

Guest control of structure in porous organic cages†

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Two porous organic cages with different thermodynamic polymorphs were induced by co-solvents to interchange their crystal packing modes, thus achieving guest-mediated control over solid-state porosity. *In situ* crystallography allows the effect of the co-solvent guests on these structural interconversions to be understood.

Porous molecular solids¹ comprise non-covalent intermolecular interactions that are weaker and often less directional than the coordination or covalent bonds that define zeolites,² metal–organic frameworks (MOFs),³ and covalent organic frameworks (COFs).⁴ Structural polymorphism is therefore common, and this raises challenges for the targeted design of solid-state function.

We have developed porous organic cage molecules *via* [4+6] cycloimination reactions of 1,3,5-triformylbenzene (TFB) with vicinal aliphatic diamines such as (1*R*,2*R*)-diaminocyclohexane (CC3-*R*) and (1*R*,2*R*)-diaminocyclopentane (CC4-*R*) (Fig. 1a).⁵ The molecular solubility of organic cages⁶ allows crystallisation to be decoupled from synthesis, unlike MOFs and COFs where these two processes constitute the same step.

CC3-*R* prefers to pack in a window-to-window arrangement,^{5,7} which generates an interconnected diamondoid pore network (Fig. 1b). This thermodynamic polymorph, CC3 α , has an apparent BET surface area (S_{BET}) of $\sim 410 \text{ m}^2 \text{ g}^{-1}$ in the crystalline state.⁷ Both crystal structure predictions (CSP)⁸ and DFT ‘cage dimer’ calculations⁷ rationalise this crystal packing. By contrast, CC4-*R* packs *via* a window-to-arene interaction, despite its close structural similarity with CC3-*R*, and this results in a 2-D pore structure in its thermodynamic polymorph, CC4 α , rather than a 3-D diamondoid pore structure (Fig. 1c).

In this study, we show that these two thermodynamic ‘alpha’ crystal packing modes can be interchanged by using specific co-solvents to direct the crystal packing, such that CC3-*R* packs

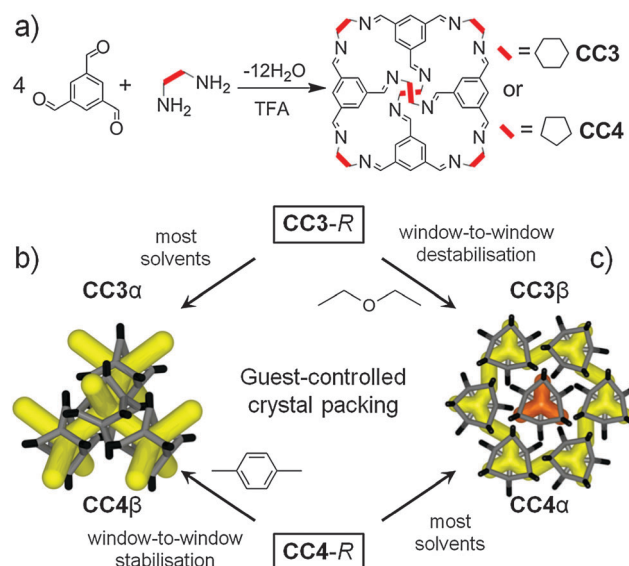


Fig. 1 (a) Synthesis of organic cage molecules **CC3** and **CC4**. Scheme showing interchangeable crystal packings for cages **CC3** and **CC4**, (b) 3-D diamondoid pore channels, and (c) 2-D layered pore structure with formally disconnected voids. Orange = disconnected voids; yellow = interconnected pores.

like CC4-*R*, and *vice versa* (Fig. 1b and c). We showed recently that 1,4-dioxane can direct three different tetrahedral cage molecules (CC1, CC2 and CC13)⁹ to crystallise isostructurally with CC3 α , providing that the cages comprise a racemic mixture.

Here we expand this concept to the homochiral cage molecules CC3-*R* and CC4-*R*. We direct these two cages into isostructural packings, and hence achieve control over pore topology, pore volume, and surface area. Computational CSP can map the energy landscape for solvated crystal structures¹¹ or for solvent-free organic cage crystals,^{8,10} but this is not yet computationally affordable for cage solvates. Our strategy, therefore, was to combine intuitive concepts of shape-direction,⁹ solvent screening, and CSP calculations for solvent-free crystals.¹⁰

CC3-*R* is readily soluble in CH_2Cl_2 but it is insoluble in Et_2O . However, the addition of excess of an antisolvent, Et_2O , to solutions

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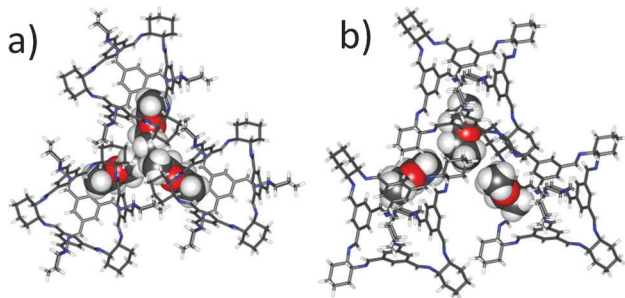


Fig. 2 Single crystal structure for **CC3-R**·(Et₂O)₃·CH₂Cl₂, viewed parallel to [001] axis (a) and off axis (b). Et₂O solvent molecules are shown in space filling representation, with the CH₂Cl₂ molecules in the cage cavities omitted for clarity.

of **CC3-R** in CH₂Cl₂ did not result in direct precipitation of the cage, even when a twentyfold volumetric excess of Et₂O was added. Instead, slow evaporation of the resultant homogenous solution affords hexagonal, needle-shaped single crystals of (**CC3-R**)·(Et₂O)₃·CH₂Cl₂ (Fig. S1, ESI[†]), which crystallise in the trigonal space group *R*3̄. The asymmetric unit comprises one third of a **CC3-R** molecule positioned on a threefold rotation axis, plus one CH₂Cl₂ molecule in the intrinsic cage cavity and one well-ordered Et₂O solvent molecule in the window site (Fig. 2). Overall, Et₂O occupies three of the four cage window sites, with one hydrophobic methyl terminus directed toward the hydrophobic cage cavity, sharing the cage void with CH₂Cl₂.

The inclusion of Et₂O during crystallisation disrupts the otherwise favourable window-to-window **CC3-R** cage pairing, as found in **CC3α**, which has a cage–cage intermolecular interaction energy calculated to be around 150 kJ mol^{−1} (DFT-D3).⁷ Hence, the three cage windows pack in a relatively inefficient manner, forming three extrinsic cavities into which the methyl termini of the Et₂O solvent molecules are directed (Fig. 2). The fourth cage window is occupied by the aromatic face of a neighbouring cage: this window-to-face interaction has a binding energy that was calculated to be 55 kJ mol^{−1} less favourable than the window-to-window interaction,⁷ but this is evidently compensated by favourable cage–solvent interactions.

Stacks of **CC3-R** molecules are formed as a result of this window-to-face packing (Fig. 3). This packing is very similar to that reported previously for crystalline **CC4-R**.¹² For **CC4-R**, we observed a de-symmetrisation upon desolvation (*R*3̄ to *P*3̄) as a result of the **CC4-R** molecules undergoing a screw-type rotation to afford what is referred to hereafter as **CC4α** (Fig. 1c). Similarly, desolvation of (**CC3-R**)·(Et₂O)₃·CH₂Cl₂ affords a new solvent-free, metastable polymorph of **CC3-R**, **CC3β**. This is also accompanied by screw-type rotation of the cage molecules in the crystal lattice (Fig. 3; Section 1.3, Fig. S2–S11, ESI[†]). Single-crystal-to-single-crystal transformations have been reported for discrete host molecules such as metallocycles,¹³ and also for extended frameworks,¹⁴ but preservation of single crystallinity is not typically observed when there is a substantial structural rearrangement. For **CC3**, loss of the guest solvent causes the cages to pack in a more frustrated arrangement in **CC3β**, causing a significant contraction of the unit cell volume (5%).

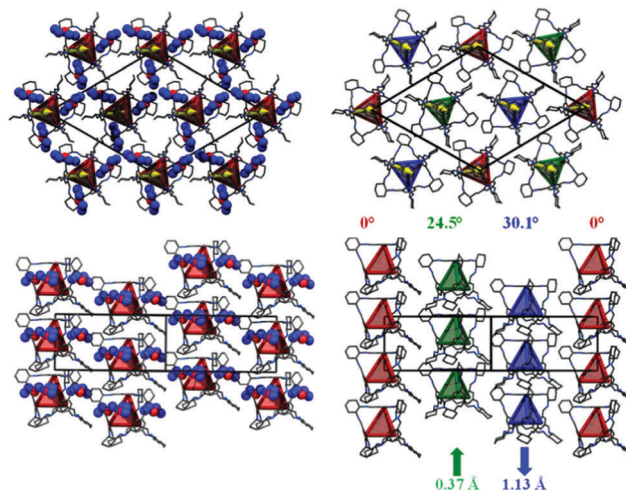


Fig. 3 Single crystal structure for **CC3-R**·(Et₂O)₃·CH₂Cl₂ collected at 100 K (left), viewed along [001] (above), along the direction of the cage stacks, and [110] (below), perpendicular to the cage stacks. Et₂O solvent molecules shown as space filling representation, carbon atoms in blue. Intra-cavity CH₂Cl₂ solvent molecules and hydrogen atoms are omitted for clarity. Red tetrahedra represent the connectivity between four aromatic faces of the crystallographically equivalent **CC3-R** molecules. Single crystal structure for desolvated **CC3β** collected at 300 K (right). Red, green and blue tetrahedra represent the connectivity between the aromatic faces of the three crystallographically distinct **CC3-R** molecules. Magnitude of rotation and translation of the cage molecules upon desolvation are indicated.

For **CC3β**, the angle of cage rotation upon desolvation was defined as the angle rotated relative to the red cage (Fig. 3), which remains static and which, for convenience, has been located on the cell origin. For **CC3**, these rotation angles were determined to be 24.5° and 30.1°, respectively, for the green and blue cages shown in Fig. 3. There is also a 0.37 Å and 1.13 Å shift, respectively, along the *z*-direction for the green and blue cages in a convergent manner. Unlike the thermodynamic desolvated polymorph **CC3α**, **CC3β** does not display any window-to-window packing of the cage molecules. **CC3α** is predicted to be the lowest energy solvent-free form of **CC3-R**,⁸ and it is the most prevalent experimentally, being obtained from most solvents tested. **CC3α** also exhibits the most efficient packing between cages and has the highest density. We suggest, therefore, that **CC3β** is a kinetically trapped polymorph, as supported by the occasional observation of the β-form when **CC3-R** is rapidly precipitated from solvent by rotary evaporation.

Gas sorption analysis shows that **CC3β** is porous to nitrogen, with an apparent *S*_{BET} of 555 m² g^{−1}, and a microporous type I adsorption isotherm (Fig. 4). The nitrogen uptake and porosity for **CC3β** is higher than for **CC3α**, which can be rationalised by its lower density (0.922 g cm^{−3} for **CC3β** vs. 0.973 g cm^{−3} for **CC3α**). Hence, **CC3β** has extrinsic lattice sites that can accommodate additional N₂ molecules. The **CC3β** polymorph is also porous to hydrogen and carbon dioxide, in both cases with a higher gas uptake than was found for the original **CC3α** polymorph (Fig. S12 and S13, ESI[†]). Even though the crystal packings for **CC4α** and **CC3β** are essentially isostructural, the gas sorption isotherms are strikingly different. Gas-induced transformations



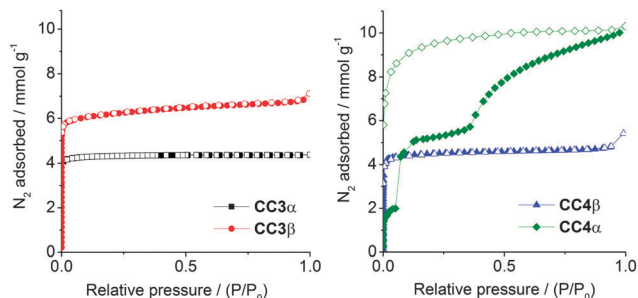


Fig. 4 Nitrogen sorption isotherms, at 77 K, for **CC3** α and β (left), and **CC4** α and β (right). Adsorption isotherm branches are shown as solid symbols, and desorption branches as open symbols.

are known for organic solids.¹⁵ For **CC4** α , there are low-pressure adsorption steps¹² that are not observed for **CC3** β . Also, **CC4** α also adsorbs significantly more N_2 than **CC3** β at higher relative pressures (Fig. 4). *In situ* powder XRD data indicates a sharp, low-pressure structural transition (Section 1.5, Fig. S14–S17, ESI†). Analogous *in situ* powder XRD data for **CC3** β did not show any low-pressure transition (Fig. S18, ESI†). The observation of a low-pressure structural transition for **CC4** α , but not for **CC3** β , might be due to the smaller cycloalkane vertex functionality in **CC4**. The more compact cyclopentyl vertices in **CC4-R** might allow small rotational rearrangements of the cages at low N_2 pressures, enabling the guest to gain more extensive access to the pore network at higher N_2 pressures. By contrast, we speculate that the analogous rearrangements are hindered in **CC3** β by the bulkier cyclohexyl substituents, and a hence a classic type I isotherm is observed with no structural transition steps (Fig. 4).

Having directed **CC3-R** to pack in the typical **CC4-R** fashion (Fig. 1c), the reverse challenge was to direct **CC4-R** to pack isostructurally with **CC3** α (Fig. 1b). A recent computational study predicted a polymorph, **CC4** β , as the global lattice energy minimum for the homochiral **CC4-R**.¹⁰ **CC4** β is isostructural with **CC3** α , suggesting initially that the same window-to-window cage pairing is also thermodynamically preferred for **CC4-R**. However, for **CC4-R**, the rigid-molecule constraint used in the CSP limits the reliability of the structure searches. Specifically, the experimental **CC4** α structure involves close intermolecular contacts that distort the molecular geometry, and hence geometry-constrained CSP calculations lead to artificially high lattice energies for **CC4** α . This was resolved by periodic DFT-D calculations, which showed the known **CC4** α polymorph to have a formation energy that is 8.19 kJ mol⁻¹ more stable than **CC4** β when molecular flexibility is taken into account. In the context of lattice energies, 8.19 kJ mol⁻¹ is a relatively small difference, suggesting that **CC4-R** might be directed to pack as **CC4** β by inclusion of an appropriate directing solvent.

A screen of 30 different crystallisation co-solvents was used in an attempt to access the **CC4** β polymorph that was suggested by CSP (Section 1.6, Table S4, ESI†). PXRD data showed that only one of the 30 co-solvents tested, *para*-xylene, produced a crystalline form that was distinct from **CC4** α (Fig. S19, ESI†), again supporting the conclusion that **CC4** α is the thermodynamically

most stable polymorph, as predicted computationally using DFT-D.¹⁰ We showed recently that *para*-xylene can occupy the interstitial site between two cage windows in the crystal lattice of **CC3-R**,¹⁶ and also that it can bridge adjacent windows in 1-D cage catenane chains.¹⁷ Here, *para*-xylene ‘pegs’ adjacent **CC4-R** cage windows together in a similar fashion. Single crystals of this new **CC4-R** phase were grown from a layered CH_2Cl_2 -*para*-xylene solution (see Section 1.7, ESI†). Single crystal X-ray diffraction revealed a new phase, **CC4-R**·(C_8H_{10})₃·(H_2O)₂ had crystallised in the chiral orthorhombic space group $P2_12_12_1$, where each of the four cage windows are penetrated by the methyl terminus of a *para*-xylene solvent molecule (Fig. 5a). The linear shape of the *para*-xylene molecule would therefore seem to be important, as observed for 1,4-dioxane in our previous study.⁹ This is supported by the fact that the structural isomers of this solvent, *meta*- and *ortho*-xylene (Table S4, ESI†), did not produce the **CC4** β polymorph (*cf.*, 1,3-dioxane in our previous work⁹).

In this structure, four *para*-xylene molecules are shared equally between two adjacent **CC4-R** molecules. Extending this window pairing arrangement in three dimensions generates a diamondoid network filled with *para*-xylene solvent molecules (Fig. S20 and S21, ESI†), reminiscent of the known **CC3-R** *para*-xylene solvate.¹⁶ However, unlike the **CC3-R** solvate, one additional *para*-xylene solvent molecule per cage unit is located in extrinsic 1-D channels in the **CC4-R** solvate (Fig. S21, ESI†). As a result of this additional solvent molecule, desolvation to a symmetrical diamondoid pore network requires significant anisotropic contraction of the unit cell parameters, potentially resulting in a transformation to an alternate pore topology or loss of crystallinity. Remarkably, gradual heating of a solvate crystal of **CC4-R**·(C_8H_{10})₃·(H_2O)₂ (Section 1.7 & Fig. S22, ESI†) resulted in preservation of single crystallinity and isolation of **CC4** β (Fig. 5b and c), which was formed after a transformation of the crystal symmetry to the chiral cubic space group, $F4_132$. This structure is isostructural both with **CC3** α and with the computationally predicted structure obtained for **CC4** in CSP studies, as reported previously.¹⁰ This single-crystal-to-single-crystal transformation was accompanied by a large contraction

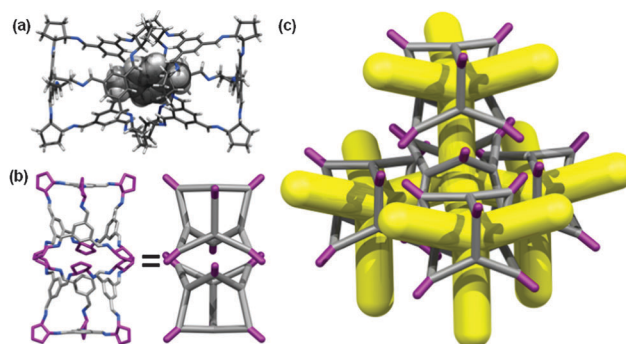


Fig. 5 (a) *para*-Xylene ‘pegged’ **CC4-R** molecules from the single crystal structure **CC4-R**·(C_8H_{10})₃·(H_2O)₂; (b) direct window-to-window pairing arrangement displayed in **CC4** β , generating (c) an interconnected diamondoid pore network upon desolvation of the xylene solvate.



of the cell volume per CC4-R molecule of around 13%. The bulk CC4 β material also remains crystalline after desolvation under heating and vacuum, and the resultant gas sorption properties very similar to those of CC3 α , as expected from the isostructural packing. Indeed, the nitrogen sorption isotherms for CC3 α and CC4 β are almost identical (SA_{BET} for CC3 α = 409 m² g⁻¹; SA_{BET} for CC4 β = 387 m² g⁻¹; Fig. 4). The preservation of single crystallinity upon desolvation is remarkable, given the large anisotropic contraction in the unit cell. The 'pegged' orientation of the CC4-R molecules in the *para*-xylene solvate probably facilitates the preservation of this window-to-window packing after desolvation and prevents the cages from rearranging to form CC4 α . Transformation to CC4 α would involve substantial and presumably high-energy reorganisation from the *para*-xylene solvate in order to generate the window-to-arene packing mode found in the thermodynamic polymorph.

Two organic cages, CC3-R and CC4-R, were induced to interchange their low-energy packing modes by using directing solvents. This shows that polymorphism in porous molecular solids can to some extent be controlled by combining CSP calculations, intuitive design, and high-throughput crystallisation screens. We can identify specific co-solvents that either reinforce (*para*-xylene in CC4 β) or disrupt (Et₂O in CC3 β) the solid-state window-to-window packing arrangement for these porous cages. Without the use of these directing co-solvents, the lowest energy α -polymorph for CC3-R is isostructural with the higher energy β -polymorph of CC4-R, and *vice versa* (Fig. 1b).

Isostructural crystal packings, however, can lead to quite different gas sorption properties: for example, CC4 α has marked steps in its isotherm, whereas CC3 β does not. By contrast, the sorption isotherms for CC3 α and CC4 β are almost identical.

This represents a further step toward controlling the functional properties of porous molecular crystals by design. In the future, the *a priori* prediction of these solvent effects might also be possible, although at present this is prohibited by the computational expense of the relevant CSP methods.

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Notes and references

- 1 J. Tian, P. K. Thallapally and B. P. McGrail, *CrystEngComm*, 2012, **14**, 1909–1919; M. Mastalerz, *Angew. Chem., Int. Ed.*, 2010, **49**, 5042–5053; L. J. Barbour, *Chem. Commun.*, 2006, 1163–1168; J. L. Atwood, L. J. Barbour, A. Jerga and B. L. Schottel, *Science*, 2002, **298**, 1000–1002; G. Zhang and M. Mastalerz, *Chem. Soc. Rev.*, 2014, **43**, 1934–1947.
- 2 C. S. Cundy and P. A. Cox, *Chem. Rev.*, 2003, **103**, 663–702.
- 3 O. M. Yaghi, H. Li, C. Davis, D. Richardson and T. L. Groy, *Acc. Chem. Res.*, 1998, **31**, 474–484; A. K. Cheetham, G. Férey and T. Loiseau, *Angew. Chem., Int. Ed.*, 1999, **38**, 3268–3292; S. Kitagawa, R. Kitaura and S.-i. Noro, *Angew. Chem., Int. Ed.*, 2004, **43**, 2334–2375; D. Bradshaw, J. B. Claridge, E. J. Cussen, T. J. Prior and M. J. Rosseinsky, *Acc. Chem. Res.*, 2005, **38**, 273–282.
- 4 A. P. Côté, A. I. Benin, N. W. Ockwig, M. O'Keeffe, A. J. Matzger and O. M. Yaghi, *Science*, 2005, **310**, 1166–1170.
- 5 T. Tozawa, J. T. A. Jones, S. I. Swamy, S. Jiang, D. J. Adams, S. Shakespeare, R. Clowes, D. Bradshaw, T. Hasell, S. Y. Chong, C. Tang, S. Thompson, J. Parker, A. Trewin, J. Bacsá, A. M. Z. Slawin, A. Steiner and A. I. Cooper, *Nat. Mater.*, 2009, **8**, 973–978.
- 6 M. Mastalerz, M. W. Schneider, I. M. Oppel and O. Presly, *Angew. Chem., Int. Ed.*, 2011, **50**, 1046–1051; M. W. Schneider, I. M. Oppel, A. Griffin and M. Mastalerz, *Angew. Chem., Int. Ed.*, 2013, **52**, 3611–3615; A. Avellaneda, P. Valente, A. Burgun, J. D. Evans, A. W. Markwell-Heys, D. Rankine, D. J. Nielsen, M. R. Hill, C. J. Sumby and C. J. Doonan, *Angew. Chem., Int. Ed.*, 2013, **52**, 3746–3749.
- 7 T. Hasell, S. Y. Chong, K. E. Jelfs, D. J. Adams and A. I. Cooper, *J. Am. Chem. Soc.*, 2012, **134**, 588–598.
- 8 J. T. A. Jones, T. Hasell, X. Wu, J. Bacsá, K. E. Jelfs, M. Schmidtman, S. Y. Chong, D. J. Adams, A. Trewin, F. Schiffman, F. Cora, B. Slater, A. Steiner, G. M. Day and A. I. Cooper, *Nature*, 2011, **474**, 367–371.
- 9 T. Hasell, J. L. Culshaw, S. Y. Chong, M. Schmidtman, M. A. Little, K. E. Jelfs, E. O. Pyzer-Knapp, H. Shepherd, D. J. Adams, G. M. Day and A. I. Cooper, *J. Am. Chem. Soc.*, 2014, **136**, 1438–1448.
- 10 E. O. Pyzer-Knapp, H. P. G. Thompson, F. Schiffmann, K. E. Jelfs, S. Y. Chong, M. A. Little, A. I. Cooper and G. M. Day, *Chem. Sci.*, 2014, **5**, 2235–2245.
- 11 A. J. Cruz-Cabeza, S. Karki, L. Fabian, T. Friscic, G. M. Day and W. Jones, *Chem. Commun.*, 2010, **46**, 2224–2226; A. J. Cruz-Cabeza, G. M. Day and W. Jones, *Phys. Chem. Chem. Phys.*, 2011, **13**, 12808–12816.
- 12 T. Mitra, X. Wu, R. Clowes, J. T. A. Jones, K. E. Jelfs, D. J. Adams, A. Trewin, J. Bacsá, A. Steiner and A. I. Cooper, *Chem. – Eur. J.*, 2011, **17**, 10235–10240.
- 13 L. Dobrzańska, G. O. Lloyd, H. G. Raubenheimer and L. J. Barbour, *J. Am. Chem. Soc.*, 2005, **128**, 698–699; G. O. Lloyd, H. G. Raubenheimer and L. J. Barbour, *J. Am. Chem. Soc.*, 2005, **127**, 13134–13135.
- 14 E. Y. Lee and M. P. Suh, *Angew. Chem., Int. Ed.*, 2004, **43**, 2798–2801; H. J. Choi and M. P. Suh, *J. Am. Chem. Soc.*, 2004, **126**, 15844–15851; S. K. Ghosh, W. Kaneko, D. Kiriya, M. Ohba and S. Kitagawa, *Angew. Chem., Int. Ed.*, 2008, **47**, 8843–8847.
- 15 P. K. Thallapally, B. Peter McGrail, S. J. Dalgarno, H. T. Schaefer, J. Tian and J. L. Atwood, *Nat. Mater.*, 2008, **7**, 146–150.
- 16 T. Mitra, K. E. Jelfs, M. Schmidtman, A. Ahmed, S. Y. Chong, D. J. Adams and A. I. Cooper, *Nat. Chem.*, 2013, **5**, 276–281.
- 17 T. Hasell, X. Wu, J. T. A. Jones, J. Bacsá, A. Steiner, T. Mitra, A. Trewin, D. J. Adams and A. I. Cooper, *Nat. Chem.*, 2012, **2**, 750–755.

