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Poly-cyclodextrin modified graphene oxide for PFAS removal from drinking water

Andrea Trifoglio,[†]^a Angela Pintus,[†]^{af} Sara Khaliha,[†]^a Francesco Blancato,^a Marco Agnes,^a Lucrezia Aversa,^b Roberto Verucchi,^b Tainah Dorina Marforio,[†]^{eg} Nina Burduja,^{cd} Giuseppe Nocito,^c Antonino Mazzaglia,[†]^c Matteo Calvaresi,[†]^{eg} and Manuela Melucci[†]^{*a}

Per- and polyfluoroalkyl substances (PFASs) are persistent organic pollutants of growing concern due to their widespread occurrence in drinking water and resistance to conventional remediation technologies. Granular activated carbon (GAC), the current benchmark adsorbent, exhibits limited efficiency toward short- and medium-chain PFASs and slow adsorption kinetics. Here, we report the synthesis of a graphene oxide (GO) material covalently grafted with an amino-functionalized β -cyclodextrin polymer crosslinked with epichlorohydrin (GO-Poly- β CD). The hybrid material was characterized and evaluated for the adsorption of a mixture of PFASs with varying chain lengths (CF)_{3–9} and functional groups under environmentally relevant conditions. GO-Poly- β CD outperformed pristine GO, showing enhanced affinity for medium-chain PFAS and removal efficiencies up to 97% for PFHpA (CF)₆ and PFHxS (CF)₆ after only 15 min of contact. At neutral pH, PFPeA (CF)₄ removal reached 44%, compared to no removal by GO, while under acidic conditions its removal increased to 80%. Notably, PFBA (CF)₃, which was not removed at neutral pH, exhibited 31% removal under acidic conditions. Molecular dynamics simulations revealed a cooperative adsorption mechanism in which PFAS molecules are stabilized through pH-responsive conformational rearrangements that strengthen interactions between the hybrid material and the contaminants, primarily via van der Waals and hydrophobic forces. Compared to GAC, GO-Poly- β CD achieved a higher overall PFAS uptake (4.0 $\mu\text{g g}^{-1}$ vs. 1.3 $\mu\text{g g}^{-1}$ for GAC) within the same contact time, demonstrating its potential for rapid and efficient PFAS removal in real drinking water treatment applications.

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Introduction

Per- and poly-fluoro alkyl substances (PFASs) have attracted global concern due to their widespread use in everyday products, their environmental and biological persistence and widely reported effects on human health and ecosystems.^{1–3} PFASs are a class of synthetic organo-fluorine compounds characterized by the presence of fully or partially fluorinated alkyl-chains with a carboxylate or sulphonate ending group.⁴ PFASs are widely

used as additives in food packaging, coatings, non-stick cookware, waterproof clothing and firefighting foams.⁵ The disposal of everyday products inevitably results in the release of PFASs into various environmental compartments and, ultimately, into drinking water sources through multiple pathways.¹ The C–F chemical bond being very stable, their persistency in the environment is exceptional making PFASs the so-called “forever chemicals”.⁶ The Environmental Working Group reported that in 2018 over 16 million people in the United States were exposed to PFAS contaminated drinking water with concentrations exceeding 10 ng L^{−1}.³ The removal of PFASs from drinking water remains one of the key challenges in water purification. Among the treatment strategies currently implemented in municipal drinking water plants, adsorption is the most effective approach. In particular, granular activated carbon (GAC) represents the most widely employed adsorbent for PFAS mitigation.^{7–9} GAC is a non-specific adsorbent that exhibits moderate affinity toward long-chain PFASs but limited effectiveness for their short- and medium-chain counterparts.^{10,11} Moreover, its adsorption process is characterized by slow, diffusion-limited kinetics, which represents a significant

^aInstitute for Organic Synthesis and Photoreactivity (ISOF-CNR), Bologna, Italy.
E-mail: manuela.melucci@cnr.it

^bInstitute of Materials for electronics and Magnetism (IMEM-CNR), c/o Fondazione Bruno Kessler, Trento, Italy

^cInstitute for Nanostructured Materials (ISMN-CNR), Messina, Italy

^dDepartment of Chemical, Biological, Pharmaceutical and Environmental Sciences (ChiBioFarAm), University of Messina, Messina, Italy

^eDepartment of Chemistry ‘Giacomo Ciamician’, University of Bologna, Italy

^fDipartimento di Physics, University of Modena e Reggio Emilia, Modena, Italy

^gIRCCS Azienda Ospedaliero – Universitaria di Bologna, Preclinical & Translational Research in Oncology Lab (PRO), Bologna, Italy

[†] These authors contributed equally.



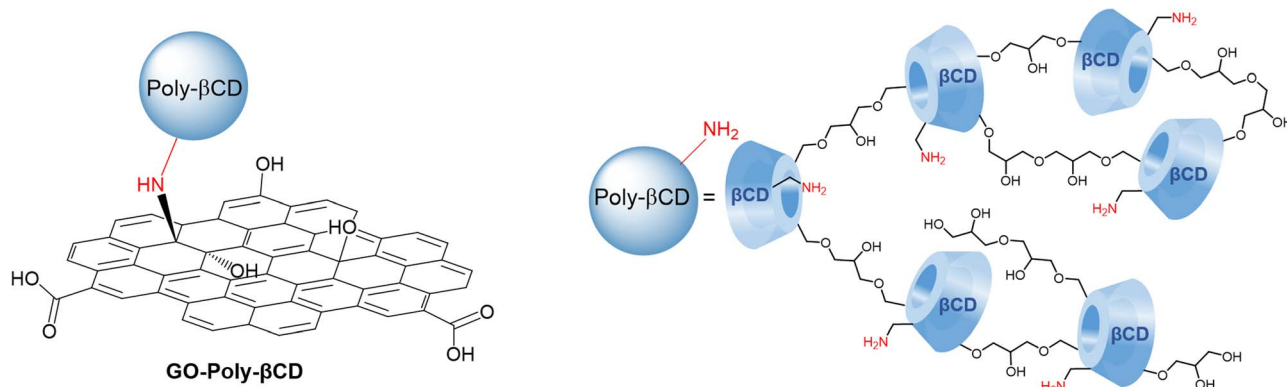


Fig. 1 Chemical structure of GO-Poly- β CD and Poly- β CD.

drawback under real plant flow conditions.^{12,13} New technologies for GAC replacement or integration are currently a matter of great interest. Among emerging materials, graphene oxide (GO) nanosheets have attracted considerable attention as promising adsorbents for the removal of emerging contaminants from drinking water, offering new opportunities for advanced treatment technologies.^{10,14} The most relevant features of GO are its large specific surface area and the abundance of tunable oxygen-containing functional groups on the nanosheet surface, which enable versatile covalent functionalization.¹⁵

GO nanosheets have previously been explored for PFAS removal using a two-step approach involving batch adsorption followed by tandem microfiltration.¹⁰ Furthermore, some of us demonstrated that incorporating GO into polysulfone hollow fiber membranes (now commercially available as Graphisulfone®¹⁶) provides a new integrated filter model combining adsorption and ultrafiltration mechanisms and allowing the removal of long-chain PFASs from drinking water.¹³ As a step forward, it has been demonstrated that the covalent functionalization of GO nanosheets enables the tuning of adsorption selectivity and significantly enhances the removal of contaminants that exhibit poor affinity for unmodified GO, including organic emerging pollutants with performances overcoming those achieved by GAC.^{10,17,18} For instance, functionalization of GO with *N,N*-dimethylethylenediamine (GO-DMEN),¹⁹ provided nanosheets with high affinity for medium- and long-chain PFASs. Studies based on theoretical modelling demonstrated that adsorption depends on multiple interactions including hydrophobic, electrostatic and van der Waals forces.¹³ In particular, the interaction between non-functionalized GO and PFASs is hindered by electrostatic repulsion between the negatively charged nanosheets and the negatively charged PFAS molecules.¹⁶ Hence, PFASs with longer hydrophobic chains are more efficiently adsorbed, whereas shorter, more hydrophilic PFASs are not removed.

Cyclodextrins are cyclic oligosaccharides made up of glucose subunits linked by α -1,4-glycosidic bonds. They have a toroidal structure with a hydrophobic inner cavity and a hydrophilic outer surface.²⁰ Cyclodextrins,^{21,22} in particular β -cyclodextrins^{23,24} (*i.e.*, consisting of seven glucose subunits), can form inclusion complexes with PFASs through non-covalent forces such as van der Waals interactions and hydrophobic effects. Some of us

recently demonstrated that covalent modification of GO with an amino-derived β CD and a C6 aliphatic linker to GO nanosheets enables the efficient removal of perfluorobutanoic acid (PFBA, (CF)₃)²⁵ thanks to the formation of a sandwich-like complex, where PFBA is stably confined between the GO nanosheet and the β CD cavity. Interestingly, β CD can serve as a monomer for the synthesis of β CD-based polymers through various types of cross-linkers.²⁶ PFAS adsorption has been demonstrated for β CD polymers containing fluorinated aromatic cross-linkers,^{27,28} which raise environmental concerns similar to those of the targeted PFASs.²⁹ Moreover, the partial water solubility of polycyclodextrins makes their direct application as sorbents under real conditions unrealistic.³⁰ To overcome this limitation, we propose here a novel polymeric material insoluble in water and based on a non-fluorinated chain (Poly- β CD from now on) and its use for PFAS removal from drinking water (Fig. 1). GO-Poly- β CD adsorption mechanism was investigated by combining adsorption experiments under different pH conditions with molecular dynamics (MD) simulations.

Experimental section

Materials

GO was purchased from Layer One (Norway) and used without further purification. Before using, GO was sonicated in ultrapure water to exfoliate the bulk material into a monolayer (>99%) with a lateral size of few micrometers. GAC was purchased from CABOT Norit Spa (Ravenna, Italy, Norit GAC 830 AF, MB index min 240 mg g⁻¹, BET surface area >1000 m² g⁻¹) and used without further purification. PFAS standards were purchased from Agilent Technologies (Santa Clara, CA, US). The experiments on PFASs were carried out by using polypropylene vials. Poly- β CD was kindly provided by Cyclolab (Budapest).

Synthesis of poly- β -cyclodextrin (Poly- β CD)

Poly- β CD was provided as the chlorohydrated salt of amino β -cyclodextrin polymer crosslinked with epichlorohydrin (average M_w = 25 kDa) and with 70% of β -cyclodextrin content. The salt was treated with NaHCO₃ and dialysed for 72 hours in ultrapure water then lyophilized and used in the next synthetic step.



Synthesis and purification of GO-Poly- β CD

The synthesis of GO-Poly- β CD was carried out *via* an epoxide ring-opening reaction.²⁵

Briefly, 100 mg of GO was dispersed in 30 mL of a 1 : 1 H₂O : EtOH solution by sonication for 2 hours to obtain a stable aqueous suspension. Separately, 200 mg of Poly- β CD-NH₂ was dissolved in 100 mL of distilled water and added to the GO suspension (GO : Poly- β CD mass ratio 1 : 2, total volume 130 mL). The resulting mixture was refluxed at 80 °C for 12 hours. Upon completion of the reaction, the crude product was purified by centrifugation (9000 rpm, 15 minutes per cycle, eight cycles), and the final material was obtained by lyophilization.

Synthesis and purification of GO-Control

The synthesis of GO-Control was carried out under the same conditions as GO-Poly- β CD, but in the absence of Poly- β CD-NH₂. Briefly, 100 mg of GO was dispersed in 130 mL of a 1 : 1 H₂O : EtOH solution by sonication for 2 h to obtain a stable suspension. The mixture was then refluxed at 80 °C for 12 h. After completion of the reaction, the crude product was purified by centrifugation (9000 rpm, 15 min per cycle, eight cycles) and lyophilized.

Characterization

ATR-FTIR spectra were recorded with an Agilent Cary 630 FTIR spectrophotometer, and the spectra are expressed by wave-number (cm⁻¹).

TGA was conducted with a PerkinElmer Thermogravimetric Analyzer TGA 4000 by PerkinElmer, in an air atmosphere, from 30 °C to 800 °C, with a scanning temperature of 10 °C min⁻¹.

Elemental analysis was performed on powder materials by using an Elementar Unicube Elemental analyser, method GRAPHITE.

XPS was performed using the Mg-K α emission at 1253.6 eV of a non-monochromatized X-ray photon source. Photoelectrons were analysed using a VSW HA100 electron energy analyser, resulting in a total energy resolution of 0.86 eV. Core level BEs were referred to the Au 4f_{7/2} core level signal (at 84.0 eV), obtained from a sputtered gold surface. The photoemission line shapes of selected core levels were analyzed using Voigt line shape deconvolution, after background subtraction of a Shirley function. The typical precision for energy peak position is ± 0.05 eV, while the uncertainty for FWHM is $< \pm 5\%$ and for area evaluation it is $\pm 5\%$. GO, GO-Control, and GO-Poly- β CD samples were mounted directly on the sample holder, while the Poly- β CD-NH₂ powder was supported on carbon tape.

Dynamic light scattering (DLS)

The mean hydrodynamic diameter (D_H), width of distribution (polydispersity index, PDI) and ζ -potential were measured with a Malvern Zetasizer Nano ZS (Malvern Instruments, Malvern, UK), equipped with a He-Ne laser ($\lambda = 633$ nm, 4 mW). Measurements were performed using the non-invasive back-scattering technique (NIBS) with an angle of 173° to the incident beam. For freshly prepared dispersions, 45 acquisitions at a temperature of 25 ± 1 °C for each measurement were collected. Deconvolution of the measured correlation curve

(distribution of relaxation times and the cumulant analysis) into a dimensional average intensity distribution was achieved using the Laplace inversion algorithm.

Colloidal stability

The colloidal stability of GO-Poly- β CD was evaluated with an adapted turbidimetric method for a 250 $\mu\text{g mL}^{-1}$ dispersion in aqueous 1 mM NaCl. The sample was placed in a 1 cm quartz cuvette inside a Jasco V-770 (Jasco Co., Tokyo, Japan) Peltier temperature-controlled UV/Vis-NIR scanning spectrophotometer immediately after preparation. Experiments were performed by recording the extinction values at 790 nm every 5 min in a time interval from 0 to 600 min at 25 °C, respectively. Results were plotted against time and fitted using the OriginPro 2021 software with the equation

$$y = y_0 + A_1 e^{\frac{-x}{\tau_1}} + A_2 e^{\frac{-x}{\tau_2}} \quad (1)$$

for a two-phase exponential decay function. The half-life parameters were calculated from the following formulae

$$(t_{1/2})_1 = \tau_1 \ln 2 \quad \text{and} \quad (t_{1/2})_2 = \tau_2 \ln 2 \quad (2)$$

Adsorption experiments

25 mg of powdered materials (GO or GO-Poly- β CD) was sonicated in 25 mL of tap water for 2 hours. A mixture containing ten PFASs (125 μL , 100 $\mu\text{g L}^{-1}$) was then added to the suspensions to achieve a final concentration of 0.5 $\mu\text{g L}^{-1}$ each in a total volume of 25 mL. Similarly, 25 mg of GAC was directly added to 25 mL of tap water, followed by the addition of 125 μL of the ten-PFAS mixture (100 $\mu\text{g L}^{-1}$), resulting in the same final PFAS concentration (0.5 $\mu\text{g L}^{-1}$). All samples were subjected to gentle agitation for 15 minutes, then centrifuged (10,000 rpm, 10 minutes) and analyzed by UPLC-MS/MS.

PFAS quantification

Samples containing PFASs were analyzed by UPLC-MS/MS (ACQUITY UPLC H-Class PLUS coupled to a XEVO TQS Micro mass detector, Waters). A 1 mL aliquot of each sample was used for automated injection. Chromatographic separation was performed on a Waters Acquity UPLC CSH Phenyl-Hexyl column (1.7 μm , 2.1 \times 100 mm) with a Waters Isolator Column (2.1 \times 50 mm). The column temperature was maintained at 34 °C, the flow rate was 0.3 mL min⁻¹, and the injection volume was 40 μL . Total run times were 8 min for PFBA and 21 min for the mixture of eight PFASs. The mobile phase consisted of a biphasic gradient: 2 mM NH₄OAc in ultrapure water : methanol (95 : 5) as phase A, and 2 mM NH₄OAc in methanol as phase B (for further details, see SI Section 4).

Computational modelling

All MD simulations were carried out using the Amber22 software suite. The PFAS molecules were parameterized using the generalized AMBER force field (GAFF), and atomic partial charges were derived using the restrained electrostatic potential (RESP) fitting procedure after geometry optimization at the HF/6-31G* level of theory. The GO-Poly- β CD model-system was generated by



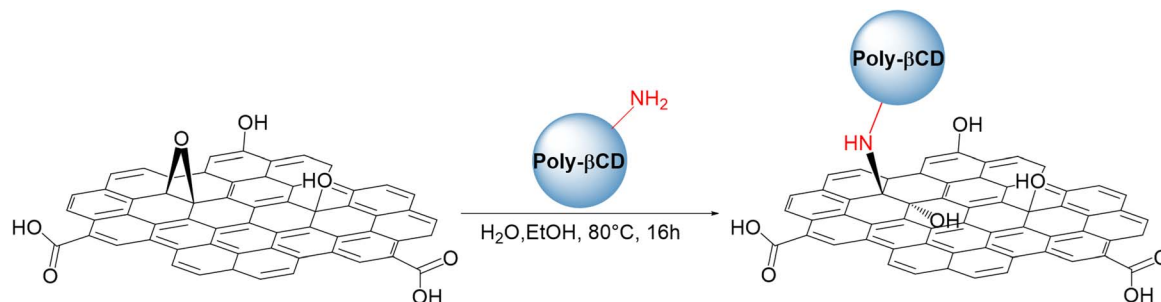
covalently grafting a representative Poly- β CD onto a GO surface, modelled according to experimental XPS data. Different protonation states of the GO-Poly- β CD system were used to mimic pH 4, 7, and 10, where surface groups were protonated or deprotonated accordingly. Complexes of GO-Poly- β CD with PFASs were solvated in an explicit TIP3P water box, ensuring a 20 Å buffer from the solute to the box edge. The systems were neutralized by adding appropriate counterions. Energy minimization was performed in two steps: first with restraints on solute atoms to relax the solvent environment, and second without restraints to minimize the entire system. Subsequently, each system was equilibrated under NPT conditions at 298 K for 10 ns using periodic boundary conditions and an Andersen thermostat. Production MD simulations were then run for 100 ns under constant temperature and pressure conditions. To evaluate binding affinities, Molecular Mechanics-Generalized Born Surface Area (MM-GBSA) calculations were performed and the total binding free energy ($\Delta G_{\text{binding}}$) was decomposed into van der Waals (VDW), electrostatic (E_{El}), non-polar solvation (E_{SURF}), and entropic ($T\Delta S$) contributions. Entropic contributions were estimated *via* normal mode analysis. The cavities reported in this study were identified using ChimeraX software (version 1.10.1) with the Cavity tool.

Results and discussion

Synthesis and characterization

The synthesis of GO-Poly- β CD was carried out *via* the epoxide ring-opening reaction of exfoliated GO nanosheets,²⁵ using Poly- β CD-NH₂ obtained by crosslinking of mono-6-azido β -CD in the presence of epichlorohydrin and subsequent reduction of azido to amino groups³¹ (Scheme 1). Under the reaction conditions (80 °C, 16 h), GO nanosheets are partially reduced, thus to clarify the role of Poly- β CD, a control material (GO-Control) was prepared by treating GO under the same conditions as those used for GO-Poly- β CD synthesis, but in the absence of Poly- β CD. The resulting materials were purified from unreacted materials through repeated washing and centrifugation steps with water until the supernatant reached neutral pH. The lyophilized products were then characterized and used for the targeted adsorption experiments. The surface chemical properties of pristine GO, GO-Control, GO-Poly- β CD, and Poly- β CD-NH₂ were investigated by X-ray photoelectron spectroscopy (XPS). Long-range spectra (Fig. S1) revealed the presence of carbon, oxygen, and nitrogen in almost all samples; therefore, the

corresponding core levels (C 1s, N 1s, and O 1s) were analyzed in detail. Fig. 2a and b reports the C 1s and N 1s core levels of all samples while O 1s core level is shown in Fig. S2. GO and GO-Control surfaces were found to be almost identical, exhibiting the typical C=C sp² component (285.0 eV), ascribed to the aromatic graphene framework, together with C-O, C=O, and COOR features (287.0–287.1 eV, 288.4–288.7 eV, and 289.3 eV, respectively). The N 1s intensity was very low (<1% atomic concentration) and was detected only in GO-Control. Its deconvolution yielded a C-NH-C signal (graphitic nitrogen) at 400.1 eV, as previously reported,³² and two additional components at 398.5 eV and ~402.0 eV, which can be assigned to pyridinic nitrogen³² and to amino groups,²⁵ respectively, although the latter attribution is not straightforward since different N-based functionalities contribute in this range. Poly- β CD-NH₂ displayed a completely different C 1s line shape, dominated by C-O bonds at 286.4 eV, and a single N 1s peak corresponding to primary amines at 399.8 eV. Upon functionalization of GO, a complete change in the C 1s line shape was observed, along with an increase in the intensity of high-binding-energy N 1s components. The relative contribution of graphitic and amino-related components in N 1s increased from 43.1% in pristine GO to 74.8% in GO-Poly- β CD. These changes are compatible with the covalent functionalization of GO with poly- β CD.^{19,25} Thermogravimetric analysis (TGA) supported the successful covalent bonding of GO with Poly- β CD. The TGA profile of GO-Poly- β CD (Fig. 2c) exhibited a distinct inflection point at 333 °C, clearly visible as a peak in the derivative thermogravimetric (DTG) curve. This thermal event was absent in pristine GO and closely matched the degradation onset of pure Poly- β CD (Fig. 2e), which showed a similar DTG peak at 342 °C. The slight shift in decomposition temperature suggested covalent interaction between GO and Poly- β CD, which altered the thermal stability of the polymer moieties. Additionally, the increased weight loss observed in the 200–400 °C range for GO-Poly- β CD compared to pristine GO (Fig. 2d) supported the presence of organic functional groups from the grafted polymer. Elemental Analysis (EA) was used to estimate the bulk composition of the synthesised materials and to evaluate the Poly- β CD loading (Table S1). GO-Poly- β CD exhibits a higher nitrogen content compared to pristine GO (0.3% *vs.* 0.08%), ascribed to the presence of amine functionalities in the glucose units. Under the reaction conditions (16 hours at 80 °C), the GO nanosheets tend to undergo partial reduction¹⁵ (GO-



Scheme 1 Synthetic route to GO-Poly- β CD.



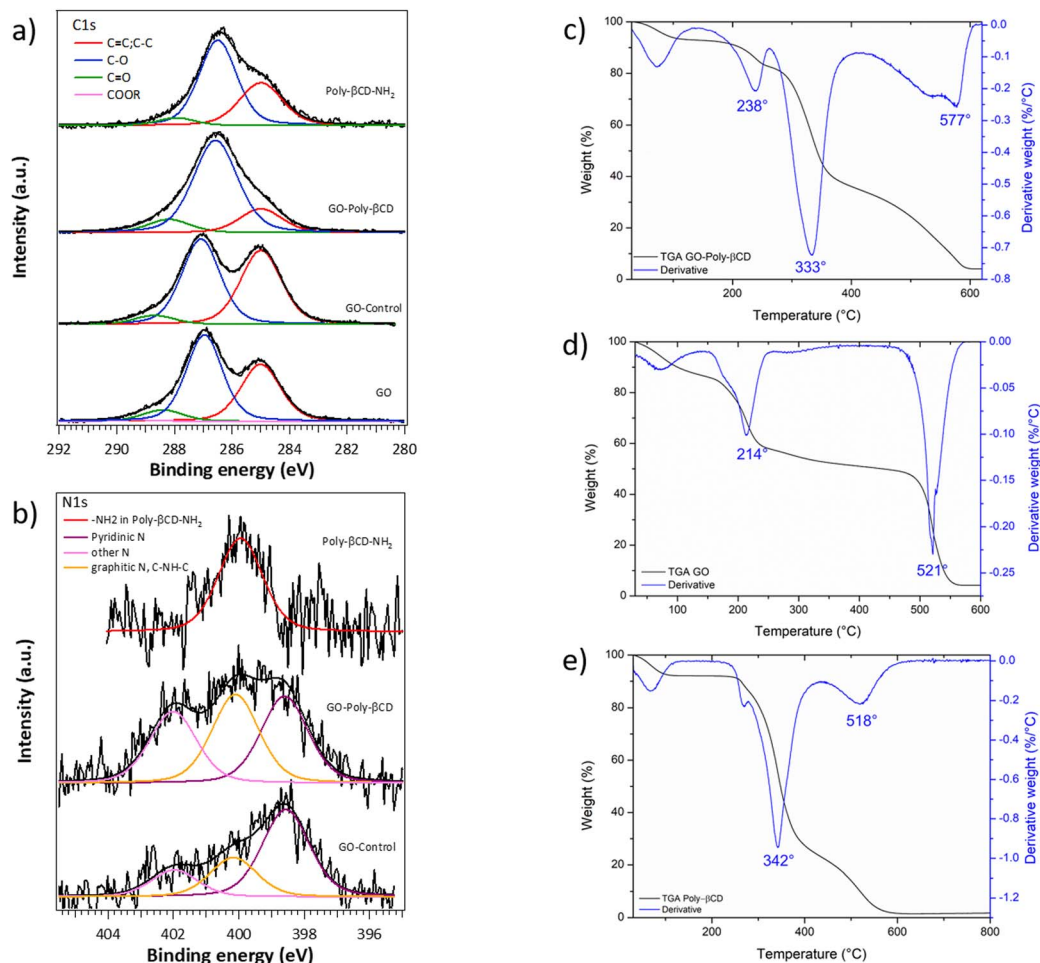


Fig. 2 XPS core level spectra of (a) C 1s and (b) N 1s of GO, GO Control, GO-Poly-βCD and Poly-βCD-NH₂ (for reference) and TGA profiles of (c) GO-Poly-βCD, (d) pristine GO and (e) Poly-βCD.

Control), leading to a decreased oxygen content (24% vs. 30.8%) and, consequently, an increased C/O ratio (1.4 vs. 1.3). By monitoring the increase in nitrogen content, it was possible to estimate the Poly-βCD loading. Starting from a βCD/epichlorohydrin ratio of 7:3 in the starting material, the molecular formula of the Poly-βCD repetitive unit was determined to be C₆₇H₁₂₉NO₄₂. Given a total of 239 atoms per repetitive unit, the theoretical nitrogen content was calculated as 0.42%, which is in excellent agreement with the experimentally determined value (Table S1). The Poly-βCD loading in GO-Poly-βCD was found to be 55% as it can be estimated using eqn (3), where nN is the number of nitrogen atoms involved (one in this case), $nTotal$ is the total number of atoms per polymer unit (239) and the % refers to that of atomic nitrogen in GO-Poly-βCD and GO-Control (see SI Section 1 for more details):

$$L\% = \left[\frac{\% N(\text{GO-Poly-}\beta\text{CD}) - \% N(\text{GO-Control})}{nN} \right] \times nTotal \quad (3)$$

Fig. 3 shows the attenuated total reflection Fourier-transform infrared spectroscopy (ATR-FTIR) spectra of GO-Poly-βCD in comparison with pristine GO and Poly-βCD. All GO-based

samples exhibited a broad band in the 3700–3000 cm^{−1} region, associated with O–H and N–H stretching vibrations (in the case of GO-Poly-βCD), and the C–H stretching bands of Poly-βCD between 2920 and 2800 cm^{−1} (in the case of Poly-βCD and GO-Poly-βCD) confirming the presence of the polymer. In the 1800–1500 cm^{−1} region, GO-Poly-βCD displayed the characteristic carbonyl stretching bands of GO at 1710 and 1610 cm^{−1}, indicative of C=O vibrations in carboxylic groups. The band of Poly-βCD at 1636 cm^{−1} was no longer visible after functionalization, consistent with the chemical modification of the amine groups. Overall, the spectral features confirm the coexistence of both GO and Poly-βCD in the hybrid material.

The colloidal stability of GO-Poly-βCD was assessed by turbidimetry and Dynamic Light Scattering (DLS) (Fig. S4, S5 and Table S4). Turbidimetric analysis of 250 μg mL^{−1} dispersions revealed a two-phase exponential decay with half-lives of ~35 and 883 min, indicating polydispersity with at least two particle populations. DLS at 62.5 μg mL^{−1} confirmed a predominant fraction with hydrodynamic diameter, $D_H \approx 100$ nm alongside larger aggregates (~570 nm), with ζ-potential ~−28 mV ensuring stability. pH-dependent studies showed enhanced stability under alkaline conditions, while at pH 2–3



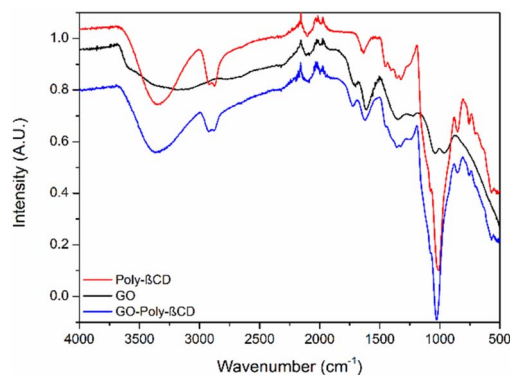


Fig. 3 ATR-FTIR spectra of GO (black), Poly- β CD (red) and GO-Poly- β CD (blue).

protonation of GO surface groups led to aggregation and increased D_H , evidencing partial destabilization.

PFAS adsorption

GO-Poly- β CD nanosheets were tested as sorbents under batch conditions for a mixture of ten PFAS molecules (Fig. 4a), varying in chain length ((CF)_{3–9}) and end-group substitution (sulfonate and carboxylate).

The concentration of the PFAS mixture was selected to reflect environmentally relevant levels commonly found in contaminated water sources (0.1–3 $\mu\text{g L}^{-1}$) and the pH of the tested solution was that of tap water (pH 7). To simulate real-field applications, the adsorption contact time was set to 15 minutes, in line with the typical contact time between water and GAC in municipal water treatment plants (approximately 10–20 minutes). The removal of each contaminant was estimated by UPLC-MS-MS analysis (details in the SI, Section 4). Fig. 4b compares the PFAS removal efficiency of pristine GO, GO-Poly- β CD, and GO-Control after 15 minutes of

contact. It can be seen that GO-Poly- β CD outperforms pristine GO for short- and medium-chain PFASs, achieving up to 97% removal for long-chain PFASs with (CF)_{6–9} length. Medium-chain PFASs ((CF)_{4–6}) are also effectively removed by GO-Poly- β CD, with removal efficiencies exceeding 80%. On the other hand, PFPeA ((CF)₄) was adsorbed by GO-Poly- β CD with removal up to 44%, compared to 0% and 11% observed for GO and GO-Control respectively, while PFBA ((CF)₃), one of the most persistent PFASs, was not removed at all. Pristine GO showed negligible adsorption for short- and medium-chain PFASs ((CF)_{3–7}), with increasing removal observed for long-chain PFASs, ranging from 14% to 76%. GO-Control, which exhibits a slightly higher degree of reduction than pristine GO (C/O = 1.4 vs. 1.3), showed modest removal of medium-chain PFASs ((CF)_{4–7}) and increasing removal for long-chain PFASs ((CF)_{8–9}), reaching up to 59% for PFDA. Overall, GO-Poly- β CD demonstrates consistently higher adsorption selectivity toward medium-chain PFASs compared to both unmodified GO and GO-Control. This is clearly evidenced by the adsorption of (CF)_{4–6} chains, which are removed by GO-Poly- β CD with efficiencies ranging from 44% to 97%, whereas GO-Control shows only limited removal, with a maximum of 20% for PFHpA ((CF)₆), and pristine GO exhibits negligible removal across this range. Poly- β CD polymers carry positive charges due to multiple protonated amine moieties, which interact with the negatively charged, deprotonated carboxylate groups of GO. As a result, pH variations can strongly influence the adsorption behavior of the GO-Poly- β CD system. We therefore compared the removal performances at pH 4, 7, and 10 (Fig. 4c). Adsorption at pH 4 is consistently higher than at pH 7, with the most pronounced improvements observed for short- and medium-chain PFASs. At pH 4, PFBS, PFHxA, and PFHpA were removed with exceptional efficiencies of 95%, 94%, and 97%, respectively. PFPeA, which exhibited 44% removal at pH 7, showed an increased removal of 80% at pH 4.

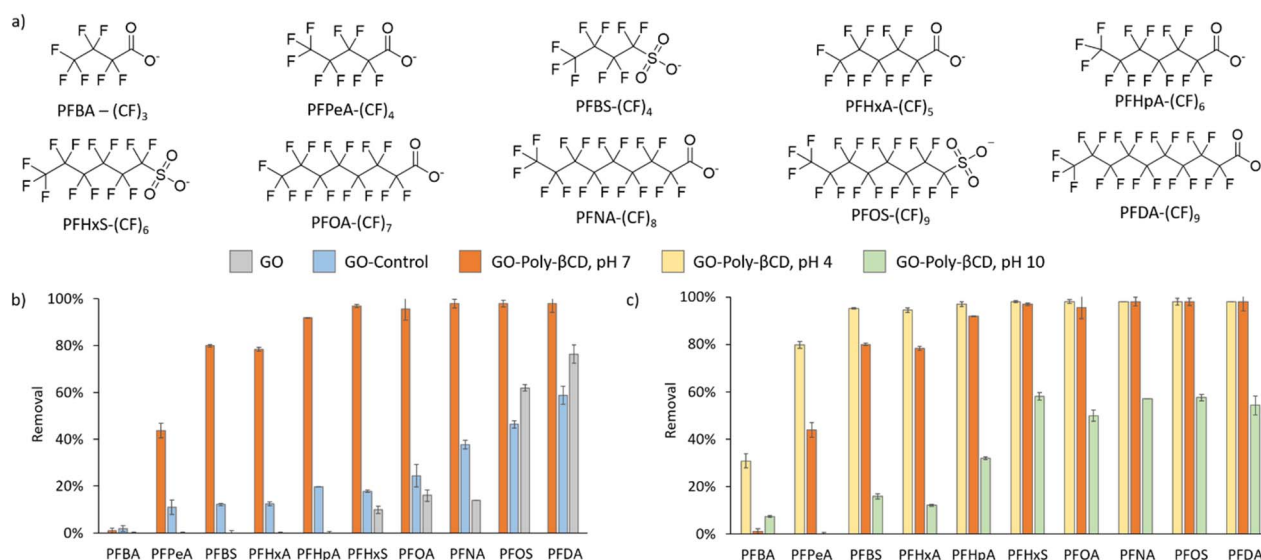


Fig. 4 (a) Molecular structure of the PFAS molecules, with different chain lengths ((CF)_{3–9}) and end-group substitution (sulfonate and carboxylate) studied in this work. (b) Removal of PFAS mixture at pH 7 (orange) from GO-Poly- β CD compared to GO-Control (blue) and unmodified GO (grey). (c) Experiments at pH 4 (yellow), pH 7 (orange) and pH 10 (green).



Notably, PFBA, which was not removed at neutral pH, was adsorbed at acidic pH with a removal efficiency of 31%. Under basic conditions, the system exhibited substantially reduced adsorption: short- and medium-chain PFASs ((CF)₃₋₆) were removed with negligible to poor efficiency, while long-chain PFASs ((CF)₇₋₉) showed only moderate adsorption, with a maximum removal of 58% for PFHxS, significantly lower than the near-quantitative removal observed under neutral and acidic conditions.

Mechanistic insights of adsorption

To elucidate the binding mechanism responsible for the enhanced removal of PFASs by GO-Poly- β CD at pH 7, molecular dynamics (MD) simulations were performed using perfluorohexanoic acid (PFHxA) as a representative case study. To investigate the specific role of Poly- β CD moieties grafted onto the GO surface, the interactions of PFHxA with both GO and GO-Poly- β CD were analyzed. Five distinct adsorption sites for PFHxA were identified in the GO-Poly- β CD system (Fig. 5), designed to capture a range of interaction environments: adsorption on GO (Case 1), interaction with Poly- β CD (Case 2), encapsulation within the cyclodextrin cavity (Case 3), and sandwich interaction with both GO and Poly- β CD (Case 4). Five independent MD simulations, each 100 ns in length, were carried out from these starting geometries to dynamically sample the interactions between PFHxA and the nanosorbent. The binding free energy between PFHxA and GO-Poly- β CD was calculated in each case using the MM-GBSA approach (Table 1). PFHxA exhibited weak binding to GO ($\Delta G_{\text{binding}} = -1.9 \text{ kcal mol}^{-1}$ for pristine GO, $-2.8 \text{ kcal mol}^{-1}$ for GO-Poly- β CD, Case 1). Similarly, interactions between PFHxA and Poly- β CD alone were very weak (Case 2, $-1.8 \text{ kcal mol}^{-1}$).

Encapsulation of PFHxA within the β CD cavity of the Poly- β CD polymer (Case 3) significantly enhanced binding ($\Delta G_{\text{binding}} = -12.4 \text{ kcal mol}^{-1}$). Notably, the strongest interaction was observed in Case 4, where PFHxA is sandwiched between Poly- β CD and GO, interacting with both components of the nanosorbent, yielding a $\Delta G_{\text{binding}}$ of $-15.6 \text{ kcal mol}^{-1}$. This value demonstrates that Poly- β CD is more effective than monomeric β CD derivatives (PFBA: β CD = $-12.3 \text{ kcal mol}^{-1}$).²⁵ The high

Table 1 Binding free energy ($\Delta G_{\text{binding}}$) and energy contributions calculated for each material. All energies are reported in kcal mol^{-1}

PFHxA@	VDW	E_{el}	E_{SURF}	$\Delta H_{\text{binding}}$	ΔS	$\Delta G_{\text{binding}}$
GO	-16.0	4.8	-1.2	-12.4	-10.5	-1.9
Case 1	-18.7	4.2	-1.4	-15.9	-13.1	-2.8
Case 2	-15.2	4.4	-2.7	-13.4	-11.6	-1.8
Case 3	-29.7	6.0	-4.3	-28.0	-15.6	-12.4
Case 4	-35.2	8.0	-5.1	-32.3	-16.7	-15.6

affinity arises from a favorable combination of interaction forces, primarily driven by van der Waals interactions, which contribute $-35.2 \text{ kcal mol}^{-1}$. These strong dispersion forces result from the close contact between PFHxA and both the GO surface and the overlying Poly- β CD. The electrostatic contribution is slightly unfavorable ($E_{\text{el}} = 8.0 \text{ kcal mol}^{-1}$), likely due to repulsion between the negatively charged head group of PFHxA and the GO surface, as well as desolvation of the PFAS head group.

In contrast, non-polar solvation contributions were favorable ($E_{\text{SURF}} = -5.1 \text{ kcal mol}^{-1}$), reflecting burial of the PFAS aliphatic chain within the cleft formed by Poly- β CD and the GO sheet. The entropy term ($\Delta S = -16.7 \text{ kcal mol}^{-1}$) represents a binding penalty due to the conformational restriction of PFHxA upon adsorption. These results indicate that stable adsorption and retention of PFASs require the combined action of both Poly- β CD and GO moieties, highlighting the necessity of both components for effective pollutant capture and supporting a cooperative “sandwich-like” mechanism as the most favorable binding mode. The binding energies of all investigated PFASs on GO-Poly- β CD were also calculated, reproducing the experimentally observed trends (Fig. S3).

To assess the influence of pH on PFAS adsorption, MD simulations were performed for a model system (PFHxA adsorption on GO-Poly- β CD) at pH 4, 7, and 10, and the corresponding binding free energies were calculated (Table 2). A net decrease in binding affinity was observed with increasing pH: $\Delta G_{\text{binding}} = -19.9$, -15.6 , and $-10.6 \text{ kcal mol}^{-1}$ at pH 4, 7, and 10, respectively, consistent with the experimental removal trend (Fig. 4c). Electrostatic repulsion (E_{el}) between PFHxA and GO-Poly- β CD increases as pH rises, since PFHxA remains deprotonated across the studied

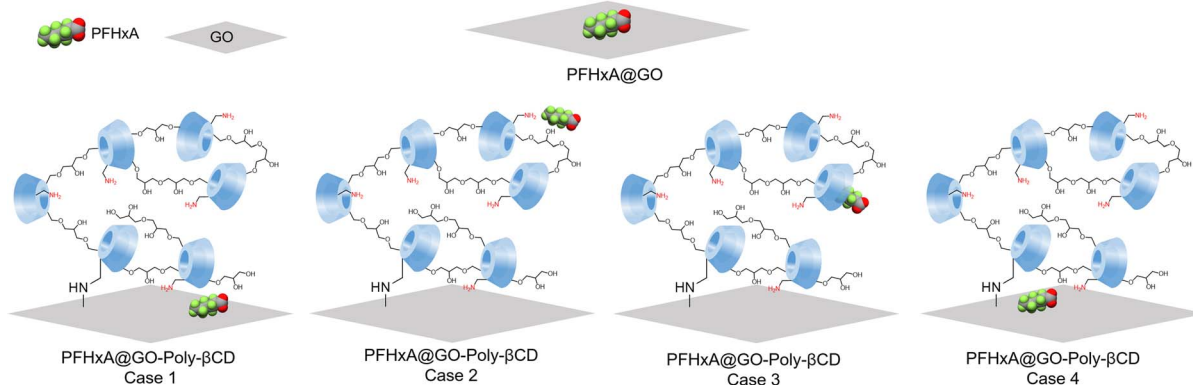


Fig. 5 Schematic representation of the interactions between PFHxA and GO-Poly- β CD, highlighting the different adsorption sites.



Table 2 Binding free energies ($\Delta G_{\text{binding}}$) obtained for PFHxA@GO-Poly- β CD at three different pH values (4, 7 and 10). All energies are reported in kcal mol⁻¹

PFHxA@ GO-Poly- β CD	VDW	E_{El}	E_{SURF}	$\Delta H_{\text{binding}}$	ΔS	$\Delta G_{\text{binding}}$
pH 4	-37.8	7.8	-5.1	-35.2	-15.3	-19.9
pH 7	-35.2	7.9	-5.1	-32.3	-16.7	-15.6
pH 10	-31.2	10.6	-3.2	-23.7	-13.1	-10.6

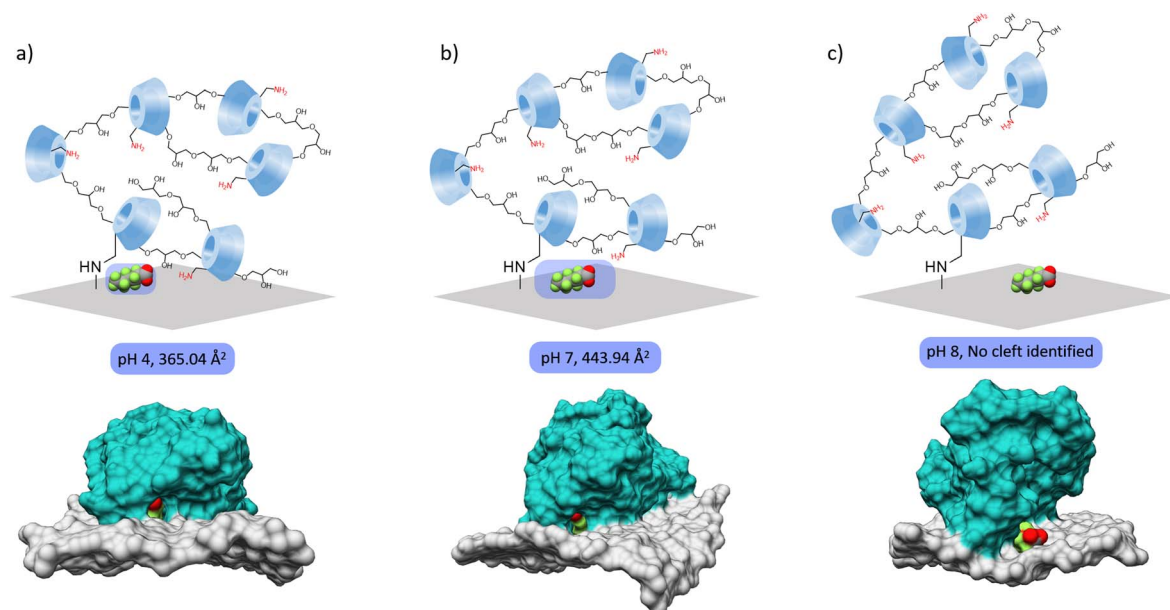
range ($\text{p}K_{\text{a}} = -0.16$). The improved adsorption under acidic conditions is primarily attributed to stronger van der Waals interactions, which are unexpectedly influenced by the pH. Structural analysis revealed a pH-dependent conformational change of the nanosystem, affecting the accessibility and stability of PFAS adsorption sites (Fig. 6). At pH 4, Poly- β CD remains closely associated with GO through hydrogen bonding, forming tight binding clefts (365.04 \AA^2) for PFHxA.

These clefts become less well-defined at pH 7 (443.94 \AA^2), while at pH 10, Poly- β CD partially detaches from the GO surface, disrupting the interaction network and dismantling the clefts responsible for PFHxA adsorption. Consequently, under basic conditions, PFHxA remains largely exposed on the GO surface and is only partially covered by the polymer, breaking the sandwich-like adsorption site, reducing the synergistic binding effect, and resulting in overall weaker interactions. This indicates that the adsorption of PFHxA—and PFASs in general—on GO-Poly- β CD is primarily hydrophobically driven. This behavior is also reflected in the non-polar solvation term (E_{SURF}), which becomes less stabilizing at higher pH ($E_{\text{SURF}} = -3.2 \text{ kcal mol}^{-1}$ at pH 10 vs. $-5.1 \text{ kcal mol}^{-1}$ at pH 4). As with van der Waals interactions, E_{SURF} depends directly on the contact area between the adsorption site and the PFAS, *i.e.*, their shape complementarity.

Although it was initially hypothesized that pH would primarily modulate electrostatic interactions between PFAS and the GO-Poly- β CD surface—particularly involving the protonated

amines of Poly- β CD—our findings indicate that the critical factor is the pH-dependent conformational rearrangement of the hybrid material. At lower pH, stronger interactions between GO and Poly- β CD generate well-defined clefts that facilitate efficient guest inclusion and encapsulation of PFAS.

To further confirm the pivotal role of Poly- β CD in modulating the adsorption behavior of the hybrid material under different pH conditions, the removal performance of pristine GO was also evaluated across pH 4, 7, and 10 (Fig. S6). Short- and medium-chain PFASs (CF_{3-6}) exhibited negligible removal at all tested pH values. A modest improvement was observed at pH 4, with removal increasing to 16% for PFHpA and 29% for PFHxS. This can be attributed to the partial neutralization of the negative surface charge of GO under acidic conditions, slightly reducing electrostatic repulsion with the anionic PFASs. For long-chain PFASs (CF_{7-9}), removal efficiency increased more substantially under acidic conditions, reaching up to 91% in the case of PFDA. At neutral and basic pH, only minor improvements were observed for PFNA, while PFOS and PFDA showed higher removal efficiencies, though they never exceeded 80%. The gradual increase in adsorption with chain length under all pH conditions reflects the stronger hydrophobic interactions of longer-chain PFASs. Overall, the negligible adsorption of pristine GO under most conditions highlights the essential role of Poly- β CD in stabilizing PFASs and corroborates the experimental and computational findings described above.

**Fig. 6** Sketch of PFHxA@GO-Poly- β CD and relative cleft formation and calculated area at (a) pH 4, (b) pH 7 and (c) pH 10.

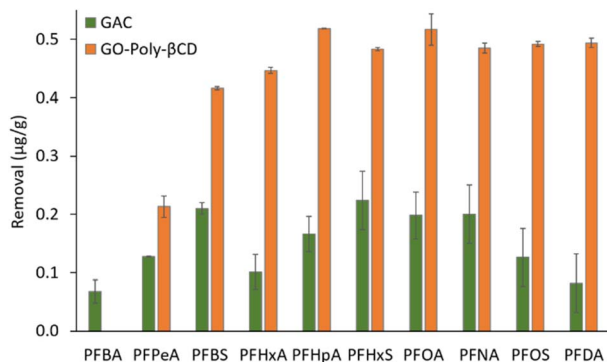


Fig. 7 PFAS removal performances of GO-Poly-βCD and GAC.

PFAS adsorption of GO-Poly-βCD vs. GAC

To evaluate the potential of GO-Poly-βCD for drinking water treatment, its adsorption performance was compared with that of GAC, representing the current industrial sorbent benchmark widely employed in treatment plants for tertiary and quaternary water treatment as well as in point-of-use purification systems.

The comparison was performed in tap water at neutral pH, with contact time of 15 minutes, conditions that closely reflect real-world applications. Fig. 7 shows the removal efficiencies of GO-Poly-βCD and GAC, expressed as micrograms of PFAS removed per gram of sorbent ($\mu\text{g g}^{-1}$). The data clearly indicate that GO-Poly-βCD outperforms GAC in PFAS adsorption. While GAC achieves a maximum removal of $0.2 \mu\text{g g}^{-1}$ for most of the selected PFASs, GO-Poly-βCD reaches up to $0.5 \mu\text{g g}^{-1}$ within the same contact time. For PFPeA ((CF)₄), GAC removes $0.1 \mu\text{g g}^{-1}$, slightly lower than the $0.2 \mu\text{g g}^{-1}$ achieved by GO-Poly-βCD. The shortest-chain PFAS, PFBA, is only minimally adsorbed by GAC ($0.07 \mu\text{g g}^{-1}$) and remains undetectable for GO-Poly-βCD. The advantage of GO-Poly-βCD becomes even more pronounced when considering total PFAS removal, which reaches $4 \mu\text{g g}^{-1}$ compared to $1.3 \mu\text{g g}^{-1}$ for GAC. In addition to higher removal efficiencies, GO-Poly-βCD exhibits rapid adsorption kinetics, achieving significant PFAS uptake within just 15 min. This feature is particularly relevant for practical drinking water purification, including both household point-of-use filters and industrial treatment systems operating at short contact times.

Conclusions

In conclusion, we report the synthesis of a novel GO material covalently grafted with poly-β-cyclodextrin crosslinked with epichlorohydrin. The resulting hybrid is insoluble in water, unlike previously reported β-cyclodextrin-based polymers that often exhibit partial solubility, making it more water-processable and suitable for practical applications in aqueous environments. GO-Poly-βCD material adsorption performances were evaluated on a mixture of PFASs with varying chain lengths ((CF)₃₋₉) and acidic and sulfonic functional groups. Batch adsorption experiments demonstrated that GO-Poly-βCD outperforms pristine GO in the removal of medium-chain PFASs, with remarkable efficiencies of 97% for PFHxA and PFHpA ((CF)₆₋₇) and 80% for PFPeA ((CF)₄), whereas pristine GO

and GO-Control exhibited negligible adsorption in this range. Compared to GAC, GO-Poly-βCD achieved higher overall PFAS uptake within 15 minutes of contact, highlighting its superior performance under field-relevant conditions. Molecular dynamics simulations revealed that PFAS adsorption is governed by a synergistic mechanism, in which contaminants are stabilized through a pH-responsive conformational arrangement of GO grafted with poly-βCD. Overall, these findings provide new insights into the design of hybrid nanomaterials for PFAS remediation and underscore the potential of poly-β-cyclodextrin-based GO materials as efficient and sustainable sorbents for drinking water treatment.

Author contributions

Andrea Trifoglio: methodology, investigation, reviewing editing, writing original draft. Angela Pintus: methodology, investigation. Sara Khaliha: methodology, investigation. Marco Agnes: methodology. Lucrezia Aversa: methodology, investigation. Roberto Verucchi: methodology, investigation. Nina Burduja: investigation. Giuseppe Nocito: methodology, investigation and review original draft. Antonino Mazzaglia: methodology and review original draft. Tainah Dorina Marforio: investigation, methodology, formal analysis. Matteo Calvaresi: methodology, investigation. Manuela Melucci: coordination, conceptualization, validation, writing original draft.

Conflicts of interest

There are no conflicts to declare.

Data availability

Supplementary information (SI): GO-Poly-βCD loading calculation, X-ray photoelectron spectroscopy (XPS), binding affinities, UPLC-MS/MS details, colloidal stability studies, and PFAS adsorption from GO at different pH values. See DOI: <https://doi.org/10.1039/d5ta08410e>.

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Notes and references

- 1 *Nat. Water*, 2023, **1**, 993.
- 2 O. S. Arvaniti and A. S. Stasinakis, *Sci. Total Environ.*, 2015, **524–525**, 81–92.
- 3 M. Sun, E. Arevalo, M. Strynar, A. Lindstrom, M. Richardson, B. Kearns, A. Pickett, C. Smith and D. R. U. Knappe, *Environ. Sci. Technol. Lett.*, 2016, **3**, 415–419.
- 4 S. Garg, J. Wang, P. Kumar, V. Mishra, H. Arafat, R. S. Sharma and L. F. Dumée, *J. Environ. Chem. Eng.*, 2021, **9**, 105784.
- 5 J. Gardiner, *Aust. J. Chem.*, 2015, **68**, 13–22.
- 6 S. Dolui, D. Kumar, S. Banerjee and B. Ameduri, *Acc. Mater. Res.*, 2021, **2**, 242–251.
- 7 Z. Du, S. Deng, Y. Bei, Q. Huang, B. Wang, J. Huang and G. Yu, *J. Hazard. Mater.*, 2014, **274**, 443–454.
- 8 T. H. Boyer, Y. Fang, A. Ellis, R. Dietz, Y. J. Choi, C. E. Schaefer, C. P. Higgins and T. J. Strathmann, *Water Res.*, 2021, **200**, 117244.
- 9 P. McCleaf, S. Englund, A. Östlund, K. Lindegren, K. Wiberg and L. Ahrens, *Water Res.*, 2017, **120**, 77–87.
- 10 S. Khaliha, A. Bianchi, A. Kovtun, F. Tunioli, A. Boschi, M. Zambianchi, D. Paci, L. Bocchi, S. Valsecchi, S. Polesello, A. Liscio, M. Bergamini, M. Brunetti, M. Luisa Navacchia, V. Palermo and M. Melucci, *Sep. Purif. Technol.*, 2022, **300**, 121826.
- 11 E. Gagliano, M. Sgroi, P. P. Falciglia, F. G. A. Vagliasindi and P. Roccaro, *Water Res.*, 2020, **171**, 115381.
- 12 A. M. Kennedy, A. M. Reinert, D. R. U. Knappe, I. Ferrer and R. S. Summers, *Water Res.*, 2015, **68**, 238–248.
- 13 S. Khaliha, F. Tunioli, L. Foti, A. Bianchi, A. Kovtun, T. D. Marforio, M. Zambianchi, C. Bettini, E. Briñas, E. Vázquez, L. Bocchi, V. Palermo, M. Calvaresi, M. L. Navacchia and M. Melucci, *Environ. Sci.:Water Res. Technol.*, 2024, **10**, 1097–1107.
- 14 S. Khaliha, T. D. Marforio, A. Kovtun, S. Mantovani, A. Bianchi, M. Luisa Navacchia, M. Zambianchi, L. Bocchi, N. Boulanger, A. Iakunkov, M. Calvaresi, A. V. Talyzin, V. Palermo and M. Melucci, *FlatChem*, 2021, **29**, 100283.
- 15 S. Guo, S. Garaj, A. Bianco and C. Ménard-Moyon, *Nat. Rev. Phys.*, 2022, **4**, 247–262.
- 16 M. Melucci, L. Bocchi, M. Zambianchi and V. Palermo, *Nat. Water*, 2025, **3**, 369–371.
- 17 S. Mantovani, S. Khaliha, L. Favaretto, C. Bettini, A. Bianchi, A. Kovtun, M. Zambianchi, M. Gazzano, B. Casentini, V. Palermo and M. Melucci, *Chem. Commun.*, 2021, **57**, 3765–3768.
- 18 S. Mantovani, S. Khaliha, T. D. Marforio, A. Kovtun, L. Favaretto, F. Tunioli, A. Bianchi, G. Petrone, A. Liscio, V. Palermo, M. Calvaresi, M. L. Navacchia and M. Melucci, *Chem. Commun.*, 2022, **58**, 9766–9769.
- 19 A. Trifoglio, S. Mantovani, S. Khaliha, A. Kovtun, T. D. Marforio, M. Calvaresi and M. Melucci, *Nanoscale*, 2025, **17**, 12124–12133.
- 20 G. Crini, *Chem. Rev.*, 2014, **114**, 10940–10975.
- 21 A. H. Karoyo, A. S. Borisov, L. D. Wilson and P. Hazendonk, *J. Phys. Chem. B.*, 2011, **115**, 9511–9527.
- 22 P. Lo Nostro, I. Santoni, M. Bonini and P. Baglioni, *Langmuir*, 2003, **19**, 2313–2317.
- 23 H. Tatsuno and S. Ando, *J. Phys. Chem. B.*, 2006, **110**, 25751–25760.
- 24 A. H. Karoyo, P. S. Sidhu, L. D. Wilson, P. Hazendonk and A. Borisov, *J. Phys. Chem. C.*, 2015, **119**, 22225–22243.
- 25 F. Tunioli, T. D. Marforio, L. Favaretto, S. Mantovani, A. Pintus, A. Bianchi, A. Kovtun, M. Agnes, V. Palermo, M. Calvaresi, M. L. Navacchia and M. Melucci, *Chem.–Eur. J.*, 2023, **29**, e202301854.
- 26 Y. Miyah, N. El Messaoudi, M. Benjelloun, J. Georgin, D. S. P. Franco, M. El-habacha, O. A. Ali and Y. Acikbas, *Carbohydr. Polym.*, 2025, **350**, 122981.
- 27 L. Xiao, Y. Ling, A. Alsbaiee, C. Li, D. E. Helbling and W. R. Dichtel, *J. Am. Chem. Soc.*, 2017, **139**, 7689–7692.
- 28 A. Alsbaiee, B. J. Smith, L. Xiao, Y. Ling, D. E. Helbling and W. R. Dichtel, *Nature*, 2016, **529**, 190–194.
- 29 J.-L. Sun, H. Zeng and H.-G. Ni, *Chemosphere*, 2013, **90**, 1751–1759.
- 30 C. Belenguer-Sapiña, E. Pellicer-Castell, A. R. Mauri-Aucejo, E. F. Simó-Alfonso and P. Amorós, *Nanomaterials*, 2021, **11**, 1.
- 31 A.-M. Hada, M. Potara, S. Astilean, A. Cordaro, G. Neri, M. Malanga, A. Nostro, A. Mazzaglia, A. Scala and A. Piperno, *Carbohydr. Polym.*, 2022, **293**, 119736.
- 32 X. Deng, A. Verdaguer, T. Herranz, C. Weis, H. Bluhm and M. Salmeron, *Langmuir*, 2008, **24**, 9474–9478.

