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Enhancing the gas adsorption capacities of UiO-66 by nanographite addition

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Abstract

New composites have been synthesized containing nanographite (nGr) particles as platforms for the deposition of UiO-66 crystals. The samples were analyzed by X-ray diffraction, thermogravimetry, N₂ adsorption and scanning electron microscopy. The addition of nGr markedly increased the porosity determined by N₂ adsorption (0.5 nm ultramicropores were formed, and the volume in pores > 1.1 nm increased). This was the result of defects formation either on UiO-66 or at the nGr/UiO-66 interface. To assess the adsorptive properties and the accessibility of the pores, adsorption of H₂, CO₂, C₂H₄ and C₂H₆ was measured. The results have shown that not only the extent of the porosity but also the interactions with the graphite phase and the accessibility to the pores affect the amount adsorbed and the strength of adsorption. Generally, the formation of composites had a positive effect on the adsorption properties owing to an increased surface heterogeneity and porosity.

Keywords: metal organic framework; nanographite; small gas molecule adsorption; defects; porosity; isosteric heat of adsorption

1. Introduction

Metal–organic frameworks (MOFs) are a group of materials represented by a variety of structures and chemistries affecting their properties and applications [1, 2]. Even though they are perfect crystals, this perfection is not always beneficial for the target performance, especially in catalysis. Therefore, in recent years, research has been carried out with focus on increasing the defects populations in otherwise perfect MOFs [3]. The defects in MOFs can arise from crystal imperfections caused mainly by partial ligand replacement [4, 5], missing clusters [6], heterogeneity of ligands [7, 8] or a use of modulators during the MOF synthesis [9, 10].

UiO-66 is a well-studied MOF, which exhibits a high surface area, chemical, mechanical and thermal stability [11-13]. The synthesis of this MOF is relatively easy and enables the tunability of its structural defects [4-10]. These defects, besides being known to increase pore volumes and surface areas, also change pores' morphology/distribution and affect the adsorption of gases such as CO₂ [14-18], H₂O [19, 20], H₂ [15, 21-23], C₂H₆ [24-27] or C₂H₄ [25].

Adsorption of CO₂ on UiO-66 has been extensively studied in search of sufficient sequestration media. Both experimental [15-17] and theoretical approaches have been explored [14, 17]. At 0.1 MPa and 0°C, adsorbed amounts of ~ 2 mmol/g have been reported [15, 16]. While the heat of adsorption measured by adsorption methods was between 22 and 28 kJ/mol [15, 16] and increased with the introduction of defects [16], results of infrared spectroscopy at variable temperature suggested that the heat of CO₂ adsorption by hydrogen bonding reaches 38 kJ/mol, and 30.2 kJ/mol - by dispersive interactions [14]. Recently, we have reported that defects in UiO-66, synthesized in the presence of nanocellulose crystals of various chemistries, not only increased the amount of CO₂ adsorbed by about 30 % at 0°C, compared to UiO-66 (3.25 mol/g

as compared to 2.5 mmol/g), but also led to an increase in the heat of adsorption from 20 to 27 kJ/mol [28].

In the case of H₂ adsorption on UiO-66, the amount adsorbed at – 196°C and 0.1 MPa increased from 1.25 % to 1.50 % upon changing the solvent from ethanol to DMF used in its synthesis [21]. This also resulted in an increase in the homogeneity of the crystals (well defined sizes of 150-200 nm in DMF) and in an increase in the surface area from 784 m²/g in ethanol to 1358 m²/g in DMF. In this case, the lower surface area of the sample synthesized in ethanol was linked to impurities due to polymerization products of benzenedicarboxylate (BDC). Nanosized UiO-66 of an enhanced porosity due to the more effective removal of solvent and unreacted BDC from the pores had a surface area of 1434 m²/g and a H₂ uptake of 1.50 % [15]. Ren et al. studied adsorption of hydrogen in UiO-66 synthesized with various amounts of formic acid as a modulator [23]. They found that the increase in the amount of modulator increased the size of the crystals and the surface area to 1367 m²/g. On this material, 1.5 % of H₂ was adsorbed at – 196°C and 0.1 MPa.

Although the separation of alkanes from alkenes is an industrially important process, UiO-66 has rather rarely been used in this application, likely owing to the lack of unsaturated metal sites, which would provide the specific interactions/ π -complexation, as in the case of Cu-BTC (benzenetricarboxylate) [24]. Nevertheless, Qian et al. [24] reported that the number of carboxyl groups in the linkers of UiO-66 affected the C₂H₆/C₂H₄ separation. When 1, 2, 4 BTC (in UiO-66-COOH) was used as a linker, there was no difference between the amount adsorbed of both species at 25°C, and about 1.3 mmol/g were reported. Increasing the number of carboxylic acid in the linker (1, 2, 4, 5-BTEC; in UiO-66-(COOH)₂) increased the amount adsorbed to 1.8 mmol/g and the surface was slightly more favorable for adsorption of C₂H₆ compared to that of

C_2H_4 (~ 0.1 mmol/g more of C_2H_6 was adsorbed at 0.1 MPa). The latter happened despite the smaller surface area of $UiO-66-(COOH)_2$ ($622\text{ m}^2/\text{g}$ compared to $713\text{ m}^2/\text{g}$ for $UiO-66-COOH$). This trend was linked to smaller pore sizes in $UiO-66-(COOH)_2$ compared to those in $UiO-66-COOH$. The heats of C_2H_4 adsorption (Q_{st}°) showed the homogeneity of the surface sites (flat) and Q_{st}° increased from 24.7 kJ/mol for $UiO-66-COOH$ to 27.4 kJ/mol for $UiO-66-(COOH)_2$ [24]. The effect of $UiO-66$ topology on the adsorption of ethane and ethylene was also studied by Li and coworkers [25]. On their $UiO-66$ with a surface area of $1014\text{ m}^2/\text{g}$, 67 % more of C_2H_6 than that of C_2H_4 was adsorbed, and the heats of adsorption were 26 and 24 kJ/mol, respectively. The difference in the heat of adsorption was linked do the difference in the polarity of these two molecules. The adsorption of ethane at 30°C on $UiO-66$ was studied by Llewellyn and coworkers [26], who measured a capacity of 1.7 mmol/g and a heat of adsorption of $\sim 32\text{ kJ/mol}$ on a material having a surface area of $\sim 1050\text{ m}^2/\text{g}$.

Considering that a small alteration in the porosity of $UiO-66$ and defects in their structure were reported to affect the gas adsorption behavior, the objective of this paper is to investigate the effects of the addition of nanographite (~ 1 to 6 wt. %) with particle sizes of about $19\text{ }\mu\text{m}$ on the porous properties and the gas adsorption behavior of the final $UiO-66$ /nanographite composites. Since the nanographite particles are much larger than crystals/aggregates of $UiO-66$ [23], they are expected to work as platforms for the deposition of the latter. Owing to the lack of functional groups on the basal planes of nanographite flakes [29], the $UiO-66$ deposition might occur mainly via dispersive forces and due to the affinity of the organic linker (BTC) for the graphite surface. This is expected to create either an interface or defects in the MOF phase and, as a consequence, to increase the porosity. Although composites containing $UiO-66$ and graphite oxide have been synthesized and used as CO_2 adsorbents, resulting in an increased storage

capacity [17, 18], their mechanism of formation was different than that expected here due to the lack of oxygen groups on nanographite used in our approach. These defects can be either of chemical or physical in nature and their geometric dimensions should be rather small and of the order of the fraction of a nanometer. Therefore, to test the extent of the approach applied, the surface accessibility of the synthesized materials has been investigated with regard to adsorption of N₂, CO₂, H₂, C₂H₆ and C₂H₄. Even though nitrogen is commonly used to evaluate the porosity [30], despite its known kinetic limitations to access very small pores, we have also measured the adsorption of hydrogen at – 196°C. CO₂ was not used for the textural characterization owing to its possible specific interaction with the surface of adsorbents [31]. The isosteric heats of adsorption were calculated for CO₂, C₂H₆ and C₂H₄ from their adsorption isotherms measured at least at two temperatures. The heats of adsorption of C₂H₆ and C₂H₄ were analyzed in details since the larger sizes of these molecules than other chosen by us as the porosity probes might affect the surface interactions. Moreover π bond of C₂H₄ might affect the strength of adsorption if acidic groups are on the accessible surface [32]. The adsorption of several small molecules on UiO-66/nanographite composites with different contents of nGr allowed understanding the effect of nanographite on the final performances of the materials.

2. Experimental

2.1. Materials

All chemicals, zirconium tetrachloride (ZrCl₄), *N,N*-dimethylformamide (DMF, 99%), terephthalic acid (C₈H₆O₄, 99+%), hydrochloric acid (HCl, 37%) and ethanol (EtOH, 96%) were supplied by either Alfa Aesar, Acros organics or VWR and used as received. Nano-sized graphite (Nano-19) was supplied by Asbury Graphite Mills Ltd., USA.

2.2. Synthesis of materials

All UiO-66 and composite samples were synthesized following the procedure reported elsewhere [11, 12], with slight modifications. The experimental steps were as follows: Step 1: A 120 mL vial was loaded with 4 mL of concentrated HCl, 0.5 g of ZrCl₄ and 20 mL of DMF. Then, the mixture was sonicated for 20 minutes until complete dissolution; Step 2: 40 mL of DMF and 0.5 g of C₈H₆O₄ (ligand) were added to the solution, sonicated for 20 minutes and then heated overnight at 80°C; Step 3: The resulting product was filtered off, washed with DMF (8 × 30 mL) and subsequently with EtOH (8 × 30 mL) to remove the excess of unreacted organic linkers. The final product was dried in a vacuum oven at 90°C for 24 h before being analyzed.

The composite materials were synthesized by following exactly the same procedure as described above but 1, 2, and 6 % of nanographite (based on the final mass of MOF) were added in the first step. The samples were labeled as UiO-66 and UiO-66-nGr1, UiO-66-nGr2 and UiO-66-nGr6 for the composites with increasing nanographite contents.

2.3. Analysis Methods

Textural and morphological characterization

Nitrogen adsorption isotherms were measured at – 196°C on a Micromeritics ASAP 2020 instrument. Prior to each measurement, each sample was outgassed overnight at 150°C until a constant vacuum of 10⁻⁷ kPa was reached. The Brunauer-Emmett-Teller (BET) method within a relative pressure range of 0.05-0.15 was used to calculate the specific area (S_{BET}) while the total pore volume (V_{T}) was calculated from the N₂ uptake at relative pressure p/p_0 of 0.998.

The Non-Local Density Functional Theory in its two-dimensional version, 2D-NLDFT, considering slit-shaped pores with energetically heterogeneous and rough walls [33], was used to

estimate the volume of micropores, V_{mic} , and ultramicropores ($V_{<0.7 \text{ nm}}$) and to obtain the pore size distributions (PSDs). The volume of mesopores, V_{mes} , was calculated as a difference between V_{T} and V_{mic} .

X-ray diffraction (XRD) patterns were collected with a Philips PANalytical X'Pert X-ray diffractometer using CuK_α (40 kV, 40 mA) radiation. All the measurements were acquired in the 2-theta (2θ) range from 5° to 70° , in steps of 0.02° , and with a time of 0.5 s per step.

Scanning electron images were recorded using a Zeiss Supra 55 VP scanning electron microscope (SEM) with an electron beam of 5 keV.

Thermal analysis was performed using an SDT Q600 of TA Instruments. All the samples were analyzed under helium and air flow (100 mL/min), heating them up to 1000°C at $10^\circ\text{C}/\text{min}$. The two thermogravimetric (TG) curves were collected, and the corresponding differential thermal gravimetric (DTG) curves were calculated in order to determine the thermal behavior of the samples.

Adsorption measurements

CO_2 adsorption isotherms at three different temperatures (0°C , 20°C and 40°C) were measured using an ASAP 2020 (Micromeritics) up to 120 kPa. Each sample was outgassed for 12 h at 150°C under vacuum ($P < 10^{-7}$ kPa) prior to each CO_2 adsorption test.

H_2 adsorption capacity was evaluated at -196°C up to 100 kPa by using a 3Flex automatic device (Micromeritics) equipped with a high-vacuum system and three 0.013 kPa pressure transducers, one per analysis port. The samples were outgassed for 24 h at 120°C under vacuum ($P < 10^{-7}$ kPa) for at least 24 h before being weighed. An additional outgassing was carried out for at least 6 hours more on the analysis ports at the same temperature and under the same final vacuum as in the previous outgassing step. High-resolution H_2 isotherms were obtained by

dosing small hydrogen amounts, as low as $3 \text{ cm}^3/\text{g}$. The determination of the warm and cold volume was carried out after H_2 adsorption-desorption analysis to avoid any possibility of a helium entrapment in the ultramicroporosity.

Measurements of C_2H_4 and C_2H_6 isotherms at -58 and -73°C were carried out using again the high-resolution Micromeritics 3Flex adsorption instrument. The latter was equipped with a high-vacuum system, three 0.1 Torr pressure transducers, and was coupled with a cryostat (Cold Edge Technologies). The isotherms of C_2H_4 and C_2H_6 as well as CO_2 at different temperatures were used to calculate the isosteric heats of adsorption. The calculations were performed using the analytical form of virial equation [34].

3. Results and Discussion

The X-ray diffraction patterns of the synthesized samples are presented in Figure 1. Although the diffraction peaks of the composite materials resemble those of UiO-66, two main diffraction peaks at 2θ less than 10° show a marked disturbance in the crystal structure, indicating some shortening of the lattice parameters. This might be related to the pores of smaller sizes than those in the parent UiO-66 materials [35].

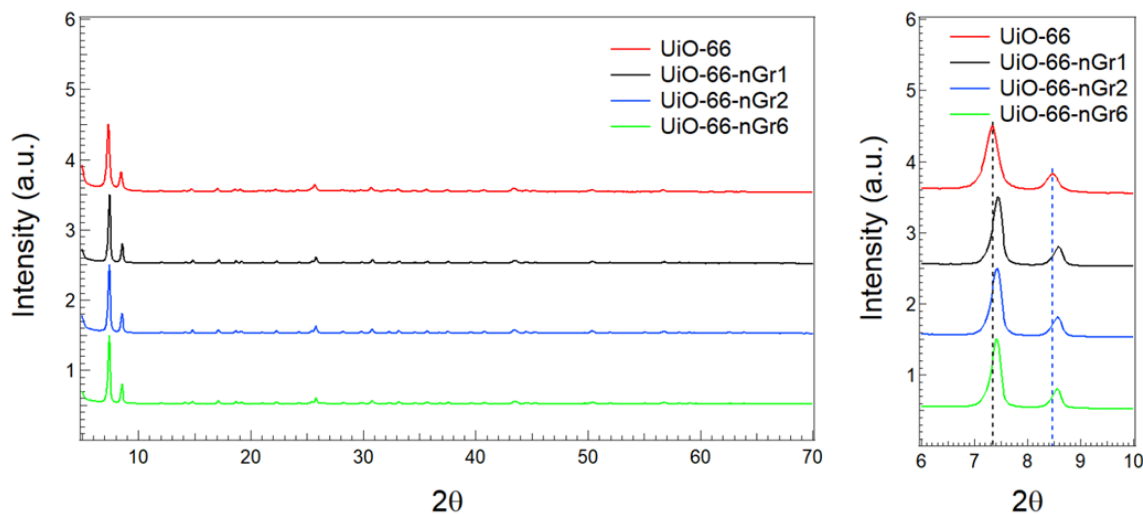


Figure 1. XRD patterns of UiO-66 and its composites with nGr.

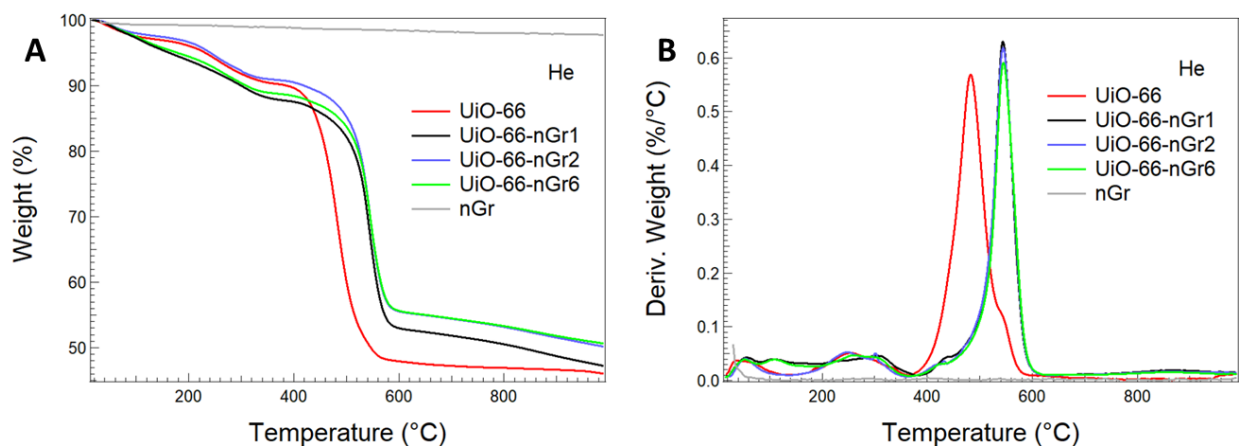


Figure 2. TG (A) and DTG (B) curves under helium flow for pure UiO-66, nGr and UiO-66-nGr composite samples.

TG and DTG curves measured in helium are collected in Figure 2. Based on the thermal stability of the composites upon addition of nGr, they bring indirect information about the change in the chemical composition of our composites. While the linkers in UiO-66 decompose at 460°C, causing the collapsing of the MOF structure [11, 36], this process for all composites takes place

at a temperature of about 80°C higher. This shift might be related to the change in a heat capacity upon the addition of the nanographite phase. This change in the heat capacity might also be due to the expected alteration in the samples' porosity. Interestingly, the decomposition event of the composites overlaps with a small shoulder seen on the UiO-66 DTG curve, which likely suggests the imperfection of our UiO-66 structure. A broad peak between 200 and 400°C is linked to the removal of the residual solvent. This process occurs over a broader temperature range for the composites and might be caused by the interaction of the solvent with the nGr phase.

The results of thermal analysis in air are presented in Figures 1S and 2S of the Supplementary Information. From these data, assuming that the missing linkers are the main defects in our materials, and based on the approach of Katz et al. [12], the number of linkers was calculated. The obtained results indicate that 8, 7.8, 7.8 and 7.6 linkers are present per node in UiO-66, UiO-66-nGr, UiO-66-nGr2 and UiO-66-nGr6, respectively. Since the ideal structure is expected to have 12 linkers, all our materials can be considered defective and the level of defects is higher in the composites and increases with an increase in the nGr content.

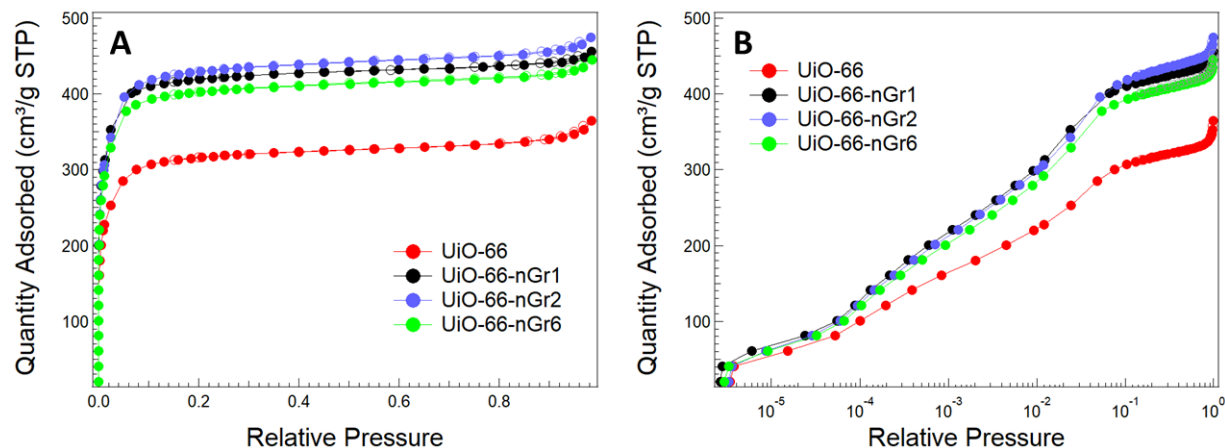


Figure 3. Nitrogen adsorption isotherms in linear (A) and logarithmic (B) scale.

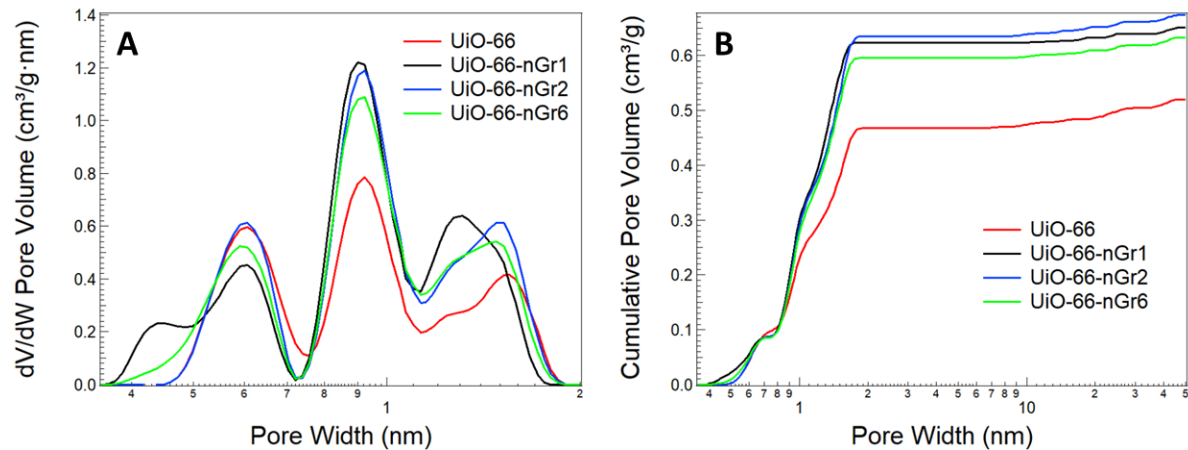


Figure 4. Differential pore size distributions calculated from N_2 isotherms using the 2D-NLDFT up to 2 nm (A) and cumulative pore volume (B).

This increase in the level of defects should be reflected in an increased porosity. The nitrogen adsorption isotherms and the pore size distributions calculated from them [33] are presented in Figures 3 and 4, respectively. The addition of nGr increased the volume of N_2 adsorbed, and the calculated surface areas of the composites increased by about 30 % compared to that of UiO-66 (Table 1). This increase in the surface area is consistent with the decrease in the numbers of linkers per node discussed above. Interestingly, no trend in the porosity development was found with an increase in the nGr content, and the S_{BET} of all composites remained at the level of about $1600 \text{ m}^2/\text{g}$. The volumes of ultra-, micro-, mesopores and the total pore volumes of the composites are also higher than those of UiO-66.

Table 1. Textural parameters calculated from N₂ adsorption isotherms at – 196°C.

Sample	S_{BET} [m ² /g]	$V_{<1\text{nm}}$ [cm ³ /g]	V_{mic} [cm ³ /g]	V_{mes} [cm ³ /g]	V_{T} [cm ³ /g]
UiO-66	1192	0.240	0.468	0.052	0.520
UiO-66-nGr1	1598	0.314	0.623	0.028	0.651
UiO-66-nGr2	1623	0.304	0.635	0.039	0.674
UiO-66-nGr6	1526	0.290	0.596	0.037	0.633

Table 2. CO₂, C₂H₄, C₂H₆ and H₂ adsorption capacity at 100 kPa.

Sample	CO ₂ uptake [mmol/g]			C ₂ H ₄ uptake [mmol/g]		C ₂ H ₆ uptake [mmol/g]		H ₂ uptake [wt. %]
	0°C	20°C	40°C	-58°C	-73°C	-58°C	-73°C	-196°C
UiO-66	2.51	1.73	1.14	6.77	7.84	6.80	7.47	1.27
UiO-66-nGr1	2.99	2.02	1.33					1.41
UiO-66-nGr2	2.87	1.99	1.25	9.62	11.14	9.72	10.66	1.48
UiO-66-nGr6	2.94	1.99	1.20					1.56

The pore size distributions calculated from the N₂ adsorption isotherms (Figure 4) bring more details about the alterations in the porous texture of our materials. As expected from the structure of UiO-66, the predominant pore sizes should be ~ 0.6 nm and ~ 0.9 nm, which represent tetrahedral and octahedral cavities [19]. Defects such as missing linkers and missing clusters are expected to increase the sizes of these latter pores [6, 12, 37]. The striking difference between

UiO-66 and the composites is the appearance, in the latter materials, of pores smaller than 0.5 nm and an increase in the volume of pores wider than 1.1 nm in their size. This effect is especially seen for the composite with the smallest amount of nGr. A detailed comparison of the change in the predominant pore sizes based on the PSDs is presented in Figure 5, where it is clearly seen that the greatest effect is an increase in pores wider than 0.7 nm upon the composite formation. This increase in the porosity is due to both, the pores of diameter between 0.7 and 1.1 nm, which represent the octahedral cavities, and to pores wider than 1.1 nm, which are assigned to defects such as missing linkers [12, 37] and clusters [6]. Pores smaller than 0.5 nm might represent the interface between nGr and UiO-crystal deposited on the nGr particles and thus, the smallest amount of nGr added might result in the highest dispersion of this phase and therefore in a more extended interface.

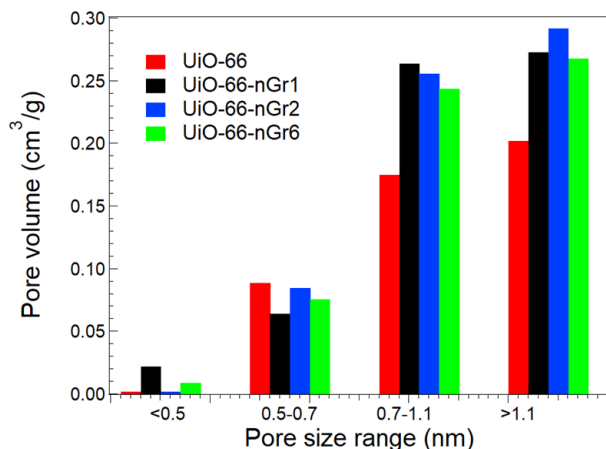


Figure 5. Pore volumes in the different diameter ranges deduced from the pore size distributions given in Figure 4.

Figure 6 shows SEM images of UiO-66 and UiO-66-nGr2, which confirmed that UiO-66 is deposited on nGr flakes and that “wall structures” of UiO-66 are assembled horizontally. This morphology might increase the accessibility of the adsorbates to the UiO-66 adsorption

sites/pores. The SEM images of nGr are presented in Figure 3S of the Supplementary Information.

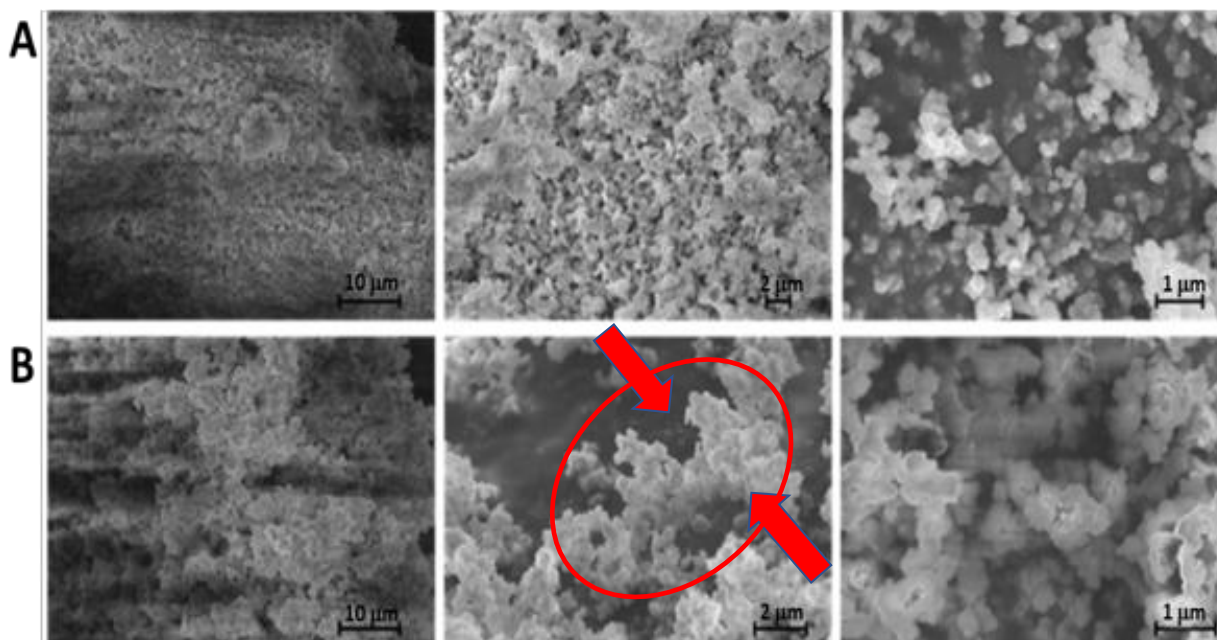


Figure 6. SEM images of pure UiO-66 (A) and UiO-66-nGr2 composite (B) samples. The assembled wall of nanoparticles on the nGr surface is marked in the bottom middle micrograph.

Figure 7 shows the CO₂ adsorption isotherms measured on the four materials at 0, 20 and 40°C. The amounts of CO₂ adsorbed at atmospheric pressure are compared in Table 2 and are in the range of those reported in the literature [14-18]. Upon the composite formation, a ~20 % increase in the amount of CO₂ adsorbed was found. The best performing sample is UiO-66-nGr1, and we link it to its highest volume of pores smaller than 0.5 nm in which the adsorption forces for CO₂ are the strongest and where, at the low relative pressure, the density of CO₂ is the highest [38]. The amount of CO₂ adsorbed on UiO-66-nGr2 and UiO-66-nGr 6 is similar to that on UiO-66-nGr1, which is in agreement with the similarities in the volume of ultramicropores in the

composite materials. The CO₂ uptake decreased with the increasing temperature, from 3 mmol/g at 0°C to 1.2 mmol/g at 40°C for UiO-66-nGr1, in good agreement with the exothermic nature of CO₂ adsorption on these materials.

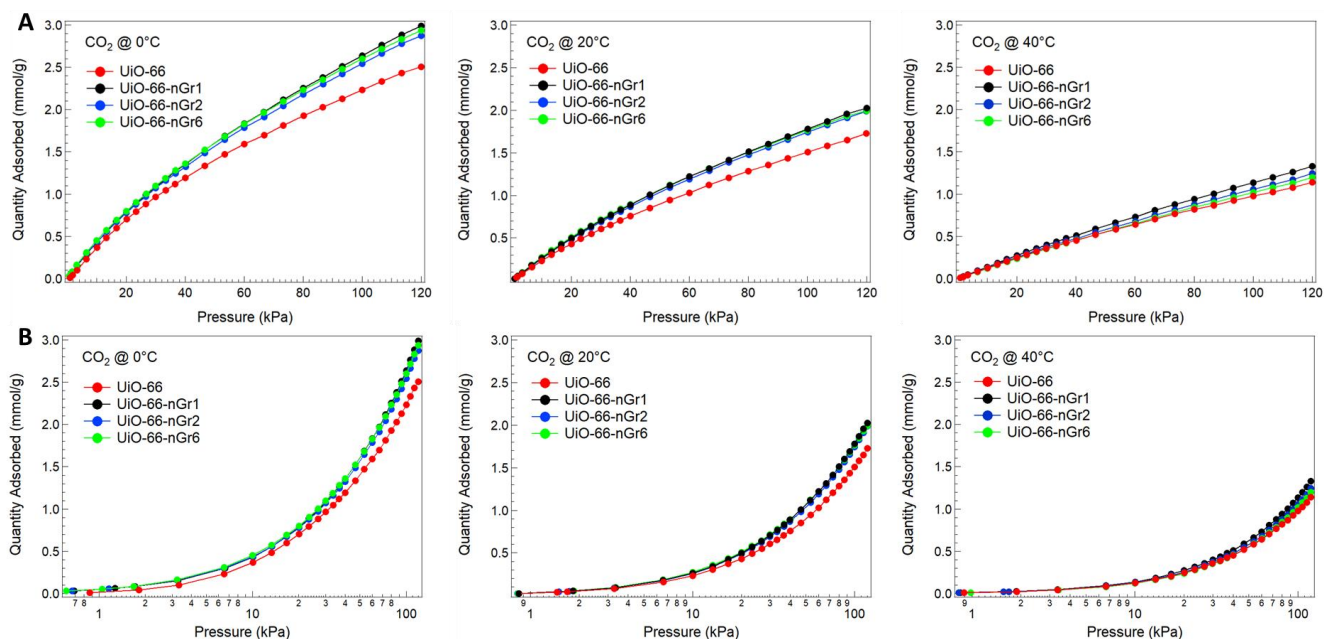


Figure 7. CO₂ adsorption isotherms on UiO-66 composites at 0°C, 20°C and 40°C in the pressure range 0–120 kPa on (A) linear and (B) logarithmic scale.

Figure 8A shows the isosteric heats of CO₂ adsorption, whose values are in the range of those reported in the literature [15, 16]. Even though there is no visible trend in the amount adsorbed with an increase in the amount of nGr, an increase in the heats of adsorption was observed with an increase in the nGr content (Figure 9A). The latter suggests that the addition of nGr not only enhanced the amount of CO₂ adsorbed but also increased the strength of CO₂ adsorption, which might be related to a denser array of carbon atoms in nGr. It is important to mention that the heats of adsorption show little variations with the surface coverage, suggesting the homogeneity of the adsorption centers occupied by CO₂ in each sample at our experimental conditions. Thus, based on the amount of CO₂ adsorbed only 0.050 cm³/g would be covered. Therefore, the

volume occupied by CO₂ represents only a small fraction of the total pore volume and, based on the N₂ adsorption results, the pores that have very homogeneous sizes contribute to this volume in all the samples.

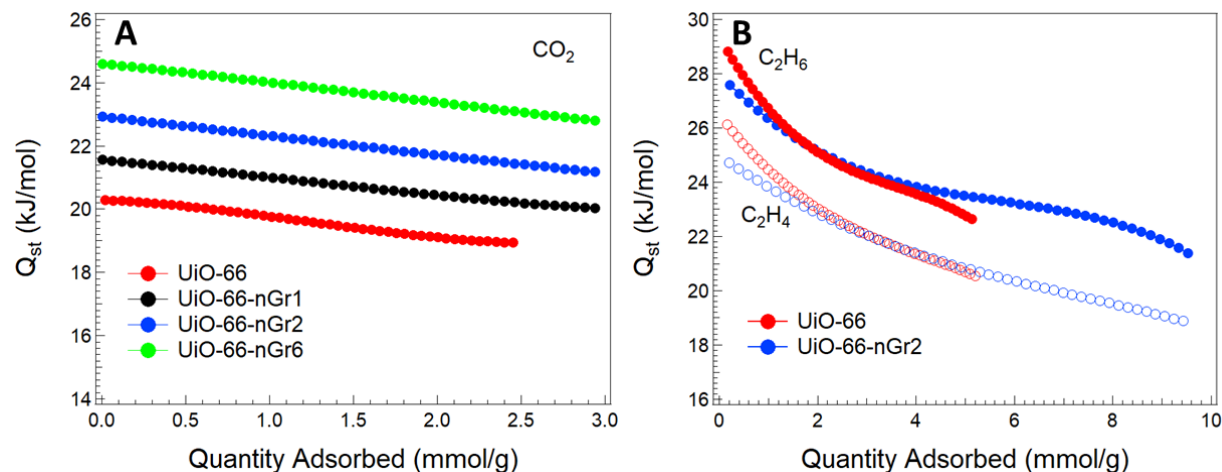


Figure 8. Isothermic heats of adsorption (Q_{st}) of CO₂ (A) and C₂H₄ (open symbols)/C₂H₆ (solid symbols) (B) on pure UiO-66 (red) and composite samples (blue).

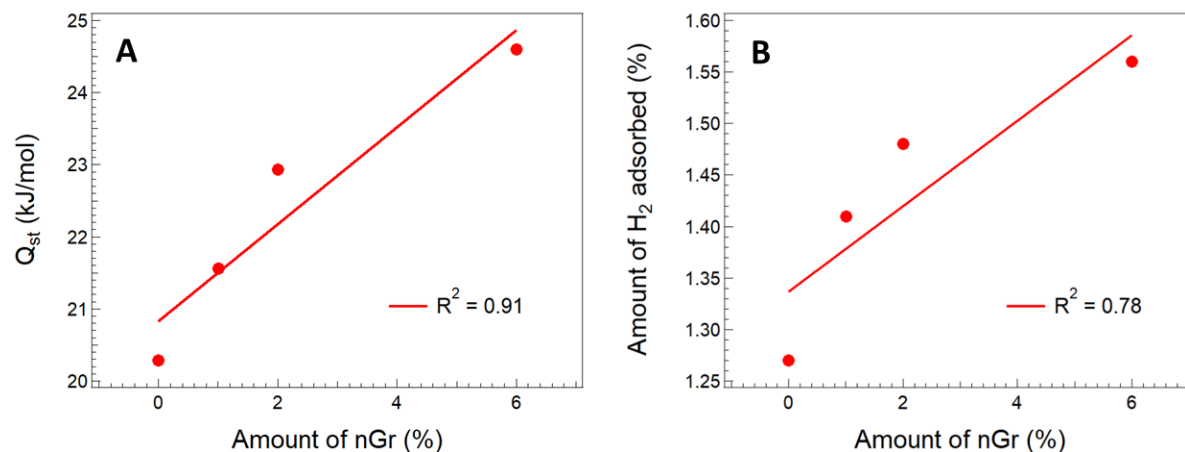


Figure 9. A) Relationship between the isothermic heat of CO₂ adsorption at zero coverage and the amount of nGr in the composites; B) relationship between the amount of H₂ adsorbed and the amount of nGr in the composites.

Figure 10 shows the H₂ adsorption-desorption isotherms measured at – 196°C on UiO-66 and its composites. Desorption was fully reversible for all materials studied. Table 2 shows the hydrogen amounts adsorbed at 100 kPa, which are similar to those reported in the literature [15, 21, 22]. As in the case of CO₂, the amount of hydrogen adsorbed in composites was higher than in pure UiO-66. However, unlike CO₂, a clear increase in the hydrogen uptake with the increase in the nGr content was found observed (Figure 9B). On UiO-66-nGr6, 23 % more of H₂ was adsorbed than on UiO-66. Even though H₂ is expected to be adsorbed in the smallest pores, no trend between the amount adsorbed and the porosity could be found taking into consideration the pore texture analyzed by N₂ adsorption. The molecular diameter of N₂ is 0.349 nm while that of H₂ is 0.296 nm, therefore H₂ molecules should have access to pore diameters that are not accessible by N₂ molecules [39, 40]. Moreover, the absolute quadrupole moment of N₂ is higher than that of H₂, 0.46 and 0.28 [41], respectively, and this could also have an effect on the differences observed [40]. Thus, the dependence on the amount of nGr added might be related to the specific morphology of the composite and to the enhanced diffusion of H₂ at – 196°C to the UiO-66 units assembled in the vertical pseudo-walls.

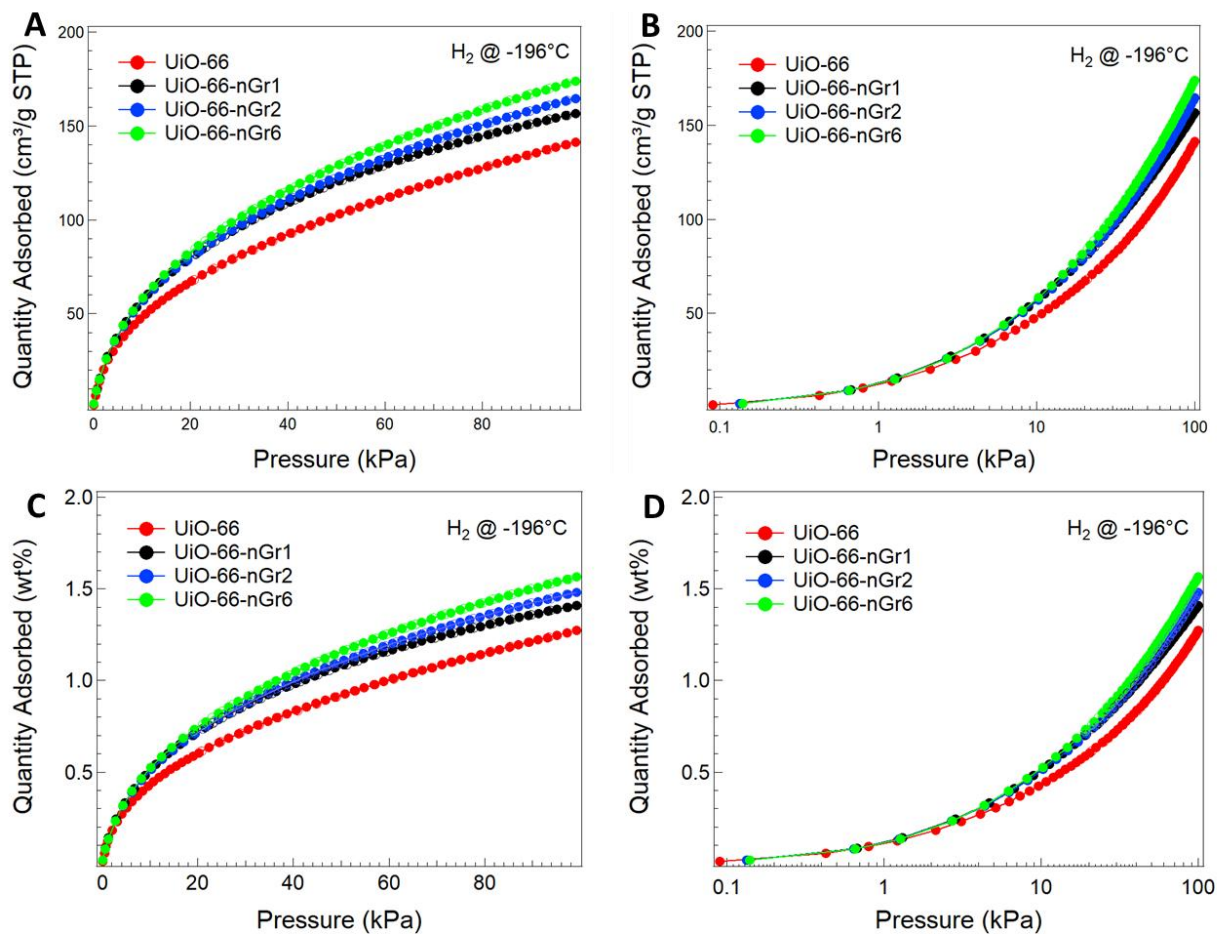


Figure 10. H₂ adsorption isotherms on pure UiO-66 and its composites at – 196°C in the pressure range 0-100 kPa, reported in cm³/g and wt. % on both (A,C) linear and (B,D) logarithmic scales.

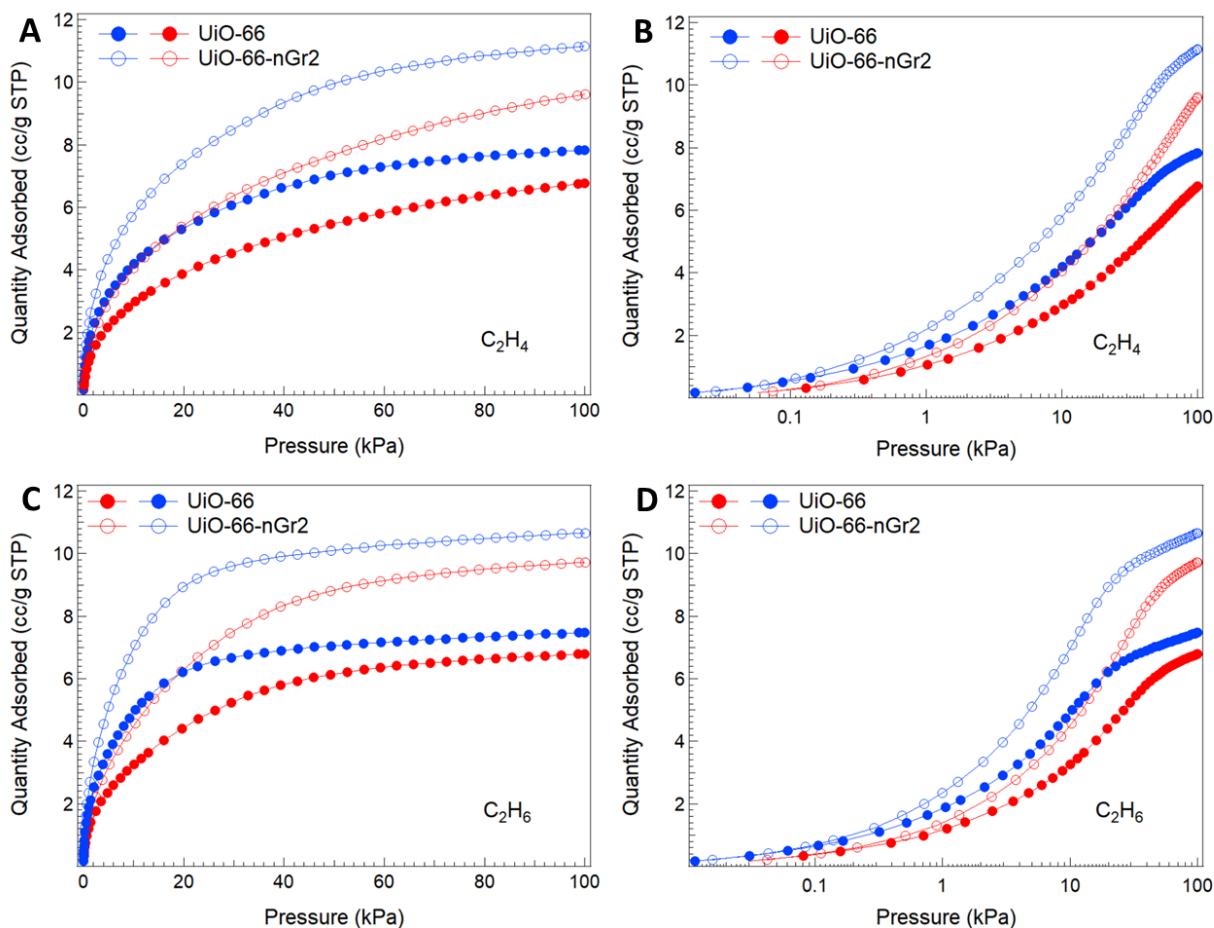


Figure 11. C_2H_4 adsorption isotherms in a linear (A) and logarithmic scale (B) and C_2H_6 adsorption isotherm in a linear (C) and logarithmic scale (D) on the UiO-66 composites at $-58^\circ C$ (blue symbols) and $-73^\circ C$ (red symbols) in the pressure range 0–100 kPa

To further explore the effects of composites texture and chemistry on the adsorption of small gas molecules, the adsorption of C_2H_4 and C_2H_6 was measured on UiO-66 and on UiO-66-nGr2 (Figure 11). Ethane and ethylene, besides having slightly different kinetic molecular sizes (σ for C_2H_4 and C_2H_6 is 0.416 and 0.444 nm, respectively [42]), also differ in chemical properties with the former being more basic due to the presence of a π -bond [32]. Their adsorption was measured at -58 and $-73^\circ C$ and the results are summarized in Table 2. In this case, building the

composite significantly increased the amounts of both C_2H_4 and C_2H_6 adsorbed compared to those on UiO-66, and, on UiO-66-nGr2, ~ 42% more of both gases were adsorbed at 100 kPa. Since the differences in the overall porosity between these two samples are not so pronounced (the porosity of the composite is only about 25 % higher), this difference might be caused by different extents of pore filling processes and in the accessibility of the pores. The affinity of hydrocarbons to adsorb on the graphite/graphite-UiO-66 interface might play a role. The biggest difference in the porosity between UiO-66 and the composite is in pores with sizes > 0.7 nm, and these pores seem to govern the amount of both gases adsorbed.

The comparison of the isosteric heats of C_2H_4 and C_2H_6 adsorption is presented in Figure 8B. Q_{st}° for C_2H_4 is 26.1 and 24.7 kJ/mol on UiO-66 and on the composite, respectively. For C_2H_6 , these values are 28.8 and 27.6 kJ/mol, respectively. The heats are consistent with those reported by Wang and co-workers [25]. The decrease in Q_{st}° with the coverage indicates that pores of various sizes are involved in the adsorption process, especially for ethane, those that are first occupied being the narrowest, for which the adsorption potential is the strongest. Therefore, the slightly higher heats at low surface coverage on UiO-66 might be due to the volume of pores with size 0.5-0.7 nm that is higher in pure UiO-66 than in the composite. Then, with the progress of the adsorption process, the octahedral pores and these related to defects are filled and the composite has higher volume of such pore than has UiO-66.

The addition of nanographite had a positive effect on the gases adsorbed either by an increase in the narrow porosity or by an increase in the specific interactions with the nanographite phase, and could be a strategy to increase the gas adsorption capacity of MOFs. An exceptional hydrogen storage has indeed been achieved with SNU-70, UMCM-9, and PCN-610/NU-100, but analysis of trends revealed the existence of a volumetric ceiling at ~ 40 g H_2 L^{-1} [43]. Surpassing

this ceiling is proposed as a new capacity target for hydrogen adsorbents. In the same study, it was concluded that usable capacities in the highest-capacity materials are negatively correlated with density and volumetric surface area. Instead, the H₂ capacity is maximized by increasing gravimetric surface area and the porosity, which have been enhanced by nanographite addition in the present study.

4. Conclusions

The adsorption of different small gas molecules allowed to clarify that the addition of even small amounts of nanographite phase, targeting 1, 2 and 6 wt. % in the composites during the synthesis of UiO-66, resulted in a marked (~ 30 %) increase in the porosity. Ultramicropores narrower than 0.5 nm and pores wider than 0.7 nm were formed and they probably have their origin in defective UiO-66 deposited on the nanographite particles. These defects increased markedly the adsorbed amounts of H₂, CO₂, C₂H₄ and C₂H₆ on the composites compared to those on the parent UiO-66. Even though no direct dependence of the porosity on the amount of nanographite in the composites was found when the texture was analyzed by N₂ adsorption, the increase in the quantity of hydrogen adsorbed with an increase in the nanographite content might indicate the existence of ultramicropores only accessible to H₂. The isosteric heat of CO₂ adsorption at zero surface coverage almost linearly increased with the nanographite content, suggesting the strong adsorption of CO₂ on the interface. In the case of H₂, its amount depended on the amount of nanographite suggesting the extensive effects of the interface or the improved diffusion of H₂ to the smallest pores owing to the assembly of UiO-66 particle “walls” on the nanographite components. In the case of ethane and ethylene adsorption, besides an increased porosity of the composite, the interactions of these hydrocarbons with nanographite phase likely contributed to

the marked increase in the amount adsorbed on the composite (42%). The trends in the heats of adsorption (higher on pure UiO-66 than on the composite) suggested that these molecules adsorbed in tetrahedral, octahedral, and defects-related pores. In particular, the addition of nanographite appears as an especially interesting strategy to increase H₂ adsorption capacity of MOFs.

References

- [1] S. Yuan, L. Feng, K. Wang, J. Pang, M. Bosch, C. Lollar, Y. Sun, J. Qin, X. Yang, P. Zhang, Q. Wang, L. Zou, Y. Zhang, L. Zhang, Y. Fang, Y. Li, H-C. Zhou, Stable metal–organic frameworks: design, synthesis and applications, *Adv. Mater.* 2018; 30: 170430.
- [2] A.E. Baumann, D.A. Burns, B. Liu, V.S. Thoi, Metal-organic framework functionalization and design strategies for advanced electrochemical energy storage devices, *Comms Chem.* 2019; 2; 86-99.
- [3] Z. Fang, B. Bueken, D.E. De Vos, R.A. Fischer, Defect-Engineered Metal-Organic Frameworks, *Angew. Chem. Int. Ed.* 2015; 54, 7234-7254.
- [4] G.C. Shearer, J.C. Vtillo, S. Bordiga, S. Svelle, U. Olsbye, K.P. Lillerund, Functionalizing the defects: postsynthetic ligand exchange in the metal organic framework UiO-66, *Chem. mater.* 2016; 28: 7190-7193.
- [5] M. Taddei, R.J. Wakeham, A. Koutsianos, E. Andreoli, A.R. Barron, Post-synthetic ligand exchange in zirconium-based metal-organic frameworks: beware of the defects!, *Angew. Chem. Int. Ed.* 2018; 57: 11706-11710.
- [6] B. Shan, S.M. McIntyre, M.R. Armstrong, Y. Shen, B. Mu, Investigation of Missing-Cluster Defects in UiO-66 and Ferrocene Deposition into Defect-Induced Cavities, *Ind. Eng. Chem. Res.* 2018; 57, 14233-14241.
- [7] L. Yuan, M. Tian, J. Lan, X. Cao, X. Wang, Z. Chai, J.K. Gibson, W. Shi, Defect engineering in metal–organic frameworks: a new strategy to develop applicable actinide sorbents, *Chem. Comm.* 2018; 54: 370-373.

- [8] L. Shen, R. Liang, M. Luo, F. Jing, L. Wu, Electronic effects of ligand substitution on metal-organic framework photocatalysts: the case study of UiO-66, *Phys. Chem. Chem. Phys.* 2015;17:117-121.
- [9] S. Dissegna, K. Epp, W.R. Heinz, G. Kieslich, R.A. Fischer, Defective metal-organic frameworks, *Adv. Mater.* 2018; 30: 1704501.
- [10] J. M. Taylor, S. Dekura, R. Ikeda, H. Kitagawa, *Chem. Mater.* 2015, 27, 72286
- [11] J.H. Cavka, S. Jakobsen, U. Olsbye, N. Guillou, C. Lamberti, S. Bordiga, K.P. Lillerud, A New Zirconium Inorganic Building Brick Forming Metal Organic Frameworks with Exceptional Stability, *J. Am. Chem. Soc.* 2008, 130, 13850-13851.
- [12] M.J. Katz, Z.J. Brown, Y.J. Colon, P.W. Siu, K.A. Scheidt, R.Q. Snurr, J.P. Hupp, O.K. Farha, A facile synthesis of UiO-66, UiO-67 and their derivatives, *Chem. Commun.* 2013, 49, 9449-9451.
- [13] M.R. De Stefano, T. Islamoglu, S.J. Garibay, J.T. Hupp, O.K. Farha, Room-temperature synthesis of UiO-66 and thermal modulation of densities of defects, *Chem. Mater.* 2017;9: 1357-1361
- [14] T.G. Grissom, D.M. Driscoll, D. Troya, N.S. Sapienza, P.M. Usov, M.J. Morris, J.R. Morris, Molecular-level insight into CO₂ adsorption on the Zirconium-based metal-organic framework, UiO-66: a combined spectroscopic and computational approach, *J. Phys. Chem. C.* 2019; 123: 13731-13738.
- [15] H.R. Abid, H. Tian, H-M. Ang, M.O. Tade, C.E. Buckley, S. Wang, Nanosize Zr-metal organic framework (UiO-66) for hydrogen and carbon dioxide storage, *Chem. Eng. J.* 2012; 187: 415-420.

- [16] H. Wu, Y.S. Chua, V. Krungleviciute, M. Tyagi, P. Chen, T. Yildirim, W. Zhou, Unusual and highly tunable missing-linker defects in zirconium metal-organic frameworks UiO-66 and their important effects on gas adsorption, *J. Am. Chem. Soc.* 2013; 135: 10525-10532.
- [17] Y. Cao, Y. Zhao, Z. Lv, F. Song, Q. Zhong, Preparation and enhanced CO₂ adsorption capacity of UiO-66/graphene oxide composites, *J. Ind. Eng. Chem.* 2015;27:102-107.
- [18] A.R. Noorpoor, S. Nazri Kudahi, Analysis and study of CO₂ adsorption on UiO-66/graphene oxide composite using equilibrium modeling and ideal adsorption solution theory (IAST), *J. Environ. Chem. Eng.* 2016;4:1081-1091.
- [19] H. Furukawa, F. Gandara, Y.-B. Zhang, J. Jiang, W.L. Wueen, M.R. Hudson, O.M. Yaghi, Water adsorption in porous metal-organic frameworks and related materials, *J. Am. Chem. Soc.* 2014; 136: 4369-4382.
- [20] P. Ghosh, Y.J. Colon, R.Q. Snurr, Water adsorption in UiO-66: the importance of defects, *Chem. Comm.* 2014; 50:11329-11331.
- [21] Q. Zhao, W. Yuan, J. Liang, J. Li, Synthesis and hydrogen storage studies of metal organic framework UiO-66, *Int. J. Hydrogen Energy.* 2013; 38: 13104-13109.
- [22] B.E.G. Lucier, Y. Zhang, K.J. Lee, Y. Lu, Y. Huang, Grasping hydrogen adsorption and dynamics in metal-organic frameworks using ²H solid-state NMR, *Chem. Comm.* 2016; 52: 7541-7544.
- [23] J. Ren, H.W. Langmi, B.C. North, M. Mathe, D. Bessarabov, Modulated synthesis of zirconium-metalorganic framework (Zr-MOF) for hydrogen storage applications, *Int. J. Hydrogen Energy.* 2014; 39: 890-895.
- [24] L. Zhang, L. Li, E. Hu, L. Yang, K. Shao, L. Yao, K. Jiang, Y. Cui, Y. Yang, B. Li, B. Chen, G. Qian, Boosting ethylene/ethane separation with copper(I)-chelated metal-organic

- frameworks through tailor-made aperture and specific p-complexation, *Adv. Sci.* 2020, 7:1901918.
- [25] X. Wang, L. Li, Y. Wang, J-R. Li, J. Li, Exploiting the pore size and functionalization effects in UiO topology structures for the separation of light hydrocarbons, *Cryst. Eng. Comm.* 2017;19: 1729-1737.
- [26] N. Chanut, A.D. Wiersum, U-H. Lee, Y.K. Hwang, F. Ragon, H. Chevreau, S. Bourrelly, B. Kuchta, J-S. Chang, C. Serre, P.L. Llewellyn, Observing the effects of shaping on gas adsorption in metal organic frameworks, *Eur. J. Inorg. Chem.* 2016; 27:4416-4423.
- [27] N.A. Ramsahye, J. Gao, H. Jobic, P.L. Llewellyn, Q. Yang, A.D. Wiersum, M.M. Koza, V. Guillermin, C. Serre, C.L. Zhong, G. Mauron, Adsorption and diffusion of light hydrocarbons in UiO-66(Zr): a combination of experimental and modeling tools, *J. Phys. Chem. C* 2014; 118: 27470-27482.
- [28] A. Policicchio, M. Florent, M.F. Attia, D.C. Whitehead, J. Jagiello, T.J. Bandoz, Effect of the incorporation of functionalized cellulose nanocrystals into UiO-66 on composite porosity and surface heterogeneity alterations, *Adv. Mater. Interfaces.* 2020, 1902098.
- [29] D.A. Giannakoudakis, T.J. Bandoz, Defectuous UiO-66 MOF Nanocomposites as Reactive Media of Superior Protection against Toxic Vapors, *Chem. Eng. J.* 2020, 3984, 123280
- [30] M. Thommes, K. Kaneko, A.V. Neimark, J.P. Olivier, F. Rodriguez-Reinoso, J. Rouquerol, K.S.W. Sing, *Pure Appl. Chem.* 2015; 87: 1051-1069.
- [31] M. Seredych, J. Jagiello, T.J. Bandoz, Complexity of CO₂ adsorption on nanoporous sulfur-doped carbons: Is surface chemistry an important factor?, *Carbon* 2014; 74: 207-2017

- [32] T.J. Bandoz, K. Putyera, J. Jagiello, J.A. Schwarz, Application of inverse gas chromatography to the study of the surface properties of modified layered minerals, *Micro. Mater.* 1993; 1: 73-79.
- [33] J. Jagiello, J.P. Olivier, Carbon slit pore model incorporating surface energetical heterogeneity and geometrical corrugation, *Adsorption* 2013; 19:777-783.
- [34] L. Czepirski, J. Jagiello, Virial-type thermal equation of gas—solid adsorption, *Chem. Eng. Sci.* 1989; 44: 797-801
- [35] R. Heck, O. Shekhah, O. Zybaylo, P.G. Weidler, F. Friedrich, R. Maul, W. Wenzel, C. Woll, Loading of two related metal-organic frameworks (MOFs), [Cu₂(bdc)₂(dabco)] and (Cu₂(ndc)₂(dabco)], with ferrocene, *Polymers* 2011; 3L 1565-1574.
- [36] Y. Han, M. Liu, K. Li, Y. Zuo, Y. Wei, S. Xu, G. Zhang, C. Song, Z. Zhang, X. Guo, Facile synthesis of morphology and size-controlled zirconium metal–organic framework UiO-66: the role of hydrofluoric acid in crystallization, *Cryst. Eng. Comm.* 2015; 17: 6434-6440.
- [37] C.A. Clark, K.N. Heck, C.D. Powell, M.S. Wong, Highly Defective UiO-66 Materials for the Adsorptive Removal of Perfluorooctanesulfonate, *ACS Sustainable Chem. Eng.* 2019; 7: 6619-6628.
- [38] D.H. Everett, J.C. Powl, Adsorption in slit-like and cylindrical micropores in the Henry's law region, *J. Chem. Soc. Faraday Trans.* 1984; 619-621.
- [39] J. Jagiello, J. Kenvin, A. Celzard, V. Fierro, Enhanced resolution of ultra micropore size determination of biochars and activated carbons by dual gas analysis using N₂ and CO₂ with 2D-NLDFT adsorption models, *Carbon* 2019; 144: 206–215.
- [40] J. Jagiello, J. Kenvin, C.O. Ania, J.B. Parra, A. Celzard, V. Fierro, Exploiting the adsorption of simple gases O₂ and H₂ with minimal quadrupole moments for the dual gas

characterization of nanoporous carbons using 2D-NLDFT models, *Carbon* 2020; 160: 164–175.

[41] A.D. Buckingham, R.L. Disch, D.A. Dunmur, Quadrupole moments of some simple molecules, *J. Am. Chem. Soc.* 1968; 90: 3104-3107.

[42] S. Aguado, G. Bergeret, C. Daniel, D. Farrusseng, Absolute molecular sieve separation of ethylene/ethane mixtures with silver zeolite A, *J. Am. Chem. Soc.* 2012, 134, 14635–14637

[43] A. Ahmed, S. Seth, J. Purewal, A.G. Wong-Foy, M. Veenstra, A.J. Matzger, D.J. Siegel, Exceptional hydrogen storage achieved by screening nearly half a million metal-organic frameworks, *Nat. Commun.* 2019; 10: 1–9.