Capacity and Modeling of Acid Blue 113 Dye Adsorption onto Chitosan Magnetized by Fe₂O₃ Nanoparticles

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Capacity and modeling of acid blue 113 dye adsorption onto chitosan magnetized by Fe₂O₃ nanoparticles

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Abstract

A chitosan polymer was magnetized by coating with magnetite Fe_2O_3 nanoparticles, and the resultant material ($\text{C-Fe}_2\text{O}_3$) was first characterized through scanning electron microscopy equipped with energy-dispersive X-ray spectroscopy, transmission electron microscopy, atomic force microscopy, thermogravimetric, X-ray diffractometry, Fourier transform infrared spectroscopy, Brunauer–Emmett–Teller, and point of zero charge analyses. $\text{C-Fe}_2\text{O}_3$ was then employed as a separable and efficient adsorptive agent to remove acid blue 113 (AB113) dye from aqueous solution. The removal efficiency was optimized at different environmental parameter values (pH: 3–11, $\text{C-Fe}_2\text{O}_3$ dose: 0.1–1 g/L, initial AB113 dye concentration: 10–100 mg/L, adsorption time: 0–300 min, and temperature: 300–318 K). Under optimum conditions, an AB113 dye removal efficiency of 99.68% was achieved. In addition, the effect of the presence of NaCl , NaNO_3 , Na_2CO_3 , and MgSO_4 ions on the AB113 dye removal efficiency could be ranked as $\text{NaCl} > \text{NaNO}_3 > \text{MgSO}_4 > \text{Na}_2\text{CO}_3$. The statistical analysis using the coefficient of determination, root mean square error, chi-square test, sum of squared errors, and average relative error showed that the Freundlich and pseudo-second-order equations were the best mathematical models for fitting the isothermal and kinetics data. Further kinetics analyses showed that the adsorption of AB113 molecules on $\text{C-Fe}_2\text{O}_3$ active sites was dominated by the intraparticle diffusion process. Thermodynamic parameters indicated that the AB113 dye adsorption process was favorable, endothermic, and spontaneous. Furthermore, an increase in temperature had a positive impact on AB113 dye removal. The regeneration study confirmed the excellent shelf life of $\text{C-Fe}_2\text{O}_3$, with only a slight loss in the removal efficiency (<7%) being detected after six operational cycles of AB113 dye adsorption. Compared with other adsorbents,

C-Fe₂O₃ was more effective for the adsorption of AB113 dye, with an adsorption uptake up to 128 mg/g.

Keywords: Acid blue 113; Chitosan; Fe₂O₃ nanoparticles; Modeling; Adsorption study

1. Introduction

Currently, dye pollutants are among the most dangerous compounds that are pumped into the environment via the effluents of different factories, such as those for paper, textiles, paints, leather, and cosmetics (Wu et al., 2013; Elgin et al., 2008). Azo dyes are widely applied in various dyeing processes due to their high activity; thus, these types of dyes generally make up approximately 50–70% of the total dye quantity employed in industry (Shan et al., 2015; Bazrafshan et al., 2013; Giustetto et al., 2011). On the other hand, simple azo dyes (e.g., picric acid and anionic and cationic dyes) consist of hydroxyl groups in their structure and do not establish a strong bond with the texture of fabrics and linen fibers; thus, for dyeing, a large quantity of simple azo dyes are consumed (Crini et al., 2008; Balarak et al., 2020). Azo dyes contain natural and synthetic organic compounds and are characterized in nature as refractory and stable organic pollutants (Savic et al., 2014; Zhang et al., 2013). For example, acid blue 113 (AB113) dye has been selected as the target dye due to its wide application in industry, especially in the paints and textile industries. Furthermore, studies have shown that azo dyes have two types of auxochrome (-OH, -NH₂, and NR₂) and chromophore (C=C, N=N, and C=O) functional groups (Zhang et al., 2013; Eren et al., 2010). These two groups have a high tendency to attach to benzene and naphthalene rings that may be found in wastewater and, in some cases, can also be attached to aliphatic and heterocyclic groups, leading to the generation of hazardous byproducts (Sanghi et al., 2013; Huang et al., 2011; Ma et al., 2011; Li et al., 2010). In addition to their toxicity to humans, the presence of dyes in water bodies can disturb aquatic life via their

ability to reduce the penetration of sunlight, which is essential for photosynthesis (Hao et al., 2013; Balarak et al., 2020). Thus, wastewater containing dyes must go through efficient purification processes prior to discharge to the environment. In reality, dyes cannot be lowered to the acceptable concentration via pretreatment or conventional methods (Crini et al., 2008; Anirudhan et al., 2009). Therefore, researchers are looking for advanced techniques to effectively remove these compounds prior to discharge into water bodies (Elgin et al., 2008; Wu et al., 2013).

Low portions of dye pollutants are removed by conventional treatment methods. Furthermore, some efficient methods have high costs (Baskaralingam et al., 2006; Balarak et al., 2020). Therefore, alternative techniques to traditional dye removal methods are still in demand. Currently, the efficient and low-cost method used for this purpose is separation by adsorption processes (Sillanpää et al., 2021; Balarak et al., 2016; Cheung et al., 2007). One of the most important operational and economic factors of adsorption treatment is the adsorbent used, as well as separating the spent adsorbents after treatment. To solve the separation problem and simultaneously increase the adsorption efficiency of the used adsorbent, the magnetization of used adsorbents is a promising method. Coating used adsorbents with magnetic nanoparticles has also been recommended because this modification method not only improves the removal efficiency but also improves the economic efficiency, environmental compatibility, mechanical strength, reusability and scalability of the magnetized adsorbents (Xua et al., 2012; Kyzas et al., 2013). In this direction, Fe_2O_3 nanoparticles have been widely used (Broujeni et al., 2018; Mittal et al., 2014). However, the use of pure magnetic particles also has a drawback, as these particles have a high tendency to agglomerate in aqueous solution. When these particles are combined

with other materials, such as chitosan, the agglomeration mechanism is hampered (Kavitha et al., 2012).

Natural materials are welcomed and highly recommended for application as adsorptive treatment agents (Tabak et al., 2010). Chitosan is a natural hydrophilic and cationic biopolymer obtained by the removal of acetyl-chitin groups in an alkaline environment and has been extensively studied as an adsorbent to remove a variety of contaminants (e.g., heavy metals, antibiotics, and dyes) (Bazrafshan et al., 2013; Tabak et al., 2010; Gupta et al., 2005). Chitosan has low cytotoxicity and unique chemical properties; furthermore, chitosan is obtained from chitin, which is effortlessly attained from shrimp and crab shells (these are considered waste products of the seafood industry) (Tabak et al., 2010). Chitosan has two hydroxyl groups and one amino group in the glycosidic fraction and has a suitable matrix that allows fabrication with foreign particles (Xua et al., 2012). Because chitosan dissolves in acidic and aqueous solutions, there is no need to use hazardous organic solvents (Cheung et al., 2007; Khandanlou et al., 2013). Sun et al. used a chitosan/cellulose composite to adsorb lead ions from aqueous solutions (Sun et al., 2009). Furthermore, chitosan magnetized by Fe_2O_3 nanoparticles has also been employed for eradicating heavy metals from solutions (Chokami et al., 2017; Broujeni et al., 2018).

Through our review of previous research, it was found that very few studies have been performed on the removal of organic pollutants, such as dyes using chitosan magnetized by Fe_2O_3 nanoparticles (hereafter referred to as C- Fe_2O_3). Therefore, the present study offers a comprehensive account of the application of C- Fe_2O_3 as an efficient and recyclable adsorbent for the removal of AB113 dye from aqueous solutions. Initially, the surface and structural characterizations of used C- Fe_2O_3 are obtained by advanced characterization analyses. The adsorption process of the AB113 dye and C- Fe_2O_3 couple is reported with different

environmental parameters. Isotherm and kinetics studies, which are the focus of this study, are conducted using linear and nonlinear models. The thermodynamics parameters are also determined. Finally, the recyclability of C-Fe₂O₃ is evaluated in several AB113 dye adsorption–desorption cycles.

2. Materials and methods

2.1 Materials

Chitosan with a chemical formula (C₆H₁₁O₄N)_n and purity of 98% was purchased from Sigma–Aldrich (Germany). Acetic acid (≥99.7%), FeCl₃.6H₂O (≥98%), FeCl₂.4H₂O (≥99%), HCl (36.5%), and NaOH (≥98%) were purchased from Merck (Germany). In addition, a stock solution of AB113 dye (1000 mg/L) was bought from the Alvan Sabet factory (Hamadan, Iran).

2.2 Preparation of C–Fe₂O₃

Initially, 2 g of chitosan was gradually dissolved in deionized water (100 mL) containing 2% acetic acid and magnetically stirred at 120 rpm for 1 h. To ensure the dissolution of all used chitosan quantities, the prepared chitosan solution was homogenized by means of ultrasonic waves for 20 min using an ultrasonic device (Sunshine ultrasonic cleaner, model SS6508T). A mixture of FeCl₃.6H₂O (6.1 g) with FeCl₂.4H₂O (4.2 g) at a molar ratio of 2:1 was mixed with the chitosan solution with the simultaneous dropwise addition of a 30% NaOH solution (this process was carried out through the employment of a nitrogen gas stream at a temperature ≥70 °C, while being mixed at 1000 rpm for 2 h). Subsequently, the mixture was homogenized by ultrasonic waves for 20 min. After that, the temperature of the homogenized mixture was increased to 90 °C, and when reaching this temperature, the stirring process was further continued for 2 h. Next, the resulting mixture was deoxygenated 4 times with distilled water and rinsed with ethanol. The supernatant (C–Fe₂O₃) was then collected in a container and

neutralized. Afterward, a centrifuge (3600 rpm for 10 min) was used to separate C-Fe₂O₃ from the solution, followed by drying in an oven (70 °C for 5 h). Eventually, after crushing and crystallization, C-Fe₂O₃ was placed in a desiccator in the presence of a magnetic field for dehumidification.

2.3 Characterization analyses

Surface and structural characterizations of the C-Fe₂O₃ sample were performed using XRD with a Cu K α radiation source (ESCALAB250 Thermo Fisher), SEM (Model JSM6510LV, JEOL Ltd., Tokyo, Japan) connected with an EDX attachment (SEM/EDX), TEM (Philips CM120, Netherlands), FTIR spectroscopy (PerkinElmer, Spectrum GX), and UV-Vis spectrophotometry (DR5000, HACH). The magnetization curves of Fe₂O₃ nanoparticles and C-Fe₂O₃ were measured using a vibrating sample magnetometer (Lake Shore Cryotronics, 7407-S). The improvement in the thermal stability of chitosan after magnetization with Fe₂O₃ nanoparticles was determined using thermogravimetric analysis (TGA, TA Instruments SDT Q600, USA). The surface area and pore size were obtained by employing Brunauer-Emmett-Teller (BET) analyses. The changes in the topography of the C-Fe₂O₃ surface due to AB113 dye adsorption were explored using atomic force microscopy (AFM, Nano Wizard II, JPK Germany). The pH_{pzc} value of C-Fe₂O₃ was evaluated using a previously reported methodology (Khodadadi et al., 2019). In addition, the magnetic properties of chitosan and C-Fe₂O₃ were determined at ambient temperature using a vibrating-sample magnetometer (VSM, MDKB).

2.4 Batch experiments

The AB113 dye adsorption experiments were carried out in several 200 mL Erlenmeyer flasks with a dye solution of 100 mL. AB113 dye removal by C-Fe₂O₃ was optimized by varying the pH (3–11), C-Fe₂O₃ dose (0.1–1 g/L), initial AB113 dye concentration (10–100 mg/L),

adsorption time (0–300 min), and temperature (388–318 K). In addition, the effect of the presence of interference ions (NaCl, NaNO₃, Na₂CO₃, and MgSO₄) on the adsorptive performance of C–Fe₂O₃ was examined. All flasks were placed in a shaker incubator at 120 rpm. Except for the thermodynamic experiments, which were performed from 288 to 318 K, the flasks were maintained at room temperature (25±2 °C) throughout the adsorption experiments. For the kinetic study, the experimental data of the analysis of the effects of the initial AB113 dye concentration on the C–Fe₂O₃ adsorption capacity were used. Furthermore, for the isotherm study, an experiment was conducted to determine the relationship between the adsorption capacity of C–Fe₂O₃ at equilibrium and the AB113 dye concentration. During the adsorption process, a 3 mL liquid sample was taken from each flask at regular adsorption time intervals and subjected to centrifugation at 1509 x g for 15 min. The concentration of AB113 dye in the separated supernatant was determined using a UV–Vis laboratory spectrophotometer (DR 5000, HACH). From the determined AB113 dye concentrations, the dye removal efficiency (R (%)) and adsorption capacity (or uptake) (q_t) of C–Fe₂O₃ for AB113 dye were calculated using Equations (1) and (2), respectively (Khodadadi et al., 2019; Mittal et al., 2014).

$$RE (\%) = \frac{(C_0 - C_t)}{C_0} \times 100 \quad (1)$$

$$q_t = \frac{(C_0 - C_t)}{M/V} \quad (2)$$

where C_0 is the initial AB113 dye concentration (mg/L), C_t is the AB113 dye concentration after a specific time of the adsorption process (t , min), M is the C–Fe₂O₃ mass (g), and V is the volume of the AB113 dye solution (L) (100 mL). Note that the AB113 dye concentration and C–Fe₂O₃ adsorption capacity at equilibrium time are denoted in Equation (1) by C_e (mg/L) and q_e (mg/g), respectively.

2.5 Error function analysis

The purpose of conducting the error function analysis in the present study is to assess the compatibility of the theoretical isotherm and kinetics model with the experimental data. However, the experimental data were fitted with both linear and nonlinear equations of these models. Depending on the values of the coefficient of determination (R^2), sum of squared errors (SSE) (Equation 3), root mean square error ($RMSE$) (Equation 4), chi-square test (χ^2) (Equation 5), and average relative error (ARE) (Equation 6), the most appropriate model for the experimental data was evaluated (Hu et al., 2015; Fontana et al., 2016).

$$\text{sum of squared errors (SSE)} = \sum_{i=1}^n (q_{e \text{ exp}} - q_{e \text{ cal}})^2 \quad (3)$$

$$\text{root mean square error (RMSE)} = \sqrt{\frac{1}{n-p} \sum_{i=1}^n (q_{e \text{ exp}} - q_{e \text{ cal}})^2} \quad (4)$$

$$\text{chi-square test } (\chi^2) = \sum_{i=1}^n \left[\frac{(q_{e \text{ exp}} - q_{e \text{ cal}})^2}{q_{e \text{ cal}}} \right] \quad (5)$$

$$\text{average relative error (ARE)} = \frac{100}{n} \sum_{i=1}^n \left[\frac{q_{e \text{ exp}} - q_{e \text{ cal}}}{q_{e \text{ exp}}} \right] \quad (6)$$

where $q_{e \text{ exp}}$ is the experimental uptake determined based on Equation (2), and $q_{e \text{ cal}}$ is the calculated uptake determined from the application of the relevant model with the experimental data.

3. Results and discussion

3.1 Characterization analyses

BET analysis revealed that the surface area of C-Fe₂O₃ is 112.9 m²/g. In addition, the pore volume and mean diameter are 0.286 cm³/g and 17.2 nm, respectively. Compared with the documented surface area value for a sample of pure chitosan (91.4 m²/g) (Broujeni et al., 2018), it can be concluded that the coating of chitosan with Fe₂O₃ significantly enhances the surface

area of this polymer. Notably, solid materials with a large surface area are favorable as adsorbents in adsorption treatment systems, as this parameter reflects the number of adsorption sites for pollutant molecules.

The SEM images of the C-Fe₂O₃ before and after AB113 dye adsorption are shown in Fig. 1a and 1b, respectively. It appears that C-Fe₂O₃ exhibits an extremely rough surface and porous structure, and many fragments are randomly spread onto this material (Fig. 1a). In reality, these morphological characterizations of C-Fe₂O₃ indicate that this adsorbent has a large reaction area for adhering AB113 dye molecules. After AB113 dye adsorption (Fig. 1b), the surface and pores of C-Fe₂O₃ are completely covered, which provides an indication about the ability of C-Fe₂O₃ to adsorb AB113 molecules. The EDX analysis (equipped on an SEM device) performed on the C-Fe₂O₃ sample shows the presence of C (39.95%), N (11.31%), O (45.72%), and Fe (3.02%). C and N are essential elements in the structure of chitosan (Cheung et al., 2007; Xua et al., 2012). On the other hand, the presence of Fe and O in the structure of C-Fe₂O₃ is the result of the successful magnetizing process of chitosan by Fe₂O₃ nanoparticles.

Fig. 1c is a TEM image of the C-Fe₂O₃ sample taken at a scale of 50 nm. From this figure, it is perceived that the Fe₂O₃ nanoparticles are entrenched into the chitosan structure, where the light-colored area belongs to chitosan and the dark spots represent the Fe₂O₃ nanoparticles. Furthermore, the TEM image reveals that the synthesized C-Fe₂O₃ particles are almost spherical in shape, and no aggregation is observed between these particles. The size of the C-Fe₂O₃ particles is approximately 50 nm, as also confirmed by XRD.

The X-ray diffraction pattern of C-Fe₂O₃ is depicted in Fig. 1d. The C-Fe₂O₃ has major peaks at $2\theta=63^\circ$ and 36° . The average size of C-Fe₂O₃ is calculated by substituting the main peak phase

value of the C-Fe₂O₃ in the Scherrer formula (Equation 7). Based on this equation, the synthesized nanoparticle size in the main peak at 2θ=36° (with an intensity of approximately 120 a.u.) is determined to be 35.5 nm.

$$D = \frac{K\lambda}{\beta \cos \theta} \quad (7)$$

where D is the mean size of C-Fe₂O₃, K is a fixed number equal to 0.89, λ is the wavelength of the Cu-K α radiation (nm), β is the width of the main peak at half its intensity (radians) and θ (diffraction angle, radians) is the peak position in the XRD pattern (Hilal et al., 2012).

Fig. 1. SEM images of C-Fe₂O₃ before (a) and after (b) AB113 adsorption. TEM image of C-Fe₂O₃ (c). X-ray diffraction pattern of C-Fe₂O₃ (d).

Fig. 2 shows the 2D and 3D AFM images, which are used to compare the topographical changes on the C-Fe₂O₃ surface as a result of AB113 dye adsorption. Before adsorption (Fig. 2A), the C-Fe₂O₃ topography is characterized by many scattered bumps and small masses. It is clear that these masses cluster around large blocks, which may be due to the Fe₂O₃ nanoparticles attracting them. However, in Fig. 2B, it can be seen that after AB113 dye adsorption, remarkable changes occur in the surface topography of C-Fe₂O₃. The previously diagnosed bumps are completely covered, and more masses are dispersed onto the C-Fe₂O₃ surface, which denotes the existence of AB113 dye molecules on the C-Fe₂O₃ surface. Similar results relevant to the AFM analysis in the present study have been presented in previous works (Broujeni et al., 2018; Kyzas et al., 2013).

Fig. 2. 2D and 3D AFM images of C-Fe₂O₃ before (A) and after (B) adsorption of AB113 dye.

The TGA graph of chitosan and C-Fe₂O₃ is presented in Fig. 3a. The first weight loss detected up to 130 °C is related to the dehydration of water molecules from all the chitosan and C-Fe₂O₃. The second thermal step occurs in the temperature range of 130–320 °C. The further dehydration, depolymerization, and acetylation process of chitosan, which often occurs at this temperature range, can be the reason for the detected weight loss. The third step is observed at temperatures >320 °C. is the weight loss at this step is ascribed to thermal decomposition in the chitosan structure at high temperatures. These thermal steps have also been reported in other studies

(Tabak et al., 2010; Xua et al., 2012). In the case of C-Fe₂O₃, decomposition and weight loss are observed at higher temperatures compared to chitosan, which indicates an improvement in the thermal stability of chitosan by the incorporation of Fe₂O₃ nanoparticles; this has also been reported by other studies (Broujeni et al., 2018).

The FTIR spectra of chitosan (C), C-Fe₂O₃, and C-Fe₂O₃ after reaction with AB113 dye (C-Fe₂O₃/AB113) samples are shown in Fig. 3b. In the chitosan spectrum, peaks related to C-H, N-H, and C-O-C bands are observed at 2875 cm⁻¹, 1650 cm⁻¹, and 1066 cm⁻¹, respectively. All of the above peaks can be detected in the spectrum of C-Fe₂O₃. Peaks at 1570 cm⁻¹ and 1260 cm⁻¹ are observed, and the peak at 2420 cm⁻¹ in chitosan becomes the larger peak at 2340 cm⁻¹ in C-Fe₂O₃. The resulting changes in the spectrum of C-Fe₂O₃ indicate the combination of Fe₃O₄ with chitosan (Broujeni et al., 2018). Additionally, the peak of the Fe-O band can be spotted at 590 cm⁻¹, which indicates the existence of magnetite Fe₂O₃ nanoparticles in the produced samples.

The magnetization curves of chitosan and C-Fe₂O₃ provided by the VSM analysis are presented in Fig. 3c and d, respectively. From this figure, the saturation magnetization value of C-Fe₂O₃ is 24.5 emu/g, which shows a 46% reduction in magnetization compared to 45.5 emu/g for Fe₃O₄ nanoparticles. This decrease in magnetic properties is due to the fabrication of chitosan with Fe₃O₄ magnetic nanoparticles; thus, this material can be separated from aqueous solutions using a magnet (Kyzas et al., 2013). Notably, C-Fe₂O₃ is a highly magnetic material, despite containing nonmagnetic chitosan (Kavitha et al., 2012). Based on the VSM analysis, the superparamagnetic structure of the Fe₃O₄ nanoparticles is confirmed in the present study. The absence of a residual loop in the VSM diagram indicates the supermagnetic behavior of the Fe₃O₄ nanoparticles, although the interaction with chitosan leads to a decrease in their intensity (Broujeni et al., 2018).

Fig. 3. TGA graph of chitosan and C-Fe₂O₃ (a). FTIR spectra of chitosan, C-Fe₂O₃ and C-Fe₂O₃ after reaction with the AB113 dye (C-Fe₂O₃/AB113) (b). Magnetization curves of the Fe₂O₃ nanoparticles (c) and C-Fe₂O₃ (d).

3.2 Effects of the environmental parameters on AB113 adsorption

Several adsorption studies have demonstrated that the adsorption process of organic pollutants is significantly affected by various environmental parameters (Nasseh et al., 2021; Al-Musawi et al., 2018; Rostamian and Behnejad, 2016). Generally, the solution pH and temperature have a considerable effect on the adsorption process, as the ionization degree of the adsorbent surface and the dissolution rate of the organic pollutant molecules mainly depend on these two parameters. From an economic perspective and for designing large-scale adsorption treatment

systems, the determination of the optimum adsorbent dose for the removal of pollutants is considered a key factor for such studies. Furthermore, the most important analyses in the isotherm study depend on the relationship between the equilibrium uptake of the adsorbent for the target pollutant at a constant temperature. Investigation of the effects of adsorption time at different initial pollutant concentrations is essential for kinetic studies. Additionally, Samarghandi et al. (2015) reported that the presence of some ions in aqueous solution may compete with the absorption of the target pollutant. In the present work, the effects of the abovementioned environmental parameters on AB113 dye adsorption and removal efficiency were studied, and the results are depicted in Fig. 4. Moreover, the temperature effect data are presented in the thermodynamic study.

3.2.1 Solution pH

The solution pH of the solution greatly affects the rate of adsorption; under acidic conditions, the adsorbent surface becomes protonated and has a more positive charge. This charge causes an electrostatic interaction between the adsorbent and negatively charged ions in the water. Conversely, in an alkaline state, the adsorbent surface becomes negatively charged, and the interaction between the adsorbent and the positive ions increase (Fontana et al., 2016). In the present study, the effect of solution pH on the removal efficiency of AB113 dye by C-Fe₂O₃ was investigated from pH 3 to 11, and the results are depicted in Fig. 4a (AB113 concentration: 10 mg/L, adsorption time: 120 min, and C-Fe₂O₃ dose: 0.6 g/L). As seen, with an increasing pH, the dye adsorption efficiency decreases significantly. At low pH values, more protons exist, so the amine groups in chitosan are protonated; thus, the electrostatic force between the negatively charged dye molecules and the positively charged adsorbent sites increases and more adsorption occurs (Tabak et al., 2010). Additionally, the high adsorption under acidic conditions can be explained by a pH_{pzc} analysis. In the present study, the pH_{pzc} of C-Fe₂O₃ is determined to be

approximately 7 (inset picture in Fig. 4a). Based on this, at solution pH values greater than 7, the surface of C-Fe₂O₃ is negatively charged, while for pH values lower than 7, the surface has a positive charge (Hilal et al., 2012). In contrast, the AB113 dye molecule (having two sulfonated (-SO₃⁻) groups) is a disulfonate acid dye (with a pKa of 0.5), and at pH values higher than 0.5, AB113 tends to have a negative charge. Thus, under acidic conditions, increasing the density of positive charges on the C-Fe₂O₃ surface may be behind the enhanced AB113 removal. At higher pH values, OH⁻ in the environment competes with the studied pollutant for combining with chitosan, thus reducing the adsorption rate of AB113 dye on the cationic adsorbent (Bazrafshan et al., 2013).

3.2.2 C-Fe₂O₃ dose

The effect of the C-Fe₂O₃ dose was investigated by varying the dose from 0.1 g/L to 1 g/L with the following experimental conditions: initial concentration of AB113=10 mg/L, pH=3, and adsorption time: 120 min. As shown in Fig. 8, it is revealed that with an increasing quantity of C-Fe₂O₃, the removal efficiency also increases. This result is because of the increase in total surface area by increasing the C-Fe₂O₃ dose, which in turn increases the availability of active adsorption sites for AB113 adsorption. The maximum adsorption of AB113 is observed at 0.6 g/L with an adsorption capacity of 124.2 mg/g for C-Fe₂O₃. Therefore, a dose of 0.6 g/L was selected for further batch experiments. Notably, the removal of AB113 onto C-Fe₂O₃ is found to be constant with an increasing dose. The reduction in AB113 removal with a further increase in the adsorbent dose might be due to a decrease in the available sites and surface area and due to the agglomeration of C-Fe₂O₃ particles that occurs at high doses (Dotto et al., 2011). The results of this experiment are in accordance with the results of previous studies (Yao et al., 2010).

3.2.3 AB113 concentration and contact time

Fig. 4c shows the kinetics curves, which represent the removal efficiency of AB113 dye as a function of adsorption time (0–300 min) with different concentrations (10–100 mg/L). The other parameters were fixed as the following: pH =3 and C-Fe₂O₃ dose =0.6 g/L. The plotted curves of Fig. 4c present similar profiles at all analyzed concentrations of AB113 dye. These curves show a very steep increase in the first 60 min of adsorption, and after 90 min of adsorption, the slopes of these curves gradually level off until reaching equilibrium at 120 min. Clearly, the AB113 removal efficiency increases from 77.93% to 99.68% when the AB113 dye concentration is decreased from 100 mg/L to 10 mg/L. The decrease in the removal efficiency with an increasing initial AB113 concentration is because of the increase in the competition rate among AB113 molecules to occupy the available active sites of a fixed quantity of C-Fe₂O₃. In addition, the increase in the AB113 dye concentration can lead to the rapid exhaustion of the used adsorbent; therefore, the adsorbent can become saturated (Joshi et al., 2019). On the other hand, Fig. 4c reveals that the AB113 removal efficiency is faster in the first 30 min, and the removal efficiency beyond 30 min is slightly slower. This result is because in the first 30 min of adsorption, the active adsorption sites on C-Fe₂O₃ are easily available for AB113 molecules (Madrakian et al., 2011). Afterward, the number of free sites decrease; hence, the AB113 dye removal efficiency is slightly slower, particularly after 60 min of adsorption.

3.2.4 Interference ions

Several studies have shown that different ions can interfere with the adsorption process by affecting the electrostatic interactions between the adsorbent and pollutant molecules (Balarak et al., 2016; Samarghandi et al., 2015). In fact, the reason for this phenomenon is that these ions have a higher ability to adsorb or neutralize the positive or negative charges on the adsorbent surface; thus, they inhibit the attraction between the pollutant molecules and adsorbent particles

(Balarak et al., 2016). In the present study, the competition of NaCl, NaNO₃, Na₂CO₃, and MgSO₄ ions on AB113 dye adsorption was investigated and compared with a control sample. This experiment was performed under the optimized conditions found from the previous analyses (pH =3, AB113 concentration = 10 mg/L, C-Fe₂O₃ dose = 0.6 g/L, and adsorption time = 120 min), and the results are presented in Fig. 4d. In addition, the initial concentration of these interference ions was fixed at 10 mg/L. The presence of NaCl has the greatest impact on the AB113 removal efficiency. From this experiment, the effect of the presence of interference ions on the AB113 removal efficiency can be ranked as NaCl> NaNO₃> MgSO₄> Na₂CO₃.

Fig. 4. Effect study of pH and the pH_{pzc} analysis (a), C-Fe₂O₃ dose (b), Initial Ab113 dye concentration (c), and Interference ions (d).

3.3 Kinetics study

Analyses of the kinetic reaction are essential to evaluate the applicability of the used adsorbent, as it is useful for understanding the rate and type of adsorption (chemisorption, physisorption, or mixed). In addition, the determination of a kinetic model that can describe the experimental data of the kinetic reaction of an adsorbent-adsorbate system is necessary for the precise design of large-scale adsorption treatments (Al-Musawi et al., 2018; Balarak et al., 2016). Therefore, the

experimental data shown in Fig. 4c are first treated using Equation 2 and then modeled with the pseudo-first-order (PFO) (Equation 8) and pseudo-second-order (PSO) (Equation 9) kinetic models (Ferrero et al., 2018; Somasekhara et al., 2016).

$$q_t = q_e(1 - \exp(-k_1 t)) \quad (8)$$

$$q_t = \frac{q_e^2 k_2 t}{1 + q_e k_2 t} \quad (9)$$

where K_1 (1/min) and K_2 (g/mg·min) are the PFO and PSO rate constants, respectively.

Nonlinear regression methodology using the MATLAB program was applied to model the kinetic data. Table 1 lists the results of the kinetic models and regression level parameters of AB113 dye adsorption onto C—Fe₂O₃. Note that the fitting of each kinetic model with the experimental data is estimated in accordance with the R^2 , SSE , $RMSE$, χ^2 , and ARE values as well as the convergence between the calculated uptake ($q_e(cal)$) and experimental uptake ($q_e(exp)$). The results show that high regression levels are obtained for the fitting of the PSO model compared to those for the fitting of the PFO model (Table 1). In addition, the values of $q_e(cal)$ determined from the application of the PSO model are close to the $q_e(exp)$ values. Therefore, the adsorption kinetics of the AB113 dye on C—Fe₂O₃ follows PSO kinetics, suggesting a chemisorption process (Mohammed et al., 2019; Khodadadi et al., 2019).

By plotting the kinetics data according to the intraparticle diffusion model (IPD, Equation 12), it is found that the kinetics adsorption of AB113 dye onto C—Fe₂O₃ consists of three consecutive phases, as shown in Fig. 5: bulk diffusion, film diffusion, and pore diffusion. In fact, the first phase observed at $3 < t^{0.5} < 5.5 \text{ min}^{0.5}$ represents the surface and IPD processes, the second phase observed at $5.5 < t^{0.5} < 11 \text{ min}^{0.5}$ represents liquid film diffusion, and the third phase observed at $11 < t^{0.5} < 17.5 \text{ min}^{0.5}$ represents the diffusion of AB113 dye molecules through

pores to the active sites of C-Fe₂O₃; then, equilibrium conditions are achieved. In the present study, the first phase is modeled as a term in the IPD model. This is an important step in adsorption studies, as the IPD model provides information about the role of the IPD rate in controlling adsorption (Balarak et al., 2016).

$$q_t = K_b t^{0.5} + C \quad (12)$$

where K_b (mg/g·min^{0.5}) is the rate constant of the IPD model and C (mg/g) is the IPD constant that provides information about the thickness of the boundary layer.

The values of K_b and C represent the slope and intercept of the linear plot equation of the first phase in Fig. 5, respectively. The results show a high compatibility level ($R^2 > 0.99$) of the AB113 kinetic data with the IPD model (Table 1). Therefore, it can be concluded that compared to other diffusion types, the IPD process is the dominant rate-controlling step during the adsorption of AB113 dye onto C-Fe₂O₃. This result is because the linear plot of q_t versus $t^{1/2}$ does not pass through the origin; therefore, boundary layer diffusion occurs during the adsorption process. The positive values of C for all AB113 concentrations are indicative of involving the IPD in the adsorption process; nevertheless, the adsorption process is governed not only by IDP as the rate-limiting step but also by other factors controlling AB113 adsorption on C-Fe₂O₃.

Table 1. Results of the kinetic analyses of AB113 dye adsorption on C-Fe₂O₃

PFO kinetics model								
C_0 (mg/L)	$q_e(exp)$	$q_e(cal)$	K_1	R^2	χ^2	RMSE	SSE	ARE
10	16.61	9.24	0.026	0.977	11.20	17.10	9.25	13.84
25	38.86	21.18	0.023	0.975	13.95	11.84	9.38	13.02
50	71.21	42.84	0.022	0.969	14.48	11.95	18.41	9.28
100	129.80	11.71	0.021	0.978	8.46	6.45	4.71	8.73
PSO kinetics model								
C_0 (mg/L)	$q_e(exp)$	$q_e(cal)$	K_2	R^2	χ^2	RMSE	SSE	ARE
10	16.61	17.36	0.0053	0.999	1.48	2.15	1.09	2.71
25	38.86	40.98	0.0018	0.999	1.76	2.95	1.49	3.11
50	71.21	75.75	0.0008	0.998	2.35	1.79	4.18	1.48
100	129.80	140.84	0.0003	0.998	3.78	4.29	2.18	1.73
Intraparticle diffusion model								
C_0 (mg/L)	K_b (mg/g·min ^{0.5})		C (mg/g)	R^2				
10	2.63		1.35	0.999				
25	6.25		4.85	0.999				
50	12.1		13.9	0.998				
100	13.4		32.9	0.999

Fig. 5. Three stages of the kinetics adsorption of AB113 onto C-Fe₂O₃.

3.4 Adsorption isotherms

To clarify the interaction between the adsorbent and target pollutant, adsorption isotherms, i.e., the Langmuir (Equation 13), Freundlich (Equation 14), Dubinin-Radushkevich (D-R, Equation 15), and Temkin (Equation 16) equations, were employed. The occurrence of adsorption in monolayers at homogeneous sites on the adsorbent is hypothesized by the Langmuir model. In contrast, the Freundlich isotherm describes adsorption on a heterogeneous surface and reversible adsorption (Akar et al., 2010). Moreover, by the D-R model, the chemical or physical mechanism is appraised (Arami et al., 2006). Through employment of the Temkin isotherm, the heat of adsorption and the adsorbent–adsorbate interaction are defined (Fiorentin et al., 2010; Lakshmi et al., 2009).

$$q_e = \frac{Q_m K_L C_e}{1 + K_L C_e} \quad (13)$$

$$q_e = K_F C_e^{1/n} \quad (14)$$

$$q_e = q_m \exp(-\beta \varepsilon^2) \quad (15)$$

$$q_e = B \ln(K_T C_e) \quad (16)$$

where Q_m (mg/g) is a very important adsorption parameter in the Langmuir model, denoting the maximum q_t of the used adsorbent for the target pollutant; K_L (L/mg) is an equilibrium constant reflecting the affinity level of the active sites of the adsorbent; K_F is the Freundlich constant, indicative of the binding energy (mg/g)(L/mg)^{1/n}; $\frac{1}{n}$ is the heterogeneity parameter; q_m is the theoretical adsorption capacity (mg/g) of the D-R model; β (mol²/kJ²) is the activity coefficient constant associated with the mean free sorption energy, which is denoted by $E = \frac{1}{\sqrt{-2\beta}}$ (kJ/mol); $\varepsilon = RT \ln(1 + \frac{1}{C_e})$ is the Polanyi potential (kJ/mol); K_T and $B = \frac{RT}{b}$ are Temkin model constants

that provide information about the heat of sorption (J/mol); b is the Temkin isotherm constant (L/g); R is the ideal gas constant (8.314 J/(mol. K)); and T is the thermodynamic absolute temperature (K).

In addition to the Langmuir model (Equation 13), the characteristics of adsorption can be defined based on the dimensionless separation factor (R_L) (Equation 17). Based on the value of this factor, the favorable adsorption case can only be detected in the case of $0 < R_L < 1$.

$$R_L = \frac{1}{(1+K_L C_0)} \quad (17)$$

Table 2 lists the determined values of the isotherm model parameters. The results show that the fitting parameters are high for the Freundlich model compared to those for the other isotherm models; thus, the isothermal data of AB113 dye adsorption on C-Fe₂O₃ is consistent with the Freundlich model. This finding indicates that the AB113 dye molecules demonstrate multilayer adsorption at the heterogeneous adsorption sites of C-Fe₂O₃. In addition, as the value of the Freundlich parameter n is herein considered greater than one, the AB113 dye on the C-Fe₂O₃ adsorption process is suitable (Akar et al., 2009). The maximum adsorption capacity of AB113 by C-Fe₂O₃ based on the fitting results of the Langmuir model is 128.2 mg/g. Furthermore, the value of the R_L parameter of the Langmuir model is between zero and one, indicating the favorable adsorption process of AB113 dye on C-Fe₂O₃ (Zazouli et al., 2013). Since the E obtained from the D-R isotherm is less than 8 kJ/mol, the adsorption of AB113 on C-Fe₂O₃ is physical in nature (50). The theoretical models used for fitting the experimental isothermal data of AB113 dye adsorption on C-Fe₂O₃ are graphically presented in Fig. 6.

Considering the results of the isotherm study, the adsorption capacity of C-Fe₂O₃ was compared with that of other adsorbents in dye removal. As reported in Table 3, the adsorption capacity of

C-Fe₂O₃ is good and shows high adsorption compared to other adsorbents. Therefore, C-Fe₂O₃ is more effective than other adsorbents for dye removal.

Table 2. Isotherm model parameters of AB113 dye adsorption on C-Fe₂O₃ (conditions of this experiment were fixed at the following optimized values: pH = 3; C-Fe₂O₃ dose = 0.1–1 g/L; AB113 dye concentration = 10 mg/L; adsorption time = 120 min, and temperature = 25±2 °C)

Freundlich		Langmuir		Dubinin–Radushkevich		Temkin	
K_F (mg/g)(L/mg) ^{1/n}	6.961	Q_m (mg/g)	128.200	q_m (mg/g)	58.710	B (J/mol)	47.401
$1/n$	0.392	R_L	0.209	β (mol ² /kJ ²)	0.008	K_T (J/mol)	19.411
n	2.551	K_L (L/mg)	0.322	E (kJ/mol)	7.491	b (L/g)	52.260
R^2	0.994	R^2	0.941	R^2	0.674	R^2	0.839
χ^2	2.920	χ^2	4.714	χ^2	11.650	χ^2	4.762
$RMSE$	2.490	$RMSE$	3.961	$RMSE$	4.792	$RMSE$	9.890
SSE	1.843	SSE	6.721	SSE	9.460	SSE	14.811
ARE	2.080	ARE	4.340	ARE	10.380	ARE	19.371

Fig. 6. Experimental and theoretical isothermal data of AB113 dye adsorption on C-Fe₂O₃.

Table 3. Comparison between the maximum Langmuir adsorption capacities (Q_m) of C-Fe₂O₃ and of other adsorbents used for dyes removal

Adsorbents	Anionic dye	Q_m (mg/g)	Reference
Multi-walled carbon nanotubes	Acid Blue 225	10.2	(Balarak et al., 2016)
Cyperus rotundus	Acid Orang 7	31.98	(Balarak et al., 2016)
Azolla filiculoides	Acid Blue 92	36.98	(Balarak et al., 2016)
Orange bagasse	Reactive Blue 5	40.71	(Fiorentin et al., 2010)
Hazelnut shell	Acid Blue 25	60.2	(Ferrero et al., 2007)
Jujuba seed	Congo Red	55.56	(Somasekhara et al., 2012)
Canola	Acid Blue 113	56.9	(Zazouli et al., 2013)
Durian peel	Acid Green 25	63.29	(Hameed et al., 2008)
Rice husk	Acid Red 66	65.1	(Balarak et al., 2018)
Risk husk	Acid Blue 74	97.06	(Lakshmi et al., 2009)
Pyracantha coccinea	Acid Red 44	105.1	(Akar et al., 2010)
Thuja orientalis cone	Acid Blue 40	114.9	(Akar et al., 2008)
Meal hull	Acid Blue 92	114.9	(Arami et al., 2010)
C-Fe ₂ O ₃	Acid Blue 113	128.2	This study

3.5 Effect of temperature and the thermodynamic study

The effect of temperature on the AB113 dye removal efficiency was examined in the range of 288 – 318 K as a function of adsorption time (0–300), and the results are depicted in Fig. 7. This figure clarifies the positive effect of an increasing temperature on the removal efficiency of AB113 dye on C-Fe₂O₃. This behavior of improved removal efficiency with an increasing

temperature is due to in the enhanced diffusion rate or kinetic energy of AB113 dye molecules across the bulk and internal boundary layers of the C-Fe₂O₃ particle (Mohammed et al., 2019).

The thermodynamic parameters, including the enthalpy (ΔH^o) (kJ/mol), entropy (ΔS^o) (kJ/mol·K), and Gibbs free energy (ΔG^o) (kJ/mol), were calculated after determining the q_e values of the data in Fig. 7 using Equation 2. The above three parameters were determined using Equations 18–20 (Meziti et al., 2012; Hameed et al., 2008). From these equations, ΔH^o and ΔS^o can be directly calculated from the linear plot of $\ln K_o$ vs. $1/T$, where K_o (L/mg) is the equilibrium constant ($= \frac{q_e}{c_e}$). The slope and intercept with the y-axis of the obtained trend line equation represent $(-\Delta H^o/R)$ and $(\Delta S^o/R)$, respectively. Thereafter, ΔG^o can be calculated using Equation 20.

$$\Delta G^o = -RT(\ln K_o), \quad (18)$$

$$\ln K_o = \frac{\Delta S^o}{R} - \frac{\Delta H^o}{RT}, \quad (19)$$

$$\Delta G^o = \Delta H^o - T\Delta S^o, \quad (20)$$

Table 4 presents the results of the thermodynamic analysis. A positive ΔS^o indicates the high randomness of AB113 dye adsorption and approves the stability of the adsorption process. The negative values of ΔG^o suggest that the adsorption of AB113 dye on C-Fe₂O₃ is a spontaneous reaction (Chu et al., 2013; Han et al., 2009). Moreover, the increase in the negative values of ΔG^o with an increase in temperature may be because of the dehydration phenomena of both adsorbent particles and pollutant molecules at high temperatures, which simplifies the reaction between them and ultimately makes the adsorption of AB113 dye on C-Fe₂O₃ more favorable at high temperatures (Gok et al., 2010). Furthermore, a positive ΔH^o refers to the endothermic adsorption process of the AB113 dye on C-Fe₂O₃, which means the formation of strong

chemical bonds between the adsorbed AB113 molecules and the C-Fe₂O₃ surface (Suna et al., 2010).

Fig. 7. Effects of temperature on the AB113 dye removal efficiency by C-Fe₂O₃ (pH = 3; C-Fe₂O₃ dose = 0.6 g/L; AB113 dye concentration = 10 mg/L; and adsorption time = 0–300 min)

Table 4. Thermodynamic parameters for the adsorption process of AB113 dye on C-Fe₂O₃

T (K)	ΔG^o (kJ/mol)	ΔH^o (kJ/mol)	ΔS^o (kJ/mol·K)
288	-2.78	50.20	0.182
298	-3.92		
308	-5.25		
318	-8.51		

3.6 Regeneration study

In the present study, the recyclability of C-Fe₂O₃ was examined for six consecutive AB113 dye adsorption–desorption cycles under the following optimal conditions: pH = 3; C-Fe₂O₃ dose =

0.6 g/L; AB113 dye concentration = 10 mg/L; adsorption time = 120 min; and temperature = 25 ± 2 °C. The spent quantity of C-Fe₂O₃ was separated from the aqueous solution using a magnet, rinsed with ethanol and deionized water, dried at 75 °C for 4 h, and then reused in the next AB113 dye adsorption cycle. The results of this experiment are presented in Fig. 8, revealing that C-Fe₂O₃ can be recycled six times to adsorb AB113 dye. This figure also demonstrates that the removal efficiency of AB113 dye decreases by only 7% from the 1st to the 6th adsorption cycle. This decrease may be because of the loss of the adsorption ability of C-Fe₂O₃ due to the washing process (Osma et al., 2007). Based on these findings, it can be concluded that C-Fe₂O₃ has an excellent shelf life for use in an AB113 dye adsorption treatment system.

Fig. 8. Regeneration analysis of C-Fe₂O₃ used for six consecutive AB113 dye adsorption–desorption cycles

4. Conclusion

The current work represents the first study on the application of chitosan magnetized by Fe_2O_3 nanoparticles (C- Fe_2O_3) for removing acid blue 113 (AB113) dye from aqueous solutions. Characterization analyses using advanced techniques, including SEM/EDX, TEM, XRD, FTIR, TGA, BET, and pH_{pzc} , revealed that the adsorptive specifications of chitosan, such as the surface morphology, magnetization value, and thermal stability, were greatly improved after magnetization with Fe_2O_3 nanoparticles. In addition, the 2D and 3D AFM images evidenced the adsorption phenomenon of AB113 dye on C- Fe_2O_3 . The AB113 dye removal efficiency was 99.68% and occurred under the following conditions: $\text{pH} = 3$, C- Fe_2O_3 dose = 0.6 g/L, AB113 dye concentration = 10 mg/L, and adsorption time = 120 min. The presence of interference ions, including NaCl, NaNO_3 , Na_2CO_3 , and MgSO_4 , led to a negative effect on the AB113 dye removal efficiency but at different levels. The kinetic data obtained at different AB113 dye concentrations (10, 25, 50, and 100 mg/L) were consistent with the pseudo-second-order model. Based on the determined statistical goodness values of nonlinear fitting, the experimental data relevant to the isotherm and kinetic studies could be represented by the Freundlich and pseudo-second-order kinetic equations, respectively. In addition, the determined values of the thermodynamic parameters (positive ΔH° and ΔS° values and negative ΔG° values) indicated that the process of AB113 dye adsorption on C- Fe_2O_3 was favorable, spontaneous, and endothermic. Compared with other adsorbents used for dye removal, it was shown that C- Fe_2O_3 was an efficacious adsorbent and good alternative for treating wastewater containing AB113 dye in acidic media. Additionally, C- Fe_2O_3 could be recycled for six consecutive AB113 dye adsorption-desorption cycles with little loss in its efficacy. The key finding of this study was that

C-Fe₂O₃ has a high ability to remove AB113 dye from aqueous solutions; thus, it has potential for possible application in tertiary treatment units of dye effluents.

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Declaration of competing interest

There are no conflicts of interest to declare.

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Figures

Figure 1

SEM images of C-Fe₂O₃ before (a) and after (b) AB113 adsorption. TEM image of C-Fe₂O₃ (c). X-ray diffraction pattern of C-Fe₂O₃ (d).

Figure 2

2D and 3D AFM images of C-Fe₂O₃ before (A) and after (B) adsorption of AB113 dye.

Figure 3

TGA graph of chitosan and C-Fe₂O₃ (a). FTIR spectra of chitosan, C-Fe₂O₃ and C-Fe₂O₃ after reaction with the AB113 dye (C-Fe₂O₃/AB113) (b). Magnetization curves of the Fe₂O₃ nanoparticles (c) and C-Fe₂O₃ (d).

Figure 4

Effect study of pH and the pHpzc analysis (a), C-Fe₂O₃ dose (b), Initial Ab113 dye concentration (c), and Interference ions (d).

Figure 5

Three stages of the kinetics adsorption of AB113 onto C-Fe₂O₃.

Figure 6

Experimental and theoretical isothermal data of AB113 dye adsorption on C-Fe2O3.

Figure 7

Effects of temperature on the AB113 dye removal efficiency by C-Fe2O3 (pH = 3; C-Fe2O3 dose = 0.6 g/L; AB113 dye concentration = 10 mg/L; and adsorption time = 0–300 min)

Figure 8

Regeneration analysis of C-Fe₂O₃ used for six consecutive AB113 dye adsorption–desorption cycles

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