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## Pore space partition of a fragile Ag(I)-carboxylate framework *via* post-synthetic linker insertion†

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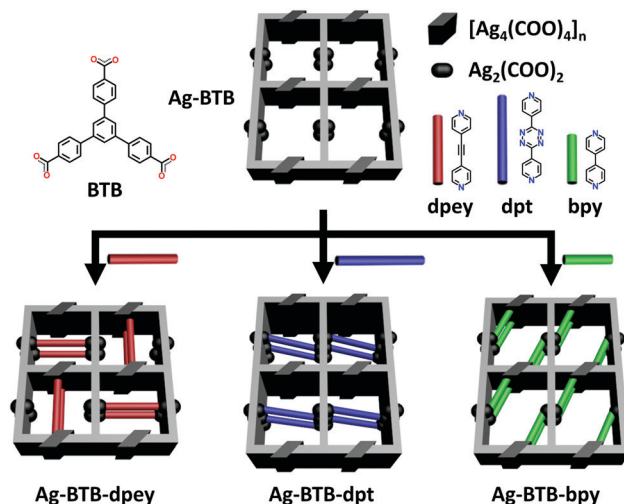
**The pore space partition approach *via* post-synthetic linker insertion was used to modulate the porosity of a fragile Ag(I)-carboxylate framework with potentially large pore space. The resulting Ag(I)-MOFs with partitioned pores showed enhanced permanent porosity compared with a nonpartitioned Ag(I)-carboxylate framework.**

Metal-organic frameworks (MOFs) are porous crystalline networks primarily composed of coordination bonds between metal ions and organic ligands.<sup>1</sup> Their structural diversity represents a significant advantage for many applications based on the intrinsic properties of the constituent metals and functionalized ligands.<sup>2</sup> After surveying metal ions for novel porous architectures, silver(I) has become a promising candidate due to its tolerance for a multitude of coordination numbers and geometries.<sup>3</sup> Over the past decades, significant efforts have been devoted to preparing porous Ag(I)-based MOFs, but only a few have been investigated for their gas sorption properties<sup>4</sup> because of the relatively weak Ag(I)-ligand bonds. These weak bonds often result in framework collapse during the activation process. Therefore, the synthesis of Ag(I)-based MOFs with permanent porosity remains a challenge in the expansion of MOF diversity and the utilization of silver. Recently, a pore space partition (PSP) strategy<sup>5</sup> has been developed as a tool for improving the structural stability<sup>6</sup> and gas separation abilities of MOFs.<sup>7</sup> PSP utilizes the division of large pore spaces into smaller segments by using secondary metal<sup>8</sup> or ligand insertion.<sup>9</sup> However, most MOFs with partitioned pore space are obtained *via* one-pot synthesis using all framework building blocks including pore partitioning agents.

Herein, a rare example of permanently microporous Ag(I)-MOFs achieved by the PSP approach *via* post-synthetic insertion<sup>10</sup> of auxiliary linkers is reported. The parent Ag(I)-carboxylate framework

(AgBTB) contains a large one-dimensional (1D) solvent channel. However, AgBTB is fragile and practically nonporous for gas sorption. After partitioning the 1D channel *via* post-synthetic insertion of dipyriddy linkers, the resulting Ag(I)-MOFs showed improved structural stability upon guest molecule removal. In addition, the framework porosity could be adjusted by the amount and type of linkers added (Scheme 1).

The solvothermal reaction of AgNO<sub>3</sub> with 4,4',4''-benzene-1,3,5-triyl-tri-benzoic acid (H<sub>3</sub>BTB) afforded AgBTB crystals. A three-dimensional (3D) coordination network was formed based on a finite [Ag<sub>2</sub>(COO)<sub>2</sub>] dinuclear cluster and an infinite [Ag<sub>4</sub>(COO)<sub>4</sub>]<sub>n</sub> 1D ribbon with inorganic nodes and BTB as an organic linker (Fig. 1 and Fig. S1, ESI†). Ag1 of the finite [Ag<sub>2</sub>(COO)<sub>2</sub>] secondary building unit (SBU) adopted a square planar geometry with two *syn-syn* bridging bidentate carboxylate oxygen atoms in a *trans*-configuration, an Ag atom of an inter-silver bond, and a ligated water molecule. Ag2 and Ag3 of the infinite [Ag<sub>4</sub>(COO)<sub>4</sub>]<sub>n</sub> structure adopted distorted square



**Scheme 1** Pore space partition of AgBTB *via* post-synthetic insertion of dipyriddy linkers.

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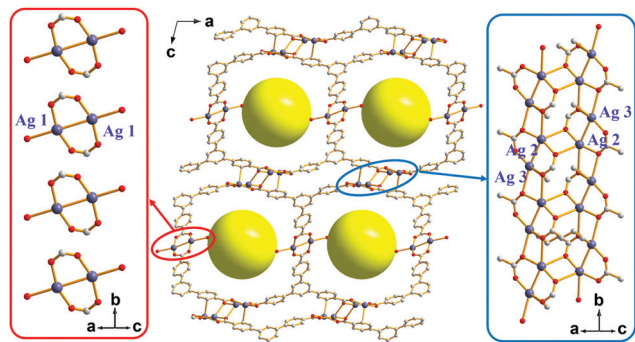


Fig. 1 The framework structure of **AgBTB** with an array of finite  $[\text{Ag}_2(\text{COO})_2]$  dinuclear clusters and an infinite  $[\text{Ag}_4(\text{COO})_4]_n$  1D ribbon.

pyramidal geometries and were interconnected *via* two carboxylates to form a 1D ribbon along the crystallographic *b*-axis. **AgBTB** contained large 1D solvent channels that were  $10.4 \times 13.0 \text{ \AA}^2$  in size along the crystallographic *b*-axis. The total pore volume of **AgBTB** calculated by PLATON<sup>11</sup> was determined to be  $3,508 \text{ \AA}^3$  per unit cell, corresponding to 45.4% of the total unit cell volume. Dye adsorption results show that the **AgBTB** can absorb dye molecules such as methylene blue and Nile red in ethanol solutions, which suggests the void space of the **AgBTB** is well preserved in liquid medium (Fig. S2, ESI<sup>†</sup>). However, the  $\text{N}_2$  adsorption experiment of activated **AgBTB** showed no  $\text{N}_2$  uptake at 77 K. This nonporous behavior is due to pore collapse upon solvent removal from the channel. Interestingly, at least six potential linkage sites of three different categories existed between the Ag atoms in the framework (Fig. 2). Moderate-length linkage sites 1 and 2 of the first category could interconnect in-plane  $\text{Ag1-Ag1}$  ( $\sim 14.7 \text{ \AA}$ ) and off-plane  $\text{Ag1-Ag1}$  ( $\sim 15.8 \text{ \AA}$ ), respectively, in the dinuclear Ag clusters facing each

other across the 1D channel. However, the longer linkage sites 3 and 4 of the second category interconnect  $\text{Ag2-Ag2}$  ( $\sim 17.3 \text{ \AA}$ ) and  $\text{Ag3-Ag3}$  ( $\sim 17.4 \text{ \AA}$ ), respectively, in the tetranuclear Ag clusters. The short linkage sites 5 and 6 of the third category interconnect  $\text{Ag1-Ag2}$  ( $\sim 11.6 \text{ \AA}$ ) and  $\text{Ag1-Ag3}$  ( $\sim 10.7 \text{ \AA}$ ), respectively, in the **AgBTB** framework.

Given the above information, the PSP strategy was used to rationally partition the original 1D channel into smaller channels, leading to improved porosity. The PSP was achieved *via* post-synthetic insertion of carefully selected dipyriddy linkers, 1,2-di(pyridin-4-yl)ethyne (dpey), 3,6-di(pyridin-4-yl)-1,2,4,5-tetrazine (dpt), and 4,4'-bipyridine (bpy), with lengths approximately matching those of the potential linkage sites (Fig. S3, ESI<sup>†</sup>). Powder X-ray diffraction (PXRD) patterns of the resulting crystals indicated that the dpey linker induced significant reorganization of the **AgBTB** framework (Fig. S4, ESI<sup>†</sup>). In contrast, dpt and bpy linkers resulted in only moderate structural changes. Single-crystal structural analyses revealed that the inserted dipyriddy linkers acted as pore-partition agents that interconnect the potential linkage sites in the **AgBTB** framework (Fig. 3 and Fig. S5–S7, ESI<sup>†</sup>). The insertion and partitioning modes of the dipyriddy linkers depend on their length. For **AgBTB-dpey**, since the length of dpey ( $\sim 14.6 \text{ \AA}$ , including two  $2.4 \text{ \AA}$  Ag–N bonds) matches with the potential linkage site 1 ( $\sim 14.7 \text{ \AA}$ ), the fit into linkage site 1 was expected. Interestingly, insertions of dpey occurred at the preferred linkage site 1 and at the long linkage site 3 ( $\text{Ag2-Ag2}$  pair; Fig. 3b and Fig. S5, ESI<sup>†</sup>). The original 1D channels ( $10.4 \times 13.0 \text{ \AA}^2$ ) along the crystallographic *b* axis in **AgBTB** were partitioned by the dpey with horizontal and vertical modes, affording two rectangular 1D solvent channels (types A and B; Fig. 3b). Channel A ( $3.7 \times 11.6 \text{ \AA}^2$ ) was generated by the insertion of dpey into linkage site 1, resulting in horizontally partitioned channels. In contrast, the vertically partitioned channel B ( $3.8 \times 9.8 \text{ \AA}^2$ ) was created by embedding dpey in linkage site 3. Notably, embedding of the length-mismatched dpey linker at linkage site 3

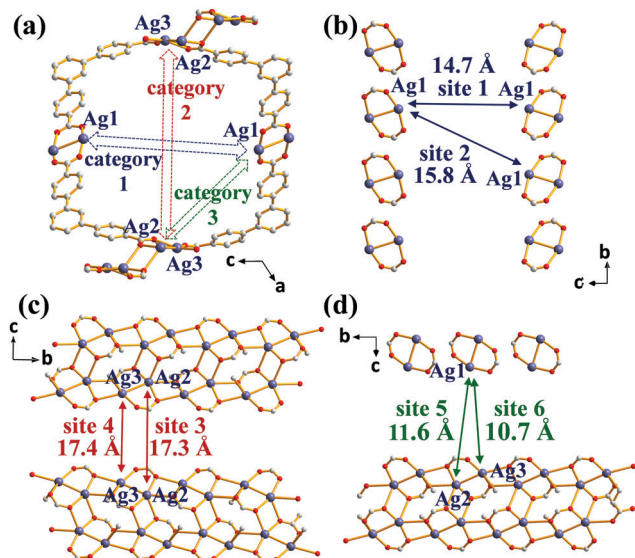


Fig. 2 (a) Potential linkages across the 1D solvent channel in **AgBTB**. Inter-silver distances (b) between dinuclear clusters, (c) between tetranuclear clusters of the 1D ribbons and (d) between the dinuclear and tetranuclear clusters of the 1D ribbon.

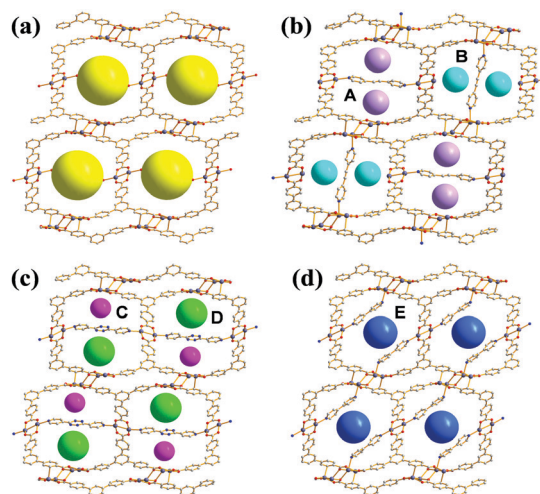


Fig. 3 (a) 1D channels of **AgBTB** and the PSPs through the insertions of the dipyriddy linkers in (b) **AgBTB-dpey**, (c) **AgBTB-dpt**, and (d) **AgBTB-bpy**.

induced a significant contraction in the unit cell along the *c*-axis and a  $\sim 5\%$  reduction in **AgBTB-dpey** unit cell volume compared to that of **AgBTB** (Table S1, ESI†). It should be noted that the total amount of dpey inserted was  $\sim 75\%$  of the maximum amount. While linkage site 1 was almost fully occupied, only 55% of linkage site 3 was occupied in the single crystal structure. The inserted dpey linker content in the **AgBTB-dpey** framework was estimated to be  $\sim 74\%$  from the  $^1\text{H}$  NMR spectrum of the digested sample (Fig. S8, ESI†). This value is in good agreement with the average site occupancy of the dpey linkers in the single crystal structure.

The length of dpt is  $\sim 16.0$  Å (with two  $\sim 2.4$  Å Ag–N bonds); thus, it was expected that linkage site 2 would be the preferred insertion site. Structural analysis of **AgBTB-dpt** revealed that post-synthetic insertion of the dpt linker occurred exclusively at linkage site 2 with a site occupancy of 0.74 (Fig. 3c and Fig. S6, ESI†). In contrast to **AgBTB-dpey**, insertion of the dpt linker into linkage site 2 showed little effect on the unit cell parameters and volume of **AgBTB-dpt** (Table S1, ESI†). The inserted dpt linker partitioned the original 1D channel into two 1D rectangular channels— $3.6 \times 11.9$  and  $5.2 \times 12.3$  Å<sup>2</sup> in size for channels C and D, respectively. The  $^1\text{H}$  NMR spectrum of digested **AgBTB-dpt** suggested that the inserted dpt linker content was as high as 63% of the theoretical maximum, slightly lower than the site occupancy of dpt linkers determined *via* single crystal structure refinement (Fig. S9, ESI†).

The length of bpy ( $\sim 12.0$  Å, including two  $2.4$  Å Ag–N bonds) is well-matched with the short-length linkage site 5 ( $\sim 11.6$  Å). As expected, the insertion of bpy was found to occur only at linkage site 5 with a site occupancy of  $\sim 0.8$  (Fig. 3d and Fig. S7, ESI†). The insertion of bpy linkers into all linkage site 5 locations within the 1D channel of **AgBTB** formed the 1D channel E with dimensions of  $5.2 \times 12.6$  Å<sup>2</sup>. Similar to dpt insertion, bpy induced no significant rearrangement of the framework. Therefore, the unit cell parameters and volume of the **AgBTB-bpy** single crystal were very similar to those of **AgBTB** (Table S1, ESI†). The bpy content estimated from the  $^1\text{H}$  NMR spectrum of digested **AgBTB-bpy** was  $\sim 1.06$  (Fig. S10, ESI†), indicating that all the linkage sites were fully occupied by bpy linkers, as determined *via* crystal structure analysis.

It should be noted that the length of the inserted linker is essential for framework rearrangement during linker insertion. As demonstrated, only the dpey linker causes significant contraction of the 1D channel and subsequent reorganization of the framework due to insertion of the length-mismatched linker into linkage site 3 (Table S1 and Fig. S5b, ESI†). In contrast, both dpt and bpy linkers localized to length-matched linkage sites in the framework, exerting negligible effects on the structure reorganization.

**AgBTB-dpey**, **AgBTB-dpt**, and **AgBTB-bpy** prepared in *N,N*-dimethylformamide (DMF) released substantial amounts of dipyriddy linkers during solvent exchange with methylene chloride. Therefore, **AgBTB-dpey(x)**, **AgBTB-dpt(x)**, and **AgBTB-bpy(x)** crystals were obtained *via* immersing **AgBTB** crystals into *x* mL of either 1 or 2 mM methylene chloride solution containing the dipyriddy linkers, where *x* denotes the volume of methylene

chloride solution. It should be noted that the amounts of inserted linkers were lower than those of crystals prepared in DMF (Fig. S11, ESI†). Interestingly, the PXRD pattern of **AgBTB-dpey(40)** differs from that of **AgBTB-dpey** prepared in DMF (Fig. S4, ESI†). This structural discrepancy may be attributed to the lower site occupancies of the linkage sites 1 and 3 in **AgBTB-dpey(40)**. Because the total amount of the dpey inserted in **AgBTB-dpey(40)** is much smaller than that of **AgBTB-dpey**, both linkage sites 1 and 3 on **AgBTB-dpey(40)** are likely to have lower site occupancies. In particular, the low site occupancy of length-mismatched linkage site 3 did not cause the framework contraction, resulting in a PXRD pattern similar to that of the parent framework **AgBTB**. Meanwhile, the **AgBTB-dpt(40)** and **AgBTB-bpy(40)** PXRD patterns are similar to those prepared in DMF. For the gas sorption experiments, activated **AgBTB-dpey(x)a**, **AgBTB-dpt(x)a**, and **AgBTB-bpy(x)a** were prepared in vacuum at 298 K for 1 day to minimize pore collapse.

Activated **AgBTB** (**AgBTB-a**) showed no  $\text{N}_2$  uptake at 77 K and minor  $\text{CO}_2$  uptake (up to  $\sim 10$  cm<sup>3</sup> g<sup>−1</sup>) at 195 K (Fig. S12, ESI†). Meanwhile, the  $\text{N}_2$  uptake of **AgBTB-dpey(10)a** with  $\sim 25\%$  site occupancy was  $\sim 98$  cm<sup>3</sup> g<sup>−1</sup> at relative pressure  $P/P_0 = 1$ , reaching  $\sim 137$  cm<sup>3</sup> g<sup>−1</sup> for **AgBTB-dpey(40)a** with  $\sim 48\%$  site occupancy (Fig. 4a). The BET specific surface areas were calculated as 503 and 342 m<sup>2</sup> g<sup>−1</sup> for **AgBTB-dpey(40)a** and **AgBTB-dpey(10)a**, respectively. At 195 K, the  $\text{CO}_2$  isotherms exhibited stepwise adsorption and hysteresis during desorption, which becomes evident for **AgBTB-dpey(40)a** (Fig. 4b). The  $\text{CO}_2$  adsorption amounts at 1 bar increased from  $\sim 118$  cm<sup>3</sup> g<sup>−1</sup> for **AgBTB-dpey(10)a** to  $\sim 160$  cm<sup>3</sup> g<sup>−1</sup> for **AgBTB-dpey(40)a**. However, at 273 and 298 K the  $\text{CO}_2$  isotherms exhibited neither stepwise adsorption nor hysteresis (Fig. S13, ESI†). The stepwise adsorption is likely due to stability differences between partitioned channels A and B in the **AgBTB-dpey** framework. Highly populated dpey linkers in channel A effectively prevented pore collapse during activation to a larger extent than the less populated dpey linkers in channel B. The stepwise

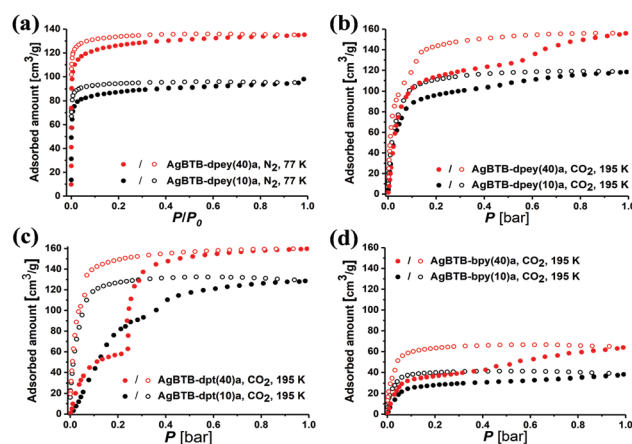


Fig. 4 (a)  $\text{N}_2$  adsorption isotherms of **AgBTB-dpey(x)a** at 77 K. (b)  $\text{CO}_2$  adsorption isotherms of **AgBTB-dpey(x)a**, (c) **AgBTB-dpt(x)a**, and (d) **AgBTB-bpy(x)a** at 195 K (filled circles: adsorption; empty circles: desorption).

adsorption observed at  $\sim 0.6$  bar is likely a result of the recovery of the partially collapsed pore upon  $\text{CO}_2$  adsorption. The  $\text{N}_2$  and  $\text{CO}_2$  adsorption results demonstrated that the PSP approach accomplished *via* post-synthetic dpey insertion enhanced the structural stability of **AgBTB-dpey(x)a** over that of **AgBTB**.

For **AgBTB-dpt(x)a**, neither **AgBTB-dpt(10)a** nor **AgBTB-dpt(40)a** showed any  $\text{N}_2$  uptake at 77 K (Fig. S14, ESI<sup>†</sup>). Even though the site occupancies of the dpt linkers,  $\sim 35\%$  and  $\sim 48\%$  for **AgBTB-dpt(10)a** and **AgBTB-dpt(40)a**, respectively (Fig. S11, ESI<sup>†</sup>), are comparable to those of the corresponding **AgBTB-dpey(x)a** moieties, the frameworks were not sufficiently stable upon activation. The site 2 linkage between the off-plane Ag1–Ag1 pair across the 1D channel resulted in inefficient partitioning of the 1D channel, leading to pore collapse in **AgBTB-dpt(x)** during activation. However, at 195 K **AgBTB-dpt(x)** exhibited stepwise  $\text{CO}_2$  adsorption behavior with hysteresis, which is pronounced for **AgBTB-dpt(40)a** (Fig. 4c). The  $\text{CO}_2$  adsorption at 1 bar increased from  $\sim 130 \text{ cm}^3 \text{ g}^{-1}$  for **AgBTB-dpt(10)a** to  $\sim 160 \text{ cm}^3 \text{ g}^{-1}$  for **AgBTB-dpt(40)a**. This stepwise adsorption can be attributed to the degree of recovery of the collapsed channels. The smaller channel C ( $3.6 \times 11.9 \text{ \AA}^2$ ) is likely restored at lower  $\text{CO}_2$  pressures, followed by the recovery of the larger channel D ( $5.2 \times 12.3 \text{ \AA}^2$ ) at higher  $\text{CO}_2$  pressures. At 273 and 298 K, the recovery of both channels is ineffective, and  $\text{CO}_2$  uptakes were  $\sim 35$  and  $17 \text{ cm}^3 \text{ g}^{-1}$  at 273 and 298 K, respectively, for **AgBTB-dpt(40)a** (Fig. S15, ESI<sup>†</sup>).

Similarly, the bpy linkers in **AgBTB-bpy(10)a** and **AgBTB-bpy(40)a** occupied 22% to 36% of the potential linkage sites, showing no  $\text{N}_2$  uptake at 77 K (Fig. S16, ESI<sup>†</sup>). The  $\text{CO}_2$  adsorption of **AgBTB-bpy(10)a** measured at 195 K exhibited a type I isotherm curve, with total  $\text{CO}_2$  uptake of  $\sim 40 \text{ cm}^3 \text{ g}^{-1}$  (Fig. 4d). The total  $\text{CO}_2$  uptake for **AgBTB-bpy(40)a** reached  $\sim 65 \text{ cm}^3 \text{ g}^{-1}$  due to the increased linkage site occupancy of the bpy linkers. Stepwise adsorption was observed along with hysteresis, which can be rationalized using the same argument discussed earlier. The presence of the bpy linker in the framework of **AgBTB-bpy(40)a** allowed for initial  $\text{CO}_2$  adsorption up to  $\sim 0.35$  bar. At pressure of  $>0.35$  bar, recovery of the collapsed solvent channels can occur. At 273 and 298 K, gradually increasing  $\text{CO}_2$  uptake is observed with increasing amounts of bpy linkers. The maximum  $\text{CO}_2$  uptakes for **AgBTB-bpy(40)a** were  $\sim 15$  and  $10 \text{ cm}^3 \text{ g}^{-1}$  at 273 and 298 K, respectively (Fig. S17, ESI<sup>†</sup>).

In summary, rare examples of permanently porous Ag(I)-based MOFs were prepared herein using the PSP approach *via* post-synthetic insertion of auxiliary dipyrpyridyl linkers. The dipyrpyridyl linkers determined the partitioning modes of the original 1D channel in the parent **AgBTB** framework and enhanced framework stability. Consequently, permanent porosities of the prepared MOFs (**AgBTB-dpey**, **AgBTB-dpt**, and **AgBTB-bpy**) were

achieved. The stability and porosity of the frameworks could be adjusted by the amount and type of linkers added.

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## Conflicts of interest

There are no conflicts to declare.

## Notes and references

- (a) N. W. Ockwig, O. Delgado-Friedrichs, M. O'Keeffe and O. M. Yaghi, *Acc. Chem. Res.*, 2005, **38**, 176; (b) 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.
- H. Furukawa, K. E. Cordova, M. O'Keeffe and O. M. Yaghi, *Science*, 2013, **341**, 1230444.
- (a) L.-Y. Du, W.-J. Shi, L. Hou, Y.-Y. Wang, Q.-Z. Shi and Z. Zhu, *Inorg. Chem.*, 2013, **52**, 14018; (b) M. A. Haja, C. B. Aakeröy and J. Desper, *New J. Chem.*, 2013, **37**, 204; (c) B. S. Fox, M. K. Beyer and V. E. Bondybey, *J. Am. Chem. Soc.*, 2002, **124**, 13613.
- (a) C. R. Murdock and D. M. Jenkins, *J. Am. Chem. Soc.*, 2014, **136**, 10983; (b) M. Handke, H. Weber, M. Lange, J. Möllmer, J. Lincke, R. Gläser, R. Staudt and K. Krautscheid, *Inorg. Chem.*, 2014, **53**, 7599; (c) I. Bassanetti, F. Mezzadri, A. Comotti, P. Sozzani, M. Gennari, G. Calestani and L. Marchiò, *J. Am. Chem. Soc.*, 2012, **134**, 9142.
- Q.-G. Zhai, X. Bu, X. Zhao, D.-S. Li and P. Feng, *Acc. Chem. Res.*, 2017, **50**, 407.
- Y.-P. Wu, J.-W. Tian, S. Liu, B. Li, J. Zhao, L.-F. Ma, D.-S. Li, Y.-Q. Lan and X. Bu, *Angew. Chem., Int. Ed.*, 2019, **58**, 12185.
- (a) S.-T. Zhang, J. T. Bu, Y. Li, T. Wu, F. Zuo, P. Feng and X. Bu, *J. Am. Chem. Soc.*, 2010, **132**, 17062; (b) Y.-X. Tan, Y.-P. He and J. Zhang, *Chem. Commun.*, 2011, **47**, 10647; (c) Y. Ling, M. Deng, Z. Chen, B. Xia, X. Liu, Y. Yang, Y. Zhou and L. Weng, *Chem. Commun.*, 2013, **49**, 78; (d) Y. Ye, Z. Ma, R.-B. Lin, R. Krishna, W. Zhou, Q. Lin, Z. Zhang, S. Xiang and B. Chen, *J. Am. Chem. Soc.*, 2019, **141**, 4130; (e) H. Yang, Y. Wang, R. Krishna, X. Jia, Y. Wang, A. N. Hong, C. Dang, H. E. Castillo, X. Bu and P. Feng, *J. Am. Chem. Soc.*, 2020, **142**, 2222.
- (a) Z. Lu, J. Zhang, J. Duan, L. Du and C. Hang, *J. Mater. Chem. A*, 2017, **5**, 17287; (b) H.-X. Zhang, M. Liu, G. Xu, L. Liu and J. Zhang, *Chem. Commun.*, 2016, **52**, 3552; (c) X. Zhao, X. Bu, E. T. Nguyen, Q.-G. Zhai, C. Mao and P. Feng, *J. Am. Chem. Soc.*, 2016, **138**, 15102.
- (a) X. Zhao, X. Bu, Q.-G. Zhai, H. Tran and P. Feng, *J. Am. Chem. Soc.*, 2015, **137**, 1396; (b) S.-T. Zheng, J. J. Bu, T. Wu, C. Chou, P. Feng and X. Bu, *Angew. Chem., Int. Ed.*, 2011, **50**, 8858; (c) X. Wang, W.-Y. Gao, J. Luan, L. Wojtas and S. Ma, *Chem. Commun.*, 2016, **52**, 1971.
- (a) S. M. Cohen, *J. Am. Chem. Soc.*, 2017, **139**, 2855; (b) P. Müller, F. M. Wisser, V. Bon, R. Grüner, I. Senkovska and S. Kaskel, *Chem. Mater.*, 2015, **27**, 2460; (c) S. Yuan, Y.-P. Chen, J.-S. Qin, W. Lu, L. Zou, Q. Zhang, X. Wang, X. Sun and H.-C. Zhou, *J. Am. Chem. Soc.*, 2016, **138**, 8912; (d) C.-X. Chen, Z. Wei, J.-J. Jiang, Y.-Z. Fan, S.-P. Zheng, C.-C. Cao, Y.-H. Li, D. Fenske and C.-Y. Su, *Angew. Chem., Int. Ed.*, 2016, **55**, 9932; (e) H. Kim, D. Kim, D. Moon, Y. N. Choi, S. B. Baek and M. S. Lah, *Chem. Sci.*, 2019, **10**, 5801.
- A. L. Spek, *Acta Crystallogr., Sect. C: Struct. Chem.*, 2015, **71**, 9.