Tetracycline removal enhancement with Fe-saturated nanoporous montmorillonite in a tripartite adsorption/desorption/photo-Fenton degradation process

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Abstract
The adsorption and photo-Fenton degradation of tetracycline (TC) over Fe-saturated nanoporous montmorillonite was analyzed. The synthesized samples were characterized using XRD, FTIR, SEM, and XRF analysis, and the adsorption and desorption of TC onto these samples, as well as the antimicrobial activity of TC during these processes, were analyzed at different pH. Initially, a set of adsorption/desorption experiments was conducted, and surprisingly, up to 50% of TC adsorbed was released from Mt structure. Moreover, the desorbed TC had strong antibacterial activity. Then, an acid treatment (for the creation of nanoporous layers) and Fe saturation of the montmorillonite were applied to improve its adsorption and photocatalytic degradation properties over TC. Surprisingly, the desorption of TC from modified montmorillonite was still high up to 40% of adsorbed TC. However, simultaneous adsorption and photodegradation of TC were detected and almost no antimicrobial activity was detected after 180 min of visible light irradiation, which could be due to the photo-Fenton degradation of TC on the modified montmorillonite surface. In the porous structures of modified montmorillonite high, ‘OH radicals were created in the photo-Fenton reaction and were measured using the Coumarin technique. The ‘OH radicals help the degradation of TC as proposed in an oxidation process. Surprisingly, more than 90% of antimicrobial activity of the TC decreased under visible light (after 180 min) when desorbed from nanoporous Fe-saturated montmorillonite compared to natural montmorillonite. To the best of our knowledge, this is the first time that such a high TC desorption rate from an adsorbent with the least residual antimicrobial activity is reported which makes nanoporous Fe-saturated montmorillonite a perfect separation substance of TC from the environment.

Keywords Montmorillonite · Photo-Fenton · Nanoporous · Tetracycline · Adsorption · Desorption · Fe-saturated

Introduction
The release of active metabolite species of antibiotics in the ecosystem cycle and the prevalence of a new generation of antibiotic-resistant bacteria and genes have caused a global concern over their threats to mankind (Imanipoor et al. 2021; Liu et al. 2021a; Mosaleheh and Sarvi 2020; Wang et al. 2020a; Wu et al. 2018). Tetracycline (TC) is recognized as the second most widely consumed antibiotic, which is extensively distributed in the environment due to its high water solubility and low absorption rate by men, livestock, and poultry (i.e. about 10 to 40%) (Jiao et al. 2008; Norvill et al. 2017; Shen et al. 2020; Zhang et al. 2017). TC was considered a potential agent able to change and evolve genetic features of bacteria and thereby resulting in the emergence of a new generation of antibiotic-resistant bacteria (Ramazani Afarani et al. 2018; Verma and Haritash 2020). The successful application of TC for recently emerged viruses might result in an unexpected increase of TC in the environment, and consequently, new strategies have to be developed for its removal from the environment. Different techniques have been developed for the separation of antibiotics from the environment such as adsorption...
and catalysis degradation (Jain et al. 2020; Niu et al. 2021). Adsorption, as one of the most effective separation techniques, has been developed by many scientists, and numerous adsorption substances have been synthesized and modified for this purpose (Ahsan et al. 2018; Wang et al. 2021; Zhao et al. 2015). When looking at the adsorption technique as a practical method, the development of environmentally friendly adsorbents with the least residual environmental side effects, low secondary environmental pollution, high adsorption efficiency, and low-cost production becomes crucial (Mosaleheh and Sarvi 2020; Veiskarami et al. 2016). Besides, due to the abundance of antibiotics and possible residual antimicrobial activity of adsorbed antibiotics, providing a low release tendency or employing a degradation capability on the adsorbent to reduce the chance of secondary pollution has been the goal of many researchers (Bugueño-Carrasco et al. 2021; Lu et al. 2019a, b).

The degradation of bio-contaminants with a photocatalytic process has recently been converted to a promising method having high efficiency, high degradation rate, and the least side effects (Hassan et al. 2019; Huang et al. 2020; Li et al. 2021). Photocatalyst substance accelerates the decomposition of biomolecules through the production of reactive species (Thomas et al. 2021). Hence, the development of photocatalysts with the features mentioned above has become a controversial topic related to environmental problems. Hence, the development of a combined adsorption and photodegradation process could help to increase the efficiency of the separation method practically. The photocatalytic process is a promising process for the degradation of antibiotics, such as TC (Han et al. 2020; Tong et al. 2021; VM Starling et al. 2021; Wang et al. 2020b). This process, as an advanced oxidation process, is widely used for the treatment of antibiotics contaminated media (Ameta et al. 2018). This is one of the most environmentally friendly processes with the least secondary pollution and residual side effects (Wang et al. 2020b).

Montmorillonite has been used for the adsorption of different biomolecules (Akbari Alavijeh et al. 2017; Ambrogi et al. 2014; Imanipoor et al. 2021; Jayrajsinh et al. 2017). Moreover, it has such an extent potential to use as a barrier for halting the spread of pollutants in the environments due to its high cation exchange and adsorption capacity, eco-friendly nature, and non-toxicity (Carretero et al. 2013; Zhao et al. 2015). Nevertheless, modifying the montmorillonite-layered structure into a photocatalyst was proposed to decompose the adsorbed bio-components (Chen et al. 2009). Although montmorillonite was reported to have great adsorption characteristics, the release of the adsorbed antibiotics after saturation of the layered structure (Akbari Alavijeh et al. 2017; Ramazani Afarani et al. 2018) reduces the chance of using that for adsorption. In addition, its layered structure which prevents the penetration of light beams in the structures (Ambrogi et al. 2012) reduces its efficiency when used as a photocatalyst. These problems limit the application of montmorillonite and photocatalyst-loaded montmorillonite as a potential separation media for different antibiotics (Maroga Mboula et al. 2012).

In this research, initially, the role of montmorillonite as adsorbent of TC for the separation of TC from the environment was challenged. Then, a nanoporous structure was formed on the montmorillonite, and furthermore, its surface was modified to improve its mutual function for adsorption and photodegradation of the tetracycline. For this purpose, an acid treatment was applied to create nanopores onto its layered structure with more active sites for the adsorption as well as photoreaction. Besides, the layered structure of the montmorillonite was saturated with the Fe ions to improve the photo-Fenton degradation process of the adsorbed TC. The montmorillonite-layered structure is used to support the Fe ions and help the degradation of TC which is the host of adsorbed TC. In the separation process, the adsorption and desorption of TC from montmorillonite were characterized and the photodegradation after the adsorption/desorption cycle was analyzed.

Materials and methods

Materials

Bentonite as a source of montmorillonite was purchased from a bentonite mine located in Salafchegan, Iran with a CEC of 73 mEq/100 gr. Tetracycline hydrochloride (> 98%, Sigma-Aldrich), hydrochloric acid (HCl, 37% w/w, Merck), and hydrogen peroxide (H2O2, 30% w/w, Merck) were used as received. Coumarin was provided by Sigma-Aldrich and used as received. Deionized water was used for all experiments. A visible halogen light lamp was used as the light source for photocatalyst experiments.

Montmorillonite modification

The particles of montmorillonite smaller than 2.5 microns were separated using centrifuge force (Veiskarami et al. 2016). Typically, a 3% (w/v) solution of bentonite in water was prepared and mixed overnight at 25 °C. Afterward, the obtained suspension was centrifuged at 1000 rpm, 25 °C for 260 s. The supernatant was collected and dried at 60 °C overnight and named Mt. To prepare a nanoporous montmorillonite sample, the Mt sample was then reacted with 1 M HCl at boiling temperature for 60 min. After completing the acid treatment, the mixture was centrifuged at 5000 rpm, 25 °C for 5 min. The precipitates were washed with plenty of water several times, dried at 60 °C overnight, and named NP-Mt.
The two montmorillonite samples (Mt and NP-Mt) were then saturated with Fe$^{3+}$ (Hong et al. 2019). For this purpose, the montmorillonite samples were added to a 0.1 M FeCl$_3$ solution for 2 h, and this process was repeated several times. An excess amount of iron was washed with water, and then, the samples were then separated using centrifuge force and dried in an oven at 60 °C overnight. All samples were kept in a desiccator containing silica gel before use.

Kinetic of adsorption and desorption of TC

Initially, the adsorption and desorption characteristics of different montmorillonite samples were analyzed and it was proved that the montmorillonite in all forms (natural or modified) releases the desorbed TC to the solution with the least changes in its antimicrobial activity characteristics. For this purpose, the kinetic adsorption experiments were carried out in 12 intervals of 1, 2, 4, 8, 15, 30, 60, 90, 120, 180, 240, and 400 min at 25 °C and a dark condition. Typically, 100 mg of montmorillonite sample was mixed with 20 mL of TC solution (1 mg/mL) at three different pH of 5, 7, and 9 (the reason why such high TC concentration was used is described in the supporting information section). Afterward, the mixture was centrifuged at 5000 rpm, 25 °C for 7 min, and the TC concentration of the supernatant was measured using a UV–Visible spectrophotometer at wavelength 276 nm. Finally, the amount of adsorbed TC was calculated based on the mass balance, according to a prepared standard curve.

Desorption kinetic experiments were designed to investigate the amount of TC desorption after adsorption on montmorillonite samples. Typically, after adsorption, the solution was centrifuged and the precipitates (montmorillonite-TC hybrid) were added to a rotating 500 mL glass container containing 250 mL of the receptor solution (with the same pH of adsorption) and covered with aluminum foil (the temperature was set at 25 °C using a water bath). The amount of desorbed TC was monitored by measuring the concentration of the balloon solution at different intervals (9, 18, 27, 36, 45, 54, 63, 72, 81, 90, 99, 108, 117, 126, 150, 174, 198, and 225 min) using a UV–Visible spectrophotometer at a wavelength of 276 nm. The test vessel was refilled with the equivalent amount of sample taken out with the fresh solution.

Tetracycline photo-Fenton degradation

In this stage, the montmorillonite surfaces were used as a suitable place for photodegradation of TC. The photo-induced changes in TC by the montmorillonite samples were then studied. Generally, all the experiments were performed using a halogen light lamp, which was located at a 20 cm distance from samples in a quartz container, and the temperature was set at 25 °C using a refrigerated incubator. For the photo-Fenton reaction, hydrogen peroxide was added to the mixture before the light irradiation. For each montmorillonite sample, simultaneously, two similar series of adsorption tests, as aforementioned, were conducted. Typically, 20 mL of 1 mg/mL TC solution was added to 100 mg of montmorillonite sample, and the mixture was blended at 25 °C for 8 h (according to adsorption kinetic results) in a dark condition to reach the maximum adsorption. The mixture was then centrifuged and the precipitates were converted to a quartz container containing 250 mL of black solution with the same pH of adsorption test. One set of samples was then exposed to irradiation under visible light for three durations of 60, 120, and 180 min while shaking and the other set of the sample were converted to the same solution and shaken in dark. Finally, the antimicrobial activity of the solution was tested for all samples.

Antibacterial test

The antibacterial activity of TC was evaluated using the agar-disk diffusion method. Muller Hinton agar plates were inoculated with $1.5 \times 10^{-8}$ CFU/mL inoculum of Escherichia coli. Then, filter paper disks, containing the 300 μL of the TC solutions, which were filtered by a syringe filter (0.22-micron), were placed on the agar surface. The Petri dishes were incubated at 37 °C for 24 h. Finally, the inhibition growth zones were analyzed. For all samples, two sets of the test were done, one is dark and the other under light irradiation, and in both, all chemicals were kept the same to characterize the photodegradation characteristics of the montmorillonite over TC.

Characterization

A sunlight simulator lamp (Philips 150 W) with a UV cut-off filter ($\lambda > 420$ nm) was used as a visible-light source in all photocatalytic experiments. Low angle XRD analysis was done in 2θ ranging from 4 to 10 degrees using Rigaku D/max-1200 with Cu Kα irradiation. Ultraviolet–visible spectroscopy liquid analysis was performed by the Rey-light-1600 UV–Visible spectrophotometer to measure the TC concentration in the solution. Scanning electron microscopy (SEM) analysis was used to characterize the morphology of the montmorillonite after treatment using a Zeiss Sigma model (samples were gold-sputtered before analysis). FTIR analysis was conducted using a tensor27 Bruker Fourier spectrometer. Also, the chemical composition of the solid material was determined by Bruker: S4PIONEER X-ray fluorescence (XRF) spectrometer. The nitrogen sorption analysis was done using a Bel sorp-mini II Japan, at 77 °K (the samples were initially degassed for 5 h at 150 °C). The coumarin method was used to
evaluate the potential for \( \cdot \text{OH} \) radical production (Czili and Horváth 2008; Louit et al. 2005; Maezono et al. 2011). In a typical analysis, 2 mL of coumarin solution (2 mM) was added to the 200 µl sample solution. To ensure almost complete trapping of the \( \cdot \text{OH} \) radical by coumarin, more coumarin solution was added. The \( \cdot \text{OH} \) radicals react with the coumarin molecules, resulting in a highly luminescent 7-hydroxy coumarin. The cell was turned into a detection spectrophotometer equipped with a fluorescent lamp (7 watts near-UV fluorescent lamp). The fluorescent intensity was measured with a spectrophotometer at a wavelength of 470 nm.

**Results and discussion**

Figure 1 shows the nitrogen sorption results of different montmorillonite samples. All isotherms correspond to H3 hysteresis adsorption associated with capillary condensation in nanopore structures (Sing 1985). The type H3 is normally corresponding to plate-like particles which cause slit-shaped pores (Maged et al. 2020). After acid treatment of montmorillonite, an increase in the surface area of montmorillonite samples was observed (from 28 up to 92 cm\(^2\)/g, Table 1), indicating an increase in the porous structures on the montmorillonite layers. This is due to the destruction of the octahedral structure of the montmorillonite (Bieseki et al. 2013; Franco et al. 2016). Besides, the pore volume increased hugely after acid treatment, indicating the formation of nanopores (Table 1). The Fe saturation hardly affected the nitrogen sorption results of the montmorillonite or the acid-treated one. The Fe is enriched on the surfaces of the montmorillonite; however, it is proposed that the Fe is saturated in the removed octahedral sheets of the nanoporous montmorillonite. After acid treatment, the porosity of the montmorillonite (such as specific surface area and pore volume) was increased. Besides, according to XRF analysis (Table 2), most cations (Ca\(^{2+}\) and Na\(^{+}\)) were leached out from the structure. In addition, the amount of Mg and Fe was slightly reduced, indicating the effect of acid treatment on the montmorillonite.

The SEM images of samples confirm the layered morphology of montmorillonite and the acid-treated ones (Fig. 2). Cracked aggregates, segmented and flaky pieces, and reduced particle size are some of the noteworthy signs of changes in the montmorillonite structure via acid treatment. On both Mt and NP-Mt samples, no significant changes in the structure of montmorillonite occurred after Fe saturation.

FTIR analysis was performed to investigate the changes made to montmorillonite structure through acid treatment (Fig. 3). After acid treatment (to provide nanoporous montmorillonite), the intensity of bands at 835, 875, and 914 cm\(^{-1}\) decreased, which are corresponding to the bending vibration of AlMgOH, AlFeOH, and Al\(_2\)OH, respectively (Franco et al. 2016). This could be due to the leaching of the cations of octahedral sheets by order of Al, Mg, and Fe. In addition to cation removal, and consequently their hydroxyl groups (3620 cm\(^{-1}\)), some changes were detected in the intensity of the bands at 1113, 1035, 792, 610, 522, 461, 3620 cm\(^{-1}\).

### Table 1 Results of nitrogen sorption analysis

| Sample | BET surface area (m\(^2\)/g) | Pore size (nm) | Total pore volume (cm\(^3\)/g) |
|--------|-------------------------------|----------------|-----------------------------|
| Mt     | 28                            | -              | 0.12                        |
| Mt-Fe  | 26                            | -              | 0.11                        |
| NP-Mt  | 92                            | 1.5            | 0.40                        |
| NP-Mt-Fe | 89                           | 1.5            | 0.46                        |

### Table 2 XRF analysis of montmorillonite samples

| Sample  | Mass (%) | SiO\(_2\) | Al\(_2\)O\(_3\) | CaO | Fe\(_2\)O\(_3\) | K\(_2\)O | MgO | TiO\(_2\) | Na\(_2\)O | LOI |
|---------|----------|-----------|----------------|-----|----------------|--------|-----|-----------|----------|-----|
| Mt      | 54.5     | 16.4      | 1.9            | 4.1 | 0.4            | 3.7    | 0.4 | 0.6       | 18.0     |     |
| Mt-Fe   | 51.2     | 13.5      | 1.9            | 11.1| 0.3            | 3.5    | 0.4 | 0.5       | 17.6     |     |
| NP-Mt   | 60.8     | 13.3      | 0.2            | 2.1 | 0.5            | 2.1    | 0.4 | 0.1       | 20.5     |     |
| NP-Mt-Fe| 56.7     | 10.6      | 0.2            | 11.8| 0.5            | 2.2    | 0.2 | 0.1       | 17.7     |     |
and 462 cm⁻¹ (Fig. 3). These decreases showed the acid has influenced tetrahedral and octahedral sheets (Si–O–Si, Al–O–Si), and Si–O bands were decomposed partially (Madejová 2003; Maged et al. 2020). Comparing the spectrum of montmorillonite with Fe-saturated samples, a new peak emerged at 700 cm⁻¹, which represents the FeOOH bending vibration. This peak suggested that Fe ions are hydrolyzed into the interlayer spaces of the montmorillonite by pillaring (Yuan et al. 2008).

The kinetics of adsorption and desorption of TC onto different montmorillonite samples were plotted in Fig. 4 as well as the results of two kinetic models (pseudo-second-order and intraparticle diffusion models) fitted to the data (details of kinetic models are presented in Table 3). For all samples, the equilibrium of adsorption was reached after 60 min, and the Mt and Mt-Fe samples adsorbed more TC than the nanoporous montmorillonite samples to some extent at all pH which is due to the partial destruction of the nanoporous montmorillonite after acid treatment and lower cation exchanges.

The pseudo-second-order model fitted to all data (Fig. 4 and Table 3) indicated a similar trend and the adsorption capacity was reduced by increasing the pH. A higher amount of TC adsorbed at lower pH (5 and slightly at 7)
which is due to the higher positive charge of TC at this condition (Figueroa et al. 2004) and resulted in a facile cation exchange mechanism (Akbari Alavijeh et al. 2017; Ramazani Afarani et al. 2018). By increasing the pH to 9, higher negative charges of TC resulted in the lower tendency in the adsorption onto the montmorillonite. In addition, lower interlayer adsorption was detected at this condition.

The intraparticle diffusion model results indicated that the adsorption of TC on the external surfaces of all montmorillonite particles was almost the same at all pH of the solution (Fig. 4 and Table 3 the first linear region of the plots). At acidic and neutral pH, the intraparticle diffusion was another important mechanism of adsorption for all samples which implies the adsorption of TC in the interlayer spaces which will be discussed in the XRD results as well. However, by
increasing the pH of adsorption, the intraparticle diffusion was reduced for nanoporous samples (acid-treated ones) indicating a lower tendency of TC to be diffused in the interlayer spaces due to the higher negative charges of the TC and lower chance of cation exchange mechanism with montmorillonite interlayer cations (Chahardahmasoumi et al. 2019).

The changes in the layered structure of montmorillonite after treatments and adsorption of TC were analyzed using XRD analysis (Fig. 5). After acid treatment, the basal spacing of the montmorillonite was increased from 11.6 to 13 Å which could be due to the destruction of the octahedral structure of the montmorillonite upon acid treatment (Temuujin et al. 2004). The saturation of both montmorillonite and nanoporous montmorillonite samples with Fe ion did not affect the basal spacing very much which is following what was reported previously (Hong et al. 2019; Zhao et al. 2020). The XRD results indicated that at two acidic and neutral pH, the TC was adsorbed mainly on the interlayer spaces of all montmorillonite samples (the basal spacing was increased for 6 to 7 Å, Fig. 5). However, at pH 9 for two samples that were not acid-treated, no changes in the interlayer spaces were detected showing no adsorption of TC between interlayer spaces, while for the acid-treated (nanoporous montmorillonite samples) at pH of 9, a twin peak was detected showing mutual adsorption of TC in the interlayer spaces and on the external surfaces of the samples (this trend is in accordance with the intraparticle diffusion models, Fig. 4 and Table 3). The reason for adsorption of TC in the interlayer spaces of the nanoporous montmorillonite at pH 9 is the increase in the cation charges (H+) in the interlayer spaces created during the acid treatment which allows TC to diffuse in the interlayer spaces and increase the basal spacing (TC is negatively charged at this pH) (Chahardahmasoumi et al. 2019). Enrichment of the montmorillonite with Fe ion did not change the trend of TC adsorption on the external surface or into the basal spacing of different samples (Fig. 5).

Surprisingly, more than 30% of adsorbed TC was desorbed from all samples with a linear trend at all pH conditions (Fig. 4), showing that the montmorillonite (natural or nanoporous one) could not fully immobilize TC in its structure and release it back to the environment in high amount. Higher amounts of TC were desorbed from the montmorillonite samples at a pH of 9 (Fig. 4) indicating more repulsion forces between the montmorillonite and the TC. The observed desorption of TC from montmorillonite samples was somehow expected as reported previously, however, compared to the desorption rate from agricultural soil (Conde-Cid et al. 2020), such a high amount of desorption was contrary to expectation (Fig. 4). This phenomenon is partly due to the different nature of the adsorbents and partly due to the extremely low concentration of TC in the solution (in the range of 0.003 to 0.2 mg/mL) for adsorption onto the agricultural soil (Conde-Cid et al. 2019). However, by increasing the TC concentration in the solution, the amount of TC adsorbed was increased (Fig. 4) and the desorption rate onto the montmorillonite samples increased as well, which is a realistic prediction of the empirical situation. Hence, as the montmorillonite might reach its maximum adsorption capacity, it finally releases the adsorbed TC to the environment. This could be a big challenge in the use of such environmentally friendly and cheap adsorbents for adsorption/separation of antibiotics such as TC. One possible remediation for such a problem is to apply a second capability on the montmorillonite to degrade the TC in the adsorption–desorption cycle. Hence, the desorbed pollutants do not harm the environment upon desorption. A possible remediation process is to apply a photo-Fenton enhanced surface for photodegradation of TC. The Fe ions were loaded on the montmorillonite for this purpose and the properties
of montmorillonite on the degradation of TC during the desorption process were analyzed.

To check the photocatalytic properties of montmorillonite samples, after adsorption of the TC, the montmorillonite was separated and put in a solution of the same pH and the light irradiation was applied under shaking. Then, the antibacterial activity test on the solution was done in different duration and the results were compared with the results of the antibacterial activity of the same sample in which the adsorption–desorption cycle was carried out in dark. Antibacterial activity results and the inhibition zone summary results of released TC from montmorillonite samples were plotted in Figs. 6 and 7, respectively. The adsorption characteristics of the montmorillonite were employed to adsorb as much as TC on the montmorillonite surface and provide a suitable place for the photodegradation process.

For the Mt and NP-Mt sample, almost no changes in the antimicrobial activity of the TC were detected after irradiation under light for up to 180 min at all pH compared to samples in dark (the inhibition zone in Figs. 6a1, b1, c1, a3, b3, c3, and 7). By increasing the desorption duration, the inhibition zone increased (Fig. 7) up to 120 min due to the increasing amount of TC desorbed and after that almost remained constant. Comparing the samples under light and in dark (comparing disk 1 which was under light irradiation with 4 which has the same chemicals as disk 1
and just was carried out in dark, 2 with 5, and 3 with 6 in Fig. 6), the inhibition zone did not vary very much when placed under light for these two montmorillonite samples. Hence, the main conclusion is that the TC is desorbed from the montmorillonite and reasonably strong antimicrobial activity was detected even after being under light for 180 min. Considering the application of such adsorbent in the environment, montmorillonite does not perfectly separate the TC from the soil and release it back to the environment with almost no changes.

On the other hand, for the Fe-saturated samples, the antibacterial activity of TC was reduced significantly even after 60 min of light irradiation (Figs. 6a2, b2, c2, a4, b4, c4, and 7). The reduction in the antimicrobial activity of TC was higher for the nanoporous Fe-saturated montmorillonite than the normal Fe-saturated montmorillonite. The main reason could be the higher accessible site for the adsorption and possible photodegradation process. Indeed, the higher transparent nanoporous structure provides a higher chance for interlayer adsorbed TC to absorb light and be degraded. By increasing the light irradiation time from 60 to 180 min, the antimicrobial activity reduced steadily for Fe-saturated samples compared to the same samples in dark. Therefore, during the desorption, the degradation of the TC occurs for Fe-saturated samples and this process continues in an equilibrium adsorption–desorption–photodegradation cycle. Meanwhile, the degradation of TC
was facilitated at acidic conditions due to the higher amount of TC desorbed from the montmorillonite samples (Fig. 4) and a lower chance of degradation.

It is proposed that the photodegradation happens through a photo-Fenton process in which the Fe cations seemingly played a key role in the photo-Fenton deactivation of TC (Wang et al. 2020b). Fe cations degraded the TC by the production of active species via Fenton-type reaction. The production of \( \cdot \)OH radicals is the key element in the degradation of the TC via creating complexation (Brillas and Garcia-Segura 2020; Fareed et al. 2021; Liu et al. 2021b). Although the mechanism of the \( \text{H}_2\text{O}_2 \) decomposition in the photo-Fenton reaction is not well understood, the following reactions have been suggested corresponding the reduction of \( \text{Fe}^{3+} \) with the production of \( \cdot \)HO\(_2\) radicals, followed by regeneration of \( \text{Fe}^{3+} \) which resulted in the generation of the hydroxyl radicals (Ramirez et al. 2007):

\[
X - \text{Fe}^{3+} + \text{H}_2\text{O}_2 \rightarrow X - \text{Fe}^{2+} + \cdot \text{OH}_2 + \text{H}^+ \tag{1}
\]

\[
X - \text{Fe}^{2+} + \text{H}_2\text{O}_2 \rightarrow X - \text{Fe}^{3+} + \text{OH}^- + \cdot \text{OH} \tag{2}
\]

\[
\text{TC} + \cdot \text{OH} \rightarrow \text{degraded TC} \tag{3}
\]

in which the \( X \) represents the surface of the montmorillonite surface. The concentration of \( \cdot \)OH radicals was evaluated using the coumarin method in which the produced \( \cdot \)OH radicals would react with the coumarin and produce

\[
\begin{array}{c}
\text{Fluorescence spectra of the sample taken from the solution containing different montmorillonite samples after being under irradiation under UV for 5 min}
\end{array}
\]
7-hydroxycoumarin that is highly fluorescent (Louit et al. 2005) and can be detected with a spectrophotometer (Fig. 8). The proposed reactions take place on the montmorillonite surface when enriched with the Fe cations (Scheme 1). Based on the proposed mechanism, the Fe saturation of the non-porous montmorillonite mainly occurs on the external surfaces (Scheme 1). The TC adsorbed on the surfaces of the montmorillonite would be in a part of the Fenton reaction in which the created ˙OH radicals (Fig. 7) help to degrade TC. However, for the nanoporous montmorillonite, the nanopores (Scheme 1), giving rise to a protonated (H+) amorphous hydrous silica, provide a high accessible surface area for adsorption of Fe cations (Wei et al. 2017). In this structure, the absorption of light would be much higher than the non-porous montmorillonite. Hence, a great opportunity for the production of ˙OH radicals was proved by the fluorescence spectra (Fig. 8), and hence, the photo-Fenton reactions would occur.

The amount of iron leached out from the montmorillonite samples was characterized using atomic absorption. Initially, the amount of iron loaded on the montmorillonite was measured, and then, the amount of iron in the solution was measured and the released iron was calculated by mass balance. The results indicated that the amount of iron resealed was very much neglectable (below 5%) and it is higher in acidic conditions.

Conclusion

In this study, we proposed a modification method to convert montmorillonite into Fe-saturated nanoporous layered structure with high affinity in adsorption and photo-Fenton degradation properties against tetracycline. Via an acid-treatment process, the surface area of the montmorillonite was increased four times and a nanoporous structure with pore sizes around 1.5 nm was created. The nanoporous Fe-saturated montmorillonite showed a high affinity for the adsorption of TC. The antibacterial tests indicated that more than 30% of adsorbed TC was released from the normal montmorillonite with high antibacterial activity. Enrichment of the nanoporous montmorillonite with Fe ions increased its photocatalytic activity (photo-Fenton properties) and almost more than 90% of desorbed TC was degraded under visible light. The nanoporous structure of the montmorillonite provided a higher surface area for photodegradation of the TC compared to normal Fe-enriched montmorillonite by a higher amount of ˙OH radicals created in the nanoporous structure which helps the degradation of TC. The results of this study provided a novel environmentally friendly adsorbent with photocatalytic activity against TC which can be of interest to many scientists in this field.
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Author contribution All authors contributed to the study conception and design. Material preparation: [Mehdi Nasiri Sarvi], conceptualization: [Shiva Chahardahmasoumi], data collection and analysis: [Seyed Amir Hossein Jalali], methodology: [Shiva Chahardahmasoumi], writing—original draft preparation: [Shiva Chahardahmasoumi], supervision: [Seyed Amir Hossein Jalali], visualization: [Seyed Amir Hossein Jalali], validation: [Mehdi Nasiri Sarvi], reviewing and editing: [Mehdi Nasiri Sarvi], project administration: [Mehdi Nasiri Sarvi]. All authors read and approved the final manuscript. All authors whose names appear on the submission (1) made substantial contributions to the conception or design of the work; or the acquisition, analysis, or interpretation of data; or the creation of new software used in the work; (2) drafted the work and revised it critically for important intellectual content; (3) approved the version to be published; and (4) agreed to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Availability of data and materials All data are available on request.

Declarations

Consent to publish All the authors of this manuscript have agreed to its submission and we confirm that the work has not been submitted simultaneously to another journal for consideration.

Competing interests The authors declare no competing interests.

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