Thermodynamic studies of adsorption of rhodamine B and Congo red on graphene oxide

Juma Sahar,*, Abdul Naeem, Muhammad Farooq, Shah Zareen, Ata urRahman

National Centre of Excellence in Physical Chemistry, University of Peshawar, Peshawar, 25120, Khyber Pakhtunkhwa, Pakistan, email: jsahar14@yahoo.com (J. Sahar)
Institute of Chemical Sciences, University of Peshawar, Peshawar, 25120, Khyber Pakhtunkhwa, Pakistan

Received 22 January 2019; Accepted 19 May 2019

ABSTRACT

In the present work, graphene oxide was prepared by the modified Hummer’s method for the effective removal of rhodamine B and Congo red from their aqueous solution. The synthesized graphene oxide was characterized by different techniques such as thermogravimetric/differential thermal analysis, scanning electron microscopy, Fourier transform infrared spectrometry and X-ray diffraction. It was found that among different models applied, the Langmuir isotherm model fitted the data very well for both the rhodamine B and Congo red dyes adsorption onto the graphene oxide. Moreover, Van’t Hoff equation was used to calculate the various thermodynamic parameters such as, enthalpy ($\Delta H^o$), entropy ($\Delta S^o$) and Gibbs free energy ($\Delta G^o$) for both the selected dyes. The $\Delta H^o$ for rhodamine B and Congo red was found to be 4.18 kJ mol$^{-1}$ and 10.69 kJ mol$^{-1}$ whereas, $\Delta S^o$ was found to be 5.95 J mol$^{-1}$K$^{-1}$ and 8.80 J mol$^{-1}$K$^{-1}$ respectively. Similarly, the Gibbs free energy ($\Delta G^o$) was calculated to be -1.89 kJ mol$^{-1}$ and -2.78 kJ mol$^{-1}$ for rhodamine B and Congo red respectively. The thermodynamic studies indicated that the entire adsorption process was physisorption, endothermic and spontaneous in nature. The high adsorption affinity of the graphene oxide could be due to the electrostatic interaction between the positively charged surface of rhodamine B and negatively charged surface of graphene oxide. Further, this study depicts that the synthesized graphene oxide has a great potential to remove the dyes from waste water effectively.

Keywords: Graphene oxide; Rhodamine B; Congo Red; adsorption; Langmuir adsorption isotherm

1. Introduction

Water is an indispensable part of life as most of the human metabolic activities also depend upon water. Water is used for drinking, cleaning, bathing, air conditioning, washing and heating. Water containing pollutants affect the natural level of environment. Any undesirable change in the chemical, physical and biological properties of the water lead to harmful effects on the human life as well as the natural environment [1].

Approximately 3% of the entire earth surface water is fresh water. A very small proportion of this water is available for the human society [2]. The toxic effluents from industries and agriculture sector are significant threat to the environment [1]. Similarly, domestic and municipal wastes are also responsible for water pollution. In the above context, fresh water in general and particularly in Pakistan is becoming unsafe for the use of human purposes [2].

Dyes are the main cause of water pollution and becoming the major environmental problem in the universe. Dyes have low biodegradability, affects human beings and natural ecosystem as well [3]. Different dyes are used for different purposes by textiles, dyeing operations, printing, food and paper making industries [4–6]. Colored effluents coming from the textile, paper, plastic, dyeing and printing industries containing Congo red and rhodamine B [5,7].

Congo red having anionic nature in solution belongs to diazo dye and it is the first synthetic dye used for dyeing cotton. Congo red dye upon metabolization produces benzidine which is known to be human carcinogen [8,9]. It also
causes irritation to eyes, skin, gastro intestine and affects blood clotting as well as induces the respiratory problems. Congo red dye is optically, thermally and physiocoherently stable [10,11]. Therefore, biodegradability of this dye is difficult due to its complex aromatic nature. This dye is also sensitive towards the pH and its color changes from red to blue as the pH of the solution decreases below pH 5 and becomes reddish in the pH range of 5-10 [8].

Similarly, Rhodamine B belongs to xanthene dyes and is soluble both in water and organic solvents. Due to its stability, it is used as a biological stain, pathology, histology and for dye laser production [12]. It can also be used for dyeing wool, jute leather, silk and cotton in textile industries. It is highly toxic and causes irritation to eyes, skin and respiratory tract if swallowed by humans and animals [13]. Due to the presence of N-ethyl groups on the sides of xanthene ring it is highly toxic and carcinogenic due to which it is banned in many countries while using for coloring of food and cosmetics [14].

Diffrerent physico-chemical techniques such as coagulation, ozonation, flocculation, chemical oxidation, ion exchange, reverse osmosis, photo catalysis, precipitation and adsorption are used for the treatment of different industrial waste effluents [15,16]. Among these techniques, adsorption process is the most effective technique which is successfully employed for the removal of color from wastewater [17]. Adsorption is more reliable process due to its simplicity, high efficiency, regenerability [15] and recycling of the adsorbant and low production of residues [16-18].

Different adsorbents like activated carbon, fly ash, manganesse oxide, silica gel, soil, alumina, red mud, bentonite, kaolinite, hematite, biomass, polymers and clay [15,16] are used for the removal of toxins from aqueous system [19]. Yang et al. [20] studied the adsorptive removal of Congo red dye using functionalized carbon nanotube mixed metal oxide and its maximum adsorption capacity was 1250 mg/g. Jia et al. [21] used 3D hierarchical porous iron oxides for the adsorptive removal of Congo red dye from waste water. The maximum adsorption capacity was obtained for FeOx. The removal performance was reported in the order of FeOx > FeOx > γ-FeOx. Motahari et al. [22] studied the removal of rhodamine B from waste water using the NioN nano particles in the presence of H+ acacen ligand and the maximum monolayer capacity was observed to be 111 mg/g.

Graphene oxide displays excellent electronic, thermal, mechanical and optical properties. Therefore, it can be used in catalysis, electronics, sensors, and for energy conversion and storage [23]. Graphene oxide can also be modified by using nanomaterials, surfactants, biomolecules and polymers to enhance its efficiency for different applications. Graphene oxide has been used as an adsorbent for the adsorption of different dyes, aromatic wastes and heavy metal cations from water on account of its acidic nature and high surface area [24,25]. Graphene oxide can act as an excellent adsorbent for the removal of rhodamine B and Congo red. However, no systematic studies on the ion exchange removal of rhodamine B and Congo red from aqueous solutions by the graphene oxide have been reported in the literature. In the present work, graphene oxide has been utilized for the sorption of rhodamine B and Congo red from aqueous solutions. Moreover, the effect of pH has also been evaluated in order to develop an efficient process for the removal of selected dyes under suitable conditions.

2. Experimental

2.1. Materials and chemicals

Graphite powder was purchased from DaeJung chemicals, potassium permanganate, sulphuric acid, hydrochloric acid, sodium nitrate, sodium hydroxide and hydrogen peroxide, were purchased from Scharlau. Rhodamine B having a molecular weight equal to 479.01 g/mol and a molecular formula C24H24N2O6Cl was purchased from BioM laboratories Cerritos USA (Chemical Division, Malaysia) and Congo red dye having a molecular formula C31H26N4Na2O8S, and molecular weight equal to 693.665 g/mol was obtained from BDH chemical Ltd Poole England. All the chemicals used during the whole studies were of analytical grade.

2.2. Graphene oxide syntheses

The graphene oxide was synthesized by using modified Hummer’s method [26]. Typically, 3 g of graphite powder was dissolved in 70 mL of concentrated sulphuric acid (H2SO4) in the double walled cell and the temperature of the system was kept below 10°C. The mixture was stirred for 30 min and then 9 g of KMnO4 was added slowly and the temperature was maintained lower than 5°C. The mixture was stirred and the temperature was allowed to increase gradually. When the temperature reached to 40°C, about 150 mL of deionized water (H2O) was added slowly to the mixture and the temperature was raised up to 90°C. Then 500 mL of deionized water was further added which was followed by the addition of 15 mL of 30% H2O2. As a result, the color of the solution changed from dark brown to yellow and the suspension was stirred to make it homogeneous. The material was filtered and washed with 10% HCl solution to remove metal ions. The as synthesized material was again dispersed in 1000 mL of deionized water and kept overnight. The suspension was centrifuged and washed with deionized water until the pH became neutral. The obtained particles (graphite oxide) were dispersed in deionized water and sonicated to get the graphene oxide dispersion. The graphene oxide dispersion was then centrifuged and dried in oven at 100 for further studies [26,27].

2.3. Characterization of graphene oxide

Different analytical techniques such as thermogravimetric differential thermal analysis (TG/DTA), scanning electron microscopy (SEM), fourier transform infrared spectrometry (FT-IR) and X-ray diffraction (XRD) were used for the characterization of the synthesized graphene oxide before and after adsorption studies.

2.4. Adsorption studies

Batch adsorption experiments were performed in order to obtain the rate and data of equilibrium. A bout 40 mL of each rhodamine B and Congo red (1.4 × 10^-4 M) was taken into 100
mL titration flasks separately and then 0.1 g of adsorbent was added to each reaction vessel. The reaction mixtures were equilibrated for 24 h at 298 K in the shaker bath. The suspension was then filtered by using the Whatman filter paper 44 to separate the graphene oxide. The supernatant solution absorbance was then determined in order to calculate the concentration of residual dyes. UV-Visible spectrophotometer (BMS VIS-1100) was used to measure the concentration before and after the treatment of graphene oxide with selected dyes in the wavelength range of 498 to 554 nm. Initial concentration of dye and adsorbent dosage were varied to study their effect on the adsorption kinetics. In order to observe the pH effect on the dyes adsorption, the adsorption process was carried out in the pH range of 2–10. The solutions pH was adjusted by using HCl/NaOH solutions and pH meter. The studies of adsorption were performed at various temperatures, such as 298 K, 308 K, and 318 K, in order to find the effect of temperature on the thermodynamic parameters. The sorption amount at time, \( q_t \) (mg/g or mol/g) was calculated by using the following equation:

\[
q_t = \frac{(C_0 - C_t)V}{W}
\]

where \( C_t \) (mg/L or mol/L) is the dye liquid phase concentration at any time, \( C_0 \) (mg/L or mol/L) is the initial concentration of dye in the solution, \( V \) (L) is the solution volume and \( W \) (g) is the graphene oxide mass.

The equilibrium sorption capacity amount \( q_e \) (mg/g or mol/g) was determined by using the following equation:

\[
q_e = \frac{(C_0 - C_e)V}{W}
\]

The percent removal of rhodamine B and Congo red can be determined from the following equation:

\[
q_e = \frac{(C_0 - C_e)}{C_0} \times 100
\]

where \( C_0 \) and \( C_e \) (mg/L or mol/L) are the initials and an equilibrium liquid phase concentration of the rhodamine B and Congo red in the solution.

3. Results and discussion

3.1. Characterization of graphene oxide

3.1.1. X-ray diffraction analysis (XRD)

The X-ray diffraction (XRD) analysis was performed to achieve the information about the degree of oxidation. Fig. 1 shows the XRD pattern of the synthesized graphene oxide. It can be observed that only one peak appeared at the \( 2\theta = 10.5^\circ \) in the XRD pattern of the graphene oxide which corresponds to (001) plane. This confirms that the sample of graphene oxide was synthesized successfully. The present findings are in close agreements with the literature [28,29].

3.1.2. Thermal gravimetric analysis (TGA)

The thermodgravimetric analysis was used to measure the weight loss in the graphene oxide as shown in Fig. 2. It can be observed from the thermogravimetric curve that the graphene oxide is thermally unstable and starts to lose its weight upon heating. Further, it can be observed that the weight loss of graphene oxide was observed in three stages. In the first stage i.e. from 70–140°C the weight loss could be attributed to the removal of adsorbed water. A high weight loss could be observed at 200°C, which may be due to the removal of oxygen containing groups in graphene oxide in the form of CO\(_2\) and H\(_2\)O. The peak appeared at 550°C is attributed to the combustion of carbon skeleton of graphene oxide [30]. Similarly, the DTA curve also supported the three weight losses at about 97.83°C, 283.16°C and 600°C.

3.1.3. Scanning electron microscopy (SEM)

SEM is widely used for the surface and morphological characteristics of the adsorbent material. It also deals with the porosity and surface texture of the adsorbent and plays an important role in the surface availability determination for the dyes adsorption onto the adsorbent. Fig. 3a shows the SEM images of synthesized graphene oxide before adsorption while Figs. 3b and 3c show the SEM images of graphene oxide after the adsorption of rhodamine B and Congo red respectively. Before the adsorption, the graphene
oxide particles are highly porous while after adsorption the surface of graphene oxide is loaded with rhodamine B and Congo red and the graphene oxide becomes bulky and the particles are agglomerated and becomes spherical in shape [31,32].

3.1.4. Fourier transform infrared spectroscopy (FTIR)

FTIR was used to identify the functional groups that are responsible for the rhodamine B and Congo red adsorption onto the surface of graphene oxide. Fig. 4 shows the FTIR spectra of graphene oxide before and after the rhodamine B and Congo red adsorption. Different oxygen containing functional groups could be observed in the FTIR spectra. The main absorption band recorded at 3340 cm\(^{-1}\) could be assigned to the stretching vibrations of O-H groups. The peaks of absorption at 1730 and 1630 cm\(^{-1}\) could be due to the C=O stretching of carboxylic and carbonyl functional groups respectively. While the two peaks of absorption appeared at 1226 cm\(^{-1}\) and 1044 cm\(^{-1}\) are due to the stretching vibrations of C-O [31]. After the rhodamine B adsorption, strong characteristic band appeared at 821 and 1331 cm\(^{-1}\) (Fig. 4b). These absorption bands could be assigned to the adsorption of rhodamine B onto the graphene oxide surface [33]. Similarly, the absorption peaks at 1182 cm\(^{-1}\) (Fig. 4c) is associated with the benzene rings vibrations and the peak at 1045 cm\(^{-1}\) is assigned to the stretching vibration of S=O bond [34]. Thus, it is concluded from the FTIR analysis, that rhodamine B and Congo red dyes were successfully adsorbed onto the surface of graphene oxide.

3.1.5. Point of zero charge of graphene oxide

Point of zero charge (PZC) of the graphene oxide was determined by the salt addition method. 40 mL of NaNO\(_3\) was taken in various conical flasks and then the pH of these solutions was adjusted in the range of 1 to 11 using NaOH/HCl. About 0.1 g of adsorbent (graphene oxide) was then added to each flask and kept for 24 h in a shaker bath. After 24 h, the pH of each suspension was determined and finally the pH versus pHi was plotted to calculate the PZC for graphene oxide as shown in Fig. 5. The PZC for graphene oxide was calculated to be 2, which is in close agreement with the reported values [35–37]. At this pH, graphene oxide behaves neutrally. The surface is positively charged below the PZC and adsorption of anionic dyes is most favourable while the surface is negatively charged above the PZC which adsorbs preferably the cationic dyes. Graphene oxide consists of a several functional groups such as epoxy, hydroxyl, ketone and carboxyl groups. When graphene oxide is dispersed in water, a negative charge is created on its surface because of the ionization of the surface functional groups. The PZC graph demonstrates that the surface of graphene oxide remains negatively charged almost over the entire pH range (above pH\(_{\text{PZC}}\) value 2).

3.2. Adsorption studies

3.2.1. pH effect on adsorption of rhodamine B and Congo red

The pH plays an important role in the dye adsorption onto the adsorbent surface. The surface charge of the adsorbent and the degree of ionization of the dye are dependent upon the change in the pH of the solution. The solution pH...
also changes the color intensity and structural stability of
the dyes [38]. The adsorption data collected at various pH
are given in Fig. 6a. It can be seen that the adsorption of
rhodamine B increases with the increasing solution pH. The
pH adsorption curve shows that the uptake of rhodamine
B is strongly dependent on initial pH of the solution. Fur
ther, it can be observed that the rhodamine B removal
from aqueous solution increases almost linearly with the
increase in the pH. However, at pH 7 the curve was level
ed off and no further change in the adsorption capacity
of the graphene oxide was noted. This indicates the satu
ration of the graphene oxide surface. The decrease in the
rhodamine B adsorption at pH 2 could be assigned to the
electrostatic repulsion between the negatively charge sur
face and rhodamine B. While the increase in the uptake
of rhodamine B in alkaline pH region can be attributed to
the electrostatic attraction between the negatively charged
surface and cationic dye [35,37]. While the effect of pH
on Congo red dye adsorption is shown in Fig. 6b. As can
be seen that the Congo red adsorption decreases with an
increase in the solution pH. The pH adsorption curve shows
that the uptake of Congo red is strongly dependent on ini
tial solution pH. Graphene oxide contains several oxygen
functional groups such as carboxylic (-COOH), epoxy and
hydroxyl (-OH) groups. Therefore, when it is dispersed in
aqueous solution it exist in anionic form due to carboxylic
groups (-COOH) which are present in carboxylate (-COO−)
form. On the other hand, the Congo red dye is dipolar in
nature. The Congo red dye may exists either as a cationic
or anionic form depending on the pH of the solution. At
lower pH, the NH2 and –SO3– groups of the Congo red
dye are protonated and converted to the NH3+ and –SO3H
respectively and show attraction towards the negative sur
face of the graphene oxide which results in the maximum
adsorption at pH 3 while at higher pH the Congo red exist
as negatively charged –SO3– due to the deprotonation of
the dye and showing repulsion towards the negatively charged
graphene oxide surface [39,40]. As such, the adsorption of
Congo red decreases with the increase in the pH of the solu
tion. But due to its sensitivity towards the acids, the color
of Congo red changes from red to blue with the change in
solution pH i.e. at low pH and in the pH range of 5-10 it
becomes reddish [11]. Consequently, pH 7 was chosen for
further studies because at this pH the dye remains stable
and no change in the color was observed at this pH value.
3.3. Adsorption isotherms

The adsorption isotherms are used to study the adsorbent performance in the process of adsorption and define the interactions between adsorbent and adsorbate. In the present studies, various isotherms such as Langmuir, Freundlich and Dubinin-Radushkevich were used to interpret the experimental data.

3.3.1. Langmuir adsorption isotherm for rhodamine B and Congo red adsorption

The monolayer formation on the adsorbent surface is best defined by the Langmuir adsorption isotherm. According to this model, a uniform adsorption of adsorbate molecules on certain specific sites of the adsorbent takes place. Further, the adsorption sites are energetically uniform and there is no adsorbate transmigration on the plane of the adsorbent surface [41,42]. The linear form of the Langmuir adsorption isotherm, which is based on the above assumptions, can be represented as follow.

\[ \frac{C_e}{q} = \frac{1}{q_m K_L} + \frac{1}{q_m} \]

where \( C_e \) (mg/L or mol/L) is the concentration of dye at equilibrium present in the solution, \( q \) is the amount of adsorbed dye per unit mass of adsorbent (mg/g or mol/g), \( q_m \) is the area occupied by the adsorbate monolayer, indicating the adsorption capacity (mg/g or mol/g), \( K_L \) is the Langmuir isotherm constant (L/mg) and associated to the adsorption free energy. The values of \( q_m \) and \( K_L \) for the rhodamine B and Congo red adsorption were obtained from the slope and intercept of the plot of \( C_e/q \) versus \( C_e \) as presented in the Fig. 7a-b. The theoretically calculated monolayer adsorption capacities from the Langmuir isotherm and experimental results for selected dyes are shown in Table 1 and 2. The experimental adsorption capacities are in close agreement with the theoretical adsorption capacities, which indicates that Langmuir model is best fitted to the experimental data of rhodamine B and Congo red adsorption onto the graphene oxide [43,44]. The monolayer adsorption capacity value slightly increases with increase in temperature from 298 to 318 K. The values of \( q_m \) and \( K_L \) concludes that maximum removal relates to the monolayer saturation of the molecules of adsorbate on surface of adsorbent with constant energy and no adsorbate transmission occurred in the plane of the surface of adsorbent. Further, the value of \( K_L \) shows that the entire process is endothermic in nature. The separation factor (\( R_L \)) was calculated (Tables 1 and 2) to confirm the favorable nature of the adsorption process. The Langmuir adsorption isotherm important characteristic can be expressed in term of dimensionless constant referred to equilibrium parameter or separation factor \( R_L \), and the shape of isotherm can be indicated from the value of \( R_L \).

\[ R_L = \frac{1}{1 + K_L C_i} \]

where \( C_i \) is adsorbate (dye) initial concentration present in the solution and \( K_L \) is the constant of Langmuir isotherm. The value of \( R_L \) shows the type of isotherm to be linear (\( R_L = 1 \)), irreversible (\( R_L = 0 \)), unfavorable (\( R_L > 1 \)) and favorable (0 < \( R_L < 1 \)). The calculated values of \( R_L \) at all initial concentrations of both rhodamine B and Congo red were found to be in the range of 0–0.01, which confirms that the rhodamine B dye uptake was favorable on graphene oxide [45].

3.3.2. Freundlich adsorption isotherm for rhodamine B and Congo red adsorption

Freundlich isotherm is used for the heterogeneous surfaces adsorption that involves the interaction between the molecules adsorbed. This model is not limited to the monolayer formation and the sorption energy decreases exponentially on the completion of the adsorbent sorption sites [46,47]. This isotherm is used for the determination of the adsorption intensity and capacity of the adsorbent. The empirical equation of the Freundlich isotherm is given below,

\[ q_e = K_f C_e^{1/n} \]
where $q_e$ is the amount of the adsorbed dye at equilibrium, $C_e$ is the dye concentration present in the solution at equilibrium, $K_F$ is Freundlich isotherm constant related to the capacity of adsorption (mg/g or mol/g), $n$ is empirical parameter signifies the intensity of adsorption and also shows the favourability of the process of adsorption, favorable ($0 < 1/n < 1$), irreversible ($1/n = 0$) and unfavorable ($1/n > 1$) [48]. If $n = 1$ then the partitions are concentration independent between the two phases. If $1/n$ value is below one it shows a normal sorption. On the other hand, $1/n$ value above one indicates the cooperative adsorption. Moreover, if $n$ value lies between 1 and 10, then it shows that the adsorption process is favorable. The Freundlich isotherm in linearized form is given below:

$$\ln q_e = \ln K_F + \frac{1}{n} \ln C_e$$  \hspace{1cm} (7)

The value of $1/n$ and $K_F$ can be obtained from the intercept and slope of the plot of $\log q_e$ versus $\log C_e$ (Fig. 8). The values obtained for the coefficient of correlation ($R^2$) for Freundlich isotherm are found to be lesser than the Langmuir isotherm, suggesting that Langmuir isotherm is best fitted to experimental data. This model further assumes that the adsorbate concentration at the interface increases with its corresponding increase in the solution. If $n$ value is equal to unity, then the adsorption process is linear. If $n$ value is below unity, it shows the process of adsorption is unfavorable, and if the $n$ value is above unity, then the process of adsorption is favorable. Further, if $1/n$ value is less than unity so it further confirms the normal Langmuir isotherm. As in our case, the value of $1/n$ is lesser than 1, so it confirms that the process of adsorption follows the Langmuir adsorption isotherm. If the value of $n$ lies between 1 and 10 so it shows a beneficial process of adsorption. In the present work, the value of $n$ for both dyes lies within this range. This implies that graphene oxide has a high affinity towards the rhodamine B and Congo red removal. The increase in the $K_F$ value with increasing temperature indicates that the capacity of adsorption has been increased for the rhodamine B and Congo red adsorption onto the surface of graphene oxide. Further, the increase in the capacity of adsorption provides an opportunity for the development of efficient adsorbents for wastewater treatment.

### Table 2

<table>
<thead>
<tr>
<th>Temperature (K)</th>
<th>Theoretical $q_m \times 10^{-5}$ (mol g$^{-1}$)</th>
<th>Experimental $q \times 10^{-5}$ (mol g$^{-1}$)</th>
<th>$R^2$</th>
<th>$K_L$ (g/L)</th>
</tr>
</thead>
<tbody>
<tr>
<td>298</td>
<td>8.79</td>
<td>8.69</td>
<td>0.996</td>
<td>1.233</td>
</tr>
<tr>
<td>308</td>
<td>9.60</td>
<td>9.44</td>
<td>0.997</td>
<td>1.294</td>
</tr>
<tr>
<td>318</td>
<td>11.33</td>
<td>10.70</td>
<td>0.963</td>
<td>1.371</td>
</tr>
</tbody>
</table>

**Fig. 8.** Freundlich isotherm for the removal of dye (a) rhodamine B and (b) congo red onto the surface of graphene oxide at pH 7.
Freundlich adsorption with increasing temperature shows that the process of adsorption is endothermic in nature.

### 3.3.3. Dubinin–Radushkevich model for rhodamine B and Congo red adsorption

The Dubinin–Radushkevich (D-R) model can be used to estimate the adsorption free energy and characteristic porosity. It helps in the determination of the nature of the adsorption, i.e., whether the process is chemical or physical. The D–R isotherm is temperature dependent and it indicates the non-homogeneous surface of the adsorbent [48,49]. The D–R isotherm linear form is given below.

$$\ln q = \ln q_m - \beta \varepsilon^2$$  \hspace{1cm} (8)

where $q$ is the amount of adsorbed dye per unit weight of graphene oxide (mol/g), $q_m$ is the maximum capacity of adsorption (mol/g), while $\varepsilon$ is the Polanyi potential and is equal to

$$\varepsilon = RT \ln \left(1 + \frac{1}{C_e}\right)$$  \hspace{1cm} (9)

where $R$ is the general gas constant and its value is equal to $8.314 \text{ kJ/molK}$ and $T$ is the absolute temperature (K), $\beta$ is the coefficient of activity associated with the adsorption mean free energy (mol$^2$/kJ$^2$), which is defined as "when one mole of ion from an infinity of solution is transferred to the solid surface then it is called free energy of adsorption and is represented by $E$" as given by the following equation,

$$E = (2k)^{-0.5}$$  \hspace{1cm} (10)

The process of physical adsorption occurs, when the $E < 8 \text{ kJ/mol}$ while $8 < E < 16$ shows that the chemical adsorption mechanism (ion-exchange). The Dubinin-Radushkevich model supported the mechanism of monolayer to be

![Fig. 9. D-R plots for the dye (a) rhodamine B and (b) congo red onto the surface of the graphene oxide at pH 7.](image-url)
3.4. Thermodynamic studies of rhodamine B and Congo red adsorption

The temperature effect on dyes adsorption is used to determine the important thermodynamic parameters such as Gibbs free energy, entropy and enthalpy change [18, 50]. The influence of these parameters on the rhodamine B adsorption was calculated based on the following equations:

\[ \Delta G = RT \ln K_c \] (11)

\[ \Delta G^o = \Delta H^o - T \Delta S^o \] (12)

where \( \Delta G \) is the Gibbs free energy change, \( R \) is the universal gas constant and its value is equal to 8.3145 J/ molK, \( T \) is absolute temperature (K), and \( K_c \) is the constant of equilibrium and is given below:

\[ K_c = \frac{C_p}{C_s} \] (13)

where \( C_p \) is the equilibrium dye concentration and \( C_s \) is the dye concentration in solution. The standard enthalpy and entropy of the process of adsorption can be calculated by using the Van't Hoff equation:

\[ \ln K_c = \frac{\Delta S^o}{R} - \frac{\Delta H^o}{RT} \] (14)

The value of \( \ln K_c \) was calculated from the Langmuir adsorption isotherm. The values of \( \Delta H^o \) and \( \Delta S^o \) were found to be 4.18 kJ·mol\(^{-1}\) and 5.95 J·mol\(^{-1}\)K\(^{-1}\) respectively for rhodamine B, whereas, for Congo red the values of \( \Delta H^o \) and \( \Delta S^o \) were 10.69 kJ·mol\(^{-1}\) and 8.80 J·mol\(^{-1}\)K\(^{-1}\) respectively. The \( \Delta H^o \) values of both dyes show that the removal was endothermic in nature. Moreover, the \( \Delta S^o \) values clearly indicates the irregular increase in the randomness at solid-liquid interface during the rhodamine B adsorption from aqueous solution onto the graphene oxide. The \( \Delta S^o \) value further indicates an increase in the degree of freedom of species adsorbed as reported elsewhere [18].

The Gibbs free energy (\( \Delta G^o \)) for the adsorption of rhodamine B and Congo red onto the graphene oxide was calculated by using Eq. (12). The negative value of \( \Delta G^o \) (Table 7) for rhodamine B and Congo red (Table 8) confirms that the removal process is spontaneous and thermodynamically favorable at all temperatures. As can be seen, that \( \Delta G^o \) value for rhodamine B changes from −1.77 to −1.89 kJ/ mol whereas, the \( \Delta G^o \) value for Congo red changes from −2.61 to −2.78 kJ/ mol. This indicates that the removal of rhodamine B and Congo red onto the surface of the graphene oxide at higher temperature is more spontaneous. Similar results were also reported by other researchers in the literature [51].

3.5. Adsorption mechanism

The mechanism of adsorption depends upon the texture properties and chemistry of the adsorbent and adsorbate.

Table 7

<table>
<thead>
<tr>
<th>Temperature (K)</th>
<th>( \Delta G^o ) (kJ mol(^{-1}))</th>
<th>( \Delta H^o ) (kJ mol(^{-1}))</th>
<th>( \Delta S^o ) (J mol(^{-1})K(^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>298</td>
<td>−1.77</td>
<td>4.18</td>
<td>5.95</td>
</tr>
<tr>
<td>308</td>
<td>−1.83</td>
<td></td>
<td></td>
</tr>
<tr>
<td>318</td>
<td>−1.89</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Table 8

<table>
<thead>
<tr>
<th>Temperature (K)</th>
<th>( \Delta G^o ) (kJ mol(^{-1}))</th>
<th>( \Delta H^o ) (kJ mol(^{-1}))</th>
<th>( \Delta S^o ) (J mol(^{-1})K(^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>298</td>
<td>−2.61</td>
<td>10.69</td>
<td>8.80</td>
</tr>
<tr>
<td>308</td>
<td>−2.70</td>
<td></td>
<td></td>
</tr>
<tr>
<td>318</td>
<td>−2.78</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
The forces that are involved in the process of adsorption are electrostatic attraction and interaction as shown in Scheme 1. The electrostatic attraction exists between the opposite charges on graphene oxide and both dyes rhodamine B and Congo red while the interactions occur between the aromatic ring of rhodamine B and Congo red and the aromatic ring of graphene oxide [52].

3.6. Comparison with other adsorbents

The adsorption capacity of graphene oxide for adsorption of rhodamine B and Congo red was compared with other adsorbents reported in the literature. The maximum adsorption capacity of graphene oxide was calculated to be 1655 mg/g, which is higher than the adsorption capacities of other adsorbents as shown in Table 9. Postai et al. [53] investigated the removal of rhodamine B from waste water by using the waste seeds Aleurites moluccana (WAM) as an adsorbent. The maximum adsorption capacity of rhodamine B was 117 mg/g. Peng et al. [54] used the humic acid modified Fe₃O₄ nanoparticles (Fe₃O₄/HA) for the adsorption of rhodamine B from aqueous solution and the maximum adsorption capacity was reported to be 161.8 mg/g. Khan et al. [55] reported the kaolinite adsorption efficiency for the removal of rhodamine B from aqueous solution and the maximum adsorption capacity for rhodamine B adsorption was 46.08 mg/g. Zhou et al. [56] used the shrimp shell powder for the removal of Congo red from waste water. Results revealed that adsorption capacity for treated shrimp shell powder was 288.2 mg/g for the removal of Congo red. The above study suggest that graphene oxide may be used as an excellent adsorbent for the removal of rhodamine B and Congo red from aqueous solution due to the presence of different oxygen functional groups, high surface area and high adsorption capacity. Therefore, graphene oxide has a great potential to be applied for the treatment of waste water in the near future.
Graphene oxide was successfully synthesized by modified Hummer’s method. The various analytical techniques such as thermogravimetric/differential thermal analysis, scanning electron microscopy, fourier transform infrared spectrometry and X-ray diffraction were used to evaluate the physiochemical properties of the synthesized graphene oxide. The synthesized graphene oxide showed potential capacity towards adsorption of rhodamine B and Congo red. However, it was found that graphene oxide showed greater affinity towards the adsorption of rhodamine B as compared to Congo red. This may be due to the strong electrostatic interaction between the positively charged surface of rhodamine B and negatively charged surface of graphene oxide. Moreover, the thermodynamic parameters such as $\Delta H^o$, $\Delta S^o$ and $\Delta G^o$ calculated in this study, clearly indicated that the adsorption process of the selected dyes on graphene oxide was physiosorption, endothermic and spontaneous at all temperatures. Thus, graphene oxide may be used as a potential adsorbent for the removal of toxic pollutants from drinking water.

References
