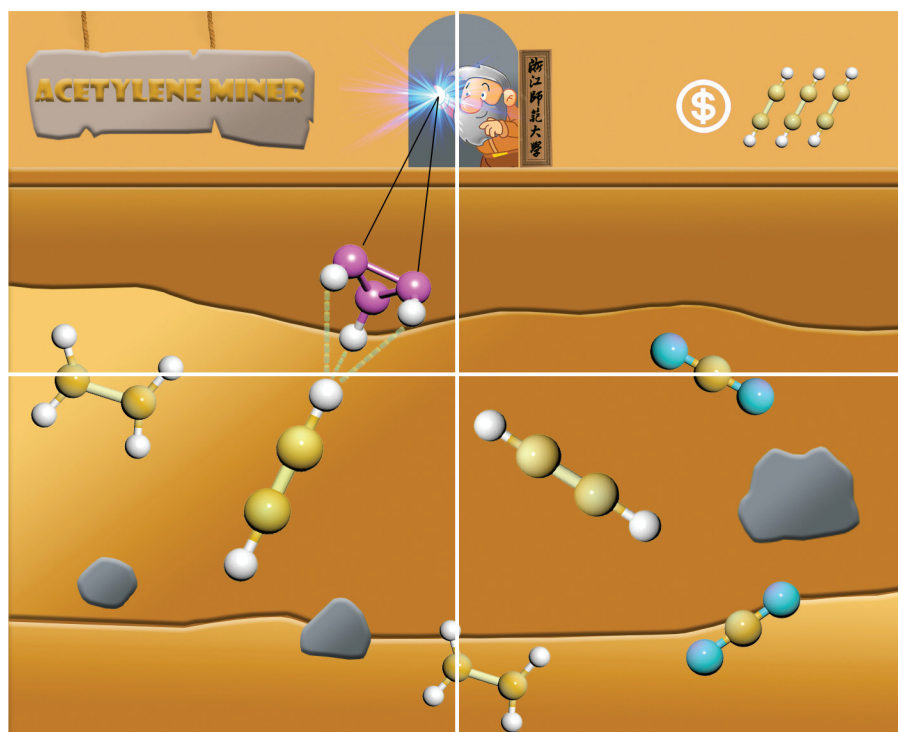




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A new boron cluster anion pillared metal organic framework with ligand inclusion and its selective acetylene capture properties†

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The separation of acetylene (C_2H_2) from carbon dioxide (CO_2) and ethylene (C_2H_4) is important in industry but challenging due to their similar physical properties. Herein, a novel microporous boron cluster pillared metal–organic framework BSF-10 was synthesized with ligand inclusion for efficient C_2H_2/CO_2 and C_2H_2/C_2H_4 adsorption separation. The free dipyriddy ligands in the large pore reduce the porosity of BSF-10 but stabilize the framework. The available narrow pores without inclusion of ligands are suited for the accommodation of C_2H_2 by cooperative dihydrogen bonding. High C_2H_2 capacity and high C_2H_2 selectivity over CO_2 and C_2H_4 are achieved. The practical separation ability was confirmed by the breakthroughs using C_2H_2/CO_2 and C_2H_2/C_2H_4 gas mixtures with good recyclability. The dynamic separation factor of 2.8 for the equimolar C_2H_2/CO_2 mixture is comparable to those of many benchmark materials.

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The separation of C_2H_2/CO_2 and C_2H_2/C_2H_4 to produce C_2H_2 and C_2H_4 in high purity is important in industry.^{1–4} C_2H_2 is a vital fundamental material for the synthesis of various organic chemicals and polymers. Produced from the cracking of hydrocarbons or partial combustion of natural gas, CO_2 is a contaminant and needs to be removed.³ On the other hand, the deep removal of C_2H_2 is a must in the process of ethylene production to obtain polymer grade C_2H_4 (>99.996%) since trace C_2H_2 can poison the ethylene polymerization catalysts by forming metal acetylides.⁴ Current state-of-the-art technologies for the separation and purification of C_2H_2 from other gases largely rely on cryogenic distillation, partial hydrogenation, or solvent extraction, which are either energy-intensive or associated with pollution. When compared, physisorption separation using porous materials is more energy-efficient and eco-friendly.^{1,2} However, due to their similar molecular diameter and polarity (Table S1†), efficient C_2H_2/CO_2 and C_2H_2/C_2H_4 separation is still difficult to achieve.

The design of advanced porous materials is vital for many applications such as gas separation⁵ and catalysis.⁶ Recent years have witnessed families of porous metal organic frameworks (MOFs) with excellent properties for physisorption separations.^{7–16} In this context, inorganic anion pillared MOFs as a unique subclass with strong Lewis basic functional anion sites, developed firstly by the Zaworotko group¹⁷ and Kitagawa group,¹⁸ have been of particular interest for selective gas separation due to their high molecular recognition competence. These inorganic anion pillared MOFs can be classified to five series to date according to the anions used: (1) MFSIX networks pillared by hexafluorometallate anions (*e.g.* SiF_6^{2-} , GeF_6^{2-} , and TiF_6^{2-}),^{19,20} (2) MO_xF_y pillared networks with anions containing octahedral metal centers bonded through both O and F atoms (*e.g.* $NbOF_5^{2-}$ and $TaOF_5^{2-}$),^{21–23} (3) networks cross-linked by tetrahedral MO_4^{2-} oxyanions (*e.g.* CrO_4^{2-} , MoO_4^{2-} , SO_4^{2-}),²⁴ (4) DICRO coordination networks that are sustained with $Cr_2O_7^{2-}$ dianions,²⁵ and (5) boron cluster anion pillared supramolecular metal organic frameworks (BSFs) featuring *closo*- $[B_{12}H_{12}]^{2-}$ or *closo*- $[B_{12}H_{11}I]^{2-}$ as the inorganic pillars.^{26–32}

The icosahedral *closo*-dodecaborate $[B_{12}H_{12}]^{2-}$ is a stable dianionic boron cluster consisting of 12 identical B–H vertices (Fig. 1A).³³ It is considered as a three-dimensional aromatic analogue to two-dimensional aromatic benzene due to the electron delocalization.³⁴ While other anions use electronegative F and O to coordinate transition metals, $[B_{12}H_{12}]^{2-}$ utilizes hydride, which is not common in coordination polymers. In 2019, we reported the first BSF material termed BSF-1, which

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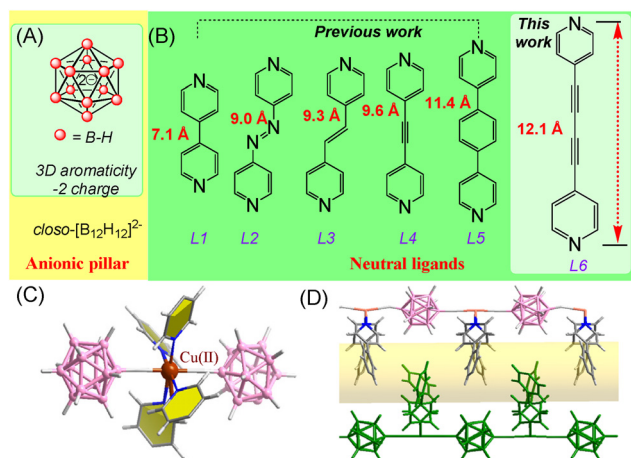


Fig. 1 (A and B) Structure of *closo*-dodecaborate $[B_{12}H_{12}]^{2-}$ and organic ligands. (C) General coordination mode in BSFs. (D) General 1D pore channels of interpenetrated BSFs.

is constructed from *closo*- $[B_{12}H_{12}]^{2-}$ dianions, Cu^{2+} ions and 1,2-di(pyridin-4-yl)ethyne (L4, Fig. 1B) with $N \cdots Cu$ and $B-H \cdots Cu$ coordination as well as $B-H \cdots H-C$ dihydrogen bonds (Fig. 1C).²⁶ Every hexacoordinate Cu center serves as a six-connected node. Four pyridyl units from different ligands comprise the equatorial plane and the axial positions of the Cu nodes coordinate to hydrogen atoms from $[B_{12}H_{12}]^{2-}$, which bridge two Cu nodes to generate an infinite Cu-dodecaborate chain (Fig. 1D). BSF-1 displays high separation selectivity for C_3H_8/CH_4 and C_2H_6/CH_4 . The extension of the organic linker to L5 (Fig. 1B) produces an isostructural MOF BSF-3 with improved porosity, which shows superior C_2H_2 capacity as well as enhanced C_2H_2/C_2H_4 and C_2H_2/CO_2 separation selectivity compared to BSF-1.²⁸ BSF-4 with L2 and BSF-9 with L3 were expected to show similar gas separation performance due to the similar length of organic linkers.^{29,30} However, the interpenetration symmetry difference endows BSF-9 with symmetrical interpenetration mode and a record high C_2H_2/CO_2 separation selectivity in robust MOFs without open metal sites, which is over 5 fold that in BSF-4 with asymmetrical interpenetration mode.³⁰ Another approach to tune the porosity of BSFs is to alter the interpenetration degree. Eliminating the interpenetration of BSFs by controlling the binary solvent system can afford enhanced theoretical porosity. However, these non-interpenetrated MOFs (eg, BSF-5, 6, 7, 8) are actually very unstable and lose most of the porosity after guest removal due to the framework collapse.³⁰ Thus, it is still very challenging to design BSFs with a desirable stable porous structure and the interpenetration mode in BSFs is still difficult to predict and control.

Herein, we would like to report another approach to tune the porosity and stability of BSFs. 1,4-di(pyridin-4-yl)buta-1,3-diyne (L6, Fig. 1B) with a $N \cdots N$ distance of 12.1 Å and negligible steric hindrance was utilized as the organic linker. A new boron cluster anion pillared metal organic framework BSF-10 with ligand inclusion was prepared. BSF-10 is asymmetrically

interpenetrated. There is one large pore with a square shaped window and two small pores with narrow rectangular shaped windows. In the large pore, L6 is included and unable to be removed by soaking into polar solvents or heating under vacuum. This ligand inclusion reduces the porosity of BSF-10 but enhances the stability for practical applications, which has never been reported in BSFs.⁶ The available narrow pores without inclusion of ligands are suited for the accommodation of C_2H_2 by cooperative dihydrogen bonding. Thus, the high capacity of C_2H_2 ($65.0 \text{ cm}^3 \text{ g}^{-1}$) as well as good separation selectivity for C_2H_2/CO_2 (13.52–5.86) and C_2H_2/C_2H_4 (4.09–2.93) under ambient conditions are achieved. The practical separation ability was further confirmed by the dynamic breakthroughs using C_2H_2/CO_2 and C_2H_2/C_2H_4 gas mixtures with good recyclability. The dynamic separation factor of 2.8 for equimolar C_2H_2/CO_2 mixtures is comparable to those of many benchmark materials.

BSF-10 was readily prepared by stirring a mixture of $Na_2[B_{12}H_{12}] \cdot 2H_2O$, $Cu(NO_3)_2 \cdot 3H_2O$ and L6 in MeOH at 35 °C for 48 h. Single crystals of BSF-10 were produced by layering a MeOH solution of L6 onto an aqueous solution of $Na_2[B_{12}H_{12}] \cdot 2H_2O$ and $Cu(NO_3)_2 \cdot 3H_2O$. X-ray structural analysis of BSF-10 revealed that it crystallizes in a three-dimensional (3D) framework in the monoclinic space group $P21/c$ (Table S2†). The asymmetric interpenetration leads to the generation of two kinds of pore channels: large Pore I and small Pores II/III (Fig. 2A). The large Pore I features a square shaped window and the small Pores II/III display narrow rec-

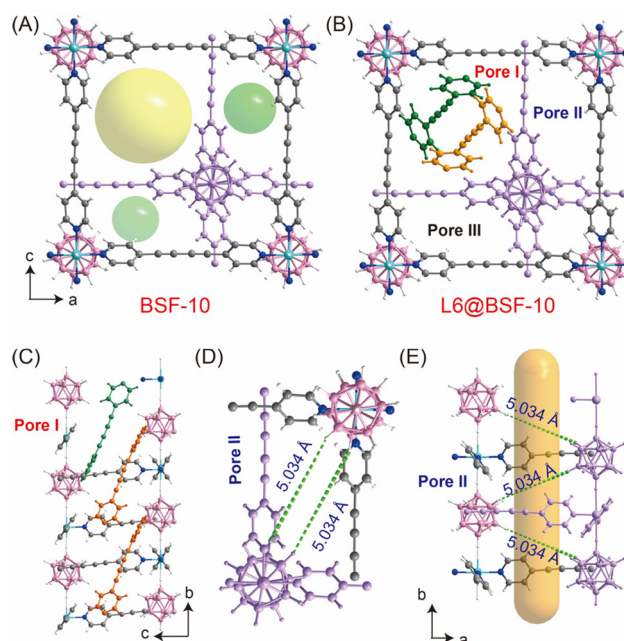


Fig. 2 Porous structure of BSF-10. (A) Generation of two kinds of pores by asymmetrical interpenetration. (B) L6 included in the large Pore I. (C) The packing mode of L6 in Pore I. (D and E) Structure of Pore II in two directions highlighting the closest $B-H \cdots H-B$ distance after subtracting the two van der Waals radii of hydrogen.

tangular shaped windows. Notably, L6 is included in the large pore, and unable to be removed by soaking BSF-10 into polar solvents or heating under vacuum (Fig. 2B). Analysis of the configuration of L6 indicates the existence of two forms of L6 in Pore I with a packing mode of AABB (Fig. 2C). Strong $\pi\cdots\pi$ interactions among guests as well as multiple weak B-H $\cdots$ H-C interaction exist, which stabilize the structure of L6@BSF-10 and prevent the release of L6. Notably, the Cu[B₁₂H₁₂]:L6 ratio has no influence on the structure of BSF-10. Even when excess Cu[B₁₂H₁₂] was used, the single crystals still show the composition of Cu[B₁₂H₁₂](L6)₃ with free L6 inside Pore I (Table S3[†]).

Pore II and Pore III are nearly identical. The closest distance between two opposite B-H units in BSF-10 is 5.034 Å (Fig. 2D and E). This distance is suited to trap an acetylene molecule by cooperative dihydrogen bonding (B-H δ^- $\cdots$ H δ^+ -CC-H δ^+ $\cdots$ H δ^- -B)²⁸ and thus be potential for C₂H₂/CO₂ and C₂H₂/C₂H₄ separation. Besides, the pore window size of *ca.* 4 Å (Fig. 3) defined by the opposite H(Py) $\cdots$ H(Py) is also close to the diameter of C₂H₂ (kinetic diameter 3.3 Å, 3D diameter: 3.32 × 3.34 × 5.70 Å³), allowing tight C₂H₂ accommodation.

Inspired by the structural analysis, we were interested to investigate the potential of BSF-10 for the selective adsorption separation of C₂H₂ from CO₂ and C₂H₄. The first step is to confirm the permanent porosity of BSF-10 after solvent removal as BSFs with larger pores are not stable upon activation.³¹ N₂ gas adsorption experiments at 77 K were conducted after activating BSF-10 under vacuum at 75 °C for 12 h. A typical Type I isotherm was observed, indicating the microporous character of BSF-10. The BET surface area was calculated to be 426.3 m² g⁻¹ with a total pore volume of 0.216 cm³ g⁻¹ at *P/P*₀ = 0.95, (Fig. 4A), which is quite close to the calculated pore volume of 0.222 cm³ g⁻¹ based on the single crystal structure of BSF-10 using a probe with the radius of 1.2 Å (Fig. 3).

The establishment of permanent microporosity in BSF-10 motivated us to study the gas separation performance for C₂H₂/CO₂ and C₂H₂/C₂H₄. First of all, single component adsorption isotherms of C₂H₂, CO₂ and C₂H₄ were collected

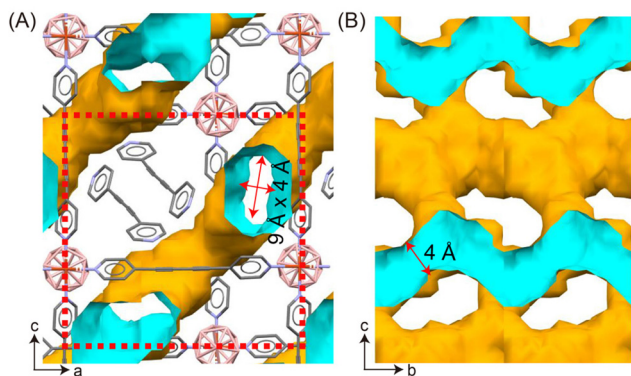


Fig. 3 Voids and pore windows of BSF-10 generated by a probe with the radius of 1.2 Å seen along (A) *b*-axis and (B) *a*-axis. The pore window dimensions are determined after subtracting the two van der Waals radii of hydrogen.

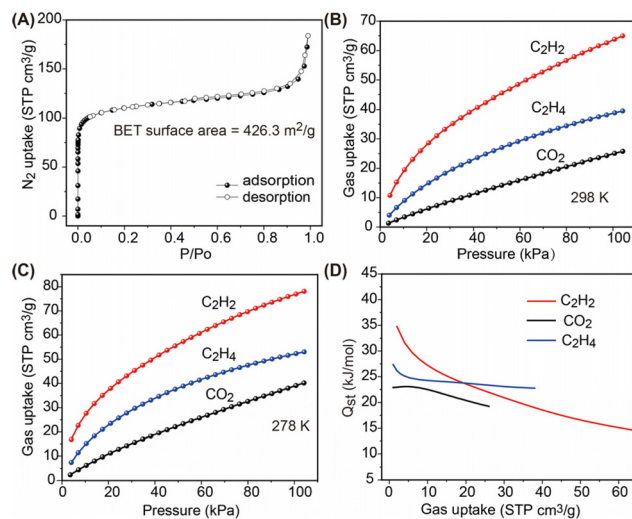


Fig. 4 (A) N₂ adsorption and desorption isotherms of BSF-10. (B and C) Single component adsorption isotherms of C₂H₂, CO₂ and C₂H₄ on BSF-10 at 298 and 278 K. (D) Adsorption heats of C₂H₂, CO₂ and C₂H₄ on BSF-10.

under 298 and 278 K. At 1.0 bar, the C₂H₂/CO₂/C₂H₄ uptakes were 65.0/25.8/39.5 cm³ g⁻¹ at 298 K and 78.1/53.0/40.2 cm³ g⁻¹ at 278 K, respectively (Fig. 4B and C). The isosteric heat of adsorption (*Q*_{st}) for BSF-10 was then calculated using the Clausius–Clapeyron equation after fitting the isotherms to the Langmuir–Freundlich equation with excellent accuracy (*R*² > 0.9999, Tables S4 and 5[†]). *Q*_{st} values at near-zero loading for C₂H₂, CO₂, and C₂H₄ were 34.8, 27.4, and 22.9 kJ mol⁻¹ (Fig. 4D). This trend of *Q*_{st} values is consistent with the slope of the adsorption isotherms, indicating the preferential adsorption of C₂H₂ over CO₂ and C₂H₄.

Additionally, the *Q*_{st} value for C₂H₂ in BSF-10 is higher than that of BSF-1 (30.7 kJ mol⁻¹), but lower than that of BSF-3 (42.7 kJ mol⁻¹), consistent with the C₂H₂ adsorption uptake trend of BSF-1 (2.35 mmol g⁻¹) < BSF-10 (2.90 mmol g⁻¹) < BSF-3 (3.59 mmol g⁻¹). Notably, this modest *Q*_{st} renders BSF-10 a suitable porous material for practical application with a low energy footprint for regeneration.

Since selectivity is as important as capacity to evaluate the separation performance, the C₂H₂/CO₂ and C₂H₂/C₂H₄ selectivity on BSF-10 at 298 K and 278 K was calculated using ideal adsorbed solution theory (IAST), which revealed that the selectivity for equimolar C₂H₂/CO₂ and C₂H₂/C₂H₄ at 298 K and 1.0 bar was 13.52–5.86 and 4.09–2.93, respectively (Fig. 5A). The selectivity is slightly increased under low pressure when the temperature reduced to 278 K, and is 17.4–5.2 for C₂H₂/CO₂ and 4.8–2.8 for C₂H₂/C₂H₄ (Fig. 5B). The C₂H₂/CO₂ selectivity with different C₂H₂ molar ratios was further calculated for BSF-10, which indicates that the decrease of the C₂H₂ molar ratio leads to the increased C₂H₂/CO₂ selectivity (Fig. 5C). The C₂H₂ capacity and the C₂H₂/CO₂ selectivity on BSF-10 are superior to those of many well-performing materials such as BSF-1 (52.5 cm³ g⁻¹, *S* = 3.4),²⁶

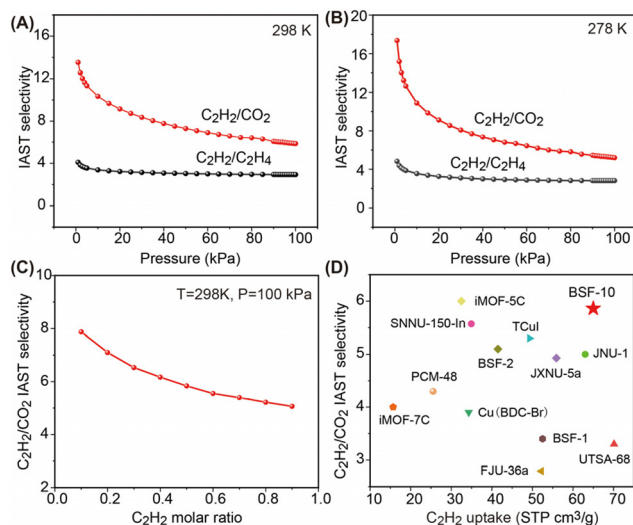


Fig. 5 (A and B) IAST selectivity of C_2H_2/CO_2 and C_2H_2/C_2H_4 at 298 and 278 K. (C) IAST selectivity of C_2H_2/CO_2 with different C_2H_2 molar ratios. (D) Comparison of the C_2H_2 uptakes and C_2H_2/CO_2 selectivity among BSF-10 and others well-performing MOFs.

BSF-2 ($41.5 \text{ cm}^3 \text{ g}^{-1}$, $S = 5.1$),²⁷ TCuI ($49.3 \text{ cm}^3 \text{ g}^{-1}$, $S = 5.3$),³⁵ JNU-1 ($63 \text{ cm}^3 \text{ g}^{-1}$, $S = 5$),³⁶ iMOF-7C ($15.7 \text{ cm}^3 \text{ g}^{-1}$, $S = 4$),³⁷ FJU-36a ($52.2 \text{ cm}^3 \text{ g}^{-1}$, $S = 2.8$),³⁸ SNNU-150-In ($34.9 \text{ cm}^3 \text{ g}^{-1}$, $S = 5.6$),³⁹ PCM-48 ($25.5 \text{ cm}^3 \text{ g}^{-1}$, $S = 4.3$),⁴⁰ and JXNU-5a ($55.9 \text{ cm}^3 \text{ g}^{-1}$, $S = 5$),⁴¹ (Fig. 5D), but inferior to those of BSF-3 ($80.4 \text{ cm}^3 \text{ g}^{-1}$, $S = 16.3$).²⁸ The C_2H_2/CO_2 uptake ratio of 2.52 at 298 K and 1 bar is also relatively high. This value is superior to those of most MOFs that separate C_2H_2/CO_2 by a thermodynamic mechanism,^{42–52} but slightly lower than those of $Cu^I@UiO66-(COOH)_2$ (2.58),⁴³ MAF-2 (3.68),⁴⁷ and SIFSIX-21-Ni (3.1)^{17c} as shown in Fig. 6 and Table S8.†

To confirm the practical separation ability of BSF-10 for C_2H_2/CO_2 and C_2H_2/C_2H_4 mixtures, dynamic breakthrough experiments were conducted. 0.233 g of BSF-10 powder was packed into a stainless column with a size of $\Phi 4.6 \text{ mm} \times 5 \text{ cm}$. After activating the sample by Ar purge at 75°C for 12 h, an equimolar C_2H_2/CO_2 mixture was introduced. CO_2 appeared at the outlet within 12 min while C_2H_2 was retained in the

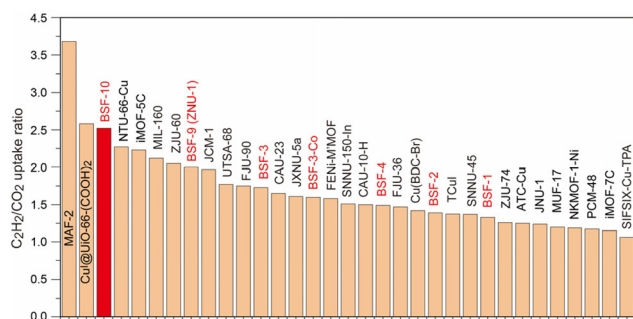


Fig. 6 Comparison of the C_2H_2/CO_2 uptake ratios at 298 K and 1 bar among BSF-10 and other well-performing materials in the context of C_2H_2/CO_2 separation by a thermodynamic mechanism.

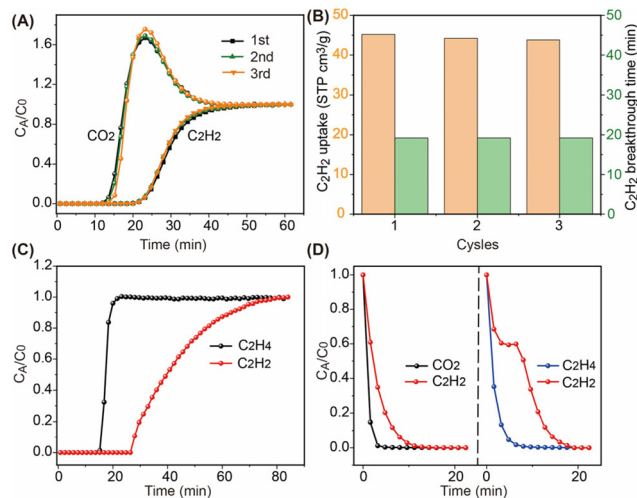


Fig. 7 (A) Experimental breakthrough curves of C_2H_2/CO_2 (50/50) on BSF-10 at 298 K. (B) Comparison of the dynamic C_2H_2 uptake from the C_2H_2/CO_2 (50/50) mixture and breakthrough time among three cycles of breakthrough experiments. (C) Experimental breakthrough curves of C_2H_2/C_2H_4 (1/99) on BSF-10 at 298 K. (D) The desorption curves of BSF-10 after breakthrough experiments of C_2H_2/CO_2 (50/50) mixtures and C_2H_2/C_2H_4 (1/99) mixtures.

column until 20 min (Fig. 7A). After complete breakthrough, Ar gas with a flow rate of 5 mL min^{-1} was used to desorb the adsorbed C_2H_2 and CO_2 gases as well as to regenerate the material. Notably, nearly all the C_2H_2 and CO_2 gases are desorbed within 20 min at 75°C and the material is regenerated for further use (Fig. 7D). Calculation of the areas of the breakthrough curves and desorption curves indicated that BSF-10 has a dynamic C_2H_2 capacity of $45 \text{ cm}^3 \text{ g}^{-1}$ with a C_2H_2/CO_2 separation factor of 2.8. This separation factor is slightly lower than that of benchmark materials such as CAU-10-H (3.4),⁴² $Cu^I@UiO-66-(COOH)_2$ (3.4),⁴³ and JCM-1 (4.4),⁴⁴ but higher than that of NKMOF-1-Ni (2.6),⁴⁵ FJU-90 (2.1)⁴⁶ and HOF-3a (2).⁵³ Besides, the dynamic C_2H_2 capacity of BSF-10 is also comparable to that of JCM-1 ($49.3 \text{ cm}^3 \text{ g}^{-1}$). The regenerated BSF-10 was further used for repeated tests. Nearly no performance reduction was observed over 3 cycles (Fig. 7B). The same sample was further used for C_2H_2/C_2H_4 (1/99) mixture breakthrough experiments. C_2H_4 was detected at 15 min and C_2H_2 at 28 min, suggesting a good capture ability of trace C_2H_2 from bulky C_2H_4 (Fig. 7C). Desorption curves further indicated that the captured C_2H_2 can be desorbed within 20 min by Ar purge at 75°C (Fig. 7D). Combining the high dynamic C_2H_2 capacity, high separation factor as well as facile regeneration conditions, BSF-10 is a good candidate for practical C_2H_2/CO_2 and C_2H_2/C_2H_4 separation.

In conclusion, we prepared a novel microporous boron cage pillared metal-organic framework BSF-10 with unprecedented ligand inclusion. Free dipyrindyl ligands occupy the large pores and interact with the framework by multiple interactions, which reduces the porosity of BSF-10 but enhances the stability of the framework. The narrow pores without inclusion of

ligands are suited for the selective accommodation of C_2H_2 by cooperative dihydrogen bonding. Thus, high C_2H_2 capacity and high C_2H_2 selectivity over CO_2 and C_2H_4 are achieved. The practical separation ability was completely confirmed by the breakthroughs using C_2H_2/CO_2 and C_2H_2/C_2H_4 gas mixtures with good recyclability and facile regeneration conditions. The dynamic separation factor of 2.8 for the equimolar C_2H_2/CO_2 mixture is comparable to those of many benchmark materials. Boron cage pillared metal-organic frameworks constructed by non-linear organic linkers are under exploration. Besides, MOFs with dual C_2H_2/C_2H_4 and C_2H_2/CO_2 separation ability will be increasingly important for separating multicomponent gas mixtures using a single adsorbent, which currently largely relies on synergistic/tandem packed adsorbents in fixed-beds.⁵⁴

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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