Wollastonite Ceramic/CuO Nano-Composite for Cadmium Ions Removal from Waste Water

Nabila Ammar¹, Ahmed Fahmy A. Youssef², S. H. Kenawy^{3*}, E.M.A. Hamzawy⁴, M.A. El-Khateeb¹

¹Water Pollution Control Department, National Research Center, ²Chemistry Department, Cairo University, ³Ceramic Department, ⁴Glass Department, National Research Center, Dokki, Cairo, Egypt.

NANO-PARTICLES, within 29-45 nano-meters, based on wollastonite were prepared with/ without CuO through wet method. The adsorption process was occurred on nano-particles powder ceramic at different conditions (pH, contact time and dose) for removing of cadmium ions from waste water. The SEM of nano-particles after treatment of waste water revealed that, surface texture of the blank adsorbents changes totally due to the adsorption of cadmium ions on the particles. At optimum operating conditions (at pH 9) cadmium ions removal reached 98.88% after measurement by ICP and EDX techniques. The obtained results showed that the produced ceramic nano-particles could be used efficiently for removal of cadmium ions with reasonable simple and low cost technique.

Keywords: Wollastonite, Ceramic, Nano materials, Water treatment, Adsorption.

Introduction

The heavy metals (HM) release into the environment has a potential threatening effect to water, soil quality as well as to plant, animal and human health, directly and indirectly. Heavy metals adversely affect the fertility of soils, production of plants, animals and the entire environment if exceed certain limits [1]. Furthermore, these metals could be soluble in aquatic media in the environment and consequently, may be easily absorbed by the cells of living creatures. Thus, by entering the food chain, HM could be bioaccumulated in higher levels of food chain. When HM absorbed above the recommended dose, could cause severe health hazards [2, 3]. Heavy metals could be bio-accumulated through different trophic levels and unlike organic toxicants considered non-biodegradable [4, 5]. Over the last two decades there was an extensive use of Cd worldwide for production of batteries compared with its use for other applications. There were several international agreements and legislation to regulate and monitor the emission of Cd to the environment consequently; minimizing human and environmental threat of Cd which could cause failure of kidney in all mammals [6, 7]. Also, Cd has particular environmental interest as one of the toxic heavy metals, because it could

be accumulates in soils via agricultural usage of fertilizers, sewage residues, disposal of Cdpolluted wastes (industrial as well as municipal). Cadmium is an outstanding human cancer-causing agent and may cause hypertension, bone injuries and lung deficiency [8, 9].

Adsorption among the available remedial technologies is the most popular because of its simplicity and low cost. Different natural (e.g. metal oxides, bentonites, natural zeolites) and synthetic (resins, modified zeolites, synthetic hydroxides /oxides/hydroxides, metal phosphates and silicates) materials have been tested in adsorption process for removal of heavy metal from wastewater with various efficiencies [10, 16]. In addition, significant research work has been done on different waste materials to improve and investigate proper sorbents for water and wastewater treatment, blast furnace slug [17, 18], biomass [19, 20], fly ash [21], and bagasse fly ash [22, 23], among other have been investigated as sorbents for removal of heavy metal with various levels of achievement.

Nano-metal oxides are a highly valuable material with various applications in catalysts [24], gas sensors [25], optical, electrical and mechanical devices (Yi et al. 2011), cosmetics

^{*}Corresponding author e-mail: ksayed6631@gmail.com DOI: 10.21608/ejchem.2017.1289.1076

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and sunscreens [26]. Nano-metal oxides such as hydrous manganese oxides, iron, aluminum, titanium oxides and zinc oxide are extensively used for removal of heavy metal ions from water and wastewater. These nano-metal oxides can be manufactured by different techniques such as microwave-assisted route [27], chemical vapor deposition (CVD), different hydrothermal techniques and sol–gel method [28].

The main objectives of this study are the synthesis a nominal nano-calcium silicate (wollastonite) ceramic doped with CuO as metal oxide using a very simple technique. As well as, investigate the adsorption capacity of the produced nano-composite as an adsorbent material to remove of Cd ions from water samples.

Materials and Methods

Materials

Nominal nano-calcium silicate ceramic (wollastonite) was prepared from calcium carbonate (El-Gomhorya Company for Chemicals, Egypt 99%), silicon hydroxide gel (Fluka) and copper carbonate (CuCO₃). The source of Cd(II) used in the adsorption studies was high purity cadmium chloride hexahydrate (CdCl₂. $6H_2O$), (Chem-Supply-Merck).

Synthesis of Nominal nano-calcium silicate ceramic

Nominal calcium silicate ceramic (wollastonite) was prepared from pure calcium carbonate, silicon hydroxide (silica gel) and copper carbonate (CuCO₂), was used as precursors for CuO was doped (0.0 - 7g/ 100 g CaSiO₃ powder) . CaCO₃ was dissolved in stochimetric amount of nitric acid (HNO₂). The freshly prepared calcium nitrate [$Ca(NO_2)_2$] was dissolved in appropriate amount of distilled water gently with slow stirring. The silicon hydroxide gel was added slowly to the calcium nitrate under vigorous stirring for 2h to ensure good mixing. The CuO dopant, freshly prepared from CuCO₂ with distilled water to gave Cu(OH), was added under continues vigorous stirring for 1 hr. The homogenous composite was dried at 100° C overnight followed by calcined at 550° C/2h. The following equations explain the steps of produce CaSiO₂ with and without CuO dopant :

CaCO₃ + HNO₃ <u>Cooling</u> Ca(NO₃) + H₂CO₃ Ca(NO₃) _{sub.} + Si (OH)₄ <u>survive</u> <u>String /24</u> Ca SiO_{3 starty}

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X-ray investigations of the CaSiO₃ with /CuO calcined at 550°C in air for 2h was conducted using XRD (BRUKER, D8 ADVANCED Cu target, Germany). XRD was carried out in the range from 20 to 60 20°. The morphology of the sample was checked via scanning electron microscopy (SEM) coupled with energy-dispersive spectroscopy EDX, (SEM Model Quanta FEG 250).

Removal of the metal ions using the nano-ceramic sorbent

A batch sorption method was used to study the efficiency of the nano-ceramic for the removal of Cd ions. The influences of initial metal ion concentration, adsorbent dose, contact time and pH of solution on the sorption of Cd ions were examined. A Jar test procedure was used for ensuring homogeneity of the metal ions salts in solutions. The metal ions were estimated in the samples after filtration with filter papers Whatman[®] (No. 41). Control samples without adsorbent were used for comparison between sorption and precipitation of metal ions. All experiments were repeated three times, the percentage of metal ions removal by the nanoceramic was measured for all samples using inductive coupled plasma optical emission spectrometry (Agilent ICP-OES 5100) according to standard methods for the examination of water and wastewater [APHA, 2012].

Results and Discussion

Characterization of nano-ceramic sorbent

The X ray diffraction pattern of calcined $CaSiO_3/CuO$ powder was shown in Fig. 1. The identified phases were wollastonite (CaSiO₃, ICDD, 01-076-0925) and for tenorite (CuO, ICDD, 00-005-0661).

The SEM micrograph of the calcined $CaSiO_3/$ CuO powder before soaking in waste water was shown in Fig. 2. The photograph show cluster of nano-particles within 29–45 nm nano-meter , however some connected barrier of nano-particles which make boxes or pores were clearly visible in the all sample field (Fig.2). Micrograph of the sample, illustrate some tiny particles as well as particles of a somewhat larger size indicating some of crystallinity. Figure 2 shows that, the particles are near uniform spherical, the irregular surface of the particles increases the surface area and the particle size can be estimated in the range of 29–45 nm. The resulting microstructure and the pore size distribution of the obtained ceramic



Fig. 1. X ray diffraction patterns of CuO- doped wollastonite treated at 550°C/2hr.

are particularly important for potential adsorption applications.

The pH has significant effect on adsorption, as it affects the surface charge of adsorbent and also chemical speciation of the adsorbate. Adsorption of Cd ions was studied at different pH ranging from 3 upto 9 where, Cd^{2+} , Cd(OH), $Cd(OH)_2$, and $Cd(OH)_{2(S)}$ are the forms of cadmium present in de-ionized water . In these experiments adsorbate concentration was 25 mg/L, ceramic mass was 0.75 g at contact time varying from 0 to 60 min.

Figure 3 shows the effect of pH on adsorption



Fig. 2. SEM of the Nano-ceramic before Cadmium adsorption.

process (initial Cd concentration was 25 mg/l). It can be seen that, adsorption increases with increasing the pH of the solution. The removal efficiency increased from 43.2 % at pH 3 to 98.88 % at pH 9. Where, at lower pH values H⁺ ions compete with Cd(II) ions for binding sites on the absorbent surface. However, as the pH increases there is a reduction in competition between H^+ , Cd^{2+} and $Cd(OH)^+$ ions and as a result, metal uptake by the absorbent increases. The experimental results indicate that, Cd(II) removal was predominantly controlled by adsorption upto a pH 8. It was showed that [29], the Cd(II) ion predominates at pH below 7 and it begins to precipitate out as Cd(OH), at pH values just above 9. Based on the results, further experiments were conducted at pH 9.

Effect of contact time

Adsorption of Cd ions was studied at different contact time ranging from 0 to 120 min. with



Fig. 3. Effect of pH value on the removal efficiency of Cd ions by using the nano-ceramic (initial Cd concentration was 25 mg/l).

25 mg/L of adsorbate concentration, the nanoceramic mass was 0.75 g and pH 9. Figure 4 shows the influence of contact time on the efficiency of the nano-ceramic for the removal of Cd(II) ions. It can be noted that, the removal efficiency increased from 92.4 % to 98.88 % and almost



Fig. 4. Effect of contact time on the removal efficiency of Cd ions by using the nano-ceramic at pH: 9 and T: 298 (initial Cd concentration was 25 mg/l).

remain constant from 60 until 120 min. Therefore, the equilibrium adsorption time is 60 min.

Effect of adsorbent dosage on adsorption

The effect of adsorbent dose was studied using 25 mg/L of adsorbate concentration, at pH 9 with nano-ceramic mass varied from 0.25 to 1.0 g/L with 0.25 mg/l intervals, at contact time ranging from 0 to 60 min.

The general trend indicates that, 0.25 to 0.75 g/L of sorbent dose uptake huge percent of Cd(II) (Fig. 5) where, it can be observed that the removal efficiency increased from 96.04 - 98.88 % and nearly remain constant from 0.75 - 1.0 g/L. The reason for the increasing of removal efficiency could be attributed to the increasing of adsorbent

dose that means increase of the total available surface area of the adsorbent particles. Beyond 0.75 g/L, the removal remained almost unaffected by the adsorbent dosage so, at 0.75 g/L, the adsorbent surface becomes saturated with Cd(II).

The SEM of Cd on the nano-ceramic sorbent

The SEM micrograph of the Cd loaded on nano-calcium silicate (wollastonite) doped with CuO is shown in Fig. 6. The photos show light coated nano-particles of Cd-metal on the surface of ceramic. It could be noted from Fig.6, that the surface morphology and porosity of the nanoceramic (wollastonite) doped with CuO were drastically changed after loading of cadmium as compared with the surface shape of the blank adsorbents. The EDS microanalysis indicates the



Fig. 5. Effect of adsorbent dosage of the nano-ceramic on the removal efficiency of Cd ions at pH: 9 and T: 298 (initial Cd concentration was 27.4 mg/l).

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Fig. 6. SEM of the nano-ceramic sample after cadmium adsorption.

presence of cadmium cations after adsorption onto surface of nano-ceramic doped with CuO (Fig.7).

The aforementioned results were in agreement with previous work where, the synthesized nano-structured calcium silicates showed that the synthesized material based on an amorphous materials without a defined structure. In our work the particle size were ranged from 29-45 nano-meter synthesized using commercially inorganic materials, while for the previous work the mean particle size



Fig.7. Energy dispersive spectrum (EDS) of the Cd-loaded nano-Calcium Silicate doped with CuO.

averaging was 0.5 to 1.0 μ m with particles forming larger agglomerates.

Conclusions

- Nano-Calcium Silicate doped with nanocopper is found to beneficial for removing of heavy metal such as Cd from an aqueous solution. The studied material was synthesized using very simple technique (coprecipitation method) from low coast starting chemicals. The prepared ceramic doped with CuO was investigated using different tools.
- Higher uptakes (98.88% of Cd(II)) from aqueous solution is possible using wollastonite as an adsorbent provided the initial concentration of the Cd(II) in the wastewater is around 25 mg/l. This material could be used for industrial wastewater treatment.
- The pH has been found to be a master variable controlling the adsorption of Cd(II) by the oxides present in wollastonite. The adsorption is maximum around pH 9.0

and it has been concluded that surface complexation involving H^+ exchange and chemical precipitation contribute towards maximum uptake.

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متوالفات نانو سير اميكيات الولوستونيت/اكسيد النحاس لاز الة ايونات الكادميوم من المياة الملوثة

نبيلة عمار '، أحمد يوسف'، سيد قناوي '، عصمت حمزاوي ' و محمد الخطيب ' 'قسم تلوث المياه - المركز القومي للبحوث، 'قسم الكيمياء - جامعة القاهرة، 'قسم السير اميك و 'قسم الزجاج -المركز القومي للبحوث - القاهرة - مصر .

تم تحضير حبيبات نانو ولوستونيت مطعم بأكسيد النحاس باستخدام طريقة الترسيب. وكان حجم الحبيبات يتراوح بين ٢٩-٤٠ نانوميتر. وقد تمت عملية الادمصاص لايونات الكادميوم عند درجات مختلفة من الاس الهيدروجيني وكذلك علي فترات زمنية محددة وايضا عند تركيزات مختلفة من ايونات الكادميوم. وقد اظهر الفحص باستخدام الميكروسكوب الماسح الاليكتروني ان حبيبان النانو للعينات بعد الادمصاص تغطية كاملة لكل حبيبات النانو بأيونات الكادميوم مما ادي الي تغيير النسيج السطحي للحبيبات. وقد لوحظ انه عند الاس الهيدروجيني المثالي وهو (٩) ان حوالي ٩٨.٨٪ من ايونات الكادميوم قد ازيلت بواسطة تقنية ال ICP و EDX . ومما سبق اظهرت النتائج المتحصل عليها امكانية استخدام حبيبات السيراميك النانوميترية بكافاءة عالية لازالة ايونات الكادميوم من المياه الملوثة مع الاخذ في الاعتبار ان طريقة التحضير للسيراميك بسيطة ورخيصة للغاية.