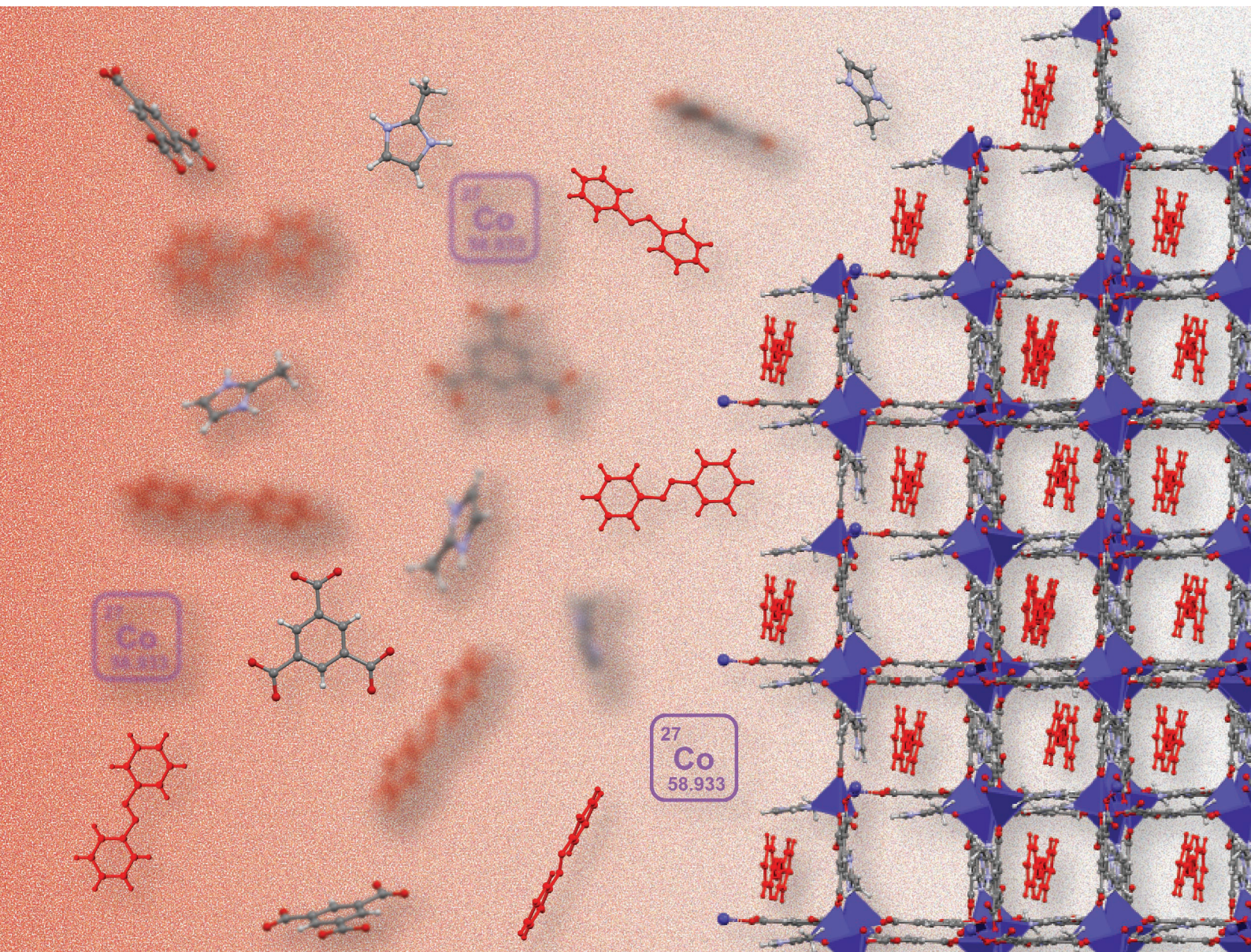


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Neutral and ionic Co(II) metal–organic frameworks with 2-methylimidazole and trimesate: design and evaluation for molecule encapsulation and slow release†

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Two Co(II) mixed-ligand metal–organic frameworks (MOFs) based on 2-methylimidazole and trimesate were synthesised at room temperature. The structure and properties of the two MOFs, named material Deutsches Elektronen Synchrotron-1 and -2 (mDESY-1 and mDESY-2), were verified by single crystal X-ray diffraction (SCXRD), powder X-ray diffraction (PXRD), SQUID magnetic susceptibility and N₂ adsorption. The structural analysis indicates that mDESY-1 is a 3D ionic framework with 2-methyl-1*H*-imidazol-3-ium counterions residing in its pores, while mDESY-2 is a 2D neutral framework isostructural to ITH-1, with water as a co-crystallising solvent. PXRD data demonstrates that mDESY-1 exhibits better crystallinity than mDESY-2. Magnetic measurements indicate that both MOFs are paramagnetic with a weak ferromagnetic transition above room temperature. Although both structures suggest the presence of voids, N₂ adsorption data confirms that these voids are not accessible in either MOF. Nevertheless, mDESY-1 was capable of encapsulating azobenzene during synthesis, which was observed *via* SCXRD. The encapsulated molecules were then slowly released in ethanol, with a release of up to 30 mg of azobenzene per g of MOF in a period of 60 days.

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Introduction

Metal–organic frameworks are porous materials constructed from metal ions or clusters and organic bridging ligands.^{1,2} The ability to fine-tune their properties by varying metal ions, ligands and synthesis conditions have made MOFs significantly attractive for a wide range of applications such as carbon capture,³ gas storage and separation,⁴ heterogeneous catalysis,⁵ luminescence,⁶ chemical sensing,⁷ biological imaging,⁸ and drug delivery.⁹

Among the various combinations of MOFs, the ionic MOFs (iMOFs) subclass remains relatively unexplored with just over 1000 publications as of 2021.¹⁰ Ionic MOFs are charged frameworks (either positive or negative) containing uncoordinated counter ions inside the framework, which maintain the overall charge neutrality. This subclass can be further divided into cationic MOFs (iMOFs-C) and anionic MOFs (iMOFs-A). While iMOFs-C bear free counter anions inside their pores, iMOFs-A contain free counter cations.¹¹ The ions inside the charged frameworks are often exchanged with other exogenous ions, which makes them ideal candidates for application in ion exchange resins.^{11–13}

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The selection of metal ions/cluster, ligands and counter ions is crucial for developing new iMOFs.¹¹ Imidazolium derivatives and benzene 1,3,5-tricarboxylic acid (H_3btc) are frequently used as ligands for the synthesis of ionic frameworks.^{10,11,14,15} For example, Hupp *et al.* utilised 1,3-bis(2,6-dimethyl-3,5-dicarboxylphenyl)imidazolium as a ligand to synthesise the cationic framework $[Cu_2(C_{23}H_{17}N_2O_4)(DMSO)_2][Cl]$,¹⁶ while Wang *et al.* used H_3btc to develop a 12-connected anionic MOF with the formula $H_2Na_4[Cu_{12}(OH)_6(pz^-)_6(btc)_6] \cdot 23H_2O$ (pz^- = pyrazolate).¹⁷ However, a 3D iMOF-A based on the common ligands 2-methylimidazole (mIm) and benzene 1,3,5-tricarboxylic acid with 2-methylimidazolium (HmIm) counter ions has not yet been reported.

Benzene 1,3,5-tricarboxylic acid, also known as trimesic acid, and 2-methylimidazole are two well-known organic compounds used in the synthesis of MOFs and other small molecules. For instance, mIm is employed in the synthesis of zeolitic imidazolate frameworks (ZIFs), such as ZIF-8¹⁸ and ZIF-67,¹⁹ while H_3btc is used in the production of HKUST-1.²⁰ The ligands have also been combined with zinc ions to synthesise a 2D mixed-ligand MOF named ITH-1.²¹ This MOF possesses a neutral network with linking btc linkers and mono coordinated mIm moieties as pendants throughout the framework. In the case of small molecules, we have previously demonstrated that the protonated HmIm and the deprotonated btc can form a hexaaquacobalt-based complex with 1*H*-imidazol-3-ium cations when mixed with $CoCl_2 \cdot 6H_2O$ in EtOH : water : DMF solution.²² Additionally, we have demonstrated that the organic ligands can form a crystal structure containing 1*H*-imidazol-3-ium as cation and the singly deprotonated 3,5-dicarboxybenzoate as anion.²³

The combination of imidazoles and polycarboxylates with metal salts to produce MOFs and coordination polymers is a well-established strategy, with many compounds synthesised through this approach.^{24–36} For example, Zhu and *et al.* reported the synthesis of a mixed-ligand MOF $[Co_7(bdc)_6(nIm^-)_2(H_2O)_6] \cdot 3H_2O \cdot 3DFM$, (H_2bdc = 1,4-benzenedicarboxylic acid, nIm^- = 2-nitroimidazole), which formed a 3D framework with polyhedral cages measuring up to 31.019 Å in diagonal length, and exhibited a CO_2 uptake capacity of 14.5 wt% at 273 K and 1 atm.³⁴ Another example has been provided by Kirillov *et al.*, who synthesised the 3D network $[Zn_2(\mu_4-Clta)(\mu_4-biim^{2-})]_n$ and the 1D coordination polymer $[Cd(\mu-Clta)(\mu_4-biim)]_n$, (H_2Clta = 2-chloroterephthalic acid, $biim$ = 2,2'-biimidazole), both of which showed catalytic potential in the coupling of CO_2 with epoxides.³⁵ Finally, Balkus and *et al.* developed two 2D mixed-linker MOFs by incorporating a secondary linker (1,4-benzenedicarboxylic acid or 2,6-naphthalenedicarboxylic acid) into the 1D coordination polymer $Cu(L1^-)(Cl)_2$, ($L1$ = 2,2'-(1*H*-imidazole-4,5-diyl)di-1,4,5,6-tetrahydropyrimidine). In these compounds, hydrogen bonding between two different linkers promotes layer stacking, resulting in a 3D structure with internal pore channels.³⁶ While these materials exemplify the versatility of the mixed-ligand strategy in constructing a variety of metal–organic structures, none of these

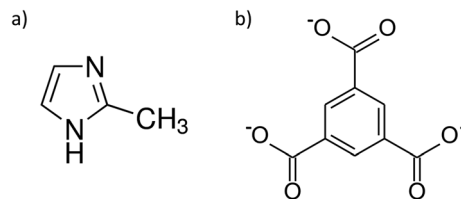


Fig. 1 Lewis structure for (a) mIm and (b) btc^- ligands.

examples form an iMOF, highlighting the rarity of this type of material.

Here, we present two Co^{II} mixed-ligand MOFs using the well-known mIm and btc ligands (Fig. 1). The structural analysis demonstrates that the iMOF designated as mDES-1 forms an anionic 3D framework with btc linkers, mono coordinated mIm as pendants and HmIm as counterions residing in the channels of the framework. On the contrary, the neutral MOF, named mDES-2 (also Co-ITH-1), is a 2D network isostructural to the previously reported ITH-1.²¹ Adsorption measurements indicate that both frameworks are non-porous. However, azobenzene molecules can be captured inside the channels of mDES-1 during synthesis, which can later be slowly released in ethanol.

Synthesis and methods

Materials

Ethanol ($\geq 99.8\%$), mIm (99%) and H_3btc (95%) were purchased from Sigma-Aldrich, while $CoCl_2 \cdot 6H_2O$ (98%) and azobenzene ($>97\%$) were bought from Alfa Aesar. The chemicals were used as received.

Synthesis of MOFs

mDES-1. In the original procedure, 2 ml of a 0.11 M ethanolic solution of $CoCl_2 \cdot 6H_2O$ was added to a flask containing 60 ml of ethanol. Then, 0.8 ml of a 1.58 M ethanolic solution of mIm and 2 ml of a 0.12 M solution of H_3btc in ethanol were added to the mixture. A purple precipitate was formed immediately. After resting the solution for a week, purple crystals of mDES-1 were obtained. The product was collected by filtration and washed 3 times in ethanol.

mDES-2. The initial mixture was prepared by fully dissolving 12.7 mg of $CoCl_2 \cdot 6H_2O$ (0.095 mmol) and 64.9 mg of HmIm in 2 ml of ethanol. Subsequently, 1.5 ml of a 0.01 M ethanolic solution of H_3btc was added to the mixture under constant stirring. Within minutes, small purple crystals of mDES-2 were obtained. The product was recovered by filtration.

Azobenzene@mDES-1 (Azo@mDES-1). The procedure to encapsulate azobenzene in mDES-1 followed a similar approach to that described for mDES-1. Typically, 2 ml of a 0.11 M ethanolic solution of $CoCl_2 \cdot 6H_2O$ was added to 60 ml of a 0.06 M solution of azobenzene in ethanol. Subsequently, 0.8 ml of a 1.58 M ethanolic solution of mIm and 2 ml of a



0.12 M solution of H₃btc in ethanol were added to the mixture. After one-week, dark purple crystals were formed. The product was separated by filtration and briefly washed with ethanol once.

Characterisation

Single crystal X-ray diffraction (SCXRD). The SCXRD experiments for mDES_Y-1 and mDES_Y-2 were performed at 100 K. The X-ray data was collected using undulator synchrotron radiation with $\lambda = 0.6199$ Å at the P11 beamline of PETRA III, DESY, Hamburg, Germany. An Eiger 2 × 16 M detector was positioned 154 mm from the sample. The data frames were collected in φ scan mode with a step size of $\Delta\varphi = 0.1^\circ$ and an exposure time of 10 ms per frame. Indexing of the X-ray diffraction pattern, unit cell refinement and spot integration were performed using the XDS program package.³⁷ Attempts to collect crystallographic data of Azo@mDES_Y-1 at 100 K using synchrotron radiation were unsuccessful since an unknown phenomenon was observed (see Fig. S1†).

The crystal structures of mDES_Y-1, azobenzene and Azo@mDES_Y-1 were determined at room temperature (~290 K) using a single-crystal diffractometer at DESY, Hamburg, Germany. The crystals were mounted on an APEX Duo Bruker diffractometer equipped with a monochromatic X-ray source (Mo K α , $\lambda = 0.71073$ Å) and an APEX CCD detector. Data frames were collected through multiple φ and ω scans of crystal orientations, with a step size of 0.5° and an exposure time of 15 seconds per frame for azobenzene and mDES_Y-1. The encapsulation of azobenzene in mDES_Y-1 reduced the crystallinity of the MOF, and therefore, the exposure time was increased to 60 seconds per frame to obtain a maximum resolution of 0.88 Å. The sample-to-detector distance was maintained at 50 mm. Data reduction was performed using the SAINT program.³⁸

The crystal structures of all collected data were solved with the SHELXT³⁹ structure solution program within the Olex2 software package.⁴⁰ Subsequently, the structures were refined against F^2 isotropically, followed by full matrix anisotropic least squares refinement by SHELXL.⁴¹ In the final cycle of the refinement, hydrogen atoms were fixed geometrically in idealised positions and allowed to ride with the respective C or N atoms to which each was bonded. The solvent mask routine incorporated in Olex2 was used to remove the disordered HmIm cations in mDES_Y-1.

Figures of the asymmetric unit and packing of the mDES_Y-1 with refinement of one counter ion laying in a symmetry element are provided in Fig. S2.† Crystal data and structure refinement parameters for mDES_Y-1 (after solvent mask) and mDES_Y-2 at 100 K are provided in Table S1,† while the corresponding data for mDES_Y-1, azobenzene and Azo@mDES_Y-1 at room temperature (RT) is presented in Table S2.† Ortep plots for all structures are available in ESI (ESI, Fig. S3–S7†) along with images of mDES_Y-1 and mDES_Y-2 view down all axes (Fig. S8 and S9†). Selected bond lengths and angles for all structures in this work are reported in Tables S3 and S4.†

PXRD. X-ray diffraction (XRD) measurements on mDES_Y-2 and Azo@mDES_Y-1, both before and after release of azobenzene, were taken on a PXRD – KIBS Bruker D8 Discover XRD with a Cu K α radiation ($\lambda = 1.5406$ Å). The tube operating mode was $U = 40$ kV, $I = 40$ mA. The range of diffraction angles 2θ varied from 5° to 80° (0.01° step); the exposure per shooting point was 1 s.

PXRD data for mDES_Y-1 sample was collected at P02.1 beamline, PETRA III.⁴² Crystals of the sample were grinded in agate mortar under solution, transferred together with the solution into 0.8 mm in diameter special glass capillary, and held vertically to let the powder sediment to increase the density of powder at the measurement point. Data was collected for capillary sput at 5000 Hz for 180 s using the VAREX CT 4343 detector (150×150 µm pixel size and 2880×2880 pixel area). The wavelength was 0.20734 Å and the detector configuration was calibrated using the LaB₆ SRM 660c standard. The calibration and integration were performed using the pyFAI software.⁴³ The LeBail fit of the data was performed using the FullProf software.⁴⁴

Magnetic measurements. The magnetic measurements were taken in a Direct Current-Magnetic Properties Measurement System (DC-MPMS) with SQUID sensor. The magnetisation was collected in Field Cooling (FC) and Field Heating (FH) modes under applied magnetic fields 0.1 and 0.2 T. Magnetic measurements at 0.1 T were performed by decreasing the temperature to 3 K in zero field cooling (ZFC) mode. The data was collected by gradually increasing the temperature from 3 to 400 K (FH mode) and then decreasing it from 400 to 3 K (FC mode). In order to check the magnetic behaviour of the samples, the measurement was repeated, this time under 0.2 T applied field and by changing the range of temperature from 282 to 3 K in FC mode, then from 3 to 350 K in FH mode and from 350 to 200 K in FC mode.

Adsorption measurements

Nitrogen physisorption measurements were performed using a 3Flex Analyzer (Micromeritics Instrument Corp.). Prior to the measurements, the samples were activated at 120 °C for 20 h under reduced pressure using a Smart VacPrep Degasser (Micromeritics Instrument Corp.). For data reduction, the software MicroActive, Version 6.00, was used. The Brunauer–Emmett–Teller (BET) method was used to determine the specific surface area (p/p_0 range from 0.1 to 0.3) as well as the total pore volume at 0.99 p/p_0 .

Release of azobenzene

An 8 ml vial containing 10 mg of Azo-mDES_Y-1 was filled with 6 ml of ethanol. The resulting heterogeneous mixture was left in the vial for a period of 2 months. UV-visible (UV-vis) spectroscopy was used to analyse the solution at specific time intervals: 1, 3, 7 and 10 hours after the addition of ethanol, and subsequently after 1, 3, 7, 10, 30 and 60 days. In each measurement, 2 ml aliquots were taken from the vial and placed in a cuvette of $12.5 \times 12.5 \times 45$ mm dimensions. The aliquots were returned to the vial after each measurement. The experiments



were conducted in triplicate. UV-vis spectra at room temperature were collected on a Varian Cary-5E UV-vis Spectrometer. All UV-vis data was collected at a scan rate of 0.2 nm s^{-1} over the wavelength range of 200–800 nm with an interval size of 0.5 nm.

Results

Structural information

The crystal structures for mDESY-1 and mDESY-2 are quite different. mDESY-1 crystallises in the tetragonal space group $I4_1/a$; the asymmetric unit contains one cobalt(II) ion, one btc ligand and one disordered mIm, which is split into two sites. The Co(II) ion adopts a tetrahedral geometry, defined by one $N_{\text{imidazole}}$ and three $O_{\text{carboxylate}}$ atoms (Fig. 2a). Based on these connections, Co(II) ions are bridged by btc ligands to generate an anionic 3D framework with square channels along the c axis (Fig. 2b). However, the counter ion HmIm resides disorderly inside of the channels, obstructing the pores (Fig. S2†). Topological features of mDESY-1 were obtained by means of the ToposPro⁴⁵ software. The analysis reveals 3-c deh net for mDESY-1 (point symbol = $8^2.10$ and vertex symbol = $[8.8(2).12]$,

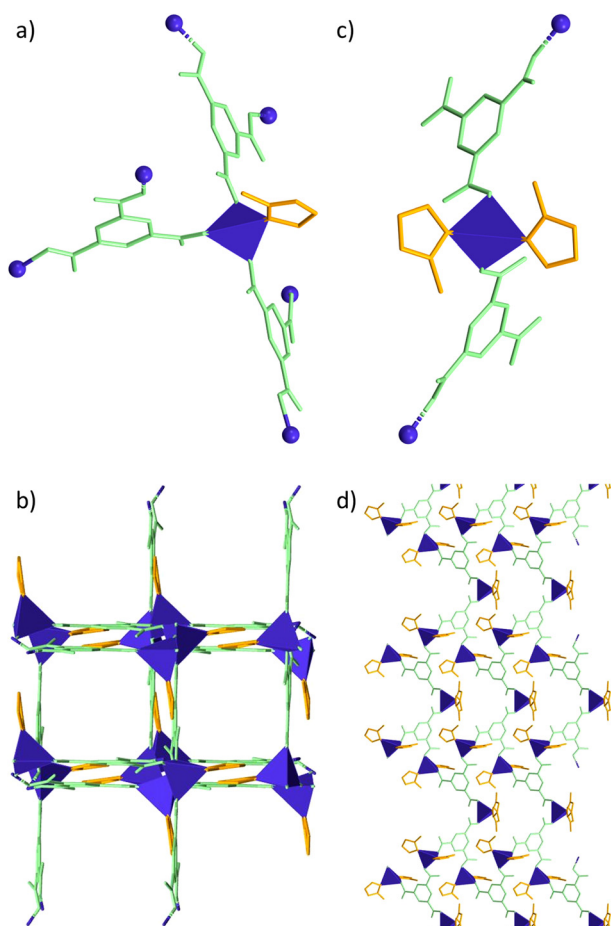


Fig. 2 Structure of mDESY-1 and mDESY-2 (Co-ITH-1). Hydrogen atoms, water and disorder molecules were omitted for clarity.

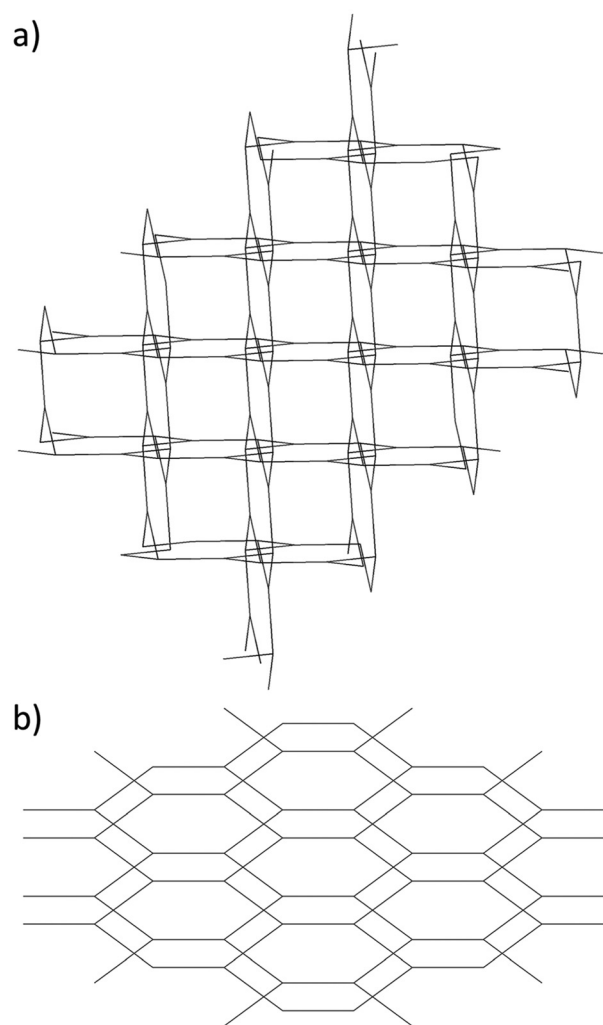


Fig. 3 Underlying nets of (a) mDESY-1 and (b) mDESY-2 calculated by the ToposPro software.

Fig. 3a). On the other hand, mDESY-2 crystallises in the $C2/c$ space group with an asymmetric unit formed by two crystallographically independent Co atoms (Co1 and Co2). Each Co atom is tetrahedrally coordinated to two nitrogen atoms from the mIm ligands and two oxygen atoms from btc linkers (Fig. 2c). A non-coordinated water molecule resides within the lattice. The compound forms neutral 2D layers parallel to the (010) plane (Fig. 2d). These layers are interconnected *via* hydrogen bonds $O_{\text{water}}-\text{H}\cdots O_{\text{carboxylate}}$, and $N_{\text{imidazole}}-\text{H}\cdots O_{\text{carboxylate}}$. Topological features for mDESY-2 obtained by the ToposPro software show a 3-c hcb topology with point symbol = 6^3 and vertex symbol = $[6.6.6]$ (Fig. 3b).

The distortion of the coordination polyhedra in both MOFs was evaluated according to the Avnir^{46,47} method, based on the continuous shape measure (CShM). The SHAPE⁴⁸ program was employed for this analysis, and the results have been summarised in Table 1. A CShM value of 0 corresponds to a perfect shape match, indicating no distortion from the ideal geometry. As the value increased, the degree of distortion or deviation



Table 1 Geometrical distortion of the cobalt ion coordination environment of all networks, calculated by SHAPE software and by τ_4 the method

Ion	mDESYS-1	mDESYS-2	
	Co(1)	Co(1)	Co(2)
Coordination	4	4	4
S(SP)	29.647–29.856	26.901	27.948
S(T)	1.161–1.400	0.842	0.361
τ_4	0.828	0.887	0.905

from the ideal shape also increased. The CShM values for mDESYS-1 and mDESYS-2 showed a slight deviation from the ideal tetrahedral shape [$S(T) = 0$] and significantly larger deviation from the ideal square planar geometry [$S(SP) = 0$]. The greatest distortion in the coordination environment of the Co(II) ion was observed in mDESYS-1, with $S(T)$ values ranging from 1.161 to 1.400.

For comparison, the τ_4 parameter was also employed to assess the distortion of the coordination polyhedra in both MOFs. This is a very simple geometry index for four-coordinate complexes, defined as:⁴⁹

$$\tau_4 = \frac{360^\circ - (\alpha + \beta)}{141^\circ}$$

where α and β are the two largest bond angles between adjacent ligands around the metal centre. The values of τ_4 can range from 1 for a perfect tetrahedral geometry [$360^\circ - 2(109.5^\circ) = 141^\circ$], to zero for a perfect square planar geometry [$360^\circ - 2(180^\circ) = 0^\circ$].

The τ_4 values were consistent with the CShM values, both indicating slight deviation from the ideal tetrahedral geometry (Table 1). The smallest τ_4 value was found in mDESYS-1 (0.828), indicating a larger distortion compared to the Co(II) centres in mDESYS-2, which exhibited τ_4 values closer to 1 (ranging from 0.887 to 0.905), suggesting a geometry closer to the perfect tetrahedron.

Fig. 4a shows the PXRD of mDESYS-1, alongside its calculated pattern obtained through Le Bail fit, based on the structural model of mDESYS-1 at 290 K (see Encapsulation of azobenzene section). The strong agreement between the observed data (black line) and the calculated pattern (red circles) demonstrates the reliability of the fit. The cell parameters obtained from the Le Bail fit (Table S5†) agrees with that observed by the structural model at 290 K (Table S2†). The final fitting parameters are $R_p = 9.08$, $R_{wp} = 8.12$, $R_{exp} = 4.22$ and $\chi^2 = 3.70$.

The difference curve reveals some unfitted peaks or changes in unit cell parameters, which confirm phase purity. These variations could result from minor impurities in the synthesis process or a decomposition product during capillary preparation for PXRD measurements. The fitted cell parameters (see Table S5†) are matching well to those from single crystal data of mDESYS-1 at 290 K.

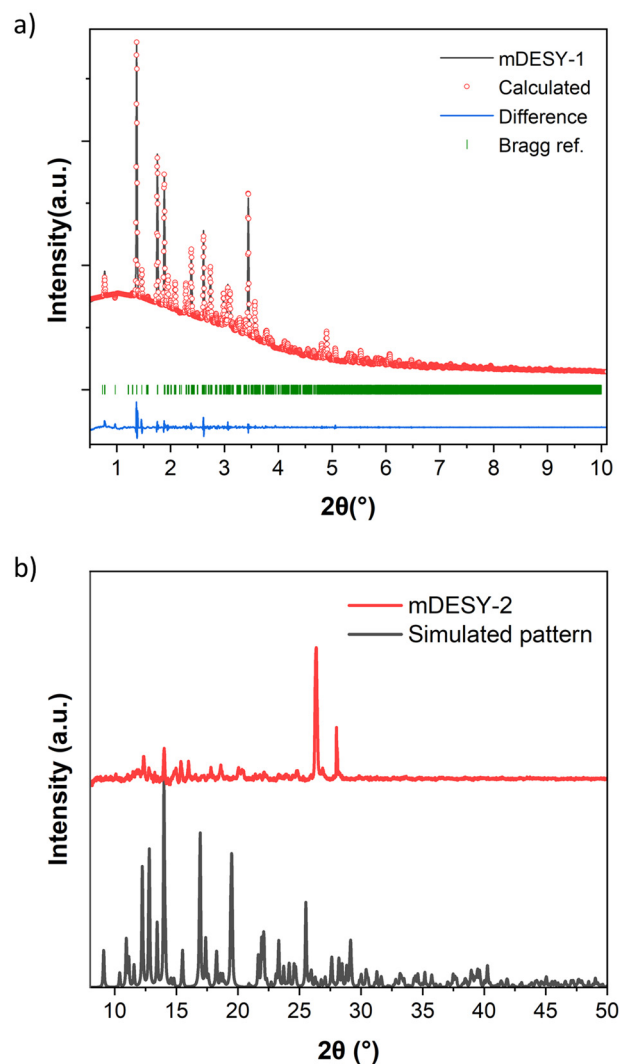


Fig. 4 (a) PXRD of mDESYS-1 and Le Bail fit of the structure model from single crystal data on PXRD data collected at P02.1, PETRA III ($\lambda = 0.20734$ Å), and (b) PXRD of mDESYS-2 collected in Bruker D8 Discover ($\lambda = 1.5406$ Å) alongside its corresponding simulated pattern generated by the software Mercury.

The PXRD pattern of mDESYS-2 and its simulated pattern generated from its crystal structure using the Mercury software^{50,51} is illustrated in Fig. 4b. In contrast to mDESYS-1, the fast precipitation of the mDESYS-2 framework results in a product with poor crystallinity. Additionally, a second phase is observed, as the prominent peaks between 2θ angles of 25° and 30° do not correspond to mDESYS-2. This suggests that further improvements in the synthesis are required to obtain a pure phase of mDESYS-2.

Adsorption properties

Crystallographic structures of mDESYS-1 and mDESYS-2 were analysed in the Mercury CSD software package using the void analysis tools to determine the limiting pore diameter.^{50,51} Disordered counter ions, if present in the pore, were removed



before void analysis. The grid spacing was set to 0.7 Å and the probe size to 1.2 Å. This probe size diameter corresponded to the largest sphere that could be inserted without overlapping the framework atoms.⁵²

Fig. 5a shows that mDESY-1 contains several interconnected square channels that run along all of the axes. The total calculated pore volume reaches 4303.85 Å³, which represents 45.6% of the unit cell volume. However, as observed by the structural analysis, the pores are blocked by the HmIm counter ions. Fig. 5b demonstrates that mDESY-2 contains three nanopores per unit cell; the pores have a volume of 260.09 Å³, which represent 5.1% of the unit cell volume.

The N₂ adsorption-desorption isotherms for both MOFs are shown in Fig. 6. The calculated BET surface areas are 2.2 m² g⁻¹ for mDESY-1 and 9.4 m² g⁻¹ for mDESY-2. These low adsorption capacities of both MOFs suggest that the voids calculated from the structural models are not accessible. For mDESY-1, the large channels found in the structure are blocked by the HmIm counter ion, while in mDESY-2, the small pores do not have possible access.

Magnetic properties

Fig. S10(a) and (b)† shows the temperature dependence of the magnetic moment for mDESY-1 and mDESY-2. In both cases, the signals are indicative of paramagnetic materials with the magnetic moment decreasing as temperature increases due to thermal vibration. Fig. S10(c) and (d)† displays the χ_M^{-1} vs T plots, where the linear response, in both FH and FC modes, intersects the origin (0,0), confirming their paramagnetic behaviour. However, above 340 K for mDESY-1 and 350 K for mDESY-2, the slopes of these linear responses change, suggesting a ferromagnetic rearrangement of the spins in both MOFs. By extrapolating the linear behaviour to the zero level, Curie temperatures (TC) of 110 K mDESY-1 and 52 K for

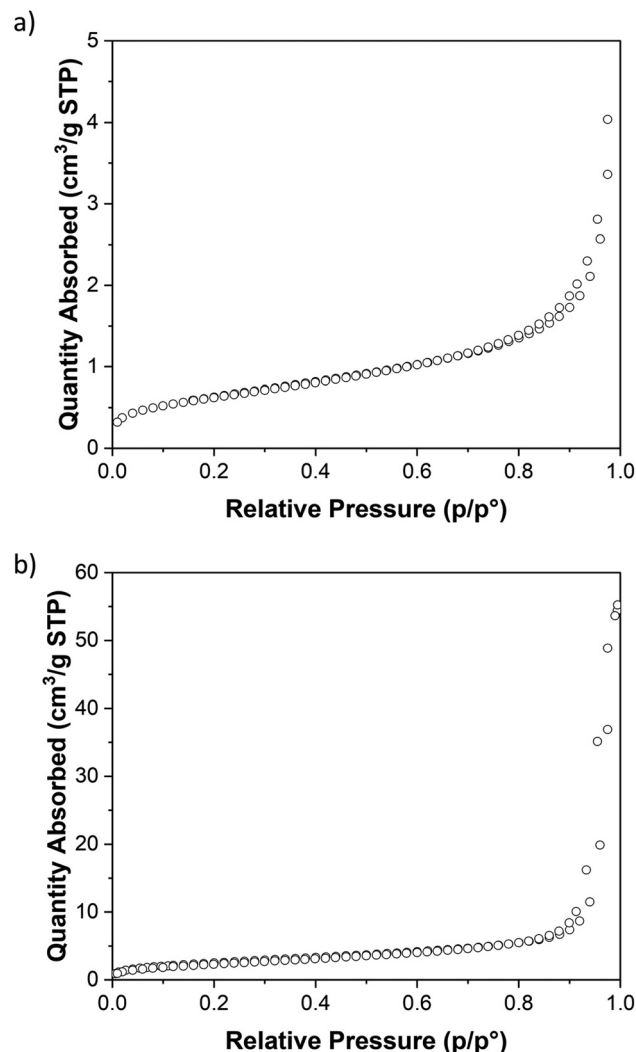


Fig. 6 N₂ adsorption measurements at 77 K of (a) mDESY-1 and (b) mDESY-2.

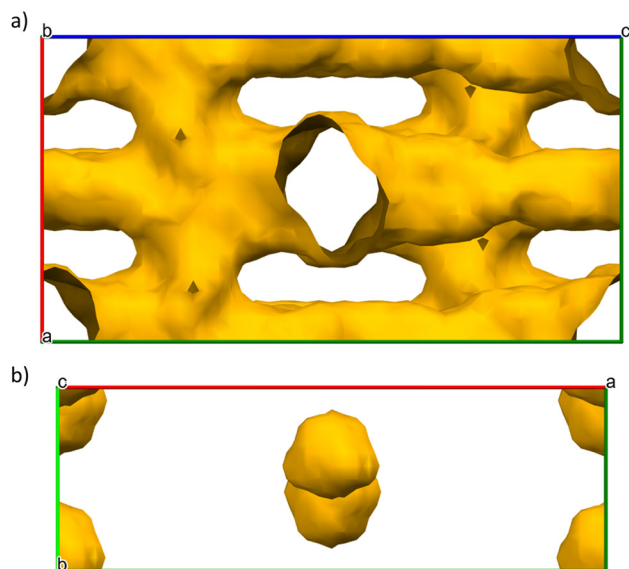


Fig. 5 Calculated voids in (a) mDESY-1 and (b) mDESY-2.

mDESY-2 were determined. Additionally, $\chi_M T$ vs T plots (Fig. S10e and f)† were constructed to better identify any irreversibility in the FH and FC loops. These plots indicate a weak ferromagnetic transition above 300 K for mDESY-1 and 320 K for mDESY-2, possibly signalling the onset alignment of some spins in the sample. Remarkably, an irreversible behaviour was noted in the FC loop, possibly due to these aligned spins.

To verify the magnetic behaviour of the samples, the measurement was repeated under a 0.2 T applied field. The temperature range was varied from 282 to 3 K in FC mode, then from 3 to 350 K in FH mode and finally from 350 to 200 K in FC mode (Fig. S11a and b)†. Fig. S11(c)† confirms that mDESY-1 retained its ferromagnetic transition with a change in the TC to 83 K. On the contrary, the inset plot in Fig. S11(d)† reveals an absent ferromagnetic transition for mDESY-2. The $\chi_M T$ vs T plots were constructed again to better differentiate if the irreversibility and magnetic transition persisted (Fig. S11e and f)†. For mDESY-1, the magnetic transition



reappears around 300 K, confirming that although most of the spins in the sample were miss-oriented, some spins realigned at this temperature. In the case of mDESY-2, the ferromagnetic alignment persisted at 320 K, but the loop became reversible, indicating a lack of thermal coercivity and that the spins in the sample were primarily affected by thermal vibrations.

Encapsulation of azobenzene

The encapsulation of azobenzene in mDESY-1 was demonstrated by single crystal diffraction at room temperature. Upon incorporating azobenzene into mDESY-1, the structure could still be solved in the $I4_1/a$ space group, albeit with a significantly higher redundancy-independent R factor (R_{int}). Additionally, azobenzene molecules were not discernible. To solve this, the structure of Azo@mDESY-1 was solved in the $P\bar{1}$ space group. In this way, the azobenzene molecules were clearly identified within the pores of the MOF. Fig. 7 (a) and (b) displays the structure of the loaded mDESY-1 and its packing in a ball-and-stick style, with the counterions HmIm omitted for clarity. Fig. 7a reveals that the azobenzene molecules exhibit orientational disorder. This disorder is dynamic, consistent with observations from structural studies of free *trans*-azobenzene at room temperature.^{53–55} This dynamic disorder is attributed to the torsional vibration of the N–Ph bonds. Fig. 7b demonstrates that the azobenzene molecules are located within the channels of the MOF, with the longest axis of the azobenzene aligned along the channel direction. The azobenzene molecule is located near the walls of the channel but does not exhibit strong interactions with the metal ions or the framework's ligands. The Hirshfeld isosurface analysis demonstrates that the primary interaction of the azobenzene is with the closest HmIm counterion (Fig. 7c). Other interaction between azobenzene molecules and other counterions or residual linkers within the pores are not visible, as the solvent mask was used for molecules that could not be modelled during the structure solution and refinement process.

A comparative analysis of bond lengths was made between encapsulated *trans*-azobenzene and free *trans*-azobenzene in single crystal form. The structure of the free *trans*-azobenzene at room temperature was determined, revealing a $P2_1/c$ space group with two crystallographically independent molecules in the asymmetric unit. In agreement with previous results in literature,^{53–55} both crystallographically independent molecules lay at inversion centres and were nearly planar. Molecules at one of the crystal sites showed no disorder (site 1), while those at the other site (site 2) exhibited orientational dynamic disorder as previously mentioned (Fig. S6†).^{53–55} The degree of misorientation was determined to be 11.3%, which is slightly smaller than the reported value in literature (17–18%).^{53,55}

Table 2 compares the main bond lengths and angles of the encapsulated azobenzene and both crystallographically independent molecules of the free azobenzene. The table highlights both the differences and similarities between the free and the encapsulated forms. The encapsulated and free azo-

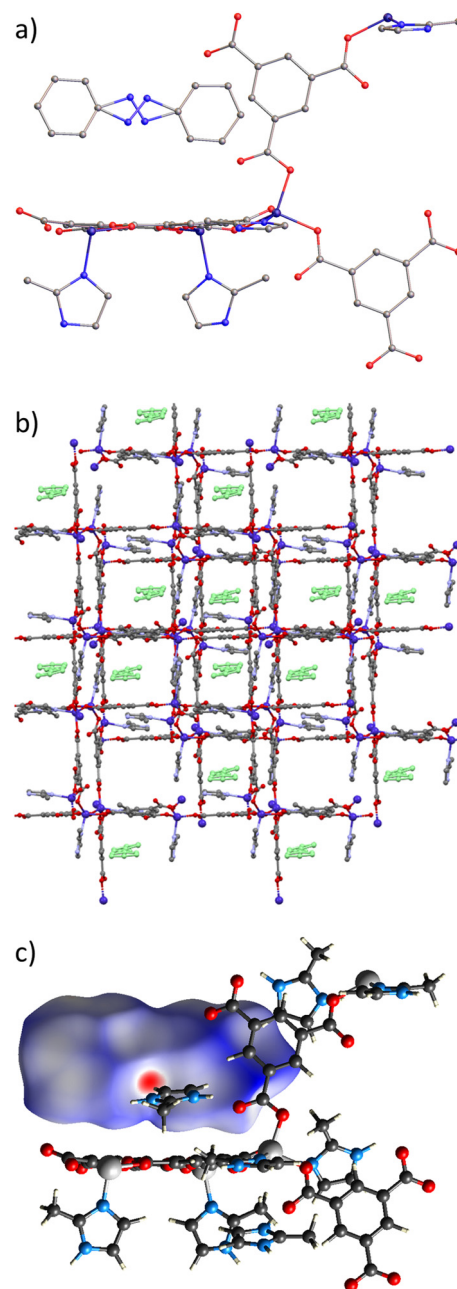


Fig. 7 (a) Structure of Azo@mDESY-1 at 290 K, counterions were omitted for clarity. (b) Packing of Azo@mDESY-1 at 290 K showing the position of the azobenzene molecule (green) in the channels of the MOF, counterions were omitted for clarity. (c) Hirshfeld isosurface of azobenzene inside of mDESY-1, showing the main interactions.

benzene exhibit similar benzene centroid-to-centroid distances. While the $-\text{C}=\text{N}=\text{N}-$ angles are shorter in the encapsulated azobenzene, this is compensated by longer $-\text{N}=\text{N}-$ bond lengths. Interestingly, the $-\text{C}-\text{N}=\text{N}-\text{C}-$ torsion angles and the angle between benzene rings indicate a non-parallel orientation of the rings in the encapsulated azobenzene, contrasting with the parallel arrangement observed in the free azobenzene molecules.



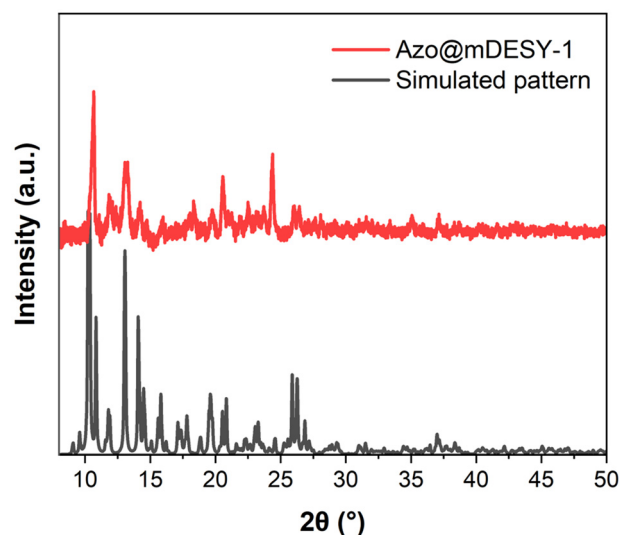
Table 2 Main distances and angles of the encapsulated and free azobenzene

	Free <i>trans</i> -azobenzene		Encapsulated <i>trans</i> -azobenzene
	Site 1	Site 2	
Benzene ring centroid to centroid distance (Å)	6.266	6.266	6.269
–N=N– (Å)	1.241 (2)	1.205 (3), 1.380 (2)	1.33 (2)
–C–N=N– (°)	114.0 (1)	112.1 (3), 105 (1)	105 (2)–106 (2)
–C–N=N–C– (°)	180.0 (1)	–180.0 (2), –180.0 (8)	–172 (2), –180 (2)
Angle between benzene rings (°)	0	0	9.91

As mentioned earlier, the best method to observe azobenzene molecules within the pores of Azo@mDES-1 involved solving the structure using a $P\bar{1}$ space group. To compare the contents and the structural change of the ionic framework with and without azobenzene, the structure of mDES-1 at room temperature was also solved using the $P\bar{1}$ space group, applying a solvent mask for the pore's contents that could not be modelled. The comparison of both structures revealed that more HmIm ions can be modelled in Azo@mDES-1 as the azobenzene molecules occupied the space between counterions, forcing more HmIm ion to align (Fig. S12†). In the structure of mDES-1, one HmIm ion was disordered across two positions, while in Azo@mDES-1, four HmIm ions were present, with two refined with 50% occupancy. The remaining 50% of those counter ions were disordered and suppressed by the solvent mask.

A comparative analysis of the metal ions' coordination environment shows a slight change in the framework. Table 3 indicates that while there is no significant difference in the average Co–N or Co–O bond lengths, the main difference lies in the distortion of the Co ion's coordination environment relative to a perfect tetrahedron. In Azo@mDES-1, this distortion is more pronounced, suggesting that the incorporation of azobenzene into the MOF induces a distortion in the overall network. This distortion also leads to a change in the unit cell dimensions, with the *a* axis of Azo-mDES-1 being 0.25 Å shorter than that of mDES-1 (Table S2†).

The PXRD pattern of Azo@mDES-1 (Fig. 8) shows some agreement with the calculated pattern obtained from the single crystal using Mercury software.^{50,51} However, Fig. 8 also highlights the sample's poor crystallinity. The inclusion of azobenzene during synthesis accelerates the compound's precipitation, which diminishes crystal quality. Additionally, the encapsulation of azobenzene distorts the network, which may introduce defects that further reduce the crystallinity. Efforts

**Fig. 8** PXRD pattern of Azo@mDES-1 alongside its simulation pattern.

to synthesise Azo@mDES-1 with lower amounts of azobenzene result in better crystal quality, but the azobenzene is not detectable in the crystal structure.

Attempts to capture the thermal isomerisation of the encapsulated azobenzene within Azo@mDES-1 using SCXRD were unsuccessful. However, optical images of the single crystal at different temperatures revealed a colour change with increasing temperature. To start, optical images of Azo@mDES-1 at room temperature (Fig. 9 a and b) were taken with the microscope in transmission mode. Unlike mDES-1 (Fig. S13a and b†), the colour of the Azo@mDES-1 crystal varied depending on its orientation, displaying a red hue in specific orientations. For temperature-dependent images, the crystal was placed on a slide over a heating plate, and the images were taken in reflection mode. Fig. 9c shows the Azo@mDES-1 crystal at room temperature (~17 °C), with the crystal exhibiting a purple colour when illuminated from above. After heating the crystal to 130 °C for 20 min to ensure the isomerisation of the azobenzene – which has been measured to occur around 68 °C (Fig. S14†) – the crystal changed colour to dark blue (Fig. 9d). This colour change was attributed to the isomerisation of azobenzene within the MOFs. After cooling the crystal to room temperature for 2 hours, it partially returned to its original colour (Fig. 9e). On the contrary, the optical images of

Table 3 Main bond length distances and distortion of the Co coordination

	mDES-1	Azo@mDES-1
⟨Co–N⟩ (Å)	2.019 (3)	2.022 (5)
⟨Co–O⟩ (Å)	1.982 (2)	1.983 (4)
S(SP)	30.140–30.266	28.459–30.368
S(T)	0.957–0.993	0.955–1.449
τ ₄	0.840–0.842	0.802–0.840



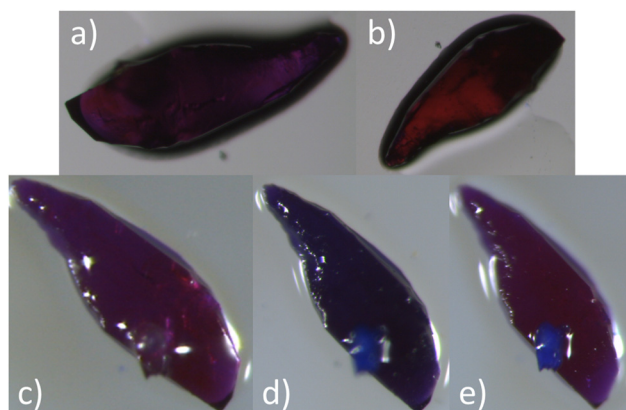


Fig. 9 (a) and (b) Optical images of Azo@mDESy-1 at room temperature at two different orientations in transmission mode. Optical images at different temperatures in reflectance mode: (c) room temperature, (d) 130 °C and (e) 2 h room temperature after heating at 130 °C.

mDESy-1, taken in reflection mode, showed a blue crystal at both room temperature and 130 °C, with the crystal degrading after more than one minute at 130 °C without any colour change (Fig. S13c–e†).

Release of azobenzene

The release of the encapsulated azobenzene from Azo@mDESy-1 was monitored over a two-month period. The UV-Vis spectra from one sample is shown in Fig. S15a,† while Fig. S15b and S15c† present the UV-Vis spectra and the corresponding calibration curve used for quantification. Fig. 10 illustrates the amount of azobenzene released per g of loaded MOF over the 2-month period. The data shows that approximately 30 mg of azobenzene per gram of Azo@mDESy-1 was released over this period. A linear fit of the first 10 hours of release in

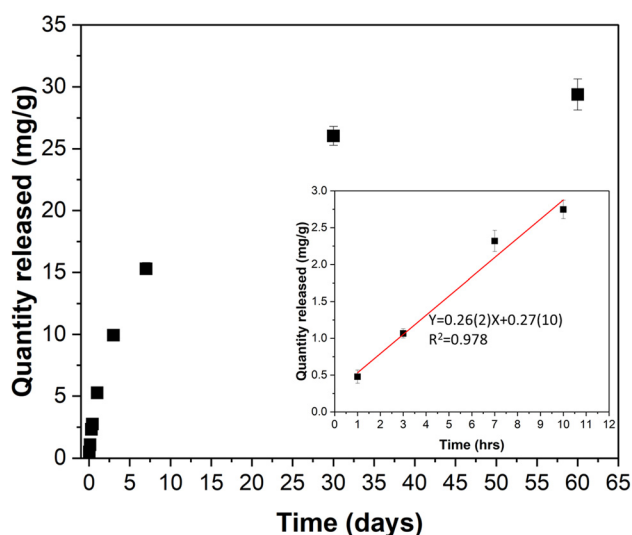


Fig. 10 Release of azobenzene from mDESy-1 in ethanol. Inset: Release of azobenzene during the first 10 h and its linear fitting.

ethanol indicates a delivery rate of 0.26 mg of azobenzene per hour per gram of Azo@mDESy-1.

The PXRD patterns of Azo@mDESy-1 before and after the release of azobenzene in ethanol are compared in Fig. S16a.† The PXRD patterns indicate that the sample retained some crystallinity, albeit poor, even prior to the release of the azobenzene. However, diffraction images from the SCXRD before and after azobenzene release (Fig. 16b and c†) exhibit a significant deterioration in the sample's quality following the release. This marked change suggests that the MOF is not suitable for reuse in subsequent applications after the release of the small molecules.

The crystallographic structure of Azo@mDESy-1 provide more information about the position of most of the cations within the framework. Therefore, the structural model of the Azo@mDESy-1 (excluding the azobenzene molecules) was examined using the void analysis tool in the Mercury CSD.

Fig. S17† shows the calculated voids in the structure, revealing constricted areas within the channel (red circle). These constrictions indicate a reduction in void space that impact the adsorption or release of molecules within the pores. Additionally, the total calculated pore volume is reduced to 1413.39 Å³, accounting for 29.4% of the unit cell volume, compared with the 45.6% obtained in the structural model of mDESy-1 without counterions.

The release of azobenzene requires that the molecule diffuse through the constricted channels of Azo@mDESy-1. This diffusion places considerable stress on the framework, leading structural damage on the crystal, as observed in the diffraction images.

Discussion

Metal–organic frameworks are among the most important classes of materials due to their exceptional tunability, making them promising candidates for a wide range of applications. However, most of the reported MOFs are electrically neutral, with only a small subset classified as ionic MOFs.¹⁰ This leaves significant room for exploration in the field of iMOFs, from the synthesis to potential application. While some strategies have been developed to obtain iMOFs, synthesising new iMOFs remains challenging and often occurs by chance.^{11,14} Therefore, it is crucial to report the synthesis, structure, properties and potential applications of new iMOFs, particularly those that are easily synthesised and do not require complex ligands.

The present anionic mDESy-1 is synthesised at room temperature, employing only ethanolic solutions containing the metal source and ligands. This iMOF also has the advantage of being synthesised from cobalt chloride, 2-methylimidazole and trimesic acid, which are well-known inexpensive chemicals. Although the synthesis is straightforward, varying the concentration of the reactants can lead to the formation of the neutral mDESy-2, which is isostructural to the reported ITH-1²¹ and may have similar applications in CO₂ fixation due



to the nitrogen-rich content. However, the potential application of mDES-1 extends beyond catalysis or the classical ion-exchange process, as it demonstrates capability of encapsulation and slow release of small molecules. This feature can be exploited for studying photoswitches in confined spaces, as well as for drug delivery.

Photoswitchable molecules are extensively studied because they can serve as building blocks for synthetic molecular machines. Molecular confinement enhances the interaction between photoswitches and their environment by physically trapping the molecules, thereby limiting their diffusion and constraining their movement.⁵⁶ However, these constraints can significantly reduce their switchability, as observed in some azo compounds within confined spaces.^{57,58}

Here, we have encapsulated azobenzene within mDES-1, and the change in colour observed through optical images at different temperatures suggests the possibility of thermal isomerisation occurring inside the iMOF. Hirshfeld isosurface analysis indicates that the primary interaction of the encapsulated molecule is with HmIm counterions, raising the possibility that the thermal or photoisomerisation of azobenzene could induce a molecular rearrangement of these counterions. Further investigation is needed to fully explore the thermal- and photoswitchability of azobenzene and its interaction with the environment within mDES-1. Nonetheless, this study underscores the utility of this ionic MOF.

Another potential application of mDES-1 could be in the biomedical field, although the cytotoxicity of the cobalt ion is an impairment for its application. In recent years, the application of MOFs in biomedicine has become a rapidly growing area of interest.^{59–61} Research into MOFs for biomedical application has included the use of iMOFs, as their charged frameworks allow for the encapsulation of ionic drugs.^{62,63} However, non-ionic drugs, such as 5-fluorouracil, can also be encapsulated by ionic MOFs as demonstrated by Zhang *et al.*⁶⁴ In this study, we have shown that mDES-1 can encapsulate and release a relatively small molecule like azobenzene. Therefore, small drugs such as 5-fluorouracil could be promising candidates for encapsulation and slow release using mDES-1. Cationic drugs are also good candidates, as they can replace the HmIm counterions inside mDES-1. However, the cytotoxicity of cobalt ions makes mDES-1 less suitable candidate for *in vivo* applications. Nevertheless, these tests with mDES-1 can serve as a valuable proof of concept. If a hydrophobic, isostructural MOF based on a less cytotoxic metal were developed, it could provide a foundation for potential application in the biomedical field.

It is important to note that in this work, we have focused on presenting the structure and some properties of the two mixed-ligand MOFs based on the well-known btc and mIm ligands. Additionally, we have explored the potential applications of the ionic mDES-1 beyond the well-known ion-exchange process. However, a more in-depth study into the isomerisation of azobenzene, as well as the encapsulation and slow release of molecules within mDES-1, is necessary and will be detailed in separate publications.

Conclusions

We have successfully synthesised and characterised two new MOFs, utilising inexpensive and well-known chemicals: 2-methylimidazole, trimesic acid and cobalt chloride. The first, mDES-1, is an iMOF with btc linkers, pendant mIm groups and HmIm counterions. Despite the pore blockage by counterions, the mDES-1 can encapsulate small molecules like azobenzene when added during the synthesis. SCXRD analysis confirms the presence of the trapped azobenzene, revealing subtle differences compared with its free form.

Moreover, the controlled release of azobenzene in ethanol was observed, achieving a release of 30 mg of azobenzene per g of loaded MOF (Azo@mDES-1) in a period of 2 months. During the first 10 h of the experiment, a delivery rate of 0.26 mg of azobenzene per hour per gram of Azo@mDES-1 was recorded.

The second MOF, mDES-2, is a 2D neutral framework that is isostructural to ITH-1 but contains cobalt ions instead of zinc ions. Similar to mDES-1, the framework in mDES-2 is formed by btc linkers and pendant mIm groups. However, the material exhibits poor crystallinity and non-accessible pores.

This work highlights the vast potential still left in the field of MOF synthesis. Despite using conventional chemicals, the successful synthesis of mDES-1 and mDES-2 demonstrate that novel MOFs with distinct properties can still be discovered. The ongoing exploration of MOF synthesis, particularly in the ionic subclass, is likely to yield further breakthroughs, advancing the capabilities of these versatile materials.

Author contributions

JJVG and ST designed the project. JJVG drafted the manuscript, performed the single crystal crystallographic experiment and analysed all results. LSV and CHWB conducted the first PXRD and magnetic measurements. LSV made the analysis of the magnetic measurements. VB performed the PXRD experiment using synchrotron radiation and the Le Bail fitting. SK and MF carried out the adsorption measurements. BK, FK, AE, FH, as interns, synthesised the MOFs under planning and supervision of JJVG. EC and TS, as interns, synthesised Azo@mDES-1 and made the calibration curve for the release of azobenzene under the planning and guidance of JJVG. LAH and WL, as summer students, re-synthesised mDES-1 and solved its structure at room temperature and captured its optical images under the guidance of JJVG. All authors provided input and agreed on the final manuscript.

Data availability

The authors declare that data supporting the findings of this study are available within the paper and its ESI.† All crystallographic data for this work has been deposited at the CCDC: 2289350, 2289467, 2377931, 2377933 and 2377934.†



Conflicts of interest

There are no conflicts to declare.

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References

- H.-C. “Joe” Zhou and S. Kitagawa, *Chem. Soc. Rev.*, 2014, **43**, 5415–5418.
- H.-C. Zhou, J. R. Long and O. M. Yaghi, *Chem. Rev.*, 2012, **112**, 673–674.
- K. Sumida, D. L. Rogow, J. A. Mason, T. M. McDonald, E. D. Bloch, Z. R. Herm, T.-H. Bae and J. R. Long, *Chem. Rev.*, 2012, **112**, 724–781.
- H. Li, K. Wang, Y. Sun, C. T. Lollar, J. Li and H.-C. Zhou, *Mater. Today*, 2018, **21**, 108–121.
- A. Bavykina, N. Kolobov, I. S. Khan, J. A. Bau, A. Ramirez and J. Gascon, *Chem. Rev.*, 2020, **120**, 8468–8535.
- M. D. Allendorf, C. A. Bauer, R. K. Bhakta and R. J. T. Houk, *Chem. Soc. Rev.*, 2009, **38**, 1330–1352.
- L. E. Kreno, K. Leong, O. K. Farha, M. Allendorf, R. P. Van Duyne and J. T. Hupp, *Chem. Rev.*, 2012, **112**, 1105–1125.
- H.-S. Wang, *Coord. Chem. Rev.*, 2017, **349**, 139–155.
- P. Horcajada, C. Serre, M. Vallet-Regí, M. Sebban, F. Taulelle and G. Férey, *Angew. Chem., Int. Ed.*, 2006, **45**, 5974–5978.
- S. Dutta, Y. D. More, S. Fajal, W. Mandal, G. K. Dam and S. K. Ghosh, *Chem. Commun.*, 2022, **58**, 13676–13698.
- A. Karmakar, A. V. Desai and S. K. Ghosh, *Coord. Chem. Rev.*, 2016, **307**, 313–341.
- J. Esteban, M. Ladero and F. García-Ochoa, *Chem. Eng. J.*, 2015, **269**, 194–202.
- M. S. Gross, B. S. Sánchez and C. A. Querini, *Appl. Catal.*, 2015, **501**, 1–9.
- S.-N. Zhao, Y. Zhang, S.-Y. Song and H.-J. Zhang, *Coord. Chem. Rev.*, 2019, **398**, 113007.
- Y. Noori and K. Akhbari, *RSC Adv.*, 2017, **7**, 1782–1808.
- J. Y. Lee, J. M. Roberts, O. K. Farha, A. A. Sarjeant, K. A. Scheidt and J. T. Hupp, *Inorg. Chem.*, 2009, **48**, 9971–9973.
- H. Zhang, Y. Lu, Z. Zhang, H. Fu, Y. Li, D. Volkmer, D. Denysenko and E. Wang, *Chem. Commun.*, 2012, **48**, 7295.
- K. S. Park, Z. Ni, A. P. Côté, J. Y. Choi, R. Huang, F. J. Uribe-Romo, H. K. Chae, M. O’Keeffe and O. M. Yaghi, *Proc. Natl. Acad. Sci. U. S. A.*, 2006, **103**, 10186–10191.
- R. Banerjee, A. Phan, B. Wang, C. Knobler, H. Furukawa, M. O’Keeffe and O. M. Yaghi, *Science*, 2008, **319**, 939–943.
- S. S.-Y. Chui, S. M.-F. Lo, J. P. H. Charmant, A. G. Orpen and I. D. Williams, *Science*, 1999, **283**, 1148–1150.
- R. R. Kuruppathparambil, T. M. Robert, R. S. Pillai, S. K. B. Pillai, S. K. Kalamblayil Shankaranarayanan, D. Kim and D. Mathew, *J. CO₂ Util.*, 2022, **59**, 101951.
- J. D. J. Velazquez-Garcia and S. Techert, *Acta Crystallogr., Sect. E:Crystallogr. Commun.*, 2022, **78**, 814–817.
- S. Baletska, S. Techert and J. D. J. Velazquez-Garcia, *Acta Crystallogr., Sect. E:Crystallogr. Commun.*, 2023, **79**, 1088–1092.
- D. Cheng, M. A. Khan and R. P. Houser, *J. Chem. Soc., Dalton Trans.*, 2002, 4555.
- C.-J. Wang, K.-F. Yue, W.-H. Zhang, J.-C. Jin, X.-Y. Huang and Y.-Y. Wang, *Inorg. Chem. Commun.*, 2010, **13**, 1332–1336.
- K. Chen, Y.-S. Kang, L. Luo, Y. Zhao, P. Wang, Q. Liu, Y. Lu and W.-Y. Sun, *Polyhedron*, 2014, **79**, 239–249.
- H.-X. Yang, Y.-Q. Yang, D. Cheng, Y.-X. Li and X.-R. Meng, *Acta Crystallogr., Sect. C:Struct. Chem.*, 2017, **73**, 828–832.
- Q.-Y. Huang, Y. Zhao and X.-R. Meng, *Acta Crystallogr., Sect. C:Struct. Chem.*, 2017, **73**, 645–651.
- G. Yuan, K.-Z. Shao, D.-Y. Du, X.-L. Wang, Z.-M. Su and J.-F. Ma, *CrystEngComm*, 2012, **14**, 1865.
- Z.-Q. Liu, K. Chen, Y. Zhao, Y.-S. Kang, X.-H. Liu, Q.-Y. Lu, M. Azam, S. I. Al-Resayes and W.-Y. Sun, *Cryst. Growth Des.*, 2018, **18**, 1136–1146.
- L. Cheng, S. Gou and J. Wang, *J. Mol. Struct.*, 2011, **991**, 149–157.
- B. Li, J. P. Vizuet, G. T. McCandless and K. J. Balkus Jr, *Polyhedron*, 2023, **230**, 116211.
- Y.-C. Cui, J.-J. Wang, B. Liu, G.-G. Gao and Q.-W. Wang, *Acta Crystallogr., Sect. E:Struct. Rep. Online*, 2007, **63**, m1204–m1205.
- X.-Q. Guo, M. Wang, Y.-F. Tang, F. Meng, G.-Q. Jiang and J.-L. Zhu, *CrystEngComm*, 2016, **18**, 1768–1774.
- J. Gu, S. Wan, X. Cheng, M. V. Kirillova and A. M. Kirillov, *Cryst. Growth Des.*, 2021, **21**, 2876–2888.
- J. P. Vizuet, T. S. Howlett, A. L. Lewis, Z. D. Chroust, G. T. McCandless and K. J. Balkus, *Inorg. Chem.*, 2019, **58**, 5031–5041.
- W. Kabsch, *J. Appl. Crystallogr.*, 1993, **26**, 795–800.
- SAINT, Bruker AXS Inc., Madison, Wisconsin, USA, 2016.
- G. M. Sheldrick, *Acta Crystallogr., Sect. A:Found. Adv.*, 2015, **71**, 3–8.
- O. V. Dolomanov, L. J. Bourhis, R. J. Gildea, J. A. K. Howard and H. Puschmann, *J. Appl. Crystallogr.*, 2009, **42**, 339–341.
- G. M. Sheldrick, *Acta Crystallogr., Sect. C:Struct. Chem.*, 2015, **71**, 3–8.
- A.-C. Dippel, H.-P. Liermann, J. T. Delitz, P. Walter, H. Schulte-Schrepping, O. H. Seeck and H. Franz, *J. Synchrotron Radiat.*, 2015, **22**, 675–687.



- 43 G. Ashiotis, A. Deschildre, Z. Nawaz, J. P. Wright, D. Karkoulis, F. E. Picca and J. Kieffer, *J. Appl. Crystallogr.*, 2015, **48**, 510–519.
- 44 J. Rodriguez-Cavajal, *Comm. Powder Diffr. Newsl.*, 2001, **26**, 12.
- 45 V. A. Blatov, A. P. Shevchenko and D. M. Proserpio, *Cryst. Growth Des.*, 2014, **14**, 3576–3586.
- 46 H. Zabrodsky, S. Peleg and D. Avnir, *J. Am. Chem. Soc.*, 1992, **114**, 7843–7851.
- 47 M. Pinsky, K. B. Lipkowitz and D. Avnir, *J. Math. Chem.*, 2001, **30**, 109–120.
- 48 M. Llunell, D. Casanova, J. Cirera, P. Alemany and S. Alvarez, *SHAPE program, version 2.1*, Barcelona, 2003.
- 49 L. Yang, D. R. Powell and R. P. Houser, *Dalton Trans.*, 2007, 955–964.
- 50 C. F. Macrae, I. Sovago, S. J. Cottrell, P. T. A. Galek, P. McCabe, E. Pidcock, M. Platings, G. P. Shields, J. S. Stevens, M. Towler and P. A. Wood, *J. Appl. Crystallogr.*, 2020, **53**, 226–235.
- 51 C. F. Macrae, I. J. Bruno, J. A. Chisholm, P. R. Edgington, P. McCabe, E. Pidcock, L. Rodriguez-Monge, R. Taylor, J. van de Streek and P. A. Wood, *J. Appl. Crystallogr.*, 2008, **41**, 466–470.
- 52 C. L. Hobday, C. H. Woodall, M. J. Lennox, M. Frost, K. Kameney, T. Düren, C. A. Morrison and S. A. Moggach, *Nat. Commun.*, 2018, **9**, 1429.
- 53 J. Harada, K. Ogawa and S. Tomoda, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1997, **53**, 662–672.
- 54 C. J. Brown, *Acta Crystallogr.*, 1966, **21**, 146–152.
- 55 J. A. Bouwstra, A. Schouten and J. Kroon, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1983, **39**, 1121–1123.
- 56 A. B. Grommet, L. M. Lee and R. Klajn, *Acc. Chem. Res.*, 2020, **53**, 2600–2610.
- 57 T. Kusukawa and M. Fujita, *J. Am. Chem. Soc.*, 1999, **121**, 1397–1398.
- 58 R. Wang, T. Iyoda, L. Jiang, D. A. Tryk, K. Hashimoto and A. Fujishima, *J. Electroanal. Chem.*, 1997, **438**, 213–219.
- 59 J. Yang and Y. Yang, *Small*, 2020, **16**, 1906846.
- 60 M. Wu and Y. Yang, *Adv. Mater.*, 2017, **29**, 1606134.
- 61 K. Lu, T. Aung, N. Guo, R. Weichselbaum and W. Lin, *Adv. Mater.*, 2018, **30**, 1707634.
- 62 Q. Hu, J. Yu, M. Liu, A. Liu, Z. Dou and Y. Yang, *J. Med. Chem.*, 2014, **57**, 5679–5685.
- 63 K. Jiang, W. Ni, X. Cao, L. Zhang and S. Lin, *Mater. Today Bio*, 2022, **13**, 100180.
- 64 Y. Zhang and J. Wang, *Inorg. Chim. Acta*, 2018, **477**, 8–14.

