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IN SITU DIFFRACTION STUDIES OF SELECTED METAL–ORGANIC FRAMEWORK MATERIALS FOR GUEST CAPTURE/EXCHANGE APPLICATIONS

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4.1 INTRODUCTION

4.1.1 Background

CO₂ is known to be the main anthropogenic contributor to the global climate change, and carbon mitigation approaches are critical for maintaining a sustainable future [1, 2]. At the current rates of energy consumption, known world coal reserves will last for more than 300 years, while known world natural gas reserves will last for about 60 years and oil reserves will last for about 40 years. Carbon capture/sequestration strategies are being developed for power plants worldwide which generally can be summarized in four steps: (1) CO₂ capture, (2) separation at point sources such as fossil-fueled power plants, followed by (3) transportation, and finally (4) long-term storage, primarily via deep underground injection [3].
Capture is the key step for the capture/sequestration process, and is the most research-intensive. Gas retention in porous solids is technically and economically feasible; therefore, development of novel solid sorbent materials could provide a cost-effective way to capture CO₂.

Porous materials offer a wide range of compositions and structures that are suitable for adsorption and capture of CO₂ [4] and other guest molecules. These materials include zeolites [5, 6], activated carbon [7], smectites [8], oxide materials such as calcium oxide [9], lithium zirconates [10], and hydrotalcites [11]. There are also a vast number of reported metal–organic frameworks (MOFs) [12–40] that show diverse capture properties and applications at different pressures and temperatures. The field of MOF research has been rapidly expanding since the past decade. According to Zhou and Kitagawa [12], the surge of MOF research in recent years has been due to five factors: (1) advances in cluster chemistry, (2) maturation of organic synthesis (ligand design and post-synthetic modification of linkers), (3) improvements in evaluation of sorption and structural properties, (4) increase in interdisciplinary MOF investigations, and (5) an expanding potential for applications.

Various techniques have been applied for characterization of MOFs and the associated guest molecules, including structural techniques, adsorption techniques, spectroscopic techniques, and modeling techniques. Up to the present, considerable knowledge of MOF structure and properties has been accumulated; however, there is still a lack of detail concerning guest adsorption mechanisms and guest–host interactions, which are necessary for designing more efficient sorptive materials.

4.1.2 In Situ Diffraction Characterization

Instrumental designs [41–46] for in situ diffraction experiments concerning adsorption of guest molecules in the cavities of MOFs and the synthesis and monitoring MOF phase formation have attracted increasing attention in recent years [32, 47–76]. These experiments continue to facilitate our understanding of adsorption/desorption properties and detailed adsorption mechanisms of various guest species. In particular, neutron and X-ray synchrotron-based diffraction have been used successfully to identify CO₂ adsorption sites in MOFs.

Time-resolved, in situ single-crystal X-ray diffraction and powder diffraction techniques provide the opportunity to study crystalline nanoporous materials under realistic ambient and non-ambient conditions. These measurements will provide experimental data to compliment data that are obtained theoretically or by spectroscopic methods. In situ single-crystal and powder diffraction techniques are being utilized in national laboratory facilities worldwide. As these special facilities have limited availability and large expenses associated with their usage, a summary of the applicability of laboratory powder X-ray diffraction to study the location of CO₂ in solid sorbents is also included here.
Example of applications of in situ diffraction experiments include detection of unstable intermediate stage during guest exchange [41]; materials screening for post-combustion CO$_2$ capture [45]; phase transformation (reversible phase transformation [47]; guest-induced structural transition [48, 49], and multiple-phase transition upon selective adsorption [57, 66]); dynamic CO$_2$ sorption behavior [50]; identification of adsorption/binding site of CO$_2$ and other gases [51, 67]; breathing modes of flexible MOF [52–56]; monitor of mechanochemical reaction-metastable intermediate [58]; gas adsorption mechanism [59]; capture of metastable intermediates in during the MOF formation [60]; internal pore location of CO$_2$ [62]; in situ synthesis and phase formation of MOFs [61, 63, 65, 70–72]; dynamic gas adsorption sites [64]; guest–host intermolecular interactions [68]; topological change/framework distortion related to solvent exchange [69]; kinetics of liquid-enhanced gas uptake [73]; high speed of transient molecular adsorption [74]; and crystal engineering of large discrete cavities [75].

The two goals of this chapter are (1) to give a brief summary of in situ studies of MOFs with regard to sorption of guests (i.e., H$_2$O, CO$_2$), for single crystals as well as powder samples, and (2) to illustrate examples emphasizing the importance of in situ techniques, particularly for unraveling the processes related to the complex behavior of MOF materials.

4.2 APPARATUS FOR IN SITU DIFFRACTION STUDIES

Since 2005, there have been a number of papers discussing in situ studies of porous materials for capture of gases and liquids [41–46]. We will briefly describe several representative cell- and chamber-based designs.

4.2.1 Single-Crystal Diffraction Applications

4.2.1.1 Environmental Control Cell For in situ single-crystal diffraction studies, it is important to have appropriate goniometers that house the single crystals and allow the presence of gas in either a closed system or an open system where the gas is allowed to flow through. Recently, an environmental control cell (ECC) has been incorporated as part of a single-crystal goniometer head [41]. This ECC can be used with any commercially available single-crystal X-ray instrument that is equipped with a flexible tubing source capable of delivering static or dynamic vacuum, liquids, or gases.

Figure 4.1 gives the description of the device and the image of the assembled ECC mounted on a standard goniometer head [41]. The base of the device is modeled after a standard magnetic base and is held by a setscrew. The components of this simple setup include a bent needle, the ECC body, a capillary retaining screw, an o-ring, a capillary, a setscrew, and a thin glass fiber for sample mounting. The cell is easy to use and is completely reusable. The device is nearly identical in size to standard single-crystal mounts, so a full unrestricted range of motion is expected for most commercial goniometers.
4.2.1.2 Environmental Gas Cell  Another environmental gas cell (EGC) which also allows a single crystal to be placed either under vacuum or exposed to a pure gas or a gas mixture was designed by Warren et al. [42] (Figure 4.2). The cell utilizes the Hampton Research short XYZ goniometer head with no modification to allow space for the gas/vacuum pipe connection. The cell body is made of borosilicate or quartz capillary tubing. The capillary tube is glued into a small recess of the EGC head. The EGC base and head were fabricated from stainless steel. A simple Swagelok stainless steel two-way valve was connected to the EGC Nalgene tube allowing vacuum or gas to be fed to the EGC head. Details are given in References [42] and [43].

4.2.1.3 Quartz Pressure Cell  A simple and inexpensive pressure cell for single-crystal study at pressures up to 1 kbar and a special attachment for mounting the cell onto a goniometer were designed by Yufit and Howard [44]. Based on a number of different previous versions, their improved design is much more convenient to use. It also takes advantage of modern diffractometer capabilities, including the use of area detectors.

Certain trade names and company products are mentioned in the text or identified in illustrations in order to specify adequately the experimental procedure and equipment used. In no case does such identification imply recommendation or endorsement by National Institute of Standards and Technology, nor does it imply that the products are necessarily the best available for the purpose.
A schematic of the **quartz pressure cell** (QPC) is shown in Figure 4.3a, where the basic components are illustrated. Figure 4.3b illustrates a QPC on the goniometer of the Bruker Smart CCD 6000 diffractometer. Monitoring the pressure was performed using a standard pressure gauge attached to the pressure line.

![Prototype environmental gas cell (EGC) (SRS on Station 9.8). Source: Adapted from Warren et al. 2009 [42] and Cernik et al. 1997 [43].](image)

![A schematic of the QPC. A, crystal; B, quartz capillary; C, brass part of quartz-metal seal; D, valve 30-12HF2; E, removable handle; F, standard 1/8" connector to a pressure line; G, stud for attachment of the QPC to the goniometer head. (b) QPC on the goniometer of the Bruker Smart CCD 6000. Source: Yufit and Howard 2005 [44]. Reproduced with permission of the International Union of Crystallography, http://ijournals.iucr.org/](image)
4.2.2 Powder Diffraction Applications

4.2.2.1 Environmental Chambers Different options for environmental chambers are provided by manufacturers such as Panalytical, Bruker, and Rikagu Inc. There are also various custom-built chambers such as those being used at national beamlines. Figure 4.4 illustrates several non-ambient chambers (Anton Paar Instruments, Inc.) used by diffractometer manufacturers. The XRK900 chamber can be used with gases such as CO$_2$ up to 10 bars (300–1175 K). The TTK450 chamber operates from 80 to 723 K, whereas the HTK16 chamber operates from 300 to 1875 K.

4.2.2.2 Simultaneous PXRD and DSC Techniques The XRD-DSC system that was developed by Woerner et al. [45] is composed of a Rigaku Ultima IV diffractometer (CuK$_\alpha$ radiation with a D/TeX high-speed linear position sensitive detector), a Rigaku XRD-DSC stage, Rigaku HUM-1 humidity generator, ULVAC vacuum pump (6.66 × 10$^{-5}$ kPa), a vacuum manifold, and a custom-built humid atmosphere swing chamber (Figure 4.5). The humid atmosphere swing chamber (HASC) was necessary to buffer between the humidity generator and a XRD-DSC stage so that both humid atmosphere and vacuum swings could be performed. The design of the XRD-DSC system allows for three types of sample treatments, namely, in situ activation, vacuum swing, and atmosphere swing.

4.2.2.3 Oxford-Diamond In Situ Cell Oxford-Diamond In Situ Cell (ODISC) [46] is a versatile, infrared-heated, chemical reaction cell developed by Moorhouse for in situ study of a range of chemical syntheses using time-resolved, energy-dispersive X-ray diffraction (EDXRD) on beamline I12 at the Diamond Light Source. A specialized reactor configuration has been constructed to enable in situ EDXRD investigation of samples under non-ambient conditions. One can use various sample vessels such as alumina crucibles, steel hydrothermal autoclaves, and

Figure 4.4 Non-ambient chambers for in situ X-ray diffraction studies. Source: Courtesy of Anton Paar Instruments, Inc.
glassy carbon tubes, at temperature up to 1200°C. Figure 4.6a gives the schematic of the ODISC furnace and Figure 4.6b shows the cross section of the lower section of the cell.

### 4.3 IN SITU SINGLE-CRYSTAL DIFFRACTION STUDIES OF MOFs

*In situ* diffraction studies as applied to porous MOF materials are illustrated with the following 12 examples, which include neutron, synchrotron, and conventional laboratory X-ray diffraction under ambient and non-ambient conditions. Crystallographic
observation of adsorbed gas molecules at high temperatures is a highly challenging task due to their rapid motion. In the selected examples, the following features are emphasized: crystal-to-crystal phase transformation, structure transformation due to the presence of guests, phase transformation induced by changes of guests (CO\textsubscript{2}/N\textsubscript{2}), intermediate stage of guest exchange, mechanism of CO\textsubscript{2} adsorption, breathing mode of flexible MOFs, multiphase-transition upon selective CO\textsubscript{2} adsorption, \textit{in situ} metastable intermediate transformation, location of CO\textsubscript{2} in the pores of MOFs, reversible gas sorption, \textit{in situ} study of framework formation, and fast screening of sorbents for CO\textsubscript{2} capture using combined \textit{in situ} XRD and DSC.

**4.3.1 Thermally Induced Reversible Single Crystal-to-Single Crystal Transformation**

Allan et al. [47] demonstrated the detailed reversible phase transformation of single-crystal Cu\textsubscript{2}(OH)(C\textsubscript{8}H\textsubscript{3}O\textsubscript{7}S)(H\textsubscript{2}O)·2H\textsubscript{2}O (Cu-SIP-3) due to the presence/absence of water using \textit{in situ} X-ray diffraction as a function of temperature. X-ray diffraction data were collected over a range of 150–500 K at the Advanced Light Source, Lawrence Berkeley National Laboratory. \textit{In situ} loading experiments were performed using the EGC goniometer head designed by Warren et al. [42].

Cu-SIP-3 undergoes a phase transformation on dehydration. As a result of the loss of coordinated water, there is a change in metal coordination which involves the breaking of several bonds in the low-temperature structure. At temperatures below a dehydration-induced phase transition ($T < 370$ K) the structure was confirmed as being hydrated (left drawings of Figures 4.7a and 4.7b). In the
temperature range where the transition takes place (370 K < T < 405 K), no discrete, sharp Bragg peaks were found in the X-ray diffraction patterns, indicating a significant loss of long-range order. At T > 405 K, the Bragg peaks return and the structure corresponds to a dehydrated phase, Cu-SIP-3 (right drawings of Figures 4.7a and 4.7b). At a temperature up to 150 K, the structure of hydrated Cu$_2$O$_{11}$C$_8$S is monoclinic P2$_1$/n (a = 7.2949(4), b = 18.2726(6), c = 10.1245(6), β = 139.172(4)°). At 500 K, the sample becomes dehydrated Cu$_4$O$_{16}$C$_{16}$S$_2$ (Cu-SIP-3). While the dehydrated form is also monoclinic P2$_1$/n, the cell parameters are different: a = 13.792(2) Å, b = 19.430(4) Å, c = 12.057(2) Å, β = 139.172(4)°. After the crystal was exposed to moisture at 293 K, the structure of the rehydrated sample (Cu$_2$O$_{11}$C$_8$S) returns to monoclinic P2$_1$/n (a = 7.3333(14) Å, b = 18.153(3) Å, c = 10.1729(19) Å, β = 94.379(4)°). Therefore, one can conclude unambiguously using in situ X-ray experiment that the loss of coordinated guest water molecules necessitates coordination changes, leading to lowering of symmetry and almost doubling the size of the asymmetric unit.

4.3.2 Structure Transformation Induced by Presence of Guests

Takamizawa et al. [48] used combined single-crystal and in situ powder X-ray to study crystal transformation from an empty host Rh$_2$(bza)$_6$(pzy) (where bza = benzoate and pzy = pyrazine) to the material after CO$_2$ molecules were adsorbed. The structure of Rh$_2$(bza)$_6$(pzy) consists of parallel one-dimensional chains and isolated cavities of 9 × 4 × 3 Å with narrow gaps of approximately 1 Å at the four corners of the cavities (Figure 4.8). The crystal structure of the host crystal without CO$_2$ and with saturated (sealed-in) CO$_2$ is rather different, namely, triclinic P-1 versus monoclinic (C2/c at 90 K and C2/m at 298 K), respectively [35]. The structure transformation was a result of the changing of interchain distances, slipping of neighboring chains, and tilting of the π–π stacked phenyl rings of the (bza) ligand.

![Figure 4.7](image-url) (a) Chains of copper tetramers in the low-temperature (left) and high-temperature (right) structures of Cu-SIP-3. (b) Copper tetramers in the low-temperature (left) and high-temperature (right) structures (Color scheme: blue balls-Cu, red balls-O, yellow balls-S, and gray balls-C). Source: Allan et al. 2010 [47]. Reproduced with permission of American Chemical Society.
(i.e., tilting of the benzene ring about 9° away from the chain). Figure 4.8b shows that the adsorbed CO$_2$ molecules were confined in the channel and interact with the phenyl rings via phenyl and CO$_2$-quadrupole interaction. In situ high-temperature XRD showed the amount of uptake of adsorbed CO$_2$ decreases as the temperatures increase. Evidence for the diffusion of gas into the crystal through the narrow gaps was obtained from sorption measurements. Interestingly, variable-temperature $^2$H NMR spectroscopy in a constant CO$_2$ pressure (0.1 MPa) showed that the “rotating door” motions of the flipping benzene rings of the host skeleton were responsible for CO$_2$ diffusion between cavities (Figure 4.9) [49].

Figure 4.8  The single-crystal structures of (a) [Rh$_2$(bza)$_4$(pyz)] and (b) CO$_2$-included [Rh$_2$(bza)$_4$(pyz)] measured at 90 K. Thermal ellipsoids are drawn at 30% probability. Source: (a) Zhang et al. 2014 [35]. Reproduced with permission of Royal Society of Chemistry. (b) Takamizawa et al. 2010 [48]. Reproduced with permission of American Chemical Society.

Figure 4.9  Schematic diagram of the “rotating door” motion of the flipping benzene rings of the host skeleton consisting of inner channel walls in [Rh$_2$(bza)$_4$(pyz)]. The light gray arrow indicates the channel direction where CO$_2$ diffuses. Source: Takamizawa et al. 2010 [49]. Reproduced with permission of American Chemical Society.
4.3.3 Dynamic CO$_2$ Adsorption Behavior

Using a bis-triazolate ligand and tetrahedral Zn(II) ion, Liao et al. [35, 50] synthesized a flexible porous coordination polymer [Zn$_2$(btm)$_2$].4H$_2$O, H$_2$btm = (bis(5-methyl-1H-1,2,4-triazol-3-yl) methane) ((MAF-23·4H$_2$O). This porous polymer is functionalized with pairs of uncoordinated triazolate N-donors that can be used as guest-chelating sites to give CO$_2$/N$_2$ selectivity. The dynamic CO$_2$ sorption behavior was monitored by single-crystal X-ray diffraction. According to Zhang et al. [35], MAF-23 showed high saturation CO$_2$ uptake (3CO$_2$ per formula unit). Single-crystal structures of MAF-23·$x$CO$_2$ ($x = 0.00, 0.07, 1.5$, and $2.8$) were measured at 195 K. In MAF-23·0.07CO$_2$, a CO$_2$ molecule was chelated by one of the two crystallographic independent chelating claws, which possessed a methyl group at the ortho-position of the N-donor, demonstrating the weak electron donating effect of the methyl group (Figure 4.10). When the CO$_2$ uptake increases, the unit cell volume of the MAF-23 continuously increases due to the distortion of coordination geometries of the Zn(II) ions and ligand shapes around the methylene groups. As CO$_2$ increases, there is a strong binding affinity due to the action of the guest-chelating claws. The cage continues to deform and there is a balance between attraction of N-donor with the CO$_2$ molecule and the repulsion between the CO$_2$ molecules. With $x = 1–4$, MAF-23·$x$CO$_2$ crystallizes in monoclinic $P2_1/n$. At 195 K, MAF-23·$x$CO$_2$ showed a Type I CO$_2$ sorption isotherm without hysteresis (Figure 4.11), giving an apparent Langmuir surface area of 622(5) m$^2$ g$^{-1}$ and a pore volume of 0.21 cm$^3$ g$^{-1}$.

4.3.4 Unstable Intermediate Stage During Guest Exchange

A successful trapping of a metastable intermediate hydrate phase was illustrated by Cox et al. using the ECC with a previously reported MOF [Co(5-NH$_2$-bdc)
In situ single-crystal X-ray diffraction experiments performed under dynamic gas flow conditions revealed this highly flexible framework is capable of exchanging a wide variety of guest molecules in a single-crystal-to-single crystal manner. It was reported that the anhydrous framework \([\text{Co(5-NH}_2\text{-bdc)(bpy)}_{0.5}]\) was prepared by heating the trihydrate phase under a vacuum at 393 K for 2 h. As the crystal structures contain multiple crystallographically unique solvent molecules, Cox et al. were interested to see if guest exchange, in particular the dehydration reaction of the crystal, occurred via the formation of discrete structures possessing an intermediate number of solvent molecules.

The initial trihydrate structure (Figure 4.12b) has one water molecule coordinated to the Co metal center and two “free” water molecules held within the pore by hydrogen bonds. The second structure, obtained after 15 min of nitrogen flow, is that of a previously unreported dihydrate phase \([\text{Co(5-NH}_2\text{-bdc)(bpy)}_{0.5}(\text{H}_2\text{O})]\cdot2\text{H}_2\text{O}\). When the flow of dry nitrogen gas is halted, the dehydrate phase (Figure 4.12a) is converted back to the starting trihydrate phase within 5 min, presumably by reabsorbing moisture from the air. Sealing the capillary effectively halts the rehydration back reaction and thereby enables the collection of diffraction data on the metastable dihydrate phase. In summary, the ECC was capable of stabilizing a novel metastable intermediate in the dehydration reaction.

Figure 4.11  \(\text{CO}_2\) and \(\text{N}_2\) adsorption (solid) and desorption (open) isotherms of MAF-23•\(\text{CO}_2\). Source: Liao et al. 2012 [50]. Reproduced with permission of American Chemical Society.
Miller et al. [51] studied the adsorption mechanism of the microporous scandium terephthalate, Sc$_2$(O$_2$CC$_6$H$_4$CO$_2$)$_3$ (ScBDC), an attractive small-pore model sorbent for small molecules relevant to CO$_2$ separation technologies. The mechanism of adsorption of CO$_2$ has been determined by single-crystal synchrotron X-ray diffraction at $\approx$230 K at the European Synchrotron Radiation Facility, Grenoble, France. A custom-made gas delivery system was used for the \textit{in situ} gas pressure studied from 1 to 9 bar of CO$_2$.

The details of the adsorption mechanisms were obtained with \textit{in situ} studies from $p_{\text{CO}_2} = (0–1)$ bar. At $p_{\text{CO}_2} = 1$ bar, the symmetry of the crystal has changed from orthorhombic \textit{Fddd} to monoclinic \textit{C2/c} through tilts in the terephthalate linkers with the presence of CO$_2$. CO$_2$ molecules take up different sites in two symmetrically different channels that result from the symmetry change. Figure 4.13a depicts the ScO$_6$ octahedra, and three different terephthalate groups, namely, group 1, group 2a, and group 2b. Group 1 in channel A was found having the orthorhombic structure. Groups 2a and 2b were a result of the rotations of the terephthalate groups. In channel A, the occupancy of CO$_2$ is close to one and the CO$_2$ molecule axis is aligned so that the O atom is pointed toward the hydrogen atoms of the framework phenyl group. In channel B, there are two symmetry-related sites for CO$_2$ but they

Figure 4.12 The reversible conversion of a trihydrate (a), single-crystal structure to dehydrate (b), is conducted by flowing dry N$_2$ through the crystal, and then by halting the flow. The appearance and disappearance of the water molecule can be seen. Source: Cox et al. 2015 [41]. Reproduced with permission of the International Union of Crystallography, http://journals.iucr.org/
cannot be simultaneously occupied. The environments of CO$_2$ in the triangular pores in channel types A and B are shown more clearly in Figures 4.13b and 4.13c.

**4.4 POWDER DIFFRACTION STUDIES OF MOFs**

**4.4.1 Synchrotron/Neutron Diffraction Studies**

The breathing behavior of MOFs for gas adsorption has been reviewed by Alhamami et al. [52]. The unique breathing behavior upon adsorption of gases or solvents underlies their potential application as host materials in gas storage for renewable energy. This unique behavior has attracted widespread attention to designing, understanding, and utilizing properties of these materials. The tools of investigation, in addition to the use of *in situ* diffraction, also include calorimetry, nuclear magnetic resonance (NMR), Fourier transform infrared spectroscopy, and theoretical modeling.
4.4.1.1 Breathing Modes of Flexible MOFs  A unique feature, which distinguishes some MOFs from conventional porous materials, is the ability to “breathe,” that is, to expand or contract in response to external stimuli such as variation in temperature. Such flexible networks are sometimes referred to as “breathing” MOFs. The best-known materials exhibiting breathing mode characteristics comprise the MIL-53 series [52–56]. Structurally, these materials are formed by connections of corner-sharing \( \text{MO}_4(\text{OH})_2 \) octahedra linked by 1,4-benzenedicarboxylic (BDC) acids. As the MIL-53 (Al, Cr) materials are synthesized hydrothermally (MIL-53as), the channels are filled with disordered BDC and \( \text{H}_2\text{O} \) molecules (narrow pore (np) structures) [53]. The narrow pore structure is formed by hydrogen bonding between the water molecule and the carboxylic and hydroxyl groups of the host molecules. Upon dehydration at high temperature, the MIL-53 gives rise to a large pore (lp) structure due to the absence of interactions. Therefore, using heat as an external stimulus, a reversible conversion between the hydrated MIL-53 (np) and the dehydrated MIL-53 form (lp) is observed (Figure 4.14a).

Figure 4.14  (a) Schematic representation of the reversible hydration–dehydration of MIL-53np and MIL-53lp. Source: Seere et al. 2002 [53]. Reproduced with permission of American Chemical Society. (b) Adsorption isotherms of MIL-53 (Cr) showing hysteresis loop. (c) Powder XRD patterns of MIL-53 (Cr) under various pressures of \( \text{CO}_2 \) at 293 K. Source: Seere et al. 2007 [54]. Reproduced with permission of John Wiley & Sons.
An interesting adsorption behavior that has been observed in MIL-53 (Cr) is the adsorption of a large amount of CO₂ molecules at room temperature [54] (Figure 4.14b). After a fast uptake at low pressure (approximately 2–3 mmol g⁻¹), the isotherm reaches its first plateau between 1 and 4 bar followed by an adsorption of more than a double amount of CO₂ at higher pressures. The desorption branch returns to coincide with the adsorption branch at approximately 2 bar. This unusual result could be interpreted as follows: while the degassed solid exhibits an expected lp structure, the first small portion of CO₂ would force the pore to be np due to host–guest interaction. A further absorption of CO₂ at higher pressure reopens the pore to accept more CO₂. This interpretation agrees with in situ diffraction experiments performed at synchrotron radiation facilities under 1–10 bar CO₂ (Figure 4.14c).

4.4.1.2 Multiple-Phase Transitions upon Selective CO₂ Adsorption

Another interesting system (provided by Henke et al. [57]) that exhibits multiple-phase transition and remarkable breathing behavior with respect to the presence/absence of polar molecules in the pores is the flexible alkyl ether-functionalized MOF [Zn₂(BME-bdc)₂(dabco)]ₙ (BME-bdc = 2,5-bis(2-methoxyethoxy)-1,4-benzenedicarboxylate, dabco = 1,4-diazabicyclo[2.2.2]octane) [57]. Apparently, the dried sample exhibits a narrow pore form with almost no accessible porosity. Upon adsorption of polar guest molecules (DMF, EtOH), the framework transforms to an open pore form, with accompanying increase of unit cell volume. This flexibility (contraction of network) is because of the strong interaction between the 2-methoxyethoxy substituents, when guest molecules are absent. This situation is opposite to the situation of MIL-53 discussed previously. When guest molecules are present in this case (CO₂), the unit cell volume increases. Due to the slow adsorption kinetics, an unexpected metastable intermediate form could be identified.

Figure 4.15 gives a sorption isotherm of [Zn₂(BME-bdc)₂(dabco)]ₙ at 195 K. The feature of this isotherm includes a stepwise adsorption and a large hysteresis, typical for flexible MOFs that undergo a structural transition upon adsorption of guest molecules. The quadrupole moment of CO₂ enables the gas to penetrate into the pore because of weak intermolecular interactions. It stays at the narrow pore region from pCO₂ = 0 to 200 mbar. The steep uptake at 200 mbar signifies a structural transformation from the narrow pore to the open pore form. On the desorption branch a large hysteresis was observed which was due to the “breathing” framework.

The in situ XRD patterns corroborate the transformation explanation. Figure 4.16 shows a close-up view of the low angle region of the X-ray patterns recorded as a function of partial pressure of CO₂. The lowest angle peak reflection (2.73° – (001)) is in roughly the same position for all three phases. Structural models of the three different states of [Zn₂(BME-bdc)₂(dabco)]ₙ upon CO₂ adsorption at 195 K are shown in Figure 4.17. The cell parameters are all rather similar to each other in the three forms. Up to 300 mbar, only “narrow pore form” is present. The unit cell is slightly expanded at 200 mbar and shrinks at 300 mbar. From 400 to 900 mbar, an intermediate pore form appears which has a mixture of the narrow, intermediate, and open pore form. At 1000 mbar, the “open pore form” almost exclusively exists.
Figure 4.15  Sorption isotherms at 195 K. Adsorption and desorption branches are shown with closed and open symbols. Source: Henke et al. 2011 [57]. Reproduced with permission of Royal Society of Chemistry.

Figure 4.16  Lower angle region of the PXRD patterns recorded at variable CO$_2$ pressure ($\lambda = 0.4592$ Å). Patterns representing the narrow pore (blue), open pore (red) and intermediate pore form (green). Source: Henke et al. 2011 [57]. Reproduced with permission of Royal Society of Chemistry.
4.4.1.3 Metastable Intermediate Transformation 

Katsenis et al. [58] used in situ, real-time synchrotron powder X-ray diffraction to capture and to monitor a metastable, novel-topology intermediate of a mechanochemical transformation. While amorphization on milling is a well-known phenomenon, spontaneous recrystallization of the amorphous phase by continued milling has not been described prior to the cited work. During the synthesis of MOF Bis (2-methylimidazolyl)-zinc (also known as ZIF-8) by applying the grinding technique, they observed the unexpected amorphization. On further milling, recrystallization into a non-porous material via a metastable intermediate new phase, katsenite (kat), took place. The sequence of solid-state transformations in the reaction of ZnO and 2-methylimidazole (HMeLm) is shown in Figure 4.18. The new topology provides evidence that milling transformation can involve short-lived, structurally

Figure 4.17 Structural models of the three different states of \([\text{Zn}_2(\text{BME-bdc})_2(\text{dabco})]_n\) upon CO\(_2\) adsorption at 195 K as a function of increasing p\(_{\text{CO}_2}\) (L \(\rightarrow\) R from 0 to 1000 mbar). Disordered 2-methoxyethoxy substituents and hydrogen atoms are not included in the model structures. Source: Henke et al. 2011 [57]. Reproduced with permission of Royal Society of Chemistry.

Figure 4.18 Sequence of solid-state transformations in the reaction of ZnO and sodalite (SOD) framework, ZIF-8. Source: Katsenis et al. 2015 [58]. Reproduced with permission of Nature Publishing Group.
unusual phases. Structure of kat-Zn(Melm)$_2$ is shown in Figures 4.19a and 4.19b. kat-Zn(Melm)$_2$ contains pores consisting of tight channels and pockets. It consists of four crystallographically independent zinc ions, each in a tetrahedral environment defined by four nitrogen atoms of four different ligands. This work was also corroborated by in situ NMR studies. The solid-state $^{13}$C NMR spectrum of kat-Zn(Melm)$_2$ indicates four symmetrically non-equivalent Melm-ligands. Samples of dia-Zn(MeIm)$_2$ are non-porous, exhibiting an almost completely flat nitrogen sorption curve.

4.4.1.4 Reversible Gas Sorption Driven by Temperature  Baek et al. (2015) [59] used combined in situ single-crystal and synchrotron powder X-ray diffraction to determine reversible CO$_2$ sorption processes in a self-assembled and apparently non-porous organic crystal, 1,3,5-tris-(4-carboxyphenyl) benzene (H$_3$BTB) (Figure 4.20a). The host material is formed by a hydrogen bond network between H$_3$BTB and N,N-dimethylformamide (DMF) and by the $\pi-\pi$ stacking between the H$_3$BTB moieties (Figure 4.20b). The material can be viewed as well-ordered array of cages which are tight-packed so that the cages are inaccessible from outside. The hydrogen bonds and $\pi-\pi$ stacking are strong enough to give relatively high thermal stability up to $\approx$373 K at 1 bar. The DMF-forming cages could be flexible, providing transient pathways for guest diffusion. The encapsulated DMF molecules are disordered around a site of a threefold rotary inversion symmetry. These molecules can be completely removed in vacuum at 323 K, resulting in very stable crystal (Figure 4.20c). The unit cell volume decreased by 3.23% after removal of DMF molecules. Each cage has about 83 Å which is large enough for capturing small gas molecules such as CO$_2$.  

Figure 4.19 Structure of kat-Zn(Melm)$_2$ viewed along the crystallographic $c$ direction: (a) ball-and-stick representation and (b) the kat framework with different coloring for each type of vertex, represented by its vertex figure. Source: Katsenis et al. 2015 [58]. Reproduced with permission of Nature Publishing Group.
In situ single-crystal X-ray diffraction experiments were done under varying pressures at 323 K. It was found that even though there are no permanent channels connecting the isolated cages, all cages are filled at 25 bar of CO₂ pressure. One cage contains one CO₂ molecule and the cage is composed of two H₃BTB and six DMF molecules, with stoichiometry (H₃BTB)₂(DMF)₆(CO₂). Therefore, the empty cages are permeable to CO₂ at high temperatures due to thermally activated transient pathways (molecular gating) between the cages. Figure 4.20d shows the cage

![Figure 4.20](https://example.com/figure4.20.png)

(a) Structure of (H₃BTB + DMF) showing the hydrogen bonding network. (b) Packing of (H₃BTB + DMF) moieties. Red color represents “O.” (c) Structure after DMF molecules have been evacuated, showing the cages. (d) Views of the cage with a CO₂ molecule (H₃BTB·CO₂). The CO₂ molecule is disordered around a site of a threefold rotary inversion symmetry and is depicted by a space-filling model. Source: Baek et al. 2015 [59]. Reproduced with permission of PNAS.
of \((\text{H}_3\text{BTB})\) with a \(\text{CO}_2\) molecule. The disordered \(\text{CO}_2\) molecule is depicted by a space-filling model.

Further variable-temperature \textit{in situ} synchrotron powder X-ray diffraction studies also confirmed that the \(\text{CO}_2\) sorption is reversible and driven by temperature increase (Figure 4.21). With combined solid-state NMR, they were also able to study the dynamical motion of \(\text{CO}_2\).

\textbf{4.4.1.5 Framework Formation in Action} In order to understand the details of the complex crystallization process of lithium tartrate MOF, Yeung et al. [60] employed high-energy \textit{in situ} synchrotron X-ray powder diffraction at the Diamond Light Source Beamline I12 to study the crystallizations and the metastable intermediates in the phase formation of a lithium tartrate MOF, \(\text{Li}_2(\text{meso-C}_6\text{H}_6\text{O}_6)\).

\(\text{Li}_2(\text{meso-C}_6\text{H}_6\text{O}_6)\) has a monodentate binding site. They observed the successive crystallization and dissolution of three competing MOF phases in one reaction. As shown in Figure 4.22, which gives the three possible products, namely, the low-density phases \(1\) (\(\text{Li}_2(\text{meso-C}_6\text{H}_4\text{O}_6)(\text{H}_2\text{O})_{0.5}\)), a metastable intermediate

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure4.png}
\caption{\textit{In situ} synchrotron PXRD patterns under varying pressures of \(\text{CO}_2\) at 323 K. The red dotted square represents the changes in diffraction peaks at the \((2-1 0)\) plane accompanying \(\text{CO}_2\) adsorption and desorption. The blue dotted square represents the intensity changes in diffraction peaks accompanying framework expansions and contractions upon reversible \(\text{CO}_2\) sorption. \textit{Source}: Baek et al. 2015 [59]. Reproduced with permission of PNAS.}
\end{figure}
2a \( \text{Li}_2(\text{meso-C}_6\text{H}_4\text{O}_6)(\text{H}_2\text{O}) \) (with gauche ligand conformation which is the preferred conformation of the mesotartaric acid), and 2b, a denser \( \text{Li}_2(\text{meso-C}_6\text{H}_4\text{O}_6) \) phase, which has a monodentate binding feature. In situ X-ray diffraction confirmed compound 2b to be the final thermodynamically stable product (Figure 4.23). They further determined the extent of crystallization as a function of reaction time, and quantified the reaction energy by determining the reaction rate constants and activation energies. The structure relationships between reactants and products were used to interpret different reaction rates (i.e., larger changes in conformation gave rise to higher activation energy).

### 4.4.1.6 Resin-Assisted Solvothermal MOF Synthesis

The \([\text{Co(NDC)}(\text{DMF})]\) (NDC = 2,6-naphthalenedicarboxylate) system is a system which is known to produce at least three distinct frameworks during the synthesis process. In order to determine the structural diversity observed in coordination polymers formed using Co(II) NDC and DMF, the system was probed by Moorhouse et al. [61] to determine, in particular, the factors influencing the preferential formation of one structure over another. The process involved the use of cation-impregnated polymer resin as both template and metal source in a relatively new technique in MOF synthesis.
Reactions were conducted using angular-disperse XRD at Beamline I12, Diamond Light Source (with the Oxford-Diamond In Situ Cell (ODISC)). Further in situ energy-disperse XRD experiments were performed by using HASYLAB Beamline F3 of the DORIS synchrotron at the Hamburg campus. It was found that the resin-assisted synthesis resulted in the preferential formation of a topology previously impossible to synthesize in bulk form. The [Co(NDC)(DMF)] framework viewed along the z-axis (showing the diamond shape channels and coordinated DMF) is shown in Figure 4.24. The stack plot in Figure 4.25 shows in situ energy-dispersive XRD data obtained during the solvothermal formation of [Co(NDC)(DMF)] at 200°C. Reflections consistent with undissolved H$_2$NDC are observed initially at 2θ = 2.48° and 3.64°, then diminished and finally disappeared before the autoclave reaches 200°C. All reflections consistent with the [Co(NDC)(DMF)] phase appear concurrently. The framework MOF was formed as a pure phase which is monoclinic with a space group C2/c ($a = 23.436(3)$ Å, $b = 8.7733(6)$ Å, $c = 7.219(1)$ Å, $\beta = 97.80(2)^\circ$).

Figure 4.23 Solvothermal conversions between lithium tartrate MOFs, showing (a, b) the extent of crystallization as a function of reaction time (t). (c, d) In situ XRD data, where 2θ is the diffraction angle. (a, c) Hydrated phase 1 (gray points) is converted into metastable phase 2a (green points), followed by (b, d) conversion of 2a into the thermodynamic product 2b (orange points). $\lambda = 0.2326$ Å. Source: Yueng et al. 2016 [60]. Reproduced with permission of John Wiley & Sons.
Figure 4.24  Structure of [Co(NDC)(DMF)] viewed along the c-axis showing coordinated DMF. Octahedral coordination spheres are shown as polyhedral. Source: Moorhouse et al. 2015 [61]. Reproduced with permission of Elsevier.

Figure 4.25 Three-dimensional-stack plot showing in situ angular-dispersive XRD data at 200°C. Source: Moorhouse et al. 2015 [61]. Reproduced with permission of Elsevier.
4.4.2 Laboratory X-ray Diffraction Studies

4.4.2.1 Location of Adsorbed CO$_2$ Molecules

The highly porous compound HKUST-1 [62, 33] is composed of dimeric copper tetracarboxylate units. HKUST-1 has a three-dimensional channel structure connecting a system of cages. The square-shaped pores are about 10 Å diameter, and the largest ones are hexagonal pores of about 18 Å in diameter. Using neutron diffraction, Wu et al. found that the unsaturated open Cu site plays an important role in enhancing binding of CO$_2$ molecules [32].

In Figure 4.26a, in situ experiments with CO$_2$ delivered at different partial pressures were conducted by Wong-Ng et al. [62] with a Panalytical X-ray diffractometer using the Anton Paar environmental chambers XRK 900 (gas flow conditions) under a series of CO$_2$ partial pressures (0.2, 0.4, 0.8, 1, and 6 bar). Several features of the X-ray patterns that are sensitive to occupancy of the cages by guest species were found. The X-ray patterns containing 311 and 222 reflections were shown: (1) refers to the as-received sample, (2) is the one that received treatment for 24 h at 150°C under flowing He, and exposure to mixture of CO$_2$/He with increasing fractions. The 311 intensity increases from near zero for hydrated sample 1 to dehydrated 2, then decreases back to small value as $p_{CO_2}$ decreases, suggesting the empty pores become occupied by CO$_2$. Figure 4.26b gives an asymmetric unit structure.

One can identify two principal adsorption sites: at “open space” A-site near Cu, and around the B-site, the “octahedral site.” Highly disordered CO$_2$ molecules were found. Using first principles calculations, Zhou et al. [76] predicted CO$_2$ to be linear with the Cu–O1–C angle of 110° (Figures 4.26c and 4.26d). The large open space in the vicinity enables the rotation of the CO$_2$ molecules. The CO$_2$ molecules in the octahedral cage are also disordered. It gives a single quadruple split site at the center of the cage (1/4, 1/4, 1/4) and four overlapped oxygen sites (unresolved double split) along the <111> directions. A plot of the occupancy of the “open space” site and the “octahedral cages” shows progressive increase as $p_{CO_2}$ increases (Figure 4.26e). The open-space site is occupied preferentially. In summary, the adsorption is mainly on two sites, one is on top of the open Cu atom and another one is in the window opening of an octahedral cage. The metal–CO$_2$ binding is due to enhanced electrostatic interaction, and the framework CO$_2$ interaction is of the van der Waals type. The highly disordered CO$_2$ molecules result from the flowing versus static gas situation.

4.4.2.2 Materials Screening for Post-Combustion CO$_2$ Capture (In Situ XRD/DSC) Combined use of in situ X-ray diffraction and simultaneous differential scanning calorimetry (DSC) is a promising technique for rapid evaluation of the suitability of microporous materials for post-combustion CO$_2$ capture in both dry and wet conditions (particularly in the presence of water vapor which is a major component of post-combustion CO$_2$ capture). Using calorimetry, the enthalpy of adsorption for a gas can be directly measured at experimental conditions. This is especially advantageous in the situations of phase transitions, or gate-opening in the presence
Figure 4.26 (a) In situ X-ray diffraction patterns under gas flow conditions under a series of partial pressures, from 0.2, 0.4, 0.8, 1 bar and higher, (1) as-received sample and (2) evacuated at 150°C under He for 24 h; (b) Atomic labeling scheme, including the disordered CO$_2$ molecules; (c) The structure of CO$_2$ molecule adsorption on the open Cu sites of HKUST-1; (d) A projection of the unit cell along ⟨110⟩ direction. Rotational disorder proposed for the CO$_2$ molecules located next to the Cu ions is illustrated by showing multiple molecule positions and further emphasized using arc arrows; (e) The occupancy of the “open space” site (round dots) and the “octahedral cages” (squares) increase progressively as $p_{CO_2}$ increases [numbers in (a); colors in (e)]. Source: Wong-Ng et al. 2016. Reproduced with permission of Elsevier.

of guests. Woerner et al. [45] used this combined technique to study the performance and structural effect of CO$_2$ adsorption under static or dynamically varying humidity, temperature, and gas on a class of potential materials for CO$_2$ capture.

One example illustrating the applications of the simultaneous XRD-DSC technique is the open-framework compound ZIF-7, Zn(bIm)$_2$ (where bIm = 2-benzimidazolate). ZIF-7 is formed by connecting Zn metal clusters through the benzimidazole (BIM) linkers. It has a sodalite topology with a crystallographic six-membered ring pore opening (Figures 4.27a and 4.27b). The position of the benzene rings in the
optimized structure in vacuum narrows the pore entrance to $\approx 0.3$ nm. However, the linkers have some freedom to rotate over a certain angle, allowing molecules larger than 0.3 nm to enter the main cavities.

In Figure 4.27c, cumulative enthalpy isotherms during CO$_2$ adsorption–desorption with XRD-DSC under dry (red) and 75% RH (blue) conditions are shown. The presence of water vapor thus reduces the measured enthalpy and therefore the capacity of ZIF-7 (85.2% of dry). Figure 4.27c shows the results of vacuum-humid atmosphere swings. However, relative to other studied sorbents [45], ZIF-7 still shows strong CO$_2$ affinity even in the presence of 75% RH. Successive XRD patterns under vacuum and under 75% RH CO$_2$ vacuum-humid 1 atm CO$_2$ swings are shown in Figure 4.28. As a summary, ZIF-7 appears to be ideally suited for post-combustion flue gas CO$_2$ capture.
In situ diffraction techniques have been shown to be effective means for study of interactions of MOF with adsorbed guest molecules. In this chapter, we illustrated these techniques with seven examples, representing the capability of in situ diffraction for study of a gas adsorption mechanism, breathing of flexible MOFs, guest exchange studies, a guest-induced structural transition, a reversible transformation, probing of adsorption sites and binding sites of CO₂ and other gases, and in situ synthesis and phase formation of MOFs. In addition, the in situ diffraction techniques can also uniquely be used for monitoring a mechanochemical reaction-metastable intermediate, to capture metastable intermediates in successive crystallizations during the formation of an MOF, and to perform a snapshot analysis of transient molecular adsorption. Some of the studies used in situ diffraction combined with other techniques such as in situ NMR, and in situ DSC. As new in situ technology continues to evolve, we will be able to further our understanding of adsorption mechanisms, to facilitate the improvement of adsorption capability and the design of new applications.

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