

Research Article

Multiwalled Carbon Nanotube-TiO₂ Nanocomposite for Visible-Light-Induced Photocatalytic Hydrogen Evolution

Ke Dai,¹ Xiaohu Zhang,² Ke Fan,² Peng Zeng,² and Tianyou Peng²

¹ College of Resources and Environment, Huazhong Agricultural University, Wuhan 430070, China

² College of Chemistry and Molecular Science, Wuhan University, Wuhan 430072, China

Correspondence should be addressed to Ke Dai; dk@mail.hzau.edu.cn and Tianyou Peng; typeng@whu.edu.cn

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Multiwalled carbon nanotube- (MWCNT-) TiO₂ nanocomposite was synthesized via hydrothermal process and characterized by X-ray diffraction, UV-vis diffuse reflectance spectroscopy, field emission scanning electron microscope, thermogravimetry analysis, and N₂ adsorption-desorption isotherms. Appropriate pretreatment on MWCNTs could generate oxygen-containing groups, which is beneficial for forming intimate contact between MWCNTs and TiO₂ and leads to a higher thermal stability of MWCNT-TiO₂ nanocomposite. Modification with MWCNTs can extend the visible-light absorption of TiO₂. 5 wt% MWCNT-TiO₂ derived from hydrothermal treatment at 140°C exhibiting the highest hydrogen generation rate of 15.1 μmol·h⁻¹ under visible-light irradiation and a wide photoresponse range from 350 to 475 nm with moderate quantum efficiency (4.4% at 420 nm and 3.7% at 475 nm). The above experimental results indicate that the MWCNT-TiO₂ nanocomposite is a promising photocatalyst with good stability and visible-light-induced photoactivity.

1. Introduction

Hydrogen resource obtained directly from water splitting using photocatalyst and solar light is under scrutiny as a clean energy resource; however it has been facing technical challenge to find a stable and efficient photocatalyst that can maximally utilize solar light [1]. In addition to oxide semiconductors, a great variety of novel photoactive semiconductors have been developed in the last few years. Among these, mixed oxides of transition metal like Nb, V, or Ta or with main group elements such as Ga, In, Sb, or Bi have been extensively investigated as attractive candidates for visible-light-induced photocatalysis [2]. Also, sulfides and nitrides of different metals have been frequently selected to obtain materials with visible-light-driven photoactivity [3]. Titania (TiO₂) is a widely used photocatalyst due to its high chemical stability, low cost, and nontoxic nature [4], but it can only absorb UV light with low quantum efficiency due to its wide bandgap (*ca.* 3.2 eV), which limits its application in visible-light-driven photocatalysis. The above problems have been partly resolved by many methods, such as surface modification via organic materials [5], semiconductor coupling [6],

bandgap modification by nonmetals [7, 8], and plasmonic metal (Ag and Au) doping [9–14], creating oxygen vacancies and disorder or dye sensitization [15–17].

Due to the special structures and extraordinary mechanical and unique electronic properties, carbon nanotubes (CNTs) have the potential to extend the photoresponse range of TiO₂ to visible-light region by modification of bandgap and/or sensitization and increase the photoactivity of TiO₂ by contribution to high surface area and inhibition of electron-hole recombination [18]. Single-walled carbon nanotubes (SWCNTs) have shown a synergy effect on enhancing photoactivity for H₂ evolution over a mixture of SWCNTs and TiO₂ [19]. Ou et al. [20] have also demonstrated that multiwalled carbon nanotubes (MWCNTs) could enhance the visible-light-driven photoactivity of TiO₂ by acting as a photosensitizer in the MWCNT-TiO₂:Ni composite. MWCNT-TiO₂ nanocomposite with visible-light-driven photoactivity was successfully synthesized via direct growth of TiO₂ nanoparticles on the surface of the functionalized MWCNTs by the hydrothermal treatment in our group [21]. However, the effects of MWCNT pretreatment, MWCNT content, and

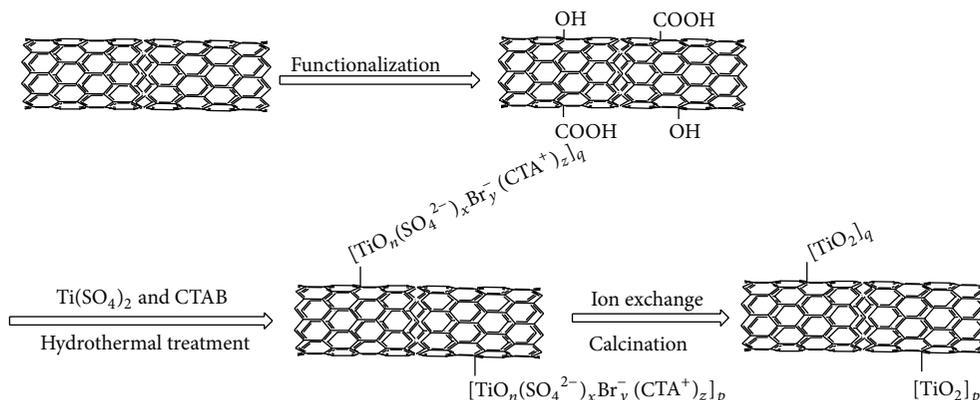


FIGURE 1: Schematic formation model for MWCNT-TiO₂ nanocomposite.

synthetic conditions of MWCNT-TiO₂ nanocomposite on its photocatalytic hydrogen evolution efficiency and quantum efficiency under monochromatic light irradiation are still beyond our knowledge.

Herein, MWCNT-TiO₂ nanocomposite was synthesized, characterized, and employed for photocatalytic H₂ production from triethanolamine (TEOA) solution. The effects of pretreatment methods, content of MWCNTs, and hydrothermal temperature on the photocatalytic H₂ evolution efficiency over the Pt-loaded photocatalysts were studied. Moreover, the apparent quantum efficiency (AQE) of 5 wt% MWCNT-TiO₂ upon incident monochromatic light is also investigated.

2. Experiment

2.1. Pretreatment Methods for MWCNTs. MWCNTs (diameter < 8 nm; length 10–30 μm; purity > 95 wt%) were purchased from Chengdu Organic Chemicals (Chinese Academy Science, China), and were functionalized by different methods: the as-received MWCNTs were chemically oxidized in a mixture of sulfuric acid and nitric acid (3/1, v/v) while being ultrasonicated for 2 h (Method A [22]); the as-received MWCNTs were treated in boiled nitrate solution (20 wt%) for 1 h for surface functionalization (Method B [21]); the as-received MWCNTs were dispersed in 5.0 M HNO₃ solution and refluxed for 48 h at 140°C (Method C). These functionalized MWCNTs were washed with water to neutral, dried under vacuum, and denoted as MWCNT (A), MWCNT (B), and MWCNT (C), respectively.

2.2. Preparation and Characterization of MWCNT-TiO₂ Nanocomposites. All other chemical reagents used in the study were of analytical grade and used without further purification, unless stated otherwise. The preparation procedure for MWCNT-TiO₂ nanocomposites is demonstrated in Figure 1. In a typical experiment, pretreated MWCNTs were ultrasonically dispersed in deionized water, and then titanium sulfate (Ti(SO₄)₂) was added into the dispersion under stirring. The obtained mixture was added into cetyltrimethylammonium bromide (CTAB) solution under the molar ratio

of Ti(SO₄)₂:CTAB:H₂O is 1:0.12:100. After stirring, the resulting mixture (pH 0.2) was aged at room temperature for 12 h and then transferred into an autoclave for 72 h hydrothermal treatment at 100°C. The resulting materials were collected using the centrifugation technique and mixed with a water and ethanol (molar ratio 1:1) solution of sodium chloride under stirring at 40°C for 5 h. Resultant sample was washed with water and ethanol, dried at 80°C overnight, and calcined at 400°C for 5 h.

X-ray diffraction (XRD) patterns were obtained on a XRD-6000 diffractometer using Cu Kα as radiation (λ = 0.15418 nm). Scanning electron microscope (SEM) observation was conducted on a JEOL-6700F electron microscope. Diffuse reflectance spectra (DRS) were recorded on a Cary 5000 UV-Vis-NIR spectrophotometer equipped with an integrating sphere (Varian, USA). Thermogravimetry (TG) curves were recorded on a STA 449C thermal analyzer (Netzsch, Germany). The Brunauer-Emmett-Teller (BET) surface areas were analyzed by nitrogen adsorption-desorption measurement using a Micromeritics ASAP 2020 apparatus after the samples were degassed at 180°C.

2.3. Photocatalytic Activity Measurement. The obtained MWCNT-TiO₂ nanocomposite was loaded with Pt and tested for its photocatalytic activity through a photocatalytic hydrogen evolution system as described in our previous publication [21, 23]. A 300 W Xe-lamp (PLS-SXE300C, Beijing Trusttech Co., Ltd., China) was applied as the light source, which was collimated and focalized into 5 cm² parallel faculae. A cut-off filter (Kenko, L-42; λ > 420 nm) was employed to obtain the visible-light irradiation (λ > 420 nm). The reaction was performed in a water suspension which contains 85 mL water, 15 mL triethanolamine (TEOA), and 40 mg photocatalyst. The suspension was irradiated from top of the system after thoroughly removing air. H₂ production rate was analyzed with a gas chromatograph (GC, SP-6800A, TCD detector, 5 Å molecular sieve columns, and Ar carrier).

The apparent quantum efficiency (AQE) was measured under the same photocatalytic reaction condition except for

TABLE 1: Summary of the physicochemical properties of MWNT and MWNT-TiO₂ nanocomposite.

Sample	Crystal size ^a (nm)	C _{TG} ^b (wt%)	S _{BET} (m ² /g)	Mean pore size ^c (nm)	Total volume ^d (cm ³ /g)
1.25 wt% MWNT-TiO ₂	12.8	1.5	110.8	10.9	0.35
2.5 wt% MWNT-TiO ₂	12.1	3.2	97.5	12.4	0.36
5 wt% MWNT-TiO ₂	12.9	5.7	79.8	12.1	0.28
10 wt% MWNT-TiO ₂	12.6	9.9	110.1	11.8	0.37
20 wt% MWNT-TiO ₂	11.8	18.5	166.7	9.1	0.42
MWNT	—	—	392.5	7.6	0.74

^aCalculated by the Scherrer equation.

^bCarbon content determined by TGA-DSC analyses.

^cAverage pore diameter calculated from BJH desorption average pore width (4 V/A).

^dSingle point total pore volume at the relative pressure of *ca.* 0.995.

the incident monochromatic light wavelength. The hydrogen yields of 1 h photocatalytic reaction under visible-light with different wavelengths of 350, 365, 380, 420, 435, 450, 475, 500, 520, and 550 nm were measured. Each run was carried out three times and the average value was taken (the relative errors are under 10%). The band-pass and cut-off filters and a calibrated Si photodiode (SRC-1000-TC-QZ-N, Oriel, USA) were used in measurement. Apparent quantum efficiencies at different wavelengths were calculated by the following equation:

$$\begin{aligned}
 \text{AQE (\%)} &= \frac{\text{The number of reacted electrons}}{\text{The number of incident photons}} \times 100 \\
 &= \frac{2 \times \text{The number of evolved H}_2 \text{ molecules}}{\text{The number of incident photons}} \times 100.
 \end{aligned} \tag{1}$$

3. Results and Discussion

Figure 2 shows the XRD patterns of MWCNT, TiO₂, and various MWCNT-TiO₂ nanocomposites. As can be seen from Figure 2(a), the main diffraction peaks of various products can be ascribed to anatase TiO₂, whereas a decrease in the crystallinity of anatase can be found after the introduction of MWCNTs, indicating the decrease in the grain size of TiO₂. The characteristic peaks for CNTs at $2\theta = 26.0^\circ$ and 43.4° were not observed for 1.25–10 wt% MWCNT-TiO₂, which is different from the previous observation [24]. This phenomenon could be attributed to the good dispersion of MWCNTs in the nanocomposite after surface functionalization in nitrate solution [21]. Average crystal sizes calculated using Scherrer equation [25] from the broadening of the (101) peaks of the anatase are 18.1, 12.8, 12.1, 12.9, 12.6, and 11.8 nm for TiO₂ and 1.25~20 wt% MWCNT-TiO₂, respectively (see Table 1). The small grain of TiO₂ nanoparticles in MWCNT-TiO₂ may be attributed to restricted direct contact of grains due to the presence of MWCNTs.

As can be observed from Figure 2(b), even for MWCNT-TiO₂ calcined at 800°C for 5 h, the XRD pattern shows that all of the crystal phases are still anatase; no peak of rutile appears. Average crystal sizes calculated from the broadening of the (101) peaks of the anatase are 11.6, 12.9, 13.6, 17.8, and 32.9 nm for the 5 wt% MWCNT-TiO₂ as-synthesized and calcined at

400, 500, 600, and 800°C, respectively. These results suggest that MWCNTs in the nanocomposite probably inhibit the phase transformation of TiO₂ from amorphous phase to anatase phase and lead to a higher thermal stability.

To estimate the real content of MWCNTs in composites, 1.25–10 wt% MWCNT-TiO₂ were analyzed by TGA technique. The results shown in Table 1 suggest that the MWCNT/TiO₂ ratios estimated before the synthesis of the MWCNT-TiO₂ were consistent with the results obtained from TGA analysis. Therefore, negligible losses of MWCNTs occurred during the nanocomposite preparation procedure, which is in accordance with the results of Raman analysis on 5 wt% MWCNT-TiO₂ [21]. The BET specific surface areas and pore volumes of the samples are summarized in Table 1. Compared with mesoporous TiO₂ nanoparticles prepared via hydrothermal processes [26], there is a rapid decrease in both BET surface area (from 318 to 111 m²/g) and total volume (from 0.61 to 0.35 cm³/g) of TiO₂ nanoparticles after modification with 1.25 wt% MWCNT, which could be due to the close-packed structure between TiO₂ nanoparticles and MWCNTs. With increasing MWCNT content, the BET surface area of MWCNT-TiO₂ decreased firstly and then increased, which can be attributed to the destruction of close-packing due to the self-agglomeration of MWCNTs or TiO₂ nanoparticles at high content.

The UV-vis spectra of MWCNT-TiO₂ display a similar absorption edge to TiO₂ (Figure 3), but an apparent enhancement of absorption throughout the visible-light region can be observed even for the nanocomposite containing 1.25 wt% MWCNTs. A correlation between the MWCNTs amount and absorption changes in the UV-vis spectra obviously features the enhancement of visible-light absorption upon increasing the MWCNT content; that is, the adsorption intensity of the present MWCNT-TiO₂ continuously increased with enhancing MWCNT content owing to its good dispersion.

Control experiments showed no appreciable H₂ evolution in the absence of either photocatalyst or irradiation under visible-light or full spectra irradiation. H₂ generation rates over 5 wt% MWCNT + TiO₂ (a simple mixture of MWCNTs and TiO₂ nanoparticles), 5 wt% MWCNT-TiO₂, TiO₂, and MWCNTs under the visible-light and full spectra irradiation are shown in Figure 4. The pristine TiO₂ and MWCNTs as well as the MWCNT + TiO₂ demonstrate no appreciable H₂ evolution under the visible-light irradiation, whereas

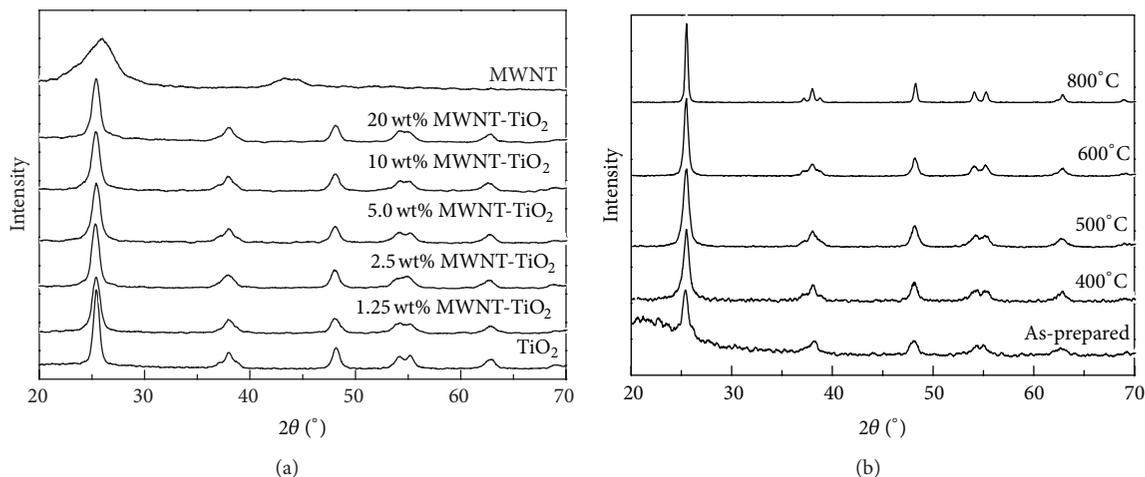


FIGURE 2: XRD patterns of MWCNT, TiO_2 , and MWCNT- TiO_2 with (a) different MWCNT content; (b) 5 wt% MWCNT and calcination at different temperatures.

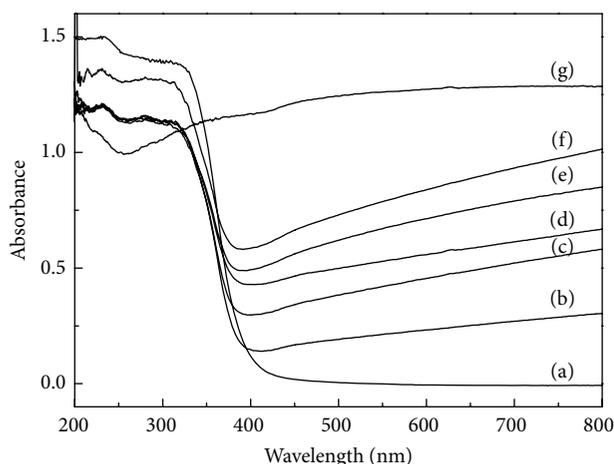


FIGURE 3: DRS patterns of MWCNT, TiO_2 , and MWCNT- TiO_2 with different MWCNT content. (a) TiO_2 ; (b) 1.25 wt%; (c) 2.5 wt%; (d) 5 wt%; (e) 10 wt%; (f) 20 wt%; (g) MWCNT.

MWCNT- TiO_2 (A, B, and C) exhibits various H_2 generation rates, suggesting the importance of chemical linking between MWCNTs and TiO_2 for their visible-light-driven photoactivity [27]. The pretreatment of MWCNTs has a strong effect on the photocatalytic H_2 evolution efficiency over MWCNT- TiO_2 . MWCNT- TiO_2 (B) demonstrates the highest H_2 evolution rate under both visible-light and full spectra irradiation.

Different oxidizing reagents possess different degrees of oxidation power, which would purify and decorate MWCNTs with oxygen-containing groups or even destroy their nanotube structure [28]. It is reported that MWCNT bundles appear exfoliated and curled after treatment with strong oxidative environment such as refluxing in nitric acid or stirring in piranha (mixture of sulphuric acid 96 wt% and hydrogen peroxide 30 wt% in ratio 70 : 30) [29]. Comparing

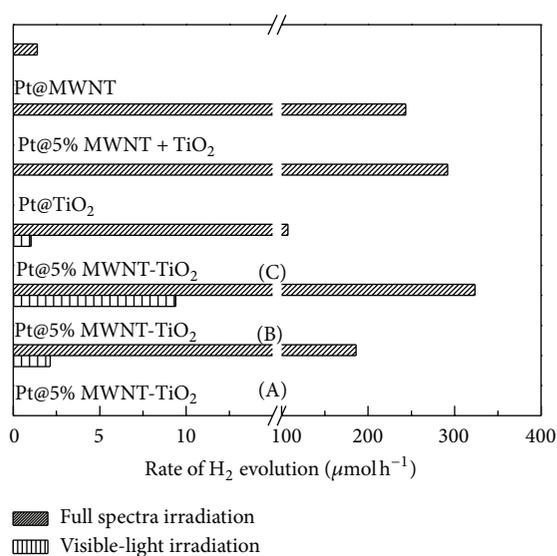


FIGURE 4: H_2 evolution efficiency over MWCNT, TiO_2 , and various photocatalysts containing MWCNT and TiO_2 under visible-light ($\lambda > 420 \text{ nm}$) and full spectra irradiation. Conditions: 1 wt% Pt-loading; 15 vol % TEOA solution (100 mL); light source, 300 W Xe-lamp. MWCNTs + TiO_2 : MWCNTs and TiO_2 nanoparticles were simply mixed together.

the method of refluxing in nitric acid with that of sonicating in a mixture of sulfuric acid and nitric acid, the method of refluxing in nitrate solution provides a moderate oxidation that would generate oxygen-containing groups but would not destroy the MWCNTs structure. Therefore, MWCNT- TiO_2 (B) exhibited the highest photoactivity.

As demonstrated in Figure 5, MWCNT- TiO_2 exhibits no photocatalytic activity until the MWCNT content enhanced to 3.5 wt% under visible-light irradiation. After that, its photocatalytic H_2 evolution efficiency increases firstly

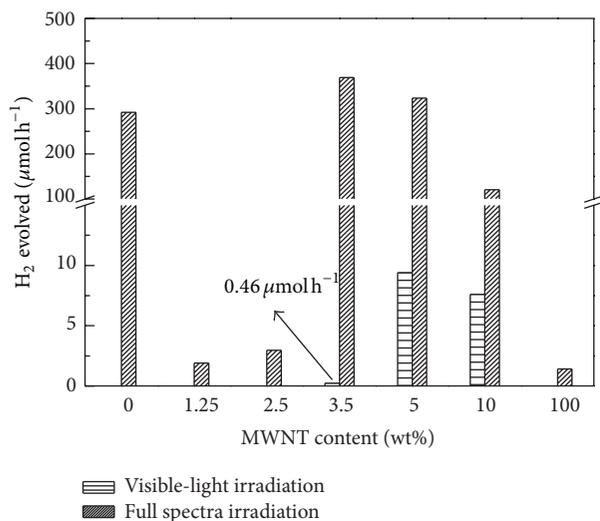


FIGURE 5: The rate of H₂ evolution on MWCNT-TiO₂ nanocomposites with different MWCNT content under visible-light ($\lambda > 420$ nm) and full spectra irradiation. Conditions: 1 wt% Pt-loading; 15 vol % TEOA solution (100 mL); light source, 300 W Xe-lamp.

and then decreases slightly with further enhancement of the MWCNT content. The maximum efficiency is achieved at 5 wt% MWCNT-TiO₂ under visible-light irradiation.

It is believed that TiO₂ nanoparticles could directly grow on the surface of the functionalized MWCNTs through the hydrothermal treatment [21]. As shown in Figure 6, no MWCNTs were found in the SEM micrograph of 2.5 wt% MWCNT-TiO₂, which can be attributed to the fact that MWCNTs are embedded inside the nanocomposite by TiO₂ nanoparticles, resulting from the direct growth of TiO₂ on the surface of MWCNTs. For 5 wt% MWCNT-TiO₂, MWCNTs can be observed on the surface of nanocomposite owing to the increase of MWCNT content. These observations can fairly explain the effect of MWCNT content on the visible-light-induced photoactivity. For the MWCNT-TiO₂ with MWCNT content less than 3.5 wt%, MWCNTs embedded inside the nanocomposite cannot be irradiated and excited by visible-light, and thus no visible-light-induced photoactivity was observed [30]. As the visible-light absorbent and sensitizer, higher MWCNT content means more efficient visible-light absorption, and more photogenerated electrons can be transferred to TiO₂; TiO₂ also plays an important role in the separation of photogenerated carriers: the electrons can be transferred from TiO₂ to the loaded Pt. It is reported [31] that the electrical conductivity in the interfacial contact between graphene and photocatalyst components is vital to the overall photocatalytic H₂ production. In addition, unintentional doping of TiO₂ may happen under annealing at 400°C [32, 33], which can increase the visible-light absorbance of TiO₂ and MWCNT-TiO₂. Therefore, there exists an optimal ratio of MWCNT to TiO₂ for achieving excellent electrical conductivity in the nanocomposites and significant photoactivity for H₂ evolution [30].

Under full spectra irradiation, MWCNT-TiO₂ with small MWCNT contents (1.25 wt% and 2.5 wt%) exhibits lower

photoactivity than the pure titania, whereas 5 wt% and 3.5 wt% MWCNT-TiO₂ demonstrate the better and the best photoactivity, respectively. Carbon materials such as graphene, C₆₀, and CNT can act as electron traps due to their high electron affinities [31, 34, 35], which would be functionalized as charge separators and enhance the photoactivity [30, 36]. However, as discussed above, for the MWCNT-TiO₂ with MWCNT content less than 3.5 wt%, MWCNTs embedded inside the nanocomposite could act as recombination center of electron and holes, resulting in a reduced photoactivity under full spectra irradiation.

It is well understood that the hydrothermal temperature plays an important role in the morphology, crystallinity, and particle size of TiO₂ [37], which are related to the photoactivity. The effect of hydrothermal temperature on the H₂ generation rate was evaluated under visible-light irradiation. As demonstrated in Figure 7, the H₂ generation rate over 5 wt% MWCNT-TiO₂ increases with the enhanced hydrothermal temperature before 140°C and decreases afterwards under both visible-light and full spectra irradiation. The highest H₂ generation rate of 15.1 μmol·h⁻¹ under visible-light irradiation was obtained with 140°C hydrothermal treatment, whereas that of 323.7 μmol·h⁻¹ under full spectra irradiation was obtained with 100°C hydrothermal treatment.

The above experimental results could be rationalized by the following discussions. As can be seen from Figure 8, the XRD patterns of 5 wt% MWCNT-TiO₂ derived from different hydrothermal temperatures confirm the fact that the crystallinity of MWCNT-TiO₂ increases with enhancing hydrothermal temperature. It can be calculated using Scherrer equation [25] that average crystal sizes of the anatase are 8.3, 12.9, 15.7, and 26.7 nm for the 5 wt% MWCNT-TiO₂ with hydrothermal temperature of 80, 100, 140, and 180°C, respectively [38]. Under visible-light irradiation, MWCNTs, as a photosensitizer, can absorb visible-light and the photo-generated electrons (e⁻) can be excited from the VB to CB of the MWCNT [39]. TiO₂, as the “bridge” between MWCNTs and loaded Pt, would transfer the photogenerated electrons from MWCNT to Pt nanoparticles to generate H₂ from water reduction [40]. As the electron acceptor and transfer station, TiO₂ with high crystallinity and tight combination with MWCNTs could benefit the electron injections and suppress the electron-hole recombination. On the other hand, high crystallinity of TiO₂ would inevitably lead to large crystal size and particle size, which would lead to a high electron-hole recombination rate owing to the long distance for electron transfer. As a result, 5 wt% MWCNT-TiO₂ derived from hydrothermal treatment at 140°C has the highest photoactivity under visible-light irradiation.

Under full spectra irradiation, photocatalytic H₂ generation over TiO₂ irradiated by UV light would make the best part comparing to that over MWCNT-TiO₂ irradiated by visible-light. 5 wt% MWCNT-TiO₂ derived from hydrothermal treatment at 100°C possesses moderate crystal size of anatase, resulting in the highest photoactivity due to the large surface area of MWCNT-TiO₂ and low electron-hole recombination rate.

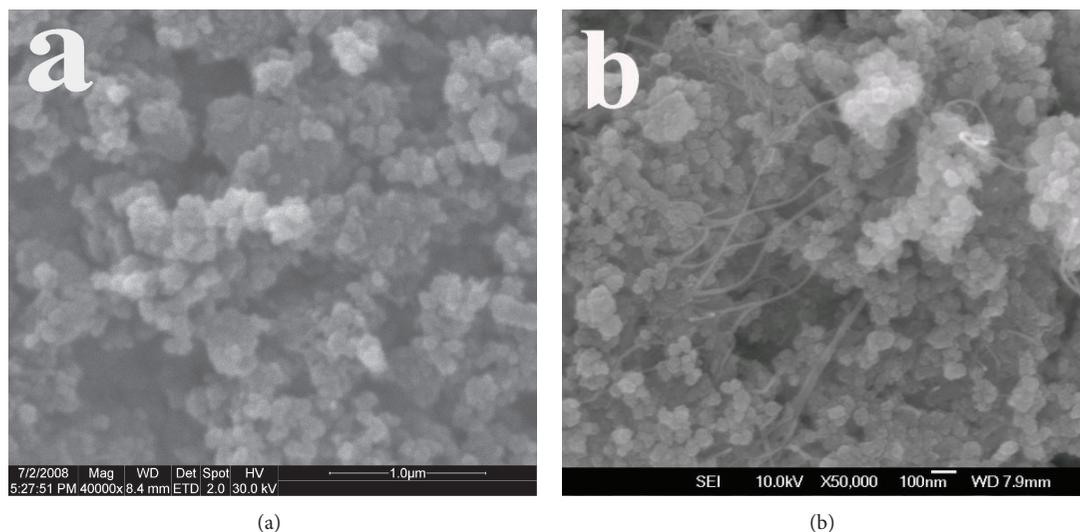


FIGURE 6: SEM micrographs of 2.5 wt% (a) and 5 wt% (b) MWCNT-TiO₂ nanocomposite.

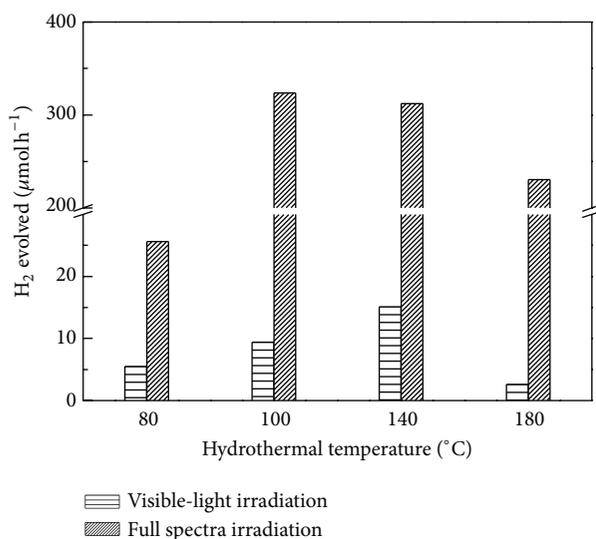


FIGURE 7: Effect of hydrothermal temperature on photocatalytic H₂ production over 5 wt% MWCNT-TiO₂.

The photocatalytic activities for H₂ production over 5 wt% MWCNT-TiO₂ upon incident light with different wavelength were investigated. MWCNT-TiO₂ shows a relatively wide photoresponse under monochromatic light of wavelength ranging from 350 to 475 nm (Figure 9), which is consistent with the experimental results obtained with visible-light irradiation (see Figure 4). The apparent quantum efficiency (AQE) of MWCNT-TiO₂ as a function of the wavelength of the incident light was calculated according to the data in Figure 9. MWCNT-TiO₂ demonstrates high quantum efficiency upon irradiation with wavelengths of 420 and 475 nm (4.4% and 3.7%, resp.). With respect to the wide adsorption band in the entire visible-light region (Figure 3), a panchromatic photocatalytic activity could be

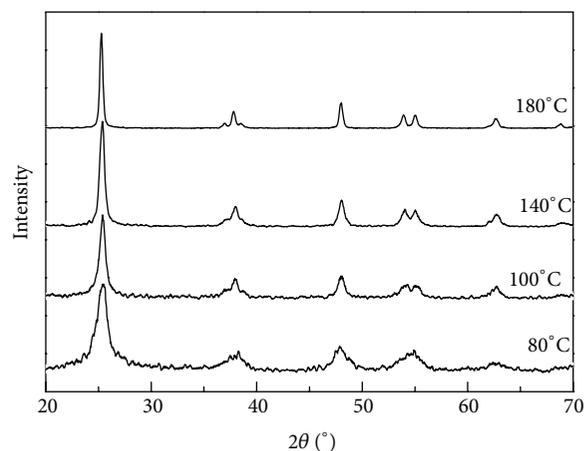


FIGURE 8: XRD patterns of 5 wt% MWCNT-TiO₂ derived from different hydrothermal temperatures for 72 h.

expected for 5 wt% MWCNT-TiO₂. However, no appreciable H₂ evolution is showed upon irradiation with wavelength longer than 500 nm. As demonstrated in our earlier study, not all MWCNTs in the nanocomposite are bound with TiO₂ nanoparticles, owing to the limited hydroxyl and carboxyl groups generated on the MWCNTs after the functionalization [21]. Therefore, the absorption band between 400 and 800 nm of MWCNT-TiO₂ nanocomposite probably is a combination of the absorption spectrum of both MWCNT-TiO₂ and uncoupled MWCNT, which would absorb light with wavelength longer than 500 nm but cannot be excited to generate electron-hole pairs. On the other hand, the energy of the incident light at longer wavelength dramatically decreased, which would inevitably depress the photoactivity over MWCNT-TiO₂. Furthermore, the photoactivity of TiO₂ under UV light irradiation is enhanced after coupling with MWCNT. The quantum efficiencies for H₂ production over

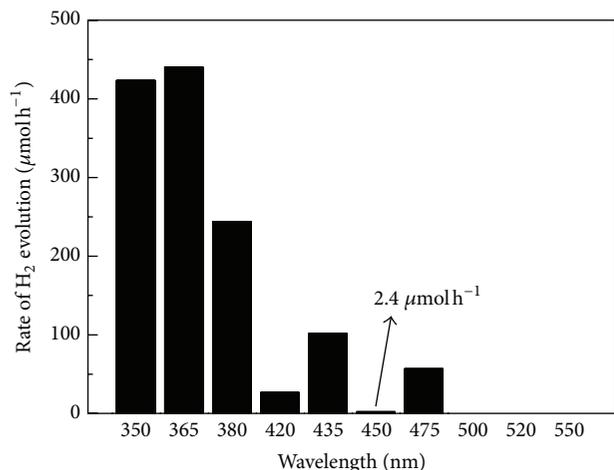


FIGURE 9: The dependence of photocatalytic activities for H₂ production over 5 wt% MWCNT-TiO₂ nanocomposite upon wavelength (controlled via cut-off filters).

P25 TiO₂, mesoporous TiO₂, and MWCNT-TiO₂ at 350 nm were measured to be 1.9, 2.5, and 3.8%, respectively. These phenomena were consistent with the previous discussion that MWCNTs could increase the carrier separation efficiency during photocatalytic process under full spectra irradiation.

4. Conclusions

A series of MWCNT-TiO₂ nanocomposites with different types of functionalized MWCNTs were synthesized through hydrothermal process and characterized by XRD, DRS, SEM, TGA, and BET techniques. The effects of pretreatment methods on MWCNT, MWCNT content, and hydrothermal temperature on the photocatalytic hydrogen evolution efficiency over MWCNT-TiO₂ nanocomposite were investigated. MWCNTs have a good dispersion in the nanocomposite, which inhibit grain growth of TiO₂ and improve its thermal stability. Appropriate pretreatment on MWCNTs could generate oxygen-containing groups, which would become the anchoring sites with TiO₂ nanoparticles in the nanocomposite. However, chemical pretreatment with strong oxidative agents would destroy the intrinsic structure of MWCNTs, which is not beneficial for a high photocatalytic activity. The best photocatalytic activity was observed for the MWCNT-TiO₂ nanocomposite with a 5% weight ratio under visible-light irradiation. The photocatalytic activity of MWCNT-TiO₂ is related to the crystallinity of TiO₂, link type between MWCNT and TiO₂, and MWCNT content. A wide range of photoresponse from 350 to 475 nm is observed with high quantum efficiency. Upon irradiation with wavelengths of 420 and 475 nm, the quantum efficiency is 4.4% and 3.7%, respectively. The above experimental results indicate that the present MWCNT-TiO₂ nanocomposite is a promising photocatalyst with good thermal stability, chemical stability under UV light irradiation, and visible-light-induced photoactivity.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

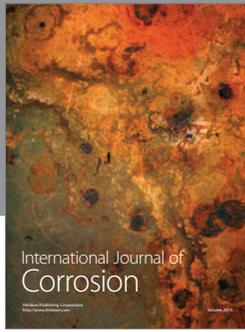
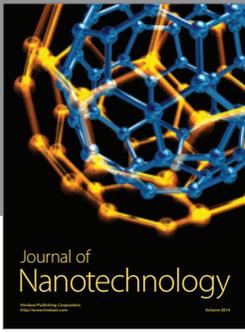
Acknowledgments

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