Enhanced adsorptive composite foams for copper (II) removal utilising bio-renewable polyisoprene-functionalised carbon derived from coconut shell waste

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A bio-renewable polyisoprene obtained from Hevea Brasiliensis was used to produce functionalised carbon composite foam as an adsorbent for heavy metal ions. Functionalised carbon materials (C-SO3H, C-COOH, or C-NH2) derived from coconut shell waste were prepared via a hydrothermal treatment. Scanning electron microscopy images showed that the functionalised carbon particles had spherical shapes with rough surfaces. X-ray photoelectron spectroscopy confirmed that the functional groups were successfully functionalised over the carbon surface. The foaming process allowed for the addition of carbon (up to seven parts per hundred of rubber) to the high ammonia natural rubber latex. The composite foams had open pore structures with good dispersion of the functionalised carbon. The foam performance on copper ion adsorption has been investigated with regard to their functional group and adsorption conditions. The carbon foams achieved maximum Cu(II) adsorption at 56.5 mg g⁻¹ foam for C-SO3H, 55.7 mg g⁻¹ foam for C-COOH, and 41.9 mg g⁻¹ foam for C-NH2, and the adsorption behaviour followed a pseudo-second order kinetics model.

Ecosystems and the environment as a whole are threatened by the continuous heavy metal contamination of natural water sources. A considerable increase in the use of heavy metals in industries such as electroplating, metal finishing, metallurgy, tanneries, chemical manufacturing, mining, and battery manufacturing have led to the release of large amounts of untreated wastewater into the environment1. Unlike organic pollutants, heavy metal ions and their compounds are not degradable and have a tendency to accumulate in living beings, often leading to toxic or carcinogenic effects. At high concentrations, Cu(II) has been linked to liver and kidney damage, and increased blood pressure and respiratory rates in humans, while Ni(II) is associated with the lung and kidney disorders, chest pain, and shortness of breath5,6. The hexavalent form of chromium is a well-known human carcinogen and can cause liver damage, pulmonary congestion, and skin irritation, which result in ulcer formation7. Therefore, these metals must be removed from wastewater before their discharge into natural water bodies, and an effective method for doing so must be developed.

Numerous conventional methods for the removal and recovery of heavy metals from contaminated industrial waste streams have been reported. These processes can be classified into three types: chemical, physical, and biological, where physical and chemical processes are more widely implemented. Chemical methods include flotation, coagulation/flocculation, chemical precipitation, electrochemical processes, and ion exchange. Membrane
Composites of inorganic or organic fillers with natural polymer matrix have received greater attention due to their adsorption ability. Special characteristics such as antibacterial activity by adding Ag nanoparticles, magnetic properties by adding ferroferric oxide particles, and mechanical/thermal resistant property by adding organoclay can be designed. Moreover, various petroleum-based polymers have been used for adsorbent preparation such as poly (ethylene glycol dimethacrylate- n-vinyl imidazole) and poly(divinylbenzene- n-vinylimidazole)6-9 for specific adsorption of metal ions. Since the composites mainly consist of fillers and polymer matrix, the functionalisation of fillers required proper modification to ensure high dispersion. In this study, composites foam of functionalised activated carbon with a natural polymer matrix was selected because they are naturally available.

Activated carbon is a versatile adsorbent owing to an extensive surface area, microporous structure, strong and universal adsorption capacity, and high degree of surface reactivity. Carbon materials have been functionalised with a wide variety of functional groups. Sulfonyl groups (−SO3H) can be added using hydrothermal carbonisation and sequential sulfonation to produce carbon with a porous functionalised surface28-30. Carbon foam is a popular alternative adsorbent because of its strength, low weight, and low cost and is typically prepared by dip coating polyurethane (PU) with pitch slurry, activated carbon slurry, or metal oxide solution, followed by carbonisation. Three-dimensional porous carbon-based materials have been used for dye adsorption, heavy metal adsorption, and electromagnetic shielding15-17. Carbon foam can remove heavy metal ions from aqueous solutions including copper, zinc, cadmium, lead, and nickel16,18. Previous reports have investigated the use of a synthetic polymer (e.g. PU foam or phenolic resin-based foam) as a support material. To prevent the use of environmentally harmful materials, this study aimed to investigate the formation of functionalised carbon and carbon foam for heavy metal adsorption using a bio-renewable material. Porous carbon obtained from coconut shells with the groups functionalised with sulfonic acid (−SO3H) and amine (−NH2) groups. Good dispersion and distribution throughout the foam was required, and the adsorption efficiency was investigated. The effects of the pH of heavy metal solution, contact time, and concentration of the adsorbate on adsorption were evaluated to better understand the adsorption kinetics.

For Cu(II) adsorption, different type of copper complex has been used, for example copper chloride, copper acetate, copper nitrate or copper sulfate. Various techniques, such as, spectrophotometer, atomic adsorption spectroscopy (AAS), and inductively coupled plasma spectrometer (ICP), have been used according to the amount of Cu(II) in aqueous media. Atomic adsorption spectroscopy (AAS) can be applied for Cu(II) concentration range of 5–400 ppm which the measurement by AAS perform higher sensitivity than spectrophotometer20-22. Trace amount of Cu(II), such as 5–200 ppm, can be studied by using ICP which perform highest sensitivity compare to spectrophotometer and AAS24.

Our research studied Cu(II) adsorption by using spectrophotometer measurement. A similar work using spectrophotometer on Cu(II) adsorption studied with 500 ppm25. The permissible limit of Cu(II) in water is less than 2.5 mg L−1. Several treatment technologies such as solvent extraction, ion exchange, and precipitation have been suggested and employed to remove heavy metals from aqueous solution, however, these technologies are costly and also create another problem with metal-bearing sludge27. Therefore, our research work focused on an economical and available adsorbent for removing copper from aqueous solution by using simple technique such as spectrophotometer.

Results and discussion

Characteristics of functionalised carbon and carbon foam. The untreated carbon particles have an irregular morphology and appear as partially flattened cylinders, as shown in the SEM images (Fig. 1a). After functionalisation via hydrothermal treatment, the particles become smaller and spherical with a rough surface (Fig. 1b). An example of the C-SO3H particles is shown in Fig. 1c, where clarity of elements distribution allows the elemental mapping. The foam surface primarily contains C, but S from the sulfonyl group is evenly distributed across the carbon surface. C-NH2 and C-COOH could not be clearly characterised using SEM-EDX, but S from the sulfonyl group is evenly distributed across the carbon surface. C-NH2 and C-COOH could not be clearly characterised using SEM-EDX, and therefore, XPS was used to evaluate the binding energy of the functionalised carbon. From Fig. 1d, the (BET) surface areas (S_{BET}) of the carbon samples decrease after hydrothermal treatment, particularly for the C-COOH samples. This decrease in the surface area is attributed to the structural deformation or collapse of the carbon framework caused by the strong reaction conditions24. SEM-EDX was used to characterise the distribution of the functional groups across the carbon surface.

The FT-IR spectra of the sylphonated carbon (C-SO3H) exhibits the absorption peaks at 1702, 1588, and 1155 cm−1 corresponding to C=O, C=C, and C=O bonds, respectively (Fig. 2a). The peak at 1030 cm−1 is attributed to the −SO3H group. Aromatic C–H bonds and O–H bonds in the carboxylic acid and phenol groups absorb strongly from 3600 to 2400 cm−1. The spectra of the carboxylated carbon (C-COOH) is characterised by an adsorption band at 1697 cm−1, attributable to the carboxylate group. The signals at 1607 cm−1 and 1176 cm−1 correspond to C=C and C–OH stretching, respectively, and indicate that the residual hydroxyl groups are present at the hydrophilic surface. The spectra of the carbon modified with ammonia (C-NH2) exhibit a broad peak at 3242–3400 cm−1, and the peak at 1591 cm−1 is attributed to the stretching and bending vibrations of –NH2, indicating that the amino groups are present at the surface. The peaks in the 1000–1400 cm−1 range correspond to the oxygen-rich groups (C–O) in the amino-functionalised carbon.

XPS was used for the analysis of the surface composition by determining the elemental composition of the material and binding energy of the elements on the surface. The high-resolution XPS survey spectra shown in Fig. 2b–f verify the composition of the samples. Wide-scan survey spectra confirm the presence and atomic concentrations of the elemental C, O, S, and N on the carbon sample surfaces (Table 1). The major elements are C (90.89%) and O (9.11%), and the atomic concentration of O increases in the carboxyl-functionalised carbon.
Figure 1. SEM images of (a) activated carbon and (b) C-SO$_3$H (inset at ×1000 magnification). (c) SEM-EDX elemental maps of C-SO$_3$H. (d) N$_2$ adsorption-desorption isotherms of the untreated carbon and functionalised carbon.

Figure 2. (a) FT-IR spectra of functionalised carbon. (b) XPS survey spectra and deconvolution of the core-level spectra of C1s of the (c) untreated carbon, (d) C-SO$_3$H (with the inset of S2p), (e) C-NH$_2$ (with the inset of N1s), and (f) C1s of C-COOH (with the inset of O1s), respectively.

Table 1. Atomic concentrations of the untreated carbon and functionalised carbon samples.
samples compared to the untreated samples due to the oxygen content of the \(-\text{COOH}\) group. Atomic concentration of 0.5% is observed for S in the C-SO$_3$H samples, confirming the presence of the \(-\text{SO}_3\text{H}\) group on the carbon surface. Similarly, the amine-functionalised carbon, C-NH$_2$, has an atomic concentration of 3.51% for N in the \(-\text{NH}_2\) group. Details of the functionalised carbon surface were determined from the C1s high-resolution XPS spectra (Fig. 2c–f). In the untreated carbon samples, the peaks corresponding to C are observed at 284.5, 284.9, and 286.8 eV, attributed specifically to $\text{sp}^2$ C (C=C), $\text{sp}^3$ C (C–C), and C–O (hydroxyl), respectively. Functionalisation leads to the appearance of new peaks. These include a peak at 289 eV corresponding to the \(-\text{COO}\) groups in C-COOH, 290 eV correlating to the C–S bond in C-SO$_3$H (confirmed by the S2p core level spectrum), and 399 eV attributable to the N1s peak for C–N bonding in C-NH$_2$. The C-COOH samples exhibit an increase in the carboxylic acid peak intensity, including O1s of O=C and O–C peaks at 533.3 and 531.9 eV. Similar XPS results have been previously reported in literatures$^{29–33}$.

Functionalised carbon slurries with concentrations ranging from 1 to 7 phr were added to the NRL. Curing agents were used to produce the carbon foam, which was subsequently steam vulcanised to yield a final bio-renewable polysisoprene-functionalised carbon composite foam with an open-cell structure. Typically, a rubber composite with a high carbon content (up to 70 phr) is produced using a conventional two-roll mill$^{34}$. However, NRL is a colloidal dispersion of cis-1,4-polyisoprene in an aqueous medium, and the amount of carbon that can be added to the latex is limited due to issues related to miscibility and phase separation of the carbon powder. Nanogrinding methods can be used to reduce the particle size in order to increase the carbon content (Fig. 3a,b). The average particle size of carbon after nanogrinding is 4–31 µm, which is smaller than that of activated carbon (3 to 229 µm). The zeta potential as a function of pH of the functionalised carbon in water suspensions indicates that the points of zero charge (pHpzc) of C-NH$_2$ and C-COOH are observed at a pH of 2 and 3, respectively, while C-SO$_3$H is negatively charged in the pH range of 1–6 (Fig. 3c). These observations are attributed to the effects of the surface functional groups on the suspension behaviour and heavy metal adsorption.

The colour of the carbon foam before and after the addition of carbon slurry is shown in Fig. 3d, which shows that the foam colour darkens with increasing carbon content. The SEM images reveal the open pore structure of the foam, with cell interconnection and fine particles distributed in the open cell wall (Fig. 3e). The open-cell wall of the unground carbon (Fig. 3f) shows agglomeration and non-uniform distribution, while the nanoground carbon slurry (Fig. 3g) has a better distribution in the rubber matrix (up to 7 phr). A pore diameter of below 200 µm is observed for the open pore structure, and the dispersion of carbon across the surface can also affect adsorption.

**Effect of contact time, pH, and initial concentration on Cu(II) adsorption.** All carbon foams were mixed with Cu(II) solutions with various pH values and initial concentrations and monitored over the full duration of the contact time to reveal that all samples reached their equilibrium adsorption within 60 min (Fig. 4). The quantitative measurement conducted via UV–Vis analysis depends on the amount of absorbed light, which varies with metal solution concentrations for bulk analysis. From UV–Vis analysis, the blank foam specimen (polysisoprene foam without functionalised carbon) showed no change in Cu(II) solution colour for all adsorption conditions (green line in Fig. 4), the calculation of the adsorption capacity ($q$) is based on the adsorption amount of Cu(II) ion per gram of carbon foam (mg $\text{Cu}^{2+}$/foam).

The used-blank and used-carbon foams were characterised by XRF and SEM-EDX to observe the amount of Cu deposition at the surface and middle of the cubic foam specimen. From Fig. 5 and Table 2, the elemental
mappings of XRF, SEM-EDX, and SRXTM show a similar trend of Cu deposition, wherein the deposition at outer surface was higher than that at the centre of the foam. Moreover, the functionalised carbon foam exhibited higher Cu amount than the blank foam, especially at the outer surface (approximately 6.9 times). This can be attributed to the efficiency of metal adsorption of the functionalised carbon foam.

For bulk analysis, the adsorption depends on the diffusion of aqueous heavy metal ions into the open cell structure of the foam and the pores of the functionalised carbon. These ions attach to the active sites within these structures, and this mechanism requires ion-exchange and chelation. Ion-exchange is driven by electrostatic attraction between the anions of the functional groups and the cations of the heavy metal, whereas chelation is a result of the binding between the lone pairs of electrons of the functional groups and cations of the heavy metal.

The C-SO$_3$H and C-COOH carbon foams adsorb Cu(II) via ion-exchange in the pH range of 1–5, with a maximum adsorption of 57 mg g$^{-1}$ foam and 56 mg g$^{-1}$ foam, respectively, for the 1500 ppm Cu(II) solution. C-NH$_2$ carbon foam adsorbs Cu(II) in a pH range of 2–5, which is mainly driven by chelation between the lone electron pairs of the –NH$_2$ group and copper ions. A maximum adsorption of C-NH$_2$ carbon foam was 42 mg g$^{-1}$ foam for the 1500 ppm Cu(II) solution. However, under highly acidic conditions of pH 1, the –NH$_2$ moieties are protonated to form ammonia ions (–NH$_3^+$) and are unable to adsorb the heavy metal cations. The results are consistent with the zeta potential calculations (Fig. 3c). The mechanism for heavy metal ion adsorption by the foams is shown in Fig. 6. Adsorption is not studied at pH > 5 due to metal hydroxide precipitation, which requires additional separation techniques for heavy metal removal.

According to literature review, adsorption capacity is mostly examined in batch experiments at room temperature (approx. 30 °C). Some literatures reported the temperature effect (4–90 °C) on the metal ion adsorption where the temperature has a slight effect on the adsorption of Cu(II) ions.$^{6,36,37}$ Therefore, the adsorption temperature was kept constant at 30 °C in our study.

Figure 4. Effects of contact time, pH, and initial concentration on the functionalised carbon foam adsorption of Cu(II): (a$_1$–a$_3$) C-SO$_3$H; (b$_1$–b$_3$) C-COOH; (c$_1$–c$_3$) C-NH$_2$, and green line (—) for the blank foam (polyisoprene foam without functionalised carbon).
Adsorption kinetics of Cu(II). The kinetic rate constant ($k$) and the correlation coefficients ($R^2$) obtained from the models indicate that a pseudo-second order kinetic model best fits the adsorption mechanism (Table 3). This suggests that the adsorption mechanism involves adsorbate-adsorbent interaction and a rate-limiting step. In most cases, rate limiting is due to the chemical reactions and diffusion mechanisms. The chemical reaction involves the sharing of valence forces or exchange of lone-pair electrons from the functional groups to the metal ions, while the diffusion mechanisms include film diffusion and intraparticle diffusion. The metal ions initially diffuse from the aqueous media to the external foam surface, from which these are transported through the carbon foam network and retained in the pore structure, where ion adsorption occurs at the active sites of the functionalised carbon.

The selective adsorption of mixed metal ions is a useful information for wastewater treatment application; however, the results could not observe by using a simple UV-spectrophotometer due to the detection limit of the instrument. From literatures, the adsorption isotherm of mixed metal ion solution have been reported. For example, the adsorption isotherm of mixed metal ions solutions (Ag(I), Cu(II), Fe(II), and Pb(II)) by an aminated polyacrylonitrile nanofiber mat using ICP technique was fitted with the Langmuir adsorption isotherm. The Langmuir constants ($K_L$) were in an order of Fe(II) > Cu(II) > Pb(II) > Ag(I) which related to the affinity of binding sites of each metal ions, the low $K_L$ values indicate low adsorption rate of metal ion. The adsorption of Cu(II)
ions in a mixture of four ion metals was achieved at 30 mg g\(^{-1}\). The selective removal capability of Cu(II) and Fe(II) by chelating resin (Rexp-501) using AAS technique indicated that the adsorption of Fe(II) was depended on the resin dosage and was not influenced by the Cu(II) concentration, as the adsorption of Cu(II) was limited to a very low level\(^{42}\). Therefore, the high sample throughput analytical process by using AAS or ICP is necessary for the selective adsorption of mixed metal ions study.

**Reusability of the composites.** The removal efficiencies of all the functionalised carbon foams decrease with increasing reuse cycles (Fig. 7). Desorption requires a sufficient amount of H\(^+\) from the desorbing agent to cover the adsorbent surface and replace the metal ions\(^{43}\); decreased removal in the third and fourth cycles is attributed to an increase in the protonated sites on the adsorbent surface\(^{38}\). The protonated groups exhibit low levels of complexation with metal ions and ion-exchange, which reduce the adsorption capacity\(^{44}\). These adsorbents are reusable, but a decreased efficiency is observed after the several cycles. The functional groups on carbon plays an important role in Cu(II) adsorption where C-SO\(_3\)H and C-COOH carbon foams adsorb Cu(II) via ion-exchange and C-NH\(_2\) carbon foams adsorb Cu(II) via chelation between the lone electron pairs, a weaker interaction. In this work, the reusability of the functionalised carbon foams was studied via repeated adsorption/desorption cycles by using 0.1 M HCl as the desorbing solution and 0.1 M NaOH as the regeneration solution. An acid desorption agent is widely used as an acidic desorption agent for desorption of heavy metal ions, however, some researchers have found that acid eluents also reduce the metal ion adsorption capabilities due to the saturation and occupation of adsorption sites with strongly adsorbed adsorbates which affect to functional groups in carbon foam by covering the active sites and reduced adsorption capacity\(^{45,46}\). Severely attenuated removal efficiencies of C-SO\(_3\)H and C-COOH probably due to the deactivation of sulfonated carbon and carboxylated carbon via proton leaching induced by ion exchange process whilst the lesser types of interactions like physical force and complexation of C-NH\(_2\) might take slighty effect on the removal performance\(^{47}\).

**Conclusions**

Three types of functionalised carbons (C-SO\(_3\)H, C-COOH, C-NH\(_2\)) were successfully incorporated into a rubber foam matrix with good distribution. UV–Vis analysis was found efficient for bulk adsorption, and the amount of Cu on the foam surface could be characterised by XRF and SEM-EDX. Sulfonyl group (−SO\(_3\)H) foam exhibited the maximum Cu(II) adsorption of 1672 mg g\(^{-1}\)\(_{\text{active carbon}}\), whereas the carbonyl group (−COOH) and amine group (−NH\(_2\)) foams showed Cu(II) adsorptions of 1705 mg g\(^{-1}\)\(_{\text{active carbon}}\) and 1293 mg g\(^{-1}\)\(_{\text{active carbon}}\), respectively. Cu(II) adsorption was optimal in the pH range of 2–5 for all functionalised carbon foams. The adsorption mechanism revealed a pseudo-second order kinetics model. The functionalised carbon foams could be reused for up to four cycles, although a gradual decrease in removal efficiency was observed. This approach offers a bio-renewable polymer material capable of adsorbing heavy metal ions from the aqueous media and shows potential for effective wastewater treatment. The selective metal ion adsorption for mixed metal ion with proper measurements, such as atomic adsorption spectroscopy (AAS), and inductively coupled plasma spectrometer (ICP), would be beneficial to further study about the selective adsorption investigation of mixed metal ions, especially for wastewater treatment application.

**Materials and methods**

**Materials.** Activated carbon produced from coconut shell waste (surface area = 768 m\(^2\) g\(^{-1}\)) was sourced from Carbokarn Co., Ltd. (Bangkok, Thailand). Acrylic acid (99%) and ammonium hydroxide were purchased from Sigma–Aldrich (St. Louis, Missouri, USA). Sulfuric acid (conc.), copper (II) acetate, and sodium hydroxide were obtained from Ajax Finechem Pty, Ltd. (Taren Point, New South Wales, Australia). Triton X-100 (Acros Organ-
ics, Geel, Belgium) was used as a surfactant. Hydrochloric acid (37%) was purchased from QRëC (Rawang, Selangor, Malaysia).

**Preparation of functionalised carbon and composite foam.** Functionalisation was conducted in a Teflon-lined acid digestion vessel. Sulfonation was achieved by a hydrothermally treating a mixture of 1.0 g of carbon with 20 mL conc. H₂SO₄ at 170 °C for 12 h. The mixture was cooled, filtered, washed with DI water until neutralised, and further washed with ethanol. The functionalised carbon material (C-SO₃H) was dried at 90 °C

| pH | Initial concentration (ppm) | qₑₒₓpₓ (mg g⁻¹ foam) | Pseudo-first-order kinetic | Pseudo-second-order kinetic |
|----|-----------------------------|------------------------|----------------------------|-----------------------------|
|    |                             |                        | k₁ (min⁻¹) qₑₒₓ (mg g⁻¹ foam) | R₁² | k₂ (g mg⁻¹ min⁻¹) qₑₒₓ (mg g⁻¹ foam) | R₂² |
| 1  | 500                         | 30.71                  | 0.035 8.64                  | 0.9824 0.981 | 0.031 29.59 | 0.9987 |
|    | 1000                        | 41.99                  | 0.026 6.97                  | 0.8860 0.023 | 0.018 40.16 | 0.9998 |
|    | 1500                        | 56.54                  | 0.042 5.86                  | 0.8840 0.018 | 0.018 56.18 | 0.9999 |
| 2  | 500                         | 27.97                  | 0.038 13.57                 | 0.9846 0.035 | 27.03 0.9949 |
|    | 1000                        | 33.01                  | 0.061 9.94                  | 0.9495 0.031 | 33.33 0.9984 |
|    | 1500                        | 42.97                  | 0.033 11.97                 | 0.9140 0.022 | 40.98 0.9960 |
| 3  | 500                         | 21.19                  | 0.012 6.80                  | 0.9149 0.039 | 17.30 0.9988 |
|    | 1000                        | 24.95                  | 0.036 14.95                 | 0.9604 0.039 | 24.15 0.9861 |
|    | 1500                        | 28.67                  | 0.041 13.25                 | 0.9853 0.034 | 27.78 0.9987 |
| 4  | 500                         | 27.21                  | 0.028 4.42                  | 0.9258 0.035 | 26.11 0.9993 |
|    | 1000                        | 30.71                  | 0.027 6.27                  | 0.9803 0.031 | 29.15 0.9994 |
|    | 1500                        | 39.86                  | 0.063 6.01                  | 0.9770 0.025 | 40.49 0.9997 |
| 5  | 500                         | 26.09                  | 0.048 2.57                  | 0.9277 0.038 | 25.97 0.9997 |
|    | 1000                        | 30.16                  | 0.037 1.30                  | 0.9185 0.033 | 30.03 1.0000 |
|    | 1500                        | 38.95                  | 0.064 5.22                  | 0.9669 0.026 | 39.06 0.9998 |

| C-SO₃H carbon foam |  | Pseudo-first-order kinetic | Pseudo-second-order kinetic |
|--------------------|---|-----------------------------|----------------------------|
|                    |   |                            |                            |
|                    |   |                            |                            |

**Table 3.** Kinetic parameters of pseudo-first order and pseudo-second order kinetic models for Cu(II) adsorption onto functionalised carbon foam.
for 4 h and stored in a desiccator. The same hydrothermal protocol was used for the production of carboxyl-functionalised carbon, C-COOH (1.0 g of carbon with 2 mL of acrylic acid), and amine-functionalised carbon, C-NH₂ (1.0 g of carbon mixed with 4 mL of conc. ammonium hydroxide). The methods reported in a previous study by the current authors were used for the preparation of a foam from natural rubber latex (NRL, Hevea Brasiliensis) and for the determination of its density and porosity. A carbon slurry was produced using a high energy ball mill (Emax, Retsch GmbH, Haan, Germany) containing tungsten carbide balls (5 mm in diameter; 180 balls per jar), whereby the carbon powder (8 g) was ground with Triton-X (1.4 g dissolved in 18.6 mL of deionized water) at 800 rpm for 30 min. The carbon slurry was added to high ammonia rubber latex at concentrations of 1–7 phr (parts per hundred of rubber). The density of the foam was controlled at 0.29 ± 0.1 g cm⁻³ and the porosity at 64 ± 4%. A static compression set test was performed following ASTM D395 (Method B, Type 2 at 70 °C for 24 h) to verify that the flexibility and elasticity of the foam fell within the range of 65–75%.

Characterisation of functionalised carbon and composite foam. The surface area of the carbon powder was characterised using N₂ isotherm (V-Sorb 2800P, Gold APP Instruments Corp., Beijing, China) through Brunauer–Emmett–Teller (BET) analysis. The morphologies of the functionalised carbon and composite foam were investigated using scanning electron microscopy (SEM, JSM-6510LV, JEOL Ltd., Tokyo, Japan) at an acceleration voltage of 20 kV. Elemental mapping of the used-functionalised carbon and used-composite foam were analysed by scanning electron microscope (SEM, FEI, Quanta 450) and energy dispersive X-ray spectrometer (EDX, Oxford, XMax) and XRF microscopy (Horiba, XGT-7200). X-ray photoelectron spectroscopy (XPS) data were recorded using an ULVAC-PHI, PHI 500 Versa Probe II (ULVAC-PHI Inc., Kanagawa, Japan) with Al Kα X-ray radiation as the excitation source. The atomic concentrations of the functionalised carbon were analysed from the C1s, O1s, N1s, and S2p spectra. The instrument sensitivity was 0.01–0.5% (atomic percentage). Tomographic volume of the samples were investigated by synchrotron radiation X-ray tomographic microscopy (SRXTM). The sample projections (10 keV, at a distance of 34 m from the source) were obtained from the detection system, which was equipped with 200-μm-thick YAG:Ce scintillator (Crytur, Czech Republic), the white-beam microscope (Optique Peter, France) and the pco.edge 5.5 sCMOS camera (2560 × 2160 pixels, 16 bits). After collecting data, it was normalised using flat-field correction algorithm and reconstructed by Octopus reconstruction, respectively. To provide the 3D representation of tomographic volume of the sample, the reconstructed images were rendered using Drishti software.

Cu(II) adsorption. Heavy metal removal was performed in a batch system in which the adsorbent was added to the freshly prepared Cu(II) solutions with varying concentrations and pH and monitored for the duration of the contact time. Standard solutions of Cu(II) in a range of concentrations (500–1500 ppm) were prepared by dissolving copper acetate (Cu(CH₃COO)₂·H₂O) in deionised water. The pH values of each Cu(II) solution were adjusted in the pH range of 1–5 using 0.1 M HCl or 0.1 M NaOH. The adsorption was performed at 150 rpm on an orbital shaker, where the adsorbent was added to the conical tubes containing 25 mL of freshly prepared Cu(II) solutions for a total contact time of 0–250 min at a constant temperature of 303 K. Ion adsorption was characterised using UV–Visible spectrophotometer (Shimadzu UV 1700, Shimadzu Corp., Kyoto, Japan). The adsorption capacities (q) were calculated according to Eq. (1).

\[ q = \frac{(C_i - C_f) \times V}{m}, \]  (1)

where \( q \) is the adsorption amount (mg g⁻¹), \( C_i \) and \( C_f \) are the initial and final concentrations (mg L⁻¹), respectively, \( m \) is the weight of active carbon (g), and \( V \) is the volume of the solution (L).

For the adsorption kinetics, the pseudo-first-order kinetic and pseudo-second-order kinetic equations are expressed as follows:
\[
\log(q_e - q_t) = \log q_e - \frac{k_1 t}{2.303}
\]

where \( q_e \) and \( q_t \) are the amounts of adsorbed copper per weight of carbon foam (mg \( g^{-1} \) foam) at time \( t \) (min) and at equilibrium, respectively. Rate constant of pseudo-first-order adsorption is \( k_1 \) (min \(^{-1}\)) and pseudo-second-order adsorption is \( k_2 \) (g \( mg \) \(^{-1} \) min \(^{-1}\)).

The reusability of the functionalised carbon foams was studied via repeated adsorption/desorption cycles. The spent adsorbents can be recovered by acid washing adsorbents by 0.1 M HCl as the desorbing solution and 0.1 M NaOH as the regeneration solution. The regenerated carbon foams were washed with distilled water up to neutral pH and then reused in a new adsorption/desorption cycle. Four consecutive adsorption/desorption cycles were performed with three replications (n = 3).

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References
1. Abas, S. N. A., Ismail, M. H. S., Kamal, M. I. & Izhar, S. Adsorption process of heavy metals by low-cost adsorbent: A review. World Appl. Sci. J. 28, 1518–1530 (2013).
2. Barati, A., Asgari, M., Mirt, T. & Eskandari, Z. Removal and recovery of copper and nickel ions from aqueous solution by poly (methacrylamide-co-acrylic acid)/montmorillonite nanocomposites. Environ. Sci. Pollut. Res. 20, 6242–6255 (2013).
3. Honda, R. & Nogawa, K. Cadmium, zinc and copper relationships in kidney and liver of humans exposed to environmental cadmium. Arch. Toxicol. 59, 437–442 (1987).
4. Demiral, H., Demiral, I., Tümsek, F. & Karabacakoğlu, B. Adsorption of chromium (VI) from aqueous solution by activated carbon derived from olive bagasse and applicability of different adsorption models. Chem. Eng. J. 144, 188–196 (2008).
5. Fu, F. & Wang, Q. Removal of heavy metal ions from wastewaters: A review. J. Environ. Manag. 92, 407–418 (2011).
6. Kara, A. & Demirbel, E. Physicochemical parameters of Cu(II) ions adsorption from aqueous solution by magnetic-poly(divinylbenzene-v-vinylimidazole) microbeads. Sep. Sci. Tech. 47, 799–822 (2012).
7. Kitano, M. Adsorption of copper from aqueous solution using chitosan - montmorillonite nanocomposites. J. Hazard. Mater. 159, 290–303 (2009).
8. Ozdemir, I., Kara, A., Tékin, N. & Oğun, A. Synthesis and characterization of polymer microspheres and its application for phenol adsorption. Desalin. Water Treat. 58, 612–623 (2015).
9. Shadrashov, M. & Bakhtiari, S. The efficiency of activated carbon/iron oxide nanoparticles composites for Cu(II) adsorption. Kinetic, isothermal and thermodynamic studies. J. Hazard. Mater. 286, 612–623 (2015).
10. Abas, S. N. A., Ismail, M. H. S., Kamal, M. L. & Izhar, S. Adsorption process of heavy metals by low-cost adsorbent: A review. World Appl. Sci. J. 28, 1518–1530 (2013).
11. Zhang, W. et al. One-pot synthesis of carbonaceous monolith with surface sulfonic groups and its carbonization/activation. Carbon 49, 1811–1820 (2011).
12. Kitano, M. et al. Preparation of a sulfonated porous carbon catalyst with high specific surface area. Catal. Lett. 131, 242–249 (2009).
13. Huang, M.-R., Lu, H.-J. & Li, X.-G. Synthesis and strong heavy-metal ion sorption of copolymer microparticles from phenylenediamine and its sulfonate: J. Mater. Chem. 22, 17685–17699 (2012).
14. Shadrashov, M. & Bakhtiari, S. The efficiency of activated carbon/iron oxide nanoparticles composites for Cu(II) adsorption. Desalin. Water Treat. 58, 612–623 (2015).
15. Kumar, R., Gupta, A. & Dhakate, S. K. Nanoparticles-decorated coal tar pitch-based carbon foam with enhanced electromagnetic radiation absorption capability. RSC Adv. 5, 20256–20264 (2015).
16. Agarwal, P. R., Kumar, R., Kumari, S. & Dhakate, S. R. Three-dimensional and highly ordered porous carbon-MnO\(_2\) composite foam for excellent electromagnetic interference shielding efficiency. RSC Adv. 6, 100713–100722 (2016).
17. Rethinabapathy, M., Kang, S.-M., Jang, S.-C. & Huh, Y. S. Three-dimensional porous graphene materials for environmental applications. Carbon Lett. 22, 1–13 (2017).
18. Lee, C. G. et al. Removal of copper, nickel and chromium mixtures from metal plating wastewater by adsorption with modified carbon foam. Carbohydr. Polym. 166, 203–211 (2017).
19. Gama, E. M., Lima, A. D. S. & Lemos, V. A. Preconcentration system for cadmium and lead determination in environmental samples using polyurethane foam/Mg-BTANC. J. Hazard. Mater. 136, 757–762 (2006).
20. Shahraashou, M. & Bakhthiar, S. The efficiency of activated carbon/ion oxide nanoparticles composites in copper removal: Industrial waste recovery, green synthesis, characterization, and adsorption—Desorption studies. Micropropor Mesoporous Mater. 13, 110692 (2020).
21. Hamid, S. A., Azha, S. F., Sellauoi, L., Bonilla-Petriciolet, A. & Ismail, S. Adsorption of copper (II) cation on polysulfone/zolite blend sheet membrane: Synthesis, characterization, and adsorption modelling. Colloids Surf. A Physicochem. Eng. Asp. 601, 124980 (2020).
22. Bashir, M., Tyagi, S. & Annachhatre, A. P. Adsorption of copper from aqueous solution onto agricultural adsorbents: Kinetics and isotherm studies. Mater. Today Proc. 28(3), 1833–1840 (2020).
23. Sahebjamee, N., Soltanieh, M., Mahmood, S. M. & Amir Heydarinasab, A. Preparation and characterization of porous chitosan-based membrane with enhanced copper ion adsorption performance. React. Funct. Polym. 154, 104681 (2020).
24. Deng, L., Li, Y., Zhang, A. & Zhang, H. Nano-hydroxyapatite incorporated gelatin/zein nanofibrilousmembranes: Fabrication, characterization and copper adsorption. Int. J. Biol. Macromol. 154, 1478–1489 (2020).
25. Zia, Q. et al. Porous poly(L-lactic acid)/chitosan nanofibres for copper ion adsorption. Carbohydr. Polym. 227, 115343 (2020).
26. Prasad, M. N. V. & Freitas, H. Removal of toxic metals from solution by leaf, stem and root phytomass of Quercus ilex L. (holly oak). Environ. Pollut. 110, 277–283 (2000).
27. Brady, D., Stoll, A. & Duncan, F. R. Biosorption of heavy metal cations by non-viable yeast biomass. Environ. Technol. 15, 419–428 (1994).
28. Hao, S.-W., Hsu, C.-H., Liu, Y.-G. & Chang, B.-K. Adsorption of copper (II) cation on produced carbonaceous monolith with surface sulfonic groups and its carbonization/activation. Carbon 49, 1811–1820 (2011).
29. Zia, Q. et al. Porous poly(L-lactic acid)/chitosan nanofibres for copper ion adsorption. Carbohydr. Polym. 227, 115343 (2020).
30. Prasad, M. N. V. & Freitas, H. Removal of toxic metals from solution by leaf, stem and root phytomass of Quercus ilex L. (holly oak). Environ. Pollut. 110, 277–283 (2000).
31. Brady, D., Stoll, A. & Duncan, F. R. Biosorption of heavy metal cations by non-viable yeast biomass. Environ. Technol. 15, 419–428 (1994).
30. Adams, L., Oki, A., Grady, T., McWhinney, H. & Luo, Z. Preparation and characterization of sulfonic acid-functionalized single-walled carbon nanotubes. *Physica E* **41**, 723–728 (2009).
31. Sun, Y.-F. et al. Sensitive and selective electrochemical detection of heavy metal ions using amino-functionalized carbon microspheres. *J. Electroanal. Chem.* **760**, 143–150 (2016).
32. Hu, Z. et al. One-pot preparation and continuous spinning of carbon nanotube/poly-(p-phenylene benzobisoxazole) copolymer fibers. *J. Mater. Chem. B* **2**, 19863–19871 (2012).
33. Soliman, A. M., Elsucary, S. A. A., Ali, I. M. & Ayesh, A. I. Photocatalytic activity of transition metal ions-loaded activated carbon: Degradation of crystal violet dye under solar radiation. *J. Water Process Eng.* **17**, 245–255 (2017).
34. Lee, E.-K. & Choi, S.-Y. Preparation and characterization of natural rubber foams: Effects of foaming temperature and carbon black content. *Korean J. Chem. Eng.* **24**, 1070–1073 (2007).
35. Gupta, V. K., Pathania, D. & Sharma, S. Adsorptive remediation of Cu (II) and Ni (II) by microwave assisted H3PO4 activated carbon. *Arabian J. Chem.* **10**, S2836–S2844 (2017).
36. Kubicay, S., Gurkan, R., Savran, A. & Sahab, T. Removal of Cu(II), Zn(II) and Co(II) ions from aqueous solutions by adsorption onto natural bentonite. *Adsorption* **13**, 41–51 (2007).
37. Hu, X., Wang, H. & Liu, Y. Statistical analysis of main and interaction effects on Cu(II) and Cr(VI) decontamination by nitrogen-doped magnetic graphene oxide. *Sci. Rep.* **6**, 34378 (2016).
38. Orozco-Guareño, E. et al. Removal of Cu (II) ions from aqueous streams using poly(acrylic acid-co-acrylamide) hydrogels. *J. Colloid Interf. Sci.* **349**, 583–593 (2010).
39. Diao, M., Li, Q., Xiao, H., Duan, N. & Xu, J. Synthesis and adsorption properties of superabsorbent hydrogel and peanut hull composite. *J. Environ. Chem. Eng.* **2**, 1558–1567 (2014).
40. Ahmad, R., Rao, R. A. K. & Masood, M. M. Removal and recovery of Cr (VI) from synthetic and industrial wastewater using bark of Pinus roxburghii as an adsorbent. *Water Qual. Res. J.* **40**, 462–468 (2005).
41. Kampalanonwat, P., Supaphol, P. The study of competitive adsorption of heavy metal ions from aqueous solution by aminated polyacrylonitrile nanofiber mats. *Energy Proc.* **56**, 142–151 (2014).
42. Li, Y., Wang, X., Xiao, Q., Zhang, X. Study on selective removal of impurity iron from leached copper-bearing solution using a chelating resin. *Minerals* **6**, 106 (2016).
43. Gupta, V. K., Pathania, D., Sharma, S., Agarwal, S. & Singh, P. Remediation of noxious chromium (VI) utilizing acrylic acid grafted lignocellulosic adsorbent. *J. Mol. Liq.* **177**, 343–352 (2013).
44. Wang, W.-B., Huang, D.-J., Kang, Y.-R. & Wang, A.-Q. One-step in situ fabrication of a granular semi-IPN hydrogel based on chitosan and gelatin for fast and efficient adsorption of Cu2+ ion. *Colloids Surf. B Biointerf.* **106**, 51–59 (2013).
45. Gedam, A. H. & Dongre, R. S. Adsorption characterization of Pb (II) ions onto iodate doped chitosan composite: equilibrium and kinetic studies. *RSC Adv.* **5**, 54188–54201 (2015).
46. Scalbert, P. et al. Simultaneous investigation of the structure and surface of a Co/alumina catalyst during Fischer-Tropsch synthesis: Discrimination of various phenomena with beneficial or disadvantageous impact on activity. *Catal. Sci. Technol.* **5**(8), 4193–4201 (2015).
47. Scholz, D., Kröcher, O. & Vogel, F. Deactivation and regeneration of sulfonated carbon catalysts in hydrothermal reaction environments. *Chem. Sus. Chem.* **11**(13), 2189–2201 (2018).

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**Author contributions**  
All authors contributed to the preparation of the manuscript and approved the final version. The following is a list of individual contributions. W.K. conceived the experiments and prepared composites foam. C.S. co-supervision and provided laboratory equipment. N.C. and P.P. characterised XPS, SEM-EDX, and SRXTM. P.R. and G.G. provided resources for characterisation. S.K. (corresponding author) supervised the concept and research methodology, wrote manuscript, prepared figures, responded to the reviewer, revised and finalised the manuscript. S.K. provided research funding, co-supervision, reviewing and scientific discussion.

**Competing interests**  
The authors declare no competing interests.

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