**TiO₂-Doped Chitosan Microspheres Supported on Cellulose Acetate Fibers for Adsorption and Photocatalytic Degradation of Methyl Orange**

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**Abstract:** Chitosan/cellulose acetate (CS/CA) used as a biopolymer system, with the addition of TiO₂ as photocatalyst (C-T/CA) were fabricated by alternating electrospinning/electrospraying technology. The uniform dispersion of TiO₂ and its recovery after the removal of methyl orange (MO) was achieved by incorporating TiO₂ in CS electrosprayed hemispheres. The effects of pH values, contact time, and the amount of TiO₂ on adsorption and photocatalytic degradation for MO of the C-T/CA were investigated in detail. When TiO₂ content was 3 wt %, the highest MO removal amount for fiber membranes (C-T-3/CA) reached 98% at pH value 4 and MO concentration of 40 mg/L. According to the data analysis, the pseudo-second-order kinetic and Freundlich isotherm model were well fitted to kinetic and equilibrium data of MO removal. Especially for C-T-3/CA, the fiber membrane exhibited multiple layers of adsorption. All these results indicated that adsorption caused by electrostatic interaction and photocatalytic degradation were involved in the MO removal process. This work provides a potential method for developing a novel photocatalyst with excellent catalytic activity, adsorbing capability and recycling use.

**Keywords:** TiO₂; chitosan; electrospinning; electrospraying; methyl orange; photocatalytic degradation

1. Introduction

With the quick progress of industrial technology, water pollution generated from organic dye contaminants has become an increasing global environmental concern in recent years [1–4]. Organic dyes are mainly generated from manufacturing, such as textile finishing, plastics, paper, cosmetics, pharmaceuticals and food processing. It can cause serious problems for both the environment and to human beings [5–10]. Up until now, several methods such as chemical oxidation, coagulation, adsorption, biodegradation, photocatalysis, and reverse osmosis, have been developed to treat dye contaminated wastewater [11–14]. Particularly, photocatalysis is considered a “green” treatment method which uses solar energy effectively for the degradation of organic pollutants [15–19]. Adsorption is also considered a valid method for the removal of dye in aqueous solutions with high efficiency, low price and easy operation [20–22].

At present, researchers have found that the combination of adsorption and photocatalysis had excellent potential applications for dye treatment [23]. TiO₂ is a type of common semiconductor photocatalyst on account of its prominent characteristics of chemical resistance, excellent catalytic activity, mechanical robustness, and low cost [24–28]. While the low dispersibility, inconvenient recovery, low adsorption performance of TiO₂ limit its use in photocatalysis applications [29]. Absorbent
materials with good adsorption performance on organic pollutants are often used as carrier materials to improve the photocatalysis capability of TiO$_2$.

Ideal carrier materials should provide stable anchoring of TiO$_2$, outstanding structure stability, selective adsorption for target dye pollutions and the transfer of organic dye molecules to the vicinity of the photocatalytic site. Multifarious materials have been developed for photocatalytic degradation carrier materials, comprising silica, ceramic, polymers, activated carbon, zeolite, and glass. As an environmentally friendly, nontoxic, biodegradable, biocompatible natural macromolecule [30–34], CS with functional groups of hydroxyl and amino groups derived from deacetylation of polysaccharide chitin, is a promising carrier material [35–38]. Li et al. produced pure CS membranes by electrospinning, exhibiting high adsorption capacity for acid blue-113 [37]. Huang et al. produced an organoclay/CS hybrid system, allowing efficient and concurrent adsorption of different dyes [38]. However, CS is easily soluble in water when amino groups are protonated, making it difficult to maintain integrity of the whole materials. As a hydrophilic, environmentally friendly, biodegradable, and renewable material [39–43], CA has characteristics of non-specific adsorption, and its hydroxyl and ester groups can form stable hydrogen bonding with certain polar functional groups [44,45]. Electrospinning technology is a common and effective method for the preparation of adsorbents [14]. Doh et al. electrospun TiO$_2$ nanofibers had a higher degradation rate of organic dye than TiO$_2$ nanoparticles because of a high specific surface area and excellent dispersibility [4]. Razzaz et al. proved that CS/TiO$_2$ electrospun membranes had higher effective surface area, structural stability and adsorption capacity for metal ions compared with TiO$_2$-coated CS nanofibers [46].

In this study, based on CA electrospun fibers, C-T/CA fiber membranes were successfully prepared via electrospaying CS/TiO$_2$ concave hemispheres on the surface of CA fibers. As a substrate material, CA was expected to improve the water flux of fiber membranes effectively so that aqueous solutions of organic pollutant, MO could rapidly penetrate into the interior of membranes. The CS hemispheres might disperse TiO$_2$ nanoparticles, have good adsorption performance of MO, and transfer MO to the vicinity of TiO$_2$ active sites for further photocatalytic degradation. The morphology of fiber membranes were characterized by SEM. Effects of TiO$_2$ content, pH values, contact time at different MO concentration for adsorption and photocatalytic degradation of MO were investigated. The MO removal mechanism was surveyed by FTIR, XPS and BET.

2. Experiment

2.1. Materials

CS (90% deacetylated, average $M_w = 200,000$ Da), TiO$_2$ nanoparticles (anatase-type nanopowder, particle size $< 25$ nm, metal basis), CA (white powder, $M_n = 30,000$ Da, acetyl content $\approx 39.7$ wt %, DS $= 2.45$) and MO were all purchased from Aladdin Industrial Corporation, Shanghai, China. Acetic acid (HAc) and acetone were purchased from Beijing Chemical Works, Beijing, China. Deionized water was obtained from the laboratory. The chemicals were all of analytical grade and used without further purification.

2.2. Preparation of Polymer Solutions

10 wt % CA ($w/w$) was dissolved in acetone/deionized water ($85/15$, $w/w$) and 2 wt % CS ($w/w$) was dissolved in 90% acetic acid ($v/v$), respectively. The solutions were stirred for 24 h at room temperature to obtain the uniform solutions. Then, the various contents of TiO$_2$ (1%, 2%, 3%, 4%, 5%, $w/w$) were added to CS solutions at room temperature and stirred for 24 h. The C-T-X solutions were obtained, where X stands for the content of TiO$_2$ (1, 2, 3, 4, 5 wt %). The solutions were placed in an ultrasonic water bath for 30 min before using. The frequency of ultrasonication was 25 kHz.
2.3. Fabrication of C-T/CA Fiber Membranes

The fabrication processes including (I) electrospinning and (II) electrospraying are shown in Figure 1. The solution was loaded into a 5 mL plastic syringe. The electrospinning conditions included the applied voltage of 20 kV, the 15 cm distance between the tip of the needle and the collector, and the flow rate of 1.0 mL/h. The condition for electrospraying was 18 kV voltage with a distance of 15 cm between the tip of the needle and the collector, and the flow rate was 0.3 mL/h. Then the membrane was dried at 25 °C for 24 h, and peeled off aluminum foil for further study.

![Figure 1. The preparation of C-T/CA fiber membranes: (I) Electrospinning, and (II) electrospraying.](image)

2.4. Characterizations

The viscosity of precursor solutions was measured by a viscosimeter (NDJ-1, Yutong, Shanghai, China). The electrical conductivity of the solutions was measured using a conductometer (DDS-11A, Shengci, Shanghai, China). All measurements were taken 3 times at room temperature. The morphology and structure of membranes were characterized by SEM (JSM-6700F, JEOL, Tokyo, Japan). All samples were sputtered with platinum under vacuum before assessment. The average microsphere diameters were measured by Image J software. Surface area, pore volume and pore size were characterized by N₂ adsorption and desorption using a Brunauer–Emmett–Teller analyzer (BET, Autosorb-iQ2, Quantachrome Instruments, Shanghai, China). The structures of the composite fibers were analyzed by FTIR spectroscopy (FTIR-4100, Jasco, Shanghai, China), within the wavenumber ranging from 400 to 4000 cm⁻¹ and a wide angle X-ray diffractometer (XRD, D/Max, Rigaku, Beijing, China), under mode of 40 kV and 50 mA, λ = 1.5406 Å radiation using copper-K alpha. The samples were scanned in the 2θ range from 5° to 60° with a scanning rate of 30 min⁻¹. The chemical composition of membrane surfaces were analyzed by XPS (ESCALab220i-XL, VG Scientific, Waltham, MA, USA) with Al Kα X-ray source, Ni-filtered radiation, 40 kV of tube electric pressure and 30 mA of tube electric current.

2.5. Investigations on MO Adsorption and Photocatalysis Study

Organic dye MO was used as the adsorbate for adsorption and photocatalysis studies of fiber mats. 20 mg of fiber mats were immersed in 40 mg/L MO aqueous solution (20 mL). The solution with fiber mats was stirred in dark conditions for 1 h to reach the adsorption equilibrium. After that, the mix solution was irradiated for 30 min in a photodegradation reactor under UV light (wavelength of 365 nm) by mercury lamp (25 W). The mixed solution (6 mL) was taken out every 5 min and centrifuged. The centrifugation was used to separate the solid particles in the suspension from the liquid to obtain a supernatant for subsequent absorbance detection. The UV spectrophotometer of the residual MO content was recorded. The process was repeated six times. The other set of photocatalytic experiments
were performed under visible light (wavelength of 600 nm) using a 350 W xenon lamp. The operation was the same as the process using UV light. The concentration of the residual MO was recorded every 20 min for 120 min. The adsorption and photocatalytic degradation R (%) of membranes to MO was calculated by the Equation (1):

$$ R(\%) = \frac{A_0 - A_t}{A_0} \times 100\% $$

(1)

where $A_0$ and $A_t$ are the original and equilibrium absorbances of the MO solutions.

The batch experiments of MO onto the fiber membranes were carried out as functions of TiO$_2$ concentration (0–5 wt %), pH (4–9), contact time (0–85 min), and initial concentration (10–80 mg/L) at room temperature. The adsorption and photocatalysis ability was computed as follows:

$$ q_t = \frac{C_0 - C_i}{m} \times V $$

(2)

where $q_t$ (mg/g) represented the removal amount at time $t$, $C_0$ (mg/L) and $C_i$ (mg/L) are the original and equilibrium concentrations of the MO solutions, $V$ (L) is the volume of MO solutions, $m$ (g) is the quantity of fiber membrane.

In the reusability experiment, the used adsorbent was washed by deionized water. Then the supernatant of the solution, after the reaction was replaced by fresh MO solution. The photodegradation reaction was carried out under UV illumination with magnetic stirring for 1 h. This experiment was cycled five times.

3. Results and Discussion

3.1. Morphology Observation

The morphology and diameter distributions of the CA substrate are displayed in Figure 2. The surfaces of the CA fibers were smooth without beads. The mean diameter of CA fibers was 0.55 ± 0.24 μm (shown in Figure 2b).

![Figure 2. (a) The morphology, and (b) the diameter distribution of the CA fibers.](image)

The morphology and the diameter distributions of the CS microspheres with different additions of TiO$_2$ are shown in Figure 3. Pure CS showed concave hemisphere morphology as shown in Figure 3a. With the content of 1, 2, 3 wt % TiO$_2$, CS still maintained the concave hemisphere morphology, and TiO$_2$ nanoparticles were evenly distributed on the surface of CS hemispheres (see Figure 3b–d). When TiO$_2$ content was 4 wt % and 5 wt %, TiO$_2$ nanoparticles begun to accumulate, and the original uniform morphology of CS hemispheres gradually disappeared (Figure 3e,f).

The particle size of pure CS hemispheres was 1.07 ± 0.24 μm. With the increase of TiO$_2$ content from 1 wt % to 3 wt % (Figure 3b1–d1), the diameters of CS hemispheres were 1.35 ± 0.26 μm,
1.40 ± 0.28 μm and 1.46 ± 0.34 μm, showing an upward trend. This was because the increase of viscosity was more important than the conductivity of the electrosprayed solution (Table 1). As the content of TiO₂ continued to increase, the diameters of CS hemispheres were 1.10 ± 0.24 μm and 1.11 ± 0.30 μm, respectively. The quick increase in the viscosity of the solutions led to the agglomeration of TiO₂ on the surface of CS hemispheres, as shown in Figure 3e₁,f₁. The size of the decrease in the CS hemispheres was due to an increase in the conductivity of the electrospray solution.

![Figure 3. SEM images and diameter distributions of the microspheres.](image)

**Table 1.** The parameters of C-T-X electrospraying solutions.

| TiO₂ Content | 0%   | 1%   | 2%   | 3%   | 4%   | 5%   |
|--------------|------|------|------|------|------|------|
| Viscosity (mPa s) | 140.7 | 158.8 | 172.5 | 183.6 | 209.5 | 220.4 |
| Conductivity (μS cm⁻¹) | 219.0 | 230.7 | 243.8 | 256.1 | 265.0 | 273.4 |
3.2. XRD Characterization

The XRD patterns of all the samples are illuminated in Figure 4. The broad peak at 20.10° and 22.01° were characteristic peaks of CS and CA, respectively [47,48]. TiO$_2$ powder showed three sharp feature peaks at 48.06°, 37.86° and 25.41° of 2θ values [45]. Compared with XRD of CA powders, the peak at 22.01° of CS/CA was weakened due to the addition of amorphous CS which inhibited crystallization of CA. Compared with XRD of CS and CA powders, the peaks at 20.10° and 22.01° of C-T-3/CA were weakened due to the addition of TiO$_2$ which inhibited crystallization of CS and CA. It indicated that there were interactions between TiO$_2$ and CS/CA, which might stabilize TiO$_2$ on the membranes [45].

![Figure 4. XRD spectra of CA, TiO$_2$, CS powders, CS/CA and C-T-3/CA fiber membranes.](image)

3.3. MO Removal Analysis

3.3.1. Effect of TiO$_2$ Content on MO Removal

The effect of the amount of TiO$_2$ on the removal of MO under visible light is demonstrated in Figure 5. Before visible light irradiation, the absorbance of MO solutions for all samples decreased due to adsorption of MO by the CS based membrane. Under visible light, the absorbance of all solutions with different fibrous membranes decreased as the reaction time increased (shown in Figure 5a). There was a significantly faster removal trend for the MO in visible light than in darkness because of the effects of both adsorption and photocatalysis. The removal of MO consisted of two stages for all samples. In the first stage, as the reaction time increased, the removal rate of MO was fast. In the second stage, the removal rate declined and got close to an equilibrium state. All samples could reach saturation state at 120 min. In combination with Figure 5b, the removal amount increased at first and then decreased with a further increase of TiO$_2$ content. When the TiO$_2$ content was higher than 3 wt %, the MO removal amount declined. This might be attributed to the reduction of the reaction sites by the TiO$_2$ agglomeration. Therefore, the best adsorption and photocatalysis performance was obtained at 3 wt % content of TiO$_2$ for CS based adsorbents.
All samples could reach saturation state at 30 min. In Figure 6b, it can be seen that the removal amount increased at first and then decreased with further increase of TiO\textsubscript{2} content. When the TiO\textsubscript{2} content was higher than 3 wt %, the MO removal amount declined. This might be attributed to the reduction of the reaction sites by the TiO\textsubscript{2} agglomeration. The removal amount of MO by adsorption and photocatalytic degradation under UV light were superior to that in visible light because of more energy for electronic transitions [23]. Therefore, C-T-3/CA was selected for further MO removal studies by UV irradiation in the following tests.

![Figure 5](image1.png)

**Figure 5.** MO removal performance under visible light. (a) The effect of reaction time on absorption amount, (b) the effect TiO\textsubscript{2} content on removal rate.

The effect of TiO\textsubscript{2} contents on the MO removal under UV light is demonstrated in Figure 6. Under UV light, the absorbance of all solutions decreased faster than samples in darkness because of concurrent adsorption and photocatalytic degradation as the reaction time increased (shown in Figure 6a). A similar situation was found as in the visible light, where the removal of MO consisted of two stages for all samples. In the first stage, the removal rate of MO was fast as reaction time increased. In the second stage, the removal rate declined and then got close to an equilibrium state. All samples could reach saturation state at 30 min [20]. In Figure 6b, it can be seen that the removal amount increased at first and then decreased with further increase of TiO\textsubscript{2} content. When the TiO\textsubscript{2} content was higher than 3 wt %, the MO removal amount declined. This might be attributed to the reduction of the reaction sites by the TiO\textsubscript{2} agglomeration. The removal amount of MO by adsorption and photocatalytic degradation under UV light were superior to that in visible light because of more energy for electronic transitions [23]. Therefore, C-T-3/CA was selected for further MO removal studies by UV irradiation in the following tests.

![Figure 6](image2.png)

**Figure 6.** MO removal performance under UV light. (a) The effect of reaction time on absorption amount. (b) The effect of TiO\textsubscript{2} content on removal rate.
3.3.2. Effect of pH Values on the MO Removal

The pH value, as a crucial parameter, controlled the MO removal ability by affecting the degree of ionization of hydrogen ions, hydroxide ions in solutions and structural stability. Figure 7 shows the removal amount of MO in the pH value range from 4 to 9 under UV light. Both samples exhibited high removal amount with pH value in the range of 4–6, and MO removal amount gradually declined as the pH value increased. C-T-3/CA had a higher removal amount than CS/CA due to both adsorption and photocatalysis for C-T-3/CA. The removal amount was greatly improved by the addition of TiO$_2$ (that allowed for simultaneous adsorption and photocatalysis) under acidic conditions. This was because under acidic conditions, the –NH$_2$ on the CS molecule was protonated, and TiO$_2$ existed in the form of TiOH$_2^{2+}$, resulting in the electrostatic attraction among—SO$_3^-$ in MO, –NH$_3^+$ and TiOH$_2^{2+}$ [48,49]. Under alkaline conditions, the –NH$_2$ on the CS molecule was deprotonated, TiO$_2$ generated the TiO$^-$, and an electrostatic repulsion appeared between fibrous membranes and MO, which resulted in the decrease of removal amount [47]. The maximum removal amount of MO for both samples was obtained at pH = 4.

![Figure 7](image)

**Figure 7.** Effect of pH values on the MO removal amount (initial concentration 40 mg/L).

3.3.3. Effect of Initial MO Concentration on the MO Removal

The effect of initial MO concentration on MO removal is indicated in Figure 8. The MO removal ratio of CS/CA and C-T-3/CA decreased with the increase of the initial concentration of MO. At a lower initial concentration of MO, fiber membranes CS/CA and C-T-3/CA provided sufficient reactive sites for removal of MO. With the increase of initial concentration of MO, the total amount of MO molecules increased. However, the amount of fiber membranes remained unchanged, meaning that the amount of reactive sites did not change. Therefore, the removal ratio of MO decreased at the higher initial concentration of MO. C-T-3/CA showed a higher removal ratio of 98.4% than CS/CA at initial concentration of 10 mg/L due to simultaneous adsorption and photocatalysis for the MO removal by C-T-3/CA.
where $q_e$ and $q_t$ (mg/g) represent the equilibrium removal amount and the removal amount at time $t$; $k_1$ and $k_2$ are the constants pseudo-first-order and pseudo-second-order rate, respectively. The experimental kinetic curves of MO removal are shown in Figure 9b,c. The relevant kinetic parameters calculated from the curves are listed in Table 2. The correlation coefficients of the pseudo-first-order model and the pseudo-second-order model for MO removal by CS/CA were nearly equal to and greater than 0.95, indicating that physical adsorption and chemical adsorption all contributed to the removal of MO. The $R^2$ of the pseudo-second-order model (0.9952) was higher than that of the pseudo-first-order model (0.9878) for the removal of MO by C-T-3/CA. This suggested that fiber membranes exhibited mainly chemical adsorption accompanied by physical adsorption via the addition of TiO$_2$ [50].
Figure 9. Effect of contact time on the MO removal amount. (a) The kinetics model for MO removal, (b) pseudo-first-order model, and (c) pseudo-second-order model.

Table 2. Kinetics parameters for MO removal.

| Sample    | \( q_e, exp \) (mg/g) | Pseudo-First-Order Model | Pseudo-Second-Order Model |
|-----------|------------------------|--------------------------|----------------------------|
|           | \( k_1 \) | \( R^2 \) | \( q_e, cal \) (mg/g) | \( k_2 \) | \( R^2 \) | \( q_e, cal \) (mg/g) |
| CS/CA     | 19.24   | 5.44 \( \times 10^{-2} \) | 0.9748 | 15.08 | 13.48 \( \times 10^{-2} \) | 0.9799 | 20.80 |
| C-T-3/CA  | 39.20   | 9.86 \( \times 10^{-2} \) | 0.9878 | 30.61 | 32.33 \( \times 10^{-2} \) | 0.9952 | 40.67 |

3.3.5. Isotherm Study

The Langmuir model and Freundlich model were used to describe the isotherm equilibrium data of MO removal by CS/CA and C-T-3/CA fiber membranes [50]. It can be expressed as Equations (5) and (6).

The Langmuir model:

\[
\frac{C_e}{q_e} = \frac{1}{q_m K_L} + \frac{C_e}{q_m}
\]  

(5)

The Freundlich model:

\[
\log q_e = \frac{1}{n} \log C_e + \log K_F
\]  

(6)

where \( q_e \) (mg/g) represents the equilibrium removal amount, \( C_e \) (mg/L) is the equilibrium concentration of MO in aqueous solution and \( q_m \) (mg/g) represents the maximum removal amount, \( K_L \) is the Langmuir isotherm constant related to free energy and \( K_F \) and \( n \) are the Freundlich isotherm constant which expressed removal capacity and removal intensity, respectively.
The experimental isotherm curves of MO removal are shown in Figure 10a,b. The relevant isotherm parameters calculated from the curves are listed in Table 3. On the basis of the correlation coefficients of MO removal by CS/CA, the Freundlich isotherm model (Figure 9b) was better fitted than the Langmuir isotherm model (Figure 9a). Therefore, the CS/CA mainly exhibited multiple layers of adsorption. After the addition of TiO\textsubscript{2}, the correlation coefficients of the Langmuir model and Freundlich model were approximately equal, indicating that C-T-3/CA exhibited multiasorption behavior. This might be due to both the effects of photocatalytic degradation and adsorption [50].

![Figure 10. Isotherm model of MO removal: (a) Langmuir model, (b) Freundlich model.](image)

### Table 3. Isotherm parameters for MO removal.

| Sample     | Langmuir Model | Freundlich Model |
|------------|----------------|-----------------|
|            | $K_L$          | $q_m$ (mg/g)    | $R^2$ | $K_F$ | $n$ | $R^2$ |
| CS/CA      | $7.28 \times 10^{-2}$ | 30.01         | 0.9714 | 2.30  | 1.41 | 0.9932 |
| C-T-3/CA   | $62.5 \times 10^{-2}$ | 76.22         | 0.9948 | 25.33 | 1.90 | 0.9997 |

3.4. Removal Mechanism Analysis

3.4.1. FTIR Spectra Results

The FTIR spectra of C-T-3/CA fibers before and after removal of MO are demonstrated in Figure 11. Before removal of MO, the peaks at 3100–3500 cm\textsuperscript{-1} were ascribed to –OH and –NH\textsubscript{2}, 1638 and 1572 cm\textsuperscript{-1} were attributed to –NH\textsubscript{3}\textsuperscript{+} and –NH\textsubscript{2} of CA and CS, respectively. The peak appearing at 1154 cm\textsuperscript{-1} was attributed to the stretching of the C–O–C band of CA. The peaks in the range of 400–800 cm\textsuperscript{-1} were ascribed to the characteristic absorption of Ti–O [47,51]. After the adsorption and photocatalytic degradation of MO, the peak of the –NH\textsubscript{2} group at 1572 cm\textsuperscript{-1} decreased and the –NH\textsubscript{3}\textsuperscript{+} at 1638 cm\textsuperscript{-1} increased [49]. It demonstrated that –NH\textsubscript{2} was protonated as –NH\textsubscript{3}\textsuperscript{+} in the process of MO removal. Several new characteristic peaks that appeared were listed as follows: 1400–1450 cm\textsuperscript{-1} were ascribed to –N=\text{N}– group, and 1039 cm\textsuperscript{-1} was attributed to –SO\textsubscript{3}\textsuperscript{−} of MO. It showed that MO was adsorbed on surfaces of CS molecules during the reaction process. The peaks at 3100–3500 cm\textsuperscript{-1} increased, which might be due to the formation of abundant active hydroxyl radicals (\text{OH}) during the photodegradation process (see Equations (9) and (10)).
3.4.2. XPS Spectra Results

As shown in Figure 12, XPS was employed to analyze the chemical element of the membrane surface before and after the removal (that allowed for cointaneous adsorption and photocatalysis) of MO with full-range and magnified spectra of S, N, O and Ti. It can be seen in Figure 12a that C-T-3/CA membranes before MO removal showed peaks of binding energy at 285.5, 397.7, 531.2 and 457.1 eV, which were assigned to carbon atom (C 1s), oxygen atom (O 1s), nitrogen atom (N 1s) and titanium atom (Ti 2p), respectively. As seen in Figure 12b, the peak of S 2p appeared at 167.7 eV after removal of MO, indicating MO adsorbed on the surface fiber membranes. For the N 1s spectrum in Figure 12c, the peak at 399.1 eV was assigned to –NH₂ group before MO removal. After MO removal, a new peak appeared at 401.7 eV and was attributed to –NH₃⁺, and the peak of –NH₂ at 399.1 eV shifted to 399.8 eV. This indicated that –NH₂ was protonated as –NH₃⁺, and electron transfer occurred between –NH₂ and MO molecules, respectively [22]. As seen in Figures 12d,e, the O 1s and Ti 2p binding energy increased after the removal of MO. Ti could exist in the form of TiOH₂⁺ in acid solutions. After the reaction with MO molecules, the groups of –OH₂⁺ as electron acceptors could interact with –SO₃⁻ of MO due to electrostatic attraction, which could influence charge density of Ti and O [20].

3.4.3. BET Analysis

The BET results of CS/CA and C-T-3/CA membranes were measured by nitrogen adsorption method (see Table 4). After the addition of TiO₂, the fiber membrane attained higher specific surface area and pore volume owing to uniform distribution of TiO₂ on CS hemispheres and an increase of the distance among C-T hemispheres (see Figure 3d). Therefore, the relative high adsorption capacity of MO on C-T-3/CA was ascribed to the high specific surface area, which should be good for the photodegradation of MO.
3.4.3. BET Analysis

The BET results of CS/CA and C-T-3/CA membranes were measured by nitrogen adsorption method (see Table 4). After the addition of TiO$_2$, the fiber membrane attained higher specific surface area and pore volume owing to uniform distribution of TiO$_2$ on CS hemispheres and an increase of the distance among C-T hemispheres (see Figure 3d). Therefore, the relative high adsorption capacity of MO on C-T-3/CA was ascribed to the high specific surface area, which should be good for the photodegradation of MO.

| Sample     | Specific Surface Area (m$^2$/g) | Pore Volume (cc/g) | Average Pore Diameter (nm) |
|------------|---------------------------------|-------------------|---------------------------|
| CS/CA      | 301.30                          | 0.33              | 4.39                      |
| C-T-3/CA   | 342.10                          | 0.48              | 4.79                      |

3.4.4. Mechanism Analysis

According to the above analysis, the high removal capacity of C-T-3/CA was mainly due to the effective dispersion of TiO$_2$ in combination with CS, which improved the specific surface area of fiber membranes and increased the sites of photocatalytic degradation and adsorption. MO molecules could be adsorbed to the vicinity of photodegradation sites by electrostatic attraction between CS and...
MO molecules, promoting the contact probability of TiO\textsubscript{2} and MO in the following photocatalytic degradation process. The reaction mechanism between MO and C-T-3/CA is illustrated in Figure 13. The mechanism of photocatalysis is summarized from Equation (7) to Equation (12). The MO removal mechanism included: (1) Adsorption, the electrostatic attraction between –NH\textsubscript{3}\textsuperscript{+} group of CS and –SO\textsubscript{3}\textsuperscript{−} of MO molecules; (2) photocatalysis, the photocatalytic degradation of the MO by UV illumination began with photoexcitation of the TiO\textsubscript{2} and then formed an electron-hole pair (Equation (7)). High oxidation valence-band holes (h\textsuperscript{+}) directly oxidized MO degradation (Equation (8)). Water decomposition produced \cdot\text{OH} (Equation (9)) or a reaction of h\textsuperscript{+} with OH\textsuperscript{−} (Equation (10)). Meanwhile, the reaction between conduction-band electrons (e\textsuperscript{−}) and proper electron acceptors (such as O\textsubscript{2}) yielded oxidative radicals as described by Equation (11). The generated hydroxyl radicals easily degraded MO to form inorganic small molecules (Equation (12)) [6]. These results agreed with BET, FTIR and XPS analyses.

\begin{align*}
\text{TiO}_2 + \text{hv} & \rightarrow \text{e}^{-} + \text{h}^{+} \quad (7) \\
\text{h}^{+} + \text{MO} & \rightarrow \text{MO}^{+} \rightarrow \text{Oxidation of MO} \quad (8) \\
\text{h}^{+} + \text{H}_2\text{O} & \rightarrow \text{H}^{+} + \cdot\text{OH} \quad (9) \\
\text{h}^{+} + \text{OH}^{−} & \rightarrow \cdot\text{OH} \quad (10) \\
\text{e}^{-} + \text{O}_2 & \rightarrow \cdot\text{O}_2 \quad (11) \\
\text{MO} + \cdot\text{OH} / \cdot\text{O}_2^{−} & \rightarrow \text{Degradation of MO} \quad (12)
\end{align*}

![Image](image.png)

**Figure 13.** The mechanism of MO removal by C-T-3/CA.

3.5. Recycling Ability

To research the recycling ability of the composite membranes, the absorbance changes were measured in five cycles of MO removal. As demonstrated in Figure 14, the MO removal of fiber membranes was almost invariable for five consecutive cycles. The membranes still had a removal ratio of 98% at the end of the fifth cycle, testifying that the membranes maintained good performance of adsorption and photocatalytic degradation. This was because the stability of CS in aqueous solutions was improved by the intermolecular hydrogen bonds between CS and CA, and TiO\textsubscript{2} could be uniformly distributed via the interactions with CS molecules for effective photocatalytic degradation of MO [45].
4. Conclusions

In this work, the C-T/CA fiber membranes were prepared via electrospinning and electrospraying. After the addition of TiO$_2$, the removal capacity of MO was improved, as a result of concurrent adsorption and photocatalytic degradation. When TiO$_2$ content was 3 wt %, the highest MO removal amount for fiber membranes (C-T/CA) was reached, 98% at pH value of 4 and MO concentration of 40 mg/L. The kinetic and isotherm model for the removal of MO by fiber membranes were well fitted pseudo-second-order kinetic and Freundlich isotherm models. Especially, C-T/CA membranes exhibited multiple layer adsorption with the addition of TiO$_2$. The involved mechanisms including adsorption and photocatalytic reaction occurred in the removal process which was demonstrated by FTIR, XPS and BET results. The recycling experiment showed fiber membrane had excellent stability and reusability.

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