

## Batch and bulk removal of hazardous colouring agent Rose Bengal by adsorption techniques using bottom ash as adsorbent

Vinod Kumar Gupta,<sup>\*ab</sup> Alok Mittal,<sup>c</sup> Damodar Jhare<sup>c</sup> and Jyoti Mittal<sup>c</sup>

Received 31st May 2012, Accepted 12th July 2012

DOI: 10.1039/c2ra21351f

Rose Bengal is a halogen-containing fluorescein water soluble dye, which is widely used for medical purposes. The dye, however, is highly toxic and can cause irritation, itching *etc.* to the human skin and eyes. It is therefore considered worthwhile to develop a systematic method for the removal of Rose Bengal by adsorption processes. The present investigation is devoted to batch as well as bulk removal of Rose Bengal. Under preliminary batch studies, adsorption isotherm measurements and kinetic studies were carried out, while for the bulk removal, a glass column was used as a fixed bed adsorber made up of Bottom Ash. Attempts were also made for the recovery of the dye from the exhausted Bottom Ash column by eluting dilute NaOH. The paper also presents a detailed procedure for activating Bottom Ash and its chemical and physical analysis. Rose Bengal showed a decrease in adsorption with increasing pH and conversely, increases in concentration, temperature, amount of adsorbent and sieve size increased the adsorption. Langmuir, Freundlich, Tempkin and D–R adsorption isotherm models were also verified, and on the basis of Langmuir constants thermodynamic parameters such as the Gibb's free energy, enthalpy and entropy of the adsorption were also calculated. A pseudo-second order process was found to operate during the adsorption. During column operations various parameters like fractional capacity of the column, mass flow rate, percentage saturation of column *etc.* were calculated. Desorption from the exhausted column gave almost 91% of dye recovery.

### 1. Introduction

Rose Bengal (Fig. 1), also known as Acid Red 94 [IUPAC name 4,5,6,7-tetrachloro-3',6'-dihydroxy-2',4',5',7'-tetraiodospiro(isobenzofuran-1(3H),9'-(9H)xanthen)-3-one], is a well known fluorescein class of halogen-containing dye. It is a water soluble dye and exhibits a pink colour. It has several medical diagnostic applications, *e.g.*, liver function testing, staining of necrotic tissue in the eyes, devitalized cells of the cornea in conjunctivitis *etc.*<sup>1</sup> The low concentration dosage of Rose Bengal is also used in the treatment of eczema and psoriasis. Apart from various clinical uses, Rose Bengal also has severe toxic effects on the human corneal epithelium.<sup>2,3</sup> The dye is extremely hazardous when it comes in contact with skin and creates irritation, itching, scaling, reddening and even blistering. It also causes inflammation, redness, watering and itching in the eyes. On ingestion by inhalation it damages mucous membranes, which leads to respiratory irritations in humans.<sup>4</sup>

For the past few years economic and eco-friendly methods have been developed for the eradication of highly toxic and hazardous dyes from their aqueous solutions by adsorption using waste materials as adsorbents.<sup>5,6</sup> It is now established that adsorption technique is one of the most powerful tools for the removal of pollutants from the water<sup>7–10</sup> and the technique has several advantages over other physico-chemical methods like electrochemical treatment<sup>11</sup> photochemical treatment,<sup>12</sup> biodegradation,<sup>13</sup>

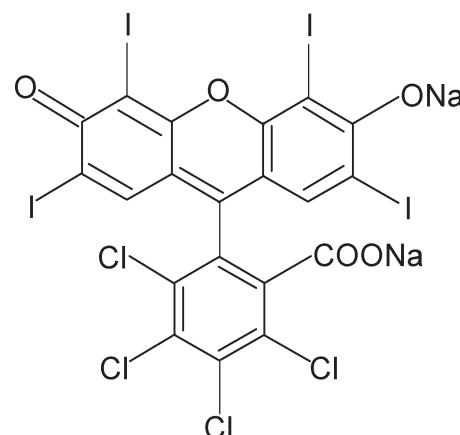


Fig. 1 Chemical structure of the dye Rose Bengal.

<sup>a</sup>Department of Chemistry, Indian Institute of Technology Roorkee, Roorkee - 247 667, India. E-mail: vinodfcy@gmail.com; Fax: 00911332286202; Tel: 00911332285918

<sup>b</sup>King Fahd University of Petroleum and Minerals, Dhahran 31261, Saudi Arabia

<sup>c</sup>Department of Chemistry, Maulana Azad National Institute of Technology, Bhopal, 462051, India

froth flotation<sup>14</sup> and reverse osmosis<sup>15</sup> etc. The abilities of the adsorption process to remove and recover the larger organic molecules, without their decomposition, have also popularized it over other modern analytical, electrochemical and photochemical methods.<sup>16</sup> Use of waste materials as potential adsorbents influences the economics of the developed process and recently, selection of cheap and convenient waste materials has become a main focus of research. Reports are available that the degradation products of the bulky dye Rose Bengal are much more toxic than the original molecule.<sup>17</sup> Therefore employing an adsorption process for the removal and recovery of a costly dye like Rose Bengal is advantageous and necessary, while the cost reduction of the developed process can be achieved by selecting a waste material as adsorbent.

Bottom ash is a waste material collected at the bottom of the furnace in the thermal power plants. It is always produced in huge quantity and disposed in nearby land. Discarding of bottom ash has always been a matter of great concern to the plant authorities as the dumped ash is highly unsuitable for agricultural uses; it makes the agricultural land barren, infertile and unproductive.<sup>18</sup> To date there is no evidence of the presence of any toxic element in the bottom ash<sup>19,20</sup> and therefore its use as an adsorbent for the removal of water soluble molecules is extremely safe. This research article presents the attempt of our laboratory to utilize bottom ash as a potent adsorbent for the efficient removal and necessitous recovery of toxic Rose Bengal dye. The developed method is easy, cheap, fast and versatile.

## 2. Experimental

### 2.1. Material and methods

Rose Bengal ( $C_{20}H_2Cl_4I_4Na_2O_5$ ,  $M_r$ : 1017.65) was procured from M/S Merck and the 0.01 M stock solution was the dye was prepared in doubly distilled water. All other reagents used were of analytical reagent grade and purchased through M/s Advance Scientific, Bhopal. Bottom ash was obtained from the Thermal Power Station (T. P. S.) of M/s Bharat Heavy Electrical Limited (B. H. E. L), Bhopal (India). The T. P. S. of B. H. E. L, Bhopal purchases sub-bituminous coal from M/s South Eastern Coal Field Limited, Chirmiri, Sarguja (India).

During adsorption experimentation all pH measurements were monitored by microprocessor based pH meter model number HI 8424, M/s Henna Instruments, Italy. The dye uptake was measured by recording the absorbance of the dye solution on a UV/VIS spectrophotometer model 117 (M/s Systronics, Ahmedabad, India) over wide wavelength range of 200 to 700 nm.

In order to measure various physical properties of activated bottom ash, different instruments were used. The surface area was determined by using Quantasorb Model QS-7, porosity by a Mercury porosimeter and density by specific gravity bottles. To characterize bottom ash, the IR spectrum was recorded on an Infra-red spectrophotometer (HP FT-IR), scanning electron microscopy was performed on a Philips SEM 501 electron microscope, while X-ray measurements were carried out on a Philips X-ray diffractophotometer employing Nickel filtered Cu- $\alpha$ -radiation.

For the bulk removal and recovery of Rose Bengal, adsorption and desorption were carried out through column operations. The

main component in column studies was a long tubular glass column of 30 cm length and 1 cm internal diameter filled with the bed of adsorbent, through which the dye solution is percolated to absorb the dye material. On exhaustion of column dilute NaOH was slowly percolated for the recovery of the dye.

### 2.2. Material development

Activation of the adsorbent material plays a very significant role in the adsorption process, since it increases the number of surface active sites of the adsorbent, thereby providing it with a wider surface available for the adsorption. To activate bottom ash, it was first ground into very small granules and then thoroughly washed with doubly distilled water. To oxidize all undesired organic impurities from it, small sized granules of bottom ash were dipped into a hydrogen peroxide solution (30% w/v) for about 24 h. The solution is then filtered and washed again several times by doubly distilled water. From the material thus obtained moisture is first removed by keeping it in an oven at 100 °C for about 15 min. Further activation of bottom ash was achieved by keeping the dried material in a furnace at 500 °C for about 30 min. Bottom ash granules were then sieved to obtain the desired particle sizes such as 0.425–0.150 mm (36 BSS Mesh), 0.150–0.088 mm (100 BSS Mesh) and  $\leq 0.088$  mm (170 BSS Mesh) (BSS: British Standard Size), stored in vacuum desiccators.

**2.2.1. Characterization of adsorbent.** The chemical characterization of the activated bottom ash was carried out by applying standard conventional chemical methods.<sup>20</sup> It was found that  $SiO_2$  is present in the highest amount of 45.4%, while  $CaO$ ,  $Al_2O_3$ ,  $Fe_2O_3$ ,  $MgO$  and  $Na_2O$  were present 15.3, 10.3, 9.7, 3.1 and 1% by weight, respectively. Physical parameters like surface area, density, porosity and loss of ignition of the activated bottom ash were obtained as  $870.5\text{ cm}^2\text{ g}^{-1}$ ,  $0.6301\text{ g mL}^{-1}$ , 46% and 1.14%, respectively.

To determine the nature of bottom ash, its weighed amount was added to 25mL of distilled water, pH = 7.0, and stirred thoroughly. The solution was kept undisturbed for 24 h in the 100 mL airtight measuring flasks and then filtered to measure the pH. The pH value was found to decrease, confirming thereby the acidic nature of bottom ash.

Different magnifications of scanning electron microscopic photographs of the activated granules of bottom ash reveal that its particles are almost spherical and porous in nature. Differential Thermogravimetric Analysis (DTA) curves indicate that the activated bottom ash is thermally stable and a very negligible weight loss occurred at high temperature. The X-ray diffraction indicates the presence of alumina ( $Al_2O_3$ ), gypsum ( $CaSO_4 \cdot 2H_2O$ ), beaverite [ $Pb(Cu,Fe,Al)_3(SO_4)_2(OH)_6$ ], borax ( $Na_2B_4O_7 \cdot 10H_2O$ ) and kaolinite [ $2\{Al_2Si_2O_5(OH)_4\}$ ] in the activated bottom ash. The infrared spectra of activated bottom ash give peaks in  $790\text{ cm}^{-1}$  region, corresponding to kaolinite [ $2\{Al_2Si_2O_5(OH)_4\}$ ]. The bands at  $3467$ ,  $2930$ ,  $2676$ ,  $1502$ ,  $1097$ ,  $470.2$  and  $790\text{ cm}^{-1}$  confirm the presence of laumontite [ $4CaAl_4Si_4O_12 \cdot 4H_2O$ ], amber, mulite [ $3Al_2O_3 \cdot 2SiO_2$ ], azurite [ $Cu_3(CO_3)_2(OH)_2$ ], bavenite [ $4Ca_4(BeAl)_4Si_9(O,OH)_{29}(OH)_2$ ], gypsum [ $3(CaSO_4 \cdot 2H_2O)$ ] and corundum [ $2(\alpha-Al_2O_3)$ ] respectively (Fig. 2).

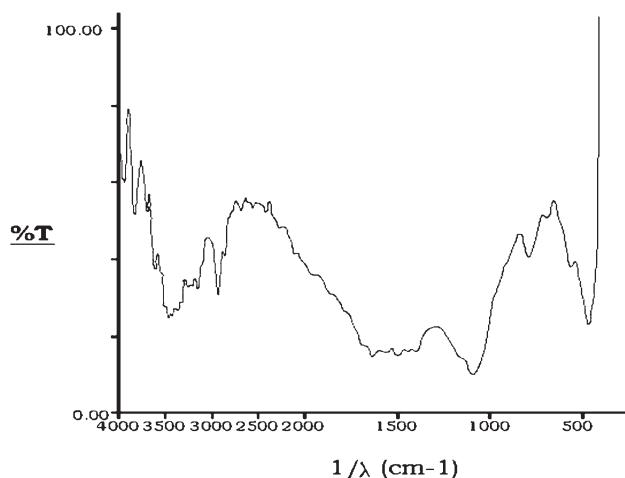


Fig. 2 Infra-red spectrum of activated bottom ash.

### 2.3. Adsorption studies

Adsorption of the halogen containing dye Rose Bengal was first achieved *via* a batch technique at temperatures 30, 40 and 50 °C; batch experiments were performed to observe the effects of pH, adsorbate concentration, amount of adsorbent, sieve size of adsorbent, temperature and time of contact.

In an airtight 100 mL volumetric flask, 25 mL of the dye solution of desired concentration was taken and a fixed amount of bottom ash of a particular sieve size was added into it. The conical flask kept in a water bath was then agitated by a mechanical shaker and then kept for 24 h for saturation. When the equilibrium was thought to have been established, the supernatant was then filtered through Whatman filter paper (No. 41) and the amount of dye up taken was monitored spectrophotometrically at the absorption maxima of Rose Bengal, *viz.*,  $\lambda_{\text{max}} = 544$  nm. It is pertinent to note that the pHs of the dye solutions were maintained by using NaOH and HCl.

### 2.4. Kinetic studies

Kinetic measurements were carried out by taking 25 mL of the dye solutions in 100 mL volumetric flasks and adding suitable amount of bottom ash. The solutions were shaken at regular intervals and after filtration, spectrophotometric analysis was performed at  $\lambda_{\text{max}} = 544$  nm. The change in the concentration of the dye before and after adsorption gave the dye uptake.

### 2.5. Column studies

The main objective of flow-through column experiments is to provide a more sensible simulation of how the bottom ash adsorbs the colour contaminant. The column was prepared by making a slurry of the bottom ash of desired size in doubly distilled water. The slurry was then slowly fed into the glass column over a glass wool support leading to the formation of adsorbent beds. In order to avoid air entrapment in the column the water of the slurry was continuously expelled through the column outlet. When the stable beds are formed, a dye solution of  $6 \times 10^{-5}$  M concentration was then eluted through the column with an appropriate flow rate of 0.5 mL min<sup>-1</sup> and

absorbance of the received dye solution were monitored by UV/VIS spectrophotometrically.

### 2.6. Column regeneration

Once the concentration of received dye solution equalised to that of the fed dye solution, the column was considered to be exhausted and at this stage column operation was stopped. In order to recover the adsorbed dye from the exhausted column, aliquots of dilute NaOH were passed with a constant flow rate of 0.5 mL min<sup>-1</sup> and the dye material was recovered at the outlet of the column. When complete dye was drained out of the adsorbent bed of the column, column was properly washed with hot water and made ready for next cycle of operation.

## 3. Results and discussion

### 3.1. Adsorption studies

**3.1.1. Effect of solution pH.** The pH of the dye solution plays an important role in the adsorption of any adsorbate. Batch studies have been performed for the adsorption of Rose Bengal over bottom ash by varying the pH from 2.2 to 11.2 at a fixed initial dye concentration of  $1 \times 10^{-4}$  M, 0.50 g and 100 BSS mesh size of adsorbent at temperatures 30, 40 and 50 °C. It was found that at low pH almost 100% removal was achieved, while with increasing pH the dye uptake decreases (Fig. 3). Since almost 50% adsorption of the dye was achieved at pH 7.2, this pH was selected for all subsequent studies.

**3.1.2. Effect of adsorbate concentration and temperature.** The adsorption behaviors of Rose Bengal on bottom ash was investigated in concentrations ranging from  $2 \times 10^{-5}$  M to  $10 \times 10^{-5}$  M, at a fixed pH of 7.2 and different temperatures (30, 40 and 50 °C). The concentration study reveals that an increase in the concentration of the dye solution increases the extent of adsorption over bottom ash almost linearly (Fig. 4). Fig. 4 also confirms that with increasing temperature the amount of dye uptake increases, indicating thereby endothermic nature of the ongoing adsorption.

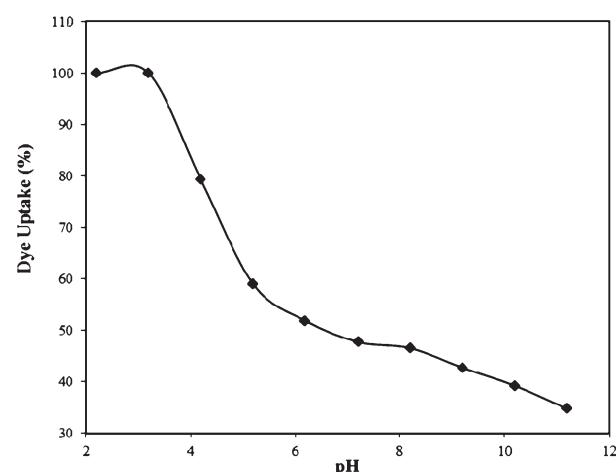
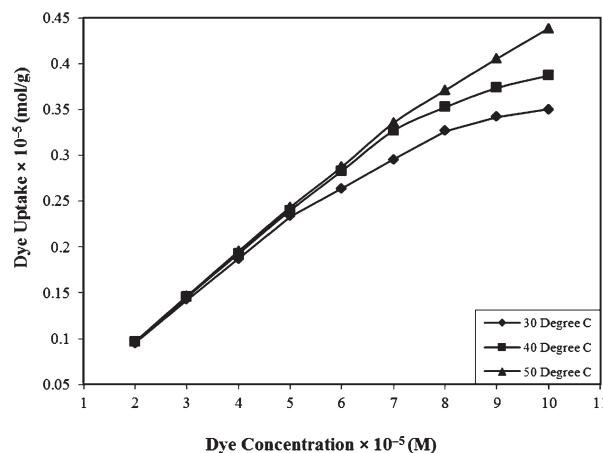


Fig. 3 Effect of pH on the Rose Bengal (concentration =  $1 \times 10^{-4}$  M)–bottom ash (amount = 0.50 g; mesh size = 100 BSS) system.



**Fig. 4** Effect of concentration on the Rose Bengal (pH = 7.2)-bottom ash (amount = 0.50 g; mesh size = 100 BSS) system at different temperatures.

**3.1.3. Effect of amount and sieve size of adsorbent.** To measure the effect of the amount of bottom ash on the adsorption of Rose Bengal, batch experiments were conducted at 30, 40 and 50 °C. 0.10 to 0.80 g of bottom ash were added at each temperature to 25 mL of  $1 \times 10^{-4}$  M Rose Bengal solutions maintained at pH 7.2. It was observed that the adsorption of the dye increases with an increase in the amount of adsorbent (Table 1). The most favourable adsorption was observed between 0.50 to 0.80 g of the adsorbent. Subsequently, adsorption studies were performed by selecting 0.50 g of adsorbent at each temperature.

The adsorbent material bottom ash used in the present research was sieved into three different mesh sizes *viz.* 36, 100 and 170 BSS Mesh. In the batch studies, 0.50 g of bottom ash of all the three mesh sizes is brought into contact with  $1 \times 10^{-4}$  M dye solution for about 24 h. It was also examined that with the decrease in the size of the particles of bottom ash, the percentage uptake of Rose Bengal increases. The amount of adsorption achieved for 36, 100 and 170 BSS mesh were  $0.333 \times 10^{-5}$ ,  $0.431 \times 10^{-5}$  and  $0.442 \times 10^{-5}$  mol g<sup>-1</sup>. This may be due to an increase in the surface area of the adsorbents.

**3.1.4. Effect of contact time.** The effect of contact time was monitored at a dye concentration of  $5 \times 10^{-5}$  M, a pH of 7.2 and 0.5 g of bottom ash. At temperatures 30, 40, 50 °C, equilibrium was established at around 90 minutes and about

**Table 1** Effect of amount of bottom ash (100 BSS mesh) on the adsorption of Rose Bengal (concentration  $1 \times 10^{-4}$  M; pH = 7.2) at different temperatures

Amount of bottom ash (g)	Amount of Rose Bengal Adsorbed ( $\times 10^{-6}$ g)		
	30 °C	40 °C	50 °C
0.1	0.140	0.487	0.698
0.2	0.565	1.150	0.146
0.3	1.278	1.855	2.015
0.4	1.838	2.220	2.243
0.5	2.120	2.347	2.380
0.6	2.312	2.358	2.390
0.7	2.393	2.400	2.412
0.8	2.403	2.410	2.425

2.41, 2.90 and 3.40 (mg per gram) of the dye was adsorbed respectively (Fig. 5). The increase in adsorption of the dye with increasing temperature once again confirms the endothermic nature of the ongoing process.

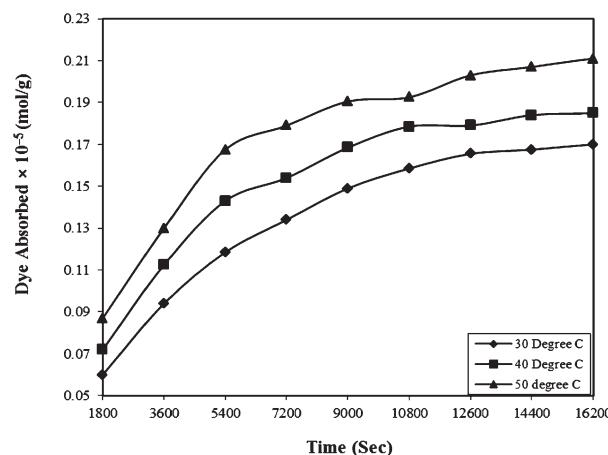
### 3.2. Kinetic studies

**3.2.1. Adsorption rate constant study.** The kinetics of Rose Bengal-bottom ash adsorption was monitored by various kinetic models and it was established that the ongoing adsorption follows a pseudo-second order process. Following Ho and McKay<sup>21</sup>, a pseudo-second order rate expression was applied to calculate the specific rate constant of the adsorption:

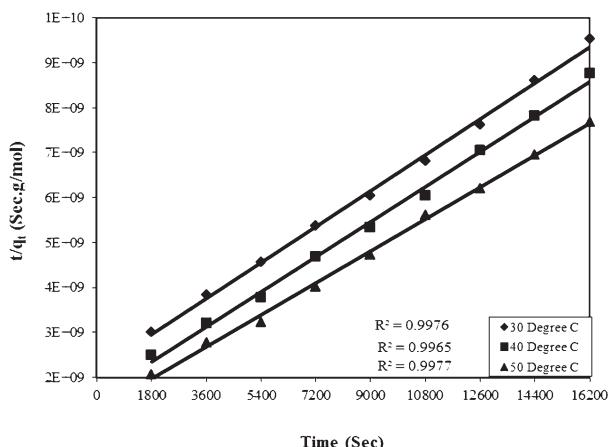
$$\frac{1}{q_t} = \frac{1}{k_2 q_e^2} + \frac{1}{q_e} \quad (1)$$

Where  $q_e$  and  $q_t$  are adsorption capacities (mol g<sup>-1</sup>) at equilibrium (e) and time  $t$  (sec), respectively, and  $k_2$  is the rate constant of the pseudo-second order rate expression (sec g mol<sup>-1</sup>). The kinetic measurements were carried out in  $5 \times 10^{-5}$  M of the dye using 0.5 g of bottom ash in the solution. The plot of  $t/q_t$  against time (Fig. 6) gives straight lines with regression coefficient values equivalent to almost unity, confirming thereby the pseudo-second order kinetics of the ongoing process at all the temperatures. The rate constant for the process was found almost  $2 \times 10^{-9}$  sec g mol<sup>-1</sup> at all temperatures.

**3.2.2. Rate expression and treatment of data.** The interpretation of the experimental data governing the overall rate of adsorption of Rose Bengal over bottom ash was carried out using the mathematical treatment recommended by Boyd *et al.*<sup>22</sup> and Reichenberg.<sup>23</sup> These mathematical treatments were employed to distinguish particle diffusion from a film diffusion process. The theoretical aspects of these models have already been presented elsewhere,<sup>24</sup> while the quantitative treatment of the adsorption can be monitored through following expressions:



**Fig. 5** Effect of contact time on the Rose Bengal (pH = 7.2, concentration =  $5 \times 10^{-5}$  M)-bottom ash (0.50 g and mesh size = 100 BSS) system at different temperatures.



**Fig. 6** Plot of time versus  $t/q_t$  for the Rose Bengal ( $\text{pH} = 7.2$ , concentration  $= 5 \times 10^{-5} \text{ M}$ )-bottom ash (0.50 g and mesh size = 100 BSS) system at different temperatures.

$$F = 1 - \frac{6}{\pi^2} \sum_1^{\infty} \left( \frac{1}{n^2} \right) \exp(-n^2 B_t) \quad (2)$$

Where,  $n$  is the Freundlich constant and  $F$  is the fractional attainment of equilibrium at time  $t$ , and is obtained by using the following equation:

$$F = \frac{Q_t}{Q_{\infty}} \quad (3)$$

Where,  $Q_t$  and  $Q_{\infty}$  are amounts adsorbed after time  $t$  and after infinite time respectively. The other parameter ' $B_t$ ' is known as time constant and obtained by following expression:

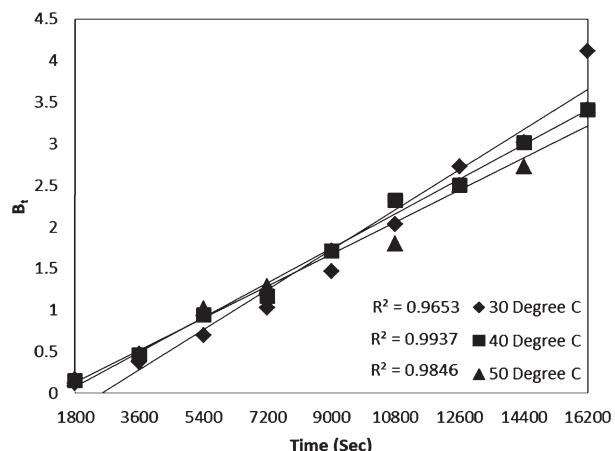
$$B_t = \frac{\pi^2 D_i}{(r_o)^2} = \text{Time Constant} \quad (4)$$

Where,  $B_t$  = time constant,  $D_i$  = effective diffusion coefficient of adsorbate in the adsorbent phase and  $r_o$  = radius of adsorbent particles assumed to be spherical.

For every observed value of  $F$ , corresponding values of  $B_t$  were derived from Reichenberg's table.<sup>23</sup> The  $B_t$  versus time plot for  $5 \times 10^{-5} \text{ M}$  solution of dye (Fig. 7) was found to be linear and straight lines obtained at different temperature were found to pass almost through the origin. This suggests that the rate determining process governs through particle diffusion at all the temperatures 30, 40, 50 °C.

For the ongoing adsorption, the values of the effective diffusion coefficient  $D_i$  were also calculated at 30, 40 and 50 °C temperatures using the time versus  $B_t$  graphs. The energy of activation ( $E_a$ ), entropy ( $\Delta S^{\#}$ ) and pre-exponential constant ( $D_o$ ), values were calculated using the following equations:

$$D_i = D_o \exp\left(-\frac{E_a}{RT}\right) \quad (5)$$



**Fig. 7** Time versus  $B_t$  plot for Rose Bengal ( $\text{pH} = 7.2$ , concentration  $= 5 \times 10^{-5} \text{ M}$ )-bottom ash (0.50 g, mesh size = 100 BSS) adsorption at different temperatures.

$$D_o = (2.72d^2 KT/h) \exp\left(\frac{\Delta S^{\#}}{R}\right) \quad (6)$$

In the equations given above,  $D_o$  is the pre-exponential constant,  $\Delta S^{\#}$  is the entropy,  $d$  gives the average distance between two successive sites of the adsorbent,  $h$  is the Planck constant,  $K$  the Boltzmann constant,  $E_a$ , the energy of activation,  $T$  is the temperature and  $R$  is the universal gas constant.

At 30, 40 and 50 °C the values of the effective diffusion coefficient ( $D_i$ ) have been found to be  $1.4833 \times 10^{-9}$ ,  $1.3121 \times 10^{-9}$  and  $1.1980 \times 10^{-9}$ , respectively. The decreasing values of effective diffusion coefficient ( $D_i$ ) with respect to increasing temperature clearly indicate that the mobility of the ions decreases due to an increased retardation force acting on the diffusing particles of the dye. The values of the pre-exponential constant ( $D_o$ ), activation energy ( $E_a$ ) and entropy of activation ( $\Delta S^{\#}$ ) for the diffusion of Rose Bengal adsorbing over bottom ash have also been calculated and found as  $4.68 \times 10^{-11}$ ,  $8.713 \text{ kJ mol}^{-1}$  and  $478.145 \text{ J K}^{-1}\text{mol}^{-1}$ , respectively. The negative values of  $\Delta S^{\#}$  obtained for both the systems reveal that the internal structure of the bottom ash do not go through any significant change during the adsorption of the dye.

### 3.3. Adsorption isotherm models

The preliminary investigations presented above determine various working conditions to achieve optimum adsorption of the dye. To develop efficient dye removal methodology the results obtained during preliminary investigations are now used to verify various isothermal models, because equilibrium sorption isotherms are fundamentally very significant for the designing of any sorption system. In the present research work, Langmuir, Freundlich, Tempkin and D-R isotherm models have been applied. The information gathered under various isothermal conditions, during the course of investigations, are utilized to evaluate useful thermodynamic parameters, which suggest the feasibility, favorability and spontaneity of the ongoing adsorption process in the dye-adsorbent system. Moreover, endothermic or

exothermic nature and chemical or physical character of the adsorption have also been investigated in the present studies for the adsorption of Rose Bengal over bottom ash.

**3.3.1. The Langmuir adsorption isotherm model.** The well known following linear form of Langmuir's adsorption isotherm equation was applied for the Rose Bengal–bottom ash system:

$$\frac{1}{q_e} = \frac{1}{q_0} + \frac{1}{bq_0C_e} \quad (7)$$

Where,  $q_e$  is the number of moles of solute adsorbed per unit weight at concentration  $C$ ,  $C_e$  is the molar concentration in the bulk-fluid phase.

For the verification of the Langmuir isotherm model,  $1/C_e$  versus  $1/q_e$  graphs were plotted for the data obtained during adsorption of Rose Bengal over bottom ash at temperatures of 30, 40 and 50 °C. Straight lines with appreciable  $R^2$  values were obtained at all the temperatures, confirming thereby the verification of Langmuir adsorption model and involvement of monolayer adsorption in each case. These straight lines are helpful in calculating the Langmuir constant  $b$  and the number of moles of the dye adsorbed per unit weight of the adsorbent ( $q_0$ ) through their slopes and intercepts, respectively (Table 2).

**3.3.2. Thermodynamic parameters.** The Langmuir constant  $b$  thus calculated was used for calculating thermodynamic parameters, like change in free energy ( $\Delta G^\circ$ ), enthalpy ( $\Delta H^\circ$ ) and entropy ( $\Delta S^\circ$ ) during the adsorption process, by applying the following equations:<sup>24–52</sup>

$$\Delta G^\circ = -RT\ln b \quad (8)$$

$$\Delta H^\circ = -R \frac{T_2 T_1}{T_2 - T_1} \times \ln \frac{b_2}{b_1} \quad (9)$$

$$\Delta S^\circ = \frac{H^\circ - G^\circ}{T} \quad (10)$$

**Table 2** Different adsorption isotherm constants for the Rose Bengal (pH = 7.2)–bottom ash (0.50 g, mesh size = 100 BSS) system at different temperatures

Langmuir constants			$b \times 10^4 \text{ L mol}^{-1}$		
$Q_0 \times 10^{-5} \text{ mol g}^{-1}$	30 °C	40 °C	50 °C	30 °C	40 °C
3.68	4.27		4.89	4.25	5.86
Freundlich constants			$K_F \times 10^{-4}$		
$n$	30 °C	40 °C	30 °C	40 °C	30 °C
3.636	3.521		3.636	3.521	3.636
Tempkin constants			$k_2 \times 10^6$		
$k_1 \times 10^{-7} \text{ (L mol}^{-1})$	30 °C	40 °C	30 °C	40 °C	30 °C
7	8		7	8	7
D–R constants			$X_m \times 10^{-5} \text{ mol/g}$		
$-\beta \times 10^{-9} \text{ mol}^2 \text{ J}^{-2}$	30 °C	40 °C	30 °C	40 °C	30 °C
2.0	2.0		2.0	2.0	2.0

Where,  $b$ ,  $b_1$  and  $b_2$  are the equilibrium constants at 30, 40 and 50 °C, respectively.

On the basis of eqn (8) the Gibb's free energy ( $\Delta G^\circ$ ) of the ongoing adsorption process have been calculated as  $-32.647$ ,  $-34.561$  and  $-36.008 \text{ kJ mol}^{-1}$  at 30, 40 and 50 °C, respectively. The negative free energy values indicate that the ongoing adsorption process is spontaneous in nature and with increasing temperature, the adsorption capacity of the bottom ash increases. On the other hand, the average enthalpy ( $\Delta H^\circ$ ) and entropy ( $\Delta S^\circ$ ) of the process have been evaluated as  $18.120 \text{ kJ mol}^{-1}$  and  $51.703 \text{ J K}^{-1} \text{ mol}^{-1}$ , respectively. The positive value of change in enthalpy suggests the endothermic nature of the process, while the positive value for the change in entropy reveals the increased randomness during the adsorption process and indicates a specific affinity of bottom ash towards the Rose Bengal molecule.

A dimensionless separation factor  $r$ , introduced by Weber and Chakravorti<sup>53</sup> was also calculated to determine the feasibility and favourability of the ongoing adsorption process, by using the following equation:

$$r = \frac{1}{1+bC_0} \quad (11)$$

Where,  $b$  is the Langmuir constant and  $C_0$  is the initial concentration. The values of  $r$  were found to be 0.19, 0.14 and 0.13 at 30, 40 and 50 °C respectively. Values of  $r$  that are  $< 1$  at all the temperatures suggest highly favourable adsorption during the ongoing process.

**3.3.3. The Freundlich adsorption isotherm model.** The following equation describing the Freundlich model for the adsorption of solutes from a liquid to a solid surface is applied for the present adsorption system:

$$\log q_e = \log K_F + (1/n) \log C_e \quad (12)$$

Here  $C_e$  denotes the equilibrium concentration ( $M$ ) of the adsorbate and  $q_e$ , the amount adsorbed ( $\text{mol g}^{-1}$ ) and  $K_F$  and  $n$  are the Freundlich constants related to the adsorption capacity and adsorption intensity of the adsorbate–adsorbent system, respectively. Thus, the Freundlich adsorption isotherm plot was sketched by taking  $\log C_e$  on the abscissa and  $\log q_e$  on the ordinate. The experimental data obtained in terms of  $\log C_e$  and  $\log q_e$ , was found to fit straight lines with regression coefficients close to unity. Values of  $K_F$  and  $n$  were derived from the intercepts and slope of these straight lines, respectively and are presented in Table 2.

**3.3.4. The Tempkin adsorption isotherm model.** To monitor the effect of indirect interactions amongst adsorbate particles, the Tempkin isotherm model was applied to the present system. The model assumes that the heat of adsorption of all the molecules in the layer decreases linearly with surface coverage due to adsorbent–adsorbate interactions. The linear form of the Tempkin isothermal model is expressed as:

$$q_e = k_1 \ln k_2 + k_1 \ln C_e \quad (13)$$

Where,  $q_e$  is the amount of adsorbate adsorbed per unit mass of adsorbent at equilibrium ( $\text{mol g}^{-1}$ ),  $C_e$  is the final concentration at equilibrium ( $\text{mol L}^{-1}$ ),  $k_1$  is the Tempkin isotherm energy constant ( $\text{L mol}^{-1}$ ) related to the heat of adsorption and  $k_2$  is the Tempkin isotherm constant. Thus, a plot of  $\ln C_e$  as a function of the amount adsorbed at equilibrium ( $q_e$ ) gave straight lines (Fig. 8) suggesting the uniform distribution of binding energy arising due to interaction of the adsorbate molecules. The straight lines obtained from the graphs were also helpful in determining the Tempkin isotherm constants presented in Table 2.

**3.3.5. The D–R adsorption isotherm model.** In order to identify the nature of the ongoing adsorption process, the following D–R adsorption isotherm model was applied:

$$\ln C_{\text{ads}} = \ln X_m - \beta \varepsilon^2 \quad (14)$$

Where,  $X_m$  is the maximum adsorption capacity,  $\beta$  is the activity coefficient related to mean sorption energy, and  $\varepsilon$  is the Polanyi potential, which is equal to:

$$\varepsilon = RT \ln \left( 1 + \frac{1}{C_e} \right) \quad (15)$$

Where,  $R$  is the gas constant and  $T$  is temperature in Kelvin. The activity coefficient ( $\beta$ ) and the adsorption capacities ( $\ln X_m$ ) were evaluated from the slopes and intercepts of the plot  $\ln C_{\text{ads}}$  versus  $\varepsilon^2$  (Fig. 9) respectively at 30, 40 and 50 °C. The results are depicted in Table 2. The mean sorption energy was calculated from the values of  $\beta$  by following expression:

$$E = \frac{1}{\sqrt{-2\beta}} \quad (16)$$

On the basis of values of  $\beta$ , obtained from the D–R isotherm graph, values of  $E$  are calculated. At all the temperatures the values have been found as 15.8 kJ indicating thereby

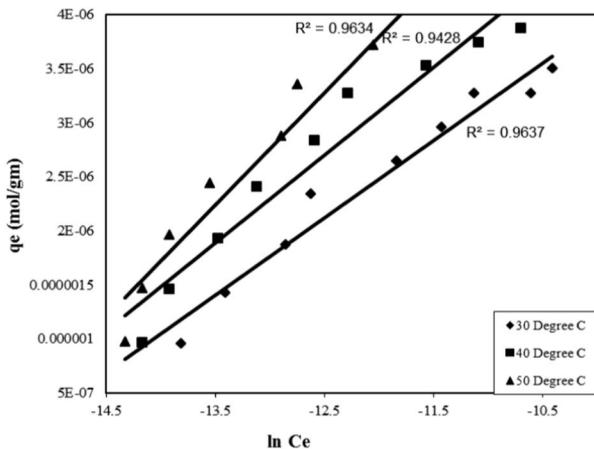


Fig. 8 Tempkin adsorption isotherm for the Rose Bengal ( $\text{pH} = 7.2$ )–bottom ash (0.50 g, mesh size = 100 BSS) system at different temperatures.

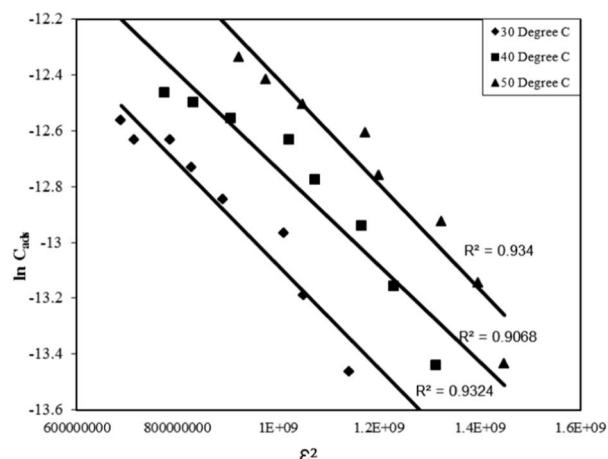


Fig. 9 D–R adsorption isotherm for the Rose Bengal ( $\text{pH} = 7.2$ )–bottom ash (0.50 g, mesh size = 100 BSS) system at different temperatures.

involvement of chemisorption in the present adsorption process at all the temperatures.

### 3.4. Column studies

The practical utility of the developed method was assessed by carrying out bulk removal of the Rose Bengal by adsorbing it over bottom ash in a glass column. A literature survey reveals that various types of columns have been suggested by different workers.<sup>54,55</sup> However, it is specified that in those cases where rate to establish equilibrium of adsorbate concentration between aqueous and adsorbed phase is known, it is advisable to use a fixed bed adsorber as this allows more efficient utilization of the adsorptive capacity than the batch process.<sup>56</sup> Moreover, by fixed bed column studies both adsorption and desorption studies can be made easily. Thus fixed bed column studies were carried out for removal as well as recovery of the Rose Bengal.

In the present studies, parameters like length of primary adsorption zone ( $\delta$ ), time involved in establishing the primary adsorption zone ( $t_x$ ), time required by primary adsorption zone to move down its length ( $t_\delta$ ), time of initial formation of primary adsorption zone ( $t_f$ ), fractional capacity of the prepared column

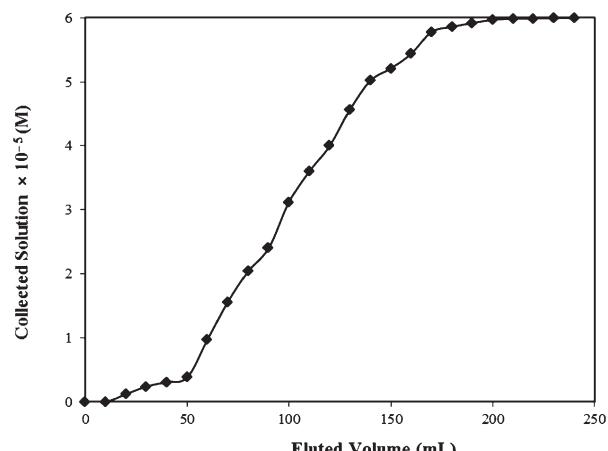


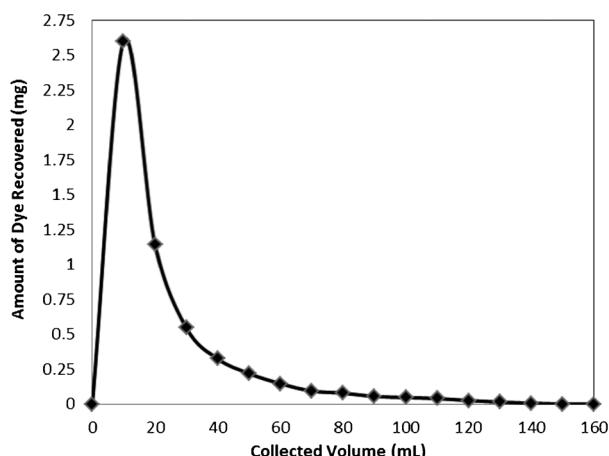
Fig. 10 Breakthrough curve for Rose Bengal–bottom ash column.

**Table 3** Fixed bed adsorber calculations for the adsorption of Rose Bengal in a bottom ash column

$C_0$ (M)	$C_x$ (M)	$C_b$ (M)	$V_x$ (mL)	$V_b$ (mL)	$(V_x - V_b)$ (mL)	$F_m$ (mg cm $^{-2}$ min $^{-1}$ )	$D$ (cm)
$6 \times 10^{-5}$	$0.38 \times 10^{-5}$	$5.78 \times 10^{-5}$	170	50	120	0.0389	1.0

**Table 4** Parameters for fixed bed adsorber–bottom ash for Rose Bengal adsorption

$t_x$ (min)	$t_\delta$ (min)	$t_f$ (min)	$f$	$\delta$ (cm)	Percentage Saturation
4372.423	3086.416	100	0.968	0.722	97.65

**Fig. 11** Recovery of the dye Rose Bengal from the bottom ash column.

( $f$ ), mass rate of flow to the adsorbent ( $F_m$ ) and percentage saturation of column at break point were calculated by using well-known equations.<sup>57</sup>

**3.4.1. Column adsorption.** In the present studies  $6 \times 10^{-5}$  M solution of Rose Bengal was passed at a flow rate of  $0.5 \text{ ml min}^{-1}$  through the bottom ash column and a breakthrough curve was obtained by collecting 240 mL drained out solution of Rose Bengal (Fig. 10). Out of 14.65 mg of the dye taken in the original solution, 5.85 mg was adsorbed over bottom ash. Different column parameters were calculated in the column studies and are presented in Tables 3 and 4.

**3.4.2. Column regeneration, dye recovery and adsorption efficiency.** The column was regenerated and the dye was recovered using dilute NaOH, maintaining a flow rate of  $0.5 \text{ ml min}^{-1}$  and a desorption curve was sketched (Fig. 11). It was observed that first three aliquots of 10 mL each of dilute NaOH was sufficient to desorb almost 80% of the dye from the column and remaining amount of dye was recovered in next 120 mL of the NaOH. Thus total 150 mL of NaOH was found sufficient to recover almost complete dye from the column and 91.24% percentage recovery was achieved.

The adsorption efficiency was determined by reloading the column with dye solution of same concentration and breakthrough capacities were found as 85, 77, 66 and 54% during first, second, third, and fourth cycles, respectively.

## 4. Conclusions

The studies presented above clearly exhibit that the removal of Rose Bengal from its aqueous solutions can be very efficiently achieved through batch and column operations by activated bottom ash at all the temperatures. It was also found that the adsorption is very much dependent upon parameters like pH, contact time, amount of adsorbent, sieve size of adsorbent adsorbate concentration and temperature. The adsorption exhibit a pseudo second order rate kinetics and the value of rate constants at all the temperatures were found almost similar, *viz.*  $2 \times 10^{-9}$ . The contact time studies also exhibit that the ongoing adsorption process follow particle diffusion mechanism and very promising results were obtained for the activation energy and entropy of the process. Other thermodynamic parameters like change in enthalpy, entropy and free energy were also calculated through Langmuir constants and it was found that the process is spontaneous and feasible. As exhibited by the straight lines obtained at different temperatures, it was also concluded that the adsorption of the Rose Bengal over bottom ash follows Langmuir, Freundlich, Tempkin and D-R adsorption isotherms. The excellent adsorption ability of bottom ash was also ascertained by the column operations and a percentage saturation of almost 97% was achieved. Desorption studies also gave 91% of adsorbed dye from the column after eluting with dilute NaOH.

## Acknowledgements

One of the authors, DJ, thanks MANIT for providing financial assistance to carry out this work.

## References

- 1 *Saunders Comprehensive Veterinary Dictionary* (2007) 3rd Edition, Elsevier, Inc.
- 2 H. M. Tabery, *Acta Ophthalmol. Scand.*, 1998, **76**(2), 142–145.
- 3 Y. C. Lee, C. K. Park, M. S. Kim and J. H. Kim, *Cornea*, 1996, **15**(4), 376–385.
- 4 <http://www.sciencelab.com/msds.php?msdsId=9924832>.
- 5 V. K. Gupta and Suhas, *J. Environ. Manage.*, 2009, **90**, 2313–2342.
- 6 I. Ali and V. K. Gupta, *Nat. Protoc.*, 2007, **1**(6), 2661–2667.
- 7 S. Karthikeyan, V. K. Gupta, R. Boopathy, A. Titus and G. Sekaran, *J. Mol. Liq.*, 2012, **173**, 153–163.
- 8 V. K. Gupta, I. Ali, T. A. Saleh, A. Nayak and S. Agarwal, *RSC Adv.*, 2012, **2**, 6380–6388.
- 9 P. Nigam, G. Armour, I. Banat, M. D. Singh and R. Merchant, *Bioprocess Technol.*, 2000, **72**, 219–226.
- 10 T. A. Saleh and V. K. Gupta, *Environ. Sci. Pollut. Res.*, 2012, **19**, 1224–1228.
- 11 Z. M. Shen, D. Wu, J. Yang, T. Yuan, W. H. Wang and J. P. Jia, *J. Hazard. Mater.*, 2006, **131**, 90–97.
- 12 R. Jain and M. Shrivastava, *J. Hazard. Mater.*, 2008, **152**, 216–220.
- 13 N. C. G. Tan, F. X. Prenafeta-Boldu, J. L. Opsteeg, G. Lettinga and J. A. Field, *Appl. Microbiol. Biotechnol.*, 1999, **51**, 865–871.
- 14 W. J. Weber Jr, *Physicochemical Processes for Water Quality Control*. Wiley-Interscience, New York, 1972.
- 15 D. Li and H. Wang, *J. Mater. Chem.*, 2010, **20**, 4551–4566.
- 16 P. Garu, *Wat. Sc. Technol.*, 2005, **24**, 97–103.

17 *Millodot: Dictionary of Optometry and Visual Science*, 7th Edition. Butterworth-Heinemann, 2009.

18 M. Ahmaruzzaman and V. K. Gupta, *Ind. Eng. Chem. Res.*, 2011, **50**, 13589–13613.

19 C. Y. Lin and D. H. Yang, *J. Environ. Sci. Health, Part A: Toxicol. Hazard. Subst. Environ. Eng.*, 2002, **37**, 1509–1522.

20 F. Goodarzi and F. E. Huggins, *J. Environ. Monit.*, 2001, **3**, 1–6.

21 Y. S. Ho and G. McKay, *Trans. Inst. Chem. Eng.*, 1998, **76**, 313–318.

22 G. E. Boyd, A. W. Adamson and L. S. Meyers, *J. Am. Chem. Soc.*, 1947, **69**, 2836–2848.

23 D. Reichenberg, *J. Am. Chem. Soc.*, 1953, **75**, 589–597.

24 V. K. Gupta, S. K. Srivastava and D. Mohan, *Ind. Eng. Chem. Res.*, 1997, **36**, 2207–2218.

25 V. K. Gupta, R. Prasad and A. Kumar, *Talanta*, 2003, **60**, 149–160.

26 V. K. Gupta, R. Mangla, U. Khurana and P. Kumar, *Electroanalysis*, 1999, **11**, 573–576.

27 A. K. Jain, V. K. Gupta, L. P. Singh and U. Khurana, *Analyst*, 1997, **122**, 583–586.

28 V. K. Gupta, S. K. Srivastava and R. Tyagi, *Water Res.*, 2000, **34**, 1543–1550.

29 V. K. Gupta, R. Mangla and S. Agarwal, *Electroanalysis*, 2002, **14**, 1127–1132.

30 A. K. Jain, V. K. Gupta, U. Khurana and L. P. Singh, *Electroanalysis*, 1997, **9**, 857–860.

31 A. K. Jain, V. K. Gupta and L. P. Singh, *Anal. Proc.*, 1995, **32**, 263–265.

32 A. K. Jain, V. K. Gupta, B. B. Sahoo and L. P. Singh, *Anal. Proc.*, 1995, **32**, 99–101.

33 S. K. Srivastava, V. K. Gupta, M. K. Dwivedi and S. Jain, *Anal. Proc.*, 1995, **32**, 21–23.

34 V. K. Gupta and A. Rastogi, *J. Colloid Interface Sci.*, 2010, **342**, 135–141.

35 R. N. Goyal, V. K. Gupta, A. Sangal and N. Bachheti, *Electroanalysis*, 2005, **17**, 2217–2223.

36 V. K. Gupta, A. Mittal and L. Krishnan, *J. Hazard. Mater.*, 2005, **117**, 171–178.

37 V. K. Gupta, A. K. Jain, P. K. S. Agarwal and G. Maheshwari, *Sens. Actuators, B*, 2006, **113**, 182–186.

38 V. K. Gupta, A. Mittal, L. Kurup and J. Mittal, *J. Colloid Interface Sci.*, 2006, **304**, 52–57.

39 V. K. Gupta, A. Mittal, L. Krishnan and J. Mittal, *J. Colloid Interface Sci.*, 2006, **293**, 16–26.

40 V. K. Gupta, R. Jain, and S. Varshney, *J. Hazard. Mater.*, 2007, **142**, 443–448.

41 R. N. Goyal, V. K. Gupta, M. Oyama and N. Bachheti, *Talanta*, 2007, **72**, 976–983.

42 V. K. Gupta, R. Jain, and S. Varshney, *J. Colloid Interface Sci.*, 2007, **312**, 292–296.

43 V. K. Gupta, I. Ali and V. K. Saini, *Water Res.*, 2007, **41**, 3307–3316.

44 V. K. Gupta, A. K. Singh and B. Gupta, *Anal. Chim. Acta*, 2007, **583**, 340–348.

45 R. N. Goyal, V. K. Gupta and N. Bachheti, *Talanta*, 2007, **71**, 1110–1117.

46 V. K. Gupta, A. Mittal, R. Jain, M. Mathur and S. Sikarwar, *J. Colloid Interface Sci.*, 2009, **163**, 396–402.

47 V. K. Gupta, R. N. Goyal, and R. A. Sharma, *Int. J. Electrochem. Sci.*, 2009, **4**, 156–172.

48 V. K. Gupta, A. Mittal, A. Malviya and J. Mittal, *J. Colloid Interface Sci.*, 2009, **335**, 24–33.

49 R. N. Goyal, V. K. Gupta and S. Chatterjee, *Biosens. Bioelectron.*, 2009, **24**, 1649–1654.

50 V. K. Gupta, M. Al Khayat, A. K. Singh and Manoj. K. Pal, *Anal. Chim. Acta*, 2009, **634**, 36–43.

51 R. N. Goyal, V. K. Gupta, Neeta Bachheti and R. A. Sharma, *Electroanalysis*, 2008, **20**, 757–764.

52 R. N. Goyal, M. Oyama, V. K. Gupta, S. P. Singh and S. Chatterjee, *Sens. Actuators, B*, 2008, **134**, 816–821.

53 T. W. Weber and R. K. Chakravorti, *AIChE J.*, 1974, **20**, 228–238.

54 H. J. Fornwalt and R. A. Hutchins, *J. Chem. Eng.*, 1966, **73**, 179–186.

55 A. S. Michaels, *Ind. Eng. Chem.*, 1952, **44**, 1922–1930.

56 V. Sarin, T. S. Singh and K. K. Pant, *Bioresour. Technol.*, 2006, **97**(16), 1986–1993.

57 W. A. Johnston, *Chem. Eng.*, 1972, **79**, 87–92.