



Cite this: *Mol. Syst. Des. Eng.*, 2018, 3, 223

Received 4th September 2017,  
Accepted 9th October 2017

DOI: 10.1039/c7me00090a

rsc.li/molecular-engineering

## A solution-processable dissymmetric porous organic cage†

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Two dissymmetric racemic analogues of the chiral porous organic cage, CC3, were isolated and unambiguously characterised as a racemate pair of the *R,R,R,S,S,S* and *S,S,S,R,R,R*-diastereomers (CC3-*RS* and CC3-*SR*). CC3-*RS*/CC3-*SR* equals the highest porosity measured for CC3 but is an order of magnitude more soluble, making it an excellent candidate for incorporation into a membrane for separation applications.

### Design, System, Application

The porous organic cage (POC), CC3, is a promising material for a variety of industrial separations, and has been shown to separate noble gases, hexane isomers, racemic alcohols, and SF<sub>6</sub> from N<sub>2</sub>. CC3 is remarkably hydrolytically stable, but is formed from expensive chiral starting materials, and has limited solubility for further processing (up to 3 mg mL<sup>-1</sup> in chloroform) – clear disadvantages when considering industrial applications. We sought to design a cheaper route to CC3 using racemic precursors (£1.36 g<sup>-1</sup> vs. £42.90 g<sup>-1</sup> for racemic vs. chiral diamine), and in the process serendipitously discovered a new dissymmetric isomer of CC3 that is sixteen times more soluble (48 mg mL<sup>-1</sup>) and retains the desirable porosity characteristics of the original analogue in the solid state. We anticipate this will make the industrial use of porous organic cages more feasible, and open new routes to processing POCs into membranes for separation technologies. This discovery also has implications for high-throughput screening workflows for designing and discovering new POCs, underlining the importance of *in situ* reaction monitoring for even seemingly simple reactions.

### Introduction

The separation of chemicals and gases from crude mixtures accounts for 10–15% of the world's energy consumption, chiefly due to distillation processes.<sup>1</sup> Membrane-based separations can use up to 90% less energy than distillation,<sup>2</sup> and hence there is a demand for new materials that can be processed into separation membranes. Molecular cage compounds<sup>3,4</sup> are attractive materials for membrane applications, in part due to their solution processability, which can be used to tune their porosity by crystal engineering,<sup>5</sup> to deposit them onto a range of substrates, and to blend them with other materials, such as polymers, to form mixed matrix membranes.<sup>6</sup> The porous organic cage (POC) CC3 (ref. 7) shows remarkable hydrolytic stability and can separate noble gases,<sup>4</sup> SF<sub>6</sub> from N<sub>2</sub>,<sup>8</sup> hexane<sup>9</sup> and xylene isomers,<sup>10</sup> and racemic alcohols and amines.<sup>4,9</sup> To date, CC3 is the subject of more than 20 publications.

CC3 is a chiral imine cage and is synthesised from 1,3,5-triformylbenzene (TFB) and either *R,R*- or *S,S*-cyclohexanediamine (CHDA), resulting in CC3-*R* or -*S*, respectively. The sorption and separation capabilities of CC3 in a chiral crystalline form arises from the window-to-window packing of the cages in the structure, which affords a diamondoid pore network, CC3- $\alpha$ , with a narrow pore size distribution (static pore diameter = 3.6 Å).<sup>4</sup> The pore topology found in chirally pure CC3 (CC3-*R* or CC3-*S*) is also found in its racemate, CC3-*R*/CC3-*S*.<sup>‡</sup> The latter racemate precipitates immediately when a solution of CC3-*R* is mixed with a solution of CC3-*S* (Fig. 1).<sup>11</sup> Previous gas phase density functional theory (DFT) dimer calculations showed that heterochiral dimer pairs were more stable than homochiral dimer pairs (–169 kJ mol<sup>-1</sup> versus –150 kJ mol<sup>-1</sup>), explaining the rapid precipitation of a stable network on the mixing of enantiomers in solution.<sup>11</sup> The marked solid state stabilization of the racemic crystalline CC3-*R*/CC3-*S* material with respect to homochiral CC3 was also demonstrated by crystal structure prediction.<sup>12</sup> More generally, this chiral window pairing extends to a range of other [4 + 6] tetrahedral imine

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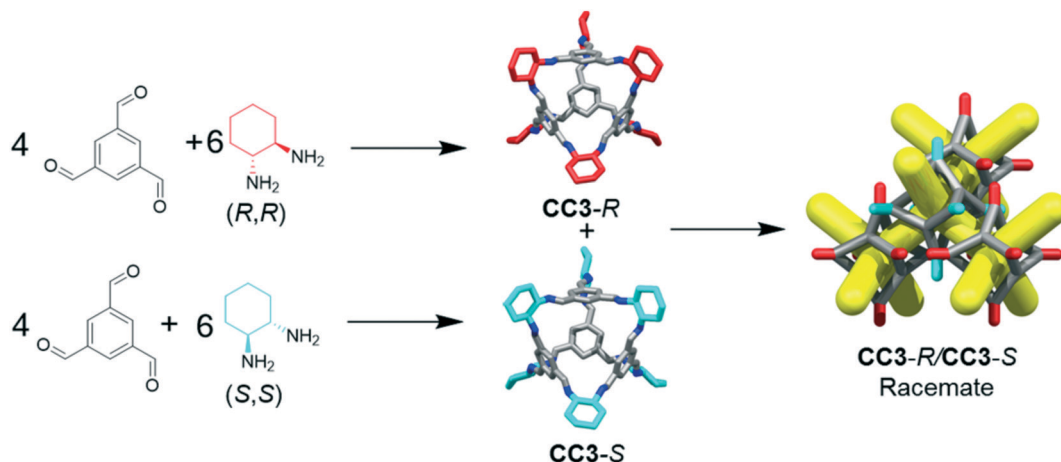
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† Electronic supplementary information (ESI) available. CCDC 1571867 and 1571866. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c7me00090a

‡ Notation used throughout the text for CC3 is as follows: CC3-*R* and CC3-*S* are enantiomerically pure cages; CC3-*R*/CC3-*S* is the racemic co-crystal of CC3-*R* and CC3-*S*; CC3-*RS* and CC3-*SR*, the new cages reported here, are a racemate pair of diastereomers of CC3; CC3-*RS*/CC3-*SR* refers to the co-crystal containing both CC3-*RS* and CC3-*SR*.





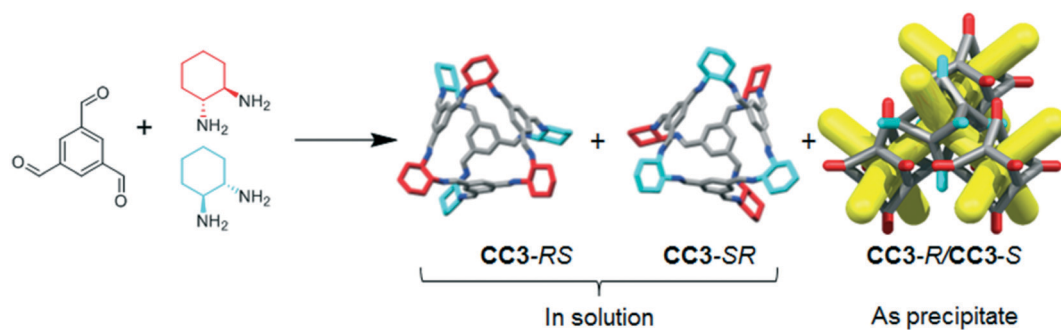
**Fig. 1** Reaction scheme for formation of CC3-*R* and CC3-*S*, which co-crystallises immediately upon mixing to form the racemate CC3-*R/CC3-S*. For CC3-*R*, the cyclohexane groups are shown in red; for CC3-*S*, in turquoise; other C, grey; N, blue; H omitted. The structure on the right is a schematic representation of the desolvated racemic CC3-*R/CC3-S* co-crystal; diamondoid pore network shown in yellow, simplified cage frame in grey, simplified cyclohexyl vertices in red (*-R*) and turquoise (*-S*).

POCs,<sup>11,13,14</sup> and also to non-tetrahedral POCs bearing the same chiral windows, such as linear, tubular POCs.<sup>15</sup>

CC3 and its racemate have good hydrolytic stability<sup>16</sup> and amination derivatives of this POC are even stable to strong acids and bases.<sup>17</sup> However, for CC3 or similar materials to find use in industrial applications, the synthesis must also be cost-efficient. Enantiomerically pure CHDA is much more expensive than racemic CHDA (£214.50 for 5 g and £12.90 for 10 mL respectively; prices obtained from Sigma-Aldrich®)<sup>18</sup> and hence the use of the racemic diamine would provide a significant cost benefit. The racemic POC CC3-*R/CC3-S* can be accessed by directly reacting TFB with ( $\pm$ )-*trans*-1,2-cyclohexanediamine (*rac*-CHDA)<sup>19</sup> and we thus sought to synthesise a porous solid in this manner (Fig. 1). The racemic co-crystal was found to precipitate rapidly from the reaction solution as a poorly-soluble crystalline solid (408 mg isolated, 47% yield, ESI<sup>†</sup> section 2.1; Fig. 4bii for PXRD) — a clear disadvantage for further processing, such as incorporation into membranes. Others have reported the isolation of new stereoisomers of CC3 by collecting the precipitate resulting from the reaction of racemic CHDA with TFB.<sup>12</sup> In our experiments, we only observed CC3-*R/CC3-S* precipitating from so-

lution (characterised by PXRD, Fig. 4bi and ii). However, analysis of the supernatant revealed the presence of two previously unreported diastereomers of CC3, CC3-*RS* and CC3-*SR*<sup>†</sup> (Fig. 2; 312 mg isolated, 36% yield, ESI<sup>†</sup> section 2.1); that is, an asymmetric racemic cage that, unlike CC3-*R/CC3-S*, retains its solution processability.

NMR analysis (Fig. 3, ESI<sup>†</sup> section 2.1.1 and 2.4) suggested that both enantiomers, CC3-*RS* and CC3-*SR*, contain three *R*, *R*-CHDA vertices and three *S*, *S*-CHDA vertices, and that these cages were the majority enantiomers present in solution. The NMR spectra of the asymmetric species, CC3-*RS* and CC3-*SR*, are markedly different to homochiral CC3, which enables identification of the two cage geometries. To probe this in more detail and to ascertain if any other species were formed, even transiently, the reaction between TFB and *rac*-CHDA was analysed by <sup>1</sup>H NMR immediately after mixing, and then at 1, 2, 4, and 24 hours (Fig. 3). A solid precipitate formed over the course of the reaction, which was identified by PXRD analysis as the known racemate, CC3-*R/CC3-S* (Fig. 4bii, red line). NMR analysis of the solution after 1 hour indicated a complex mixture of products that could not be identified unambiguously (Fig. 3 and ESI<sup>†</sup> Fig. S3 and S4).



**Fig. 2** TFB and *rac*-CHDA react to form CC3-*RS* and CC3-*SR*, which are soluble and remain in solution, as well as CC3-*R* and CC3-*S*, which immediately co-crystallise to form the racemate, CC3-*R/CC3-S*, as a white precipitate. Colours as in Fig. 1.



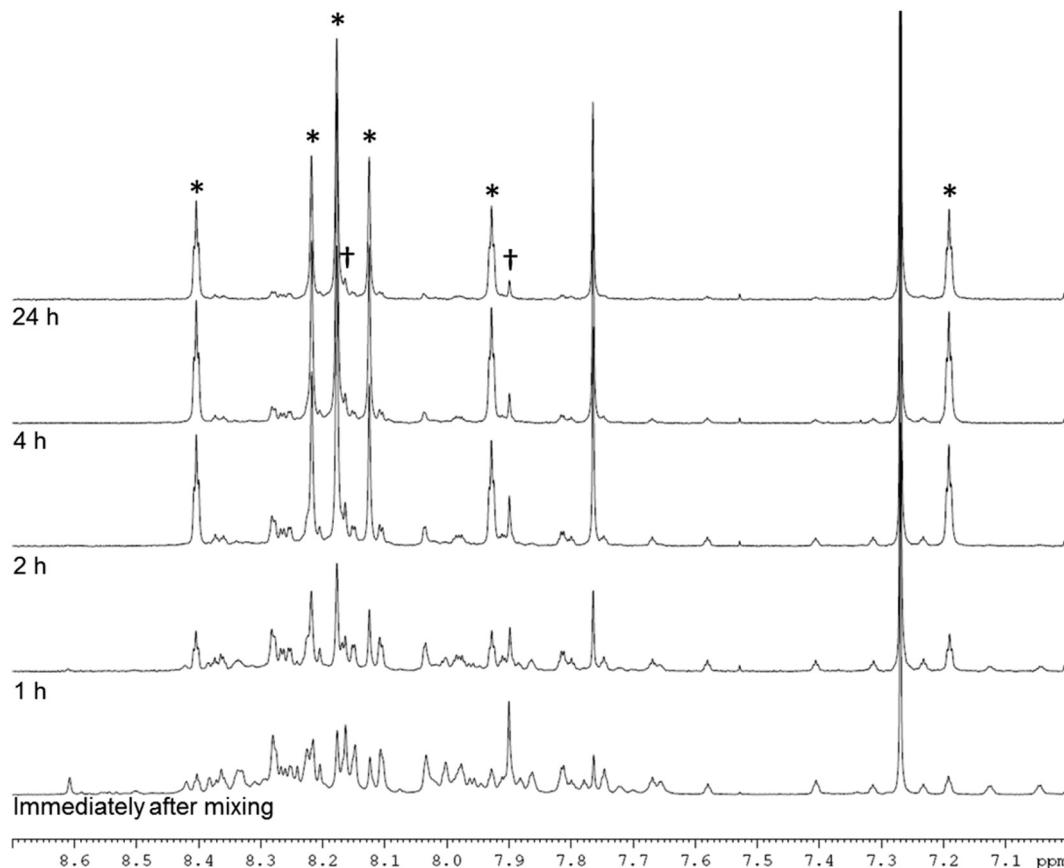


Fig. 3 NMR spectrum (400 MHz,  $\text{CDCl}_3$ ) between 7.0 and 8.7 ppm of the reaction of TFB + *rac*-CHDA immediately after mixing, and after 1, 2, 4, and 24 hours. Asterisks indicate peaks arising from CC3-*RS* and CC3-*SR*, confirmed by 2D NMR, MS, and analysis of pure cage sample (ESI† section 2); daggers indicate peaks assigned to CC3-*R*- or -*S*.

After 24 hours, an imine-containing cage-like product was the major component remaining in solution.

We have shown previously that the reaction of TFB with a mixture of diamines, such as a binary mixture of cyclohexanediamine and ethylenediamine, results in the formation of a statistical mixture of scrambled cage products.<sup>20</sup> In that example, 12 mixed cage products were formed. By contrast, only four of the possible 12 products were isolated in these experiments: CC3-*R*, CC3-*S*, CC3-*RS*, and CC3-*SR*.

DFT calculations were carried out to assess the relative stabilities of the two isomers (ESI† section 3). CC3-*RS* was calculated to be 10  $\text{kJ mol}^{-1}$  less stable than CC3-*R*. As previously reported,<sup>11</sup> precipitation of the CC3-*R*/CC3-*S* co-crystal (Fig. 1) provides an additional solid state stabilization. That CC3-*R*/CC3-*S* is not the only product suggests that both CC3-*RS* and CC3-*SR* are kinetically trapped in this configuration, and that the process is insufficiently dynamic, potentially due to the evaporation of catalytic TFA.

To explore whether the presence of CC3-*RS* and CC3-*SR* over other possible enantiomers of CC3 was a result of the 1:1 ratio of *R,R*- and *S,S*-CHDA, reactions were performed with varying proportions of each enantiomer of CHDA (5:1, 4:2, 2:4, and 1:5 of *R,R*-CHDA:*S,S*-CHDA respectively; each set repeated twice; ESI† section 2.2 and Fig. S5 and

S6). After 24 hours, all reactions resulted in either mixtures of homochiral CC3 and CC3-*RS*/CC3-*SR*, or, in one case, homochiral CC3, CC3-*RS* and -*SR*, and aldehyde starting materials. This strongly suggests that homochiral CC3 and CC3-*RS*/CC3-*SR* are more stable than other potential diastereomers such as, for example, a cage containing 1 *S,S*-CHDA and 5 *R,R*-CHDA vertices. Such chiral self-sorting has been reported recently in salicylimine cages;<sup>21</sup> in that case, both products could be retained in solution in certain solvents, thus allowing a detailed analysis of their relative energies. Such analysis is not possible in our system due to the very low solubility ( $<1 \text{ mg mL}^{-1}$ ) of the CC3-*R*/CC3-*S* racemate.

POCs with different geometries have divergent physical properties,<sup>7,15</sup> which results from their different crystal packings and the impact of this on pore connectivity, internal cavity size, window configuration, surface area, and porosity. However, it is rare to be able to study two cages with identical chemical composition, but different geometries. Here, we can identify differences that arise purely from the *shape* of the cage molecules. We therefore investigated the solid-state crystal packing and physical properties of this new CC3-*RS*/CC3-*SR* racemate compare to the previously reported racemate, CC3-*R*/CC3-*S*.





**Fig. 4** a) Four cages are shown with both arene-to-arene (leftmost cage pair) and window-to-window packing (central cage pair), taken from the single crystal structure of CC3-RS/CC3-SR (ESI† section 2.8). Hydrogen atoms removed for clarity; colouring as in Fig. 1. b) PXRD patterns of (i) known racemate CC3-R/CC3-S, (ii) CC3-R/CC3-S precipitate isolated *via* filtration; (iii) CC3-RS/CC3-SR isolated by precipitation with hexane, (iv) simulated PXRD from sc-XRD data for CC3-RS/CC3-SR and (v) CC3-RS/CC3-SR post gas sorption measurements. c) Gas sorption isotherms for CC3-RS/CC3-SR for N<sub>2</sub> (77 K, black), H<sub>2</sub> (77 K, blue), CO<sub>2</sub> (295 K, orange), Xe (273 K, purple), and Kr (273 K, green). Filled and open symbols represent adsorption and desorption isotherms respectively.

CC3-RS/CC3-SR was isolated from the reaction mixture in solution by filtration to remove the CC3-R/CC3-S precipitate. The filtrate was then reduced in volume and poured into hexane, resulting in a white precipitate that was isolable by filtration. Unlike CC3-R/CC3-S (solubility of  $<1 \text{ mg mL}^{-1}$  in CHCl<sub>3</sub>), and to a lesser extent CC3-R (solubility of  $3 \text{ mg mL}^{-1}$ ),<sup>22</sup> the solid product was highly soluble in chloroform ( $48 \text{ mg mL}^{-1}$ , ESI† section 1). The solid CC3-RS/CC3-SR precipitate was found to be crystalline and easily distinguished from CC3-R/CC3-S by PXRD (Fig. 4biii and ii, blue and red line respectively, ESI† section 2.9). Single crystals of the filtrate were grown from DCM/hexane, DCM/acetone, and DCM/MeOH by vial-in-vial crystallisation experiments. Single crystal X-ray diffraction (SCXRD) unambiguously identified the product as a racemate of CC3-RS and CC3-SR which has crystallised in the trigonal space group  $R\bar{3}$  (Fig. 4a, ESI† section 2.8) and the simulated PXRD diffraction pattern of the crystal structure matches with the PXRD pattern of the hexane precipitated filtrate (Fig. 4biii and iv, pink and blue line respectively). CC3-RS and CC3-SR both have one triangular window with an equilateral shape, identical to the windows found in CC3-R (or -S). However, unlike CC3-R (or -S), in CC3-

RS and CC3-SR there are three windows with an acute triangle shape. In addition, one aromatic ring is angled towards the centre of the cage cavity (ESI† section 2.8). During crystallisation, CC3-RS and CC3-SR self-sort into 1-D packed arrangements. In the structure, CC3-RS packs in an alternating window-to-window and arene-to-arene packing arrangement (Fig. 4a, central two cages). 1-D packed arrangements of CC3-SR are equivalent and related by inversion symmetry. By contrast, there are no arene-to-arene contacts in chiral CC3 or in the previously reported racemate, CC3-R/CC3-S.

Gas sorption measurements (Fig. 4c) showed CC3-RS/CC3-SR has an apparent BET surface area of  $800 \text{ m}^2 \text{ g}^{-1}$  (ESI† section 2.10), as compared to values of  $409\text{--}819 \text{ m}^2 \text{ g}^{-1}$  and up to  $696 \text{ m}^2 \text{ g}^{-1}$  obtained for CC3-R and CC3-R/CC3-S, respectively (the higher values were obtained from rapidly precipitated samples with a high proportion of structural defects).<sup>11</sup> As such, the dissymmetric crystalline racemate is more porous than either crystalline form of CC3 reported so far, and its surface area is similar to that reported previously for amorphous CC3.<sup>11,23</sup> As the CC3-RS/CC3-SR used for gas sorption was isolated by rapid precipitation, there is likely to be a contribution to porosity from crystal defects in the sample,



as seen for rapidly precipitated CC3.<sup>11</sup> As much higher concentrations of the dissymmetric cage can be achieved in solution than for CC3, it is likely that crystallisation conditions favouring higher defect inclusion can be achieved *via* rapid precipitation, potentially achieving even higher gas sorption capacities. The isolated crystalline solid is stable to >300 °C, and it is also stable in neutral solutions in DCM for at least 6 days, thus it can be solution processed without conversion to alternative diastereomers of CC3.

## Conclusions

A new diastereomer of the well-known homochiral POC, CC3, has been isolated and identified as CC3-*RS*. CC3-*RS* surpasses crystalline CC3-*R* and CC3-*R*/CC3-*S* in terms of both solubility and microporosity, and it does not require the use of expensive homochiral amines. An oft-cited advantage of imine condensation routes to POCs is that the reactions are one-pot in nature. This study shows, however, that the cage-forming mechanism can be more complex than implied by the majority product (Fig. 3). In turn, this suggests that it might be useful to pay more attention to *in situ* reaction monitoring for these systems in the future.

## Conflicts of interest

There are no conflicts of interest to declare.

## Acknowledgements

We acknowledge the EPSRC (EP/N004884/1) and European Research Council under the European Union's Seventh Framework Programme (FP/2007-2013) through grant agreement n. 321156 (ERC-AG-PE5-ROBOT) for funding. KEJ thanks the Royal Society for a University Research Fellowship; AGS thanks the Royal Society and the EPSRC for a Royal Society-EPSRC Dorothy Hodgkin Fellowship. The authors thank Dr Tom Hasell for useful discussions. We acknowledge Stephen Moss and the MicroBioRefinery for assistance with QTOF-MS measurements.

## Notes and references

- D. S. Sholl and R. P. Lively, *Nature*, 2016, 533, 316.
- M. Mulder, *Basic principles of membrane technology*, Kluwer Academic, Dordrecht, Boston, 2nd edn, 1996.
- M. Mastalerz, *Angew. Chem., Int. Ed.*, 2010, 49, 5042–5053.
- L. Chen, P. S. Reiss, S. Y. Chong, D. Holden, K. E. Jelfs, T. Hasell, M. A. Little, A. Kewley, M. E. Briggs, A. Stephenson, K. M. Thomas, J. A. Armstrong, J. Bell, J. Busto, R. Noel, J. Liu, D. M. Strachan, P. K. Thallapally and A. I. Cooper, *Nat. Mater.*, 2014, 13, 954–960.
- M. A. Little, M. E. Briggs, J. T. A. Jones, M. Schmidtman, T. Hasell, S. Y. Chong, K. E. Jelfs, L. J. Chen and A. I. Cooper, *Nat. Chem.*, 2015, 7, 153–159.
- Q. Song, S. Jiang, T. Hasell, M. Liu, S. Sun, A. K. Cheetham, E. Sivaniah and A. I. Cooper, *Adv. Mater.*, 2016, 28, 2629–2637.
- 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.
- T. Hasell, M. Miklitz, A. Stephenson, M. A. Little, S. Y. Chong, R. Clowes, L. Chen, D. Holden, G. A. Tribello, K. E. Jelfs and A. I. Cooper, *J. Am. Chem. Soc.*, 2016, 138, 1653–1659.
- A. Kewley, A. Stephenson, L. J. Chen, M. E. Briggs, T. Hasell and A. I. Cooper, *Chem. Mater.*, 2015, 27, 3207–3210.
- 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.
- T. Hasell, S. Y. Chong, K. E. Jelfs, D. J. Adams and A. I. Cooper, *J. Am. Chem. Soc.*, 2012, 134, 588–598.
- J. T. A. Jones, T. Hasell, X. F. 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.
- T. Hasell, M. A. Little, S. Y. Chong, M. Schmidtman, M. E. Briggs, V. Santolini, K. E. Jelfs and A. I. Cooper, *Nanoscale*, 2017, 9, 6783–6790.
- T. Hasell, S. Y. Chong, M. Schmidtman, D. J. Adams and A. I. Cooper, *Angew. Chem., Int. Ed.*, 2012, 51, 7154–7157.
- A. G. Slater, M. A. Little, A. Pulido, S. Y. Chong, D. Holden, L. Chen, C. Morgan, X. Wu, G. Cheng, R. Clowes, M. E. Briggs, T. Hasell, K. E. Jelfs, G. M. Day and A. I. Cooper, *Nat. Chem.*, 2017, 9, 17–25.
- T. Hasell, M. Schmidtman, C. A. Stone, M. W. Smith and A. I. Cooper, *Chem. Commun.*, 2012, 48, 4689–4691.
- M. Liu, M. A. Little, K. E. Jelfs, J. T. A. Jones, M. Schmidtman, S. Y. Chong, T. Hasell and A. I. Cooper, *J. Am. Chem. Soc.*, 2014, 136, 7583–7586.
- <http://www.sigmaaldrich.com/catalog/search?interface=All&term=cyclohexanediamine&N=0&focus=product&lang=en&region=GB>, accessed on 7th August 2017.
- G. Zhu, C. D. Hoffman, Y. Liu, S. Bhattacharyya, U. Tumuluri, M. L. Jue, Z. Wu, D. S. Sholl, S. Nair, C. W. Jones and R. P. Lively, *Chem. – Eur. J.*, 2016, 22, 10743–10747.
- S. Jiang, J. T. Jones, T. Hasell, C. E. Blythe, D. J. Adams, A. Trewin and A. I. Cooper, *Nat. Commun.*, 2011, 2, 207.
- D. Beaudoin, F. Rominger and M. Mastalerz, *Angew. Chem., Int. Ed.*, 2017, 56, 1244–1248.
- R. L. Greenaway, D. Holden, E. G. B. Eden, A. Stephenson, C. W. Yong, M. J. Bennison, T. Hasell, M. E. Briggs, S. L. James and A. I. Cooper, *Chem. Sci.*, 2017, 8, 2640–2651.
- S. Jiang, K. E. Jelfs, D. Holden, T. Hasell, S. Y. Chong, M. Haranczyk, A. Trewin and A. I. Cooper, *J. Am. Chem. Soc.*, 2013, 135, 17818–17830.

