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TITLE: BIOSYNTHESIS, CHARACTERISATION AND DETERMINATION OF ADSORBENT

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EXTRACT

AUTHORS: Fahriye TASKIRAN, Deniz UZUNOGLU, Ayla ÖZER

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BIOSYNTHESIS, CHARACTERISATION AND DETERMINATION OF ADSORBENT PROPERTIES OF SILVER NANOPARTICLES WITH CYPRUS ACACIA (Acacia cyanophylla) LEAF EXTRACT

Fahriye TAŞKIRAN¹, Deniz UZUNOĞLU^{1,*}, Ayla ÖZER¹

¹ Department of Chemical Engineering, Faculty of Engineering, Mersin University, Mersin, 33343, Turkey

ABSTRACT

In this work, silver nanoparticles (AgNPs) were biosynthesized using Cyprus acacia (*Acacia cyanophylla*) leaf extract as reductant agent; the biosynthesized AgNPs were characterized by DLS, FT-IR, SEM, EDX, and XRD. Then, the single and multistage batch adsorption of Basic Red 46 dye (BR46) onto AgNPs were studied and the optimum conditions were determined as follows: initial pH of 7.0, initial dye concentration of 200 mg/L, temperature of 35 °C, and adsorbent concentration of 1.0 g/L. For BR 46-AgNPs adsorption, the equilibrium was defined by Langmuir isotherm model and the adsorption kinetic data were best described by the pseudo second order kinetic model. From Weber-Morris model, it was concluded that only intraparticle mass transfer was effective on the adsorption of BR 46 by AgNPs. It was obtained from the thermodynamic studies that the studied adsorption system was exothermic, induced, and decreasing in randomness of adsorbed species.

Keywords: Adsorption, Biosynthesis, Cyprus acacia, Multi-stage adsorption, Silver nanoparticles

1. INTRODUCTION

Nanoparticles have some outstanding physicochemical properties like good electrical and thermal conductivity, high aspect ratio of ordered structure, light weight, metallic/semi-metallic feature, high mechanical endurance, large specific surface area [1]. Beginning by Faraday's study with colloidal gold in the middle of the 1800s, the unusual physicochemical properties of metallic nanoparticles have been subjects to ongoing investigation by physics and chemistry communities [2]. Metallic nanoparticles are conventionally synthesized by electro-chemical method, reverse micelles/microemulsion method, hydrothermal synthesis, sonochemical reduction, and chemical reduction [3]. Because of these methods' disadvantages such as use of hazardous chemicals and toxic materials, low conversion, difficult and costly purification requirements, high energy demand, biosynthesis methods using biological reductant agents like microorganisms, enzymes, plant extracts have attracted much attention in the recent years. The metallic nanoparticles are biosynthesized by reduction from the metal salt with the aid of the biological agents. Compared to other biological agents used in the biosynthesis method, using plant extract is more advantageous choice because plants are too available in the environment, the method does not require to sterilization, processing of plants is easier and safer. Given the diversity of plants in our country; plant leaves, unused ecofriendly resources, can be converted into economic value when they are evaluated in metallic nanoparticles biosynthesis. [4]. Recently, the metallic nanoparticles are commonly used as adsorbents in pollutant removal processes because they have remarkable advantages like only a bit of nanoparticle is sufficient for wastewater treatment, they have large specific surface area and a great number of unsaturated atoms on their surfaces resulting in comparatively high adsorption capacities [1]. In this work, the silver nanoparticles (AgNPs) were biosynthesized by using Cyprus acacia (Acacia cyanophylla) leaf extract and were characterized, and the adsorbent properties of AgNPs for BR46 dye were also determined.

^{*}Corresponding Author: <u>denizuzunoglu4@gmail.com</u>

2. MATERIALS AND METHODS

2.1. Biosynthesis of AgNPs

5.0 g of the purified and dried A. cyanophylla leaves were boiled in 100 mL of pure water at 100 °C for desired time and the leaf extract was obtained by this way. 5 mL of the leaf extract was added dropwise to 100 mL of 10⁻³ M AgNO₃, which was stirring magnetically, and then the mixture in the conical flask was stirred magnetically for 120 min. The resulting mixture left at room temperature for 1 day in an attempt to be accomplished the biosynthesis. By this way, +1 valent silver ion in AgNO₃ structure was reduced to zero valent silver, silver nanoparticles, with the aid of the leaf extract. biological reductant agent. AgNPs precipitated at the bottom of the conical flask; and then the precipitate was separated manually from the AgNPs-leaf extract mixture. The residual AgNPs-leaf extract suspension was added to petri plates and the aqueous leaf extract was evaporated at 110 °C in the stove for 6 h. The obtained AgNPs were washed with pure water; they were dried at 180 °C in the stove for 24 h, and were stored at +4 °C in the refrigerator [5]. The characterization studies of the biosynthesized AgNPs were carried out by Fourier Transform Infrared Spectrometer (FTIR- Perkin Elmer, Shelton), Scanning Electron Microscope (SEM- Zeiss/Supra 55, Germany), Energy Dispersive X-ray Spectroscopy (EDX- Zeiss/Supra 55, Germany), X-ray Diffractometer (XRD- Philips XPert, Netherlands), zetasizer (Malvern, UK) using Dynamic Light Scattering (DLS) technique, and zetapotential analyzer (Malvern, UK).

2.2. Adsorption Studies

Adsorption studies were done with artificial wastewater containing BR46 (molecular formula: C18H21BrN6, CAS Registry Number:12221-69-1, molar mass: 401.3 g/g-mole). In this work, the effects of experimental conditions such as initial pH (7.0, 8.0, 9.0, and 10), initial dye concentration (25, 50, 75, 100, 150, 200, 300, and 500 mg/L), temperature (25, 35, 45, and 55 °C), and adsorbent concentration (0.5, 1.0, 2.0, and 3.0 g/L) were investigated in a batch system. of 0.1 g the synthesized AgNPs (not suspension of AgNPs) were added to 100 mL (0.1 L) of dye solution, equal to (0.1 g adsorbent)/(0.1 L dye solution) = 1.0 g/L of adsorbent concentration, for the all adsorptionstudies except for the adsorbent concentration experiments. The other adsorbent concentrations can be calculated in the same way. In this respect, the required amount of AgNPs was blended with 100 mL of the dye solution in 250-mL conical flasks. In order to achieve the adsorption equilibrium, the flasks were shaken in a thermostated water bath at constant temperature for 180 min. At predetermined time intervals (0.0, 0.5, 2.0, 5.0, 10, 20, 30, 60, 120, and 180 min), the sample of dye solution was taken and separated from the adsorbent with centrifuge at 3000 rev/min for 5.0 min. The residual dye concentration in the supernatant was analyzed at 530 nm by UV-vis spectrophotometer (Chebios Optimum-One, Italy).

The adsorbed amount at equilibrium, q_{eq} (mg/g), and the percentage of adsorption (%) were computed as follows:

$$q_{eq} = \frac{C_{init} - C_{eq}}{X_{init}} \tag{1}$$

Percent adsorption (%) =
$$\frac{C_{init} - C_{eq}}{C_{init}} \times 100$$
 (2)

3. RESULTS AND DISCUSSION

3.1. Adsorbent Characterization

The distribution of effective hydrodynamic diameters of biosynthesized AgNPs was measured by DLS technique and the results were presented in Figure 1. As can be seen from Figure 1, single AgNPs with

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the effective hydrodynamic diameter of 32.67 nm were the most frequent ones (14.20 %) while the smallest AgNPs (15.69 nm) had the distribution of 3.50 % and the average particle size of the biosynthesized AgNPs was determined as 38.55 nm with low polydispersity index (PDI) of 0.246 by DLS technique.



Figure 1. The distribution of effective hydrodynamic diameters of AgNPs

FT-IR spectra of AgNPs before and after the adsorption were given in Figure 2. Accordingly; =C–H alkenes (2980 cm⁻¹), –C–H alkanes (2887 cm⁻¹), -COOH carbonyl group (1630 cm⁻¹), C=C aromatic group (1410 cm⁻¹), –C–H alkane (1387 cm⁻¹), C–N stretch aliphatic amines (1091 cm⁻¹), C–N amines (1005 cm⁻¹), C–H oop aromatics (818 cm⁻¹ to 660 cm⁻¹) peaks were observed from FT-IR spectra of AgNPs [6].



Figure 2. (a) FT-IR spectrum of AgNPs before adsorption; (b) FT-IR spectrum of AgNPs after adsorption

According to Figure 2. (b); the peaks of 1091 and 1005 cm⁻¹ were disappeared post-adsorption, which indicating that amines were responsible groups in the adsorption.

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XRD patterns of AgNPs before and after the adsorption were given in Figure 3. XRD pattern of AgNPs exhibited peaks at 20 angles of 38.10° , 44.28° , 64.41° , and 77.36° which corresponded to the [1 1 1], [2 0 0], [2 2 0], [3 1 1], and [2 2 2] crystal planes of a cubic lattice structure of AgNPs [7]. The unassigned peaks were also observed indicating the crystallization of bio-organic phase on the surface of the silver nanoparticles due to the leaf extract [8]. The obtained XRD patterns were consistent with the various similar studies in the literature [5, 9-12].



Figure 3. XRD patterns of AgNPs

Moreover, there was not any difference between before and after adsorption for XRD patterns of AgNPs, which was indicating the phenomena of physical adsorption.

EDX analysis results were given in Table 1 and it showed that AgNPs involved 90.43 % Ag, 3.99 % Cl, 3.18 % C, and 2.40 % O before adsorption. Accordingly, the strong signal in the silver region affirmed the formation of AgNPs, the obtained of carbon, oxygen, and nitrogen elements stemmed from the extracellular organic moieties of *A. cyanophylla* extract on the adsorbent surface. The observed 0.76% N and the increase in weight percentages of C and O post-adsorption resulted from the adsorbed BR46 ($C_{18}H_{21}BrN_6$) molecules onto AgNPs.

Table 1. EDX	analysis	results
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Elomon4	Weig	ht %
Element	Pre-adsorption	Post-adsorption
Ag	90.43	90.35
C	3.18	3.23
Ο	2.40	2.42
Cl	3.99	3.24
Ν	-	0.76

From SEM images (Figure 4), the porous structures were observed pre-adsorption (a) and these pores were covered by BR46 molecules post-adsorption (b).

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Figure 4. (a) SEM images of AgNPs before adsorption; (b) SEM images of AgNPs after adsorption

3.2. The Single and Multi-Stage Adsorption of Basic Red 46 onto AgNPs

The single stage adsorption of BR46 onto AgNPs was investigated in a batch process for various experimental conditions such as initial pH, initial dye concentration, temperature, and adsorbent concentration.

3.2.1. Effects of initial pH

The effect of initial pH on BR 46 adsorption was studied in the range of 7.0–10 at 25°C temperature, 1.0 g/L AgNPs concentration, and 100 mg/L initial BR46 concentration for 180 min contact time and the equilibrium uptake amounts at different initial pH values were presented in Figure 5.a. From 5.a, it was observed that the maximum equilibrium BR 46 uptake amount (q_{eq}) was observed at initial pH 7.0 and q_{eq} values decreased with increasing initial pH thereafter. This state can be explained by the zeta potential analysis for AgNPs (Figure 5.b). It is well-known that a suspension with zeta potential above ±30 mV is stable and above ±60 mV shows excellent stability while a suspension with zeta potential below ±20 mV has limited stability and below ±5 mV aggregates significantly [13]. Accordingly, the surface of AgNPs had the maximum negative charge (-24.17 mV) at pH 7.0; therefore, in consequence of the strong electrostatic attraction between the anionic adsorbent surface and BR46 cations, the maximum adsorbed BR46 amount was observed at this pH value. The negative charge of AgNPs surface decreased at higher pH values than 7.0 and so the adsorbed amounts also decreased thereafter as a result of the decreasing in the mentioned electrostatic attraction. For these reasons, the optimum initial pH was determined as 7.0 for BR 46 adsorption onto AgNPs.





Figure 5. (a) Initial pH effect; (b) zeta potential values of AgNPs for different pHs

3.2.2. Effects of temperature

The studies of temperature effects on BR46 adsorption onto AgNPs was done in the range of 25-55 °C at 100 mg/L initial BR46 concentration, 7.0 initial pH, and 1.0 g/L AgNPs concentration for 180 min contact time and the equilibrium uptake amounts at different temperatures were presented in Figure 6.



Figure 6. Temperature effect

As can be seen from Figure 6, q_{eq} values increased with increase in temperature down to 35 °C and then they decreased hereafter. The increasing of q_{eq} values with temperature can be explained the opening of the pores in the adsorbent surface in the range of 25-35 °C. The decrease of q_{eq} values with further increase in temperature arose from the physical and exothermic phenomena of BR46 adsorption on AgNPs. Therefore, the optimum temperature value was found as 35 °C for BR46 adsorption onto AgNPs.

To evaluate the effect of temperature on BR46 adsorption onto AgNPs in details, the thermodynamic parameters of change of Gibb's energy (ΔG), change of enthalpy (ΔH), and change of entropy (ΔS) were determined with Van't Hoff equation and the calculated parameters were given in Table 2. Table 2 showed that BR46 adsorption onto AgNPs was exothermic (ΔH <0), induced (ΔG >0), and decreasing in randomness of adsorbed species (ΔS <0).

T (°C)	ΔG (J/mole)	ΔH (J/mole)	ΔS (J/mole.K)
35	2829		
45	3477	-19518	-72.47
55	4281		

 Table 2. Thermodynamic parameters

In order to discuss the effect of the temperature on the adsorption equilibrium in details, the linear forms of Langmuir ($\frac{1}{q_{eq}} = \frac{1}{q_{m.}b_{L.}C_{eq}} + \frac{1}{q_{m}}$) and Freundlich ($\ln(q_{eq}) = \ln(K_{fr}) + \frac{1}{n}\frac{1}{C_{eq}}$) adsorption isotherm models were evaluated for different temperature values and the obtained isotherm constants along with the regression coefficients (R²) and error values ($\operatorname{Err} = \left(\frac{100}{n}\right) \sum_{1}^{n} \left|\frac{(q_{eq,cal}-q_{eq,ex})}{q_{eq,ex}}\right|$) were presented in Table 3. Accordingly, higher regression coefficients (R²>0.99) and lower Err values of Langmuir isotherm model showed that BR46 adsorption equilibrium data best described by Langmuir isotherm model indicating that BR46 adsorption onto AgNPs was monolayer in nature. Besides, it was noted that the maximum monolayer adsorption capacity was determined at 35 °C, which was the optimum temperature value for BR46 adsorption onto AgNPs.

Table 3. Isotherm constants and the related regression coefficients (R²) for the adsorption of BR46 onto AgNPs at various temperatures

Langmuir isotherm model				Freundlich isotherm model				
1 (°C)	qm (mg/g)	b _L (L/mg)	Err	R ²	${ m K_{fr}} \ (mg/g)(L/mg)^{1/n})$	1/n	Err	R ²
25	98.97	0.0124	7.12	0.995	4.27	0.526	10.41	0.956
35	126.48	0.1473	3.74	0.994	31.85	0.266	19.24	0.834
45	115.27	0.0990	5.18	0.994	26.63	0.277	19.81	0.796
55	89.81	0.1800	4.86	0.993	29.22	0.214	17.94	0.761

The adsorption mechanism of BR 46 on AgNPs progresses in accordance with the assumptions of the Langmuir isotherm model. The prevenient mechanism of BR46 adsorption onto AgNPs as follows;

$$COO^{-} + [C_{18}H_{21}N_6]^+ \rightarrow [(COO^{-})(C_{18}H_{21}N_6)^+]$$
(3)

According to Eq. 3, one cationic BR46 molecule ($[C_{18}H_{21}N_6]^+$) could adsorb onto carboxyl groups (COO⁻) of AgNPs surface (indicated by FT-IR) and further adsorption cannot occur at that site on the adsorbent surface.

Also, Table 4 represented the maximum adsorption capacities of AgNPs for different dyes in the literature. It was observed that AgNPs used in this work had higher adsorption capacity as against the other AgNPs used as adsorbents for different dyes in the literature.

Adsorbent	Synthesis Method	Particle size (nm)	Dye	qm (mg/g)	Reference
AgNPs loaded activated carbon	Reduction by NaOH	20-90	Direct yellow 12	454.54	[1]
AgNPs- colemanite ore	Thermal reduction without reductant	5-30	Reactive Yellow 86 Reactive Red 2	68.71 59.13	[14]
Polyaniline/Ag nanocomposite	Reduction by NaOH	4.97	Brilliant green	23.66	[15]
AgNPs	Biosynthesis by <i>A</i> . <i>cyanophylla</i> extract	103	Basic Red 46	126.48	This study

Table 4. The adsorption capacities of AgNPs synthesized by different methods in the literature

3.2.3. Effects of initial dye concentration

In order to evaluate the effect of initial dye concentration, the adsorption experiments were carried out in the range of 25-500 mg/L at 35°C temperature, 7.0 initial pH, and 1.0 g/L AgNPs concentration for 180 min contact time and the equilibrium uptake amounts along with the adsorption percentages for various initial concentrations of BR46 were presented in Figure 7. According to Figure 7, q_{eq} values increased with increase in the initial BR46 concentration down to 200 mg/L in consequence of the increasing driving force (Δ C) and then remained constant thereafter because at high concentrations, the available binding sites on the adsorbent become less forming the limiting factor for the adsorption. Therefore, in BR 46 adsorption onto AgNPs, the optimum initial concentration of BR46 was observed as 200 mg/L.



Figure 7. Initial dye concentration effect

The most important parameter affecting adsorption kinetics was the initial solute concentration. In this study, the experimental adsorption data determined at different initial BR46 concentrations were used to evaluate the kinetics of BR46 adsorption on AgNPs. The pseudo first order $(\log(q_{eq}-q_t)=\log(q_{eq})-\frac{k_{f}t}{2.303})$ and the pseudo second order $(\frac{t}{q_t} = \frac{1}{q_{eq}^2 \cdot k_s} + \frac{t}{q_{eq}})$ kinetic models were applied to the experimental kinetic data for various initial BR46 concentrations, and the results were shown in Table 5. Higher values of regression coefficients and the consistency between experimental $(q_{eq,ex})$ and calculated $(q_{eq,cal2})$ uptake values from kinetic models showed excellent fit of the pseudo second order kinetic model as seen from Table 5.

Cinit	q _{eq,ex}	Pseudo first order kinetic model		Pseudo first order Pseudo second order kinetic model kinetic model			Weber-Morris model		
(mg/L)	(mg/g)	q _{eq,cal1} (mg/g)	k _f (min ⁻¹)	R ²	q _{eq,cal2} (mg/g)	ks (g/mg.min)	R ²	K _{in} (mg/g.min ^{0.5})	R ²
25	24.35	17.88	0.0324	0.983	25.11	0.0061	0.999	3.766	0.990
50	49.06	41.57	0.0253	0.988	51.69	0.0017	0.991	5.949	0.980
75	73.07	65.87	0.0179	0.985	76.87	0.0008	0.995	6.777	0.990
100	92.42	71.79	0.0244	0.965	96.03	0.0010	0.990	10.98	0.986
150	107.22	88.14	0.0246	0.937	110.53	0.0008	0.987	9.295	0.971
200	116.55	98.58	0.0133	0.965	117.42	0.0005	0.976	9.431	0.983
300	117.71	114.26	0.0214	0.909	122.83	0.0011	0.997	14.120	0.980

Table 5. Kinetic parameters and the related regression coefficients (R²) for the adsorption of BR46 onto AgNPs at various initial concentrations

To interpret the effects of the external diffusion and intraparticle diffusion on BR46 adsorption by AgNPs, Weber-Morris model ($q_t=K_{in}$. \sqrt{t} + I) was applied to the experimental data at different initial BR 46 concentrations and the calculated parameters were presented in Table 5. According to Weber-Morris model, if the plot of q_t versus \sqrt{t} is linear and pass through the origin, the adsorption process follows only intraparticle diffusion. On the other side, the Weber-Morris plot can be linear and also has intercept value if both intraparticle and film diffusion are effective on the adsorption process. As a result, only intraparticle diffusion was effective on BR46 adsorption onto AgNPs due to the lack of intercept values of Weber-Morris plots as seen in Table 5.

3.2.4. Effects of adsorbent concentration: modelling of multistage adsorption

The effect of adsorbent concentration on BR 46 adsorption onto AgNPs was studied for 0.5, 1.0, 2.0, and 3.0 g/L at 7.0 initial pH, 100 mg/L initial BR46 concentration, and 35 °C temperature for 180 min contact time. The variation of the equilibrium uptake amounts and percentages of adsorption with adsorbent concentration was given in Figure 8. From Figure 8, q_{eq} values decreased while the adsorption percentages increased with increasing the adsorbent concentration. This can be explained with the availability of more number of binding sites on the adsorbent surface for lower adsorbent concentrations. As a result, an adsorption system containing low adsorbent concentration may be offered for the desired purification because higher q_{eq} values were attained at low adsorbent concentrations. For these reasons, the optimum AgNPs concentration was observed as 1.0 g/L for BR46 adsorption onto AgNPs.



Figure 8. Adsorbent concentration effect

If the adsorption process is separated in multistage small batches of adsorbent instead of in a single batch, with filtration between each stage, the higher removal capacity can be achieved with higher economy of adsorbent. n-staged removal process which involved V_{init} volume of BR46 solution was given in Figure 9, and the mass balance of dye for 1st stage represented in Eq. 4.



Figure 9. Multistage removal process

$$V_{\text{init}}.C_{\text{init}} + X_{\text{init}}.q_{\text{init}} = X_{\text{init}}.q_{\text{eq1}} + C_{\text{eq1}}.V_{\text{init}}$$
(4)

Eq. 4 can be rearranged as follows,

$$\left(\frac{V_{\text{init}}}{X_{\text{init}}}\right) = \frac{(q_{\text{eq1}} - q_{\text{init}})}{(C_{\text{eq1}} - C_{\text{init}})}$$
(5)

In the same way, Eq. 5 can be rearranged for nth stage as follows,

$$-\left(\frac{V_{\text{init}}}{X_{\text{init}}}\right) = \frac{(q_{\text{eqn}} - q_{\text{init}})}{(C_{\text{eqn}} - C_{\text{eq(n-1)}})}$$
(6)

The adsorbed BR46 mass per gram of AgNPs at the entrance of each reactor (q_{init}) is equal to zero. The operating lines, provided from Eq. (5) and Eq. (6), passing through (q_{init}, C_{init}) and (q_{eq1}, C_{eq1}) for the 1st stage and $(q_{init}, C_{eq(n-1)})$ and (q_{eqn}, C_{eqn}) for the nth stage, respectively. If the amount of AgNPs used in each stage is equal, the operating lines, each slope is $-V_{init}/X_{init}$, are parallel [16].

For BR46 adsorption onto AgNPs, the equilibrium curve plotted by the experimental equilibrium data (C_{eq}, q_{eq}) along with the operating lines whose slopes were 1.0 (- (V_{init}/X_{init}) =-(1L/1g)=-1) for four stages were given in Figure 10. From Figure 10, the unadsorbed BR46 concentrations at the 1th, 2th, 3th, and 4th stages were determined as 376, 256, 138, and 38 mg/L for initial BR46 concentration of 501.17 mg/L, respectively.



Figure 10. The equilibrium curve with the operating lines (each slope: $-V_{init}/X_{init}$ =-1) for four staged batch stirred reactor in series

The multistage adsorption system can be also described mathematically by use of Langmuir isotherm model to forecast the unadsorbed dye concentration for each stage. Substituting the Langmuir equation for $q_{eq1} \left[= \frac{q_{m} \cdot b_L \cdot C_{eq1}}{1 + b_L \cdot C_{eq1}} \right]$ and q_{eqn} , in Eqs. (7) and (8) gives:

$$y = -\left(\frac{V_{\text{init}}}{X_{\text{init}}}\right) = \frac{(q_{\text{m.b}\text{L.Ceq1}})}{[(1+b_{\text{L.Ceq1}}).(C_{\text{eq1}}-C_{\text{init}})]}$$
(7)

$$y = -\left(\frac{V_{init}}{X_{init}}\right) = \frac{(q_m.b_L.C_{eqn})}{[(1+b_L.C_{eqn}).(C_{eq(n-1)}-C_{eqn})]}$$
(8)

Rearranging Eqs. 7 and 8 give for stages 1th and nth:

$$b_{L}.C_{eq1}^{2} + (1-b_{L}.C_{init} + q_{m}.b_{L}.y)C_{eq1} - C_{init} = 0$$
 (9)

$$b_{L}.C_{eqn}^{2} + (1-b_{L}.C_{eq(n-1)} + q_{m}.b_{L}.y)C_{eqn} - C_{eq(n-1)} = 0$$
(10)

The positive root calculated from analytical solution of Eqs. 9 and 10 gives the equilibrium dye concentration for 1^{th} and $n^{th}(C_{eq1}, C_{eqn})$ stages, respectively.

At the optimum adsorption conditions, the experimental q_m and b_L values were used for obtaining C_{eq} values in the output of each stage from Eqs. 9 and 10. The unadsorbed BR 46 concentrations at the end of the 1th, 2th, 3th, and 4th stages were determined as 376.92, 253.74, 133.39 and 30.15 mg/L, respectively. As a conclusion, there is good consistency between the equilibrium BR 46 concentrations determined by the experimental equilibrium data-operating line method and calculated with the aid of Langmuir isotherm model constants.

It was clearly understood from the unadsorbed BR 46 concentrations that the required purification could not be provided by a single stage according to the discharge standards given by Water Pollution Control Regulation of T.R. Environment and Urbanization Ministry (Table 6); but it was observed that the unadsorbed BR 46 concentration could be reduced below the discharge limit values in four-staged adsorption system.

Parameter	Unit	Sample for 2 hours	Sample for 24 hours
Chemical oxygen demand (COD)	(mg/L)	350	240
Nitrogen of Ammonium (NH ₄ -N)	(mg/L)	5	-
Free chlorine	(mg/L)	0.3	-
Total chrome	(mg/L)	2	1
Sulfur (S ⁻²)	(mg/L)	0.1	-
Sulphite	(mg/L)	1	-
Oil and grease	(mg/L)	10	-
pH	-	6-9	6-9
(Additional line:RG-24/4/2011-27914)	(Pt-Co)	280	260
Color	(mg/L)	52	48

Table 6. Discharge limit values according to Water Pollution Control Regulation of T.R. Environment and Urbanization Ministry

4. CONCLUSION

AgNPs were biosynthesized using Cyprus acacia (*Acacia cyanophylla*) leaf extract and they were characterized by DLS, FT-IR, SEM, EDX, XRD. The biosynthesized AgNPs were used as adsorbents in the single stage adsorption of BR46 onto AgNPs in the batch system and the optimum conditions were determined as follows: initial pH of 7.0, initial dye concentration of 200 mg/L, temperature of

35 °C, and adsorbent concentration of 1.0 g/L. Weber-Morris model results showed that only intraparticle diffusion was effective in the removal. The BR46 adsorption kinetics was in the best agreement with the pseudo second order model. Thermodynamic studies stated that the studied adsorption process was exothermic, induced, and decreasing in randomness of adsorbed species. The BR46 adsorption equilibrium was defined by Langmuir isotherm model. The maximum monolayer coverage capacity of AgNPs was determined as 126.48 mg/g (R²=0.994, Err=3.74) at 35 °C, which was the optimum temperature value and the calculated maximum monolayer coverage capacity of AgNPs confirmed that the biosynthesized AgNPs could be used effectively for BR46 adsorption in comparison to the other nanoparticles used for different dyes. In addition, the multistage adsorption of BR46 onto AgNPs was studied; it was observed that the desired purification could be achieved by four staged adsorption system.

NOMENCLATURE

1/n	a constant of Freundlich isotherm model related to adsorption intensity
b_L	a constant of Langmuir isotherm model (L/mg)
C _{eq}	concentration of residual BR46 at equilibrium (mg/L)
Ceqn	concentration of residual BR46 at equilibrium for n th stage (mg/L)
Cinit	initial concentration of BR46 (mg/L)
Err	error value
\mathbf{k}_{f}	a rate constant of pseudo first order kinetic model (1/min)
\mathbf{K}_{fr}	a constant of Freundlich isotherm model (mg/g)/(L/mg)1/n)
Kin	Weber-Morris model constant related to intraparticle diffusion (mg/g.min ^{0.5})
ks	a rate constant of pseudo second order kinetic model (g /(mg min))
oop	out of plane
q_{eq}	adsorbed mass of BR46 per gram of AgNPs at equilibrium (mg/g)
q _{eq,cal1}	adsorbed mass of BR46 per gram of AgNPs calculated from pseudo first order kinetic
	model (mg/g)
q _{eq,cal2}	adsorbed mass of BR46 per gram of AgNPs calculated from pseudo second order kinetic
	model (mg/g)
$q_{eq,ex}$	experimental adsorbed mass of BR46 per gram of AgNPs at equilibrium (mg/g)
q_{eqn}	adsorbed mass of BR46 per gram of AgNPs at equilibrium for n th stage (mg/g)
q_m	maximum monolayer coverage capacity of AgNPs obtained from Langmuir isotherm
	model (mg/g)
q_{o}	adsorbed mass of BR46 per gram of AgNPs at the beginning in each reactor (mg/g)
Т	temperature (°C, K)
t	time (min)
V _{init}	adsorption solution volume (L)
X_{init}	AgNPs concentration (g/L)
ΔG	change of Gibb's free energy (J/mole)
ΔH	change of enthalpy (J/mole)
ΔS	change of entropy (J/mole.K)

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