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Evaluation of Treatment Strategies by Adsorption for Lead Removal from Aqueous Solution

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ABSTRACT

The contamination due to heavy metals from industrial waste water is most important issues for industrialized area. In this work removal of lead from aqueous solution was investigated using granular activated carbon modified with 1,3,5-Triazine-2,4,6-triamine. The adsorption of lead using GAC filtrasorb 300 (GAC F-300) and filtrasorb 816 (GAC F-816) were carried out at constant temperature $25 + 1^{\circ}$ C at about 500 rpm. The adsorption data were well analysed for Freundlich, Langmuir and Temkin adsorption isotherm models.

Keywords: Adsorption, Lead, Granular activated carbon (GAC), Filtrasorb 300 (F-300), Filtrasorb 816 (F-816), 1,3,5-Triazine-2,4,6-triamine

INTRODUCTION

Now days an environment and human being facing a serious problem due to the toxicity, persistence and bioaccumulation affinities of heavy metals in water bodies. Metals and their compounds are extremely important to the industrial and technological development in developed countries due to numerous of applications for commercial uses [1, 2]. Lead is one of the heavy metals found in industrial wastewater and its discharge into water bodies poses an adverse effect on aquatic and terrestrial lives. Lead poisoning causes severe damage to the nervous system, kidney, reproductive system, brain, liver cause illness or death. Severe exposure to lead has been connected with stillbirths, abortion, sterility and neonatal death [3-5].

Lead is naturally occur as an element in insoluble form in the earth crust and biologically inoffensive forms [6]. There are number of methods employed for the treatment of industrial effluents containing Lead. Some of the important methods are chemical precipitation, ion exchange, electrodialysis and carbon adsorption use for water treatment [7-19]. Many advanced techniques are available for removal of heavy metal involves high investments, which are not suitable for small scale industries, which discard comparatively low volumes of wastewater. Adsorption is an effective separation and purification technique used in industry especially in water and waste water treatments [20]. Treatable amount of lead wastes are growing need in the industrial sector to try and find solution to remove this metal from waste waters using granular activated carbon [21-33]. Lead is considered as a major threat for various human diseases once it goes beyond the permissible limit recommended by the World Health Organisation (WHO) (3-10 μ g/L) in drinking water [34].

MATERIALS AND METHODS

The Granular Activated Carbon filtrasorb 300 (F-300) and filtrasorb 816 (F-816) gifted by M/s Calgon Carbon Corporation Ltd Pittusberg, USA were selected as adsorbent for adsorption study. These adsorbents were first sieved to obtain the carbon particles of desired size. All the particles were collected in clean petridish which used as adsorbent for overall studies. The sieved GAC particles were completely washed with hot double distilled water until the supernatant was free of dirt particles and then kept in an vacuum oven at a temperature of 105°C for 5 hours for dry. It was cooled in a desiccators containing anhydrous CaCl, to verify complete removal of moisture from the pores on

surface of GAC. A synthetic solution of lead ions was prepared by dissolving required quantity of $Pb(NO_3)_2$ (S.D. Fine Chem. Limited) in freshly prepared double distilled water and all working solutions were prepared by dilution with double distilled water.

Each solution was treated with 2 ml aqueous solution of alizarin Red (S) and 1 ml NaOH (2M). The optical densities of all experimental solutions were estimated using Chemito Spectrascan UV 2700 Double beam UV Visible spectrophotometer at λ_{max} 485 nm. A Beer's law standard curve was obtained by plotting optical densities versus concentration of lead. The equation computed was used to estimate residual concentration of Pb²⁺ ions [35]. All the reagents used in this work were of AR grade. A sample of 1,3,5-Triazine-2,4,6-triamine(S.D. Fine Chem. Limited) was purified and recrystallized by standard method. The melting point of 1,3,5-Triazine-2,4,6-triamine was found to be 344.5°C compared with literature value 345°C [36]. The sample was also characterized through determination of molecular weight by the technique of pH titration against standard NaOH. To study the adsorption isotherm of Lead ions, 200 ml of 0.001 M solution of 1,3,5-Triazine-2,4,6-triamine agitated with 0.5 g of GAC in reagent bottles of 300 ml capacity. It was then shaken for about five hours using Teflon bladed stirrer at about 500 rpm. After five hours the solution was decanted and the carbon particles were washed completely with double distilled water which recognized as loaded carbon. This loaded carbon was then transferred to same reagent bottle and then 200 ml of lead solution of pH = 6 were added to it. The contents were agitated for 5 hours in a thermostat at a constant temperature of $25 \pm 1^{\circ}$ C. The initial and final concentrations of lead ions were estimated by putting the values of absorbance in the equation obtained from Beer's law curve. The experiments were repeated twice to test reproducible results.

RESULTS AND DISCUSSION

The adsorption data of Pb^{2+} on GAC were analyzed in the light of Freundlich, Langmuir and Temkin models. The relationship between the liquid phase concentration and surface concentration of adsorbate at equilibrium was obtained to describe adsorption isotherm.

The quantity of Lead on the modified GAC was estimated using the equation

 $q = (C - C) \times V/W$

Where,

q = Concentration of Lead ion on the modified GAC (mg/millimoles)

C_=Initial concentration of Lead ion (mg/L)

C_=Final concentration of Lead ion (mg/L)

W=Millimoles of the ligand actually present on GAC (0.5 g).

V=Volume of solution in liters,

The adsorption isotherms of modified F-300 and F-816 GAC obtained by plotting q, versus C, and shown in Figure 1.

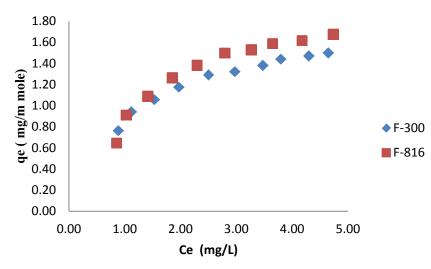


Figure 1: Adsorption Isotherm System: GAC-1,3,5-Triazine-2,4,6-triamine-Pb²⁺.

(1)

The Langmuir equation could be expressed as	
$q_e = Q^o b \times C_e / (1 + bC_e)$	(2)
Where,	
Q° = Amount adsorbed per unit weight of the GAC to form monolayer.	
b = Langmuir constant.	
Rearranging equation (2)	
$1/q_e = 1/Q^o b \times 1/C_e + 1/Q^o$	(3)
A plot of $1/q_e$ versus 1/Ce was found to be fairly linear indicate the validity of isotherm.	
The Freundlich equation express as	
$q_e = K.C_e^{1/n}$	(4)
Where, k and l/n are Freundlich constants determined experimentally. Using equation (4)	
$\log q_e = \log K + 1/n \log C_e$	(5)

The linearity in the plot of log q against log C showed validity of Freundlich equation over a range of concentrations.

Langmuir and Freundlich isotherms for F-300 and F-816 are illustrates in **Figure 2 and 3**. The plots of $1/q_e$ against $1/C_e$ were found to be linear indicating the validity of Langmuir model. The Langmuir constants relating to the sorption capacity (Q^o) and adsorption energy (b) were determined (**Table 1**).

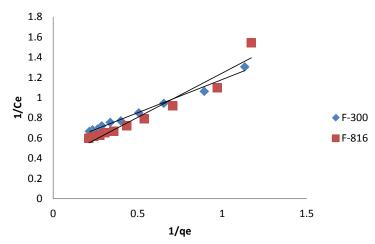


Figure 2: Langmuir Adsorption Isotherm System: GAC-1,3,5-Triazine-2,4,6-triamine-Pb²⁺.

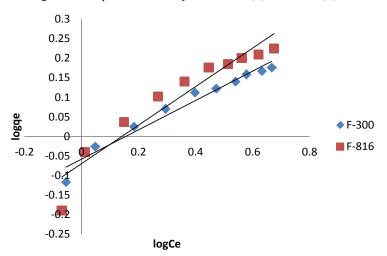


Figure 3: Freundlich Adsorption Isotherm System: GAC-1,3,5-Triazine-2,4,6-triamine-Pb²⁺.

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(6)

(7)

Sr.	Adsorption System	Langmuir Constant			Freundlich Constant		
No.		Q٥	b	R ²	K	1/n	R ²
1	F-300-1,3,5-Triazine-2,4,6-triamine-Pb ²⁺	1.9173	0.7927	0.9882	0.8744	0.3752	0.9636
2	F-816-1,3,5-Triazine-2,4,6-triamine-Pb ²⁺	2.7236	0.4193	0.9374	0.8515	0.4924	0.9090

Table 1: Equilibrium Isotherm Constants For Langmuir and Freundlich model.

 Q° obtained from graph was used to estimate the surface area occupied by lead ion on GAC. The surface area of the GAC through Lead adsorption can then be represented as

S'=Na.Qº.A

Where,

S'=Surface area of adsorbent (cm²/g)

A=Cross-sectional area of the adsorbent molecule (cm²).

Na=Avogadro number

Determination of value of S' needed the value of A the surface area occupied by a single Lead ion. The values of A were calculated using the expression given by Brunauer and Emmet.

 $A=4 \times 0.866 [M/4\sqrt{2.Na.d}]^{2/3}$

Where,

M=Atomic weight of the lead

Na=The Avogadro number

d=The density of the lead, [37]

The values of S' obtained from Q° and S obtained from $q_{e max}$ are reported in Table 2.

Sr. No.	System	A (cm ²)	q _{e max} (mg/m. mol.)	S (cm²/gm)	Q ⁰	S'(cm²/ gm)
1	F-300-1,3,5-Triazine-2,4,6-triamine-Pb ²⁺	5.4225 × 10 ⁻¹⁶	1.5000	1.8381 × 10 ³	1.9173	2.3494 x10 ³
2	F-816-1,3,5-Triazine-2,4,6-triamine-Pb ²⁺	5.4225 × 10 ⁻¹⁶	1.6765	2.0544×10^{3}	2.7236	3.3376 x10 ³

Table 2: Values of S, A, Q⁰ and S' for a system GAC-1,3,5-Triazine-2,4,6-triamine-Pb²⁺.

Temkin model considers the effect of indirect adsorbate-adsobent interactions on adsorption and suggests that the heat of adsorption of all the layer would decrease linearly with coverage due to adsorbate-adsobent interactions. It assumes that the decrease in the heat of adsorption is linear rather than logarithmic as stated in Freundlich expression. The linearised form of Temkin equation is

 $q_e = RT/b_T \ln K_T + RT/b_T \ln C_e$

Where,

K_r=equilibrium binding constant (L/mg)

b=Temkin constant related to the heat of adsorption (kJ/mol)

B=RT/b_T

Figure 4 illustrates the plot of Temkin isotherms for F-300 and F-816. Temkin constants K_T and b_T are calculated from plot of q_a versus ln C_a given in **Table 3**.

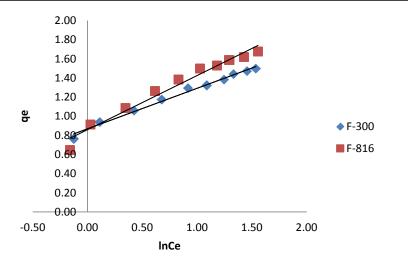


Figure 4: Temkin Adsorption Isotherm System: GAC-1,3,5-Triazine-2,4,6-triamine-Pb²⁺.

Sr. No.	Adsorption System	K _r	b _T	R ²
1	F-300-1,3,5-Triazine-2,4,6-triamine-Pb ²⁺	7.7673	5849.4436	0.9889
2	F-816-1,3,5-Triazine-2,4,6-triamine-Pb ²⁺	4.576739	4382.762	0.9680

Table 3: Equilibrium Isotherm Constants For Temkin model.

CONCLUSION

This study examined the efficiency of adsorbent GAC in the removal of Pb^{2+} ions from aqueous phase in presence of 1,3,5-Triazine-2,4,6-triamine. The main advantages of the adsorption study include cost effectiveness, simplicity and offers flexibility in design. From the plot of q_e against C_e , it is observed that initially C_e increases with q_e but at the saturation level q_e tends to be constant which indicates monolayer formation of Lead ion on the pores site of surface of adsorbent GAC. The experimental data seen to be of the favourable type and subjected for adherence to Langmuir and Temkin model better than by the Freundlich model. In adsorption study F-816 loaded with 1,3,5-Triazine-2,4,6-triamine-adsorbs lead to a remarkable extent as compared to F-300. This is probably due to presence of large active sites available on GAC surface and its porous nature.

REFERENCES

- [1] Edokpayi JN, Odiyo JO, Msagati TAM, Popoola EO (2015) A novel approach for the removal of lead (II) ion from wastewater using Mucilaginous leaves of Diceriocaryum eriocarpum plant. *Sustainability* 7: 1402-1404.
- [2] Mwangi IW, Ngila JC, Okonkwo JO (2012) A comparative study of modified and unmodified maize tassels for the removal of selected trace metals in contaminated water. *Toxicol Environ Chem* 94: 20-39.
- [3] Gercel O, Gerçel HF (2007) Adsorption of lead (II) ions from aqueous solutions by activated carbon prepared from biomass plant material of Euphorbia rigida. *Chem Eng J* 132: 289-297.
- [4] Mahmoud ME, Osman MM, Hafez OF, Hegazi AH, Elmelegy E (2010) Removal and preconcentration of lead (II) and other heavy metals from water by alumina adsorbents developed by surface-adsorbed-dithizone. *Desalination* 25: 123-130.
- [5] Imamoglu M, Tekir O (2008) Removal of copper (II) and lead (II) ions from aqueous solutions by adsorption on activated carbon from a new precursor hazelnut husks. *Desalination* 228: 108-113.
- [6] Ogunleye OO, Ajala MA, Agarry SE (2014) Evaluation of biosorptive capacity of banana (Musa paradisiaca) stalk for Lead (II) removal from aqueous solution. *J Environ Prot* 5: 1451-1465.
- [7] Juttner K, Galla U, Schmieder H (2000) Electrochemical approaches to environmental problems in the process industry. *Electrochimica Acta* 45: 2575-2594.

- [8] Bose P, Bose MM A, Kumar S (2002) Critical evaluation of treatment strategies involving adsorption and chelation for wastewater containing copper, zinc, and cyanide. Adv Environ Res 7: 179-195.
- [9] Wingenfelder U, Hansen C, Furrer G, Schulin R (2005) Removal of heavy metals from mine water by natural zeolites. *Environ Sci Technol* 39: 4606-4613.
- [10] Shamma NK, Wang LK and Hung YT (2004) Coagulation and flocculation in Physicochemical Treatment Processes. Humana Press, New Jersey, USA, pp: 103-140.
- [11] Semerjian L, Ayoub GM (2003) High-pH-magnesium coagulation–flocculation in wastewater treatment. *Adv Environ Res* 7: 389-403.
- [12] Ayoub GM, Semerjian L, Acra A, Fadel M, Koopman B (2001) Heavy metal removal by coagulation with seawater liquid bittern. *J Environ Eng* 127: 196-202.
- [13] Metcalf, Eddy (2003) Wastewater Engineering: Treatment and Reuse. McGraw Hill International Edn, New York, USA, pp: 478-483.
- [14] Eckenfelder WW (1996) Industrial Water Pollution Control. McGraw-Hill Companies. pp: 451-457.
- [15] Jokela P, Keskitalo P (1999) Plywood mill water system closure by dissolved air flotation treatment. Water Sci. Technol 40: 33-42.
- [16] Matis KA Zouboulis AI, Lazaridis NK and HancockI C (2003) Sorptive flotation for metal ions recovery. Int J Miner Process 70: 99-108.
- [17] Wojtowicz A, Jarosinski A (1996) Removal of chromium Cr(III) on smectite ion exchange column, 3rd International Conference on Environment and Mineral Processing Ostrava, pp: 217-277.
- [18] Corupeioglu MO, Huang CP (1987) The adsorption of heavy metals on to hydrous activated carbon. *J Water Res* 21: 1031-1044.
- [19] Rivera Utrilla J, Garcia MAF (1987) Study of cobalt adsorption from aqueous solution on activated carbons from almond shells. *Carbon* 25: 645.
- [20] Guo Y, Qi J, Yang S, Yu K, Wang Z (2003) Adsorption of Cr(VI) on micro and mesoporous rice husk based active carbon. *Mater Chem Phys* 78: 132-137.
- [21] Khan S, Nigam SK, Dwivedi HP, Singh PK (2004) Oxidative degradation of methyl ethyl ketone by n-chlorosaccharin. Asian J Chem 16(2): 751-754.
- [22] Olusegun KA, Otaighe JOE (2008) Adsorption Behaviour of 1-phenyl-3-methylpyrazol-5-one on Mild Steel from HCI Solution. *Int J Electrochem Sci* 3: 191-198.
- [23] Islam M, Patel RK, Mater JH (2007) Evaluation of removal efficiency of fluoride from aqueous solution using quick lime. J Hazard Mater 143: 303-310.
- [24] Daifullah AAM, Yakout SM, Yang XJ, Fane AG, MacNaughton S (2001) Removal and recovery of heavy metals from wastewater by supported liquid membranes. *Water Sci Technol* 43: 341-348.
- [25] Babel S, Kurniawan TA (2003) Lowcost adsorbents for heavy metals uptake from contaminated water: A review. J Hazard Mater B97: 219-243.
- [26] Abudaia JA, Sulyman MO, Elazaby KY, Ben-Ali SM (2013) Adsorption of Pb (II) and Cu (II) from aqueous solution onto activated carbon prepared from dates stones. *IJEST* 4.
- [27] Paajanen A, Lehto J, Sataapakka T, Morneau JP (1997) Sorption of cobalt on activated carbons from aqueous solutions. *Sep Sci Technol* 32: 813.
- [28] Netzer A and Hughes DE (1984) Adsorption of copper leas and cobalt by activated carbon. J Water Res 18: 927-933.
- [29] Elreefy SA (2007) Adsorption of fluoride in aqueous solutions using KMnO₄-modified activated carbon derived from steam pyrolysis of rice straw. *J Hazardous Mater* 62: 1-32.
- [30] Meena AK, Rajagopal C, Kiran CR, Mishra GK (2010) Removal of Heavy metal ions from aqueous solutions using chemically (Na₂S) treated granular activated carbon as an adsorbent. *J SciInd Res* 69: 449-453.

- [31] Meshram YK, Khati NT, Khope RU (2014) Evaluation of Adsorptive Capacity of Bioadsorbent in Removal of Congo Red from Aqueous Solution. *Der Chem Sin* 5: 25-29.
- [32] Gawande NJ, Chaudhari AR, Khope RU (2012) Influence of surface characteristics of adsorbent and adsorbate on competitive adsorption equilibrium. *Adv Appl Sci Res* 3: 1836-1841.
- [33] Gunjate JK (2016) Selective Adsorption of Cobalt In Aqueous Solution Using Chemically Modified Activated Carbon. IOSR-JESTFT 10: 161-165.
- [34] Needleman HL (1999) History of lead poisoning in the world. The George Foundation, Bangalore, India, pp: 17-25.
- [35] Alsamarrai KF (2011) Spectrophotometric assay of lead in human hair samples by using alizarin red (S) in samarra area. *J of university of Anbar for pure science* 5: 5-12.
- [36] Lide DR (1997) CRC Handbook of Chemistry and Physics. 78th Edn., Boca Raton, FL, CRC Press, pp: 3-323.
- [37] Wuana RA, Okieimen FE (2011) Heavy metals in contaminated soils: A review of sources, chemistry, risks and best available strategies for remediation. *ISRN Technology* 20.