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Dynamic breathing behaviour of the titanium-based metal-organic framework NTU-9 upon adsorption of water and organic solvents

What happens to the structure of a framework material when the pore content is changed or reduced? While investigating and optimising the reproducibility of synthesising the two-dimensional titanium-based MOF NTU 9, we discovered a dynamic behaviour of NTU-9 triggered by guest adsorption and desorption. After applying additional stimuli (i.e., vacuum), the material exhibits reversible pore distortion by compression in the lateral dimension, forming a new metastable form, NTU-9-d (d stands for distorted), with a reduction in unit cell volume, pore size, and crystal symmetry.

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Dynamic breathing behaviour of the titanium-based metal–organic framework NTU-9 upon adsorption of water and organic solvents†

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Understanding the structural response of framework materials to external stimuli has been of great interest. However, little is known about the stimuli responsiveness of titanium-based metal–organic frameworks. We investigate the reproducibility of the synthesis of the two-dimensional metal–organic framework NTU-9 (NTU = Nanyang Technological University) composed of Ti⁴⁺ cations and 2,5-dihydroxyterephthalate as the organic linker and the flexible response of the framework structure to external stimuli. Using acetic acid simultaneously as mediator and solvent leads to the reproducible formation of large NTU-9 crystallites after long reaction times. The MOF synthesis in isopropanol:acetonitrile (*i*-PrOH:MeCN) mixtures without a modulator is significantly faster but yields smaller NTU-9 crystallites. Pure heat treatment under ambient conditions removes a significant amount of the incorporated host solvent molecules without alteration of the framework's structure. After applying additional external stimuli (*i.e.*, vacuum), the material exhibits a pore distortion by compression in the lateral dimension, depending on the synthetic procedure. The new, distorted, metastable form NTU-9-d shows a reduction in unit cell volume, pore size, and crystal symmetry. Under humidity/air exposure or solvent resuspension, the framework reverts into its original state. The synthesis conditions significantly affect the flexibility of the MOF structure, where samples synthesized without modulator showed a lower tendency for distortion. Our results emphasise the importance of an in-depth understanding of the structure–property relationships in flexible MOFs through a detailed characterisation of the material's stimuli responsiveness process.

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Introduction

Metal–organic frameworks (MOFs) are porous reticular materials consisting of inorganic nodes (*e.g.*, metal ions, clusters) linked by organic bridging ligands through coordination bonds.^{1–3} Among them, MOFs constructed from tri- and tetra-valent metal-based clusters combined with carboxylate-based ligands show higher stability according to the hard and soft acids and bases (HSAB) principle,⁴ which makes them suitable for applications such as gas storage and separation, as well as catalysis.^{2,5}

Titanium-based MOFs are of particular interest due to the earth abundance of their metal source, high stability, low toxicity, redox activity, and promising photochemical properties.⁶ The latter pave the way for attractive applications, such as in photocatalysis, because tetravalent titanium cations exhibit an empty d-shell constituting the conduction band minimum, which can generate long-lived charge carrier, resulting in high solar-to-chemical conversion efficiencies.^{7,8} However, the synthesis of carboxylate-based Ti⁴⁺-clusters is challenging due to the high reactivity (prone to fast oxidation and oxolation) and the hydrolytic instability of titanium precursors, impeding the crystallisation process.^{9–11} Ti⁴⁺ ions are highly susceptible to hydrolysis, which results in the rapid formation of various molecular titanium-oxo-clusters and, eventually, in the precipitation of TiO₂. As a result, significantly fewer titanium-based MOFs have been reported in the literature compared to other MOFs based on transition metals such as Zn²⁺, Cu²⁺, and Zr⁴⁺.^{9,12–14} Useful synthesis strategies for enabling the formation of Ti-MOF single crystals have been introduced by the Marti-Gastaldo group, *i.e.*, by metal doping¹⁵ or using hydroxamate-

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based linkers.¹⁶ Another strategy was proposed by Li *et al.* using a chelating coordination modulation method.¹⁷

One of the most interesting features of reticular materials is the combination of robust and flexible entities in a collective structure, which is also referred to as soft porous crystals^{18,19} or flexible framework materials^{20–23} which react to external stimuli (*e.g.*, temperature, pressure, light) by reversibly changing the framework structure without bond-breaking.^{18,19} These materials can respond to incorporation and release of guest molecules with structural deformations of the framework, resulting in expansion or shrinkage of the pore space, also referred to as breathing behaviour. Since then, framework materials have been classified into three categories according to their softness:^{18,19} flexibility through synergistic effects of the metal nodes' coordination symmetry, the rotatable axis of organic linkers, and strong host–guest interactions.^{22,24–27}

Férey and co-workers first reported on the breathing/swelling behaviour in MOFs in 2002.²⁸ To date, the dynamic behaviour of titanium-based MOFs is rarely described in the literature, and only a few reports are published.^{9,29–31} In COK-69, the framework flexibility results from the dynamic conformational change of the (*cis/trans*)1,4-cyclohexanedicarboxylate linker upon solvent removal.⁹ Another example is the heterometallic (TiMn₂) MUV-35, where the single-crystal transformation is controlled by linker conformation in open, intermediate, and closed states.³⁰

In the following, we report the synthesis, activation procedure, and the dynamic behaviour of the Ti-based MOF NTU-9 (NTU = Nanyang Technological University) triggered by guest adsorption and desorption. We optimised the synthesis parameters and evaluated for reproducibility towards large single crystals and microcrystalline bulk powders. Furthermore, we discovered a new metastable form NTU-9-d (d stands for distorted), which exhibits a strong pore and unit cell distortion upon activation. We investigated the structural transitions by *in situ* powder X-ray diffraction (PXRD) and THz Raman spectroscopy. These investigations revealed the first dynamic breathing behaviour of a titanium-based metal–organic framework with a rigid linker upon solvent loss and exposure to humidity/air. Our work establishes a reproducible synthesis procedure and a detailed mechanism of the previously unknown framework dynamics.

Results and discussion

Synthesis, reproducibility and materials characterisation

In 2014, Gao *et al.* synthesised NTU-9 (C₂₄H₆O₁₈Ti₂) as a photoactive titanium(IV)-based MOF with p-type semiconductor behaviour and utilised it for the photochemical degradation of organic dyes.³² This 2D MOF (Fig. 1) results from the reaction between titanium(IV)-isopropoxide and 2,5-dihydroxyterephthalic acid (H₄dhta) in acetic acid, forming dark red hexagonal crystals composed of a single Ti ion-based hexagonal framework in an underlying **hcb** net.^{32,33} Later, investigations on NTU-9's efficient gas separation ability with an optimised synthetic procedure were reported.³⁴

We first synthesised NTU-9 by reproducing different synthesis conditions reported in the literature.^{32,35–37} These

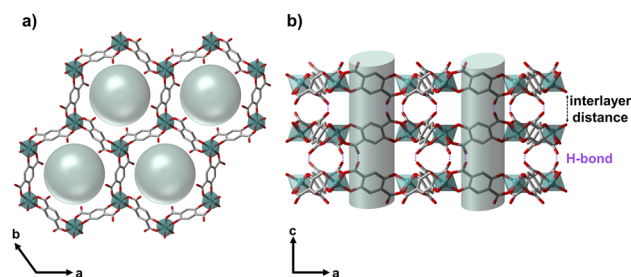


Fig. 1 Image of the crystal structure of NTU-9 (ref. 32) (a) along the pore (view along *c*-axis) and (b) visualising the layer structure (view along *b*-axis), with the pores and pore content, *i.e.*, positionally disordered solvent molecules.

synthetic routes differ in terms of solvent (acetic acid *vs.* *i*-PrOH : MeCN mixture) and reaction time (5 d *vs.* 1 d). Gao *et al.* reported the first solvothermal synthesis **S1_{5d}** (Solvent_{reaction-time}) in acetic acid (**1**), acting as both modulator and solvent.³² The acidic medium during synthesis promotes a slow seeding by shifting the deprotonation equilibrium of the linker H₄dhta toward the precursor side, thus slowing down the reaction, resulting in larger crystals.^{33,38} In contrast, Yu *et al.* used a 1 : 1 mixture of *i*-PrOH and MeCN (**2**) in an alternative synthesis **S2_{1d}** in the absence of modulator and decreased the reaction time to one day instead of five.³⁵

As the limited reproducibility of MOF syntheses is a recurring problem that is gaining attention in the field,^{39,40} we statistically studied the influence of the reaction time for each synthetic protocol by performing the reaction between 4 and 21 times. Indeed, we found different trends depending on the solvent used (Table 1 and Fig. S1, S2†).

Half of the **S1_{5d}** syntheses resulted in the formation of an amorphous product (Fig. S3†), as has been previously observed by other groups.⁴¹ The successful **S1_{5d}** attempts resulted in the formation of a heterogeneous mixture of dark red, crystalline hexagonal NTU-9 and microcrystalline powder (Fig. 2c and S4†). We confirmed the phase purity of the bulk powder by PXRD measurements (Fig. 2a). Scanning electron microscopy (SEM) revealed the formation of crystals with varying size (20–50 μm) and microcrystalline powder with average crystallite size smaller 0.5 μm (Fig. 2c). **S2_{1d}** syntheses also resulted in the expected phase and showed uniformly sized (≈ 2.5 μm) hexagonal red crystals (Fig. 2a, b and S4†). The refined lattice parameters (Fig. 2a, inset) differ slightly from the reported values for the crystallographic *c*-axis and, hence, cell volume.³²

Additionally, FT-IR data^{32,35,42} (Fig. S5†) as well as THz Raman spectroscopy⁴³ (Fig. S6†) of the synthesised solids align with the reported literature and indicate successful formation of NTU-9 for both synthesis pathways. More detailed information regarding the synthetic conditions can be found in the ESI.†

We were able to solve the reproducibility problem for **S1** by doubling the reaction time (**S1_{10d}**). SEM images of **S1_{10d}** revealed a uniform distribution of larger agglomerated hexagonal crystals (120 μm). Conversely, decreasing the reaction time to three days did not yield any significant difference (**S1_{3d}**) in terms of reproducibility, and decreasing it further did not



Table 1 Synthetic conditions for NTU-9 varying in solvent (acetic acid vs. *i*-PrOH : MeCN) and reaction time (between 15 min and 10 days). Reactions yielded an amorphous phase (red), phase mixture (yellow), or pure phase (green) of NTU-9. (successful synthesis/total reaction attempts)

Time	S1 ^a	S2 ^b
15 min ^c	Amorphous phase (0/4)	Pure phase (4/6)
1 d	Amorphous phase (0/4)	Pure phase ³⁵ (10/10)
3 d	Pure phase (2/4)	Phase mixture (0/4)
5 d	Pure phase ³² (10/21)	Phase mixture (1/4)
10 d	Pure phase (8/8)	Phase mixture (1/5)

^a Acetic acid as solvent. ^b 1 : 1 ratio of *i*-PrOH : MeCN as solvent.

^c Microwave-assisted reaction.

produce any crystalline product. The attempt to change from solvothermal to microwave-assisted synthesis to further reduce the reaction time, as shown by Yan *et al.*³⁷ with formic acid, lead to amorphous products for the acetic acid based reaction.

On the contrary, we found that a microwave-assisted synthesis could reduce the reaction time to 15 minutes for the synthesis without modulator (S2_{mw}), yielding a mixture of polycrystalline powder and different-sized hexagonal crystals. Instead, increasing the reaction time resulted in a phase mixture (S2_{3d}, S2_{5d}, S2_{10d}). Additional broad reflections in the diffraction patterns could not be assigned to one of the reported polymorphs of NTU-9^{17,44} (Fig. S7 and S2†).

S1_{3d}, S1_{5d}, S1_{10d} and S2_{mw}, S2_{1d} resulted in phase-pure NTU-9 with slight peak shifts in the PXRD patterns. The peaks of the resulting products of synthesis route S2 are slightly shifted to lower 2θ values, indicating a larger unit cell volume, compared to S1, which is confirmed by Rietveld refinement⁴⁵ for each sample (Fig. 2, S8 and Table S1†). All products crystallise in the trigonal space group $P\bar{3}1c$; but the *c*-axis increases for the S2. This variance may result from different pore fillings and solvent interactions. Despite multiple washing steps of each sample with ethanol, it is possible that a complete solvent exchange did not occur, and the



Fig. 2 (a) PXRD pattern of NTU-9 calculated from the single crystal (black)³² including the refined and reported lattice parameters of the as-synthesised S1_{5d} (red) and S2_{1d} (blue) isolated crystallites. Inset: table with corresponding space group, refined lattice parameters, and unit cell volume. SEM images of (b) S2_{1d} and (c) S1_{5d}.

reaction solvent might exhibit a stronger binding interaction with the framework. This leads to an extension or shrinking of the interlayer distance (Fig. 1b), which indicates a tendency for NTU-9 to be flexible towards different solvents.

Stability towards solvents – guest molecules for structural stabilisation

We studied the framework–solvent interaction by solvent exchange of the incorporated solvent molecules, letting the NTU-9 crystals soak in polar and non-polar solvents (hexane, H₂O, EtOH, acetone, *i*-PrOH, THF, DMF, MeCN) for three days at room temperature (Fig. 3). The overall PXRD patterns of the resulting solvent-exchanged NTU-9 samples can be indexed by the reported trigonal cell, which shows that the MOF is stable in different solvents. Some peaks, however, are significantly shifted, *e.g.*, 002 or 102. This could indicate that different-sized solvent molecules are accommodated within the pores. With different solvents incorporated, the unit cell volume slightly varies between 2174(2) Å³ and 2208(2) Å³. The volume change mainly originates from a slight change of the *c*-axis (Fig. S9 and Table S2†). This behaviour can be explained by the anisotropic structure of NTU-9 (Fig. 1): in the *ab*-plane, covalent and coordination bonds are stabilising the framework, which is chemically stiff. Perpendicular to the *ab*-plane, the 2D sheets are stabilised between each other by hydrogen bonds and van der Waals forces, which are chemically less strong and more elastic. The layered structure in the *c*-direction can thus be more easily expanded and accommodates different-sized solvent molecules, also depending on the specific solvent–framework interactions. After resuspension of NTU-9 in DMF, new, unindexed peaks



Fig. 3 PXRD patterns of S1 with different solvents used for solvent-exchange. Dotted lines highlight the peak shifts and red asterisks indicate newly formed peaks.



emerge in the PXRD patterns, indicating the very slow formation of an additional, unknown phase (Fig. 3, S9 and S10[†]).

Thermal stability at ambient pressure and under vacuum

Next, we explore the thermal and solvolytic stability of NTU-9 (**S1_{5d}**) in air and under dynamic vacuum. Thermogravimetric analysis (TGA) in synthetic air shows that the material is stable up to around 290 °C. Two main mass loss steps can be distinguished. Step I (20 wt% mass loss) in the temperature range of 30–220 °C can be assigned to the release of solvent molecules. The combustion of the linkers, which is associated with the deconstruction of the framework structure, leads to step II (48 wt% mass loss) in the range of 220–460 °C. At 460 °C, a small, additional mass loss occurs (3 wt% mass loss), and the TG-curve eventually reaches a plateau at 550 °C (Fig. 4a). *Ex situ* PXRD analyses show that combustion of NTU-9 leads to the formation of pure rutile (TG temperature 1000 °C). TGA was measured twice (5 K min⁻¹ vs. 10 K min⁻¹) for the same batch to distinguish the solvent loss, which is assigned to be around 20 wt% (19 and 21 wt% for the different heating rates). We cannot be sure that all solvent is removed in the first step and remaining coordinating solvent molecules can be expected due to diffuse, residual electron density in the pores, as established by Rietveld refinement of the dried NTU-9 (Fig. S11–S13[†]).

For structural insights, we performed temperature-dependent *in situ* PXRD using the same sample (**S1**) within the temperature range of solvent release, *i.e.*, up to 220 °C. The heating results in changes in peak intensity, attributed to the release of adsorbed solvent molecules and peak broadening, indicating a decrease in crystallinity. The absence of a strong peak shift illustrates the low thermal expansion of NTU-9 and shows its rather strong thermal stability, as the overall pattern and, therefore, the framework of NTU-9 remains unchanged. The 100 and 110 peaks do not change positions but broaden, whereas the other peaks shift to slightly higher 2θ values, indicating smaller lattice parameters as the loss of guest molecules likely leads to a smaller lattice contraction (Fig. 4b).

A typical activation procedure for MOFs employs heating under dynamic vacuum.⁴⁶ Different activation procedures for NTU-9 are reported, *e.g.*, 120 °C under vacuum overnight,³² dried at 60 °C,³⁵ without dynamic vacuum, or 80 °C under



Fig. 5 PXRD pattern of **S1** (left) and **S2** (right) at different stages of drying: as-synthesised (black), 120 °C heated (red) and exposed to dynamic vacuum at 60 °C (blue).

vacuum for 24 h.³⁶ Surprisingly, PXRD revealed a different behaviour when activating NTU-9 using heat only (under ambient pressure) (Fig. 5, red lines) compared to applying additional dynamic vacuum (Fig. 5, blue lines). For the latter, we observed a pronounced peak splitting that implies a structural distortion. The distorted structure is characterised by a peak splitting of the 100 reflection and a shift of peaks, *i.e.*, 110 or 002, in the PXRD pattern. For **S2_{1d}**, the conversion to NTU-9-d is incomplete, resulting in a phase mixture of NTU-9 and NTU-9-d. In the mechanistic study to understand the distortion, we focus on synthesis **S1**, which results in NTU-9-d upon evacuation and not in a phase mixture (Fig. 5, S14 and S15[†]).

Characterisation of the new form (NTU-9-d)

We derived a structural model of the distorted NTU-9-d from the PXRD data. Starting from the trigonal structure of non-distorted NTU-9, we gradually reduced the space group symmetry and performed global optimisation of the lattice parameters by performing multiple Rietveld refinements⁴⁵ (further details are given in the ESI[†]). Eventually, we were able to describe the diffraction data of the distorted NTU-9 by using a triclinic unit cell with space group $P\bar{1}$ (Table 2). After distortion, NTU-9 is contracted along the a axis and the β angle significantly decreases. This leads to a decrease of 9% in unit cell volume from 2192(2)Å³ to 1995(4)Å³ (activation *via* vacuum at 60 °C) and to an ellipsoidal pore distortion. (Table S3[†]) This behaviour is unusual in 2D frameworks as it is not the elastic axis c (layer distance) that changes, but the distortion occurs in the ab -plane, which results in the pore deformation. The pore channel running along the c -direction is slightly tilted and perpendicular to the ab -plane (Fig. 6). After the Rietveld refinement using only the node and the linkers, a significant amount of residual electron density is still visible within the pores (Fig. S13[†]). The broad, anisotropic peak shape of NTU-9-d indicates a significant structural disorder of the distorted framework. This led to instabilities in the final Rietveld refinements, when we attempted to refine the cation and linker position. Consequently, we can neither report reliable bond angles and lengths between the titanium nodes and the linker molecules nor quantify the residual amount of guest molecules within the pores.

We investigated the nature of this structural distortion by different activation methods: evaporation at room temperature/

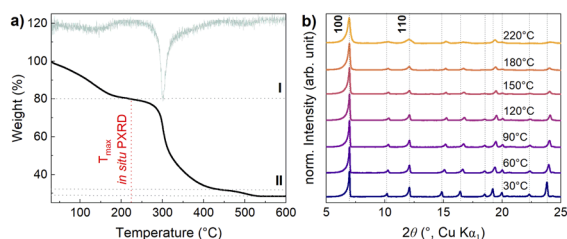


Fig. 4 (a) TGA data of **S1**, heating rate 5 K min⁻¹ under synthetic air atmosphere (flow rate 70 mL min⁻¹) (black) and the derivation of the TGA data (light green). (b) Temperature dependent (30–220 °C) *in situ* PXRD measurements including selected reflection indices using an open capillary of **S1**. Dotted lines are included to indicate the peak shifts.



Table 2 Cell parameters of NTU-9 and the resulting parameters of the model from NTU-9-d by different activation methods

Activation procedure	As-synthesised	120 °C, 7 h	120 °C, N ₂ , 7 h	scCO ₂	Vacuum, rt	Vacuum, 60 °C
Polytype	NTU-9	NTU-9	NTU-9-d (83.8 wt%) NTU-9 (16.2 wt%)	NTU-9-d	NTU-9-d	NTU-9-d
Space group	<i>P</i> $\bar{3}1c$	<i>P</i> $\bar{3}1c$	<i>P</i> $\bar{1}$	<i>P</i> $\bar{1}$	<i>P</i> $\bar{1}$	<i>P</i> $\bar{1}$
<i>a</i> /Å	14.63(1)	14.62(1)	13.24(1)	14.15(1)	13.42(1)	13.24(2)
<i>b</i> /Å	14.63(1)	14.62(1)	15.23(1)	15.00(1)	15.36(1)	15.14(2)
<i>c</i> /Å	11.82(1)	11.53(1)	11.35(1)	11.50(1)	11.37(1)	11.31(2)
α /°	90	90	94.5(1)	92.3(1)	94.2(1)	94.3(1)
β /°	90	90	79.9(1)	85.6(1)	80.6(1)	79.9(1)
γ /°	120	120	116.0(1)	118.5(1)	117.1(1)	116.6(1)
<i>V</i> /Å ³	2192(1)	2133(1)	2026(1)	2138(2)	2060(2)	1995(4)

ambient pressure (overnight), heating at 120 °C under ambient pressure (7 h), heating at 120 °C under continuous nitrogen stream (7 h), supercritical CO₂ activation (scCO₂), and under dynamic vacuum (with or without additional heating). We observed that NTU-9-d formation was triggered by dynamic vacuum, scCO₂, or heating in a continuous nitrogen flow, whereas drying the sample up to 220 °C at 1 bar (ambient pressure) did not distort the framework (Fig. 4b). The refined cell parameters of the differently activated samples **S1** show that the cell distortion varies depending on the activation process. For the sample dried with only heat under nitrogen stream a mixture of NTU-9 and NTU-9-d is present. For the scCO₂-activated sample, we observed less pronounced changes in the cell parameters compared to the vacuum-activated analogue; by dynamic vacuum exposure, we obtained a more distorted pore with an overall smaller cell volume, which can be amplified by additional heat. We can state that drying under vacuum distorts the pore most efficiently (Table 2, Fig. S16 and S17†).

Although vacuum-induced structural transformations in MOFs have been investigated,⁴⁷ this is the first time that the flexibility behaviour of NTU-9 has been reported. We hypothesise that the differences between synthesis protocols (**S1** vs. **S2**) in the activation process (NTU-9-d vs. phase mixture) can be either explained by the difference in crystallite size which is known to impact dynamic framework transitions,⁴⁸ or by the difference in the kind or position of the guest molecules within the pores. We noticed a difference by nuclear magnetic resonance (NMR) of the digested samples with and without modulator. In synthesis **S1**_{5a}, the digested sample shows residual acetic acid (Fig. S18†).



Fig. 6 Structural model of NTU-9-d's distorted crystal structure upon vacuum exposure (a) in the *ab*-plane and (b) along the *c*-axis. Green ellipsoidal cylinders represents the random distribution of solvent molecules within the channels.

To the best of our knowledge, no external stimuli-induced flexible behaviour for polymorphs of NTU-9 such as MIL-167-169 or FIR-117-119 has been reported as yet.^{17,44} However, some flexibility studies were reported for FIR-131-138, with sulfonated linker, when soaking crystals in different solvents.^{49,50}

To investigate the vacuum influence on the distortion of NTU-9 in more detail, we performed *in situ* PXRD under dynamic vacuum. Sample **S1** was first heated to 60 °C followed by a stepwise reduction in pressure, eventually reaching a final dynamic vacuum of 3×10^{-3} mbar while constantly measuring diffraction patterns (Fig. 7a). At first, NTU-9 remained undistorted when heating at 60 °C and under ambient pressure. A slight decrease of the *c*-axis (Fig. 7b, green squares) indicates the incipient release of solvent molecules. Further pressure decrease to 200 mbar initiates the formation of NTU-9-d, indicated by the degeneration of the *a*- and *b*-axes and the α - and β -angles (Fig. 7b and c). The gradual distortion of the trigonal unit cell towards triclinic metrics and the reduction of the unit cell volume continues with further reducing the pressure. Interestingly, the transition from NTU-9 to its distorted form does not affect the contraction behaviour of the elastic *c*-axis.

The formation of NTU-9-d can also be followed by *in situ* THz Raman spectroscopy in dynamic vacuum at 60 °C. Changes from the initial spectrum compared to the vacuum evacuated sample (**S1**) at different pressure values between 0 and 1100 mbar and 60 °C can be observed (Fig. S6b and c†). The vibrations of the organic H₄dhta linker dominate the Raman spectrum. Here we mainly focus on the Ti–O stretching band around 683 cm⁻¹.^{43,51} The band broadening in the process of vacuum exposure indicates changes in the Ti–O bond upon pore distortion. We can assume that, as proven for Zr-MOFs, node distortion caused by guest removal/addition can also occur in Ti-MOFs.⁵² Broadening can also be observed in the symmetrical stretching vibration mode around 700 cm⁻¹.⁴³ This spectroscopic result hints at the formation of the distorted form NTU-9-d and, in particular, indicates changes in the octahedral titanium-linker coordination (Fig. S6b and c†).

CO₂ gas adsorption experiments were conducted to analyse porosity and adsorption behaviour of NTU-9-d (**S1**). The CO₂ adsorption isotherm was collected at 273 K and resembles a type I isotherm, indicating the present micro-pores (<2 nm) (Fig. S19a†). NTU-9-d adsorbed 72 cm³ g⁻¹ of CO₂ with a total pore volume of 0.15 cm³ g⁻¹ and a calculated Brunauer–Emmett–Teller (BET) surface area of $S_{\text{BET}} = 199 \text{ m}^2 \text{ g}^{-1}$





Fig. 7 (a) *In situ* diffraction data of sample S1. The pressure was gradually reduced from ambient pressure to dynamic vacuum at 10 mbar at 60 °C. Evolution of (b) lattice parameter changes, volume changes, and (c) the unit cell angle changes during *in situ* vacuum PXRD.

(Fig. S19b†). The calculated pore size distributions is centred at 0.6 nm (Fig. S19c†). In comparison to a reported similar procedure for S2 from Cai *et al.* (2 d reaction time, without ethanol solvent exchange), the CO₂ uptake of NTU-9 (no peak splitting observed) is about 12% higher with a largely higher BET surface area (917 m² g⁻¹ from N₂ sorption data).³⁴ These differences could originate from the tendency of S1 to distort, which causes lower accessibility of the material's pores, resulting in lower uptake and BET surface area. PXRD data after gas adsorption experiments indicate the stability of the distorted pore (Fig. S19d, S20 and Table S4†).

Reversibility of the deformation of NTU-9

After applying vacuum, NTU-9-d can be transformed back into its original state by exposure to moisture. We monitored the structural changes upon water uptake of NTU-9-d from the ambient atmosphere (22 °C, 25, 5% R.H.) in an open glass capillary with an opening diameter of 0.7 mm by PXRD over a period of 42 hours (Fig. 8).

The *c*-lattice parameter is hardly influenced by the water uptake (Fig. 8b, light green), whereas the *a*- and *b*-axis exhibit significant changes. Initially, both the *a*- and the *b*-axis expand upon exposure to ambient conditions (Fig. 8d, cyan and olive), which leads to a rapid increase in the unit cell volume (Fig. 8b, black). After approximately three hours, the *b*-axis starts contracting while it keeps expanding, leading to an overall expansion of the unit cell volume. Both unit cell axes become more similar in length, and the γ -angle between *a* and *b* expands towards 120° (Fig. 8c, light green). While the α -angle slightly decreases towards 90°, the β -angle rapidly expands to 90° (Fig. 8c, cyan and olive). Consequently, water uptake from the atmosphere distorts the ellipsoidal pore back towards the round shape of NTU-9, and the tilted channel becomes more perpendicular to the *ab*-plane. Despite the fact that the cell relaxes towards the trigonal metric of NTU-9, it does not reach this state after 42 hours of exposure to



Fig. 8 (a) *In situ* PXRD measurement of open capillary of vacuum dried S1 over time (RH of 25 ± 5%). Evolution of (b) lattice parameter changes, volume changes, and (c) the unit cell angle changes during the rehydration of NTU-9-d by ambient atmosphere.

the ambient atmosphere, indicating a relatively slow water uptake from the atmosphere (Fig. S6d†).

We confirmed by *ex situ* PXRD that resuspending NTU-9-d in different solvents (ethanol, water, acetonitrile) overnight results in the recovery of NTU-9 (Fig. S21 and S22†). Different solvent molecules inside the filled channels lead to slight differences in the lattice parameters, in particular in the length of the elastic *c*-axis, of the resulting undistorted NTU-9 (Table S5), demonstrating the adaptive nature of the framework towards different guest molecules. Interestingly, we observe that the reversibility, from NTU-9-d to NTU-9, is significantly faster in the liquid phase compared to the gas phase. We demonstrated full reversibility of this desolvation-solvation process over 5 cycles by PXRD (Fig. S23†).

Conclusions

In this work, we explore the impact and reproducibility of the synthesis conditions on the formation of the layered Ti-MOF NTU-9. We observe that the selection of solvent, additive, and reaction time is crucial for a reliable, reproducible synthesis of the phase pure material. Using pure acetic acid as modulator and solvent at 120 °C leads to the reproducible formation of large NTU-9 crystals. When long reaction times of at least 10 days are applied, the presence of acetic acid in the framework can be detected, even after vigorous post-synthetic washing steps. Solvothermal synthesis in an *i*-PrOH:MeCN mixture without a modulator at 120 °C is faster (1 day) and can be further sped up using microwave-assisted synthesis (15 min), but yields smaller NTU-9 crystals.

Pure heat treatment of the solvated MOFs up to 220 °C and ambient pressure removes large amounts of the incorporated guest solvent molecules without changing the framework's structure. However, activating the material with heating under a nitrogen flow, scCO₂, or vacuum results in a previously unknown, distorted structure, abbreviated as NTU-9-d. The



structural distortion transforms the circular pore channels into an ellipsoidal shape and reduces the unit cell volume by 9%. The distortion mechanism is primarily governed by compression within the hard coordination plane (*ab*-plane, metal–ligand coordination connectivity), with only minor changes in the elastic axis *c* (layer stacking *via* hydrogen bonding). This result is particularly interesting, as it contradicts the prevailing assumption that the soft connectivity between layers predominantly drives flexibility in 2D layered frameworks. Raman spectroscopy indicates a distortion of the octahedral coordination geometry of the titanium cations. Interestingly, the synthesis conditions strongly affect the flexibility of the MOF structure, where samples synthesised without modulator yield smaller crystals and show a lower tendency for distortion. The dried and distorted MOF is able to take up water from the atmosphere and a variety of polar and non-polar solvent molecules from the liquid phase, leading to a reversible structural relaxation into the undistorted form of NTU-9 over several cycles. Incorporating different solvent molecules into the structure of NTU-9 leads to significant changes in the elastic *c*-axis lattice parameters, which correspond to the interlayer distance. Despite the observed large flexibility of the framework, the long-range order is still maintained after multiple cycles of de- and resolvation and the associated pore distortion and relaxation.

This work illustrates the often-overlooked dynamic properties of reticular materials in general and Ti-based MOFs in particular under solvent evacuation/resuspension. Our study thus suggests that thorough characterisation of reticular materials for each process is essential to deeply understand framework behaviour and its resulting applications.

Data availability

The data supporting this article have been included as part of the ESI.† The authors have cited additional references within the ESI.†^{53–59}.

Author contributions

J. E. K. and S. B. conceived the idea of the project. J. E. K. synthesised and characterised the metal–organic framework. F. H. performed CO₂ sorption measurements. K. Gj. measured the *in situ* Raman data, A. S. the *in situ* vacuum PXRD data. S. B. performed the *in situ* rehydration measurements and the Rietveld refinements. J. E. K. wrote the manuscript with contributions from all authors. All authors read and commented on the final manuscript.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 S. L. James, Metal-organic frameworks, *Chem. Soc. Rev.*, 2003, **32**, 276–288.
- 2 H. Furukawa, K. E. Cordova, M. O'Keeffe and O. M. Yaghi, The Chemistry and Applications of Metal-Organic Frameworks, *Science*, 2013, **341**, 1230444.
- 3 A. J. Howarth, A. W. Peters, N. A. Vermeulen, T. C. Wang, J. T. Hupp and O. K. Farha, Best Practices for the Synthesis, Activation, and Characterization of Metal-Organic Frameworks, *Chem. Mat.*, 2017, **29**, 26–39.
- 4 R. G. Pearson, Hard and Soft Acids and Bases, *J. Am. Chem. Soc.*, 1963, **85**, 3533–3539.
- 5 A. Dhakshinamoorthy, Z. Li and H. Garcia, Catalysis and photocatalysis by metal organic frameworks, *Chem. Soc. Rev.*, 2018, **47**, 8134–8172.
- 6 M. Dan-Hardi, C. Serre, T. Frot, L. Rozes, G. Maurin, C. Sanchez and G. Férey, A New Photoactive Crystalline Highly Porous Titanium(IV) Dicarboxylate, *J. Am. Chem. Soc.*, 2009, **131**, 10857–10859.
- 7 A. Cadiau, N. Kolobov, S. Srinivasan, M. G. Goesten, H. Haspel, A. V. Bavykina, M. R. Tchalala, P. Maity, A. Goryachev, A. S. Poryvaev, M. Eddaoudi, M. V. Fedin, O. F. Mohammed and J. Gascon, A Titanium Metal–Organic Framework with Visible-Light-Responsive Photocatalytic Activity, *Angew. Chem., Int. Ed.*, 2020, **59**, 13468–13472.
- 8 M. Sachs, L. Harnett-Caulfield, E. Pastor, B. Davies, D. J. C. Sowood, B. Moss, A. Kafizas, J. Nelson, A. Walsh and J. R. Durrant, Ligand field states control photocatalytic efficiency of transition metal oxides, in preparation, *ChemRxiv*, 2024, DOI: [10.26434/chemrxiv-2024-2wn0d](https://doi.org/10.26434/chemrxiv-2024-2wn0d).
- 9 B. Bueken, F. Vermoortele, D. E. P. Vanpoucke, H. Reinsch, C. C. Tsou, P. Valvekens, T. D. Baerdemaeker, R. Ameloot, C. E. A. Kirschhock, V. Van Speybroeck, J. M. Mayer and D. De Vos, A Flexible Photoactive Titanium Metal–Organic Framework Based on a [Ti(IV)₃(μ₃-O)(O)₂(COO)₆] Cluster, *Angew. Chem., Int. Ed.*, 2015, **54**, 13912–13917.
- 10 T. Devic and C. Serre, High valence 3p and transition metal based MOFs, *Chem. Soc. Rev.*, 2014, **43**, 6097–6115.
- 11 L. Rozes and C. Sanchez, Titanium oxo-clusters: precursors for a Lego-like construction of nanostructured hybrid materials, *Chem. Soc. Rev.*, 2011, **40**, 1006–1030.
- 12 H. L. Nguyen, Perspectives on titanium-based metal–organic frameworks, *JPhys Energy*, 2021, **3**, 021003.



- 13 L. Li, X.-S. Wang, T. F. Liu and J. Ye, Titanium-Based MOF Materials: From Crystal Engineering to Photocatalysis, *Small Methods*, 2020, **4**, 2000486.
- 14 C. Serre and G. Férey, Hybrid Open Frameworks. 8. Hydrothermal Synthesis, Crystal Structure, and Thermal Behavior of the First Three-Dimensional Titanium(IV) Diphosphonate with an Open Structure: $\text{Ti}_3\text{O}_2(\text{H}_2\text{O})_2(\text{O}_3\text{P}(\text{CH}_2)\text{PO}_3)_2 \cdot (\text{H}_2\text{O})_2$, or MIL-22, *Inorg. Chem.*, 1999, **38**, 5370–5373.
- 15 J. Castells-Gil, N. M. Padial, N. Almora-Barrios, J. Albero, A. R. Ruiz-Salvador, J. González-Platas, H. García and C. Martí-Gastaldo, Chemical Engineering of Photoactivity in Heterometallic Titanium–Organic Frameworks by Metal Doping, *Angew. Chem., Int. Ed.*, 2018, **57**, 8453–8457.
- 16 N. M. Padial, J. Castells-Gil, N. Almora-Barrios, M. Romero-Angel, I. D. Silva, M. Barawi, A. García-Sánchez, V. A. De La Pena O'Shea and C. Martí-Gastaldo, Hydroxamate Titanium–Organic Frameworks and the Effect of Siderophore-Type Linkers over Their Photocatalytic Activity, *J. Am. Chem. Soc.*, 2019, **141**, 13124–13133.
- 17 H.-Z. Li, S. Li, F. Wang and J. Zhang, A chelating coordination modulation method for the synthesis of Ti-MOF single crystals, *Inorg. Chem. Front.*, 2024, **11**, 2876–2883.
- 18 S. Horike, S. Shimomura and S. Kitagawa, Soft porous crystals, *Nat. Chem.*, 2009, **1**, 695–704.
- 19 S. Kitagawa and M. Kondo, Functional micropore chemistry of crystalline metal complex-assembled compounds, *Bull. Chem. Soc. Jpn.*, 1998, **71**, 1739–1753.
- 20 T. Loiseau, C. Serre, C. Huguenard, G. Fink, F. Taulelle, M. Henry, T. Bataille and G. Férey, A Rationale for the Large Breathing of the Porous Aluminum Terephthalate (MIL-53) Upon Hydration, *Chem. –Eur. J.*, 2004, **10**, 1373–1382.
- 21 C. Mellot-Draznieks, C. Serre, S. Surblé, N. Audebrand and G. Férey, Very Large Swelling in Hybrid Frameworks: A Combined Computational and Powder Diffraction Study, *J. Am. Chem. Soc.*, 2005, **127**, 16273–16278.
- 22 C. Serre, C. Mellot-Draznieks, S. Surblé, N. Audebrand, Y. Filinchuk and G. Férey, Role of Solvent-Host Interactions That Lead to Very Large Swelling of Hybrid Frameworks, *Science*, 2007, **315**, 1828–1831.
- 23 N. Klein, H. C. Hoffmann, A. Cadiou, J. Getzschmann, M. R. Lohe, S. Paasch, T. Heydenreich, K. Adil, I. Senkovska, E. Brunner and S. Kaskel, Structural flexibility and intrinsic dynamics in the $\text{M}_2(2,6\text{-ndc})_2(\text{dabco})$ ($\text{M} = \text{Ni}, \text{Cu}, \text{Co}, \text{Zn}$) metal–organic frameworks, *J. Mater. Chem.*, 2012, **22**, 10303–10312.
- 24 J. H. Lee, S. Jeoung, Y. G. Chung and H. R. Moon, Elucidation of flexible metal-organic frameworks: Research progresses and recent developments, *Coord. Chem. Rev.*, 2019, **389**, 161–188.
- 25 A. Schneemann, V. Bon, I. Schwedler, I. Senkovska, S. Kaskel and R. A. Fischer, Flexible metal–organic frameworks, *Chem. Soc. Rev.*, 2014, **43**, 6062–6096.
- 26 T. D. Bennett, A. K. Cheetham, A. H. Fuchs and F.-X. Coudert, Interplay between defects, disorder and flexibility in metal-organic frameworks, *Nat. Chem.*, 2017, **9**, 11–16.
- 27 M. Alhamami, H. Doan and C.-H. Cheng, A Review on Breathing Behaviors of Metal-Organic-Frameworks (MOFs) for Gas Adsorption, *Materials*, 2014, **7**, 3198–3250.
- 28 K. Barthelet, J. Marrot, D. Riou and G. Férey, A Breathing Hybrid Organic–Inorganic Solid with Very Large Pores and High Magnetic Characteristics, *Angew. Chem., Int. Ed.*, 2002, **41**, 281–284.
- 29 J. Cao, W. Ma, K. Lyu, L. Zhuang, H. Cong and H. Deng, Twist and sliding dynamics between interpenetrated frames in Ti-MOF revealing high proton conductivity, *Chem. Sci.*, 2020, **11**, 3978–3985.
- 30 C. Chinchilla-Garzón, M. Galbiati, A. Misturini, P. Gimeno-Fonquernie, N. Almora-Barrios, N. M. Padial and C. Martí-Gastaldo, Structural Control of Photoconductivity in a Flexible Titanium-Organic Framework, *Adv. Mater.*, 2025, 2412045, DOI: [10.1002/adma.202412045](https://doi.org/10.1002/adma.202412045).
- 31 Q. Yan, J. Wang, L. Zhang, J. Liu, M. Wahiduzzaman, N. Yan, L. Yu, R. Dupuis, H. Wang, G. Maurin, M. Hirscher, P. Guo, S. Wang and J. Du, A squarate-pillared titanium oxide quantum sieve towards practical hydrogen isotope separation, *Nat. Commun.*, 2023, **14**, 4189.
- 32 J. Gao, J. Miao, P.-Z. Li, W. Y. Teng, L. Yang, Y. Zhao, B. Liu and Q. Zhang, A p-type Ti(IV)-based metal–organic framework with visible-light photo-response, *Chem. Commun.*, 2014, **50**, 3786–3788.
- 33 H. L. Nguyen, The chemistry of titanium-based metal–organic frameworks, *New J. Chem.*, 2017, **41**, 14030–14043.
- 34 Y. Cai, L. Zou, Q. Ji, J. Yong, X. Qian and J. Gao, Two dimensional Ti-based metal-organic framework with polar oxygen atoms on the pore surface for efficient gas separation, *Polyhedron*, 2020, **190**, 114771.
- 35 S. Yu, Y. Xiao, Z. Liu, J.-M. Lyu, Y.-L. Wang, Z.-Y. Hu, Y. Li, M. Sun, L.-H. Chen and B.-L. Su, Ti-MOF single-crystals featuring an intracrystal macro–microporous hierarchy for catalytic oxidative desulfurization, *Chem. Commun.*, 2023, **59**, 1801–1804.
- 36 N. Crespi Sanchez, G. Turnes Palomino and C. Palomino Cabello, TiO_2 derived from NTU-9 metal organic framework as highly efficient photocatalyst, *Mater. Sci. Eng., B*, 2021, **273**, 115424.
- 37 D. Yan, J. Jia, K. Tong, J. Ouyang and Y. Pan, Metal organic framework as a matrix for laser desorption/ionisation of environmental pollutants and other small molecules, *Int. J. Environ. Anal. Chem.*, 2023, **104**, 9153–9163.
- 38 R. S. Forgan, Modulated self-assembly of metal–organic frameworks, *Chem. Sci.*, 2020, **11**, 4546–4562.
- 39 H. L. B. Boström, S. Emmerling, F. Heck, C. Koschnick, A. J. Jones, M. J. Cliffe, R. A. Natour, M. Bonneau, V. Guillermin, O. Shekhah, M. Eddaoudi, J. Lopez-Cabrelles, S. Furukawa, M. Romero-Angel, C. Martí-Gastaldo, M. Yan, A. J. Morris, I. Romero-Muñiz, Y. Xiong, A. E. Platero-Prats, J. Roth, W. L. Queen, K. S. Mertin, D. E. Schier, N. R. Champness, H. H. M. Yeung and B. V. Lotsch, How reproducible is the synthesis of Zr–porphyrin metal–



- organic frameworks? An interlaboratory study, *Adv. Mater.*, 2023, **36**, 2304832.
- 40 C. Koschnick, M. W. Terban, S. Canossa, M. Etter, R. E. Dinnebier and B. V. Lotsch, Influence of Water Content on Speciation and Phase Formation in Zr-Porphyrin-Based MOFs, *Adv. Mater.*, 2024, **36**, 2210613.
- 41 H. Chun and D. Moon, Metal–Organic Frameworks from Group 4 Metals and 2,5-Dihydroxyterephthalic Acid: Reinvestigation, New Structure, and Challenges Toward Gas Storage and Separation, *Cryst. Growth Des.*, 2017, **17**, 2140–2146.
- 42 E. Dhivya, D. Magadevan, Y. Palguna, T. Mishra and N. Aman, Synthesis of titanium based hetero MOF photocatalyst for reduction of Cr (VI) from wastewater, *J. Environ. Chem. Eng.*, 2019, **7**, 103240.
- 43 Y. Yu, S. Li, L. Huang, J. Yu, H. Zhang, S. Song and T. Zeng, Solar-driven CO₂ conversion promoted by MOF-on-MOF homophase junction, *Catal. Commun.*, 2021, **150**, 106270.
- 44 H. Assi, L. C. Padro Pérez, G. Mouchaham, F. Ragon, M. Nasalevich, N. Guillou, C. Martineau, H. Chevreau, F. Kapteijn, J. Gascon, P. Fertey, E. Elkaim, C. Serre and T. Devic, Investigating the Case of Titanium(IV) Carboxyphenolate Photoactive Coordination Polymers, *Inorg. Chem.*, 2016, **55**, 7192–7199.
- 45 H. M. Rietveld, A profile refinement method for nuclear and magnetic structures, *J. Appl. Crystallogr.*, 1969, **2**, 65–71.
- 46 J. E. Mondloch, O. Karagiari, O. K. Farha and J. T. Hupp, Activation of metal–organic framework materials, *CrystEngComm*, 2013, **15**, 9258–9264.
- 47 Y. Li, Z.-M. Hao, M.-Y. Chao, W.-H. Zhang and D. J. Young, Vacuum-Induced Guest *N,N'*-Diethylformamide Binding in a Metastable Cd₅-Based Metal–Organic Framework, *Inorg. Chem.*, 2022, **61**, 20227–20231.
- 48 C. L. Hobday, S. Krause, S. M. J. Rogge, J. D. Evans and H. Bunzen, Perspectives on the Influence of Crystal Size and Morphology on the Properties of Porous Framework Materials, *Front. Chem.*, 2021, **9**, 772059.
- 49 D.-F. Lu, S. Li, M.-X. Guan, Y.-P. Han, Y. Sun, X. Wu, F. Wang and J. Zhang, Visible-Light-Active Titanium Sulfonate Framework for Photocatalytic Organic Synthesis, *ACS Mater. Lett.*, 2023, **5**, 2836–2842.
- 50 D.-F. Lu, Y.-P. Han, Y. Sun, F. Wang and J. Zhang, Titanium Sulfonate-Based Metal–Organic Frameworks, *Cryst. Growth Des.*, 2023, **23**, 3778–3784.
- 51 R. Hassandoost, A. Khataee and E. Doustkhah, Photocatalytic performance of disordered titanium-based hollow nanosheet metal–organic frameworks in wastewater treatment, *J. Taiwan Inst. Chem. Eng.*, 2024, **157**, 105409.
- 52 T. Schertenleib, M. Asgari, B. Mourino, V. V. Karve, T. Felder, D. Stoian, V. Bon, J. Hao, A. Ortega-Guerrero, E. Oveisi, K. V. Agrawal, B. Smit, S. Kaskel, S. J. L. Billinge and W. L. Queen, Anisotropic Node Distortions in Amorphous MOFs: Low Valent Zr Sites as Catalytic Hotspots, preprint, *ChemRxiv*, 2024, DOI: [10.26434/chemrxiv-2024-56r8t](https://doi.org/10.26434/chemrxiv-2024-56r8t).
- 53 A. A. Coelho, TOPAS and TOPAS-Academic: an optimization program integrating computer algebra and crystallographic objects written in C++, *J. Appl. Crystallogr.*, 2018, **51**, 210–218.
- 54 R. W. Cheary and A. Coelho, A fundamental parameters approach to X-ray line-profile fitting, *J. Appl. Crystallogr.*, 1992, **25**, 109–121.
- 55 A. L. Bail, H. Duroy and J. L. Fourquet, *Ab initio* structure determination of LiSbWO₆ by X-ray powder diffraction, *Mater. Res. Bull.*, 1988, **23**, 447–452.
- 56 A. A. Coelho, Whole-profile structure solution from powder diffraction data using simulated annealing, *J. Appl. Crystallogr.*, 2000, **33**, 899–908.
- 57 L. Zhang and Y. H. Hu, Structure distortion of Zn₄O₁₃C₂₄H₁₂ framework (MOF-5), *Mater. Sci. Eng., B*, 2011, **176**, 573–578.
- 58 J. W. M. Osterrieth, J. Rampersad, D. Madden, N. Rampal, L. Skoric, B. Connolly, M. D. Allendorf, V. Stavila, J. L. Snider, R. Ameloot, J. Marreiros, C. Ania, D. Azevedo, E. Vilarrasa-Garcia, B. F. Santos, X.-H. Bu, Z. Chang, H. Bunzen, N. R. Champness, S. L. Griffin, B. Chen, R.-B. Lin, B. Coasne, S. Cohen, J. C. Moreton, Y. J. Colón, L. Chen, R. Clowes, F.-X. Coudert, Y. Cui, B. Hou, D. M. D'Alessandro, P. W. Doheny, M. Dincă, C. Sun, C. Doonan, M. T. Huxley, J. D. Evans, P. Falcaro, R. Ricco, O. Farha, K. B. Idrees, T. Islamoglu, P. Feng, H. Yang, R. S. Forgan, D. Bara, S. Furukawa, E. Sanchez, J. Gascon, S. Telalović, S. K. Ghosh, S. Mukherjee, M. R. Hill, M. M. Sadiq, P. Horcajada, P. Salcedo-Abraira, K. Kaneko, R. Kukobat, J. Kevin, S. Keskin, S. Kitagawa, K.-i. Otake, R. P. Lively, S. J. A. DeWitt, P. Llewellyn, B. V. Lotsch, S. T. Emmerling, A. M. Pütz, C. Marti-Gastaldo, N. M. Padial, J. García-Martínez, N. Linares, D. Maspoch, J. A. S. d. Pino, P. Moghadam, R. Oktavian, R. E. Morris, P. S. Wheatley, J. Navarro, C. Petit, D. Danaci, M. J. Rosseinsky, A. P. Katsoulidis, M. Schröder, X. Han, S. Yang, C. Serre, G. Mouchaham, D. S. Sholl, R. Thyagarajan, D. Siderius, R. Q. Snurr, R. B. Goncalves, S. Telfer, S. J. Lee, V. P. Ting, J. L. Rowlandson, T. Uemura, T. Iiyuka, M. A. van derVeen, D. Rega, V. Van Speybroeck, S. M. J. Rogge, A. Lemaire, K. S. Walton, L. W. Bingel, S. Wuttke, J. Andreato, O. Yaghi, B. Zhang, C. T. Yavuz, T. S. Nguyen, F. Zamora, C. Montoro, H. Zhou, A. Kirchon and D. Fairen-Jimenez, How Reproducible are Surface Areas Calculated from the BET Equation?, *Adv. Mater.*, 2022, **34**, 2201502.
- 59 M. Brunelli and A. N. Fitch, A glass capillary cell for *in situ* powder X-ray diffraction of condensed volatile compounds. Solid HCFC-123a and HCFC-124, *J. Synchrotron Radiat.*, 2003, **10**, 337–339.

