

Comparison of experimental and quantum chemical studies of the effect of different functionalities of graphene oxide/polymer composites onto selective CO₂ capture

Branislav Stankovic

University of Belgrade

Iranzu Barbarin

University of the Basque Country UPV/EHU

Oihane Sanz

University of the Basque Country UPV/EHU

Radmila Tomovska (✉ radmila.tomovska@ehu.eus)

IKERBASQUE, Basque Foundation for Science

Fernando Ruipérez

University of the Basque Country UPV/EHU

Article

Keywords:

Posted Date: April 8th, 2022

DOI: <https://doi.org/10.21203/rs.3.rs-1522884/v1>

License:   This work is licensed under a Creative Commons Attribution 4.0 International License.

[Read Full License](#)

Abstract

There is a constant need of versatile technologies to reduce the continuously increasing concentration of CO₂ in the atmosphere, able to provide effective solutions under different conditions (temperature, pressure) and composition of the flue gas. In this work, a combination of graphene oxide (GO) and functionalized waterborne polymer particles was investigated, as versatile and promising candidates for CO₂ capture application, with the aim to develop an easy scalable, inexpensive and environmentally friendly CO₂ capture technology.

There are huge possibilities of different functional monomers that can be selected to functionalize the polymer particles and to provide CO₂-philicity to the composite nanostructures. Density functional theory (DFT) was employed to gain a deeper understanding of the interactions of these complex composite materials with CO₂ and N₂ molecules, and build a basis for efficient screening for functional monomers. Estimation of the binding energy between CO₂ and a set of GO/polymer composites, comprising copolymers of methyl methacrylate, n-butyl acrylate, and different functional monomers, shows that it depends strongly on the polymer functionalities. In some cases, there is a lack of cooperative effect of GO. It is explained by a remarkably strong GO-polymer binding, which induced less effective CO₂-polymer interactions. When compared with experimental results, in the cases when the nanocomposite structures presented similar textural properties, the same trends for selective CO₂ capture over N₂ were attained. Besides novel functional materials for CO₂ capture and deeper understanding of the interactions between CO₂ molecules with various materials, this study additionally demonstrates that DFT calculations can be a shorter route toward efficient selection of the best functionalization of the composite materials for selective CO₂ capture.

1. Introduction

Significant and continuous augmentation of the concentration of greenhouse gases in the atmosphere has turned into one of the most fundamental and persistent problems nowadays because fossil fuel reserves are still affordable and developing countries are in the process of economic growth. Between various greenhouse gasses, carbon dioxide (CO₂) is a key player in the global-warming scenario [1]. Even though the global CO₂ capture capacity has reached 40 million tons by 2020, gigatonnes per year of CO₂ have to be captured to have a significant impact on climate change [2]. Since green energy technologies are far away from the point to replace fossil fuels energy sources, the reduction of CO₂ emissions and, thus, the reduction of global warming, is one of the most challenging environmental issues nowadays. Therefore, the development of efficient, selective, and low-cost carbon-capture technologies is crucial [2]. Strategies such as chemical/physical adsorption [3], enzymatic conversion [4], and membrane separation [5] have emerged as potential solutions. Various adsorbents have been proposed for CO₂ capture [6–8], namely, porous polymers, ion-exchange resins, covalent- and metal-organic frameworks, zeolites, silica- and alumina- based materials, metal oxides, etc. However, most of them suffer from low adsorption capacities (or long time is required for saturation), lack of good chemical/thermal stability and/or

selectivity relative to other gases, or they have reduced activity in the presence of moisture, such as zeolite-based adsorbents [9, 10].

Carbon-based adsorbents are arising as a promising alternative to overcome most of the mentioned drawbacks, owing to one of the highest adsorption capacities and relatively low energy requirements for regeneration [6–8]. Furthermore, features such as large surface area, stability in cycle operations, porous structure that can be easily functionalized, and fast adsorption kinetics endorse them as one of the most promising adsorbents. Among these materials, due to lower production costs, graphene and its derivatives have been considered for commercial use [11]. Aimed to further improve the adsorption capacity and separation ability, functionalization of graphene surface with heteroatoms (N, S, O, etc.) has been widely investigated, as well as the production of composites with polymers such as polypyrrole [12], polyaniline [13], polyindole [14], polythiophene [15], mono-, di-, and triethylene-triamine [16], tetraethylenepentamine [17], poly (diallyldimethylammonium chloride) / polystyrene sulfonate [18], poly(dimethylsiloxane) [19], polyether block amide [20], polyethylene-imine [21], and also with metal-organic frameworks [22].

The addition of polymer nanoparticles onto graphene platelets, together with the functionalization of the surface, improves the physicochemical properties and provides easier handling. Such composite platelets are more durable and present improved stability in cycle operations. The polymer particles are usually produced by emulsion polymerization of different combinations of (meth)acrylic monomers, functionalized onto the surface by using a minor amount of functional monomers during the synthesis. The selection of highly CO₂-philic functional monomers may be decisive towards the development of efficient composite platelets for CO₂ selective adsorption and, thus, the screening and evaluation of different monomers is a fundamental step [23, 24].

Computational chemistry has been revealed as a powerful tool in materials science [25, 26]. In particular, density functional theory (DFT) calculations may provide a detailed insight into the interactions between CO₂ and different molecules, giving useful information for the pursuit of potential new adsorbents. One of the first computational studies, regarding the interaction of CO₂ and graphene was performed by Cabrera-Sanfeliu [27]. Author found that CO₂ can be better physisorbed on top of vacancies in the graphene structure. Liu and Wilcox [28] performed more detailed calculations and stated that the CO₂ binding energy on a monovacancy is four-fold higher than that on the defectless graphene. The same authors also showed that hydroxyl and carbonyl groups in graphene establish stronger interactions with CO₂ due to higher electron densities [29]. Sun et al. [30] showed improved adsorption near nitrogen atoms in N-doped graphene and, by means of energy decomposition analyses, demonstrated the relevance of dispersion interactions in this process. Wang et al. [31] performed a similar study on N and O co-doped graphene and established that the heteroatoms bind CO₂ more strongly. They also highlighted the relevant role of dispersion interactions. Dasgupta et al. [32] used higher levels of theory and small molecular models to estimate the binding energy between graphene functionalized with different groups and CO₂. Seema et al. [33] synthesized S-doped microporous carbon materials by chemical activation of

reduced graphene oxide/polythiophene (rGO/PTh) and showed, using DFT calculations, that the interaction energy of CO₂ with thiophene is higher than that with pyrrole, in agreement with the observed higher adsorption capacity of rGO/PTh. These authors suggested that the reason for this stronger attraction is due to a larger negative charge of S in thiophene than that of N in pyrrole. Finally, polymeric systems have also been considered. Patel et al. [34] calculated the binding energies for CO₂ adsorption on several azo-bridged covalent organic polymers. Although simple models were used, accurate trends in binding energy were obtained. In summary, quantum chemical calculations confirm the well-known fact that carbon materials doped with heteroatoms show enhanced CO₂ adsorption. Das et al. [35] investigated the role of surface OH groups in triazine-based N-rich porous organic polymers for enhancing CO₂ capture. Using the ωB97XD density functional method, capable for a reliable description of both short-range and long-range interactions, with, relatively small, 3-21G* basis set, they found that CO₂ mainly interact through the N-N···O and O-N···O hydrogen bonds, but also through the weak N···C, N-O···C, and C-H···O interactions. Ullah et al. [36] performed a combined experimental and theoretical study of CO₂ adsorption by amine and amide porous polymers. Using the B3LYP-D3/6-311++G** level of theory they obtained a moderate agreement with experiments due to the different morphologies of samples.

As it has been previously mentioned, GO/polymer composites are very promising candidates for commercial applications. However, to the best of our knowledge, there is no data in the literature on the theoretical estimation of binding energy trends for the CO₂ adsorption on these composites. Thus, in this work we have performed quantum chemical calculations to estimate the binding energy of CO₂ with a set of GO/polymer composites. More precisely, in the pursuit of new materials for CO₂ capture, we investigated copolymers of methyl methacrylate (MMA), n-butyl acrylate (BA), and different functional monomers. Furthermore, the functional monomers, which, according to the theoretical study provided the strongest interactions with CO₂, have been used for the synthesis of functionalized MMA/BA particles and combined with graphene oxide platelets. Finally, the synthesized composites have been evaluated experimentally for CO₂ capture. Besides, a similar study has been performed for N₂ to evaluate the CO₂/N₂ selectivity of these materials. The observed experimental trends are in accordance with the computational predictions, demonstrating that the DFT calculations are a useful tool for the development of new materials for CO₂ capture.

2. Computational Details

All geometry optimizations were performed within density functional theory (DFT) by using the long-range corrected ωB97XD functional [37] together with the 6-31+G(d) basis set. For the calculations including the GO model, the smaller 6-31G(d) basis set was used to relieve the computational effort. The zero-point vibrational energies (ZPVE) were evaluated within the harmonic oscillator approximation at the same level of theory. All optimized structures showed no imaginary frequencies, meaning that all structures are indeed minima on the potential energy surface. In order to obtain global minima, CO₂ and

N₂ were placed near each of the heteroatom of polymer (i.e. composite) and the structure with the lowest energy (highest binding energy) is presented. Also, when the structure of composites was optimized, the polymer was placed so that as many interactions between functional groups of GO and polymer are achieved. No less than six rearrangements were checked. The electronic energies were then refined by single-point calculations performed on the optimized structures using the 6-311 + + G(2df,2p) basis set. All calculations were carried out with the Gaussian 16 package [38].

The model of GO is constructed to be the smallest one which can guarantee that the polymer remains within the line which connects centroids of outer benzene rings. Graphene is randomly functionalized with two carboxyls, two epoxies, and one hydroxyl group in order to approximate the average functionalization of the real GO, that is, to achieve an adequate ratio between carbon and other atoms, as well as the uniformity of the distribution. In particular, the molecular formula for the model of the GO platelet was C₆₉H₂₂O₇ (Figure S1).

3. Experimental Part

3.1. Materials

An aqueous dispersion of graphene oxide sheets of 4 mg·mL⁻¹ (Graphenea) was used as supplied. The monolayer content in the dispersion was > 95% and in a pH range between 2.2–2.5. The elemental analysis of graphene oxide layers was provided in the technical data sheet from Graphenea: C (49–56%), H (0–1%), N (0–1%), S (2–4%), and O (41–50%). Technical monomers, methyl methacrylate (MMA, Quimidroga) and butyl acrylate (BA, Quimidroga), were used as supplied without any further purification. Sodium 4-vinylbenzenesulfonate (NaSS, Sigma-Aldrich), glycidyl methacrylate (GMA, Acros Organics), 2-hydroxyethyl methacrylate (HEMA, Sigma-Aldrich), and 2-aminoethyl methacrylate hydrochloride (AEMH, Sigma-Aldrich) were used as functional monomers. Tert-butyl hydroperoxide solution (TBHP, Sigma-Aldrich) and L-ascorbic acid (AsA, Sigma-Aldrich) were employed as redox initiators. Furthermore, sodium dodecyl sulfate (SDS, Sigma-Aldrich) and hexadecyltrimethyl ammonium chloride (HAC, Sigma-Aldrich) were employed as emulsifiers. Sodium bicarbonate (NaHCO₃, Sigma-Aldrich) was used as a buffer. Deionized water was used throughout the experimental work.

3.2. Synthesis of polymer nanoparticles

The batch emulsion polymerization process was used for the synthesis of functionalized polymer nanoparticles in aqueous dispersion (polymer latex). As main monomers, MMA and BA in 50/50 weight ratio were used, to which 3 wt% of functional monomer was added, i.e. NaSS, GMA, HEMA, and AEMH. In all cases, the same formulation was employed as described in Table 1, with a solids content of the final aqueous dispersions of 20 wt%. Two different types of surfactants were used, SDS in GMA, HEMA, and NaSS systems and HAC in AEMH system.

Table 1
Formulation for synthesis of
latexes.

Compounds	Amount (g)
MMA	16.49
BA	16.49
FM	1.02
Surfactant	1.02
NaHCO ₃	0.85
TBHP	0.34
AsA	0.34
Water	136

The reactions were performed in a 250 mL jacketed glass reactor equipped with a reflux condenser, temperature probe, nitrogen and feeding inlet, and stainless-steel stirrer rotating at 200 rpm. Pre-emulsion (monomers, surfactant, buffer, and water) was charged in the reactor and then the temperature was raised at 70°C, after which an aqueous solution of TBHP initiator was added as a shot, whereas AsA was fed during 120 min into the reactor. The reaction mixture was then kept at 70°C for an additional 30 min before cooling to room temperature.

3.3. Synthesis of composite GO/polymer platelets

Composite platelets were prepared by simple blending of aqueous dispersions of GO and polymer particles (1:0.5 weight ratio of GO:polymer), which was left agitated overnight at room temperature. Afterward, the hybrid dispersion were subjected to freeze-drying process, in Telstar LyoQuest 55 at -49 °C and 0.2 mbar for 3 days. To check how much polymer was incorporated onto the GO platelets, the aqueous phase of dispersion was analysed gravimetrically. The agitation of GO and polymer mixture was stopped at different time periods. When the agitation was stopped, the platelets precipitated and the residual aqueous phase was analysed. 2 mL of the dispersion extracted at different times was dried in oven. The polymer quantity present in the aqueous phase was calculated from the difference in weight of wet and dry sample. It was found that after 3h agitation of the dispersion mixture of GO and polymer, there was no any polymer left in the dispersion, or with other word, all added polymer was completely incorporated onto the GO platelets.

This method of synthesis was selected in order to prevent development of complex, hierarchical porous morphology, typical for this systems, as we reported previously [23, 24], expecting to produce composite platelets that differ just in the functionalization, introduced by the selected functional monomers. In this way a solid platform for comparison with the theoretical models used for the calculations was provided.

3.4. Characterization

The z-average particle size (d_z) of the polymer particles was measured by Dynamic Light Scattering Spectroscopy (DLS), using the Malvern Zetasizer Nano ZS. Before the measurement, a fraction of the latex was sufficiently diluted with deionized water in order to avoid multiple scattering. Analyses were carried out at a temperature of 25°C. The reported particle sizes are the average of three repeated measurements per sample.

The gel contents of the polymer particles (fraction of polymer insoluble in THF due to the presence of crosslinked and branched polymer chains) were determined by Soxhlet extraction. A few drops of latex were placed on glass fiber square pads and dried overnight at 60°C. Then, the filter, together with the dried polymer, was subjected to a continuous extraction with THF under reflux in the Soxhlet for 24 hours. After that, the filter was dried overnight at 60°C in order to weigh the gel polymer fraction.

The molecular weights corresponding to the soluble fraction of the polymers were determined by gel permeation chromatography (GPC). The soluble part from the Soxhlet extraction was first dried, redissolved in GPC grade THF at a concentration of 2 mg mL⁻¹, and finally, the solution was filtered (polyamide Φ = 45 μ m) before injection into the GPC instrument via an autosampler (Waters 717). The GPC consisted of a pump (LC-20A, Shimadzu), a differential refractometer (Waters 2410), and three columns in series (Styragel HR2, HR4, and HR6, with pores sizes ranging from 10² to 10⁶ Å). The chromatograms were obtained at 35°C using a THF flow rate of 1 mL min⁻¹. The equipment was calibrated using narrow polystyrene standards, thus, the presented molecular weights are relative to this standard.

In terms of graphene-polymer composite materials, thermal stability and the amount of oxygen-containing functional groups presented in the monolithic structure were studied by the TGA500 apparatus (TA instrument). Samples of around 2 mg were heated under a nitrogen atmosphere (90 mL min⁻¹) from 25 to 800°C, at a rate of 10°C min⁻¹.

The surface morphology of the composites was analyzed by scanning electron microscopy (SEM) using Hitachi TM3030 scanning electron microscope at 15 kV after the samples were coated with a thin gold layer, whereas the structure of the composites was analyzed by transmission electron microscopy (TEM) using Tecnai TM G2 20 Twin device at 200 kV (FEI Electron Microscopes). Prior to analysis, the samples were embedded in epoxy resin, from which ultra-thin sections (80 nm) were cut with a diamond knife on Leica EMFC6 ultramicrotome device and places on a 200 mesh copper grid.

The textural properties of the monoliths were characterized by N₂ adsorption-desorption, performed at -196°C in a Micromeritics ASAP 2020. Prior to the measurements, the samples were degassed at 110°C for 8 h under vacuum. From N₂ adsorption-desorption isotherms, the specific surface area and the adsorption average pore width (4V/A) were calculated from the Brunauer–Emmett–Teller (BET) equation. Moreover, the t-plot method was used to estimate the micropore volume. Finally, the pore volume was

calculated using the method proposed by Barrett–Joyner–Halenda. The CO₂ adsorption capacities of the 2D graphene-polymer composites were determined using a TGA analyzer. Prior to the adsorption measurements, the samples were heated to 100°C in N₂ atmosphere at a flow of 50 mL·min⁻¹ and held at that temperature 30 min. The samples were then allowed to cool to 25°C. Once the temperature reached 25°C, the gas was changed to pure CO₂ at a flow rate of 50 mL·min⁻¹ until a constant weight was observed. The weight change of the sample was interpreted as the CO₂ adsorption capacity. The same measurements were performed for N₂ adsorption. These values were corrected taking into account the buoyancy effect of gas change during the measurements.

4. Results And Discussion

In this work, a set of functional monomers, namely, acrylamide (Am), 2-aminoethyl methacrylate hydrochloride (AEMH), 4-bromostyrene (BS), 2-hydroxyethyl methacrylate (HEMA), glycidyl methacrylate (GMA), methyl 2-chloro acrylate (MCIA), and sodium 4-vinylbenzenesulfonate (NaSS), see Fig. 1, have been computationally studied in the pursuit of new materials to improve the CO₂ adsorption capacity of GO/polymer composites. To this end, the binding energy of CO₂ in these materials have been estimated using the theoretical methods of quantum chemistry and the most promising composites have been evaluated experimentally.

The results are organized as follows: (i) the computational studies comprise of the analysis of the binding energy of CO₂ with the bare functional monomers, then with MMA/BA/monomer copolymers, and finally with GO/copolymer composites. These composites have also been used to analyze the binding energy for N₂. (ii) the experimentally evaluated CO₂ and N₂ capture abilities of selected composites.

4.1. Computational analysis

The theoretical investigation was divided into four parts. In the first part, we calculated the binding energies between a set of functional monomers and CO₂. In the second one, it was examined how the introduction of MMA and BA monomers affects the values of binding energy and the trends between them. In the third part, for some of the functional monomers, we estimated the binding energy between CO₂ and GO/MMA/BA/monomer composites, as well as the interaction between the GO platelet and the copolymer. Finally, in the last part, the previously selected composites were used to study the binding energy with N₂.

4.1.1. Interaction between CO₂ and functional monomers

In this subsection, the CO₂ affinity of the functional monomers represented in Fig. 1 was analyzed in terms of binding energies and geometrical features. In Table 2 were collected binding energies and selected geometrical parameters of the complexes formed by CO₂ and the functional monomers. The

optimized geometries are represented in Fig. 2. The binding energy was estimated, using the ZPVE-corrected energies (E_0), as:

$$\Delta E_{\text{CO}_2} = E_0(\text{complex}) - E_0(\text{monomer}) - E_0(\text{CO}_2) \quad (1)$$

These energies include the correction of the basis set superposition error (BSSE) by means of the counterpoise method [39, 40].

Inspecting Table 2, it is observed that the weakest interactions with CO_2 correspond to BS and MCIA monomers (-5.46 and -5.54 $\text{kJ}\cdot\text{mol}^{-1}$, respectively). As a consequence, the C = O bond lengths of CO_2 remain almost unchanged compared to the free molecule (1.165 Å) and the O = C = O angle deviates slightly from linearity (179.0° in both cases). Also, the CO_2 -monomer distances (D) are longer than for other monomers.

Table 2

The binding energy of CO_2 with functional monomers (ΔE_{CO_2}), in $\text{kJ}\cdot\text{mol}^{-1}$. Distance between CO_2 and the monomers (D), in Å. C = O bond length (R_i), in Å, and O = C = O angle (α), in degrees, of CO_2 molecule.

	Am	AEMH	BS	GMA	HEMA	MCIA	NaSS
ΔE_{CO_2}	-16.06	-7.19	-5.46	-10.68	-12.73	-5.54	-29.06
D	2.241	2.934	3.328	3.111	3.037	3.292	3.072
R_1	1.169	1.172	1.164	1.163	1.166	1.166	1.174
R_2	1.161	1.158	1.166	1.166	1.164	1.164	1.156
α	176.3	176.0	179.0	178.5	177.5	179.0	176.2

The highest binding energy corresponds to NaSS (-29.66 $\text{kJ}\cdot\text{mol}^{-1}$), which is reflected in a larger deformation of CO_2 molecule ($R_1 = 1.174$ Å, $R_2 = 1.156$ Å and $\alpha = 176.2^\circ$). Also, the distance between S = O groups and CO_2 is relatively small considering the presence of the sodium cation. Furthermore, the relative position of this cation with respect to the S = O groups is modified after the introduction of CO_2 .

The remaining four functional monomers show intermediate binding energies. Again, for larger binding energies, larger deformations of CO_2 and, in general, smaller distances between this molecule and the monomer are observed. The only clear exception is AEMH, which shows one of the lowest binding energies (-7,19 $\text{kJ}\cdot\text{mol}^{-1}$), but a rather small CO_2 -monomer distance ($D = 2.934$ Å) and remarkable changes in CO_2 geometry ($R_1 = 1.172$ Å, $R_2 = 1.158$ Å and $\alpha = 176.0^\circ$). This monomer comprises a molecular complex between the HCl and the NH_2 moiety. After the adsorption of CO_2 , the molecular geometry of AEMH is significantly changed; the NH_2 -HCl interaction is remarkably affected, in such a way

that the acidic proton is almost transferred to the amino group (the N-H distance is shortened from 1.612 Å to 1.460 Å, while the H-Cl distance is enlarged from 1.377 Å to 1.717 Å). Thus, the capacity of the nitrogen atom to interact with CO₂ is decreased, which is reflected in the low binding energy. Finally, AEMH and NaSS are the only functional monomers for which geometry is changed with adsorption of CO₂, due to the presence of weakly bonded molecule or ions.

4.1.2. Interaction between CO₂ and copolymers containing MMA/BA

In this subsection, the CO₂ affinity of copolymers composed of methyl methacrylate (MMA), butyl acrylate (BA), and the functional monomers are evaluated. The introduction of MMA/BA in the system notably increases the binding energy for CO₂ in all cases (between 4 and 12 kJ·mol⁻¹, depending on the functional monomer), except for Am, see Table 3. This may be explained by the presence of the carbonyl groups of the MMA/BA copolymer, which allows additional interactions with CO₂. The molecular structures of the copolymer-CO₂ complexes are in the Supporting Information (**Figures S1-S7**). It is convenient to say that values of calculated energies are in the range of those which can be found in the literature for CO₂ adsorption by polymers [7, 34, 35], which indicates the size of the model and level of theory are adequate.

Table 3
The binding energy of CO₂ with MMA/BA/monomer copolymers (ΔE_{CO_2}), in kJ·mol⁻¹. Distance between CO₂ and the polymer (D), in Å. C = O bond length (R_i), in Å, and O = C = O angle (α), in degrees, of CO₂ molecule.

	Am	AEMH	BS	GMA	HEMA	MCIA	NaSS
ΔE_{CO_2}	-13.35	-14.68	-9.52	-18.34	-24.24	-14.70	-33.42
D	3.008	2.230	3.697	3.038	2.208	3.074	3.072
R_1	1.163	1.164	1.165	1.164	1.167	1.167	1.174
R_2	1.167	1.150	1.165	1.166	1.163	1.163	1.156
α	176.7	176.5	179.2	177.0	177.4	178.2	175.8

The decrease observed in Am-containing copolymer with respect to the free Am (from -16.06 to -13.35 kJ·mol⁻¹) can be ascribed to the interaction of the NH₂ of Am with the C = O of BA (see **Figure S2**). Now, the amino group is less available to interact with CO₂ and, consequently, CO₂ is displaced closer to the carbonyl group of Am. For AEMH, the presence of MMA/BA allows CO₂ to locate in a position closer to the amino group, and several bonds in AEMH change significantly (see **Figure S2**). For BS, the introduction of MMA and BA also changes the positioning of CO₂, which is now located in a parallel

plane over the benzene ring of BS, allowing for π - π interactions that increase the binding energy (see **Figure S4**). In the case of MClA, CO_2 is now located relatively far from the Cl atom and closer to the carbonyl group, enhancing notably the affinity (see **Figure S5**). For the GMA system, CO_2 is located between the C = O group of BA and the epoxide group of GMA (see **Figure S6**). The CO_2 -BA distance is smaller than the CO_2 -GMA one in the free monomer. As a consequence, the binding energy is increased. Similar to the previously discussed cases, in copolymer with HEMA, CO_2 is adsorbed both by BA and HEMA. In the free monomer, CO_2 mainly interacts with the carbonyl group; however, after including the MMA and BA, CO_2 is located close to both OH group of HEMA and the carbonyl group of BA (see **Figure S7a**) and, thus, the largest increment of the binding energy ($11.51 \text{ kJ}\cdot\text{mol}^{-1}$) is observed. The introduction of CO_2 notably alters the molecular structure of the copolymer (see **Figure S7b**). Finally, in the copolymer including NaSS, the presence of MMA/BA allows the CO_2 to locate in a position where π - π interactions with the benzene ring of functional monomer (i.e. NaSS) are favored, as in the copolymer with BS (see **Figure S8**).

4.1.3 Interaction of CO_2 with GO/copolymer composites

In this section, the performance of the GO/copolymer composites in CO_2 capture is analyzed for the four copolymers that showed the highest affinity with CO_2 , concretely, those which include AEMH, GMA, HEMA, and NaSS functional monomers. In this case, the interaction energy between the GO platelet and the copolymer was estimated, as in the previous section, by using the ZPVE-corrected energies (E_0):

$$\Delta E_{\text{int}} = E_0(\text{composite}) - E_0(\text{GO}) - E_0(\text{copolymer}) \quad (2)$$

The results were collected in Table 4. As in the case of the values from Table 3, calculated energies are similar to those measured and calculated for GO and its composite with polymers [15, 30, 32], which indicates the reliability of the results. It is remarkable how the binding energy is increased for the composite including GMA (from -18.34 to $-53.96 \text{ kJ}\cdot\text{mol}^{-1}$), while a softer increase is found for AEMH (from -14.68 to $-19.32 \text{ kJ}\cdot\text{mol}^{-1}$). However, the binding energy is decreased for the composites including NaSS (from -33.42 to $-23.08 \text{ kJ}\cdot\text{mol}^{-1}$) and HEMA (from -24.24 to $-17.03 \text{ kJ}\cdot\text{mol}^{-1}$). These last two cases are striking since it is well known that GO is a good CO_2 adsorbent [11] and, therefore, a cooperative effect is expected, enhancing the capacity of the copolymer.

Table 4

The binding energy of CO₂ with GO/copolymer composite (ΔE_{CO_2}), in kJ·mol⁻¹. Distance between CO₂ and the composite (D), in Å. C = O bond length (R_i), in Å, and O = C = O angle (α), in degrees, of CO₂ molecule. Interaction energy between GO and the copolymer (ΔE_{int}), in kJ·mol⁻¹.

	AEMH	GMA	NaSS	HEMA
ΔE_{CO_2}	-19.32	-53.96	-23.08	-17.03
D	2.470	1.920	2.402	2.700
R_1	1.171	1.176	1.165	1.165
R_2	1.160	1.158	1.165	1.165
α	175.5	176.7	176.1	177.6
ΔE_{int}	-172.18	-137.71	-224.83	-115.72

The composite with GMA presents the highest binding affinity for CO₂ (-53.96 kJ·mol⁻¹). Without CO₂, the copolymer is oriented in such a way that the carbonyl group of MMA is close to one of the carboxyl groups of GO. The interaction energy of GO and the copolymer is -137.71 kJ·mol⁻¹. However, the introduction of CO₂ significantly changes the positioning of the copolymer and, thus, after the adsorption, the epoxy group of GMA is now able to interact with the acid group of GO, with an O_{epox} - H_{COOH} distance of only 1.721 Å. The CO₂ is placed between the carboxyl group of GO and the carbonyl group of MMA and, thus, a cooperative effect between GO and the copolymer is observed (see **Figure S9**).

When the AEMH-containing copolymer is placed in the GO platelet, the binding energy is increased to a lower extent than in the case of GMA. AEMH is placed in such a way that the Cl⁻ ion interacts with the OH group of the GO surface, while the amino group is close to the carboxyl group. This makes the copolymer-GO interaction energy to be notably higher (-172.18 kJ·mol⁻¹) than for the GMA copolymer. The molecule of CO₂ interacts mainly with the amino group of AEMH, as in the copolymer without GO, weakening the GO-copolymer interaction (the Cl⁻ ...HO distance increases from 2.259 Å to 2.284 Å). Thus, in the presence of CO₂, the copolymer is attached to GO mainly by the interaction between the carbonyl group of AEMH and the nearby carboxyl group. The small increase observed in the binding energy can be due to a rather small model of GO. In **Figure S10** can be observed how the copolymer is covering the whole GO platelet and, therefore, the CO₂ is able to interact mainly only with the copolymer, in such a way that the effect of GO is almost absent. A larger model probably would increase this binding energy.

In the case of NaSS, there is a strong interaction between the copolymer and GO. Na^+ is placed near the epoxy group and one of the oxygens from the sulfonate group interacts with the carboxyl group of GO. Also, BA moiety is oriented so that the carbonyl group is in the vicinity of another epoxy group of the GO surface. The distance between Na^+ and the epoxy group is increased after CO_2 adsorption, i.e. Na^+ moves towards the carboxyl group. Besides, the sulfonate anion moves further from the COOH and $\text{C}=\text{O}$ groups of GO. In the composite, CO_2 is placed closer to the sulfonate group, while in the isolated copolymer there is an interaction with the carbonyl of BA and the aromatic ring of NaSS. Thus, the increase in the interaction energy provided by the GO platelet is compensated by a weaker interaction with the copolymer, and the net effect is a small decrease in binding energy (see **Figure S11**).

Finally, the HEMA composite presents the lowest binding affinity toward CO_2 . Also, of all four investigated composites, one with HEMA has the lowest interaction energy between the GO platelet and the copolymer. The copolymer is placed in such a way that OH groups of HEMA and GO form hydrogen bonds, while oxygen atom from ether moiety of the carboxyl group of MMA forms another with the carboxyl group of GO. Molecule of CO_2 is positioned close to the hydroxyl group of HEMA and in relatively close vicinity of the carbonyl group of BA (see **Figure S12**). After the adsorption, the copolymer moves further from GO. For instance, two mentioned hydrogen bonds change the length from 2.050 Å to 2.124 Å and from 1.831 Å to 1.843 Å, respectively. As in the case of NaSS, CO_2 is positioned far from the GO and interacts mainly with the copolymer. Therefore, as a result, binding energy decreases.

4.1.4. Interaction of N_2 with GO/copolymer composites

In this subsection CO_2/N_2 selectivity of investigated materials was estimated. More precisely, energies by which four composites selected in the previous subsection bind N_2 were calculated. The results were collected in Table 5. The composite with AEMH binds N_2 with the lowest energy. The nitrogen molecule is positioned close to the butyl group of BA and the carbonyl group of MMA (see **Figure S13**). Similarly, in the case of NaSS, N_2 is placed between the butyl group of BA and one of the oxygens from the sulfonate group, to which it is close (see **Figure S14**). Therefore, the binding energy is only slightly higher than in the previously discussed composite. In the composite with HEMA, bonding occurs through the ether group of HEMA and carbonyl group of BA (see **Figure S15**), and thus, since N_2 is placed closely to the two heteroatoms (i.e. oxygens), the energy of adsorption is higher than in the case of the AEMH and the NaSS. Lastly, composite with GMA has the highest binding energy, and nitrogen is placed close to the oxygens of the ether group of GMA and carbonyl group of BA (see **Figure S16**). For any of the composites, the introduction of N_2 does not remarkably change the relative position of the copolymer in the composite. The position at which CO_2 is primarily bound differs from that where N_2 is adsorbed. Moreover, in the case of GMA and AEMN, these two molecules bind CO_2 at significantly different positions.

Table 5
 Binding energy of N₂ with GO/copolymer composite (ΔE_{N_2}), in kJ·mol⁻¹. Distance between N₂ and the composite (D), in Å. N \equiv N bond length (R) of N₂ molecule.

	AEMH	GMA	NaSS	HEMA
ΔE_{N_2}	-6.84	-9.29	-6.99	-8.58
D	2.737	2.809	3.003	2.741
R	1.101	1.101	1.101	1.101

5. Experimental Results

For the experimental work, GMA, AEMH, HEMA, and NaSS were selected as functional monomer added in small amount (3%) to the main monomer mixture made of MMA/BA, to produce functionalized polymer nanoparticles. Table 6 presents characteristics of the polymer particles. Despite that the syntheses were performed under identical conditions, the polymer dispersions have distinct features, which means that the type of functional monomer affected the polymerization process and the polymer properties. NaSS-, GMA- and HEMA- functionalized particles have similar average particle size of about 80 nm, whereas AEMH functionalized particles are much larger and polydispersed. Obviously, the last system was less colloiddally stable during emulsion polymerization, likely due to the cationic surfactant employed for colloidal stabilization in order to avoid possible ionic interactions between the surfactant and AEMH. As a consequence of larger average size of AEMH functionalized particles, shorter polymer chains were produced due to the lower number of radicals per particle that reduced the possibility of bimolecular termination. THF insoluble polymer fraction (gel fraction) that indicates presence of cross-linked and branched structures was very low. The gel fraction of NaSS-containing polymer was slightly increased, which is probably the result of the incorporation of sulfonate moieties within MMA/BA chains, lowering their solubility in THF, especially high molar mass chains. GMA- and HEMA- containing polymers presented an increased amount of insoluble fraction, likely corresponding to crosslinked and branched structures.

Table 6
Characteristic of polymer particles.

	Molar mass M_w (Da)	Polydispersity	THF insoluble fraction (gel) (%)	Particle size (nm)
MMA/BA/NaSS	796524	3.58	12	81.5 ± 0.2
MMA/BA/GMA	1284206	3.6	24	86.7 ± 1.2
MMA/BA/AEMH	428966	4.31	< 5	161.6 ± 1.2
MMA/BA/HEMA	648523	2.08	46	93.5 ± 0.61

The nanocomposites structures were created by mixing of the functionalized polymer particles dispersion with GO nanoplatelets dispersion, during which process the polymer particles were adsorbed onto the platelet surface. By gravimetric analysis of resultant water after the composite platelets were recuperated, it was shown that the whole amount of polymer from the dispersion was incorporated within the composite structure. Therefore, the resulting composites contain GO and polymer in a 1:0.5 weight ratio. In Fig. 3, the morphology of the nanocomposites determined by SEM is presented under two magnifications. Porous materials were obtained in all cases, probably due to hydrophobic interaction of the composite pGO-polymer platelets placed in aqueous dispersion. Namely polymer nanoparticles interacted with the oxygen functional groups at GO surface, creating H bondings (H donor groups such as COOH, OH in GO interacted with H accepting groups in polymers, such as carbonyl, sulfonate or amino). As the oxygen functional groups on GO provide amphiphilic character to GO, by decreasing the number of these functionalities, GO platelets became more hydrophobic. This induced aggregation, wrinkling and crumpling of the platelets, forming the porous structure, which processes were even promoted during drying. However, the presence of different functional monomers, even in as low amounts as 3 wt% with respect to polymer, influences the structure and morphology. Placed onto the polymer particles surface, the different functionalities affected polymer – GO interactions. While nanocomposites functionalized with NaSS, GMA, and HEMA (Figs. 3a c, and g, respectively) have fluffy structures with very well-developed pores of about 5–10 μm , the nanocomposites functionalized with AEHM is less porous (Fig. 3e). The cationic nature of the last probably induced ionic complexing with the numerous anionic oxygen-containing functional groups of GO, giving rise to more compact composite structures. Under higher magnification (Figs. 3b, d, f, and h), no important differences between the four nanocomposites may be noticed, which creates a stable platform for comparison of the interaction forces between CO₂ and the respective functionalities in the nanocomposites.

The structure of the nanocomposites was evaluated by TEM imaging (Fig. 4), where the cross-section of the composite platelets may be observed. The black areas represent the GO platelets, whereas the white areas correspond to the polymer. A peculiar combination of these two phases may be observed, in which the platelets wrap the single polymer particle or aggregates of few of them, creating composite honeycomb-like structures. The presence of GO platelets probably prevented the fulls particle

coalescence and formation of large polymer areas. The thickness of the composite platelets depends on the functional monomer, therefore, the composite platelets functionalized with NaSS present a thickness of about 200 nm, those functionalized with GMA and HEMA have a thickness of 250–300 nm, whereas the AEMH functionalized platelets, with a thickness of 500–1000 nm, are the thickest. Two possible causes can affect the formation of thicker AEMH-containing composite platelets. On one hand, the size of AEMH functionalized polymer particles is double on average in comparison to other functionalized particles, and on the other, the ionic interactions between cationic polymer particles and anionic GO increased the likelihood of platelets aggregation.

TGA curves of the nanocomposites, shown in **Figure S17** in Supporting Information, were used to determine the content of oxygen functionalities in each composite. Thermal degradation occurred in three steps, assigned as follow. The humidity is lost until 100°C, the weight loss between 100 and 260°C corresponds to a loss of oxygen functionalities distributed onto GO, whereas the polymer is degraded between 300-400°C [23]. The advantage that the oxygen functionalities over GO are lost in distinct region than the polymer itself (including the functionalities containing oxygen within the polymer chains), provided possibility to calculate their relative contents. The content of oxygen functionalities (originating from GO) within the composites and their textural properties are presented In Table 7. The quantity of the oxygen functional groups is similar in all composites.

Relatively modest specific BET surface area was observed for all nanocomposites (Table 7), which is not surprising, as the synthesis of the nanocomposites was altered in order to limit the development of the porous structures and to provide a base to investigate the effect of functionalities on the CO₂ selective capture. The observed porosity in SEM images is surface morphology on micron level, while the textural properties from Table 7 demonstrate that no deep meso- and micropores were developed. NaSS functionalized composite presents the lowest BET surface area and total pore volume, indicating that this composite is less porous and more compact than others, which might be due to aromatic ring from NaSS functional monomer that interact more tightly with GO than other polymers. Despite this, only NaSS functionalized composite is characterized by microporosity, although in small quantity. It might provide compensation in case of CO₂ adsorption, as it is known that microporosity is one of determining characteristics for the CO₂ adsorption capacity. [23] The textural properties of GMA and AEMH composites are similar.

Table 7
Textural properties of the composite platelets

Material	% O- functionality	BET Surface area (cm ² g ⁻¹)	Total pore volume (cm ³ g ⁻¹)	Micropore volume (cm ³ g ⁻¹)	Average pore width (nm)
GO/pol- NaSS	15.4	22	0.0364	0.0027	13
GO/pol- GMA	13.3	48	0.0878	< 0.001	12
GO/pol- AEMH	12.5	50	0.081	< 0.001	10
GO/pol- HEMA	14.7	77	0.1249	< 0.001	12

The CO₂ and N₂ adsorption capacity of nanocomposites is given in **Table S1** in Supporting information and in Fig. 5. For comparison, the CO₂ adsorption of neat polymers is also shown. Taking into account that the gas adsorption was studied at atmospheric conditions, CO₂ adsorption is within the range typical for carbon-based nanomaterials (0.5–1 mmol g⁻¹), but it is still much lower than those we previously achieved by similar materials (3.7 mmol g⁻¹) [24]. Surprisingly, contrary to the previous study [24], N₂ adsorption was very similar to that of CO₂ adsorption, indicating that nanocomposites have very low selectivity. The modest capacity for selective CO₂ capture of the present nanocomposites might be due to a few reasons. The most important for this study is that the composite synthesis was directed to obtain possible similar materials and thus to provide a good basis for the comparison using theoretical study.

The amount of CO₂ adsorbed by neat polymer materials is approximately one order of magnitude lower than that adsorbed by nanocomposites. (**Table S1**, Supporting information). Considering that the polymers are not porous, this result is not surprising. Nevertheless, it is clear that a combination of polymers with GO might be a useful way to increase their CO₂ adsorption performance.

The comparison of the experimental results with the theoretical prediction of the binding energy between nanocomposites and CO₂ is not straightforward. Even with the intention to eliminate the effect of the textural characteristics of the synthesized nanocomposites, only GMA and AEMH functionalized nanocomposites presented similar porous morphology, providing a basis for comparison. According to Table 4, GMA functionalized composite presents the highest binding energy, much higher than that of AEMH (54 kJ·mol⁻¹ versus 19 kJ·mol⁻¹). According to **Table S1**, GMA composite adsorbs 0.7 mmol·g⁻¹, whereas AEMH composite adsorbs 0.5 mmol·g⁻¹, thus, the same trend is followed. On the other hand, Table 5 presents that GMA-based composite has N₂ binding energy of almost 10 kJ·mol⁻¹, whereas AEMH composite only of 7 kJ·mol⁻¹. The same trend is followed by experimental results, thus GMA composites adsorbed 0.61 mmol g⁻¹ of N₂, and AEMH adsorbed 0.34 mmol·g⁻¹. Therefore if, by

appropriate design of synthesis procedure, the effect of morphology over the gas absorption capacity of the material can be eliminated, comparison with the theoretical prediction of the adsorption can be performed. In such a case, the presented theoretical study seems to be an excellent tool to predict the interaction of functionalized composite structures with CO₂ and N₂, which can be useful for the selection of functionalization of composites for application in gas adsorption.

Moreover, the comparison of the theoretical prediction and experimental results for HEMA composites provides evidence on the importance of the porous structures' morphology for the selective CO₂ capture. According to Table 4, HEMA functionalized composite presented three-time lower binding energy for CO₂ and slightly lower N₂ binding energy, as shown in Table 5. Experimental results show that this composite presents the highest CO₂ adsorption (1 mmol·g⁻¹) and the highest selectivity over N₂ (Fig. 5). Considering the textural properties of HEMA composites presented in Table 7, i.e. the highest BET surface area and total pore volume, this composite is clearly the most porous when compared to the other studied nanocomposites in this work, and accordingly, it achieved the highest performance for selective CO₂ capture.

6. Conclusions

We have performed a computational and experimental study in the pursuit of new GO/copolymer composites that may be promising candidates for CO₂ capture materials. In particular, the interactions between CO₂ and seven functional monomers, namely, acrylamide (Am), 2-aminoethyl methacrylate hydrochloride (AEMH), methyl 2-chloro acrylate (MClA), hydroxyethyl methacrylate (HEMA), glycidyl methacrylate (GMA), sodium 4-vinylbenzenesulfonate (NaSS), and 4-bromostyrene (BS), have been studied by means of density functional theory (DFT) calculations. The binding energies with CO₂ were analyzed first for the isolated functional monomers and then for the copolymers with MMA/BA, in terms of the interactions with the different moieties and structural rearrangements. The incorporation of MMA/BA enhances the CO₂ affinity due to the presence of the carbonyl groups of the MMA/BA copolymer, allowing for new interactions with CO₂. Only exception is Am, in which case CO₂ do not interact with carbonyl groups. From this study, four copolymers including AEMH, GMA, HEMA, and NaSS have been selected for analysis of adsorption of the GO/polymer composites.

The calculations show that the binding energy for CO₂ is increased for the composites including AEMH and GMA, especially for the latter, while for NaSS and HEMA is decreased. This striking result, where the cooperative effect of the GO seems to be absent, may be explained by a remarkably strong interaction between the copolymer and GO, in such a way that the CO₂, while having limited interaction with GO, now shows a less effective interaction with the copolymer.

These four composites have also been synthesized experimentally and the CO₂ selective adsorption over N₂ has been evaluated. Even with a synthesis procedure designed to decrease the development of highly porous structures, only GMA and AEMH functionalized composites presented very similar textural

structures allowing us to study the effect of different functionalities on the selective CO₂ capture. The trends observed experimentally were exactly predicted by the theoretical study. Moreover, the difference between the theoretical and experimental results in the case of HEMA functionalized composite confirmed the importance of the effect of morphology. In summary, this work shows that DFT is a useful tool for screening of functional monomers for the design and synthesis of new GO-based materials for CO₂ capture.

Declarations

Conflicts of Interest

There are no conflicts of interest to declare.

Data Availability

Some of the datasets used and/or analyzed during the current study are presented in Supporting information file.

The datasets used and/or analyzed during the current study, which are not shown in Supporting information file, are available from the corresponding author on reasonable request.

Acknowledgements

Spanish Government (BES-2017-080221) is gratefully acknowledged for their financial support. The authors would like to acknowledge the contribution of the COST Action CA 15107. The authors thank SGIker (UPV/EHU, ERDF, EU) for technical and human support.

References

1. IPCC, 2018: Global Warming of 1.5°C. An IPCC Special Report on the impacts of global warming of 1.5°C above pre-industrial levels and related global greenhouse gas emission pathways, in the context of strengthening the global response to the threat of climate change, sustainable development, and efforts to eradicate poverty [V. Masson-Delmotte, P. Zhai, H. O. Pörtner, D. Roberts, J. Skea, P.R. Shukla, A. Pirani, W. Moufouma-Okia, C. Péan, R. Pidcock, S. Connors, J. B. R. Matthews, Y. Chen, X. Zhou, M. I. Gomis, E. Lonnoy, T. Maycock, M. Tignor, T. Waterfield (eds.)].
2. B. Smit, J. R. Reimer, C. M. Oldenburg, I. C. Bourg, Introduction to Carbon Capture and Sequestration; Imperial College Press: London, 2014.
3. C.-H. Yu, C.-H. Huang, C.-S. Tan "A review of CO₂ capture by absorption and adsorption", *Aerosol Air Qual. Res.* **12**, 745–769 (2012).
4. J. Shi, Y. Jiang, Z. Jiang, X. Wang, X. Wang, S. Zhang, P. Han, C. Yang "Enzymatic conversion of carbon dioxide", *Chem. Soc. Rev.* **44**, 5981–6000 (2015).

5. G.-X. Dong, H.-Y. Li, V. Chen "Challenges and opportunities for mixed-matrix membranes for gas separation", *J. Mater. Chem. A* **1**, 4610–4630 (2013).
6. A.-H. Lu, G.-P. Hao "Porous materials for carbon dioxide capture" *Annu. Rep. Prog. Chem., Sect. A: Inorg. Chem.* **109**, 484–503 (2013).
7. J. Wang, L. Huang, R. Yang, Z. Zhang, J. Wu, Y. Gao, Q. Wang, D. O'Hare, Z. Zhong "Recent advances in solid sorbents for CO₂ capture and new development trends" *Energy Environ. Sci.* **7**, 3478–3518 (2014).
8. S.-Y. Lee, S.-J. Park "A review on solid adsorbents for carbon dioxide capture" *J. Ind. Eng. Chem.* **23**, 1–11 (2015).
9. T. L. Chew, A. L. Ahmad, S. Bhatia "Ordered mesoporous silica (OMS) as an adsorbent and membrane for separation of carbon dioxide (CO₂)" *Adv. Colloid Interface Sci.* **153**, 43–57 (2010).
10. J.-R. Li, Y. Ma, M. C. McCarthy, J. Sculley, J. Yu, H.-K. Jeong, P. B. Balbuena, H. C. Zhou "Carbon dioxide capture-related gas adsorption and separation in metal-organic frameworks" *Coord. Chem. Rev.* **255**, 1791–1823 (2011).
11. A. T. Najafabadi "Emerging applications of graphene and its derivatives in carbon capture and conversion: Current status and future prospects" *Renew. Sust. Energ. Rev.* **41**, 1515–1545 (2015).
12. V. Chandra, S. U. Yu, S. H. Kim, Y. S. Yoon, D. Y. Kim, A. H. Kwon, M. Meyyappan, K. S. Kim "Highly selective CO₂ capture on N-doped carbon produced by chemical activation of polypyrrole functionalized graphene sheets" *Chem. Commun.* **48**, 735–737 (2012).
13. K. C. Kemp, V. Chandra, M. Saleh, K. S. Kim "Reversible CO₂ adsorption by an activated nitrogen doped graphene/polyaniline material" *Nanotechnology* **24**, 235703 (2013).
14. M. Saleh, V. Chandra, K. C. Kemp, K. S. Kim "Synthesis of N-doped microporous carbon via chemical activation of polyindole-modified graphene oxide sheets for selective carbon dioxide adsorption" *Nanotechnology* **24**, 255702 (2013).
15. H. Seema, K. C. Kemp, N. K. Le, S.-W. Park, V. Chandra, J. W. Lee, K. S. Kim "Highly selective CO₂ capture by S-doped microporous carbon materials" *Carbon* **66**, 320–326 (2014).
16. Y. Zhao, H. Ding and Q. Zhong "Preparation and characterization of aminated graphite oxide for CO₂ capture" *Appl. Surf. Sci.* **258**, 4301–4307 (2012).
17. S. Gadipelli, Y. Lu, N. T. Skipper, T. Yildirim, Z. Guo "Design of hyperporous graphene networks and their application in solid-amine based carbon capture systems" *J. Mater. Chem. A* **5**, 17833–17840 (2017).
18. J. Heo, M. Choi, J. Chang, D. Ji, S. W. Kang, J. Hong "Highly permeable graphene Oxide/polyelectrolytes hybrid thin films for enhanced CO₂/N₂ separation performance" *Sci. Rep.* **7**, 456 (2016).
19. H. Ha, J. Park, S. Ando, C. B. Kim, K. Nagai, B. D. Freeman, C. J. Ellison "Gas permeation and selectivity of poly(dimethylsiloxane)/graphene oxide composite elastomer membranes" *J. Membr. Sci.* **518**, 131–140 (2016).

20. J. Shen, G. Liu, K. Huang, W. Jin, K.-R. Lee, N. Xu "Membranes with fast and selective gas-transport channels of laminar graphene oxide for efficient CO₂ capture" *Angew. Chem. Int. Ed. Engl.* **127**, 588–592 (2015).
21. F.-Q. Liu, W. Li, J. Zhao, W.-H. Li, D.-M. Chen, L.-S. Sun, L. Wang, R.-X. Li "Covalent grafting of polyethyleneimine on hydroxylated three-dimensional graphene for superior CO₂ capture" *J. Mater. Chem. A* **3**, 12252–12258 (2015).
22. R. Kumar, D. Raut, U. Ramamurty, C. N. Rao "Remarkable improvement in the mechanical properties and CO₂ uptake of MOFs brought about by covalent linking to graphene" *Angew. Chem. Int. Ed.* **55**, 7857–7861 (2016).
23. N. Politakos, I. Barbarin, L. Serrano Cantador, J.A. Cecilia, E. Mehravar, R. Tomovska "Graphene-based monolithic nanostructures for CO₂ capture" *Ind. Eng. Chem. Res.* **59**, 8612–8621 (2020)
24. N. Politakos, I. Barbarin, T. Cordero-Lanzac, A. Gonzalez, R. Zangi, R. Tomovska "Reduced graphene oxide/polymer monolithic materials for selective CO₂ capture" *Polymers* **12**, 936, (2020)
25. J. A. Elliott "Novel approaches to multiscale modelling in materials science" *Int. Mater. Rev.* **56**, 207–225 (2011)
26. F. Ruipérez "Application of quantum chemical methods in polymer chemistry" *Int. Rev. Phys. Chem.* **38**, 343–403 (2019)
27. P. Cabrera-Sanfeliix "Adsorption and reactivity of CO₂ on defective graphene sheets" *J. Phys. Chem. A* **113**, 493–498 (2009).
28. Y. Liu and J. Wilcox "CO₂ adsorption on carbon models of organic constituents of gas shale and coal" *Environ. Sci. Technol.* **45**, 809–814 (2011).
29. Y. Liu, J. Wilcox "Effects of surface heterogeneity on the adsorption of CO₂ in microporous carbons" *Environ. Sci. Technol.* **46**, 1940–1947 (2012).
30. F. Sun, X. Liu, J. H. Gao, X. X. Pi, L. J. Wang, Z. B. Qu, Y. K. Qin "Highlighting the role of nitrogen doping in enhancing CO₂ uptake onto carbon surface: A combined experimental and computational analysis" *J. Mater. Chem. A* **4**, 18248–18252 (2016).
31. Dasgupta, S. N. Punnathanam, K. G. Ayappa "Effect of functional groups on separating carbon dioxide from CO₂/N₂ gas mixtures using edge functionalized graphene nanoribbons" *Chem. Eng. Sci.* **121**, 279–291 (2015).
32. Y. Wang, X. Hu, J. Hao, R. Ma, Q. Guo, H. Gao H. Bai "Nitrogen and oxygen codoped porous carbon with superior CO₂ adsorption performance: A combined experimental and DFT calculation study" *Ind. Eng. Chem. Res.* **58**, 13390–13400 (2019).
33. H. Seema, K. C. Kemp, N. H. Le, S.-W. Park, V. Chandra, J. W. Lee, K. S. Kim "High selective CO₂ capture by S-doped microporous carbon materials" *Carbon* **66**, 320–326 (2014).
34. H. A. Patel, S. H. Je, J. Park, Y. Jung, A. Coskun, C. T. Yavuz "Directing the structural features of N₂-phobic nanoporous covalent organic polymers for CO₂ capture and separation" *Chem. Eur. J.* **20**,

772–780 (2014).

35. S. K. Das, P. Bhanja, S. K. Kundu, S. Mondal, A. Bhaumik " Role of surface phenol-OH groups in N-rich porous organic polymers for enchancing the CO₂ uptake and CO₂/N₂ selectivity: Experimental and computational study" *ACS Appl. Mater. Interfaces* **10**, 23812–23824 (2018).
36. R. Ullah, H. Patel, S. Aparicio, C. T. Yavuz, M. Atilhan" A combined experimental and theoretical study on gas adsorbtion performance of amine and amide porous polymers" *Micropor. Mesopor. Mat.* **279**, 61–72 (2021).
37. J.-D. Chai, M. Head-Gordon "Long-range corrected hybrid density functionals with damped atom–atom dispersion corrections" *Phys. Chem. Chem. Phys.* **10**, 6615–6620 (2008).
38. Gaussian 16, Revision C.01, M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, G. A. Petersson, H. Nakatsuji, X. Li, M. Caricato, A. V. Marenich, J. Bloino, B. G. Janesko, R. Gomperts, B. Mennucci, H. P. Hratchian, J. V. Ortiz, A. F. Izmaylov, J. L. Sonnenberg, D. Williams-Young, F. Ding, F. Lipparini, F. Egidi, J. Goings, B. Peng, A. Petrone, T. Henderson, D. Ranasinghe, V. G. Zakrzewski, J. Gao, N. Rega, G. Zheng, W. Liang, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, K. Throssell, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, T. A. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, J. M. Millam, M. Klene, C. Adamo, R. Cammi, J. W. Ochterski, R. L. Martin, K. Morokuma, O. Farkas, J. B. Foresman, and D. J. Fox, Gaussian, Inc., Wallingford CT, 2016.
39. S. F. Boys, F. Bernardi "The calculation of small molecular interactions by the difference of separate total energies. Some procedures with reduced errors" *Mol. Phys.* **19**, 553–566 (1970).
40. S. Simon, M. Duran, J. J. Dannenberg "How does basis set superposition error change the potential surfaces for hydrogen bonded dimers?" *J. Chem. Phys.* **105**, 11024–11031 (1996).

Figures

Figure 1

Molecular structures of the monomers: acrylamide (Am), 2-aminoethyl methacrylate hydrochloride (AEMH), methyl 2-chloro acrylate (MCIA), hydroxyethyl methacrylate (HEMA), glycidyl methacrylate (GMA), sodium 4-vinylbenzenesulfonate (NaSS), 4-bromostyrene (BS), butyl acrylate (BA), and methyl methacrylate (MMA).



Figure 2

Optimized geometries of the CO₂-monomer complexes.

Figure 3

Morphology of the GO/polymer composites with different functional monomers under different magnification: (a) and (b) NaSS; (c) and (d) GMA; (e) and (f) AEMH; (g) and (h) HEMA.



Figure 4

TEM images of (a) NaSS; (b) GMA; (c) AEMH; and (d) HEMA functionalized composites. Scale bars in all images are of 500 nm.

Figure 5

Adsorption of CO₂ and N₂ of the functionalized GO/polymer composites

Supplementary Files

This is a list of supplementary files associated with this preprint. Click to download.

- [Supportinginformation.docx](#)