disorder in hybrid glasses. The addition of network-modifier ions (that is, the creation of network defects) to lower the melting point of silicates is an established technique in the glass domain⁴². A related route has already been employed by Kato and Kasuga⁴³ in the pursuit of proton-conductive hybrid glasses. Here, taking advantage of the known transfer of protons between imidazolate-type ions⁴⁴, a zinc phosphate glass was heated with a benzimidazolate additive to 160°C , resulting in the formation of a hybrid, proton-conducting material. The building in of defects such as lanthanide or transition metal ions into crystalline structures prior to melting — and subsequently carrying it through into the glass phase — would appear to be a viable and attractive route to functional disordered materials.

Given the propensity for defects in other MOF families, the strategy of utilizing defects to induce melting behaviour may hence be applicable to other MOF systems, aside from ZIFs. Alternatively, the hot-pressing method utilized by Horike *et al.* to form a glass from $[\text{Cd}(\text{H}_2\text{PO}_4)_2(1,2,4\text{-triazole})_2]$ could also be employed for glass formation⁴⁵.

Connecting flexibility, defects and disorder

The occurrence of flexibility, defects and disorder in MOFs are inevitably interconnected, though the intricacies of the relationship are only starting to emerge and all aspects are not yet fully understood.

Although the chemical advantages of combining inorganic and organic components in one system are well established, the physical consequences of this duality are often believed to be detrimental. However, this unique fusion allows flexibilities and disorder — more commonly associated with organic polymers — to be linked with defects reminiscent of classical oxide chemistry. We illustrate here some examples of the complexity arising from the presence of flexibility, defects and disorder.

Although bimetallic MOFs have been reported early in the advancement of MOF research, more complex heterometallic MOFs containing larger numbers of cations are now starting to appear, such as Yaghi's family of mixed-metal MOF-74, which have up to ten different divalent metal cations (Mg, Ca, Sr, Ba, Mn, Fe, Co, Ni, Zn and Cd) incorporated in their structure⁴⁶. These heterometallic MOFs can have an impact on the performance of the material due to the addition of the functions of metal-based functionalities or through synergistic effects of the heterometals. There is probably a strong effect of the presence of ordering, or disorder, in the metals' spatial distribution, although this has been barely studied so far.

Increases in performance can also be elicited from homometallic materials. For example, MIL-47(v) can exhibit remarkably different breathing behaviours (that is, unit-cell volume change), depending on the oxidation state of the metal centre. The structure, in which terephthalate ligands bridge corner sharing VO_6 octahedra to yield interconnected chains, was previously demonstrated to exist in a rigid open form, regardless of solvent or guest content. This is very different to other related structures, for example, MIL-53, where adsorption of solvent elicits a shrinking of the unit cell⁴⁷. However, careful evacuation of the as-synthesized MIL-47 material appeared to prevent oxidation of V^{III} centres to V^{IV} , yielding an evacuated material of composition $\text{V}^{\text{III}}(\text{OH})(\text{bdc})$. Breathing behaviour in this material was observed due to solvent interactions with OH groups on metal centres. An interesting extension on the work was the preparation of mixed-valence MIL-47 derivatives by partial

oxidation of $V^{III}(OH)(bdc)$. Conductivity measurements performed upon samples showed that although single-oxidation-state derivatives were very poorly conducting, mixed-oxidation-state samples demonstrated appreciably greater conductivities. In this case, differences were ascribed to the homogeneous distribution of oxidation site defects throughout the material, though the possibility of domain formation was not investigated further⁴⁷.

There is now a growing realization that, in addition to ligand and metal substitutions, other types of defects can exist in MOFs, and in particular UiO-66 (Hf). Metal cluster vacancies in the material were shown to be present by Goodwin *et al.* and controlled by altering the quantity of synthetic modulator. However, it was also demonstrated that the structural disorder introduced by such vacancies was correlated, with defective sites effectively dispersed unevenly into nanoregions of 7 nm in size within a non-defective framework matrix⁴⁸. The existence of these correlated defects had a significant impact upon the elastic, shear and bulk moduli of the material, with decreases of ~50% noted in each case. Such increases in flexibility raise the prospect of tuning the materials' performance in charge transport, catalysis and mechanical and optical responses, and will also necessitate the development of tools — both experimental and theoretical — to investigate the correlations between defects and their spatial distribution across the MOF family.

These phenomena of defects, disorder and flexibility in MOFs present significant challenges. It has often been the case that they are systematically compared to other porous inorganic or organic systems. A hitherto prevailing thought has been that their principal advantages (chemical versatility and pore size/tunability) are severely tempered by structural fragility, which manifests itself in flexibility, defects and disorder. A wealth of studies are now emerging that look to seize upon these assumed weaknesses and utilize them for new applications and directions in the field, rather than try and ameliorate or 'solve' the structural fragilities that are now recognized as quintessentially characteristic of MOFs. It is worth noting that a very similar approach was pioneered by de Gennes, in coining the term fragile materials, and thus launching a new field of exploration into soft polymeric matter. Areas where these design principles might be expected to play a particularly beneficial role in the future include proton and ion conductors⁴⁹, electrical semiconductors⁵⁰ and tuneable optical systems.

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References

- Schneermann, A. *et al.* Flexible metal–organic frameworks. *Chem. Soc. Rev.* **43**, 6062–6096 (2014).
- Dubbeldam, D., Walton, K. S., Ellis, D. E. & Snurr, R. Q. Exceptional negative thermal expansion in isoreticular metal–organic frameworks. *Angew. Chem. Int. Ed.* **46**, 4496–4499 (2007).
- McKellar, S. C. & Moggach, S. A. Structural studies of metal–organic frameworks under high pressure. *Acta Cryst. B* **71**, 587–607 (2015).
- Li, W. *et al.* Negative linear compressibility of a metal–organic framework. *J. Am. Chem. Soc.* **134**, 11940–11943 (2012).
- Cairns, A. B. *et al.* Giant negative linear compressibility in zinc dicyanoaurate. *Nat. Mater.* **12**, 212–216 (2013).
- Kaye, S. S., Dailly, A., Yaghi, O. M. & Long, J. R. Impact of preparation and handling on the hydrogen storage properties of $Zn_4O(1,4\text{-benzenedicarboxylate})_3$ (MOF-5). *J. Am. Chem. Soc.* **129**, 14176–14177 (2007).
- Cooper, A. I. & Rosseinsky, M. J. Metal–organic frameworks improving pore performance. *Nat. Chem.* **1**, 26–27 (2009).
- Fang, Z. L., Bueken, B., De Vos, D. E. & Fischer, R. A. Defect-engineered metal–organic frameworks. *Angew. Chem. Int. Ed.* **54**, 7234–7254 (2015).
- Li, L. *et al.* A synthetic route to ultralight hierarchically micro/mesoporous Al(III)-carboxylate metal–organic aerogels. *Nat. Commun.* **4**, 1774 (2013).
- Sholl, D. S. & Lively, R. P. Defects in metal–organic frameworks: challenge or opportunity? *J. Phys. Chem. Lett.* **6**, 3437–3444 (2015).
- Hoskins, B. F. & Robson, R. Design and construction of a new class of scaffolding-like materials comprising infinite polymeric frameworks of 3D-linked molecular rods. A reappraisal of the $Zn(CN)_2$ and $Cd(CN)_2$ structures and the synthesis and structure of the diamond-related frameworks $[N(CH_3)_4][Cu^{II}Zn^{II}(CN)_4]$ and $Cu^{II}[4, 4', 4'', 4''']\text{-tetracyanotetraphenylmethane} BF_4 \cdot xC_6H_5NO_2$. *J. Am. Chem. Soc.* **112**, 1546–1554 (1990).
- Bennett, T. D. *et al.* Hybrid glasses from strong and fragile metal–organic framework liquids. *Nat. Commun.* **6**, 8079 (2015).
- Cairns, A. B. & Goodwin, A. L. Structural disorder in molecular framework materials. *Chem. Soc. Rev.* **42**, 4881–4893 (2013).
- Ohrstrom, L. Let's talk about MOFs — topology and terminology of metal–organic frameworks and why we need them. *Crystals* **5**, 154–162 (2015).
- Loiseau, T. *et al.* A rationale for the large breathing of the porous aluminum terephthalate (MIL-53) upon hydration. *Chem.-Eur. J.* **10**, 1373–1382 (2004).
- Bennett, T. D. *et al.* Melt-quenched glasses of metal–organic frameworks. *J. Am. Chem. Soc.* **138**, 3484–3492 (2016).
- Horike, S., Shimomura, S. & Kitagawa, S. Soft porous crystals. *Nat. Chem.* **1**, 695–704 (2009).
- Wu, H., Yildirim, T. & Zhou, W. Exceptional mechanical stability of highly porous zirconium metal–organic framework UiO-66 and its important implications. *J. Phys. Chem. Lett.* **4**, 925–930 (2013).
- Valenzano, L. *et al.* Disclosing the complex structure of UiO-66 metal–organic framework: a synergic combination of experiment and theory. *Chem. Mater.* **23**, 1700–1718 (2011).
- Wu, H. *et al.* Unusual and highly tunable missing-linker defects in zirconium metal–organic framework UiO-66 and their important effects on gas adsorption. *J. Am. Chem. Soc.* **135**, 10525–10532 (2013).
- Vermoortele, F. *et al.* Synthesis modulation as a tool to increase the catalytic activity of metal–organic frameworks: the unique case of UiO-66(Zr). *J. Am. Chem. Soc.* **135**, 11465–11468 (2013).
- Thornton, A. W., Babarao, R., Jain, A., Trouselet, F. & Coudert, F.-X. Defects in metal–organic frameworks: a compromise between adsorption and stability? *Dalton Trans.* **45**, 4352–4359 (2016).
- Deng, H. X. *et al.* Multiple functional groups of varying ratios in metal–organic frameworks. *Science* **327**, 846–850 (2010).
- Fukushima, T. *et al.* Solid solutions of soft porous coordination polymers: fine-tuning of gas adsorption properties. *Angew. Chem. Int. Ed.* **49**, 4820–4824 (2010).
- Kleist, W., Jutz, F., Maciejewski, M. & Baiker, A. Mixed-linker metal–organic frameworks as catalysts for the synthesis of propylene carbonate from propylene oxide and CO_2 . *Eur. J. Inorg. Chem.* **24**, 3552–3561 (2009).
- Cohen, S. M. Postsynthetic methods for the functionalization of metal–organic frameworks. *Chem. Rev.* **112**, 970–1000 (2012).
- Karagiari, O. *et al.* Opening ZIF-8: a catalytically active zeolitic imidazolate framework of sodalite topology with unsubstituted linkers. *J. Am. Chem. Soc.* **134**, 18790–18796 (2012).
- Deria, P. *et al.* Beyond post-synthesis modification: evolution of metal–organic frameworks via building block replacement. *Chem. Soc. Rev.* **43**, 5896–5912 (2014).
- Karagiari, O., Bury, W., Mondloch, J. E., Hupp, J. T. & Farha, O. K. Solvent-assisted linker exchange: an alternative to the *de novo* synthesis of unattainable metal–organic frameworks. *Angew. Chem. Int. Ed.* **53**, 4530–4540 (2014).
- Bouéssel du Bourg, L., Ortiz, A. U., Boutin, A. & Coudert, F.-X. Thermal and mechanical stability of zeolitic imidazolate frameworks polymorphs. *APL Mater.* **2**, 124110 (2014).
- Zhang, Z. Y., Li, W., Carpenter, M. A., Howard, C. J. & Cheetham, A. K. Elastic properties and acoustic dissipation associated with a disorder–order ferroelectric transition in a metal–organic framework. *CrystEngComm* **17**, 370–374 (2015).
- Hobday, C. L. *et al.* A computational and experimental approach linking disorder, high-pressure behavior, and mechanical properties in UiO frameworks. *Angew. Chem. Int. Ed.* **55**, 2401–2405 (2016).
- Kitagawa, S. & Kondo, M. Functional micropore chemistry of crystalline metal complex-assembled compounds. *Bull. Chem. Soc. Jpn.* **71**, 1739–1753 (1998).
- Tan, J. C. *et al.* Exceptionally low shear modulus in a prototypical imidazole-based metal–organic framework. *Phys. Rev. Lett.* **108**, 095502 (2012).
- Cook, T. R., Zheng, Y. R. & Stang, P. J. Metal–organic frameworks and self-assembled supramolecular coordination complexes: comparing and contrasting the design, synthesis, and functionality of metal–organic materials. *Chem. Rev.* **113**, 734–777 (2013).
- Tsuruoka, T. *et al.* Nanoporous nanorods fabricated by coordination modulation and oriented attachment growth. *Angew. Chem. Int. Ed.* **48**, 4739–4743 (2009).
- Xiang, S. *et al.* Porous organic–inorganic hybrid aerogels based on Cr^{3+}/Fe^{3+} and rigid bridging carboxylates. *J. Mater. Chem.* **22**, 1862–1867 (2012).
- Lohe, M. R., Rose, M. & Kaskel, S. Metal–organic framework (MOF) aerogels with high micro- and macroporosity. *Chem. Commun.* 6056–6058 (2009).

39. Umeyama, D., Horike, S., Inukai, M., Itakura, T. & Kitagawa, S. Reversible Solid-to-liquid phase transition of coordination polymer crystals. *J. Am. Chem. Soc.* **137**, 864–870 (2015).
40. Umeyama, D. *et al.* Glass formation via structural fragmentation of a 2D coordination network. *Chem. Commun.* **51**, 12728–12731 (2015).
41. Zhao, Y., Lee, S.-Y., Becknell, N., Yaghi, O. M. & Angell, C. A. Nanoporous transparent MOF glasses with accessible internal surface. *J. Am. Chem. Soc.* **138**, 10818–10821 (2016).
42. Shelby, J. E. *Introduction to Glass Science and Technology* 2nd edn (Royal Society of Chemistry, 2005).
43. Kato, H. & Kasuga, T. Preparation of proton-conducting hybrid materials by reacting zinc phosphate glass with benzimidazole. *Mater. Lett.* **79**, 109–111 (2012).
44. Kim, J. & Honma, I. Anhydrous solid state proton conductor based on benzimidazole/monododecyl phosphate molecular hybrids. *Solid State Ionics* **176**, 979–984 (2005).
45. Chen, W. Q. *et al.* Glass formation of a coordination polymer crystal for enhanced proton conductivity and material flexibility. *Angew. Chem. Int. Ed.* **55**, 5195–5200 (2016).
46. Wang, L. J. *et al.* Synthesis and characterization of metal–organic framework-74 containing 2, 4, 6, 8, and 10 different metals. *Inorg. Chem.* **53**, 5881–5883 (2014).
47. Leclerc, H. *et al.* Influence of the oxidation state of the metal center on the flexibility and adsorption properties of a porous metal organic framework: MIL-47(v). *J. Phys. Chem. C* **115**, 19828–19840 (2011).
48. Cliffe, M. J. *et al.* Correlated defect nanoregions in a metal–organic framework. *Nature Commun.* **5**, 4176 (2014).
49. Taylor, J. M., Dekura, S., Ikeda, R. & Kitagawa, H. Defect control to enhance proton conductivity in a metal–organic framework. *Chem. Mater.* **27**, 2286–2289 (2015).
50. Tominaka, S. *et al.* Topochemical conversion of a dense metal–organic framework from a crystalline insulator to an amorphous semiconductor. *Chem. Sci.* **6**, 1465–1473 (2015).

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Author contributions

All authors initiated this discussion and designed the paper, T.D.B. and F.-X.C. wrote the manuscript, all authors revised it.

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Competing financial interests

The authors declare no competing financial interests.