Synthesis of High-Reactive Facets Dominated Anatase TiO₂

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Received (in XXX, XXX) Xth XXXXXXXXXX 200X, Accepted Xth XXXXXXXXXX 200X

First published on the web Xth XXXXXXXXXX 200X

DOI: 10.1039/b000000x

As an important metal oxide, anatase titanium dioxide has been widely investigated because of its many promising properties in catalysis and photocatalysis. The properties of anatase TiO₂ crystals are largely determined by external surfaces exposed. Many efforts have been made to improve the percentage of high-reactive facets like {001} facets of anatase TiO₂ to enhance its catalytic properties. This review reports the recent progress in designing and fabricating high-reactive facets dominated anatase TiO₂ via various strategies including traditional vapor phase epitaxial processes, hydrothermal/solvothermal methods, nonhydrolytic alcoholysis methods and high temperature gas phase reactions. Furthermore, focusing on (001) surface, the review also covers advances in the theoretic simulations of various high-reactive facets of anatase TiO₂ crystals. Finally, we offer a summary and some perspectives on the challenges and new directions for future research in this emerging frontier.

1. Introduction

Metallic and semiconducting nanocrystals with tailored facets have always attracted intense interests in the last decade due to their many intrinsic shape-dependent properties, such as water splitting for hydrogen, solar cells, CO₂ reduction, gas sensors etc.¹⁻²¹ As one of the commonly available and most studied metal oxides, titanium dioxide (TiO₂) has been widely used as a pigment and in sunscreens, paints, ointments, toothpaste, and catalysis.⁹ we have witnessed an exponential growth of research activities since Fujishima and Honda discovered the phenomenon of photocatalytic splitting of water on a TiO₂ electrode under ultraviolet (UV) light in 1972.²² Many promising applications have been developed

including photovoltaic cells, photo-/electrochromics, photocatalysis, photonic crystals, and smart surface coatings.^{5,23,24} Synthesis of highly reactive TiO₂ is always desired to improve its properties and extending its applications.

TiO₂ has three major crystal structures: rutile, anatase and brookite. Of the three phases of TiO₂, anatase is the most widely investigated polymorphs because of its high activity in catalysis and photocatalysis. The average surface energies of anatase TiO₂ (0.90 J/m² for {001}, 0.53 J/m² for {100}, 0.44 J/m² for {101}) have been demonstrated extensively.^{2,25} Unfortunately, {101} surfaces, normally the majority of the external surface of anatase TiO₂ (more than 94% percent, according to the Wulff construction), are thermodynamically stable with a low surface energy (0.44 J/m²), which determines its low reactivity.²⁶ To improve its reactivity, the preparation of shape-controlled anatase TiO₂ crystals with specific reactive facets exposed is greatly desired, and the minority surfaces such as {001} ones with a higher surface energy (0.90 J/m²) attract extensive interest.

Well-defined anatase single crystals with a high percentage

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of {001} facets had not been synthesized until we successfully prepared them with 47% of the highly reactive {001} facets by using hydrofluoric acid (HF) as a capping agent under hydrothermal conditions.²⁷ Following this breakthrough, extensive efforts have been made in the preparation, modification, and application of TiO₂ with {001} facets. The percentage of {001} facets has been improved steadily and reached as high as ~ 100% currently.²⁸⁻³¹ Meanwhile, the thickness of 2-dimensional (2D) anatase TiO₂ nanosheets has reduced to ~ 1.6 nm along [001] crystallographic direction, which means that the nanosheets have only two layers of anatase TiO₂ crystal units.²⁹ By introducing a newly developed gas phase process, anatase TiO₂ crystals with a large percentage of {001} facets were also fabricated.³² Concerning the modification of anatase TiO₂ crystals with {001} facets, in situ nitrogen doped anatase TiO₂ crystals were achieved by employing TiN as the titanium source.³³ Aside from {001} facets, other high-reactive facets such as {110} facets were observed recently.³⁴ Considering so many

breakthroughs reported and the fast advances of this field in the past few years, we believe that a comprehensive review of this important functional material will further promote the research in this field. In this feature article, we will focus on the recent synthesis advances, chiefly discussing the various construction methods and the diverse range of raw materials used to prepare anatase TiO₂ with reactive facets, and will provide some invigorating perspectives as to future directions.

2. Theoretic predictions on the morphology of anatase TiO₂

Inorganic single crystals with highly reactive surfaces have long been studied, owing to their importance in scientific research and technological applications.³⁵⁻³⁷ The effect of surface chemistry has been thought to be crucial for the equilibrium morphology of inorganic single crystals and thereby is critical to the synthesis of single crystals with high reactivity.^{2,26,35-46} Anatase TiO₂, compared to the other two crystal polymorphs, has been proved to be more reactive in some cases, especially for photocatalysis and catalysis.^{2,24,47} It was found that the most abundant {101} facets are not very reactive, and experimental evidence shows that this surface has few defects after annealing in ultrahigh-vacuum conditions (UHV) and thus {101} facets of anatase TiO₂ are more difficult to reduce in comparison with the widely investigated {110} facets of rutile TiO₂.^{47,51} Also, many studies have proven that the minor {001} facets with high density of surface undercoordinated Ti atoms exhibit higher reactivity in contrast with {101} facets.^{26,35,44,52} So anatase TiO₂ crystals with a large percentage of {001} facets are very desirable. Thus, as an important environmental and energy material, uniform and highly pure anatase TiO₂ single crystals with controllable crystallographic {001} facets is vital because of its good performance. For this purpose, efforts have been taken using various adsorbate atoms to change the relative stabilities of different crystal facets and thus alter the

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ally and internationally including the Orica Award, RK Murphy Medal, Le Fevre Prize, ExxonMobil Award, IUMRS Young Scientist Award, Top 100 Most Influential Engineers in Australia (2004, 2010).

equilibrium shapes of anatase TiO₂.^{9,46,48-50}

In an early report, Oliver et al. presented some interesting polymorphs of rutile and anatase TiO₂ with the evaluation of surface energies.⁵³ {001} facets of anatase TiO₂ with five-coordinate Ti atoms were found to be adjustable by changing the surface adsorbates. Considering the effect of hydration on equilibrium morphologies, Arrouvel et al. deduced the equilibrium morphologies of anatase TiO₂ as a function of the working conditions including temperature and partial pressure of water.⁵² Their idea is that the capability of water adsorptions on (101) or (001) planes varies with the changes of pressure and temperature, thereby resulting in variations of area proportions of the two surfaces.

Barnard et al. established a shape dependent thermodynamic model for a nanoparticle X (X represents anatase or rutile) on the basis of Gibbs free energy G_x^o in which the contributions of particle bulk and surface were taken into account.⁴⁶

$$G_x^o = G_x^{bulk} + G_x^{surface}$$

Then they further introduced the surface energy γ_i for each crystallographic surface i . Hence,

$$G_x^o = \Delta_f G_x^o + \frac{M}{\rho_x} (1-e) [q \sum_i f_i \gamma_{xi}]$$

where e is the volume dilation induced by surface tension. The introduction of the effect of surface tension could revise the deviation generated from Wulff construction and the shape of nanoparticles was optimized as a function of size, or other parameters such as temperature. Degree of truncation is another meaningful concept proposed, which can be defined as the ratio of the side of "truncation" facet (B, see Fig.1a) with respect to the side of bipyramid (A). A plot of the optimized B/A ratio of anatase nanocrystals according to the variation of surface chemistry was subsequently provided (Fig. 1b). The most impressive part of this work is the final shapes

Fig. 1 (a) An anatase tetragonal {101} bipyramid is shown in outline, and the side lengths labelled **A** and **B** are used to define the degree of truncation. (b) Plot of the optimized ratio **B/A** for anatase nanocrystals with each of the various surface chemistries, with a side length **A** = 2 to 100 nm. The facet edges **A** and **B** are defined in (a).⁴⁶

Fig. 2 Morphology predicted for anatase TiO₂ with (a) hydrogenated surfaces, (b) with hydrogen-rich surface adsorbates, (c) hydrated surfaces, (d) hydrogen-poor adsorbates, and (e) oxygenated surfaces.⁴⁶

Fig. 3 (a) Calculated energies of the {001} and {101} surfaces surrounded by X atoms. (b) Plots of the optimized value of **B/A** and percentage of {001} facets for anatase single crystals with various adsorbate atoms X.²⁷

of anatase TiO₂ crystals with different surface chemistry (Fig. 2). From Fig. 2, it can be concluded that surface adsorbates containing a large fraction of hydrogen played a major role in stabilizing anatase nanoparticles.

Thus, inorganic anions acting as important adsorbed species, have attracted great research attention to investigate their role in the shape control of anatase TiO₂.⁵⁰ In order to obtain the {001} facets dominated anatase TiO₂, it was both predicted by first-principle calculations and confirmed through experimental results that termination of F atoms can yield the lowest surface energy and make {001} facets more stable, compared to {101} facets.²⁷

The most stable crystal facets of anatase TiO₂ among oxygenated surfaces are {100} facets, while under clean and hydrogenated conditions the {101} facets are the most stable.^{2, 26, 46} However, the high surface energies (γ) generated by H- or O-terminated anatase surfaces give rise to the barrier to the formation of large single-crystal TiO₂. High values of γ are the main reason for high bonding energies (D_0) of H-H (436.0

Fig. 4 (a) Morphology evolutions of anatase TiO_2 crystals. Representation of atomic schematic structures of {101}, {001}, {100}, {103}, {105} and {107} surfaces, respectively. O in red and Ti in grey.

kJ/mol) and O-O (498.4 kJ/mol).⁵⁴ Therefore, to find a low D_0 element with strengthened bonding to Ti was considered to be an effective solution for stabilizing the faceted surfaces. Fortunately, F is such an element, as $D_0^{F-F} = 158.8$ kJ/mol and $D_0^{F-Ti} = 569.0$ kJ/mol.^{54, 55} To further explore the effects of various adsorbate atoms, a systematic investigation of 12 non-metallic atoms X (X = H, B, C, N, O, F, Si, P, S, Cl, Br, I) was carried out using first-principle calculations. The calculated γ values for different adsorbates are illustrated in Fig. 3a, from which two conclusions can be drawn: (i) among the 12 non-metal-terminated surfaces and the clean surfaces, F-terminated anatase surfaces have the lowest γ for both {001} and {101}; and (ii) for F-terminated anatase surfaces, {001} surfaces are preferential and more stable, compared to {101} surfaces. These results indicate that a high percentage of anatase {001} facets may be achievable if their surfaces are terminated by F atoms. Furthermore, based on the shape dependent thermodynamic model proposed by Barnard et al.,⁵⁶ the optimized ratio B/A and the percentage of {001} facets can be predicted if γ is known. As shown in Fig. 3b, the F-terminated surfaces have the highest degree of truncation and, in turn, the F-terminated surfaces of anatase TiO_2 should

be dominated by {001} facets.

We have verified these theoretical predictions experimentally, which also include a control experiment in the absence of hydrofluoric acid (see Fig. 5). As to the stabilization mechanism, this can be explained by the balance of O-O/O-F repulsions and Ti-O/Ti-F attractions, which could stabilize Ti and O atoms on the surfaces. As a summary, some possible morphologies as well as corresponding surface atomic structures of anatase TiO_2 crystals are illustrated in Fig. 4, where some of morphologies have been confirmed experimentally, while the others are expected to be realized in the future.

3. Synthetic strategies of {001} facets dominated anatase TiO_2

The different facets of inorganic nanocrystals are largely determined by different surface energies and the surfaces with high reactivity usually diminish rapidly during the growth process as a result of the minimization of total surface energy. Surfaces adsorption of capping agents resulting in different surface energy has been expected to be an effective approach in fabricating faceted nanocrystals, the shape of which are

hence controlled. The choice of capping agent remains empirical due to the lack of experimental and theoretical evidence on the adsorption energy between capping agents (organic surfactants or non-metallic ions) and nanocrystal surfaces. But undoubtedly, the fact that some organic surfactants and non-metallic ions adsorb selectively on specific facets widely exists.^{27,53} Selective adsorption of capping agents reduces the surface energy of materials with more active sites, and thus retards or inhibits the crystal growth along the corresponding direction, which results in the preservation of high-reactive or high-index facets. As to synthesizing anatase TiO₂ with highly reactive {001} facets, the high density of undercoordinated Ti atoms on the surface and large Ti-O-Ti bond angles make some organic surfactants or non-metallic ions adsorb on {001} facets preferentially, leading to truncated bipyramid morphology.

Various methods have been developed to synthesize anatase TiO₂ crystals dominated with {001} facets. Herein, the representative synthetic processes by traditional vapor phase epitaxial processes with SrTiO₃ and LaAlO₃ as substrates, hydrothermal methods, nonhydrolytic approaches, and high temperature gas phase reactions to prepare anatase TiO₂ crystals are reviewed.

3.1 Traditional vapor phase epitaxial processes by using SrTiO₃ and LaAlO₃

Epitaxial growth is defined as oriented overgrowth of a crystalline phase on the surface of a substrate that is also crystalline. The vapor processes such as chemical vapor deposition (CVD), molecular beam epitaxy (MBE), and laser ablation are usually involved in epitaxial growth of (001) anatase films. Chen et al. reported the fabrication of epitaxial anatase films grown on SrTiO₃ (001) substrate by metal-organic chemical vapor deposition (MOCVD), which is significant due to the different crystallographic structures and different lattice systems.⁵⁷ The reason for selecting SrTiO₃ (001) as a substrate was attributed to the good lattice matching between SrTiO₃ (001) (0.3905 nm × 0.3905 nm) and anatase (001) (0.3785 nm × 0.3785 nm).⁵⁸ And it was because of the lack of a good structural match between (001) SrTiO₃ and all of the major low-index planes of rutile TiO₂ that the heteroepitaxial growth of anatase was achieved. Furthermore, the result demonstrated the important role played by the substrate structure on the heteroepitaxial deposition of films and their orientation. Using the same method, Herman et al. prepared the anatase thin film and investigated their electronic and structural properties by X-ray photoelectron spectroscopy (XPS), low energy electron diffraction (LEED), and X-ray photoelectron diffraction (XPD).⁵⁹ The results showed that these thin films had good short-range order on the surface and exhibited high stability. Another investigation using oxygen plasma assisted MBE to grow (001) anatase films on SrTiO₃ (001) substrate revealed that Sr was segregated to the surface during growth with substrate temperature above 823 K. The surface had a (1 × 4) reconstruction during the growth and this reconstruction remained after cooling to room temperature, confirmed by the high-energy electron diffraction patterns.⁶⁰ Recently, with the

assistance of ultrahigh vacuum scanning electron microscopy, Marshall and Castell studied the temperature-dependent shape transitions of epitaxial islands of (001) anatase TiO₂ on SrTiO₃ (001) substrate.⁶¹

In order to obtain anatase films with higher quality, (001) face of LaAlO₃ substrate was also employed because of its smaller lattice mismatch (-0.2%) compared to that of SrTiO₃ (-3.1%). The as-obtained films exhibited higher crystallinity as expected. Kennedy et al. prepared (001) oriented anatase thin films on both (001) LaAlO₃ and (001) SrTiO₃ substrates by pulsed laser deposition.⁶² The thickness dependence of the lattice parameter and the crystalline quality of the product were investigated. Considering a wider band gap energy of LaAlO₃ substrate (~5 eV), Murakami et al. elucidated the possibility of evaluating the optical properties of anatase films and indicated that high-quality anatase thin film was expected to be a potential material for optical devices.⁶³ Phase-pure (001) oriented anatase films on (001) LaAlO₃ substrate were also prepared using water vapor as the oxidizing species by reactive agent sputter deposition.⁶⁴ Cheung et al. studied the visible-light driven photocatalytic properties of N-doped TiO₂ (001) anatase prepared by plasma-assisted MBE on LaAlO₃ (001) substrate, and a substantial red shift in the optical band gap was observed.⁶⁵

3.2 Hydrothermal/Solvothermal processes

Hydrothermal method, one of the traditional methods for crystal growth of semiconductors, is frequently used to prepare TiO₂ exposed with {001} facets. After anatase TiO₂ crystals with 47% of {001} facets were successfully synthesized (Fig. 5),²⁷ the concept of synergistic functionality was proposed in which 2-propanol was employed as a synergistic capping agent and reaction medium together with HF to synthesize high-quality anatase TiO₂ single-crystal nanosheets with 64% of the {001} facets.⁶⁶ As protecting agent, 2-propanol heterolytically dissociated to form alkoxy groups ((CH₃)₂CHO⁻) under acidic conditions and then the groups bound to coordinatively unsaturated Ti⁴⁺ cations on {001} and {101} facets. Selective adsorption of 2-propanol

Fig. 5 Typical SEM image of anatase TiO₂ nanosheets exposed with 47% {001} facets.²⁷

Fig. 6 Typical SEM/TEM images (a-c) of the anatase TiO₂ nanosheets synthesized with 2-propanol as a synergistic capping agent.⁶⁶

was favored on {001} facets with higher density of 5-fold Ti. As a result, the growth of anatase TiO₂ single crystals along [001] direction was retarded. The as-synthesized product exhibited superior photoreactivity 5 times greater than P25, a benchmarking material. Fig. 6 shows the typical TEM/SEM images of the as-synthesized TiO₂ single crystalline nanosheets.

In previous studies, large particle size of the synthesized products resulted in a relatively small specific surface area, limiting the improvement in photocatalytic activities. Han et al. improved the percentage of {001} facets greatly by using tetrabutyl titanate, Ti(OBu)₄, as a precursor and 47% hydrofluoric acid solution as a capping agent.^{28,67,68} Both high percentage (89%) of {001} facets and relatively small crystal size (side length ~40 nm, thickness ~6 nm) of anatase TiO₂ were beneficial for the photocatalytic activity. Then the thinnest anatase TiO₂ nanosheets with dominant {001} facets (~1.6 nm) was prepared by Yang et al. recently through a modified hydrothermal route (Fig. 7),²⁹ and the efficiency for H₂ evolution was achieved as high as 7381 μmol h⁻¹ g⁻¹,

Fig. 7 (a), (b) High-resolution TEM images of ultra-thin anatase TiO₂ nanosheets and (c) the corresponding structure model.²⁹

under UV-vis light irradiation. This 2D anatase TiO₂ nanosheets is novel because it only contains 2 layers of crystal units along [001] crystallographic direction. Another study which realized a shape controlled synthesis of anatase TiO₂ with high-reactive facets employ both oleic acid (OA) and oleylamine (OM) as capping agents in the presence of water vapor.⁶⁹ By increasing the amount of oleic acid, anatase TiO₂ with {001} facets could be preserved (Fig. 8a). It was demonstrated that OM selectively adsorbed on {101} facets while OA on {001} facets, resulting in anatase TiO₂ crystals exposed with {001} facets in the presence of a relatively large amount of OA. Dai et al. introduced electrospinning technique into the synthesis process for the first time to generate {001} facets exposed anatase TiO₂.⁷⁰ Before hydrothermal treatment was carried out, nanofibers containing Ti sources were prepared by electrospinning process, in which small and uniform amorphous TiO₂ particles were prepared with the help of poly(vinyl pyrrolidone) (PVP) matrix. The researchers speculated that titanium tetraisopropoxide hydrolyzed with trace moist air in the electrospinning process. The small diameter of the as-spun nanofibers benefited for the diffusion of moist air into the interior of these fibers. The relative slow hydrolysis rate and the PVP matrix resulted in small amorphous TiO₂ particles which was beneficial for the conversion from amorphous TiO₂ to anatase TiO₂ nanocrystals through hydrothermal treatment. The as-obtained nanocrystals exhibited a truncated tetragonal bipyramidal shape with 9.6% of {001} facets exposed

Fig. 8 (a) Truncated rhombic shape anatase TiO₂ nanocrystals by Dinh et al.⁶⁹ (b) Truncated tetragonal bipyramid by Dai et al. using electrospinning technique.⁷⁰

(Fig. 8b). Recently, Wang et al. synthesized anatase TiO₂ films with oriented {001} facets from the pretreated Ti foils.⁷¹ By using a different titanium source, Liu et al. reported a new method by the hydrothermal synthesis of titanium sulfate (Ti(SO₄)₂) and hydrofluoric acid (HF) to produce nanosized TiO₂.⁷² Compared to the micron-sized anatase with 72% of {001} facets synthesized in a similar way, these new TiO₂ nanoparticles exhibited significantly greater level of photocatalytic activity both for generating ·OH radicals and for watersplitting. In situ nitrogen doped anatase TiO₂ crystals with dominant {001} facets were also successfully synthesized for the first time (Fig. 9),³³ which demonstrated excellent photooxidation and photoreduction activity due to the significantly enhanced visible light absorption ability. One-pot synthesis of highly crystalline TiO₂ with oxygen deficiency without post-treatment under extreme conditions was realized recently.⁷³ The researchers attributed the formation of oxygen-deficient anatase TiO₂ sheets to the synergistic functions of HF and H₂ from acid hydrolysis process of TiB₂. {001} facets were stabilized by surface Ti-F bonds while oxygen vacancies were generated by H₂ reduction of Ti⁴⁺ on the surface. Oxygen-deficient TiO₂ sheets loaded with Pt nanoparticles shows substantially enhanced photoactivity in hydrogen evolution.

Interestingly, ionic liquid was also proved to be an ideal solvent for the synthesis of TiO₂ nanosheets exposed by high-reactive {001} facets. 1-butyl-3-methylimidazolium tetrafluoroborate was introduced to create a fluorine-rich TiO₂ crystal surfaces through a microwave-assisted process and the as-obtained micrometer-sized TiO₂ single crystals with remarkable 80% of {001} facets exhibited much higher catalytic activities for photodegradation of 4-chlorophenol.⁷⁴ Using similar method, biocompatible anatase TiO₂ single-crystals with 27-50% of chemically reactive {001} facets were fabricated with another ionic liquid, 1-methyl-imidazolium tetrafluoroborate, and the as-synthesized products were shown to be nontoxic by the experiment using Zebrafish.⁷⁵

Furthermore, some hydrothermal experiments resulted in the anatase TiO₂ with special morphologies. Using ammonium bifluoride (NH₄HF₂) as a morphology controlling agent and Ti(SO₄)₂ as the Ti source, Yu et al. synthesized tabular-shaped anatase single crystals exposed in micron size and with highly

Fig. 9 The SEM/TEM images, and SAED patterns of in situ nitrogen doped anatase TiO₂ nanosheets acquired by Liu et al. with TiN as titanium source.³³

energetic {001} facets.⁷⁶ By adjusting the molar ratio of fluorine to titanium, the morphology, crystallinity, and crystal size could be controlled. The chemical reaction was demonstrated as follows:

Similar morphology was also achieved by Liu et al.⁷⁷ They employed ethanol as a stabilizing agent and successfully synthesized hollow TiO₂ microspheres containing building units exposed with {001} facets. Hu et al. adopted a hydrothermal method to fabricate single-crystalline anatase TiO₂ with unique structures using anodized TiO₂ nanotubes as the starting materials and the assembled microspheres were composed of hollow structured particles exposing one {001} facet and eight {101} facets.⁷⁸ The effect of HF made the wall

Fig. 10 Multi-twinned anatase TiO₂ crystals by Ma et al.⁸⁰

of anodized TiO₂ nanotubes thinner and meanwhile, higher oxygen pressure was generated inside these nanotubes, which resulted in the fracture of TiO₂ nanotubes. Wang et al. employed pretreated tetragonal ZrO₂ polycrystalline fibers as a template to synthesize TiO₂ particles in (NH₄)₂TiF₆ aqueous solution.⁷⁹ After hydrothermal process, remarkable tubular structured materials composed of micro-sized anatase TiO₂ crystals were observed. It was speculated that anatase TiO₂ particles were deposited and grew on the surfaces of fibers while ZrO₂ fibers were gradually corroded, which can be examined from the evolution of XRD patterns of the products synthesized at different reaction time. Besides, Ma et al. synthesized uniform anatase TiO₂ crystals with multi-twinned {001} facets by using disodium ethylene diamine tetraacetate (EDTA) and F as morphology controlling agents (Fig. 10).⁸⁰ Both F and EDTA played an important role in stabilizing {001} facets. It was hypothesized that EDTA⁴⁻ should exist in the form of stable [Ti(EDTA)]²⁻ in the reaction medium of pH value close to 7.0. EDTA⁴⁻ adsorbed on {001} facets preferentially due to higher density of under-coordinated 5c-Ti on them in contrast with that on {101} facets. So this unique structure was obtained.

3.3 Nonhydrolytic alcoholysis approaches

Nonhydrolytic approaches can also produce TiO₂ dominated with {001} facets. Shan and Demopoulos prepared aqueous well-dispersed and phase-pure anatase TiO₂ nanoplatelets with the size of 20-40 nm via controlled hydrolysis of titanium tetrachloride (TiCl₄) in ethylene glycol.⁸¹ The reaction mechanism was as follows:

Water molecules released from reaction (1) can coordinate to Ti⁴⁺ cations and then titanium oxy/hydroxychloride complexes ([Ti(OH)_xCl_n]^{4-(n+x)} or [TiO_yCl_n]^{4-(n+x)}) were formed. Finally, TiO₂ was produced from the polymerization/condensation of these complexes. Another new nonhydrolytic solvothermal alcoholysis approach was developed by Zhu et al. and anatase TiO₂ crystals with dominant {001} facets were prepared successfully. This process only involved TiF₄ as source of Ti and F capping agent.⁸² This is a relatively green route compared to the traditional hydrolysis process in HF solution. In addition, TiO₂ microspheres composed of anatase TiO₂ nanosheets with dominant {001} facets (up to 83%) were synthesized by a simple alcoholthermal method, and these TiO₂ microspheres exhibited an excellent photocatalytic activity for the degradation of organic contaminants.⁸³ Chen et al. synthesized hierarchical structured spheres by the self-organization of anatase TiO₂ nanosheets dominated with {001} facets through a simple solvothermal method.³⁰ Apart from the large percentage of {001} facets (nearly 100%), small thickness (~3 nm) and a high specific surface area (~170 m²) rendered the material unique ability for lithium ion battery applications. Fang et al. also reported a new solvothermal process recently giving rise to three dimensional(3D) hierarchical structures of single-crystalline

Fig. 11 Three dimensional hierarchical structures of anatase TiO₂ nanosheets by Fang et al.³¹

anatase TiO₂ nanosheets dominated with well-faceted {001} facets (Fig. 11).³¹ Based on these experimental results, Fang et al. concluded that this unique structure was attributed to isobutyl alcohol acting as the reaction media, which might change the growth behavior of anatase TiO₂ nanosheets units and their mutual interactions as well.

3.4 High temperature gas phase oxidation techniques

A special gas-phase process developed by Amano et al. successfully fabricated faceted decahedral single-crystalline anatase particles with sizes of 50-250 nm for the first time (Fig. 12).³² The as-synthesized products exhibited extremely high photocatalytic activity for H₂ evolution from an aqueous methanol solution and the oxidative decomposition of organic compounds in an aqueous solution. It was postulated that the scarcity of crystalline defects might have induced these superior properties.

Fig. 12 Anatase TiO₂ crystals dominated with {001} facets by a gas phase process.³²

3.5 Other methods

Alivov and Fan have realized the transformation of TiO₂ nanotubes to truncated pyramid-like-shaped TiO₂ nanoparticles with a high percentage of photocatalytically active {001} facets after thermal annealing in ambient fluorine.⁸⁴ This method was quite different from the hydrothermal route employed by Hu et al.⁷⁸ and the size of nanoparticles was controllable, ranging from 20 nm to 500 nm by changing fluorine concentration during annealing process. More details of this transformation were further provided by studying the initial stages of the process.⁸⁵ Liu et al. prepared a unique yolk-shell structured particle with {001} faceted anatase TiO₂ as shells and Mo nanoparticles as cores by the arc-discharge method.⁸⁶ The as-obtained sample displayed enhanced photocatalytic efficiency, which was consistent with the result of UV-visible absorption spectra.

4. Synthetic strategies of other high-reactive facets dominated anatase TiO₂

Interestingly, besides {001} facets, other unusual reactive facets such as {110} and {100} facets of anatase TiO₂ were also observed. The highest-surface-energy {110} facets were observed by Liu et al. in the controlled synthesis of anatase TiO₂ single crystals from Ti powders (Fig. 13a).³⁴ This was the first time that {110} facets were observed on anatase TiO₂ single crystals where a high percentage of {001} facets was also concurrently observed. Wu et al. synthesized nanostructured anatase TiO₂ with high-energy {001} facets via a nonaqueous synthetic route.⁸⁷ Simultaneously, they tailored the reaction process by introducing a trace amount of water and then the pure {010} facets exposed rhombic anatase nanocrystals were acquired on the basis of a hydrolysis mechanism. Most recently, anatase TiO₂ crystals with photocatalytic active {100} facets were also synthesized using solid Na-titanates as a Ti source under hydrothermal condition (Fig. 13b).⁸⁸ At the beginning, Ti(OH)₄ fragments, hydroxyl and sodium ions were produced on the surface of Na-titanates, and then anatase TiO₂ nuclei was formed by the arrangement of Ti(OH)₄ fragments through dehydration reaction. Then the hydroxyl ions selectively adsorbed on anatase {100} facets in the growth process of anatase TiO₂ nuclei and thus {100} facets were preserved. These reports are

Fig. 13 (a) Anatase TiO₂ single crystals with {110} facets and a large percentage of {001} facets.³⁴ (b) Anatase TiO₂ single crystals exposed with {100} facets.⁸⁸

very significant and may open the pathway to deeper understanding the synthetic strategies for anatase TiO₂ exposed with other high-reactive facets.

5. Summary and outlook

Understanding and controlling the morphology of single crystals plays a key role for the synthesis of high-quality materials with defined surface properties. This article summarizes the synthetic chemistry of anatase TiO₂ with high-energy facets, especially with the first-principle calculations predicting the thermodynamically stable crystal morphology and various novel synthesis strategies developed to fulfill the high-energy facets in the past few years.

In spite of the rapid development in the recent past of the theoretical predictions as well as the experimental results for the anatase TiO₂ with high-energy facets, many challenges exist in several aspects. For example, Herman et al. reported the reconstruction of (001) surface under ultrahigh-vacuum conditions (UHV), which strongly stabilizes the high-energy surface and largely reduces its reactivity as well.⁴⁵ In this respect, examining the status of possible reconstruction of these high-energy facets is necessary and avoiding the reconstruction through novel synthesis method is highly desirable. Furthermore, considering TiO₂ crystals are being generally used as a model system to explore numerous important surface phenomena on atomic scale, synthesis of high-quality large size anatase TiO₂ single crystals is also very useful for surface science study. Importantly, almost all the synthesis methods reported to prepare anatase TiO₂ crystals with high-energy facets used F-containing compounds as capping agents to control the morphology, which is quite toxic and corrosive. Thus it is highly desirable to develop a new synthesis strategy with reduced or no fluorine containing compounds as capping agents. More importantly, organic surfactants adsorbing onto the surfaces can also remarkably change the free energy of these surfaces and interfacial properties, but the choice of surfactant remains empirical due to many unsolved issues including a quantitative description of the adsorption behaviour and in-depth understanding of selective surface-adsorption energies on nanocrystals. The more complex capping agents might be also a practical solution to create anatase TiO₂ crystals with highly-reactive facets. High-index facets can be also expected to be prepared in the future through theoretical predictions and experimental design. Apart from these challenges, liquid phase epitaxial process might be a useful technique to generate the 2-dimensional structured anatase TiO₂ exposed by {001} facets using substrates with suitable lattice spacing such as SrTiO₃. In conclusion, it could be expected that both opportunities and challenges exist for the synthesis of anatase TiO₂ with high-energy facets, and a better understanding and controlling of the synthesis process will be the key to further breakthroughs in this emerging front of research.

Acknowledgements

This work was financially supported by Scientific Research

Foundation of East China University of Science and Technology (YD0142125), Pujiang Talents Programme and Major Basic Research Programme of Science and Technology Commission of Shanghai Municipality (09PJ1402800, 10JC1403200), Shuguang Talents Programme of Education Commission of Shanghai Municipality (09SG27), National Natural Science Foundation of China (20973059, 91022023, 21076076), Fundamental Research Funds for the Central Universities (WJ0913001), Program for New Century Excellent Talents in University (NCET-09-0347) and the Australian Research Council (ARC) through Discovery Project program (DP1095861, DP0987969).

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