

Biosorption Studies on Cu (II) Ions Using Powdered Leaves of *Tabernaemontana divaricata* as Adsorbent

R. Rabena¹, R. Ajitha²

¹Mphil Scholar, Department of Chemistry, Women's Christian College, Afflicted to ManonmaniamSundaranar University, Tirunelveli, Tamil Nadu, India 629 001

²Assistant professor, Department of Chemistry, Women's Christian College, Afflicted to ManonmaniamSundaranar University, Tirunelveli, Tamil Nadu, India 629 001.

Abstract—The commonly used process of heavy metal removal is chemical precipitation, electro winnowing etc., in this study low-cost biosorbent such as the leaf powders of Tabernaemontana divaricata were used for the study of Cu(II) metal ions removal. The experiment was performed by using batch experiments. The effect of concentration, dosage, and time variations was checked. According to the results highest percentage of adsorption attained by Tabernaemontana divaricata was 75% at 750mg dosage of adsorbent and time of contact at 30 min. The regression value is indicating that the order of all used isotherm models such as Freundlich, Langmuir and Temkin. The characterization of adsorbents was done by using FT-IR spectral analysis

Keywords—- Biosorption, Biosorbent, Batch experiment, characterization, isotherm.

I. INTRODUCTION

Water contamination by trace metals is rapidly growing due to increasing human activities in the mining, agriculture and manufacturing industries. The trace metals are nonbiodegradable, highly toxic and cannot be detoxified biologically [1]. Biologically copper is an essential trace element for the terrestrial green plants and algae under the concerned limits and involves in photosynthesis, it is a cofactor of oxidase, oxygenase and enzymes such as superoxide dismutase and acrobat oxidase [2]. Researchers have reported that water discharged from copper electroplating industries, textile industries or washing effluents may contain up to 500mg-¹ copper [3]. Therefore, it is necessary to remove Cu ions from industrial wastewater before discharging them into natural water bodies. The conventional methods used for the removal of heavy metals include chemical precipitation, lime coagulation, ion exchange etc.., [4]. Biosorption is an alternative process for the uptake of heavy metals from aqueous solutions by biological materials. The biosorption of metals by biomass has been much explored in recent years. Different forms of inexpensive, non-living plant materials such as agricultural waste materials and seaweeds, molds, yeasts and other dead microbial biomass have been widely investigated as a potential biosorbents for heavy metals [5,6]. In this study, we used some of the leaf powder of Tabernaemontana divaricata is used as the biosorbent. It is considered economic and ecofriendly due to its unique chemical composition, availability in abundance, renewable, low in cost and more efficient for heavy metal remediation.

II. MATERIALS AND METHODS

A. Preparation of adsorbent

Tabernaemontana divaricata leaves were selected as adsorbents for the removal of Cu(II) ions in this study. The leaves were obtained from the concerned plants located in nearby areas of Nagercoil. *Tabernaemontana divaricata* leaves

were collected and each leaf was washed thoroughly under running tap water to remove dust and any adhering particles. The leaves were then dried under sunlight for a few days and then in an oven at 80 °C until they become crisp. The driedleaves were crushed and blended to powder form using a blender. The *Tabernaemontana divaricata* leaves powder were boiled and washed several times until it was free of color and turbidity. Then the powder was dried in an oven at 80 °C overnight and stored in airtight plastic for further use toavoid contact with moisture in the atmosphere.

B. Preparation of adsorbate

A stock solution of Cu (II) ions was prepared from the salts of copper sulfate (CuSO₄.5H₂O) in double distilled water. The amount of 3.921 g of CuSO₄.5H₂O is making up to 1000 mg/L. The amount of copper sulfate before and after biosorption is estimated by using Iodometric titration.



Fig. 1. Leaves of Tabernaemontana divaricata

Fig. 2. Powdered leaves of Tabernaemontana divaricata

C. Adsorption experiment

Batch experiments were carried out by shaking a series of bottles containing various amounts of the different adsorbent such as *Tabernaemontana divaricata* in heavy metal ion solutions like Cu(II) and separately. The experiment is varied



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with the effect of initial concentration, the effect of adsorbent dosage and the effect of contact time. The solution bottles were agitated for one hour until equilibrium was attained, at the end of the mixing of the adsorbent particles with the adsorbate, the whole mixture was separated from the suspension by filtration through a Whatman filter paper. The percentage of metal removal was calculated by using the formula,

% Biosorption = $(Co - Ce)/Co \times 100$ Where, C_o=Initial concentration of metal ion; C_e= Concentration of metal ion at equilibrium state. Freundlich, Langmuir and Temkin isotherms were also applied to explain the relationship between adsorbent and adsorbate. The characterization of the adsorbent was studied by using FT-IR analysis.

III. RESULT AND DISCUSSION

A. Effect of initial concentration

The effect of the initial concentration of adsorbate ion is studied by taking 20mg, 40mg, 60mg, 80mg & 100mg with an adsorbent dosage of 150mg, constant room temperature (27.6 °C), Constant time (60mins) and constant rotation per minute (150rpm). The adsorption percentage of Cu(II) ion on *Tabernaemontana divaricata* leaves was decreased by 45%, 32.5%, 25%, 15% & 12% while the initial concentration increased were depict in Fig: 3.

The percentage of adsorption of the metal ion by *Tabernaemontana divaricata* leaves was decreased in every varying concentration of metal ion this is because at lower concentration the number of Cu(II) ion is low when compared to the available adsorbent active sites, therefore, adsorption is more frequent.



Fig. 3. Plot of percentage of adsorption Vs initial concentration for *Tabernaemontana divaricata* leaves

B. Effect of adsorbent dosage

In this study, five different dosages were recommended 150mg, 300mg, 450mg, 600mg, and 750mg respectively. The process of adsorption was carried out under the identical initial metal ion concentration condition (20mg), constant room temperature (27.6°), constant contact time (60 min) and constant rotation per minute (150 rpm). The adsorption percentage of Cu(II) ions on Tabernaemontana divaricata leaves was increased by 45%, 60%, 65%, 70%, & 75% at beginning adsorption increases rapidly after a particular dosage 300mg the

percentage of adsorption increases in a gradual manner is depicted in Fig:4.

The above results concluded that the adsorbent dosage increase and the adsorption of Cu(II) ions on *Tabernaemontana divaricata* leaves were also increases. The equilibrium value of the amount adsorbed (q_e) also decreases. This is because of the increase in the amount of adsorbent and also there is an increase in the number of available adsorption sites which directly increases the rate of adsorption. The number of metal ions adsorbed to the adsorbent unit mass decreases and causes q_e value to decrease as the adsorbent dosage increases. The maximum percentage removal of Cu(II) ion was achieves as 75% by Tabernaemontanadivaricata leaves as adsorbent.



Fig. 4. Plot of percentage of adsorption Vs adsorbent dosage for *Tabernaemontana divaricata*leaves

C. Effect of contact time

The recommended contact time for this study ranged from 30-150 minutes, at time intervals of 30 minutes. In this study, the effect of the contact time of the adsorption of Cu(II) ions was carried out at identical initial metal concentration conditions (20mg), constant temperature (27,6 C), constant rotation per minute (150rpm) and constant adsorbent dosage (150mg).

The percentage of adsorption on *Tabernaemontana divaricata* leaves is maximum at 90min at 60% but from the initial time at 30min adsorption is only 55% after that its tends to decrease till 35% this may be due to the availability of the active sites and uncovered surface on the adsorbent (Here dosage keeps constant as 300mg). After 60minutes adsorption was not rapid and become about constant due to the saturation of available binding sites with the metal ions depict in Fig:5.



Fig. 5. Plot of percentage of adsorption Vs contact time for *Tabernaemontana* divaricata leaves

From the above results, the contact time increase there is a variable percentage of adsorption is observed that it reaches the saturation the variation occurs after the maximum percentage of adsorption it remains constant by decreasing or increasing



manner the highest percentage of adsorption is achieved by *Tabernaemontana divaricata* leaves as 75% at 30min contact time which may due to availability of saturation sites and higher dosage.

D. Freundlich Isotherm

The mathematical form of the Freundlich equation is given by the following equation,

$$\log qe = \log kf + \left(\frac{1}{n}\right)\log Ce$$

Where, C_o =Initial concentration of metal ion; C_e =Concentration of metal ion at equilibrium state; K_f = Freundlich constant; n = order of reaction; qe = amount of adsorbate adsorbed per unit mass of adsorbent. The value of K_F is 2.4434 and 1/n is 0.135 for *Tabernaemontana divaricata* leaves is calculated from slope and intercept of the plot log qe vs log C_e in Fig 6. The value 1/n is less than 1 and $R^2 = 0.364$ shows favorable adsorption by *Tabernaemontana divaricata* leaves.



Fig. 6. Freundlich plot for Cu(II) loaded Tabernaemontana divaricata leaves

E. Langmuir Isotherm

Mathematically the linear form of the Langmuir equation is given below.

$$\frac{Ce}{qe} = \frac{1}{aKL} + \frac{Ce}{KL}$$

Where, C_o =Initial concentration of metal ion; C_e =Concentration of metal ion at equilibrium state; qe = amount of adsorbate adsorbed per unit mass of adsorbent; a, K_L = adsorption capacity and rate of adsorption.

When C_e/q_e plot against C_e a straight line with a slope of $1/K_L$ was obtained. The characteristics of the Langmuir isotherm model are expressed in terms of the dimensionless parameter (RL). This parameter is mathematically defined as $R_L = 1 + aCo$ where, a = Langmuir constant, $C_o =$ initial concentration of copper ions the value of R_L indicates the type of isotherm is either favorable ($0 < R_L < 1$), unfavorable ($R_L > 1$), linear ($R_L = 1$) or irreversible ($R_L = 0$).

The value of K_L and a for *Tabernaemontana divaricata* leaves were found to be 4.0816 and -1.75. The value of R_L was found to be less than one and this confirms that Langmuir isotherm model is fitted for the adsorption of Cu (II) on *Tabernaemontana divaricata* leaves. The value of regression R^2 is found as 0.921, 0.977 and indicating the copper favors the Langmuir isotherm model and monolayer adsorption onto the surface of *Tabernaemontana divaricata* leaves depict in Fig: 7



Fig. 7. Langmuir plot for Cu(II) loaded Tabernaemontana divaricata leaves

F. Temkin Isotherm

The Temkin isotherm model explains the influence of metal-metal interaction on the surface of biosorbent. Therefore, the heat of biosorption of metal sorbate on the surface decreases linearly with the occupation due to metal-metal interactions. This linear form of Temkin isotherm is given by the following expression:

$$q_e = a + b \ln C_e$$

Where, C_e =Concentration of metal ion at equilibrium state; q_e = amount of adsorbate adsorbed per unit mass of adsorbent; a,b = isothermal constants related to binding capacity and Temkin constant. The value of 'a' and 'b' can be obtained from the plot of qe vs ln Ce.



Fig. 8. Temkin isotherm plot for Cu (II) loaded Tabernaemontana divaricata leaves

G. Characterization of adsorbent

FT-IR spectral analysis is used to study the charactrestic functional group which is responsible for adsorbing metal ions. The FT-IR spectra of *Tabernaemontana divaricata* before and after adsorption on Cu(II) ions were studied below.

According to Figure 8 before biosorption of Cu (II) ions on *Tabernaemontana divaricata* leaves shows a broad band near 3406 cm^{-1} due to O-H vibration in the hydroxyl group. The peak at 2921 cm⁻¹ and 2851 cm⁻¹ are assigned to C-H stretching. The band at 1619 cm⁻¹ represents N-H stretching. The peak at 1374 cm⁻¹ and 1318 cm⁻¹ is due to the deformation of CH₃ and the stretching of C=O. The band at 1248 cm⁻¹ and 1062 cm⁻¹ is due to C-O stretching & bending and C-OH stretching & bending. After the biosorption of Cu (II) ions avariation of the FT-IR spectrum is observed in Fig8 broad intense band near 3416 cm⁻¹ due to O-H vibration in the hydroxyl group the peak becomes



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more intense due to the presence of copper ions in salt used for the analysis. C-H stretching peak shifted and become intense at 2923 cm⁻¹ and 2854 cm⁻¹. The N-H stretching band shifted at 1640 cm⁻¹ represents. The deformation of CH₃ and stretching of the C=O peak shifted to 1383 cm⁻¹ and 1318 cm⁻¹. C-O stretching and C-OH stretching and bending bands shifted at 1239 cm⁻¹ and 1105 cm⁻¹.



Fig. 9. FT-IR spectrum of *Tabernaemontana divaricata* leaves before biosorption of Cu (II) ions



Cu(II) ions IV. CONCLUSION

This work aim is to find the possible use of *Tabermontana divaricata* leaves powder as adsorbent for biosorption of Cu(II) ions from an aqueous solution. All the equilibrium concentrations perfectly fitted with Langmuir, Freundlich and Temkin adsorption isotherms the highest percentage of adsorption attained by *Tabermontana divaricata* was 75% at constant time and a high adsorbent dosage of 750mg. According to the variation in the FT-IR spectra before and after adsorption considered that ion-exchange was a major adsorption mechanism and a certain amount of surface complexation mechanism coexisted. The overall results indicate that *Tabermontana divaricata* leaves are an effective,

low cost and renewable biosorbents for the biosorption of Cu(II) ions from aqueous solution.

REFERENCES

- Mambo Moyo., Admire Chirinda., Tichaona Niharingo., (2016). "Removal of Copper from Aqueous solutionusing Chemically Treated Potato (Solanum Tuberosum) Leaf Powder". *Journal ofClean SoilAir Water*, Vol 44(5), pp 488-495.
- [2]. Naveen Chandra Joshi, Vivekanand Bhaguna., (2018). "Biosorption of copper (II) on the waste leaves of Kafal (*Myrica esculenta*)". Journal of Rasayan chem, Vol 11, pp 142-150.
- [3]. Dermantzis, K., Davidis, A., Papadopoulou, D., Christoforidis, A., Ouzounis, K., (2009). "Copper Removal from industrial waste waters by means of Electrostatic shielding Electrondeionization". *Journal of Engineering science and technology*, Vol 2, pp 131-136.
- [4]. Suhanya.M., Saranya.K., Thirumurugan.M., Kannadasan.T., (2014). "Reduction of Hexavalent Chromium by using Immobilized Hibiscus Rosasinensis leaves". *International Journal of Science & Technology*, Vol 2(4), pp 77-84.
- [5]. Hashem, A., Abdel-Halim, E.S., El-Tahlawy, K.F. & Hebeish, A. (2005), "Enhancement of the adsorption of Co(II) andNi(II)ionsontopeanuthullsthroughesterificationusingcitricacid". *Journ* al of AdsorptionScience & Technology, Vol 23, pp367-380.
- [6]. Hashem, A., Akasha, R.A., Ghith, A. & Hussein, D.A. (2007), "Adsorbent based on agricultural wastes for heavy metaland dyeremoval: Areview". *Journal of EnergyEducationScienceand Technology*, Vol 19, pp 69-86.
- [7]. Bhatti, H. N., Nasir, A. W., and Hanif, M. A., (2010) "Efficacy of Daucus carota L waste biomass for the removal of chromiumfromaqueoussolutions". *Journal of Desalination*, vol.253, pp 78-87.
- [8]. Amanda E Martins, Milene S. Pereira, Alexandre O. Jorgetto, Macro A.U. Martines, Rafel I.V. Silvia, Margarida J.Saeki, Gustavo R.Castro., (2013). "The reactive surface of Castor leaf [*Ricinuc communis L.*] powder as a green adsorbent for the removal of heavy metals from natural river water". *Applied surface science, vol*276, pp 24-30.
- [9]. Anitha.A., Kohilavani.k., Murugalakshmi.R., (2018). "Removalof Copper (II) by adsorption on Biomass Carbon derived from (*Pongamia pinnata*) Leaf". *International Journal of Applied Engineering Research*, vol 13, pp 14669-14674.
- [10]. Benaïssa, H. and Elouchdi, M.A., (2006). "Removal of copper ions from aqueoussolutions by dried sunflower leaves". *Chemical Engineering and Processing*, vol 46, pp 614-622.
- [11]. Suhanya, M., Saranya, K., Thirumurugan, M., Kannadasan, T., (2014). "Reduction of Hexavalent Chromium by using Immobilized Hibiscus Rosasinensis leaves". *International Journal of Science & Technology*, vol 2(4), pp 77-84.
- [12]. Reena Malik., Suman Lata., Suhila Singhal., (2015). "Removal of heavy metal from waste water by the use of modified aloe vera leaf powder". *International Journal of Basic and Applied Chemical Science*, vol 5(2), pp 6-17.
- [13]. Parisa Ziarati., SherminehNamvar., Barbara sawicka., (2018). "Heavy metal bio-adsorption by *Hibiscus Sabdariffa L*. From contaminated water". *Journal ofscientific and technicals*, vol 4, pp 22-32.
- [14]. Langmuir, I., (1916). "The constitution and fundamental properties of solids and liquids". *Journal of the American Chemical Society*, vol 38, pp 2221-2295.
- [15]. Freudlich,H.T.M.,(1906). "Removal of Cd(II) from aqueous solution by activatedcharcoalderivedfromcoconutshell".*ElectronicJournalofEnviron mental*, Agricultureand Food Chemistry, vol 3(4), pp 702-709..