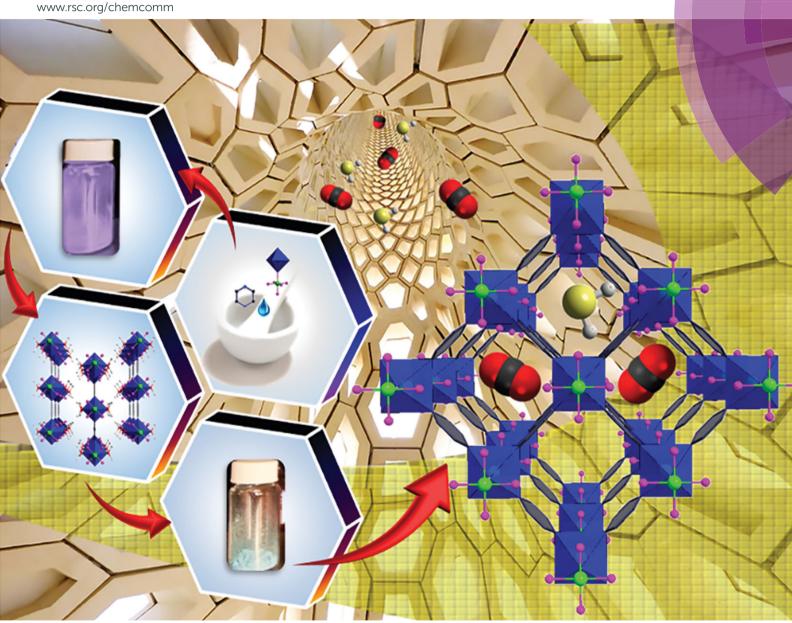
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#### COMMUNICATION

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# A facile solvent-free synthesis route for the assembly of a highly CO<sub>2</sub> selective and H<sub>2</sub>S tolerant NiSIFSIX metal-organic framework†

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The development of materials for CO2 capture with high selectivity and high tolerance to H<sub>2</sub>S is of prime importance for various industrially relevant gas streams (e.g. natural gas and biogas upgrading as well as pre-combustion capture). Here, we report the successful fabrication of a MOF with combined exceptional CO2 capture properties and H<sub>2</sub>S tolerance, namely the NiSIFSIX-based MOF using both solvothermal and solvent-free methodologies.

Metal-organic frameworks (MOFs), a special class of solid-state materials, have been considered to be one of the most burgeoning porous materials in the last two decades. MOFs are very attractive due to their modular nature, e.g. large porosity, chemical and topological versatility and structural diversity, which offer potential in various applications such as catalysis, gas storage, gas sensing, greenhouse gas emission mitigation and many other challenging gas separations.2,3

Recently, MOFs were intensively investigated for CO2 capture from various gas streams such as post-combustion, pre-combustion capture, natural gas and biogas upgrading.4 In spite of the large number of studies, only few reports focused on traces or low CO<sub>2</sub> concentration removal applications (i.e. CO<sub>2</sub> removal from air, hydrogen purification, and medical applications such as anaesthesia machines).5-7 Furthermore, development of materials capable of selective CO2 removal, in CO2 diluted streams, with high tolerance to H<sub>2</sub>S is of prime practical importance.

Our quest for made-to-order materials that can address efficiently the separation and capture of CO2 at different concentrations has prompted us to explore the potential of various MOFs with different structural properties and chemical compositions. 4d,7-9

Recently, the successful practice of reticular chemistry allowed us to fabricate a series of isoreticular very stable MOFs with

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periodically arrayed hexafluorosilicate (SIFSIX) pillars, called SIFSIX-3-Zn<sup>4d</sup> and SIFSIX-3-Cu.<sup>7</sup>

These aforementioned porous MOFs disclose unique features for carbon capture, namely uniformly aligned strong CO2 adsorption sites in an idealized and contracted pore system with high localized charge density. Particularly the use of a relatively small cation, i.e. Cu2+, led to an isoreticular analogue with a very high selectivity toward CO<sub>2</sub> adsorption at very low CO<sub>2</sub> partial pressures when compared to other physical adsorbents reported in the open literature.4d,7

These unique materials offer great potential for air-capture applications due to the periodically aligned very high (nonreactive) CO<sub>2</sub> adsorption energetic sites, resulting in a fully reversible physically driven adsorption-desorption operation under very mild conditions. The contracted pore system associated with a localized high charge density had afforded the selective CO2 adsorption with relatively much faster rates and stronger interactions than H2, CH4 and N2, and subsequently the attainment of unprecedented separation performances based on a unique synergistic mechanism involving both thermodynamics and kinetics in a wide range of CO2 concentrations.

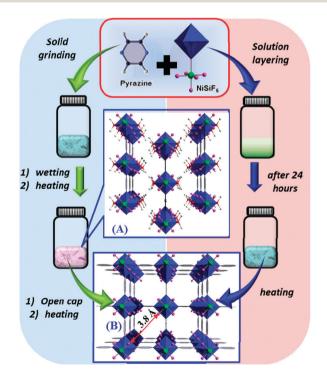
It is worth noting that commonly hybrid porous solid-state materials are synthesized using solvo/hydrothermal routes. 10,11 Various other synthetic approaches have been explored and reported for the construction of MOF materials such as (i) microwaveassisted solvothermal methods, 11 (ii) micro-fluidics, 11c (iii) ionic liquids,11d and (iv) electro-chemistry.11e Nevertheless, all these synthetic practices are found to be practical for the lab-scale research but are hardly implemented, and not often economically applicable, at the pilot or larger industrial scales. Mainly, the scale-up of MOF materials is hindered by the expensive cost, safety and the negative environmental impact associated with the use of organic solvents. Furthermore, MOF solution-based synthesis approaches suffer from additional drawbacks such as the plausible toxic and/or corrosive character associated with dissolved metal salts, generating acids as byproducts and often susceptible to solvolysis and must be disposed. 11 Additionally, the need to dispose or recycle large quantities of solvents is not

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cost effective. All the above mentioned downfalls have hindered the timely progress in the scale-up of MOFs and their successful deployment in industrial applications. 12 Accordingly, the successful use of the solvent-free synthesis route for the construction of MOFs can be regarded as a plausible route toward the requisite cost-effective scale-up of MOFs, a critical step for MOF industrial deployment.<sup>11a</sup>

Here, we report the solvent-free synthesis of a new H<sub>2</sub>S tolerant MOF material, namely SIFSIX-3-Ni, isostructural to the Zn and Cu based SIFSIX-3-M, 4d-7 with very high CO2 selectivity in a wide range of CO<sub>2</sub> concentrations. After countless failed attempts to develop solvent-free procedures for the synthesis of the previously isolated Zn and Cu SIFSIX analogues (SIFSIX-3-Zn and SIFSIX-3-Cu),  $^{4d-7}$  we extended our focus and efforts to other metal sources that can also potentially form SIFSIX-3-M isostructures. Accordingly, first we opted to target and confirm the construction of the nickel analogue using the conventional solvothermal route and then subsequently explore the ability to fabricate this Ni-analogue using a free-solvent process.

Indeed, the SIFSIX-3-Ni analogue (B in Scheme 1) was prepared successfully by layering a methanol solution of pyrazine (pyz) in a glass tube onto a methanol solution of NiSiF<sub>6</sub>·xH<sub>2</sub>O. Upon layering, a slow formation of light blue powder was observed, and the powder was left for 24 hours in the mother solution. Before the gas adsorption study, the powder was first collected, washed extensively with methanol and then heated to 105 °C, dried under vacuum and characterized using powder X-ray diffraction (PXRD) (Fig. S1, ESI†). The structure of this expected new analogue is in very good agreement with those of the Zn and Cu analogues



Scheme 1 Schematic representation of the two different preparation procedures for SIFSIX-3-Ni. (right) Solvothermal route based on layering methanol solutions of both precursors. (left) Solvent-free route that involves the grinding of precursors, wetting and heating

reported previously, 4d,7 but with a slightly smaller unit cell (378 Å<sup>3</sup>) than the Zn (388 Å<sup>3</sup>) analogue which is attributed to the relatively strong bonding between the Ni<sup>2+</sup> and the pyrazine as compared to Zn<sup>2+</sup> (Table S1, ESI†).<sup>7,8</sup> The unit cell size of the Ni analogue was found to be between the Zn and the Cu analogues, in good agreement with the relatively sharp pore size distribution (PSD) centred at 3.8 Å (average pore size), as determined from the CO<sub>2</sub> adsorption isotherm analysis using the NLDFT model (Fig. 1b). This result is in good agreement with the associated crystal structure and as anticipated is situated between the observed values for SIFSIX-3-Zn (3.8–4 Å) and SIFSIX-3-Cu (3.5 Å).

The steepness of variable temperature adsorption isotherms at low pressure for the newly isolated Ni analogue (Fig. 1a) is another indication of the relatively small pore size as compared to the Zn analogue. The intermediate steepness of CO<sub>2</sub> adsorption isotherms is also an indication of stronger/weaker CO2-SIFSIX-3-Ni interactions as compared to the Zn/Cu analogues. Accordingly, the  $Q_{st}$  of  $CO_2$  adsorption for SIFSIX-3-Ni (Fig. 2) is 7% and 8% higher/lower (48 kJ mol<sup>-1</sup>) than the Zn and the Cu analogues, respectively. The Q<sub>st</sub> of CO<sub>2</sub> adsorption is an intrinsic property that dictates the affinity of the MOF toward CO<sub>2</sub>, which in turn plays an important role in determining the necessary energy to release CO2 during the regeneration step. The Q<sub>st</sub> for CO<sub>2</sub> is in the range of fully reversible CO<sub>2</sub> adsorption  $(30-50 \text{ kJ mol}^{-1})$ , <sup>4d</sup> in agreement with the full evacuation of SIFSIX-3-Ni at 298-323 K in a vacuum. As in the case of SIFSIX-3-Zn and SIFSIX-3-Cu, the Q<sub>st</sub> for CO<sub>2</sub> adsorption was relatively

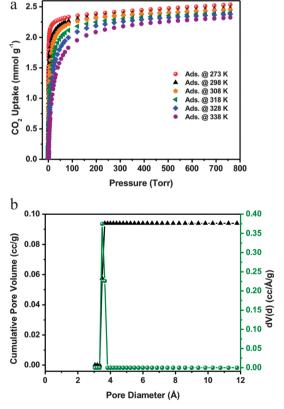


Fig. 1 (a) Variable temperature adsorption isotherms of CO<sub>2</sub> for SIFSIX-3-Ni, (b) pore size distribution from CO<sub>2</sub> sorption isotherms at 77 K for SIFSIX-3-Ni.

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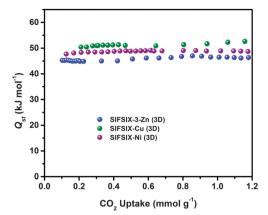


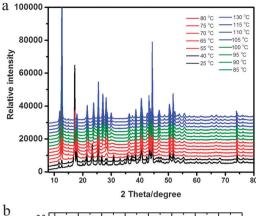
Fig. 2 Isosteric heats of adsorption at low coverage for SIFSIX-3-Cu, SIFSIX-3-Zn and SIFSIX-3-Ni

constant indicating homogenous energetic sites over the full range of CO<sub>2</sub> loading (Fig. 2). 4d,7 By analogy and based on the full mixed gas adsorption and kinetics study carried out on the Zn and Cu analogues, 4d,7 it is logically expected that SIFSIX-3-Ni will perform at least similarly as the Zn analogue<sup>4d</sup> in terms of CO<sub>2</sub> separation from N<sub>2</sub>, O<sub>2</sub>, CH<sub>4</sub> and H<sub>2</sub> containing mixtures. The anticipated high selectivity toward CO2 in the case of SIFSIX-3-Ni is projected to be driven by the unique optimal combination between thermodynamics and kinetics, i.e., the occurrence of strong/homogenous CO2-MOF interactions and the relatively fast CO<sub>2</sub> adsorption than those of other relevant gases. 4d,7

The successful isolation of the new Ni analogue solvothermally had encouraged us to explore its plausible isolation using a solvent-free route. After few attempts, and in contrast to the case of Zn and Cu analogues, we successfully discovered the appropriate conditions that permit a solvent-free route synthesis of SIFSIX-3-Ni. This extremely facile methodology is based first on mechanically mixing both the organic and inorganic solid precursors i.e. pyrazine and NiSiF<sub>6</sub> with a molar ratio of 4:1 respectively, followed by a wetting procedure with few drops of water and then a careful heating procedure (Scheme 1).

Indeed, after dry mechanical mixing and wetting, the examination of the resultant powder mixture using powder X-ray diffraction data showed that a very less crystalline product was obtained (Fig. 3a). The resultant powder mixture was heated to 65 °C for 4 hours in a closed vial which resulted in a colour change to light violet. The observed colour change is associated with the formation of the 2D structure of the SIFSIX-3-Ni analogue (A in Scheme 1). Interestingly, upon heating (drying) at 105 °C for an additional 4 hours, we observed a structural phase transformation. The PXRD diagram of the final material prepared using the solvent-free route method after the drying process was found to match and correspond to the 3D SIFSIX-3-Ni structure (B in Scheme 1) prepared using the solvothermal method (Fig. S1, ESI†). It is worth mentioning that the application of a similar solvent-free procedure to the Zn and Cu analogues did not permit the observation of the same phase transformation from the 2D to the 3D structure.

Essentially, the Rietveld refinement allowed us to clearly confirm the crystal structure of the resultant crystalline material



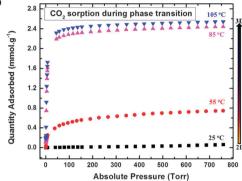


Fig. 3 (a) Variable temperature PXRD for phase transition of the 2D (NiSiF<sub>6</sub>·xH<sub>2</sub>O) to the 3D SIFSIX-3-Ni structure, and (b) CO<sub>2</sub> sorption at 298 K during phase transition at different temperatures.

from the solvent-free synthesis as SIFSIX-3-Ni (Fig. S1, ESI†). In order to better understand the crystallisation process affording the formation of SIFSIX-3-Ni, variable temperature PXRD was conducted (Fig. 3a) from 25 °C to 130 °C. At low temperature (25-40 °C), a first product with a very low crystallinity appeared which corresponds to a 2D structural analogue that was previously reported (CuSiF<sub>6</sub>(pyz)(H<sub>2</sub>O)<sub>2</sub>) (A in Scheme 1).<sup>7</sup> A further increase in the temperature resulted in a phase transformation that afforded the formation of targeted 3D SIFSIX-3-Ni starting at around 55 °C and completed at 70 °C. At higher temperatures, PXRDs confirm the good crystallinity of the 3D material and its associated noticeable stability up to 130 °C.

It is worth noting that the phase transformation was accompanied by a discernible colour change from light violet to light blue when full converted.

Delightfully, the Ni analogue prepared by the solvent-free route showed the same variable temperature adsorption isotherms at low pressure as the Ni analogue prepared by the solvothermal methods which further supports the conformity of the SIFSIX-3-Ni materials resulting from both approaches (Fig. 3b). Fig. 3b illustrates the evolution in the CO<sub>2</sub> adsorption uptake, recorded at 25 °C, as a function of the evacuation temperature for the SIFSIX-3-Ni analogue prepared by the solvent-free route method. The CO<sub>2</sub> adsorption showed an increase in the CO<sub>2</sub> uptake with increased evacuation temperatures up to 105 °C, corresponding to the optimal CO2 adsorption isotherm which also correlates with the phase change observed in the VT-PXRD and the Communication

Adsorbed amount (mmol g<sup>-1</sup> H<sub>2</sub>S sorption 10 Concentration (%)

Fig. 4 CO<sub>2</sub> and H<sub>2</sub>S sorption isotherms for SIFSIX-3-Ni. H<sub>2</sub>S uptake remains same after repeated sorption and activation cycles.

complete conversion to 3D SIFSIX-3-Ni. The Q<sub>st</sub> for CO<sub>2</sub> was found to be in good agreement with the associated values for SIFSIX-3-Ni prepared using the solvothermal method (Fig. S3, ESI†).

In light of the exceptional CO2 capture properties of SIFSIX-3-Zn and SIFSIX-3-Cu<sup>4d,7</sup> analogues, it is expected that the high CO<sub>2</sub> selectivity of SIFSIX-3-Ni at intermediate and high CO<sub>2</sub> concentrations will be similarly high. For the particular case of SIFSIX-3-Ni we found that the structure is stable up to 80% RH both in CO2-humidity and CO2-air and undergo phase change to the 2D structure (Scheme 1A) at relative humidity higher than 80% RH (Fig. S2, ESI†).

In addition to favourable adsorption properties for CO<sub>2</sub> capture, an ideal CO<sub>2</sub> adsorbent should also be tolerant to impurities such as H<sub>2</sub>S, since natural gas contains acidic gases in various amounts that can reach high concentration up to 10-20% in the case of H<sub>2</sub>S. Accordingly, we have investigated the stability of SIFSIX-3-Ni after exposure and adsorption of H2S. It is worth noting that MOFs known to be stable upon exposure to H<sub>2</sub>S are scarce.<sup>13</sup>

The H<sub>2</sub>S sorption isotherm was collected for the SIFSIX-3-Ni sample and revealed an uptake of H<sub>2</sub>S around 1.3 mol g<sup>-1</sup> at 10% concentration (Fig. 4). The H<sub>2</sub>S adsorption isotherm was repeated on the same sample in order to check/confirm the material's stability; the second H<sub>2</sub>S adsorption isotherm was found to be identical to the first one as shown in Fig. S4 (ESI†), and thus confirming the stability and regeneration/recycling of SIFSIX-3-Ni in the presence of H<sub>2</sub>S. This unique stability of the SIFSIX-3-Ni material in the presence of H<sub>2</sub>S was also supported by the similarities between the PXRD patterns before and after the H<sub>2</sub>S exposure (Fig. S2, ESI†). The resultant H<sub>2</sub>S exposed SIFSIX-3-Ni sample was then used for another cycle of CO<sub>2</sub> sorption isotherm studies. Comfortably, the new collected CO<sub>2</sub> adsorption isotherm matched exactly with the adsorption isotherm for the fresh sample not exposed to H2S as shown in Fig. S5 (ESI†). Certainly, the repeated H<sub>2</sub>S exposure did not affect/alter the subsequent capacity of SIFSIX-3-Ni for CO<sub>2</sub> and H<sub>2</sub>S. It is worth stating that the Cu and Zn analogues are not tolerant to H<sub>2</sub>S, unlike the H<sub>2</sub>S tolerant SIFSIX-3-Ni, and lose their crystallinity and CO2 uptake capacity upon exposure to H<sub>2</sub>S. This further supports the uniqueness of SIFSIX-3-Ni and its prospective use as an excellent adsorbent for CO2 removal from a gaseous mixture containing acidic gases for natural gas upgrading application.

In conclusion, we showed herein that the successful practice of MOF chemistry had permitted for the first time the fabrication of an H<sub>2</sub>S-tolerant MOF, SIFSIX-3-Ni, for practical CO<sub>2</sub> capture in a wide range of CO2 concentrations, using a solventfree synthetic approach. The exceptional CO<sub>2</sub> capture and high tolerance to H<sub>2</sub>S, in addition to the simplified synthesis procedure, qualify the SIFSIX-3-Ni analogue as a prospective candidate for scale-up and testing at the pilot scale. This projected accomplishment will facilitate the smooth transition to commercialization and will pave the way for this technology technically and economically to be viable.

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