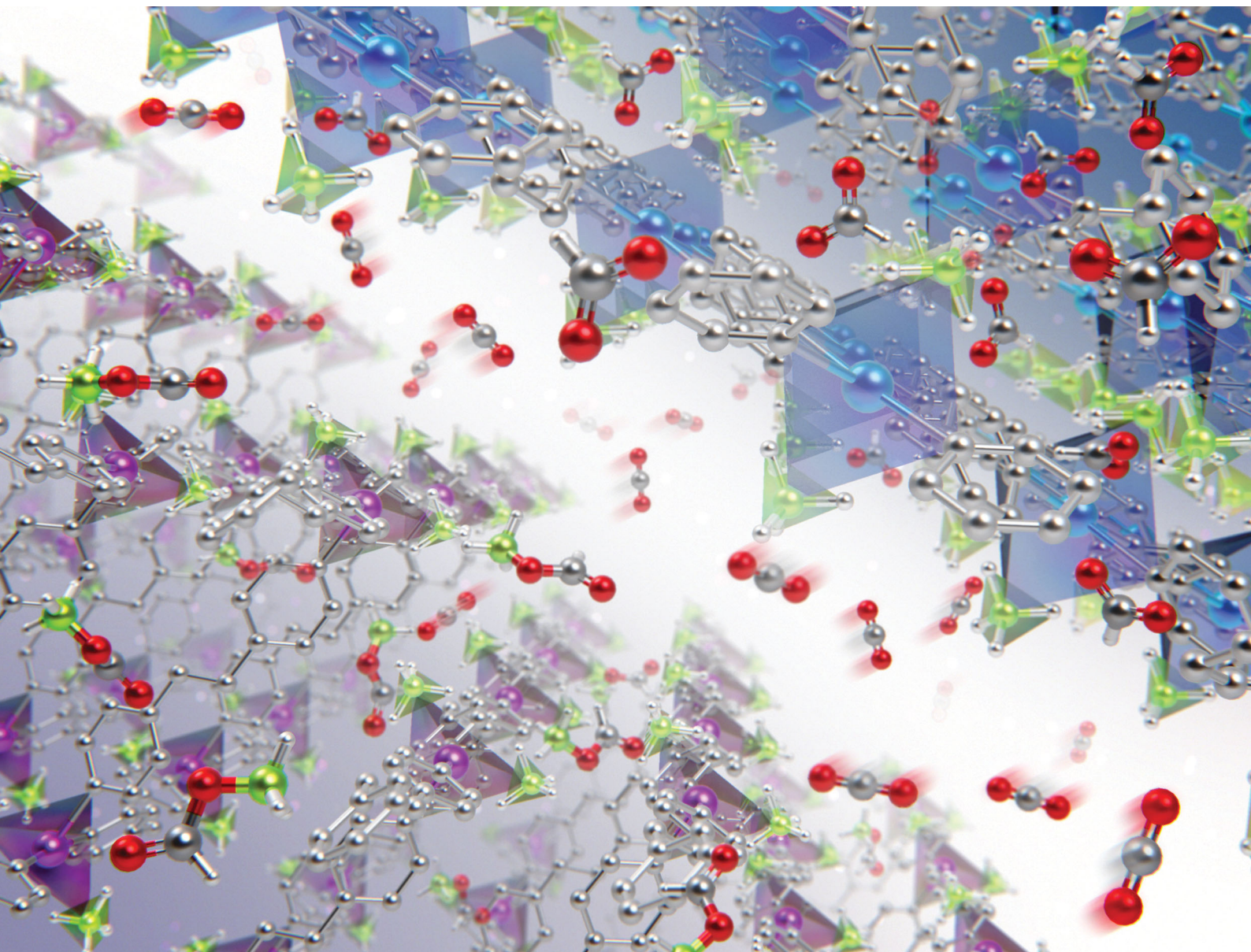


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Reactivity of borohydride incorporated in coordination polymers toward carbon dioxide†

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Borohydride (BH₄⁻)-containing coordination polymers converted CO₂ into HCO₂⁻ or [BH₃(OCHO)]⁻, whose reaction routes were affected by the electronegativity of metal ions and the coordination mode of BH₄⁻. The reactions were investigated using thermal gravimetric analysis under CO₂ gas flow, infrared spectroscopy, and NMR experiments.

Conversion of carbon dioxide (CO₂) into valuable chemicals is a key to realize a sustainable society.^{1,2} In particular, it is essential to establish chemical reactions that transform CO₂ into various types of chemical moieties under mild conditions.³ However, the inherent inertness of CO₂ has hampered the utilization of CO₂ in transformation reactions. To overcome the inertness, various catalytic and stoichiometric reactions have been widely studied in both solution and the solid-state, including metals, metal oxides,⁴ metal complexes,^{5,6} and metal-free organic molecules.⁷

Borohydride (BH₄⁻), a hydride-based complex anion, has been commonly utilized as a reducing agent. In the solution phase, metal borohydrides (MBHs) stoichiometrically react with CO₂ at ambient temperatures and pressures.^{8–11} BH₄⁻ in solution is able to convert CO₂ into chemical species such as formate (HCO₂⁻) and formylhydroborate ([BH_{4-x}(OCHO)_x]⁻, *x* = 1, 2 and 3) depending on the reaction conditions, *e.g.* counter cations, temperatures,

solvents, and pressures.^{8,9,12} The solid-state reactivity of BH₄⁻ toward CO₂ is also interesting from the viewpoint of heterogeneous catalysts and CO₂ scrubbers. Nevertheless, limited studies have been performed on the solid-state reactivity of MBHs toward CO₂.^{13,14} This is because slow diffusion of CO₂ in dense MBHs results in low reactivity under mild conditions.¹⁴ Although porous structures are advantageous for the diffusion of CO₂, MBHs with the porous structure are limited except for a few examples, *e.g.* γ-Mg(BH₄)₂.^{14,15}

Coordination polymers (CPs) and metal–organic frameworks (MOFs) are crystalline solids constructed from metal ions and bridging organic linkers.^{16–18} Their open structures have offered an attractive platform for various gas–solid reactions, such as CO₂ sorption^{19–21} and post-synthetic modification.^{22,23} In addition, rich structural and chemical tunability of CPs demonstrated the controlled reactivity of reactive species, *e.g.* radicals,^{24,25} imines,²⁶ and photoactive metal complexes.²⁷ CPs are a promising platform for solid–gas reactions between BH₄⁻ and CO₂. BH₄⁻-containing CPs are constructed from metal ions (*e.g.* Mg²⁺, Ca²⁺, Mn²⁺, Zn²⁺, and Th⁴⁺) and N-based neutral linkers show various types of the chemical environment of BH₄⁻.^{28–30} Here, we investigate the reactivity of BH₄⁻-containing CPs to convert CO₂ into HCO₂⁻ or [BH₃(OCHO)]⁻ under mild conditions depending on their structures.

[M(BH₄)₂(pyz)₂] (**M-pyz**, M = Mg²⁺, Ca²⁺, pyz = pyrazine)^{28,29} were selected to investigate the influence of metal ions on the reactivity of BH₄⁻ toward CO₂. The metal ion center shows an octahedral geometry and the two BH₄⁻ ions coordinate in the axial positions (Fig. 1A). The extended structure of **M-pyz** comprises a 2D square grid constructed by [M₄pyz₄] units (Fig. 1B). The electronic properties and reactivity of BH₄⁻ are influenced by the electronegativity of counter metal ions.^{31,32} Attempts at the synthesis of isostructural **M-pyz** were made using a Mn²⁺-based MBH precursor. [Mn(BH₄)₂·3THF]·NaBH₄ was prepared following the literature methods.³³ The general synthetic method involves mechanochemical milling of the MBH precursor and pyz under Ar. **Mg-pyz** was previously

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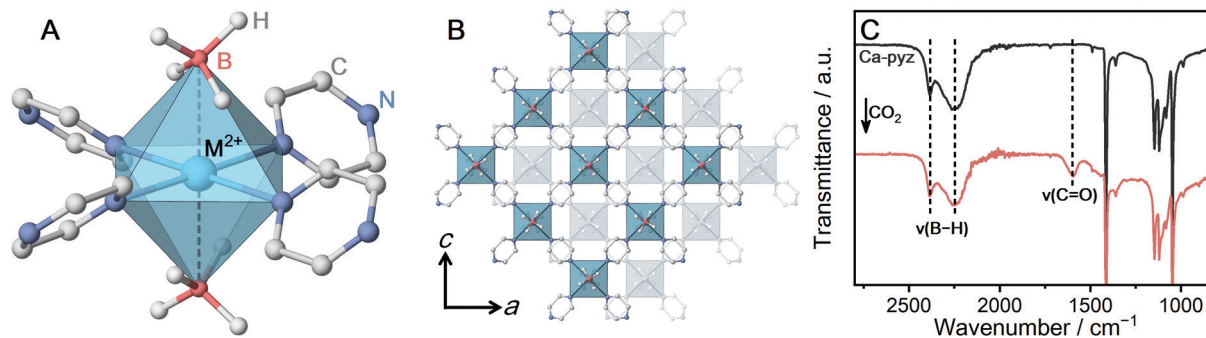


Fig. 1 (A) Local coordination geometry of **M-pyz** ($M = \text{Mg}^{2+}$, Ca^{2+} , Mn^{2+}). (B) ABAB stacking structure of the extended 2D layers of **M-pyz** ($M = \text{Mg}^{2+}$, Ca^{2+} , Mn^{2+}). (C) IR spectra of **Ca-pyz** before and after CO_2 adsorption at 25 °C.

synthesized in the solution phase, whereas the solvent-free conditions afford a highly crystalline product as well (Fig. S1, ESI[†]). The powder X-ray diffraction (PXRD) pattern of **Mn-pyz** shows a good agreement with that of **Mg-pyz** (Fig. S1, ESI[†]).

The solid-state synthesis of **M-pyz** proceeds without solvents at 25 °C within 30 min. The fast reaction kinetics in the solid-state is ascribed to the low melting point of pyz (52 °C). The lower melting point of reactants leads to higher molecular mobility, enhancing the reactivity in the solid-state.³⁴ Mechanical milling is useful to synthesize CPs from MBHs because most of the MBHs are poorly soluble in common organic solvents. The thermal properties were characterized by thermal gravimetric analysis (TGA) under N₂ (Fig. S2, ESI[†]). Each compound exhibits a weight loss at relatively low temperatures; 50, 70, and 70 °C for **Mg-**, **Mn-**, and **Ca-pyz** due to the low boiling point of pyz (115 °C). Isothermal TGA measurements at 40 °C under N₂ indicate that **Ca-pyz** shows higher thermal stability than **Mn-pyz** (weight loss after 6 hours; 0.2 vs. 3.2 wt%, Fig. S3, ESI[†]). In the case of MBHs, electropositive metal ions construct MBHs with higher thermal stability.³⁵ Meanwhile, in the case of BH₄⁻-containing CPs, the strength of the coordination bonds is also essential. The Hard and Soft Acids and Bases (HSAB) theory reveals that electropositive metal ions (hard acids) form weaker coordination bonds with nitrogen-based linkers (soft bases) such as pyz. Therefore, the trend of thermal stability for **M-pyz** does not simply follow the electronegativity of metal ions (thermal stability: Mg < Mn < Ca, Pauling electronegativity: Ca < Mg < Mn).

To characterize the chemical environment of BH₄⁻ in the CP, solid-state ¹¹B magic angle spinning (MAS) nuclear magnetic resonance (NMR) was carried out on non-paramagnetic **Ca-pyz**. The ¹¹B NMR spectrum of **Ca-pyz** displays a peak at -36 ppm corresponding to the signal of BH₄⁻ (Fig. S4, ESI[†]). The total charge on BH₄⁻ is correlated with the chemical shift of ¹¹B NMR: electron-rich BH₄⁻ shows a peak in a lower frequency.³¹ The low-frequency shift of the ¹¹B peak indicates that BH₄⁻ in **Ca-pyz** is more electron-rich than Ca(BH₄)₂. In the framework of **Ca-pyz**, the Lewis acidity of Ca²⁺ was reduced by electron donation from the coordinating pyz molecules, which leads to the formation of electron-rich BH₄⁻.²⁹

CO₂ adsorption measurement was carried out to evaluate the reactivity of **Ca-pyz** in gas–solid equilibrium. The CO₂

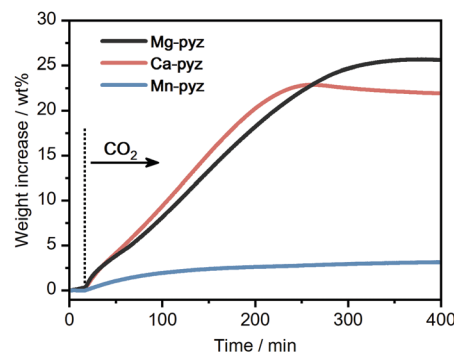


Fig. 2 Isothermal TGA profiles of **M-pyz** ($M = \text{Mg}^{2+}$, Ca^{2+} , Mn^{2+}) under CO_2 flow (0.1 MPa, 30 mL min⁻¹) at 40 °C.

isotherm at 25 °C displays irreversible adsorption (7 mL g⁻¹ at 100 kPa), which is characteristic of chemisorption behavior (Fig. S5, ESI[†]).³⁶ The IR spectrum of **Ca-pyz** after CO₂ adsorption displays a new peak at 1600 cm⁻¹, corresponding to C=O stretching (Fig. 1C). The solid-state ¹H-¹³C cross-polarization (CP) MAS NMR spectrum of **Ca-pyz** after CO₂ adsorption shows peaks at 170 and 145 ppm. The peaks correspond to the signals of HCO₂⁻ and pyz, respectively (Fig. S6, ESI[†]). The results indicate that BH₄⁻ in **Ca-pyz** reduces CO₂ to HCO₂⁻ with the release of diborane (B₂H₆) as a by-product.¹⁰

The kinetic reactivity of **M-pyz** toward CO₂ was evaluated using isothermal TGA under CO₂ flow. Fig. 2 displays the TGA profiles of each powder sample (10 mg) under CO₂ flow (0.1 MPa, 30 mL min⁻¹) at 40 °C. **Mg-pyz** and **Ca-pyz** exhibit higher weight increases than **Mn-pyz** (25.5, 21.9, and 3.2 wt% after 400 min, respectively). **Ca-pyz** was amorphous after the CO₂ reaction, as confirmed by PXRD (Fig. S7, ESI[†]). To identify the chemical species after the CO₂ reaction, solution NMR was carried out on **Ca-pyz** dissolved in DMSO-*d*₆. The solution ¹³C NMR spectrum of **Ca-pyz** after the CO₂ reaction displays peaks at 167, 146, 53, 50, 47 and 44 ppm (Fig. S8, ESI[†]). The peaks at 167 and 146 ppm correspond to the ¹³C signals of HCO₂⁻ and pyz, respectively. The peak at 47 ppm is assigned to piperazine formed by the reduction of pyz by B₂H₆, whereas the rest of the peaks are not able to be assigned.^{29,37} The higher reactivity of **Ca-pyz** toward CO₂ is attributed to the preferable electronic

interaction between Ca^{2+} (hard acid) and HCO_2^- (hard base) rather than BH_4^- (soft base).

The formation of $[\text{BH}_{4-x}(\text{OCHO})_x]^-$ from BH_4^- and CO_2 was investigated at a BH_4^- -containing CP. Given that $[\text{BH}_{4-x}(\text{OCHO})_x]^-$ is bulky than HCO_2^- , $[\text{Mn}(\text{BH}_4)_2(\text{dpe})_{1.5}]$ (**Mn-dpe**, dpe = dipyr-dithane) having voids was selected.²⁸ The two BH_4^- ions coordinate to the Mn^{2+} center in a bidentate manner, which was confirmed by single-crystal X-ray diffraction (SC-XRD) in Fig. 3A. The extended structure of **Mn-dpe** comprises a 1D ladder constructed from $[\text{Mn}_4\text{dpe}_4]$ units (Fig. 3B). The coordination mode of BH_4^- was confirmed by IR spectroscopy as well. The IR spectrum of **Mn-dpe** displays two stretching peaks in the B–H stretching region at 2378 and 2127 cm^{-1} , respectively (Fig. 4B). The peak at 2378 cm^{-1} corresponds to the B–H bond coordinating to the Mn^{2+} center, whereas the peak at 2127 cm^{-1} corresponds to the non-coordinating B–H.³⁸ In contrast to the broadened B–H stretching peak of **Ca-pyz** (Fig. 1C), **Mn-dpe** displays distinct two peaks of B–H stretching, which is originated from a stronger binding interaction between Mn^{2+} (soft acid) and BH_4^- (soft base).

The kinetic curve of the CO_2 reaction with **Mn-dpe** was collected using the same procedure as **M-pyz** (Fig. 3B). **Mn-dpe** demonstrates a weight increase of 26.2 wt% after 600 min at 40 °C, which corresponds to a value of the 1.1:1 molar ratio of reacted CO_2 per BH_4^- . After the CO_2 reaction, **Mn-dpe** shows small diffraction peaks different from the original peaks (Fig. S9, ESI[†]). Solution ^{11}B NMR measurement was carried out to determine the chemical species after the CO_2 reaction. The $^{11}\text{B}\{^1\text{H}\}$ NMR spectrum of digested **Mn-dpe** after the CO_2 reaction displays the peaks at –33, –11, and 2.2 ppm in Fig. 4A. The broad peaks were observed due to the paramagnetic effect of Mn^{2+} . The ^{11}B peaks correspond to BH_4^- , $[\text{BH}_3(\text{OCHO})]^-$ and $[\text{BH}_2(\text{OCHO})_2]^-$, respectively.^{9,39} Successive CO_2 insertions into the B–H bond of BH_4^- produce $[\text{BH}_{4-x}(\text{OCHO})_x]^-$, and the number of reacted CO_2 molecules is affected by the reaction conditions such as pressure and temperature in the solution phase.^{8,9} The reaction of NaBH_4 in acetonitrile with 0.1 MPa of CO_2 for 10 minutes produces $[\text{BH}(\text{OCHO})_3]^-$ as a major product, and $[\text{BH}_3(\text{OCHO})]^-$ is not observed.⁹ This is because all the hydrogen atoms of BH_4^- dissociated in acetonitrile are available for the reaction with CO_2 . On the other hand, in the case of **Mn-dpe**, two of the hydrogen atoms of BH_4^- are pinned with the Mn^{2+} center by

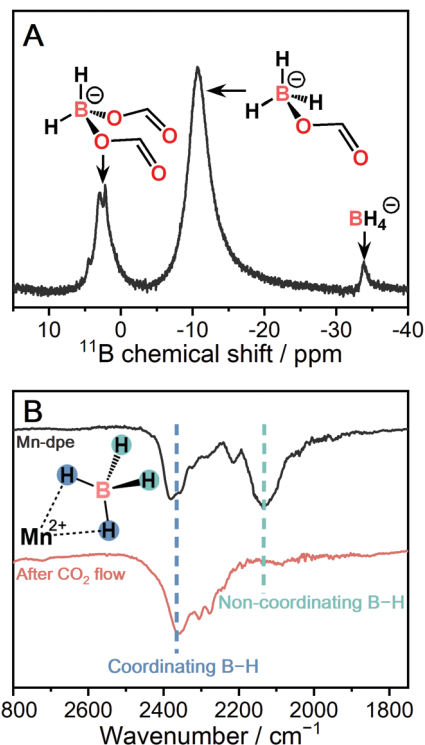


Fig. 4 (A) Solution $^{11}\text{B}\{^1\text{H}\}$ NMR of digested **Mn-dpe** after CO_2 reaction. (B) IR spectra of **Mn-dpe** before and after the CO_2 reaction.

a coordination bond as confirmed by SC-XRD and IR spectroscopy. After the CO_2 reaction, a non-coordinating B–H stretching peak was not observed, and this is because of the reaction with CO_2 to form $[\text{BH}_3(\text{OCHO})]^-$ and $[\text{BH}_2(\text{OCHO})_2]^-$ in Fig. 4B. The coordinating B–H stretching peak is preserved after the CO_2 reaction, indicating the coordinating bonds between Mn^{2+} and $[\text{BH}_3(\text{OCHO})]^-$ or $[\text{BH}_2(\text{OCHO})_2]^-$. A sluggish kinetics of dense NaBH_4 in the solid-state toward CO_2 indicates that the open structure of **Mn-dpe** is essential for the diffusion of CO_2 (Fig. 3C). Based on the results, the reaction between **Mn-dpe** and CO_2 to produce $[\text{BH}_3(\text{OCHO})]^-$ and $[\text{BH}_2(\text{OCHO})_2]^-$ is proposed (Fig. S11, ESI[†]). The results indicate that the anisotropic coordination geometry of BH_4^- in **Mn-dpe** affects the reaction route with CO_2 .

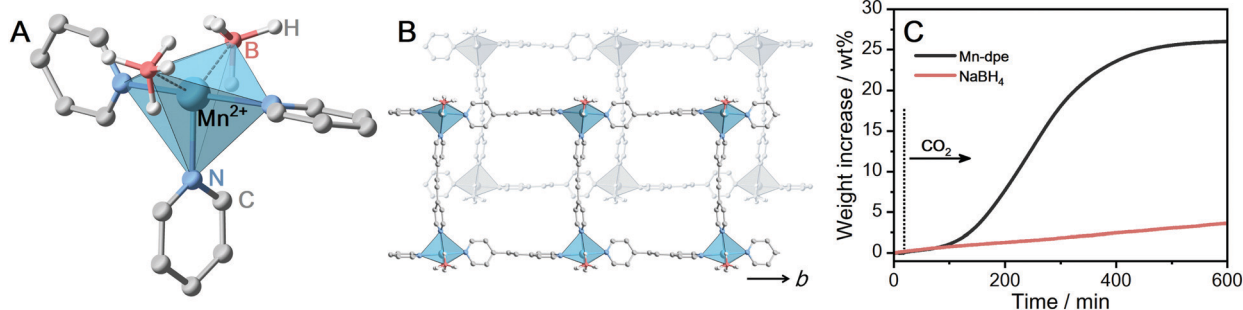


Fig. 3 (A) Local coordination geometry of **Mn-dpe**. (B) Packing structure of the extended 1D ladders of **Mn-dpe**. (C) Isothermal TGA profiles of **Mn-dpe** and NaBH_4 under CO_2 flow (0.1 MPa, 30 mL min^{-1}) at 40 °C.

In conclusion, we demonstrated the reactivity of BH_4^- toward CO_2 which is correlated with the crystal structures of BH_4^- -containing coordination polymers. The reactivity of $[\text{M}(\text{BH}_4)_2(\text{pyrazine})_2]$ ($\text{M} = \text{Mg}^{2+}, \text{Mn}^{2+}, \text{Ca}^{2+}$) and $[\text{Mn}(\text{BH}_4)_2(\text{dipyridylethane})_{1.5}]$ toward CO_2 at 40°C was investigated by using isothermal TGA under CO_2 flow, IR and NMR. BH_4^- in $[\text{Ca}(\text{BH}_4)_2(\text{pyrazine})_2]$ converted CO_2 into HCO_2^- . The BH_4^- pinned by coordination bonds with Mn^{2+} in $[\text{Mn}(\text{BH}_4)_2(\text{dipyridylethane})_{1.5}]$ regulated the successive CO_2 insertion reaction and produced $[\text{BH}_3(\text{OCHO})]^-$ as a major species. The structural diversity of coordination polymers provides a new approach to regulate the reaction routes between BH_4^- and CO_2 in the solid-state.

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Conflicts of interest

The authors declare no conflict of interest.

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