Enhanced photocatalytic activity of Se-doped TiO$_2$ under visible light irradiation

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Anatase TiO$_2$ is a typical photocatalyst, and its excellent performance is limited in ultraviolet light range due to its wide band gap of 3.2 eV. A series of Se-doped TiO$_2$ nanoparticles in anatase structure with various Se concentrations up to 17.1 at.% were prepared using sol-gel method. The doped Se ions are confirmed to be mainly in the valence state of $+4$, which provides extra electronic states in the band gap of TiO$_2$. The band gap is effectively narrowed with the smallest gap energy of 2.17 eV, and the photocatalytic activity is effectively improved due to the extended absorption range. The photocatalytic activity was evaluated by the degradation of Rhodamine B (RhB) in aqueous solution under visible light irradiation. The results show that Se doping significantly improves the photocatalytic activity of TiO$_2$ and 13.63 at.% Se-doped TiO$_2$ has the best performance.

In recent years, global environmental problems are becoming more and more concerned due to the severe pollutions, especially organic pollutants. A new solution for the degradation of pollutants is photocatalysis which is very effective and environmental friendly, compared with many other methods. Common photocatalysts usually include metal oxides and sulfides, such as, TiO$_2$, ZnO, SnO$_2$, ZrO$_2$, CdS, and so on. Among them, the first discovered photocatalyst, TiO$_2$, has been intensively studied$^{1,2}$. However, the band gap of TiO$_2$ is 3.2 eV for anatase phase and 3.0 eV for rutile phase, respectively. Thus, TiO$_2$ is nearly inactive under visible light irradiation so that sustainable solar energy can’t be used adequately.

Many efforts have been devoted to overcome the shortcomings and enhance the photocatalytic properties of TiO$_2$, especially in the visible light range. For example, ion doping$^{3–7}$, anatase-rutile phase coexisting$^8$, or p-n heterojunction formation$^9$, were explored. Among them, ion doping was an efficient approach$^{10}$. Many elements were doped into TiO$_2$, such as Ag$^{11}$, Fe$^{12,13}$, N$^{14–16}$, S$^{17}$, C$^{18–20}$, and B$^{21}$, etc. New energy levels can be introduced by the doping ions to decrease the band gap energy and enhance the photocatalytic activity of TiO$_2$, especially under visible light irradiation. Furthermore, electron-hole recombination rate can be effectively reduced due to the formation of charge trapping sites by foreign ions$^{22,23}$.

Selenium (Se) is in the same group element with oxygen (O) in the periodic table of elements. Se possesses cationic (Se$^{4+}$, Se$^{6+}$) and anionic (Se$^{2−}$) characters, which was suggested to introduce extra electronic states in the band gap by calculations$^{24}$. However, TiO$_2$ doped with Se has not been widely studied. Zhang et al. prepared the SeO$_2$/TiO$_2$ nanocomposites, in which the recombination of electron-hole pairs might be effectively suppressed due to the formation of heterostructure and absorption might be extended to visible light due to the narrow band gap of 2.0 eV of Se$^0$ by the reduction of Se$^{4+}$$^{25}$. Se was doped into anatase TiO$_2$ by various methods, with narrowed band gap energy and effectively extended absorption to the visible light range. Therefore, significantly enhanced photocatalytic activity was achieved under visible irradiation$^{24,26–28}$. However, the Se doping concentration was rather low, due to the easy sublimation (315–317 °C) of SeO$_2$ in the calcining processes$^{28}$. To further narrow the band gap and improve the photocatalytic activity under visible light irradiation, higher concentration of Se should be doped into TiO$_2$. In this paper, we significantly increase the doping concentration of Se up to 17.1 at.% in TiO$_2$ by lowering the calcining temperature. The band gap is effectively narrowed and the photocatalytic activity under visible light irradiation is improved.

Experimental
Sample preparation. The chemicals used in this study were absolute ethanol (C$_2$H$_5$OH), tetrabutyl titanate (C$_{16}$H$_{36}$O$_4$Ti), nitric acid (HNO$_3$), selenium dioxide (SeO$_2$), which were of analytical grade and used as received.
without further purification. The distilled water was produced using a Direct-Q Millipore filtration system with resistivity of 18.2 MΩ·cm (Millipore Limited, Watford, UK).

A series of Se-doped TiO$_2$ nanoparticles with various Se concentration were prepared by sol-gel method. The Se-doped TiO$_2$ powders were synthesized as following steps: solution A was the mixture of C$_{16}$H$_{36}$O$_4$Ti and C$_2$H$_5$OH which was stirred to transparent solution in dropping process of nitric acid, solution B was ethanol solution of different concentration of SeO$_2$. The amount of SeO$_2$ used was based on the designed atomic concentration of Se in TiO$_2$ (Ti$_{1-x}$Se$_x$O$_2$), which were 0 at.% (pure TiO$_2$), 5 at.% , 10 at.% , 15 at.% , 20 at.% and 25 at.% respectively. Solution B was dropped into solution A by violent stirring. The uniform sol solution was kept at room temperature for 16 h till it became a gel solution. The gel solution was dried at 120°C until the xerogel was generated, then the xerogel was calcined at 300 °C for 3 h to get the final powders. The obtained products are denoted as TiO$_2$, TSe5, TSe10, TSe15, TSe20, TSe25, respectively, and the number denotes the designed atomic percentage of Se in Ti$_{1-x}$Se$_x$O$_2$.

**Table 1.** Lattice constant $a$ and $c$, Crystallite size, Band gap and EDX determined Se concentration of Se-doped TiO$_2$.

| Catalyst powder | $a$ (nm) | $c$ (nm) | Crystallite size (nm) | Band gap (eV) | EDX Se (at.%) |
|-----------------|----------|----------|----------------------|---------------|---------------|
| TiO$_2$         | 3.801    | 9.542    | 7.4                  | 2.82          | 0             |
| TSe5            | 3.793    | 9.505    | 10.9                 | 2.39          | 2.2           |
| TSe10           | 3.800    | 9.617    | 7.7                  | 2.19          | 6.76          |
| TSe15           | 3.789    | 9.567    | 6.8                  | 2.73          | 11.77         |
| TSe20           | 3.789    | 9.567    | 6.4                  | 2.58          | 13.63         |
| TSe25           | 3.789    | 9.592    | 6.3                  | 3.03          | 17.1          |
| Degussa P25     |          |          | 0.89                 | 0             |               |

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**Structural characterization.** The crystal structure of Se-doped TiO$_2$ nanoparticles was analyzed by X-ray diffraction (XRD, Rigaku SmartLab3) using Cu Kα radiation ($\lambda = 0.15418$ nm). The UV-Vis diffuse reflectance spectra (DRS) were recorded using a UV-visible spectrophotometer (JASCO, UV-670) with a wavelength range of 200–800 nm. The photoluminescence (PL) and PL excitation (PLE) spectra were recorded on a PL spectrofluorometer (HJY-FL3–211–TCSPC) at an excitation wavelength of 365 nm. The morphologies were studied by a scanning electron microscope (SEM, FEI Inspect F50), equipped with an energy dispersive X-ray spectroscope (EDX). EDX was performed for the chemical analysis of the doped samples, and the measured concentrations are listed in Table 1. As can be seen, the measured concentrations of Se are smaller than the designed values, which might be due to the partial sublimation of SeO$_2$ during the calcining processes$^{29}$. Raman studies were carried out on a Raman spectrometer (Horiba Jobin Yvon Lab RAM HR 800) under the backscattering geometric configuration at room temperature. The valence states of each element were studied by X-ray photoelectron spectroscopy (XPS, Thermo Fisher Scientific) with Al Kα X ray source (hv = 1486.6 eV). Binding energies were referenced to the C1s peak at 284.5 eV.

**Photocatalytic experiments.** The visible-light-induced photocatalytic activity was evaluated in aqueous solution, by using Rhodamine B (Rhb, Tianjin China Chemical Reagent Ltd.) as a model contaminant in this study. The light irradiation system contained a 500 W Xe lamp (Beijing Trusttech Co Ltd, CHF-XM), equipped with visible light pass filter (400–800 nm). In photocatalytic experiment: 0.4 g photocatalyst was mixed with a 150 mL Rhb solution (7.5 mg L$^{-1}$), an adsorptive experiment primarily was proceeded in the dark for 30 min to achieve absorption-desorption equilibrium of Rhb onto the catalyst surface before light irradiation. After interval of scheduled time, 2–3 mL Rhb solution was collected for concentration analysis, which was monitored by using a UV-Vis spectrophotometer (Hitachi U-3900).

**Results and Discussion**

Figure 1 shows the XRD patterns of TiO$_2$ and Se-doped TiO$_2$ powders. As can be seen, the main diffraction peaks of TiO$_2$, which is due to the smaller radius of Se$^{4+}$ (0.64 Å) than Ti$^{4+}$ (0.745 Å) (The valence state of Se is determined to be +4 by XPS, which will be discussed later$^{30}$). With further increasing Se doping concentration, the lattice constant $a$ is still smaller than that of TiO$_2$, but lattice constant $c$ becomes larger than that of TiO$_2$. The effect of Se doping on the crystallite size can be evaluated according to the Scherrer’s equation, $D = K\lambda/\beta \cos \theta$, where $K$ is the constant depending how the half height width of selected diffraction peak is determined (here we use 0.89), $\lambda$ is the wavelength of X ray (0.15418 nm), $\beta$ is the half height width, and $\theta$ is the Bragg angle. Here we
select (101) peak to estimate the crystallite size, which is the strongest diffraction peak for all the samples. As can be seen, the peak becomes narrower first with small doping concentration of TSe5, and then becomes broader with further increasing the doping concentration. The calculated crystallite size is listed in Table 1. Similar phenomenon was observed by Khan et al., though the doping concentration was much smaller in their work. Due to the small concentration of Se in TSe5, the less formation of TiO$_6$ octahedra might facilitate the polycondensation process of TiO$_6$ octahedra in corner sharing arrangement, leading to the larger crystallite size of anatase phase. However, with further increasing the doping concentration, the doped Se ions might not occupy the lattice site, and thus the growth of the crystallite size was prevented. The similar rule but much higher Se concentration in our work compared with Khan et al.’s work might be due to the much lower calcining temperature in our work, leading to the much smaller crystallite size and much higher tolerable doping concentration. The morphologies of the samples are further studied by SEM, and the images are shown in Fig. S1 (Supplementary materials). The particle sizes of all samples are all smaller than 10 nm, which is consistent with XRD results.

Raman spectroscopy is a powerful technique for the investigation of crystalline and defect structure, and to elucidate the various phases of TiO$_2$. The Raman spectra of Se-doped TiO$_2$, with various Se concentrations are shown in Fig. 2. The peak positions are observed at 147 cm$^{-1}$ (1-$E_g$), 398 cm$^{-1}$ (A$_{1g}$), 514 cm$^{-1}$ (B$_{1g}$), 639 cm$^{-1}$ (3-$E_g$) for TiO$_2$, corresponding to the vibration modes of anatase phase. All the Se-doped TiO$_2$ exhibit approximately the same bands, which is consistent with the pure anatase phase observed by XRD. TSe5 shows an extra peak at 195 cm$^{-1}$, which corresponds to the 2-$E_g$ band. This confirms the better crystalline structure of TSe5. TSe5 shows a distinct behavior, with much narrower peak and significant shift to smaller wavenumbers. This can be understood by the better crystalline structure with much larger crystallite size for TSe5.

Figure 1. XRD patterns of Se-doped TiO$_2$. “o” labels the diffraction peak from orthorhombic phase.

Figure 2. Raman spectra of Se-doped TiO$_2$, inset shows the enlarged view of 1-$E_g$ peak.
XPS was used to study the chemical states and electronic structure of Se-doped TiO$_2$. Figure 3 shows the XPS spectra of Ti, O and Se for TSe20. The Ti2p$_{3/2}$ and Ti2p$_{1/2}$ peaks can be deconvoluted into two sets of peaks, the stronger peaks at 458.7 eV and 464.3 eV correspond to Ti$^{4+}$, confirming the main valence state of +4 for Ti in Se-doped TiO$_2$. A set of weaker peaks at 456.4 eV and 461.8 eV can be attributed to the Ti$^{3+}$. This might be due to the low calcining temperature, oxygen is not active enough to fully oxidize Ti, resulting in the formation of O vacancies and Ti$^{3+}$ sites in TiO$_2$ lattice. To confirm the formation of Ti$^{3+}$, we further annealed the TiO$_2$ powders at 500 °C for 1 hour, and the XPS spectrum is shown in Fig. 3(b). As can be seen, only Ti2p$_{3/2}$ peak at 458.5 eV and Ti2p$_{1/2}$ peak at 464.2 eV can be observed, indicating the valence state of +4. In Fig. 3(c), O1s peak can be deconvoluted into two peaks, one at 529.8 eV, and the other at 528.2 eV. The peak at 529.8 eV is attributed to the Ti-O bond. The observation of peak at 528.2 eV is quite abnormal, since the bonding energy of Se-O in SeO$_2$ should be higher. It is known that the O2p peak in La$_2$O$_3$ is 528.8 eV, which is due to the higher ionic nature of the La-O bonding. Thus O ions will have lower binding energy when they attract more electrons. In the Se-doped TiO$_2$ with much higher Se concentration, Se may not occupy the lattice sites due to the large lattice distortion. O may attract electrons not only from the neighboring Ti, but also from Se ions due to the formation of Ti-O-Se structure, leading to lower binding energy, in comparison with O ions locating at regular lattice site. Figure 3(d) shows the Se3d XPS peak, which can be deconvoluted into two peaks: one at 58.5 eV, and the slightly smaller value might be due to the doping in TiO$_2$ lattice environment. The peak at 55 eV indicates the existence of small amount of Se$^{0}$, which might be due to the reduction from C and H in the sources and further confirms the less chemical activity of O at such low calcining temperature.

To investigate the optical absorption properties of Se-doped TiO$_2$ nanoparticles, the UV-vis DRS were recorded and the results are shown in Fig. 4. The absorption edges of TiO$_2$ shift to longer wavelength in the visible region, which is consistent with its brown-colored appearance, as shown in the inset of Fig. 4(a). This can be understood by the existence of Ti$^{3+}$ due to the large concentration of O vacancies and disorder in TiO$_2$ with very low calcining temperature. With Se doping, the TiO$_2$ powders first exhibits darker color, with the darkest color for TSe5. Then the color becomes pale with further increasing the Se doping concentration. TSe5 shows the highest absorption coefficient in the visible range, which decreases with further increasing Se doping concentration. TSe25 shows almost the same absorption coefficient as TiO$_2$. The optical band gap of Se-doped TiO$_2$ can be evaluated by its absorption spectrum. The optical absorption near the band edge of a semiconductor often obeys the Kubelka-Munk equation: $(\alpha h \nu)^n = A(h \nu - E_g)$, where $A$ is a constant, $h \nu$ is corresponding to photon energy, $E_g$ is the band gap of the semiconductor, $\alpha$ is the absorption coefficient, and $n$ is 0.5 for indirect band gap materials, such as TiO$_2$. The approximate values of the band gap can be obtained from the intercept of the
tangent to X-axis, as shown in Fig. 4(b). The calculated band gap values are listed in Table 1. As can be seen, TSe10 shows the smallest $E_g$ of 2.19 eV. There are two competing effects which may influence the final $E_g$ of Se-doped TiO$_2$. The Se doping may introduce additional electronic energy levels inside the band gap, which will effectively narrow the band gap$^{24}$. The suppressed PLE peaks from TiO$_2$ host lattice, shown in Fig. S2 (Supplementary materials), give an indirect evidence for the introduction of extra energy levels from doped Se ions. However, with further increasing Se concentration higher than that of TSe5, the crystallite size becomes smaller. The band gap increases with a decrease in crystallite size due to the quantum size effect$^{27}$.

PL spectra have been widely used to investigate the change of surface states of TiO$_2$, the efficiency of charge carrier trapping, immigration and transfer to understand the fate of electron-hole pairs in semiconductor particles$^{39}$. Figure 5 shows the PL spectra of Se-doped TiO$_2$. A strong peak at 394 nm can be found for Se-doped TiO$_2$, but lacking in TiO$_2$, indicating the significantly suppressed electron-hole recombination in TiO$_2$, which might be due to the high defects concentration$^{40}$. Slight Se doping improves the crystalline structure, which may suppress the defects concentration and increase the electron-hole recombination rate. Clear PL peaks can be observed in Se-doped TiO$_2$. However, with increasing Se doping concentration, more defects are formed, which provides more electron trapping centers. Interestingly, no PL peaks can be observed for TSe20, suggesting the strongly suppressed electron-hole recombination rate$^{40}$. Further increasing Se doping concentration to TSe25, strong PL peak can be observed again. This indicates that there must be an optimum level up to which the addition of doping ions can help to lower electron-hole recombination rate. Otherwise, the excess doping ions act as recombination centers, which can dramatically decrease the photocatalytic activity of the doped catalyst$^{27}$.

The typical time-dependent UV-Vis spectra of RhB solution by TiO$_2$ and TSe20 in photochemical reaction are shown in Fig. 6. The intensity of the characteristic absorption peak of RhB solution decreases with time. It can be seen that TiO$_2$ can effectively decompose RhB under the visible light irradiation. TSe20 shows much faster decomposition of RhB under visible irradiation, confirming the promotion of Sedoping on the photocatalytic activity of TiO$_2$. The absorption peak position shifts to shorter wavelength, revealing that RhB is de-ethylated in a stepwise manner (i.e., ethyl groups are removed one by one as confirmed by the gradual peak wavelength shifts toward the blue region)$^{42}$. However, after 45 minute, for RhB under irradiation in the presence of TSe20, the shift of wavelength reaches maximum. This indicates the decomposition mechanism of RhB changes to destroy its conjugated structure$^{45}$.

Figure 4. (a) UV-vis DRS of Se-doped TiO$_2$. Insets show the corresponding photos of Se-doped TiO$_2$ powders, (b) Curve-fitting by using the Kubelka-Munk function method for the absorption curves.
Photocatalytic reactions for the degradation of RhB aqueous solutions using Se-doped TiO$_2$ are shown in Fig. 7. For comparison, the photocatalytic activity of Degussa P25 was also measured. For the first 30 min in dark, it can be seen that Se-doped TiO$_2$ shows higher adsorption ability than TiO$_2$. Since the crystallite size of all the samples is comparable, the surface area should not be the main reason. It is supposed to be due to more active sites on the surface of Se-doped TiO$_2$. TiO$_2$ shows superior photocatalytic performance than Degussa P25. With Se doping, the photocatalytic capability is significantly improved. After irradiation for 30 min, nearly 91.3% of RhB was degraded by the sample TSe20, while only 60.3% Rhb was degraded by TiO$_2$. Experimental dependencies of the molar concentration of RhB in the presence of undoped/doped TiO$_2$ and Degussa P25 powders in visible irradiation exhibit pseudo first order kinetics, as shown in Fig. 7(b) by plotting ln(C/Co) versus irradiation time.
The apparent reaction rate constant ($k_{\text{app}}$) is calculated from the slope of the curve, as illustrated in Fig. 7(c). With small concentration of Se doping, TSe5 shows much smaller $k_{\text{app}}$ value. With further increasing Se doping concentration, $k_{\text{app}}$ continuously increases. TSe20 exhibits the highest $k_{\text{app}}$. With further increasing Se doping concentration, $k_{\text{app}}$ of TSe25 drastically decreases. Comparing the rate constant values under the visible light irradiations, the powder TSe20 attains the highest $k_{\text{app}}$ value of 0.088 min$^{-1}$, which is much higher than that of undoped TiO$_2$ (0.030 min$^{-1}$) and Degussa P25 (0.0004 min$^{-1}$). TiO$_2$ exhibits better photocatalytic activity than Degussa P25, mainly because of the synergetic effect of narrow band gap in the visible light range and mixed phases (orthorhombic and anatase) which hinder the recombination of generated electron-hole pairs$^{44}$.

As shown in Fig. 8, the photoinduced electrons and holes play important role in the photocatalytic reaction. The electrons are adsorbed by O$_2$ to produce superoxide radical anions of $\cdot$O$_2^{-}$, while holes can oxidize OH$^{-}$ to produce $\cdot$OH. Both can further react with RhB which is decomposed to CO$_2$ and H$_2$O. The improved photocatalytic activity of Se-doped TiO$_2$ under visible light can be first understood by the decreasing band gap into the visible range with increasing Se doping concentration, which might increase the absorption efficiency of the visible light, as shown in Fig. 8. It has been calculated that Se$^{4+}$ doping into TiO$_2$ may induce additional mid-gap electronic levels close to the conduction band edge$^{24}$. These intermediate electronic states are determined to be mainly originated from the
Se 3p states and not populated by electrons. They are not donor states but allowed energy states hybridized with the Ti 3d states in the conduction band. The increase in the concentration of dopant Se⁴⁺ introduces more electronic states into the band-gap, enhancing the density of electronic states in the gap. These intermediate energy levels offer additional steps for the absorption of low energy photons through the excitation of valence band electrons to these intermediate energy levels, from where they can be excited again to the conduction band. Furthermore, the photocatalytic capability also depends on the separation efficiency of generated electron-hole pairs. To improve the photocatalytic activity, the photoinduced electrons and holes inside the particles should survive during the transportation to the surface of particles, and the recombination should be avoided. As can be seen, no obvious peaks can be observed in the PL spectrum of TSe20, indicating the strongly suppressed recombination rate of electron-hole pairs. The defects, such as O vacancies, Ti⁴⁺, etc. may play significant role, since they can trap the electrons and suppress the recombination of electron-hole pairs. With slight concentration of Se doping, the crystal structure is improved, thus defects concentration is decreased, leading to the worse photocatalytic capability for TSe5. With further increasing Se doping concentration, more defects will be introduced in Se-doped TiO₂, the photo-generated electrons can be effectively trapped, and the recombination of electron-hole pairs is effectively suppressed, leading to the improved photocatalytic performance. As can be seen from PL measurements, TSe20 exhibits the negligible PL intensity, indicating the lowest recombination rate of electron-hole pairs, as can be seen the significant PL intensity for TSe25. The photocatalytic performance of TSe25 decreases drastically. Thus, it can be concluded that the improved photocatalytic activity in Se-doped TiO₂ with optimum concentration (TSe20) is a synergetic contribution from the narrow band gap by Se⁴⁺ doping and suppressed electron-hole pairs recombination due to the optimum defects concentration.

Conclusions
A series of Se-doped TiO₂ nanoparticles of anatase structure with various Se concentrations up to 17.1 at.% were prepared using sol-gel method. Slight concentration of Se doping has been confirmed to improve the crystalline structure of TiO₂, while higher concentration of Se doping deteriorates the crystalline structure. The doped Se ions are confirmed to be mainly in the valence state of +4, which provides extra electronic states in the band gap of TiO₂. The band gap is effectively narrowed with the smallest gap energy of 2.17 eV by Se doping of concentration of 6.76 at.%. With further increasing Se doping concentration, Se doping significantly improves the photocatalytic activity of TiO₂ and 13.63 at.% Se-doped TiO₂ has the highest photocatalytic activity from the photo degradation of RhB in aqueous solution under visible light irradiation, which is attributed to the synergetic contribution of narrowed band gap by Se doping and suppressed electron-hole recombination due to the optimum defects concentration.

References
1. Mills, A. & Hunte, S. L. An overview of semiconductor photocatalysis. *Journal of Photochemistry and Photobiology A* **108**, 1–35 (1997).
2. Fu, W. *et al.* F, Ca co-doped TiO₂ nanocrystals with enhanced photocatalytic activity. *Dalton Transactions* **43**, 16160–16163 (2014).
3. Behnjady, M. A. & Eskandarloo, H. Silver and copper co-impregnated onto TiO₂-P25 nanoparticles and its photocatalytic activity. *Chemical Engineering Journal* **226**, 1207–1213 (2013).
4. Wang, Q. Q., Xu, S. H. & Shen, F. L. Preparation and characterization of TiO2 photocatalysts co-doped with iron (III) and lanthanum for the degradation of organic pollutants. *Applied Surface Science* 257, 7671–7677 (2011).
5. Iliev, V., Tomova, D. & Rakovsky, S. Nanosized N-doped TiO2 and gold modified semiconductors – photocatalysts for combined UV-visible light destruction of oxalic acid in aqueous solution. *Desalination* 260, 101–106 (2010).
6. Pan, L. et al. Ti3+-defected and V-doped TiO2 quantum dots loaded on MCM-41. *Chemical Communications* 50, 988–990 (2014).
7. Monteiro, R. A. R., Lopes, F. V. S., Bouaventura, R. A. R. & Silva, A. M. T. V. J. P. Synthesis and characterization of N-modified titania nanoparticles for photocatalytic applications. *Environmental Science and Pollution Research* 22, 810–819 (2015).
8. Zhang, J., Xu, Q., Feng, Z. C., Li, M. J. & Li, C. Importance of the relationship between Surface Phases and Photocatalytic Activity of TiO2. *Angewandte Chemie International Edition* 47, 1766–1769 (2008).
9. Chen, Y. S., Crittenenden, J. C., Hackney, S., Sutter, L. & Hand, D. W. Preparation of a Novel TiO2-Based p–n Junction Nanotube Photocatalyst. *Environmental Science & Technology* 39, 1201–1208 (2005).
10. Herrmann, J. M. Heterogeneous photocatalysis: fundamentals and applications to the removal of various types of aqueous pollutants. *Catalysis Today* 53, 115–129 (1999).
11. Arabatzis, I. M. et al. Silver-modified titanium dioxide thin films for efficient photodegradation of methyl orange. *Applied Catalysis B: Environmental* 42, 187–201 (2003).
12. Wang, C. Y., Bottcher, C., Baehnemann, D. W. & Dohrmann, J. K. A comparative study of nanometer sized Fe(III)-doped TiO2 photocatalysts: synthesis, characterization and activity. *The Journal of Materials Chemistry* 13, 2322–2329 (2003).
13. Litter, M. I. & Navfo, J. A. Photocatalytic properties of iron-doped titania semiconductors. *Journal of Photochemistry and Photobiology A* 98, 171–181 (1996).
14. Asahi, R., Morikawa, T., Ohwaki, T., Aoki, K. & Taga, Y. Visible-light photocatalysis in nitrogen-doped titanium oxides. *Science* 293, 269–271 (2001).
15. Borda, G. et al. Enhanced nitrogen doping in TiO2 nanoparticles. *Nano Letts* 3, 1049–1051 (2003).
16. Cong, Y., Zhang, J. L., Chen, F., Anpo, M. & He, D. Preparation, photocatalytic activity, and mechanism of nano-TiO2 co-doped with nitrogen and iron (III). *The Journal of Physical Chemistry C* 111, 10618–10623 (2007).
17. Luo, H. et al. Photocatalytic activity enhancing for titanium dioxide by co-doping with bromine and chlorine. *Chemistry of Materials* 16, 846–849 (2004).
18. Wu, Y. M., Xing, M. Y., Zhang, J. L. & Chen, F. E. Effective visible light-active boron and carbon modified TiO2 photocatalyst for degradation of organic pollutant. *Applied Catalysis B: Environmental* 97, 182–189 (2010).
19. Xu, C., Killmeyer, R., Gray, M. L. & Khan, S. Photocatalytic effect of carbon-modified n-TiO2 nanoparticles under visible light illumination. *Applied Catalysis B: Environmental* 64, 312–317 (2006).
20. Li, Y., Hwang, D. S., Lee, N. H. & Kim, S. J. Synthesis and characterization of carbondoped titania as an artificial solar light sensitive photocatalyst. *Chemical Physics Letters* 404, 25–29 (2005).
21. Lu, N. et al. Fabrication of boron-doped TiO2 nanotube array electrode and investigation of its photoelectrochemical capability. *The Journal of Physical Chemistry C* 111, 11836–11842 (2007).
22. Choi, W., Termin, A. & Hoffmann, M. R. The Role of Metal Ion Dopants in Quantum-Sized TiO2: Correlation between Photoreactivity and Charge Carrier Recombination Dynamics. *The Journal of Physical Chemistry* 98, 13669–13679 (1994).
23. Akpan, U. G. & Hameed, B. H. The advancements in sem−gel method of doped−TiO2 photocatalysts. *Applied Catalysis A: General* 375, 1–11 (2010).
24. Gurkan, Y. Y., Kasapbasi, E. & Cinar, Z. Enhanced solar photocatalytic activity of TiO2 by selenium(VI) ion-doping: Characterization and DFT modeling of the surface. *Chemical Engineering Journal* 214, 34–44 (2013).
25. Zhang, S. Y., Chen, X. J., Tian, Y. P., Jin, B. K. & Yang, J. X. Preparation and characterization of novel SeO2/TiO2 nanocomposite. *Journal of Crystal Growth* 304, 42–46 (2007).
26. Rockafellow, E. M., Haywood, J. M., Witte, T., Houk, R. S. & Jenks, W. S. Selenium-Modified TiO2 and Its Impact on Photocatalysis. *Langmuir* 26, 19052–19059 (2010).
27. Khan, H. & Berk, D. Selenium Modified Oxalate Chelated Titania: Characterization, Mechanistic and Photocatalytic Studies. *Applied Catalysis A: General* 305, 285–301 (2011).
28. Stengl, V., Bakardjeva, S. & Bludска, J. Se and Te-modified titania for photocatalytic applications. *J. Mater. Sci.* 46, 3523–3536 (2011).
29. Shannon, R. D. Revised effective ionic radii and systematic studies of interatomic distances in halides and chalcogenides. *Acta Crystallographica Section A* 32, 751–767 (1976).
30. Khade, G. V., Suwarnkar, M. B., Gavade, N. L. & Garadkar, K. M. Sol−gel microwave assisted synthesis of Sm-doped TiO2 nanoparticles and their photocatalytic activity for the degradation of Methyl Orange under sunlight. *Journal of Materials Science: Materials in Electronics* 27, 6425–6432 (2016).
31. Santara, B., Pal, B. & Giri, P. K. Signature of strong ferromagnetism and optical properties of Co doped TiO2 nanoparticles. *Journal of Applied Physics* 110, 114322–114327 (2011).
32. Xia, Y. et al. Effect of calcined atmosphere on the photocatalytic activity of P-doped TiO2. *Applied Surface Science* 289, 306–315 (2014).
33. Rudyk, B. W., Blanchard, P. E. R., Cavell, R. G. & Mar, A. Electronic structure of lanthanum copper oxycarbonogenides LaCuOCh (Ch=S, Se, Te) by X−ray photoelectron and absorption spectroscopy. *Journal of Solid State Chemistry* 184, 1649−1654 (2011).
34. Hashimoto, M., Hayashi, K. & Kitaoka, S. Enhanced apatite formation on Ti metal heated in PO2-controlled nitrogen atmosphere. *Materials Science and Engineering: C* 33, 4153−4159 (2013).
35. Chen, X., Liu, Y. L., Yu, P. Y. & Mao, S. S. Increasing Solar Absorption for Photocatalysis with Black Hydrogenated Titanium Dioxide Nanocrystals. *Science* 331, 746−750 (2011).
36. Liu, L., Yu, P. Y., Chen, X., Mao, S. S. & Shen, D. Z. Hydrogenation and Disorder in Engineered Black TiO2. *Physical Review Letters* 111, 065505 (2013).
37. Yen, C. C., Wang, D. Y., Shih, M. H., Chang, L. S. & Shih, H. C. A combined experimental and theoretical analysis of Fe-implanted TiO2, modified by metal plasma ion implantation. *Applied Surface Science* 256, 6865–6870 (2010).
38. Mo, L. et al. Band gap engineering of TiO2 through hydrogenation. *Applied Physics Letters* 105, 202114 (2014).
39. Yamashita, H. et al. Photocatalytic decomposition of NO at 275 K on titanium oxide catalysts anchored within zeolite cavities and framework. *Applied Surface Science* 121, 305–309 (1997).
40. Yu, J., Ma, T. & Liu, S. Enhanced photocatalytic activity of mesoporous TiO2 aggregates by embedding carbon nanotubes as electron-transfer channels. *Journal of Physical Chemistry C* 113, 3491−3501 (2009).
41. Wu, T., Liu, G., Zhao, J., Hidaka, H. & Serpone, N. The Journal of Physical Chemistry B 102, 5845−5851 (1998).
42. Zhu, S. et al. Effect of Au supported TiO2 with dominant exposed [001] facets on the visible-light photocatalytic activity. *Applied Catalysis B: Environmental* 119–120, 146−155 (2012).
43. Luo, L. et al. Preparation of multiferroic Co substituted BiFeO3 with enhanced coercive force and its application in sorption removal of dye molecules from aqueous solution. *Journal of Alloys and Compounds* 558, 73−76 (2013).
44. Huang, S. et al. 3D hierarchical FeSe, microspheres: Controlled synthesis and applications in dye-sensitized solar cells. *Nano Energy* 15, 205−215 (2015).
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Author Contributions
Q.X. conceived the experiment idea and designed the research. W.X. and R.L. carried out the experiment and did most measurement. Q.X. and W.X. wrote the paper.

Additional Information
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