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Preparation of CoFe$_2$O$_4$@Carboxymethylcellulose/HZSM-5 as a Novel Nano magnetic Adsorbent for Metronidazole Removal: Isotherm, Kinetic and Thermodynamic Study

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Abstract

In this research CoFe$_2$O$_4$@CMC/HZSM-5 synthesized in the presence of Carboxymethylcellulose as a biopolymer in the green conditions as a new nanomagnetic adsorbent for metronidazole removal from aqueous media. Physical and chemical structure of adsorbent was investigated by FTIR, FESEM, EDS, Mapping, TEM, XRD, VSM, BET and TGA techniques. The results indicated that 94% of the metronidazole concentration removed by CoFe$_2$O$_4$@CMC/HZSM-5 in the optimal conditions including pH 6, temperature 20°C, metronidazole concentration 50 mg/L, contact time 60 min and adsorbent dose 2 g/L. The resulting data from adsorption experimental experiments had better correlated with the Freundlich isotherm and pseudo-second order kinetic. Also, the thermodynamic study demonstrated that the adsorption process was an endothermic process and had a physical mechanism. As well as, the achieved findings demonstrated that following six adsorption runs, the adsorbent chemical structure had no change and the nanomagnetic adsorbent efficiency in the removal process had a slight decrease. CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanocomposite had effective adsorption capacity for metronidazole removal.

Keywords: Metronidazole, Carboxymethylcellulose, Adsorption
1. Introduction

About 30-90% of antibiotics are not metabolized in the animal and human bodies. Eventually, they enter the environment through the urine and stool in the form of active compounds. The hospital wastewater is the main source of antibiotics in the environment [1, 2]. Medications and antibiotics are the most important groups of emerging pollutants. These organic compounds enter the water sources in two ways. One of these pollution sources is the pharmaceutical industry wastewater and the other one is excretion of the medicines compounds metabolites due to their incomplete metabolism during treatment. Because of the ability of these compounds to alter ecosystems and reduce biodiversity, removal of these compounds from the aquatic environments is so important [3].

Metronidazole is one of the nitroimidazole antibiotics group which has antibacterial and anti-inflammatory properties and it can be used to treat diseases that caused by anaerobic bacteria, protozoa, and bacteroides. Metronidazole is one of the most commonly used antibiotics in the world which is in the WHO's list of essential medicines. Metronidazole, like other antibacterial and antioxidant medicines which have nitroimidazole ring structure, is suspicious to causes carcinogenic and mutagenic [4]. By reason of the non-biodegradability, toxicity, mutagenic and carcinogenic potential of metronidazole, the wastewater containing this antibiotic causes harmful effects on the humans and ecological environment so, removal of metronidazole has environmental, technical and economic importance [5].

Removal of complex organic compounds such as antibiotics from water resources with conventional filtration methods due to their sustainability in the environment and non–biodegradable structure is so difficult. Purification processes including ion exchange [6], membrane filtration [7], solvent extraction [8], electrochemical methods [9], chemical precipitation [10], biological methods [11], electrochemical oxidation [12], catalytic [13, 14] and fenton oxidation [15, 16] are used for water treatment.

The mentioned methods have some disadvantages. For example disadvantages of the membrane filtration process are the membrane blockage owing to the chemical sediment growth formation and the physicochemical properties changes of the membrane surface [17]. In the biological methods, during the treatment process, high concentrations antibiotics could make toxicity. The other problem is resistance of some antibiotics against biological treatment. In addition, the chemical methods because of sludge production and need to chemical materials are costly and not popular [18].

The adsorption process compared to the other mentioned methods has more high efficiency, simple design and operation, low cost and non-production of hazardous materials, which are basic features of the adsorption process. Therefore, many researchers focus on its use in the environmental pollutants removal [19].

Some of adsorbents such as activated carbon, ash, coal, metal oxides, agricultural wastes, kaolinite, bagasse, resins, and compost are used as adsorbents to removal of heavy metals and some organic compounds from the aquatic environments. But, some of these adsorbents show low adsorption efficiency since their low surface areas and adsorption capacities. Also, separate those from the environment by filtration and centrifugation methods is time-consuming. In
most cases, the adsorbent passes through the filter and leads to the secondary pollution production in the environment [20-24].

To solve this problem, magnetic nanoadsorbents are used. One of the most widely used adsorbents is magnetic nanocomposites [25-27]. Magnetic nanocomposites due to their good magnetic property, high area surface and adsorption capacity have attracted the many researchers attention. In a lot of researches, nanocomposites which contain some of biopolymers such as cellulose, methylcellulose, carboxymethylcellulose, and chitosan in their structure, are used as bioadsorbents in water treatment processes [28, 29].

The mentioned polysaccharide compounds owing to cross-links and functional groups such as OH, NH$_2$, COOH, COONa in their structure can be easily dissolved in water and cause electrostatic interaction between the adsorbent and the contaminant [30]. These polysaccharide derivatives have some advantages such as availability, biodegradability and non-toxicity. Therefore, they are widely used as adsorbents for the pollutants removal from aquatic environments [31]. Cellulose derivatives such as Methylcellulose (MC), hydroxypropyl methylcellulose (HPMC), ethylcellulose (EC) and carboxymethylcellulose (CMC) are used in the various nanocomposites preparation for water treatment process [32]. Using raw cellulose because of some disadvantages such as lower solubility, inability to reuse, poor interaction with pollutants and low surface area is limited. So, to solve this problem, the nanocomposites combined with ionic functionalization, ester, and the etheric cellulose derivatives are used which can produce advanced material based on the cellulose derivatives [33, 34]. Among these derivatives, CMC is a frequently used cellulose derivative and soluble in the water, which is used as an adsorbent in the water purification process. CMC as a biopolymer is sensitive to pH changes [35]. Use of these adsorbents could make some limitations such as filtration, dispersion, turbidity, secondary contamination and high cost of regeneration. Therefore, create conditions for optimal use of CMC and other nanoscale or powdered adsorbents could make magnetic property in them through compression with metal ferrite compounds [36]. Polysaccharide-based composites owing to their impressive properties such as high surface area, chemical stability and mechanical strength have attracted great attention to themselves [37].

To enhance properties of these composites including physiochemical and adsorption capability, some various materials such as montmorillonite [38], bentonite [39], alumina [40], kaolin [41], and zeolite [42] have been used in their preparation. HZSM-5 zeolite by reason of some properties like low priced, diversity of structural properties, good mechanical firmness, high surface area, and high adsorption valence towards cation species was utilized as a solo adsorbent or composite in the water pollutants removal [43]. In the field of pollutants removal from water and wastewater HZSM -5 zeolite composites are receiving increased attention. In previous research about pollutants removal from aqueous, various ferritic cellulose derivatives such as ZnFe$_2$O$_4$@CMC, ZnFe$_2$O$_4$@CMC, CuFe$_2$O$_4$@MC and CoFe$_2$O$_4$/AC@chitosan due to the renewability, cellulose stable nature reasonable price, and availability have been prepared and applied as sorbents or photocatalysts [33, 34]. Also, several cellulosic adsorbents such as attapulgite/chitosan (ATP/CS) [44], chitosan-cellulose enwrapped magnetic carbon foam (CCMF), carboxylated cellulose/MnFe$_2$O$_4$ aerogel [45], magnetic-PEI-cellulose [46],
Fe$_3$O$_4$@SiO$_2$@Cellulose [47], Fe$_3$O$_4$@CMC@PDA [48][49], grafted magnetic cellulose with thiol/amine (GMC–N/S)[50], magnetic carboxylated cellulose nanocrystal composite (CCN-Fe$_3$O$_4$) [51], Magnetic chitosan/cellulose [52], magnetic hydrogel beads (m-CS/PVA/CCNFs) [53], amino-functionalized magnetic cellulose composite [49], magnetic cellulose-chitosan hydrogels have been used for the pollutants removal from aqueous solutions [48].

In this research CMC that applied in the synthesis of the HZSM, played an essential role to improve the CoFe$_2$O$_4$@CMC/HZSM-5 nanoadsorbent structural features. CMC in the CoFe$_2$O$_4$@CMC/HZSM-5 composition caused increasing the adsorbent surface area. As well as, carboxyl and hydroxyl groups in CMC create the electrostatic attraction between metronidazole and CoFe$_2$O$_4$@CMC/HZSM-5 composite surface. CoFe$_2$O$_4$@CMC/HZSM-5 was prepared with highly efficient microwave-assisted co-precipitation method by iron and cobalt salts on CMC in the presence of HZSM-5 and was used in highly efficient adsorption of the metronidazole. CoFe$_2$O$_4$@CMC/HZSM-5 that was used as a nanomagnetic adsorbent in the adsorption process to remove contaminants was easily separated from aqueous solutions as well as recycled and reused. According to the literature, the purpose of this investigation is preparation and characterization of CoFe$_2$O$_4$@CMC/HZSM-5 as a new nanomagnetic adsorbent for metronidazole removal.

2. Materials and Methods

2.1. Materials and equipment

All the required materials for this study, including carboxymethylcellulose (CMC), iron chloride (FeCl$_3$.6H$_2$O), cobalt chloride (CoCl$_2$.6H$_2$O) and nitric acid (HNO$_3$), were purchased from Merck (Germany). All used solutions in the study were prepared from high purity deionized water (DW). The solutions pH were adjusted using chlorinated hydrochloric acid (HCl) and 0.1 molar sodium hydroxide (NaOH) and were measured by using pH meter (HACH-HQ-USA). In order to adsorbent magnetic separation from the solution, a cubic magnet with a magnetic intensity of 1.3 Tesla was used. The metronidazole was bought from DarouPakhsh Holding Company (Iran).

2.2. Preparation of CoFe$_2$O$_4$@CMC/HZSM-5

The magnetic nanocomposite was prepared using iron chloride (FeCl$_3$.6H$_2$O), cobalt chloride (CoCl$_2$.6H$_2$O) salts, carboxymethylcellulose (CMC) and /HZSM-5 zeolite. At first, salts of iron chloride FeCl$_3$.6H$_2$O and cobalt chloride CoCl$_2$.6H$_2$O (2:1) were dissolved in 100 mL of distilled water (DW) then, 1 g of CMC and finally, 1 g of HZSM-5 zeolite were added to the solution. Subsequently, NaOH was added to the resulting suspension for one hour until the solution pH reached 11. The reaction container color changed and became black after one hour of stirring. Then, the reaction container was transferred to the microwave and was irradiated by microwave waves. The radiation process was performed in three periods of five minutes with 50% power of the microwave and 30 seconds with the off mode. The obtained product was a light and black powder that was washed many times using DW and dried in the oven (100 °C within 24 h) (Figure 1).
2.3. Characterization techniques of CoFe$_2$O$_4$@CMC/HZSM-5

Fourier-transform infrared spectroscopy (FTIR) was done using WQF-510 FT-IR spectrometer to determine the adsorbent’s functional groups. Characterization of the specimens was done using a field-emission scanning electron microscope-energy dispersive X-ray spectroscopy-Mapping (FESEM-EDS-Mapping) (FE-SEM TESCAN MIRA3) and Transmission electron microscopy (TEM-Philips EM 208S). To find out the presence of the cobalt ferrite crystalline structure in the adsorbent, X-Ray diffraction (XRD) using Philips X-Pert device (the Netherlands) was employed. The magnetic property of the CoFe$_2$O$_4$@CMC/HZSM-5 were characterized by VSM (LakeShore Cryotronics-7404) at room temperature. To measure the porosity of the adsorbent surface area, Brunner-Emmett-Taller (BET) method and micrometrics model 021LN2 transfer device was used. Also, thermal gravimetric analysis (TGA) was applied to evaluate the adsorbent thermal resistance (TA Co. Model Q600-USA).

2.4. Adsorption Method

The metronidazole adsorption process experiments on the CoFe$_2$O$_4$@CMC/HZSM-5 nanocomposite were performed in the batch reactor and inside the Erlenmeyer flasks with a volume of 100 mL. At first, metronidazole stock solutions with a concentration of 1000 mg/L were prepared. Then, in order to investigate the synthesized adsorbent capabilities, 5 g of adsorbent was added to 50 mL metronidazole solution 50 mg/L and was placed in the shaker at 240 rpm for a certain period of time. After this period, by using magnetic separation, the adsorbent was separated from the solution and the metronidazole remaining concentrations were measured.

To recognize and measure the remained concentration of metronidazole, a HPLC device (YL 9100 Waters, USA) was used. Also, for adjusting the device Standard metronidazole with a purity of 99% was utilized. As well as, deionized water and acetonitrile were applied as mobile phases with volume ratios of 30:70. The utilized column was C18 with 5 μm particles, length of 250 mm, and internal diameter of 4.6 mm. By using a UV absorbance detector within the wavelength of 348 nm with an injection volume of 20 μL and flow rate of 1 mL/min, the metronidazole remained concentration was measured.

As control samples, the metronidazole samples with similar concentrations without adsorbent were utilized in the all steps of the experiment. The metronidazole initial concentration (50-300 mg/L), pH (3 -11), amount of adsorbent (0.5 -3 g/L), contact time (0 -300 min), and temperature (20-50°C) were investigated, to optimize the adsorption conditions and determine the optimal values of each of the adsorption process affecting factors. By solid addition method, the CoFe$_2$O$_4$@CMC/HZSM-5 point of zero charge (pHpzc) was evaluated. At first, 10 different pH (2–11) values, 100 mL of KCl solution (0.1 mol/L) was prepared. Then, 0.01 g of the magnetic nanobiocomposite was added to each solution. After that, the obtained mixture was stored at room temperature overnight. In the next step, by using pH meter (Hanna), the initial and final pH (pH$_i$) values of solutions were determined. To obtain the pH$_{pzC}$, ΔpH = pH$_f$ – pH$_i$
and pH, plot was drawn. At last, the intersection point of ΔpH = 0 was taken as pH\textsubscript{pzc}. By using NaOH and HCl 0.1 N, the solutions pH was adjusted.

The adsorption experiments were repeated three times in each step and their mean values as the results were used in the calculations. At the end of each experiment, the amount of adsorbed metronidazole on adsorbent (\(q_e\)) and the percentage of adsorption for each sample was calculated according to equations (Eq. 1) and (Eq. 2), respectively.

\[
q_e = \frac{(C_0 - C_e)}{W} \quad \text{(Eq. 1)}
\]

\[
\text{Adsorption(\%) = }\left(1 - \frac{C_e}{C_0}\right) \times 100 \quad \text{(Eq. 2)}
\]

Where \(C_0\) (mg/L) is the adsorbate initial concentration (metronidazole), \(C_e\) (mg/L) is the pollutants equilibrium concentration and \(W\) is the adsorbent mass in the solution volume (g/L).

### 2.5. Adsorption kinetic equations

The kinetic equations are used to describe the behavior of the adsorbate molecules transfer per time. The adsorption process maybe takes place in several stages with different speeds. In the present study, four kinetic models, Lagergren (pseudo-first order) (Eq. 3), HO (pseudo-second order) (Eq. 4), Elovich model (Eq. 5) and intraparticle diffusion model (Eq. 6) were used.

\[
\log (q_e-q_t) = \log q_e - k_1 t/2.303 \quad \text{(Eq. 3)}
\]

\[
t/q_t = 1/k_2 q_e^2 + 1/q_e t \quad \text{(Eq. 4)}
\]

\[
q_t=\beta \ln (\alpha/\beta) + \beta \ln t \quad \text{(Eq. 5)}
\]

\[
q_t = k_i t^{1/2} + C_i \quad \text{(Eq. 6)}
\]

\(k_1\) (1/min) and \(k_2\) (mg/g min) are the pseudo-first order and pseudo-second order rate constant, respectively. \(q_e\) (mg/g) and \(q_t\) (mg/g) are adsorption capacity at equilibrium time and the specified time, respectively and \(\alpha\) (mg/g/min) and \(\beta\) (g/mg) are also the initial adsorption rate and the surface coating size (or activation energy for chemical adsorption).

### 2.6. Adsorption equilibrium equations

In the adsorption systems design, the equilibrium isotherms are used as important parameters. In fact, they describe relationship between the adsorbent adsorption capacity and adsorbate concentration. In the present study, two common models, the Freundlich (Eq. 7) and Langmuir (Eq. 8) model, have been used for this purpose.

\[
\log q_e = \log k_F + 1/n \log C_e \quad \text{(Eq. 7)}
\]

\[
\frac{C_e}{q_e} = \frac{C_e}{q_0} + 1/k_F q_0 \quad \text{(Eq. 8)}
\]

\(k_F\) (mg/g(L/mg) and \(n\) are the Freundlich constants that \(1/n\) is intensity of the adsorption process and \(k_F\) is adsorbent adsorption capacity. As well as, \(k_L\) (L/mg) is the Langmuir constant and \(q_0\) (mg/g) is the maximum amount of adsorbed materials per unit mass of adsorbent. \(R_L\) is
used to find out the adsorption process desirability in the Langmuir model and determinant by using the following equation:

\[
R_L = \frac{1}{1 + kLC_0}
\]  
(Eq. 9)

3. Results and Discussion

3.1. CoFe_2O_4@CMC/HZSM-5 characterization

3.1.1. FT-IR spectroscopy

Comparative FT-IR of CMC, HZSM-5, and CoFe_2O_4@CMC/HZSM-5 are demonstrated in Fig. 2. In the FT-IR spectrum of carboxymethyl cellulose the wide absorption band at 3424 cm\(^{-1}\) belonged to the stretching vibrations of the OH groups. The absorption band at 2920 cm\(^{-1}\) belonged to the stretching vibrations of the CH\(_2\) groups. At 1111 and 1160 cm\(^{-1}\), the CMC typical vibration was representing the stretching vibration of the C-OH groups. The stretching vibrations of the intra molecular hydrogen bond were related to the strong broad band at 3444 cm\(^{-1}\). The band at 2920 cm\(^{-1}\) showed the stretching vibrations of the C-H bonds. Moreover, the strong peak at 1648 cm\(^{-1}\) was associated with the stretching vibrations of the carboxyl group, and the absorption bands in the range 1100 to 1150 cm\(^{-1}\) belong to C-O stretching vibrations in carboxymethyl cellulose. In the FT-IR spectrum of HZSM-5 the wide absorption band at 3424 cm\(^{-1}\) belonged to the stretching vibrations of the OH groups. The absorption bands at 799 cm\(^{-1}\) (external symmetric stretch), 1102 cm\(^{-1}\) (internal asymmetric stretch), 450 cm\(^{-1}\) (T-O bending vibration of the Si-O and Al-O internal tetrahedral) and 548 cm\(^{-1}\) (double ring vibration) were referred to the HZSM-5 typical characteristics. Also, an asymmetric stretch vibration of the band at 1221 cm\(^{-1}\), was confirmed the HZSM-5 zeolite presence [30]. Meanwhile, this band can used to determine the crystallinity of HZSM-5 zeolites and show the absence of amorphous silica. In the FT-IR of CoFe_2O_4@CMC/HZSM-5, the wide absorption band at 3431 cm\(^{-1}\) was associated with the stretching vibrations of hydroxyl groups that overlaid with acidic OH group of carboxyl group. The absorption band at 1623 cm\(^{-1}\) belonged to the stretching vibrations of the carboxyl group, confirming the presence of carboxymethyl cellulose in the structure of CoFe_2O_4@CMC/HZSM-5. Also, Strong absorption bonds at 1349 cm\(^{-1}\) and 1381 cm\(^{-1}\) indicates the presence of HZSM-5 zeolite in the structure of CoFe_2O_4@CMC/HZSM-5. In addition, the absorption bands around 400-600 cm\(^{-1}\) in CoFe_2O_4@CMC/HZSM-5 is related to vibrations of metal-oxygen groups (Co-O, Fe-O) in the cobalt ferrite and zeolite structure (Si-O, Al-O).

(Figure 2)
3.1.2. FESEM-EDS-Mapping and TEM of CoFe$_2$O$_4$@CMC/HZSM-5

The surface morphology of the adsorption material was assessed by FESEM and TEM. The morphological result of CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanocomposite is illustrated in Fig. 3a-c. The CMC attendance in the CoFe$_2$O$_4$@CMC/HZSM-5 synthesis causes the smoothly, uniformly, and loosely aggregated sphere-shaped of magnetic nanocomposite form. The average particle size of CoFe$_2$O$_4$@CMC/HZSM-5 is about 27 nm.

(Figure 3)

The synthesized CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanoadsorbent purity and chemical structure were evaluated by EDS analysis (Fig. 4). The results obtained 46.44% O, 26.82% Fe, 14.16% Co, 8.56% C, 3.88% Si, and 0.15% Al which are following the expected values.

(Figure 4)

Mapping is the observation of how the elements are distributed at high resolution. It was used to investigate the CoFe$_2$O$_4$@CMC/HZSM-5 elements distribution. Based on the obtained results which is showed in Fig. 5, Al, Si, Co, Fe, O, and C had a homogeneous distribution that indicates high uniformity of the synthesized CoFe$_2$O$_4$@CMC/HZSM-5 (Fig. 5).

(Figure 5)

3.1.3. XRD

The XRD patterns were prepared separately from CMC, HZSM-5, CoFe$_2$O$_4$, and CoFe$_2$O$_4$@CMC@HZSM-5 and were compared with each other (Figure 6). In the CMC XRD pattern, the index peak can be seen in the area of 18.28°, 23.15° and 24.06° peaks are belong to HZSM-5 and the values which are seen in $2\theta$ =30.26°, 35.70°, 43.35°, 53.74°, 57.29°, and 62.81° are related to cobalt ferrite. The XRD results of CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanocomposite in the range from 10 to 80 degrees ($2\theta$ = 10°- 80°) are depicted in figure 6. The sharp diffractions in regions 18.28°, 23.15°, 24.06°, 30.26°, 35.70°, 43.35°, 53.74°, 57.29°, and 62.81° indicated the crystalline structure in the spinel shape, which was consistent with the Joint Committee on Powder Diffraction Standards (no. 96-900-5813). The presence of sharp and strong peaks in this pattern and comparison of peak locations with reference data indicated the crystalline structure of cobalt ferrite with complete crystallization that has preserved the crystalline structure of CoFe$_2$O$_4$@CMC/HZSM-5, after being composite with carboxymethylcellulose and zeolite.

(Figure 6)

3.1.4. VSM

The results of vibrating-sample magnetometer (VSM) analysis of CoFe$_2$O$_4$@CMC/HZSM-5 are demonstrated in figure 7. Magnetic coercive force (Hc), saturation magnetization (Ms), and
Remnant magnetization (Mr) were equal to 0.73 Qₑ, 41.36 emu/g, and 50 emu/g, in that order. The results revealed that the CoFe₂O₄@CMC/HZSM-5 magnetic nanocomposite had a ferromagnetic effect with elevated saturation magnetization and high magnetic coercive force. It could be quickly separated and retrieved from the reaction container using a magnet and could be reused in subsequent periods.

(Figure 7)

3.1.5. BET

The obtained information from the BET test showed that surface area, pore volume, and pore size equaled 235.21 m²/g, 0.207 cm³/g, and 3.52 nm, respectively. The BET analysis results of CoFe₂O₄@CMC/HZSM-5 magnetic nanocomposite can be considered as a mesoporous material

3.1.6. TGA

TGA was used to evaluate CoFe₂O₄@CMC/HZSM-5 thermal decomposition. The results are demonstrated in Fig. 8. TGA analysis was done in the room temperature to 600 °C, to evaluate the CoFe₂O₄@CMC/HZSM-5 nanocomposite thermal stability (Fig. 8). The mass loss was happened in the temperature range of 20–100 °C which is linked with the adsorbed water loss. Due to the breakdown of CMC glycosidic bonds and also, its decomposition into lower fatty acids, the weight loss was reported at 100–600 °C. According to the results, the adsorbent total weight loss was seen 10.18% until 600 °C which shows the CoFe₂O₄@CMC/HZSM-5 magnetic nanocomposite high thermal stability

(Figure 8)

3.2. Adsorption experiment optimization

3.2.1. pH effect

Environmental acidity where the adsorption process takes place is largely effective in adsorption and removal efficiency. A part of this issue is related to hydrogen ions (H⁺), which adsorbed on the adsorbent surface. The contaminant adsorption on the adsorbent surface leads to reduce the positive ions adsorption. The other part refers to the pH effect, which is effective in some compounds ionization. In order to evaluate effect of pH changes on the metronidazole removal by the CoFe₂O₄@CMC/HZSM-5 nanocomposite, 50 mL solution containing 2 g/L adsorbent and the metronidazole initial concentrations 50 mg/L were used. The results are presented in figure 9. The obtained results showed that the pollutant removal efficiency increased with increasing pH from 2 to 6 and decreased at pH values more than 6. The highest metronidazole removal efficiency took place at pH 6 and was equal to 94 %.

(Figure 9)
The effect of pH could be explain in terms of the adsorbent pH<sub>zpc</sub> and MNZ pK<sub>a</sub>. pH<sub>zpc</sub> of CoFe<sub>2</sub>O<sub>4</sub>@CMC/HZSM-5 was determined by solid addition method and was obtained 6.8. At pHs less than pH<sub>zpc</sub>, the CoFe<sub>2</sub>O<sub>4</sub>@CMC/HZSM-5 surface charge was positive; thereby the chance of MNZ ions adsorption increased. At pHs more than pH<sub>zpc</sub>, the surface charge of the CoFe<sub>2</sub>O<sub>4</sub>@CMC/HZSM-5 was negative. As a result, the repulsion between the interfaces increased in the solution [54]. On the other hand, high pH leads to higher concentrations production of -OH in the solution and also, inhibit of MNZ ions scattered, thereby the chance of MNZ adsorption decreased [54]. Therefore, pH 6 was selected as the optimum pH.

3.3.2. Equilibrium time

The equilibrium time for the metronidazole removal process was evaluated over a period of 300 min, metronidazole initial concentrations 50 mg/L and at the optimum pH (Figure 10). The above figure shows that the adsorbates adsorption rate in the initial times was much higher than the next times or in other words, by increasing contact time, the adsorption capacity increased. But after 60 minutes, no significant changes were observed in adsorption capacity. For example, by increasing the contact time from 60 to 300 minutes, the metronidazole adsorption capacity increased from 22.5 and 23.5 mg/g to 22.6 and 23.7 mg/g, respectively. These results indicate that the metronidazole removal process is balanced in 60 minutes and therefore, this time was considered as the equilibrium time for the adsorption experiments continuation. Significant increase in adsorption capacity at the beginning of the contact time (Figure 10) could be relate to the presence of many active and unsaturated sites on the CoFe<sub>2</sub>O<sub>4</sub>@CMC/HZSM-5 external surfaces. However, by increasing contact time, the adsorption capacity is stabilized. The stabilization of the adsorption capacity or reaching the equilibrium point in more than 60 minutes may be due to the active sites occupation on the adsorbent surface or the adsorbent capacity completion. In general, by increasing contact time, the adsorbate ions access to the empty active sites on the adsorbent surface is reduced and eventually these sites are saturated and the process reaches equilibrium.

(Figure 10)

3.3.3. Influence of different concentrations of adsorbent and adsorbate

Figures 11a and b are showing the results of the adsorbent (CoFe<sub>2</sub>O<sub>4</sub>@CMC/HZSM-5) and adsorbate different amounts concentration (metronidazole) effects under optimal conditions, respectively. As shown in figure 11a, by increasing the amount of adsorbent from 0.5 to 3 g/L, the metronidazole adsorption increased from 73 to 94%. Therefore, considering that in the amount of 2g/L of adsorbent, the removal efficiency was 94%, this value was selected as the adsorbent optimal dose. But, in figure 11b, it can be seen that the changes in the initial concentration had a negative effect on the adsorption rate, such that by increasing the metronidazole initial concentration from 50 to 300 mg/L, the removal efficiency decreased from 95 to 59%, respectively. The removal efficiency increasing in the study of the amount of adsorbent changes (Figure 11a) maybe due to the adsorbent surface increasing or increase the access of the adsorbate molecules to the pores on the magnetic nanocomposite surface. On the other hand, figure 11b shows that by increasing the metronidazole initial concentrations, the
removal efficiency decreased. Probably, constant numbers of active sites on the adsorbent against the increasing in the number of adsorbate molecules or the saturation of the adsorbent surface at high concentrations of contaminants was the main reason for the decrease in removal efficiency by increasing the initial contaminant concentration.

(Figure 11)

3.3.4. Adsorption kinetic

The kinetic parameters of the metronidazole removal process are presented in table 1. The computational adsorption capacity values \( q_e, \text{cal} \) for pseudo-first order and pseudo-second order models were obtained 22 and 24.2 mg/g, respectively. In addition, the experimental adsorption capacity \( q_e, \text{exp} \) from the obtained experiments was 23.5 mg/g. The lowest correlation coefficient was obtained for the Elovich equation and intraparticle diffusion model. The curves of the studied metronidazole removal kinetic models compared with the obtained experimental data from the experiments are presented in figure 12. The results of the metronidazole removal kinetic study by CoFe\(_2\)O\(_4@CMC/HZSM-5\), according to the table 1 showed that based on the calculated correlation coefficients, the adsorption behavior per unit time follows pseudo-first order and pseudo-second order models. However, it is observed that the metronidazole correlation coefficient in the pseudo-second order kinetic was 0.997 that is higher than the obtained values for the pseudo-first order kinetic model. Therefore, it can be admitted that the model in the present study has more validity than the other three models. Tang et al. (2012) [55] in the study on aniline and caprolactam simultaneous adsorption by the granular activated carbon found that the pollutants adsorption behavior per unit time follows the pseudo-second order model. As well as, valderrama in the study of the phenol and aniline simultaneous adsorption by the activated carbon, reported both pseudo-first order and pseudo-second order kinetic models as suitable models for the data describing. According to table 1, the boundary layer thickness values \( Ci \) in the intraparticle diffusion model were not equal to zero, that indicates the intraparticle diffusion model was not the only controlling step of the process and perhaps the external adsorption mechanism may also had been effective as a control factor in the adsorption process.

(Figure 12)

(Tabla 1)

3.3.5. Adsorption isotherm

Table 2 shows the Langmuir and Freundlich equilibrium isotherms values for the metronidazole adsorption in the temperature range of 20-50 °C. The above table indicates that the correlation coefficient in Langmuir and Freundlich models is \( R^2>0.97 \). In this research, the maximum adsorption capacity \( q_0 \) based on the Langmuir model at 50 °C was obtained 206.6 mg/g. In figure 13, the curves of the Langmuir and Freundlich equilibrium isotherm models with experimental data were compared. In the metronidazole adsorption isotherms examination (Table 2) it can be seen that the obtained regression coefficients in Freundlich and Langmuir models for all three temperatures are more than 0.97. In fact, these results indicate that both
models are suitable for describing experimental test data. In the above table, it is also noteworthy that the pollutant maximum adsorption capacity values based on the Langmuir isotherm model for the increased by increasing the solution temperature. For example, by increasing temperature from 20 to 50 °C, the maximum adsorption capacity increased from 201.1 to 206.6 mg/g. This trend indicates that the increase in the temperature had a positive effect on the adsorption capacity. In addition, the Langmuir model RL values of the adsorbent were studied at all temperatures and was obtained less than one (1<RL<0). Therefore, it is concluded that the CoFe2O4@CMC/HZSM-5 had high efficiency in the metronidazole removal. Table 2 also shows that the 1/n values related to the Freundlich isotherm were less than one in all three temperatures, which indicate the high adsorption of the pollutant.

(Figure 13)
(Table 2)

3.3.6. Adsorption thermodynamic

The results of measuring Lnk_c values versus temperature changes based on the Vant Hoff curve to determine the values of the metronidazole adsorption process thermodynamic parameters are shown in figure 14. Also, table 3 presents the obtained values of the thermodynamic parameters. Table 3 demonstrates that the standard enthalpy (ΔH°) and standard entropy values (ΔS°) are positive and the standard gibbs free energy (ΔG°) is negative. Due to the presented results in table 3, it can be seen that the standard enthalpy values (ΔH°) is positive, which indicates that the metronidazole removal process is endothermic and adsorption at the higher temperatures is desirable. Standard entropy value (ΔS°) was obtained positive which indicates the adsorbent tendency to adsorb the adsorbate and increase the efficiency by increasing the temperature in the solid-liquid interface during the adsorption process. Nevertheless, the Gibbs free energy values (ΔG°) were negative in all three studied temperatures. These negative values indicate that the metronidazole removal by CoFe2O4@CMC/HZSM-5 is spontaneous reaction. Since the metronidazole ΔH° value was less than 40 kJ/mol, it can be concluded that the adsorption process had a physical mechanism. In addition, the ΔG° obtained values at all the studied temperatures were less than -20 kJ/mol, which confirm that the adsorption process was a physical type.

(Figure 14)
(Table 3)

3.3.7. Effect of CMC on the removal efficiency

The obtained metronidazole removal efficiencies by CoFe2O4, CMC, HZSM-5, and CoFe2O4@CMC/HZSM-5 were compared (Figure 15). Based on the findings, the removal efficiencies by using HZSM-5, CMC, CoFe2O4, and CoFe2O4@CMC/HZSM-5 were 75%, 83%, 87%, and 94%, respectively. CoFe2O4@CMC/HZSM-5 showed a significantly higher
removal efficiency than the other adsorbents after 60 min contact time. It can be concluded that CoFe$_2$O$_4$@CMC/HZSM-5 composed of carboxymethylcellulose as a polysaccharide with high carboxyl and hydroxyl groups. Consequently, the presence of these functional groups causes the interaction of electrostatic attraction between the adsorbent surface and metronidazole [56, 57]. As well as, CMC plays an important role in improving the structural properties of CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanocomposite. The presence of CMC during the preparation of CoFe$_2$O$_4$@CMC/HZSM-5, increases the adsorbent surface area and provides more active sites for the metronidazole removal.

(Figure 15)

3.4. Mechanism

The metronidazole adsorption efficiency depends upon to the surface groups of the magnetic nanoadsorbent protonation and non-protonation status. The pH of solution is one of the most important parameters which can qualifies the CoFe$_2$O$_4$@CMC/HZSM-5 charge and has effect on the interaction between metronidazole molecules. To determine the amount of CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanoadsorbent surface charge and its type, at different pHs, pH$_{ZPC}$ was investigated. The synthesized magnetic nanoadsorbent surface charge was negative at pHs more than 6.8, although at the lower pHs, the adsorbent surface charge was positive. Metronidazole due to its electron-rich aromatic rings has desire to adsorb on the adsorbent positively charged surface. Metronidazole pK$_a$ was equal to 2.55, it created an electrostatic gravity energy which leaded to more metronidazole removal by the CoFe$_2$O$_4$@CMC/HZSM-5. The amount of metronidazole molecules which have negatively charged increased slowly by increasing the pH value in the solution [58]. With attention to the CoFe$_2$O$_4$@CMC/HZSM-5 adsorption behavior at different pH values and its characteristics, the main mechanism for the metronidazole removal perhaps makes a complex between metronidazole anions and the adsorbent surface. The metronidazole removal mechanism by CoFe$_2$O$_4$@CMC/HZSM-5 is shown in figure 16.

(Figure 16)

3.5. Regeneration, reusability and chemical stability of CoFe$_2$O$_4$@CMC/HZSM-5

Regeneration and recycling are two important items in the adsorbent functional. After metronidazole removal, CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanocomposite, were recycled for six consecutive adsorption runs. To regenerate CoFe$_2$O$_4$@CMC/HZSM-5, this magnetic nanocomposite, following each cycle, was separated from the solution by a magnet and was washed with HCl/DW for several times. At last, the magnetic nanoadsorbent was dried in an oven at 60°C. The dried magnetic nanoadsorbent was reused to start the next cycle of metronidazole adsorption. Base on the figure 17a, the metronidazole removal efficiency after six recycling cycles, reduced from 94% to 86%. On the other hand, the pollutant adsorption percentage from the first to last cycles demonstrated solely 12% reduction that can be because of the magnetic nanocomposite mass over the several cycles. Based on the obtained results
from the CoFe$_2$O$_4$@CMC/HZSM-5 regeneration, a slight decrease in the removal efficiency can be related to the occupation of active adsorbent sites by the metronidazole and decreasing in the amount of CoFe$_2$O$_4$@CMC/HZSM-5 during the washing and regeneration processes. Therefore, due to the high recyclability of CoFe$_2$O$_4$@CMC/HZSM-5, its application is economical in terms of operational costs. The CoFe$_2$O$_4$@CMC/HZSM-5 stability was investigated by FESEM (Fig. 17b) band XRD (Fig. 17c) analysis. Indicates the CoFe$_2$O$_4$@CMC/HZSM-5 XRD pattern after the sixth run of the magnetic nanoadsorbent recycling. The XRD analysis demonstrated that the CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanoadsorbent, did not have obvious changes except in the intensity of diffraction peaks following the sixth run of magnetic nanoadsorbent recycling. Also, the morphology of the magnetic nanoadsorbent has been preserved and did not have significantly change. The results showed the chemical stability of the magnetic nanoadsorbent after sixth run recycling.

(Figure 17)

3.6. Removal of metronidazole from real wastewater

The optimal conditions were determined. After that, the experiment was done on the campus wastewater at Kerman University of Medical Sciences. The real sample physicochemical characterizations were obtained and its specifications were achieved COD (415 mg L$^{-1}$), BOD (247 mg L$^{-1}$), TSS (20 mg L$^{-1}$), TDS (545 mg L$^{-1}$), TKN (78 mg L$^{-1}$), Phosphate (16.7 mg L$^{-1}$), Nitrate (16.5 mg L$^{-1}$), Sulfate (352 mg L$^{-1}$), and Metronidazole (45 mg L$^{-1}$). The metronidazole maximum removal efficiency was 85% in the adsorbent 2 g/L, pH 6, initial metronidazole concentration 45 mg/L and contact time 60 min. Due to the presence of cations, anions, TDS and TSS in the real wastewater, the metronidazole removal efficiency decreased.

The performance of CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanoadsorbent in the pharmaceutical compounds removal was compared with the other magnetic nanoadsorbent. It can be concluded that CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanoadsorbent in comparison with the other magnetic nanoadsorbent, had a higher pollutants removal efficiency in a shorter time, a lower dose of adsorbent and at the relatively neutral pH. Also, CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanoadsorbent had a high specific surface area and the higher adsorption capacity.

4. Conclusion

The results of the present study showed that the using magnetic nanoparticles with the aim of magnetic property induction in widely used adsorbents, especially CMC, is a simple, fast and practical method due to the lack of problems belong to separation, filtration and centrifugation. CMC plays an important role in improving the structural properties of CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanocomposite. The presence of CMC during the preparation of this nanocomposite, increased the adsorbent surface area and provided more active sites for the metronidazole removal. The optimal conditions in the present study were obtained at pH 6, contact time 60 min, adsorbent 2 g/L and temperature 20 °C. The metronidazole removal efficiency and maximum adsorption capacity based on the Langmuir
equilibrium model were obtained 94 and 206.6 mg/g, respectively. The adsorption efficiency increased by increasing contact time, the adsorbent amount and solution temperature, while the metronidazole initial concentrations had an indirect effect on the adsorption efficiency. Magnetized nanocomposite due to its high specific surface area can be used as an effective and cost-effective adsorbent to remove environmental pollutants from water sources. Therefore it is expected it use for quickly and easily separate different types of adsorbents from solutions. According to the obtained results, it could be concluded that the adsorption process was endothermic and had a physical mechanism. Finally, the chemical stability and reusability of the synthesized CoFe$_2$O$_4$@CMC/HZSM-5 magnetic nanoadsorbent were evaluated in the metronidazole removal. The achieved findings demonstrated that following six adsorption runs, he adsorbent chemical structure had no change and the magnetic nanoadsorbent efficiency in the removal process had a slight decrease.

**Credit Author Statement**

Alireza Nasiri: Project administration, Supervision, Mohammad Reza Heidari: Investigation, Writing original draft. Ghazal Yazdanpanah: Investigation, Validation, Writing, review & editing.

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Figure and Table Captions:

**Figure 1.** Schematic representation of CoFe₂O₄@CMC/HZSM-5 synthesis

**Figure 2.** The FT-IR spectrum of CMC, HZSM-5, and CoFe₂O₄@CMC/HZSM-5 magnetic nanoadsorbent

**Figure 3.** The FESEM (a, b) and TEM (c) images of CoFe₂O₄@CMC/HZSM-5 magnetic nanoadsorbent

**Figure 4.** The EDS of CoFe₂O₄@CMC/HZSM-5 magnetic nanoadsorbent

**Figure 5.** The Mapping of CoFe₂O₄@CMC/HZSM-5 magnetic nanoadsorbent

**Figure 6.** The XRD patterns of CMC, HZSM-5, CoFe₂O₄, and CoFe₂O₄@CMC/HZSM-5 magnetic nanoadsorbent

**Figure 7.** Magnetic hysteresis curve of CoFe₂O₄@CMC/HZSM-5 magnetic nanoadsorbent

**Figure 8.** The TGA analysis of CoFe₂O₄@CMC/HZSM-5 magnetic nanoadsorbent

**Figure 9.** The effect of pH changes on the metronidazole removal rate by MNZ = 50 mg/L, adsorbent dose = 2 g/L, T = 20 °C

**Figure 10.** The effect of contact time changes on the metronidazole removal rate by CoFe₂O₄@CMC/HZSM-5 (pH = 6, MNZ = 50 mg/L, adsorbent dose = 2 g/L, T = 20 °C)

**Figure 11.** (a) The effect of adsorbent value changes and (b) the effect of adsorbates concentration changes on the removal efficiency by CoFe₂O₄@CMC/HZSM-5 (pH = 6, T = 20 °C)

**Figure 12.** The kinetic model curves of the metronidazole adsorption process by CoFe₂O₄@CMC/HZSM-5 (pH = 6, MNZ = 50 mg/L, adsorbent dose = 2 g/L, T = 20 °C)

**Figure 13.** Metronidazole adsorption process equilibrium models by CoFe₂O₄@CMC/HZSM-5 (pH = 6, MNZ = 50 mg/L, adsorbent dose = 2 g/L, t = 60 min)

**Figure 14.** Metronidazole adsorption process Vant Hoff curve

**Figure 15.** Comparison of metronidazole adsorption by CoFe₂O₄, CMC, HZSM-5, and CoFe₂O₄@CMC/HZSM-5 adsorbents (pH = 6, MNZ = 50 mg/L, adsorbent dose = 2 g/L)

**Figure 16.** Proposed mechanism of metronidazole removal by CoFe₂O₄@CMC/HZSM-5

**Figure 17.** Recycling of CoFe₂O₄@CMC/HZSM-5 (pH = 6, MNZ = 50 mg/L, adsorbent dose = 2 g/L) (a), FESEM (b) and XRD pattern (c) of CoFe₂O₄@CMC/HZSM-5 magnetic nanoadsorbent after the sixth run of recycling

**Table 1.** Metronidazole removal kinetic parameters by CoFe₂O₄@CMC/HZSM-5

**Table 2.** Metronidazole adsorption process equilibrium isotherms of by CoFe₂O₄@CMC/HZSM-5

**Table 3.** The thermodynamic parameters of metronidazole adsorption process by CoFe₂O₄@CMC/HZSM-5
Figure 1

Schematic representation of CoFe2O4@CMC/HZSM-5 synthesis
Figure 2

The FT-IR spectrum of CMC, HZSM-5, and CoFe2O4@CMC/HZSM-5 magnetic nanoadsorbent.
Figure 3
The FESEM (a, b) and TEM (c) images of CoFe2O4@CMC/HZSM-5 magnetic nanoadsorbent

Figure 4
The EDS of CoFe2O4@CMC/HZSM-5 magnetic nanoadsorbent

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Figure 6

The XRD patterns of CMC, HZSM-5, CoFe2O4, and CoFe2O4@CMC/HZSM-5 magnetic nanoadsorbent
Figure 7

Magnetic hysteresis curve of CoFe2O4@CMC/HZSM-5 magnetic nanoadsorbent
Figure 8

The TGA analysis of CoFe2O4@CMC/HZSM-5 magnetic nanoadsorbent
Figure 9

The effect of pH changes on the metronidazole removal rate by MNZ = 50 mg/L, adsorbent dose =2 g/L, T=20 oC)
Figure 10

The effect of contact time changes on the metronidazole removal rate by CoFe2O4@CMC/HZSM-5 (pH=6, MNZ= 50 mg/L, adsorbent dose=2 g/L, T=20 oC)

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(a) The effect of adsorbent value changes and (b) the effect of adsorbates concentration changes on the removal efficiency by CoFe2O4@CMC/HZSM-5 (pH = 6, T = 20 °C)

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The kinetic model curves of the metronidazole adsorption process by CoFe2O4@CMC/HZSM-5 (pH = 6, MNZ = 50 mg/L, adsorbent dose = 2 g/L, T = 20 °C)
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Figure 14

Metronidazole adsorption process Vant Hoff curve

\[ y = -1.1223x + 5.5481 \]

\[ R^2 = 0.9941 \]
Figure 15

Comparison of metronidazole adsorption by CoFe2O4, CMC, HZSM-5, and CoFe2O4@CMC/HZSM-5 adsorbents (pH = 6, MNZ= 50 mg/L, adsorbent dose=2 g/L)
Figure 16

Proposed mechanism of metronidazole removal by CoFe2O4@CMC/HZSM-5
Recycling of CoFe2O4@CMC/HZSM-5 (pH = 6, MNZ= 50 mg/L, adsorbent dose=2 g/L) (a), FESEM (b) and XRD pattern (c) of CoFe2O4@CMC/HZSM-5 magnetic nano adsorbent after the sixth run of recycling