Adsorptive/desorptive potential of cationic basic yellow 28 (BY28) dyes onto natural untreated clay from aqueous phase

Abstract

The adsorption of basic dye “C.I. basic yellow 28” (BY28) onto Tunisian smectite-rich clay in aqueous solution was studied in a batch system. Tunisian raw clay sampled in Jebel Romana (meridional Atlas of Tunisia) composed mainly of smectite (52%) has shown efficiency for removal of BY28 from aqueous solutions. The results showed that basic pH is favorable for the adsorption of dye and adsorption. Adsorption isotherm was described by both Freundlich and Langmuir models. The maximum monolayer adsorption capacity was found to be 76.92 mg/g at 25°C (at room temperature). Adsorption kinetics was studied and modeled. The maximum BY28 desorbed percentage was 17% and it was obtained for strong acidic conditions (pH 2). Smectitic clay showed to be an advantageous adsorbent in terms of price and versatility, being effective under a wide pH range, for cationic dye. Its use as adsorbent does not require further purification or chemical modifications, which is beneficial for the environment and for the economy of the treatment.

Keywords: adsorption, smectitic clay, textile dye, equilibrium, kinetics, desorption

Abbreviations: FTIR, fourier-transform infrared; X-RD, X-ray diffraction; BY28, dye-basic yellow 28; MB, methylene blue; CEC, cation exchange capacity

Introduction

Textile industries are one of the largest consumers of water and hence producers of liquid effluents. The dyeing and finishing processes generate volumes of wastewaters in the range 45-450 m³ per ton of product, containing salts, acids, bases, additives and unfixed dyes. A suitable treatment of these complex wastewaters should lead to a strong decline on the organic load, and desirable to a complete elimination of the residual color. There are several types of commercial dyes, and some of these dyes posses a high resistance to biodegradation and persist a long time in the environment. These compounds and their degradation by products are generally believed to be carcinogenic or toxic and become harmful to humans or to the environment. Adsorption processes have been proposed as a fast alternative to remove contaminants from water. Several natural adsorbents have been tested to remove synthetic dyes. In particular, smectite clays have been broadly studied as adsorbents of cationic dyes. This type of clay has a layered structure where each layer is composed by tetrahedral silicate units fused into an edge-shared octahedral plane of aluminum. The isomorphic substitution of Al³⁺ for Fe²⁺ or Mg²⁺ and Si⁴⁺ for Al³⁺ generates charge deficiencies on the surface and interlayer space.

The intrinsic negative charge is compensated with inorganic captions in the interlayer that can be exchanged by other inorganic or organic captions, being suitable for adsorption of cationic synthetic dyes. The aim of the present work was to investigate the efficiency of AR towards the adsorption of Basic Yellow 28 from aqueous solution. This work describes the feasibility of removal of a cationic dye-Basic Yellow 28 (BY28) from aqueous solutions by using raw clay as a new low cost adsorbent. The clay was characterized using methylene blue adsorption for determination of total specific surface area, cation exchange capacity, point of zero charge, Fourier-transform infrared (FTIR) and X-ray diffraction (XRD). Adsorption experiments were achieved out as a function of pH, contact time, initial dye concentration, adsorbent dose and temperature. Kinetics, thermodynamic, isotherms and mass transfer mechanisms have also been discussed for the adsorption of BY28 from aqueous solutions.

Experimental

Adsorbent

The BY28 was obtained from textile industry (Tunisia) with 99.99% purity. The dye solutions used in the experiments (1000 mg/L) were prepared by dissolving the required amount of dye in distilled water. The characteristics and chemical structure of BY28 are (Table 1).

| CI name | C.I basic yellow 28 |
|---------|---------------------|
| Commercial name | Maxilon Golden Yellow GL 200% |
| Chemical formula | C21H27N3O5S |
| Class | Azo |
| Molecular weight | 433,52 |
| Max. wave length (λ max) | 440 nm |
| pH stability | 3–10 |
| Molecular Structure | |

Table 1 Basic properties of the investigated dye
Adsorbent and its characterization

In this study, natural untreated clay (AR) was collected from the meridional Atlas of Tunisia (Figure 1). It was sampled in Jebel Romana in El Hamma area. Then, it was crushed, ground, and sieved through a 63 μm sieve and samples were collected from under the sieve and dried at 70 °C in an oven for 2h prior to use. It was used as the adsorbent for the removal of BY28 from aqueous solutions in a batch treatment process. Clay was used without any treatment. Fourier-transform infrared (FTIR) spectra were measured using Perkin-Elmer Spectrum One model FTIR spectrometer. The methylene blue (MB) adsorption method was used in this work to assess the surface area of AR. Cation exchange capacity (CEC) was estimated using the copper ethylenediamine complex method. The X-ray diffraction (XRD) patterns were obtained using phillips X'pert diffractometer The zero point charge of the raw AR (pH$_{zpc}$) defined as being the pH for which there is absence of positive or negative charge on the surface of the AR was determined by the method describe by.

Figure 1 Localization of the studied clay deposit.

Experimental adsorption isotherms

Adsorption experiments were evaluated in batch equilibrium mode. Sample clay of the volume 100 mL were taken after mixing the adsorbent and BY28 solution bearing solution and at predetermined time intervals (1-120 min) for the determination of residual BY28 concentration. pH was adjusted by the addition of dilute aqueous solutions of HCl or NaOH (0.01 M). The initial and equilibrium BY28 concentrations of dye was estimated using Shimadzu-1650UV-Visible spectrophotometer at the wavelength of 440 nm. The kinetics of adsorption was determined by analyzing adsorptive uptake of the dye from the aqueous solution at different time intervals. For adsorption isotherms, dye solutions of different concentrations were agitated with the known amount of adsorbent until the equilibrium was achieved. Various experimental parameters such as pH (2-11), temperature (25-45 °C), adsorbent mass (0.1-2 g/L), initial BY28 concentration (25-300 mg/L) and contact time were investigated. The amount of dye removed and the amount of dye adsorbed on per unit of adsorbent ($q_e$) at the sampling time were calculated using the following equations:

$$\text{Dye removal} \% = \frac{(C_0 - C_e)}{C_0} \times 100 \rightarrow (1)$$

$$q_e = \frac{V (C_0 - C_e)}{m} \rightarrow (2)$$

Where $C_0$ (mg/L) is the initial concentration of dye, $C_e$ (mg/L) is the concentration of dye at equilibrium, $V$ (L) is the volume of dye solution, and $m$ (g) is the mass of adsorbent.

Desorption studies

In the desorption studies, the raw adsorbent that was used for

the adsorption of dye solution was separated from solution by centrifugation and then dried. The dried clay loaded with BY28 dye, was shaken (1.0 g/L) with solutions at different pH (range 2-11) using NaOH and HCl solutions as eluents. After 5h of contact time, the dye concentration in the liquid phase was analyzed and the percentage of the dye desorbed from the AR was calculated according to:

$$\text{Desorption} \% = \frac{\text{Mass desorbed}}{\text{Mass adsorbed}} \times 100 \rightarrow (3)$$

Results

Characterization of AR clay

The FTIR spectrum of AR sample Figure 2 shows that absorption bands at 3627 and 917 cm$^{-1}$ are attributable to Al-Al-OH stretching and bending modes of vibrations. The band appeared at 1439 cm$^{-1}$ corresponds to that of carbonate [calcite (CaCO$_3$) or dolomite (Ca, Mg (CO$_3$)$_2$)]. The bands appeared at 537, 425 and 474 cm$^{-1}$ correspond to Si-O-Al, Si-O-Si and Si-O-Mg, respectively. The specific surface area and CEC of AR were about 397 m$^2$/g and 49.5 meq/100g, respectively. The X-Ray diffraction analysis indicated that the raw AR clay is mainly composed of smectite (52%) associated to kaolinite (6%), illite (11%), quartz (10%), calcite (19%), and feldspars (2%) (Table 2). Results show that pH$_{zpc}$ of AR clay was 8.9 (Figure 3).

Table 2 Mineralogical analysis of AR

| Sample | Clay minerals | Non clay minerals |
|--------|--------------|------------------|
| AR     | Smectite | Illite | Kaolinite | Quartz | Calcite | Feldspars |
|        | (S)    | (I)   | (K)     | (Q)    | (C)     | (F)      |
|        | 52     | 11    | 6       | 10     | 19      | 2        |

Citation: Islem C, Afef C, Jamoussi F et al. Adsorptive/desorptive potential of cationic basic yellow 28 (by28) dyes onto natural untreated clay from aqueous phase. Open Access J Sci. 2018;2(1):74–79. DOI: 10.15406/oajs.2018.02.00048
Effect of solution pH on adsorbate adsorption

The effect of pH on adsorption of BY28 onto AR was studied in the range of 2-11 which is presented in Figure 4. It was found that the removal efficiency of dye increased from 73.23% to 99.5% due to change in pH from 2 to 11 for a fixed initial dye concentration of 50 mg L\(^{-1}\) at equilibrium. However, pH 11 appears an optimal level due to high adsorption efficiency (99.5%). Clay minerals comprises of various functional groups such as aluminol and silanol groups which are affected by the pH of the solution. The point of zero charge, pH\(_{pzc}\) of clay particles in aqueous solution is 8.9. The predominant charges on the AR at acidic pH are positive and because of cationic dye species, it seems that the dominant mechanism of the adsorption is electrostatic repulsion. With the increasing pH values, the adsorption of BY28 on clay tends to increase, which can be explained by the increasing electrostatic attraction between the cationic dye adsorbate species and negatively charged adsorbent surfaces. Also lower adsorption at acidic pH due to the presence of excess H\(^+\) ions destabilizing cationic dye and competing with the dye captions for the adsorption sites.

Figure 4 Effect of initial solution pH on the adsorption of BY28 onto AR.

Effect of contact time

From Figure 5, it can be seen that adsorption of BY28 on the clay appears to be rapid during the initial phase of the contact time due to the fact that a large number of surfaces are available for adsorption. It then gradually slows down until it reaches equilibrium at 120 min with an adsorption rate of 92%. During this phase, the surface of the solid becomes saturated.

Figure 5 Effect of contact time on the adsorption of BY28 onto AR.

Effect of adsorbent dose

Initial adsorbent dose requires proper selection because it controls adsorption via available surface area and binding sites.\(^{10}\) The results revealed that adsorption efficiency decreased from 96 to 76% (Figure not shown here). This behavior is ascribed to a decrease in adsorption surface area and the number of active sites for adsorption.

Effect of initial dye concentration

To evaluate the effect of dye initial concentration adsorption, 20-150 mg/L concentration range was shown in Figure 6. By increasing initial dye concentration, the percentage removal increased from 20 to 80mg/L then decreases. At lower initial dye concentration, the adsorption sites were available for dye adsorption. At higher concentration, however the available sites might occupied and subsequently, the dye removal depends on the initial concentration.

Figure 6 Effect of initial dye concentration on the adsorption of BY28 onto AR.

Equilibrium modelling: In our present study two different adsorption isotherms, Langmuir and Freundlich have been used to fit the experimental adsorption data.

Langmuir isotherm: The Langmuir equation is probably the best known and most widely applied adsorption isotherm and is based on the assumption that adsorption takes place on homogeneous surface. It is represented as follows:\(^{11}\)

\[
q_e = \frac{q_m b C_e}{1 + b C_e} \quad (4)
\]

A linear form of this expression is

\[
\frac{C_e}{q_e} = \frac{1}{q_m b} + \frac{C_e}{q_m} \quad (5)
\]

Where \(C_e\) is the equilibrium concentration of BY28 in solution (mg/L), \(q_e\) is the adsorption capacity of the adsorbent at equilibrium (mg/g), \(q_m\) is the maximum adsorption capacity of the adsorbent (mg/g), \(b\) is the Langmuir equilibrium constant related to the affinity of binding sites (L/mg). The essential characteristic of the Langmuir isotherm can be expressed in terms of a dimensionless constant separation factor (Eq. (6)).

\[
R_L = \frac{1}{1 + C_0 b} \quad (6)
\]

Where \(C_0\) is the initial concentration of sorbet (mg/L) and the \(R_L\) value indicates the unfavorable (\(R_L>1\)), linear (\(R_L=1\)), favorable (\(0<R_L<1\)) or irreversible (\(R_L = 0\)) nature of adsorption process. Value of \(R_L\) for adsorption of BY28 is 0.14, which reveal the adsorption behavior of dye onto AR was favorable with \(R_L\) value less than 1.

Freundlich isotherm

The Freundlich isotherm is expressed by the following empirical equation (Freundlich 1906):

\[
q_e = k F C_e^{1/n} \quad (7)
\]

Where \(k\) and \(n\) are Freundlich constants, \(k\) indicating the adsorption capacity of the adsorbent (mg/g) and \(n\) the adsorption intensity (dimensionless).

\[
k = \frac{1}{q_m n} \quad (8)
\]

\[
F = \frac{1}{n} \quad (9)
\]
where \( K_q \) (mg/g) and \( 1/n \) are characteristic constants representing the adsorption capacity and adsorption intensity of the system respectively. The equation may be liberalized by taking the logarithm on both sides:

\[
lnq_e = lnK_F + \frac{1}{n}lnC_e \rightarrow (8)
\]

The isotherm constants for the isotherms studied and the correlation coefficient, \( R^2 \) with the experimental data are listed in Table 3. Based on the \( R^2 \) values, the adsorption of dye by clay fitted the two models. The maximum amount of adsorption of Basic Yellow 28 onto Romana clay was about 76.92 mg/g (Table 3) which is close to that found during the adsorption of methylene blue onto kaolinite-smectite rich clay where \( q_e \) value was 65.1 mg/g.\(^{12} \) While it is very important to that found during the adsorption of Methyl Orange by untreated clay collected from the region of Maghnia (Algeria) where the adsorption capacity was 35 mg/g.\(^{15} \) Similarly, it is very important to that found during the adsorption of ethyl violet on kaolinite where the adsorption capacity was 30.911 mg/g.

### Table 3 Langmuir and freundlich isotherm constants

| Langmuir | Freundlich |
|----------|------------|
| \( q_m \) (mg/g) | b (L/mg) | \( R^2 \) | \( K_F \) | \( n \) | \( R^2 \) |
| 76.92 | 0.119 | 0.14 | 0.996 | 2.863 | 1.773 | 0.981 |

### Adsorption kinetics

In order to determine the rate constants, pseudo-first order, pseudo-second order and intra-particle diffusion were employed to model the adsorption data during adsorption process (Table 4).

### Table 4 Kinetic parameters for the adsorption of BY28 onto AR

| Model type | Parameters | Values |
|------------|------------|--------|
| Pseudo-1\(^{st}\) order | \( k_1 \) (min\(^{-1}\)) | 0.032 |
| | \( q_e \) (calc.) (mg/g) | 1.72 |
| | \( q_e \) (exp.) (mg/g) | 15.33 |
| | \( R^2 \) | 0.9 |
| Pseudo-2\(^{nd}\) order | \( k_2 \) (min\(^{-1}\)) | 0.035 |
| | \( q_e \) (calc.) (mg/g) | 15.62 |
| | \( q_e \) (exp.) (mg/g) | 15.33 |
| | \( R^2 \) | 0.99 |
| Intra-particle diffusion | \( K_{st} \) (mg/g min0.5) | 0.48 |
| | \( R^2 \) | 0.95 |

### Table 5 Kinetic parameters for the adsorption of BY28 onto AR

| Adsorbent | \( \Delta H^\circ \) (Kj mol\(^{-1}\)) | \( \Delta S^\circ \) (J K\(^{-1}\) mol\(^{-1}\)) | \( \Delta G^\circ \) (Kj mol\(^{-1}\)) |
|-----------|------------------------|------------------------|------------------------|
| AR | 298K | 313 K | 333 K |
| | -15.68 | -0.041 | 3.46 | 2.23 | 1.822 |

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**Citation:** Islem C, Aref C, Jamoussi F et al. Adsorptive/desorptive potential of cationic basic yellow 28 (by28) dyes onto natural untreated clay from aqueous phase. *Open Access J Sci.* 2018;2(1):74–79. DOI: 10.15406/oajs.2018.02.00048
Thermodynamic studies

Thermodynamic parameters, i.e., the enthalpy change, $\Delta H^\circ$, and the entropy change, $\Delta S^\circ$ ads, are obtained by writing the van’t Hoff equation in the form:

$$ \ln \frac{q_e}{C_e} = \frac{\Delta H^\circ}{2.303RT} - \frac{\Delta S^\circ}{2.303R} \tag{12} $$

Where $q_e$ and $C_e$ are the adsorption capacity of the adsorbent and the equilibrium dye concentration in solution, respectively. The values of $\Delta H^\circ$ and $\Delta S^\circ$ were determined from the slope and intercept, respectively, of the van’t Hoff plots [$\log (q_e/C_e)$ vs. $1/T$] (Figure 8). The Gibbs change in free energy of the adsorption, $\Delta G^\circ$, is calculated using the following equation (Eq. 13):

$$ \Delta G^\circ = \Delta H^\circ - T\Delta S^\circ \tag{13} $$

The negative values of $\Delta G^\circ$ and negative value of $\Delta H^\circ$ (Table 6) indicated the spontaneous and exothermic nature of the adsorption. The magnitude of $\Delta H^\circ$ may be used to distinguish between physisorption (2.1-20.9 kJ/mol) and chemisorption (84 to 420 kJ/mol). The $\Delta H^\circ$ value at optimized conditions was 15.68 kJ/mol. This value is in the range indicative of physisorption. The negative value of entropy change ($\Delta S^\circ$) indicated decreased randomness at the solid/solution interface.

Desorption studies

Desorption studies were however conducted for BY28-loaded AR, as a function of pH, and results are shown in Figure 9. As it can be seen from Figure 9, that percent desorption decreased with increase in pH of the aqueous solution. The maximum BY28 desorbed percentage was 17% and it was obtained for strong acidic conditions (pH 2). This is just opposite to the pH effect. For higher pH values, desorption was very low and limited to 2-3%, which reflects the establishment of strong chemical bonds between the basic dye and the raw clay in the adsorption.

Adsorption mechanism of BY28 by AR

The smectite can adsorb cationic dye by different mechanisms: (i) exchange captions in the interlayer resulting from the interaction between dye and permanent negative charge and (ii) formation of internal sphere complexes by the silanol (Si-O-) and aluminol (Al-O-) groups of the clay. This mechanism is pH-dependent charge (Figure 10).

Conclusion

In summary, AR is found to be very effective in the removal of a basic dye from aqueous solution. Adsorption profile was highly pH dependent and was sought to be electrostatic in nature. The adsorption followed the second order kinetics with simultaneous occurrence of the intra particle diffusion of the dye. The adsorption obeyed to the Langmuir and Freundlich isotherms. The adsorption was spontaneous in nature.

Acknowledgements

None.
Conflict of interest

The authors declare that there is no conflict of interest.

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Citation: Islem C, Afef C, Jamoussi F et al. Adsorptive/desorptive potential of cationic basic yellow 28 (by28) dyes onto natural untreated clay from aqueous phase. Open Access J Sci. 2018;2(1):74–79. DOI: 10.15406/oajs.2018.02.00048