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Conversion of CO₂ into porous metal–organic framework monoliths†

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We demonstrate the one-pot conversion of CO₂ into amorphous formate-based metal–organic frameworks (MOFs) that form grain-boundary-free monoliths with permanent porosity through hot-pressing. The local coordination geometries of metal ions are characterized using solid-state NMR and synchrotron total X-ray scattering analyses. Hot-pressing decreases the pore sizes of monoliths, enhancing the adsorption selectivity toward H₂. The key for the formation of microporous monoliths is the coordination network in which formate, capable of adopting various coordination modes, is connected via stable metal–oxygen bonds.

1. Introduction

The conversion of CO₂ into functional materials is essential for realizing a carbon-neutral society.^{1–3} The synthesis of organic polymers and carbon materials from CO₂ as a feedstock has been extensively studied in the past few decades.^{4,5} In recent years, the synthesis of metal–organic frameworks (MOFs) from CO₂ has gained attention.⁶ CO₂ is converted into bridging linkers, such as formate (OCHO[−]), formylhydroborate,⁷ carbamate,⁸ and carboxylate.⁹ On the other hand, the inherent inertness of CO₂ has restricted the structural diversity of CO₂-derived linkers and functionality of the resultant MOFs. An approach to functionalize CO₂-derived MOFs that does not rely on the structural diversity of CO₂-derived linkers is highly demanded.

Tailoring structural disorder offers a powerful route for functionalizing solid materials. Amorphous materials, such as glasses and gels, lack long-range structural order and exhibit various features, *e.g.*, defects, isotropy, transparency, and high mechanical strength.¹⁰ Amorphous MOFs exhibit unique functionalities that are not achieved by the crystalline analogues. For example, grain-boundary-free monoliths with permanent porosity are formed *via* thermal and mechanical treatment, *e.g.*, melt quenching and hot-pressing.^{11,12} The formability and processability of porous monoliths is attractive for enhanced volumetric gas storage capacity and recyclable heterogeneous catalysts.^{13,14} Meanwhile, few studies focus on amorphous CO₂-derived MOFs and none of them show the formability of porous monoliths.^{7,15}

In this work, we attempted to synthesize amorphous CO₂-derived MOFs showing permanent porosity as the monolithic form. OCHO[−] was selected as a CO₂-derived bridging linker. Borohydride (BH₄[−]) readily converts CO₂ into OCHO[−] by

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CO₂ capture and utilization.

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hydride transfer.⁷ The small steric hindrance of OCHO^- allows for various coordination geometries, making it suitable for constructing an isotropic grain-boundary-free structure.¹⁶ Oxo-philic Al^{3+} and Ga^{3+} ions were employed to form a strong coordination bond with OCHO^- that is essential to preserve a stable porous structure through hot-pressing treatment.¹⁷

2. Experimental section

2.1. Synthesis of OCHO^- -based MOFs from CO_2

All chemicals were obtained from commercial suppliers and used without further purification. The powder samples of OCHO^- -based MOFs, denoted as M-CO_2 ($\text{M} = \text{Al}^{3+}$ and Ga^{3+}), were synthesized from CO_2 . Sodium borohydride (4.0 mmol) in anhydrous acetonitrile (MeCN, 20 mL) was reacted with CO_2 (99.99%) at 2.0 MPa at 25 °C for 1.5 hours in a high-pressure reaction vessel. The resulting suspension was mixed with metal nitrate salts (1.0 mmol) in anhydrous methanol (MeOH, 20 mL) at 100 °C for 48 hours. The white precipitate was isolated by centrifugation, washed with MeOH, and dried under vacuum (45 and 26% yields for Al-CO_2 and Ga-CO_2).

2.2. General characterization

Thermogravimetric analysis (TGA) was carried out using a Rigaku Thermo plus TG 8122 under N_2 flow or air with a heating rate of 10 °C min^{-1} . Differential scanning calorimetry (DSC) was carried out using a Hitachi High-Tech DSC 7200 instrument under N_2 flow with a heating/cooling rate of 10 °C min^{-1} . Scanning electron microscopy (SEM) and energy-dispersive X-ray spectroscopy (EDX) was carried out using a JEOL JSM-7610F operated with an acceleration voltage of 200 kV. Transmission Electron Microscopy and Energy-Dispersive X-ray Spectroscopy (TEM-EDS) was carried out using a JEOL JEM-ARM 200F. Fourier transform infrared (FT-IR) spectra were collected using a Bruker ALPHA II FT-IR spectrometer with a Universal ATR accessory under a N_2 atmosphere. Inductively coupled plasma-optical emission spectrometry (ICP-OES) was carried out using an Agilent 700 series. The supernatant (200 μL) was dried and digested in a solution of 2% nitric acid with a total volume of 5 mL before a measurement. The average static water contact angle was measured at 25 °C, using a contact angle goniometer by dropping 10 μL of deionized water on three different locations on a monolith. The contact angle of the droplet was analyzed using the Ossila contact angle software.

2.3. X-ray analysis

Powder X-ray diffraction (PXRD) patterns were collected on a Rigaku SmartLab SE X-ray type with $\text{CuK}\alpha$ radiation ($\lambda = 1.54059$ Å). X-ray photoelectron spectroscopy (XPS) spectra were collected using a JEOL XPS instrument (JPS-9010 MC) with a Mg $\text{K}\alpha$ and Al $\text{K}\alpha$ source (1253.6 eV and 1486.6 eV) under high vacuum (10^{-7} Pa). All binding energy values were referenced to the C 1s peak (284.70 eV). Synchrotron variable-temperature (VT) PXRD patterns were collected using synchrotron radiation ($\lambda = 0.99927$ Å) employing a large Debye-Scherrer camera with semiconductor detectors on the BL02B2 beamline at the Super Photon

Ring (SPring-8, Hyogo, Japan). Pair distribution function (PDF) analysis was performed using synchrotron scattering data. Total X-ray scattering was collected at 30 °C with two 2D CdTe detectors at the BL04B2 beamline in SPring-8. The incident energy was 112.9232 keV. $G(r)$ was obtained from the Fourier transform of $S(Q)$ with a Lorch modification function by using IgorPro software. X-ray absorption spectroscopy (XAS) including X-ray absorption near edge structure (XANES) and extended X-ray absorption fine structure (EXAFS) regions was performed in the transmission mode at the BL14B2 beamline at SPring-8.

2.4. NMR analysis

^1H , ^{11}B , and ^{13}C solution NMR spectra were collected using a Bruker Avance III HD 600 MHz. The powder samples were digested in 1.0 M $\text{DCl}/\text{D}_2\text{O}/\text{DMSO}-d_6$ for NMR measurement. CHN elemental analysis was performed using a LECO TruSpec Micro Element Series with cystine standard. Solid-state ^1H , ^{13}C , and ^{27}Al magic-angle spinning (MAS) NMR experiments were conducted on a JEOL JNM-ECZ600R spectrometer at a 14.1 T superconductor magnet at room temperature. ^{13}C cross-polarization (CP)/MAS, ^1H - ^{13}C 2D CP-heteronuclear correlation (HETCOR), ^{27}Al single-pulse, Hahn-echo, and 3QMAS measurements were performed using a JEOL 3.2 mm double resonance MAS probe at a MAS rate of 20 kHz. In the CP/MAS sequence, a ^1H 90° pulse width of 2.3 μs and ^1H and ^{13}C spin-lock pulse strengths of 70 and 50 kHz with a contact time of 3 ms were used. A ramped-amplitude spin-lock pulse was used for ^{13}C . ^{13}C signals were acquired under ^1H TPPM heteronuclear dipolar decoupling pulse irradiation with a pulse strength of 100 kHz. ^{27}Al single-pulse and Hahn-echo MAS spectra were measured with 90° and 180° pulse lengths of 1.15 and 2.3 μs . The ^{27}Al 3QMAS spectrum was obtained using the z-filter 3QMAS sequence. 0 quantum (Q)-3Q excitation, 3Q-0Q reconversion, and weak 90° pulse lengths of 3.3, 1.1, and 8.0 μs were used. ^{27}Al quadrupolar line-shape analysis was performed using ssNake software.¹⁸

2.5. Catalytic activity of M-CO_2 for CO_2 cycloaddition

Epichlorohydrin (ECH, 12.5 mmol), M-CO_2 (0.3–1.0 mol%), and tetrabutylammonium bromide (TBAB, 2.0 mol%) were added to the reaction vial (20 mL). CO_2 (99.99%) was introduced into the reaction mixture under stirring at 0.1 MPa at 30 °C for 48 hours. The reaction using styrene oxide was carried out at 80 °C. The conversion of ECH into chloropropylene carbonate (CPC) was calculated based on ^1H NMR analysis. The recovered catalyst was collected by centrifugation, followed by washing with fresh MeOH and dried in a vacuum.

2.6. Preparation of M-CO_2 monolith by hot-pressing

The powder sample of M-CO_2 (20 mg) was introduced into a 7 mm diameter stainless steel die set. A monolith was prepared under air by hot-pressing at 145 °C and 50–55 kN for 2 hours. After pressure release, the die set was taken out to cool down to room temperature.



2.7. Gas adsorption analysis

Gas adsorption isotherms were collected using a MicrotracBEL BELSORP-mini X for N₂ at 77 K and CO₂ at 195, 273, and 298 K. Brunauer–Emmett–Teller surface areas (S_{BET}) were calculated from the N₂ adsorption isotherms. H₂ adsorption isotherms at 77 K were recorded on a MicrotracBEL BELSORP-mini II. The pore size distribution (PSD) was calculated using the non-local density functional theory (NLDFT) method based on the N₂ adsorption isotherms. NH₃ temperature-programmed desorption (TPD) was carried out using a MicrotracBEL BELCAT. NH₃ was captured with 50 mg of powder sample at 40 °C for 30 minutes and the desorption profile was recorded using a TCD detector.

3. Results and discussion

3.1. Synthesis and structural characterization of M-CO₂

The synthesis is a one-pot, two-step procedure involving the insertion of CO₂ into BH₄[−] to afford formylhydroborate ([BH(OCHO)₃][−]) and subsequent solvothermal reaction that converts [BH(OCHO)₃][−] into OCHO[−] (Fig. 1).⁷ This method is classified as a type-II (one-pot) synthesis of CO₂-derived MOFs according to the literature.⁶ The choice of metal ions and solvents was essential for obtaining amorphous porous structures. A synthetic attempt using Ce³⁺ ion instead of Al³⁺/Ga³⁺ resulted in a non-porous crystalline product (Fig. S1 and S2†). The larger ionic radius of Ce³⁺ led to a higher coordination number, forming a dense structure. Methoxide (MeO[−]), derived from MeOH, serves as a linker in both M-CO₂ (details below). Control experiments using ethanol (EtOH) instead of MeOH as a synthetic solvent provided amorphous non-porous products

consisting of OCHO[−] and Al³⁺ without EtO[−] (Fig. S3–S5†). This indicates that the smaller steric hindrance of MeO[−] is suitable for constructing the porous structure of M-CO₂.

Acid-digested solution NMR was carried out to confirm the formation of OCHO[−]. The ¹H NMR spectra of M-CO₂ show peaks at 3.1 and 8.1 ppm corresponding to MeO[−] and OCHO[−], respectively. The ratios of OCHO[−] and MeO[−] were calculated as 1 : 1 and 1 : 1.8 for Al-CO₂ and Ga-CO₂ (Fig. S6†). The assignment was also confirmed by ¹³C NMR (Fig. S7†). The ¹¹B NMR spectra showed that M-CO₂ contained no boron species derived from NaBH₄ (Fig. S8†). The FT-IR spectra exhibit the C=O stretching vibration of OCHO[−] at 1581 and 1652 cm^{−1} for Al-CO₂ and Ga-CO₂, respectively (Fig. S9†).¹⁹ The broad peak at 3420 cm^{−1} in Ga-CO₂ was attributed to the O–H stretching vibration of OH[−].²⁰

The metal content of M-CO₂ was determined by pyrolysis under air. TGA heating up to 900 °C under a flow of air fully converted M-CO₂ into corresponding metal oxides (Fig. S10 and S11†). The gravimetric metal contents were calculated as 19.3 and 59.1 wt% for Al-CO₂ and Ga-CO₂, respectively. Given the results of CHN elemental analysis, the chemical compositions were determined as [Al₂(OCHO)₃(OMe)₃] for Al-CO₂ and [Ga₂O_{1.8}(OHCO)_{0.5}(OMe)_{0.9}(OH)] for Ga-CO₂. The gravimetric CO₂ content was calculated as 46.8 and 9.3 wt% for Al-CO₂ and Ga-CO₂ (Table S1†).

The PXRD patterns exhibit broad features at 8.1, 10.8, and 12.3° for Al-CO₂ and at 5.1° for Ga-CO₂, which do not match the reported patterns of metal formate compounds (Fig. 2A and S12†).^{19,21,22} SEM-EDX exhibits spherical particles with a diameter of 5 and 2 μm for Al-CO₂ and Ga-CO₂ respectively and a homogeneous distribution of each metal element (Fig. 1 and



Fig. 1 Schematic illustration of the synthesis and proposed structures of M-CO₂ (M = Al³⁺, Ga³⁺). Carbon and oxygen atoms derived from CO₂ are highlighted. The symbol X in the structure of Ga-CO₂ indicates either O^{2−}, OH[−], or OMe[−]. SEM images of M-CO₂ are displayed, respectively. The scale bar indicates 1 μm.





Fig. 2 Structural characterization of **M-CO₂**. (A) PXRD patterns of **Al-CO₂** and **Ga-CO₂**. (B) ¹H-¹³C CP-HETCOR NMR spectra of **Al-CO₂**. The correlation between ¹H of OCHO[−] and ¹³C of MeO[−] is highlighted as a green box. (C) ²⁷Al 1D sliced 3QMAS NMR spectrum (black) and fitting (red) of **Al-CO₂**. (D) PDF profiles of **Al-CO₂** and **Ga-CO₂**. (E) Histograms of M-M distances for [M-(R-COO)_n(X)_{3-n}-M] with Gaussian distribution. (F) Proposed M-M building units of **M-CO₂**. X = O^{2−}, OH[−], or OMe[−].

S13, S14†). TEM-EDS also confirmed the homogeneous distribution of each element in **M-CO₂** (Fig. S15 and S16†). Solid-state ¹H-¹³C CP-HETCOR NMR experiment was carried out to examine the intermolecular distance correlation between OCHO[−] and MeO[−]. The ¹H-¹³C HETCOR NMR spectrum of **Al-CO₂** exhibits a clear correlation peak between ¹H of OCHO[−] at 8.5 ppm and ¹³C of MeO[−] at 51.5 ppm (Fig. 2B). **Ga-CO₂** also exhibits the correlation between OCHO[−] and MeO[−] (Fig. S17†). The observed correlations indicate that both OCHO[−] and MeO[−] are homogeneously distributed to form the structure of **M-CO₂**.

The local coordination geometry of Al³⁺ was characterized by ²⁷Al MAS NMR. The ²⁷Al NMR spectrum of **Al-CO₂** exhibits an asymmetric line shape due to the ²⁷Al quadrupolar interaction (Fig. S18†). To analyze the coordination structure, we performed quadrupolar line-shape fitting to the ²⁷Al 1D spectrum sliced at the peak center of the isotropic dimension (12.31 ppm) of the 3QMAS spectrum (Fig. 2C and S19†). The isotropic chemical shift $\delta_{\text{iso}} = 8.21$ ppm, quadrupolar coupling constant $C_Q = 5.00$ MHz, and asymmetry parameter $\eta = 0.48$ were obtained. The δ_{iso} value suggests that Al³⁺ forms an octahedral six-coordinated geometry.^{23,24} The ²⁷Al MAS spectrum of [Al(OCHO)₃], possessing the highly symmetric Al³⁺-6O coordination structure, shows a narrow peak with $C_Q \sim 0$ (Fig. S20†). The larger C_Q value of **Al-CO₂** reflects an asymmetric coordination structure of Al³⁺ bound with both OCHO[−] and MeO[−]. The non-zero η suggests a low axial-symmetry structure around Al³⁺ due to the random distribution of the two linkers. The coordination geometry of Ga³⁺ in **Ga-CO₂** was characterized by synchrotron XAS. The EXAFS fitting on the first coordination shell confirmed that the coordination number of Ga³⁺ in **Ga-CO₂** was 5.6 ± 0.6 , which is indicative of octahedral geometry (Fig. S21 and Table S2†). The formation of octahedral geometry in **M-CO₂** was also confirmed by XPS (Fig. S22 and S23†). The binding energies of O (1s), C

(1s), and Al (2p_{3/2}) were found to be 530.1, 288.0 and 71.6 eV in **Al-CO₂** which correspond to Al-O coordination and the presence of OHCO[−].²⁵ **Ga-CO₂** shows binding energies of O (1s) and Ga (2p_{3/2}) at 531.5 eV and 1118.9 eV which belong to Ga-O bond and Ga³⁺, respectively.

PDF analysis was performed to characterize the structural periodicity of **M-CO₂** (Fig. 2D). The peaks below 2 Å were assigned as the metal-oxygen bonds (1.85 and 1.95 Å for **Al-CO₂** and **Ga-CO₂**).^{26,27} The peaks between 2.0 and 4.0 Å are assigned as metal-metal (M-M) correlations (2.87 Å for **Al-CO₂** and 3.05 and 3.38 Å for **Ga-CO₂**). To figure out the bridging mode of M-M, we surveyed M-M distances of an Al³⁺/Ga³⁺-based coordination compound with octahedral geometry and R-COO[−] groups in the CCDC database. The histograms of M-M distances were plotted for the building units of [M-(R-COO)_n(X)_{3-n}-M] ($M = \text{Al}^{3+}, \text{Ga}^{3+}$; $X = \text{O}^{2-}/\text{OH}^{-}/\text{OMe}^{-}$; $n = 1, 2$, Fig. 2E). We assign the peak at 2.87 Å in **Al-CO₂** as [Al-(OCHO)(OMe)₂-Al] and 3.38 Å in **Ga-CO₂** as [Ga-(OCHO)₂(X)-Ga] (Fig. 2E). The peak at 3.05 Å in **Ga-CO₂** is assigned as [Ga-(X)₂-Ga], edge-sharing octahedral geometry (Fig. 2F and Tables S3, S4†). The mixed coordination of OCHO[−] and OMe[−] in **Al-CO₂** is consistent with the low axial-symmetry structure of Al³⁺ observed by ²⁷Al MAS NMR. The longer-range periodicity was observed up to around 8 and 12 Å for **Al-CO₂** and **Ga-CO₂**. The extended network is formed by connecting the M-M building units as proposed in Fig. 1.

3.2. Thermal and chemical stability

The TGA profiles under N₂ indicate that there was no significant weight loss up to 190 and 180 °C for **Al-CO₂** and **Ga-CO₂** (Fig. S24†). The thermal stability is slightly lower than those of [M(OCHO)₃] (decomposition temperatures: 200 and 230 °C for Al³⁺ and Ga³⁺).²¹ The thermal stability of **Al-CO₂** was studied by



synchrotron variable-temperature PXRD (Fig. S25†). The crystallinity of **Al**-CO₂ was maintained up to 200 °C and the intensity reduced when heated to 320 °C. The stability of common organic solvents was also studied. The PXRD pattern of **Al**-CO₂ was intact after soaking in common organic solvents, *e.g.*, tetrahydrofuran, chloroform, and toluene, for 24 h at 25 °C, respectively (Fig. S26†).

3.3. Characterization of the defective open metal site (OMS) for CO₂ cycloaddition

We evaluated the Lewis acidity of defective OMS in **M**-CO₂ by NH₃-TPD (Fig. 3). NH₃ was completely released below 190 °C within 2 hours, indicating that the defective OMS in **M**-CO₂ serves as a weak acid site.²⁸ The total acidic sites were calculated as 0.59 and 2.25 mmol g⁻¹ for **Al**-CO₂ and **Ga**-CO₂, respectively. The larger amount of acidic sites in **Ga**-CO₂ was attributed to a defective OMS in the amorphous structure.¹⁵ Defective OMS serves as a catalytically active site for CO₂ cycloaddition with epoxides. The catalytic activity of **M**-CO₂ was evaluated with ECH as a model reaction. The solvent-free reaction of ECH and CO₂ in the presence of 1.0 mol% of **M**-CO₂ and TBAB as a co-catalyst at 30 °C yielded CPC with a high conversion of 94.4 and 91.6% for **Al**-CO₂ and **Ga**-CO₂ (Fig. 3 and S27†). The catalyst amount and reaction time were optimized in the range of 0.3–1.0 mol% and 12–72 h (Fig. S28 and Table S5†). The control experiment using only TBAB without **M**-CO₂ catalysts resulted in a conversion of 67.5%. The catalytic activity of **M**-CO₂ maintained the conversion exceeding 90% after three cycles, which is comparable to those of MOFs with catalytically active OMS (Table S7†).^{29–31} ICP-OES on the supernatant after the reactions confirms negligible leaching of metal ions from **M**-CO₂ over three cycles (**Al**-CO₂: 0.14–0.33 ppm and **Ga**-CO₂: 0.07 ppm, Table S7†).^{32,33} **Al**-CO₂ also exhibited catalytic activity toward the CO₂ cycloaddition of styrene oxide into styrene carbonate at 80 °C with conversion of 76.3% (Fig. S29 and Table S6†). Although the diffraction intensity of recycled **Al**-CO₂ decreased in PXRD, N₂ adsorption isotherms and *S*_{BET} values were comparable after three cycles (Fig. S30 and S31†). In contrast, the recycled **Ga**-CO₂ exhibited a large decrease in N₂ adsorption, whereas the catalytic activity was preserved. This

indicates that the catalytic reaction in **Ga**-CO₂ mainly occurred at the particle surface rather than internal pores, which is consistent with the catalytic CO₂ cycloaddition at the surface of dense MOFs.³⁴ This is also supported by the smaller pore size of **Ga**-CO₂ than **Al**-CO₂ confirmed by N₂ adsorption (Fig. S32†). The results indicate that CO₂-derived MOFs serve as a catalyst for CO₂ cycloaddition under ambient conditions.

3.4. Formability of transparent monoliths

Hot-pressing provided a transparent monolith of **Al**-CO₂ (Fig. 4A). In contrast, the **Ga**-CO₂ monolith prepared by hot-pressing under the same conditions was not transparent. The PXRD pattern of the **Al**-CO₂ monolith exhibits broad features in the same peak positions before hot-pressing, indicative of preservation of the long-range order (Fig. S33†). The differential scanning calorimetry (DSC) profile of **Al**-CO₂ ground powder does not display any features assignable to the glass transition in the temperature range from –50 to 150 °C (Fig. S34†). The SEM image of the **Al**-CO₂ monolith displays a grain-boundary-free, smooth surface, resulting in the formation of a transparent monolith (Fig. 4B and S35†). The surface hydrophobicity of the **M**-CO₂ monolith was tested by a contact angle analysis (Fig. 4C and S36†). The monolithic samples of **Al**-CO₂ and **Ga**-CO₂ exhibit water contact angles of 121 ± 0.4 and 116 ± 0.8°, which are categorized as hydrophobic surfaces as the angles are greater than 90°. The hydrophobic feature of **M**-CO₂ is attributed to the presence of the methyl group of MeO⁻. The lower hydrophobicity of **Ga**-CO₂ was due to the OH⁻ group which can form hydrogen bonding with water molecules.³⁵

3.5. Porous properties as powder and monoliths

The porous properties of MOFs as a monolithic form are also essential for wide applications such as catalytic processes and volumetric gas adsorption capacity.^{36–38} The gas adsorption measurements were carried out for both powder and monolith samples (N₂ at 77 K, CO₂ at 195 K, and H₂ at 77 K, Fig. 4D, E and S37†). The gas uptake and *S*_{BET} values calculated from the N₂ adsorption isotherms are displayed in Table S8.† The N₂ and CO₂ uptake of **M**-CO₂ monoliths largely decreased by 94 and 81% for **Al**-CO₂ and 85 and 83% for **Ga**-CO₂, respectively (Table S8†). This is consistent with the decrease of *S*_{BET} values of **M**-CO₂ monoliths, which are reduced by 91% for **Al**-CO₂ and 80% for **Ga**-CO₂. In contrast, **M**-CO₂ monoliths exhibit a smaller decrease in H₂ uptake compared to the reduction in N₂ and CO₂ uptake (22 and 63% for **Al**-CO₂ and **Ga**-CO₂, respectively). As the kinetic diameters increase (2.89, 3.30, and 3.65 Å for H₂, CO₂, and N₂),³⁹ larger decreases in gas uptake for both **M**-CO₂ monoliths were observed. This indicates the pore size reduction of **M**-CO₂ through hot-pressing, which is consistent with mechanical pressure-induced pore size reduction observed in flexible zeolitic imidazolate frameworks (ZIFs).⁴⁰ The key for the preservation of the microporosity of **M**-CO₂ as the monolithic form is attributed to the coordination network in which OCHO⁻, capable of adopting various coordination modes, is connected *via* stable Al/Ga–O bonds. Hybrid perovskites composed of OCHO⁻ exhibit structural transformation in

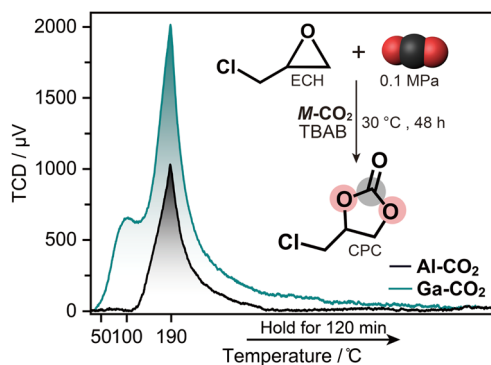


Fig. 3 NH₃-TPD profiles of **M**-CO₂. Inset: conversion of ECH into CPC via CO₂ cycloaddition reaction using **M**-CO₂ catalysis with TBAB as a co-catalyst.





Fig. 4 (A) Optical image of the transparent Al-CO₂ monolith with a diameter of 7.0 mm. (B) SEM image of the Al-CO₂ monolith. (C) Image of a water droplet on the surface of the Al-CO₂ monolith. (D) N₂ and (E) H₂ adsorption isotherms at 77 K for M-CO₂ powder and monoliths. Solid/open symbols indicate adsorption/desorption isotherms, respectively.

response to mechanical pressure, driven by alternation in the coordination modes of OCHO[−].⁴¹ The flexibility of the coordination geometry of OCHO[−] originates from its low steric hindrance.¹⁶ Due to the coordination flexibility and stable Al/Ga-O bonds, hot-pressing does not collapse M-CO₂ into a non-porous structure, and the microporous structure is maintained while the pore size decreases.

4. Conclusions

We demonstrated the one-pot synthesis of amorphous MOFs with permanent porosity, [Al₂(OCHO)₃(OMe)₃] (Al-CO₂) and [Ga₂O_{1.8}(OHCO)_{0.5}(OMe)_{0.9}(OH)] (Ga-CO₂), from CO₂ by converting CO₂ into OHCO[−]. The local coordination geometries of the metal center were revealed by solid-state NMR and synchrotron total X-ray scattering analysis. Al-CO₂ formed a grain-boundary-free transparent microporous monolith *via* hot-pressing. Hot-pressing treatment mechanically reduced the pore size of M-CO₂ monoliths, enhancing adsorption selectivity toward H₂. The key for the formation of the porous monolith was attributed to the coordination flexibility of OHCO[−] and stable metal-oxygen bonds. These findings offer a new design guideline for CO₂-derived porous materials by incorporating structural disorder.

Data availability

The data supporting this article have been included as part of the ESI.†

Author contributions

K. K., S. H. and K. S. designed the project. K. S., K. K., T. K., T. T., H. Y., and K. K. contributed to data collection and formal analyses. K. K., S. H. and K. S. wrote the manuscript and all the authors approved the final version.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

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