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Influences of Doping on Photocatalytic Properties of TiO$_2$ Photocatalyst

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Abstract

As a kind of highly effective, low-cost, and stable photocatalysts, TiO$_2$ has received substantial public and scientific attention. However, it can only be activated under ultraviolet light irradiation due to its wide bandgap, high recombination, and weak separation efficiency of carriers. Doping is an effective method to extend the light absorption to the visible light region. In this chapter, we will address the importance of doping, different doping modes, preparation method, and photocatalytic mechanism in TiO$_2$ photocatalysts. Thereafter, we will concentrate on Ti$^{3+}$ self-doping, nonmetal doping, metal doping, and codoping. Examples of progress can be given for each one of these four doping modes. The influencing factors of preparation method and doping modes on photocatalytic performance (spectrum response, carrier transport, interfacial electron transfer reaction, surface active sites, etc.) are summed up. The main objective is to study the photocatalytic processes, to elucidate the mechanistic models for a better understanding of the photocatalytic reactions, and to find a method of enhancing photocatalytic activities.

Keywords: TiO$_2$, doping, photocatalytic properties, mechanism, carrier transfer

1. Background

As the best known photocatalyst, TiO$_2$ has attracted more attention and interest of many researchers due to its exceptional properties, such as high refractive index and ultraviolet (UV) absorption, excellent incident photoelectric conversion efficiency and dielectric constant, good photocatalytic activity, photostability, chemical stability, and long-time corrosion resistance as well as nontoxicity [1–4]. It has been widely used to solve a variety of environmental problems for the water-based solution utilization.
However, there are three key drawbacks for TiO$_2$ materials to limit their practical application. Firstly, one shortcoming is related to their large bandgap (3.0 eV for rutile and 3.2 eV for anatase TiO$_2$, respectively). It is well known that the photon absorption of semiconductors depends greatly on their bandgap energy. The photons can only be absorbed by the photocatalyst if the photon energies are higher than the semiconducting bandgap energy. Consequently, its surface photoactivation can be exclusively done under UV radiation ($\lambda \leq 390$ nm) or it only can respond and generate electron-hole pairs under UV light [5]. However, solar light consists of 5% UV light (300–400 nm), 43% visible light (400–700 nm), and 52% infrared light (700–2500 nm). That is, the UV light only occupies a small portion of the sunlight and a large part of solar energy cannot be utilized. In the future, one of the most urgent tasks is finding or modifying the photocatalysts with proper semiconducting bandgaps to maximize the absorption of solar energy. Secondly, a high recombination rate of electron-hole pairs is another disadvantageous effect on the photocatalytic efficiency for TiO$_2$ photocatalysis, resulting in a low quantum yield rate and a limited photooxidation rate. Thirdly, the weak separation efficiency of photocarries results in low photocatalytic activity. All three limitations induce great influence for a wide practical application.

To overcome above-mentioned problems, many studies have been conducted in the past 5 years based on the idea of extending the wavelength range of the photoactivation of TiO$_2$ photocatalysts towards visible light region and enhancing the utilization efficiency of solar energy. That is, an increased amount of energy from the solar light spectrum is well utilized. Recently, a path to achieve above-mentioned goal is represented by alien ion doping to improve the adsorption capacity and photocatalytic activity, such as self-doping [6–8], nonmetal doping [9–11], transitional metal doping [12, 13], and rare-earth metal doping [14, 15]. Usually, the doped ions introduce additional energy levels into the band structure, which can be used to trap electrons or holes to separate carriers from the bands, thus allowing more carriers to successfully diffuse to the surface. It has also been suggested that the required red shift of the absorption edge might be done by increasing the impurity ion concentration or increasing the oxygen defects in TiO$_2$. That is, the purpose of doping is relatively straightforward: modifying its large bandgap and electronic structure to optimize its optical properties for visible light harvest, improving each step in the charge kinetics to reduce the massive recombination of photogenerated carriers, and improving the interface and surface characteristics [16]. On the contrary, the photocatalytic activity depends strongly on the exposed crystalline faces [17, 18]. Given that different crystal surfaces have different surface energy levels for the conduction band (CB) and valence band (VB), such differences in the energy levels will drive the electrons and holes to different crystal faces. Obviously, doping can effectively modulate the lattice face structures.

To make clear the original question about the photocatalytic mechanism for doped TiO$_2$, it is important to explain and understand the doping process, doping method, and defect chemistry, especially the physics of the energy levels induced by the incorporation of dopants. This chapter comprises the following four sections to elaborate above concerns and recent progresses:

1. Ti$^{3+}$ self-doping in TiO$_2$ crystals and its photocatalytic mechanism,
2. Nonmetal doping in TiO$_2$ crystals and its photocatalytic mechanism,
3. Metal doping in TiO$_2$ crystals and its photocatalytic mechanism, and

2. Self-doping

It is widely accepted that the degradation of pollutants in water proceeds by oxidation either reacting directly with generated holes or indirectly with OH free radicals [19, 20]. Moreover, the photocatalytic activity of TiO$_2$ is often dependent on the nature and density of surface defect sites. Usually, incorporating dopants into TiO$_2$ crystals at the oxygen (O) and/or titanium (Ti) sites can generate midgap states [21]. Furthermore, interactions of photogenerated carriers with impurities in TiO$_2$ can also alter the electric structure and energy band structure, which enhances the photocatalytic performance [22]. Of course, excess doping also increases crystal defects (for example, oxygen vacancies, titanium vacancies, and interstitial titanium), thermal instability, carrier trapping, and carrier recombination centers [23, 24]. In other words, system-charge equilibria and/or geometric structure optimization should be considered during the doping process, especially the doping level or doping concentration and dopant distribution.

Recently, both experimental results and theoretical predictions have demonstrated that Ti$^{3+}$ self-doped TiO$_2$ could obviously increase concomitant intrinsic oxygen vacancies in TiO$_2$ crystals because the surface chemistry of nonstoichiometric TiO$_2$ containing Ti$^{3+}$ differs markedly from that of perfect TiO$_2$. Different from traditional impurity incorporation, Ti$^{3+}$ self-doping has been reported as an effective way to extend the visible light absorption of TiO$_2$, which can avoid the mismatching of atomic diameters with other foreign elements [25–28]. Self-doping can easily modulate and realize the system-charge equilibria just through synthesis method, process control, and raw material selection. Theoretical calculations evidence that interstitial Ti can cause impurity energy levels at 1.23 to 1.56 eV below the CB, whereas vacant Ti just makes impurity energy levels above the VB [29, 30]. Usually, the two defects can result in extra shoulder absorption or a tail absorption, which is the root cause for the enhancement of photocatalytic performance [31, 32].

2.1. Preparation of Ti$^{3+}$ self-doping

The basic process of self-doping includes that the electrons can be trapped and tended to reduce Ti$^{4+}$ cations to Ti$^{3+}$ state [33], the holes oxidize O$^2-$ anions for the formation of O$^-$ trapped hole or even O$_2$ gas, and the charge transfer steps are as follows:

$$\text{TiO}_2 + \text{h}^+ \rightarrow e^-_{\text{CB}} + h^+_{\gamma \beta}$$  \hspace{2cm} (1)

$$e^-_{\text{CB}} + \text{Ti}^{4+} \rightarrow \text{Ti}^{3+} \text{(trapped electron)}$$  \hspace{2cm} (2)
Subsequently, many methods are developed according to the above mechanism, such as hydrothermal method [25, 34–37], solvothermal method [38, 39], metallic reduction method [28, 40], solution-based oxidative method [41, 42], solution-based reduction method [43, 44], ionothermal method [26], vapor-fed aerosol flame synthesis [27], combustion method [33], and evaporation-induced self-assembly (EISA) method [45].

The hydrothermal method is widely used to synthesize Ti$^{3+}$ self-doped TiO$_2$ compared to other methods. Wang et al. changed hydrothermal conditions to treat a gel precursor and obtained Ti$^{3+}$ self-doped TiO$_2$ nanoparticles [34]. They found that the contents of Ti$^{3+}$ and oxygen vacancy (O$_v$) in TiO$_2$ crystals could be reasonably adjusted. Xin et al. demonstrated that the oxidation-based solvothermal synthesis of Ti$^{3+}$ self-doped anatase TiO$_2$ was an effective strategy to prepare uniform Ti$^{3+}$ self-doped anatase TiO$_2$ nanocrystals. Both the concentration and location of the Ti$^{3+}$ defects could be well managed by simply controlling the annealing temperature. This temperature-mediated management of the location and concentration of Ti$^{3+}$ defects was achieved through a Ti$^{3+}$ reversible diffusion mechanism [38]. The metallic reduction method was also used to synthesize Ti$^{3+}$ self-doped TiO$_2$ with dominant (001) facets, and the presence of Zn obviously caused the formation of Ti$^{3+}$ ions coming from the reduction of Ti$^{4+}$ [40].

Liu et al. successfully synthesized anatase Ti$^{3+}$ self-doped TiO$_2$$_x$ nanoparticles by a simple interface ion diffusion-redox reaction, and the resulting Ti$^{3+}$ self-doped TiO$_2$$_x$ had high crystallinity and showed enhanced visible light-driven photocatalytic oxidation [41]. Tian et al. used NaBH$_4$ as a reduced source and prepared TiO$_2$ nanobelts, and the theoretical calculations and experimental results indicated that the oxygen vacancies and Ti$^{3+}$ ions were successfully formed by reduction [43].

2.2. Mechanism of enhanced photocatalytic

It is well known that the Fermi level is much closer to the CB tail at a high oxygen vacancy concentration. Therefore, it is reasonably deduced that the higher oxygen vacancies can lead to an enhanced absorption of photon energy below the direct bandgap. The midgap states below the CB edge turn broad at an enhanced oxygen vacancy concentration (Figure 1). Meanwhile, the band of defect states resulting from oxygen vacancies is close to the CB edge, allowing photogenerated electrons to easily exchange between two bands. Namely, the electrons from the VB can easily transfer to the oxygen vacancy level under visible light irradiation. In other words, the electron transfer takes place from both VB and oxygen vacancy level localized states to the tailed CB. Consequently, the onset of optical absorption of the TiO$_2$ nanocrystals is lowered to 900 to 1100 nm [46–48].

\[
\begin{align*}
h_{vb}^+ + O^{2-} &\rightarrow O^- \text{ (trapped hole)} \quad (3) \\
4h_{vb}^+ + 2O^{2-} &\rightarrow O_2 \quad (4)
\end{align*}
\]
Many theoretical data from calculation and simulation also confirm that the local electrostatic balance is broken when host Ti⁴⁺ ions are reduced to Ti³⁺ ions, and oxygen vacancies are introduced because of charge compensation [49–51]. Because the effective charge of the oxygen vacancy is positive, the central Ti³⁺ is expected to shift away from the oxygen vacancy, forming a special sublevel electric state. Conversely, the shift Ti³⁺ also forces the four O²⁻ ions to move towards the oxygen vacancy to keep the electrostatic balance. The electrons can be photoexcited to CB under the irradiation of visible light. Meanwhile, the oxygen vacancies can inhibit the photogenerated electron-hole recombination. Furthermore, those Ti³⁺ ions act as hole traps and suppress the recombination of carriers and therefore extend the lifetime of the charges. That is, the higher light absorption in Ti³⁺ self-doping may come from the strong distortion of the outer orbitals of the Ti³⁺ ions.

In other words, Ti³⁺ self-doped TiO₂ exhibits a remarkable activity and enhanced performance as a photocatalyst. However, it is difficult to implant Ti³⁺ into TiO₂ crystals in practical application because Ti³⁺ species are usually unstable and can be easily oxidized by O. Therefore, it is still challenging to develop a simple and phase-controlled method to synthesize stable Ti³⁺ self-doped TiO₂ photocatalysts in the future. Moreover, it is important to have a comprehensive understanding of the methods and the techniques of Ti³⁺ generation and monitoring as well as Ti³⁺ property exploration.

**Figure 1.** Schematic diagram of a proposed photocatalytic mechanism of Ti³⁺ self-doped TiO₂ for the visible-light response [27].
3. Nonmetal doping

Nonmetal elements with high ionization energies and high electronegativity, such as nitrogen (N) [52–54], carbon (C) [55, 56], boron (B) [57–59], sulfur (S) [60, 61], fluorine (F) [62–64], and chlorine (Cl) [65, 66], are an efficient strategy to enhance the visible light photocatalytic activity, which results in higher photocatalytic activity in the visible light region owing to the bandgap narrowing and the shift of absorption edge.

The basic process is that the nonmetal dopants influence the VB through interaction with the O 2p electrons. The localized states or p states of nonmetallic dopants generally form the impurity levels and lie above VB, which extends the optical absorption edge of TiO2. On the contrary, nonmetal dopants within a surface can exist as isolated atoms rather than clusters. Consequently, the distribution of dopant states is above the VB maximum, which has greater potential for realizing visible light photoactivity.

3.1. N doping

Among all nonmetal elements, N element has been proven to be one of the most efficient dopants for visible light-responsive TiO2 photocatalyst. Since Asahi et al. made a breakthrough work in 2001 and found that doped TiO2 with N could enhance its photocatalytic activity for the photodegradation of methylene blue under visible light irradiation [67]. Many theoretical calculations and experiments have demonstrated and confirmed that N is one of the most promising dopant candidates for red shift of the absorption edge so far. Some authors suggested a model in which the incorporation of N via O substitution results in bandgap narrowing due to the mixing of the N 2p and O 2p states, which shows a remarkable red shift of the spectrum onset [68, 69]. In most of the reported studies, only N dopant concentration below 1 at.% are mentioned, and surface photoactivation with visible light is less effective compared to UV [70–72]. In other words, N-doped TiO2 is a promising candidate photocatalyst for enhanced light harvest in the visible region because it has strongly localized N 2p states (0.3–0.5 eV) at the VB maximum [73].

3.1.1. Preparation

The formation mechanism of N-doped catalysts obtained via different preparation methods is usually different. Moreover, N doping plays an important role on the exposed high-energy facets to a certain extent, which is apparently influenced by different preparation methods.

Generally speaking, there are two kinds of processes to prepare N-doped TiO2. One process can be ascribed as one-step direct incorporation of N atoms into TiO2 lattice, such as sol-gel method [74–76], chemical vapor deposition (CVD) [77, 78], atomic layer deposition (ALD) [79–81], hydrothermal method [82–84], solvothermal method [85–88], sol-hydrothermal process [89], hydrolysis-precipitation process [90], bioprocess-inspired method [91], electrochemical method [92–94], ion implantation [95, 96], combustion method [97–99], mechanochemical method [100, 101], low-temperature direct nitridization method [102], and microwave-assisted method [103].
Samsudin et al. synthesized undoped and N-doped TiO$_2$ via a sol-gel technique using Ti(IV) isopropoxide and triethylamine as the Ti and N precursors, respectively [74]. N was doped interstitially forming Ti-O-N or Ti-N-O linkages, and induced local states 0.23 to 0.26 eV above the VB, which was responsible for the visible light response between 400 and 550 nm. Gao et al. prepared N-doped TiO$_2$ films by the dielectric barrier discharge enhanced CVD method using Ti tetraisopropoxide and NH$_3$ as Ti precursor and doping gas [77]. It was found that N doping sources changed the growth orientation, affected the surface microstructure, narrowed the bandgap, and improved the photocatalytic activity in the visible light region. Liu et al. conducted an ALD method to fabricate N-doped TiO$_2$ hollow fibers with polysulfone fibers as a template [79]. The results showed that N was successfully inserted into the anatase TiO$_2$ lattice to form impurity levels above the VB top that narrowed the bandgap.

Therein, the hydrothermal method and solvothermal method are most frequently used because of their low cost, perfect crystallinity, and good repeatability. Wang et al. prepared N-doped TiO$_2$ nanoparticles by a facile one-pot hydrothermal treatment in the presence of L-lysine, and the results showed that N-TiO$_2$/C nanocomposites increased absorption in the visible light region and exhibited a higher photocatalytic activity than pure TiO$_2$, commercial P25, and previously reported N-doped TiO$_2$ photocatalysts [82]. Li et al. prepared a series of N-doped anatase TiO$_2$ samples using a solvothermal method in an organic amine/ethanol-water reaction system [85]. Both the degree of N doping and oxygen vacancies made contributions to the visible light absorption of the sample.

Recently, Cheng et al. synthesized N-doped TiO$_2$ nanoparticles through a hydrolysis-precipitation process using ammonia water as the doping species [90]. They found that the light absorbance edge of N-doped TiO$_2$ nanoparticle was obviously red-shifted to visible light region and the separation rates of photogenerated carriers were greatly improved. Further analysis implied that the VB maximum of O 2$\pi$ was 2.3 eV. Hu et al. also developed a facile low-temperature direct nitridization method to synthesize colloidal N-doped TiO$_2$ nanocrystals in triethylamine solution during the hydrolysis of tetrabutyl titanate followed by acidic peptization at 70°C [102]. The N-doped TiO$_2$ exhibited higher photocatalytic activity both in the UV and visible light regions in contrast to the undoped TiO$_2$ because of the improved light response in the range of 400 to 500 nm, narrowing bandgap, more production of e$^-$-h$^+$ pairs, and inhibiting recombination of the photo-induced carriers.

The other important preparation process of N-doped TiO$_2$ can be ascribed to two-step oxidation of Ti nitride, such as sputtering method [104, 105], thermal annealing [106–108], and plasma-enhanced microarc oxidation [109]. As for two-step methods, there are less research compared to one-step methods due to its higher cost and more complex technique process. Abadias et al. thought that the visible light activity of these materials mainly depended on the location of this impurity states and the microstructure of the N-doped TiO$_2$ materials [104]. They synthesized N-TiO$_2$ films through reactive magnetron sputtering method under a mixture gas of argon (Ar), N, and O. N diffusion was suggested to be responsible for the more complex crystallization and a better photocatalytic activity. Ha et al. prepared a new type of N-doped TiO$_2$ mesoporous inverse opal structure via heat treatment in the presence of an N-rich precursor, and the N doping with 9.4 wt.% concentration narrowed the bandgap from 3.2
to 2.4 eV, which corresponded to light absorption at wavelengths as long as 520 nm [106]. Jiang et al. developed a novel plasma-enhanced microarc oxidation process for preparing a high concentration substitutional N-doped TiO$_2$, compared to the traditional thermal annealing, and the process provides a possibility to increase the N doping concentration up to 3.21 at.% in TiO$_2$N$_x$, which exhibits a significant red shift in the bandgap transition and narrows bandgap to 2.6 eV [109].

3.1.2. Photocatalytic mechanism of N-doped TiO$_2$

The mechanism, especially the origin of visible light photoactivity for N-doped TiO$_2$, is still in debate up to now. Valentin et al. [111] and Gao et al. [101] studied substitutional and interstitial N impurities in bulk anatase TiO$_2$ lattice (Figure 2). They thought that the N atom replaced lattice O in TiO$_2$ for the substitutional case and showed positive oxidation state ranging from hyponitrite species (NO$^-$), nitrite (NO$_2^-$), to nitrate species (NO$_3^-$). The visible light responses for substitutional N-doped anatase TiO$_2$ came from occupied N 2$p$ localized states, which were slightly above the VB maximum. The N-O bond showed localized π-bond states. The two bonding states were found in deep energy level and lay below the top of the O 2$p$ band. The other two antibonding states were found located over the O 2$p$ band. In other words, the localized states were responsible for the excellent optical adsorption when the N atom was located at the substitutional site. On the contrary, interstitial site for N doping had some disadvantageous effects on the photocatalytic reaction because of the hole trapping.

![Figure 2. Supercells of (a) pure, (b) substitutional, and (c) interstitial TiO$_2$ [111].](image)

Energy bandgap models also predict that substitutional (interstitial) N generates shallow (deep) midgap states above the top of the VB due to the mixing of N 2$p$ and Ti 3$d$ orbitals (Figure 3). Usually, N doping is associated with the formation of oxygen vacancies and electrons to maintain lattice neutrality. From this perspective, it has been suggested that the presence of extra oxygen vacancies might be the key factor allowing for better photocatalytic performances to be achieved, which causes a red shift of the absorption band edge due to a positive shift of VB and improves electron injection [112–114]. Under visible light irradiation, electrons of the localized N 2$p$ states can be excited up to the individual CB, leaving holes on
the localized states. The energy barrier between midgap states would suppress electron transfer. This can also inhibit the recombination of photoelectrons and holes to some extent [115]. In other words, the efficient separation of photo-induced carriers under visible light excitations enables N-doped TiO$_2$ to have higher photocatalytic activity in a wide wavelength ranging from UV to the visible region.

![Figure 3. Schematic illustration of electron migration in anatase/brookite structures: under visible light irradiation (path a) and under UV light illumination (path b) [114].](image)

3.2. C doping

Among the nonmetal elements, C doping has also been proposed as one of the best candidates [116]. The origin of C doping is mainly incorporated interstitially, narrowed bandgap and resulted in a red shift in the UV-visible (UV-vis) spectrum, resulting in an increase in adsorption of near-infrared (NIR) spectrum [117, 118]. Khan et al. first reported that C-doped TiO$_2$ was visible light active, and found that that the three kinds of C species could influence the photocatalytic activity, namely, elemental C, carbonate species (C-O bond) as interstitial dopant, and C substituted for O (Ti-C bond) in TiO$_2$ lattice [119].

Subsequently, a lot of researches were conducted to synthesize C-doped TiO$_2$ photocatalysts [120–124]. Shi et al. prepared nanometer-sized C-doped TiO$_2$ nanoplates with exposed [001] facets via the hydrothermal treatment of TiC powder in an HF-HNO$_3$ mixed aqueous solution and found that C doping also obviously presented red shift absorption edge towards visible
Further analysis found that the electrons in the localized C states could be excited to the CB under visible light irradiation and effectively scavenged by molecular O to produce the superoxide radical anion (•O$_2^-$) and hydrogen peroxide (H$_2$O$_2$), which could interact to produce hydroxyl radicals (•OH). Recently, Shao et al. prepared amorphous C-doped TiO$_2$ with visible light photocatalytic activity by a facile sol-gel route for the first time. The results also indicated that the most active sample with oxygenic groups had a narrower bandgap and lower recombination of electron hole, significantly broadening its potential for many practical applications [122].

About the mechanism of photocatalytic activity for C-doped TiO$_2$ catalysts, it is similar to N-doped TiO$_2$ catalysts, except some little differences. Liu et al. investigated that C doping in the TiO$_2$ lattice had graphite- and carbonate-like species at interstitial positions [125]. They thought that the two C states could narrow the bandgap of anatase TiO$_2$, serving as a photosensitizer to absorb visible light and promote the charge carrier separation, which enhanced the visible light photocatalytic activity remarkably. Zhang et al. insisted that the substitutional C (C$_s$) at O sites modified the electronic band structure of TiO$_2$ by mixing C 2p orbitals with O 2p orbitals, resulting in bandgap narrowing [123]. Under visible light irradiation, electrons can be excited directly into the CB and transferred to the adsorbed O molecule to produce •O$_2^-$ and subsequently •OH with strong oxidation power. Recently, Sun and Zhang investigated the C doping of anatase TiO$_2$ in detail using density functional theory (DFT) calculation [126]. They found that C$_{\text{Ti}}$ and C$_i$ with a shallow donor had no effect on the bandgap, whereas the (CO)$_{2V}$, C$_{\text{V}}$-V$_{\text{O}}$, and (C$_2$)$_{2O}$ defect reduced the bandgap to about 1.1, 1.7, and 1.4 eV, respectively. Based on the electronic structure analysis of the C defects, they excluded the possibility of C occupying Ti sites or interstitial sites to be responsible for the enhancement of photoactivity in the visible light region.

### 3.3. S doping

S doping in TiO$_2$ structure also shows bandgap narrowing effects [127]. However, its large ionic radius makes it difficult to incorporate into TiO$_2$ crystals due to the large formation energy. Therefore, the key issues are to find a facile, low-cost, and stable process to synthesize S-doped TiO$_2$.

Goswami and Ganguli developed a novel approach to synthesize sulfated TiO$_2$ nanoparticles using 15% Ti trichloride and thiophene in the presence and absence of oxalic acid [128]. They found that S-doped samples had higher surface area, smaller crystallite size, greater thermal stability, and better photocatalytic performance. Li et al. demonstrated a facile nonhydrolytic thermolysis route for monodisperse S-doped TiO$_2$ nanocatalysts in hot boiling organic solvents of oleic/oleyl-amine/1-octadecene [129]. Compared to the undoped TiO$_2$ nanocatalysts, S-doped TiO$_2$ nanocatalysts presented obviously enhanced visible light activation for the degradation of rhodamine B and methylene blue dyes under the artificial visible light irradiation. Recently, Ramacharyulu et al. synthesized S-doped TiO$_2$ catalysts by a sol-gel process followed by hydrothermal treatment at low temperature and tested for catalytic activity by natural sunlight photocatalytic degradation of a toxic chemical warfare agent [130]. It was observed that S-doped TiO$_2$ exhibited superior photocatalytic activity under sunlight irradiation.
tion. Further analysis indicated that the superior photocatalytic activity could be attributed to the presence of $S^{4-}/S^{6+}$ or $N$ impurity levels. Lin et al. [131] and Sharotri and Sud [131, 132] further found the importance of $S^{6+}$ for $S$-doped TiO$_2$ photocatalysts, which could form $h^+/e^-$ trapping centers, delay the phase transition from anatase to rutile, promote photocatalytic activity, and prevent $h^+/e^-$ recombination.

3.4. Halogen doping

Halogen doping in TiO$_2$ crystals is another important approach to improve photocatalytic performance, such as F doping [133–135], Cl doping [136–138], and iodine (I) doping [139–141]. Pan et al. developed an effective bottom-up synthesis strategy to prepare monodisperse F-doped TiO$_2$ mesoporous spheres by integrating sol-gel and solvothermal processes [133]. The photocatalytic experiments showed that the formation of surface fluorination was helpful to improve and enhance light harvesting in the UV-vis range. Fang et al. also synthesized F-doped rutile TiO$_2$ with tunable solar absorption via one-pot hydrothermal method [142]. They found that the optical bandgap of the catalyst could be easily manipulated from 3.05 to 2.58 eV through altering the initial F:Ti molar ratio.

The origin of the visible light activity of F-doped TiO$_2$ has been systematically investigated by many researchers via comprehensive theoretical and experimental studies [134, 143–145]. It is widely accepted that the three-coordinated surface F atoms with higher 1s binding energy are identified to be the origin of the visible light activity. The surface group can also trap the CB electrons by tightly holding electrons due to the strongest electronegativity of F. The calculated results using DFT principle show that F implantation resulted in Ti$^{3+}$ self-doping and contributes to the enhancement photocatalytic activity. Moreover, the strong electron-withdrawing ability of the surface three-coordinated surface F also reduces the recombination of photogenerated electrons and holes.

Similarly, Cl doping has the same role in TiO$_2$ crystals with F doping. Wang et al. prepared Cl-doped TiO$_2$ nanocrystalline via a simple single-step method by sonicating a solution of tetraisopropyl titanate and sodium chloride in water/ethanol at 70°C and found that the Cl doping of TiO$_2$ shifts the absorption edge toward a higher wavelength [136]. The photodegradation rates of butyl benzyl phthalate reached 92% under visible light irradiation for 240 min, which was much higher than undoped TiO$_2$.

Doping with I element in TiO$_2$ crystals also can enhance light absorption and decrease recombination [146–149]. Lin et al. synthesized I-modified TiO$_2$ nanocrystallities through a combination of sol-gel process and solvothermal process in the presence of HI solution [146]. The results showed that the I in the form of I$_2$ was responsible for the visible light response. Liu et al. developed a new strategy for homogeneous doping of I molecules (I$_2$) to achieve bandgap narrowing of TiO$_2$ nanosheets and investigated the extension of the intrinsic absorption edge into the visible light region through a shifting of the VB maximum [149]. Importantly, the geometric structure of the host retained its integrity. The experiment together with first-principles calculations revealed the molecular nature of adsorbed I atoms and implied that the mechanism of electronic structure modulation in the TiO$_2$ layers changed depending on the concentration of I$_2$ molecules.
3.5. Other doping

B$^{3+}$ is incorporated both substitutionally and interstitially in TiO$_2$ crystals. Usually, the replacement of B$^{3+}$ for Ti$^{4+}$ generates one hole in O $2p$ orbital, which can also be accompanied by a blue shift of the UV-vis absorption spectrum and an increase in oxygen vacancies [150, 151]. Xu et al. developed a new method allowing a clean one-step synthesis to obtain B-doped TiO$_2$ for the first time [150]. B doping resulted in a shift of the absorption edge up to 460 nm with a concomitant reduction of the bandgap energy. The narrowing bands were attributed to ionized oxygen vacancies and defect states in anatase for the presence of interstitial B tricoordinate and tetracoordinate to O.

Another concerned nonmetal dopant is phosphorus (P). Many papers indicated that P dopant could also improve the photocatalytic activity [152, 153]. Zheng et al. synthesized a novel thermally stable P-doped TiO$_2$ by liquid hydrolysis of TiCl$_4$ using hypophosphorus acid as the precursor of the dopant [152]. They found that increased surface P content led to a linear enhancement of the specific adsorption capacity of methylene blue because of the Coulombic attractive force between the cationic dye and the negatively charged P-doped TiO$_2$ surface.

4. Metal dopants

It is well known that the photoexcitation of TiO$_2$ catalysts involves excitation, diffusion, and surface transfer of photogenerated carriers. Therefore, the carrier’s lifetime is vital in determining the photocactivity during the photodegradation process. Usually, the surface properties of TiO$_2$ intrinsically could be influenced by the preparation method, process, and doping, which determines the surface separation and transfer of charge carriers by generating surface states where electrons and holes are spatially trapped and transferred for subsequent redox reactions.

Compared to nonmetal dopant, substitution of metal ions can introduce an intraband state close to CB edge, which results in an obvious red shift in bandgap adsorption due to sub-band gap energies, such as transition metal doping [154–156], rare-earth metal doping [157–159], and other metal doping [160–162].

However, the diffusion of metal atoms is difficult in solid materials under low temperature. Consequently, it leads to inhomogeneous distributions of dopants and limited depth near a subsurface region. To obtain metal-doped TiO$_2$ nanoparticles with good homogeneity, sintering has to be conducted at a high temperature, which leads to particle agglomeration. Moreover, the metal dopants also provide more trapping sites for electrons and holes compared to nonmetal dopants. Furthermore, electron trapping occurs at a much faster process compared to hole trapping. In other words, trapping an electron or a hole is always ineffective for carrier separation because immobilized charge species rapidly recombine with its mobile counterparts. Last, metal doping has also some other drawbacks, especially thermal instability, which reduces the repeatability of TiO$_2$ photocatalysts. That is to say, it has a detrimental effect if the process is not under the control. Therefore, many researchers focused on the resolution of the above four problems in recent several years.
4.1. Transition metal element doping

Various properties of transition metal with 3\textit{d} or 4\textit{d} electron structure are influenced by many factors, such as the number of \textit{d}-electrons on transition metal ions, crystalline structures, oxygen defects, and preparation methods [163–165]. Usually, the bandgap energy and band positions, Fermi level, and \textit{d}-electron configuration of the electronic structure in TiO\textsubscript{2} can be effectively modulated when transition metal ions were introduced into TiO\textsubscript{2} lattice. Subsequently, it forms a wide range of new energy levels below CB arising from their partially filled \textit{d}-orbitals, which results in an obvious red shift in bandgap and increases its visible light harvest [166, 167]. In addition, transition metal ions alter the carrier equilibrium concentration by serving as electron-hole trapping, suppress the recombination rate of electron-hole pairs, and enhance the degradation rates. That is, the photocatalytic performance for TiO\textsubscript{2} can be effectively improved through transition metal doping.

However, transition metal ions-doped TiO\textsubscript{2} appears to be a complex function of dopant concentration, energy level of dopant within the lattice, the \textit{d} electron configuration, distribution of dopants, electron donor density, and incident light intensity. Every transition metal ion has different \textit{d} electron configurations and its own characteristics. Therefore, it is important to further elaborate the process for every transition metal ion.

4.1.1. Iron (Fe) doping

Fe-doped TiO\textsubscript{2} shows superior activity due to its unique half-filled electronic configuration and shallow trapping compared to other metal dopants with closed shell electronic configuration, which can be more effective to influence the photoactivity [168–176]. Theoretical and experimental studies show that Fe doping can effectively reduce the trapping density and charge recombination, resulting in drastically improved adsorption.

Manu and Khadar synthesized Fe-doped TiO\textsubscript{2} nanocrystals at different atomic ratios through the hydrothermal method [168]. They found that the concentration of Fe dopants was more near the grain boundary because the dopant atoms were incorporated into the lattice at substitutional positions. The energy level associated with the peak at 2.63 eV was the deepest defect level. Consequently, the photocatalytic activity was enhanced greatly. Yan et al. developed a facile fast hydrolysis route to prepare a three-dimensional flow-like Fe-doped rutile TiO\textsubscript{2} nanostructure and investigated the relation between Fe doping and crystal planes [169]. Because the ionic radius of Fe was smaller than that of Ti, and the corresponding interplanar spacing distance of TiO\textsubscript{2}, (110) was reduced with the replacement of Ti atoms by Fe atoms, as reflected by the shift of the (110) diffraction peak, which was similar with Liu and Zhang’s result [170]. With Fe species doping into both the bulk phase and the surface, the bandgap narrowing of rutile TiO\textsubscript{2} was realized and the dissociative adsorption of water on the surface was promoted, which accordingly led to greatly enhanced activity in visible light-driven water oxidation.
4.1.2. Chromium (Cr) doping

Cr doping of TiO$_2$ also leads to a clear red shift in the UV-vis absorption spectrum, evidencing a decrease in bandgap and VB shift. Moreover, the Fermi level is also shifted to a higher energy by $\sim 0$ to $1$ eV. With the change of preparation method and doped concentration, Cr-O antibond orbitals located slightly below the CBM in the region of $\sim 3.0$ to $2.2$ eV and the unsaturated nonbonding $d$ orbitals located in the middle of the gap region of $\sim 2.5$ to $1$ eV. It also is suggested that $\text{Cr}^{3+}$ doping is attributed to the increase of conductivity due to more free charges [177–180]. Li et al. synthesized Cr-doped TiO$_2$ via a hydrothermal method and found that $\text{Cr}^{3+}$ ions could replace the Ti atoms in the lattice with oxygen vacancy compensation [177]. It was interesting that Cr doping also could prolong the lifetime of photogenerated carriers because the doped $\text{Cr}^{3+}$ ions might act as the recombination centers of carriers.

4.1.3. Niobium (Nb) doping

Over the past few years, Nb-doped TiO$_2$ has received special attention due to its excellent electrical conductivity at room temperature. The ionic radius of Nb$^{5+}$ of 0.064 nm is slightly larger than Ti$^{4+}$ of 0.0605 nm. Therefore, Nb$^{5+}$ works as an n-type dopant in TiO$_2$ lattice and generates additional carriers in its CB, which can notably increase photocatalytic performance [181–186]. Joshi et al. successfully prepared Nb-doped TiO$_2$ transparent films on glass substrates using a nonaqueous sol-gel spin coating technique [181]. Photocatalytic experiments showed that the films with 12 at.% Nb doping had excellent photocatalytic activity with 97.3% degradation of methylene blue after 2 h of UV irradiation. Archana et al. reported the high electron mobility and optical transparency of Nb-doped TiO$_2$, giving rise to the enhancement in charge transport behavior after 2 at.% Nb doping, which could assist the speedy initial reaction of the organic decomposition process and enhance the overall photocatalytic activity [182].

4.1.4. Other transition metal doping

Except above transition metal dopants, other transition metal atoms, including tungsten (W) [187–191], silver (Ag) [192–197], copper (Cu) [198, 199], cobalt (Co) [200–202], tantalum (Ta) [203–205], molybdenum (Mo) [206–208], zinc (Zn) [209–213], manganese (Mn) [214, 215], nickel (Ni) [216, 217], and vanadium (V) [218, 219], are also investigated by many researchers. Recently, considerable enhancements of photocatalytic activity with W-doped TiO$_2$ in aqueous systems for the degradation of organic compounds have also been reported [187–191]. Liu et al. synthesized ordered mesoporous crystalline TiO$_2$ with various W doping level using SBA-15 as a hard template [187]. The existence of W ions expanded the range of useful excitation light to the visible spectra, greatly inhibited the recombination of electron-hole pairs on mesoporous TiO$_2$, and played an important role in improving the photocatalytic activity in the visible light region. The enhanced photocatalytic activity for W-doped TiO$_2$ could also be especially attributed to the presence of much higher Lewis surface acidity of a W-doped TiO$_2$ surface with a higher affinity for chemical species having unpaired electrons than pure TiO$_2$. 
Ag is another important dopant in TiO₂ crystals and Ag doping can also cause an obvious shift towards narrowing the bandgap. Zhang et al. successfully synthesized Ag-doped TiO₂ with a novel hierarchical architecture via a combination of an electrospinning method and a hydrothermal process [192]. They found that Ag doping played a great role in the photocatalytic activity. Electron transfer to the Ag sites reduced the carrier recombination rate and allowed a more effective reaction between the surface trapped holes and electrons. However, excessive Ag dopant could also cause a decrease in the activity of TiO₂ because they occupied the active sites on the surface of the TiO₂. Moreover, the photogenerated electrons on the Ag sites attracted holes and recombined together.

Cu, as a kind of transition metal, is also found to be an effective dopant for TiO₂ to enhance the photocatalytic activity. Sajjad et al successfully prepared mesoporous Cu-doped TiO₂ via a sol-gel method at low temperature using water-immiscible room-temperature ionic liquid organic materials as a template and an effective additional solvent [198]. The results showed that Cu-doped TiO₂ samples exhibited superior visible light photocatalytic activities compared to undoped TiO₂ and P-25 [199, 200]. Further characterization indicated that Cu⁺/Cu²⁺ sites at interfacial Ti-O-Cu surface linkages reduced bandgap energy as well as efficient charge separation due to the formation of Ti-O-Cu bonding at the surface and the associated appearance of oxygen vacancies, resulting in higher degradation rate.

Usually, Co dopant concentrations show weak influence on photocatalytic performance, whereas the oxygen vacancy concentration and distribution in the system show much stronger influence on the optical performance due to the shift of Fermi level up by ~0.75 eV [201]. Cai et al. presented a controllable and reliable method to synthesize Co-doped TiO₂ nanowires through a combining versatile solution phase chemistry and rapid flame annealing process [202]. They found an enhanced catalytic activity in Co-doped TiO₂ crystals. However, Co doping was also shown to drastically deteriorate performance at high doping concentration due to the formation of sub-band gap states that act as recombination centers.

Sengele et al. obtained Ta-doped TiO₂ via a sol-gel route and found that the Ta doping could induce significant modifications on the structural, morphological, surface, electronic, and optical properties of TiO₂ [203]. Total diethylsulfide elimination could be reached for 100 min under continuous contaminant flux before deactivation and the conversion maintained to 80% of degradation after 200 min, which was higher than undoped TiO₂ catalysts [204–208].

Recently, many researchers found that transition metal could effectively modulate the crystal face, which could improve photocatalytic performance [209–213]. Saad et al. synthesized Zn-doped TiO₂ nanowall with a (001) facet and porous structure [209]. They found that the samples with a (001) facet exhibited enhanced photocatalytic activity and Zn-doped TiO₂ had better degradation rates compared to other samples.

4.2. Rare-earth metal doping

Another path to achieve the abovementioned goal is represented by rare-earth doping [214–219]. Compared to transition metals, rare-earth metals with 4f, 5d, and 6s² states are considered as the ideal dopants to modify the crystal structure, electronic structure, and optical properties
of TiO$_2$, which can effectively influence the positions, widths, and density of states of CB and VB [220, 221]. Furthermore, rare-earth metals can form complexes with various Lewis-based organic compounds through interaction of the functional groups with their $f$ orbital, thereby improving the photoactivity. Last, the functional integration of upconversion luminescent rare-earth ions with photocatalyst provides a potential for wavelength conversion and efficient utilization of solar energy for this purpose [222, 223]. This approach appears to be a completely new alternative for enhancing the efficiency of the photocatalytic process. In other words, the applications of the upconversion process by phosphor-like systems can optimize the photocatalytic performance of traditional UV active photocatalysts.

Therein, lanthanide (La) has been widely used and investigated in the field of optical application (LED light, laser, photocatalytic, solar cell, etc.). Particularly, the ability of these nanomaterials to release high-energy photons after NIR laser-light stimulation allows deep tissue imaging and luminescent nanoparticles [224, 225]. Du et al. prepared pure and La-doped TiO$_2$ thin films via a sol-gel method using tetrabutyl titanate as Ti precursors [224]. The results showed that the content of La was the key factor for hydrophilic and photocatalytic activity. LaTiO$_3$ could be formed in La-doped TiO$_2$ thin films, which caused the TiO$_2$ lattice distortion and restrained the transition from anatase to rutile. By adding 0.3 wt.% La to the TiO$_2$ thin films, 92.02% methylene blue was finally degraded.

Cerium (Ce), as a typical rare-earth metal, is widely used to dope TiO$_2$ because of its unique electronic structure. The main feature of Ce ion doping is the different electronic structures of 4$f$ states, such as Ce$^{3+}$ with 4$f^15d^0$ and Ce$^{4+}$ with 4$f^05d^0$. The visible light photoactivity of Ce-doped TiO$_2$ nanoparticles is mainly due to the presence of 4$f$ level in the mid-bandgap of TiO$_2$ crystals, which leads to the optical absorption between 400 and 500 nm. The possible transition of $d$ and $f$ orbital electrons can also reduce the recombination rate of electron-hole pairs, thereby making TiO$_2$ more feasible for photocatalytic response under visible light [226–229]. Maddilla et al. prepared Ce-doped TiO$_2$ catalysts using a wet impregnation method [226]. The results indicated that Ce-doped TiO$_2$ exhibited an obvious red shift, reducing the bandgap and improving the photocatalytic efficiency. Photocatalyzed ozonation with 1at.% Ce/TiO$_2$ yielded 100% degradation in 2 h under basic pH conditions.

Among various upconverting nanomaterials, erbium (Er) can be excited by NIR or visible light [230–232]. Obregon and Colon synthesized Er-doped TiO$_2$ through a surfactant free hydrothermal method, which exhibited good photoactivities under sun-like excitation for the degradation of phenol [231]. The presence of Er$^{3+}$ did not affect the structural and morphological features of the TiO$_2$ significantly, whereas photocatalytic experiments clearly evidenced that Er introduction into TiO$_2$ matrix would promote the profiting of NIR photons, enhancing the photoactivity of the catalyst.

Gadolinium (Gd) doping raises VB maximum with respect to the Fermi level and thus turns the intrinsic virgin phase into a p-type semiconductor. Choi et al. synthesized three-dimensional Gd-doped TiO$_2$ nanofibers using a simple electrospinning technique [233]. The pristine Gd-doped TiO$_2$ nanofibers showed a higher photocatalytic activity than the TiO$_2$ nanoparticles, which could be attributed to the fast electron transport. In addition, Gd-doped TiO$_2$ nanofibers...
showed nearly five-fold enhancement in the photocatalytic degradation rate due to synergistic higher electron transport.

Europium (Eu) is also a good candidate for upconversion materials and is widely used in the field of photocatalyst, laser, solar cell, etc. Recently, many researchers also attempted to dope Eu into TiO$_2$ lattice [234–236]. Leroy et al. prepared periodic mesoporous Eu-doped TiO$_2$ through the EISA process [234]. They found that strong fluctuations existed in the intensity of the $^5\text{D}_0 \rightarrow ^7\text{F}_2$ transition under UV light exposure. Correlation of the emission with the photocatalytic activity of the semiconductor for the photodegradation of an organic molecule could also be confirmed.

Yttrium (Y) doping usually shows weak influence on bandgap energy and does not alter the Fermi level. However, it can increase hole mobility to generate p-type TiO$_2$, which is helpful for the carrier transport. Wu et al. prepared a novel mesoporous Y-doped TiO$_2$ nanosheet array via a low-cost, facile, and template-free solvothermal method by employing waste tricolor fluorescent powder (WTFP) as a dopant Y source and acetic acid as a mesopore template [237]. The 2.5 and 5 wt.% Y-doped TiO$_2$ films showed higher photocatalytic activity for the methyl orange, with 82.6% and 81.3% reduction within 6 h irradiation, respectively. The enhanced photocatalytic activity of Y-doped TiO$_2$ film was attributed to the large surface area and the low electron-hole recombination rate as well as enhanced absorption of organic pollutants on the semiconductor surface. Li et al. also confirmed that ytterbium (Yb) doping was also a kind of route to improve the NIR photocatalytic degradation of rhodamine B [238].

4.3. Other metal dopants

Recently, many researchers found that lithium (Li) doping plays an important role in the inhibition of activity in the TiO$_2$ network acting as an electron trapping or hole trapping [239–241]. Bouattour et al. reported Li-doped TiO$_2$ nanoparticles through a sol-gel process and investigated their potential application as a photocatalyst for degradation of different organic compounds [240]. Results showed that an inhibition of activity is presented in the Li doping network. However, the surface accumulation of Li$^+$ on TiO$_2$ particles could favor better interfacial charge transfer.

Long et al. calculated the electronic structures of silicon (Si)-, germanium (Ge)-, tin (Sn)-, and lead (Pb)-doped anatase and rutile TiO$_2$ systematically using DFT calculations [242]. Doping with Si, Ge, Sn, and Pb elements also narrowed the bandgap of rutile TiO$_2$ due to a shift far away from the CB minimum. The reduction of the bandgap is 0.20 and 0.15 eV in Si- and Ge-doped anatase TiO$_2$, respectively. However, there were enlargements of 0.06 and 0.02 eV in the bandgap of Sn- and Pb-doped anatase TiO$_2$, respectively. They predicted that Ge-doped TiO$_2$ was efficient for the overall water splitting using visible light irradiation.

Doping with bismuth (Bi) ions can also decrease the bandgap of TiO$_2$ and thereby extend its absorption into the visible light region, enhancing its photocatalytic efficiency [243–245]. The impurity energy level from the hybrid of Bi 6s, O 2p, and Ti 3d orbitals is the cause of the red shift of the absorption edge. Therefore, Bi-doped TiO$_2$ systems are more effective for the photodegradation of organic pollutants under visible light irradiation, such as methyl blue.
and methyl orange. Wu et al. developed a facile method to prepare Bi-doped TiO₂ through hydrothermal synthesis followed by thermal annealing treatment [243]. Bi doping caused the formation of BiₓTiOₓ and reduced bandgap widths. Moreover, some special structural defects created by the migration of Bi³⁺ ions were also responsible for the high photoactivity. Bi-doped TiO₂ catalysts with high doping concentration (such as 5 and 10 mol%) showed the highest activity for the catalyzed photodegradation of methyl orange under visible light irradiation.

Sn doping is another important supplementary form to improve photocatalytic performance [246, 247]. Oropeza et al. investigated the influence of Sn doping on the anatase to rutile phase transition, as well as the photocatalytic performance [246]. They found that Sn-doped TiO₂ exhibited enhanced visible light region photocatalytic activity compared to undoped TiO₂ in dye degradation experiments, even it was higher than that of N-doped TiO₂. This was attributed to the narrowing of the bulk bandgap at low doping levels. The Sn surface states laid above the VB top and could therefore act as trapping sites for holes.

5. Codoping

Although the monodoped nonmetal or metal atoms can obviously enhance photocatalytic performance, they always act as the recombination centers because of the partially occupied impurity bands. It has been recognized theoretically that codoping using two or more foreign atoms can passivate the impurity bands and decrease the formation of recombination centers by increasing the solubility limit of dopants. Furthermore, codoping can also modulate the charge equilibrium. Consequently, codoping can effectively enhance the photocatalytic activity. Based on the research of doping effects, two or more elements are introduced into TiO₂ lattice to check the changes of electronic structure and bandgap energy, including nonmetal and nonmetal atoms [248–250], nonmetal and metal atoms [251–253], and metal and metal atoms [254–256].

Regardless of whether doping is based on single heteroatoms or coupled heteroatoms, the spectral distribution and localized states in the bandgap essentially determine the visible light absorbance and redox potential of the photo-induced charge carriers. Therefore, it is a challenge and hot topic to introduce different dopants, which can realize substantial synergetic effects.

5.1. Nonmetal-nonmetal doping

5.1.1. N-nonmetal codoping

To improve the photocatalytic performance of N-doped TiO₂ catalysts, the modification strategy has been extensively adopted and investigated by many researchers, such as N-nonmetal codoping, C-nonmetal codoping, and other nonmetal-nonmetal codoping. The modified N-doped TiO₂ usually showed favorable effects for improving the photocatalytic activity in the range of visible light compared to N-doped TiO₂.
Therein, the (N, S) codoping TiO$_2$ can play a vital role in significantly improving photocatalytic activity because of the strong synergistic interaction between S and N, in which the surface separation of photoexcited electron-hole pairs is promoted [257–259]. Consequently, the VB of TiO$_2$ shifts to a positive direction, which leads to higher oxidative ability and degradation ability towards pollutant under visible light irradiation. Etacheri et al. further proved that the formation of isolated S 3p, N 2p, and π* N-O states between the VB and CB was responsible for the visible light absorption [260].

Chung et al. synthesized the (N, S) codoped TiO$_2$ with an anatase phase using a simple solvothermal treatment and investigated their visible light photocatalytic activity associated with the thermal behavior [257]. They found that the (N, S) codoped TiO$_2$ had better visible light photocatalytic activity and adsorptivity than the commercially available P25. The S dopants effectively assisted the surface reaction by adsorbing cations of organic dyes on the codoped TiO$_2$ surface. The N dopants formed a delocalized state in the bandgap, which led to the enhanced solution bulk reaction by increasing visible light absorbance (Figure 4).

![Figure 4. Schematic illustration of the suggested effects of N and S dopants on the enhanced visible light photocatalytic activity of TiO$_2$.](http://dx.doi.org/10.5772/63234)

It is noteworthy that (N, S) codoping can also effectively modulate the preferential growth plane of TiO$_2$ crystals. Xiang et al. prepared (N, S) codoped TiO$_2$ nanosheets with exposed [001] facets by a simple mixing-calcination method using the hydrothermally prepared TiO$_2$ nanosheets as a precursor and thiourea as a dopants [261]. The first-principles DFT proved that the electrons could be easily excited in the impurity states and then migrated between VB and intermediate. These resulted in stronger absorption with a red shift in the bandgap transition.

Recently, Samsudin et al. used (N, F) codoping to improve the intrinsic properties of the TiO$_2$ catalyst and found that the (N, F) codoping not only introduced activity in the visible light region but also improved the performances of intrinsic TiO$_2$ itself. The high activity could be attributed to the presence of N and F, which resulted in the change in the morphology and increasing presence of [001] facets [262]. Subsequently, a great number of experiments and theoretical calculations were carried out to further discuss (N, F) codoping in TiO$_2$ crystals [263–267]. Rahul and Sandhyaranai synthesized three-dimensionally ordered (N, F) codoped
TiO$_2$ by templating with polystyrene colloidal photonic crystals using nitric acid and trifluoroacetic acid as raw materials and found an obvious red shift due to (N, F) codoping [263]. Further investigation showed that the enhancement of photocatalytic activity could be attributed to the bandgap scattering effect and the slow photon effect, leading to a significant improvement in solar light harvesting.

(N, B) codoped TiO$_2$ is also an effective approach to improve photocatalytic activity [268–270]. Usually, N and B atoms are interstitial species connected to the same O lattice. The energy level of the [NOB] species is located close to edge of the VB and lies below the corresponding level of [N,O] because of the electrostatic stabilization. In other words, the photocatalytic activities are influenced not by [NOB] center but by [NOB]$^-$ center located at higher energy level. Moreover, the presence of N and B species can effectively narrow the bandgap and inhibit the transformation of anatase TiO$_2$ to rutile phase [269].

Other N-nonmetal codoping includes (N, C) codoping [271, 272] and (N, H) codoping [273, 274]. Liu et al. exploited (N, C) codoped porous TiO$_2$ nanofibers by a combination of electrospinning and controlled calcination technologies [271]. The codoping of N and C in TiO$_2$ not only led to a shift of the absorption edge to lower energy by inducing new band levels but also created a large amount of single electron-trapped oxygen vacancies. Recently, Wei et al. demonstrated a new chemical approach to prepare black anatase TiO$_2$ and yellow anatase TiO$_2$ nanoparticles by doping with N and H [273]. The substitutional nonmetal doping of N and H, especially the $p$ states of N, contributed to the bandgap narrowing by mixing with O $2p$ states and formed the new VB. Theoretical work suggested that the adequately high concentration of Ti$^{3+}$ could induce a continuous oxygen vacancy of electronic states just below the CB edge of TiO$_2$, which strongly enhanced visible light absorption and the photocatalytic performance of the catalysts under visible light. The bandgap energy of the samples also decreased substantially, which was narrowed to about 2.0 eV.

5.1.2. C-nonmetal codoping

C-nonmetal codoping is also a kind of common approach to improve photocatalytic activity, such as (C, B) codoping [275, 276], (C, F) codoping [277], and (C, S) codoping [278].

Yu et al. investigated the geometry structures, formation energies, and electronic properties of the C, B and (C, B)-doped anatase TiO$_2$ using DFT calculations [275]. The results implied that (C, B)-codoped anatase TiO$_2$ could markedly influence the photocatalytic activity and light adsorption due to the change of the energy gaps ($E_g$) and Fermi levels ($E_F$). Moreover, the separation efficiency of carriers could be improved because of the existence of Ti$^{3+}$ ions. Lin et al. systematically investigated the electronic and optical properties of several possible (C, B) codoped models of anatase and rutile TiO$_2$ using DFT calculations [276]. The further calculation indicated that B 2$p$ and C 2$p$ decreased the bandgap by about 0.8 eV and the couples of the two hybridized states could also result in a downward red shift for spectrum response.

Deng et al. prepared F-modified C-doped TiO$_2$ composites via a simple sol-gel method using NaF as the F source followed by heat treatment at 700°C in N atmosphere [277]. The C atoms and F species interacted in the TiO$_2$ lattice and resulted in the narrowed bandgap (2.50 eV),
which could inhibited the recombination of photo-induced irradiation and resulted in a superior photocatalytic activity.

5.1.3. Other nonmetal-nonmetal codoping

Except for the above N-nonmetal codoping and C-nonmetal codoping, other nonmetal atoms, including (B, P) codoping [279], (B, F) codoping [280], and tridoping [281–284], are also investigated by many researchers. Yu et al. prepared a series of TiO$_2$ hollow sphere catalysts with or without nonmetal (B, P) dopants through a sol-gel process with styrene-methyl methacrylate copolymer microspheres as the template [279]. They found that both B and P dopants could also narrow the bandgap of the TiO$_2$ catalyst.

Recently, many researchers found that tridoping in TiO$_2$ crystals is an effectively supplemental tool. Ramanathan and Bansal synthesized (N, C, F) tridoped rutile TiO$_2$ nanorods that not only displayed enhanced photocatalytic performance in UV light but also allowed a significant level of visible light photocatalytic activity [281]. In visible light, pure TiO$_2$ only got 6% photodegradation of Congo red over 30 min, whereas (N, C, F) tridoped TiO$_2$ caused 56% dye degradation.

5.2. Metal-metal codoping

Although monometal doping can improve the bandgap structure, the serious recombination centers also deteriorate carrier transport due to its partially occupied impurity bands. Hence, many researchers want to improve carrier transport through codoping with two different metal elements, such as (Ti, Ni) codoping [285], (Ti, Fe) codoping [286], (Ag, W) codoping [287], (Ag, Zr) codoping [288], (Zn, Mn) codoping [289], (Cu, V) codoping [290], and (Fe, Ce) codoping [291].

Zhang et al. developed a novel Ni$^{2+}$ and Ti$^{3+}$ codoped porous anatase TiO$_2$ via a facile sol-gel technique combined with an in situ solid-state chemical reduction approach followed by mild calcinations (350°C) in Ar atmosphere [285]. The results showed that the doping of Ti$^{3+}$ and Ni$^{2+}$ species for TiO$_2$ could result in the tail states and significantly narrowed the bandgap of

![Figure 5](http://dx.doi.org/10.5772/63234)
anatase TiO$_2$ due to the formation of midgap states. Namely, Ti$^{3+}$ state below CB minimum and Ni$^{2+}$ state above VB maximum would be helpful in light adsorption and slowed the carrier recombination (Figure 5). In other words, (Ti, Ni) codoped porous black anatase TiO$_2$ resulted higher photocatalytic performance for methyl orange and rhodamine B.

Chen et al. found that Fe$^{3+}$/Ti$^{3+}$ codoping was also a good way to further improve Ti$^{3+}$ self-doping TiO$_2$ photocatalyst [286]. Fe$^{3+}$ doping could result in a blue shift for the VB edge by 1.0 eV. The synergistic effects between Ti$^{3+}$ and Fe$^{3+}$ dopants in the samples significantly narrowed the bandgap, leading to efficient photocatalytic performance in the visible light range.

Khan et al. synthesized (Ag, W) codoped TiO$_2$ with different W doping concentrations using the hydrothermal method [287]. The codoped TiO$_2$ displayed pure anatase phase with strong absorption in visible light region. Among all the codoped samples, the one with W doping concentration of 3.5 at.% possessed the best photocatalytic activity, which was attributed to the synergistic effect between the dopants and optimal doping concentration.

Benjwal and Kar investigated (Zn, Mn) codoped TiO$_2$ photocatalysts through varying dopant concentrations and found that the highest photocatalytic degradation of methyl blue was obtained with 1.0 at.% (Zn, Mn) codoped TiO$_2$ [289]. Further analysis showed that the incorporation of Zn$^{2+}$ and Mn$^{2+}$ ions into TiO$_2$ lattice caused a decrease of the energy bandgap due to the formation of impurity levels below VB, which resulted in an enhancement of light absorbance in the UV region. With further increasing codoping concentration, the density of new energy levels also increased, which ultimately enhanced the recombination possibilities of electron and holes because they could be easily trapped on these recombination centers.

Christoforidis and Fernández-García prepared two series of Cu$^{2+}$ and V$^{4+}$ codoped anatase TiO$_2$ samples using the microemulsion synthetic route by varying the metal/Ti ratio [290]. They found that isolated and highly dispersed Cu$^{2+}$ and V$^{4+}$ species favored the hole formation while at the same time decreased the amount of Ti$^{3+}$ centers under irradiation. Therefore, it could be deduced that (Cu, V) codoping contributed to the better photocatalytic performance due to the faster carrier separation.

Visible light-sensitive Fe$^{3+}$ and Ce$^{4+}$ codoped nano-TiO$_2$ photocatalyst was also investigated by Jaimy et al. [291]. They concluded that Fe doping and Ce doping could improve the light adsorption through trapping carriers and bending the VB and CB, respectively. In other words, the recombination of photoexcited electrons and holes was obviously prevented. Consequently, the codoped TiO$_2$ compositions exhibited higher photocatalytic activity than that of pure TiO$_2$ and commercial Degussa P25 under visible light.

5.3. Nonmetal-metal doping

5.3.1. N-metal doping

Recently, many researches about metal and N comodified TiO$_2$ indicate that metal assist N doping into the lattice TiO$_2$ and enhanced the shift of absorption edge to the visible light range, such as La, Ti, Zn, W, Cu, Fe, Ni, V, Cr, Ce, and Mn. Usually, N doping can form new states lying just above the VB, whereas the metal ion incorporated into the lattice TiO$_2$ or dispersed...
on the surface of TiO$_2$ can enhance the separation rate of photogenerated charges. Under visible light irradiation, the electron can be excited from the N impurity level to the metal ion impurity level or from the N impurity level to the CB or from the VB to the metal ion impurity level. This strategy has been extensively adopted to design visible-active photocatalysts with high efficiency.

Sun et al. [292] and Yu et al. [293] investigated the interaction between substitutional N and implanted La at the (101) and [001] facets of TiO$_2$ using first-principles DFT calculations and analyzed the origin of enhanced visible light photocatalytic activity of (N, La) codoped TiO$_2$. They found that substitutional probability of N atom obviously decreased due to La implantation. However, oxygen vacancies were greatly enhanced because of the synergistic effects of La doping and N doping. For the substitutional La/N codoped surface, the charge compensation between the substitutional La and substitutional N led to the formation of two isolated occupied N(s)-O $\pi^*$ impurity levels in the gap, forming the acceptor-donor-acceptor compensation pair and providing a reasonable mechanism for the enhanced visible light photocatalytic activity of (N, La) codoped TiO$_2$ anatase.

Li et al. developed a facile solvothermal strategy to simultaneously realize N doping and Ti$^{3+}$ self-doping in TiO$_2$ crystals [294]. N doping and concomitant Ti$^{3+}$ incorporation accounted for the reduction of the bandgap and realized photocatalytic activity in the visible light region. In other words, the introduced Ti$^{3+}$ ions could form a new sublevel state below the CB and thus generate more electrons and holes, which combined with the midgap state induced by N doping, synergistically reduced the bandgap to a lower level and finally improved the response to visible light. Consequently, (N, Ti) codoped hierarchical anatase TiO$_2$ catalysts manifested an excellent photocatalytic activity, deriving from its superior light harvesting ability, narrowing bandgap from the codoping strategy, enhancing adsorption capacity, and accelerating carrier transport rate.

A new type of (N, Zn) codoped TiO$_2$ photocatalyst was prepared by Wang et al. via a simple sol-gel technique, which exhibited a higher photocatalytic activity than pure TiO$_2$, N-doped TiO$_2$ and Sn-doped TiO$_2$ under both visible and UV light irradiation [295]. This implied that (N, Zn) codoping was a more efficient way to improve the photocatalytic activity than doping with just one type of ion. Hu et al. further confirmed the above results through nitridation and hydrogenation of a zinc titanium precursor [296].

(N, W) codoping is also an efficient way to improve the recombination of photogenerated carriers [297, 298]. Lai and Wu prepared (N, W) codoped TiO$_2$ nanobelt through a facile and low-temperature route followed by a subsequent calcination [297]. The achieved nanobelt film possessed a reduced bandgap of approximately 2.3 eV at approximately 2.6 at.% W and 3.1 at. % N. Moreover, the single N-doped, W-doped, and (N, W) codoped TiO$_2$ induced photodegradations of 88%, 73%, and 95% rhodamine B molecules, respectively. The higher activity of the (N, W) codoped TiO$_2$ further supported the positively synergetic effects arising from the codoping.

Recently, some researchers develop a novel (N, Cu) codoped TiO$_2$ photocatalysts and also find that the photocatalytic effect was good. For example, Wang et al. found that 7.7% of methyl
blue were adsorbed on the surface of (N, Cu) codoped TiO$_2$ nanosheet when the adsorption reached equilibrium in the dark, whereas 5.9% of methyl blue were adsorbed on the control sample [299]. The relatively low adsorption of methyl blue might be attributed to the low surface area of the N-doped TiO$_2$ nanosheet, and the slightly enhanced adsorption for (N, Cu) codoped TiO$_2$ nanosheet might be attributed to its modified surface by the Cu doping.

Zhang et al. prepared TiO$_2$ nanotubes codoped with Fe and N using a one-step hydrothermal method [300]. Compared to the commercial TiO$_2$ powders and pure TiO$_2$ nanotubes, (N, Fe) codoped TiO$_2$ nanotubes exhibited a stronger visible light absorption capability and an enhanced photocatalytic activity under visible light irradiation. The increase in photocatalytic activity could be ascribed to the narrowing of the bandgap and the promoted separation of the photogenerated electrons and holes. Fe$^{3+}$ ions were thought to occupy the active sites on the surface of (N, Fe) codoped TiO$_2$ as recombination centers for the electrons and holes.

Other N and metal codoping also includes (N, Ni) codoping [301], (N, Bi) codoping [302], (N, V) codoping [303], and (N, Mo) codoping [304]. Liu et al. reported the facile synthesis of reduced (N, Ni) codoped TiO$_2$ nanotubes and their photocatalytic activity application [301]. The narrowed bandgap of TiO$_2$ due to the doping of N and Ni elements could enhance the light absorption effectively. The electrochemical characterization revealed that photo-induced carriers were more efficient charge separation and transportation in reduced (N, Ni) codoped TiO$_2$ nanotubes photoanodes.

5.3.2. S-metal codoping

Recently, S-metal codoping has also been widely researched, such as (S, Fe) codoping [305], (S, Mo) codoping [306], and (S, Cu) codoping [307]. He et al. found that S and Fe could be easily implanted into the lattice of TiO$_2$ via a precipitation method [305]. Compared to undoped TiO$_2$ (S, Fe) codoped TiO$_2$ showed a higher photocatalytic activity under both UV and visible light irradiation, and the optimal methyl blue degradation level was 96.92%.

Zhang also developed for the first time a one-step hydrothermal process to synthesize (S, Mo) codoped TiO$_2$ mesoporous nanospheres [306]. The photodegradation ability towards rhodamine B was 3.8 times higher than that of the only S-doped samples.

5.3.3. Other nonmetal-metal doping

(F, Ca) codoped TiO$_2$ also caused several beneficial effects, including the enhancement of surface acidity, creation of oxygen vacancies, and increase of active sites as well as an impurity energy state (2.0 eV) below the CB of TiO$_2$, which could shorten the excitation path of electrons and reduce the apparent bandgap [308]. The smaller crystal size caused by doping with Ca could exhibit more powerful redox ability and the efficient separation of photogenerated electron-hole pairs. This implied that (F, Ca) codoping might be a kind of efficient way to improve the photocatalytic activity of TiO$_2$.

(B,Co) codoped TiO$_2$ was also investigated by Jaiswal et al. via DFT calculations [309]. They found that B occupied the interstitial site at low concentration (1 at.%), whereas it occupied
substitutional O position as the concentration increased (2 and 3 at.%). Both these B-doped TiO$_2$ showed improved photocatalytic activity due to the formation of shallow energy level, whereas higher visible light absorption was achieved owing to the presence of two deep energy levels in the bandgap according to DFT calculations.

Recently, a series of tridoped has also been developed [310, 311]. For example, Li et al. prepared (Cu, Ce, B) tridoped TiO$_2$ nanotubes via a hydrothermal method assisted by cetyl trimethyl ammonium bromide [310]. They found that Cu, Ce, and B could be intercalated into the interlayer spacing of the nanotubes besides the substitution for the Ti$^{4+}$ or O$_2^-$ . The synergetic effect of narrowing the bandgap of (Cu, Ce, B) tridoping exhibited the highest photocatalytic activity, which greatly inhibited the recombination of electrons and holes and enhanced the concentration of photogenerated carriers.

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