A close view of the organic linker in a MOF: structural insights from a combined $^1$H NMR relaxometry and computational investigation

Silvia Pizzanelli, D, E Susanna Monti, D, A Larisa G. Gordeeva, B, C Marina V. Solovyeva, B, C Angelo Freni a and Claudia Forte D, A

The organic linker in a metal organic framework (MOF) affects its adsorption behavior and performance, and its structure and dynamics play a role in the modulation of the adsorption properties. In this work, the combination of $^1$H nuclear magnetic resonance (NMR) longitudinal relaxometry and theoretical calculations allowed detailed structure and dynamics of the organic linker in the NH$_2$-MIL-125 MOF to be obtained. In particular, fast field cycling (FFC) NMR, applied here for the first time on MOFs, was used to disclose the dynamics of the amino group and its electronic environment through the analysis of the $^{14}$N quadrupole relaxation peaks, observed in the frequency interval 0.5–5 MHz, at different temperatures from 25 to 110 °C. The line width of the peaks allowed a lower boundary on the rotational correlation time of the N–H bonds to be set, whereas relevant changes in the amplitudes were interpreted in terms of a change in the orientation of the $^{14}$N averaged electric field gradient tensor. The experimental findings were complemented by quantum chemistry calculations and classical molecular dynamics simulations.

Introduction

Over the last two decades, metal organic frameworks (MOFs), also known as coordination polymers, have aroused increasing interest because of their high potential for applications in diverse fields, such as gas separation and storage, energy storage, catalysis, and thermal energy conversion. These hybrid crystalline solids are made of metal clusters linked by organic multidentate ligands that form extended three-dimensional porous frameworks with tunable pore size and shape through suitable chemical functionalities. Distinctive features of these versatile materials are extraordinarily large porosity and surface area, and unique adsorption properties.

The functionalization of the organic ligand may induce a modulation of the adsorption properties. Although the role of the functionalization has been disclosed in many applications, very little attention has been paid to the dynamics and the environment of the functional groups, which might contribute to rationalizing the adsorption behavior.

This study is focused on NH$_2$-MIL-125, a microporous MOF composed of Ti$_2$O$_4$(OH)$_7$(BDC-NH$_2$)$_6$ (BDC = benzene-1,4-dicarboxylate) units, with a cyclic inorganic octamer structure built from TiO$_6$(OH) octahedra and connected to other 12 octamers through NH$_2$-BDC linkers. This MOF exhibited promising properties in the field of gas separation, liquid phase separation of hydrocarbons, adsorption heat transformation, and storage. A previous Nuclear Magnetic Resonance (NMR) investigation on the rotational motion of the amino group bound to the BDC ligand and of the aromatic ring in a Zn-based MOF, IRMOF-3, showed that the correlation time for both processes is on the order of microseconds in the 25–100 °C temperature range. Values of the same order of magnitude were found for the rotational motion of the aromatic ring in NH$_2$ functionalized UIO-66(Zr).

Here, we applied $^1$H FFC relaxometry to characterize the dynamics and the electronic environment of the amino group bound to the BDC unit, exploiting the quadrupolar relaxation enhancement (QRE) of the $^1$H longitudinal relaxation rate. Indeed, when the dynamics is relatively slow, the $^1$H spin lattice relaxation rate increases at $^1$H frequencies matching the $^{14}$N quadrupole spin transition. This effect was observed in some solids, and in biological systems but, to the best of our knowledge, it was never reported for a MOF. The positions of the...
quadrupolar peaks were used to derive the quadrupolar coupling constant and the asymmetry parameter of the electric field gradient (EFG) acting on the $^{14}$N nucleus, whereas the relative intensities and widths of the peaks were related to the orientation of the N–H vectors with respect to the averaged EFG principal axes and to their rotational correlation and $^{14}$N relaxation times through simple analytical expressions. Differently from $^2$H NMR spectroscopy, which is a well established technique for probing molecular mobility, applied by Kolokolov et al. to linkers and guest molecules adsorbed in MOFs, the method used here does not require deuteration of the sample, but it is restricted to systems containing a quadrupolar nucleus, such as $^{14}$N. The flip of the BDC units was excluded to obtain a picture of the amino group geometry and dynamics, and of the EFG tensor at the nitrogen centers.

### Materials and methods

#### NMR measurements

NH$_2$-MIL-125 was synthesized according to the procedure reported in ref. 14. The as-synthesized product was dried at 150°C in vacuo for 20 hours.

$^1$H longitudinal relaxation times, $T_1$, were measured in the 0.01–35 MHz Larmor frequency range on a SpinMaster FFC-2000 Fast Field-Cycling NMR relaxometer (Stelar srl, Mede, Italy) using the pre-polarized and non-polarized pulse sequences below and above 12 MHz, respectively. The polarization and detection fields were 0.6 T and 0.5 T, which correspond to $^1$H Larmor frequencies of 25.0 and 21.5 MHz, respectively. The switching time was 3 ms and the 90° pulse duration was 9.8 μs. A single scan was acquired. All the other experimental parameters were optimized for each measurement. All the $^1$H magnetization curves vs. time were mono-exponential within experimental error and the errors in the relaxation rate $R_1 = 1/T_1$ fitting were always less than 1%. The experiments were performed at three temperatures, i.e. at 25, 80 and 110°C; the temperature was controlled within ±0.1°C with a Stelar VITC90 variable temperature controller.

$^3$H–$^{13}$C dipolar interactions were estimated at room temperature using a Bruker Avance Neo-300 WB spectrometer equipped with a 4 mm probe. The operating frequencies were 300.13 and 75.47 MHz for $^1$H and $^{13}$C, respectively. The 90° pulses on $^1$H and $^{13}$C were 3.0 and 4.0 μs, respectively. The 2D LG-CP experiment was conducted using the sequence reported in Fig. 2a of ref. 29. For the LG-CP period, the $^1$H effective field strength was 50 kHz and the $^{13}$C spin-lock field strength was adjusted to the first order sideband conditions. The LG-CP contact time was incremented in 41.7 μs steps with 100 steps. The spinning speed was 12 kHz, the recycle delay was set to 1.2 s and the number of scans was 2000. Only cosine-modulated data were collected. The $t_e$ time signals were processed using a baseline correction mode “qfil” in Bruker TopSpin 3.5 software, which subtracts a constant intensity before performing a real Fourier transformation, to eliminate the zero-frequency peak.

#### Analysis of the $R_1$ profile: theoretical basis

$^{14}$N is a quadrupolar nucleus with spin quantum number equal to 1. In the low field regime, i.e. when the magnetic field $B_0$ is smaller than 0.1 T, the Zeeman coupling of the $^{14}$N spin is much weaker than its electrostatic interaction with the local EFG and can be neglected. Therefore, the $^{14}$N energy levels are determined by the interaction between the nuclear quadrupole moment and the EFG. The frequencies associated to the possible transitions between these levels are $\nu_\pm = \frac{e^2 q Q}{h} \left(\frac{1}{2} \pm \eta\right)$, and $\nu_0 = \frac{e^2 q Q}{h} \frac{\eta}{2}$ with $\eta$ representing the $^{14}$N quadrupolar coupling constant and the asymmetry parameter, respectively. In the $^1$H $R_1$ profile a local maximum may occur when the $^1$H Zeeman transition energy matches one of the transition frequencies $\nu_\pm$, $\nu_0$, or $\nu_\parallel$. In the case of a $^{14}$N nucleus belonging to a slowly tumbling entity (i.e., $2\pi\frac{e^2 q Q}{h}\tau_{rot} \gg 1$, with $\tau_{rot}$ representing the N–H rotational correlation time), the following simple analytical expression of the $^1$H longitudinal QRE, $R_1^{\text{HIN}}$, was reported:

$$R_1^{\text{HIN}}(\nu_H) = 2c \times \left\{ A_1 \left[ \frac{\tau_{\text{eff}}}{1 + (2\pi\tau_{\text{eff}})^2(\nu_H - \nu_\pm)^2} + \frac{\tau_{\text{eff}}}{1 + (2\pi\tau_{\text{eff}})^2(\nu_H + \nu_\pm)^2} \right] + A_2 \left[ \frac{\tau_{\text{eff}}}{1 + (2\pi\tau_{\text{eff}})^2(\nu_H - \nu_\parallel)^2} + \frac{\tau_{\text{eff}}}{1 + (2\pi\tau_{\text{eff}})^2(\nu_H + \nu_\parallel)^2} \right] \right. $$

$$\left. + A_0 \left[ \frac{\tau_{\text{eff}}}{1 + (2\pi\tau_{\text{eff}})^2(\nu_H - \nu_0)^2} + \frac{\tau_{\text{eff}}}{1 + (2\pi\tau_{\text{eff}})^2(\nu_H + \nu_0)^2} \right] \right\}$$

(1)

with $c = \frac{1}{3} \left( \frac{\mu_0 \gamma_H \gamma_N N_H}{4\pi} \right)^2 r^6$, where $r$ represents the internuclear N–H distance, $\gamma_H$ and $\gamma_N$ are the gyromagnetic ratios of $^1$H and $^{14}$N nuclei, respectively, and $\mu_0$ is the permeability. In eqn (1) $\tau_{\text{eff}}$ is an effective correlation time dependent on $\tau_{rot}$ and on the $^{14}$N relaxation time $T_{1\text{N}}$ due to spin–flip fluctuations, according to the equation:

$$\frac{1}{\tau_{\text{eff}}} = \frac{1}{\tau_{\text{rot}}} + \frac{1}{T_{1\text{N}}}$$

(2)

At $\nu_H$ frequencies close to $\nu_\pm$, $\nu_\parallel$, or $\nu_0$, the terms in eqn (1) depending on $\nu_H + \nu_\pm$, $\nu_H + \nu_\parallel$, and $\nu_H + \nu_0$ are negligible, and $R_1^{\text{HIN}}$ is described by three Lorentzian peaks centered at $\nu_\pm$, $\nu_\parallel$, and $\nu_0$ with amplitudes $A_1$, $A_2$, and $A_0$, respectively; their full width at half-height (FWHH), expressed in frequency units, is related to $\tau_{\text{eff}}$ according to the following equation:

$$\text{FWHH} = \frac{1}{\pi\tau_{\text{eff}}}$$

(3)
In principle, one should also consider a contribution to the linewidths due to the dependence of the resonance condition on the orientation of the external magnetic field in the EFG principal axis system (PAS). However, this contribution is estimated to be less than 10% and can be safely neglected. It was shown that the amplitudes of the peaks depend on the polar and azimuthal angles \( \theta \) and \( \phi \) defining the orientation of the \(^{1}H^{14}N\) inter-spin vector in the PAS of the averaged EFG, according to the following equations:

\[
A_{+} = \frac{1}{3} + \sin^{2} \theta \cos^{2} \phi \quad (4a)
\]

\[
A_{-} = \frac{1}{3} + \sin^{2} \theta \sin^{2} \phi \quad (4b)
\]

\[
A_{0} = \frac{1}{3} + \cos^{2} \theta \quad (4c)
\]

In our system two N–H bonds are present, each one contributing to \( \kappa_{\text{HIN}} \); it can be assumed that a common \( \tau_{\text{eff}} \) value is associated with the two bonds. These, however, are characterized by different orientations in the EFG PAS, expressed by the angles \( \theta_{1}, \phi_{1} \) and \( \theta_{2}, \phi_{2} \), pictured in Fig. 1. Hence, the two hydrogen atoms contribute to a given peak with different amplitudes related to their different orientations. In order to take this into account, eqn (4) were modified as follows:

\[
A_{+} = \sum_{i=1}^{2} \left( \frac{1}{3} + \sin^{2} \theta_{i} \cos^{2} \phi_{i} \right) \quad (5a)
\]

\[
A_{-} = \sum_{i=1}^{2} \left( \frac{1}{3} + \sin^{2} \theta_{i} \sin^{2} \phi_{i} \right) \quad (5b)
\]

\[
A_{0} = \sum_{i=1}^{2} \left( \frac{1}{3} + \cos^{2} \theta_{i} \right) \quad (5c)
\]

Analysis of the \( R_{1} \) profile: data treatment

The measured \(^{1}H\) longitudinal relaxation rate consists of a homonuclear dipolar contribution, \( R_{1}^{\text{HH}} \), due to \(^{1}H^{1}H\) dipole-dipole interactions, and a QRE contribution, \( R_{1}^{\text{QRE}} \), due to \(^{1}H^{14}N\) dipole-dipole interactions. The overall relaxation \( R_{1}(\nu) \) is the sum of the two contributions:

\[
R_{1}(\nu) = R_{1}^{\text{HH}}(\nu) + R_{1}^{\text{QRE}}(\nu) \quad (6)
\]

with the first term contributing to a regular decrease with increasing frequency and the second term giving rise to the quadrupolar peaks.

In order to obtain \( R_{1}^{\text{QRE}} \), \( R_{1}^{\text{HH}} \) was subtracted from the measured \( R_{1} \) at each frequency using the baseline correction tool in the Origin software. We did not attempt to analyze the \( R_{1}^{\text{HH}} \) contribution, as carried out by other authors, because the data did not show distinctive features capable of unambiguously defining the dynamic processes modulating the \(^{1}H^{1}H\) dipolar interactions. For a given temperature, the experimental \( R_{1}^{\text{QRE}} \) data were fitted using eqn (1) within the Mathematica 10 programming environment. The fitting parameters relative to each peak \( i \), with \( i = +, -, 0 \), were \( \nu_{i} \) and \( \Delta \nu_{i} \) besides the common \( \tau_{\text{eff}} \).

The values of the four angles defining the orientation of the two N–H bonds, namely \( \theta_{1}, \phi_{1}, \theta_{2}, \phi_{2} \), were derived from the relative amplitudes of the \( \nu_{+} \) and \( \nu_{-} \) peaks with respect to that of the \( \nu_{0} \) peak, using eqn (3). We imposed the constraint that the angle between the two N–H bonds was between 105 and 125 degrees, the most populated region according to MD simulations (Fig. S1 of ESI†), and looked for numerical solutions by varying the polar and the azimuthal angles on a grid in the range 0–\( \pi \) and 0–2\( \pi \), respectively. We rejected those angular sets that produced deviations of the calculated rescaled amplitudes from the experimental values larger than 5%. It should be noticed that, given the symmetry of the trigonometric functions appearing in eqn (5), we were not able to distinguish among the possible solutions \( \theta_{i}, -\theta_{i}, \pi - \theta_{i} \) and \( -\pi + \theta_{i} \), the same holding for \( \phi_{i} \).

Quantum chemistry calculations and classical simulations

The atomic coordinates and unit cell parameters (18.7028 × 18.7028 × 18.2028 Å³) of NH₂-MIL-125 were obtained from the Cambridge Crystallographic Data Centre and optimized at the density functional (DFT) level, after adding the hydrogen atoms, using the Quantum Espresso package, ultra-soft pseudopotentials and the generalized gradient approximation (GGA) with the Perdew–Burke–Ernzerhof (PBE) functional. Periodic boundary conditions were applied in all directions and a cutoff energy of 600 eV was employed to obtain convergent results. The Gauge Including Projector Augmented Wave (GIPAW) approach was used to calculate and fully characterize the traceless EFG tensor for each nitrogen nucleus (that is, the quadrupolar coupling constant, the asymmetry parameter and the orientation of the PAS with respect to the molecular frame).

To explore the dynamics of the MOF at 25 and 110 °C and generate a reasonable variety of geometries for predicting and explaining the experimental data, classical reactive molecular
dynamics simulations based on the ReaxFF methodology were carried out. This approach was selected because an appropriate parametrization for this type of system is available in the literature. The DFT optimized geometry was energy minimized at the classical level and then gradually heated to 25 °C (or 110 °C). Equilibration was carried out for about 50 ps. After equilibration, the sampling simulations were performed in the NV T ensemble for about one nanosecond. System configurations were collected every 0.1 ps. The temperature was controlled through the Berendsen’s thermostat with a relaxation constant of 0.05 ps, and the time step was set to 0.2 fs. The analysis of the production trajectories was focused on the orientation of the NH2 group relative to the ring system, the degree of pyramidalization of the nitrogen atom, the H–N–H angles, the C–C–N–H dihedral angles, and the length of the hydrogen bond between the amino and carboxyl group. Representative configurations were extracted from the two trajectories and EFG-GIPAW calculations were carried out to predict the orientations of the EFG tensors in the molecular frame (Fig. S2 of ESI†).

Results and discussion

The $R_1$ dispersion curves of dry NH$_2$-MIL-125, recorded at 25, 80 and 110 °C, are shown in Fig. 2a. The profiles at 25 and 80 °C are very similar, and only minor changes could be observed upon further heating to 110 °C. This indicated that, in the investigated temperature domain, no major dynamic process was activated in the 10 kHz–35 MHz frequency range. Moreover, the profiles display three quadrupolar peaks (Fig. 2b), due to the level crossing with the $^{14}$N quadrupole frequencies, which indicated that all the $^{14}$N nuclei are characterized by very similar EFGs, in line with the theoretical predictions, and provided important information on the very slow dynamics of the NH$_2$ group.

The quadrupolar contribution to $R_1$, after subtraction of the background curve mainly due to $R_{1HH}$, is shown in Fig. 3. The small frequency shifts of the maxima suggested that the quadrupole parameters $\frac{e^2 qQ}{\hbar}$ and $\eta$ slightly changed with the temperature. Fig. 3 clearly shows that the peaks sharpened upon increasing the temperature from 25 °C to 80 °C, whereas a further temperature increase to 110 °C did not result in a significant variation. Furthermore, increasing the temperature determined a change in the relative amplitudes of the three signals, which was most evident for the 25 °C to 80 °C variation.

For a quantitative analysis, the shape of the peaks, after subtraction of the background, was reproduced using the fitting function reported in eqn (1), with fitting parameters $\tau_{\text{eff}}$, $\nu_i$ and $cA_i$ relative to each peak $i$, with $i = +, -, 0$.
Fig. 3 shows the fitting curve at the investigated temperatures and Table 1 reports the best fit values. It is worth noting that the quadrupolar parameters are close to those determined for anthranilic acid with $^{14}$N quadrupole resonance spectroscopy.\(^{24}\)

A first theoretical estimate of the nuclear quadrupole constants and anisotropy parameters was obtained through the full optimization of the experimental crystal structure of the MOF at the QC level and subsequent calculations of the EFG tensor at the nitrogen nuclei. Inspection of the QC optimized geometry indicated that in NH$_2$-ML-125 all the $^{14}$N nuclei occupy positions characterized by similar electric field gradients with sp$^2$ hybridized nitrogen atoms hydrogen-bonded to the carboxyl oxygens. The best agreement of the theoretical nuclear quadrupole coupling constant with the experimental result was obtained with a $^{14}$N nuclear quadrupole moment of $1.94 \times 10^{-30}$ m$^2$, chosen in line with the studies of Bailey,\(^{44}\) that produced a quadrupolar coupling constant of $-3.79 \pm 0.04$ MHz and an asymmetry parameter around 0.31 ($\pm 0.03$) for all the nitrogen nuclei. This ideal situation satisfactorily compares with the experimental results.

Information on the dynamics of the amino group could be obtained from the $\tau_{\text{eff}}$ values determined from the fitting of the experimental data (Table 1). Of relevance is the fact that $\tau_{\text{rot}}$ should decrease with increasing temperature, the observed increase in $\tau_{\text{rot}}$ upon heating from 25 to 80 °C, as determined from the linewidth of the peaks using eqn (3); a further increase in temperature up to 110 °C did not give rise to any additional change. Since $\tau_{\text{rot}}$ should decrease with increasing the temperature, the observed increase in $\tau_{\text{rot}}$ suggests that $\tau_{\text{rot}}$ is mainly determined by $T_{1N}$ (see eqn (2)). A similar behavior was previously reported for the solid [C(NH$_2$)$_3$]$_3$Bi$_2$I$_9$ and was indeed explained taking into account the longitudinal relaxation time of $^{14}$N, $T_{1N}$, associated with a fast fluctuation of an EFG component, $i.e.$, on the time scale of 1 ns.\(^{18}\) In immobilized proteins, $T_{1N}$ values on the order of microseconds were ascribed to fast fluctuations of the EFG about its average value, probably due to a rocking motion on a 10 ns time scale.\(^{23}\) In our case, we could suppose that $T_{1N}$ significantly contributed to $\tau_{\text{rot}}$ at all the investigated temperatures, with $T_{1N} < \tau_{\text{rot}}$. Indeed, the detection of the quadrupolar peaks is evidence for a slow N–H bond dynamics, with the condition that the slow dynamics is characterized by a correlation time $\tau_{\text{rot}} \geq 1/2\pi/\nu_0 \approx 0.2 \mu$s.\(^{23}\) On the other hand, $T_{1N}$ values can be very short, especially at low fields, and in the extreme narrowing limit,\(^{23,45}\) the extreme narrowing regime predicts an increase in $T_{1N}$ upon heating, which results in the sharpening of the peaks with increasing the temperature. The observed temperature trend of $\tau_{\text{rot}}$ and the values determined indicated that $\tau_{\text{rot}}$ must be longer than 2.2 μs at 110 °C, a condition which also holds at 25 °C a fortiori. Therefore, the lower boundary obtained for $\tau_{\text{rot}}$ is longer than 0.2 μs, the low limit which was determined based on the conditions $\tau_{\text{rot}} \geq 1/2\pi/\nu_0$. Long $\tau_{\text{rot}}$ values were consistent with the absence of BDC flips at room temperature within a time scale of tens of microseconds, as revealed by the estimate of the $^{1}$H–$^{13}$C dipolar interactions by means of the 2D LG-CP experiment (Fig. S3 of ESI†). In fact, aromatic carbon atoms carrying a hydrogen atom were found to be subject to a $^{1}$H–$^{13}$C dipolar interaction close to the static value (see caption of Fig. S3 of the ESI†).

Other features of relevance for characterizing the changes that occur in the system upon heating were the absolute and relative peak amplitudes (see Table 1). The significant decrease in the amplitudes of the quadrupolar peaks, observed upon heating from 25 to 80 °C, indicated that the effective H–N dipole–dipole interaction leading to the QRE effect became weaker on increasing the temperature; this can be attributed to the pre-averaging of this interaction by a relatively fast dynamic process. Also, the relative values of the amplitudes changed upon heating from 25 to 80 °C, with the peaks centered at $\nu_0$ and $\nu_s$ assuming larger and smaller amplitude values, respectively. Only minor changes were observed upon further heating up to 110 °C, with a slight increase in amplitude of the $\nu_s$ and $\nu_0$ peaks and a decrease for the $\nu_z$ peak. We ascribed the variation of the relative amplitudes to a change in the orientation of the EFG tensor, according to eqn (5). These equations were used to fit the three amplitudes $A_x$, $A_y$, $A_z$ with the constraint that the angle between the two N–H bonds could assume values between 105 and 125 degrees, which is the most populated region according to the MD simulations discussed afterwards, and with variable parameters the angles $\theta_1$, $\phi_1$, $\theta_2$, $\phi_2$ defining the direction of the two N–H bonds in the averaged EFG PAS (Fig. 1).

At 25 °C, it was found that the set of angles {$(\theta_1 = 90 \pm 6, \phi_1 = 115 \pm 3, \theta_2 = 90 \pm 6, \phi_2 = 0 \pm 10$ degrees) was compatible with the observed relative amplitudes. This set implies that the EFG $z$-axis is approximately perpendicular to the H–N–H plane. This is in line with other studies,\(^{46,47}\) which found that the principal axis associated with the largest EFG component, $V_{zz}$, is oriented along the direction of the nitrogen lone pair. The two hydrogen atoms do not occupy symmetric positions with respect to the EFG $x$ and $y$ axes, and this could be ascribed to the formation of a hydrogen bond between the carboxylate oxygen and one of the hydrogen atoms.

At higher temperatures, since no major differences were observed between the amplitudes at 80 and 110 °C, we focused only on those at 110 °C. Several sets of angles were compatible with the experimental amplitudes, with $\theta$ values spread in the range 80–150 degrees and $\phi$ values clustered around –10 and

| $T$ (°C) | $\frac{CQ}{h}$ (MHz) | $\eta$ | $\tau_{\text{eff}}$ (μs) | $\frac{cA_x}{cA_y}$ | $\frac{cA_y}{cA_z}$ (s$^{-1}$) | $\nu_s$ (MHz) | $\nu_0$ (MHz) |
|----------|---------------------|-------|------------------------|-------------------|------------------------|-----------------|----------------|
| 25       | 3.52 (1)            | 0.41 (1) | 0.90 (2) | 25 (1); 3.00 (1); 21 (1); 2.28 (1); 8 (1); 0.67 (4) | 14 (1); 3.00 (1); 6 (1); 2.33 (1); 7 (1); 0.65 (2) | 13 (1); 3.00 (1); 8 (1); 2.31 (1); 10 (1); 0.64 (1) |
| 80       | 3.55 (1)            | 0.38 (1) | 2.2 (1) | 14 (1); 3.00 (1); 6 (1); 2.33 (1); 7 (1); 0.65 (2) | 13 (1); 3.00 (1); 8 (1); 2.31 (1); 10 (1); 0.64 (1) |
| 110      | 3.54 (1)            | 0.39 (1) | 2.2 (1) | 14 (1); 3.00 (1); 6 (1); 2.33 (1); 7 (1); 0.65 (2) | 13 (1); 3.00 (1); 8 (1); 2.31 (1); 10 (1); 0.64 (1) |

Values obtained by fitting $R_{1N}$ data, as described in the section Materials and methods. Uncertainty on the last digit is given in parenthesis.
130 degrees (Fig. 4). For the sake of clarity, the figure shows only four compatible sets, out of the possible values found. Among these sets, the corresponding value of the H–N–H angle was in the interval 105–125 degrees.

In order to better understand the experimental data and, possibly, confirm the analysis, an MD study was performed on the system at 25 and 110 °C. Examination of the conformational tendency of the BDC-NH₂ units at the two extreme temperatures revealed that the nitrogen atom could be both trigonal planar (sp² hybridization) or tetrahedral (sp³ hybridization) due to a rapid umbrella inversion that became more marked at a higher temperature. Given the inequivalence of the nitrogen substituents, the motion was slightly asymmetric and influenced by the relatively tight intramolecular hydrogen bond between one of the amine hydrogens and the nearby carboxyl oxygen. The MD results indicated that this hydrogen bond is strong enough to induce a moderate pyramidalization with a preferential direction, which is apparent in Fig. S1 of the ESI† where the distributions of the distance of the nitrogen from the C–H(N)–H(N) plane are displayed. This distance represents the displacement of the nitrogen above and below the substituents plane and can be correlated with the degree of pyramidalization of the atom. From inspection of these data it is apparent that the peaks of both curves are shifted to negative values and centered at approximately −0.03 Å. Further evidence of the nature of the nitrogen was provided by the H–N–H angle distribution (Fig. S1 of ESI†) which indicated that the most probable values were in the 110°–120° range, corresponding to characteristic tetrahedral and planar arrangements of the hydrogen atoms. This angular range was indeed the one considered, although slightly extended, in the analysis of the experimental data.

The asymmetry of the motion at both temperatures is clearly visible in Fig. 5 which shows the probabilities of the dihedral
angles defining the positions of the amine hydrogens relative to the ring plane at the two temperatures investigated. At 110 °C the higher propensity of the amino group to explore the outer regions of the range mentioned above was confirmed by the widening of the contours. Furthermore, the vibrations of the various portions of the MOF were faster and accompanied by modest deformations of the BDC-NH₂ rings and occasional rotations of both the NH₂ group (around the C–N bond) and the COO moiety (around the C–C bond). In particular, in the MD data displayed in Fig. 5, the NH₂ rotation occurred only once over a trajectory of 1 ns.

Representative configurations at the two temperatures were extracted for the calculation of the EFG tensor through the GIPAW approach; the one at 25 °C is shown in Fig. 6a where the BDC-NH₂ unit and the local EFG PAS are highlighted. It is apparent that the calculated ¹⁴N EFG tensor orientations are identical for all the planar BDC-NH₂ units, with Vₓᵧ perpendicular to the C–H(N)–H(N) plane (in this case, the plane of the molecule), oriented as the N lone pair, V𝑧ₓ in the plane of the molecule almost aligned to the H–N–H angle bisector and Vₓ𝑧 in the plane of the molecule perpendicular to the Vₓᵧ axis. This static picture became more complex when the effect of the temperature was applied and the local configurations of the BDC-NH₂ units were sampled as a function of time. In fact, a great variety of nitrogen pyramidalizations were identified and the EFG descriptors estimated in each case. The simplest cases were those where the hybridization of the nitrogen was not accompanied by a rotation of the NH₂ group around the C–N axis. In these cases, the hydrogens remained close to the molecular plane and the EFG PAS maintained an orientation typical of the flat sp² arrangement. Fig. 6b shows the EFG tensor orientation for pyramidalized nitrogen with hydrogens almost perpendicular to the O–C–O plane. Vₓᵧ is again in the direction of the lone pair, Vₓ𝑧 is aligned with the bisector of the H–N–H angle and Vₓ𝑧 is roughly perpendicular to the molecular plane. Therefore, the trait common to all the situations is the alignment of Vₓᵧ with the lone pair direction. Moreover, local effects did not drastically perturb the environment of the nitrogen. Even though the values of $\frac{e^2\eta Q}{h}$ and η were different in each unit, their average was close to the values found for the QC optimized structure and from the experiments.

At 25 °C, where the main tendency of the NH₂ group was to adopt an almost flat arrangement with only a slight pyramidalization, the GIPAW calculations on the DFT-optimized structure and on geometries extracted from the MD simulations gave the angular values $\{\theta₁ = 85 \pm 12, \phi₁ = 132 \pm 7, \theta₂ = 85 \pm 12, \phi₂ = 19 \pm 14 \text{degrees}\}$ for the two amine hydrogen atoms in the EFG PAS, in fairly good agreement with the values determined from the fitting of the experimental peak amplitudes. MD simulations did not predict significant changes in the static EFG tensor when the temperature was increased from 25 to 110 °C. Therefore, the change in the relative amplitudes of the QRE peaks could not be ascribed to a modification of the orientation of the static EFG tensor. However, the observed amplitudes were compatible with sets of angles $\{\theta₁, \phi₁, \theta₂, \phi₂\}$ (see Fig. 4) implying a significant deviation of the Vₓᵧ component from the lone pair direction. Such deviation was quantified through the angle between the normal to the plane defined by H–N–H and the EFG z-axis for our compatible sets, which was $65 \pm 10^\circ$, to be compared with the values of 0 and $35^\circ$ for an sp² and an sp³ hybridization, respectively. We thus speculated that the tensor is subject to a fast motion that modifies its orientation, and consequently the angles $\{\theta₁, \phi₁, \theta₂, \phi₂\}$ at 110 °C. Considering the relative amplitudes of the peaks and the observed $\frac{e^2\eta Q}{h}$ and η values, we excluded that the motion could be simply modelled as a two-site rotation of the NH₂ group around the C2 axis with jump angles ranging from 0 to $180^\circ$ or as a rocking motion about an axis perpendicular to the NH₂ plane. A more complex motional process might occur, possibly involving a different dynamic behavior of the two H–N bonds, as hinted by the MD data. At 25 °C this motion is probably more restricted so that the static tensor satisfactorily accounts for the amplitude values. This motion would be responsible for the decrease of $T_{1N}$ and for the slight change in the quadrupolar parameters upon heating.

**Conclusions**

For the first time we have applied ¹H FFC NMR relaxometry to study a metal organic framework, disclosing important details of the amino group dynamics in the specific environment constituted by NH₂-MIL-125, and of its EFG tensor at different temperatures. From the linewidth of the QRE peaks we could set a lower boundary on the rotational correlation time of the N–H bonds, which was 2.2 µs at 110 °C, whereas from the relative amplitudes we could estimate the orientation of the N–H bonds with respect to the averaged EFG principal axes system. It was found that at 25 °C the averaged EFG tensor orientation coincides with the static picture, identified by QC calculations, where the z EFG component, aligned with the lone pair, is perpendicular to the BDC-NH₂ unit and the dynamics is

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**Fig. 6** Principal components of the ¹⁴N EFG tensor (blue lines) in two different local structures of the BDC-NH₂ unit of NH₂-MIL-125. (a) Planar unit extracted from the MOF structure optimized at the QC level. (b) Deformed unit, extracted from the MD trajectory at 110 °C, where the nitrogen is pyramidalized and rotated (before the exchange of the NH₂ hydrogens). The plane of the three N substituents is depicted with a cyan disk (on the right-hand side, the transparent effect mitigates the color of the disk) to highlight the perpendicular orientation of Vₓᵧ (lone pair direction). The representation of EFG tensors as surfaces obtained by means of TensorView is shown in Fig. S5 of the ESI†.
quite restricted. This was not the case at 110 °C, where larger fast fluctuations modulate the static interaction and are responsible for a change in the orientation of the averaged EFG.

Besides the structural and dynamic details obtained, this study demonstrates the applicability and usefulness of FFC NMR, hitherto applied to aqueous solutions, soft matter or fluids in porous materials, also in the study of MOFs, unveiling information of relevance for their technological application. Furthermore, the combination with theoretical calculations, on the one hand, guides the data analysis, and, on the other, validates and at the same time complements the information obtained.

Conflicts of interest
There are no conflicts of interest to declare.

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Notes and references
1 R.-B. Lin, S. Xiang, H. Xing, W. Zhou and B. Chen, Coord. Chem. Rev., 2019, 378, 87–103.
2 H. Li, K. Wang, Y. Sun, C. T. Lollar, J. Li and H.-C. Zhou, Mater. Today, 2018, 21, 108–121.
3 V. Shrivastav, S. Sundriyal, P. Goel, H. Kaur, S. K. Tuteja, K. Vikrant, K.-H. Kim, U. K. Tiwari and A. Deep, Coord. Chem. Rev., 2019, 393, 48–78.
4 C. Xu, R. Fang, R. Luque, L. Chen and Y. Li, Coord. Chem. Rev., 2019, 388, 268–292.
5 M. F. de Lange, K. J. F. M. Verouden, T. J. H. Vlugt, J. Gascon and F. Kapteijn, Chem. Rev., 2015, 115, 12205–12250.
6 S. K. Henninger, F. Jeremias, H. Kummer and C. Janiak, Eur. J. Inorg. Chem., 2012, 2625–2634.
7 S. Cui, M. Qin, A. Marandi, V. Steggles, S. Wang, X. Feng, F. Nouar and C. Serre, Sci. Rep., 2018, 8, 15284.
8 H. Furukawa, K. E. Cordova, M. O’Keeffe and O. M. Yaghi, Science, 2013, 341, 1230444.
9 C. Zlotea, D. Phanon, M. Mazaj, D. Heurtaux, V. Guillerm, C. Serre, P. Horcajada, T. Devic, E. Magnier and F. Cuevas, et al., Dalton Trans., 2011, 40, 4879–4881.
10 M. Dan-Hardi, C. Serre, T. Frot, L. Rozes, G. Maurin, C. Sanchez and G. Férey, J. Am. Chem. Soc., 2009, 131, 10857–10859.
11 S. Vaesen, V. Guillerm, Q. Yang, A. D. Wiersum, B. Marszalek, B. Gil, A. Vinmont, M. Daturi, T. Devic, P. L. Llewellyn, C. Serre, G. Maurin and G. De Weireld, Chem. Commun., 2013, 49, 10082–10084.
12 F. Vermoortele, M. Maes, P. Z. Moghadam, M. J. Lennox, F. Ragon, M. Boulhout, S. Biswas, K. G. M. Laurier, I. Beurroles, R. Denoyel, M. Roëffaers, N. Stock, T. Düren, C. Serre and D. E. De Vos, J. Am. Chem. Soc., 2011, 133, 18526–18529.
13 F. Jeremias, V. Lozan, S. K. Henninger and C. Janiak, Dalton Trans., 2013, 42, 15967–15973.
14 L. G. Gordeeva, M. V. Solovyeva and Y. I. Aristov, Energy, 2016, 100, 18–24.
15 W. Morris, R. E. Taylor, C. Dybowski, O. M. Yaghi and M. A. Garcia-Garibay, J. Mol. Struct., 2011, 1004, 94–101.
16 S. Devautour-Vinot, G. Maurin, C. Serre, P. Horcajada, D. P. da Cunha, V. Guillerm, E. de Souza Costa, F. Taulelle and C. Martineau, Chem. Mater., 2012, 24, 2168–2177.
17 W. Masierak, M. Florek-Wojciechowska, I. Ogłodék, R. Jakubas, A. F. Privalov, B. Kresse, F. Fujara and D. Kruk, J. Chem. Phys., 2015, 142, 204503.
18 M. Florek-Wojciechowska, M. Wojciechowski, R. Jakubas, Sz. Brym and D. Kruk, J. Chem. Phys., 2016, 144, 054501.
19 M. Florek-Wojciechowska, R. Jakubas and D. Kruk, Phys. Chem. Chem. Phys., 2017, 19, 11197–11205.
20 F. Winter and R. Kimmich, Biochim. Biophys. Acta, 1982, 719, 292–298.
21 E. P. Sunde and B. Halle, J. Magn. Reson., 2010, 203, 257–273.
22 D. Kruk, E. Masiewicz, A. M. Borkowska, P. Rochowski, P. H. Fries, L. M. Broche and D. J. Lurie, Biomolecules, 2019, 9, 652.
23 P. H. Fries and E. Belorizky, J. Chem. Phys., 2015, 143, 044202.
24 A. E. Khudozhnikov, S. S. Arzumanov, D. I. Kolokolov, D. Freude and A. G. Stepanov, Phys. Chem. Chem. Phys., 2020, 22, 5976–5984.
25 D. I. Kolokolov, A. G. Stepanov and H. Jobic, J. Phys. Chem. C, 2015, 119, 27512–27520.
26 A. Knebel, B. Geppert, K. Volgmann, D. I. Kolokolov, A. G. Stepanov, J. Twiefel, P. Heitjans, D. Volkmer and J. Caro, Science, 2017, 358, 347–351.
27 D. I. Kolokolov, H. Jobic, A. G. Stepanov, V. Guillerm, T. Devic, C. Serre and G. Férey, Angew. Chem., Int. Ed., 2010, 49, 4791–4794.
28 D. I. Kolokolov, A. G. Stepanov and H. Jobic, J. Phys. Chem. C, 2014, 118, 15978–15984.
29 B.-J. van Rossum, C. P. de Groot, V. Ladizhansky, S. Vega and H. J. M. de Groot, J. Am. Chem. Soc., 2000, 122, 3465–3472.
30 M. Hong, X. Yao, K. Jakes and D. Huster, J. Phys. Chem. B, 2002, 106, 7355–7364.
31 S. L. Gould, D. Tranchemontagne, O. M. Yaghi and M. A. Garcia-Garibay, J. Am. Chem. Soc., 2008, 130, 3246–3247.
32 J. Gonzalez, R. N. Devi, D. P. Tunstall, P. A. Cox and P. A. Wright, Microporous Mesoporous Mater., 2005, 84, 97–104.
33 D. Kruk, A. Herrmann and E. A. Rössler, Prog. Nucl. Magn. Reson. Spectrosc., 2012, 63, 33–64.
34 Origin(Pro), Version 8.6.0. OriginLab Corporation, Northampton, MA, USA.
35 Wolfram Research, Inc. MATHEMATICA, (Version 10.0), Wolfram Research Inc., Champaign, IL, 2014.
36 A. P. Smalley, D. G. Reid, J. C. Tan and G. O. Lloyd, CrystEngComm, 2013, 15, 9368–9371.
37 P. Giannozzi, S. Baroni, N. Bonini, M. Calandra, R. Car, C. Cavazzoni, D. Ceresoli, G. L. Chiarotti, M. Cococcioni and I. Dabo, et al., J. Phys.: Condens. Matter, 2009, 21, 395502.
38 J. P. Perdew, K. Burke and M. Ernzerhof, Phys. Rev. Lett., 1996, 77, 3865–3868.
39 C. J. Pickard and F. Mauri, Phys. Rev. B: Condens. Matter Mater. Phys., 2001, 63, 245101.
40 J. R. Yates, C. J. Pickard and F. Mauri, Phys. Rev. B: Condens. Matter Mater. Phys., 2007, 76, 024401.
41 S. Monti, A. C. T. van Duin, S.-Y. Kim and V. Barone, J. Phys. Chem. C, 2012, 116, 5141–5150.
42 H. J. C. Berendsen, J. P. M. Postma, W. F. van Gunsteren, A. DiNola and J. R. Haak, J. Chem. Phys., 1984, 81, 3684–3690.
43 D. A. d’Avignon and T. L. Brown, J. Phys. Chem., 1981, 85, 4073–4079.
44 W. C. Bailey, Chem. Phys., 2000, 252, 57–66.
45 A. Carrington and A. D. McLachlan, Introduction to Magnetic Resonance: With Applications to Chemistry and Chemical Physics, Chapman & Hall, London, UK, 1979.
46 J. Autschbach, S. Zheng and R. W. Schurko, Concepts Magn. Reson., 2010, 36a, 84–126.
47 B. Kleibömer and D. H. Sutter, Z. Naturforsch., 1988, 43a, 561–571.