

Batch Adsorption Technique for the Removal of Janus Green B Dye from Industrial Waste Water by Using Walnut Kernel Shell as Adsorbent

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Abstract

Batch experiments were carried out for the adsorption of Janus Green B (JGB) onto walnut kernel shell particle. walnut kernel shell treated with burn in 300 C⁰ was tested as a low cost adsorbent for the removal of Janus Green B from aqueous solution. The effects of various experimental parameters, such as adsorbent dosage and particle size, initial dye concentration, pH and contact time, were investigated in batch mode. An amount of 1.5 g/l of (WKSC) could remove 99.36 % of the dye from an aqueous solution of 50 ppm with the agitation time 150 min and particle size 75 mm and with acid medium (pH=2). The well known Langmuir and Freundlich isotherm models were applied for the equilibrium adsorption data and the various isotherm parameters were evaluated. The results indicate that activated walnut kernel shell carbon (WKSC) could be employed as a low cost alternative to commercial activated carbon in wastewater treatment for the removal of color and dyes.

Keywords

adsorption capacity, walnut kernel shell, equilibrium isotherm models, Janus Green B.

1.Introduction:-

The wastewater treatment for long time has been a main problem of the textile industry. Dyes are widely used in industries such as textiles, rubber, paper, plastics, cosmetics, etc. to color their products. Pollution of water due to the discharge of effluents from dyeing industries affects the environment due to its toxicity these effluents contain many harmful chemical that pose serious problems to human beings and aquatic life [1][2].

Due to their molecular structure, dyes are resistant to light, heat. The common method have been used for dye removal from wastewater include biological methods (anaerobic treatment) and physicochemical methods such as coagulation, electro coagulation, floating, filtration, ion exchange, membrane filtration and advanced oxidation[3-7]. However, many of these technologies are expensive, especially when they are used for treatment of large wastewater streams. Consequently, adsorption methods using low cost adsorbents have the most potential for application in industrial wastewater treatment, because of their efficiency is proven in the removal

of organic and mineral pollutants and economic considerations[8][9] . Natural materials that are available in abundance, or certain waste products from industrial or agricultural operations, may have great potential as an inexpensive sorbents. Due to their low cost, after these materials have been expended, they can be disposed of without expensive regeneration. The abundance and availability of agricultural by products make them good sources of raw materials for activated carbons. Some of the materials which are used for the preparation of activated carbon in the recent past are, oil palm shell[10], almond shell[11], yam peels[12], coconut shell[13][14], coconut coir[15], pistachio shells[16], hazelnut shell[17], walnut shell[18], palm shell[19], apple pulp[20], chickpea husks[21], rice husk[22], Banana shells[23].

This study aimed at evaluating the adsorbent potential of walnut kernel shell as a sorbent for the removal of dyes, using Janus Green B (JGB) as a dye model.

2.Experimental

2.1. Materials and Methods

2.1.1. Preparation of Sorbent

walnut kernel shell(WKS) was obtained from local pistachio mills and was washed several times with tap water followed by filtration. The cleaned walnut kernel shells was oven burning completely at 300 C^0 , then cooled, The Activated carbon (WKSC) used in this study was washed with distilled water to remove water soluble materials present in the carbon prior to the adsorption study and then sieved to (0.075-0.85) mm size which was used without further treatment. Figure(1(a1-2, b)) shows the walnut kernel carbon before and after burn it.

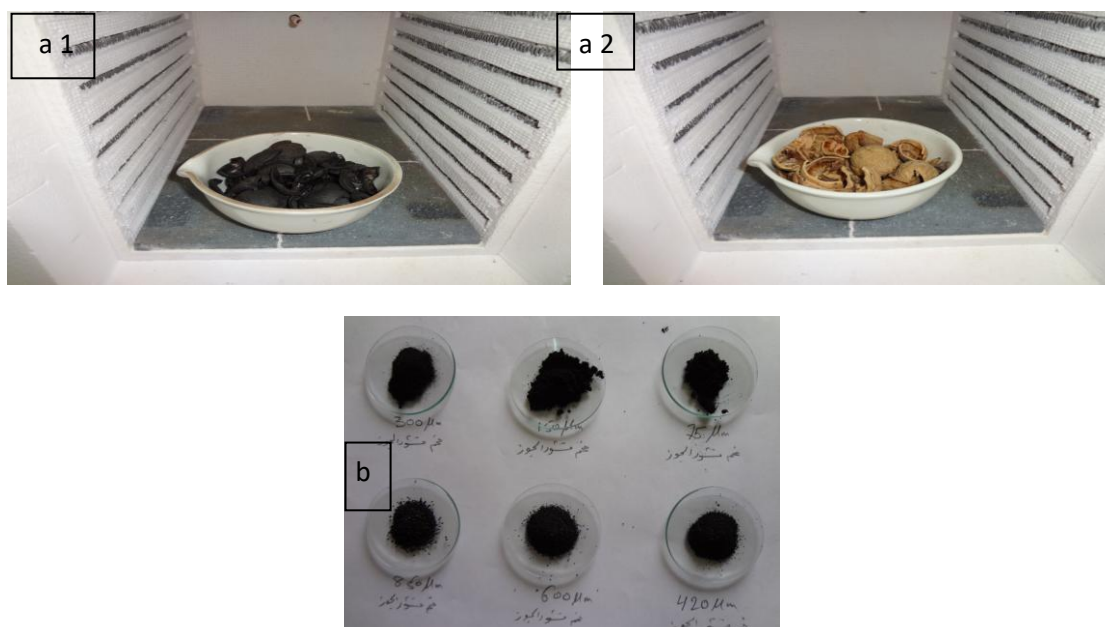


Fig.(1):Iraqi walnut kernel shells used in this study, (a-1)The raw walnut kernel shells before bur

(a-2) The raw walnut kernel shells after burn, (b)-Granular walnut kernel shells

carbon

2.1.2. Preparation of Adsorbate

Preparation of dye solution

Janus Green B (JGB) dye is [Synonyms: 3-diethylamino-7-(4-dimethylaminophenylazo)-5-phenylphenaziniumchloride, 3-(diethylamino)-7-((p-(dimethylamino)phenyl)azo)-5-phenylphenazinium chloride] . The chemical structure of dye used in this study is described in Figure(2).

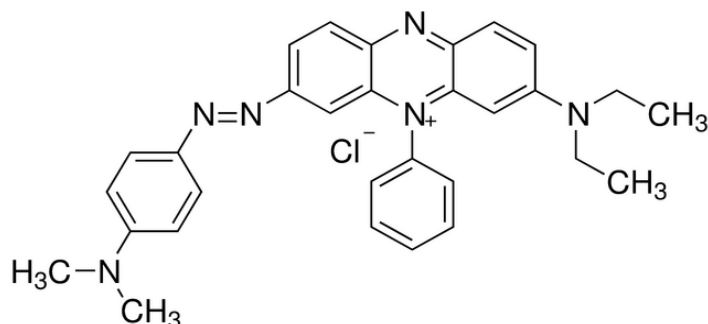


Fig.(2): The chemical structure of Janus Green B dye studied in this work

The physiochemical properties of the JGB dye can be shown by Table (1)[24]

Table (1): Physiochemical properties of Janus Green B dye

Parameter	Value
Molecular formula	$C_{30}H_{31}N_6Cl$
Molecular weight	511.07g/mol
C.I. Name	11050
Absorption maxima	611nm
Nature	Cationic dye

An accurately weighed quantity of the dye was dissolved in distilled a stock solution of the dye was prepared by dissolving 1 gram of dye in 1000 ml distilled water to make a stock solution of 1000(mg / L) . The experimental solution was prepared by diluting definite volume of the stock solution to get the desired concentration .

For absorbance measurements a UV - VIS spectrophotometer (UV / VIS – 1650 PC

SHIMATZU) was employed .

The maximum wavelength λ max. for JGB was measured at 611 nm . Concentrations during experimental work were determined from a standard calibration curve .

The pH of each solution was adjusted with 0.1 M HCL or NaOH using pH –meter to its effective adsorption pH value.

Batch adsorption studies

Adsorption experiments were carried out by agitating 0.5 to 3 g of adsorbent with a dsorbate solution of 50 ppm concentration at pH from 2.0,7.0 to 12, at temperature room in a shaker for 20,30 and 180 min. .The pH was measured using pH meter. The pH of the solutions was adjusted by means of 0.1M HCl and 0.1M NaOH solutions. The samples were withdrawn from the shaker at predetermined time intervals.

The concentration of final sample is measured by spectrophotometric determination. The amount of Janus Green B dye JGB adsorbed was calculated from the following equation:

$$q_e = \frac{V}{W} (C_0 - C_e) \dots\dots\dots(1)$$

Where q_e is the amount of dye adsorbed per unit weight of activated walnut kernel shell (mg/g); C_0 the initial concentration of (JGB) (ppm); C_e the concentration of (JGB) in solution at equilibrium time (ppm); V the solution volume (l); W is the activated carbon dosage (g).

The adsorption behaviors of the samples were studied by evaluating the percentage removal efficiency of (JGB) from the relation

$$\text{Removal efficiency (\%)} = \frac{C_0 - C_e}{C_0} * 100\% \dots\dots\dots(2)$$

Where C_0 is the initial concentration of (JGB), C_e is the solution concentration after adsorption at any time. Equilibrium studies give the capacity of the adsorbent

A desorption isotherm models:-

Langmuir and Freundlich model:-

The analysis of isotherm data is useful for design purpose. In present study the equilibrium data were treated by Langmuir and Freundlich isotherms. The Langmuir isotherm can be represented by the following equation[25].

$$q_e = \frac{q_m K_a C_e}{1 + K_a C_e} \dots\dots\dots (3)$$

Where, q_e is the amount adsorbed per unit mass of sorbent at equilibrium (mg/g), q_m is the maximum adsorption capacity (mg/g), C_e is the equilibrium dye concentration (mg/L) and K_a is the adsorption equilibrium constant. The plot of C_e/q_e versus C_e (eq .4) is linear which show that the adsorption of dye onto walnut kernel shell carbon follows Langmuir isotherm model .

$$\frac{C_e}{q_e} = \frac{1}{q_m K_a} + \frac{C_e}{q_m} \dots\dots\dots (4)$$

The essential characteristics of Langmuir isotherm can be express by a dimensional constant called equilibrium parameter, R_L [26] that is defined by:

$$R_L = \frac{1}{1 + bC_o} \dots\dots\dots (5)$$

where, b is the Langmuir constant and C_o is the initial concentration. The value of R_L indicates the shape of the isotherm to be either un favorable ($R_L > 1$), linear ($R_L = 1$), favorable ($0 < R_L < 1$) or irreversible ($R_L = 0$).

The Freundlich isotherm was also applied for the adsorption of dye by walnut kernel shell carbon [27].

$$\log q_e = (1 / n) \log C_e + \log k_f \dots\dots\dots(6)$$

where, q_e is the amount adsorbed per unit mass of adsorbent at equilibrium (mg/g), C_e is the equilibrium dye concentration of the solution (mg/L). k_f and n are the Freundlich constants, n gives an indication of the favorability and k_f [mg/g(L/mg)^{1/n}], The values of K_f and n can be obtained from the plate of $\log q_e$ versus $\log C_e$ and they equal to the intercept and slop of the plate respectively. The value of n lies between 2 and 10, which implies good adsorption .

Result and Discussion:

Adsorption of dye:-

The adsorption of dye were investigated in the study using different parameters such as adsorbent dosage, contact time, pH, initial dye concentration and particles size of adsorbent.

Effect of adsorbent dosage:

The effect of adsorbent dose was also investigated for the removal of dyes from aqueous solution. The experiments were carried out with adsorbent dose varied from (0.5 - 3) gm with keeping other parameters are constant (initial concentration of 50 mg/l and 0.3mm particle size). The percentage removal of dye was found (31.42 – 99.15)% Figure (3a).

The increase in removal of dyes with adsorbent dose due to the introduction of more binding sites for adsorption. Similar results have been reported by the other

investigators[28-29] .

However, the adsorption capacity showed a decreasing trend with increasing adsorbent dosage. The amount of JGB adsorbed per unit weight of adsorbent decreased with increase in adsorbent dosage (Figure 3b). The reduction in the value of adsorption capacity (q_e) attributed to make a large number of sites available for a fixed dye concentration[30], These sites remaining unsaturated during the adsorption process.

By this study, it was observed that the economical dose with good removal occur at the dose of 1.5 g / 100 ml for activated, walnut kernel shell carbon (WKSC) and that is 90.82 % .

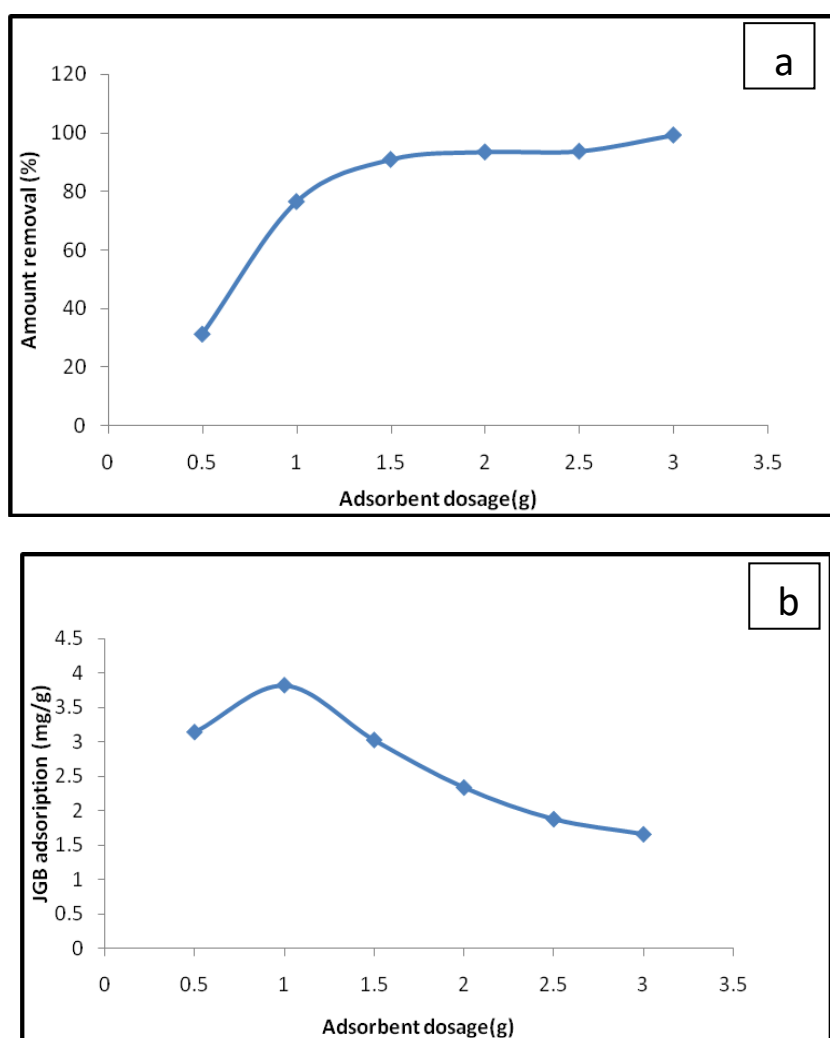


Figure (3) Effect of adsorbent dosage in the removal of JGB by (WKSC)
(a) Adsorption percentage. (b) Amount of dye adsorbed(mg/g).

Effect of contact time :-

The experiments were carried out by taking 100 ml sample of dye (concentration 50 mg/L) in conical flasks and treated with 1.5 gm dosage of adsorbent with several time (20, 30, 45, 60, 80, 120, 150, 180). The variation in percent removal of dye with the elapsed time has been shown in Figure (4). It is evident from the figure that WKSC treatment resulted in 81.18% removal of Janus Green B dye in first 20 min., which increased up to 92.78% in 120-180 min. . It is due to saturation of active sites which do not allow further adsorption to take place [31,32].

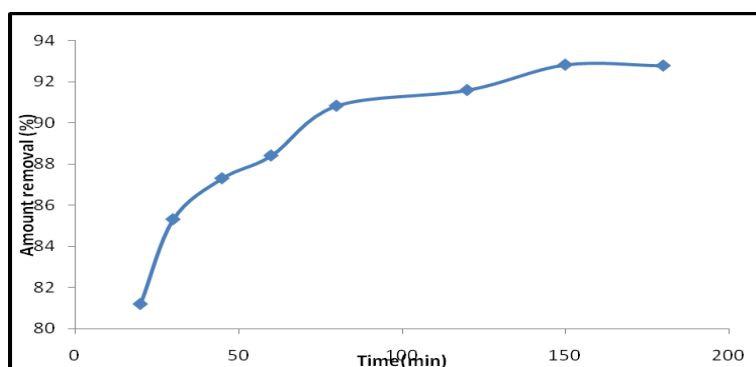
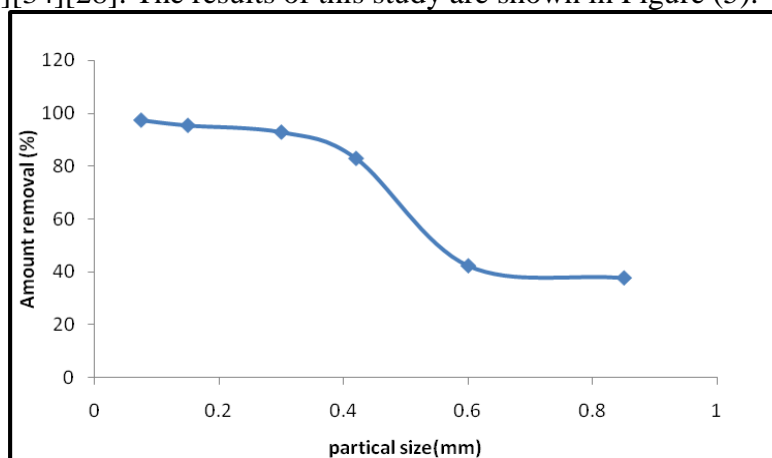


Figure (4) : Effect of contact time for adsorption of (JGB) onto (WKSC) at initial dye concentration of 50 mg/l, adsorbent dose of 1.5 g, and particle size of 0.3mm

Effect of Particle size of adsorbent :-

The effect of Particle size of adsorbent (WKSC) on adsorption of Janus Green B dye has been studied on walnut kernel shell particle of varying size (0.075, 0.15, 0.3, 0.42, 0.6, 0.85) mm. The experimental data show that amount of Janus Green B adsorbed decreases with increase in Particle size of the adsorbent. This indicates that the smaller the (WKSC) particle size for a given mass of (WKSC), the more surface area is available and as a consequence the greater the number of binding sites available [33][34][28]. The results of this study are shown in Figure (5).

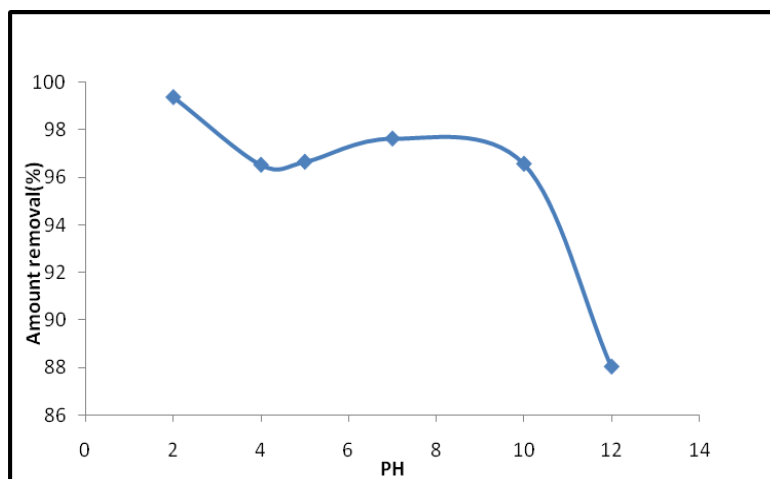


Figure(5) : Effect of particle size on adsorption capacity of (JGB) onto (WKSC)

at initial dye concentration of 50 mg/l and adsorbent dose of 1.5gm

Effect of pH :-

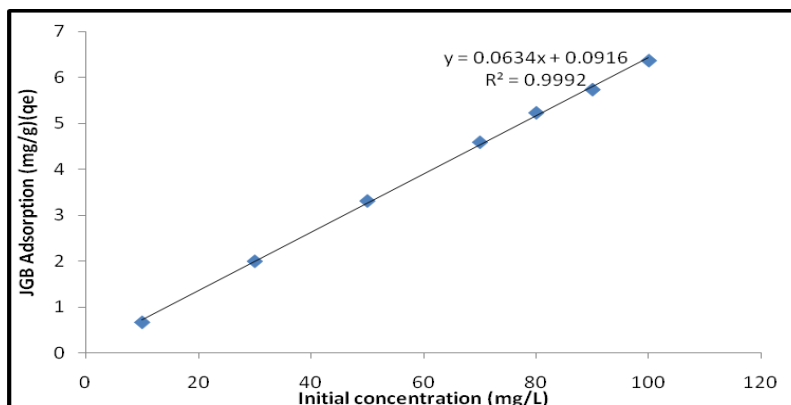
The aqueous solution of dye (JGB) having concentration of 50 mg/L was treated by 1.5 gm dosage of adsorbent with pH 2 to 12 . The pH was maintained with help of 0.1 N (HCL) and 0.1 N (NaOH) solution. Figure (6) it is evident the results obtained are presented in this Figure which describes that it was no significant change in the percentage removal of dye uptake over the entire pH range of (4-10) [35]and the best percentage in (pH=2) .



Figure(6) : Adsorption of (JGB) by (WKSC) as a function of solution pH at initial concentration of 50 mg/l and adsorbent dosage of 1.5 g with 0.075 mm particle size

Effect of initial dye concentration:-

The effect of concentration of dye (JGB) (10-100 mg/L) have been also tested with constant dosage of adsorbent for 150 min. . The removal of dye decreased from 100% to 95.54% . The results indicated that the adsorption of dye are much dependent on concentration of solution .Figure (7) illustrated the effect of initial dye concentration on adsorption of (JGB) onto the adsorbent and can be seen the adsorption capacity increased from (0.66 -6.36) mg/g when (JGB) concentration increase from 10 -100 mg/L for the adsorbent of (WKSC). Sites for adsorption becomes fewer for adsorption. Also the formation of second layer of the dye molecules is highly hindered at higher initial concentration of the dye, due to the repulsive interaction between adsorbed and unadsorbed dye molecules present on the solid surface and in solution, respectively[28][30].



Figure(7) : Effect of initial concentration on adsorption capacity of (JGB) onto (WKSC) (0.075mm particle size and 1.5gm adsorbent dose)

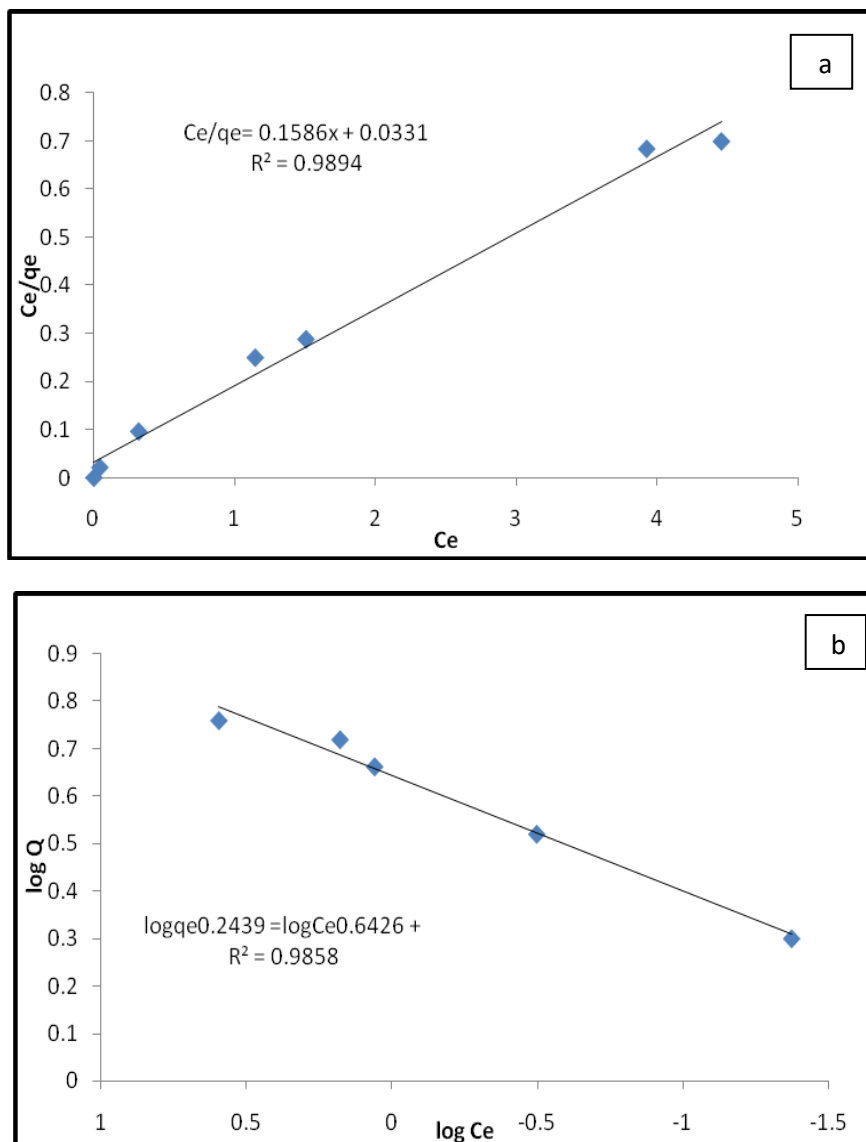
Isothermal analysis :-

The adsorption isotherm were developed from the data collected . Equilibrium adsorption data were fitted to the linear form of Langmuir's and freundlish equation (4,6). Table (2) shows that the adsorption of (JGB) dye using (WKSC) carbon both satisfies of Langmuir's and freundlish isotherm .

Table (2) : Adsorption isotherm parameters for (JGB) dye removal

	Langmuir		Freundlich
q_m (mg/g)	6.3051	$K_F(mg/g)(l/mg)^{1/n}$	4.9271
K_a (l/mg)	4.7938	1/n	0.2439
R^2	0.9894	R^2	0.9858
R_L	(0.0204-0.0020)		

The plats of liberalized from of Langmuir and Freundlich are shown in figure (8a-b). The Langmuir equilibrium adsorption curves relating solid and liquid phase concentration of (JGB). and the Freundlich equilibrium adsorption curves relating solid and liquid phase concentration of (JGB).



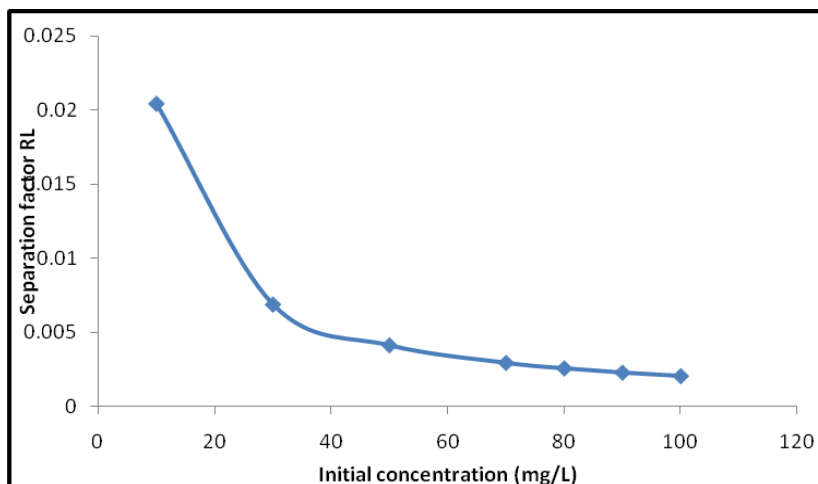
Figure(8): Linearized adsorption isotherm model of (JGB) onto (WKSC)

(a) Langmuir model

(b) Freundlich model

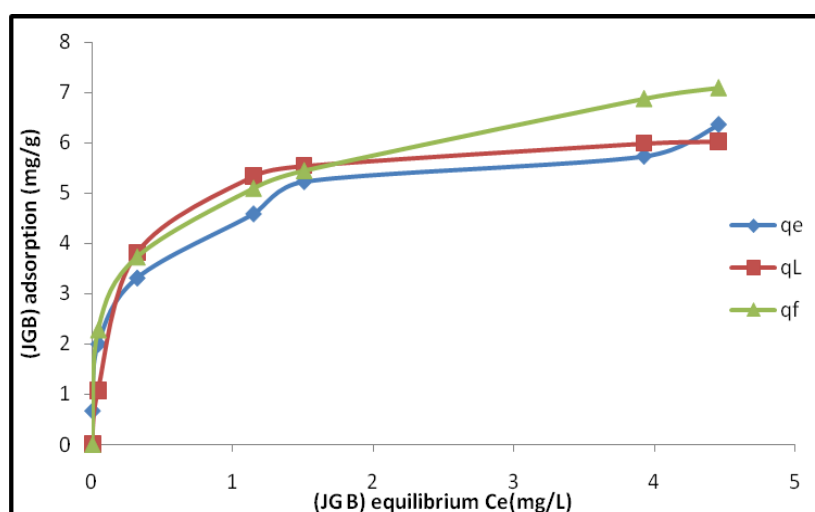
The Langmuir isotherm model assumes monolayer coverage of adsorbate on a homogeneous adsorbent surface. The well fitting of data with Langmuir isotherm indicates to the homogenous distribution of active sites on the adsorbent surface.

The variation of separation factor (R_L) with initial (JGB) concentration is shown in Figure (9). The (R_L) values for the adsorption of (JGB) onto (WKSC) are observed to be in the range of (0 – 1), indicating that the adsorption was favorable process.



Figure(9) : Separation factor versus initial (JGB) concentration on to (WKSC)

Figure (10) shows the deviation of these models from the experimental data. It appears that the adsorption of (JGB) dye on activated (WKSC) could be well fitted by the two isotherms. clearly, the Langmuir equation provided better fitting in terms of R_L .



Figure(10): Comparison of experimental and calculated data by Langmuir and Freundlich equilibrium isotherms for the system (JGB) – (WKSC)

This results indicate homogenous nature of (WKSC) surface, which means each (JGB) molecule (WKSC) has equal adsorption activation energy. The results also demonstrate the formation of monolayer coverage of (JGB) molecule at the outer surface of (WKSC).

Conclusion :-

The findings of the present work reveal that the WKSC which is easily and

abundantly available agro waste in our country can be easily converted into good adsorbent by using simple methods of activation. A suitable amount (1.5g/l) of the WKSC adsorbent could decolorize as much as 99.36% of the dye from an aqueous solution (50 ppm) if agitated for 150 min. demonstrated sufficient potential of WKSC as an adsorbent for the removal of the dye (JGB), from water solutions. The adsorption of the dye was maximum around the (pH=2) of the aqueous solution of (JGB). On applying both Langmuir and Freundlich isotherm .

References:-

- [1] Szygula, A.; ruiz, M.; guibal, E.; M.sastre, A., Removal of an anionic reactive dye by chitosan and its regeneration ,2nd international conference on Management, water pollution corfu, Greece, (2008).
- [2] Jyoti and Beena janveja , A study on removal of congo red dye from the effluents of textile industry using rice husk carbon activated by steam, Rasayan J. Chem. 1, (4)936-942 (2008).
- [3] Patel, R. and S. Suresh, kinetic and equilibrium studies on the biosorption of reactive black 5 dye by aspergillusfoetidus Bioresource Technology, 99 (1) 51-58 (2008) .
- [4] Robinson, T.; Chandran, B. ,Removal of dyes from a synthetic textile dye effluent by biosorption on apple pomace and wheat straw, Water Research 36, 2824–2830 (2001).
- [5] Crini, G. ,Non-conventional low-cost adsorbents for dye removal: a review, Bioresource Technology 97, 1061–1085(2006).
- [6] Eren, Z. and Acar, F. N. ,Adsorption of reactive black 5 from an aqueous solution equilibrium and kinetic studies, Desalination 194, 1–10(2006).
- [7] Mondal, S. ,Methods of dye removal from dye house effluent – an overview, Environmental Engineering Science 25 383–396(2008).
- [8] Calvete, T.; Lima, E.C.; Cardoso, N.F.; Dias, S.L.P. and Pavan, F.A. , application of carbon adsorbents prepared from the brazilian pine-fruit-shell for the removal of procion red mxb from aqueous solution-kinetic, equilibrium ,and thermodynamic studies. Chemical engineering Journal, 155(3) 627-636 (2009).
- [9] Liang, S.; Guo, X.; Feng, N. and Tian, Q., Isotherms, kinetics and thermodynamic studies of adsorption of Cu^{2+} by Mg^{2+} from aqueous solutions k⁺ type orange peel adsorbents. Journal of hazardous materials, 174(3)-756 (2010). 762

- [10] Arami- Niya A.; Daud, W.M.A. Wan, and Mjalli, F.S., Production of palm shell based activated carbon with more homogeneous pore size distribution , J. Applied Sciences ,10(24),3361 -3366 (2010)
- [11] Gergova, K.; Petrov, N. and Eser,S., Adsorption properties and microstructure of activated carbons produced from agricultural by –products by stream pyrolysis, carbon. 32, 693 -703 (1994).
- [12] Hilary, I. Owamah; Ilabor S. Chukwujindu and Augustine, K. Asiagwu, Biosorptive capacity of yam peels waste for the removal of dye from aqueous solutions Civil and Environmental Research 3(1)(2013).
- [13] Hu, Z.; Srinivasan, MP., Preparation of high-surface-area activated carbons from coconut shell, Micropor. Mesopor .Mater. 27 ,11-18(1999).
- [14] Tan, I.A.W.; Ahmad, A.L. and Hameed, B.H. , Preparation of activated carbon from coconut husk: Optimization study on removal of 2,4,6-trichlorophenol using response surface methodology, J. Hazard. Mater. 153 ,709-717 (2008) .
- [15] Kadirvelu, K. and Namasivayam, C., Activated carbon from coconut coir pith as metal adsorbent: adsorption of Cd (II) from aqueous solution, Adv. Environ. Res. 7, 471-478(2003).
- [16] Ravanpaykar, A.; Asfaram, M. R.; Fathiemadaba, Removal of Dye (Blue 56) From Aqueous Solution via Adsorption onto Pistachio A. Shell: kinetic and isotherm study of removal process Journal of Chemical Health Risks 2(1)41-48 (2012).
- [17] Balsi, S.; Dogu, T. and Yucel, H., Characterization of activated carbon produced from almond shell and hazelnut shell , Journal Chemical Technology And Biotech. 60, 419-426 (1994).
- [18] Juan F. Gonza ´lez; Silvia Roma ´n; Carmen M. Gonza ´lez-Garci ´a; Valente Nabais, J. M. and A. Luis Ortiz Porosity Development in Activated Carbons Prepared from Walnut Shells by Carbon Dioxide or Steam Activation Ind. Eng. Chem. Res., 48, 7474–7481 (2009).
- [19] Daud WMAW; Ali WSW; Suleiman MZ., The effects of carbonization temperature on pore development in palm-shell-based activated carbon, Carbon 38, 1925-1932 (2000).
- [20] Sua ´rez-Garci ´a F.; Mart ´nez-Alonso, A.; Tascon ,JMD., Pyrolysis of apple pulp: chemical activation with phosphoric acid ,J.Analy. Appl. Pyrol. 63, 283-301 (2002).

- [21] Hayashi, J.; Kazehaya, A.; Muroyama, K.; Watkinson AP., Preparation of activated carbon from lignin by chemical activation, *Carbon* 38 ,1873-1878(2000).
- [22] Sharma, Y.C.; Singh, B. and Uma Fast Removal of Malachite Green by Adsorption on Rice Husk Activated Carbon *The Open Environmental Pollution & Toxicology Journal* 1, 74-78(2009)
- [23] Sugumaran, P.; Priya Susan, V.; Ravichandran, P. and Seshadri, S., Production and Characterization of Activated Carbon from Banana Empty Fruit Bunch and Delonixregia Fruit Pod *Journal of Sustainable Energy & Environment* 3 ,125-132 (2012).
- [24] Bagher, H.; Saber, M.; Shishehbore, M. R.; Shahvazian, M., Pro. Color colorants coat. 3(11)58-65(2010).
- [25] Langmuir, I., The adsorption of gases on plan surface of glass, mica and platinum, *J. Am. Chem. Soc.*, 40, 1361-1403(1918).
- [26] Weber, T.W.; Chakravorti, R.K., Pore and solid diffusion models for fixed bed adsorbers, *Amer Inst Cheml Engrs' J*, 20,228(1974).
- [27] Freundlich, H.Z., Over the adsorption in solution, *J. Phys. Chem.*,57A:385-470(1906).
- [28] Taha, D.N.; Samaka, I.S.; Khalil, A.H., Comparative adsorption study for the basic dye wastewater treatment using natural Iraqi clay minerals and various adsorbents, *J. Education College-University of Babylon-Iraq*, 2(2)229-246(2009).
- [29] Theivarasu, C.; Mysamy, S. ,Equilibrium and kinetic adsorption studies of Rhodamine –B from aqueous solutions using cocoa (*Theobroma cocoa*) shell as anew adsorbent *International Journal of Engineering sciencee and Technology* 2 (11) 6284-6292 (2010).
- [30] Ho, Y.S.; Chiang, T.H.; Hsueh, Y.M., Removal of basic dye from aqueous solution using tree fern as a biosorbent , *Process Biochem.*, 40,119-124(2005).
- [31] Banat, F.A.; AL-Bashir, B.; AL-Asheh, S.; Hayaijneh, O., Adsorption of phenol by bentonite, *Environ. Poluut.*, 107,391-398(2000).
- [32] Senthikumar, S.; Varatharajan, PR.; Porkodi, K.; Subburaam, C.V., Adsorption of methylene blue carbon onto jute fiber carbon, *Colloid Interface Sci*, 284,79(2005).

- [33] Mckay, G; Otterburn, M.S.; Sweeny, A.G., The removal of colour from effluent using various adsorbents-IV silica: equilibria and column studies, *Water Res.*, 14, 21-27 (1980).
- [34] Al-Degs, Y.; Khraisheh, M.A.; A.S.J.; Ahmed, M.N., Evaluation of activated carbon adsorbents for the removal of textile reactive dyes from wastewater. *Jordan International Chemistry Engineering Conference*, 1, 159-167 (2000).
- [35] Mohan, D.; Singh, K.P.; Sinha, S.; Gosh, D., Removal of pyridine from aqueous solution using low cost activated carbons derived from agricultural waste materials, *Carbon* 42, 2409 – 2421 (2001)