



Cite this: *RSC Adv.*, 2017, 7, 11827

Green solid-state synthesis and photocatalytic hydrogen production activity of anatase TiO₂ nanoplates with super heat-stability†

Jindou Hu, Yali Cao,* Kun Wang and Dianzeng Jia

A simple and environmentally friendly new method, a low-heating solid-state chemical method (LSCM), has been used to prepare uniform NH₄TiOF₃ nanoplates with a thickness of about 100 nm in the absence of surfactants. The solid–solid transformation was used to convert NH₄TiOF₃ nanoplates to TiO₂ nanoplates with an exposed (001) facet *via* a simple sintering process at different temperatures. The anatase TiO₂ nanoplates obtained in this work have super heat-stability, and can withstand high temperatures and maintain highly active anatase up to 1000 °C. The obtained compact TiO₂ nanoplates have high photocurrent density and photocatalytic hydrogen-production activity, better than as-prepared TiO₂ porous nanoplates and reported values in the literature, which is due to the exposed (001) facet and the high degree of crystallinity. This work provides a simple, efficient and environmentally friendly way to prepare TiO₂ nanoplates and nanoparticles, and also offers a new approach for preparing anatase TiO₂ with super heat-stability and high reactivity, which can be applied to H₂-production *via* photocatalytic water splitting.

Received 22nd November 2016
Accepted 5th February 2017

DOI: 10.1039/c6ra27160j

rsc.li/rsc-advances

1. Introduction

The energy crisis has attracted worldwide attention, due to the extensive use and storage limitations of fossil fuels.^{1–4} The development and application of new energy sources have become important research areas. Among various new energy sources, and offering the benefits of high energy density and environmental friendliness, hydrogen obtained from the photocatalytic splitting of water is regarded as one of the ideal candidates.^{5,6} Developing and researching semiconductor photocatalysts for hydrogen production has been recognized as a focus of research in inorganic chemistry, physical chemistry and material chemistry.⁷

To date, among various semiconductor photocatalysts that can produce hydrogen *via* the photocatalytic splitting of water, TiO₂ has the features of superior photocatalytic activity, chemical stability, low-cost and nontoxicity,^{8–13} so it is regarded as a suitable material for use in photocatalytic water splitting for hydrogen production.^{14–18} The photocatalytic performance of TiO₂ is influenced by many factors, such as the band gap, the active facet, the degree of crystallinity, the size and the morphology. In particular, for anatase TiO₂, the high activity of

the (001) facet has been verified to influence the photocatalytic activity, through both theory and experiment.^{19,20} However, the preparation of anatase TiO₂ materials with an exposed (001) facet is still a difficult task. Yang *et al.*¹⁹ used HF in a hydrothermal synthetic process to restrain the growth of the (001) facet, and obtained TiO₂ with an exposed (001) facet with high photocatalytic activity to produce H₂.^{21,22} Thereafter, a number of methods were used to prepare TiO₂ with an exposed (001) facet.^{19,23–27} These methods usually involved hydrothermal or solvothermal processes at high pressure, and extremely corrosive and toxic HF or NH₄F were used as supporting materials for the growing of crystal facets. Consequently, a facile, green and efficient method is needed for the preparation of anatase TiO₂ with an exposed (001) facet.

The low-heating solid-state chemical method (LSCM), needing only room temperature or a temperature below 100 °C, is a facile, green, simple and environmentally friendly method to synthesize nanomaterials.^{28,29} The LSCM has been used to prepare many inorganic nanomaterials, including oxides, sulfides, metals and composites, in our research group.^{30–36} In this work, the LSCM was applied to the controlled synthesis of crystal surface exposed semiconductors to further develop the simple and effective solid-state method. We designed and prepared fluoro-chemical NH₄TiOF₃ nanoplates using the LSCM without any surfactant, and then converted the NH₄TiOF₃ nanoplates to TiO₂ nanoplates. In the process of transformation, the presence of fluorine in the NH₄TiOF₃ nanoplates restrained the growth of the (001) facet. Thus, the anatase TiO₂ exposed (001) facet was obtained simply, avoiding the use of

Key Laboratory of Energy Materials Chemistry, Ministry of Education, Key Laboratory of Advanced Functional Materials, Institute of Applied Chemistry, Xinjiang University, Autonomous Region, Urumqi, Xinjiang 830046, China. E-mail: caoyali523@163.com; Fax: +86-991-8588883; Tel: +86-991-8583083

† Electronic supplementary information (ESI) available: Supplementary figure (Fig. S1–S3) and tables (Tables S1–S3). See DOI: 10.1039/c6ra27160j



toxic substances that have been reported in the literature, such as HF and NH₄F. The photocatalytic activity of the as-obtained anatase TiO₂ with an exposed (001) facet for H₂ production was investigated in detail.

2. Experimental

2.1. Synthesis

The various chemicals used in this work were purchased from commercial sources and used without further purification. In a typical experiment, 8 mmol of ammonium hexafluorotitanate was weighed and ground for 10 min in an agate mortar. Subsequently, 6 mmol of boric acid was added. After grinding at room temperature for 0.5 h, the resulting solid samples were heated in a water bath at 60 °C to perform the entire reaction. The as-prepared samples were washed with distilled water and dried under natural conditions for 12 h. The obtained samples were assigned as the precursor and denoted as T1.

The final products were fabricated through the calcination of precursor T1. The precursor T1 was annealed at 200 °C, 300 °C, 400 °C, 500 °C, 600 °C, 700 °C, 800 °C, 900 °C or 1000 °C for 2 h under an air atmosphere. The final products were denoted as T2, T3, T4, T5, T6, T7, T8, T9 and T10, respectively, and were collected for further characterization and testing. The detailed description of the obtained samples and the corresponding preparation conditions are provided in Table S1.†

2.2. Characterization

A Bruker D8 Advance diffractometer was used to measure the X-ray diffraction (XRD) patterns of the products, with Cu K α radiation ($k = 1.54184 \text{ \AA}$) under operating conditions of 40 kV and 40 mA, over the range of 10–80° at a scanning rate of 2° min⁻¹. The Scherrer formula is used to calculate the average size of the as-obtained products:

$$D = K\lambda/B \cos \theta$$

where K is usually taken as 0.89, λ is the X-ray wavelength (1.5418 Å in this study), θ is the diffraction angle for the corresponding plane and B is the full width at half maximum (FWHM). Note that XRD analysis gives the average sizes of the as-prepared products; the original sizes and morphologies of the products were confirmed using electron microscopy. A Hitachi S-4800 scanning electron microscope was used to obtain the scanning electron microscopy (SEM) image at 5 kV. A JEOL JEM-2010F transmission electron microscope was used to obtain the transmission electron microscopy images and selected area electron diffraction (SAED) patterns at 200 kV. X-ray photoelectron spectroscopy (XPS) measurements and analyses were conducted on an ESCALAB 250Xi photoelectron spectroscope. Porosity measurements and specific surface areas were obtained using a Quantachrome surface area and porosity analyzer (Autosorb-iQ2). The specific surface areas were estimated using the Brunauer–Emmett–Teller (BET) method. The UV-vis absorption spectra of the as-obtained products were obtained using a Hitachi U-3010 spectrophotometer over the range 200–800 nm. An electrochemical analyzer (CHI 660E) was

used to measure the photocurrent intensities of the products using a standard three-electrode system at room temperature. The counter electrode was a platinum wire and the reference electrode was an Ag/AgCl electrode. The working electrodes were prepared by pasting uniform slurries consisting of the products on 1.2 cm × 3.6 cm F-doped SnO₂-coated glass (FTO glass); the paste area on the FTO was controlled at 4.32 cm². The electrolyte was a 0.5 M Na₂SO₄ aqueous solution.

2.3. Photocatalytic tests

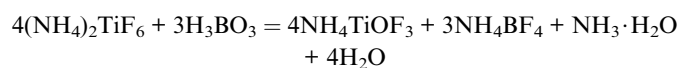
A photocatalytic H₂ preparation system (CEL-SPH2NP, CHINA) was used to measure the photocatalytic hydrogen production. Typically, 50 mg of photocatalyst was put into a glass reaction cell containing 74 mL of distilled water, 1.2 mL of H₂PtCl₆ (1.91 mmol L⁻¹) and 25 mL of methanol. The system was then placed under vacuum for 20 min to remove the O₂ in the system prior to irradiation. A 300 W Xe lamp with a different optical reflector or filter (VISREF: 350–780 nm, QD365: 365 nm, QD400: 400 nm, QD600: 600 nm) was used to provide the light source. The reaction temperature was controlled at *ca.* 6 °C by means of cooling water. The evolved H₂ was analyzed every 1 h using an online gas chromatograph with a TCD detector. The following equation was used to calculate the apparent quantum efficiency (QE) value:

$$\begin{aligned} \text{QE} &= \frac{\text{number of reacted electrons}}{\text{number of incident photons}} \times 100\% \\ &= \frac{\text{number of evolved H}_2 \text{ molecules} \times 2}{\text{number of incident photons}} \times 100\% \end{aligned}$$

3. Results and discussion

3.1. Crystal structures and compositions of the as-obtained samples

Fig. 1 shows the XRD patterns of the as-obtained products from solid-state synthesis. Typical peaks at $2\theta = 14.039^\circ$, 23.476° , and 27.449° from NH₄TiOF₃ (PDF#52-1647) are observed in T1 (Fig. 1a), and no peaks from the reactants (NH₄)₂TiF₆ and H₃BO₃ are observed, indicating that NH₄TiOF₃ is successfully prepared using the LSCM.³⁷ For the fabrication, environmentally friendly (NH₄)₂TiF₆ and H₃BO₃ were used as the reactants. The following equation is suggested to account for the reaction:



In addition, no surfactant was used as an auxiliary material during the process of synthesis, which illustrates the simple, mild, environmentally friendly and low-cost features of the LSCM and indicates the superiority and practical value of green methods in the field of chemical synthesis.

Fig. 1b shows the XRD patterns of the as-obtained products following the calcination of the precursor T1 at different temperatures. The information from Fig. 1b indicates that NH₄TiOF₃ can successfully transform to TiO₂ (PDF#04-0477) *via* a simple heating process at 300–1000 °C. Specifically, the



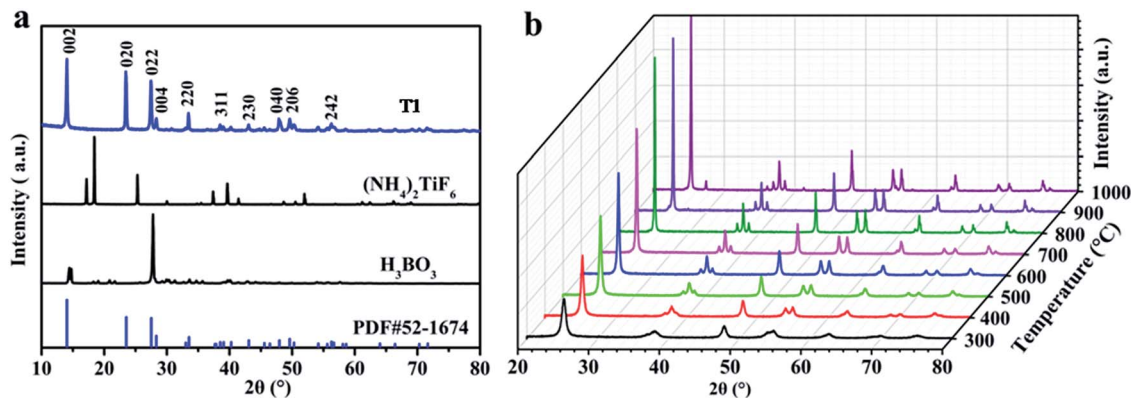


Fig. 1 XRD patterns of the samples: (a) raw materials H_3BO_3 , $(\text{NH}_4)_2\text{TiF}_6$ and the precursor T1; and (b) samples T3–T10.

precursor starts to convert to anatase TiO_2 at 200 °C (see Fig. S1†); pure anatase TiO_2 (PDF#04-0477) was obtained after treating the precursor at 300 °C for 2 h. The obtained anatase TiO_2 always exists over the annealing temperature range of 300–900 °C. When the temperature reaches 1000 °C, a typical small peak for rutile TiO_2 at $2\theta = 27.479^\circ$ is observed.³⁸ This indicates that TiO_2 obtained from annealing NH_4TiOF_3 was able to withstand high temperatures and maintained a stable anatase phase until 1000 °C; similar results are also obtained from the thermal analysis of T1 in Fig. S3,† which indicates that the TiO_2 obtained in this work has super heat-stability. With a rise in treatment temperature, the degree of crystallinity of the samples progressively improves. A high degree of crystallinity may be beneficial for H_2 production from the splitting of water in sunlight using photocatalysts.³⁹

The elemental compositions and bonding structures were characterized using XPS techniques. The sharp photoelectron peaks appearing at binding energies of 458.9 eV (Ti 2p), 531.1 eV (O 1s) and 564.3 eV (Ti 2s) in Fig. 2a indicate that sample T8 contains the elements Ti and O, and there are no obvious

signals from the elements B and F in the sample T8 (Fig. S2†). The high-resolution XPS spectra (HR-XPS) of the Ti 2p peaks are shown in Fig. 2b, and four peaks at 455.18 eV, 458.98 eV, 461.58 eV and 464.68 eV can be observed, which can be assigned to the peaks of $\text{Ti}^{3+} 2p_{3/2}$, $\text{Ti}^{4+} 2p_{3/2}$, $\text{Ti}^{3+} 2p_{1/2}$ and $\text{Ti}^{4+} 2p_{1/2}$.^{40,41} Fig. 2c shows the HRXPS spectra of O 1s; the peak at 530.6 eV in Fig. 2c can be assigned to oxygen in the form of Ti–O–Ti, and the peak at 531.8 eV can be ascribed to the Ti–O–H form.⁴² The hydroxy on the surface of TiO_2 may be beneficial for the photocatalytic performance through improving the hydrophilicity of the material. EDS techniques were used to further analyze the elemental compositions. Fig. 2d shows the EDS spectrum of sample T8; the inset is the element content of sample T8. The results from the EDS spectrum and the inset indicate that there is no F element in the samples. The high-resolution XPS spectra of F 1s and B 1s in Fig. S2† also indicate that there is no F or B element in sample T8, because F can easily be removed by heating for 30 min. The results are consistent with those obtained using XPS. The results from XPS and EDS indicate that the simple LSCM has not introduced any impurity elements into the target during the synthesis process, and extra purification after the LSCM is also unnecessary, which indicates the superiority and feasibility of the LSCM in the field of chemical synthesis.

3.2. Morphologies of the products

Fig. 3a shows that the morphology of the as-prepared precursor T1 consists of uniform platelets with a thickness of about 100 nm. The uniform morphology and the simple synthesis process indicate the simple features of the LSCM. The morphology of sample T2 is the same as for the precursor T1 (Fig. 3b). But the morphologies of samples T4 (Fig. 3c) and T6 (Fig. 3d) consist of nanoplates composed of abundant nanoparticles, and the thickness of the single-layer nanoplates is about 50 nm. When the temperature is at or below 800 °C, the morphology of the obtained sample T8 consists of smooth nanoplates with a thickness of about 50 nm (Fig. 3e). But when the temperature is above 800 °C, the samples quickly grow to bigger particles and reunification phenomena occur (Fig. 3f). The average size of the obtained samples and the increment in

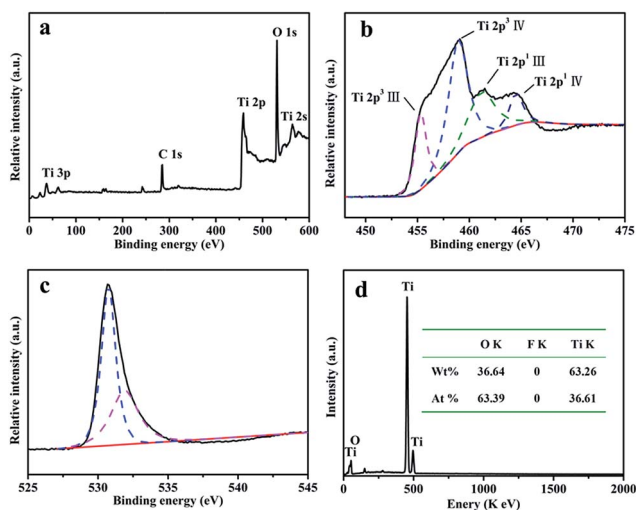


Fig. 2 (a) An XPS spectrum of sample T8; the high-resolution XPS spectra of (b) Ti 2p and (c) O 1s in sample T8; and (d) the EDS spectra and element content of sample T8.



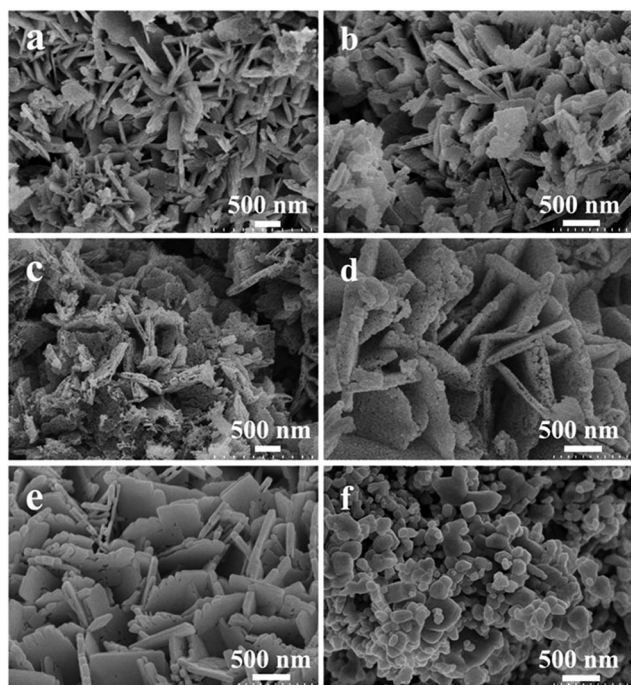


Fig. 3 SEM micrographs of the obtained samples: (a) T1; (b) T2; (c) T4; (d) T6; (e) T8; and (f) T10.

average size during the calcination process at different temperatures were further calculated using the Scherrer formula and are shown in Table S2.†

In order to explore the specific formative processes of the TiO₂ nanoplates under high temperatures, the progressive changes in the TiO₂ nanoplates during the heating process were observed and a simulated diagram is presented in Fig. 4a. Typically, F⁻ and NH₄⁺ will disappear from the NH₄TiOF₃ nanoplates during the heating process, and the precursor will start to convert to TiO₂ during calcination above 200 °C. During the heating process, the particles in the middle of the nanoplates may migrate to the surface of the nanoplates. Thus, the middle of the nanoplates will become hollow; when the treatment temperature is up to 300 °C, the morphology of the as-prepared TiO₂ will involve double-layer nanoplates consisting of particles.

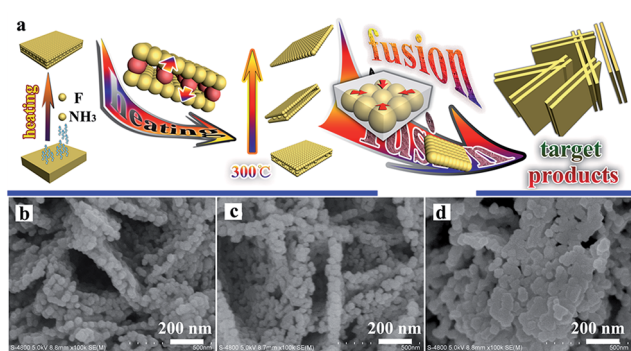


Fig. 4 A simulated diagram of the progressive change in the morphology of TiO₂ during the heating process (a), and SEM images of TiO₂ sintered at 800 °C for different times: 30 min (b), 60 min (c), and 90 min (d).

With an increase in the treatment temperature, the double-layer nanoplates will be decomposed to single-layer nanoplates or even nanoparticles with a size of about 50 nm. At 800 °C, the nanoparticles will become nanoplates with a smooth surface, with orientated fusion between nanocrystals.⁴³ This orientated fusion phenomenon is affirmed from the SEM images (Fig. 4b–d). With prolongation of the annealing time at 800 °C, the morphology of TiO₂ will change from porous nanoplates (Fig. 4c) to compact nanoplates (Fig. 4d). Finally, after 2 h at 800 °C, the nanoplates composed of nanoparticles will generate smooth nanoplates. If the temperature keeps increasing above 800 °C, the samples will quickly grow to become large particles (Fig. 3f), which is in accordance with the regularity seen in Table S2,† and the reunification phenomenon was also observed when the temperature was higher than 900 °C. During the process of growing the products, no auxiliary materials were used to control the formation of the nanoplates, which also indicates the simple and low-cost features of the solid-state method.

Fig. 5 shows micrographs of sample T8. The abundant nanoplates are displayed in the low-resolution SEM (Fig. 5a) and high-resolution SEM (Fig. 5b and c) images. The TEM image of T8 (Fig. 5d) also indicates that the morphology of sample T8 is nanoplates with a thickness of about 50 nm. The lattice rules from the SAED in Fig. 5e indicate that the as-prepared TiO₂ is monocrystalline. The interplanar spacing in Fig. 5f and g indicates that the as-prepared samples are anatase TiO₂ with an exposed (001) facet.^{44–46} Zhou *et al.* also reported that heating at 700–900 °C will lead to TiO₂ with an exposed (001) facet.⁴³ The surface energy and reactivity of the (001) facet of TiO₂ is relative higher than for the other ones.²³ Therefore, the exposed (001) facet may be beneficial for the photocatalytic performance. The results from electron microscope observation indicate that TiO₂ with a high-activity (001) facet can also be synthesized using the simple solid-state method without any additional surfactant to control the process of growing the products.

3.3. UV-vis absorption spectra

The absorbance properties of the as-obtained products are displayed in Fig. 6a. The absorption intensities of the samples

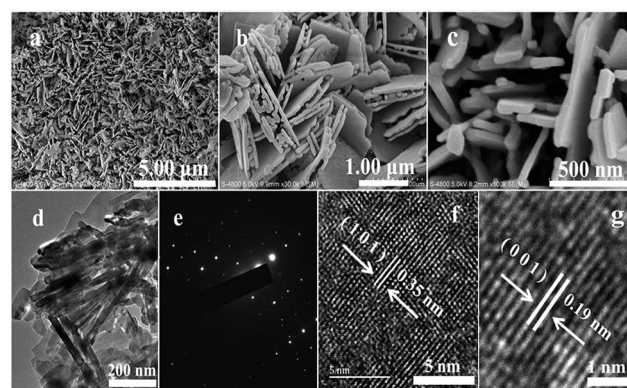


Fig. 5 Micrographs of sample T8: (a) SEM; (b and c) high-resolution SEM images; (d) TEM; (e) SAED; and (f and g) HRTEM.



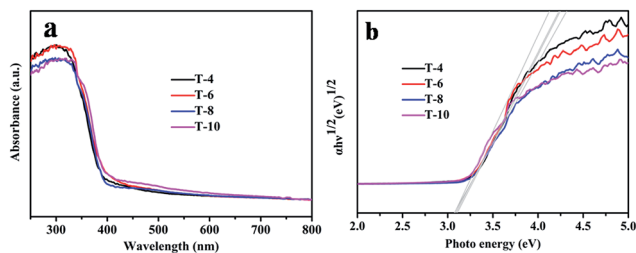


Fig. 6 (a) UV-vis absorption spectra of samples T4, T6, T8 and T10; and (b) Tauc plots of samples T4, T6, T8 and T10.

obtained at different temperatures all start to increase at 400 nm (Fig. 6a), which is similar to the intrinsic band gap of anatase TiO₂ (about 3.2 eV (ref. 47)). A similar band gap of about 3.2 eV for all the samples is also shown in Fig. 6b. This indicates that the absorbance properties and band gaps of the samples have nothing to do with the sintering temperature. There are no absorption peaks caused by N or B elements observed in Fig. 6a, which also indicates that pure TiO₂ can be obtained using the solid-state method.

3.4. Photocatalytic activity and photocurrent density

Photocatalytic H₂ production from water splitting was used to evaluate the photocatalytic activities of all samples, and the results are shown in Fig. 7. The information in Fig. 7a indicates that the rate of H₂ generation firstly increases, and then decreases, with a rise in the sintering temperature, and the rate using sample T8 is 13 mmol h⁻¹ g⁻¹, which is the highest rate from all the as-prepared photocatalysts in this work and better than reported values in the literature.^{48,49} But the rate of H₂ generation decreases when the sintering temperature is above

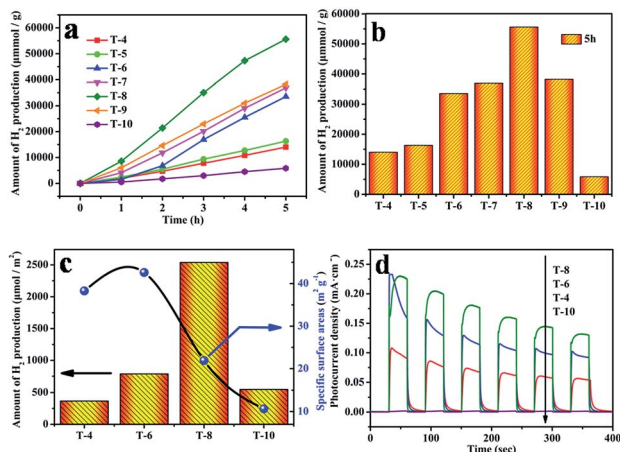


Fig. 7 (a) Photocatalytic hydrogen evolution using the obtained TiO₂; (b) the amount of photocatalytic hydrogen production after 5 h of irradiation; (c) the amount of photocatalytic hydrogen production relative to the specific surface area – the histogram is the amount of photocatalytic hydrogen production for each specific surface area after 5 h of irradiation and the line is the specific surface area of the samples T4, T6, T8, and T10; and (d) the photocurrent density for samples T4, T6, T8, and T10 under UV irradiation at 0.36 V vs. Ag/AgCl.

800 °C. In comparison, the amount of H₂ production after 5 h is shown in Fig. 7b; it can clearly be seen that the amount of H₂ generation using sample T8 is 4 times that of sample T4, and 9.5 times that of sample T10. The corresponding quantum efficiency of sample T8 under 365 nm, 400 nm and 600 nm light is shown in Fig. 8. The reason behind the high photocatalytic activity may be the comprehensive impact of the size of the photocatalyst (see Table S2†), the degree of crystallisation, the specific surface area (see Table S3†) and the exposed (001) facet. The amount of photocatalytic hydrogen production relative to the specific surface area is shown in Fig. 7c. This demonstrates that the photocatalytic performance of sample T8 is also excellent without the effect of specific surface area. Thus, the high photocatalytic activity of T8 may be ascribed to the exposed (001) facet and the high degree of crystallinity.

The semiconductor's ability to generate and transfer photo-generated charge carriers will be indirectly reflected by the value of the photocurrent under irradiation.⁵⁰ In order to confirm the active photocatalytic performance of the photocatalysts obtained using the simple solid-state method, the as-obtained photocatalysts were coated on FTO electrodes to evaluate the photocurrent, and the results are displayed in Fig. 7d. Prompt generation of photocurrent was observed when the electrodes were illuminated. With an increase in the annealing temperature, the photocurrent density of the as-prepared TiO₂ became higher. In particular, sample T8 shows the highest photocurrent density. It reached a value as high as 0.22 mA cm⁻², which is twice that of sample T4, and 20 times that of sample T10. The photocurrent density decreases quickly when the sintering temperature is above 800 °C. Note that sample T10 shows a low photocurrent density, which may be caused by a larger size of crystal and a rapid recombination of the photo-generated electrons and holes in TiO₂. The highest photocurrent density from sample T8 may be a consequence of the exposed (001) facet and the more perfect crystalline nature of the sample. Zhao *et al.* also reported that with an exposed (001) facet the slow recombination rate of photo-induced charges will lead to more favorable photocatalytic activity for TiO₂.^{23,51} The more perfect crystalline nature would reduce obstacles to electronic transmission and allow more electrons to reach the surface of the photocatalyst to assist the catalytic reactions.

The QE values for the photocatalyst T8 were also calculated, using the amount of H₂ production under light of different wavelengths (365 nm, 400 nm and 600 nm). The QE is 0.93%,

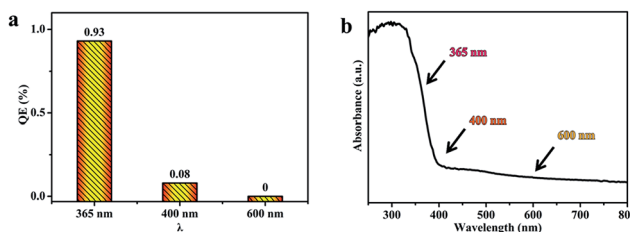


Fig. 8 (a) The quantum efficiency of sample T8 under 365, 400 and 600 nm light; and (b) the UV-vis absorption spectrum of sample T8.



0.08% and 0% under 365 nm, 400 nm and 600 nm, respectively. The results in Fig. 8a show that the QE decreases with an increase in the wavelength, which is consistent with the absorption intensity in the UV-visible absorption spectra (Fig. 8b).

4. Conclusions

In summary, we have successfully prepared uniform NH_4TiOF_3 nanoplates with a thickness of about 100 nm using a green, solvent-free LSCM in the absence of a template. The simple solid–solid transformation was used to convert NH_4TiOF_3 to TiO_2 nanoplates with an exposed (001) facet, with a single layer thickness of about 50 nm. The TiO_2 obtained in this work has good heat-stability, and is able to withstand high temperatures and maintain a stable anatase phase up to 1000 °C. The anatase TiO_2 nanoplates exhibit a high rate of H_2 generation, due to the exposed (001) facet and the high crystallinity. This work provides an efficient and environmentally friendly method to prepare TiO_2 nanoplates with good heat-stability and high photocatalytic activity for hydrogen production.

Acknowledgements

This work was financially supported by the National Natural Science Foundation of China (No. 21361024, 21471127 and U1503392) and the Natural Science Foundation of Xinjiang Province (No. 2014711004 and 2014211A013).

Notes and references

- S. Landsmann, Y. Surace, M. Trottmann, S. Dilger, A. Weidenkaff and S. Pokrant, *ACS Appl. Mater. Interfaces*, 2016, **8**, 12149–12157.
- W. Zhou, Z. Yin, Y. Du, X. Huang, Z. Zeng, Z. Fan, H. Liu, J. Wang and H. Zhang, *Small*, 2013, **9**, 140–147.
- T. Montini, V. Gombac, L. Sordelli, J. J. Delgado, X. Chen, G. Adami and P. Fornasiero, *ChemCatChem*, 2011, **3**, 574–577.
- X. An, Y. Wen, A. Almuji, D. Cheng, J. Li, X. Jia, J. Zou and Y. Ni, *RSC Adv.*, 2016, **6**, 89457–89466.
- X. Zou and Y. Zhang, *Chem. Soc. Rev.*, 2015, **44**, 5148–5180.
- H. X. Han and C. Li, *Natl. Sci. Rev.*, 2015, **2**, 143–145.
- D. Lang, T. Shen and Q. Xiang, *ChemCatChem*, 2015, **7**, 943–951.
- J. Yu, T. Ma and S. Liu, *Phys. Chem. Chem. Phys.*, 2011, **13**, 3491–3501.
- B. Kenens, M. Chamtour, R. Aubert, K. Miyakawa, Y. Hayasaka, H. Naiki, H. Watanabe, T. Inose, Y. Fujita, G. Lu, A. Masuhara and H. Uji-i, *RSC Adv.*, 2016, **6**, 97464–97468.
- S. A. Bakar and C. Ribeiro, *RSC Adv.*, 2016, **6**, 89274–89287.
- U. Mandi, N. Salam, S. K. Kundu, A. Bhaumik and S. M. Islam, *RSC Adv.*, 2016, **6**, 73440–73449.
- G. B. Soares, R. A. P. Ribeiro, S. R. de Lazaro and C. Ribeiro, *RSC Adv.*, 2016, **6**, 89687–89698.
- H. Zhang, Q. Lin, S. Ning, Y. Zhou, H. Lin, J. Long, Z. Zhang and X. Wang, *RSC Adv.*, 2016, **6**, 96809–96815.
- J. Zhang, Q. Xu, Z. Feng, M. Li and C. Li, *Angew. Chem., Int. Ed.*, 2008, **47**, 1766–1769.
- F. Cao, J. Xiong, F. Wu, Q. Liu, Z. Shi, Y. Yu, X. Wang and L. Li, *ACS Appl. Mater. Interfaces*, 2016, **8**, 12239–12245.
- J. M. Macak, M. Zlamal, J. Krysa and P. Schmuki, *Small*, 2007, **3**, 300–304.
- I. Paramasivam, H. Jha, N. Liu and P. Schmuki, *Small*, 2012, **8**, 3073–3103.
- Y. Tang, R. Hao, Y. Fu, Y. Jiang, X. Zhang, Q. Pan and B. Jiang, *RSC Adv.*, 2016, **6**, 96803–96808.
- H. G. Yang, C. H. Sun, S. Z. Qiao, J. Zou, G. Liu, S. C. Smith, H. M. Cheng and G. Q. Lu, *Nature*, 2008, **453**, 638–641.
- M. Maisano, M. V. Dozzi, M. Coduri, L. Artiglia, G. Granozzi and E. Selli, *ACS Appl. Mater. Interfaces*, 2016, **8**, 9745–9754.
- J. G. Yu, L. F. Qi and M. Jaroniec, *J. Phys. Chem. C*, 2010, **114**, 13118–13125.
- G. Liu, H. G. Yang, X. Wang, L. Cheng, J. Pan, G. Q. Lu and H. M. Cheng, *J. Am. Chem. Soc.*, 2009, **131**, 12868–12869.
- Y. Zhao, Q. Zhao, X. Li, Y. Hou, X. Zou, J. Wang, T. Jiang and T. Xie, *Mater. Lett.*, 2012, **66**, 308–310.
- Q. Xiang and J. Yu, *Chin. J. Catal.*, 2011, **32**, 525–531.
- C. X. Lei, X. L. Jiang, X. Huang, X. Liu, D. Q. Zeng, Y. T. Ma, L. S. Wang and D. L. Peng, *Appl. Surf. Sci.*, 2015, **359**, 860–867.
- H. Li, T. Li, H. Liu, B. Huang and Q. Zhang, *J. Alloys Compd.*, 2016, **657**, 1–7.
- S. Liu, J. Yu and M. Jaroniec, *J. Am. Chem. Soc.*, 2010, **132**, 11914–11916.
- H. Yu, Q. F. Xu, Z. R. Sun, S. J. Ji, J. X. Chen, Q. Liu, J. P. Lang and K. Tatsumi, *Chem. Commun.*, 2001, 2614–2615.
- C. Zhang, Y. Song, X. Wang, F. E. Kühn, Y. Wang, Y. Xu and X. Xin, *J. Mater. Chem.*, 2003, **13**, 571–579.
- L. Wang, Y. D. Huang, R. R. Jiang and D. Z. Jia, *J. Electrochem. Soc.*, 2007, **154**, 1015–1019.
- J. X. Guo, L. Liu, G. F. Liu, D. Z. Jia and X. L. Xie, *Org. Lett.*, 2007, **9**, 3989–3992.
- Y. L. Cao, P. F. Hu, W. Y. Pan, Y. D. Huang and D. Z. Jia, *Sens. Actuators, B*, 2008, **134**, 462–466.
- Y. L. Cao, W. Y. Pan, Y. Zong and D. Z. Jia, *Sens. Actuators, B*, 2009, **138**, 480–484.
- Y. L. Qi, Y. D. Huang, D. Z. Jia, S. J. Bao and Z. P. Guo, *Electrochim. Acta*, 2009, **54**, 4772–4776.
- R. Y. Wang, D. Z. Jia, L. Zhang, L. Liu, Z. P. Guo, B. Q. Li and J. X. Wang, *Adv. Funct. Mater.*, 2006, **16**, 687–692.
- Y. Z. Li, Y. L. Cao and D. Z. Jia, *J. Mater. Chem. A*, 2014, **2**, 3761–3765.
- M. Hojamberdiev, G. Zhu, P. Sujaridworakun, S. Jinawath, P. Liu and J. P. Zhou, *Powder Technol.*, 2012, **218**, 140–148.
- B. Liu, X. Li, Q. Zhao, J. Ke, M. Tadé and S. Liu, *Appl. Catal., B*, 2016, **185**, 1–10.
- L. Yan, X. He, Y. Wang, J. Li and D. Wang, *J. Mater. Sci.: Mater. Electron.*, 2016, **27**, 4068–4073.
- X. Liu, S. Gao, H. Xu, Z. Lou, W. Wang, B. Huang and Y. Dai, *Nanoscale*, 2013, **5**, 1870–1875.
- D. Kumar, M. S. Chen and D. W. Goodman, *Thin Solid Films*, 2006, **515**, 1475–1479.



- 42 W. Zhao, Z. Ai, J. Dai and M. Zhang, *PLoS One*, 2014, **9**, 103671–103680.
- 43 L. Zhou, J. Chen, C. Ji, L. Zhou and P. O'Brien, *CrystEngComm*, 2013, **15**, 5012–5015.
- 44 J. Q. Yan, H. Wu, H. Chen, Y. X. Zhang, F. X. Zhang and S. Z. F. Liu, *Appl. Catal., B*, 2016, **191**, 130–137.
- 45 S. Yu, Y. Q. Zhong, B. Q. Yu, S. Y. Cai, L. Z. Wu and Y. Zhou, *Phys. Chem. Chem. Phys.*, 2016, **18**, 20338–20344.
- 46 W. Lin, H. Zheng, P. Zhang and T. Xu, *Appl. Catal., A*, 2016, **521**, 75–82.
- 47 M. Miyauchi, A. Ikezawa, H. Tobimatsu, H. Irie and K. Hashimoto, *Phys. Chem. Chem. Phys.*, 2004, **6**, 865–870.
- 48 L. Li, L. Yu, Z. Lin and G. Yang, *ACS Appl. Mater. Interfaces*, 2016, **8**, 8536–8545.
- 49 G. L. Chiarello, E. Selli and L. Forni, *Appl. Catal., B*, 2008, **84**, 332–339.
- 50 Z. A. Huang, Q. Sun, K. L. Lv, Z. H. Zhang, M. Li and B. Li, *Appl. Catal., B*, 2015, **164**, 420–427.
- 51 K. Takanahe and K. Domen, *ChemCatChem*, 2012, **4**, 1485–1497.

