Synthesis of Multi Wall Carbon Nanotubes based on zirconium oxide as supported material and its application as nanosorbent for copper ions removal

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GRAPHICAL ABSTRACT



Abstract

Water pollution caused by various toxic contaminants has become one of the world's most serious issues. Nanotechnology has received a lot of attention in recent decades, and various nanomaterials for water remediation have been developed. For this purpose, MWCNTs were directly synthesized using a chemical vapor deposition process, which was then purified and functionalized using an acid treatment. At various temperatures, the synthesis of MWCNTs by chemical vapor deposition (CVD) of acetylene is investigated. The wet impregnation method is used to prepare catalysts using Fe/Co as a catalyst in the presence of zirconium oxide and aluminum oxide as support

materials. Zirconium oxide was found to be a good support material for the deposition of highpurity MWCNTs with high yield 88.9 %. The effect of growing time and temperature on carbon yield was investigated, and it was discovered that the amount of MWCNTs deposited increased as reaction time increased, with the optimum temperature for the reaction being 700 ^oC. The effect of temperature on the crystal size of synthesized MWCNTs was investigated using XRD, and it was found that the crystallite size declined as the temperature goes up. SEM and TEM were used to examine the structure and purity of synthesized MWCNTs at various temperatures. The MWCNTs that were synthesized have a web-like network structure, a homogeneous distribution on the catalyst surface, and a smooth surface. The prepared CNTs were purified by chemical oxidation, and the effect of acid treatment on the surface of the CNTs was investigated using XRD and FTIR. The functionalized CNTs were used for copper ion adsorption, achieving an adsorption potential of 238.09 mg/g and a high regeneration efficiency.

Keywords: Aluminum oxide, Carbon nanotube, Chemical vapor deposition, Copper ions, Zirconium oxide

1. Introduction

Carbon nanotubes (CNTs) play a role in a number of scientific and technical fields (Burakova et al. 2018). Nanoelectronics, nanosensors, photovoltaics, and chemical sensing are examples of these, since they have special electrical and thermal conductivity properties. Because of their high surface area and porous nature, CNTs have the potential to play a role in water purification (Anzar et al.2020).

CNTs are made up of graphene sheets that have been rolled up to form hollow cylindrical tubes (Arora and Attri 2020). There are three types of carbon nanotubes (CNTs). Single-walled carbon nanotubes (SWCNTs) have a single-cylinder with a standard diameter of 1.4 nm, while double-walled carbon nanotubes (DWCNTs) and multi-walled carbon nanotubes (MWCNTs) have concentric cylinders with an inter-tube spacing of 0.34 nm and a typical diameter ranging from 2 nm to 30 nm, according to the rolling up of graphene sheets. The lengths of SWCNTs and MWCNTs vary from a few micrometres to several centimetres. CNTs have a strength of thirty to one hundred times that of steel, although their density is around one-sixth that of steel. CNTs have an elastic modulus of around 1.2 TPa, which is identical to the diamond's elastic modulus. Otherwise, SWCNTs have a density of about 1.33–1.40 g/cm³, which is about half that of

aluminium, and their thermal conductivity (6000 W/m K at room temperature) is roughly double that of diamond. Furthermore, their electrical current has a larger magnitude than copper (Hussein et al. 2014 & Patel et al. 2021).

For the development of CNTs, various synthetic methods have been established, such as laser vaporisation, which operates at room temperature but yields low yields and is otherwise a high-cost technique. The second approach for producing CNTs is arc discharge, which produces high-quality and low-cost CNTs but necessitates a high-temperature environment (Dhorea et al. 2018). Although both arc discharge and laser ablation methods produce CNTs, purification is required (Sari et al. 2018). The chemical vapor method (CVD) is the most suitable method for producing CNTs with high yield, high purity, and regulated growth. It is a simple technique that produces CNTs of high quality and quantity, with a realistic yield of (>75 percent) (Yang et al. 2019& Tchudan et al. 2019). The most important catalysts for the CVD system of CNT growth are cobalt (Co), iron (Fe), and nickel (Ni) (Wanga et al. 2019).

All of the parameters and conditions of CNT fabrication, including the supporting materials that contribute to the catalyst for CNT development, are still being investigated. During the CVD process, the presence of support materials has a significant effect on the high quality of the grown CNTs. Peigney et al. (1997), for example, used Fe-alumina nanocomposite as a catalyst to produce single and multiwall carbon nanotubes (Peigney et al. 1997). The use of porous MgO as a support material greatly improved the yield of SWCNTs grown using silica/alumina supported material with Fe, Co, or Ni oxide as catalysts, according to the researchers (Cassell et al. 1999& Qingwen et al. 2002).

Heavy metal contamination in water, such as cadmium, zinc, and copper, is a serious concern to human health and the environment. As a result, many attempts are made to reduce water contamination. Toxin removal strategies include chemical precipitation, ion exchange, and electrolytic reduction, reverse osmosis, advanced oxidation, membrane technology, and biological treatments (Mwafya and Mostafa 2020). The methods listed are costly and produce a lot of sludge, while the adsorption method is a good option for water treatment because it is both cost-effective and reliable (Mwafya and Mostafa 2020). Adsorbents such as activated carbon, zeolites, clays, and carbon nanotubes are used to eliminate toxic contaminants from water (Ihsanullah et al. 2020). Activated carbon, zeolites, and clays (Ihsanullah et al. 2020).

Carbon nanotubes have been found to be among the most successful materials in the treatment of water and the adsorption of heavy metals such as copper, cadmium, and lead in many published studies. It was also effective in both polar and nonpolar organic molecules (Hanbali et al. 2020). This was due to the various physical and chemical interactions that MWCNT would have such as Covalent bonding, hydrogen bonding (Du et al. 2014), and electrostatic interactions (Chen et al. 2014).

CNT functionalizations are important for increasing the performance and compatibility of CNTs as adsorbents. As compared to non-functionalized CNTs, the acid modified CNTs have a higher adsorption capacity in almost all studies. This may be due to an electrostatic attraction between the divalent heavy metal ions and the negative charge on the CNTs surface after acid treatment (Abbas et al. 2016).

The effects of zirconium oxide as a supporting material in the synthesis of MWCNTs using Fe/Co catalysts by CVD of acetylene gas are described in this paper. The morphological features of the grown MWCNTs were analyzed using TEM, SEM, and XRD to determine the optimum conditions for MWCNT fabrication. Furthermore, the MWCNTs were functionalized and used as a nanosorbent to remove copper ions from aqueous solutions.

2. Materials and methods

2.1 chemicals

The used chemicals were iron (III) nitrate, cobalt nitrate, aluminum oxide, and zirconium oxide were obtained as the analytical grade from Sigma-Aldrich and used without further purification. Nitric acid, sulfuric acid, sodium hydroxide, copper sulfate pentahydrate were purchased from El-Nasr Co. Egypt, and were used without further purification.

2.2 Preparation and characterization of Carbon Nanotube catalysts.

The support materials were impregnated in salt solutions of Fe $(NO_3)_3$ ·9H₂O and Co $(NO_3)_2$ ·6H₂O to create catalysts. Aqueous solutions were made by dissolving a 1:1 molar ratio in distilled water in a limited volume. After stirring until the salts in the solution were homogeneous, zirconium oxide ZrO₂ was added in a molar ratio of Fe: Co: ZrO₂ of 2.5 %, 2.5 %, and 95 %, respectively. The resulting catalyst (Fe/Co/ZrO₂) was dried in a 125°C oven for 12 hours. The dried material

was then ground in a mortar to make a fine powder, which was then calcined in a tube furnace at 600° C under N₂ gas flow.

In the same way, Fe/Co supported on Al_2O_3 was prepared to compare the grown of MWCNT on new supported material zirconium oxide with that grown on aluminum oxide, which is commonly used in CNT preparation.

X-ray Diffractometer (XRD-7000, Schimadzu Corp., Columbia, MD) with Cu-Ka radiation (D 0.154060 nm) produced at 30 kV and 30 mA was used to examine the catalyst samples. Scans were performed at 2°C min⁻¹ for 2 values ranging from 10 to 90 degrees. The morphology of the catalysts was investigated using a scanning electron microscope (SEM) (JEOL, Model JSM 6360LA, Japan). The surface area of both catalysts was measured using the Belsorp-mini-II from Japan.

2.3 Optimization of MWCNT synthesis.

Using the two prepared catalysts, MWCNTs were produced by pyrolysis of acetylene gas in a horizontal tube furnace at different temperatures (700, 800, and 900 °C), different nitrogen flow rates (75, 120 ml. min⁻¹), and different reaction times (40 min, 60 min).

The reactor (consisting of a 40 mm radius 70 cm length of quartz tube heated by an electrical tube furnace with a temperature controller at a rate of 50 °C min⁻¹ to reach the target temperature and an additional 10 minutes to enable the furnace to regulate the flow of nitrogen at (75, and 120 ml min⁻¹). The reactor tube was then filled with acetylene (C₂H₂) at rates of 30 and 50 ml min⁻¹ for 40 and 60 minutes, respectively. A mass flow controller was used to monitor the flow of gases. The samples were then cooled to room temperature under a 40 ml min⁻¹ nitrogen flow and defined as stated below.

The yield of carbon deposits was calculated by measuring the catalysts before and after MWCNT deposition. For accurate measurements, the same weight of catalyst used in MWCNT growth was treated in the tube furnace without acetylene gas.

Equation 1 (Kathyayini et al. 2004 & El-Maghraby et al. 2014) was used to measure the weight gain during the reaction.

Carbon deposit % =
$$\frac{M_3 - (M_1 - M_2)}{M_3} \times 100$$
 (1)

Where M_1 is the initial weight of the catalyst, M_2 is the weight loss of the catalyst at the reaction temperature and M_3 is the weight of the carbon deposit and the catalyst.

Scan electron microscope (SEM) (JEOL, Model JSM 6360LA, Japan) and transmission electron microscopy (TEM) (JEOL-JEM2100) were utilized for displaying the distribution of MWCNTs in the carbon deposit.

2.4. Functionalization of the synthesized MWCNTs

Carboxylation of MWCNTs (c-MWCNTs) was carried out according to (Pham et al. 2011), where MWCNTs were dispersed through ultrasonication in a mixture solution of concentrated sulfuric acid H₂SO₄ and nitric acid HNO₃ with a volume ratio of 3:1, and the carboxylic function groups on the MWCNTs surface were investigated using Fourier transform infrared spectroscopy FTIR and X-Ray diffraction XRD (Schimadzu-7000).

2.5. Removal studies of copper ions by c-MWCNT

Different quantities of c-MWCNTs were applied to aqueous copper ion solutions, and the solution was shaken for the required duration at 200 rpm on a rotary shaker. The effect of pH, contact time, adsorbent, and copper ion concentrations on the copper ion removal process was investigated. The concentrations of copper ions were calculated using a spectrophotometer set to 550 nm (Sarker and Ullaha 2013). Equation 2 (Ajitha et al. 2017) was used to quantify the extracted copper ions at equilibrium qe (mg/g).

$$\mathbf{q}_{e} = \frac{(Co-Ce)}{m} \times V$$

(2)

Where C_0 is the initial copper ion concentrations (mg/L), Ce is equilibrium copper ions Concentrations (mg/L), V is the volume (L) of the adsorbate solution and m represents The adsorbent mass in g/L.

2.6. Adsorption isotherm of Cu ions

The relationship between the adsorbed copper ions per unit mass of c-MWCNT and their equilibrium concentrations was evaluated using Langmuir, Freundlich.

The Langmuir isotherm model describes the formation of a uniform monolayer adsorbate on the outer surface of c-MWCNTs (Langmuir 1916).

The Langmuir isotherm equation can be described in equation (3):

 $\frac{C_e}{q_e} = \frac{1}{k_a Q_m} + \frac{1}{Q_m} C_e$

 C_e is the equilibrium concentration of Cu ions in mg/L, while qe (mg/g) is represents the number of Cu ions adsorbed at equilibrium, Q_m (mg/g) is the maximum capacity of monolayer coverage and k_a (L/mg) is the constant of Langmuir isotherm.

On the otherhand Freundlich isotherm model describes the adsorption on a heterogeneous surface with un-uniform energy (Freundlich 1906). The Freundlich isotherm equation can be described in equation (4):

$$\log q_e = \log k_f + \frac{1}{n} \log c_e \tag{4}$$

 K_f (mg/g) is the Freundlich isotherm constant and n is the adsorption intensity.

2.7. Kinetic parameters for Cu ions adsorption

At an initial concentration of 200 ppm for copper ion and 10 mg of c-MWCNT, the kinetics of adsorption were investigated for contact times of 10 to 90 minutes. During the entire contact time, the samples were agitated at 180 rpm. Following agitation, the samples were centrifuged at 3900 rpm for 20 minutes before being filtered (pore size 0.45 mm) and the remaining ion concentrations measured. Two kinetic models are tested to match experimental results, including the pseudo-first-order equation (5) (Lagergren 1898), the pseudo-second-order equation (6) (Ho cKay 1999).

$$\log(q_e - q_t) = \log q_e - \frac{K_1}{2.303} (t)$$

$$\frac{t}{q_t} = \frac{1}{K_2 q_e^2} + \frac{1}{q_e} (t)$$
(5)
(6)

Where q_e and q_t are the adsorption capacities at equilibrium and time t, respectively (mg/g), and k_1 (min⁻¹) is the pseudo first order adsorption rate constant, and k_2 (g/mg min) is the pseudo second order adsorption rate constant.

2.9. Regeneration test

One-step desorption experiments were performed to assess the possibility of c- MWCNT reusability for future use. HCl was chosen as the eluent agent in this study because it is the most efficient eluent for desorbing loaded ions from adsorbent surfaces (Smiciklas et al., 2006; Gautam et al., 2014). To test the sorbents' reusability, copper ions at an initial concentration of 200 ppm were used.

Following batch adsorption tests, each sample's supernatant was decanted, and the metal-loaded samples were dried at 100 ^oC for 24 hours. A total of 50 mL of 0.01 M HCl (pH 2) was added to

the dried samples, which were then shaken for 24 hours at 180 rpm and centrifuged for 20 minutes at 3900 rpm. The next adsorption cycle began with the used sorbents being dried in an oven at 60 $^{\circ}$ C for 24 hours. The adsorption-desorption tests were carried out three times. Each test was run twice, and the mean was calculated.

3. Results and discussion

3.1 Characterization of the prepared catalysts & carbonaceous materials

The XRD pattern of catalysts Fe/Co/Al₂O₃, and Fe/Co/ZrO₂ are shown in **Figure 1a, b.** The peaks of the ZrO₂ are clearly detected in the range (27-32) 2Θ corresponding to lines (111, and 110), the presence of both tetragonal and monoclinic ZrO₂ phases was suggested. At room temperature, pure ZrO₂ exists as a monoclinic phase (mZrO₂), but as the temperature rises, it transforms into tetragonal zirconia (t-ZrO₂) and cubic zirconia (c-ZrO₂). On the other hand, the addition of iron and cobalt stabilised the teteragonal phase of zirconia and prevented the transition from the tetragonal to the monoclinic phase during calcination, demonstrating the good dispersion of iron and cobalt oxides on the surface of ZrO₂ (Ohtsuka 2003 & Legorreta et al. 2008). The corresponding (511) line defines the peak of Cobalt oxide at 59.6 2Θ Otherwise, the characteristic peaks for Iron oxide at 24.9 and 40.8 2Θ corresponding to lines (012 and 113), and 67.3 2Θ corresponding to line (522) for γ -Al₂O₃.

The scale of both catalysts was confirmed to be in the nanometer range via SEM observation as seen in Figure 2, the average size of Fe/Co/Al₂O₃ around 19 nm \pm 0.84, while the size of Fe/Co/ZrO₂ around 78±0.63 nm.

Figure 3 shows that the surface area of the Fe/Co/ZrO₂ catalyst is greater than that of the Fe/Co/Al₂O₃ catalyst, which agrees with the XRD result because the transformation of zirconia from tetragonal to monoclinic phase during catalyst calcination was accompanied by a change in surface area (Calafat, 1998). Besides that, both catalysts obey IV behavior, with pore diameters ranging from 2 to 50 nm, indicating that they were mesoporous materials (Table 1).

The key factors that influenced the yield of MWCNTs were catalyst-metal support, synthesis period, temperature, carbon source (acetylene) flow, and nitrogen gas flow, according to the regression study. The experiments were carried out at two different acetylene flow rates (25 and 50 mL min⁻¹) and for different reaction times (40 and 60 minutes), pointing to a better conversion of acetylene into MWCNTs at a flow rate of 50 mL min⁻¹ for 60 minutes, as reported in (Table 2). It was observed that the MWCNTs yield was increased with the increase of synthesis duration.

The highest yield of prepared MWCNTs on both catalysts was obtained by increasing the reaction time from 40 to 60 minutes and performing the reaction at 700 °C with a low acetylene flow rate (25 ml min⁻¹). For all of the catalysts used, the yield of MWCNTs decreased as the rate of flow of N₂ increased from 75 to 120 mL min⁻¹. This is due to the fact that as the carrier gas flow rate increases, the contact time between the acetylene and the catalyst decreases. However, when the flow rate of acetylene gas was increased, the change in the amount of N₂ flow did not show significant change in the percentage of carbon deposit.

It has previously been stated that high temperatures are associated with higher yields of carbon nanotubes, and that high temperatures have a significant impact on metal activity and the decomposition of carbon-containing gases. High temperature, on the other hand, can cause agglomeration of catalytic metal particles, which is a negative factor in CNT synthesis (El-Maghraby et al. 2014). The carbon content (MWCNT yield) which obtained from the proposed Fe/Co/ZrO₂ and Fe/Co/Al₂O₃ is described in Table 3. It was also found that the best temperature for growing MWCNTs was 700°C (Table 3). The results in this section of study are compared with that of literature in Table 4.

SEM observation of the synthesized MWCNT at different temperatures in figure 4 reveals that the synthesized MWCNT at 700 °C with a web-like network structure and homogeneous distribution on the catalyst surface, however, as the temperature of the reaction increase from 700 to 900°C the amorphous carbon appeared with the synthesized MWCNT as well as the diameter of the synthesized carbonaceous material increase by elevation of the reaction temperature, these results similar to that obtained by (Mohammed et al. 2017).

TEM observations for the grown MWCNTs on both catalyst at optimum conditions (700°C in Acetylene for 60 min) are in general agreement with the above results figure 5, and 6. The tubes formed are multiwall with a smooth surface, with only a few showing roughness, suggesting weak graphitization. On the outside of the bundle, no amorphous carbon could be found. Bamboo-like structures or carbon nanoparticles were not found in the produced samples. The high quality of the synthesized MWCNTs could be explained by the good interactions between the metal phase and the support. However, the presence of catalyst particles encapsulated with grown MWCNTs suggested that MWCNTs grew from catalyst particles via a tip growth mechanism. The MWCNTs

synthesized on Fe/Co/ZrO₂ are well deformed tubes with a thick network, with an average diameter of 30-40 nm. This is explained by the fact that the high surface area of Fe/Co/ZrO₂ allows for more contact between the catalyst and the carbon source, as well as a decrease in metal nanoparticle migration along the metal support surface, resulting in uniform metal distribution on the supporting material (Monshi et al. 2012).

3.2. Characterization of the Functionalized MWCNT (c-MWCNT) for copper ions adsorption

Figure 7a showed the XRD pattern of oxidized MWCNTs which were synthesized at the optimum conditions (at 700 0 C, acetylene gas flow 50 ml/min, for 60 min, and grown on Fe/CO/ZrO₂. From the pattern, it was found downshifting of graphite peak at 2 θ = 26.5 which corresponding to the (002) reflection into 2 θ = 25 as a result of increasing in sp², C=C layer spacing (Gupta and Saleh 2011). The diffraction peaks at 2 θ of 43 and 56.8 represent the (100), (101) reflections.

Figure 7b exhibits FTIR of the oxidized MWCNT, as observed the wide peak at 3433 cm⁻¹ associated with the O-H stretch of the hydroxyl group while the peak at 1702.24 cm⁻¹ Refer to the stretch of the C-O bond of the carbonyl group (El Nemr et al. 2019, Hole et al. 2021). This assigned to the functionalization of MWCNT with the carboxyl groups. On the other side the peaks at 16433.41, and 1422.55 cm⁻¹ assigned to C=O, and C-C bonds (Yan et al. 2007, Wang et al. 2021) FTIR confirmed the functionalization of MWCNT with (COOH) as the main function groups on the MWCNT surface represented by the carbonyl groups, and the hydroxyl groups that are attached to the same carbon atom.

Adsorption studies

3.3.1. Effect of pH and contact time on copper ions adsorption

The adsorption of the copper ion at various pH (3-7) was studied by using 10 mg of c-MWCNTs. As shown in figure. 8a, changes in pH influence adsorption by inducing a change in the charge on the surface of MWCNTs. The optimal pH for adsorption is 6. At low pH, H⁺ inhibits carboxyl group dissociation and prevents the formation of O-, COO-. As the pH rises, the H⁺ content of the

solution decreases, causing deprotenation of the c-MWCNT surface and an increase in adsorption sites, as well as facilitating the coordination of carboxyl groups with metal ions, resulting in an increase in adsorption capacity. (Jamnongkan et al. 2014). However, as the pH rises, the solution becomes increasingly alkaline. According to the metal ion dissolution precipitation curve, when pH>5, some metal ions in the solution begin to precipitate, even though they are still mostly adsorbed. The precipitation trend intensifies as the pH rises, which is also the reason for the sharp drop in the adsorption effect of c-MWCNTs as the pH rises. (Najam et al. 2016).

Figure 8b shows the effect of contact time (from 10-90 minutes) on adsorption efficiency, with c-MWCNTs dose (10 mg), optimum pH=6. the adsorption potential increased steadily as the contact time was increased until it reached a constant of 90 minutes. At the start of the reaction, the binding sites on c-MWCNTs were available to bind with copper ions, but as the contact time was increased, the binding sites were occupied and the equilibrium was reached (Oyetade et al.2016).

3.3.2. Effect of adsorbent dose and initial copper ions concentrations on the adsorption process

The adsorbent dose is an important parameter to determine the free binding site that can bind with initial concentrations of ions. Figure 8c exhibits the effect of c-MWCNT dose (5-20 mg) on the adsorption capacity of copper ions at optimum pH=6, and contact time 90 min. The binding sites for copper ions increased by increasing the dose leading to adsorption capacity enhancement.

At optimum conditions, Figure 8d shows the effect of initial Cu ions concentrations (50-500 ppm aqueous solution) on adsorption capability. According to the findings, increasing the concentration of Cu ions increased the adsorption capacity, demonstrating that the high affinity of c-MWCNT towered the adsorption of Cu ions and that as a result of their surface area.

3.3.3. The adsorption isotherm of copper ions on c-MWCNT

Figure 9a, b show the adsorption isotherms resulting from the two models Langmuir and Freundlich. The correlation coefficient of the Langmuir model is greater than the correlation coefficient of the Freundlich model. As a result, the Langmuir model fits the Cu (II) adsorption data better than the Freundlich model, and the maximum adsorption capacity was 238.09 mg/g. This is consistent with the literature results, which show that the adsorption of Cu (II) on the other materials best fits the Langmuir model (Table 5).

3.3.4. The adsorption kinetic of copper ions by c-MWCNT

As confirmed by many studies, the adsorption kinetic data of heavy metals on CNTs are generally well described by the pseudo-second-order model (Sun et al. 2014). Also from this study the adsorption mechanism can be better described by pseudo-second-order mechanism with good correlation, as shown by the high value of correlation coefficients (0.989) and the well fit to the measured q with experimental results (Figure 10a, b, and Table 6). This suggests that chemisorption, in which valence forces are involved through electron sharing or exchange between the carboxyl (COOH) group and the Cu (II) ions, is the rate-limiting step for Cu (II) ions onto c-MWCNTs (Gupta et al. 2017).

The proposed mechanism of Cu ions adsorption:

Electrostatic attraction, adsorption–precipitation, and chemical interaction between metal ions and CNT surface functional groups are some of the mechanisms for metals adsorbed by CNTs (Rao et.al, 2007). The functionalization of the carbon surface can result in a surface structure that is both more hydrophilic and has a greater number of oxygen-containing functional groups. The complexation and decomplexation reactions are highly dependent on the functionalized groups and solution pH (Rao et.al, 2007), and the chemical reaction between metal ions and the surface functional groups of CNTs is widely accepted as the major adsorption mechanism (Bystrzejewski & Pyrzyńska, 2011, and Rosenzweig et.al, 2013). In the current study, Cu ions exists mainly as a divalent ion (Cu²⁺) at pH 6.0, whereas the carboxyl groups of c-MWCNT are deprotonated. Cu ions is adsorbed by c-MWCNTs by forming a complex with the carboxyl groups on their surface. Chemical complexation, rather than electrostatic attraction, appears to be the most important adsorption mechanism for Cu ions and c-MWCNT binding (scheme 1).

Regeneration of c-MWCNT

After using c-MWCNT as a Cu ion adsorbent, it was centrifuged and washed with diluted HCl (0.1M) to extract Cu ions, washed with distilled water to neutralize the acid, and dried at 100°C to be ready for the second cycle of adsorption. The efficiency of c-MWCNT was found to be 83.5 % after the first cycle, 63.1 % after the second cycle, and 48.9 % after the third cycle. This shows the effectiveness of the synthesized c-MWCNTs regeneration.

4. Conclusion

Because of their importance in a variety of uses and applications, there is a lot of interest in developing low-cost, high-yield, and high-quality MWCNTs. The catalyst Fe/Co supported on aluminum oxide and zirconium oxide was used to synthesis MWCNTs, and the results were compared. Aluminum oxide and zirconium oxide were reported to be suitable supports for the growth of high-quality MWNTs. The optimization of reaction parameters such as the reaction temperature, the gas flow rate, and the catalysts used and their supporting materials could result in MWCNTs with low impurities and high yield. The carbon nanotubes were synthesized in a web-like structure with a homogeneous distribution on the surface of the catalysts, as shown by SEM images. MWCNT grew from catalyst particles using a tip growth method, with an average diameter of 20-30 nm for MWCNT grown on alumina and 30-40 nm for MWCNT grown on zirconia, as seen in TEM images. The carboxylic groups on the MWCNTs were introduced to increase their adsorption efficiency for removing Cu ions from an aqueous solution as part of waste water treatment. With a k_a value of 0.038 and an R² value of 0.995, the data fit best in the Langmuir model, and the adsorption of Cu (II) ion adopted the pseudo-second order model rather than the pseudo-first-order model. The overall adsorption capacity of c-MWCNTs was 238.09 mg/g.

Declarations

Conflicts of interest: There are no conflicts to declare.

Reference

- Abbas A., Al-Amer A. M., Laoui T., Al-Marri M J, Nasser M S, Khraisheh M, Atieh M A (2016), Heavy metal removal from aqueous solution by advanced carbon nanotubes: critical review of adsorption applications. Separation and Purification Technology 157,141-161.
- Abdel Salam M. (2013), Removal of heavy metal ions from aqueous solutions with multiwalled carbon nanotubes: Kinetic and thermodynamic studies. International Journal of Environmental Science and Technology 10, 677–688.
- Ajitha P., Vijayalakshmi K., Saranya M., Gomathi T., Rani K., Sudha P. N., Sukumaran,A. (2017), Removal of toxic heavy metal lead (II) using chitosan oligosaccharide-graft-

maleic anhydride/polyvinyl alcohol/silk fibroin composite. International Journal of Biological Macromolecules, 104, 1469-1482.

- Anzar N., Hasan R., Tyagi M., Yadav N., Narang J. (2020), Carbon nanotube A review on Synthesis, Properties and plethora of applications in the field of biomedical science, Sensors International: 100003.
- Arora B., Attri P. (2020), Carbon nanotubes (CNTs): a potential nanomaterial for water purification, Journal of Computer Science, 4(3), 135.
- Atchudan R., Cha B. G., Lone N., Kim J., Joo J. (2019), Synthesis of high-quality carbon nanotubes by using monodisperse spherical mesoporous silica encapsulating iron oxide nanoparticles, Korean Journal of Chemical Engineering, 36, 157-165.
- Banerjee D., Nguyen T., Chuang T–J. (2016), Mechanical properties of single-walled carbon nanotube reinforced polymer composites with varied interphase's modulus and thickness:
 A finite element analysis study, Computational Materials Science, 114, 209–218.
- Burakova E. A., Dyachkova T. P., Rukhov A. V., Tugolukov E. N., Galunin E. V., Tkachev A. G., Basheer A. A., Ali I. (2018), Novel and economic method of carbon nanotubes synthesis on a nickel magnesium oxide catalyst using microwave radiation, Journal of Molecular Liquids, 253, 340-346.

Bystrzejewski M., Pyrzyńska K., (2011), Kinetics of copper ions sorption onto activated carbon, carbon nanotubes and carbon-encapsulated magnetic nanoparticles, Colloids and Surfaces A: Physicochemical and Engineering Aspects, 377, 402-408.

- Calafat A. (1998), The influence of preparation conditions on the surface area and phase formation of zirconia. In Studies in Surface Science and Catalysis, 118, 837-843. Elsevier.
- Cassell A. M., Raymakers J. A., Kong J., Dai H. (1999), Large scale CVD synthesis of single- walled carbon nanotubes. Journal of Physical Chemistry. B, 103, 6484-6492.
- Chen L., Xin H., Fang Y., Zhang, C., Zhang, F., Cao, X., ... & Li, X. (2014), Application of metal oxide heterostructures in arsenic removal from contaminated water. Nanomaterials 2014 .
- Colomer J., Stephan C., Lefrant S., Van Tendeloo G., Willems I., Onya Z. K., Fonseca A.C. J. B. (2000), Large-scale synthesis of single-wall carbon nanotubes by catalytic chemical vapor deposition (CCVD) method. Chemical Physics Letters, 317, 83-89.

- Dhorea V. G., Rathodb W. S., Patil K. N. (2018), Synthesis and Characterization of High Yield Multiwalled Carbon Nanotubes by Ternary Catalyst. Material Today, 5, 3432-3437.
- Du R., Wu J., Chen L., Huang H., Zhang X., Zhang J. (2014), Hierarchical Hydrogen Bonds Directed Multi-Functional Carbon Nanotube-Based Supramolecular Hydrogels. Small, 10(7), 1387-1393.
 - El-Maghraby A., El-Deeb H. A., Khattab M. A. (2014), Influence of FeNi/Al₂O₃ Catalyst Compositions on the Growth of Carbon Nanotubes. Fullerenes Nanotubes and Carbon Nanostructures, 23, 27–34.
- Ertan B., Eren T., Ermiş I., Saral H., Atar N., Yola M. (2016), Sensitive analysis of simazine based on platinum nanoparticles on polyoxometalate/multi-walled carbon nanotubes, Journal of Colloid and Interface Science, 470, 14-21.
- Fischer J. E., Carbon nanotubes: structures and properties, in: Y. Gogotsi (Ed.), Nanotubes and Nanofibers, Taylor and Francis, CRC Press, New York, (2006) 1–36.
- Fonseca A., Hernadi K., Piedigrosso P., Colomer J –F, Mukhopadhyay K, Doome R, Lazarescu S, Biro L P, Lambin P, Thiry P A, Bernaerts D, Nagy J B (1998) Synthesis of single- and multi-wall carbon nanotubes over supported catalysts. Appl Phys A 67(1): 11-12.
- Freundlich H. M. F. (1906), Uber die adsorption in losungen, Z. Phys. Chem. (Leipzig) 57, 385–470.
- G.P. Rao, C. Lu, F. Su, Sorption of divalent metal ions from aqueous solution by carbon nanotubes: a review, Sep. Purif. Technol. 58 (2007) 224–231.
 - GOPALAN R., CHANG C -H, LIN Y S (1995) Thermal stability improvement on pore and phase structure of sol-gel derived zirconia, Journal of Materials Science, 30. 3075- 3081.
 - Grover, V., Shukla, R., & Tyagi, A. K. (2007), Facile synthesis of ZrO₂ powders: Control of morphology. Scripta materialia, 57(8), 699-702.
 - Gupta V., Saleh T. A. (2011), Synthesis of Carbon Nanotube-Metal Oxide Composite: Adsorption and Photo Degradation; DOI: 10.5772/18009.
 - Hanbali G., Jodeh S., Hamed O., Bol R., Khalaf B., Qdemat A., ... & Dagdag, O (2020),Magnetic multiwall carbon nanotube decorated with novel functionalities: synthesis and application as adsorbents for lead removal from aqueous medium. Processes, 8(8), 986.

Hole R., Munde A., Jaybhaye S. (2021), Functionalization of multiwalled carbon nanotubes with active pharmaceutical ingredient via carboxylation. Materials Today: Proceedings, 45, 3860-3862.

- Hussain M., Begum S. S., Kalita M. K., Ahmed K. U., Nath R. (2018), Additives used in semen preservation in animals: A short review. International Journal of Chemical Studies, 6(5), 354-361.
- Hussein M. Z., Jaafar A. M., Yahaya A. H., Masarudin M. J., Zainal Z. (2014), Formation and Yield of Multi-Walled Carbon Nanotubes Synthesized via Chemical Vapour Deposition Routes Using Different Metal-Based Catalysts of FeCoNiAl, CoNiAl and FeNiAl-LDH, International Journal of Molecular Sciences 15, 20254-20265.
- Ihsanullah I., Sajid M., Kabeer M., Shemsi A. M., Atieh M. A. (2020), First investigations on the removal of tungsten species from water using multi-walled carbon nanotubes. Water, Air, & Soil Pollution, 231(3),1-11.
- Jamnongkan, T., Kantarot, K., Niemtang, K., Pansila, P. P., & Wattanakornsiri, A. (2014), Kinetics and mechanism of adsorptive removal of copper from aqueous solution with poly (vinyl alcohol) hydrogel. Transactions of Nonferrous Metals Society of China, 24(10), 3386-3393.
- Karimi-Maleh H., Tahernejad-Javazmi F., Atar N., Yola M., Gupta V., Ensafi A. (2015) A Novel DNA Biosensor Based on a Pencil Graphite Electrode Modified with Polypyrrole/Functionalized Multiwalled Carbon Nanotubes for Determination of 6-Mercaptopurine Anticancer Drug, Industrial & Engineering Chemistry Research, 54, 3634-3639.
- Kathyayini H.N., Nagaraju N., Fonseca A., and Nagy J.B. (2004), Catalytic activity of Fe, Co and Fe/Co supported on Ca and Mg oxides, hydroxides and carbonates in the synthesis of carbon nanotubes, Journal of Molecular Catalysis A: Chemical 223: 129-136.
- Lagergren S. (1898), About the theory of so-called adsorption of soluble substances. Kungliga Svenska Vetenskapsakademiens Handlingar, 24: 1–39.
- Langmuir I (1916). The constitution and fundamental properties of solids and liquids. Journal of the American Chemical Society, 38: 2221–2295.

- Maruyama S., Murakami Y., Shibuta Y., Miyauchi Y., Chiashi S. (2004) Generation of Single-Walled Carbon Nanotubes from Alcohol and Generation Mechanism by Molecular Dynamics Simulations, Journal of nanoscience and nanotechnology, 4(4):360-367.
- Mohammed I. A., Bankole M. T., Abdulkareem A. S., Ochigbo S. S., Afolabi A. S., Abubakre O. K. (2017), Full factorial design approach to carbon nanotubes synthesis by CVD method in argon environment, South African Journal of Chemical Engineering, 24, 17-42.
- Mwafy E. A. & Mostafa A. M. (2020), Tailored MWCNTs/SnO₂ decorated cellulose nanofiber adsorbent for the removal of Cu (II) from waste water. Radiation Physics and Chemistry, 177: 109172.
- Najam R., Muzaffar S., Andrabi A. (2016), Removal of Cu(II), Zn(II) and Cd(II) ions from aqueous solutions by adsorption on walnut shell-Equilibrium and thermodynamic studies: treatment of effluents from electroplating industry. Desalination and Water Treatment, 57, 27363-27373.
- Oyetade O. A., Nyamori V. O., Martincigh B. S., Jonnalagadda S. B. (2016), Nitrogenfunctionalised carbon nanotubes as a novel adsorbent for the removal of Cu(II) from Aqueous solution. RSC Advances, 6, 2731.
- Patel J., Parikh S., Patel S., Patel R., & Patel P. (2021), Carbon Nanotube (CNTs): Structure, Synthesis, Purification, Functionalisation, Pharmacology, Toxicology, Biodegradation and Application as Nanomedicine and Biosensor: Carbon Nanotube (CNTs). Journal of Pharmaceutical Sciences and Research, 1(02), 017-044.
- Peigney A, Laurent Ch, Dobingeon F, Rousset A (1997) Carbon nanotubes grown in situ by a novel catalytic method, Journal of Material Research, 12, 613-615.
- Pham Q. L., Haldorai Y., Nguyen V. H., Tuma D., Shim J. (2011), Facile Synthesis of Poly(p-Phenylenediamine)/MWCNT Nanocomposites and Characterization for Investigation of Structural Effects of Carbon Nanotubes. Bulletin of Materials Science, 34, 37–43.
- Qingwen L., Hao Y., Yan C., Jin Z., Zhongfan L. (2002), A scalable CVD synthesis of highpurity single-walled carbon nanotubes with porous MgO as support material. Journal of Materials Chemistry, 12, 1179–1183.

Rosenzweig S., Sorial G.A., Sahle-Demessie E., Mack J., (2013), Effect of acid and alcohol network forces within functionalized multiwall carbon nanotubes bundles on adsorption of copper (II) species, Chemosphere, 90, 395–402.

- Sari A. H., Khazali A., Parhizgar S. S. (2018), Synthesis and characterization of long-CNTs by electrical arc discharge in deionized water and NaCl solution, International Nano Letters 8, 19-23.
- Tofighy M. A., Mohammadi, T. (2016), Copper ions removal from aqueous solutions using acid-chitosan functionalized carbon nanotubes sheets. Desalination and Water Treatment, 57(33), 15384-15396.
- Wang J., Shen B., Lan M., Kang D., Wu C. (2020), Carbon nanotubes (CNTs) production from catalytic pyrolysis of waste plastics: The influence of catalyst and reaction pressure. Catalysis Today, 351, 50-57.

Wang Z., Xu W., Jie F. *et al.* (2021), The selective adsorption performance and mechanism of multiwall magnetic carbon nanotubes for heavy metals in wastewater. Scientific Reports **11**, 16878.

- Yan Y. H., Cui J., Chan-Park M. B., Wang X., Wu Q. Y. (2007), Systematic Studies of Covalent Functionalization of Carbon Nanotubes via Argon Plasma-Assisted UV Grafting. Nanotechnology, 18, 24–42.
- Yang Y., Zhang H., Yan Y. (2019), Synthesis of CNTs on stainless steel microfibrous composite by CVD: effect of synthesis condition on carbon nanotube growth and structure. Composites Part B: Engineering, 160, 369-383.
- Yola M., Necip Atar N., (2014a), A novel voltammetric sensor based on gold nanoparticles involved in p-aminothiophenol functionalized multi-walled carbon nanotubes: Application to the simultaneous determination of quercetin and rutin. Electrochimica Acta, 119, 24-31.
- Yola M., Eren T., Atar N., (2014b), Molecularly imprinted electrochemical biosensor based on Fe/Au nanoparticles involved in 2-aminoethanethiol functionalized multi-walled carbon nanotubes for sensitive determination of cefexime in human plasma, Biosensor Bioelectronics, 60, 277-285.

Fig. 1. XRD patterns of (a) Fe\Co / ZrO₂ (b) catalyst Fe/Co / Al₂O₃.

Fig. 2. Scanning electron micrograph (SEM) of (a) Fe/Co /Al₂O₃ and (b) Fe/Co /ZrO₂.

Fig. 3. Plot of the isotherms $Fe/Co/Al_2O_3$, and $Fe/Co/ZrO_2$ catalysts powders measured at temperature 77K.

Fig. 4. Scanning electron micrograph (SEM) of carbonaceous nonomaterial obtained with Fe/Co /ZrO₂ (a) at 700 °C, (b) at 800 °C, (c) at 900 °C, SEM of Fe/Co/Al₂O₃ (d) at 700 °C, (e) at 800 °C, (f) at 900 °C.

Fig. 5. TEM images at magnification 200 nm (A) and 500 nm (B) for MWCNTs grown on Fe/CO/ Al_2O_3 at 700 °C in acetylene for 60 min.

Fig. 6. TEM images at magnification 200 nm (A) and 500 nm (B) for MWCNTs grown on Fe/CO /ZrO₂ at 700 °C in acetylene for 60 min.

Fig 7. (a) XRD pattern of the oxidized MWCNT, (b) FTIR analysis of the oxidized MWCNT

Fig. 8 (a) the effect of pH on the adsorption of copper ions, (b) the effect of contact time on the adsorption process, (C) effect of the MWCNTs dose on the adsorption process, (d) the adsorption capacity of MWCNTs at different concentrations of Copper ions in the aqueous solutions.

Fig. 9a, Langmuir isotherm model for Cu (II) ion adsorption on 10 mg c-MWCNTs, (b) Freundlich isotherm model for Cu (II) ion adsorption on 10 mg c-MWCNTs.

Fig. 10a, Pseudo-first-order kinetic plot for Cu (II) ion adsorption on 10 mg c-MWCNTs, (b) Pseudo-second-order kinetic plot for Cu (II) ion adsorption 10 mg c-MWCNTs.

Sechem1. The adsorption mechanism for Cu ions by c-MWCNT.

Table 1. BET results of Fe/Co/Al₂O₃, and Fe/Co/ZrO₂ catalysts

Catalyst	$S_{BET} (m^2 g^{-1})$	Pore volume (cm ^{2} g ^{-1})	Pore diameter (nm)	
Fe/Co/Al ₂ O ₃	2.59	7.164 ×10 ⁻³	11.026	
Fe/Co/ZrO ₂	11.67	63.4 ×10 ⁻³	21.729	

Table 2. The yield % of carbon deposit from acetylene decomposition on ZrO_2 and Al_2O_3 supported materials at reaction temperature 700°C.

Catalyst	Nitrogen flow (ml min ⁻¹)	Acetylene flow (ml min ⁻¹)	The yield (%) of Carbon deposit depending on the reaction time	
			40 min	60 min
Fe/Co on ZrO ₂	75	25	71.5	82.21
		50	86.3	88.9
	120	25	52.1	60.3
		50	55.0	61.1
Fe/Co on Al ₂ O ₃	75	25	59.1	64.0
		50	67.4	73.5
	120	25	48.0	56.3
		50	55.9	71.34

Table 3. The yield % of MWCNTs at different reaction temperature.

Catalyst	Temperature °C	Yield %
Fe/Co on ZrO ₂	700	76.0
	800	64.3
	900	60.0
Fe/Co on Al ₂ O ₃	700	61.1
	800	59.1
	900	55.0

Table 4. Literature reaction conditions and yield % of obtained MWCNTs.

Catalyst	Carbon	Reaction		Reaction		Yield	Ref.
	source	Conditions		%			
		Temp.	Flow Rate				
		°C	ml min ⁻¹				
Fe/Co/Al ₂ O ₃	Acetylene	700	50	73.5	Present work		
Fe/Co/ZrO ₂	Acetylene	700	50	88.9	Present work		
Fe/Ni/Al ₂ O ₃	Ethanol	900	-	68	(Colomer et al. 2000)		
Co/Fe/Cu on SiO ₃	acetylene	700	8	71-76	(Fonseca et al. 1998)		
Fe/Co/Zeolite	Ethanol	800	5 Torr	42	(Maruyama et al. 2004)		
powder							

Table (5): Maximum adsorption capacity of different carbon materials.

Adsorbent	Adsorption capacity (mg/g)	Adsorption technique	Ref	
c-MWCNT	238.09	10 mg	This work	
acid–chitosan- functionalized CNTs sheets	57.34	50 mg	Tofighy & Mohammadi (2016).	
P-MWCNTs	0.398	125 mg	Abdel Salam (2013)	
MWCNTs-COOH	49.1	30 mg	Esmail & Ebrahimi (2014)	
F-MWCNT	118.41	10 mg	Gupta et al. (2017)	
MWCNTs-COOH	19.44	50 mg	Oyetade et al.(2013)	

Table (6): Fitting parameters for first and second order kinetic for adsorption of Cu (II) ion onto c-MWCNT.

Adsorbent	q exp	Pseudo-first order			I	Pseudo-seco	nd order
c- MWCNT		K 1	qe (cal)	R ² %	K ₂	qe (cal)	R ² %
	133.3	0.034	74.47	0.876	0.0699	131.31	0.989



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(a)

(b)



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(c)

(d)



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