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# Green synthesis of nanoscale anion exchange resin for sustainable water purification<sup>†</sup>

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The challenge of providing safe and reliable drinking water is being exacerbated by accelerating population growth, climate change, and the increase of natural and anthropogenic contamination. Current water treatment plants are not effective at the removal of pervasive, hydrophilic, low molecular weight contaminants, which can adversely affect human health. Herein, we describe a green all-aqueous synthesis of an ion exchange resin comprised of short chain polyelectrolyte brushes covalently bound to single walled carbon nanotubes. This composite material is incorporated onto a membrane and the active sites are tested against analyte adsorption. Our control studies of water or brine pushed through these materials, found no evidence of single-walled carbon nanotubes (SWCNTs) or carbon/polymer coming out of the membrane filter. We have measured the adsorption capacity and percentage removal of ten different compounds (pharmaceuticals, pesticides, disinfection byproducts and perfluoroalkylated substances). We have measured their removal with an efficiency up to 95–100%. The synthesis, purification, kinetics, and characterization of the polyelectrolytes, and the subsequent nanoresin are presented below. The materials were tested as thin films. Regeneration capacity was measured up to 20 cycles and the material has been shown to be safe and reusable, enabling them as potential candidates for sustainable water purification.

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#### Water impact

Improving water quality significantly impacts human health, the ecosystem, and economic logistics in all parts of the world. Centralized water treatment systems are ineffective in delivering water free from many persistent contaminants which arise due to natural and anthropogenic activities. We report on a fast, reliable and sustainable system that efficiently removes a wide varieties of these contaminants from water.

## Introduction

The US Environmental Protection Agency (USEPA) is focused on the monitoring and removal of many classes of compounds from drinking water and wastewater including disinfection byproduct (DBP) precursors, pharmaceuticals, personal care products, heavy metals, and per- and polyfluoroalkyl substances (PFAS).<sup>1</sup> These pervasive substances are related to adverse human health conditions and they are persistent in the environment. Under environmental conditions, PFAS do not hydrolyze, photolyze or biodegrade and hence, are extremely persistent in the environment.<sup>2</sup> It is becoming increasingly difficult for water treatment facilities to remove molecular contaminants at the limits set by the USEPA (e.g., PFAS 70 ppt, 80 ppb for trihalomethanes (THM) and 60 ppb for haloacetic acids (HAA)).<sup>3</sup> Possible health risks from prolonged exposure include kidney, liver, and central nervous system issues, as well as cancer.4,5 Many water utilities rely solely on coagulation as the means of lowering the levels of molecular contaminants; however, this is not as effective a method for low molecular weight and hydrophilic compounds. Activated carbon is the most widely used adsorbent material, but it comes with a high operating cost.<sup>6</sup> With natural and anthropogenic contaminant levels rising in drinking water sources worldwide, there is a significant increase in demand for more efficient removal.7-11 In general, there are a large number of compounds, across many chemical classes, that are naturally occurring in surface water. Many others are anthropogenic, and released into the environment due to improper treatment of wastewater. All drinking water sources contain natural organic matter (NOM), which can become DBPs during required water treatment

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processes. In our previous work we have demonstrated that our ion exchange nanomaterials can remove (NOM) faster, and at a higher capacity than other methods and materials currently deployed.<sup>12</sup> Since the publication of those results, we have optimized the materials and processes and here show that we can also remove other classes of compounds from water. Specifically, we demonstrate the removal of several types of pharmaceuticals (viz., tetracycline and carbenicillin), pesticides (viz., bentazon, terbacil, and bromacil), DPBs (viz., bromoacetic acid and chloroacetic acid) and PFAS (viz., perfluorooctanoic acid (PFOA) and perfluorooctanesulfonate (PFOS)), from water. We have synthesized these nanoresin materials using a green all-aqueous process.

Use of pristine nanostructured carbons in a synthesis typically requires a good dispersant to stabilize the nanoparticles. Often these dispersants are aprotic polar solvents like N,Ndimethylformamide (DMF) or N-methyl-2-pyrrolidone (NMP) which are both toxic and expensive. We have eliminated all organic solvents from our green all-aqueous synthesis of functionalized nanostructured carbon using the activators regenerated by electron transfer-atom transfer radical polymerization (ARGET-ATRP) method. Aqueous ARGET-ATRP reactions are complicated by multiple issues. The activation constants for initiators tend to be over an order of magnitude higher in water than in organic solvents.<sup>13,14</sup> This leads to an increase in ATRP polydispersity which is inversely proportional to the ratio of active to inactive chains.<sup>15</sup> In addition, changes in pH can lead to further complications such as the protonation of the catalyst. We have optimized a process that maintains low polydispersity during this all-aqueous synthesis of shortchain strong-base ion-exchange polymers. We have covalently functionalized these linear non-crosslinked polyelectrolytes to pristine SWCNTs using the same allaqueous ARGET ATRP process. These short brush-like polyelectrolytes form a conformal coating around the SWCNT resulting in a nanoresin with a ~10 nm cross-section, allowing access to all of the binding sites without being diffusion limited. The cationic nature of the polymer allows the nanoresin to be readily deployed in aqueous systems. This then promotes rapid adsorption of a broad spectrum of contaminants from drinking water or wastewater.

We have synthesized these nanoresins, using the ARGET-ATRP method where the reaction follows a living radical propagation mechanism. This method is preferable to traditional ATRP because of the reduced transition metal catalyst required and the increased control of the polydispersity.<sup>16</sup> The catalyst concentration can be kept to a minimum by using a reducing agent. A common ATRP catalyst is the Cu(1)– Br/TPMA complex (TPMA is tris(2-pyridylmethyl)amine). This is a very reactive complex in water and the use of a reducing agent is required for control of the reaction. A low ratio of the Cu(1) to Cu(1) form of the catalyst (activator/deactivator) is required for ATRP in water to reduce radical concentrations and maintain control of the polymerization.<sup>16</sup> Rapid addition of a reducing agent pushes the catalyst too far toward its activator form leading to poor polymerization control. Often ascorbic acid is added slowly to regenerate the catalyst. Here we describe a way to maintain the reducing agent concentration by using tin( $\pi$ ) 2-ethylhexanoate in the aqueous reaction. Sn( $\pi$ ) in water exists in many species but the Sn(OH)<sub>2</sub>(aq) form is prevalent at the pH range and ionic strength of our reaction,<sup>17,18</sup> and the  $K_{sp}$  of Sn(OH)<sub>2</sub>(s) is 5.45 × 10<sup>-27</sup>. As the reducing agent reacts to reactivate the catalyst, the Sn<sup>2+</sup> oxidizes to Sn<sup>4+</sup>, but the concentration of the reducing agent stays constant throughout the reaction due to the slow dissolution of the Sn(OH)<sub>2</sub>(s) back into the reaction solution. This enables a constant regenerator concentration, which in turns maintains a constant activator/catalyst concentration and thereby a constant rate of polymerization.

This paper describes a green synthesis of a polyelectrolyte modified nanostructured carbon material. We show outstanding adsorption loading and fast adsorption kinetics for several class of water contaminants. These materials are also regeneratable and reusable such that they enable a sustainable new material for water purification.

# Synthetic methods

#### Materials and methods

All reagents were used as purchased without additional purification or modification. HiPCO (Grade P CNT, 0.8-1.2 nm diameter, 100-1000 nm length; Lot #P0276) SWCNTs were used. Polymer synthesis was based on: vinylbenzyl trimethylammonium chloride (vbTMAC) (Fisher, 97%; Lot #A0311318) monomer, copper(II) bromide (Acros, 99+%; Lot #A0344238), tris(2-pyridylmethyl)amine (TPMA) (TCI, >98.0%; Lot #Z8GMO-AD) for the catalyst, tin( $\pi$ ) 2-ethylhexanoate (Sigma-Aldrich, 92.5-100%, Lot #SLBP5072V) for the reducing agent, 2-hydroxyethyl 2-bromo-isobutyrate (HEBiB) (Sigma-Aldrich, 95%, Lot # MKBW2607) initiator. Other analytes include: D-(+)-Glucose (Sigma, Lot #89H0150), toluene (HPLC grade, lot #), perfluorooctanoic acid (Aldrich, Lot #MKCC6736), potassium perfluorooctanesulfonate, 98% (Matrix Scientific, Lot #M22Q), tetracycline hydrochloride (Fisher BioReagents®, Lot #106895-36), carbenicillin disodium (Fisher Scientific, Lot #148020), bentazon (Ultra Scientific, Lot #NT059639), bromacil (Ultra Scientific, Lot #NT053779), terbacil (Ultra Scientific, Lot #NT059638), bromoacetic acid (99%: Acros Organics, Lot #106572500), and chloroacetic acid (99%: Lot B0143934) and sodium fluorescein (Sigma, Lot# BCBR1213V).

Workup of the synthesis required: polypropylene membrane filters (0.45  $\mu$ m pore size, 47 mm wide, Lot #2075-5), nitrocellulose mixed ester (MCE) (Sterlitech Corp., Lot. # 31127255) with 0.45  $\mu$ m pore diameter and 13 mm width, Whatman Anotop sterile syringe filter (25 mm, 0.02 micron pore size, Cat. No. #68092102), 12 kDa molecular weight cutoff (MWCO) dialysis membrane (Thermo Fisher Scientific, Lot. #AD-8099-2) 45 mm width and 28.6 mm diameter, 2 kDa MWCO dialysis membrane (Spectrum Labs, Lot. #3294218) 45 mm flat width and 29 mm diameter, and 50 kDa MWCO dialysis membrane (Spectrum Labs, Lot. #3292110) 34 mm flat width and 22 mm diameter. Membranes were activated by incubating for 10 minutes in Milli-Q water with constant stirring.

#### Aqueous synthesis of vbTMAC polyelectrolytes

Polymerization was performed using a Schlenk apparatus under a positive pressure of Ar(g). TPMA (6.32 mg, 21.8 µmol) and an aqueous CuBr<sub>2</sub> solution (17.24 µL, 20.6 mM, 0.35 µmol) were added into 1.0 mL of Milli-Q water to form the catalyst complex. Additional Milli-Q water (15.1 mL), vbTMAC (1.60 g, 7.57 mmol) were added and fully dissolved. The reaction mixture was sparged with Ar(g) for 20 minutes. HEBiB (10.3 µL, 71.0 µmol) was then added. Tin(II) 2-ethylhexanoate (6-60 µL) was added and the reaction flask heated in a 110 °C oil bath under reflux conditions for the duration of the reaction. 300 µL aliquots were taken for <sup>1</sup>H NMR analysis throughout the reaction. To reach 95% conversion of the monomer, the reaction proceeds for 4158 min and is then cooled and exposed to air to terminate the polymerization. The mixture was centrifuged at 100 000 RCF for 1 h at 20 °C to remove the thick polymer pellet. The polymer was dissolved in water and then dialyzed against Milli-Q water through a 2 kDa MWCO cellulose dialysis membrane to remove remaining monomer and other reagents. Up to eight dialysis iterations were performed until the monomer was no longer detected in the dialysate. vbTMAC is easily detected to  $<1 \text{ mg L}^{-1}$  by measuring the UV-vis absorption peak at 254 nm.

#### Aqueous synthesis of functionalized SWCNT nanoresin (Aq-SNR)

Polymerization was performed using a Schlenk apparatus under Ar(g). TPMA (6.59 mg, 22.7  $\mu$ mol) and CuBr<sub>2</sub> solution in water (17.3  $\mu$ L, 20.6 mM, 0.35  $\mu$ mol) were dissolved into 1.0 mL of Milli-Q water. Additional Milli-Q water (15.1 mL), vbTMAC (1.61 g, 7.62 mmol) and were added and fully dissolved. The reaction mixture was degassed with Ar(g) for 20 minutes. HEBiB (10.3  $\mu$ L, 71.0  $\mu$ mol) was then added, tin( $\pi$ ) 2-ethylhexanoate (6–60  $\mu$ L) was added and the reaction flask heated in a 110 °C oil bath for the duration of the reaction. 300  $\mu$ L samples were taken for <sup>1</sup>H NMR analysis throughout the reaction. DMF is added to the reaction (1% by volume) as an internal standard to quantify the remaining vinylic protons measured in the <sup>1</sup>H NMR spectra.

After 10 min, SWCNT (30.3 mg HIPCO) were added to a 5.0 mL aliquot of the reaction mixture. This mixture contained very short ionomers that were then sonicated with the SWCNTs to disperse the tubes into the polymer solution (15 min, 10  $W_{RMS}$ ) followed by ten min sparge under Ar(g). The SWCNT dispersion was added back into the polymerization reaction (40 min from the start of the polymerization, and a 33% conversion of monomer). The reaction was kept at reflux under Ar(g) until completion (typically from 2 h or up to three days for 99% conversion of any remaining vbTMAC

monomer). The reaction was cooled and exposed to air to quench the catalyst. SWCNTs do not disperse into water without the polymer present. The polyelectrolyte strands act as a surfactant and stabilizes the SWCNT dispersion which enable subsequent covalent functionalization of polymer to the nanotubes in an aqueous reaction.

Purification of as-synthesized Aq-SNR is required for the removal of unreacted monomer, other reaction reagents, and all non-covalently bound polymer strands. We have shown previously that these polyelectrolytes will wrap around the SWCNTs leaving them stable in water.<sup>12</sup> By increasing the ionic strength of the solution, the surfactant is screened from the SWCNT surface and the SWCNTs aggregate as the noncovalently bound polymer is removed. NaCl(s) was added to the reaction mixture to bring the concentration to 4 M. The reaction mixture was sonicated to disperse the product and mechanically disrupt the physisorbed polymer from the SWCNTs (15 min, 10 W). The attachment processes has not been optimized for atom efficiency. Moreover, since this is a fundamental study, only a small amount of SWNCTs are used for each batch and most,  $\sim$ 95% of the polymer is not used. Future studies with less expensive nanostructured carbons will optimize the atom efficiency. HCl(aq) was added to dissolve residual  $Sn(OH)_2(s)$  from the material. The dispersion was centrifuged (100 000 RCF, 60 min) in the high ionic strength solution. Desorbed polymer strands remain in the supernatant while the Aq-SNR collects in the sediment. Unbound polymer and other residual molecular and ionic species, are decanted and properly disposed of in hazardous waste (see ESI<sup>†</sup> Fig. S1). This procedure was repeated until the unbound polymer concentration was below our detection limit as measured by UV-vis absorption where  $\varepsilon_{254}$  = 0.0102 ± 0.0007 mg<sup>-1</sup> L cm<sup>-1</sup>, yielding a minimum detection limit of  $<0.3 \text{ mg L}^{-1}$  as shown in Fig. 1. Dialysis was used to remove the remaining NaCl(aq) from the sample, (multiple



**Fig. 1** Removal of vbTMAC monomer and unbound polymer after successive centrifugation and dispersal into brine cycles. UV-vis determination of concentration.

replicates, 75 mL internal volume, into 1000 mL external volume, through a 50 kDa membrane) (Fig. S2<sup>†</sup>). Conductivity of the dialysate was used to monitor removal of the brine until the conductivity was  $\kappa < 1 \ \mu\text{S cm}^{-1}$ . The final Aq-SNR product was dispersed into Milli-Q water. The concentration of SWCNTs was measured by UV-vis-NIR. The total mass concentration was measured by filtering a known volume of the Aq-SNR dispersion using a 0.45 µm pore size polypropylene membrane filter, which retained all of the nanoparticles. The thin film was washed with methanol then dried under vacuum at T = 65 °C overnight. The mass of the Aq-SNR deposited onto the membrane was measured by subtracting off the initial mass of polypropylene membrane. Concentrations are reported in mg Aq-SNR L<sup>-1</sup> of dispersion. Synthesis and cleanup of Aq-SNR did not use any organic solvents (Green Chemistry Principle 3, 4, and 5).<sup>19</sup>

## Characterization methods

#### Kinetics of polymer growth

Polymerization kinetics were measured by monitoring monomer concentration versus time as determined by <sup>1</sup>H NMR analysis. Standard polymer synthesis (1.50 g, 7.10 mmol vbTMAC) were monitored by pulling 300 µL aliquots as a function of time. For each aliquot of reaction mixture, 200  $\mu$ L was mixed into 500 µL D<sub>2</sub>O. Proton spectra taken on a 500 MHz <sup>1</sup>H NMR (32 scans, 40 second relaxation time) and baseline corrected. To quantify the relative concentration of monomer, the vinylic protons of the vbTMAC at 5.8 ppm and 5.3 ppm were compared to the singlet at 7.77 ppm (internal standard DMF aldehyde proton). Integrated areas decrease in value relative to the internal standard as the monomer was polymerized. The percent conversion for each aliquot was calculated accordingly. Loss of vinylic protons during polymerization is illustrated in Fig. S3.† Percent conversion versus time is illustrated in spectra in Fig. S4A<sup>+</sup> and pseudo first order  $\ln([M]_o/[M])$  vs. time in Fig. S4B.<sup>†</sup>

#### UV-vis-NIR spectroscopy

Concentration and yield of the Aq-SNR was measured using UV-vis-NIR spectroscopy. The concentration of SWCNT was measured using the absorption at 925 nm, with an extinction coefficient of 0.0206 mg<sup>-1</sup> L cm<sup>-1</sup>. From the total volume of dispersion and the concentration of SWCNTs determined from characteristic UV-vis-NIR absorption, the mass of the functionalized SWCNTs was calculated. The polyelectrolyte strands do not absorb at this wavelength. All mass and concentrations are reported as mg Aq-SNR or mg Aq-SNR L<sup>-1</sup> of dispersion respectively.

# Raman spectroscopy for covalent functionalization of SWCNTs

Samples of pristine SWCNT and of Aq-SNR were deposited onto a 0.45  $\mu$ m pore size polypropylene membrane filter as a thin film. A Kaiser Raman spectrometer was used to measure

the integrated area under the D band (sp<sup>3</sup> character C) relative to the integrated area under the G band (sp<sup>2</sup> character C). The D:G ratio of the Aq-SNR increased relative to the D: G ratio of the pristine sample which indicates covalent binding between the polyelectrolyte strands and the SWCNT surface.<sup>20</sup> Typical results of a 2 h functionalization are shown in Fig. 2, with the D:G ratio increasing from 0.16 to 0.30.

#### Total organic carbon (TOC) analysis of aqueous analyte

Concentrations of control and sample solutions were determined by total organic carbon (TOC) using a Shimadzu TOC-LCPN instrument following standard methods.<sup>21</sup> All amber glass collection vials were prepared to remove residual carbon. The vials were pre-cleaned by rinsing ten times in Milli-Q water then placed in acid bath (10% HCl(aq)) overnight. They were rinsed multiple times in Milli-Q water to remove the acid solution, then covered with aluminum foil, and heated at 400 °C for at least 2 hours in air to remove any residual carbon in the vials. The TOC system was calibrated using standard dilutions of a stock aqueous glucose solution. The response was linear over the range 0.00 mg L<sup>-1</sup> to 10.00 mg L<sup>-1</sup> with an accuracy of ±0.03 mg L<sup>-1</sup>. All samples were diluted by a factor of 15 before analysis.

#### Dynamic light scattering (DLS) and zeta potential

The effective diameter and zeta potential of poly(vbTMAC) and Aq-SNR dispersion was measured using Malvern DLS Zetasizer instrument.<sup>22</sup> DLS samples of 10  $\mu$ g mL<sup>-1</sup> of poly(vbTMAC) and 7  $\mu$ g mL<sup>-1</sup> of Aq-SNR were dispersed in Milli-Q water just before measurement. All zeta potential measurements used 2 mg L<sup>-1</sup> samples. The temperature of the cell was kept at 25 °C and the equilibrium time was 120



Fig. 2 Normalized Raman spectra of a 2 h pristine SWCNT and Aq-SNR after centrifugation at 20 000 RCF for 1 h. Increase in the ratio of the area under the D and G bands (the D:G ratio) indicates covalent functionalization of ionomer strands to SWCNT wall.

s. Multiple runs were averaged after maximum and minimum outliers were discarded.

#### Scanning electron microscopy (SEM)

Dispersions of Aq-SNR were diluted and resonicated then drop cast and dried onto a Si chip surface. Drying induced aggregation was kept to a minimum. A LEO 1550 column with field emission gun and 2 nm beam width was used for imaging at 10 kV and a 30  $\mu$ m aperture. Fig. 3 compares pristine SWNCTs (3A) to the Aq-SNR material. Some drying induced aggregation is found in all samples. Individual SWCNTs appear as wide as the FE-SEM beam width (2–3 nm). Polymer coated nanotubes aggregated (3B), but even the narrowest tubes show conformal coating of polymer with 8– 10 nm diameter. Morphology of thin membrane surfaces was also measured (3C) and is discussed below.

#### Analyte adsorption capacity studies

Multiple analytes were tested to evaluate their broad spectrum removal by the Aq-SNR. Sodium fluorescein (NaFL) was used as a low MW (376.27 g mol<sup>-1</sup>) surrogate for fulvic acids.

Fig. 3 FE-SEM images of pristine SWCNTs (A) and Aq-SNR (B) drop cast from a dilute dispersion onto a Si chip. Scale bar 200 nm and column conditions identical for all images. Survey of Aq-SNR shows no residual free polymer or salts. In 3(C) we show typical thin film of Aq-SNR deposited on an MCE membrane.

Tetracycline hydrochloride and carbenicillin disodium were used to evaluate removal of pharmaceutical compounds. PFOA and PFOS were used to test the removal of perfluorinated compounds. Bromoacetic acid and choloroacetic acid were used to evaluate removal of halogenated compounds (*e.g.*, DBP). Bentazon, terbacil, and bromacil were used to evaluate removal of pesticides from water. NaFL sample concentrations were measured using UVvis spectroscopy ( $\varepsilon_{490} = 0.358 \pm 0.011 \text{ mg-C}^{-1} \text{ L cm}^{-1}$ ). All other analyte concentrations were measured using TOC analysis.

#### Adsorption capacity: fast-filtration

A known mass of **Aq-SNR** was deposited on an MCE membrane by filtration. Control studies were performed to determine adsorption of the analyte by the apparatus without any **Aq-SNR** present. 3.0 ml of each analyte solution, of known initial concentration, were pushed manually through the film using a syringe. Low backpressure allowed the filtration to be completed in less than ten seconds. The concentration of analyte in the filtrate was measured using UV-vis spectroscopy or TOC analysis.

#### **Regeneration and reuse studies**

A known mass of **Aq-SNR** was deposited onto an MCE support membrane and placed into a glass vial. A NaFL solution of known concentration was added to the vial. The sample was vortexed for 2 h at 500 rpm while small aliquots were taken at intervals to test adsorption kinetics. Fig. S5† shows the loading q of NaFL adsorbed onto the outer interface of the membrane as a function of time. To simulate prebreakthrough conditions we tested the outer interface of the thin film. The adsorption data are fit to a pseudo-second order diffusive kinetics model with a rate constant k = 0.0048 s<sup>-1</sup> (mg g<sup>-1</sup>)<sup>-1</sup>.

For regeneration studies, a known mass of **Aq-SNR** was deposited onto MCE membrane. The film was placed inside a vial and was allowed to incubate in brine solution (1.5 ml, 2.0 M NaCl(aq)) for 5 min. After removal of the brine, Milli-Q water was used to rinse the remaining brine from the vial and resin (3 replicates, 1.5 mL). A known volume and concentration of NaFL(aq) was then added and allowed to incubate for 5 min. The concentration of the NaFL(aq) solution after incubation was measured by UV-vis spectroscopy and the process was repeated for 20 cycles.

## **Results and discussion**

#### **Characterization of Aq-SNR**

Kinetics of polyelectrolyte growth were measured by <sup>1</sup>H NMR as described above. We have not yet optimized the performance of the Aq-SNR as a function of polyelectrolyte length. We use a 33% conversion (40 min polymerization) for all polymer synthesis before adding SWCNTs for functionalization. Since we use a 100:1 monomer to initiator

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ratio these polyelectrolytes should have a molecular mass of about  $33 \times 211.73$  g mol<sup>-1</sup> or 7 kD. Using DLS we measured an average hydrodynamic diameter of 1.7 nm three separate syntheses of 25% converted polyelectrolyte. Data shown in Fig. S6A;† These data are consistent with radius of gyration data on similar polyelectrolytes. Moreover, since there is only one peak in the DLS analysis of the effective diameters we conclude that there is a minimal amount of bi-molecular radical termination during the polymer growth. Any polymer end termination would result in loss of atom economy since those strands could not then react with the SWCNTs. The multi angle light scattering analysis by Takahara *et al.* show a ~2 diameter  $(2 \times R_{\rm H})$  for 7 kD polyelectrolyte in low salt solution.<sup>23</sup> Their scattering model is consistent with a wormlike cylinder or brush polymer morphology. Zeta potential data (shown in Fig. S6B<sup>†</sup>) measured a +8.8 mV which is consistent with a strong base polyelectrolyte.

Aq-SNR was characterized by UV-vis-NIR to get the concentration of functionalized SWCNT as a stable dispersion, reported in mg-SWCNT L<sup>-1</sup>. Centrifugation of 200 000g is required to completely sediment the polyelectrolyte functionalized particles. The total concentration of Aq-SNR includes covalently bound polymer and SWCNT mass, mg L<sup>-1</sup> and the total mass to SWCNT mass ratio was used to get total polymer mass for adoption testing. Energy dispersive X-ray spectroscopy (EDX) on thin films of the Aq-SNR found ~2% Sn by mass (Fig. S7†). Residual tin from the Sn(OH)<sub>2</sub> equilibrium was removed from the dispersion by sedimentation and then by washing with HCl(aq) and then Milli-Q water. There may be some SnO<sub>4</sub>(s) that we cannot dissolve easily. This residual contaminant will be addressed in subsequent scale-up of our reactor design.

Conformal coating of polymer around the SWCNTs is shown in the FE-SEM image Fig. 3B. It is important to note that there is no free polymer aggregates or salt on these samples. To use the Aq-SNR for analyte removal we form a thin film on an MCE membrane. These films show very smooth surface topography on top of the MCE membrane (surface morphology due to MCE membrane). Slightly aggregated polymer coated SWCNTs are shown in Fig. 3C. The thin film exhibits a pin hole free, mesoporous architecture. A survey of the film as measured by SEM is shown in Fig. S8.† Fig. S9† shows cross-sectional AFM data. The film thickness is ~140 nm for this sample with a total mass of 0.17 mg and mass density of 0.046 mg cm<sup>-2</sup>. The film is not vacuum dried before AFM and the probe tip is interacting very strongly with the surface charges of the polyelectrolytes.

Water samples must pass through a tortuous morphology such that the analyte can easily access the very high surface area Aq-SNR material. The open polymer microstructure and thin film morphology enables this "contact resin" behavior. This method of analyte removal was chosen over incubation so as to simulate ultrafiltration (UF) water purification technology. Intrinsic membrane resistance of these films was calculated by measuring the flux of water at a constant transmembrane pressure of  $1.0 \times 10^5$  Pa, according to Darcy's law.<sup>24,25</sup> The measured membrane resistance  $\kappa_m = 2.1 \times 10^{11}$  m<sup>-1</sup> is smaller than typical UF membranes with a  $\kappa_m$  = 2 ×  $10^{12}$  m<sup>-1</sup>. A modest loading value  $q_e = 20$  mg g<sup>-1</sup> with a 1 m<sup>2</sup> membrane at 0.9 mg cm<sup>-2</sup> mass density, would effectively remove 95% PFAS from  $10^5$  L of 1 mg-C L<sup>-1</sup> effluent. Our results are competitive with other functionalized nanostructured materials. A recent study of PFAS removal by fluorographene found an equilibrium loading  $q_{\rm e}$  < 5 mg-C  $L^{-1}$  of PFOA and PFOS at equilibrium concentrations <3 mg  $L^{-1}$  analyte.<sup>26</sup> In Table S14<sup>+</sup> we show the equilibrium loading data for our Aq-SNR membranes of  $q_e = 44$  mg-C L<sup>-1</sup> and  $q_e =$ 65 mg-C  $L^{-1}$  at equilibrium concentration ~1.5 mg-C  $L^{-1}$  for PFOA and PFOS respectively. The larger loading onto the Ag-SNR materials, event at lower analyte concentration than the fluorographene work, is consistent with the larger binding constants we have reported (116  $\pm$  3 (mg-C g<sup>-1</sup>) (mg-C L<sup>-1</sup>)<sup>-1/n</sup> for similar anionic analytes.<sup>12</sup>

We monitor covalent functionalization of the polymer to the SWCNTs by measuring the D:G ratio from Raman spectroscopy. The larger the D:G ratio, the more defects we introduce into the SWCNT's wall. The living radical end of the polymer reacts easily with  $\pi$  conjugated electron density on the SWCNTs and other nanostructured carbons.<sup>27</sup> We have not fully optimized the functionalization process. However, the Raman spectroscopy suggests that the attachment of polyelectrolytes to the SWCNTs is quite fast. Baseline-normalized Raman spectra in the D and G band region are overlaid in Fig. 2 and in Fig. S10.<sup>†</sup> The area under each band is calculated and the D: G ratio for pristine versus functionalized Aq-SNR is compared. The D:G ration for the 2 h, 12 h, and 3 day functionalization are similar, 0.30, 0.33, and 0.33 respectively. While the D:G ratio does indicate covalent attachment of polyelectrolyte strands to the SWCNT it does not give a quantitative measure of the degree of functionalization of the Aq-SNR. We use the supernatant of the centrifuged samples to partially purify the samples. It is clear that the Aq-SNR is not uniformly coated with polymer. During purification of the material some of the nanostructured carbon material sediments at low <10 000g. The D: G ratio of the sediment material after a 20 000g treatment is 0.26. Additionally, we have analyzed the Raman D:G ratio of Aq-SNR from the supernatant of a 200 000g centrifugation. The material that stays in the supernatant at that high a RCF, is stabilized with polyelectrolyte attachment. The spectra in Fig. S10B<sup>†</sup> shows a D:G ratio of 0.93 from material collected from the supernatant only. So this ratio is a measure of how stable the functionalized tubes are during centrifugation. Moreover, supernatant from samples after centrifugation at 20 000g have a higher zeta potential, 38.4 mV versus the zeta from the sample before any centrifugation, 22.8 mV as shown in Fig. S11<sup>+</sup> for the 2 h functionalized Aq-SNR. Changes to our reactor design should allow us to optimize functionalization density.

DLS of 2 h, 12 h and 3 day Aq-SNR samples are shown in Fig. S12.† Calculated hydrodynamic effective diameters for 2 h, 12 h, and 3 day are 213  $\pm$ 5 nm, 205  $\pm$  5 nm, and 213  $\pm$ 7 nm respectively. The effective diameter is calculated from the measured diffusion coefficient and assumes a spherical

model. These particles are more rod like. From SEM we estimate the particle width to be w = 10 nm. Then for a flexible rod length L = 1000 nm, the calculated hydrodynamic diameter is  $D_{\rm h} = L/(\ln(L/w) + 0.32) = 203$  nm, which is consistent with our measured values.<sup>28</sup>

Fully characterized materials are then deposited as thin films and used to remove analyte from water samples. The ratio of the total mass of Aq-SNR (measured by filtering out the purified material) to the SWCNT mass only (measured by the NIR absorption at 1025 nm) increases as functionalization time increases. These data are listed in Fig. S10A.<sup>†</sup> The 12 h sample has a higher amount of polymer attachment per SWCNT mass than the 2 h (3.73 ratio to 2.40 ratio respectively). For this fundamental study we only add a small amount, ~30 mg of SWCNTs to the polymer solution during the attachment reaction. The SWCNTs are added to the ARGET ATRP reaction at the same time (after ~33% conversion). As the reaction proceeds, the radical end of the 33-mer attacks the  $sp^2$  carbon on the SWCNT walls. But, many of the polyelectrolyte chains continue to react with the other  $\sim 60\%$ of the vbTMAC monomer resulting in longer chains, with more mass, which then subsequently attach to the SWCNTs. After removal of unbound polyelectrolyte the mass of polymer attached to the SWCNTs is 2.4-4.8 times greater than the mass of the SWCNTs.

#### Adsorption capacity testing

To compare different adsorbent materials with various adsorbate compounds we measure the equilibrium loading capacity  $q_{\rm e}$  in mg analyte per g adsorbent. In our previous works we published and analyzed the adsorption isotherm analysis and reported that these thin film nanoresin materials behave like "contact" resins which reach a pseudo-equilibrium in only a few seconds of exposure. We have shown that we can reproducibly synthesize the Aq-SNR, We measured the adsorption capacity of each batch of Aq-SNR against NaFL, which we use as a surrogate for natural water DBP precursors. A known volume and concentration of analyte is passed through a known mass of Aq-SNR as a thin film membrane. All fast-filtration experiments take less than 10 seconds. From control studies of water or brine passed through the Aq-SNR/MCE filters, we found no evidence of SWCNTs or carbon/polymer coming out of the membrane filter. Results for  $q_e$  of NaFL on 2 h and 12 h Aq-SNR are shown in Fig. 4. We compare  $q_e$  using an initial concentration  $C_0 = 5.8 \text{ mg-C} \text{ NaFL L}^{-1}$ . The SWCNT mass only, was used so that we can directly compare the material based on number of nanotubes in the film. We show the comparison of an addition four syntheses of 2 h and 12 h Aq-SNR binding capacity in Fig. S13.<sup>†</sup> All ten nanoresin samples showed a  $q_e$  of ~100 mg-C NaFL per g SWCNT, a 10% loading capacity in <10 seconds of contact time.

#### Percentage removal of analyte

The USEPA is evaluating systems for PFAs contamination reduction by challenging them with an influent of  $1.5 \pm 30\%$ 



**Fig. 4** Equilibrium loading capacity of NaFL onto **Aq-SNR** using an initial concentration  $C_{\rm o} = 5.8$  mg-C L<sup>-1</sup>. The mass of the adsorbent is the SWCNT mass only so we can directly compare the material based on number of nanotubes in the film.

 $\mu$ g L<sup>-1</sup> (total of both PFOA and PFOS)<sup>4</sup> and must reduce this contamination by more than 95% to 0.07  $\mu$ g L<sup>-1</sup> or less. This Aq-SNR also removes other classes of analyte from water. In Fig. 5 we show percent removal data for PFOS, PFOA and the antibiotic carbenicillin disodium, and the pesticide bentazon. Membrane mass was increased until at least 95% of the analyte was removed in one fast-filtration pass through the Aq-SNR film. The mass density for 95% removal of carbenicillin disodium, and bentazon was 0.13 mg cm<sup>-2</sup>. The required mass density for the PFAS compounds was 0.90 mg cm<sup>-2</sup>. The higher mass required for the PFAS is consistent with the challenge of removing these hydrophilic and persistent contaminants from water. The initial concentration of all analytes ranged between 2-4 mg-C L<sup>-1</sup>. The concentration was measured using TOC analysis (mg-C L<sup>-1</sup>). The limit of detection of our TOC analyzer and method is only ~0.2 mg-C  $L^{-1}$ . Further studies, at the lower EPA detection limits, will require tandem mass spectrometry.29,30

Relative equilibrium adsorption capacity of the Aq-SNR against several other compounds are listed in Table S14.<sup>†</sup> In addition to the surrogate compound there are four classes of compounds that these materials can remove from water effectively. The equilibrium loading  $q_e$  in mg-C analyte per g Aq-SNR are from fast-filtration experiment where the sample was pressed through the membrane in just a few seconds. In its current form, all of the Aq-SNR are anion exchange resins so they are most effective at removing compounds with some negative charge character. With the exception of the tetracycline HCl, all of these compounds have negative zeta potentials at natural water pH 6.5–7.5. The absolute value of the  $q_e$  is not the subject of this study. Percent removal is the most important quality and we have shown near complete removal



Fig. 5 Percent removal of analyte versus Aq-SNR membrane mass. Membrane area for PFOA and PFOS is 0.58 cm<sup>2</sup> and is 3.78 cm<sup>2</sup> for carbenicillin and bentazon removal.

of most of these contaminants from water test samples. Future work will test our cation exchange resins and our materials in which we optimize for a specific analyte, like short chain PFAS.

#### Regeneration and reuse: a sustainable material

Similar to our previous work, we show here that these materials can be regenerated and reused many times without loss of adsorption capacity. To simulate pre-breakthrough behavior we did not push the analyte through the membrane, rather we allowed it to diffuse into the outer membrane-solution interface for a 5 min incubation while stirring. Adsorbed surrogate mass was measured by UV-vis of remaining NaFL in solution after incubation. The film was then briefly incubated with brine, rinsed with water and then re-exposed to analyte solution. This was repeated for 20 cycles and the data shown in Fig. 6. The slope of these data is -0.2% per cycle and the 95% confidence interval error in the

slope is 0.2% per cycle. The loss of performance from a single cycle of use is statistically indistinguishable from zero. We do not measure any residual polymer degradation when we run control regeneration studies. The regeneration study was done in triplicate. Other results have a slope of  $-0.13 \pm 0.3\%$  per cycle. Extrapolating out, this material should maintain up to 50% of its effectiveness for over 300 cycles. Subsequent studies of these materials will not use SWCNTs because of their expense. This polyelectrolyte attachment chemistry is applicable to many other, less expensive, nanostructured carbon materials. This is also a scalable synthesis leading to green and sustainable manufacturing process.

## Conclusions

We have demonstrated an all-aqueous synthesis and purification of functionalized SWCNTs with short chain polyelectrolyte resins using a modified facile ARGET ATRP mechanism. The synthesis and purification procedure did not involve any



**Fig. 6** Regeneration and reusability study. Surrogate NaFl used to test adsorption by incubating against a thin film of Aq-SNR. 5 min brine incubation followed by water rinse before next exposure to surrogate adsorbate. Process repeated for 20 cycles. Experiment repeated in triplicate. 0.23% decrease in *q* per cycle is close to the 95% confidence interval error in the slope.

organic or hazardous reagents and control studies found no traces of SWCNTs or unbound polymer released from the materials during water treatment. Polyelectrolyte resin chains enable fast equilibration with analyte compounds in the influent solution, while the SWCNTs form a mesoporous scaffolding and provide a tortuous, but low resistance, path for the analyte to flow through. The covalent attachment of resins to SWCNTs is also confirmed with Raman spectroscopy and zeta potential measurements of highly stabilized dispersions. The DLS, zeta potential and the adsorption capacity of various Aq-SNRs synthetic batches show consistency in hydrodynamic diameters, high surface charge, and adsorption performance. We have shown that these materials successfully target and electrostatically bind to several classes of pervasive compounds. We showed the effectiveness of these thin films by testing them against several compounds and reducing the analyte concentration by 95-100%. We have demonstrated a sustainable material that can be regenerated and reused many times. We have shown that the nanostructured carbon materials do not leach or escape from the membrane and therefore are safe to the environment<sup>31,32</sup> and drinking water applications (Green Principle 12).<sup>19</sup> Implementing these materials into existing membrane filtration systems should increase removal capabilities and lower operating costs of water treatment facilities. This fundamental proof-ofprinciple study will be extended to other nanostructured carbon materials that are much less expensive than SWCNTs, thereby lowering the cost of scaling this methodology.

# Safety

Personal protective equipment should be used when doing any of the experiments described in this paper. Special care should be taken when working with evacuated glassware and Schenk line apparatus. All chemical waste should be disposed of as hazardous waste in accordance with regulatory agency oversight.

# Conflicts of interest

There are no conflict of interests to declare.

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