



Research article

Carbon-cryogel hierarchical composites as effective and scalable filters for removal of trace organic pollutants from water



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ABSTRACT

Effective technologies are required to remove organic micropollutants from large fluid volumes to overcome present and future challenges in water and effluent treatment. A novel hierarchical composite filter material for rapid and effective removal of polar organic contaminants from water was developed. The composite is fabricated from phenolic resin-derived carbon microbeads with controllable porous structure and specific surface area embedded in a monolithic, flow permeable, poly(vinyl alcohol) cryogel. The bead-embedded monolithic composite filter retains the bulk of the high adsorptive capacity of the carbon microbeads while improving pore diffusion rates of organic pollutants. Water spiked with organic contaminants, both at environmentally relevant concentrations and at high levels of contamination, was used to determine the purification limits of the filter. Flow through tests using water spiked with the pesticides atrazine (32 mg/L) and malathion (16 mg/L) indicated maximum adsorptive capacities of 641 and 591 mg pollutant/g carbon, respectively. Over 400 bed volumes of water contaminated with 32 mg atrazine/L, and over 27,400 bed volumes of water contaminated with 2 µg atrazine/L, were treated before pesticide guideline values of 0.1 µg/L were exceeded. High adsorptive capacity was maintained when using water with high total organic carbon (TOC) levels and high salinity. The toxicity of water filtrates was tested *in vitro* with human epithelial cells with no evidence of cytotoxicity after initial washing.

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1. Introduction

A wide range of contaminants are continuously released into waterways as a result of anthropogenic activity, many of which are not effectively removed by current waste water treatment processes (Wilkinson et al., 2016; Loos et al., 2013). Problem

contaminants commonly comprise agrochemicals, pharmaceuticals, hormones, plasticisers, flame retardants, personal care products and food additives (Homem et al., 2015; Kokotou et al., 2012; Richardson and Ternes, 2014). These organic compounds are distributed across different environmental systems (e.g. soils, sediments, surface and ground waters), and the most polar (e.g. amines, alcohols or carboxylic acids and derivatives with small hydrocarbon skeletons and low partition coefficients) are bioavailable and may persist in the aqueous phase. These contaminants are now ubiquitous, and even at very low environmental concentrations (e.g. below 0.5 µg/L in surface and ground water, Cabezam et al., 2012; Loos et al., 2013; Sorensen et al., 2015) can potentially cause ecological changes and contaminate water

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supplies with (in some cases) unknown consequences for human health (Fent et al., 2006; Jeong et al., 2012). There is therefore a pressing need to purify water to safe (i.e. sub-ppb or ppt) levels. A range of technologies have been proposed to remove trace organic contaminants from waste and other waters, including chemical oxidation and improved adsorption methods, often using activated carbons. Indeed, in advanced waste water treatment processes, Granulated Activated Carbon (GAC) is used as a relatively low cost bulk adsorbent and is recognised as an industry standard. However, the adsorptive capacity of GAC diminishes over time, and it shows relatively poor adsorptive uptake for a range of more polar organic contaminants such as estrogens, pharmaceuticals and some pesticides (Busquets et al., 2014; Cabezam et al., 2012; Iñechebuegu, 2011). This has led to a number of studies examining the potential of “designer” activated carbons, where control over the pore architecture is attempted to target particular dissolved species, to improve polar contaminant removal (Busquets et al., 2014; Gueudré et al., 2014; Ji et al., 2010; Unur, 2013; Zhang et al., 2011). In addition, use of hierarchical porous materials has been suggested to enhance mass transfer properties (Gueudré et al., 2014) and provide more effective adsorption of trace contaminants from water.

The present work explores the design of a flexible and scalable hierarchical composite filter comprised of phenolic resin-derived carbon microbeads with optimised micro-, meso- and macroporous structure embedded in a monolithic, flow permeable poly(vinyl alcohol) cryogel for the removal of small size, polar pesticides. Atrazine, (a herbicide of the triazine class, 2-chloro-4-ethylamino-6-isopropylamino-1,3,5-triazine, $\log K_{ow}$ 2.75; US EPA, 2006), and malathion, (an organophosphate insecticide, diethyl[(dimethoxyphosphinothioyl)thio]butanedioate, $\log K_{ow}$ 2.36–2.89; WHO, 2004), which enter waterways via runoff from agricultural land and have been frequently detected in surface or ground water (Fadaei et al., 2012; Nogueira et al., 2012; Sorensen et al., 2015; Varca, 2012), are used as model compounds to assess and optimise the water filters. Application and scalability are also discussed.

2. Materials and methods

2.1. Materials

The sources and purities of the chemicals used are provided in the Supporting Information S1. GAC was kindly provided by a UK water company.

2.2. Synthesis of carbon microbeads

Spherical activated carbon microbeads were obtained from Novolac phenol-formaldehyde resin with an average molecular weight 700–800 D (Hexion Specialty Chemicals Ltd., UK) following the technology patented by Tennison et al. (Tennison et al., 2000). Briefly, micro-, meso- and macroporous phenolic resin-derived carbon beads were prepared by pyrolysis and physical activation in carbon dioxide. Beads were tailored by dissolving Novolac resin (100 weight parts) and cross-linking agent hexamethylenetetramine (20 weight parts) in varying quantities of ethylene glycol (180–300 weight parts for micro-, meso- (TE3), and macroporous (TE7) resins respectively). The resin suspension was then heat-cured in mineral oil. After removing the ethylene glycol (which acts as a solvent and pore former) by hot water washing or vacuum drying, the resin beads were ready for processing into carbons. The amount of carbon loss (burn-off) during activation with CO₂ was 34% and 55% in carbon beads denoted in this work as TE3 and TE7, respectively and the particle size studied ranged from 40 to 250 µm.

The structural features of the carbon beads are summarised in Supporting information S2.

2.3. Preparation of polymer scaffold and composite filter

Monolithic macroporous polymer “scaffolds” (Macroporous Polymer Structures (MPPSTM)) produced by the authors (Protista), also referred to as cryogels, were prepared by chemical cross-linking of polyvinyl alcohol (PVA) at –12 °C and used to support the carbon beads. PVA was dissolved at a concentration of 5% w/v in boiling water and cooled to room temperature. Hydrochloric acid (1 M, 0.8 mL) was mixed with the PVA solution (7.2 mL) and kept in an ice bath for 30 min. Glutaraldehyde solution (75 µL, glutaraldehyde: water 50:50 v/v) and carbon beads (40–60 µm diameter, 320 mg) were added to the cooled PVA solution with stirring. The mixture was poured into Ø3 × 100 mm glass tubes. The sealed tubes were placed into a cryostat (Arctest, Finland) at –12 °C overnight. The cryogels were rinsed with water, then placed in sodium cyanoborohydride solution (0.1 M, pH 9.2) overnight to reduce the residual aldehyde groups, and finally rinsed with 4 bed volumes of water. The cryogels containing carbon beads were Ø3 × 85 mm (0.6 mL) in size when non-hydrated and were used in this configuration for flow through studies.

2.4. Structural characterisation

The structure of the prototypes and GAC was analyzed using low temperature nitrogen adsorption-desorption isotherms and scanning electron microscopy (Supporting Information S2, 3). The mechanical properties of the composites were tested with a T.A. XTPplus texture analyser equipped with a 1 cm diameter cylindrical probe (Stable Micro Systems, UK). The composite samples were compressed with steadily increasing force of 5 N/min from 0.01 N to 20 N at room temperature. The stiffness was assessed as the stress necessary to achieve 75% compression. The Young's modulus of elasticity E , which is the slope of the stress-strain curve in the range of linear proportionality of stress to strain, has been calculated using equation (1), where F is the applied force; A , the cross-sectional area; L , the unstressed height; and ΔL , the change in height due to F ; F/A (in compression or tension) is the stress and $\Delta L/L$ is the strain.

$$E = (F/A)/(\Delta L/L) \quad (1)$$

The stress-strain curve obtained for the composite carried out immediately after a filtration test is shown in Supporting Information S4.

2.5. Adsorption experiments

Adsorption tests were carried out in batch mode under equilibrium conditions using an orbital shaker at 90 rpm, 25 °C for 48 h or magnetic stirring. To assess breakthrough curves in the high capacity carbon-cryogel prototypes using realistic volumes of water at laboratory scale, it was necessary to filter highly contaminated water. Hence, pesticides concentrations were high (32 mg atrazine and 16 mg malathion per L) relative to environmental concentrations. Ethanol (1.4%) was added to stabilise the contaminants in aqueous solution and prevent their precipitation, which would cause overestimation of the removal capacity of the developed prototypes. Batch adsorption tests were also carried out at more environmentally relevant contamination levels of 2 µg atrazine/L. For determination of the diffusion coefficient of atrazine and kinetics of its uptake, carbon beads (15 mg), or carbon-cryogel composites cut into cubes of 2 mm sides containing an equivalent

mass of carbon beads (15 mg), were incubated with spiked water (40 mL). Details on the quantification of adsorption capacity and pore diffusion kinetics are given in [Supporting Information S3](#).

2.6. Flow through adsorption studies

Contaminated water was filtered continuously through the carbon-cryogel composite or a column packed with only carbon beads until saturation, using a peristaltic pump equipped with silicone tubing. Fractions of the filtrate were collected for analysis. Selectivity for removal of atrazine (32 mg/L) and malathion (16 mg/L) was studied with: (a) ultrapure water containing ethanol (1.4%); (b) a “model” water with high total carbon prepared by dissolving potassium hydrogen phthalate in aqueous solution (to mimic high-COD water) and ethanol (1.4%); and (c) water from the Ouse estuary (collected from Newhaven harbour, East Sussex, UK). The composition of the estuarine water (c) before spiking (and before adding 1.4% ethanol) was 19.2 mg/L total carbon; 12.7 mg/L inorganic carbon; 6.5 mg/L total organic carbon (TOC); Na^+ 9320 mg/L; K^+ 514 mg/L, Mg^{2+} 1261 mg/L and Ca^{2+} 339 mg/L as major ions. The pH was 8.1. The methodology used for the water analysis is given in the [Supporting information S5](#). To assess adsorption at low (more environmentally relevant) concentrations, breakthrough curves were also determined using 2 μg atrazine/L spiked ultrapure water.

2.7. Analysis of atrazine and malathion

Atrazine and malathion were quantified using d_5 -atrazine (atrazine with fully deuterated *N*-ethyl moiety) as an internal standard. Quantification was by LC-MS. Chromatographic and MS operating conditions are provided in the [Supporting Information S5](#).

2.8. Modelling

The dimensions of atrazine and the minimum pore volume in a 3D carbonaceous structure for its adsorption were modelled using the Build interface in Spartan '10 software. Geometry optimisation was achieved using the semi-empirical quantum chemistry method PM6 from an initial structure calculated using molecular mechanics (Merck Molecular Force Field - MMFF).

2.9. Cytotoxicity study

The toxicity of ultrapure water filtered through the carbon-cryogel composites was tested on heterogeneous human epithelial colorectal adenocarcinoma cells (Caco-2). Cell viability was assessed *in vitro* using the lactate dehydrogenase (LDH) ([Smith et al., 2011](#)) and MTS assays ([Malich et al., 1997](#)). The cell culture methods and materials used are described in [Supporting Information S6](#).

2.10. Data analysis

Statistical treatment: a t-student significance test, with $P < 0.05$, was used to assess the adsorption of atrazine by PVA scaffolds without carbon beads; compare the content of carbon surface functional groups in the different carbons synthesized; and compare the viability of cells when treated with fractions of water filtrate.

3. Results and discussion

3.1. Synthesis and application of phenolic carbon beads for water treatment

A graphite-like lattice of activated carbon contains a cloud of delocalised electron density that favours the adsorption of organic compounds, especially if they are aromatic, via π – π interaction. Pores of appropriate size can add additional interaction between the pollutant and the carbon, and can significantly enhance removal, which is particularly poor for polar and small organic molecules. With atrazine as an example of a ubiquitous micro-pollutant in surface water, we have modelled and synthesised porous carbon sorbents for its optimal removal from water. The dimensions of the optimal pores in the carbonaceous material that would present the highest interaction with atrazine and produce a state of minimal energy were derived using molecular mechanics (MMFF force field) and semi-empirical (PM6) methods and are shown in [Fig. 1](#). Large micropores (1.0–1.7 nm width) and mesopores (2–50 nm) in the low size range are predicted to be the preferential sites for the adsorption of the study contaminant.

For standard GAC used in the waste water industry, the maximum adsorptive capacity for atrazine was 52.2 ± 6.8 mg/g (20 mg of GAC was incubated with 40 mL of 32 mg/L atrazine containing 1.4% ethanol in water, in batch conditions until equilibrium). The surface area of GAC determined from the nitrogen adsorption-desorption isotherm was $552 \text{ m}^2/\text{g}$. Given the dimensions of atrazine estimated in [Fig. 1](#), approximately 18% of the surface area of GAC was covered with the herbicide at equilibrium conditions.

The nitrogen adsorption-desorption isotherm also revealed that the GAC was mainly a microporous sorbent, with pores of 1.1–2.0 nm width, and hence appropriate for the removal of the herbicide according to the model shown in [Fig. 1](#). This model assumed that a size-exclusion effect found for bulky compounds ([Ji et al., 2010](#)) would not occur. However solvation could increase the effective solute dimensions and hinder access to micropores.

Aside from well-developed microporosity, the GAC has very low mesoporosity, in the range of 2.1–3.8 nm, and macropores were not detected. The absence of (meso- and macroporous) transport pores hinders the accessibility of the micropores, which can explain in part the low adsorptive capacity observed under high concentrations of atrazine.

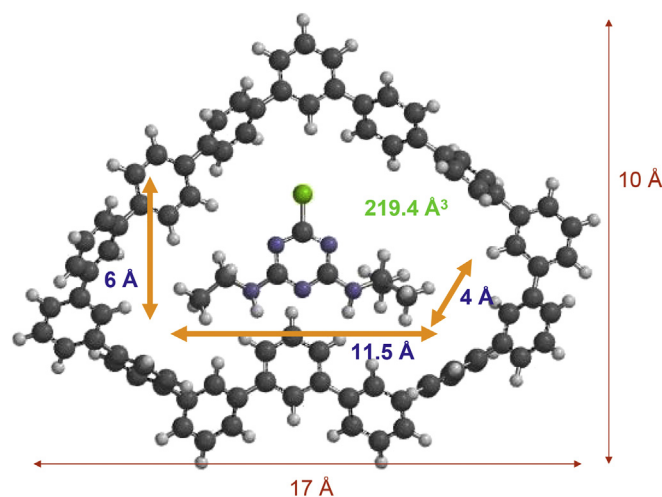


Fig. 1. Diagram showing the dimensions of an optimal carbon structure for adsorption of atrazine.

To maximise pesticide removal, porous carbon microbeads with larger surface area and higher meso- and macropore volume compared to GAC were prepared. The porosity was primarily controlled by the amount of the pore former, ethylene glycol, which is entrapped in the liquid form within the phenolic resin when it solidifies in contact with hot mineral oil. Washing out the entrapped ethylene glycol creates the porous structure in the polymeric precursor of activated carbon. In contrast, in the absence of the pore former, the oligomer chains cross-link, increase in weight and precipitate. Secondly, the activation stage, which was carried out at 900 °C in a CO₂ atmosphere, caused an increase of the number of micropores and the specific surface area and a slight widening of the meso- and macropores, depending on the activation degree, starting from 30% burn-off. The phenolic resin-derived carbon obtained by the described process has a S_{BET} of 534 m²/g when non activated, for a particle size of 125–250 μm. This increases to 1057 m²/g at 30% activation; 1432 m²/g at 47% activation; 1703 m²/g at 52% activation and 2018 m²/g at 65% activation. Phenolic carbons freshly activated in carbon dioxide had a point of zero charge of 10, indicating that at typical environmental water pH the surface of the carbon would be positively charged. Analysis of the surface chemistry of the carbon beads was carried out using Boehm titrations (Valente Nabais and Carrott, 2006). The range of phenolic carbons beads prepared did not have significant differences in their surface chemistry. Total carboxylic functions (mainly free carboxylic acid + anhydride + lactone + lactol) were 0.58 mmol/g as determined by back titration after reaction with Na₂CO₃. Total acidic functionalities were 1.16 mmol/g as determined by back titration after reaction with NaOH. Hence, the phenolic OH content was 0.58 mmol/g.

From the range of phenolic carbon prototypes synthesised, those with clear differences in their mesoporosity were tested for the adsorption of atrazine in a column flow through experiment (conditions and carbon prototype properties are given in Materials and methods). Phenolic carbons beads, named as TE3 and TE7 (prepared with 180 and 300 wt parts of ethylene glycol per 100 parts of the resin), had abundant microporosity, low and high meso- and macroporosity, and an S_{BET} of 1057 and 1703 m²/g, respectively. The phenolic carbon beads with the highest active surface area and pore volume (TE7) gave the greatest uptake of atrazine: 419 mg atrazine/g upon saturation. We previously showed that TE3 and TE7 effectively removed the emerging contaminant metaldehyde (Busquets et al., 2014).

To examine the effects of S_{BET} on the uptake of atrazine, TE7 carbons were synthesised with smaller particle size (40–60 μm), which had a 15% larger S_{BET} : 1942 m²/g. In this case, the maximum adsorptive capacity for atrazine increased to 641 mg atrazine/g carbon. This maximum adsorptive capacity would give a coverage of 1238 m²/g with atrazine according to the dimensions of the herbicide shown in Fig. 1. This represents 64% of the surface area of the carbon covered, and indicates that there are specific sites on the carbon where the pesticide adsorbs. This contrasts with the 18% surface coverage found for GAC. The TE7 carbons were found to be optimal: they have higher porosity and smaller particle size than GAC, both characteristics are considered important for the removal of atrazine and potentially for micropollutants of similar size. High adsorptive capacities (i.e. the phenolic carbon beads adsorbed atrazine to more than half their own weight) were found. Fig. 2 shows the structural properties of the carbons and their adsorption capacity under flow-through conditions. The pore size distribution diagrams (Fig. 2, right) show that while the size of the micropores is similar in all the carbons, the size range of meso- and macropores increases from TE3 to TE7 mainly due to the larger amount of the pore former used.

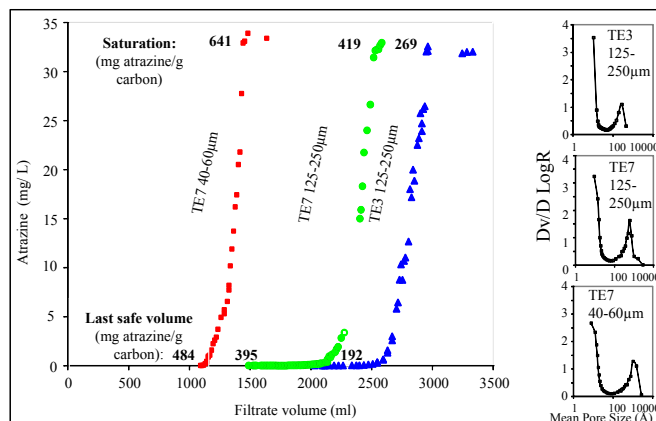


Fig. 2. Breakthrough curves for the adsorption of atrazine on different phenolic carbons packed in a column with dimensions 5 × 70 mm TE7 40–60 μm particle size (73 mg of carbon) (■); TE7 125–250 μm particle size (195 mg of carbon) (●); TE3 125–250 μm particle size (352 mg carbon) (▲). The adsorption capacities at the last safe water filtrate fraction (below 0.1 μg/L) and saturation point of the curves are given. The pore size distribution of the carbons is also shown.

3.2. Incorporation of phenolic carbon beads in a macroporous polymer matrix

Although the prepared carbon beads were highly efficient for atrazine removal compared to conventional GAC, their relatively small particle size (particularly for the 40–60 μm microbeads with higher active surface area) makes their use in conventional large scale water treatment problematic. In addition, use of small packed beads may significantly increase back pressures in column filtration mode. To overcome this, we examine here the effects of incorporating the beads into a macroporous polymer scaffold, specifically a PVA-derived cryogel or macroporous polymer structure (MPPS™ - see experimental). This scaffold allows the preparation of elastic filters of various geometries and facilitates their repeated use for water filtration. The large macropores of the polymer gel matrix, approximately 100–200 μm in diameter, allow flow through with low flow resistance, as well as keeping the active carbon surface accessible for adsorbing contaminants from water. Under the filtration conditions used, a 7% reduction of the flow rate was observed when using the composite. In contrast,

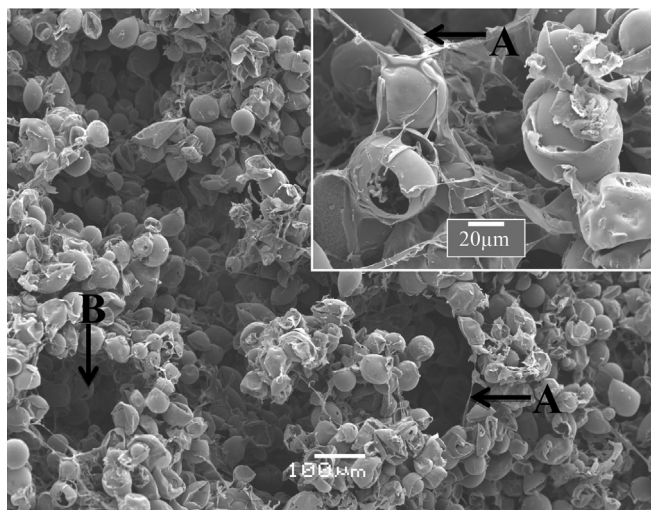


Fig. 3. SEM image of the water purification device based on carbon microbeads embedded in a macroporous PVA cryogel.

no reduction was observed when the filtration was carried out through a microbead-packed column with the same dimensions as the composite. Fig. 3 shows a SEM image of the composite, and illustrates (i) the spherical carbon beads held by a thin PVA film (Fig. 3, arrow A) and (ii) the macropores in the gel (Fig. 3, arrow B). Hence, the filter is a hierarchical material; it contains carbon beads with a range of macro-, meso- and micropores that play an important role in contaminant removal within a macroporous scaffold.

The effect of coating part of the carbon surface with a polymer film on the adsorption kinetics was investigated by comparing the adsorption of atrazine on “free” carbon beads with that on the carbon-cryogel composite. The adsorbent and the atrazine solution were put in contact using two approaches: (a) manual shaking for 10 s, with samples then left to incubate at room temperature without further agitation; (b) under magnetic stirring. While the adsorption was faster in the free carbon beads in the absence of agitation, the adsorption capacity of the carbon beads embedded in PVA was not noticeably suppressed by the PVA film since the final concentration of atrazine in solution was approximately the same in both free carbon and carbon-filled PVA systems (Fig. 4a). These static conditions were assayed to observe the maximum difference in diffusion between the “free” beads and the PVA-embedded beads. When the atrazine diffusion experiment was carried out under magnetic stirring, conditions which did not modify the structure of the particles, the adsorption was approximately twice as fast (Fig. 4b1, Fig. 4b2) compared to the non-agitated samples (Fig. 4a1, Fig. 4a2). There was no significant adsorption of atrazine

by PVA scaffolds without carbon beads (not shown) ($P > 0.05$).

The fractional equilibrium (F) was calculated at different adsorption times to allow the treatment of the kinetic data of Fig. 4 in dimensionless form. The figure in Supporting Information S7 shows F as a function of the square root of time (according to equation (4), Supporting information), for carbon beads (replicates: *, o, Δ in the figure) and for beads embedded in PVA cryogel (replicates: \square , +, \diamond). The data indicate markedly faster uptake values for the carbon beads but with at least one order of magnitude lower pore diffusion coefficient (\bar{D} of $4.2 \times 10^{-10} \text{ cm}^2/\text{s}$) compared to the diffusion when the beads were embedded in the cryogels (\bar{D} of $2.5 \times 10^{-8} \text{ cm}^2/\text{s}$). This could be a consequence of the greater wettability of the PVA-carbon cryogels compared to the carbon microbeads and therefore, a higher accessibility of the adsorbent for atrazine in the aqueous solution. \bar{D} was calculated using a model proposed by Samsonov and Kuznetsova, 1992 (see equations (6) and (7) in Supporting information). The experiment performed under stirring conditions (which exhibited fast adsorption) had insufficient early data points to be analyzed by the first model (equation (4) in the Supporting Information). Only the data points that were taken after >6 min were used to fit a curve (equation (7) Supporting Information), and in that case \bar{D} was $3.3 \times 10^{-9} \text{ cm}^2/\text{s}$, i.e. higher than \bar{D} for the static carbon beads, which could be expected because magnetic stirring provides better mass transport and results in faster external diffusion of the solute to the carbon surface. Notably, the \bar{D} for the stirred carbon sample remained lower than the diffusion coefficient for the carbon-PVA under static conditions.

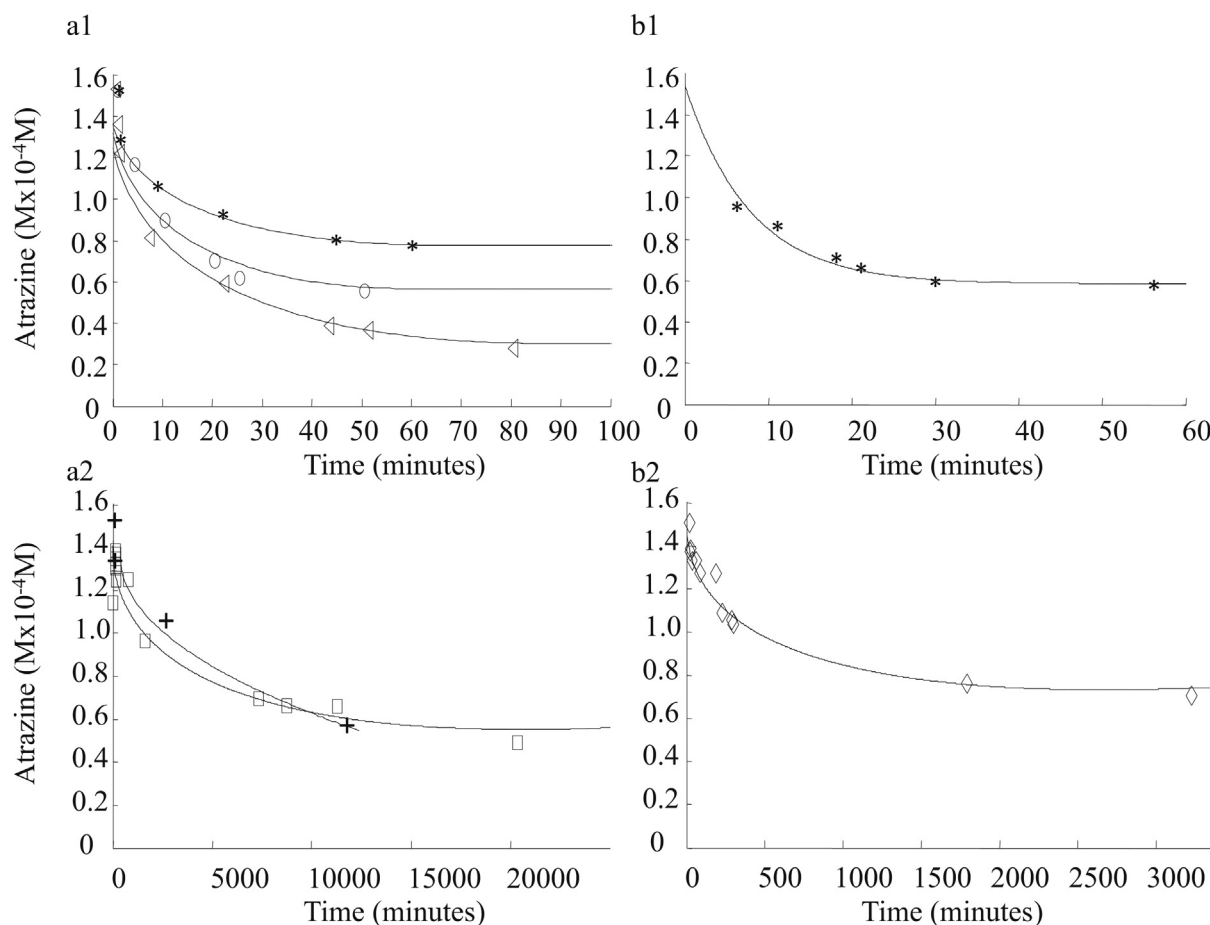


Fig. 4. Uptake of atrazine by carbon beads (a1, b1), replicates shown as *, o, Δ , and beads embedded in cryogel (a2, b2), replicates shown as \square , +, \diamond , under static (a1, a2) and dynamic (magnetically stirred) conditions (b1, b2).

3.3. Mechanical properties

The Young's modulus of elasticity of the composites studied was 13.1 ± 0.8 MPa ($n = 8$), and the loading achieved at 75% compression was 19.6 ± 2.1 N ($n = 8$), conditions that did not break the monolith. Fig. 4S, Supporting Information, displays the stress-strain curve of the composite, showing elastic behaviour under compression at moisture conditions representative of its state during filtration. Under the pressure exerted at 39 h^{-1} flow in the flow-through experiments (where the flow rate is expressed as liquid hourly space velocity), the carbon-cryogel composites were compressed by up to 23%. In contrast no compression was observed in a column packed with beads only.

3.4. Flow-through experiments using carbon-cryogel composite filters

The purification performance of cryogel-carbon composites ($\varnothing 3 \times 85$ mm dimensions, containing 30 mg TE7 carbon beads) was tested at 39 h^{-1} (flow rate expressed as liquid hourly space velocity; liquid flow rate/bed volume) for the purification of atrazine ($2 \mu\text{g/L}$ spiked in ultrapure water and 32 mg/L spiked in: ultrapure water; ultrapure water containing 32 mg potassium hydrogen phthalate/L as a model of water with high organic carbon content; and estuarine water (Ouse river, Newhaven harbour, East Sussex, UK) the composition of which before spiking is described in Materials and methods). The 39 h^{-1} flow rate used was selected to exceed flows typically experienced in water treatment plants, to provide a more realistic flow-through scenario. Results are shown in Fig. 5 and Table 1.

Testing the developed filter materials with water containing a high content of organic matter, and also inorganic substances (i.e.

high dissolved salts) in the case of estuarine water, mimics the effect of possible competing species at high concentration on atrazine removal. Tests with spiked water had TOC values an order of magnitude greater than typical surface water to determine filter performance under extreme conditions. Fig. 5 shows that a total water filtrate of >200 bed volumes contained concentrations of atrazine $<0.1 \mu\text{g/L}$ compared to the inlet water containing 32 mg/L of atrazine. This was observed in the estuarine water sample and the water sample spiked at TOC levels an order of magnitude greater than typical surface water. The purification achieved when using concentrations of atrazine a million times greater than some environmental levels reported (ca. 3 ng/L) (Loos et al., 2010), or in the same order of magnitude (Nogueira et al., 2012), illustrates the high capacity of the composite. For ultrapure water contaminated at $2 \mu\text{g/L}$, 27,400 bed volumes were filtered before the concentration of atrazine exceeded $0.1 \mu\text{g/L}$. When these levels of atrazine were tested in batch mode, concentrations were below the limit of detection (see Supporting information S4) at equilibrium, hence the direct quantification of adsorption capacity was not possible at these concentrations.

Notably, the purification of a more complex matrix than ultrapure water, with high concentrations of soluble organic matter or high salinity, caused a relatively small drop in adsorption capacity, i.e. the amount of filtered water with safe levels of pesticide decreased by less than 30% (Table 1). This suggests that the sites adsorbing atrazine may have some size exclusion selectivity for the adsorption of small molecules such as atrazine, as the sites in the phenolic carbon retain their capacity to remove atrazine in the presence of high concentrations of potentially competing substances. This highlights their potential capacity for contaminant removal from other environmental, industrial or biological media. The slight decrease in adsorptive capacity observed when filtering estuarine water could be due to the blockage of transport pores under these high ionic strength conditions.

Removal of malathion, another polar pesticide with chemical structure different to atrazine, was also tested. In this case ultrapure water containing 16 mg malathion/L was filtered through the composite filter. The adsorptive capacity reached at the last safe filtrate fraction ($0.1 \mu\text{g/L}$) (Council of the European Communities, 1998) was 244 mg malathion/g carbon and 591 mg malathion/g carbon at the saturation of the carbon-cryogel composites, at 745 and 2533 bed volumes, respectively. Supporting information Fig. S8 shows the breakthrough curve for the purification of malathion. This result indicates that the high purification capacity of the developed material is not specific to atrazine but also may apply to other polar pollutants.

3.5. Cytotoxicity assays

Studies of the effect of carbon-cryogel filtered water on the viability of Caco-2 cells (Fig. 6), which are the first type of cells that would be exposed to filtered water when drinking, indicated a significant reduction in cell viability associated with the first 10 mL aliquot filtered through the composite ($P < 0.05$). No significant reduction in viability was observed however with subsequent aliquots (Fig. 6 A) ($P > 0.05$). While the MTS assay applied has been reliably used to assess cell viability and toxicity (Malich et al., 1997), the assay only assesses metabolic activities of mitochondria (Wang et al., 2010; McGowan et al., 2011). As such, the reduction in the conversion of MTS into formazan may not necessarily mean cell death. An LDH assay was therefore carried out to complement the MTS assay results. LDH is a cytoplasmic enzyme found in all cells and it can only be found extracellularly, such as in culture medium, in the event of cell membrane damage and/or cell death, where the amount of LDH released is proportional to the amount of dead cells

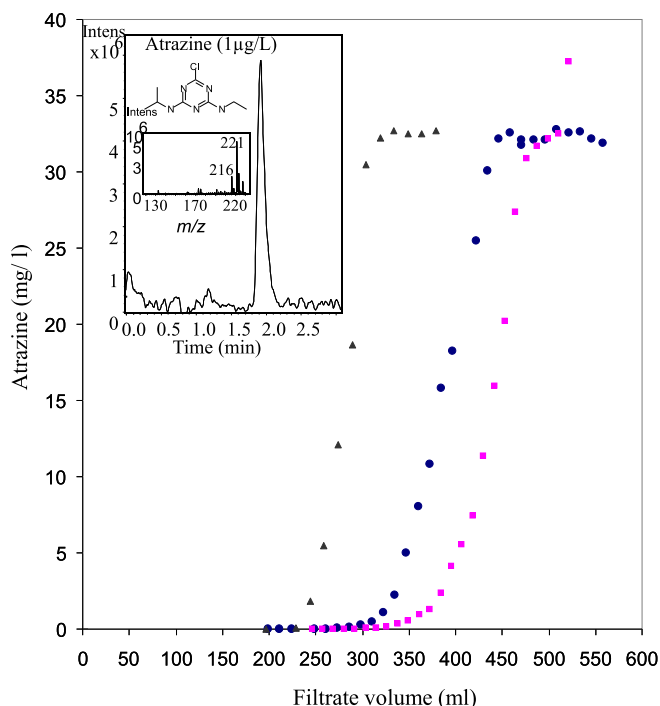


Fig. 5. Removal of atrazine spiked at 32 mg/L in ultrapure water with 1.4% ethanol (■); ultrapure water with potassium hydrogen phthalate added at the same concentration as atrazine (32 mg/L) (●); estuarine water (▲) by a carbon-cryogel composite ($1.5 \text{ i.d.} \times 85 \text{ mm}$, 0.6 mL bed volume). A chromatogram of atrazine and full scan mass spectra of atrazine m/z $[\text{M}+\text{H}]^+$ 216 and D_5 -atrazine m/z $[\text{M}+\text{H}]^+$ 221 are shown.

Table 1

Adsorptive capacities of the developed cryogel-carbon composite filters when filtering different waters spiked with atrazine at 32 mg/L (1.4% ethanol). The values given correspond to the last filtrate volume with atrazine below the statutory concentration limit (0.1 µg/L) (Council of the European Communities, 1998) and to the volume where the carbon beads reach saturation.

Filtration stage	Ultrapure water		Ultrapure water with added potassium hydrogen phthalate		Estuarine water	
	Adsorptive capacity (mg atrazine/g carbon)	b.v	Adsorptive capacity (mg atrazine/g carbon)	b.v	Adsorptive capacity (mg atrazine/g carbon)	b.v
At the last safe water volume (<0.1 µg atrazine/L)	295	428	214	435	229	327
Saturation of the filter	539	832	457	762	390	557

b.v: bed volume.

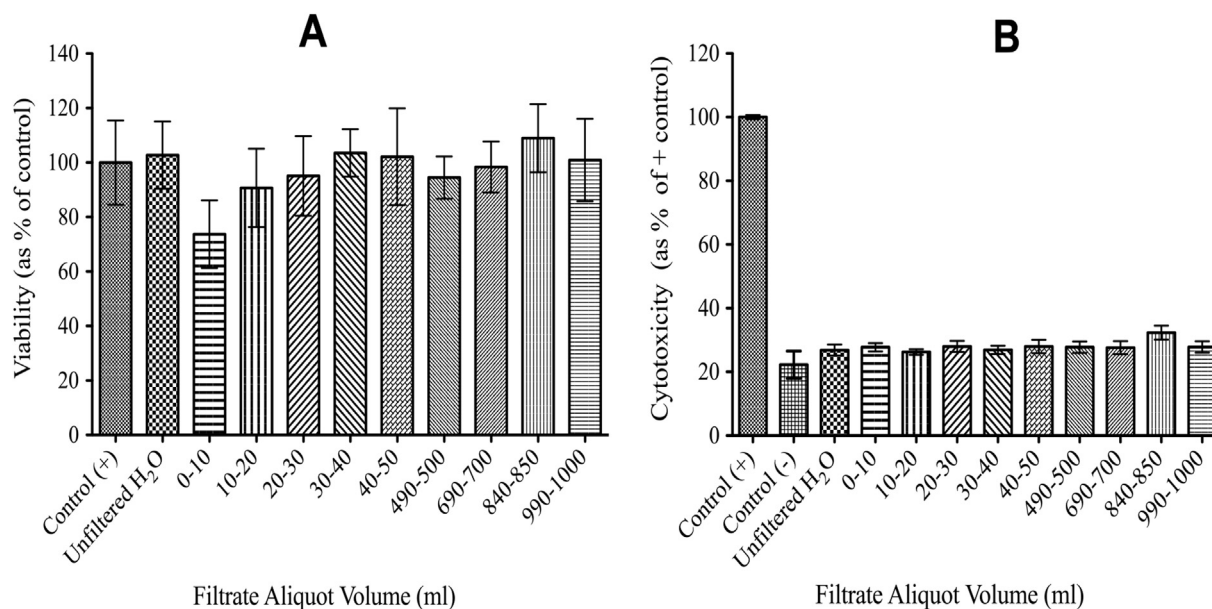


Fig. 6. Viability of Caco-2 cells exposed to (ultrapure) water passed through a PVA-carbon column. (A) Cell viability as determined by MTS assay and (B) cytotoxicity as determined by LDH assay. In the MTS assay, control (+) corresponds to cells that did not receive treatment with PVA-C filtrate water. In the LDH assay, control (+) and (–) represent total cell death and cells that did not receive PVA-C filtrate water, respectively.

(Jurisic, 2003; Smith et al., 2011). In this work, LDH assay results show no significant toxicity associated with water filtrate aliquots across all volumes, including the first 10 mL (Fig. 6B) ($P > 0.05$). This suggests that the reduction in cell viability seen in the MTS assay is not due to cell death but rather to a compromised mitochondrial activity. Crucially however, this problem of reduced metabolic activity can be addressed by an initial pre-wash of the PVA-carbon membrane/column with a volume of water that is 5 times the bed volume prior to use. Previous work has indicated that the carbon beads used in the composite filter do not produce a cytotoxic response against a V79 cell line (Barnes et al., 2009).

3.5.1. Potential implementation for water clean-up

The developed composites have shown high purification capacity at lab scale at liquid hourly space velocities (39 h^{-1}) which are greater than typical flow rates applied in water treatment plants. This makes them potentially useful both for low flow rate applications and for larger scale applications with higher flow rates and larger water volumes. Recent work by the authors (Savina et al., 2016) shows that cryogel-based composites of the type described here can be produced in volumes of 400 mL or greater in a monolithic (i.e. column filter) form with uniform pore size and carbon particle distribution, indicating the potential scalability of the filters. The use of a flexible polymer scaffold produced via

cryogelation also means that the composite filters can be produced in a variety of shapes: sheets for reactive barriers, discs, beads or monoliths (Savina et al., 2011), or within robust plastic carriers (Le Noir et al., 2009; Plieva et al., 2006) for more aggressive physical settings (e.g. in settlement tanks, fluidised beds, etc.). This provides significant flexibility in terms of device configuration.

4. Conclusions

Effective filters with superior adsorptive capacity for the removal of polar micropollutants, based on PVA macroporous cryogels with embedded phenolic carbon microbeads with controllable porosity, have been developed and tested for the treatment of contaminated water with high TOC levels. Embedding the carbon beads in a polymer support retains the adsorptive capacity of the carbons while improving the pore diffusion. The structure of the phenolic carbons, their micro- and mesopores, high S_{BET} and small particle size were shown to play an important role in the high adsorptive capacity of the filter. Adsorptive capacities of up to 641 mg atrazine/g carbon and 591 mg malathion/g carbon were obtained when filtering water containing 32 mg atrazine/L and 16 mg malathion/L (and 1.4% ethanol) in flow through experiments at 39 h^{-1} . The cytotoxicity of water passed through the filter at the same flow rate was tested with human epithelial cells, and no

toxicity was found from 17 to 1667 bed volumes. Prototypes developed at the lab scale are a promising water purification technology with superior adsorptive capacity.

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Appendix A. Supplementary data

Supplementary data related to this article can be found at <http://dx.doi.org/10.1016/j.jenvman.2016.07.061>.

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