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# Removal of chromium(vi) from polluted wastewater by chemical modification of silica gel with 4-acetyl-3-hydroxyaniline

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In the current study, a new adsorbent that is insoluble in water and many acid solutions and has a high adsorption capacity for Cr(vi) metal ions was synthesized. In the synthesis process, 3-chloropropyltrimethoxysilane (CPTS) was first modified on a silica gel (Si) surface. Secondly, 4-acetyl-3-hydroxyaniline (AHAP) was immobilized on the modified silica gel compound (Si-CPTS). As a result of the immobilization process, a new adsorbent compound named Si-CPTS-AHAP (silica gel-3-chloropropyltrimethoxy silane 4-acetyl-3-hydroxyaniline) was obtained, which was used to separate Cr(vi) ions from aqueous solution ( $K_2Cr_2O_7$ ) and industrial wastewater. The material was characterized using scanning electron microscopy and Fourier-transform infrared spectroscopy. The amount of chromium adsorbed was detected by ultraviolet-visible spectroscopy. The adsorption was evaluated using batch methods. The effects of temperature, pH, concentration, adsorbent amount and interaction time on the adsorption of Si-CPTS-AHAP were also investigated. The adsorption of Cr(vi) ions on Si-CPTS-AHAP was investigated *via* adsorption kinetics, isotherm and thermodynamic studies. The value of the isotherm parameters and the highest adsorption yields were calculated from the Dubinin–Radushkevich, Freundlich and Langmuir isotherm equations. Thermodynamic features such as entropy ( $\Delta S$ ), enthalpy ( $\Delta H$ ) and free energy ( $\Delta G$ ) were also calculated from the experimental results. The experimental results showed that the best recoveries of Cr(vi) metal ions are under the conditions of 180 min (interaction time), 0.05 g (adsorbent amount) and 323.15 K (temperature) at pH 2. Si-CPTS-AHAP can be used for the removal of poisonous pollutants in wastewater.

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## 1. Introduction

The scarcity of clean drinking water in the world has been a problem for decades. Some poisonous organic and inorganic contaminants, especially heavy metal ions such as Co(II), Ni(II), Cd(II), Pb(II), Hg(II), Cr(III) and dyes, enter the food chain *via* wastewater. As a result, people, animals, and aquatic organisms are adversely affected.<sup>1</sup> In humans, toxic metals cause Alzheimer's, Parkinson's, blood pressure problems, stuttering, psychological disorders, hypersensitive reactions, immune system diseases, embolism and amnesia.<sup>2</sup> These problems can be solved by the removal of domestic wastes and water contaminated from industrial wastes. For a long time, several methods including reverse osmosis,<sup>3</sup> ion-exchange,<sup>4</sup> chemical precipitation,<sup>5</sup> membrane filtration<sup>6</sup> and adsorption<sup>7</sup> have been used to remove heavy metal pollution from wastewaters. Among these methods, adsorption is one of the most interesting, because it is high yielding and so is a low-cost method and is used the most widely in the treatment of contaminated water.<sup>8</sup>

Many kinds of resin such as active carbon, zeolites, silica gel, wood bark, walnut shell, and rice husk have been used as adsorbents in the removal of toxic heavy metal ions from wastewater.<sup>9</sup>

Chromium and its compounds are broadly used in a range of industrial enterprises such as the chemical industry, metallurgy, tanneries, textile dyeing, wood preserving and metal plating preparation of chromate solution. Hence, chromium pollution has frequently occurred in many industrial zones. Random leaks or insufficient care taken in the storage of waste are the main factors behind chromium contamination.<sup>10–14</sup> In nature, chromium compounds are present as Cr(vi) and Cr(III) ions.<sup>15</sup> Cr(vi) ions are highly soluble as oxyanions,<sup>16</sup> however, Cr(III) ions can be easily precipitated as Cr(OH)<sub>3</sub> and are less soluble.<sup>17</sup> Cr(vi) ions are 1000 times more toxic than Cr(III) ions.<sup>18</sup> In contrast, Cr(III) exhibits lower toxicity in many organisms<sup>19</sup> and is breathed in through the mouth, but does have carcinogenic properties.<sup>15</sup> Cr(vi) ions in wastewater polluted by industrial foundations and other sources cause serious problems and toxic and non-biodegradable substances tend to accumulate in the vital organs of humans.<sup>15</sup> For this reason, Cr(vi) ions must be removed from contaminated waters before expulsion into natural bodies of water.

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Silica gel modification has been used in metal ion pre-concentration,<sup>20</sup> exchange,<sup>21</sup> biotechnology,<sup>22</sup> catalysis,<sup>23</sup> adsorption,<sup>24</sup> clean technology, and green chemistry.<sup>25</sup> Adsorption is one of the most important methods used in the treatment of heavy metal ions from waste water.

Silica gel (with a micropore size of 6 nm, a surface area of 480–540 m<sup>2</sup> g<sup>-1</sup> and low cost) is a good solid support for adsorption because it has chemical, thermal and mechanical stability.<sup>26,27</sup> On the surface of silica gel there are two different types of functional groups; siloxane (Si–O–Si) and silanol (Si–OH).<sup>28,29</sup>

The silica gel, used as an adsorbent, has the advantages of being robust against organic solvents, thermal stability and is not capable of swelling. The bonding of a substance to the silica gel (Si) surface is easier than bonding to an organic support surface, and the silica gel (Si) is the most popular adsorbent for surface studies. Modified silica gels usually have a higher adsorption capacity than other organic and inorganic supports. Thus, the use of immobilized molecules in the removal of heavy metal ions from industrial wastewater gives better results.<sup>30,31</sup> In this study, a new surface was formed with the chemical immobilization of suitable organic groups on the silica gel surface. This new silica gel surface can be used as an adsorbent to remove Cr(vi) heavy metal ions, which show a toxic effect to nature and living things, from industrial wastewater and aqueous solutions.

There are many studies on the removal of Cr(vi) ions.<sup>7,32–34</sup> Unlike these studies, we expanded the surface area of silica gel, which is used as a support agent. This was accomplished by modification and immobilization of 3-chloropropyltrimethoxysilane (CPTS) and 4-acetyl-3-hydroxyaniline (AHAP) compounds on silica gel, respectively. The synthesized support material is insoluble in water and many acids due to the properties of the silica gel, does not swell easily due to cross-linking, is cheap and shows reproducibility in experiments. For the first time, the modification of CPTS onto silica gel and the immobilization of AHAP onto Si-CPTS were performed. Using the newly synthesized Si-CPTS-AHAP adsorbent in the removal of Cr(vi) ions in wastewater treatment systems could potentially provide both low-cost and high-efficiency heavy metal removal. Therefore, the current study has great potential to contribute to the world economy, water and environmental cleanliness. In addition, the adsorption method is cheap, easy to apply and has a short test period.

In this work, the chelating molecule 4-acetyl-3-hydroxyaniline (AHAP) was attached onto a silica gel surface through an immobilization reaction and characterized using Fourier-transform infrared spectroscopy (FTIR) and scanning electron microscope (SEM). This new adsorbent was utilized to remove Cr(vi) metal ions in industrial wastewater taken from a valve factory in the organized industrial zone in Konya, Turkey. In addition, this new adsorbent was applied to remove aqueous solutions containing trace amounts of Cr(vi) metal ions.

## 2. Materials and methods

### 2.1. Materials

Silica gel 60 (diameter of 0.063–0.200 mm, micropore size of 6 nm, high surface area of 480–540 m<sup>2</sup> g<sup>-1</sup>), used as the support

material of the new adsorbent material, was purchased from Merck (Darmstadt, Germany, www.merck.de). 3-Chloropropyltrimethoxy silane (CPTS), 4-acetyl-3-hydroxyaniline (AHAP) and potassium dichromate (K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>) were purchased from Sigma. Toluene (99.8%) was purchased from Sigma-Aldrich. A standard Cr(vi) metal ion solution was freshly prepared daily. Adjustment of the pH of the industrial wastewater and aqueous solutions was carried out using dilute NaOH and HNO<sub>3</sub> solutions. All the chemicals used in the study were used without further purification unless otherwise stated. Industrial wastewater sample (82.21 mg L<sup>-1</sup> Cr(vi), pH = 2.0, temperature: 318.15 K, current density: 12–20 A dm<sup>-2</sup> and H<sub>2</sub>SO<sub>4</sub>: 8.05 mg L<sup>-1</sup>) containing Cr(vi) ions was obtained from a valve factory in the organized industrial zone in Konya, Turkey.

### 2.2. Instruments

SEM images of activated silica gel (Si), 3-chloropropyl-bonded silica gel (Si-CPTS) and the Si-CPTS-AHAP adsorbent were obtained using a Zeiss LS-10 device equipped with an Inca Energy 350 X-Max (Oxford Instruments) spectrometer.

FT-IR spectra of the activated silica gel (Si), 3-chloropropyl bonded silica gel (Si-CPTS) and the Si-CPTS-AHAP adsorbent compounds were scanned using a PerkinElmer 100 FTIR spectrometer (ATR; 21 °C temperature and 1 atm pressure) in the range of 650–4000 cm<sup>-1</sup>. The pH values of Cr(vi) metal ion solutions were measured using a Jenway 3010 digital pH meter. Amounts of metallic Cr(vi) cations remaining in solution were determined by ultraviolet-visible (UV) spectroscopy (Optizen brand 3220 model UV-vis).

### 2.3. Functionalization of silica gel and immobilization of AHAP

The preparation of the Si-CPTS-AHAP adsorbent was carried out in three stages. In the first stage, silica gel (10 g) was refluxed with 6 M hydrochloric acid (100 mL) for 8 h to prepare conventional activated silica to remove metal ions and maximize the number of silanol groups on the surface.<sup>35,36</sup> The resulting suspension was filtered under vacuum and the resulting activated silica gel was washed 3 times with distilled water and twice with ethanol.

In the second stage, the activated silica gel (5 g) was treated with 5 mL of 3-chloropropyltrimethoxysilane (CPTS) in 50 mL of anhydrous toluene<sup>37</sup> at 96 °C for 12 h.<sup>38,39</sup> The resulting Si-CPTS compound was filtered through the mixture and washed with toluene, ethanol and diethyl ether and dried at 60 °C for 6 h.<sup>38</sup> The possible structure of the 3-chloropropyl-bonded silica gel (Si-CPTS) is shown in Fig. 1.

In the third stage, the dried 3-chloropropyl-bonded silica gel (5 g) was mixed with 33 mL of 25% 4-acetyl-3-hydroxyaniline (AHAP) solution in 50 mL of anhydrous toluene<sup>37</sup> and was refluxed under nitrogen for 24 h at 90 °C.<sup>38,39</sup> The crude product obtained by the filtration of the suspension was washed with water and ether and then dried at 40 °C for 72 h under vacuum to afford Si-CPTS-AHAP.<sup>28–30</sup> The possible structure of the synthesized Si-CPTS-AHAP adsorbent is shown in Fig. 1.





Fig. 1 Possible structure of the synthesized Si-CPTS-AHAP adsorbent.

#### 2.4. Adsorption studies

Adsorption experiments were conducted in batch mode, using 20 mL centrifuge tubes through the contacting of 10 mL of a  $1 \times 10^{-4}$  mol L<sup>-1</sup> metal ion solution and wastewater with 20 mg of Si-CPTS-AHAP adsorbent at 200 rpm, for definite time intervals in a temperature controlled shaker incubator.

After the experimental studies and measurements, the amount of Cr(vi) ions adsorbed onto the Si-CPTS-AHAP adsorbent was calculated using eqn (1):

$$q_e = (C_0 - C_e) \times V/m \quad (1)$$

where  $q_e$  (adsorption capacity) is the amount of adsorbed Cr(vi) ions at equilibrium (mg g<sup>-1</sup>),  $C_e$  and  $C_0$  (mg L<sup>-1</sup>) are the equilibrium and initial concentrations of Cr(vi) ions in an aqueous solution,  $V$  (L) is the volume of the aqueous chromium solution and  $m$  (g) is the mass of the Si-CPTS-AHAP adsorbent.<sup>40,41</sup>

#### 2.5. pH studies

pH studies were carried out at various pH values (2.0, 3.0, 4.0, 5.0, 6.0, 7.0). 0.05 g of the Si-CPTS-AHAP adsorbent at 200 rpm was mixed with a 10 mL aqueous solution containing  $1 \times 10^{-4}$  mol L<sup>-1</sup> K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> and wastewater. The obtained solutions were shaken for 180 min at 323.15 K.<sup>42</sup>

#### 2.6. Concentration studies

The effect of Cr(vi) concentration on adsorption was investigated using a batch method for potassium dichromate (K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>) solution and industrial wastewater at  $25 \pm 1$  °C. For these adsorption measurements, 0.05 g of the Si-CPTS-AHAP adsorbent was suspended in 10 mL of aqueous solution containing different amounts of potassium dichromate (K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>) solution and industrial wastewater. These suspensions were shaken in various Cr(vi) concentrations (8, 12, 20, 40 mg L<sup>-1</sup>) in a shaker thermostat for 180 min at 323.15 K.<sup>43</sup> After adsorption equilibrium was established, the amounts of Cr(vi) metal ion residues in solution were detected by ultraviolet-visible (UV) spectroscopy.

#### 2.7. Temperature studies

0.05 g of the Si-CPTS-AHAP adsorbent and 10 mL of a  $0.0001$  mol L<sup>-1</sup> Cr(vi) metal ion solution at an optimum pH of 2.0 were added into a series of conical flasks. All flasks containing the Si-CPTS-AHAP adsorbent and Cr(vi) metal solution were kept in a water bath equipped with a thermostat.

All suspensions in the flasks were stirred for 150 min at ( $20 \pm 0.01$ ), ( $30 \pm 0.01$ ), ( $40 \pm 0.01$ ) and ( $55 \pm 0.01$ ) °C, respectively. The amounts of the adsorbed Cr(vi) metal ion residues in



solution were detected by UV spectroscopy and calculated from the change in Cr(vi) metal concentrations in aqueous solution.

## 2.8. Contact time studies

In this study, the solution concentration, pH value and temperature parameters were kept constant, 0.05 g of the Si-CPTS-AHAP adsorbent against 10 mL of  $1 \times 10^{-4}$  mol L<sup>-1</sup> stock Cr(vi) metal solutions were activated separately at different (30, 60, 90, 120, 150 and 180 minutes) contact times. In addition, this method was applied to an industrial waste solution. The amount of adsorbed Cr(vi) metal ion residues in solution was detected by UV spectroscopy and the maximum contact time was determined from the obtained data.

## 2.9. Studies of the effect of the adsorbent quantity on adsorption

In order to investigate the effect of the amount of adsorbent, the relationship between the amount of Cr(vi) metal ions adsorbed on the adsorbent surface using different adsorbent amounts (0.01, 0.015, 0.025, 0.05 and 0.075 g) was investigated.

## 2.10. Adsorption isotherm models

Adsorption equilibrium is an essential feature in adsorption studies. Considering this, numerous investigations were conducted to determine the amount of metal ions adsorbed under certain conditions. To optimize the design of an adsorption system to remove heavy metal ions from aqueous solutions, it is important to find the most appropriate correlation for the balance curve.<sup>47</sup> Adsorption isotherms are the most appropriate way to evaluate the adsorption process. In this study, Langmuir, Freundlich and Dubinin–Radushkevich (D–R) adsorption isotherm models were used to process the experimental equilibrium data to verify which model gives the best fit. Adsorption

isotherms can be investigated by changing the initial concentrations of pollutant chromium, keeping the other process parameters constant in terms of pH, temperature, processing time and agitation speed.<sup>48</sup> In this study, the models used in Table 1 are shown with mathematical expressions.

## 2.11. Adsorption thermodynamics

In the adsorption studies, thermodynamic parameters such as free energy change ( $\Delta G$ ), entropy change ( $\Delta S$ ) and enthalpy change ( $\Delta H$ ) are very important. These parameters provide information about how the adsorption process takes place. The effect of temperature on the adsorption of the newly synthesized Si-CPTS-AHAP adsorbent was investigated at temperatures between 20 and 50 °C under optimized conditions of pH, contact time and concentration values, for each of the Cr(vi) ion solutions. The values of the thermodynamic parameters, including the Gibbs free energy ( $\Delta G$ ), enthalpy ( $\Delta H$ ) and entropy ( $\Delta S$ ) were calculated from the following equations:<sup>49,50</sup>

$$\Delta G = -RT \ln K_c \quad (6)$$

where  $R$  is the universal gas constant (8.314 J mol<sup>-1</sup> K<sup>-1</sup>) and  $K_c$  is the ratio between the  $C_e$  adsorbent and  $C_e$  solution

$$\Delta G = \Delta H - T\Delta S \quad (7)$$

$$\ln K_c = \Delta S/R - \Delta H^{\circ}/RT \quad (8)$$

# 3. Results and discussion

## 3.1. Characterization

**3.1.1. FT-IR analysis.** The FT-IR spectra of the activated silica gel (Si), 3-chloropropyl bonded silica gel (Si-CPTS) and Si-

Table 1 Mathematical expressions used in the adsorption isotherm models

Model	Linear equation	Plot	Nomenclature	Ref.
Langmuir	$\frac{C_e}{q_e} = \frac{C_e}{q_o} + \frac{1}{q_o K_L}$	$C_e/q_e$ vs. $C_e$	$q_o$ is the maximum surface density at monolayer coverage, $C_e$ is the equilibrium ion concentration in the solution, $q_e$ is the amount of solute adsorbed on the surface of the adsorbent, $K_L$ is the energy constant for Langmuir	44
Freundlich	$\ln q_e = \ln K_F + \frac{\ln C_e}{n}$	$\ln q_e$ vs. $\ln C_e$	$q_e$ is the equilibrium solute concentration on the adsorbent, $n$ is a constant reflecting the adsorption intensity, $C_e$ is the equilibrium solute concentration on the adsorbent, $K_F$ is the Freundlich constant	45
Dubinin–Radushkevich (D–R)	$\ln q_e = q_m - k\varepsilon^2$ $E = (2k)^{-1/2}$	$\ln q_e$ vs. $\varepsilon^2$	$q_e$ is the amount of solute adsorbed per unit weight of adsorbent, $\varepsilon$ (Polanyi potential) is $[RT \ln(1 + (1/C))]$ , $q_m$ is the adsorption capacity (mol g <sup>-1</sup> ), $k$ is a constant related to the adsorption energy, $E$ is the value of the mean free energy	46



CPTS-AHAP adsorbent were recorded by a PerkinElmer 100 spectrometer using the potassium bromide (KBr) disc method and are shown in Fig. 2.

The FT-IR spectrum of the activated silica gel shows a band at  $1058\text{ cm}^{-1}$  due to the asymmetric stretching of Si–O–Si and corresponding symmetric stretching was observed at  $795\text{ cm}^{-1}$ .<sup>51</sup> The characteristic peaks of the Si–O–H groups of the activated silica gel were observed at  $967\text{ cm}^{-1}$  (ref. 52) and also strong and broad peaks at around  $3465\text{ cm}^{-1}$  can be attributed to the hydroxyl group (O–H) stretching vibrations.<sup>53</sup> The peak observed at  $1637\text{ cm}^{-1}$  demonstrates the presence of physically adsorbed water in the silica gel samples.<sup>54</sup>

In the FTIR spectrum of Si-CPTS, C–H peaks<sup>51</sup> were observed at approximately  $2977\text{ cm}^{-1}$  and C–Cl peaks<sup>55,56</sup> were observed at around  $684\text{ cm}^{-1}$ . The OH peak observed for active Si at  $3465\text{ cm}^{-1}$  shifted to  $3487$  from  $3465\text{ cm}^{-1}$  in the Si-CPTS compound. The OH peak also shifted from  $3487$  to  $3485\text{ cm}^{-1}$  for the Si-CPTS-AHAP.<sup>54</sup>

For the Si-CPTS compound, the observed Si–O–Si tensile vibration was  $1049\text{ cm}^{-1}$ , whereas in the Si-CPTS-AHAP adsorbent this stretching vibration was observed to be  $1039\text{ cm}^{-1}$ . C–H asymmetric and symmetrical stretching vibrations were seen in the range of  $2923$ – $2860\text{ cm}^{-1}$  in the spectrum of the Si-CPTS-AHAP adsorbent, respectively.<sup>55</sup>

The FT-IR spectrum of Si-CPTS-AHAP shows the presence of a C=C peak at  $1431\text{ cm}^{-1}$  and the appearance of a new band at  $1479\text{ cm}^{-1}$ , both characteristic of the vibrations of the C=C bonds of aromatic rings and also exhibits new peaks such as C–N at  $1204\text{ cm}^{-1}$  and C–O–H at  $1150\text{ cm}^{-1}$ . These indicate that AHAP is fixed to the surface of the silica gel.

**3.1.2. Elemental composition analysis.** Elemental composition analyses of the activated silica gel ( $\text{SiO}_2$ ), 3-chloropropyl-bonded silica gel (Si-CPTS) and Si-CPTS-AHA were carried out to determine the percentages of nitrogen (N), carbon (C) and hydrogen (H) in the samples,<sup>57</sup> the results of which are shown in Table 2.

Percentages of 34.25, 2.31 and 5.04% for carbon (C), hydrogen (H) and nitrogen (N), respectively, confirmed the surface modification of silica with AHAP.

**3.1.3. SEM images.** Adsorption takes place on a surface. Therefore, the speed and degree of adsorption depend on the surface functional groups of the adsorbent, the pore size and the surface area. SEM is one of the most widely used surface identification methods,<sup>58</sup> therefore; SEM micrographs were acquired (by applying a 50 kV electron acceleration voltage) to observe the surface morphologies of the activated silica gel (Si) and Si-CPTS-AHAP adsorbent. An SEM micrograph of activated silica gel (Si) is shown in Fig. 3(a). As can be seen from Fig. 3(a),

Table 2 Percentages (%) of nitrogen (N), carbon (C) and hydrogen (H) in the activated silica gel (Si), 3-chloropropyl-bonded silica gel (Si-CPTS) and Si-CPTS-AHAP adsorbent

Surface	Nitrogen (%)	Carbon (%)	Hydrogen (%)
$\text{SiO}_2$	0	0	0
Si-CPTS	0	7.22	0.82
Si-CPTS-AHAP	5.04	34.25	2.31

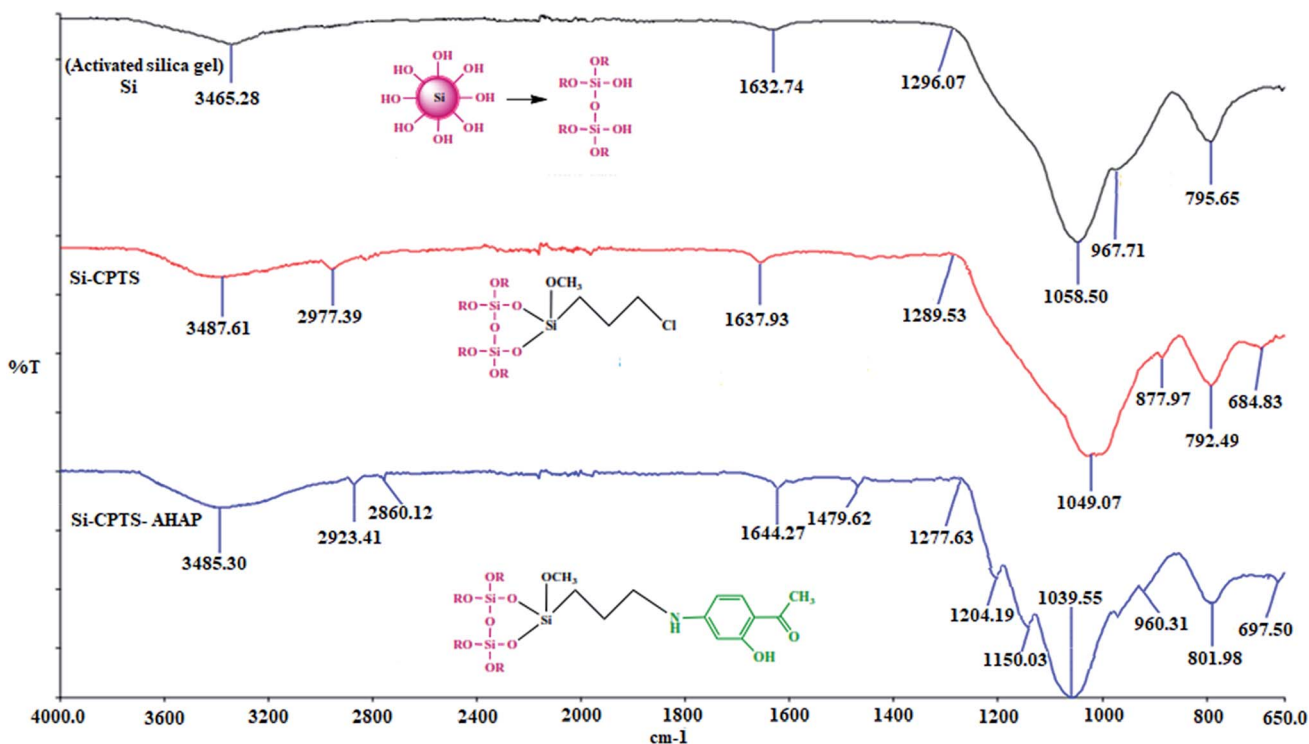


Fig. 2 FTIR spectra of the activated silica gel (Si), 3-chloropropyl-bonded silica gel (Si-CPTS) and the Si-CPTS-AHAP adsorbent.



the active silica gel surface has a very smooth morphology. An SEM micrograph of the Si-CPTS adsorbent is shown in Fig. 3(b), which exhibits an irregular morphology covered by CPTS. The structure of the active Si surface changed. However, no clear change was observed as much as the change observed in the Si-CPTS-AHAP adsorbent, as shown in Fig. 3(c). The SEM micrograph of the Si-CPTS-AHAP adsorbent obtained after the immobilization of AHAP on the active silica gel surface is shown in Fig. 3(c). It shows an irregular morphology covered by impurities, *i.e.* AHAP and CPTS. The presence of attached particles of AHAP and CPTS onto the surface of the activated silica gel verified the immobilization.



Fig. 3 SEM images of the (a) activated silica gel (Si), (b) Si-CPTS adsorbent and (c) Si-CPTS-AHAP adsorbent.

### 3.2. Adsorption studies

**3.2.1. Effect of pH on adsorption.** It is well-known that pH is one of the most significant factors that effects the adsorption process.<sup>59</sup> The effects of pH on the adsorption of chromium ions in aqueous solution and wastewater were investigated between pH 2.0 and 7.0 using Si-CPTS-AHAP as the adsorbent. Experiments were not carried out under extreme alkaline conditions, as at higher pH values precipitation of chromium hydroxide occurs. Potassium dichromate ( $K_2Cr_2O_7$ ) was used to prepare a standard solution of Cr(vi). In acid solutions, Cr(vi) becomes negatively charged ( $HCrO_4^-$  and  $Cr_2O_7^{2-}$ ). A graphical representation of the effect of pH is shown in Fig. 4(a). As seen in Fig. 4(a), the adsorption capability of the Si-CPTS-AHAP adsorbent decreases with increasing pH. Experimental results show that the adsorption of Cr(vi) ions in aqueous solution and wastewater is reduced at high pH.<sup>60</sup> The adsorption of Cr(vi) ions in aqueous solution and wastewater reached a maximum in pH 2.<sup>61</sup>

It can be seen that the adsorption mechanism of Cr(vi) ions on the adsorbent (Si-CPTS-AHAP) is based on chelation and the donor nitrogens and OH groups are coordinated to the chemisorbed heavy metal ions on the surface. It is possible to say that  $HCr_2O_7^-$  and  $Cr_2O_7^{2-}$  anions are electrostatically attracted by the acetophenone (positive charge effect) groups on the Si-



Fig. 4 (a) The effect of pH on the sorption of Cr(vi) in aqueous solution and wastewater. (b) Si-CPTS-AHAP Cr(vi) metal ion combination and a schematic of the electrostatic attraction of acetophenone to  $HCr_2O_7^-$  and  $Cr_2O_7^{2-}$  anions.



CPTS-AHAP adsorbent. A schematic diagram of the Si-CPTS-AHAP Cr(vi) metal ion complex is shown in Fig. 4(b). The data obtained were consistent with the literature.<sup>48,59,60</sup>

**3.2.2. Effect of concentration on the adsorption.** Adsorption takes place on a surface. Therefore, the size of the surface area and the concentration of the substance to be attached to the surface are important. As the surface area of the obtained adsorbent is large, the adsorption percentage increases with the increasing Cr(vi) ion concentration that is to be adsorbed, as shown in Fig. 5. As can be seen from the graph,<sup>62</sup> the adsorption increases with an increase in the concentration of the metal ions. After a while, the adsorption reached steady state values. The data obtained were consistent with the literature.<sup>48,61,63</sup>

**3.2.3. Effects of temperature on the adsorption.** Temperature is a parameter that affects adsorption. If adsorption occurs endothermically, the temperature increases the amount of adsorption. The situation is reversed for exothermic reactions.

In the experimental study, a batch adsorption method was used to see the influence of varying the temperature ( $20\text{--}50 \pm 0.001\text{ }^\circ\text{C}$ ) on the adsorption of Cr(vi) ions onto<sup>60</sup> Si-CPTS-AHAP. The obtained results can be seen in Fig. 6. These results demonstrate that adsorption takes place *via* an endothermic reaction and increases with temperature. The data obtained were consistent with the literature.<sup>48,59–61</sup>

**3.2.4. Effects of contact time on the adsorption.** An adsorbent that has hydrophilic groups is defined as the most effective adsorbent. In our study, the reason for the rapid adsorption kinetics may be related to the hydroxy groups in the silica gel and nitrogen of the imine bonds<sup>64</sup> in the adsorbent (C=N). The experimental data are shown in Fig. 7. As can be seen from Fig. 7, the Cr(vi) adsorption of the Si-CPTS-AHAP adsorbent shows a very rapid increase with the adsorption time at first<sup>62</sup> and after a certain contact time, steady state was reached. The data obtained are consistent with the literature.<sup>48,59–62</sup>

**3.2.5. Effect of the adsorbent quantity on the adsorption.** Several steps were observed (Fig. 8) from the results of experiments on varying the Si-CPTS-AHAP adsorbent quantities (0.01, 0.015, 0.025, 0.05 and 0.075 g). The study was carried out



Fig. 6 The effect of temperature on the adsorption of Cr(vi) from aqueous solution and wastewater.

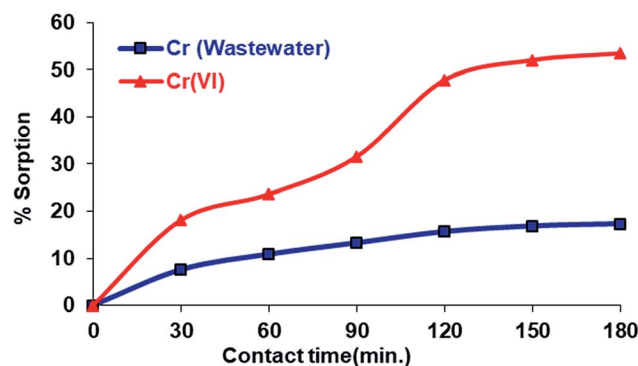


Fig. 7 The effect of the contact time on the adsorption of Cr(vi) from aqueous solution and wastewater.

separately for Cr(vi) ions in aqueous solution and industrial wastewater. As can be seen from Fig. 8, the change in the amount of adsorption depends on the increase in the amount of the Si-CPTS-AHAP adsorbent. As shown in Fig. 8, when the amount of Si-CPTS-AHAP adsorbent used was increased, the amount of Cr(vi) ions adsorbed increased.

As a result, the maximum quantity of the Si-CPTS-AHAP adsorbent needed for adsorbing Cr(vi) ions from aqueous



Fig. 5 The adsorption effect depends on the concentrations of Cr(vi) in aqueous solution and wastewater.



Fig. 8 The effect of the amount of sorbent on the sorption of Cr(vi) from aqueous solution and wastewater.



solution and industrial wastewater was found to be 0.05 g. Using 0.05 g of adsorbent, all active areas on the face of the material were occupied and an increase in the amount of adsorbent did not result in a higher percentage of sorption. The sorption capacity of Si-CPTS-AHAP was found to be Cr(VI): 0.32 and 3.22 for aqueous solution ( $K_2Cr_2O_7$ ) and industrial wastewater, respectively (eqn (1)).<sup>65</sup> The data obtained were consistent with the literature.<sup>48,59–63</sup>

### 3.3. Isotherm studies

Adsorption isotherms reflect the specific relationship between the adsorbate concentration and degree of deposition on the adsorbent surface at constant temperature.<sup>66,67</sup> Equilibrium studies can be carried out to determine the capacity of an adsorbent and define the adsorption isotherm with constants that expressing the values, surface properties and affinity of the adsorbents.<sup>68</sup> In addition, the adsorption isotherm data was determined using the three most widely used models: the Dubinin–Radushkevich (D–R), Langmuir, and Freundlich models. The parameters obtained from these three models provide important information about the adsorption mechanism and surface properties, the adsorption free energy and the affinity of the adsorbent.

**3.3.1. Langmuir adsorption isotherm.** Langmuir adsorption isotherms have been successfully applied to many real adsorption processes<sup>69</sup> and the Langmuir (1918) theory relies on the following assumptions:<sup>70</sup>

- The adsorbate forms a monolayer on the adsorbent surface.
- The active sites are the same.
- The presence of interactions between the molecules that are being absorbed.
- The adsorbent has a limited adsorption capacity ( $q_{max}$ ).

Briefly, the Langmuir adsorption isotherm shows the equilibrium distribution of the metal ions between the solid and liquid phases.<sup>65</sup> The Langmuir isotherm equation is shown in Table 1. Fig. 9 presents the linear plots of the Langmuir isotherm equation. The values of the parameters in the Langmuir adsorption isotherm equation are summarized in Table 3, as analyzed from the areas shown in Fig. 9.

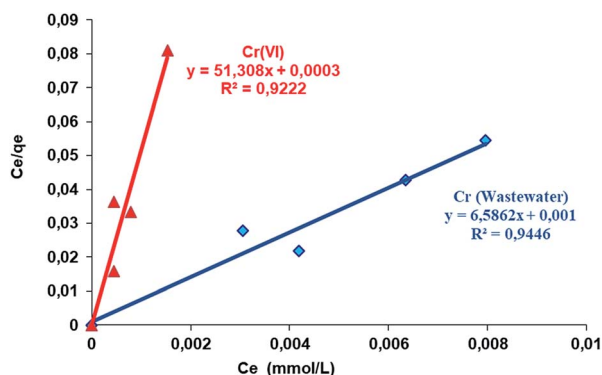


Fig. 9 Langmuir isotherms of the removal of Cr(VI) from aqueous solution and wastewater by Si-CPTS-AHAP.

Table 3 Parameters of the Langmuir isotherms for Cr(VI) adsorption onto the Si-CPTS-AHAP adsorbent

Parameters of the Langmuir isotherms					
Metal	pH	T (K)	$q_0$ (L mol <sup>-1</sup> )	$K_L$ (mol <sup>2</sup> (kJ <sup>2</sup> ) <sup>-1</sup> )	$R^2$
Cr(VI)	2	323.15	0.174	92.800	0.922
Cr (wastewater)	2	323.15	0.060	144.940	0.945

**3.3.2. Freundlich adsorption isotherm.** The Freundlich adsorption isotherm model is an empirical formula<sup>71</sup> and is usually applied to multi-layer adsorption systems. The Freundlich adsorption isotherm equation is explained in Table 1. The Freundlich adsorption isotherm equation in Table 1 provides a  $\ln q_e - \ln C_e$  graph, a straight line (Fig. 10), and the  $K_F$  and  $n$  values can be calculated from the intercept and slope of this straight line.<sup>72</sup> The values of the parameters in the Freundlich adsorption isotherm equation are summarized in Table 4, as analyzed from the areas shown in Fig. 10.

The  $1/n$  values of the Si-CPTS-AHAP adsorbent are shown in Table 3, where a value of  $<1$  is an indication of high adsorption density.<sup>73</sup> The  $K_F$  values as shown in Table 3 show that the newly synthesized Si-CPTS-AHAP adsorbent shows high adsorption of Cr(VI) metal ions from aqueous solution and industrial wastewater (0.50 and 0.10 mol L<sup>-1</sup>).<sup>74</sup> Values of  $n > 1$  are indicative of suitable adsorption conditions.<sup>75</sup>

**3.3.3. Dubinin–Radushkevich (D–R) adsorption isotherm.** The Dubinin–Radushkevich (D–R) isotherm is useful for

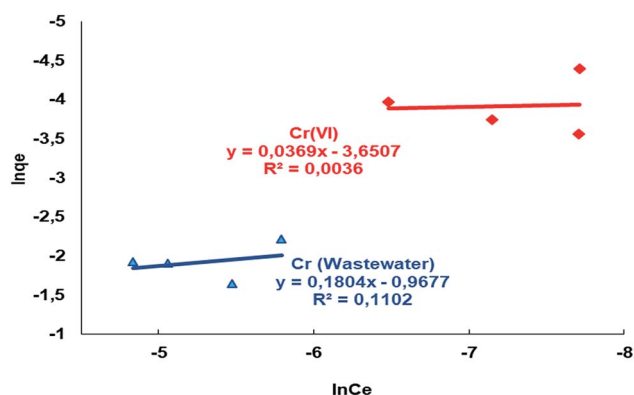


Fig. 10 Freundlich isotherms for the adsorption of Cr(VI) from aqueous solution and wastewater by the Si-CPTS-AHAP adsorbent.

Table 4 Parameters of the Freundlich isotherms for Cr(VI) adsorption onto the Si-CPTS-AHAP adsorbent

Parameters of the Freundlich isotherms					
Metal	pH	T (K)	$1/n$	$K_F$	$R^2$
Cr(VI)	2	323.15	0.0369	0.50	0.004
Cr (wastewater)	2	323.15	0.1804	0.10	0.110



estimating the characteristic porosity of the adsorbent and deducing the apparent energy of adsorption.<sup>76</sup> The data was fitted to the D–R isotherm model (calculated using the formulae mentioned in Table 1). From the data obtained using the D–R adsorption isotherm equation in Table 1, a graph of  $\epsilon^2 - \ln q_e$  was plotted (Fig. 11). The  $q_m$  value was obtained from the intercept of the graph and the  $k$  value was obtained from the slope.<sup>77</sup>

The mean free energy ( $E$ ) values were calculated using the free energy ( $E$ ) equation in Table 1, while the mean free energy ( $E$ ), calculated using the D–R adsorption isotherm, is reported in Table 5.

The  $E$  values were found to be 50.00 and 20.41 kJ mol<sup>-1</sup> for Cr(vi) metal adsorption from aqueous solution and wastewater, respectively, as shown in Table 5. The adsorption of Cr(vi) from aqueous solution and industrial wastewater took place *via* chemisorption, where an energy of between 8 and 16 kJ mol<sup>-1</sup> is sufficient for carrying out the chemical adsorption.<sup>78</sup> The values of  $n$  were  $n > 1$  at all temperatures, indicated the high affinity of Cr(vi) ions for the Si-CPTS-AHAP adsorbent.

### 3.4. Thermodynamic studies

The values of the thermodynamic parameters, including the Gibbs free energy ( $\Delta G$ ), enthalpy ( $\Delta H$ ) and entropy ( $\Delta S$ ), were calculated using eqn (6)–(8) and the value of the logarithmic plot of the distribution coefficient  $K_D$  against  $1/T$  is shown in Fig. 12. The  $\Delta G$ ,  $\Delta H$  and  $\Delta S$  values are presented in Table 6.



Fig. 11 The D–R isotherms for the adsorption of Cr(vi) from aqueous solution and wastewater by the Si-CPTS-AHAP adsorbent.

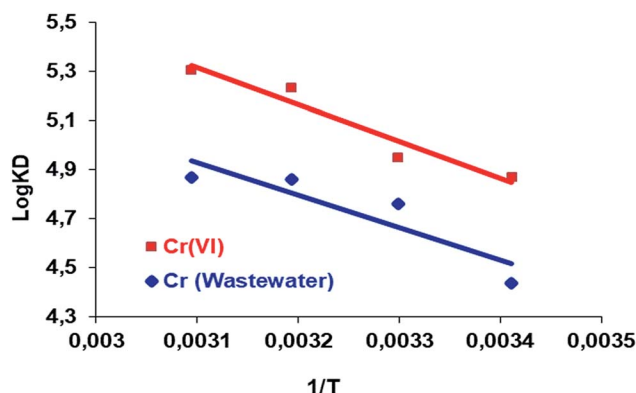


Fig. 12 Plots of  $\log K_D$  versus  $1/T$  for the removal of Cr(vi) from aqueous solution and wastewater by the Si-CPTS-AHAP adsorbent.

As shown in Table 6, the positive presence of the entropy ( $\Delta S$ ) values indicates that the adsorption reaction is endothermic and the negative values of the Gibbs free energy ( $\Delta G$ ) indicate that adsorption onto the adsorbent is possible and is a spontaneous process.<sup>28</sup> The Si-CPTS-AHAP adsorbent has complex forming ability and therefore complex formation must be the predominant mechanism. The positive enthalpy ( $\Delta H$ ) values also support this notion.

In addition, positive  $\Delta S$  values indicate the increased randomness at the solid/liquid interface during the sorption of Cr(vi) on the selected Si-CPTS-AHAP adsorbent,<sup>79</sup> which also indicates that ion exchange reactions occur. Metal ions coordinate with the water molecules in solution *via* covalent bonds and bind to the adsorbent. Due to the release of water molecules, the degree of release increases the magnitude of  $\Delta H$  depending on the adsorption energy, indicating the type of binding mechanism, *i.e.* physical or chemical adsorption.<sup>65</sup> This process is very rapid and usually reversible due to the small amount of energy required in the physical adsorption. The  $\Delta H$  values for Cr(vi) adsorbed from aqueous solution and wastewater were calculated as 28.75 and 25.59 kJ mol<sup>-1</sup> in the temperature range of between 20 and 50 °C, respectively. These values are lower than 40 kJ mol<sup>-1</sup>, indicating the weak interactions of the Cr(vi) metal ions with the Si-CPTS-AHAP adsorbent surface in the specified temperature range.<sup>28,77</sup>

### 3.5. Mechanism

This adsorption mechanism can be explained by an ion exchange process. However, it is thought that the AHAP functional group undergo a chelating effect with the Si-CPTS-AHAP adsorbent during the adsorption process. It is possible to

Table 5 Parameters of the D–R isotherms for Cr(vi) adsorption onto the Si-CPTS-AHAP adsorbent

Parameters of the Dubinin–Radushkevish (D–R) isotherms					
Metal	pH	$T$ (K)	$k$ (mol <sup>2</sup> K <sup>-1</sup> J <sup>-1</sup> )	$q_m$ (mol g <sup>-1</sup> )	$E$ (kJ mol <sup>-1</sup> )
Cr(vi)	2	323.15	0.0002	0.133	50.00
Cr (wastewater)	2	323.15	0.0012	0.061	20.41



Table 6 Thermodynamic parameters for the sorption of Cr(vi) from aqueous solution and wastewater by Si-CPTS-AHAP

Metal	$\Delta H$ (kJ mol <sup>-1</sup> )	$\Delta S$ (J K <sup>-1</sup> mol <sup>-1</sup> )	$-\Delta G$ (kJ mol <sup>-1</sup> )			
			298 K	303 K	313 K	323 K
Cr(vi)	28.75	142.19	12.93	13.19	12.87	13.03
Cr (wastewater)	25.59	124.15	10.81	11.99	12.65	13.07

Table 7 Maximum sorption capacity of different silica-based materials used in the removal of Cr(vi) ions

Adsorbent	pH	Sorption capacity (mg g <sup>-1</sup> )	Reference
PANI/SiO <sub>2</sub> composite	4.2	63.41	81
Cyphos [A336] [C272 functionalized silica (SG-5)]	0–2	15.29	82
Cyphos IL 104 functionalized silica (SG-2)	0–2	19.31	82
Silica	4	1.08	83
Silica gel chitosan composite	4	3.5	83
La(III) encapsulated silica gel chitosan composite	4	240.0	83
DMAEMA-g-silica	2.5–5	68.0	84
DMAEMA-g-silica	1.5–5.4	51.9	85
Aniline formaldehyde condensate coated on silica gel	3	17.0	86
PEI-silica nanoparticles	4	183.7	87
Zeolite NaX	4	6.414	88
Si-CPTS-AHAP	2	65.0	Present study

interpret that the hydroxyl groups in the Si-CPTS-AHAP adsorbent and the surface coordination of the donor nitrogens are responsible for the chemisorption of Cr(vi) ions.

The sorption mechanism of Cr(vi) on Si-CPTS-AHAP can be explained *via* the binding of oxygens linked to benzene groups. Because Cr(vi) is a strong hard acid it can easily coordinate with hard bases such as oxygen. Both the carbonyl and oxygen on the benzene groups on Si-CPTS-AHAP, and oxygens which did not react with CPTS on the activated silica gel can be used in the coordination.

## 4. Conclusions

The Si-CPTS-AHAP adsorbent was successfully synthesized and the FT-IR and SEM results confirmed the synthesis of Si-CPTS-AHAP adsorbent. Five different parameters were used and based on these results, the best chromium removal was carried out at pH = 2.0, with 0.05 g of adsorbent, at a temperature of 323.15 °C, at a concentration of 20 mg L<sup>-1</sup> for a contact time of 180 min.<sup>80</sup> The Langmuir, Freundlich and D-R adsorption isotherm models were calculated and the adsorption was found to fit to the Langmuir adsorption isotherm model. The mean adsorption energies for immobilized silica gel (Si-CPTS-AHAP) were found to be 50.00 and 20.41 kJ mol<sup>-1</sup> for Cr(vi) metal ions from aqueous solution and industrial wastewater, respectively.

Thermodynamic parameters calculated for the adsorption of Cr(vi) metal ions to the Si-CPTS-AHAP adsorbent surface are endothermic and spontaneous. The values of the free energy change ( $\Delta G$ ) are negative in the range of 20–50 °C for spontaneous

processes. Free energy change ( $\Delta G$ ) values decrease as the temperature increases since the adsorption reaction is endothermic. The adsorption of heavy metal ions on the Si-CPTS-AHAP adsorbent is higher at higher temperatures, meaning that it has a high adsorption capacity to remove Cr(vi) ions from wastewater.

The novelty of this study is the synthesis of the Si-CPTS in our laboratory and the fixing of organic AHAP to the Si-CPTS surface. Since the new material obtained is not soluble in water, this aids in the separation and purification methods and will be helpful for use in other studies.

This work is important in ensuring the effective use of Si-CPTS-AHAP in the removal of Cr(vi) ions in effluent, as well as providing the precise characterization of the synthesized compound. If organizations with heavy metal waste used Si-CPTS-AHAP in their waste water treatment systems, they would experience both low-cost and high-efficiency heavy metal removal. Thus, this study contributes to the world economy, water and environmental cleanliness.

The Si-CPTS-AHAP adsorbent was compared with the silica-based adsorbents reported in the literature. Table 7 shows the maximum adsorption capacity and pH values for the removal of Cr(vi) ions by Si-CPTS-AHAP compared with various silica-based adsorbents.<sup>81</sup> It can be seen from the table that the adsorption capacity of Si-CPTS-AHAP in the present study is comparable to the capacity of other adsorbents.<sup>81</sup>

## Conflicts of interest

There are no conflicts to declare.



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## References

- J. M. Gong, T. Liu, X. Q. Wang, X. L. Hu and L. Z. Zhang, *Environ. Sci. Technol.*, 2011, **45**, 6181–6187.
- M. F. Yardim, T. Budinova, E. Ekinici, N. Petrov, M. Razvigorova and V. Minkova, *Chemosphere*, 2003, **52**, 835–841.
- B. C. Ricci, C. D. Ferreira, A. O. Aguiar and M. Amaral, *Sep. Purif. Technol.*, 2015, **154**, 11–21.
- G. Sharma, B. Thakur, M. Naushad, H. Ala'a, A. Kumar, M. Sillanpaa and G. T. Mola, *Mater. Chem. Phys.*, 2017, **193**, 129–139.
- F. Fu, L. Xie, B. Tang, Q. Wang and S. Jiang, *Chem. Eng. J.*, 2012, **189–190**, 283–287.
- J. Song, H. Oh, H. Kong and J. Jang, *J. Hazard. Mater.*, 2011, **187**, 311–317.
- M. Naushad, S. Vasudevan, G. Sharma, A. Kumar and Z. AlOthman, *Desalin. Water Treat.*, 2016, **57**, 18551–18559.
- U. Guyo, T. Makawa, M. Moyo, T. Nharingo, B. Nyamunda and T. Mugadza, *J. Environ. Chem. Eng.*, 2015, **3**, 2472–2483.
- H. Kalavathy, I. Regupathi, M. G. Pillai and L. R. Miranda, *Colloids Surf., B*, 2009, **70**, 35–45.
- R. Shrivastava, R. Upreti and U. Chaturvedi, *FEMS Immunol. Med. Microbiol.*, 2003, **38**, 65–70.
- B. Flury, U. Eggenberger and U. Mäder, *J. Appl. Geochem.*, 2009, **24**, 687–696.
- M. Ragosta, R. Caggiano, M. D'Emilio and M. Macchiato, *Atmos. Environ.*, 2002, **36**, 3071–3087.
- J. A. Zazo, J. S. Paull and P. R. Jaffe, *Environ. Pollut.*, 2008, **156**, 29–35.
- D. E. Kimbrough, Y. Cohen, A. M. Winer, L. Creelman and C. Mabuni, *Crit. Rev. Environ. Sci. Technol.*, 1999, **29**, 1–46.
- S. E. J. G. Fendorf, *Geoderma*, 1995, **67**, 55–71.
- D. Rai, B. M. Sass and D. A. J. I. C. Moore, *Inorg. Chem.*, 1987, **26**, 345–349.
- A. Pechova and L. Pavlata, *Vet. Med.*, 2007, **52**, 1.
- M. Costa, *Toxicol. Appl. Pharmacol.*, 2003, **188**, 1–5.
- P. d. Magalhães Padilha, L. A. de Melo Gomes, C. C. Federici Padilha, J. C. Moreira and N. L. Dias Filho, *Anal. Lett.*, 1999, **32(9)**, 1807–1820.
- J. G. Espinola, S. F. Oliveira, W. E. Lemus, A. G. Souza, C. Airolidi and J. C. Moreira, *Colloids Surf., A*, 2000, **166**, 45–50.
- K. M. Kallury, W. E. Lee and M. J. A. C. Thompson, *Anal. Chem.*, 1993, **65**, 2459–2467.
- H. E. Fischer, S. A. King, J. B. Miller, J. Y. Ying, J. B. Benziger and J. J. I. C. Schwartz, *Inorg. Chem.*, 1991, **30**, 4403–4408.
- I. C. Ostroski, M. A. S. D. Barros, E. A. Silvab, J. H. Dantas, P. A. Arroyo and O. C. M. Lima, *J. Hazard. Mater.*, 2009, **161**, 1404–1412.
- J. H. Clark and D. J. Macquarrie, *Chem. Commun.*, 1998, **8**, 853–860.
- B. Gao, F. An and K.-K. Liu, *Appl. Surf. Sci.*, 2006, **253**, 1946–1952.
- O. G. Silva, E. C. S. Filho, M. G. Fonseca, L. N. H. Arakaki and C. Airolidi, *J. Colloid Interface Sci.*, 2007, **301**, 376–381.
- A. G. S. Prado and C. Airolidi, *Anal. Chim. Acta*, 2001, **432**, 201–211.
- A. Çimen, M. Torun and A. Bilgiç, *Desalin. Water Treat.*, 2015, **53**, 2106–2116.
- Y. Cui, X. Chang, X. Zhu, H. Luo, Z. Hu, X. Zou and Q. He, *Microchem. J.*, 2007, **87**, 20–26.
- İ. H. Gübbük, R. Güp and M. Ersöz, *J. Colloid Interface Sci.*, 2008, **320**, 376–382.
- A. R. Cestari, E. F. S. Vieira, A. J. P. Nascimento, F. J. P. Rodrigues, R. E. Bruns and C. Airolidi, *J. Colloid Interface Sci.*, 2001, **241**, 45–51.
- Z. A. Al-Othman, R. Ali and M. Naushad, *Chem. Eng. J.*, 2012, **184**, 238–247.
- G. Sharma, M. Naushad, H. Ala'a, A. Kumar, M. R. Khan, S. Kalia, M. Bala and A. Sharma, *Int. J. Biol. Macromol.*, 2017, **95**, 484–493.
- S. M. Alshehri, M. Naushad, T. Ahamad, Z. A. AlOthman and A. Aldalbahi, *Chem. Eng. J.*, 2014, **254**, 181–189.
- X. Zhu, X. Chang, Y. Cui, X. Zou, D. Yang and Z. Hu, *Microchem. J.*, 2007, **86**, 189–194.
- Y. E. Dolaksiz, F. Temel and M. Tabakci, *React. Funct. Polym.*, 2018, **126**, 27–35.
- S. Schmitz, A. C. Dona, P. Castignolles, R. G. Gilbert and M. Gaborieau, *Macromol. Biosci.*, 2009, **9**, 506–514.
- S. Sivrikaya, M. Imamoglu, S. Z. Yildiz and D. Kara, *Anal. Lett.*, 2016, **49**, 943–957.
- F. Cao, P. Yin, X. Liu, C. Liu and R. Qu, *Renewable Energy*, 2014, **71**, 61–68.
- S. Guo, H. Xu, F. Zhang, X. Zhu and X. Li, *Colloids Surf., A*, 2018, **546**, 244–253.
- P. Yin, Q. Xu, R. Qu, G. Zhao and Y. Sun, *J. Hazard. Mater.*, 2010, **173**, 710–716.
- J. A. A. Sales and C. Airolidi, *Thermochim. Acta*, 2005, **427**, 77–83.
- I. H. Gubbuk, *J. Hazard. Mater.*, 2011, **186**, 416–422.
- M. M. Kalhor, A. A. Rafati, L. Rafati and A. A. Rafati, *J. Mol. Liq.*, 2018, **266**, 453–459.
- D. Jiang, Y. Deng, G. Gao, L. Wu and H. Yang, *Colloids Surf., A*, 2018, **538**, 526–533.
- C. Xiong, S. Wang, W. Sun and Y. Li, *Microchem. J.*, 2019, **146**, 270–278.
- J. S. Piccin, G. L. Dotto and L. A. A. Pinto, *Braz. J. Chem. Eng.*, 2011, **28**, 295–304.
- M. N. Abbas, *Eur. Acad. Res.*, 2014, **2**, 8660–8690.
- A. Q. Selim, E. A. Mohamed, M. Mobarak, A. M. Zayed, M. K. Seliem and S. Komarneni, *Microporous Mesoporous Mater.*, 2018, **260**, 84–92.
- M. Auta and B. H. Hameedd, *Chem. Eng. J.*, 2011, **171**, 502–509.
- A. K. Kushwaha, N. Gupta and M. C. Chattopadhyaya, *Arabian J. Chem.*, 2017, **10**, S81–S89.



- 52 Y. Yang, Q. Li, G. Fang and S. Wang, *RSC Adv.*, 2016, **6**, 54510–54517.
- 53 H.-M. Shen, G.-Y. Zhu, W.-B. Yu, H.-K. Wu, H.-B. Ji, H.-X. Shi, Y.-B. She and Y.-F. Zheng, *Appl. Surf. Sci.*, 2015, **356**, 1155–1167.
- 54 P. Vejayakumaran, I. A. Rahman, C. S. Sipaut, J. Ismail and C. K. Chee, *J. Colloid Interface Sci.*, 2008, **328**, 81–91.
- 55 F. Adam, H. Osman and K. M. Hello, *J. Colloid Interface Sci.*, 2009, **331**, 143–147.
- 56 J. F. De Conto, M. R. Oliveira, M. M. Oliveira, T. G. Brandão, K. V. Campos, C. C. Santana and S. M. Egues, *Chem. Eng. Commun.*, 2018, **205**, 533–537.
- 57 A. H. Gemeay, E. F. Aboelfetoh and R. G. El-Sharkawy, *Water, Air, Soil Pollut.*, 2018, **229**, 16.
- 58 W. Ngeontae, W. Aeungmaitrepirom and T. Tuntulani, *Talanta*, 2007, **71**, 1075–1082.
- 59 V. S. O. Ruiz and C. Airolidi, *Thermochim. Acta*, 2004, **420**, 73–78.
- 60 K. Anbalagan, P. S. Kumar and R. Karthikeyan, *Desalin. Water Treat.*, 2016, **57**, 12585–12607.
- 61 G. De la Rosa, J. R. Peralta-Videoa and J. L. Gardea-Torresdey, *J. Hazard. Mater.*, 2003, **97**, 207–218.
- 62 R. Coşkun and Y. Dilci, *J. Macromol. Sci., Part A: Pure Appl. Chem.*, 2014, **51**, 767–782.
- 63 T. A. Khan, M. Nazir, I. Ali and A. Kumar, *Arabian J. Chem.*, 2017, **10**, S2388–S2398.
- 64 A. Çimen, A. Bilgiç, A. N. Kursunlu, İ. H. Gübbük and H. İ. Uçan, *Desalin. Water Treat.*, 2014, **52**, 4837–4847.
- 65 A. Çimen, E. Karakuş and A. Bilgiç, *Desalin. Water Treat.*, 2016, **57**, 7219–7231.
- 66 C. S. T. Araújo, I. L. S. Almeida, H. C. Rezende, S. M. L. O. Marcionilio, J. J. L. Léon and T. N. de Matos, *Microchem. J.*, 2018, **137**, 348–354.
- 67 Z. He, J. Yang, H. Li and H. J. E. Song, *Environment, Energy and Sustainable Development*, 2013, **4**, 177.
- 68 C. Srilakshmi and R. Saraf, *Microporous Mesoporous Mater.*, 2016, **219**, 134–144.
- 69 W. Liu, L. Yang, S. Xu, Y. Chen, B. Liu, Z. Li and C. Jiang, *RSC Adv.*, 2018, **8**, 15087–15093.
- 70 I. Enniya, L. Rghioui and A. Jourani, *Sustainable Chem. Pharm.*, 2018, **7**, 9–16.
- 71 R. Foroutan, H. Esmaeili, M. Abbasi, M. Rezakazemi and M. Mesbah, *Environ. Technol.*, 2018, **39**, 2792–2800.
- 72 B. Hameed, J. Salman and A. L. Ahmad, *J. Hazard. Mater.*, 2009, **163**, 121–126.
- 73 H. A. Akalin, U. Hicsonmez and H. Yılmaz, *J. Turk. Chem. Soc., Sect. A*, 2018, **5**, 85–116.
- 74 H. Dumrul, A. N. Kursunlu, O. Kocuyigit, E. Guler and S. Ertul, *Desalination*, 2011, **271**, 92–99.
- 75 Y. Wang, X. Tang, Y. Chen, L. Zhan, Z. Li and Q. Tang, *J. Hazard. Mater.*, 2009, **172**, 30–37.
- 76 L. Oubekka, N. Djelali, V. Chaleix and V. Gloaguen, *Rev. Roum. Chim.*, 2016, **61**, 175–185.
- 77 S. A. Chaudhry, T. A. Khan and I. Ali, *J. Mol. Liq.*, 2017, **236**, 320–330.
- 78 R. A. Anayurt, A. Sari and M. Tuzen, *J. Chem. Eng.*, 2009, **151**, 255–261.
- 79 H. Uçun, Y. K. Bayhan and Y. Kaya, *J. Hazard. Mater.*, 2008, **153**, 52–59.
- 80 M. Naushad, T. Ahamad, B. M. Al-Maswari, A. A. Alqadami and S. M. Alshehri, *Chem. Eng. J.*, 2017, **330**, 1351–1360.
- 81 R. Karthik and S. Meenakshi, *Journal of Water Process Engineering*, 2014, **1**, 37–45.
- 82 Y. Liu, L. Guo, L. Zhu, X. Sun and J. Chen, *Chem. Eng. J.*, 2010, **158**, 108–114.
- 83 M. R. Gandhi and S. Meenakshi, *J. Hazard. Mater.*, 2012, **203**, 29–37.
- 84 J. Y. Qiu, Z. Y. Wang, H. B. Li, L. Xu, J. Peng, M. Zhai, C. Yang, J. Q. Li and G. S. Wei, *J. Hazard. Mater.*, 2009, **166**, 270–276.
- 85 L. Zhao, J. Sun, Y. Zhao, L. Xu and M. Zhai, *Chem. Eng. J.*, 2011, **170**, 162–169.
- 86 P. A. Kumar, M. Ray and S. Chakraborty, *J. Hazard. Mater.*, 2007, **143**, 24–32.
- 87 K. Choi, S. Lee, J. O. Park, J.-A. Park, S.-H. Cho, S. Y. Lee, J. H. Lee and J.-W. Choi, *Sci. Rep.*, 2018, **8**, 1438.
- 88 P. K. Pandey, S. K. Sharma and S. S. Sambhi, *Int. J. Environ. Sci. Technol.*, 2010, **7**, 395–404.

