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Effect of dispersed hydrophilic silicon dioxide nanoparticles on batch adsorption of benzoic acid from aqueous solution using modified natural vermiculite: An equilibrium study

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Dedicated to the memory of our dear colleague Prof. Seyed Jalealudin Hashemi, who passed away before the submission of this article.

Abstract

The equilibrium adsorption of benzoic acid from an aqueous medium on a natural vermiculite-based adsorbent was studied in the presence and absence of hydrophilic silicon dioxide nanoparticles in batchwise mode. The adsorbent was prepared through grinding natural vermiculite in a laboratory vibratory disk mill and the surfactant modification of ground vermiculite by cetyltrimethylammonium bromide, subsequently. The equilibrium isotherm in the presence and absence of nanoparticles was experimentally obtained and the equilibrium data were fitted to the Langmuir, Freundlich, Dubinin–Radushkevich and Temkin models. The results indicated that the dispersion of silicon dioxide nanoparticles at optimum concentration in the liquid phase remarkably increases the removal efficiency. Furthermore, it yields a more favorable equilibrium isotherm and changes the compatibility of equilibrium data from the Langmuir and Temkin equations to just the Langmuir equation. A quadratic polynomial model predicting the equilibrium adsorbent capacity in the presence of nanoparticles as a function of the adsorbate and initial nanoparticle concentrations was successfully developed using the response surface methodology based on the rotatable central composite design. A desirability function was used in order to optimize the values of all variables, independent and dependent ones, simultaneously.

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Keywords: Adsorption; Benzoic acid; Vermiculite; Cetyltrimethylammonium bromide; Silicon dioxide nanoparticles; Response surface methodology; Desirability function

1. Introduction

The burgeoning trend of industrialization has contributed to the discharge of a great deal of pollutants into our environment. Aromatic carboxylic acids (ACAs) are a sort of organic

compounds whose occurrence in industrial waste effluents originates not only from the disposal of finished products and undesirable by-products, but also from the microbial oxidation of hydrocarbons. Because of the poor biodegradability of ACAs arising out of the electron-withdrawing carboxyl group on their phenyl ring, the wastewaters involved must be purified prior to discharge, as recommended by the European Environment Agency (Huang, Wang, & Huang, 2011; Knepper et al., 1999). Benzoic acid (BA) as the simplest ACA with widespread applications in many industrial sectors can serve as a suitable model compound to make a study of eliminating ACAs

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Nomenclature

a_0, a_i and a_{ij}	adjustable parameters of RSM-based model
A_T	Temkin isotherm equilibrium constant (L/mg)
b_T	Temkin isotherm constant (kJ/mol)
C_0	initial concentration of BA (mg/L)
C_e	equilibrium concentration of BA (mg/L)
C_i	initial concentration of nanoparticles (mg/L)
D	overall desirability function
d_i	individual desirability function
k	number of manipulated variables
K_{D-R}	Dubinin–Radushkevich isotherm constant (mol^2/kJ^2)
K_F	Freundlich isotherm constant ($(\text{mg/g})(\text{L/mg})^{1/n}$)
K_L	Langmuir isotherm constant (L/mg)
L_i	lower limit
M	adsorbent mass (g)
N	number of data points
n	number of variables
n_0	number of central point replications
n_F	Freundlich adsorption intensity constant
p, q	importance weight
q_{D-R}	Dubinin–Radushkevich adsorption capacity (mg/g)
q_e	equilibrium adsorption capacity (mg/g)
$q_{e,\text{cal}}$	calculated adsorption capacity at equilibrium (mg/g)
$q_{e,\text{exp}}$	experimental adsorption capacity at equilibrium (mg/g)
q_L	Langmuir adsorption capacity (mg/g)
R	universal gas constant (kJ/mol K)
T	temperature (K)
T_i	target value
U_i	upper limit
V	solution volume (L)
X_i	coded manipulated variable
x_i	uncoded manipulated variable
$x_{i,\text{max}}$	upper level of an uncoded manipulated variable
$x_{i,\text{min}}$	lower level of an uncoded manipulated variable
Y	predicted response
z_i	variable

from aquatic media (Giannakas, Antonopoulou, Deligiannakis, & Konstantinou, 2013).

The conventional processes that have been thus far utilized for removing BA from aqueous solution include photocatalytic degradation (Gandhi et al., 2011), electrochemical oxidation (Garg & Prasad, 2015), microbial filtration (Zachritz, Lundie, & Wang, 1996) and adsorption (Xin et al., 2011; Yan et al., 2007; Yıldız, Gönülşen, Koyuncu, & Çalımlı, 2005). From the practical perspective, the adsorption process is approved as a highly selective, facile and economic in terms of design and operation and recyclable method for the removal of a vast array of organic and inorganic contaminants (Qu, 2008).

Nowadays, natural clay minerals, like vermiculite, because of being inexpensive, abundant and eco-friendly with a high capability for ion exchange, have drawn much attention either as direct adsorbent, for example, heavy metals (Gupta & Bhattacharyya, 2011) and a variety of dyes (Yagub, Sen, Afroze, & Ang, 2014), or as host matrices to fabricate effective adsorbents concerning low and high molecular weight organic acid species like BA (Borisover & Davis, 2015). The modification of clay minerals by substituting their inorganic interlamellar cations with quaternary ammonium cations using cationic surfactants like cetyltrimethylammonium bromide (CTAB) is one of the most common methods to decrease the high degree of hydrophilicity and the high concentration of negative structural charge of clay minerals, thereby improving their performance for the adsorption of organic acids from aqueous solution (Anirudhan & Ramachandran, 2006; De Araújo Medeiros, Sansiviero, Araújo, & Lago, 2009; Su, Lin, Wang, Xie, & Chen, 2011; Xin et al., 2011; Yan et al., 2007; Yıldız et al., 2005). There are also reports on the increase of cation exchange capacity, porosity and surface chemical reactivity of clay minerals after grinding by a vibratory disk mill, which favors their susceptibility to be modified organically (Filipović-Petrović, Kostić-Gvozdenović, & Erić-Antonić, 2002; Hongo, Yoshino, Yamazaki, Yamasaki, & Satokawa, 2012; Jiménez De Haro et al., 2004).

Over the past decade, the nanotechnology revolution has opened up a promising era in numerous fields of science (Choolaei, Rashidi, Ardjmand, Yadegari, & Soltanian, 2012) (Banaei, Vojoudi, Karimi, Bahar, & Pourbasheer, 2015). In this regard, several studies have been recently carried out to increase the efficiency of separation processes like liquid–liquid extraction (Bahmanyar, Khoobi, Mozdianfard, & Bahmanyar, 2011; Khoobi et al., 2013; Raji-Asadabadi, Abolghasemi, Maragheh, & Davoodi-Nasab, 2013) and liquid chromatography (Nilsson, Birnbaum, & Nilsson, 2007; Zhang, Wang, Liao, & Liu, 2006) in the presence of dispersed nanomaterials (typically nanoparticles). Although these studies are in their infancy, they have shown the potential of nanomaterials as multifunctional components to improve the separation efficiency. All the foregoing studies are focused on the increase of mass transfer in nanofluids. The enhanced mass transfer of several substances in water-based nanofluids in comparison with pure water has been reported by a number of scholars (Ashrafmansouri & Nasr Esfahany, 2014; Krishnamurthy, Bhattacharya, Phelan, & Prasher, 2006; Veilleux & Coulombe, 2011).

To our best knowledge, the present study is one of few investigations on the effect that the dispersion of nanoparticles in the adsorption medium has on the removal efficiency and the adsorbent capacity. In 2010, Jasper et al. (2010) evaluated the effect of dispersed silicon dioxide, titanium dioxide and iron(III) oxide nanoparticles on the batch adsorption of trichloroethylene from contaminated water by activated carbon. They found that silicon dioxide nanoparticles do not have any remarkable effect on the adsorption efficiency whereas the two other nanoparticles remarkably increase the adsorption efficiency. Silicon dioxide nanoparticles are one of the most common nanoparticles by reason of their low cost and commercial availability. The good

biocompatibility of these nanoparticles makes them applicable to processes dealing with environmental issues (Sun, Yan, Yang, & Sun, 2006; Sun et al., 2007).

Today, multivariate statistical techniques are favored because they are time and cost efficient for experimentation and because they facilitate probing in the effect of inputs, alone and in combination, on the outputs in a multivariable system. Response surface methodology (RSM) is one of the most useful multivariate statistical techniques, particularly when a few variables are involved. This methodology is on the basis of fitting a polynomial model to a series of experimental data with the aim of making statistical previsions (Bezerra, Santelli, Oliveira, Villar, & Escaleira, 2008; Jain, Garg, & Kadirvelu, 2011). Since the most successful development of an RSM-based model is intertwined with the most detailed knowledge of process, an experimental design is essential to ascertain which experiments need to be performed in the experimental region being studied. The central composite design is one of the most standard experimental designs applied for establishing mathematical models estimating the first- and second-order effects in a multivariable system (Montgomery & Runger, 2010; Murugesan et al., 2014). The application of desirability function approaches in applied and analytical chemistry in order to find the best operating conditions and obtain the best result has attracted much attention in recent years (Bezerra et al., 2008).

In brief, this research is aimed at: (1) optimizing the concentration of dispersed hydrophilic silicon dioxide nanoparticles for the batch adsorption of BA from aqueous solution on mechanically and chemically modified natural vermiculite, (2) comparing the equilibrium isotherm in the presence and absence of nanoparticles, (3) analyzing the equilibrium data by four frequently used isotherm models (the Langmuir, Freundlich, Dubinin–Radushkevich and Temkin models), (4) developing an RSM-based model to study how the initial concentrations of adsorbate and nanoparticles influence the adsorption capacity and, lastly, (5) finding the best local conditions using a desirability function optimization approach.

2. Materials and methods

2.1. Materials

Natural vermiculite was collected from a mineral deposit in Kaleybar County (East Azerbaijan Province, Iran). BA ($C_7H_6O_2$) and CTAB ($C_{19}H_{42}BrN$) of analytical grade were purchased from Merck and Sigma–Aldrich companies. Highly pure hydrophilic silicon dioxide (SiO_2) nanoparticles were supplied by TECNAN Company. The field emission scanning electron microscopy (FE-SEM) analysis was conducted using a scanning electron microscope (TESCAN, MIRA3 LM) in order to investigate the surface morphology of silicon dioxide nanoparticles. The X-ray diffraction (XRD) analysis was performed using an X-ray diffractometer (PANalytical, X'Pert Pro MPD) in order to investigate the crystalline/amorphous structure of nanoparticles. The inductively coupled plasma mass spectroscopy (ICP-MS) analysis was performed by means of a mass spectrometer (Varian, OES-730) in order to determine the quantitative chemical

Table 1

Some characteristics of the silicon dioxide nanoparticles.

Morphology	Purity	Average particle size	BET surface area	Total pore volume	Average pore diameter
Spherical	>99.9%	12.80 nm	162.9 m ² /g	0.549 cm ³ /g	11.01 nm

composition of impurities in the sample of silicon dioxide nanoparticles. The dynamic light scattering (DLS) analysis was conducted by means of a dynamic light scattering analyzer (Microtrac, Nanotracc Wave) in order to determine the average particle size and also the size distribution of nanoparticles. The textural characteristics (the BET surface area, the total pore volume and the average pore diameter) of silicon dioxide nanoparticles were obtained by the N_2 adsorption/desorption technique at $-196^\circ C$ using a gas adsorption analyzer (BEL Japan, BELSORP-mini II). Some characteristics of the used nanoparticles are listed in Table 1. The silicon dioxide nanofluid was prepared by dispersing a desired amount of nanoparticles in distilled water using an ultrasonicator at a frequency of 20 kHz for 2 hr.

2.2. Preparation of adsorbent

Initially, the natural vermiculite was reduced in size with a knife mill, passed through a 120-mesh sieve, and subsequently ground in a vibratory disk mill at 2000 rpm for 10 min. Thereafter, 10 g of ground vermiculite was gently contacted with 1000 mL of 23 mM CTAB solution in a beaker at $60^\circ C$ for 24 h using a heater/stirrer. Finally, the solid phase was removed by centrifugation, rinsed with enough deionized water until no detection of bromide ions by 0.1 M silver nitrate solution, dried at $60^\circ C$ for 48 h, activated at $120^\circ C$ for 1 h, pulverized in an agate mortar and packed for further use. The characteristics of the resultant adsorbent, hereinafter designated as CTAB-V, has been thoroughly described in our earlier works (Sadeghi Pouya et al., 2015a; Sadeghi Pouya et al., 2015b).

2.3. Batch adsorption experiments

In general, the adsorption experiments were conducted by placing an optimum amount of CTAB-V samples into 100-mL Erlenmeyer flasks filled with 50 mL of BA solutions at an optimum initial pH value and then shaking the flasks at 200 rpm at a temperature of $30^\circ C$ for 12 h using a thermostatic shaking incubator. After attaining the equilibrium state, the suspensions were centrifuged at 4000 rpm for 15 min and the supernatants were spectrophotometrically analyzed for the remaining BA concentration at a λ_{max} of 227 nm (corresponding to the maximum absorbance) using a Unico UV-2100 UV/vis spectrophotometer. Our preliminary experiments demonstrated that in the studied range of BA and nanoparticles initial concentrations, the optimum adsorbent dosage is 0.5 g/50 mL and the initial solution pH is optimum when not adjusted (i.e. the initial solution pH of 3.5 ± 0.1). The upper and lower limit of BA and nanoparticles initial concentrations was also chosen as 100–1000 mg/L and

1–10 mg/L, respectively. The removal efficiency of BA and the adsorption capacity of CTAB-V at equilibrium were calculated by Eqs. (1) and (2):

$$\text{Equilibrium removal efficiency (\%)} = \frac{C_0 - C_e}{C_e} \times 100, \quad (1)$$

$$q_e = \frac{(C_0 - C_e)V}{M}, \quad (2)$$

where C_0 is the initial concentration of BA; C_e is the equilibrium concentration of BA; q_e is the equilibrium adsorption capacity; V is the solution volume; M is the adsorbent mass.

2.4. RSM modeling approach

To explore an RSM-based correlation between the equilibrium adsorption capacity of CTAB-V in the presence of nanoparticles (as predicted response) and the initial concentrations of BA and nanoparticles (as manipulated variables), the design-expert software package (v. 7.0.0) was employed. A rotatable central composite design was implemented to create an experimental framework totally consisting of 13 runs, as calculated by the expression of $2^n + 2n + n_0$ using five central point replications that is the software default value for a system with two manipulated variables (Arulkumar, Sathishkumar, & Palvannan, 2011; Cho & Zoh, 2007). Given that the manipulated variables in this study were too different in scale, they were normalized in advance of modeling. In this regard, the upper level was coded as +1, the center level as 0 and the lower level as -1. For any value that lies either between the lower limit and the center level or between the center level and the upper limit, Eq. (3) was used as follows (Cho & Zoh, 2007; Karnik, Gaitonde, & Davim, 2007):

$$X_i = \frac{2x_i - (x_{i,\max} + x_{i,\min})}{(x_{i,\max} - x_{i,\min})}, \quad (3)$$

where X_i is the coded manipulated variable; x_i is the uncoded manipulated variable; $x_{i,\max}$ is the upper level of an uncoded manipulated variable; $x_{i,\min}$ is the lower level of an uncoded manipulated variable.

The response surface was related to the coded manipulated variables by a full second-order polynomial equation suggested by Box and Wilson in 1951 as follows (Bezerra et al., 2008; On the Experimental Attainment of Optimum Conditions on JSTOR, n.d.):

$$Y = a_0 + \sum_{i=1}^k a_i X_i + \sum_{i=1}^k \sum_{j=1}^k a_{ij} X_i X_j, \quad (4)$$

where Y is the predicted response; a_0 , a_i , a_{ij} is the adjustable parameters.

Regarding the experimentally obtained data, the model adjustable parameters were calculated by the multiple linear regression analysis using the least-squares optimization method. The model validity was assessed using the analysis of variance (ANOVA) subroutine and by plotting internally studentized

residuals versus run number and also by constructing a normal probability plot of internally studentized residuals.

2.5. Desirability function optimization approach

The desirability function optimization approach is a highly popular optimization technique used to simultaneously determine the optimum values of manipulated variables that can determine the optimum performance levels of one or more responses. During the optimization of a multivariate system, every single variable is converted into an individual desirability function with a value that varies from 0 to 1, the lowest desirability value to the highest desirability value (Islam, Sakkas, & Albanis, 2009; Mourabet et al., 2012). The individual desirability functions are then combined in an overall desirability function as follows (Bezerra et al., 2008; Paterakis, Korakianiti, Dallas, & Rekkas, 2002):

$$D = \left(\prod_{i=1}^n d_i \right)^{1/n}, \quad (5)$$

where D is the overall desirability; d_i is the individual desirability; n is the number of variables.

Depending on whether the target value of a particular variable is to be maximized, minimized or located between the lower limit and the upper limit, the individual desirability function is defined by Eqs. (6)–(8) (Bezerra et al., 2008; John, 2013):

$$d_i = \begin{cases} 0 & \text{if } z_i \leq L_i, \\ \left| \frac{z_i - L_i}{T_i - L_i} \right|^p & \text{if } L_i \leq z_i \leq T_i, \\ 1 & \text{if } z_i \geq T_i, \end{cases} \quad (6)$$

$$d_i = \begin{cases} 1 & \text{if } z_i \geq U_i, \\ \left| \frac{z_i - U_i}{T_i - U_i} \right|^q & \text{if } T_i \leq z_i \leq U_i, \\ 0 & \text{if } z_i \leq T_i, \end{cases} \quad (7)$$

$$d_i = \begin{cases} 1 & \text{if } z_i \leq L_i, \\ \left| \frac{z_i - L_i}{T_i - L_i} \right|^p & \text{if } L_i \leq z_i \leq T_i, \\ \left| \frac{z_i - U_i}{T_i - U_i} \right|^q & \text{if } T_i \leq z_i \leq U_i, \\ 0 & \text{if } z_i \geq U_i, \end{cases} \quad (8)$$

where z_i is the variable; L_i is the lower limit; p , q is the importance weight; U_i is the upper limit; T_i is the target value.

3. Results and discussion

3.1. Characterization analysis of silicon dioxide nanoparticles

The surface morphology of silicon dioxide nanoparticles at two magnification levels (100.0 kx and 150.0 kx) is illustrated in

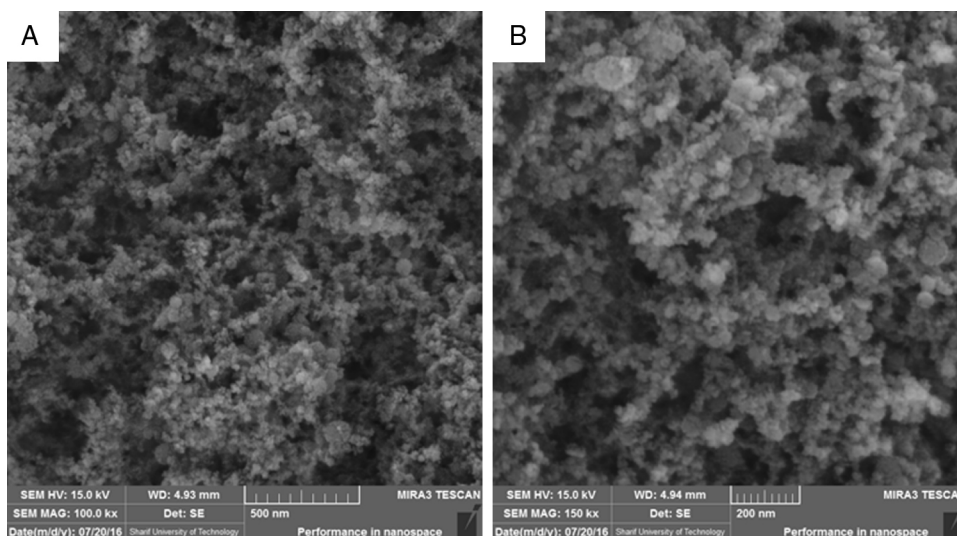


Fig. 1. FE-SEM images of silicon dioxide nanoparticles at: (A) 100.0kx and (B) 150.0kx.

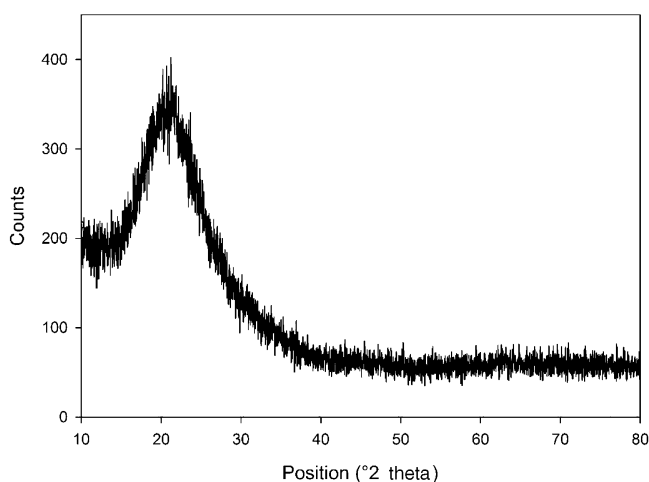


Fig. 2. XRD pattern of silicon dioxide nanoparticles.

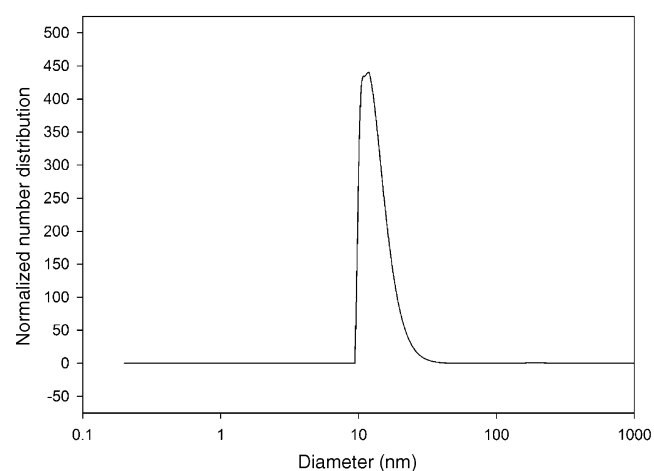


Fig. 3. Size distribution of silicon dioxide nanoparticles dispersed in deionized water.

Fig. 1. As this figure shows, silicon dioxide nanoparticles have a spherical and non-crystalline (or amorphous) morphology.

The XRD pattern of silicon dioxide nanoparticles in the range of $2\theta = 10\text{--}80^\circ$ is shown in Fig. 2. The low-intensity broad peak detected around 20° indicates the presence of a non-crystalline silicon dioxide phase in the sample that is totally in agreement with the FE-SEM images. Similar observations have been reported in previous research works (Fang et al., 2008).

The size distribution of silicon dioxide nanoparticles in semi-logarithmic coordinates is plotted in Fig. 3. This figure clearly shows that most of the nanoparticles lie in the range of 10.0–15.0 nm in diameter. The results of the DLS analysis indicated that half of the silicon dioxide nanoparticles have an average diameter of about 12.80 nm.

The quantitative chemical composition of impurities the sample of silicon dioxide nanoparticles determined by the ICP analysis and is listed in weight percentage in Table 2. A perusal of this table indicates that the purity of silicon dioxide in the sample is higher than 99.9 wt.%.

Table 2

Quantitative chemical composition of impurities in the sample of silicon dioxide nanoparticles (by weight percent).

Cu	0.0000000
Zn	0.0005884
Fe	0.0000000
Mg	0.0000042
Na	0.0000000
Al	0.0000000
As	0.0000000
Sb	0.0000031
Pb	0.0000000
Hg	0.0000000

Fig. 4A and **B** illustrates the N_2 adsorption/desorption isotherm and the pore size distribution of the sample based on Barrett–Joyner–Halenda (BJH) model. The isotherm clearly shows that the hysteresis loop does not form even at high

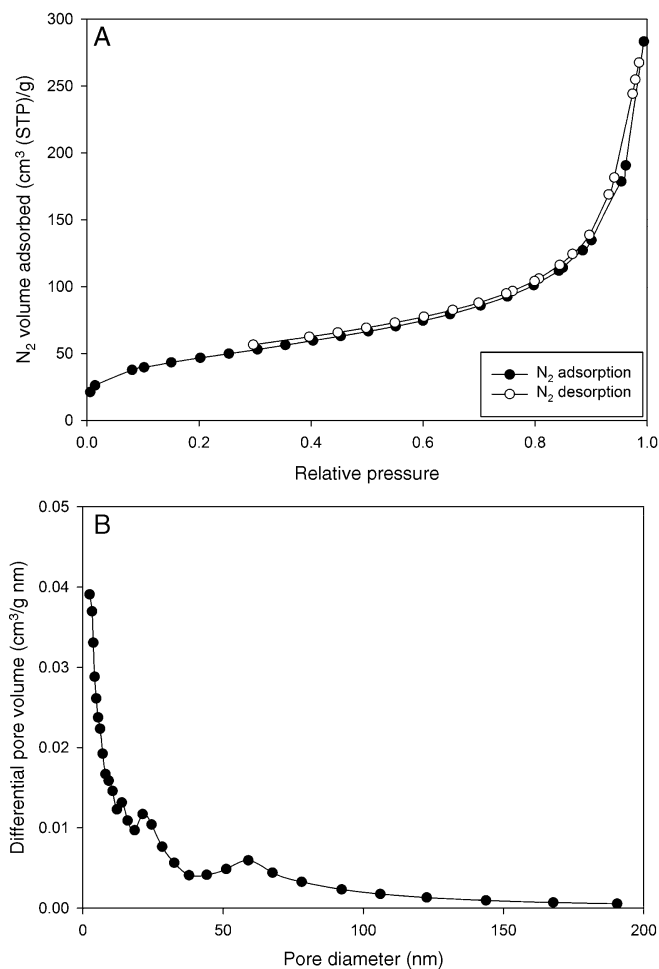


Fig. 4. (A) N_2 adsorption/desorption isotherm and (B) BJH pore size distribution of silicon dioxide nanoparticles.

relative pressures, which suggests that the sample is not mesoporous (Naumov, 2009; Sing, 2001).

3.2. Optimization of nanoparticles initial concentration

The initial concentration of silicon dioxide nanoparticles was optimized by shaking 0.5 g of the CTAB-V samples with 50 mL of 100–1000 mg/L BA solutions at the initial pH value of 3.5 in the presence of various concentrations of dispersed nanoparticles (1, 2, 3, 5, 7.5 and 10 mg/L). Fig. 5 shows the equilibrium removal efficiency as a function of the initial concentration of silicon dioxide nanoparticles and BA. It can be observed from Fig. 5 that the equilibrium removal efficiency peaks at a certain initial concentration of nanoparticles and completely levels off afterwards. The initial concentration of nanoparticles was found to be optimum at 5 mg/L when the initial concentration of BA ranges between 100 and 400 mg/L and to be optimum at 7.5 mg/L when the initial concentration of BA in the range of 400–1000 mg/L. Accordingly, the nanofluid of 7.5 mg/L of the initial concentration was used in order to equalize the conditions for the equilibrium adsorption isotherm experiments. Fig. 6 shows the equilibrium removal efficiency in the presence and absence of nanoparticles. As it displayed in Fig. 6,

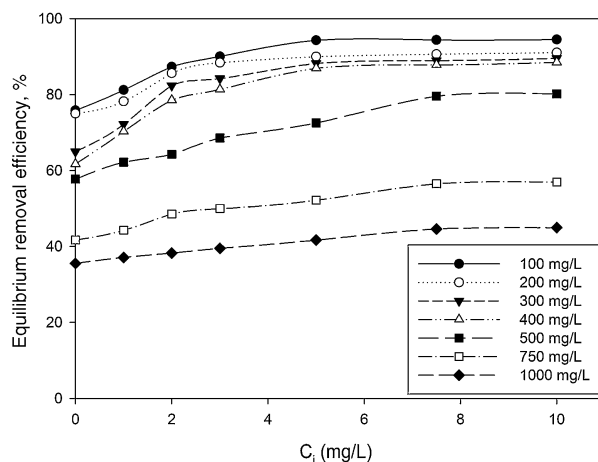


Fig. 5. The effect of initial concentration of silicon dioxide nanoparticles on the equilibrium removal efficiency at different initial concentrations of BA.

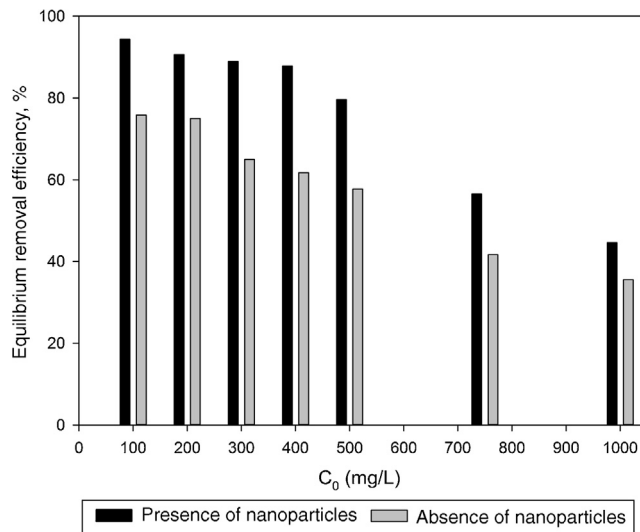


Fig. 6. The equilibrium removal efficiency of BA adsorption onto CTAB-V in the presence of 7.5 mg/L silicon dioxide nanoparticles and in their absence.

the equilibrium removal efficiency increases in the range of about 9–31.5% in the presence of 7.5 mg/L silicon dioxide nanoparticles in comparison with their absence when the initial concentration of BA varies from 100 to 1000 mg/L. Since the colloidal stability of nanofluid has an extremely important effect on the performance of nanoparticles, the relative stability of 7.5 mg/L nanofluid was investigated as a function of the elapsed time using the UV/vis spectrophotometer method, as stated by Bahmanyar et al. (2011). For this purpose, the absorbency of nanofluid with an initial concentration of 7.5 mg/L was continually determined at a λ_{max} of 325 nm (corresponding to the maximum absorbance) over 12 h that is the required contact time for the equilibrium establishment. As the results are illustrated in Fig. 7, the relative concentration of suspended nanoparticles (the concentration at time t divided by the initial concentration) reduces less than 7% within 12 h, which is indicative of an adequate stability of nanofluid during the adsorption experiments.

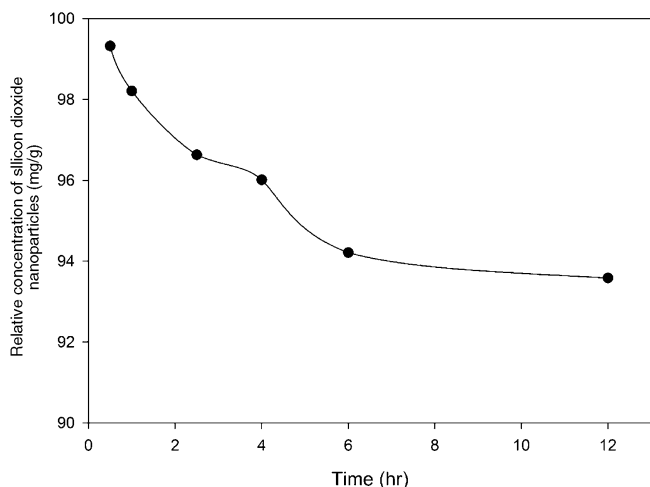


Fig. 7. The relative concentration of suspended silicon dioxide nanoparticles versus elapsed time.

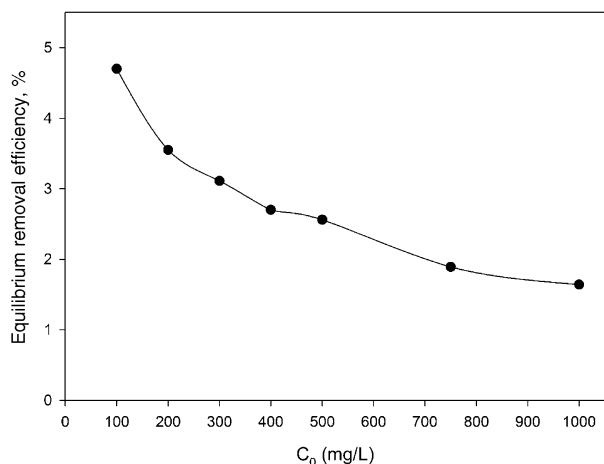


Fig. 8. The effect of 7.5 mg/L silicon dioxide nanoparticles on the equilibrium removal efficiency in the absence of CTAB-V.

The first probable explanation for the enhanced removal efficiency in the presence of silicon dioxide nanoparticles was the adsorption of BA by the mentioned nanoparticles. But this hypothesis was rejected because our additional experiments revealed that the dispersed hydrophilic silicon dioxide nanoparticles at the aforementioned concentrations do not have any remarkable affinity toward BA without introducing CTAB-V into the adsorption system. As shown in Fig. 8, the equilibrium removal efficiency of 100–1000 mg BA/L in a 7.5 mg/L silicon dioxide nanofluid in the absence of CTAB-V was found to be up to a maximum of approximately 4.7%. In contrast, Jasper et al. (2010) have found from their adsorbent-free experiments that both titanium dioxide and iron(III) oxide nanoparticles alone act as adsorption sites for up to 60% of trichloroethylene. Hence, it was concluded that the presence of nanoparticles certainly has a positive side effect on the adsorptive behavior of BA. It can be proposed that the increase in BA removal efficiency may correspond to the increase in BA mass diffusion rate in the liquid phase arising from the Brownian motion of nanoparticles that creates micro/nanoscale convection through the interstices

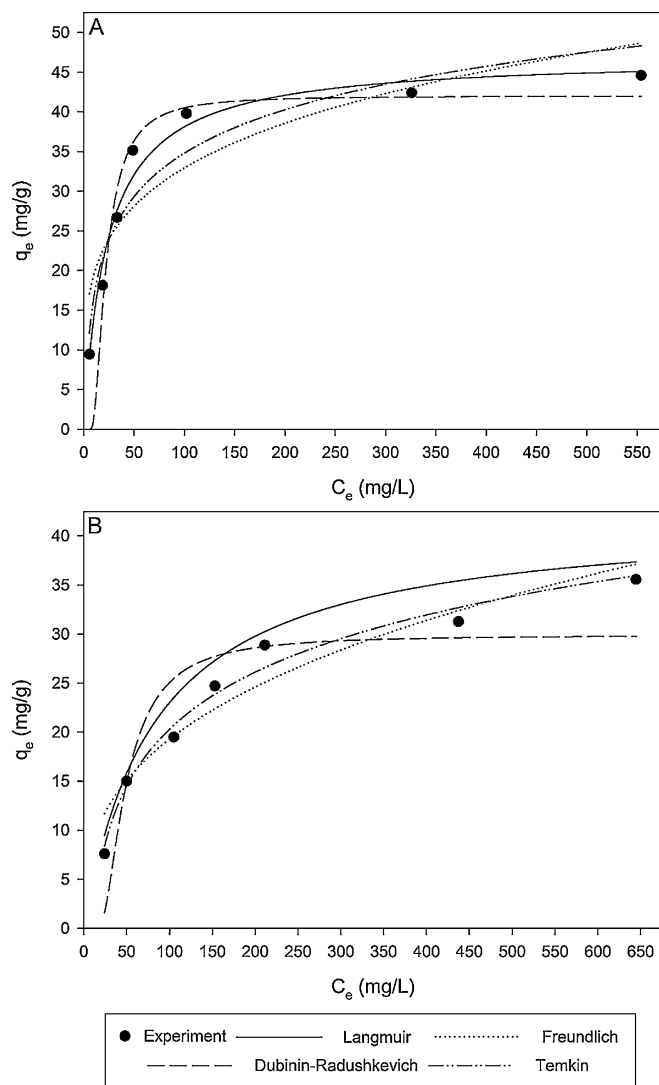


Fig. 9. The equilibrium isotherm of BA adsorption onto CTAB-V at 30 °C: (A) in the presence of 7.5 mg/L silicon dioxide nanoparticles and (B) in the absence of nanoparticles.

of adsorbent particles (Ashrafmansouri & Nasr Esfahany, 2014; Krishnamurthy et al., 2006).

3.3. Equilibrium adsorption isotherm

The Langmuir, Freundlich, Dubinin–Radushkevich and Temkin models were used to analyze the equilibrium data. The mathematical expression of these models is given in Eqs. (5)–(8) (Foo & Hameed, 2010; Ho, Porter, & McKay, 2002):

$$q_e = \frac{q_L K_L C_e}{1 + K_L C_e}, \tag{5}$$

$$q_e = K_F C_e^{1/n_F}, \tag{6}$$

$$q_e = q_{D-R} \exp \left(-K_{D-R} \left[RT \ln \left(1 + \frac{1}{C_e} \right) \right]^2 \right), \tag{7}$$

$$q_e = \frac{RT}{b_T} \ln(A_T C_e), \tag{8}$$

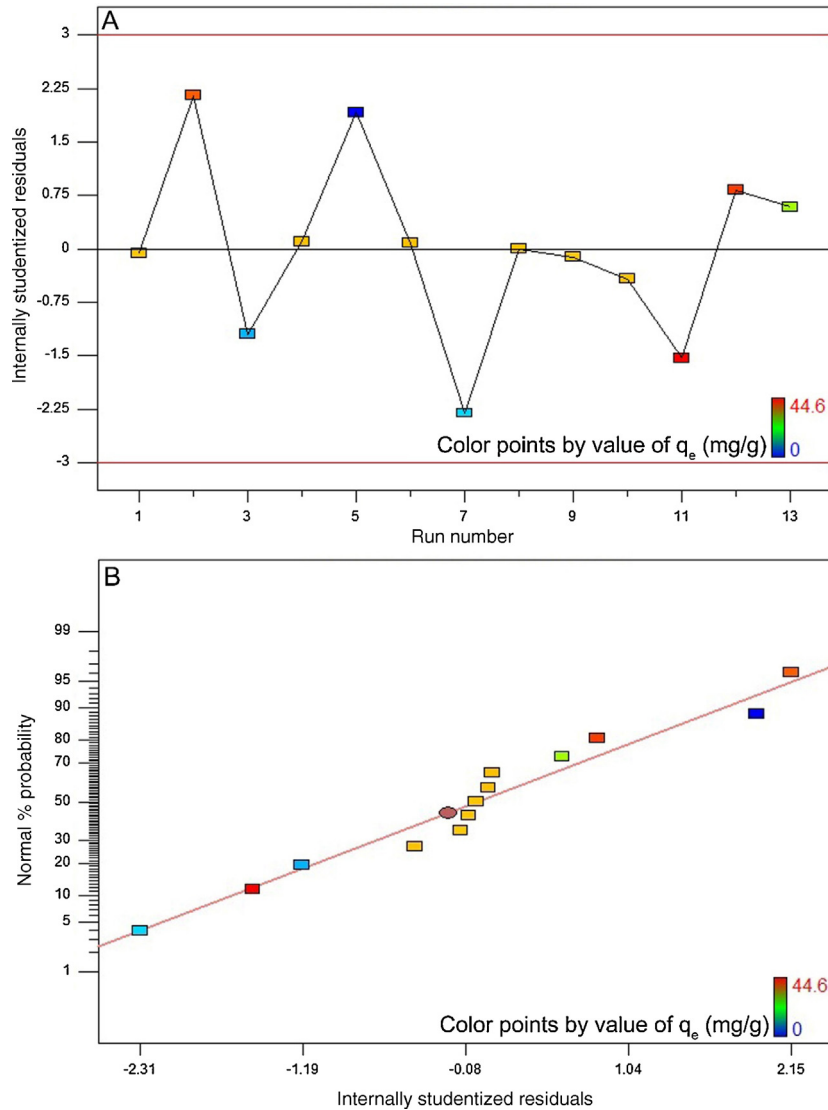


Fig. 10. (A) The internally studentized residuals versus run number and (B) the normally probability plot of the internally studentized residuals.

where q_L is the Langmuir adsorption capacity; K_L is the Langmuir isotherm constant; K_F is the Freundlich isotherm constant; n_F is the Freundlich adsorption intensity constant; q_{D-R} is the Dubinin–Radushkevich adsorption capacity; K_{D-R} is the Dubinin–Radushkevich isotherm constant; R is the Universal gas constant; T is the temperature; b_T is the Temkin isotherm constant; A_T is the Temkin isotherm equilibrium constant.

The adjustable parameters together with the coefficient of determination for each isotherm model was calculated using the MATLAB nonlinear least-squares optimization routine. The values of q_e from the experiments and from the models were also compared based on the average absolute relative deviation (AARD) defined as follows (Foo & Hameed, 2010):

$$\text{AARD (\%)} = \frac{100}{N} \sum_{i=1}^N \left| \frac{q_{e,\text{exp}} - q_{e,\text{cal}}}{q_{e,\text{exp}}} \right|, \quad (9)$$

where N is the number of data points; $q_{e,\text{exp}}$ is the experimental adsorption capacity at equilibrium; $q_{e,\text{cal}}$ is the calculated adsorption capacity at equilibrium.

Fig. 9A and B shows the experimental equilibrium data and the predicted equilibrium isotherms for BA adsorption onto CTAB-V in the presence and absence of nanoparticles. According to Alberti, Amendola, Pesavento, and Biesuz (2012), it is apparent that the shape of experimental isotherm alters from the Langmuir type (without strict plateau) in the absence of silicon dioxide nanoparticles to the high affinity type in the case of nanoparticles presence. This in turn manifests that the adsorption of BA onto CTAB-V takes place more favorably in the presence of nanoparticles (Kyzas & Matis, 2015). Table 3 tabulates the fitting results together with the AARD values for each isotherm model. As can be seen, the equilibrium data are better fitted to the Langmuir and Temkin models in the absence of nanoparticles whereas in the presence of nanoparticles, they are satisfactorily fitted by just the Langmuir model.

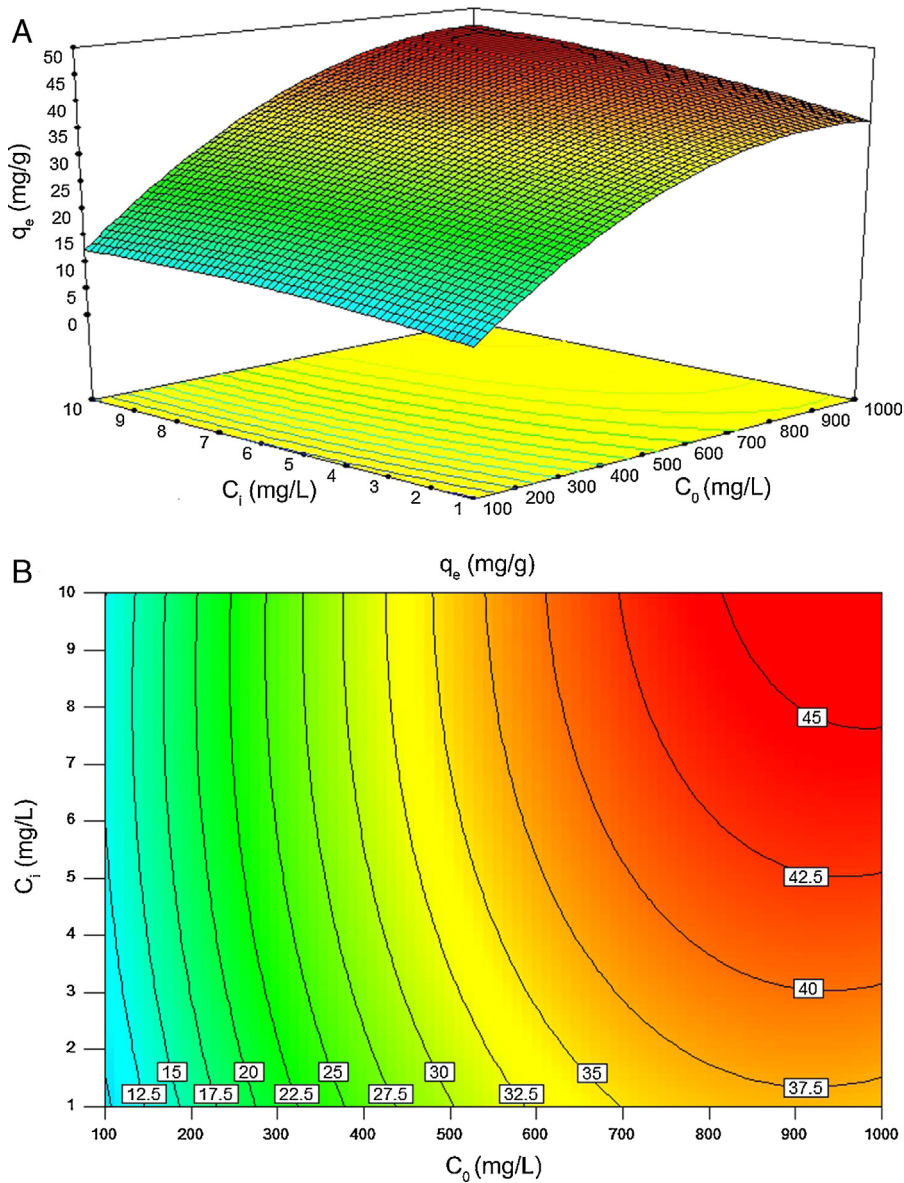


Fig. 11. The effect of BA and silicon dioxide nanoparticles on the equilibrium adsorption capacity of CTAB-V: (A) three dimensional response surface plot and (B) two dimensional contour plot.

3.4. RSM-based model

By fitting the results from the designed experiments to Eq. (4), the model predicting the equilibrium capacity of CTAB-V in terms of coded manipulated variables was derived as follows:

$$q_e = 36.01 + 15.32C_0 + 3.21C_i + 1.74C_0C_i - 8.38C_0^2 - 1.36C_i^2. \quad (10)$$

The high coefficient of determination (0.988) and Fisher value (115.1) verified the model adequacy. Additionally, the probability value was about 0.000. By definition, if the probability value is less than 0.05, the model is statistically considered

significant and the Fisher value is high enough. The latter signifies that the most of variations in the response can be predicted by the model (Jain et al., 2011; Murugesan et al., 2014). Fig. 10A depicts the internally studentized residuals against the run number. It is evident from this figure that the differences between the actual and the model-predicted responses fall within the acceptable range, i.e. from -3 to $+3$ in the context of internally studentized residual (Dopar, Kusic, & Koprivanac, 2011; Jadhav, Surwase, Phugare, & Jadhav, 2012). Therefore, it is possible to conclude that there were no outliers among the experimental data set. Fig. 10B depicts the normal probability plot of internally studentized residuals. It is clear that the points in this figure are appropriately located near to the diagonal straight line. This satisfies the normality assumption. In other words, it corroborates the normal distribution of errors around the mean value and

Table 3
Isotherm model parameters, coefficients of determination and AARD values for BA adsorption onto CTAB-V in the presence of 7.5 mg/L silicon dioxide nanoparticles and in their absence.

Isotherm model	Isotherm parameters	
	Presence of nanoparticles	Absence of nanoparticles
Langmuir	$q_L = 47.32$ $K_L = 0.036$ $R^2 = 0.952$ AARD (%) = 7.021	$q_L = 42.18$ $K_L = 0.012$ $R^2 = 0.991$ AARD (%) = 6.051
Freundlich	$K_F = 10.90$ $n = 4.237$ $R^2 = 0.788$ AARD (%) = 22.03	$K_F = 3.818$ $n = 2.844$ $R^2 = 0.920$ AARD (%) = 12.34
Dubinin–Radushkevich	$q_{D-R} = 43.02$ $K_{D-R} = 93.68$ $R^2 = 0.891$ AARD (%) = 16.14	$q_{D-R} = 29.9$ $K_{D-R} = 280.6$ $R^2 = 0.791$ AARD (%) = 20.72
Temkin	$A_T = 0.754$ $b_T = 0.318$ $R^2 = 0.885$ AARD (%) = 11.72	$A_T = 0.112$ $b_T = 0.303$ $R^2 = 0.984$ AARD (%) = 5.232

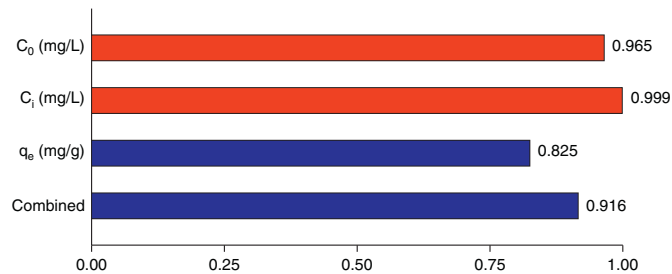


Fig. 12. The individual desirability value of each variable and their combination.

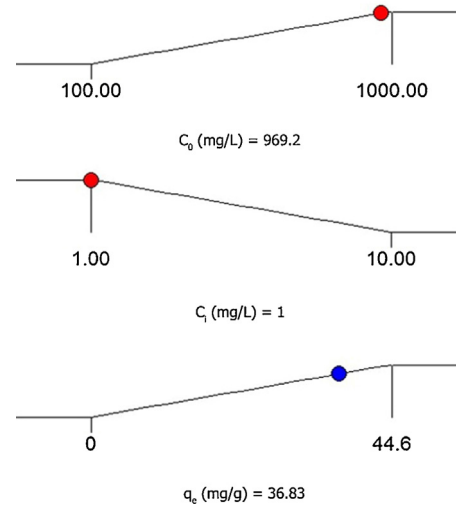


Fig. 13. Desirability value ramps for optimization of variables.

their independency from each other (Dopar et al., 2011; Jadhav et al., 2012). The mean and variance values of the internally studentized residuals in this study were found to be 0.044 and 1.342, respectively. Statistically, as the mean value of a group of data tends to zero and their variance value tends to unity, the distribution of those data is taken to be standard normal (Montgomery & Runger, 2010). Fig. 10A and B further confirms the validity of the suggested model.

Fig. 11A and B illustrates the individual and cumulative impacts of the initial concentrations of BA and nanoparticles on the equilibrium adsorption capacity in the choice of test range in three-dimensional response surface plot and two-dimensional contour plot, respectively. The plots represent that the adsorption capacity increases by increasing the initial concentrations of both BA and nanoparticles. Fig. 11A shows that the initial concentration of BA has a more profound effect on the response amplitude than the initial concentration of nanoparticles. The

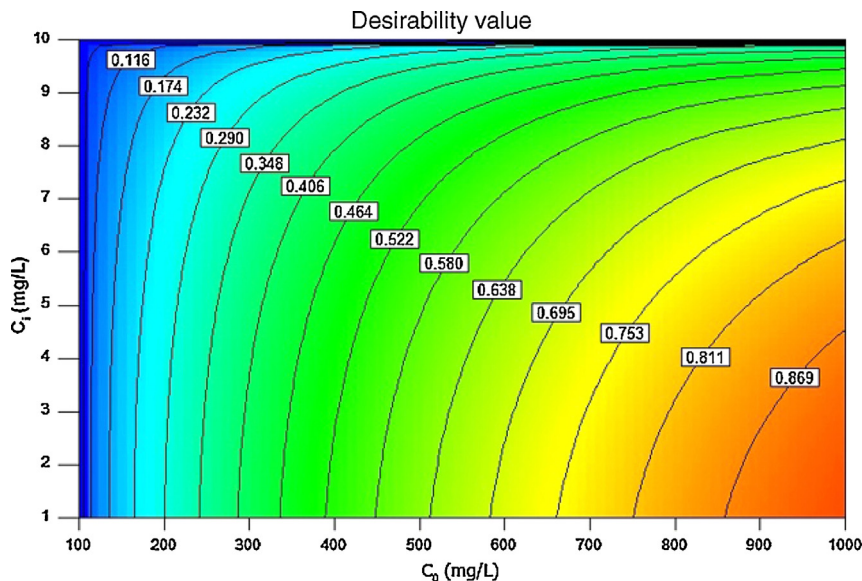


Fig. 14. Effect of BA and silicon dioxide nanoparticles initial concentration on combined desirability value.

curvature of contour lines denotes to what extent the independent variables have communal interactions. If the shape of the contour lines is more elliptical rather than circular, the interactive effect of independent variables on the predicted response is more significant. As it illustrated in Fig. 11B, the contour lines are of elliptical type in this study, which means that the interactions between C_0 and C_i are likely to be of great significance to the equilibrium adsorption capacity (Jadhav et al., 2012; Zhong & Wang, 2010).

3.5. Desirability function optimization

A maximum level of BA initial concentration, minimum level of silicon dioxide nanoparticles initial concentration and maximum level of CTAB-V adsorption capacity were set for the maximum desirability. By seeking from 30 points per optimizations that is the software default and considering the importance of each variable the highest value that can vary from 1 to 5, the best local conditions were found to be the initial concentration of BA at 969.2 mg/L, the initial concentration of silicon dioxide nanoparticles at 1 mg/L and the adsorbent capacity at about 36.83 mg/g. Fig. 12 shows the optimum values for all variables suggested by the software. The effect of BA and silicon dioxide nanoparticles initial concentration on the combined desirability values is shown in Fig. 13. Fig. 14 also shows the values of individual desirability values as well as the combined desirability value that was found to be 0.916, which indicates the reliability of the optimizations process.

4. Conclusions

To sum up, the batch adsorption of BA from aqueous solution onto CTAB-V at equilibrium was compared in the presence and absence of hydrophilic silicon dioxide nanoparticles in this research. An RSM-based model was also developed for investigating the equilibrium adsorption capacity by varying the initial concentrations of BA and nanoparticles. The main conclusions can be summarized as below:

- (I) A 5 and 7.5 mg/L initial concentration of nanoparticles optimizes the equilibrium efficiency of BA adsorption onto CTAB-V when the initial concentration of BA is 100–400 and 400–1000 mg/L, respectively.
- (II) The equilibrium removal efficiency of BA in the presence of 7.5 mg/L nanoparticles increases by between about 9.0% and 31.5% when the initial concentration of BA varies from 100 to 1000 mg/L.
- (III) The equilibrium adsorption isotherm changes from the Langmuir type (without strict plateau) in the absence of nanoparticles to the high affinity type in the presence of nanoparticles, which implies the more favorable adsorptive behavior of CTAB-V when the nanoparticles are present.
- (IV) In the presence of nanoparticles, the equilibrium data are better fitted to the Langmuir model compared to the Freundlich, Dubinin–Radushkevich and Temkin models. In the absence of nanoparticles, the equilibrium data are

better fitted to the Langmuir and Temkin models compared to the two other models.

- (V) The RSM-based model reliability was confirmed by the high coefficient of determination and Fisher value as well as the probability value of less than 0.05. The range of internally studentized residuals was acceptable (between -3 and $+3$) and they were normally distributed owing to their mean and variance values very close to zero and unity, respectively.
- (VI) The interactive effect of the initial concentrations of BA and nanoparticles on the equilibrium adsorption capacity was found to be noticeable resulting from the elliptical form of contours. Individually, it was observed that the initial concentration of BA has a larger effect on the response than the initial concentration of nanoparticles.
- (VII) The desirability function optimization approach showed that the best local conditions are to be the initial concentration of BA at 969.2 mg/L, the initial concentration of silicon dioxide nanoparticles at 1 mg/L and the adsorbent capacity at about 36.83 mg/g.

Conflict of interest

The authors have no conflicts of interest to declare.

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