

**INTERNATIONAL JOURNAL OF CURRENT RESEARCH IN
CHEMISTRY AND PHARMACEUTICAL SCIENCES**

(p-ISSN: 2348-5213; e-ISSN: 2348-5221)

www.ijcrpcs.com

DOI: 10.22192/ijcrpcs

Coden: IJCROO(USA)

Volume 5, Issue 4 - 2018

Research Article



DOI: <http://dx.doi.org/10.22192/ijcrpcs.2018.05.04.001>

**Adsorption of Nickel (II) Ion from Aqueous Solution onto
sulphuric acid Activated Carbon Prepared from
Mimusops elengi Leaves**

A. Elavarasan^{a*} and V. Nandhakumar^b

^{a*} Department of Chemistry, Sengunthar College of Engineering, Tiruchengode-637205
Namakkal, Dt – Tamil Nadu, India.

^b Department of Chemistry, A.V.V.M Sri Pushpam College, Poondi – 613 503, Tamil Nadu, India.

*Corresponding Author: kingofchemistry1983@gmail.com

Abstract

Adsorption of Nickel (II) ion onto Activated Carbon prepared from *Mimusops elengi* leaf (SATMEAC) under various experimental conditions was studied. Batch mode adsorption experiments were conducted. The characteristics of the (SATMEAC) were determined by SEM, XRD, BET, FTIR, analysis and pH_{ZPC}. The maximum removal efficiency of Ni (II) ion is 85% at pH 6. Under the chosen conditions, Experimental data obtained were fitted with linearised forms of Lagergren and Ho kinetic models. The Sum of Error Squares Percentage (SSE %) for first order and second order kinetics were 1.58 & 0.75 respectively. Thus this adsorption followed second order kinetics. The equilibrium adsorption data were analyzed with four isotherm models. Order of best fitting isotherm models were found to be Freundlich >Tempkin>Langmuir >Dubinin Raduskevich. The results in this study indicated that (SATMEAC) could be employed as an adsorbent for the removal of Ni (II) ion from an aqueous solution.

Keywords: Adsorption; Nickel (II) ion removal, Kinetics, Isotherms, Sulphuric acid activation.

Introduction

The progressive increase of industrial technology results in a continuous increase of pollution, so that a great effort has been dedicated to minimizing these dangerous pollutants and therefore, avoiding their unsafe effects on animals, plants and humans. Metals can present health hazard if their concentrations exceed allowable limits. Even when the concentrations of metals do not go beyond these limits, there is still a potential for long-term contamination, since heavy metals are known to be accumulative within biological systems [1, 2, 3]. These heavy metals are not eco-friendly and their presence in streams and lakes leads to bioaccumulation in living organisms, causing health

problems in animals, plants and human beings [1, 4]. Heavy metal ions are reported as priority pollutants, due to their mobility in natural water ecosystems and due to their toxicity [5]. Cadmium, zinc, copper, nickel, mercury, and chromium are often detected in industrial wastewaters, which are originated from metal plating, mining activities, smelting, battery manufacture, pesticides, nuclear industry, etc. Nickel salts are commonly used in metal plating and its concentration in industrial wastewater varies between 6 to 12 mg/L, which is above the safe limit (0.15 mg/L) [6]. The chronic toxicity of nickel to humans and the environment is well known and high nickel concentration causes gastrointestinal irritation and lung and bone cancers [7,8].

Among several chemical and physical methods, the adsorption onto activated carbon has been found to be superior to other techniques because of its capability of adsorbing a broad range of different types of adsorbates efficiently and its simplicity of design [9]. However, commercially available activated carbons are still considered expensive [10]. As a result, many researchers have studied cheaper substitutes, which are relatively inexpensive and at the same time endowed with reasonable adsorption capacity. These studies include the use of coal [11], fly ash [8, 12-13],

activated clay [14], palm fruit bunch [15], rice husk [16], hazelnut shell [17], coconut shell [18] and peat [19, 20].

The objective of this study is to evaluate the feasibility of using the adsorbent prepared from *Delonix Regia* pods for the removal of Nickel (II) ion from aqueous solution. The influence of experimental parameter such as contact time, adsorbent dosage and initial Ni (II) ion concentrations were studied. Kinetic and equilibrium aspects of the adsorption process were studied.

Table: 1 Nomenclature

Nomenclature	
C_i	Liquid phase initial concentrations of the adsorbate (mg/L)
C_t	Liquid phase concentrations of the adsorbate at time "t" (mg/L)
C_e	Liquid phase concentrations of the adsorbate at equilibrium (mg/L)
V	Volume of the adsorbate solutions in liter (L)
W	Mass of the adsorbent (g)
q_e	Quantity adsorbed at equilibrium (mg/g)
q_t	Quantity adsorbed at time "t" (mg/g)
T	Time in minutes
Q_e	Amount of solute adsorbed per unit weight of adsorbent (mg/g)
C_e	Equilibrium concentration of solute in the bulk solution (mg/L)
Q_0	Langmuir adsorption efficiency
b	Langmuir adsorption energy
R_L	Langmuir separation factor
C_0	Initial concentration of the adsorbate
K_f and n	Freundlich constants incorporating all factors affecting the adsorption capacity and intensity of adsorption respectively
B_1	Tempkin constant related to heat of sorption (J/mol)
K_T	Tempkin equilibrium binding constant
qD	D-R isotherm theoretical saturation capacity (mg/g)
B	D-R isotherm constant related to the mean free energy
	D-R isotherm polanyi potential
E	D-R isotherm mean free energy of adsorption
R	Gas Constant
T	Temperature (K)
k1	Rate constant of adsorption (l/min)
k2	Second-order constants
h	Initial adsorption rate (mg/g min)
N	Number of data points

Materials and Methods

All the chemicals used for these experiments are of analytical grade. Activated Carbon was prepared from *Mimusops elengi leaf*. $(\text{NH}_4)_2 \cdot \text{Ni} \cdot (\text{SO}_4)_2 \cdot 6\text{H}_2\text{O}$, complexing reagent such as Bromine water, Ammonia and Dimethyl glyoxime purchased from Merck Company.

Preparation of Stock solution

Stock solution 500mg/L of Ni(II) ion was prepared by dissolving 0.3360g of $(\text{NH}_4)_2 \cdot \text{Ni} \cdot (\text{SO}_4)_2 \cdot 6\text{H}_2\text{O}$ in double distilled water. The solution was further diluted to the required concentrations. Before mixing the adsorbent, the pH of each Ni (II) ion solution was adjusted to the required value by 0.1 M NaOH or 0.1 M HCl solution. The Ni(II) ion optical density of the solution measuring at 450 nm.

Preparation of adsorbent using H_2SO_4 :

The dried *Mimusops elengi* leaf powder was carbonized by treating with concentrated sulphuric acid in the weight ratio of 1:1 (w/v). Thus obtained charred masses were heated for 8 hours in a muffle furnace at 400°C to complete carbonization and activation. The resulting carbons were washed with distilled water until a constant pH of the slurry was reached. Then the carbon was dried for four hours at 100°C in a hot air oven. The dried material was ground well to a fine powder and sieved well using the sieves of desired particle size range. The carbon obtained from the *Mimusops elengi* leaf powder was after this chosen as Sulphuric acid Treated *Mimusops elengi* leaf Activated Carbon (SATMEAC).

Physico-chemical Characterization of SATMEAC

The properties of (SATMEAC) were given in Table 2. The total surface areas were calculated using the BET equation ^[21].

Table: 2 Physico-chemical characteristics of SATMEAC

S.No.	Properties	SATMEAC Values
1.	PHzpc	6.0
2.	Particle size, mm	96-110
3.	Surface area (BET), m^2/g	917.1235
4.	Pore volume, cm^3/g	0.3887
5.	Pore size (Pore width), nm	2.8174
6.	Bulk density, g/ml	0.55
7.	Fixed Carbon, %	71.10
8.	Moisture content, %	4.38

Table: 3 Data processing Tools

S. No.	Parameters	Formulae	
1.	Kinetic Models & SSE %	Pseudo First order kinetics (Legergren equation)	$\log (q_e - q_t) = \log q_e - k_1 / 2.303 \times t$
		Pseudo Second order kinetics (Ho equation)	$t/q_t = 1/k_2 \cdot q_e^2 + 1/q_e t$
		The initial adsorption rate h	$h = k_2 q_e^2$
		Sum of error squares	$\text{SSE} (\%) = \frac{[(q_e)_{\text{exp}} - (q_e)_{\text{cal}}]}{N}$
2.	Isotherms	Langmuir Separation factor	$C_e / Q_e = 1/Q_0 b + C_e / Q_0$ $R_L = 1 / (1 + b C_0)$
		Freundlich	$\log Q_e = \log K_f + 1/n \log C_e$
		Tempkin	$q_e = B_1 \ln K_T + B_1 \ln C_e$
		Dubinin – Raduskevich, Polanyi potential Mean free energy of adsorption	$\ln q_e = \ln q_D - B^2$ $= RT \ln (1 + 1/C_e) E = 1 / (2B)^{1/2}$

Batch Adsorption procedure:

Batch adsorption studies were carried out in 250 mL iodine flasks with 50 mL of the working Ni (II) ion solution of different concentrations ranging from 10 mg/L to 20 mg/L. Known amount of adsorbent was added to the solution. The flasks were agitated at a constant speed of 180 rpm. Samples were collected from the flasks at predetermined time intervals for analyzing the residual Ni (II) ion concentration in the solution. The amount of Ni (II) ions adsorbed in milligram per gram of adsorbent was determined by using the following mass balance equation:

$$q_e = (C_i - C_e)V/W$$

Where C_i and C_e are Ni (II) ion concentrations (mg/L) before and after adsorption, respectively, V is the volume of adsorbate in litre and m is the weight of the adsorbent in grams. The percentage of removal of Ni (II) ions was calculated using the following equation:

$$\text{Removal (\%)} = (C_i - C_e)/C_i \times 100$$

Result and Discussion

Effect of Contact time and initial concentration

The effect of contact time on percentage removal of Ni (II) ion for different initial concentration has been shown in figure 1. Adsorption of Ni (II) ion from the solution increases with the time and finally attains equilibrium at 60, 80 and 100 minutes for the initial concentrations 10, 15 and 20 mg/L respectively. The percentage of removal increased with the increase in contact time and decreased with the increase of initial concentration of the Ni (II) ion. However the amount of metal ion adsorbed on the adsorbent increased with the increase of initial concentration of the Ni (II) ion solution [22]. Which is depicted in figure 1 and given in table 4.

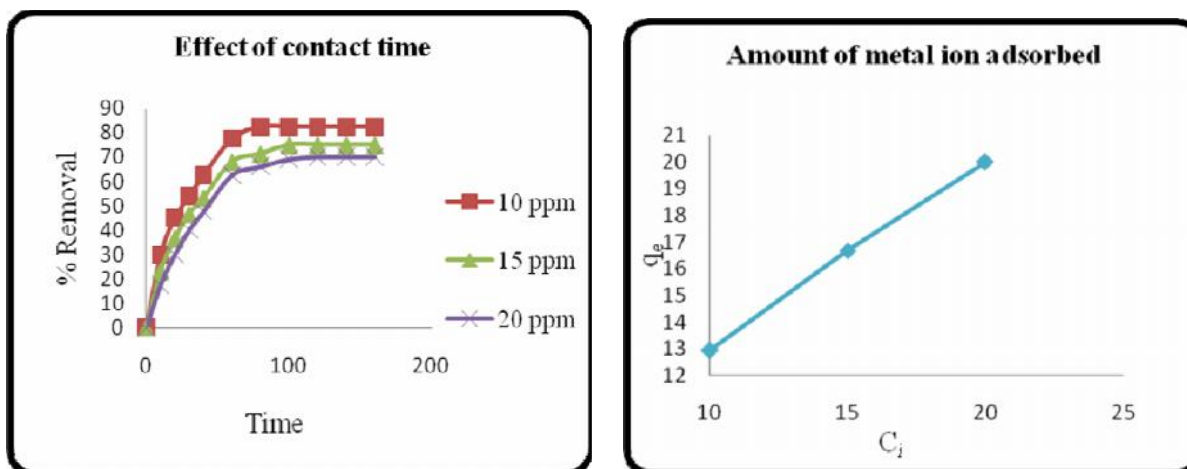


Figure: 1 Effect of contact time & Amount of Ni (II) metal ion adsorbed at equilibrium

Table: 4 Percentage of removal of metal ion and amount of metal ion adsorbed

C_i (mg/L)	% of Removal of metal ion at equilibrium	Adsorbed amount of Ni(II) ion at equilibrium (mg/g)
10	82.55	12.90
15	74.00	16.50
20	70.10	20.10

Effect of pH:

The pH of the adsorbate solutions has been identified as the most important parameter governing sorption of metal ion on different adsorbents. This is partly due to the fact that hydrogen ions themselves are a strong competing sorbate and partly to the fact that the solution pH influences the chemical speciation of

metal ion. The effect of pH on adsorption of Ni(II) ion onto SATMEAC is shown in Figure 2. As shown in this figure, at low pH values, the adsorption percentage is low due to the increase in positive charge density (protons) on the surface sites, resulting in electrostatic repulsion between the Ni(II) ion and positive charge on the surface. Electrostatic repulsion decreases with increasing pH because of the reduction of positive

charge density on the sorption edges. In an alkaline medium, percentage of removal was not good. This may be due to the role of inter ionic attraction between the OH⁻ ion and the metal ion. According to Figure 2, the maximum adsorption of metal ion was found at pH

6.0. Which is the pHzpc of the carbon where the surface of the adsorbent is neutral? Therefore, an optimum pH 6.0 was selected for further experiments [23].

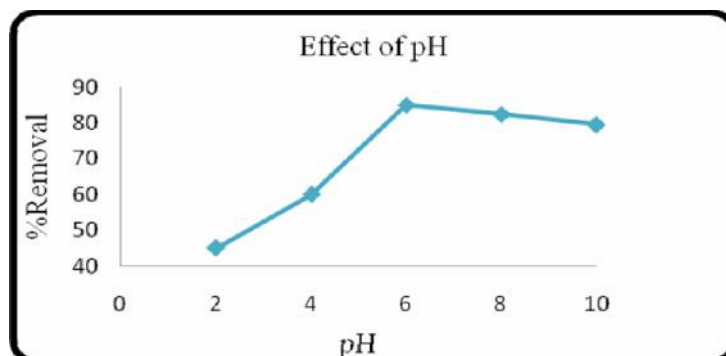


Figure: 2 Effect of pH

Effect of Temperature:

It is well known that temperature plays an important role in the adsorption process. The metal ions removal increase rapidly from 303K, 313K and 323K this result suggests that the experimental

Temperature had a greater effect on the adsorption process implying that the surface coverage increased at higher temperatures. This may be attributed to the increased penetration of metal ions inside micro pores or the creation of new active sites at higher temperatures. This indicates the endothermic nature of the controlled adsorption process. Similar result has been reported in the literature.

Adsorption Kinetics:

The adsorption kinetics shows the evolution of the adsorption capacity through time and it is necessary to identify the types of adsorption mechanism in a given system. Plots of different kinetic models applied were given in the Figure 3 and the kinetic parameters calculated were given in the Table

5. Between the first order and second order, second order kinetic model seems to best describe the above adsorption system as its R² values were very close to unity. Moreover, difference between q_e (cal) and q_e (exp) values of second order is small when compared to first order kinetic model. Statistically it is tested with the tool Sum of error squares (SSE %) [25]. The q_e and SSE % values were given in the Table 5 from which it was concluded that second order kinetic model was more appropriate rather than first order kinetic model.

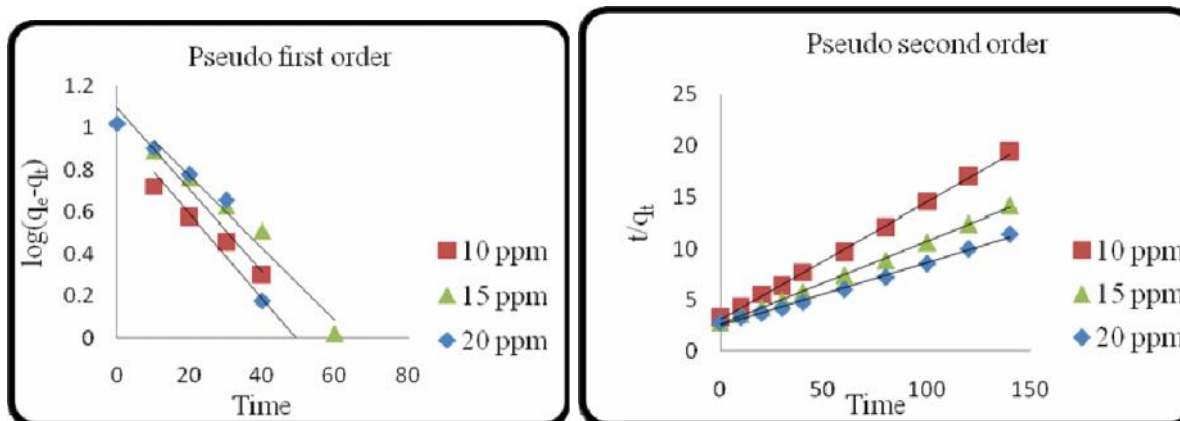


Figure: 3 Pseudo first and second order kinetics

Table: 5 Kinetic parameters

C _i mg/L	Rate constants		q _e (cal) mg/g		q _e (exp) mg/g		q _e		R ²		(SSE %)	
	k ₁ (10 ⁻²) (min ⁻¹)	k ₂ (10 ⁻³) (gmg ⁻¹ min ⁻¹)	First Order	Second order	First order	Second Order	First order	Second Order	First order	Second order	First order	Second Order
10	4.61	5.08	8.35	8.35	9.89	8.75	1.55	0.54	0.955	0.999	1.59	0.75
15	4.54	2.29	11.35	11.35	12.83	12.4	1.58	1.28	0.967	0.999		
20	4.38	1.47	14.0	14.0	12.42	14.39	1.58	0.39	0.881	0.995		

Adsorption Isotherm Studies:

The existence of equilibrium between the liquid and solid phase is well described by adsorption isotherms. Equilibrium data collected at different temperatures were fitted in Langmuir, Freundlich, and Tempkin and Dubinin-Raduskevich adsorption isotherm models [24]. These isotherms are depicted in Figure 4.

R² values of these isotherm plots reveal that Freundlich isotherm well describes the present system that is the possibility of multilayer adsorption. R² value of Dubinin-Raduskevich isotherm is very low. In Dubinin-Raduskevich isotherm, the very low value of the constant 'B' related to the mean free energy of adsorption per mole of the adsorbate and the adsorption is physical in nature. Results of various isotherms are presented in Table 6.

Table: 6 Results of various isotherms plots for the adsorption of Ni(II) ion onto SATMEAC

Isotherm Models	Temp (K)	Parameters and their results			
		Q ₀ (mg/g)	b	R _L	R ²
Langmuir	303	12.06	2.08	0.1767	0.989
	313	15.38	1.86	0.2351	0.989
	323	34.47	1.00	0.4347	0.997
Freundlich		K _f (mg/g ⁻¹)	n		R ²
	303	9.74	2.92		0.999
	313	11.11	2.66		0.999
Tempkin		B ₁ (J/mol)	K _T (L/g)		R ²
	303	458.02	4.53		0.997
	313	395.22	4.48		0.994
Dubinin Raduskevich		q _D (mg/g)	E (kJ/mol)	B × 10 ⁻⁴ (mol ² /J ²)	R ²
	303	19.95	6.86	0.0013	0.934
	313	22.39	5.66	0.0025	0.945
	323	26.91	3.16	0.0038	0.993

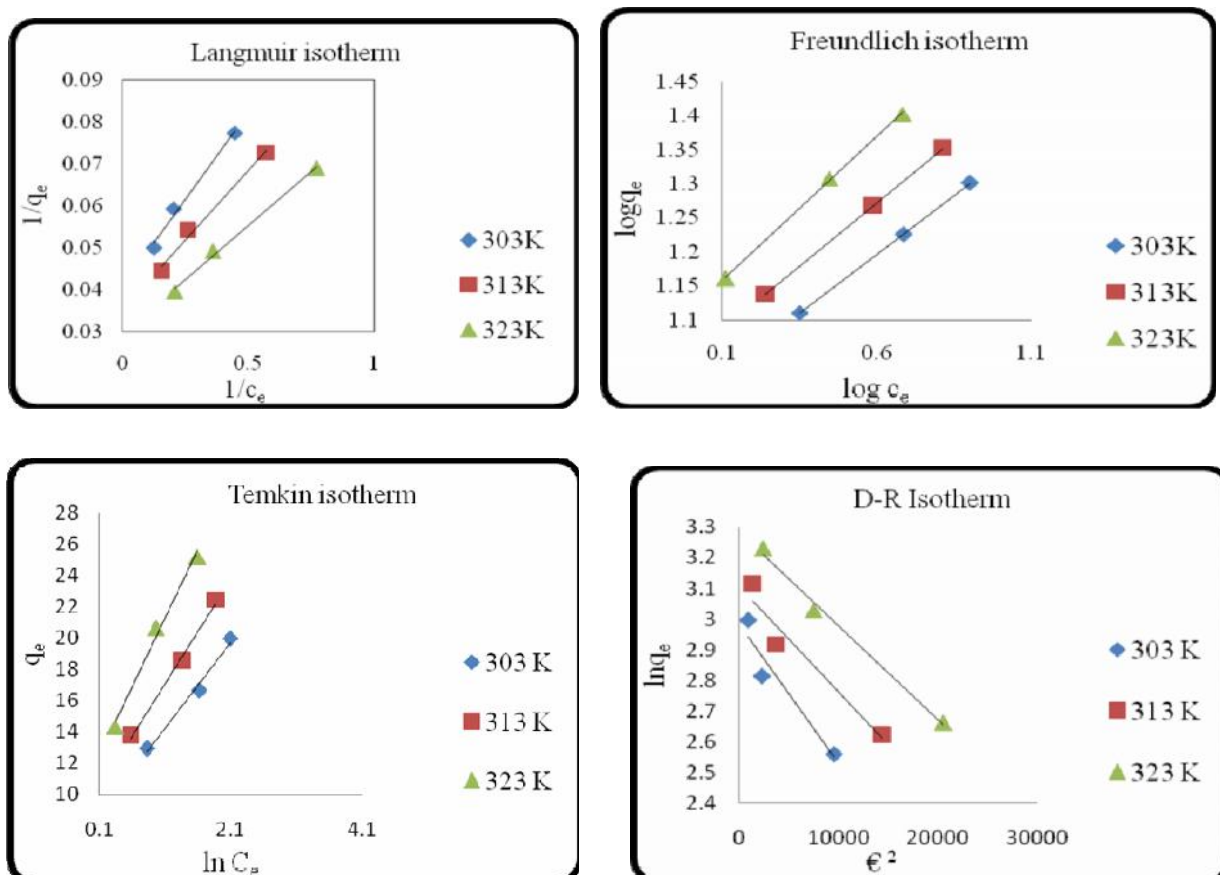


Figure: 4 Langmuir, Freundlich, Temkin and D-R isotherms

Analysis of Isotherm

Langmuir isotherm

In the present study Q_0 value ranges from 12.06 to 34.47, as the temperature increases the monolayer adsorption capacity also found to increase. These kinds of results were obtained in various similar studies [25]. The separation factor R_L values were in between 0 to 1 which indicates the favourable adsorption. R^2 values of isotherm ranged between 0.989 to 0.997.

Freundlich isotherm

The values of n were between 1 and 10 which indicates cooperative adsorption [26]. The R^2 values were close to unity which revealed the good fitting into Freundlich isotherm.

Temkin Isotherm

B_1 -Temkin constant is related to the heat of adsorption. This B_1 value increased from 306.79 to 458.02 as the temperature of adsorption increased. The Temkin parameter K_T value gives an idea about nature of adsorption [27]. In our present study the K_T values ranged from 3.77 to 4.53 which indicate the adsorption is physical nature. The R^2 value was low compared to Freundlich isotherm.

Dubinin-Raduskevich

The activation energy E value ranges from 3.16 to 6.86 and B value from 0.0013 to 0.0038 indicates the physisorption. The R^2 value was very low when compared to other three isotherms.

In general the fitting data in isotherm equation were in the following order: Freundlich > Temkin > Langmuir > Dubinin-Raduskevich.


Conclusion

The present investigation shows that (SATMEAC) is an effective adsorbent for the removal of Ni (II) ion from aqueous solutions. From the kinetic studies, it is observed that adsorption of Ni (II) ion is very rapid in the initial stage and decreases while approaching equilibrium. The equilibrium time increases with initial concentration. The percentage removal of Ni (II) ion increases with the increase in adsorbent dosage and decreases with increase in initial concentration. Experimental results are in good agreement with Freundlich isotherm when compared to other three isotherm models. Adsorption of Ni (II) ion obeys pseudo-second order equation with good correlation.

References

- [1] S.Ong, C.Seng, and P.Lim, Kinetics of adsorption of Cu (II) and Cd (II) from aqueous solution on rice husk and modified rice husk. *Electronic J. Environ. Agric. Food Chem.*, 2007, 6(2), 1764-1774.
- [2] A.J.Marais Barros, S.Prasad, V.Duarte Leite and A.Gouveia Souza, The process of biosorption of heavy metals in bioreactors loaded with sanitary sewage sludge. *Braz.J.Chem. Eng.*, 2006, 23, 153-162.
- [3] E.A.Silva, E.S.Cossich, C.G.Tavares, L.Cardozo, R.Guirardello, Biosorption of binary mixtures of Cr(III) and Cu(II) ions by *Sargassum* sp. *Braz. J. Chem. Eng.*, 2003, 20, 213-227.
- [4] H.I.Ceribas, and U.Yetis, Biosorption of Ni (II) and Pb (II) by *Phanaerochate chryso sporium* from binary metal system-kinetics. *Water SA*, 2001, 27(1), 15-20.
- [5] B.Volesky, and Z.R.Holan, Biosorption of heavy metals. *Biotechnol. Progress*, 1995, 11(3), 235-250.
- [6] R.C.Bansal, and M.Goyal, *Activated Carbon Adsorption*. CRC Press, New York, 2005, 346.
- [7] S.Erdogan, Y.Onal, C.Alanil-Basar, S.Bilmez-Erdemoglu, C.Sarici-ozdemir, E.Koseoglu and G.Lcduygu. Optimization of nickel adsorption from aqueous solution by using activated carbon prepared from waste apricot by chemical activation. *Applied Surface Sci.*, 2005, 252(5), 1324-1331.
- [8] V.K.Gupta, C.K.Jain, I.Ali, M.Sharma, and V.K.Saini, Removal of cadmium and nickel from wastewater using bagasse fly ash-a sugar industry waste. *Water Research*, 2003, 37(16), 4038-4044.
- [9] A.A.Ahmad, B.H.Hameed, and N.Aziz, Adsorption of direct dye on palm ash: Kinetic and equilibrium modeling. *Journal of Hazardous Materials*, 2007, 141(1), 70-76.
- [10] De.S.Chakraborty, S.DasGupta and J.K.Basu. Adsorption study for the removal of a basic dye: Experimental and modeling. *Chemosphere*, 2005, 58(8), 1079-1086.
- [11] S.V.Mohan, N.C.Rao and J.Karthikeyan, Adsorptive removal of direct azo dye from aqueous phase onto coal based sorbents: A kinetic and mechanistic study. *Journal of Hazardous Materials*, 2002, 90(2), 189-204.
- [12] H.Nollet, M.Roels, P.Lutgen, P.V.Meeren and W.Verstraete, Removal of PCBs from wastewater using fly ash. *Chemosphere*, 2003, 53(6), 655-665.
- [13] V.K.Gupta, I.Ali, Removal of lead and chromium from wastewater using bagasse fly ash-sugar industry waste. *Journal of Colloid and Interface Science*, 2004, 271(2), 321-328.
- [14] P.Ricou, I.Lecuyer and P.L.Cloirec, Experimental design methodology applied to adsorption of metallic ions onto fly ash. *Water Research*, 2003, 35(4), 965-976.
- [15] F.C.Wu, R.L.Tseng and R.S.Juang, Kinetics of colour removal by adsorption from water using activated clay. *Environmental Technology*, 2001, 22(6), 721-729.
- [16] M.M.Nassar, The kinetics of basic dye removal using palm fruit bunch. *Adsorption Science and Technology*, 1997, 15(8), 609-617.
- [17] K.S.Low and C.K.Lee, Quaternized rice husk as sorbent for reactive dyes. *Bioresource Technology*, 1997, 61(2), 121-125.
- [18] M.Koby, Removal of Cr (VI) from aqueous solutions by adsorption onto hazelnut shell activated carbon: Kinetic and equilibrium studies. *Bioresource Technology*, 2004, 91(3), 317-321.
- [19] P.A.Brown, S.A.Gill and S.T.Allen, Metal removal from wastewater using peats. *Water Research*, 2000, 34(16), 3907-3916.
- [20] Y.S.Ho and G.McKay, The kinetics of sorption of divalent metal ions onto sphagnum moss flat. *Water Research*, 2000, 34(3), 735-742.
- [21] S.P.Brunauer, H.Emmett, E.Teller, Adsorption of gases in multimolecular layers. *J. Am. Chem. So.* 1938, 60,309-319.
- [22] V.K.Gupta, A.I.Suhas, V.K.Saini. Removal of rhodamine B fast green and methylene blue from waste water using red mud, an aluminum industry waste. *Ind. Eng. Chem. Res.* 2004; 43:1740-1747.
- [23] M.Torab-Mostaedi, H.Ghassabzadeh, Removal of cadmium and nickel From aqueous solution using Expanded perlite. *Brazilian Journal of Chemical Engineering.* 2010, Vol. 27, No. 02, pp. 299 - 308,
- [24] K.Ramesh, A.Rajappa, V.Nandhakumar. Adsorption of hexavalent chromium on tomicrowave assisted zinc chloride activated carbon prepared from delonix regia pods. *Inter. Jour. Res Chem. Envirt.* 2014, Vol 4 Issue (3). Pages: 3-9.

- [25] I.Langmuir, the Journal of the American Chemical Society, 1918, 40, 1361-1403.
[26] K.Fytianos, E.Voudrias, E.Kokkalis.Chemosphere, 2000, 40-43.
[27] M.J.Temkin, V.Pyzhev, Acta physiochem USSR, 1940, 12-217

Access this Article in Online	
	Website: www.ijcrops.com
	Subject: Chemistry
Quick Response Code	
DOI: 10.22192/ijcrops.2018.05.04.001	

How to cite this article:

A. Elavarasan and V. Nandhakumar. (2018). Adsorption of Nickel (II) Ion from Aqueous Solution onto sulphuric acid Activated Carbon Prepared from *Mimusops elengi* Leaves. Int. J. Curr. Res. Chem. Pharm. Sci. 5(4): 1-9.

DOI: <http://dx.doi.org/10.22192/ijcrops.2018.05.04.001>