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Enhanced Rhodamine B Dye Adsorption by Groundnut Shell Activated Carbon Coated With Fe₃O₄

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ABSTRACT

In this present study, adsorption capacity of groundnut shell activated carbon coated with Fe_3O_4 as a low cost adsorbent for removal of cationic dye, Rhodamine B (RDB) from aqueous solution was explored. The surface morphology and functional groups present were investigated by scanning electron microscope (SEM) and Fourier transform infrared (FTIR) spectroscopy. The effect of various parameters such as contact time, pH, adsorbent dosage and concentration of adsorbate were studied at ambient temperature. Equilibrium data were analysed using Langmuir, Freundlich and Temkin isotherm models. Kinetic data were studied using Pseudo-first and Pseudo-second order kinetic models and the mechanism of adsorption was described by intraparticle diffusion model.

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Introduction

Nowadays as a result of growing environmental awareness, the use of natural dye stuff obtained from vegetable and other animal sources have decreased substantially. They are replaced by synthetic dye stuffs produced on an industrial scale with its largest consumer being the textile industries ^[1]. More than 10,000 different types of dyes and pigments, with an annual production of more than 0.7 million tons are commercially available ^[2]. Unfortunately, during the dyeing process, a large percentage of the synthetic dye does not bind and is lost to the waste stream as residual dyes, approximately 5-10% dyes are released into the environment making the effluent highly coloured with breakdown products that are toxic, carcinogenic or mutagenic. Thus clearly, dyes are a potential hazard to living organisms ^[3].

Conventional waste water treatment methods for removing dyes include physicochemical, chemical and biological methods, but are very expensive and advance in technology ^[4]. Adsorption onto activated carbon process is one of the effective techniques that have been successfully employed for dyes removal from waste water because of its capability for adsorbing a broad range of different types of adsorbates efficiently, and its simplicity of design ^[4]. However, commercially available activated carbon is still considered expensive ^[5]. Groundnut shells are sometimes used for stock feed, but not as popular as cereal straws and legume stovers. Therefore, they can be used for the commercial production of activated carbon without much competition from livestock industry. Moreover, large quantities of groundnut shells can be obtained very cheaply from companies that are involved in shelling groundnut.



Preparation of activated carbon

Groundnut shells were obtained from shelling units. The shells were washed with deionized water and dried in a hot air oven at temperature of 120°C for 3 hours. The dried shells were then grounded into fine particles for easy activation. The activation procedure was carried out in a 2L beaker by thoroughly soaking the powdered material in concentrated sulphuric acid 1:2 (W/V) heated to 200°C with continuous agitation for 1h. During this period, the water was evaporated and the mixture became slurry and started to solidify. Agitation was then stopped while heating continues until the material became a carbon like material. The resulting carbon was then allowed to cool to room temperature, washed with distilled water, soaked in dilute sodium hydroxide for 30 minutes; washed a second time with distilled water and dried in an oven. The product was crushed, sieved into sizes at 100µm and then stored in a tight bottle ready for use.

Preparation of Fe₃O₄-AC

The chemical precipitation technique has been used to prepare particles with homogeneous composition and narrow size distribution ^[5]. A complete precipitation of Fe₃O₄ was achieved under alkaline condition, while maintaining a molar ratio of Fe²⁺ to Fe³⁺, 1:2, under inert environment. To obtain 2g of Fe₃O₄, 2.1g of FeSO₄.7H₂O and 3.1g of FeCl₃.6H₂O were dissolved under inert atmosphere in 80ml of double distilled water with vigorous stirring using a mechanical stirrer. While the solution was heated to 80°C, 10ml of NH₄OH solution (25%) was added. To ensure complete growth, 10g of the prepared groundnut shell activated carbon was added and reaction continued for 30 minutes. The resulting suspension was cooled down to room temperature and then repeatedly washed with distilled water to remove unreacted chemicals. The product was dried in an oven at 50°C for 2h, tested using a magnetic rod and then stored in a tight bottle ready for use. The reactions that occur in the production are shown in chemical equations (1) and (2).

 $\begin{array}{ccc} FeSO_4.7H_2O + 2FeCl_3.6H_2O + 8NH_4OH \longrightarrow Fe_3O_4 + 6NH_4Cl + \\ (NH_4)_2SO_4 + 17H_2O & (1) \\ Fe_3O_4 + AC + 17H_2O \longrightarrow AC-Fe_3O_4 & (2) \end{array}$

Batch Adsorption Experiments

The batch adsorption experiments were conducted to study optimum removal of colour from the dye solution. 0.2g of AC-Fe₃O₄ was added into a glass stopper conical flask containing 50ml of dye solution at 25°C. All adsorption experiments were conducted at temperature of 303K and agitation was provided at constant rate of 100rpm. The contents were filtered using a whatman filter paper no. 42. Analysis was done using UV-Visible spectrophotometer (Elico Model No: SL210). Dye concentration in the supernatant was estimated by measuring absorbance at maximum wavelength 545nm and computing from calibration curve. The kinetic studies were carried out using a mechanical stirrer. The samples were withdrawn at predetermined time intervals and filtered. The percentage removal and amount of RDB adsorbed was calculated by the following equations.

Percentage removal of RDB	$= \frac{C_o - C_e}{C_o} X$	100(1)
Amount of RDB adsorbed (q_e)	$= \frac{(C_o - C_e) V}{m}$	(2)

Where C_0 and C_e are the initial and equilibrium concentration of RDB (mgL^{-1}) respectively, m is the mass of the adsorbent (g) and V is the volume of RDB solution (L).

Results and Discussion

Adsorbent Analysis Method

The Fe₂O₃ enriched with groundnut shell was characterized by SEM and FT-IR. The Scanning Electron Microscopy (SEM) measurements were carried out using SEM-Quanta. The images were taken with an emission current = $100 \mu \text{Å}$ by the Tungsten filament and an accelerator voltage =10kV. Fourier transform infrared spectroscopy of the adsorbent was done by using an FT-IR spectrophotometer (Model: FT-IR Bruker IFS 66V). Spectra obtained in the range of 400 to 4000 cm^{-1} were analysed.

SEM Studies

SEM is a primary tool for characterizing the surface texture, morphology and fundamental physical proportion of the adsorbent. It is useful for determining particle shape, porosity, and appropriate size distribution of the adsorbent ^[6-10]. SEM images for the parent activated carbon (AC), Fe₃O₄ coated activated carbon (AC- Fe₃O₄) and after RDB adsorption (AC-Fe₃O₄-RDB) are shown in figure 2. The parent activated carbon (AC) as shown in figure 2a has more active sites, while figure 2b shows the parent activated carbon completely covered with iron oxide. Figure 2c shows the SEM image, after the adsorption of RDB onto AC- Fe₃O₄. The iron oxide particles exhibit magnetic behaviour, which is depicted by more particles being adsorbed as shown in figure 2c.



Figure 2. SEM image of (a) parent AC, (b) AC- Fe₃O₄ and (c) AC- Fe₃O₄-RDB

Fourier Transform Infrared Spectroscopy

Fourier transform infrared spectroscopy of the adsorbent was done by using an FTIR spectrophotometer (Model: FTIR Bruker IFS 66V). Spectra obtained in the range of 400 to 4000 cm⁻¹ were analysed. The FTIR spectra of both Parent AC and AC- Fe₃O₄ are shown in figures 3a and 3b. The spectrum of the Parent AC displayed the following bands: 3412cm⁻¹: OH and NH stretching; 2938cm⁻¹: C-H stretching in alkanes, 1631cm⁻¹: C=O stretching in amides, 1621cm⁻¹: C=C stretching in alkenes, 1208cm⁻¹: C-O stretching in ethers; $798cm^{-1}$ CH₂ bending in benzene derivatives ^[10-18]. The main oxygen groups were suggested to be carbonyl groups (such as, ketone, quinones), and phenol, ether in the parent AC. The band at 468 and 534 cm⁻¹ which were associated with the in-plane and out-of-plane aromatic ring deformation vibrations common that is quite common for activated carbon. When comparing the two peaks in figures 3a and 3b, figure 3b shows that there were various functional groups detected on the surface of AC- Fe₃O₄ and a peak is observed in the Fe-O group. There are some peaks that were shifted, disappeared and new peaks were produced. Significant band decrease of functional group on the AC-Fe₃O₄ were detected at bands 1631, 1402, 1209 and 1034 cm⁻¹ which corresponded to the bonded C=O stretching, stretching in quinones, stretching alkanes and C-O stretching in ethers respectively. The newer peak at 589cm⁻¹ observed in figure 3b is related to the Fe-O group, and the peak around 3432 cm⁻¹ in curve a was assigned to the -OH group on the surface of the magnetite. Figure 3c shows the FTIR spectrum after adsorption of RDB onto AC- Fe₃O_{4.} When comparing figure 3c with 3b, it can be clearly seen that most of the functional groups present on the surface get destroyed. The presence of polar functional groups on the surface is likely to confer considerable anion exchange capacity on the adsorbents [16-21].







Figure 3. FTIR image of (a) parent AC, (b) AC- Fe₃O₄ and (c) AC- Fe₃O₄-RDB

Effect of Time

The percentage of dye adsorbed at different time interval is calculated by keeping the concentration as 100 mgL^{-1} and the mass of the adsorbent as 0.2g. From figure 4, it can be seen that the rate of adsorption was very rapid at initial period of contact time. After 120 minutes, the solution attains equilibrium ^[7]. The quick adsorption in the first 60 minutes is due to the high availability of adsorption sites, there after the available adsorption sites become fewer which decrease the rate of adsorption and an equilibrium state was attained. This time was taken as optimum contact time for all subsequent studies ^{[6-}





Effect of pH

pH of the aqueous solution is a controlling parameter in the adsorption process. The influence of pH was studied using a portable pH meter (Systronics made) by adjusting the reaction mixture to different initial pH (2 to 6) value using 0.1M Sodium Hydroxide or Hydrochloric Acid before adding the adsorbent. The pH meter was calibrated using 4.0 buffer tablet. As clearly seen in figure 5, there is high uptake at low pH values compared to the high pH values. This may be attributed to the fact that at pH value below 6, the RDB ions can enter into the pore structure, while at high pH values above 6, the zwitter ions of RDB may increase the aggregation of RDB to form a bigger molecular form (dimer) and become unable to enter the into the pore structure ^[8-14].



Effect of AC- Fe₃O₄ Dose

This factor is used to determine the uptake capacity of an adsorbent for a given initial concentration of an adsorbate. This was studied by varying the AC- Fe₃O₄ dose (0.1 – 0.5 g/50ml) for RDB of concentration 100 mgL⁻¹. The percentage adsorption (as shown in figure 6) increases with increase in the AC- Fe₃O₄ dose. This is attributed to the increase in surface area and more adsorption sites ^[6]. This observation is consistent with Langmuir hypothesis of an increase competition among adsorbent particles for organic substances with increasing number of adsorbent particles per unit volume ^[9-10].





This was studied by varying the initial concentration of the dye, while keeping other parameters constant. It was observed (as shown in figure 7) that the percentage removal decreases with increase in dye concentration. This is due to the fact that, at low concentration, there are much available sites for adsorption whereas, with increase in initial concentration, the available sites for adsorption becomes fewer, thereby reducing the percentage removal of RDB^[10-16].



		Tab	le 1. Isother	m Parameters	for the remo	val of RDB	onto AC-Fe ₃	O ₄		
_	Langmuir isotherm			Freundlich isotherm			Temkin isotherm			
	q _m (mgg ⁻¹)	K _L (mgL ⁻¹)	\mathbf{R}^2	K _f (mgg ⁻¹)	1/n	\mathbf{R}^2	Α	В	\mathbf{R}^2	
	47.61	1.3291	0.9982	21.38	0.2155	0.9678	1.06	11.28	0.9781	
		Ta	ble 2. Kineti	c Parameters fo	or the remov	al of RDB o	nto AC-Fe ₃ C) ₄		
	Pseu	Ta do-first order	ble 2. Kineti	c Parameters fo Pseu	or the remov do-second ord	al of RDB o ler	nto AC-Fe ₃ C Intr) ₄ ra-Particle	diffusion	
k_1(Pseu (min ⁻¹) x10 ⁻⁵	Ta do-first order q _e (mg g ⁻¹)	$\frac{1}{R_1^2}$	$\frac{\text{Parameters fo}}{\frac{\text{Pseu}}{q_e(\text{mg}}}$	or the remove do-second ord k ₂ (g ⁻¹ mg min)x10 ⁻⁴	al of RDB o ler R ₂ ²	nto AC-Fe ₃ C Intr K _{int} (mg g min ^{-0.5})	D ₄ ra-Particle	diffusion	R ²

Adsorption Isotherm

The adsorption isotherm indicates equilibrium relationship between the adsorbent and adsorbate ^[17]. The analysis of the isotherm data by fitting them to different isotherms models is an important step to find the suitable model that can be used for the designed purpose. Its applicability was judged with the regression coefficients (\mathbb{R}^2). The isotherms used include:

Langmuir Isotherm

The Langmuir isotherm was chosen for the estimation of maximum adsorption capacity corresponding to complete monolayer coverage on the adsorbent surface ^[18]. It applies to the cases of adsorption on completely homogeneous surfaces where interactions between adsorbed molecules are negligible. The linearized Langmuir equation was used to analyse the data. The equation is given as:

Where, C_e is the equilibrium concentration of the adsorbate (mgL^{-1}) , q_e is the amount of adsorbate adsorbed per unit mass of the adsorbent (mg/g), q_m and K_L is the Langmuir constants related to adsorption capacity and energy of adsorption, respectively. The more the R^2 value approaches unity, the more fitted the adsorption is to Langmuir isotherm ^[16].



Figure 8. Langmuir Isotherm for RDB adsorption onto AC-Fe₃O₄

The essential features of the Langmuir isotherm may be expressed in terms of equilibrium parameter R_L which is a dimension less constant referred to as separation factor or equilibrium parameter.

$$R_{L} = \frac{1}{1 + K_{L}C_{o}}.....(4)$$

Where, C_o is initial dye concentration (mg/L) and K_L is Langmuir constant related to the energy of adsorption. R_L value indicates the adsorption nature to be either unfavourable if $R_L >$ 1, linear if $R_L =$ 1, favourable $0 < R_L <$ 1 and irreversible if $R_L =$ 0. The R_L was 0.204 suggesting isotherm to be favourable at concentrations studied as the value lies between 0 and 1 ^[16-18].

Freundlich Isotherm

The Freundlich isotherm model was chosen to estimate the adsorption intensity of the adsorbent. It assumes that uptake of dyes occurs on a heterogeneous adsorbent surface that has unequal available sites with different energies of adsorption. The linear form of Freundlich equation is given as follows:

$$\ln q_e = \ln K_f + \ln C_e \frac{1}{n}.....(5)$$

Where, Kf is a constant indicative of the adsorption capacity of the adsorbent (mg/g) and the constant 1/n indicates the intensity of the adsorption. Figure 9 shows linearized Freundlich isotherm for RDB adsorption onto AC-Fe3O4 respectively. The more the R2 value approaches unity, the more fitted the adsorption is to Freundlich isotherm [16].



Figure 9. Freundlich Isotherm for RDB adsorption onto AC- Fe₃O₄

The magnitude of the exponent, n gives an indication of the favourability and capacity of the adsorbate/adsorbent system, where (1<n<10), represent favourable adsorption process and new adsorption sites are generated, while if n =1, then the partition between the two phases are independent of concentration ^[23]. In this work, the value of n was found to be 4.64, indicating favourable adsorption.

Temkin Isotherm

The Temkin isotherm model predicts a uniform distribution of binding energies over the population of surface binding adsorption ^[17]. The linear form of Temkin equation is given as follows:

$$q_e = B \ln A + B \ln C_e....(6)$$

where A and B are the Temkin isotherm constant (L/g) and heat of sorption (J/mol) respectively. R is the gas constant (J/mol/K), b is the Temkin isotherm constant linked to the energy parameter B as shown in equation 7:

$$B = \frac{RT}{b}....(7)$$

Figure 10 shows linearized Temkin isotherm for RDB adsorption onto AC-Fe₃O₄ respectively. The more the R^2 value approaches unity, the more fitted the adsorption is to Temkin isotherm ^[17-18].



Figure 10. Temkin Isotherm for RDB adsorption onto AC- Fe_3O_4

Adsorption Kinetics

The kinetics of sorption describes the solute uptake rate, which in turn governs the residence time of adsorption reaction. In order to investigate the adsorption process of RDB onto AC- Fe_3O_4 , three kinetic models were used including pseudo-first order, pseudo-second order, and intra-particle diffusion models. **Pseudo-First Order Model**

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Kinetic data were treated with the pseudo- first order kinetic model ^[19]. The differential equation has the following forms:

$$\frac{dq_t}{dt} = k_1 \left(q_e - q_t \right)....(8)$$

Where q_e and q_t refer to the amount of the dye adsorbed (mg/g) at equilibrium and at time, t (min) respectively and k_1 is the equilibrium rate constant of pseudo- first- order adsorption (min⁻¹), Integrating equation (1) for the boundary conditions t = 0 to q_t gives:

$$\log \frac{q_e}{q_e - q_t} = \frac{k_1}{2.303} t...(9)$$

Which is integrated rate law for a pseudo- first- order equation. Equation (9) can be rearranged to obtain a linear form:

$$\log (q_e - q_t) = \log q_e - \frac{\kappa_1}{2.303} t...(10)$$

In order to obtain rate constants, the straight line plot of log (qe-qt) against t as shown in figure 11 was used to determine the first order rate constant k_1 .



Figure 11. Pseudo-First Order Kinetics for RDB adsorption onto AC- Fe_3O_4

Pseudo-Second Order Model

Kinetic data were further applied to the pseudo second order kinetic model .The differential equation has the following form:

$$\frac{dq_t}{dt} = k_2 (q_e - q_t)^2 \dots (11)$$

Where k_2 is the equilibrium rate constant of the pseudosecond order adsorption (g/mg min⁻¹) Integrating equation 11 for the boundary condition t = 0 to q_t, gives:

This is integrated rate law for a pseudo –second order reaction. Equation.12 can be rearranged to obtain a linear form:

$$\frac{t}{q_t} = \frac{1}{k_2 {q_e}^2} + \frac{t}{q_e}....(13)$$

The kinetic plots of t/q_t versus t for RDB removal is presented in figure 12. The relationship is linear, and the correlation coefficient (R²) suggests a strong relationship between the parameters and also explains that the process of the sorption of RDB follows pseudo- second order kinetics. The product k₂q_e² is the initial sorption rate represented as h = k₂q_e²



Figure 12. Pseudo-Second Order Kinetics for RDB adsorption onto AC- Fe₃O₄

Intra-particle Diffusion Study

The intra-particle diffusion model was applied to identify important of pore diffusion in the adsorption process. The mathematical expression of intra-particle diffusion model is: $q_t = K_{int}t^{0.5} + C$ (14) Where, K_{int} is the intra-particle diffusion constant (mg g⁻¹ min^{0.5}) and the intercept (C) reflects the boundary layer effect. If the rate limiting step is intra-particle diffusion, then the graph drawn between q_t vs. t^{0.5} should yield a straight line passing through the origin. The slope give the value of the intra-particle diffusion constant (K_{int}) and the value of C gives an idea about the thickness of the boundary layer. The values of K_{int} were calculated from the slope of as shown in figure 13 and presented in table 2.



Figure 13. Intra-Particle Diffusion Plot for RDB adsorption onto AC- Fe₃O₄

Conclusion

The use of groundnut shell activated carbon coated with Fe_2O_3 has better potential than raw groundnut shell activated carbon as a low cost adsorbent for the removal of RDB from aqueous solutions. The effect of various parameters such as contact time, pH, adsorbent dosage and dye concentration was

studied. Maximum adsorption of RDB from aqueous solutions occurred at pH of 6. Adsorption increased with increasing dose of adsorbent and decreased with increasing initial RDB concentration. The experimental data shows high association with Langmuir but not Freundlich isotherm model and Temkin isotherm model. Kinetic study reveals that, the reaction order fits with pseudo second order. The process is economically feasible and easy to carry out.

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