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# TiO<sub>2</sub>-Au composite nanofibers for photocatalytic hydrogen evolution

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TiO<sub>2</sub>-based materials for photocatalytic hydrogen (H<sub>2</sub>) evolution have attracted much interest as a renewable approach for clean energy applications. TiO<sub>2</sub>-Au composite nanofibers (NFs) with an average fiber diameter of ~160 nm have been fabricated by electrospinning combined with calcination treatment. *In situ* reduced gold nanoparticles (NPs) with uniform size (~10 nm) are found to disperse homogeneously in the TiO<sub>2</sub> NF matrix. The TiO<sub>2</sub>-Au composite NFs catalyst can significantly enhance the photocatalytic H<sub>2</sub> generation with an extremely high rate of 12 440 μmol g<sup>-1</sup> h<sup>-1</sup>, corresponding to an adequate apparent quantum yield of 5.11% at 400 nm, which is 25 times and 10 times those of P25 (584 μmol g<sup>-1</sup> h<sup>-1</sup>) and pure TiO<sub>2</sub> NFs (1254 μmol g<sup>-1</sup> h<sup>-1</sup>), respectively. Furthermore, detailed studies indicate that the H<sub>2</sub> evolution efficiency of the TiO<sub>2</sub>-Au composite NF catalyst is highly dependent on the gold content. This work provides a strategy to develop highly efficient catalysts for H<sub>2</sub> evolution.

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## Introduction

Photocatalytic hydrogen (H<sub>2</sub>) evolution has drawn great attention as a promising approach to overcome the energy crisis by converting solar energy to H<sub>2</sub> fuel directly.<sup>1-3</sup> On the basis of Fujishima's pioneering work,<sup>4</sup> numerous semiconductor-based photocatalysts have been developed to produce H<sub>2</sub> (such as TiO<sub>2</sub>,<sup>5</sup> CdS,<sup>6</sup> ZnO,<sup>7</sup> Cu<sub>2</sub>O,<sup>8</sup> etc.). TiO<sub>2</sub> has attracted much attention owing to its excellent properties including low cost, environmental friendliness, strong oxidizing power and high stability.<sup>9</sup> However, the photocatalytic efficiency of TiO<sub>2</sub>-based materials is still limited, with a low solar light harvesting capability due to its wide band gap and high charge recombination rate.<sup>10</sup> Therefore, it is important to develop TiO<sub>2</sub>-based photocatalysts for increasing the light harvesting and promoting the photoinduced charge separation during the H<sub>2</sub> generation process.

Numerous attempts have been devoted to improving the H<sub>2</sub> generation efficiency of TiO<sub>2</sub> materials including doping,<sup>11</sup> metal deposition,<sup>10,12</sup> and heterostructures formation.<sup>13</sup> Chiu<sup>14</sup> has designed a quaternary Ti-Nb-Ta-Zr-O mixed-oxide nanotube arrays; Hsieh<sup>15</sup> has conducted compact deposition of In<sub>2</sub>S<sub>3</sub> nanocrystals on the TiO<sub>2</sub> nanowire array; Li<sup>16</sup> has prepared by electrodepositing a Cu<sub>2</sub>O layer on the surface of Au particle-coated TiO<sub>2</sub> arrays; Chang<sup>17</sup> has synthesized CdS/CdSe co-sensitized brookite TiO<sub>2</sub> nanostructures with hydrogen doping

(H:TiO<sub>2</sub>/CdS/CdSe) in a facile solution reaction. All these TiO<sub>2</sub>-based materials were developed to enhance the amount of accessible charge carriers, modify the electronic structure, improve the hole injection kinetics for the photoelectrochemical water splitting.

Combining TiO<sub>2</sub> with highly dispersed noble metal nanoparticles (NPs) as cocatalyst is a feasible approach for enhancing the photocatalytic performance.<sup>18</sup> Among the noble metal NPs (*i.e.* Au,<sup>19,20</sup> Ag,<sup>21</sup> Pt,<sup>22</sup> Pd<sup>23,24</sup>), gold with finite NPs size can spatially separate the photogenerated charge carriers.<sup>3,25</sup> Much efforts have been focused on TiO<sub>2</sub> NPs as matrix and different size or shape of gold NPs, to understand the fine-tuning the surface plasmon resonance effect.<sup>26</sup> Pu and his colleagues<sup>27</sup> have reported that the photocatalytic ability of TiO<sub>2</sub> nanowires can be effectively enhanced in the entire UV-visible region by adopting different shapes of the decorated Au nanostructures. Compared to other nanostructures, TiO<sub>2</sub> nanofibers (NFs) possess additional advantages due to its specific morphologies with large aspect ratio and large specific surface area for photocatalytic activity.<sup>28</sup> Yoshikawa<sup>29</sup> has proved that the TiO<sub>2</sub> NFs catalyst can promote the opportunities for photocatalytic H<sub>2</sub> evolution. Xia and Li<sup>30,31</sup> successfully fabricated the TiO<sub>2</sub> NFs and its composite deposited with gold NPs. However, there are seldom literatures studying the TiO<sub>2</sub>-Au composite NFs as photocatalyst for H<sub>2</sub> generation and the effect of gold content on photocatalytic performance.

Herein, TiO<sub>2</sub>-Au NFs composites with different gold content were fabricated *via* electrospinning combined with subsequent calcination. The gold NPs are *in situ* reduced during the calcination process while the polymer components are decomposed and removed to form TiO<sub>2</sub> NFs.<sup>32</sup> The gold content is tuned by controlling the amount of gold precursor added during

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synthesis process, and all the chemicals are mixed homogeneously in solution with molecular level. The  $H_2$  evolution with  $TiO_2$ -Au NFs catalyst has been investigated.

## Methods

### Materials

Polyvinylpyrrolidone (PVP,  $M_w \sim 1\,300\,000$ ) and chloroauric acid ( $HAuCl_4 \cdot 3H_2O$ ,  $\geq 99.9\%$ ) were purchased from Aldrich (St. Louis, MO, USA). Titanium isopropoxide (TTIP,  $C_{12}H_{28}O_4Ti$ ,  $\geq 99.9\%$ ), ethanol (EtOH) and acetic acid (HAC) were obtained from Chengdu Kelong Chemical Reagent Factory (Chengdu, China). All chemical reagents were analytically pure and used directly without further purification.

### Fabrication of $TiO_2$ -Au composite NFs with different gold content

$TiO_2$ -Au composite NFs catalysts were synthesized through electrospinning (ES) combined with subsequent calcination process as shown in the schematic diagram (Fig. 1). In a typical process, PVP were dissolved in EtOH to obtain 7 wt% PVP-EtOH polymer solution, and then 1.5 g of TTIP were added into a mixture of 3 mL HAC and 3 mL EtOH, mixing with 6.43 g of the previous PVP-EtOH solution to form PVP-TTIP ES solution.<sup>33</sup> A given amount of gold precursor ( $HAuCl_4 \cdot 3H_2O$ ) were added directly into the PVP-TTIP solution under vigorous stirring to form PVP-TTIP-Au ES solution and the mass ratio of Au/ $TiO_2$  in the final products were 0 wt%, 3 wt%, 6 wt%, 9 wt%, and 12 wt%, respectively. The as-spun precursor NFs were obtained after a typical ES process, and then calcined at 500 °C for 2 hours in air to get the final products of  $TiO_2$  NFs and  $TiO_2$ -Au composite NFs with different gold content, which were denoted as  $TiO_2$ -Au-0 wt% (equaled to bare  $TiO_2$  NFs),  $TiO_2$ -Au-3 wt%,  $TiO_2$ -Au-6 wt%,  $TiO_2$ -Au-9 wt%, and  $TiO_2$ -Au-12 wt%,

respectively. The commercial titania catalyst of Degussa P25 is chosen as the benchmark.

### Photocatalyst characterization

The morphologies and structures of the  $TiO_2$ -Au composite NFs were characterized by scanning electron microscope (SEM, Inspect F50, FEI, USA) and X-ray diffractometer (XRD, DX-2000, HAOYUAN, China), using Cu  $K\alpha_1$   $\lambda = 1.54056$  Å radiation. The average fiber diameter and particle size of samples were determined with software of Nano Measurer performed on SEM images. Transmission electron microscope (TEM) and high-resolution transmission electron microscope (HR-TEM) images, as well as energy-dispersive X-ray spectra (EDS) were obtained on a JEOL 2100F microscope to evaluate sample morphology and further confirm crystalline structures and elemental compositions. Photoluminescence (PL) spectra were performed with an excitation wavelength of 320 nm by a spectrofluorometer (FluoroMax-4, Horiba, Japan). UV-visible diffuse reflectance spectra (UV-vis) were obtained on a Shimadzu UV 3600 spectrophotometer.

### Photocatalytic activity evaluation

The photocatalytic  $H_2$  generation performances of all the samples were evaluated under full spectrum irradiation (a 300 W xenon arc lamp) for 3 hours. In a typical procedure, 30 mg of each prepared catalysts was dispersed in 150 mL of water/methanol mixture with the volume ratio of 9 : 1 under ultrasonic. In order to avoid the influence of air, argon was purged into the system half an hour to remove the air. And then the as-prepared suspension solution was irradiated under light (xenon lamp, 300 W) with magnetic stirring, and a water jacket was used to remove the thermal catalytic effect. The irradiation intensity for photocatalytic experiment was 380  $mW\ cm^{-2}$ . During the photocatalytic  $H_2$  generation process, the evolved gas was analyzed by using a gas chromatography (GC, 8A, Shimadzu) with an interval of 30 min.<sup>34</sup> Apparent quantum yield (AQY) of  $H_2$  evolution were determined by using the equation of  $AQY (\%) = (\text{number of reacted electrons}/\text{number of incident photons}) \times 100\%$ .<sup>35</sup>

## Results and discussion

### Gold content effects on morphologies and structures

The morphologies of  $TiO_2$  NFs and  $TiO_2$ -Au composite NFs with different gold content were investigated by SEM as shown in Fig. 2. It can be seen that the  $TiO_2$  NFs (average diameter of 160 nm) possess a rough surface while the  $TiO_2$ -Au composite NFs still have fibrous morphology, but with well dispersed bright dots, that were identified as gold NPs by XRD and TEM results. The gold NPs disperse homogeneous in all the  $TiO_2$ -Au composite NFs with different gold content. In order to check the effects of gold content on the particle size, the average size of gold nanoparticles for each sample were determined from measurements of SEM images performed on 100 particles using the software Nano Measurer. The gold nanoparticles can be divided in two types: one is uniform gold particles, another is

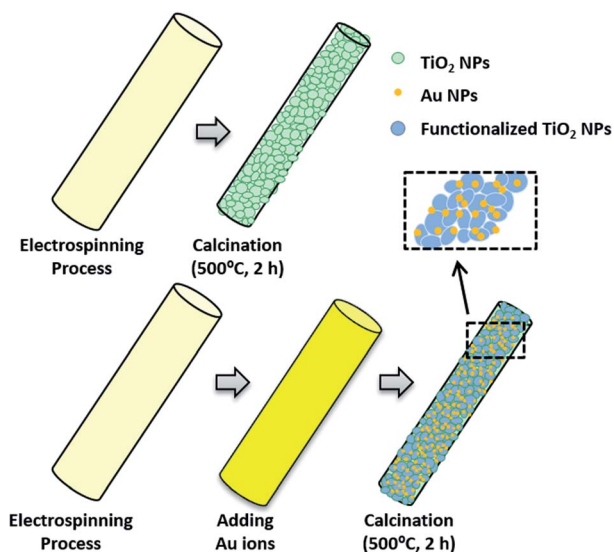


Fig. 1 Schematic diagram illustrating the formation of  $TiO_2$ -Au composite NFs.



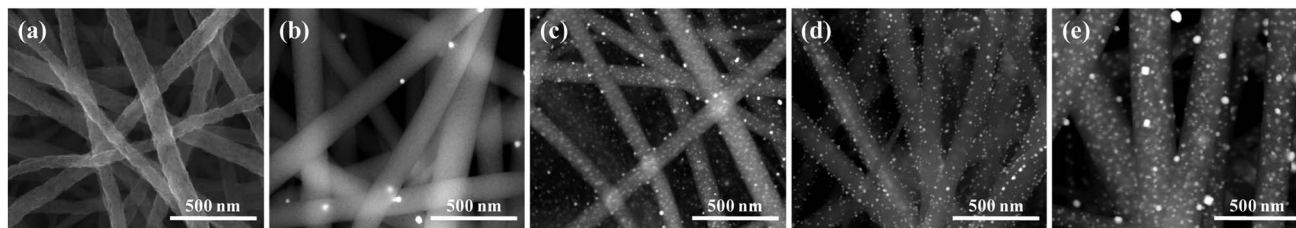


Fig. 2 SEM images of  $\text{TiO}_2$ -Au composite NFs with different gold content calcined at  $500^\circ\text{C}$  in air for 2 h: (a) 0 wt%; (b) 3 wt%; (c) 6 wt%; (d) 9 wt% and (e) 12 wt%.

the nonuniform gold nanoparticles on the surface of  $\text{TiO}_2$  fiber. Firstly, by analyzing the SEM images carefully, it can be seen that the gold nanoparticles with uniform size were of  $\sim 10$  nm, which is contributed to the crystal of  $\text{TiO}_2$  that preventing the growth of gold. And this comments also are confirmed by the related TEM images as shown in Fig. 4.

Secondly, the gold nanoparticles with nonuniform size and disturbing by the gold contents on the surface of  $\text{TiO}_2$  NFs. The nanoparticles are more easily to migrate and aggregate to form larger nanoparticles with increasing of the gold content, that can overcome the inhibition from  $\text{TiO}_2$  crystals. So the higher gold content the larger particle size should make sense in logical. However, the size of particle on the surface didn't have the direct relationship with gold content, it was random form tens of nanometer. The seed formation and particle growth are complicated and influenced by many factors of thermodynamics and kinetics, especially during the calcination process. Thus the size of gold nanoparticle on the surface of  $\text{TiO}_2$ -Au NFs with different contents were nonuniform. As increasing the gold content, the number of gold NPs increase, and the gold content in the final product can be varied precisely by the amount of gold precursor added in the ES solution.

It has been figured out by Wang *et al.*<sup>34</sup> that the intensity of the diffraction peaks of gold grows as increasing the gold content. Thus the sample with highest gold content of 12 wt% has been chosen and analyzed to check the difference between  $\text{TiO}_2$  NFs and  $\text{TiO}_2$ -Au composite NFs more clearly. The crystal structures of P25, pure  $\text{TiO}_2$  NFs and  $\text{TiO}_2$ -Au composite NFs (with gold content of 12 wt%) were determined by XRD as shown in Fig. 3. The XRD patterns of P25 and  $\text{TiO}_2$  can be well-indexed to anatase (JCPDS card no. 00-021-1272). To check carefully, there are three others peaks for P25 could be found at  $2\theta = 27.4^\circ$ ,  $36.1^\circ$ , and  $41.2^\circ$  indexed as the (110), (101), and (111) planes of rutile (JCPDS card no. 00-021-1276), respectively.<sup>32</sup> Comparing to pure  $\text{TiO}_2$ , the XRD patterns of  $\text{TiO}_2$ -Au composite NFs are also well-indexed to anatase and there is no peaks related to rutile phase. Meanwhile, there are three additional peaks located at  $2\theta = 44.4^\circ$ ,  $64.6^\circ$ , and  $77.5^\circ$ , that correspond to the (200), (220), and (311) planes of the gold cubic phase (JCPDS card no.00-004-0784).<sup>32</sup> The commercial Titania Degussa P25 is the composite of anatase and rutile, while the sample of  $\text{TiO}_2$  NFs and  $\text{TiO}_2$ -Au composite NFs are mainly of anatase. These results confirm that the  $\text{TiO}_2$ -Au composite NFs are composed of anatase and gold, and the crystal sizes were

determined to be 20 nm and 8 nm based on the XRD patterns by using Scherrer's equation.

From TEM images (Fig. 4a), it can be seen that the NFs is well decorated with spherical NPs, where the average size is  $\sim 10$  nm. The crystallographic (111) plane of gold and (101) plane of anatase are observed clearly with the corresponding lattice distance of  $2.2 \text{ \AA}$  and  $3.3 \text{ \AA}$  in the HRTEM images shown in Fig. 4b. Hence, the HRTEM results of  $\text{TiO}_2$ -Au composite NFs confirm that the gold NPs are well dispersed into  $\text{TiO}_2$  NFs matrix, which is in accordance with the XRD results. Furthermore, the added elemental composition of gold and titanium (during the preparation of the photocatalysts) were matching with the composition of elements analyzed by EDS shown in Fig. 4c. Murdoch *et al.*<sup>36</sup> have reported that gold NPs with the size range of 3–30 nm deposited on  $\text{TiO}_2$  are highly active, whereas there is no influence on the  $\text{H}_2$  generation rate over the 3–12 nm range. Moreover, gold NPs with similar size has better functionalization effect on anatase than rutile NPs. Thus, the  $\text{TiO}_2$ -Au composite NFs with gold NPs ( $\sim 10$  nm) and anatase matrix of  $\text{TiO}_2$  NFs are supposed to show good photocatalytic performance for  $\text{H}_2$  evolution.

### Gold content effects on optical properties

Photoluminescence (PL) measurement was conducted on all the samples with an excitation wavelength of 320 nm, to evaluate

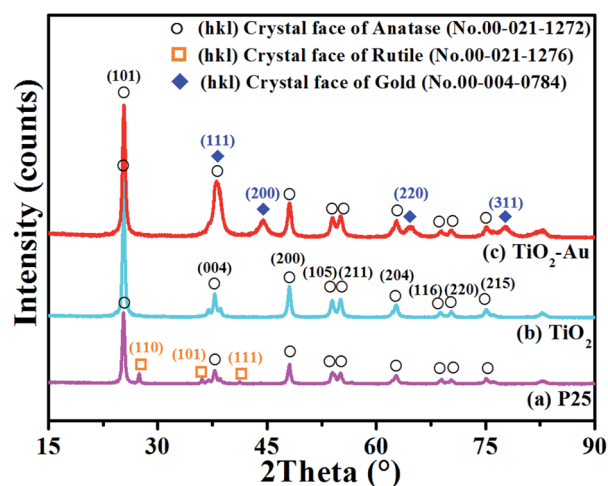


Fig. 3 XRD patterns of P25,  $\text{TiO}_2$  NFs and  $\text{TiO}_2$ -Au-12 wt% composite NFs.



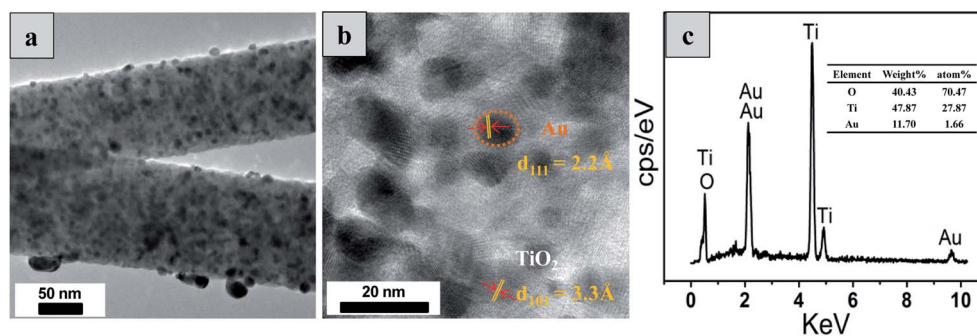


Fig. 4 (a) TEM image of  $\text{TiO}_2$ -Au-12 wt% composite NFs; (b) high-resolution TEM image of Au NPs embedded  $\text{TiO}_2$  NFs showing the Au and  $\text{TiO}_2$  lattices; (c) EDS spectrum of  $\text{TiO}_2$ -Au-12 wt% composite NFs.

the effects of gold content on the photoinduced charge transfer and charge recombination loss.<sup>37</sup> As shown in Fig. 5, the PL spectra of each sample have three main emission peaks located at 394 nm, 413 nm and 467 nm within a broad emission band ranging from 375 nm to 500 nm. Comparing the PL intensity of all the samples, P25 possesses the highest intensity which is followed by  $\text{TiO}_2$ . After doping with gold NPs, the PL intensity of  $\text{TiO}_2$ -Au is lower than  $\text{TiO}_2$ , and the intensity gradually decreases as the gold content increasing. Au as an electron acceptor had a quantitative effect on the mediation of interfacial charge transfer. When the Au content increased, the emission lifetime was reduced, suggesting the significant decrease in the charge carrier separation at the interface.<sup>38</sup> It is reported that a higher PL peak intensity corresponds to a higher

incidence of photogenerated carriers recombination within titanium powders.<sup>39</sup> The lower PL intensity for  $\text{TiO}_2$ -Au samples is attributed to the electrons transfer from  $\text{TiO}_2$  conduction band to gold NPs; in this case the electron-hole recombination is restricted. Thus the photoexcited electrons are confined within gold NPs while holes remain at the  $\text{TiO}_2$  valence band.<sup>40</sup> Consequently,  $\text{TiO}_2$ -Au composite NFs catalyst can significantly inhibit the recombination of photogenerated charge carriers.

The UV-vis absorption spectra of all the samples are shown in Fig. 6. Comparing with the pure  $\text{TiO}_2$  NFs and P25 that can only absorb UV light, the  $\text{TiO}_2$ -Au composite NFs have significant light absorption in the visible region ( $\sim 380$ – $780$  nm) due to the surface plasmon resonance (SPR) effect of gold. In general, the semiconductor band gap is related to the light absorption capability where a decreased band gap indicates the increasing of absorption edge.<sup>41</sup> The absorbance curve of  $\text{TiO}_2$ -Au composite NFs exhibited an absorption edge red shift from ultraviolet to visible region compared with that of  $\text{TiO}_2$  NFs. As increasing the gold content, the intensity of the SPR absorption peak in the visible region is dramatically increased, while the position of the SPR absorption peak shifted non-monotonically. The position and intensity of SPR absorbance peak are strongly depended on multiple factors, such as environment dielectric properties, the particle size, shape, size distribution and contents of gold NPs.<sup>42</sup> The gold NPs of  $\text{TiO}_2$ -Au composite NFs with different contents have different average size and distribution that was discussed in detail at the SEM parts. This could be the possible explanation for the non-monotonically shift.

As shown in Fig. 6b, the Tauc plots based on the UV-vis curves determine the band gaps, that are 3.15 eV and 2.75 eV for  $\text{TiO}_2$  NFs and  $\text{TiO}_2$ -Au composite NFs, respectively.<sup>43,44</sup> The reduced band gap of  $\text{TiO}_2$ -Au composite NFs suggests that the presence of gold NPs can effectively increase the absorption edge and enhance the photocatalytic efficiency. It is also beneficial for extending the light absorption to the visible light range, that is one of the vital factors to improve the photocatalytic  $\text{H}_2$  generation performance.<sup>45</sup> Therefore,  $\text{TiO}_2$ -Au composite NFs photocatalyst can be excited by visible light, resulting in an enhancement of  $\text{H}_2$  generation.

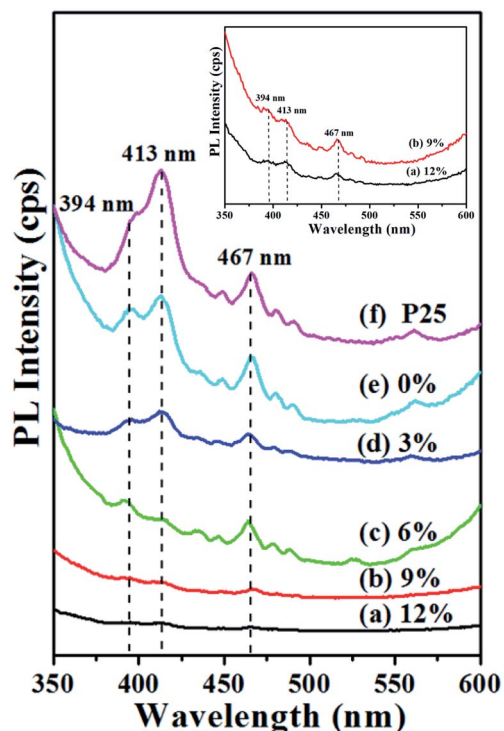


Fig. 5 Photoluminescence spectra of P25,  $\text{TiO}_2$  NFs, and  $\text{TiO}_2$ -Au composite NFs with different gold content.



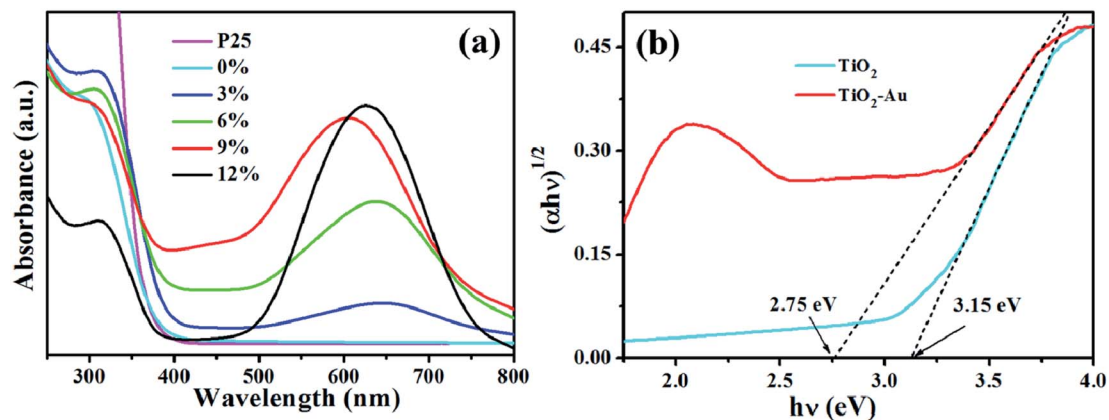


Fig. 6 UV-visible absorption spectra of (a) P25, pure  $\text{TiO}_2$  and  $\text{TiO}_2$ -Au composite with different gold content; (b) Tauc plots of  $\text{TiO}_2$  and  $\text{TiO}_2$ -Au-9 wt%.

### Gold content effects on photocatalytic $\text{H}_2$ evolution

The  $\text{H}_2$  evolution activity was compared among the P25,  $\text{TiO}_2$  NFs,  $\text{TiO}_2$ -Au composites NFs with different gold content, and the  $\text{H}_2$  evolution rate is compared by normalized as per gram of catalyst and per hour of irradiation time (as shown in Fig. 7). The linear increase of all the samples in the evolved  $\text{H}_2$  indicates the stability of the catalyst. The lowest  $\text{H}_2$  evolution activity of  $584 \mu\text{mol g}^{-1} \text{h}^{-1}$  was obtained from commercial P25 with the specific surface area of  $53.2 \text{ m}^2 \text{ g}^{-1}$ , while the activity increases more than 2 times of  $\text{TiO}_2$  NFs ( $1254 \mu\text{mol g}^{-1} \text{h}^{-1}$ ) with the specific surface area of  $18.7 \text{ m}^2 \text{ g}^{-1}$ . The photocatalytic efficiency were promoted for  $\text{TiO}_2$ -Au composite NFs, attributing to the excitation of surface plasmon band by Au NPs. The excess of Au NPs loading on  $\text{TiO}_2$  can enhance the local electromagnetic field and generate high-energy hot electrons.<sup>46</sup> As the gold content increasing from 3 wt% to 6 wt%, while the specific surface area decreased from  $11.9 \text{ m}^2 \text{ g}^{-1}$  to  $9.5 \text{ m}^2 \text{ g}^{-1}$ , leading to a higher  $\text{H}_2$  generation. The  $\text{TiO}_2$ -Au composite NFs with the gold content of 9 wt%, showing a specific surface area of  $8.4 \text{ m}^2 \text{ g}^{-1}$ , possesses the best photocatalytic performance of  $12440 \mu\text{mol g}^{-1} \text{h}^{-1}$ , which is almost 21 times higher than P25 and nearly 10 times of  $\text{TiO}_2$ . However, further increase of the

gold content to 12 wt% resulted in the  $\text{H}_2$  production of  $1605 \mu\text{mol g}^{-1} \text{h}^{-1}$ , which is comparable with  $\text{TiO}_2$ -Au-3 wt% ( $1602 \mu\text{mol g}^{-1} \text{h}^{-1}$ ). Au with relatively high content may function as recombination centers to invoke interfacial charge recombination instead of promoting charge separation.<sup>46</sup>

In order to exclude the contribution of specific surface area, the  $\text{H}_2$  evolution rate was normalized for each sample. As shown in Table 1, the specific surface areas were decreased from  $18.7 \text{ m}^2 \text{ g}^{-1}$  to  $7.2 \text{ m}^2 \text{ g}^{-1}$ , as the gold contents increased from 0 wt% to 12 wt%. It had the same results that  $\text{TiO}_2$ -Au-9 wt% was the optimal sample of  $1481 \mu\text{mol h}^{-1} \text{ m}^{-2}$ , which was 135 times higher than P25 ( $11 \mu\text{mol h}^{-1} \text{ m}^{-2}$ ) and 22 times better than  $\text{TiO}_2$  ( $67 \mu\text{mol h}^{-1} \text{ m}^{-2}$ ). It indicates that gold content is the predominant factor rather than the specific surface area for the photocatalytic efficiency. Meanwhile, the apparent quantum yield (AQY) of  $\text{H}_2$  evolution for all the catalysts have been determined and provided in Table 1. The highest hydrogen generation rate of  $12440 \mu\text{mol g}^{-1} \text{h}^{-1}$ , corresponding to the AQY of 5.11% at 400 nm, is achieved for  $\text{TiO}_2$ -Au composite NFs with the gold content of 9 wt%. And the AQY are 0.24%, 0.52%, 0.66%, 1.19%, and 0.66% for P25, 0%, 3%, 6%, and 12%, respectively. When Au NPs with a suitable amount (9 wt%) were loading on  $\text{TiO}_2$ , the interfacial charge transfer of NFs was

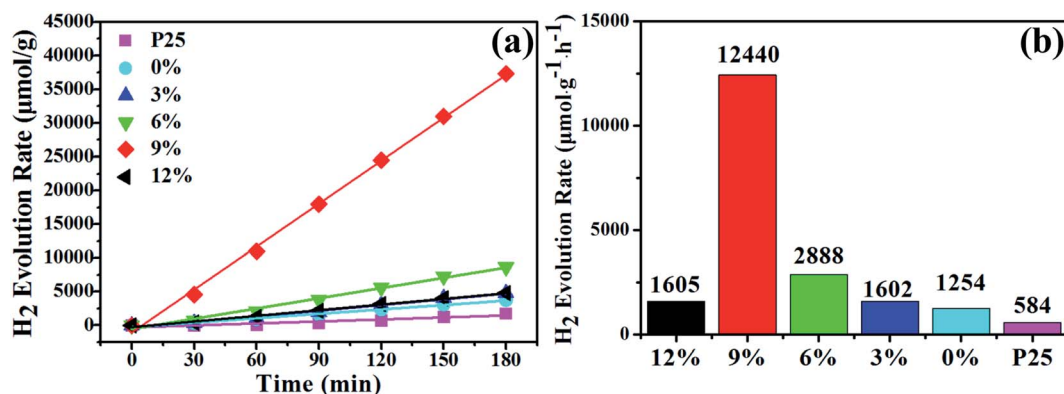
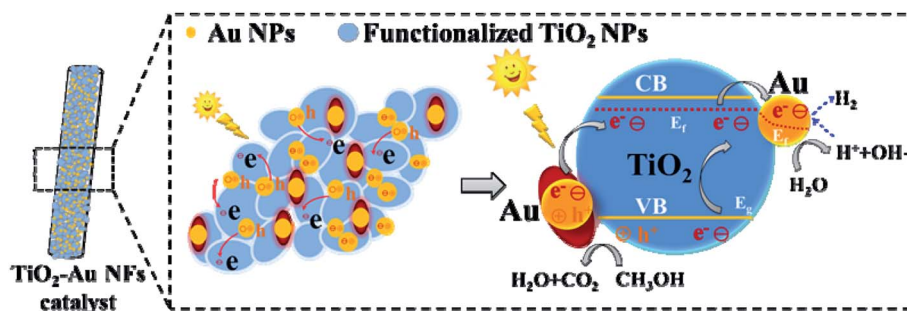


Fig. 7  $\text{H}_2$  evolution of P25,  $\text{TiO}_2$  NFs and  $\text{TiO}_2$ -Au composite NFs catalyst with different gold content under light irradiation.



**Table 1** H<sub>2</sub> evolution rate, apparent quantum yield at 400 nm, specific surface area and H<sub>2</sub> evolution rate normalized with specific surface area of each sample

Sample	P25	0%	3%	6%	9%	12%
H <sub>2</sub> evolution rate/ $\mu\text{mol g}^{-1} \text{h}^{-1}$	584	1254	1602	2888	12 440	1605
Apparent quantum yield at 400 nm	0.24%	0.52%	0.66%	1.19%	5.11%	0.66%
Specific surface area/ $\text{m}^2 \text{g}^{-1}$	53.2	18.7	11.9	9.5	8.4	7.2
H <sub>2</sub> evolution rate normalized with specific surface area/ $\mu\text{mol h}^{-1} \text{m}^{-2}$	11	67	135	304	1481	223



**Fig. 8** Scheme of the photocatalytic H<sub>2</sub> generation mechanism of TiO<sub>2</sub>-Au composite NFs.

highly mediated to render the facilitated electron scavenging from TiO<sub>2</sub>, giving rise to an increased amount of photoexcited free electrons that could be extracted for further utilization.<sup>47</sup> These results indicate that the doping of gold NPs plays a positive role in increasing the H<sub>2</sub> generation efficiency, and the photocatalytic performance also has direct relation with the gold content.

### Photocatalytic mechanism

The possible photocatalytic H<sub>2</sub> generation mechanism of TiO<sub>2</sub>-Au composite NFs from water/methanol under light irradiation is proposed (Fig. 8). The TiO<sub>2</sub>-Au composite NFs catalyst is photoexcited under light irradiation to generate the electron-hole pairs. The excited electrons from the valence band are transferred to conduction band, and then be more easily immigrate to the gold NPs band, due to the positive potential of gold compared with the conduction band level of TiO<sub>2</sub>. Thus, the gold NPs act as efficient sinks for the excited electrons, whereby it can prolong the lifetime of the charge carriers.<sup>19</sup> Meanwhile, the gold NPs can also generate hot electrons owing to the surface plasmon resonance effect.<sup>32</sup> The hot electrons can be transferred to the adjacent TiO<sub>2</sub>, greatly enhancing the potential light-harvesting capabilities of photocatalytic devices. Ratchford has been reported that the electron-injection efficiency from Au NPs to the TiO<sub>2</sub> ranges from about 25% to 45%, and the hot electrons spatial distribution is affected by the plasmonic field, with a higher generation rate for stronger electric field intensities.<sup>48</sup> Due to electromagnetic wave is transverse wave, the mostly incident light propagating approximately along the interface normal that are absorbed, so the optical electric field will be almost parallel to the interface. The relative weak light absorption efficiency and near field

enhancement may be not beneficial to generation of a large number of hot electrons.<sup>49</sup> Thus the content of gold should have an optimal value. The adsorbed H<sub>2</sub>O molecules react with the electrons on gold NPs to form the hydroxyl radicals (OH<sup>•</sup>) and hydrogen ion (H<sup>+</sup>), and then the hydrogen ions (H<sup>+</sup>) are reduced and converted into hydrogen molecules (H<sub>2</sub>) by the photo-generated electrons trapped on the gold NPs. Meanwhile, the methanol (CH<sub>3</sub>OH) molecular as hole scavenger, react with the holes left on the valence band of TiO<sub>2</sub> and to be further oxidized to CO<sub>2</sub> and H<sub>2</sub>O.<sup>34</sup> The gold NPs play two roles during the photocatalytic H<sub>2</sub> evolution process, one is to enhance the light harvesting of TiO<sub>2</sub>-Au composite materials from UV region to visible region and generate more charge carriers under light irradiation by surface plasmon resonance effect, and another one is to work as the electron sink to reduce the recombination rate of electron-hole pairs. This proposed mechanism demonstrates that the TiO<sub>2</sub>-Au composite NFs catalyst can enhance the photocatalytic performance for H<sub>2</sub> production.

### Conclusions

In summary, TiO<sub>2</sub>-Au composite NFs were successfully fabricated by electrospinning combined with subsequent calcination. Gold NPs with uniform size (~10 nm) have been dispersed homogeneously in TiO<sub>2</sub> NFs matrix (with average fiber diameter of 160 nm). The TiO<sub>2</sub>-Au composite NFs photocatalyst showed excellent photocatalytic H<sub>2</sub> generation from a water/methanol mixture under light irradiation. The H<sub>2</sub> production rate of the TiO<sub>2</sub>-Au composite NFs catalyst with the gold content of 9 wt% was dramatically enhanced to 12 440  $\mu\text{mol g}^{-1} \text{h}^{-1}$ , corresponding to an adequate apparent quantum yield of 5.11% at 400 nm, which is 21 times and 10 times higher than that of P25 (584  $\mu\text{mol g}^{-1} \text{h}^{-1}$ ) and TiO<sub>2</sub> NFs (1254  $\mu\text{mol g}^{-1} \text{h}^{-1}$ ),



respectively. These results illustrate that gold NPs can facilitate the generation and transfer of photoinduced electron-hole pairs and play a vital role in boosting the H<sub>2</sub> production efficiency, which could be achieved by tuning the gold content to an optimal value. Furthermore, such TiO<sub>2</sub>-Au composite NPs catalysts have significant potential for photocatalytic H<sub>2</sub> evolution and also provide a new strategy in solar energy conversion.

## Conflicts of interest

There are no conflicts to declare.

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