Doping of Chlorine from a Neoprene Adhesive Enhances Degradation Efficiency of Dyes by Structured TiO$_2$-Coated Photocatalytic Fabrics

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Received: 22 November 2019; Accepted: 31 December 2019; Published: 2 January 2020

Abstract: We demonstrate that using neoprene as a binder during the fabrication of TiO$_2$-coated fabrics enhances the rates of photodegradation of dyes by the fabrics. The neoprene binder simultaneously modifies the surface of the TiO$_2$ particles with Cl and dopes the TiO$_2$ with Cl, without requiring high temperatures or other harsh laboratory treatments. The adsorption of chlorine on the surface and doping of the lattice with chlorine were confirmed by X-ray photoelectron spectroscopy. The chloride ions adsorbed onto the TiO$_2$ surface introduces a negative surface charge that enhances electrostatic adsorption of cationic dyes, and greatly improves the self-sensitizing degradation performance of the dyes. Chloride ions replace lattice oxygen atoms in TiO$_2$, inducing lattice oxygen vacancies, that reduce the apparent band gap of the TiO$_2$ particles, enhancing its absorption of visible light, and further increasing the photocatalytic activity of the composite-coated fabric. The degradation rates of RhB and MB over 50 min were 95.2% and 96.0%, respectively. The degradation rate for MO reached 95.4% after 180 min. We also show that $\cdot$OH and $\cdot$O$_2$$^{-}$ are active agents in the dye-degradation mechanism. Moreover, the photocatalytic performance of the composite-coated fabric was unchanged after eight cycles of photocatalytic degradation of RhB, demonstrating that the photocatalyst-coated fabrics was highly recyclable.

Keywords: TiO$_2$-coated fabrics; self-sensitizing degradation; neoprene; chlorine doping; oxygen vacancies

1. Introduction

Continued industrial development has caused water pollution to become an increasingly serious problem. Organic dyes discharged during industrial activities, including textile production, printing, tanning, and papermaking, have damaged the environment and have even threatened human health [1]. Moreover, most organic dyes used in industries contain aromatic conjugated structures that are chemically stable and difficult to degrade [2]. Therefore, the development of effective methods for the removal of dyes from the industrial wastewater is important for environmental sustainability [3–5]. The main methods used to remove dyes from wastewater include adsorption, electrochemical oxidation, filtering, degradation by microbes [6], and photocatalytic degradation. Although adsorption is effective in removing dyes from water, adsorption does not eliminate contamination because the process does not convert the dye molecules to harmless chemicals. Photocatalytic degradation is a more recently developed method that uses sunlight to stimulate semiconducting catalysts to generate strongly
oxidative free radicals that attack most organic substances in water. Photocatalytic degradation has low operating costs and is a promising energy-saving technology for wastewater treatment.

TiO$_2$ is currently the photocatalyst that is most widely used for wastewater treatment; TiO$_2$ is chemically stable, non-toxic, inexpensive, and exhibits high photocatalytic activity [7–10]. However, TiO$_2$ has a wide band gap that limits its light absorption to ultraviolet wavelengths, and exhibits high rates of recombination for photogenerated electrons and holes. Furthermore, TiO$_2$ has limited capacity for adsorption of pollutants, and small particles of TiO$_2$ agglomerate easily [11–13]. In combination, these drawbacks limit the ability of TiO$_2$ to use sunlight efficiently, and result in low photocatalytic efficiencies [14]. Therefore, increasing the photocatalytic activity of TiO$_2$ requires the development of methods that either enhance the absorption spectrum of TiO$_2$ or inhibit the recombination of photogenerated electron-hole pairs. Recently, a variety of methods have been used to increase the photocatalytic activity of TiO$_2$, such as doping with non-metals or combining TiO$_2$ with another semiconductor or with a photosensitizer. TiO$_2$ has been doped with N, C, F, S, and I [15–19]; these dopants enhanced the photocatalytic activity by increasing the absorption spectrum of the TiO$_2$.

In recent years, efforts to extend the TiO$_2$ absorption spectra into the visible range have focused on inducing the formation of Ti$^{3+}$ defects. For example, Ti$^{3+}$ defects can be induced by generating oxygen vacancies either through a reduction reaction that occurs under high pressures of H$_2$ [20–22] or through vacuum activation [23,24]. However, these methods are expensive and require harsh conditions, and therefore are rarely suitable for practical applications.

Dye sensitization is a well-known and frequently used method of expanding the effective absorption spectra of TiO$_2$-based materials to visible wavelengths. Absorption of light by dye molecules on the surface of TiO$_2$ excites electrons in the dye that are then injected into the TiO$_2$ conduction band. Over time, the dye molecules become self-sensitized and degraded. Modification of the TiO$_2$ surface can enhance the adsorption of dyes and improve the photocatalytic performance; for example, Jincai Zhao et al. showed that the surface of TiO$_2$ can be modified with F ions through HF etching [25]. Ji-Jun Zou et al. showed that the electronic structure of the titanium dioxide surface can be regulated by aqueous media [26] using a method that changes surface charge that can be held by the TiO$_2$, and thereby enhancing the adsorption of cationic dyes. However, selective adsorption of dyes has not yet substantially enhanced the photocatalytic activity of TiO$_2$, and modifications of TiO$_2$ that can be performed under standard laboratory conditions for maximum compatibility with large-scale catalyst recovery from dye-contaminated wastewaters have not yet yielded significant improvements in photocatalytic activity.

Herein, a small amount of neoprene is used as an adhesive, which is an excellent polymer adhesive. It has stable physical and chemical properties, strong adhesion, aging resistance, heat resistance, and excellent chemical resistance [27]. It is widely used in many fields such as wires and cables, tapes, adhesives etc. [28]. The crosslinking ability of neoprene when heated has been reported. Metal oxide e.g., ZnO can be used as an agent for the crosslinking reaction of neoprene. In the crosslinking process, the C=C double bond on the polarity of chlorine can react with a crosslinking agent ZnO [29].

We describe a one-step method for modifying the TiO$_2$ surface and doping the TiO$_2$ with Cl. In this method, P25 nanoparticles are affixed to fabric using a neoprene (chloroprene) binder. The preparation modifies the TiO$_2$ surface and dopes the lattice with Cl without requiring high-temperature calcination or other subsequent treatments. Chloride ions adsorbed to the surface enhance electrostatic adsorption of cationic dyes, improving the self-sensitized degradation of the dye, while chloride ions in the lattice replace oxygen leaving oxygen vacancies that enhance the absorption of visible light by the TiO$_2$. The photocatalyst can be recycled and stabilized while on the surface of the fabric. The method is simple, low-cost, compatible with mass production, and applicable to treatment of wastewaters from printing and dyeing industries.
2. Results

Figure 1 compares the SEM micrographs of samples of pure and photocatalyst-coated fabrics. The surface of the pure fabric was smooth, and silky fibers were clearly visible; the surface of the photocatalyst-coated fabric was rough, and TiO$_2$ particles were clearly uniformly dispersed on, and closely adhered to, fabric fibers. Thus, the simple spraying method yielded a photocatalyst-coated fabric. Figure 2 compares transmission-electron micrographs of Cl-TiO$_2$ and P25 at low and high resolutions. Cl-TiO$_2$ was more readily dispersed than P25, and the particle size of Cl-TiO$_2$ was slightly smaller than that of P25. We hypothesize that the Cl-TiO$_2$ is more readily dispersed than P25 because the Cl ions on the surface of Cl-TiO$_2$ replace the hydroxyl group on the surface, reducing the degree of agglomeration of TiO$_2$. The lattice fringes of Cl-TiO$_2$ and P25 were clearly visible in the high-resolution TEM images. An interplanar spacing of 0.353 nm was measured and is attributable to the (101) crystal plane of the anatase crystal form of TiO$_2$ (JCPDS 21-1272). HRTEM images confirmed the presence of defective structures in Cl-TiO$_2$, most likely because of doping of the TiO$_2$ by chloride ions. The EDS mapping of Cl-TiO$_2$ is shown in Figure 2e. The results showed that Ti, O, and Cl elements was uniformly dispersed in the catalyst, confirming the presence of chlorine in Cl-TiO$_2$.

Figure 1. Scanning-electron micrographs of pure fabrics (top row) and photocatalyst-coated fabrics (bottom row).

Figure 3 shows X-ray diffraction patterns obtained for composite-coated and control samples and for P25 and Cl-TiO$_2$. The untreated pure fabric exhibited diffraction peaks at 18.0° and 25.8°. These peaks remained present in the spectra after coating the fabric with both neoprene and TiO$_2$, indicating that the fabric was not damaged. The fabric peak was relatively weaker for composite-coated fabrics because the fabric surface was covered by neoprene and TiO$_2$ coatings. In addition, the photocatalyst-coated fabric exhibited additional peaks at 25.3°, 27.3°, 37.8°, 48.0°, 53.9°, 55.1°, 62.7°, corresponding to that of P25 [30]. The XRD data further support that TiO$_2$ (P25) successfully dispersed onto the fabric surface, consistent with the SEM observations. Moreover, the XRD pattern for the Cl-TiO$_2$ obtained by pulverizing and filtering the coated fabric was consistent with the crystal form of P25, indicating that doping TiO$_2$ with chloride did not affect the crystal structure.
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Figure 2. Low-resolution and high-resolution transmission-electron micrographs of (a,c) Cl-TiO$_2$ and (b,d) P25. (e) Energy-dispersive X-ray spectrum EDX maps of Ti, O, and Cl in Cl-TiO$_2$. 

Figure 3. XRD patterns of pure fabric, neoprene-coated fabric, and photocatalyst-coated fabric (in the left). XRD patterns of P25 and Cl-TiO$_2$ (in the right). 

Figure 4 shows the zeta potentials of suspensions of TiO$_2$ and Cl-TiO$_2$ at different pHs. The TiO$_2$ suspension is amphoteric due to the presence of surface hydroxyl groups that undergo the following acid-base equilibration [31,32]: 

$$\text{TiOH}^+ \leftrightarrow \text{TiOH} + \text{H}^+ (1)$$

$$\text{TiOH} \leftrightarrow \text{TiO}^{2-} + \text{H}^+ (2)$$

As shown in the Figure 4, the isoelectric point of the TiO$_2$ particles is at pH = 6.7. In an acidic environment with pH < pH zpc, the zeta value of TiO$_2$ is determined by the adsorption of hydrogen ions on the surface hydroxyl groups to form TiOH$^+$. Similarly, in an alkaline environment with pH > pHzpc, the hydroxyl groups on the surface of TiO$_2$ are deprotonated and form a TiO$^{2-}$ structure with a negative zeta value. However, the zeta value of the Cl-TiO$_2$ suspension was negative in the range of pH 5–11, indicating that the adsorption of chloride ions on the surface of TiO$_2$, promotes a negative surface charge on the TiO$_2$. The negative surface charge is beneficial for the selective adsorption of cationic dyes in the photocatalytic reaction process, and improves the degradation performance of the TiO$_2$ for cationic dyes.
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Figure 5 compares the UV-Vis absorption spectra for the composite-coated and control samples. The pure fabric and the chloroprene-coated fabric had weak absorption intensities across the 200 nm–800 nm wavelength range, indicating that neoprene does not substantially enhance light absorption by the fabric. The fabric coated with P25 exhibited a high absorption value in the ultraviolet region, and weak absorption in the visible light region, consistent with the absorption of TiO$_2$ (P25). The absorption intensity of the photocatalyst-coated fabric was enhanced in the visible region across the 400 nm–800 nm wavelength range, possibly because of chloride-ion doping of the TiO$_2$ by the neoprene rubber. The absorption of chlorine-doped TiO$_2$ is red shifted relative to P25, because the chlorine doping reduces the apparent band gap of titanium dioxide [33]. The band gaps of P25 and Cl-TiO$_2$ were estimated to be 2.94 eV and 2.53 eV, respectively, based on the analyses of diffuse reflectance spectra using the relationship between band gap and $(ahv)^{1/2}$ for indirect-transition semiconductors. The narrower band gap of Cl-TiO$_2$ is beneficial for the absorption of light, and significantly enhances the photocatalytic performance relative to P25.
The C 1s peak at 284.5 eV can be attributed to carbon incorporated into TiO₂ after dechlorination of chloroprene rubber, and to carbon bonded to the chloroprene rubber on the surface of TiO₂. Figure 6a shows a survey scan for Cl-TiO₂, showing the presence of Ti, C, O, and Cl. The C 1s peak at 284.5 eV can be attributed to carbon incorporated into TiO₂ after dechlorination of chloroprene rubber, and to carbon bonded to the chloroprene rubber on the surface of TiO₂. Figure 6b shows the high-resolution XPS data in the region of the Ti 2p peaks, located at 464.43 eV and 458.65 eV, corresponding to the Ti⁴⁺2p₁/₂ and Ti⁴⁺2p₃/₂ spin-orbit-split peaks, respectively [7]. As shown in Figure 6c, the O 1s peak can be fit to three different peaks: the peaks at 529.9 eV and 533.2 eV correspond to the lattice oxygen and surface hydroxyl oxygen of titanium dioxide, respectively [17]; the peak at 531.9 eV can be attributed to oxygen vacancies in TiO₂. Figure 6d shows the XPS high-resolution spectrum in the Cl region, which shows three forms of chlorine in the sample. The low-binding-energy peak at 197.9 eV, likely corresponds to chlorine adsorbed on the surface of TiO₂ [34], either by physical adsorption or chemisorption in place of hydroxyl groups on the surface of TiO₂. For F-TiO₂, the F⁻ adsorbed on the surface can be readily replaced with OH⁻ after washing with concentrated sodium hydroxide, but F in the TiO₂ lattice is not readily replaced [35,36]. We hypothesized that the behavior of Cl here likely would be similar to that of fluorine, so we washed the Cl-TiO₂ with 1 M NaOH, and characterized the sample using XPS. As shown in Figure 6e, the peak of 197.89 eV was greatly weakened by the washing, but the peaks at 200.09 eV and 201.70 eV were not substantially weakened. These results are consistent with replacement of chloride ions on the TiO₂ surface by OH⁻, and suggest that the peak at 197.9 eV is attributable to chloride ions adsorbed on the surface of TiO₂, while the peak at ≈200.0 eV [37] is attributable to chlorine doped into the TiO₂ lattice in place of lattice oxygen. Meanwhile, the peak at 200.70 eV can be attributed to chlorine in the neoprene rubber adhesive on the surface of titanium dioxide; this peak was not affected by NaOH washing because neoprene is an alkali-resistant material.

Figure 7 compares the EPR spectra for Cl-TiO₂ and P25. P25 exhibits no EPR signal value under vacuum at 100 K, indicating that P25 does not contain solitary electrons of paramagnetic species. An EPR signal can be clearly seen for Cl-TiO₂, corresponding to the capture of a pair of electrons by oxygen vacancies, consistent with the presence of oxygen-vacancy defects in Cl-TiO₂. Again, these results support that chlorine replaces oxygen in the TiO₂ lattice, generating oxygen vacancies.
Figure 6. XPS spectra of Cl-TiO₂: (a) survey spectrum, (b) Ti 2p spectrum, (c) O 1s spectrum, (d) Cl 2p spectrum, and (e) Cl 2p spectrum after washing with 1M NaOH.

The efficiency of separation of photogenerated electrons and holes plays an important role in the photocatalytic reaction [38]. Photoluminescence measurements can reveal the photochemical characteristics of semiconductors, and the emission intensity of a PL spectrum represents the efficiency of electron-hole separation in the semiconductor. Figure 8 compares the PL spectrum of P25 with that for Cl-TiO₂ at an excitation wavelength of 380 nm. The PL emission intensity of Cl-TiO₂ was lower than that of P25, indicating a lower probability of electron-hole recombination Cl-TiO₂ [39]; therefore, the separation efficiency of electron-hole pairs on the surface of Cl-TiO₂ during the photocatalytic process might be higher than that of P25 [12]. The lattice oxygen vacancies in Cl-TiO₂ facilitate the capture of photogenerated electrons and promote the separation of electrons and holes [40,41]. The Nyquist diagram (Figure 8b) for P25 exhibited a large radius of curvature, while the plot for Cl-TiO₂ exhibited a small radius of curvature, indicating that doping TiO₂ with Cl causes a large decrease in resistance, which is beneficial to reducing the probability of electron-hole recombination in the composite and improves light utilization.
water to generate •OH, and the oxygen vacancies enhance the adsorption of oxygen and promote the generation of •O2− [43]. Therefore, it is possible that two kinds of free radicals play a major role in the degradation of MO. As shown in Figure 9f, the degradation exhibits characteristics of first-order kinetics. Kinetic constants for photodegradation of RhB, MB, and MO dyes by using photocatalyst-coated fabrics are $3.793 \times 10^{-2}$ min$^{-1}$, $5.191 \times 10^{-2}$ min$^{-1}$, and $1.803 \times 10^{-2}$ min$^{-1}$, respectively, which are higher than P25-coated fabrics (Figure 9g). In order to eliminate the effect of pure fabric and neoprene

Figure 7. EPR spectra of P25 and Cl-TiO$_2$.

Figure 8. (a) PL spectra, (b) EIS spectra of P25 and Cl-TiO$_2$.

Figure 9 shows the photocatalytic activity of photocatalyst-coated fabrics for the degradation of RhB, MB, and MO dyes under visible light. The photocatalyst-coated fabric exhibited excellent adsorption for the two cationic dyes; the adsorption mechanism is shown in Scheme 1. The 30-min adsorption-degradation rates reached 66.1% and 46.0%, respectively (Figure 9d). After 50 min, the photodegradation performances for RhB and MB were 95.2% and 96.0, respectively. As shown in Figure 9a, during the photodegradation of RhB, the solution gradually changed from pink to pale yellow and finally became colorless at 50 min. The maximum absorption peak shifts from 555 nm to 511 nm, which is caused by RhB de-ethylation [42]. The anionic dye methyl orange was very weakly adsorbed by the composite-coated fabric. However, the photocatalyst-coated fabric also exhibited excellent properties for the degradation of methyl orange under visible light, with the degradation rate reaching 95.4% after 180 min, far greater than the degradation rate for P25 (Figure 9d). We attribute the increased rate of degradation to Cl-TiO$_2$ on the surface of the coated fabric being rich in oxygen vacancies, reducing the band gap and enhancing the absorption of visible light. At the same time, the oxygen vacancies are favorable for capturing photogenerated electrons to improve electron-hole separation efficiency and thereby improve photocatalytic efficiency. The holes can oxidize water to generate •OH, and the oxygen vacancies enhance the adsorption of oxygen and promote the generation of •O2− [43]. Therefore, it is possible that two kinds of free radicals play a major role in the degradation of MO. As shown in Figure 9f, the degradation exhibits characteristics of first-order kinetics.

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on the degradation of adsorbed dyes, photocatalytic degradation of RhB by the pure fabric and neoprene-coated fabric control samples was measured. RhB only very weakly adsorbed to the pure fabric with neoprene. The lifetime of the catalyst plays a key role in the photocatalytic reaction; it is especially important that the photocatalyst strongly adheres to the fabric. We found that the performance of the composite-coated fabric was not reduced after eight cycles of photodegradation of RhB (Figure 9g), indicating that the photocatalyst-coated fabric has good recycling properties.

Figure 9. UV-vis absorption spectra over time during the degradation of RhB (a), MB (b), and MO (c) by photocatalyst-coated fabric. (d) Photodegradation of three dyes by photocatalyst-coated fabric. (e) Degradation of RhB by pure fabric, neoprene-coated fabric, P25, and photocatalyst-coated fabric. (f) Pseudo first-order kinetic curve and (g) kinetic constants for photodegradation of three dyes using photocatalyst-coated fabric and P25, (h) recycling of photocatalyst-coated fabric for the photodegradation of RhB.
Active intermediates such as h\(^+\), •OH, •O\(^{2-}\) are involved in the photocatalytic oxidation of dyes [44–46]. The composite-coated fabric exhibited excellent degradation performance for the cationic dyes because Cl-TiO\(_2\) electrostatically adsorbs cationic dyes, enhancing the photosensitization and promoting dye degradation (Scheme 1). However, the reason for the enhancement of degradation of the anionic dye MO on the composite-coated fabric remained unclear.

We therefore examined the degradation of MO by the composite-coated fabric in the presence of ammonium oxalate (AO), benzoquinone (BQ), and isopropanol (IPA) to quench h\(^+\), •O\(^{2-}\), and •OH, respectively. Figure 10b shows the photodegradation rate of MO was essentially unchanged by the addition of AO. This result is consistent with the inability of h\(^+\) to oxidize MO, because electrostatic repulsion between MO and Cl-TiO\(_2\) prevents strong adsorption of dye molecules to the catalyst surface. After the addition of BQ, the efficiency of catalytic degradation of MO was greatly reduced, with the degradation rate after 180 min being reduced from 92.2% to 45%. This result shows that •O\(^{2-}\) plays a major role in photocatalytic degradation of MO. When IPA was added, the efficiency of photocatalytic degradation of MO also decreased, with the degradation rate over 180 min being reduced from 92.2% to 78.6%, showing that •OH also oxidizes MO, but the catalytic effect is weaker than •O\(^{2-}\). Therefore, we conclude that •O\(^{2-}\) and •OH free radicals play a key role in the photodegradation of MO. As shown in Figure 10a, the valence band spectrum of Cl-TiO\(_2\) was obtained by XPS spectrum, and the energy level position can be calculated by combining the DRS spectrum. The mechanism of photocatalytic degradation of dyes is shown in Scheme 2. RhB or MB was stably adsorbed on the surface of Cl-TiO\(_2\), under visible light excitation, electrons transition from the ground state to the excited state, and then transfer to the conduction band position of Cl-TiO\(_2\), thereby achieving photosensitization of Cl-TiO\(_2\), which underwent self-sensitized degradation. For the MO system, Cl-TiO\(_2\) was excited by the visible light to produce separation of electron holes, the hole oxidized water to generate •OH, and molecular oxygen was reduced to form •O\(^{2-}\), thereby realizing degradation of MO.
3. Materials and Methods

3.1. Materials

Flumes fabric filler was obtained from Hebei Yisheng Environmental Protection Co., Ltd. (Beijing China). Titanium dioxide (P25) was purchased from Degussa. NeopreneSN244X was bought from Shanxi Datong Yujie Chemical Co., Ltd. (Beijing China). Absolute ethanol was supplied by Beijing Chemical Factory (Beijing China). Xylene purchased from Beijing Chemical Factory. Rhodamine B, methyl orange, and methylene blue were acquired from Aladdin Ltd (Beijing China). The chemicals used were of analytical grade and were used as received. The acronyms are as follows: rhodamine B (RhB), methylene blue (MB), methyl orange (MO), photocatalyst-coated fabric (PCF), P25-coated fabric (P25F), fabric-only control samples (PF), neoprene-coated fabric (NCF).

3.2. Preparation of the TiO2 Suspension and Neoprene Adhesive

Total of 1.0 g of TiO2 (P25) was stirred in 100 mL of absolute ethanol for 2 h, until the TiO2 particles were completely dispersed. A total of 10 g of neoprene (chloroprene) was cut into small pieces and was stirred in 60 mL of xylene for 12 h until the neoprene was completely dissolved.

3.3. Preparation of Photocatalyst-Coated Fabric

Flumes fabric (20 cm × 20 cm) was immersed in a mixture containing 100 mL ethanol and 100 mL acetone and was sonicated for 60 min. The fabric was then dried at 60 °C in an oven. The neoprene

Figure 10. (a) VB XPS spectra of P25 and Cl-TiO2. (b) Photodegradation of MO by composite-coated fabrics in the presence of different capture agents.
adhesive was then applied evenly to the surface of the fabric, and the titanium dioxide suspension was sprayed evenly onto both sides of the fabric using a high-pressure spray gun. The coated fabric was dried at 80 °C for 4 h. After drying, the coated fabric was taken out and placed at room temperature (25 °C) for one week to obtain a photocatalyst-coated fabric (PCF) (Scheme 1). Neoprene-coated fabric (NCF) control samples were made by coating pieces of fabric with only the neoprene adhesive, it was treated by the same drying/ageing procedure. P25-coated fabric (P25F) control samples were made by spraying the fabric with only the P25 dispersion. Fabric-only control samples (PF) were prepared from cleaned pieces of fabric without the neoprene adhesive or P25 particles. The photocatalyst-coated fabric was crushed and dispersed in an ethanol solution, ultrasonicated, washed, and filtered to obtain a Cl-TiO₂ powder. The preparation procedure of photocatalyst-coated fabrics was shown in Scheme 3. The doping of Cl in TiO₂ was schematically shown in Scheme 4.

![Scheme 3. Preparation procedure of a photocatalyst-coated fabrics.](image)

![Scheme 4. Evolution of pristine P25 to Cl-doped TiO₂.](image)

3.4. Characterization

The dispersion of TiO₂ on the surface of catalyst-coated fabrics was characterized using scanning-electron microscopy (Hitachi S-4700, Japan). The morphology of the powdered catalyst was characterized using transmission-electron microscopy (JEOL JEM-2010F, Japan). Lattice parameters and elemental distributions of powdered catalysts were characterized using a transmission-electron microscope (JEOL JEM-3010). X-ray diffraction was performed using a Shimadzu XRD-6000 (Shimadzu, Japan). The powdered catalyst was also characterized by X-ray photoelectron spectroscopy (XPS, Thermo VG ESCALAB MK II, USA). UV-visible spectroscopy (UV-3600, Shimadzu, Japan) was used to characterize the absorption spectra of the catalyst particles. The fluorescence emission of the powdered catalyst was analyzed using an FL-7000 spectrometer from Tianmei (China) Scientific Instrument Co. Ltd. (Beijing, China), and an excitation wavelength of 360 nm. The zeta potential of the solid powder dispersion was measured by a Shanghai Zhongchen JS94J microelectrophoresis
apparatus, and the pH range was 3–11. EPR was carried out using a Bruker A200 spectrometer under normal temperature conditions. The photoelectrochemical properties of solid powdered catalysts were determined using a Shanghai Chenhua CHI660E electrochemical workstation, a Ag/AgCl reference electrode and platinum counter electrode in 0.2 M Na₂SO₄ solution.

3.5. Photocatalytic Experiments

The chemical structures of the three dyes RhB, MB and MO were shown in Scheme 5. In order to test the photocatalytic activity of the prepared composite-coated fabrics, the three dyes were irradiated with visible light (λ > 425 nm) at an intensity of 100 mW/cm². The source of illumination was a 300 W xenon lamp. The composite-coated fabric (5 cm × 5 cm) was loaded into a sandwich cell, with liquid water between the interlayers to exclude thermocatalytic degradation. The composite-coated fabric was then placed in contact with a solution containing 100 mL of rhodamine B (RhB) (20 mg/L), methylene blue (MB) (20 mg/L), or methyl orange (MO) (20 mg/L), and was left in the dark for 30 min to allow the sample to reach an adsorption equilibrium with the mixture. The sample was then illuminated from a direction normal to the surface using a light source 10 cm above the liquid surface. During the photocatalytic reaction, 3 mL RhB solution or MB solution was extracted from the cell every 10 min; 3 mL MO solution was extracted every 30 min. Photocatalytic degradation of RhB by control samples (pure fabric, neoprene-coated fabric, and P25-coated fabric) was measured under the same conditions as for the composite-coated fabrics. The concentration of dye in each extracted aliquot was measured by ultraviolet absorption spectrometry. The rate constant for the degradation of the dye was calculated according to the following formula [42,47]:

\[
\frac{DRhB(DMB, DMO)}{A0} = \frac{(A0 - At)}{A0} \times 100\%
\]

\[
-ln(A0/At) = Kt
\]

where A0 is the initial absorbance of the dye, At is the absorbance of the dye at different time points, t is the irradiation time, and K is the rate constant for the photocatalytic reaction.

![Scheme 5. Chemical structures of the three types of dyes used for photocatalytic degradation.](image)

4. Conclusions

We demonstrated a one-step method for producing a photocatalyst-coated fabric for the self-sensitized photodegradation of dyes. In this method, TiO₂ particles are bound by a neoprene adhesive to a fabric. The neoprene not only modifies the surface of the TiO₂ with Cl but also dopes the TiO₂ lattice with Cl atoms that replace O in the lattice, introducing Ti³⁺ defects into the lattice. Cl on the surface of the TiO₂ particles introduces a negative surface charge to the particles that improves the adsorption of cationic dyes relative to undoped P25 TiO₂, while Cl doping in the lattice extends the absorption spectra of the TiO₂ particles into the visible region. In combination, these two effects enhance the photocatalytic degradation rates of dyes by composite-coated fabrics relative to degradation rates for P25-coated fabrics. We also showed that the photocatalyst can be recycled while on the surface...
of the fabric. These results suggest that this simple one-step method may prove to be a promising low-cost way for the mass manufacturing of photocatalysts for the treatment of industrial wastewaters.

Author Contributions: X.X. conceived and designed the experiments; Z.C. and T.Z. performed the experiments; X.X., Z.C. and T.Z. analyzed the data; X.X. and Z.C. wrote the manuscript. X.X., T.Z., P.R., D.C., Y.L., L.W. and B.Z. discussed the manuscript. Z.C. and T.Z. contributed equally. All authors have read and agreed to the published version of the manuscript.

Funding: This research was funded by the National Natural Science Foundation of China (Grant 21576016, 21521003), and the Fundamental Research Funds for the Central Universities.

Conflicts of Interest: The authors declare no competing financial interest.

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