Simplifying and expanding the scope of boron imidazolate framework (BIF) synthesis using mechanochemistry†

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Mechanochemistry† has emerged as a versatile methodology for the synthesis and discovery of advanced materials, including nanoparticle systems and metal–organic frameworks (MOFs), giving rise to materials that are challenging to obtain using conventional solution-based techniques. Mechanochemical techniques such as ball milling, twin screw extrusion and acoustic mixing have simplified and advanced the synthesis of a wide range of MOFs, permitting the use of simple starting materials such as metal oxides, hydroxides or carbonates, at room temperature and without bulk solvents, yielding products of comparable stability and, after activation, higher surface areas than solution-generated counterparts. The efficiency of mechanochemistry in MOF synthesis was recently highlighted by accessing zeolitic imidazolate frameworks (ZIFs) that were theoretically predicted, but not accessible under conventional solution-based conditions.

The advantages of mechanochemistry in MOF chemistry led us to address the possibility of synthesizing boron imidazolate frameworks (BIFs), an intriguing but poorly developed class of microporous materials analogous to ZIFs, comprising equimolar combinations of tetrahedrally coordinated boron(III) and monovalent Li⁺ or Cu⁺ cations as nodes (Fig. 1A–C). Although BIFs offer an attractive opportunity to access microporous MOFs with lower molecular weights, particularly in the case of “ultralight” systems based on Li⁺ and B(III) centers, this family of materials has remained largely unexplored—potentially due to the stability of the open SOD-framework relative to the non-porous dia-polymorph.

Fig. 1 Structures of previously reported BIFs with: (A) zni-, (B) dia-, or (C) SOD-topology (M = Li, Cu); (D) tetrakis(imidazolyl)boric acids used herein for mechanochemical BIF synthesis; and (E) schematic representation of the herein developed mechanosynthesis of dia- and SOD BIF polymorphs based on Li, Cu or Ag metal nodes.
to the need for harsh synthetic conditions, including the use of n-butyllithium in a solvothermal environment.23–34

We now show how switching to the mechanochemical environment enables lithium- and copper(i)-based BIFs to be prepared rapidly (i.e., within 60–90 minutes), without elevated temperatures or bulk solvents, and from readily accessible solid reactants, such as hydroxides and oxides (Fig. 1D and E). While the mechanochemically-prepared BIFs exhibit significantly higher surface areas than the solvothermally-prepared counterparts, mechanochemistry allows for expanding this class of materials towards previously not reported $\text{Ag}^{+}$ nodes. The introduction of BIFs isostructural with those based on $\text{Li}^{+}$ or $\text{Cu}^{+}$ but comprising of $\text{Ag}^{+}$ ions, enables a periodic density-functional theory (DFT) evaluation of their stability. This reveals that switching to heavier elements as tetrahedral nodes improves the stability of sodalite topology (SOD) open BIFs with respect to close-packed diamondoid (dia) topology polymorphs.

As a first attempt at mecanchochemical synthesis of BIFs, we targeted the synthesis of previously reported zni-topology $\text{LiB}[\text{Im}]_4$ and $\text{CuB}[\text{Im}]_4$ frameworks (Li-BIF-1 and Cu-BIF-1, respectively, Fig. 1A) using a salt exchange reaction between $\text{LiCl}$ or $\text{CuCl}$ with commercially available sodium tetrakis(imidazolyl)borate ($\text{Na[B[Im]}_4]$) (Fig. 2A). Milling of $\text{LiCl}$ and $\text{Na[B[Im]}_4]$ in a 1 : 1 stoichiometric ratio for up to 60 minutes led to the appearance of Bragg reflections consistent with the target Li-BIF-1 (CSD MOXJEP) and the anticipated NaCl byproduct. The reaction was, however, incomplete, as seen by X-ray reflections of $\text{Na[B[Im]}_4]$ starting material. In order to improve reactant conversion, we explored liquid-assisted grinding (LAG), i.e. milling in the presence of a small amount of a liquid phase (measured by the liquid-to-solid ratio $\eta$ in the range of $0.25–2 \text{ mL g}^{-1}$). Using LAG conditions with acetonitrile ($\text{MeCN, 120 }\mu\text{L, }\eta = 0.5 \text{ mL mg}^{-1}$) led to the complete disappearance of reactant X-ray reflections, concomitant with the formation of Li-BIF-1 alongside NaCl within 60 minutes.

Next, we explored an alternative synthesis approach, analogous to that previously used to form ZIFs and other MOFs: an acid–base reaction between a metal oxide or hydroxide and the acid form of the linker: tetrakis(2-methylimidazole)boric acid, $\text{HB}[\text{Im}]_4$ (Fig. 3A).36–40 Neat milling $\text{LiOH}$ with one equivalent of $\text{HB}[\text{Im}]_4$ in a stainless steel milling assembly led to the partial formation of Li-BIF-1, as evidenced by PXRD analysis (see ES†). Complete conversion of reactants into Li-BIF-1 was achieved in 60 minutes by LAG with $\text{MeCN (}}\eta = 0.25 \text{ mL mg}^{-1}$), as indicated by PXRD analysis (Fig. 3B–E), Fourier transform infrared attenuated total reflectance spectroscopy (FTIR-ATR), thermogravimetric analysis (TGA) in air, and analysis of metal content by inductively-coupled plasma mass spectrometry (ICP-MS) (see ES†).

Neat milling of H[B[Meim]]4 with Cu2O under similar conditions gave a largely non-crystalline material, as evidenced by PXRD (see ES†). Switching to the ion- and liquid-assisted grinding (ILAG) methodology, in which the reactivity of a metal oxide is enhanced by a small amount of a weakly acidic ammonium salt, and which was introduced to prepare zinc and cadmium ZIFs from respective oxides,37–40 enabled the synthesis of Cu-BIF-1 from Cu2O. Specifically, PXRD analysis revealed complete disappearance of the oxide in samples obtained by ILAG with either $\text{MeOH}$ or $\text{MeCN (}}\eta = 0.5 \text{ mL mg}^{-1}$) in the presence of $\text{NH}_4\text{NO}_3$ additive (5% by weight, see ES†). Notably, achieving complete disappearance of Cu2O reactant signals also required switching from stainless steel to a zirconia-based milling assembly, presumably due to more efficient energy delivery.41 After washing with MeOH, the material was characterized by FTIR-ATR, TGA in air, and analysis of metal content by ICP-MS (see ES†). Whereas both the metathesis and acid–base approaches can be used to mecanchochemically generate Li- and Cu-BIF-1, the latter approach has a clear advantage of circumventing the formation of the NaCl byproduct. Consequently, in order to further the development of mecanchochemical routes to other BIFs, we focused on the acid–base strategy. As next targets, we turned to MOFs based on tetrakis[2-methylimidazole]boric acid $\text{H[B[Meim]}_4]$,46 previously reported to adopt either a non-porous diamondoid (dia) topology (BIF-2) or a microporous sodalite (SOD) topology (BIF-3) with either $\text{Li}^{+}$ or $\text{Cu}^{+}$ as nodes.
Attempts to selectively synthesize either Li-BIF-2 or Li-BIF-3 by neat milling or LAG (using MeOH or MeCN as liquid additives) with LiOH and a stoichiometric amount of HB(Meim)$_4$ were not successful. Exploration of different milling times and $\eta$-values produced only mixtures of residual reactants with Li-BIF-2, Li-BIF-3, and/or not yet identified phases (see ESI†). Consequently, we explored milling in the presence of 2-aminobutanol (amb), which is a ubiquitous component of solvent systems used in the solvothermal syntheses of BIFs.\textsuperscript{32,33} Gratifyingly, using a mixture of amb and MeCN in a 1 : 3 ratio by volume as the milling liquid led to an effective strategy for the selective synthesis of both the dia-topology Li-BIF-2 (CSD code MOXKUG), and the SOD-topology Li-BIF-3 (CSD code MUCLOM).\textsuperscript{§} The selective formation of phase-pure samples of Li-BIF-2 and Li-BIF-3 was confirmed by PXRD analysis, which revealed an excellent match to diffractionograms simulated based on the previously reported structures (Fig. 4B–G). Systematic exploration of reaction conditions, including time (between 15 and 60 minutes) and $\eta$ value (between 0.25 and 1 $\mu$L mg$^{-1}$) revealed that the open framework Li-BIF-3 is readily obtained at $\eta$ either 0.75 or 1 $\mu$L mg$^{-1}$ after milling for 45 minutes or longer (Fig. 4B–G, also see ESI†).\textsuperscript{†} Lower $\eta$-values of 0.25 and 0.5 $\mu$L mg$^{-1}$ preferred the formation of the dia-topology Li-BIF-2, which was obtained as a phase-pure material upon 60 minutes milling at $\eta = 0.5 \mu$L mg$^{-1}$, following the initial appearance of a yet unidentified intermediate. The preferred formation of Li-BIF-2 at lower $\eta$-values is consistent with our previous observations that lower amounts of liquid promote mechanochemical formation of denser MOF polymorphs.\textsuperscript{37}
samples made solvothermally (762.5 m² g⁻¹) (Table 1). Analysis of Li-BIF-3 by scanning electron microscopy (SEM) revealed particles with sizes of hundreds of nanometres, forming larger aggregates several micrometres across (Fig. 5A, inset).

The analogous copper(i)-based BIF-2 and BIF-3 frameworks were readily accessible by ILAG, by controlling the volume of the liquid additive and milling time (Fig. 4H–L, also see ESI†). Similarly to our previous studies of ZIFs,¹⁷,²⁴,³⁷,³⁹ increased milling times preferred the formation of the close-packed polymorph, dia-topology Cu-BIF-2. While the PXRD pattern of the reaction mixture after 60 minutes ILAG with MeOH (η = 0.5 µL mg⁻¹) and NH₄NO₃ (5% wt/wt) indicated the presence of the SOD-topology Cu-BIF-3, longer milling led to the appearance of the dia-phase (see ESI†). The materials were identified through comparison of experimental PXRD patterns to those simulated from published structures (CSD codes MUCLIG and MOXJOZ for Cu-BIF-2 and Cu-BIF-3, respectively).³² Quantitative synthesis of Cu-BIF-2 from Cu₂O was readily accomplished by ILAG for 90 minutes (Fig. 4H–L). Following washing and drying, the products were characterized by PXRD, FTIR-ATR, TGA in air and ICP-MS elemental analysis of metal content.

In order to achieve the synthesis of phase-pure microporous Cu-BIF-3, reaction conditions were modified by increasing η to 1 µL mg⁻¹. This modification enabled the reproducible and quantitative synthesis of Cu-BIF-3 in 60 minutes milling (Fig. 4H–L), confirmed by PXRD, FTIR-ATR, TGA and elemental analysis of metal content (see ESI†). Analyses by SEM and nitrogen sorption were performed on the mecanochemically prepared product after washing and drying in vacuo at 85 °C, revealing that the sample consists of sub-micron particles and exhibits a high BET surface area of 935 m² g⁻¹, which is close to the theoretically expected value of 1100 m² g⁻¹ (Table 1 and Fig. 5B).

To verify whether the improvement in surface area was attributable to the mecanochemical synthesis or the different activation procedure, a sample of Cu-BIF-3 was synthesized according to the published solvothermal methodology and activated in the same way as the mecanochemically prepared

Table 1 Experimental Brunauer–Emmett–Teller (BET) and Langmuir surface area (in m² g⁻¹) of mecanochemically synthesized SOD-topology BIFs, compared to previously measured and theoretically calculated values, along with average particle sizes (in nm) established by SEM and calculated energies (in eV) for all Li-, Cu-, and Ag-BIF polymorphs. The difference between calculated energies for SOD- and dia-polymorphs in each system is given as ΔE (in kJ mol⁻¹)

| Material      | Surface area (m² g⁻¹) | Particle size (nm) | Electronic energy per formula unit (eV) | ΔE (kJ mol⁻¹) |
|---------------|------------------------|--------------------|-----------------------------------------|-------------|
|               | Mechochemical, BET      | Mechochemical, Langmuir | Prior work, Langmuir ¹² | Theoretical² |                        |
| dia-Li-BIF-2  | —                      | —                  | —                                       | —           | —                      |
| SOD-Li-BIF-3  | 1010                   | 1060               | 762.5                                   | 1200        | 217 (π = 24)           | —2679.174 | 14.25          |
| dia-Cu-BIF-2  | —                      | —                  | —                                       | —           | —                      | —3417.091 | 9.67           |
| SOD-Cu-BIF-3  | 935                    | 1196               | 182.3                                   | 1100        | 611 (π = 500)          | —3416.991 |               |
| dia-Ag-BIF-2  | —                      | —                  | —                                       | —           | —                      | —4738.959 | 8.66           |
| SOD-Ag-BIF-3  | 1020                   | 1205               | —                                       | 1170        | 500 (π = 25)           | —4738.869 |

Caclculated using MOF Explorer (see ESI). Determined from SEM measurements, where n corresponds to number of particles observed.

Fig. 5 BET adsorption plots for: (A) Li-BIF-3, showing a surface area of 1010 m² g⁻¹ and (B) Cu-BIF-3, showing a surface area of 935 m² g⁻¹. The insets in (A) and (B) are representative SEM images of the mecanochemically prepared BIF samples, with scale bars corresponding to 4 µm and 5 µm shown in white.
material. The resulting Cu-BIF-3 exhibited a BET surface area of 616 m² g⁻¹ (see ESI†), i.e. 34% lower compared to the mechanochemically synthesized sample, illustrating a clear benefit of mechanochemistry in providing a simpler, more efficient synthesis, as well as materials of improved porosity.32,42

The mechanochemical approaches to Li- and Cu-based BIFs are surprisingly simple compared to previously reported solvothermal methods,32,41,42 not only avoiding bulk solvents and high temperatures (85 °C for Li-based, 120 °C for Cu-based BIFs), but also enabling the use of simple, easily handled solids LiOH and Cu₂O as starting materials compared to, for example, n-BuLi.42 Notably, while the reported solvothermal synthesis of these materials also requires the use of amb for the preparation of both Li- and Cu-BIFs, the use mechanochemical conditions enabled amb-free synthesis of copper-based BIFs. Such simplifications of the synthetic procedure encouraged us to explore the possibility to extend this family of materials towards previously not reported silver(i) derivatives.

As a starting material for the synthesis of Ag(i)-based BIFs we focused on Ag₂CO₃, generated in situ from readily accessible AgNO₃ and K₂CO₃. One-pot milling reaction of HB[Meim]₄, AgNO₃, and K₂CO₃ in the respective stoichiometric ratios 1:1:1/2, using MeCN as the milling additive (n = 0.25 μL mg⁻¹) readily produced the targeted AgB[Meim]₄ material along with the side product KNO₃ (Fig. 5, also see ESI†). Specifically, analysis of the reaction mixtures by PXRD revealed that, similar to the lithium and copper(i) analogues,32 the silver-based BIF appears in two polymorphs which could be selectively synthesized by varying the milling time. The BIF products were readily separated from the KNO₃ by-product after sequential washing with cold MeOH and acetone, and their respective structures were further validated by structure determination from PXRD data measured on washed and dried materials.

Specifically, milling for 30 minutes led to the formation of a material (Ag-BIF-3) which, based on PXRD analysis, was isostructural to the SOD-topology Li-BIF-3 and Cu-BIF-3. Consequently, the crystal structure of Ag-BIF-3 (Fig. 5A) was determined through Rietveld refinement of a structural model based on the Cu-BIF-3 structure, in which the copper(i) sites have been replaced by Ag(i), giving rise to a cubic unit cell (space group P43n as in the analogous Cu-BIF-3 and Li-BIF-3 structures) with a = 16.6659(3) Å. Composition of Ag-BIF-3 was verified by TGA/DSC and elemental analysis of metal content (see ESI†). The previously published crystal structures of Li- and Cu-BIFs with Meim⁻ links, as well as the structures of Ag-BIFs herein determined, were geometry-optimized using the PBE⁷ functional combined with many-body dispersion (MBD)⁴* correction scheme. The PBE + MBD* approach has previously shown excellent agreement with experimental calorimetric measurements of ZIF polymorphs,²⁴ therefore we expected the same approach to perform reliably for the structures of BIFs. In addition to calculating the relative energies of SOD-and dia-polymorphs, we have performed Gauge Including Projector Augmented Waves (GIPAW)⁴³ simulation of the solid-state NMR spectra of Ag-BIFs to compare the simulated spectra with their experimental counterparts, confirming the low symmetry Ag-BIF-2 structure derived from PXRD data (Fig. 6D–G).

Comparison of calculated energies reveals that increasing the atomic number of the metal node results in increased stabilization of the SOD-topology open framework with respect to the close-packed dia-polymorph. The energy differences (ΔE) between SOD- and dia-topology polymorphs for each pair of Li-, Cu-, and Ag-based frameworks are shown in Table 1. Specifically, whereas Li-BIF-3 structure is calculated to be ca. 14 kJ mol⁻¹ less stable compared to its dia-counterpart, the corresponding differences for Cu-BIF-3 and Ag-BIF-3 are calculated to be considerably smaller, at 9.7 kJ mol⁻¹ and 8.7 kJ mol⁻¹, respectively. The calculated energies are in a similar range to the calculated (15.2 kJ mol⁻¹) and measured (10.6 kJ mol⁻¹) ones for the analogous zinc-based ZIFs, indicating that switching between transition metal and main group-based nodes, or switching between using only a divalent node and a combination of monovalent and trivalent nodes, does not significantly influence the difference in stability of SOD- and dia-topology frameworks. The improvement in the relative
stability of open BIF structure upon switching from Li+ to Cu+ and Ag+ nodes is notable, considering the recent interest in the development of MOFs based on heavy-atom nodes.52–54

The simulated ssNMR spectra of Ag-BIF-2 and Ag-BIF-3 showed excellent agreement with the experiment (Fig. 6) in terms of overall chemical shift and the number of distinct NMR signals arising from the crystallographic symmetry. The spectrum of the SOD polymorph is consistent with a single symmetrically unique Meim linker, while the signal splitting found in the spectrum of the dia-polymorph corresponds to four distinct 2-methylimidazolate units. The NMR simulation fully supports the structural models derived from PXRD data, with calculated chemical shifts underlining the accuracy of the herein used theoretical approach.

Conclusions

In summary, we have shown that the application of mechanochemical methodologies can greatly simplify the synthesis of zeolitic boron imidazolate frameworks, providing a rapid, room-temperature approach to this class of materials. Contrary to previously reported materials obtained by solvothermal synthesis, the open framework SOD-topology BIFs made mechanochemically show higher surface areas, that are very close to the theoretically calculated ones. The use of mechanochemistry also expanded the scope of the zeolitic BIF class of materials, by enabling the synthesis of previously not reported silver-based systems. Theoretical investigation of isostructural BIFs based on Li+, Cu+ and Ag+ reveals that the use of increasingly heavier tetrahedral metal nodes leads to the stabilization of open-framework SOD-topology structure with respect to the corresponding close-packed dia polymorph. Overall, the presented results and synthetic methodologies should provide simpler, faster access to an intriguing, but so far poorly developed, class of ultralight zeolitic MOFs.

Data availability

Details of experimental and theoretical procedures, and selected PXRD, TGA, FTIR-ATR, ICP-MS, and nitrogen sorption data are provided as part of the ESL† Crystallographic data for compounds BIF-2-Ag, BIF-3-Ag and [Hamb]B(Meim)4−·2H2O have been deposited at the Cambridge Crystallographic Data Centre (CCDC) under deposition codes 2081806, 2081807, and 2084944, respectively.

Author contributions

All authors have contributed to the writing of this manuscript. Development of mechanochemical synthesis procedures and

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Fig. 6 (A) Rietveld refinement of Ag-BIF-3 with difference plot shown in grey. (B) Rietveld refinement of Ag-BIF-2 with difference plot shown in grey. (C) BET adsorption plot Ag-BIF-3 showing a surface area of 1020 m² g⁻¹ and a SEM image of a representative sample (scale-bar 1 μm). Comparison of measured and simulated 13C CP-MA ssNMR spectra for silver-based BIFs: (D) calculated for Ag-BIF-3, (E) measured for Ag-BIF-3, (F) calculated for Ag-BIF-2 and (G) measured for Ag-BIF-2.
characterization of new materials was performed by C. B. L., J. G. C., J.-L. D. The materials characterization was done by H. M. T., A. J. H., and theoretical studies by M. A. The research was organized and coordinated by T. F.

Conflicts of interest
There are no conflicts to declare.

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Notes and references
§ The use of amb in the synthesis of bifurcates has not been explained. Based on the herein reported mechanocatalytic studies, we believe amb provides a means to increase the reactivity of the HB[Meim], ligand, potentially through salt formation, and also hinders the re-arrangement of open-structure Li-BIF-3 into Li-BIF-2. For example, 30 minutes milling of LiOH and HB[Meim] in presence of MeCN at $\eta = 1 \mu M$ leads to a mixture of Li-BIF-2, Li-BIF-3 and HB[Meim], which is evident by PXRD analysis. Milling in presence of a mixture of MeCN and amb (3 : 1 by volume, $\eta = 1 \mu M$), however, leads to complete disappearance of PXRD signals of HB[Meim] and Li-BIF-3 as the only product.

† In some cases during screening reaction conditions, the PXRD pattern of the product exhibited Bragg reflections that could not be assigned to neither Li-BIF-2, Li-BIF-3, or any of the reactants. These were subsequently found to result from the parasitic formation of the hydrated salt [Hamb][B(Meim)]$^+$ $2H_2O$, whose structure was determined from separately prepared single crystals (see ESI† CCDC deposition code 208494). The formation of [Hamb][B(Meim)]$^+$ $2H_2O$, however, can be avoided by drying both MeCN and amb over molecular sieves before use.

‡ Preliminary attempts to synthesize the Ag-BIFs from acetonitrile solution, using the methodology that was reported for Cu-BIF-3, gave a yet unidentified material. Replacing the solvent with MeOH gave a mixture of phases containing Ag-BIF-2, along with significant amount of metallic silver, evident by the black color of the material.

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