


 Cite this: *Phys. Chem. Chem. Phys.*,
 2024, 26, 18276

Exploring the effect of molecular size and framework functionalisation on transport in metal–organic frameworks using pulsed-field gradient nuclear magnetic resonance†

 Shima Zainal,^a Ahmed Alsudani,^b Ralph W. Adams,^b Mathias Nilsson,^b
 Xiaolei Fan^{*ac} and Carmine D'Agostino^{id *ad}

Molecular transport is an important aspect in metal–organic frameworks (MOFs) as it affects many of their applications, such as adsorption/separation, drug delivery and catalysis. Yet probing the fundamental diffusion mechanisms in MOFs is challenging, and the interplay between the MOF's features (such as the pore structure and linker dynamics) and molecular transport remains mostly unexplored. Here, the pulsed-field gradient nuclear magnetic resonance (PFG NMR) technique is used to probe the diffusion of several probe molecules, *i.e.*, water, xylenes and 1,3,5-triisopropylbenzene (TIPB), within the UiO-66 MOF and its derivatives (UiO-66NH₂ and UiO-66Br). Exploiting differences in the size of probe molecules we were able to probe the diffusion rate selectively in the different pore environments of the MOFs. In particular, when relatively small molecules, such as water and small hydrocarbons, were used as probes, the PFG NMR log attenuation plots were non-linear with two distinctive diffusion regions, suggesting faster diffusion in the inter-crystalline space and slower diffusion within crystal aggregates, the latter occurring mostly inside the framework of the MOFs. Conversely, experiments with a larger probe molecule, *i.e.*, TIPB, with a kinetic diameter of 0.95 nm, which makes it unable to access the framework windows of the MOF crystals, showed linear PFG NMR log attenuation plots, which indicates diffusion occurring in a single environment, most likely in the inter-crystalline space. Analysis of the apparent tortuosity values of the systems under investigation highlights the role of linker functionalisation in influencing the molecular diffusion of the probe molecules, which affects both intra-molecular interactions and pore accessibility within the MOF crystals. The findings of this work demonstrate that the diffusion behaviour of probe molecules within MOFs is influenced by the pore size, structure, functionalisation of the MOF linker and molecular interactions. Our study contributes to further advance the understanding of mass transport in MOFs by PFG NMR and provides insights that can inform the design and optimisation of MOF-based materials for various applications.

 Received 30th January 2024,
 Accepted 31st May 2024

DOI: 10.1039/d4cp00447g

rsc.li/pccp

1. Introduction

Metal–organic frameworks (MOFs) are porous, crystalline solids made by combining organic linkers and metal-oxide clusters.

In recent years, MOFs have gained prominence as versatile materials with applications ranging from gas storage to catalysis. Among these, UiO-66 (*i.e.*, University of Oslo 66)¹ and its derivatives have attracted significant attention due to their interesting properties and potential use.^{2,3} UiO-66 typically incorporates zirconium metal nodes into its structure forming strong bonds with organic linkers of terephthalic acid. These organic linkers create linkages between the metal nodes forming a crystalline three-dimensional framework. UiO-66 derivatives such as UiO-66NH₂ and UiO-66Br are the linker modified versions of the original UiO-66, where specific functional groups are introduced to the structure (by functionalisation of the organic linker). These modifications can alter the properties of the UiO-66 MOFs, and thus their applications. In UiO-66NH₂, the amino (–NH₂) group is grafted to the organic linker in UiO-66, serving as additional binding sites for specific molecules,

^a Department of Chemical Engineering, The University of Manchester, Oxford Road, Manchester, M13 9PL, UK. E-mail: carmine.dagostino@manchester.ac.uk, xiaolei.fan@manchester.ac.uk

^b Department of Chemistry, The University of Manchester, Oxford Road, Manchester, M13 9PL, UK

^c Ningbo China Beacons of Excellence Research and Innovation Institute, University of Nottingham Ningbo China, 211 Xingguang Road, Ningbo 315048, China

^d Dipartimento di Ingegneria Civile, Chimica, Ambientale e dei Materiali (DICAM), Alma Mater Studiorum – Università di Bologna, Via Terracini, 28, 40131 Bologna, Italy

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4cp00447g>



for example, adsorption of sulphur species and other contaminants in wastewater treatment applications.⁴ Incorporation of bromine (Br) atoms in UiO-66Br, altering its structure, can impact its structural properties significantly, thereby influencing chemical reactivity, adsorption capacity, and interactions with guest molecules. For instance, Br-modified MOFs were shown to be capable of removing elemental mercury (Hg⁰) from flue gas.⁵ Additionally, the introduction of -Br functional groups can induce steric hindrance, consequently modifying the MOF's electronic properties.^{6,7}

Although the applications of UiO-66 and its derivatives are well documented,^{8–10} insights into the microscopic mass transport of guest molecules within these materials remain mostly unexplored, and knowledge of mass transfer within the UiO-66 family is crucial for optimising their properties and applications as it enables the design of more efficient and tailored materials. Molecular diffusion in MOFs is influenced by a number of parameters, including the structure of the MOFs (such as pore size), functional groups on the linker, presence of defects and chemical interaction between guest molecules.¹¹ Jacob *et al.* simulated acetone diffusion within pristine UiO-66 using molecular dynamics (MD) simulation, aiming to explore the effect of framework flexibility and hydrogen bonding (between acetone molecules and the hydroxyl groups present on the nodes of the UiO-66 framework) on the diffusion behaviour of acetone within UiO-66.¹² The results demonstrate the critical significance of linker dynamics and provide insight into how hydrogen bonding affects acetone diffusivity under different loading conditions. Ramsahye *et al.* combined quasi-elastic neutron scattering (QENS) with MD simulation to investigate the diffusion of light hydrocarbons in UiO-66.¹³ These findings show a clear decrease in diffusion with an increase in the molecular size of the hydrocarbons, indicating the enhanced confinement effect within the UiO-66 framework due to the increase in the kinetic diameter of the guest molecules. Although simulation methods offer molecular insights, validation of the simulated results is challenging.

In addition to MD simulations, several experimental techniques can be used to measure diffusion in MOFs, such as gravimetric techniques and spectroscopy methods.¹¹ Gravimetric techniques measure the mass changes, which are then correlated with the adsorption/desorption of guest molecules on/from MOFs, and hence diffusion rates.¹⁴ Although the bulk information obtained by gravimetric methods are useful for exploring diffusion behaviour in MOFs, they are slow and sensitive to variations in temperature and humidity. Spectroscopic methods, such as infrared spectroscopy (IR) and NMR, provide molecular-level information about the interaction between MOF components and diffusing species.^{15–17} These techniques offer insight into molecular interactions, diffusion pathways and guest–host interactions within MOFs.

Pulsed-field gradient nuclear magnetic resonance (PFG NMR) is a non-invasive and non-destructive technique which can probe the movement of molecules within porous materials. In addition, it allows quantitative measurement of diffusion coefficients,^{15,18} which is essential for understanding and comparing the transport phenomena within different porous materials. This method can

also differentiate diffusion in different environments simultaneously, providing unique insights into the dynamics of molecular transport within porous materials.¹⁹ PFG NMR has been applied to various porous materials, such as diffusion of mixtures in microporous and mesoporous materials, including zeolites and heterogeneous catalysts. For example, PFG NMR diffusion studies were performed to investigate intra-crystalline diffusion in USY zeolites, and the findings challenge existing assumptions about the role of intra-crystalline mesopores in USY zeolites.²⁰ Experimental evidence suggests that these mesopores do not significantly contribute to intra-crystalline diffusion and do not form a connected network within the crystals. PFG NMR was also applied to mesoporous SBA-15 and KIT-6 silicas to unravel mass transport and the effect of mesopore size on molecular diffusion and catalytic performance.^{21,22} The findings provide insights into how the morphology of SBA-15 silica (such as isolated fibres and bundles) influences molecular displacements and diffusion behaviour. In particular, the results show that the measured intra-particle NMR diffusivity agrees with the charge carrier diffusivity determined using the Nernst–Einstein equation.

In the context of MOFs, previous studies employing PFG NMR to study molecular self-diffusion within ZIFs and MOF-3 helped to establish a correlation between the molecular size and pore characteristics of the MOFs.²³ These results demonstrate that light hydrocarbons can readily exchange between the intra-crystalline phase in a MOF and the surrounding environment. The findings above shed light on the diffusion dynamics of hydrocarbons within the MOF structure, which is essential for applications such as gas storage, separation, and catalysis. However, unlike zeolites and heterogeneous solid catalysts, the number of diffusion studies in MOFs using PFG NMR remains limited.

Herein, we present the results of a direct investigation into the molecular transport in UiO-66 and its linker functionalised derivatives by measuring the diffusion behaviour of relevant probe molecules using the PFG NMR technique. To characterise the type of diffusion, the experimental measurements were performed with probe molecules of different sizes focusing on investigating the effect of steric hindrance/intra-framework interaction over the length scale probed by PFG NMR.

2. Experimental

2.1 Synthesis of materials

All chemicals were of analytical grade and used without further purification. Anhydrous zirconium(IV) chloride, terephthalic acid (BDC), 2-amino terephthalic acid (BDC-NH₂), bromoterephthalic acid (BDC-Br), dimethylformamide (DMF), hydrochloric acid (HCl), *para* (*p*-), *meta* (*m*-), *ortho* (*o*-)xylenes, methanol and 1,3,5-triisopropylbenzene (TIPB) were purchased from Sigma Aldrich and used without further purification. Deionised water was purified to a resistivity of 18.2 MΩ cm in a Millipore Milli-Direct Q water purification system.

The UiO-66 MOFs were synthesized using the solvothermal method² by reacting BDC (0.09 g, 0.5 mmol) with zirconium(IV)



chloride (0.12 g, 0.5 mmol) in a ratio of 1 : 1 in 25 ml of DMF. The solution was stirred for two hours, then transferred to a stainless-steel autoclave, sealed and placed in a preheated oven at 120 °C for 24 hours. After the synthesis, the reactor was cooled down, and the resulting white solid precipitate was washed with DMF and solvent exchanged with methanol before being dried under vacuum at 120 °C overnight. The synthesis procedure was then repeated for making UiO-66NH₂ and UiO-66Br by changing the BDC linker to BDC-NH₂ and BDC-Br, respectively.

2.2 Material characterisation

Prior to characterisation all MOF materials were activated at 150 °C under vacuum. The powder X-ray diffraction (PXRD) patterns were acquired on a PANanalytical Empyrean XRD system with Cu-K α 1 radiation ($\lambda = 1.54056 \text{ \AA}$) in the 2θ range of 3° to 50° with a step size of 0.026° per second. The CryAlisPro software program was used for the integration of diffraction data.

Nitrogen (N₂) adsorption–desorption measurements were carried out using a Micromeritics ASAP 2020 analyser at 77 K. Before N₂ physisorption, approximately 100 mg of the solid sample was degassed at 180 °C under vacuum for 12 h. The specific surface areas of the materials were determined using the Brunauer–Emmett–Teller (BET) method, and the pore size distribution was obtained by using the Barrett–Joyner–Halenda (BJH) method (based on the adsorption branch of the isotherms). Infrared spectroscopy was performed at 25 °C using an FTIR on a PerkinElmer Spectrum 2000 Spectrometer. Scanning electron microscopy (SEM) of the samples was conducted using a Tescan Mira small chamber instrument with a LaB6 emitter (operated at 10 kV).

2.3 PFG NMR measurements

PFG NMR experiments were performed using a Bruker Avance NEO 500 NMR spectrometer operating at a ¹H frequency of 500 MHz with a diffusion probe (DiffBB). UiO-66, UiO-66NH₂ and UiO-66Br particles were first dried under vacuum then soaked for 48 h in 2 mL of the probe molecules (*i.e.*, water, *p*-xylene, *m*-xylene, *o*-xylene, *n*-octane and TIPB). The samples were then dried gently with filter paper to remove excess solvent and placed in a 5 mm NMR tube filled to a height of approximately 10 mm. Samples were placed in the spectrometer probe and left to equilibrate for 15 minutes before experiments. All experiments were performed at room temperature (approximately 298 K) using a 15 μs ¹H 90° pulse, a 1.4 ms diffusion encoding time using half-sine gradient pulse shape with 16 to 32 amplitude increments of the gradient pulses between 2 and 15% of 17 T m⁻¹, with a 100 ms diffusion time (Δ) and gradient recovery delays, τ of 0.2 ms. 64 scans per increment were acquired, with an inter-scan recycle delay of 5 s used for all experiments with each experiment taking between 45–90 minutes. The relaxation times T_1 and T_2 of the probe molecules in all three MOFs were measured to aid the choice of the acquisition parameters.

NMR data was collected using a bipolar pulse pair STE pulse sequence and they were subsequently fitted to the Stejskal–Tanner model²⁴ shown in eqn (1):

$$\frac{E}{E_0} = \sum_{i=1}^n p_i \exp \left[-D_i \cdot \gamma_H^2 \cdot g^2 \cdot \delta^2 \left(\Delta - \frac{5\delta}{16} - \frac{\tau}{2} \right) \right] \\ = \sum_{i=1}^n p_i \exp(-bD_i) \quad (1)$$

where E is the signal in the presence of gradient, E_0 is the NMR signal in the absence of gradient, γ_H is the gyromagnetic ratio of the nuclei being study ($\gamma_H = 42.577 \text{ MHz T}^{-1}$), g is the strength of the gradient pulse of duration δ , τ is the gradient recovery delay and Δ is the diffusing time (*i.e.*, the time interval between the leading edges of the gradient pulses). The $\gamma_H^2 \cdot g^2 \cdot \delta^2 \left(\Delta - \frac{5\delta}{16} - \frac{\tau}{2} \right)$ term is often referred to as the “ b -factor”. In eqn (1), the number of different diffusion components in the system under study is represented by the value of n . The PFG NMR signal attenuation was analysed using the general NMR analysis toolbox (GNAT) software (Matlab, version R2021b).²⁵

3. Results and discussion

3.1 Characterisation of UiO-66, UiO-66NH₂ and UiO-66Br

The characterisation of UiO-66, UiO-66NH₂ and UiO-66Br in this study builds upon previous investigations.^{2,3,7} The resulting MOFs exhibit different colours depending on the ligand used in the synthesis. UiO-66NH₂ is yellow, UiO-66 and UiO-66Br are white, consistent with the results reported elsewhere.²⁶ The bulk crystalline features of the MOFs were evaluated by PXRD to gain information on the crystallographic planes and structural similarity of these MOFs. As shown in Fig. 1, the diffraction peaks of UiO-66NH₂ and UiO-66Br were consistent with that of the UiO-66 MOF, suggesting that all the MOFs under investigation are topologically identical. Sharp diffraction peaks at $2\theta = 7.42$, 8.6 and 25.7°, corresponding to the 111, 002 and 006 planes, are identified with comparable intensities in the XRD patterns of all three samples, suggesting that the linker functionalisation did not interfere with the crystallisation of the UiO-66 MOFs during the solvothermal synthesis.³

In addition, the PXRD patterns of the three MOFs were compared with the simulated patterns (based on the CIF file from the Cambridge Crystallography Open Database²⁷). The comparison shows a very close resemblance between the XRD patterns of the synthesised MOFs and the simulated ones. Following these results, the UiO-66 structure is shown in Fig. 2. The structure consists of UiO-66 tetrahedral and octahedral cages, with sizes of 0.8 nm and 1.4 nm, respectively.²⁸

N₂ adsorption–desorption isotherms of the MOFs are shown in Fig. S1 in the ESI,† showing that UiO-66 and UiO-66Br are microporous, whilst the isotherm of UiO-66NH₂ shows the presence of a hysteresis loop. The presence of the hysteresis loop in the range of 0.9–1.0 p/p_0 indicates that the sample may



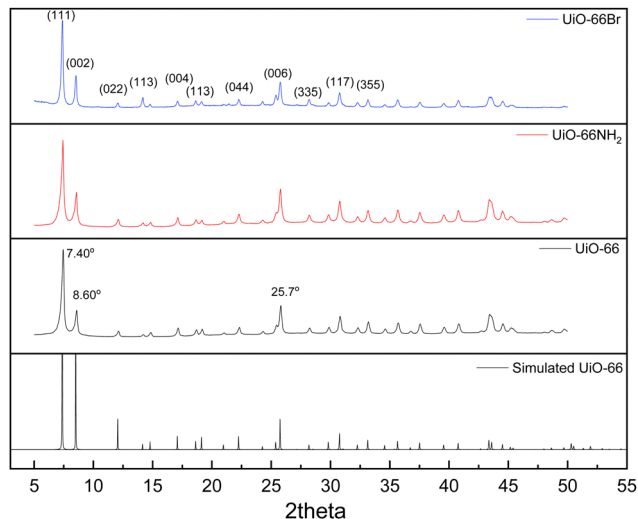


Fig. 1 PXRD patterns of the synthesised UiO-66 and its derivatives.

contain mesoscopic structural defects and/or inter-crystalline mesopores. Regarding the calculated specific surface area and pore volume, UiO-66 has the highest value of $953 \text{ m}^2 \text{ g}^{-1}$ and $0.45 \text{ cm}^3 \text{ g}^{-1}$, respectively (Table S1, ESI[†]). The functionalised UiO-66 maintains the same crystalline structure and topology of UiO-66 but with lower specific surface areas, as shown in Table S1 (ESI[†]), proving that the introduction of larger and heavier amine groups and bromine atoms to the linker decreased the surface area and the total pore volume for UiO-66NH₂ and UiO-66Br. The calculated BET surface areas for all three samples in this study are consistent with those reported in the literature.²

The presence of –NH₂ and –Br within the UiO-66 derivatives was confirmed by FTIR analysis (Fig. S2 and Table S2, ESI[†]). For UiO-66NH₂ (red solid line in Fig. S2, ESI[†]), the two small peaks at 3507 cm^{-1} and 3384 cm^{-1} reflect the characteristic vibrations of the –NH₂ group. Another two characteristic bands of the amino group can be identified at 1600 cm^{-1} and 1210 cm^{-1} , respectively, for the N–H bending vibration and the C–N stretching of aromatic amines.³⁰ Regarding UiO-66Br, its FTIR spectrum exhibits a specific peak at 680 cm^{-1} , consistent with known vibrational modes associated with the presence of

bromine functionalities. The appearance of these peaks support the effective introduction of bromine moieties onto the UiO-66 framework.³⁰ The morphology of the as-synthesised MOFs was characterized by SEM (Fig. S3, ESI[†]), showing the typical polyhedral crystals for UiO-66 MOFs,³¹ and the particle sizes of the MOFs are comparable at $< 500 \text{ nm}$ and $< 2 \mu\text{m}$. The crystals clearly display irregularities and tend to form large aggregates of the order of several microns.

3.2 PFG NMR results

For the systems under investigation, diffusion may potentially occur inside the crystals (and within crystal aggregates), between the crystals (inter-crystalline) or as a combination of both.³² Diffusion within crystal aggregates, and in general intra-particle diffusion, may further be affected by the pore size distribution (*i.e.*, presence of distinctive pore sizes within the same particle), which can lead to the observation of more than one diffusion environment. This has been previously reported for diffusion in activated carbon catalysts.³³ Fig. 3 shows the PFG NMR attenuation plots for different probe molecules diffusing through the porous structures of UiO-66 (Fig. 3(a)), UiO-66NH₂ (Fig. 3(b)) and UiO-66Br (Fig. 3(c)). The data are plotted on a semi-log scale; hence a single exponential decay would appear on the graph as a straight line. In order to determine the effect of observation time, diffusion experiments were also carried out using observation times of 300 ms and 600 ms. The PFG attenuation plots and values of diffusivities (Fig. S5 and Table S6, ESI[†]) were in all cases very similar suggesting that the observation time does not significantly affect diffusion coefficients.

The plots for all MOFs when using small probe molecules, *i.e.*, water (kinetic diameter, $K_D = 0.26 \text{ nm}$),³⁴ *n*-octane ($K_D = 0.48 \text{ nm}$)¹⁷ and xylenes ($K_D = 0.58, 0.68$ and 0.68 nm for *p*-/*m*-/*o*-xylenes)³⁵ all show non-linear behaviour. In more detail, the experimental data can be fitted well using a two-component diffusion model. Conversely, when the bulky molecule of TIPB ($K_D = 0.95 \text{ nm}$)³⁶ was used, a linear and single-component diffusion behaviour was observed for the three MOFs under investigation. The results suggest that small molecules experience two distinct diffusion environments in slow-exchange with each other, whilst the larger TIPB molecule only experiences a

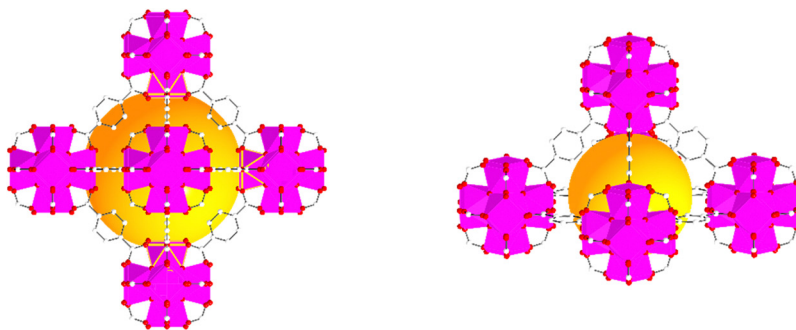


Fig. 2 Simulated UiO-66 represented by balls and sticks showing an enlarged view of its octahedral (left) and tetrahedral (right) cages. Zr atoms are depicted as octahedra (Zr in pink, C in white, O in red). For clarity, guest molecules and hydrogen atoms have been removed from the structural diagrams. The figure was created using atomic coordinates from the literature.²⁹



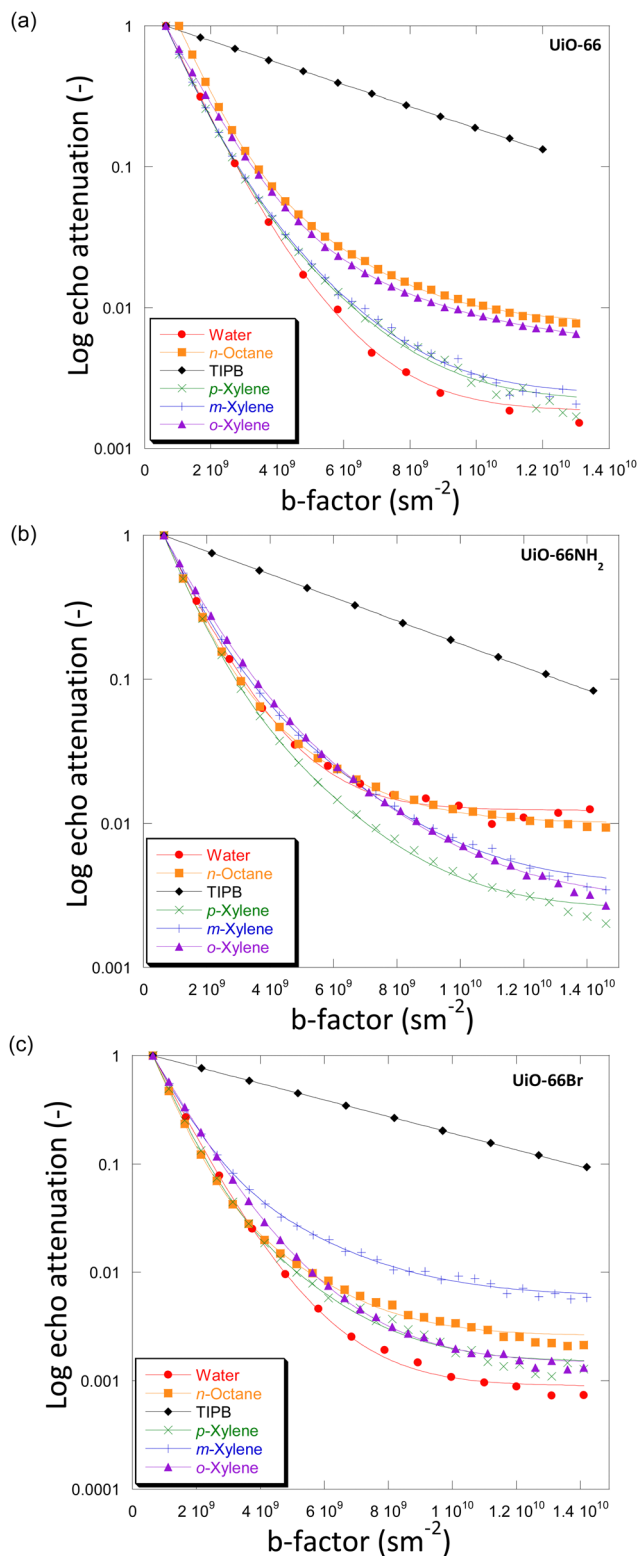


Fig. 3 Diffusion attenuation plots for different probe molecules within the porous network of (a) UiO-66, (b) UiO-66NH₂ and (c) UiO-66Br. The BBP-STE sequence was used in all experiments with a diffusion interval (Δ) of 100 ms. The solid lines are exponential fits using eqn (1).

single diffusion environment. Based on these observations, we attribute the slow-diffusion component of small probe

molecules to diffusion within crystal aggregates, whilst the fast-diffusion component can be attributed to inter-crystalline diffusion (diffusion between crystals).

To understand the difference in the diffusion behaviour of different probe molecules, it is necessary to consider their molecular sizes, as well as the typical pore dimension of the MOFs under investigation. The small probe molecules of water, *n*-octane and xylenes with K_d values of 0.26–0.68 nm, possess lower K_d values than both the size of tetrahedral and octahedral cages (0.8 nm and 1.4 nm, respectively). This allows them to access the microporous networks within the UiO-66 MOF and its derivatives. Consequently, these molecules are expected to show a slow diffusivity component within the micropore network of the MOFs. Conversely, TIPB, with $K_d = 0.95$ nm, is too large to access the inner space of the MOFs through the tetrahedral windows, hence only inter-crystalline diffusion could be probed, which is what we observed in this study.

Using the bulk self-diffusion coefficients, $D_{\text{self,bulk}}$, of the probe molecules (Fig. S4 and Tables S3–S5, ESI[†]), which are consistent with those reported in the literature,^{20,37} we can make a relative comparison between those bulk values of diffusivity and the diffusivity values measured inside the MOFs. Compared to the $D_{\text{self,bulk}}$ values of the probe molecules, the values of the fast-diffusion component within the MOFs, $D_{1,\text{self}}$, are of the same order of magnitude but nearly halved, suggesting that the first diffusion regime of the probe molecules in the systems is that of bulk molecules hindered by the presence of the walls of the MOF crystals. Such drops in diffusivity are usually observed in mesoporous/macroporous materials³⁸ and given the typical dimension of the MOF crystals studied in this work (approximately 100 nm), it is reasonable to assume that the inter-crystalline space will be formed of voids of similar dimension, hence explaining the values of $D_{1,\text{self}}$ observed. Conversely, the values of the slow-diffusion component, $D_{2,\text{self}}$, of the small probe molecules are markedly smaller than that of the corresponding $D_{\text{self,bulk}}$, indicating molecular diffusion possibly occurring inside the microporous frameworks of the MOF. This assignment is supported by previous findings on PFG NMR diffusion coefficients in microporous materials.^{33,39,40} These findings further support our assignment, whereby we can assign $D_{1,\text{self}}$ to inter-crystalline diffusion and $D_{2,\text{self}}$ to diffusion within crystal aggregates, with the diffusion process being a slow-exchange between the two different diffusion environments.

It is important to observe that in Tables S3–S5 (ESI[†]), values of root mean square displacement ($\text{RMSD} = \sqrt{2D\Delta}$, where D is the self-diffusion coefficient and Δ is the observation/diffusion time) calculated for both $D_{1,\text{self}}$ and $D_{2,\text{self}}$ are in the order of a few to tens of microns, which is much larger than the typical size of the MOF crystals. We note however that the RMSD probed for the $D_{2,\text{self}}$ component is similar to the size of the crystal aggregates, as shown by the SEM images in Fig. S3 (ESI[†]). This implies that such a component is representative of diffusion inside such crystal aggregates rather than purely intra-crystalline diffusion. As molecules diffuse, they will traverse through crystal pores in a random walk pattern.¹⁸



As shown in Tables S3–S5 (ESI[†]), the $D_{1,\text{self}}$ values of all the probe molecules within UiO-66Br are slightly higher than those measured in UiO-66 and UiO-66NH₂, whilst the $D_{2,\text{self}}$ values for all probe molecules in all three MOFs differ significantly. The slightly higher $D_{1,\text{self}}$ observed in UiO-66Br may be caused by the differences in particle size of MOF crystals due to linker functionalisation. Although the SEM data (Fig. S3, ESI[†]) indicate that the average crystal size yielded by all three MOFs is approximately 100 nm, there may be some variation in crystallinity that affects its diffusivity.

As for the inter-crystalline diffusion process, the $D_{2,\text{self}}$ values can be affected by (i) the molecular size of a specific probe molecule, (ii) the pore size and topology of the MOF and (iii) the interaction between a probe molecule and the MOF framework, the latter affected by the presence of functional groups. As for aspect (ii), it is reasonable to assume that pore size and topology is similar across all MOFs studied, whereby we neglect the effect of additional linkers on pore size.

To further discuss (i) and (iii), we introduce the concept of apparent tortuosity, which is the ratio of the bulk diffusivity of a certain species to the diffusivity of the same species within the pore space, that is, a reduction in diffusivity inside the pore space compared to the bulk. In the MOF frameworks studied here we can identify two apparent tortuosities, $\xi_1 = D_{\text{self,bulk}}/D_{1,\text{self}}$ and $\xi_2 = D_{\text{self,bulk}}/D_{2,\text{self}}$. Values of ξ_1 and ξ_2 are shown in Fig. 4 and Tables S3–S5 (ESI[†]). Fig. 4(a) shows that the ξ_1 values for all probe molecules within the three MOFs is between 1.38–2.40. These values are consistent with diffusion occurring in a mesoporous network,¹⁷ hence this indeed suggests that this is diffusion occurring in the mesoscale space between the MOF crystals.

Values of the apparent tortuosity ξ_2 are much higher, on average ten times higher than ξ_1 . Such values are consistent with molecules diffusing in much smaller micropores, which supports the assignment of $D_{2,\text{self}}$ to diffusion within crystal aggregates in MOF frameworks, whereby the typical measured pore sizes are less than 2 nm.⁴¹

We now compare ξ_2 values across the different probe molecules used in the pristine UiO-66. TIPB is too big to access

through the UiO-66 framework's triangular windows, hence no ξ_2 can be defined in this case. Water and *n*-octane exhibit higher ξ_2 (slower $D_{2,\text{self}}$) compared to xylenes. The higher ξ_2 values could be attributed to either entanglement effects during the diffusion process of these molecules within the internal pore environment of the MOFs or to the presence of interactions with the framework affecting the diffusion process. In MOFs, entanglement effects may arise as diffusion processes are affected by various factors of the framework materials such as pore geometry/topology and functional groups on linkers.⁴² Entanglement in this context arises from interactions between MOFs and diffusing probe molecules. MOFs can exhibit various types of interaction with probe molecules, including van der Waals, hydrophobic/hydrophilic and π - π interactions.¹¹ These factors determine the guest–host interactions in MOFs, such as adsorbate-pore surface interaction and surface-to-pore diffusion, which play a significant role in determining the rate of diffusion.

For the case of water, the high values in ξ_2 can be explained by considering the preferential interactions of water molecules within the frameworks. Given the geometry of UiO-66, the dimensions of its cages are suitable for confining water. Wang *et al.*, conducted a study investigating water behaviour and confinement in UiO-66 using molecular modelling and simulations. Their findings indicate that water molecules tend to diffuse through tetrahedral cages due to stronger binding compared to octahedral cages, leading to confinement within.⁴³ In this study, while we cannot definitively confirm through which cages water initially diffuses, we observe a similar behaviour characterised by low $D_{2,\text{self}}$ values (Table S3, ESI[†]). The slower diffusion suggests the presence of hydrogen bonding interactions between water and the UiO-66 framework influencing water diffusion within the MOFs. Similar phenomena were previously reported on water adsorption in UiO-66, where the presence of a metal oxyhydroxy cluster, which contains Zr–OH moieties, promotes coordination with small molecules, hence slowing down water diffusion.⁴⁴

For straight-chain alkanes, like *n*-octane, the high ξ_2 values can be attributed to the linear arrangement of the carbon chain, which may become entangled within the framework,

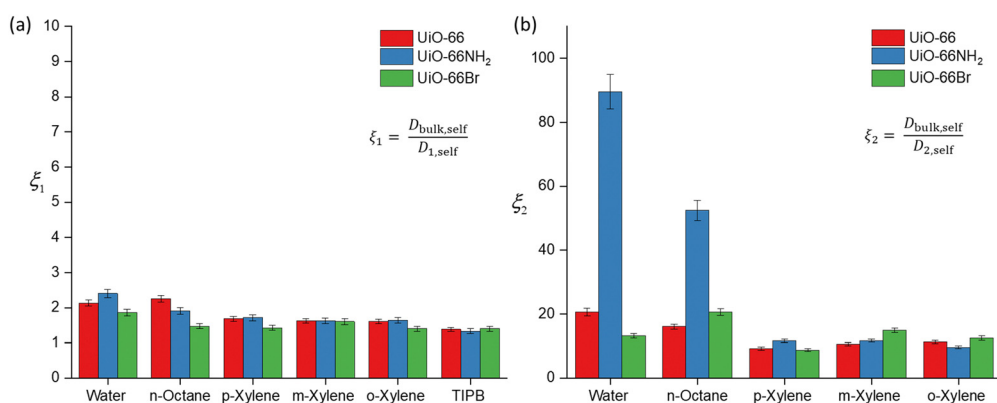


Fig. 4 Apparent tortuosity values of the probe molecule–MOF systems under investigation obtained from PFG NMR. (a) Apparent tortuosity ξ_1 values derived from the ratio of $D_{\text{bulk,self}}/D_{1,\text{self}}$. (b) Apparent tortuosity ξ_2 obtained from ratio of $D_{\text{bulk,self}}/D_{2,\text{self}}$. The relative error is approximately 5–6%.



leading to slower diffusion.⁴⁵ As shown in Fig. 2, within UiO-66, each octahedral cage is surrounded by eight tetrahedral cages. This interconnected network of tetrahedral and octahedral cages enables the diffusion of *n*-octane from one cage to neighbouring cages. However, due to its linear arrangement, *n*-octane is prone to experiencing entanglement effects as it diffuses through the tetrahedral and octahedral cages.

Regarding the xylene isomers, the trend in ζ_2 values indicates that *o*-xylene exhibits a slightly higher value (11.3), followed by *m*-xylene (10.6) and *p*-xylene (9.22). Although the differences in apparent tortuosity values are small, this indicates different preferential arrangements of these isomers within the MOFs. This discrepancy can be attributed to steric effects arising from the position of the two methyl groups on the benzene ring, leading to differences in the measured ζ_2 values. UiO-66 also shows a reverse shape selectivity pattern,⁴⁶ whereby the less bulky *p*-xylene is the least retained isomer, contrary to the typical shape selectivity based on molecular dimensions. Here, we observe a trend consistent with this pattern, with $D_{2,\text{self}}$ having the following trend, *p*-xylene > *m*-xylene > *o*-xylene. This interaction has been previously studied in the xylene isomers separation process.⁴⁷

The calculated $D_{2,\text{self}}$ and ζ_2 in UiO-66NH₂ and UiO-66Br for all probe molecules differ from those obtained for the parent UiO-66. Observing the xylenes isomers, the ζ_2 values for UiO-66NH₂ show similar values to the pristine UiO-66. Differences are more marked for UiO-66Br when considering *m*-xylene and *o*-xylene, which show slightly higher values of ζ_2 than those found for UiO-66, whilst for *p*-xylene there is no significant difference. These differences can be ascribed to changes in pore accessibility due to the introduction of the bromine group. Pristine UiO-66 showed ζ_2 values of 11.3 for *o*-xylene and 10.6 for *m*-xylene. Following linker functionalisation with -Br, these values increased. This suggests that the presence of -Br in UiO-66 alters pore accessibility by modifying the materials structure and properties. When attached to the linker, -Br molecules undergo rotational movement within the pore system, hindering pore access. This rotational motion leads to steric hindrance, which obstructs the entry of probe molecules into the pores. This rotational linker has been previously investigated.⁴⁸ As -Br linkers rotate, they can either enhance or restrict pore openings, consequently influencing the overall porosity of these materials. This effect is evidenced by the N₂ physisorption results, showing that UiO-66Br exhibits a reduced specific surface area compared to pristine UiO-66.

The ζ_2 value of water in functionalised UiO-66NH₂ increases markedly to 83.9 from 20.6 (in UiO-66) but decreases to 13.2 in UiO-66Br. These changes are attributed to the changes in guest molecule-MOF interaction due to linker functionalisation. The grafting of hydrophilic -NH₂ functional groups enhances the hydrogen bonding interaction with water, which can affect its diffusivity.⁴⁹ While the geometry of UiO-66 is intrinsically conducive to water confinement, functionalising it with -NH₂ groups intensifies the confinement effect not only due to geometry but also due to increased guest-host interactions.⁵⁰ For

UiO-66Br, the decrease in ζ_2 is due to the steric hindrance caused by the -Br group. As discussed earlier, the introduction of -Br would hinder the pore access. While UiO-66 is inherently hydrophilic,⁵¹ functionalising it with -Br will alter its hydrophilicity, resulting in lower exchange of water within the framework.

The ζ_2 value for *n*-octane increases from 16.1 in UiO-66 to 20.6 in UiO-66Br and 52.4 in UiO-66NH₂. This trend is expected because the functionalisation of MOFs introduces steric hindrance, consequently slowing down the movement of molecules. Previous studies have demonstrated that grafting a polar functionality on UiO-66 reduces the linker's rotational ability, which hinders the accessibility of guest molecules.⁵² Our results highlight a significant impact of functionalising MOFs with -NH₂ and -Br groups on the dynamics and framework flexibility within UiO-66.

4. Conclusions

In this work, we demonstrate the usefulness of the PFG NMR technique in unravelling the diffusion behaviour within UiO-66 and its derivatives. Furthermore, we investigate the impact of functionalisation on mass transport in these materials. The results show the presence of two diffusion environments, which are accessible to smaller probe molecules, $D_{1,\text{self}}$ associated with inter-crystalline diffusion and $D_{2,\text{self}}$ associated with diffusion within crystal aggregates. When molecules such as TIPB with a diameter larger than the diameter of the cage windows of UiO-66 and its derivatives are used as a probe for diffusion, only a single diffusion environment in the inter-crystalline space is observed, which is attributed to the inability of TIPB molecules to access the MOF framework. Functionalisation of the pristine MOF with amino and bromine groups leads to a significant change in diffusion within crystal aggregates, $D_{2,\text{self}}$, due to changes in intra-framework interactions, particularly for water and *n*-octane, whereas does not significantly affect transport of molecules in the inter-crystalline space. This indicates that the diffusion properties within MOF structures can be influenced by factors such as rotational barriers of functionalised linker molecules and interactions between probe molecules and the MOF structure.

Conflicts of interest

The authors declare no competing financial interest.

Acknowledgements

This project has received funding from the European Union's Horizon 2020 research and innovation program under grant agreement no. 872102 and the EPSRC (EP/N033949/1 and EP/R018790/1). S. Z. would like to thank Majlis Amanah Rakyat (MARA) for funding her PhD research.



References

- J. H. Cavka, S. Jakobsen, U. Olsbye, N. Guillou, C. Lamberti, S. Bordiga and K. P. Lillerud, *J. Am. Chem. Soc.*, 2008, **130**, 13850–13851.
- M. Kandiah, M. H. Nilsen, S. Usseglio, S. Jakobsen, U. Olsbye, M. Tilset, C. Larabi, E. A. Quadrelli, F. Bonino and K. P. Lillerud, *Chem. Mater.*, 2010, **22**, 6632–6640.
- L. Valenzano, B. Civalieri, S. Chavan, S. Bordiga, M. H. Nilsen, S. Jakobsen, K. P. Lillerud and C. Lamberti, *Chem. Mater.*, 2011, **23**, 1700–1718.
- J. Guo, X. Fan, J. Wang, S. Yu, M. Laipan, X. Ren, C. Zhang, L. Zhang and Y. Li, *Chem. Eng. J.*, 2021, **425**, 130588.
- S. Zhao, W. Huang, J. Xie, Z. Qu and N. Yan, *Fuel*, 2021, **289**, 119807.
- W. Xiang, Y. Zhang, Y. Chen, C.-J. Liu and X. Tu, *J. Mater. Chem. A*, 2020, **8**, 21526–21546.
- A. Dhakshinamoorthy, A. Santiago-Portillo, A. M. Asiri and H. Garcia, *ChemCatChem*, 2019, **11**, 899–923.
- R. Vakili, R. Gholami, C. E. Stere, S. Chansai, H. Chen, S. M. Holmes, Y. Jiao, C. Hardacre and X. Fan, *Appl. Catal., B*, 2020, **260**, 118195.
- R. Vakili, E. K. Gibson, S. Chansai, S. Xu, N. Al-Janabi, P. P. Wells, C. Hardacre, A. Walton and X. Fan, *ChemCatChem*, 2018, **10**, 4238–4242.
- R. Vakili, S. Xu, N. Al-Janabi, P. Gorgojo, S. M. Holmes and X. Fan, *Microporous Mesoporous Mater.*, 2018, **260**, 45–53.
- C. H. Sharp, B. C. Bukowski, H. Li, E. M. Johnson, S. Ilic, A. J. Morris, D. Gersappe, R. Q. Snurr and J. R. Morris, *Chem. Soc. Rev.*, 2021, **50**, 11530–11558.
- J. J. Wardzala, J. P. Ruffley, I. Goodenough, A. M. Schmidt, P. B. Shukla, X. Wei, A. Bagussetty, M. De Souza, P. Das, D. J. Thompson, C. J. Karwacki, C. E. Wilmer, E. Borguet, N. L. Rosi and J. K. Johnson, *J. Phys. Chem. C*, 2020, **124**, 28469–28478.
- N. A. Ramsahye, J. Gao, H. Jobic, P. L. Llewellyn, Q. Yang, A. D. Wiersum, M. M. Koza, V. Guillerm, C. Serre, C. L. Zhong and G. Maurin, *J. Phys. Chem. C*, 2014, **118**, 27470–27482.
- Z. Zhao, Z. Li and Y. Lin, *Ind. Eng. Chem. Res.*, 2009, **48**, 10015–10020.
- F. Stallmach and P. Galvosas, *Annu. Rep. NMR Spectrosc.*, 2007, 51–131, DOI: [10.1016/S0066-4103\(07\)61102-8](https://doi.org/10.1016/S0066-4103(07)61102-8).
- C. D'Agostino, R. C. Harris, A. P. Abbott, L. F. Gladden and M. D. Mantle, *Phys. Chem. Chem. Phys.*, 2011, **13**, 21383–21391.
- C. D'Agostino, J. Mitchell, L. F. Gladden and M. D. Mantle, *J. Phys. Chem. C*, 2012, **116**, 8975–8982.
- J. Kärger, D. Freude and J. Haase, *Processes*, 2018, **6**, 147.
- F. Stallmach and J. Kärger, *Adsorption*, 1999, **5**, 117–133.
- S. V. P. Kortunov, J. Kärger, R. Valiullin, P. Gottschalk, M. FéElía, M. S. C. M. Perez, B. Drescher, G. McElhiney, C. Berger, R. Gläser and J. Weitkamp, *J. Am. Chem. Soc.*, 2005, **10**, 13055–13059.
- S. Naumov, R. Valiullin, J. Kärger, R. Pitchumani and M.-O. Coppens, *Microporous Mesoporous Mater.*, 2008, **110**, 37–40.
- N. Dvoyashkina, C. F. Seidler, M. Wark, D. Freude and J. Haase, *Microporous Mesoporous Mater.*, 2018, **255**, 140–147.
- F. Stallmach, S. Gröger, V. Künzel, J. Kärger, O. Yaghi, M. Hesse and U. Müller, *Angew. Chem., Int. Ed.*, 2006, **45**, 2123–2126.
- D. Sinnaeve, *Concepts Magn. Reson., Part A*, 2012, **40A**, 39–65.
- L. Castañar, G. D. Poggetto, A. A. Colbourne, G. A. Morris and M. Nilsson, *Magn. Reson. Chem.*, 2018, **56**, 546–558.
- D. Zou and D. Liu, *Mater. Today Chem.*, 2019, **12**, 139–165.
- C. R. Groom, I. J. Bruno, M. P. Lightfoot and S. C. Ward, *Acta Crystallogr., Sect. B: Struct. Sci., Cryst. Eng. Mater.*, 2016, **72**, 171–179.
- M. Kim and S. M. Cohen, *CrystEngComm*, 2012, **14**, 4096–4104.
- K. Momma and F. Izumi, *J. Appl. Crystallogr.*, 2011, **44**, 1272–1276.
- S. Zhao, W. Huang, J. Xie, W. Liu, Z. Qu and N. Yan, *Fuel*, 2021, **289**, 119807.
- R. Vakili, S. Xu, N. Al-Janabi, P. Gorgojo, S. M. Holmes and X. Fan, *Microporous Mesoporous Mater.*, 2018, **260**, 45–53.
- J. Kärger, D. M. Ruthven and D. N. Theodorou, *Diffusion in nanoporous materials*, Wiley Online Library, 2012.
- C. D'Agostino, Y. Ryabenkova, P. J. Miedziak, S. H. Taylor, G. J. Hutchings, L. F. Gladden and M. D. Mantle, *Catal. Sci. Technol.*, 2014, **4**, 1313–1322.
- T. C. Bowen, R. D. Noble and J. L. Falconer, *J. Membr. Sci.*, 2004, **245**, 1–33.
- A. Kurganov, S. Marmé and K. Unger, in *Zeolites and Related Microporous Materials: State of the Art 1994 – Proceedings of the 10th International Zeolite Conference, Garmisch-Partenkirchen, Germany, 17–22 July 1994*, 1994, pp. 1299–1306, DOI: [10.1016/S0167-2991\(08\)63671-X](https://doi.org/10.1016/S0167-2991(08)63671-X).
- A. Bazyari, A. A. Khodadadi, N. Hosseinpour and Y. Mortazavi, *Fuel Process. Technol.*, 2009, **90**, 1226–1233.
- T. M. O. Popp, A. Z. Plantz, O. M. Yaghi and J. A. Reimer, *ChemPhysChem*, 2020, **21**, 32.
- Y. Jiao, L. Forster, S. Xu, H. Chen, J. Han, X. Liu, Y. Zhou, J. Liu, J. Zhang and J. Yu, *Angew. Chem., Int. Ed.*, 2020, **59**, 19478–19486.
- J. Kärger and S. Vasenkov, *Microporous Mesoporous Mater.*, 2005, **85**, 195–206.
- P. Kortunov, S. Vasenkov, J. Kärger, R. Valiullin, P. Gottschalk, M. Fé Elía, M. Perez, M. Stöcker, B. Drescher and G. McElhiney, *J. Am. Chem. Soc.*, 2005, **127**, 13055–13059.
- Q. Zhu, G. D. Moggridge, M. Ainte, M. D. Mantle, L. F. Gladden and C. D'Agostino, *Chem. Eng. J.*, 2016, **306**, 67–76.
- O. M. Yaghi, *Nat. Mater.*, 2007, **6**, 92–93.
- S. Wang, G. Zhou, Y. Sun and L. Huang, *AIChE J.*, 2021, **67**, e17035.
- T. G. Grissom, C. H. Sharp, P. M. Usov, D. Troya, A. J. Morris and J. R. Morris, *J. Phys. Chem. C*, 2018, **122**, 16060–16069.
- D. Schuring, A. Jansen and R. Van Santen, *J. Phys. Chem. B*, 2000, **104**, 941–948.
- P. S. Bárcia, D. Guimarães, P. A. Mendes, J. A. Silva, V. Guillerm, H. Chevreau, C. Serre and A. E. Rodrigues, *Microporous Mesoporous Mater.*, 2011, **139**, 67–73.



- 47 P. Silva, S. M. Vilela, J. P. Tome and F. A. Almeida Paz, *Chem. Soc. Rev.*, 2015, **44**, 6774–6803.
- 48 J. T. Damron, J. Ma, R. Kurz, K. Saalwächter, A. J. Matzger and A. Ramamoorthy, *Angew. Chem., Int. Ed.*, 2018, **130**, 8814–8817.
- 49 C. Caratelli, J. Hajek, F. G. Cirujano, M. Waroquier, F. X. L. i Xamena and V. Van Speybroeck, *J. Catal.*, 2017, **352**, 401–414.
- 50 J. Zhang, F. Paesani and M. Lessio, *J. Mater. Chem. C*, 2023, **11**, 10247–10258.
- 51 X. Liu, X. Wang and F. Kapteijn, *Chem. Rev.*, 2020, **120**, 8303–8377.
- 52 S. Devautour-Vinot, G. Maurin, C. Serre, P. Horcajada, D. Paula da Cunha, V. Guillermin, E. N. de Souza Costa, F. Taulelle and C. Martineau, *Chem. Mater.*, 2012, **24**, 2168–2177.

