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# Titanate and Titania Nanostructured Materials for Environmental and Energy Applications: A Review

Cite this: DOI: 10.1039/x0xx00000x

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Received 00th October 2014,  
Accepted 00th October 2014

DOI: 10.1039/x0xx00000x

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Nanosized TiO<sub>2</sub>-based materials with unique structural and functional properties have already led to breakthroughs in various applications including photocatalysis, adsorption, lithium-ion batteries, etc. In this review, we present the state-of-the-art development of fabrication strategies of titanate/titania nanostructures and their corresponding environmental and energy applications. First, the structural features of titanate and titania and their correlation are explained in great details. After which, recent research efforts on the development of multiple dimensional titanate materials are summarized. Following that, the applications of titanate/titania nanomaterials in the fields of adsorbent, photocatalysis, lithium-ion batteries, photovoltaics, electrochromic device, self-cleaning and oil-water separation are reviewed. Finally, the future perspectives for the nanostructured titanate and TiO<sub>2</sub> are discussed. Continuous development in this area is essential to endow TiO<sub>2</sub>-based materials with advanced functionality and improved performance for practical applications.

## 1. Introduction

Problems concerning energy depletion and environmental pollution have become hot topics across the globe in the recent decades.<sup>1,2</sup> Our rapacious consumption of carbon-based fossil fuel energy sources has led to carbon dioxide accumulation in the atmosphere, with the outcome of global warming and climate change. Although in the danger of depletion, fossil fuels have continued to be the major source of energy driving human civilization.<sup>3</sup> The energy and environmental concerns call upon the reformation of our energy structure to rely on green and sustainable sources, such as solar energy. In addition, air and water, the most fundamental matter that life depends on, have become threat to humans due to the discharge of waste chemicals. For instance, 1.4 million km<sup>2</sup> of China was covered by a hazardous dense haze in January 2013, with more than 800 million victims.<sup>4</sup> Effluent water is often contaminated with organic dyes that are hardly bio-degradable. The Fukushima Incident in Japan during March 2011 released radioactive ions to the environment. With increased social impact and general public awareness, new materials or approaches are in urgent demand to solve or mitigate these problems.

Titanium dioxide is one of the most heavily investigated oxide materials in addressing energy and environmental crises. Due to its favorable electronic and optoelectrochemical properties, it has been widely applied to solar cells, photocatalysts, lithium ion battery electrodes, smart coatings, *etc.*<sup>5-13</sup> However, improved properties are necessary to meet high demand and complex requirements in the markets. Numerous efforts has been made to develop nanostructured titania materials with engineered compositions, morphologies or

heterostructures. The prosperous development of titanium dioxide nanomaterials has also thrived the investigation of a class of TiO<sub>2</sub>-based structures: layered titanate materials.<sup>14</sup>

Layered titanate has close structural resemblance to titanium dioxide, both composed of TiO<sub>6</sub> octahedra units connected by sharing corners and edges. In fact, the distinction between layered titanate and TiO<sub>2</sub> is not very clear, especially in the early literature, and the term titanium dioxide is still occasionally used to name the material that should more formally be called layered titanate. Layered titanate materials have attractive features of their own, including extremely large ion-exchange capacity, fast ion diffusion and intercalation, and high surface charge density. These favorable properties of layered titanate are endowed by their unique crystal structure, where the negatively charged two-dimensional titanium-containing sheets are separated to a large distance by exchangeable cations and molecules in the interlayer. Another attractive feature of titanate is its ability to transform to titanium dioxide. This allows the use of titanate as flexible precursors for TiO<sub>2</sub> nano-engineering. The ability of fabricating titanate in a variety of shapes allows the morphology to be inherited by the subsequent TiO<sub>2</sub> formation. The open crystal structure of layered titanate permits easy and uniform doping of atoms or molecular assembly, which is hard to achieve if working directly with the more compact TiO<sub>2</sub> polymorphs. Several reviews summarized the recent development of titanate nanotubes or the derived titania for its photocatalytic application.<sup>15, 16</sup> In this review, we discuss the recent development of multi-dimensional titanate materials by the newly emerged technologies and their related applications, and give the readers a full picture in terms of the history and technology evolution trends of titanate materials in

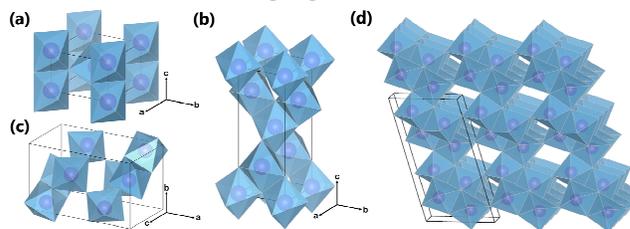
this area. We first provided a detailed discussion on the crystal structures of this class of layered titanate materials. The synthesis method for titanate formation is then reviewed, ranging from conventional solid state synthesis, to the recently developed approaches that cover titanate with one-dimensional (1D) linear (tube, wire, or belt), 2D sheet, and hierarchical 3D structures. Lastly, the applications of layered titanate materials dealing with energy and environment problems are summarized. Special focus has been given to their use as adsorbent, photocatalyst, battery electrode, photovoltaics cell, electrochromic device, and in self-cleaning, oil-water separation and biomedical applications.

## 2. Crystal Structure of Titania and Titanate

Titania ( $\text{TiO}_2$ ) can exist in a large number of different crystalline structures, but it is generally described as having three distinct polymorphs: rutile, anatase, and brookite (Figure 1). In all polymorphs, titanium cations are six-fold coordinated to oxygen anions, forming distorted  $\text{TiO}_6$  octahedra, and all polymorphs have  $\text{TiO}_6$  octahedra joined by sharing the octahedral edges (some have corner sharing as well). The different crystal structures of titania differ by the spatial arrangement of the  $\text{TiO}_6$  octahedra building blocks. Rutile is the thermodynamically stable form of bulk titania, which has tetragonal structure with a space group  $P4_2/mnm$  (136) (Figure 1a). Although being meta-stable at the bulk form, anatase and brookite can be stable when the crystal is in nano-size, due to their smaller surface energy. Comparing among equal sized particles, Zhang et al.<sup>17</sup> employed thermodynamic analysis and suggested that anatase, brookite, and rutile are stable when the size is less than 11 nm, from 11 nm to 35 nm, and greater than 35 nm, respectively. Obviously, the shape and size of nanoparticles will be affected by the experimental conditions (synthesis method, temperature, duration, precursor, etc.). In the work by Zhang et al.<sup>17</sup>, they found that when the heating temperature was at 450 °C, only anatase and brookite were present in the mixed phase particles, and their sizes varied from around 10 to 14.5 nm, and from around 12 to 17 nm respectively, when heating time increased from 2 to 24 h. Rutile phase began to form after heating the anatase and brookite mixture at 580 °C, and the rutile particle grew to around 60 nm after 21 h.<sup>17</sup> Tay et al.<sup>18</sup> hydrothermally synthesized cubic shaped anatase nanoparticles in size ranging from 18 to 42 nm using  $\text{TiS}_2$  precursor in 0.5 M of NaOH at 200 °C for 24 hours. Anatase also possesses a tetragonal structure, with space group  $I4_1/amd$  (141) (Figure 1b). Brookite, on the other hand, is orthorhombic with space group  $Pbca$  (61) (Figure 1c). Through hydrothermal synthesis, Tay et al.<sup>18</sup> obtained pure brookite nanoplates after treating  $\text{TiS}_2$  in 1.2 M of NaOH at 200 °C for 24 hours. The length of the nanoplates was around 100-400 nm, the width and thickness were in the range of 40-110 nm and 10-30 nm, respectively.

In addition to rutile, anatase, and brookite, there is one exotic phase of pure titania,  $\text{TiO}_2(\text{B})$ . This phase is often formed via solution synthesis. Although having the same chemical formula  $\text{TiO}_2$ ,  $\text{TiO}_2(\text{B})$  possesses a very open crystal structure (Figure 1d) and shares more similarity with titanate (to be elaborated later) than other  $\text{TiO}_2$  phases. Its space group  $C2/m$  (14) is the same as monoclinic titanate (Figure 2a), and can be imagined of as titanate sheet having structural entity composed of only two  $\text{TiO}_6$  octahedra, with adjacent sheets joined together by edge-sharing. Other than precipitating directly from solution,  $\text{TiO}_2(\text{B})$  was identified as an intermediate product in the calcination of titanate, following the transformation sequence of protonated titanate  $\rightarrow \text{TiO}_2(\text{B}) \rightarrow \text{anatase TiO}_2$ .<sup>19</sup> Anatase and rutile are the most studied polymorphs of titania for solar driven applications such as photocatalysis and photovoltaics.

Both of them have indirect band gap, which is 3.0 eV for rutile and 3.2 eV for anatase, corresponding to ultra-violet (UV) light absorption. It is generally accepted that anatase is the most active phase for photocatalysis applications, due to its better electronic and surface chemistry properties. Their properties can be further improved via doping or forming heterojunctions with other phases with favorable electronic coupling.

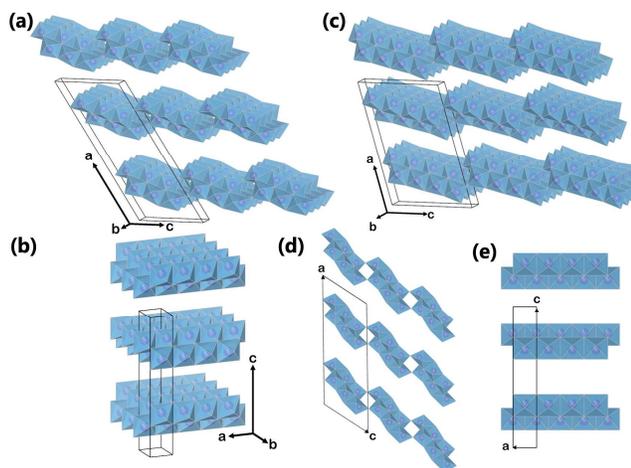


**Figure 1.** Crystal structures of  $\text{TiO}_2$  polymorphs: (a) Rutile; (b) Anatase; (c) Brookite; and (d)  $\text{TiO}_2(\text{B})$ . Purple spheres represent Ti atom, and the blue octahedra represent  $\text{TiO}_6$  blocks. Oxygen atoms at the corner of the octahedra are omitted for clarity.

Other forms of  $\text{TiO}_2$  and  $\text{TiO}_2$ -based phases exist. In their early discovery, these  $\text{TiO}_2$ -based materials have been mistakenly assigned as anatase  $\text{TiO}_2$ .<sup>20, 21</sup> However, later investigations confirmed that they were actually a class of layered titanate material, with a general formula  $M_x\text{Ti}_y\text{O}_{x/2+2y}\cdot z\text{H}_2\text{O}$  ( $M = \text{H, Li, Na, K, etc.}$ ).<sup>22, 23</sup> These titanate structures have 2D layers formed by connected  $\text{TiO}_6$  octahedral blocks, with cations and neutral molecules inserted into the inter-layer space. Based on crystal symmetry, the titanate materials can be further divided into two subclasses: monoclinic and orthorhombic (Figure 2a-b). All monoclinic forms of titanate are found to belong to space group  $C2/m$  (12). The 2D sheets in monoclinic titanate are zigzag in nature. Each of the zigzag layers is built upon the structural entity composed of several collinear edge-sharing  $\text{TiO}_6$  octahedra lying in the (010) planes. Every single structural entity is joined by corner to the other structural entities related to it by c-translation, and is further joined by edge to other structural entities related to it by  $2_1$  screw operation (at  $a = \pm 1/4$ ,  $c = 1/2$ ), forming a continuous 2D sheet. The various stoichiometry of monoclinic titanate is related to the number of  $\text{TiO}_6$  octahedra composing the structural entity. For example, in  $\text{H}_2\text{Ti}_3\text{O}_7$ , the structural entity is composed of 3 octahedra blocks (Figure 2a), whereas in  $\text{H}_2\text{Ti}_4\text{O}_9$ , the structure entity has 4 octahedra blocks (Figure 2c). The layering nature monoclinic titanate can also be influenced by the size and amount of the other cations relative to titanium. When there is not enough cations to support the large interlayer spacing, the separate interlayers will join together by sharing  $\text{TiO}_6$  octahedra corners. This can be taken advantage of for the treatment of radioactive ions, as will be discussed later. The other form of layered titanate has an orthorhombic crystal structure with space group  $Immm$  (71), which is frequently termed lepidocrocite titanate (for their structural analogy to  $\gamma\text{-FeOOH}$ ) (Figure 2b).<sup>24</sup> Although being in a different crystal system, the  $\text{TiO}_6$  octahedra arrangement in orthorhombic titanate is closely related to monoclinic titanate. Indeed, the orthorhombic titanate can be visualized as the monoclinic titanate with the structural entity composed of infinite number of  $\text{TiO}_6$  octahedra. In effect, orthorhombic titanate layer is straight and continuous in the a-direction. Because of the absence of corner-sharing, charge balance have to be maintained from partial occupancy of Ti sites, leading to chemical formula  $\text{H}_x\text{Ti}_{2-x/4}\square_{x/4}\text{O}_4\cdot z\text{H}_2\text{O}$  (where  $\square$  denotes vacancy, and  $x$  is close to 0.7).<sup>25</sup> Distinction between the monoclinic and orthorhombic titanate is usually difficult, because their crystallinity is usually low, and their structural similarity resulted in close characteristic peak positions in XRD. Projections of monoclinic

titanate  $H_2Ti_3O_7$  and orthorhombic titanate onto their (010) plane is illustrated in Figure 2d-e.

Thanks to the open crystal structure and large interlayer spacing in titanates, they can accommodate many varieties of cations and neutral molecules in the interlayer spaces, endowing them unique physicochemical properties. Large dipole moment exists in titanate materials, where the negatively charged titanate layer is balanced by positively charged interlayer cations. Layered titanate has very good cation exchange capability. Because of the negative charge on the titanate sheet, it can adsorb positively charged molecular species effectively. These effects have been applied in waste water remediation for the treatment of radioactive ions and organic dyes, which is the subject of Section 4.1. Although titanate in its native form is not very effective for solar-driven applications, it can be transformed to titania via calcination with minimal morphological distortions. Therefore, titanate can serve as precursor and scaffold for more active materials. The ability to modify the content within the interlayer of titanate provides a means for doping or forming hetero-structures of titania. Since the dopant is mixed at an atomic level through the interlayer, the formed  $TiO_2$  by transformation of modified titanate is expected to have uniform doping, in contrast with the shallow surface doping by diffusion in bulky materials. The potential of titanate in the area of photocatalysis will be discussed in Section 4.2. The open framework of titanate also provides fast Li intercalation pathways, making it a candidate cathode material for Li-ion battery, which will be addressed in Section 4.3. Section 4.4 highlights applications of  $TiO_2$  for photovoltaic and electrochromic devices. Other functionalities of titanate stem from its surface hydrophilicity and the ability to turn to hydrophobicity through chemical modification, making them useful in biomedical application, self-cleaning and oil-water separation applications, are to be discussed in Section 4.5.



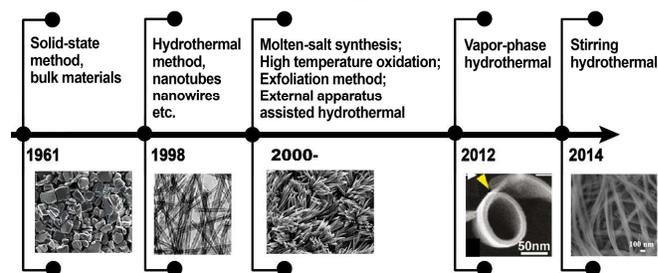
**Figure 2.** Crystal structures of (a) monoclinic titanate  $H_2Ti_3O_7$ ; (b) orthorhombic titanate  $H_xTi_{2-x/4}\square_{x/4}O_4$ ; (c) monoclinic titanate  $H_2Ti_4O_9$ . The (010) plane projection of (d) monoclinic titanate  $H_2Ti_3O_7$ , and (e) orthorhombic titanate. Purple spheres represent Ti atom, and the blue octahedra represent  $TiO_6$  blocks. Oxygen atoms at the corner of the octahedra, other cations and molecules in the interlayers are omitted for clarity.

### 3. Development of Multi-dimensional Titanate Materials

In this section, the technological development for the formation of multi-dimensional titanate materials will be described. The formation conditions of titanate include the followings: 1) having titanium precursor sources ( $TiO_2$ , Ti, etc.); 2) being in alkaline reaction environment; 3) negative free energy change (i.e. thermodynamically spontaneous) for the reaction.



Based on this guidance, the experimental designs toward multi-dimensional (bulk, 0D, 1D, 2D, and 3D) titanate materials by solid state method, hydrothermal reaction, electrochemical spallation, high-temperature oxidation, and other approaches have been developed, and the history of key development of titanate materials are summarized in Figure 3. Aiming at achieving better functional properties as well as improved performance, there has been a strong incentive for the size and morphology control of titanate materials.<sup>26</sup>



**Figure 3.** The history of technological development in the synthesis of titanate materials.

#### 3.1 Fabrication of bulk titanate materials by solid-state reaction

Solid state reaction refers to a chemical process without the usage of solvent. The reaction usually takes place at a high temperature for several days to ensure the complete reaction. The typical titanate formula  $A_2Ti_nO_{2n+1}$  consists of an alkali metal oxide ( $A_2O$ ) and titanium dioxide ( $nTiO_2$ ). Therefore, different kinds of bulk titanate materials can be obtained by mixing  $TiO_2$  with the metal carbonate ( $A_2CO_3$ ,  $A = Cs, K$  or  $Na$ , etc.) with certain molar ratio. Due to evaporative loss of the alkaline component at the calcination temperature, the final product will not be a single phase of  $A_2Ti_nO_{2n+1}$  if starting from a stoichiometric mixture. Therefore, the experimental molar ratio of the alkali oxide and the titanium dioxide should be higher than calculated stoichiometry.

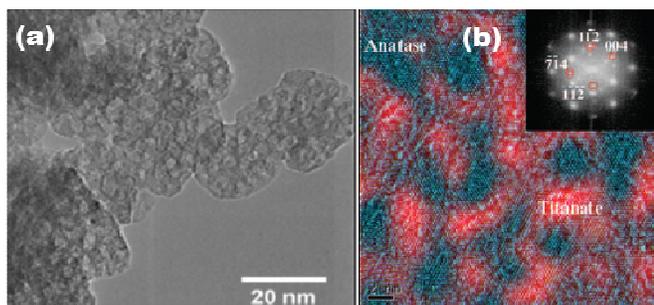
Researchers have synthesized different kinds of alkali metal titanates  $A_2Ti_nO_{2n+1}$ . For example, the  $Na_2Ti_3O_7$  and  $K_2Ti_4O_9$  were synthesized by heating mixtures of  $Na_2CO_3$  and  $TiO_2$  mixture ( $\leq 1:3$  molar ratio),  $K_2CO_3$  and  $TiO_2$  mixture ( $1:3.5$  molar ratio) at  $800^\circ C$  for 20 h.<sup>27</sup> With the same approach, Sasaki et al.<sup>28</sup> used  $Cs_2CO_3$  and  $TiO_2$  ( $1:4.5$  molar ratio) as starting reagents, synthesized the bulk  $Cs_2Ti_5O_{11}$  material. The protonation process (hydrogen-exchange) of  $Cs_2Ti_5O_{11}$ ,  $Na_2Ti_3O_7$  and  $K_2Ti_4O_9$  were performed in the acid solution. The acid solution was refreshed frequently to completely remove alkali ion from the compounds, leading to the final products of protonated titanate  $H_2Ti_5O_{11}$ ,  $H_2Ti_3O_7$  and  $H_2Ti_4O_9$  respectively. Moreover, the starting material of the alkali carbonate ( $A_2CO_3$ ) can also be replaced by  $ANO_3$ ,  $AOH$  or  $A_2O$  ( $A = Cs, K$  or  $Na$ , etc.) and so on. For example, Tornaux and co-workers<sup>29</sup> heated  $KNO_3$  and  $TiO_2$  at  $1000^\circ C$  for 48 h and the product was  $K_2Ti_4O_9$ . The resultant powder was then hydrolyzed with  $HNO_3$  at room temperature to yield a solid product  $H_2Ti_4O_9 \cdot H_2O$ . Wang et al.<sup>30</sup> also adopted a

similar method to synthesize  $K_2Ti_4O_9$  using  $K_2O$  and  $TiO_2$  as the starting materials.

Although different kinds of bulk titanate materials have been prepared through solid state reaction, the resultant particle sizes of these prepared materials are large, and the specific surface areas are small, which limits the materials' practical application in photocatalysis and a number of other areas. In order to obtain high photocatalytic activity to or improve ion-exchange/intercalation performance, it is necessary to synthesize nano-sized materials possessing large specific surface areas and with special morphological features. On the other hand, the bulk titanate materials can be precursor for the formation of nanosheets by exfoliation method, which is discussed in the following sections.

### 3.2 Controlled growth of 0D titanate/titania nanoparticles

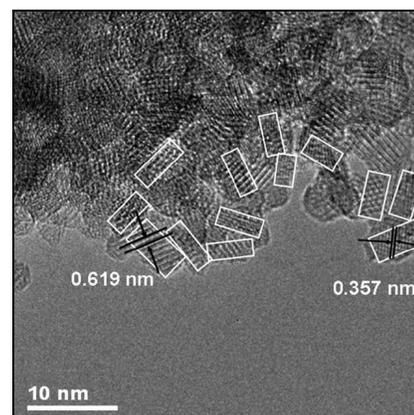
0D nanoparticle refers the ones with spherical or nearly equal-axial dimensions within the nanosize range (<100 nm). 0D titanate nanoparticles are rarely reported in the past decades, probably due to the difficulty in experimental control. Our group<sup>31</sup> reported a sol-gel method combined with solvothermal approach to fabricate titanate/titania hybrid nanoparticles. These particles have a high surface area ( $180 \text{ m}^2 \text{ g}^{-1}$ ) and the transmission electron microscopy (TEM) image (Figure 4a) shows cubic particles of averaging 20 nm in size, with an internal structure of tessellation suggestive of intragranular pores with a size around 3.0~3.5 nm. HRTEM image is shown in Figure 4b of a single particle consisting of both anatase (cyan color) and titanate (red color) nanocrystallites that share intimate interface. Each phase has an average crystal size around 5nm. A fast Fourier transformation (FFT) study of the entire area shows two sets of diffraction patterns (see inset of Figure 4b). The detailed formation mechanism is not fully understood. We suspect that the sol-gel approach can provide homogenous titanium source precursors, and the following solvothermal approach promotes crystallization process of titanate nanoparticles in alkaline condition (in triethylamine) via a bottom-up strategy.



**Figure 4.** (a) TEM and (b) HRTEM image of the as-synthesized photocatalyst with two phases distinguishable by color. The inset shows the diffraction pattern obtained from the FFT study of the monoclinic titanate (weak spots) and tetragonal anatase phases (strong spots).<sup>31, 32</sup>

Recently, titanate-derived  $TiO_2(B)$  nanoparticles<sup>33</sup> were prepared using a hydrothermal method followed by thermal treatment. Firstly, a titanium glycolate complex was formed by dissolving Ti metal in a mixture of  $H_2O_2$  and  $NH_3$  in water. Then the mixture is subjected to hydrothermal treatment, forming titanate nanoparticles. Then the as-prepared product is calcined in dry air to form  $TiO_2(B)$ . The TEM data demonstrates that the material is composed of nanoparticles of ca. 2.5~4.3 nm size (based on analysis of 100 nanoparticles), with a

relatively narrow size distribution (Figure 5). These nanoparticles form agglomerates of 0.3~3.0  $\mu\text{m}$  in size. BET surface area determined from  $N_2$  adsorption is  $251 \text{ m}^2 \text{ g}^{-1}$  (pore volume  $0.12 \text{ cm}^3 \text{ g}^{-1}$ ).

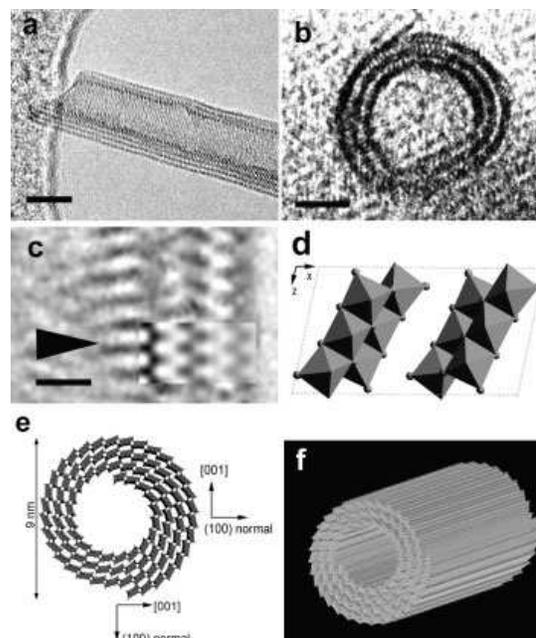


**Figure 5.** HRTEM image of  $TiO_2(B)$  nanoparticles. White boxes delineate primary (nano) particles within the agglomerates.<sup>33</sup>

### 3.3 Rational synthesis of 1D titanate nanostructures

#### 3.3.1 Conventional hydrothermal method

The first literature on the formation of  $TiO_2$ -based materials in the nanotubular form using hydrothermal method was reported by Kasuga et al.<sup>20, 21</sup> in 1998. In this process, amorphous  $TiO_2$  was treated at elevated temperatures in a solution that was highly concentrated in hydroxide ions (10 M NaOH), and no sacrificial templates was needed. A traditional reaction vessel for this purpose would be a poly (tetrafluoroethylene) (PTFE) lined autoclave, which has good chemical resistance and are able to tolerate the temperature and corrosive alkaline environment to be encountered. A clear advantage for this technology is the capability of near 100% conversion of the precursors to titanate nanotubes in one single process stage that does not involve excessively high temperature. As shown in Figure 6, the hollow structure of the nanotubes is formed, and the walls of nanotubes usually contain several layers.



**Figure 6.** Titanate nanotubes: (a, b) HRTEM image; (c) Enlarged HRTEM image of a part in (a); (d) Structure model of one unit cell of  $\text{H}_2\text{Ti}_3\text{O}_7$  on the [010] projection; (e) Schematic drawing of the structure of nanotubes; (f) Three-dimensional drawing of a nanotubes.<sup>34</sup>

However, whether the crystal structure of titanate nanotubes belongs to monoclinic or orthorhombic system is still under the debate. This is caused by a few reasons: (1) its crystallite size is very small, which leads to the broadening of peaks in the XRD patterns; (2) the proposed crystal structures share close similarities; (3) the precise positions of hydrogen atoms inside the crystals is difficult to be located.<sup>15</sup> The trititanate acid ( $\text{H}_2\text{Ti}_3\text{O}_7$ ), lepidocrocite type titanate ( $\text{H}_x\text{Ti}_{2-x/4}\square_{x/4}\text{O}_4$ ,  $\square$  represents a vacancy), dititanate acid ( $\text{H}_2\text{Ti}_2\text{O}_5\cdot\text{H}_2\text{O}$ ), and tetratitanate acid ( $\text{H}_2\text{Ti}_4\text{O}_9\cdot\text{H}_2\text{O}$ ) had been proposed as shown in the Table 1. Peng and co-workers<sup>34</sup> proposed that the crystal structure of titanate nanotubes corresponded to the layered  $\text{H}_2\text{Ti}_3\text{O}_7$  with a monoclinic crystal structure, which is obtained by rolling several (100) planes around axis [010] or [001]. However, Ma et al.<sup>24</sup> suggested that the titanate nanotube possessed a lepidocrocite-type ( $\text{H}_{0.7}\text{Ti}_{1.827}\square_{0.175}\text{O}_4\cdot\text{H}_2\text{O}$ , where  $\square$  indicates a vacancy) structures, which is supported form the selected area electron diffraction study. According to the Raman study, Gao et al.<sup>35</sup> believed that the crystal structure of titanate nanotube could be better described by protonated titanate  $\text{H}_{0.7}\text{Ti}_{1.827}\square_{0.175}\text{O}_4\cdot\text{H}_2\text{O}$ , as it has been demonstrated that the different types of polymerization of  $\text{TiO}_6$  octahedra in these compounds result in the different vibrational features. Lepidocrocite titanate contains “flat” host layers in which  $\text{TiO}_6$  octahedra are combined with each other via edge-sharing, while the crystal structure of  $\text{H}_2\text{Ti}_3\text{O}_7$  consists of corrugated “ribbons” of edge-sharing  $\text{TiO}_6$  octahedra, where the octahedra join at the corners to form a “stepped” layered structure, as shown in Figure 2.

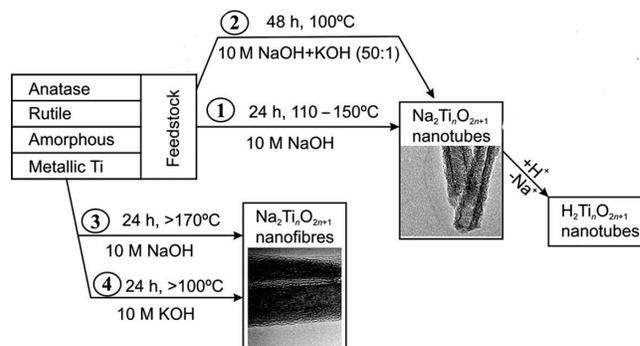
**Table 1.** Comparison of Structural Parameters of the Titanate Nanotubes to Those of the Protonic Titanates (Å)<sup>35</sup>

$\text{H}_2\text{Ti}_3\text{O}_7^a$			$\text{H}_2\text{Ti}_4\text{O}_9\cdot\text{H}_2\text{O}^b$			$\text{H}_2\text{Ti}_2\text{O}_5\cdot\text{H}_2\text{O}^c$			$\text{H}_{0.7}\text{Ti}_{1.825}\square_{0.175}\text{O}_4\cdot\text{H}_2\text{O}^{d,e}$		
<i>d</i>	<i>hkl</i>		<i>d</i>	<i>hkl</i>		<i>d</i>	<i>hkl</i>		<i>d</i>	<i>hkl</i>	
7.87	2 0 0		9.05	2 0 0		9.04	2 0 0		9.272	0 2 0	
3.65	1 1 0		3.672	1 1 0		3.696	1 1 0		3.707	1 1 0	
3.05	3 1 0		3.185	3 1 0		3.204	3 1 0		3.226	1 3 0	
2.67	3 1 -2		2.668	2 1 -3		2.684	3 0 1		2.686	0 3 1	
2.37	1 1 -3		2.298	2 1 -4		2.306	5 0 1		2.324	0 5 1	
1.88	0 2 0		1.876	0 2 0		1.893	0 2 0		1.891	2 0 0	
						1.478	2 0 2		1.491	0 0 2	

<sup>a</sup>JCPDS 47-0561 monoclinic; lattice constants  $a = 16.023$ ,  $b = 3.749$ ,  $c = 9.191$ , and  $\beta = 101.45^\circ$ . <sup>b</sup>JCPDS 36-0655 monoclinic;  $a = 18.77$ ,  $b = 3.75$ ,  $c = 11.62$ , and  $\beta = 104.6^\circ$ . <sup>c</sup>JCPDS 47-0124 orthorhombic;  $a = 18.03$ ,  $b = 3.783$ , and  $c = 2.998$ . <sup>d, e</sup>Calculated from JCPDS 83-0702 orthorhombic;  $a = 3.783$ ,  $b = 18.545$ , and  $c = 2.982$ .

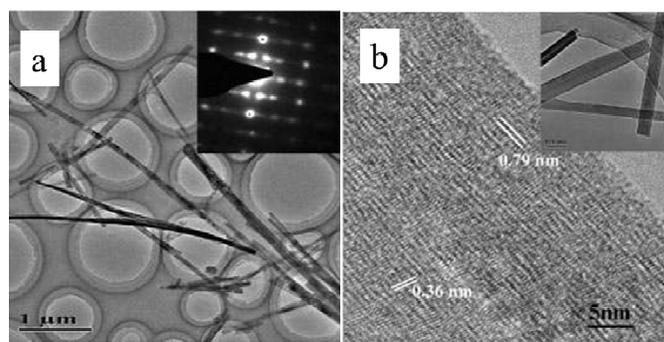
Since its introduction,<sup>20</sup> the alkaline hydrothermal production of titanate nanotube has been engineered in a direction of attaining the ease of operation, diminishing process cost, and facilitating scale-up. Further development of this alkaline hydrothermal strategy has revealed that all  $\text{TiO}_2$  polymorphs (including anatase, rutile, amorphous  $\text{TiO}_2$ , or even Ti metal) can be employed as the precursor in the formation of nanotubular or nanofibrous  $\text{TiO}_2$  (Figure 7).<sup>14</sup> Some classic preparation processes for the fabrication of titanate nanotubes are shown in Figure 7, which includes the original method (route 1 in Figure 7) of dispersing  $\text{TiO}_2$  raw material in solutions concentrating 10 M NaOH at 110–150 °C temperature range for a

duration of 24 h. Recently, it has been reported that using alternative solvents, the synthesis of titanate nanotube can be achieved at lower temperatures. For instance, using a mixture of NaOH and KOH<sup>36</sup> (route 2 in Figure 7), the conversion can be brought into near completion at a temperature as low as 100 °C with the aid of refluxing atmospheric pressure. Additional improvement in this reflux strategy includes further optimization of the aqueous solution composition and the addition of additives in order to synthesize titanate nanotubes at even lower temperatures for shorter durations.



**Figure 7.** Respective routes to the manufacture of titanate nanotubes and nanofibers.<sup>37</sup>

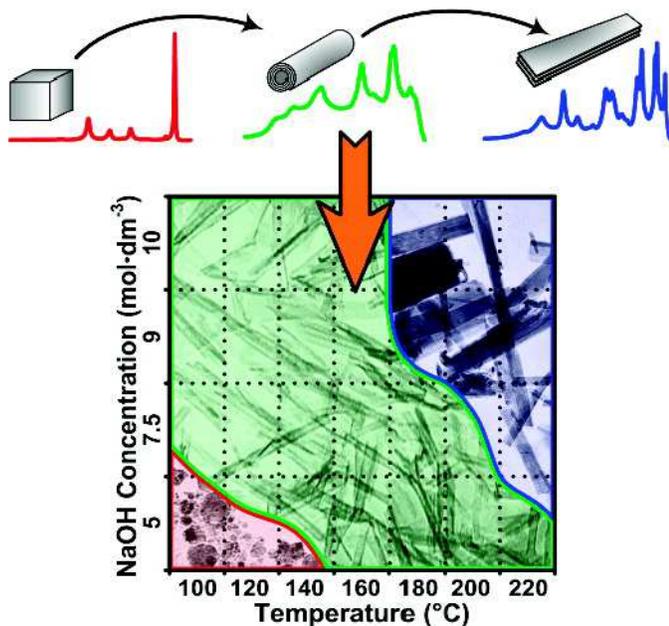
The formation of titanate nanofibers, wires or belts usually employs hydrothermal treatment of  $\text{TiO}_2$  in 10 M NaOH solution at higher temperatures than the case of nanotube ( $> 170^\circ\text{C}$ ) (route 3 in Figure 7).<sup>38</sup> For example, as shown in Figure 8, Zhu’s group<sup>39</sup> reported that the titanate nanowires were formed at 180 °C for 48 h by using the rutile titania as the starting materials. The formation of nanofibers can also be achieved if 10 M NaOH is substituted by 10 M KOH solution (route 4 in Figure 7).<sup>40</sup> The decrease of the reaction temperature tend to form nanotubes / nanofibers mixture.<sup>41</sup> Our group<sup>42</sup> has synthesized titanate nanobelts via a hydrothermal method using titanium disulfide ( $\text{TiS}_2$ ) as the precursor. Impressively, sodium titanate nanobelts could be formed under a relatively low alkaline concentration (1 M NaOH) and short duration (6 h).



**Figure 8.** (a) Low- and high- magnification TEM images of the nanowires synthesized by treating rutile with 10 M NaOH for 48 h at 180°C.<sup>39</sup>

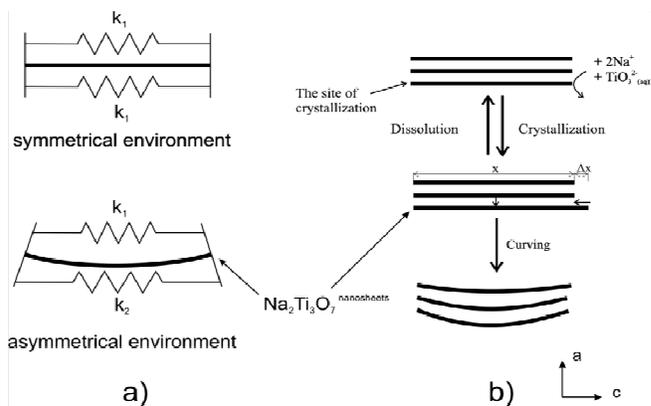
In brief, hydrothermal conditions, which include the type and concentration of the solution, temperature, and time, play a significant role for the controlled synthesis of titanate nanostructures. For example, Morgan’s group<sup>43</sup> systematically investigated caustic concentration and temperature on nanostructure

formation from Degussa P25 through alkaline hydrothermal treatment. The morphological phase diagram is provided in Figure 9, which is consistent with the above results and it is a useful guideline for nanostructure morphology control for the titanate materials.



**Figure 9.** Morphological phase diagram of Degussa P25 evolution indicating regions of nanostructure formation after 20 h of hydrothermal treatment.<sup>43</sup>

It is evident that  $\text{TiO}_2$  with different phase and morphology can be transformed to titanate nanotubes under specific hydrothermal conditions, but its formation mechanism and the detailed transformation for the nanotubular titanates is still unclear. Under hydrothermal condition, the intermediate single-layer or multilayered titanate nanosheets are observed and they play an important role during the formation of nanotubular morphology. The nanotubes can be formed by the scrolling and folding action of the sheets, and the driving force for this curving is proposed by several groups. Zhang's group<sup>44</sup> proposed that an asymmetrical chemical environment (Figure 10a) caused by the  $\text{H}^+$  or  $\text{Na}^+$  ion concentration imbalance on the different locations of the titanate nanosheets resulted in the bending of the nanosheets, generating excess surface energies. Bavykin et al.<sup>38</sup> proposed the bending of multilayered nanosheets is due to the mechanical tensions (Figure 10b) resulted from the nanosheets dissolution/crystallization process. The spontaneous crystallization and rapid growth of the layers may result in various layer widths, and it is likely that the tendency of releasing the excess surface energy, which has its root in the layer width imbalance, drives the movement of layers within the multiwall nanosheet. The consequence of this movement is the bending of multilayered nanosheets.<sup>14</sup> It is noted that the nanotube axis is along the [010] direction and thus the nanosheets bend around the b-axis due to the imbalance in the nanosheet along the c-axis. When the hydrothermal temperature is increased to 170°C, the titanate nanofibers or nanowires are formed and axis of the long wire is the [001] direction. At this high temperature, it is proposed that no nanosheets bending should occur and the resulting nanowires grow along the c direction ([001]), which is due to the maximization of dissolution/crystallization rate along this axis.



**Figure 10.** The driving forces for bending titanate nanosheets under the alkaline hydrothermal conditions. (a) Asymmetrical chemical environmental resulting in different surface tensions. (b) Imbalance in layer widths resulting in shifting of the layer and bending the nanosheets.<sup>14</sup>

### 3.3.2 Modified hydrothermal method

#### 3.3.2.1 External-apparatus assisted hydrothermal method

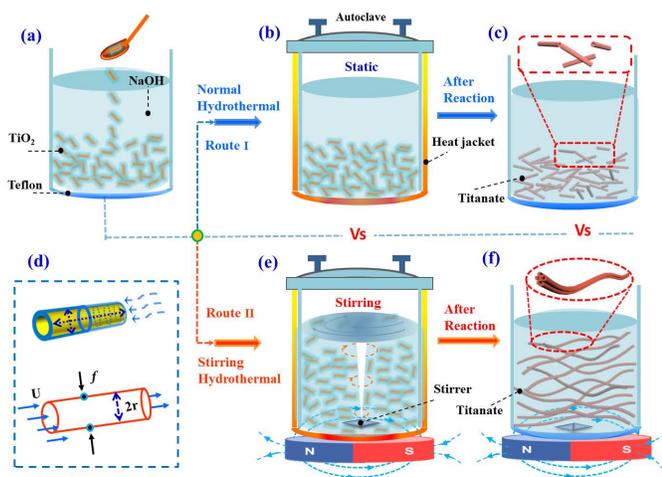
Although the discussed hydrothermal techniques have demonstrated the potential for the formation of titanate nanostructures, there are still some limitations: 1) slow reaction kinetics, which results in long reaction time; 2) short length of the nanotubes; 3) non-uniformity at large-scale. Various approaches, such as ultrasonication-assisted, microwave-assisted, and rotation-assisted hydrothermal methods have been explored to solve these problems. For example, the sonication and irradiation treatment can influence the morphology of titanate/titania nanostructured materials. The length of titanate nanotube becomes longer with the application of sonication pretreatment.<sup>45</sup> Zhu et al.<sup>46</sup> found that titanate whiskers and nanotubes were obtained at a sonication power of 580 W and 280 W, respectively. The titanate whiskers are obtained as a slender sheet with a length of about 1  $\mu\text{m}$  and a width of 60 nm, and the nanotubes with a diameter of about 5 nm, and a length of 200–300 nm are obtained. By this sonication-hydrothermal combination approach, Ma et al.<sup>47</sup> obtained nanosheet-, nanofiber- and nanotube-like morphologies at a sonication power of 100, 280 and 380 W, respectively. In particular, titanate nanotubes with outer diameters of 5–10 nm and lengths up to 600 nm were obtained. Their study<sup>46, 47</sup> revealed that the sonication with sufficient power might play an important role that promotes intercalating  $\text{Na}^+$  into titania lattices and breaking the Ti-O-Ti bonds. This induces easy peeling of large nanosheets, which consequently rolled into longer titanate nanotubes or other morphologies. Moreover, the level of applied microwave irradiation during the fabrication process is responsible for both the intercalation intensity of Na atoms into titanate nanotubes and the type of crystallization phase within nanotubes.<sup>48</sup> Interestingly, Dufour et al.<sup>49</sup> found that microwave treatment significantly reduces the heating time and generally produces smaller  $\text{TiO}_2$  nanoparticles, and it also affect the phase composition probably due to the modification of the kinetics of formation of the different phases.

Development of long nanotubes with relatively high surface area would be of great interest to tailor properties for applications.<sup>50, 51</sup> Recently, Horvath et al. reported conversion of nanotube to nanowire with revolving autoclave (60 rpm) at temperature of 130°C,

which is 20°C lower than that used in previous literature.<sup>52</sup> They suggested that flow pattern created directional mass transfer and facilitated parallel alignment of nanotubes to be further transformed into nanowires. Inspired by this setup, Torrente et al. synthesized titanate nanotubes with length scale one order of magnitude higher at a lower revolving speed of 20 rpm,<sup>53</sup> but agglomeration of nanotubes into secondary linear structure became pronounced. In addition, there are intrinsic limitations associated with such approach including high energy consumption, sophisticated apparatus setup and limited rotation speed achievable (< 100 rpm).

### 3.3.2.2 Stirring hydrothermal method

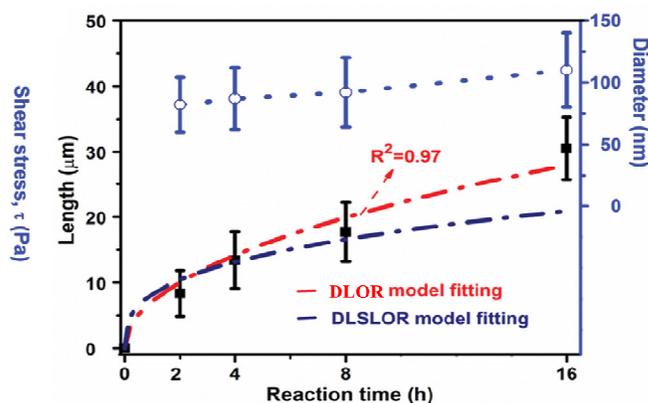
Although the external-apparatus assisted hydrothermal method has been developed to control the titanate nanotubular structures, the length of the nanotubes was still limited to one-micrometer. To overcome this limitation, our group reported<sup>10</sup> a protocol to rationally grow elongated titanate nanotubes with length up to tens of micrometers by a stirring hydrothermal method (Figure 11). Our detailed study confirmed that the mechanical force-driven stirring process synchronously improved the diffusion and surface reaction rate of titanate nanocrystal growth in solution phase, which is the reason for lengthening of the titanate nanotubes via an oriented attachment mechanism. Besides, such setup also possesses advantageous features including less energy consumption, easier scaling up, and more flexible stirring speed control. Thus it represents a more viable and efficient approach.



**Figure 11.** Schematic illustration of the formation process of short and elongated nanotubular structures under normal (a-c) and stirring hydrothermal processes (e-f) at 130°C for 24 h respectively. (d) The growth model of nanotube along axial and radial direction (top) and the force analysis (down) of an individual nanotube formed in (e).<sup>10</sup>

Since the growth process of nanotubular titanate via dissolution – crystallization step is similar to the Ostwald ripening process, we first considered the diffusion-limited Ostwald ripening (DLOR model, Figure 12), according to Lifshitz-Slyozov-Wagner theory, to be the sole contributor in the growth mechanism. However, the experimental length ( $L$ ) data of the titanate nanotube fits well with the DLOR model for the short reaction times only (Figure 12). A remarkable accuracy ( $R^2 = 0.97$ ) of fits over the entire range of experimental data was given by diffusion-limited and surface reaction limited growth (DLSLOR) model. Thus, the growth of the

titanate nanotube deviates significantly from the diffusion-limited Ostwald ripening model and follows a mechanism involving both diffusion-control and surface reaction control under the stirring condition.

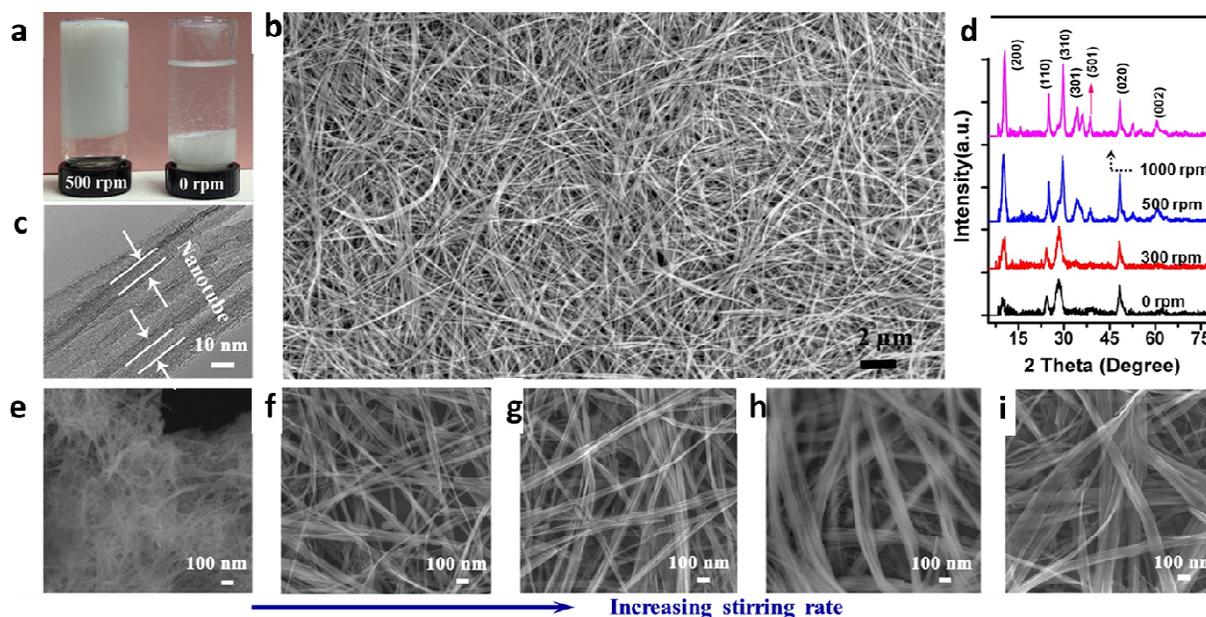


**Figure 12.** The kinetic study (the nanotubular length, diameter) of the nanotubular sample obtained under 500 rpm. The red curve and navy blue curve fit data  $L$  using the mixed diffusion- and surface reaction limited model (DLSLOR model) and diffusion-limited Ostwald ripening (DLOR) control growth model respectively.<sup>10</sup>

Based on zero-shear viscosity theory, it is known that in a suspension, solution viscosity ( $\eta_0$ ) is the sum of the solvent viscosity ( $\eta_s$ ) and a contribution from the suspended material:

$$\eta_0 = \eta_s + \frac{2}{15} v \zeta_r \quad (2)$$

where  $v$  is the number of nanotubes per unit volume and  $\zeta_r$  is the rational friction coefficient. In the traditional hydrothermal reaction for the preparation of titanate nanotubes, a low viscosity solution is formed due to the settlement of short titanate nanotubes and the weak interaction between them. To achieve high viscosity, it is essential to increase the suspended nanotube concentration ( $v$ ) and improve its interaction ( $\zeta_r$ ). In this regard, the controlled concentration of  $\text{TiO}_2$  precursor in solution and the formation of cross-linking network through elongating nanotubular structure are desired. Our study proved that the nanotube aspect ratio can be controlled by tuning the solution viscosity through “polymerization” of starting precursor, which is strongly related to the stirring rate (Figure 13).

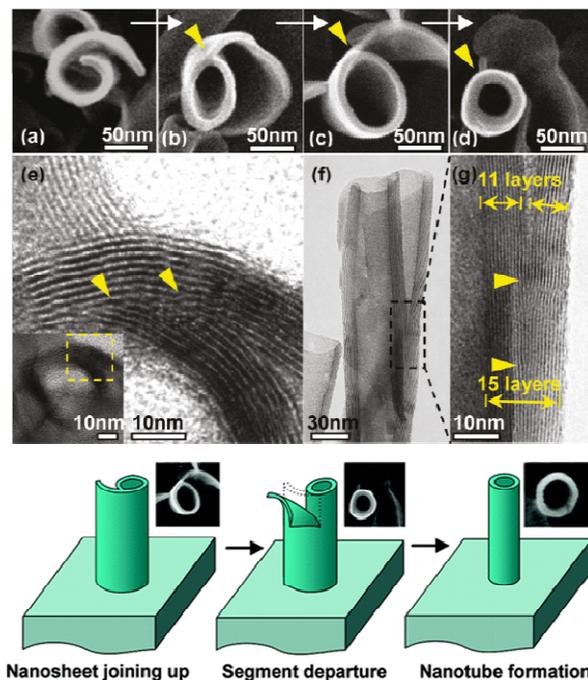


**Figure 13.** Fabrication and characterization of titanate nanotubes with different aspect ratios. (a) digital photos of resulted titanate solution obtained by hydrothermal method at 500 (left) and 0 rpm (right) after sedimentation; (b) FESEM images of titanate nanotubes obtained at 500 rpm; (c) TEM image of (b), the arrow indicating the formation of nanotubular structure; (d) XRD pattern of the products synthesized at different stirring speeds; and (e, f, g, h, i) FESEM images of the titanate nanotubes obtained by hydrothermal reaction at 130°C for 24 h in at a stirring rate of 0, 300, 400, 500 and 1000 rpm respectively.<sup>54</sup>

### 3.3.2.3 Vapor-phase hydrothermal approach

High-temperature NaOH flux and water vapor were found to play important roles in the growth of uniformly sized titanate nanorods. Recently, vertically aligned titanate nanotubes (Figure 14) were directly grown on a titanium foil substrate using a vapor-phase hydrothermal approach.<sup>44</sup> The resultant nanotubes displayed outer diameters in the range of 50 to 80 nm and average wall thickness of 10 nm, which corresponds to more than 10 titanate layers. It was observed that a distinctive nanosheet rollup mechanism dominated the nanotubes formation process, which differs remarkably from the mechanism frequently encountered in conventional liquid-phase hydrothermal reactions.

Beyond the modified hydrothermal method, various approaches have been applied to prepare 1D titanate nanostructures. For examples, the titanate nanowires can also be synthesized from reactions of TiO<sub>2</sub> nanoparticles with molten salt in the presence of a nonionic surfactant by a molten-salt synthesis method.<sup>55</sup> By this approach, large quantity of nanowires with uniform diameter around 100 nm was obtained with their length ranged from several to a few tens of microns. Moreover, a high-temperature oxidation method to obtain titanate nanorods was reported by Liu et al.<sup>56</sup> In this method, NaOH coated Ti metal chips was annealed at 800 °C in the presence of oxygen and water vapor, producing novel tabular titanate nanorods.

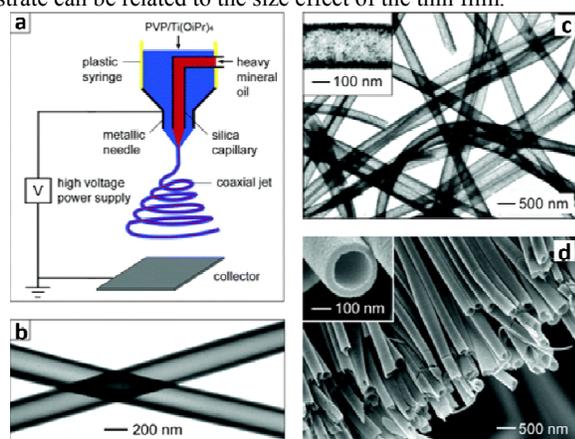


**Figure 14.** SEM and TEM images of the morphological evolution from a nanosheet into a nanotube; The proposed nanosheet roll-up mechanism is by vapor-phase hydrothermal approach.<sup>44</sup>

### 3.3.2.4 Sol-gel Approach

As a representative of wet chemistry method to fabricate 1D TiO<sub>2</sub> nanotubes or nanofibers, sol gel inherits the merit of low preparation

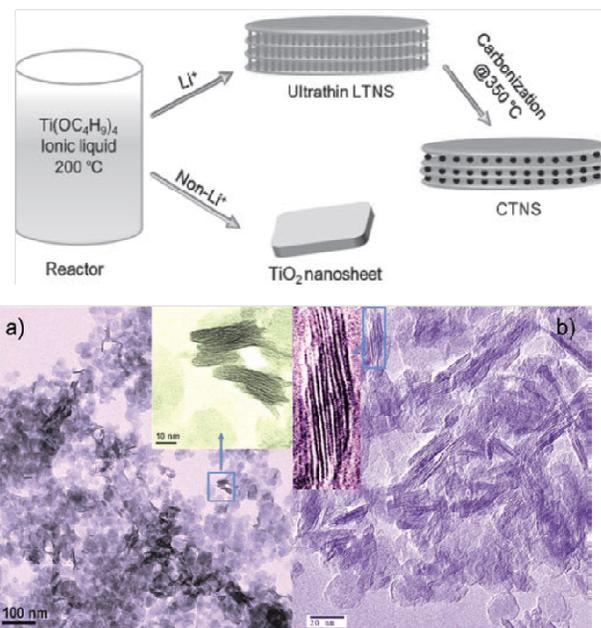
temperature and ease of morphology control. In fact, sol-gel remains the most commonly used method in fabrication of  $\text{TiO}_2$  (including doped  $\text{TiO}_2$ ) for the application of photocatalyst.<sup>57</sup> A generic routine to prepare nanocrystalline  $\text{TiO}_2$  sol gel formulations is to mix titanium tetraisopropoxide in acetic acid. The mixed acidic solution is further subjected to heating with vigorous stirring to undergo hydrolysis and condensation reaction with the aid of acetic acid to obtain the nanostructured  $\text{TiO}_2$ . The employment of sol gel titania in corporation with electrospinning was not exploited until 2003, when Xia's group introduced electrospinning to fabricate titania nanofibers.<sup>58</sup> Electrospinning is a typical organic fiber fabrication process where polymer solution is injected from a small nozzle under the influence of an electric field. A jet is formed due to the continuous accumulation of electrostatic charges on the surface of the solution droplet. The jet is then subjected to stretching to form ultrathin fiber. By mixing titanium tetraisopropoxide with poly(vinyl pyrrolidone) (PVP) in alcohol solution, Xia's group prepared  $\text{TiO}_2$ /PVP composite nanofibers with a uniform diameter of  $78 \pm 9$  nm. The composite nanofibers were further burned in air at  $500^\circ\text{C}$  to remove PVP residual, leaving titania scaffold nanofibers with a reduced diameter of  $53 \pm 8$  nm. Further modification on the electrospinning has enabled tailoring the nanofibers with core-sheath, hollow or porous structures.<sup>59</sup> This design relies on the incorporation of mineral oils loaded into the core capillary while the sol gel precursor is loaded into the sheath capillary,<sup>60</sup> as shown in Figure 15a. By extracting the mineral oils in the core and removal of PVP in the sheath, robust titania nanotubes were obtained, as shown in Figure 15b-d. Despite that the combination of sol-gel and electrospinning exhibits straightforward and versatile strategies for synthesizing 1D titania fibers, the fibers are usually randomly aligned. By incorporating recently reported nonelectrospinning spinneret-based tunable engineered parameters (STEP) technique with sol gel precursor, Wang et al<sup>61</sup> managed to fabricate aligned 1D titania structures including fibers, tubes, nanowires in microtubes. A solution of polystyrene (PS) and titanium tetraisopropoxide was utilized as the precursor. A rotating substrate was employed to pull out the solution filament from the extruded pendent solution droplet and stretch it into micro/nanoscale fibers. After hydrolysis and calcination procedure, highly porous titania hollow tubes with diameters of approximately 250 nm were obtained. Apart from titania, sol gel process has also seen significant application in fabrication of titanate thin films, such as sol gel derived lead-zirconate-titanate (PZT) ferroelectric thin film. Interestingly, study has revealed that the PZT thin films on sapphire substrates was able to complete the pyrochlore to perovskite transformation at  $650^\circ\text{C}$ . As a comparison, free-standing PZT films showed a much higher transformation temperature. The discrepancy in the behavior of free-standing sol gel derived titanate films versus titanate films on substrate can be related to the size effect of the thin film.<sup>62</sup>



**Figure 15.** (a) Schematic of the setup for electrospinning nanofibers with a core/sheath structure. (b) TEM image of two as-spun hollow fibers after the oily cores had been extracted with octane. The walls of these tubes were made of a composite containing amorphous anatase  $\text{TiO}_2$  and PVP. (c) TEM image of  $\text{TiO}_2$  hollow fibers that were obtained by calcining the composite nanotubes in air at  $500^\circ\text{C}$ . (d) SEM image of a uniaxially aligned array of anatase hollow fibers that were collected across the gap between a pair of electrodes.<sup>60</sup>

### 3.4 Formation of 2D titanate/titania nanosheets

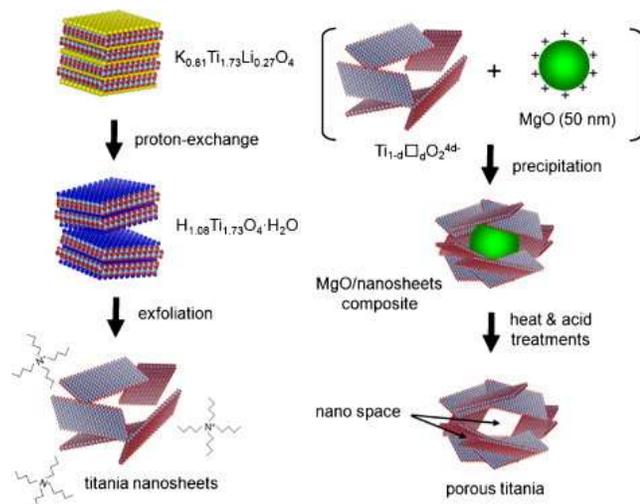
As mentioned above, titanate nanosheet is the intermediate during the hydrothermal process for the formation of the titanate nanotube and nanowire. Therefore, by precise control of the experimental condition, we can obtain the nanosheet morphology of the titanate materials. For example, titanate nanosheet was obtained when NaOH concentration is decreased to 5~8 M at  $140^\circ\text{C}$  for 48 h under hydrothermal conditions.<sup>63</sup> This special structure could also be obtained by organic-stabilizer-free synthesis,<sup>64</sup> continuous hydrothermal route (flow  $450^\circ\text{C}$  superheated water through a crystallizing medium),<sup>65</sup> room temperature synthesis,<sup>66</sup> and ionic liquid synthesis approach.<sup>67</sup> For example, the layered titanate nanosheets (LTNSs) and carbon-supported stacked  $\text{TiO}_2$  nanosheets (CTNS) obtained from the thermal treatment of LTNSs are obtained by ionic liquid synthesis approach via the scheme in Figure 16.<sup>67</sup> The typical morphology of LTNSs and CTNS is shown in Figure 16. The LTNSs have a lateral size of 20~50 nm, and layered interlamellar spacing is around ~1 nm. The disc-like nanostructure of CTNS obtained after  $350^\circ\text{C}$  annealing for 2 h is clearly evident in its high-resolution TEM image showing that several ultrathin nanosheets stack together.



**Figure 16.** The formation scheme of layered titanate nanosheets (LTNSs) and carbon-supported stacked  $\text{TiO}_2$  nanosheets (CTNS) with a sandwich-like multilamellar structure; TEM images of the (a) as-synthesized LTNSs and the (b) CTNSs obtained after annealing at  $350^\circ\text{C}$  for 2 h.<sup>67</sup>

On the other hand, the single graphene-like layered titania sheet can be prepared by delaminating a lepidocrocite-type layered protonic titanate.<sup>67, 68</sup> In this approach,  $\text{TiO}_2$  and  $\text{Cs}_2\text{CO}_3$  are calcined first to

form the Cs-titanate. Then the  $\text{Cs}^+$  is ion exchanged by  $\text{H}^+$ , and finally the titanate nanosheet is exfoliated in the presence of tetrabutylammonium hydroxide (TBAOH). Since the titania sheets possess a negative charge, the positive charge of the metal and metal oxide nanoparticles can be re-assembled with the titania sheet to form the porous nanohybrids.<sup>67, 69-73</sup> An example is shown in Figure 17, the exfoliation-precipitation method enables the construction of a new porous material through the exfoliation of the titanate sheets and through precipitation with MgO fine particles.<sup>35</sup> Autoclave is not required for this synthesis route, and ambient condition is sufficient for most of the stages. Nevertheless, this process suffers from some drawbacks, e.g. the need for multistage processing and the overall long processing time.

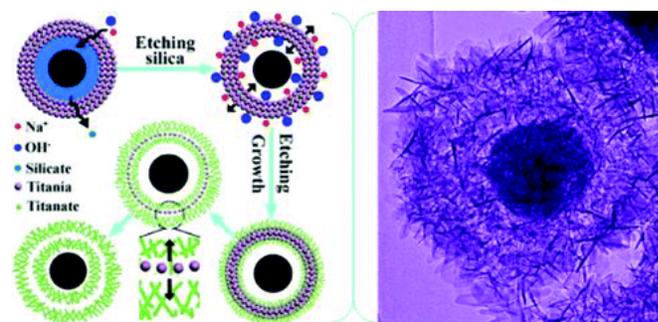


**Figure 17.** The synthesis route of titanate nanosheets by exfoliation method.<sup>35</sup>

### 3.5 Preparation of 3D titanate hierarchical materials

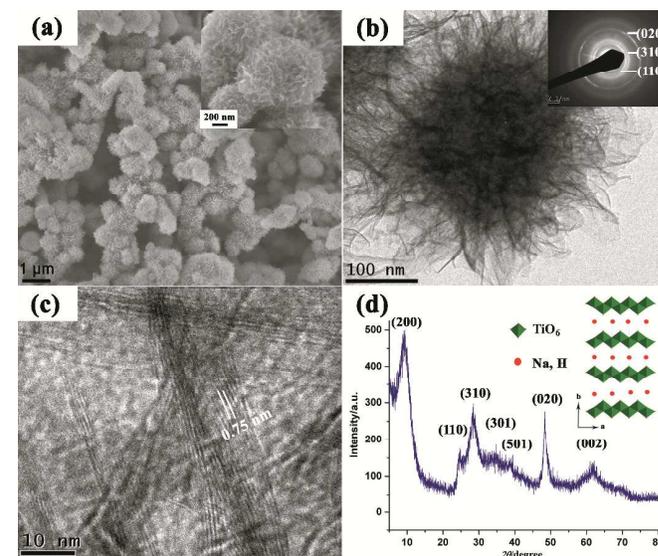
The synthesis of 1D and 2D nano-titanate has been widely investigated and well developed. However, multi-scale 3D hierarchically shaped titanate remained a challenge for many years and the synthesis is usually not straightforward. For examples, Cao et al.<sup>74</sup> synthesized titanate flowers via a template-assisted approach through hydrothermally reacting NaOH with the mixture of  $\text{TiO}_2$  and  $\text{Zn}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ , followed by the removal of ZnO flower-like nanorods framework. Yoshikawa et al.<sup>75</sup> reported a two-step method combining hydrolysis of titanium tetraisopropoxide with hydrothermal treatment of amorphous  $\text{TiO}_2$  spheres to prepare spherical titanate nanosheet. Imai's group<sup>76</sup> systematically explored various kinds of hierarchical ammonium titanate nanosheet morphologies grown in the agar gel medium by a bottom-up route. Mao et al.<sup>77</sup> demonstrated the growth of micrometer scale sea-urchin-like structures by the oxidizing  $\text{H}_2\text{O}_2$  assisted-hydrothermal method in autoclave at different temperatures and durations, and they proposed that a two-stage ("growth-then-assembly") growth mechanism for the formation of titanate spheres. The self-assembled titanate nanosheets can also be synthesized via chimie-douce method by refluxing  $\text{TiO}_2$  powder in 15 M NaOH at 423 K for more than 24 h.<sup>78</sup> Xie's group used the Kirkendall effect to construct 3D hollow titanate tubular hierarchical structures via treating  $\text{TiCl}_4$  precursor in other alkaline solutions (e.g., ethylenediamine) at 225 °C for 12 h.<sup>79</sup> Moreover, Zhao's group reported a hydrothermal etching assisted crystallization route to synthesize  $\text{Fe}_3\text{O}_4$ @titanate yolk-shell

microspheres with ultrathin nanosheets-assembled double-shell structure (Figure 18).<sup>80</sup>



**Figure 18.** Formation scheme and TEM image of the hierarchical titanate microspheres by hydrothermal method.<sup>80</sup>

Inspired by these works, we developed a novel one-pot approach to produce highly porous hierarchical 3D titanate micro-spherulite (TMS) particles (Figure 19) via a simple, high throughput method employing simultaneously electrochemical anodization and spark discharge of the anodized oxide layer into a reactive solution by carefully adjusting the applied electrical spark parameters in a traditional ambience setup.<sup>81, 82</sup> Electrochemical spark discharge spallation (ESDS) process is led by the concurrent anodic reaction oxidation and the electrical breakdown of the formed oxide layer into the electrolyte in precipitate form. The spallation is driven by a continuous spark discharge that simultaneously heat up the solution. The  $\text{TiO}_2$  precipitates then react immediately with the heated NaOH solution to form the TMS.

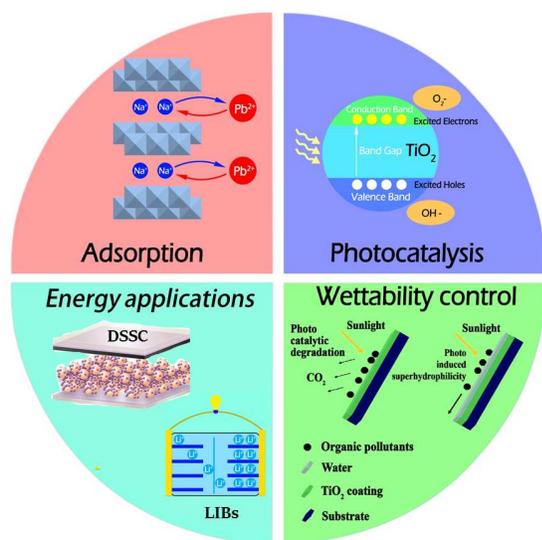


**Figure 19.** (a) FESEM image, (b) TEM image, (c) high resolution TEM image, (d) XRD pattern of the fabricated TMS. The TMS powder is collected after anodization of Ti foils in 10 M NaOH aqueous solution for 20 min. Inset shows the crystal structure for lepidocrocite titanates.<sup>81</sup>

### 4. Application of Titanate/Titania Nanomaterials

Regarding this special crystal structure with layered sheets, significant attention has been paid in the field of photon-mediated water decomposition,<sup>83</sup> photocatalysis,<sup>84</sup> fuel cell electrolytes<sup>85</sup>,

adsorbent,<sup>86</sup> lithium ion batteries (LIBs),<sup>87</sup> dye-sensitized solar cell (DSSC) and wettability control related applications owing to their excellent ion-exchange/intercalation activities, as well as adsorption/photocatalytic properties. In the following sections, the titanate and titania crystal structure and surface property related applications will be discussed in detail, which are summarized in Figure 20.



**Figure 20.** Titanate and titania nanomaterials for environmental and energy applications.

#### 4.1 Adsorbent application

Adsorption is an important process in wastewater treatment, because it does not only encompass the physical removal of pollutants from water, but also is a crucial step in photocatalysis.<sup>31, 88-90</sup> During this process, pollutant molecules are adsorbed to the surface of the adsorbent. To understand the adsorption mechanisms governing titania and titanate nanostructures, the equilibrium adsorption isotherms can be fitted to two widely-used models: the Langmuir and the Freundlich equations:

$$\text{Freundlich: } q = K_F C^n \quad (3)$$

$$\text{Langmuir: } q = \frac{q_m K C}{1 + K C} \quad (4)$$

where  $q$  ( $\text{mg g}^{-1}$ ) is the amount of adsorbed pollutant,  $C$  ( $\text{mg L}^{-1}$ ) is the concentration of pollutant at equilibrium,  $q_m$  ( $\text{mg g}^{-1}$ ) is the maximum adsorption capacity,  $K_F$  and  $n$  are Freundlich constants, and  $K$  is the Langmuir constant. The adsorption mechanism is monolayer if the adsorption isotherm exhibits Langmuir model. Conversely, a good fit with the Freundlich model indicates a heterogeneous surface binding. The amount of pollutant adsorbed by the adsorbent,  $q$ , is calculated from the difference between the initial and the final pollutant concentration as:

$$q = \frac{(C_0 - C_f)V}{M} \quad (5)$$

where  $q$  is the amount of pollutant adsorbed ( $\text{mg g}^{-1}$ ),  $C_0$  and  $C_f$  are the initial and final concentrations of pollutant remaining in the

solution ( $\text{mg L}^{-1}$ ),  $V$  is the volume of the solution containing the pollutant (L), and  $M$  is the mass of the adsorbent added (g).

To investigate the rate at which the pollutant is adsorbed on the nanostructures, it is useful to employ the pseudo-first-order kinetic model and the pseudo-second-order kinetic model. The pseudo-first-order kinetic model can be expressed in its integral form as:

$$\ln(Q_{e1} - Q_t) = \ln Q_{e1} - k_1 t \quad (6)$$

where  $Q_{e1}$  and  $Q_t$  are the amount of pollutant adsorbed ( $\text{mg g}^{-1}$ ) on the adsorbent at equilibrium and at time  $t$ , respectively,  $k_1$  is the rate constant for the pseudo-first-order kinetic model ( $\text{min}^{-1}$ ), and  $t$  the contact time (minutes). The values of  $\ln(Q_{e1} - Q_t)$  were calculated from kinetic data.

Similarly the pseudo-second-order kinetic model can be expressed in its integral form as:

$$\frac{t}{Q_t} = \frac{1}{k_2(Q_{e2})^2} + \frac{t}{Q_{e2}} \quad (7)$$

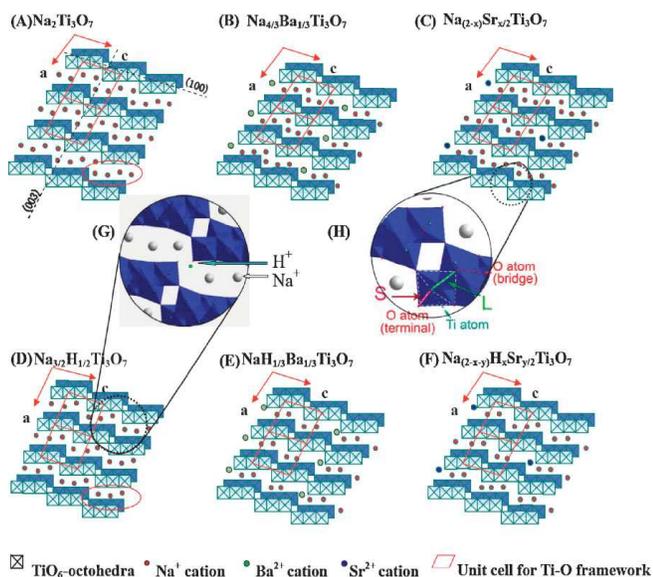
where  $k_2$  ( $\text{g mg}^{-1}\text{min}^{-1}$ ) is the rate constant for the pseudo-second-order kinetic model, and  $Q_{e2}$  the equilibrium amount of pollutant adsorbed ( $\text{mg g}^{-1}$ ).

In view of the crystal structure of titanate materials, sodium ( $\text{Na}^+$ ) or proton ( $\text{H}^+$ ) occupy the inter-layer cavities between the  $\text{TiO}_6$  octahedral sheets. The cations between the layers are exchange-able and titanate layer structure is stable during the ion-exchange process. For example, the interlayer distance of multilayered titanate nanotubes are not significantly altered when alkaline ions ( $\text{Li}^+$ ,  $\text{Na}^+$ ,  $\text{K}^+$ ,  $\text{Rb}^+$ ,  $\text{Cs}^+$ )<sup>91</sup> or some transition-metal ions ( $\text{Cu}^{2+}$ ,  $\text{Co}^{2+}$ ,  $\text{Cd}^{2+}$ ,  $\text{Zn}^{2+}$ ,  $\text{Ni}^{2+}$ )<sup>92</sup> are intercalated between the layers, demonstrating the rigidity of the titanate lattice towards ion-exchange. For the specific case of sodium ions, the maximum ion-exchange capacity, defined as the ratio between the atomic percent of sodium and titanium in the nanotubes (Na/Ti ratio), was reported to vary between 0.67<sup>93</sup> and 1.1.<sup>94</sup> The ion-exchange capacity is crystal structure dependent. For example, in an idealized crystal structure of trititanate<sup>34</sup>, complete ion exchange of protons in the titanate to sodium ions will give a theoretical value of 0.67 according to the reaction:



Very recently, 1D titanate nanowires<sup>95, 96</sup> and nanotubes<sup>86</sup> were shown to have excellent adsorption ability for the removal of organic molecules and radioactive toxic metal ions. According to Zhu's work,<sup>95, 96</sup> the radioactive cations ( $\text{Sr}^{2+}$  and  $\text{Ba}^{2+}$ ) after the ion exchange were trapped in the titanate nanofibers permanently due to the structural change, as shown in Figure 21. This capability of trapping radioactive cations permanently allows us to isolate them from contaminated water. Thus, the titanate nanowire adsorbents do not have the issue of disposal concern regarding the risk of secondary contamination, in which the adsorbed cations are released from the adsorbents. The titanate nanotubes uptake of the inorganic arsenic (As) is efficient in terms of rate and effective in terms of capacity. With the aid of NaOH solution, the adsorbed arsenic in the titanate is reported to be able to desorb quickly.<sup>86</sup> The reversible ion-exchange process is mainly dependent to the metal ion concentration, which drives the ion-exchange direction when the diameter of metal ion is small enough to insert into the layered distance of titanate structures. Moreover, the synthesized titanate

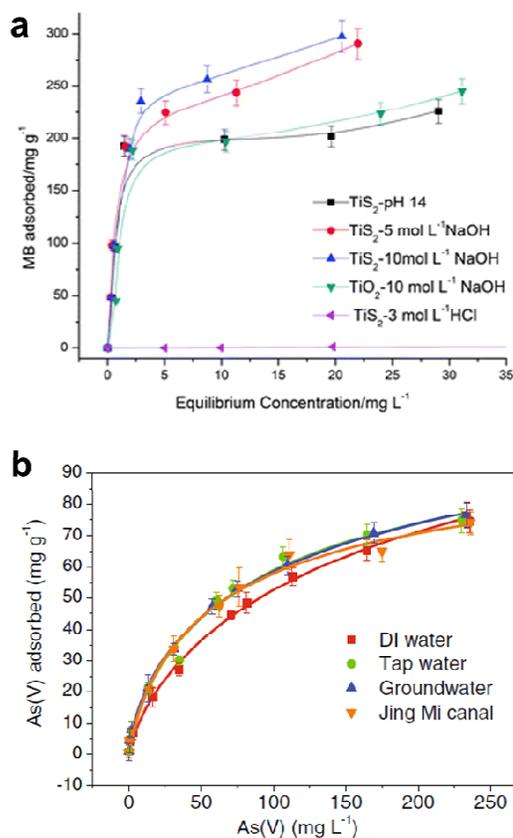
nanotube showed an excellent adsorption properties to the different organic dyes, including basic green 5, basic violet 10 and methylene blue through an ion-exchange mechanism or electrostatic force between the dye and the titanate material at certain conditions.<sup>97</sup> According to the crystal structure of titanate materials, alkali metal or proton ions ( $H^+$ ) in the  $TiO_6$  octahedral layers are exchangeable and the titanate layer structure is stable during ion-exchange process. Therefore, the different valence of metal ions ( $Co^{2+}$ ,  $Ni^{2+}$ ,  $Cu^{2+}$ ,  $Cd^{2+}$ ,  $Fe^{2+}$ ,  $Cu^{2+}$ ,  $As^{n+}$ ,  $Pb^{2+}$ , etc.)<sup>37, 92, 98-100</sup> between the layers in the titanate materials can be exchanged with alkali metal or proton ions ( $H^+$ ,  $Li^+$ ,  $Na^+$ ,  $K^+$ ,  $Rb^+$ ,  $Cs^+$ ).<sup>91</sup>



**Figure 21.** Structure of titanate nanowires made for removing radioactive ions from water.<sup>95</sup>

The adsorption mechanisms and kinetics are mainly dependent on the phases of  $TiO_2$  and titanate nanostructures and the type of pollutants. It has been widely reported that the adsorption of oxalic acid on Degussa P25 follows Langmuir model.<sup>88, 101, 102</sup> Similarly, the adsorption mechanisms of various titanate nanostructures also follow closely to the Langmuir model.<sup>32, 86, 101, 103-106</sup> Lim et al. synthesized sodium titanate nanobelts and nanotubes via hydrothermal synthesis using  $TiO_2$  and  $TiS_2$  as precursors respectively, and discovered that while both nanostructures follow the Langmuir model in the adsorption of methylene blue (MB) (Figure 22a), the  $TiS_2$ -derived nanobelts exhibit much higher adsorption capacity ( $312.50 \text{ mg g}^{-1}$ ) than the  $TiO_2$ -derived nanotubes ( $256.41 \text{ mg g}^{-1}$ ).<sup>42</sup> The dual phase anatase/titanate nanoparticles reported by Cheng et al.<sup>32</sup> also follow the Langmuir model in the adsorption of MB with a capacity of  $162.19 \text{ mg g}^{-1}$ . It was also reported that the removal of MB by anatase-covered titanate nanotubes follow the Langmuir model with a capacity of  $199.6 \text{ mg g}^{-1}$ , and that the removal of methyl orange is mostly attributed to photodegradation.<sup>89</sup> Titanate nanotubes were also employed in the adsorption of heavy metals, such as thorium<sup>103</sup>, chromium,<sup>104</sup> and arsenic (Figure 22b)<sup>86</sup> and lead.<sup>81</sup> While the adsorption of both thorium and chromium follow the Langmuir model,<sup>103, 104</sup> interestingly, the adsorption of arsenic by protonated titanate nanotubes synthesized via hydrothermal synthesis fits very well with both the Langmuir and Freundlich models.<sup>86</sup> The morphology dependence of titanate nanostructures (nanotubes, nanowires and nanorods) on the adsorption of formic acid was also investigated, and it was shown that the adsorption capacity of the titanate

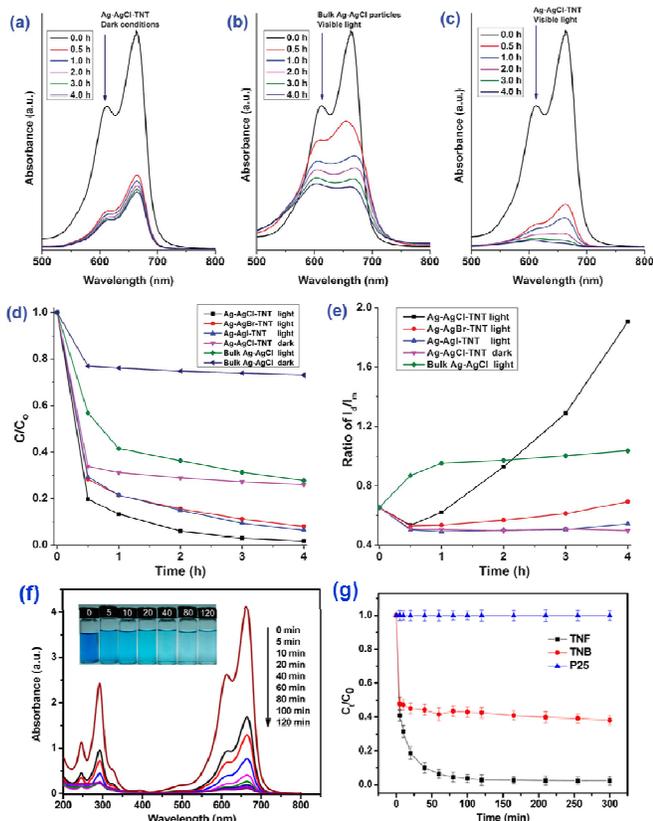
nanotubes calcined at  $400 \text{ }^\circ\text{C}$  exhibits the highest capacity following the Langmuir adsorption model.<sup>105</sup>



**Figure 22.** Adsorption isotherms of various titania and titanate nanostructures in MB. The y-axis displays the amount of MB adsorbed by the nanostructures, the x-axis displays the equilibrium concentration of MB.<sup>42</sup> Langmuir isotherms for (a) As (V), and (b) As (III) adsorption on titanate nanotube in several real water samples (pH 7.0,  $1.0 \text{ g L}^{-1}$ ).<sup>86</sup>

It is useful to carry out adsorption kinetics tests to investigate the rate of adsorption of the pollutant by the various titania and titanate nanostructures. It has been observed the MB blue concentration decreased rapidly in the first 30 minutes for various titanate nanostructures, including sodium titanate nanoflowers<sup>107</sup> and Ag-AgCl anchored on titanate nanotubes,<sup>108</sup> which is attributed to the strong electrostatic attraction forces between MB and the negatively-charge titanate nanostructures.<sup>109, 110</sup> For example, the Ag-AgX-titanate nanotubes hybrid (Ag-AgCl-TNT) with a high surface area can synergistically adsorb and degrade methylene blue (MB) under visible light, viz., the titanate nanotubes adsorb MB onto their surfaces, which aids the degradation of MB by Ag-AgX nanoparticles (Figure 23a-e). The excellent degradation performance of Ag-AgX under visible light is originated from its localized surface plasmonic resonance effect. The kinetics of adsorption of MB appears to be independent of morphology, as adsorption of MB by hydrothermally-synthesized sodium titanate nanobelts,<sup>42</sup> titanate nanotubes<sup>106</sup> and sodium titanate nanoflowers (Figure 23f-g)<sup>107</sup> all follow the pseudo-second-order kinetic model. In the case of heavy metals, it was reported that the adsorption of thorium (Th) by  $TiO_2$ -titanate nanotubes follows the pseudo-second-order kinetic model.<sup>103</sup> For the photocatalytic degradation of chromium (Cr) with titanate

nanotubes, the synergistic process of chromium removal was found dependent on pH: adsorption of Cr (III) is promoted when  $\text{pH} < 5$ , while  $\text{pH} > 5$  promotes Cr (IV) reduction on  $\text{TiO}_2$ .<sup>104</sup> Similarly, the adsorption and photocatalytic reaction of basic violet 10 (BV-10) dye on Co-doped titanate nanotubes also match well with the pseudo-second-order kinetic model, with the adsorption constant  $k_2$  being one order of magnitude higher than the photocatalytic rate constant. This implies that the photocatalysis of BV-10 dye is the rate-determining step.<sup>111</sup>

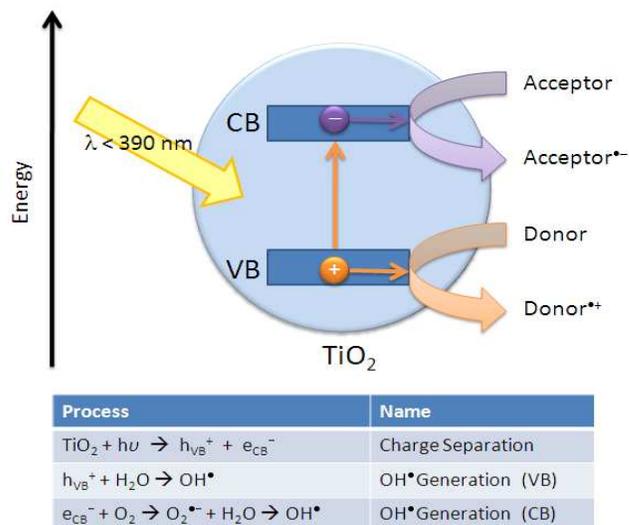


**Figure 23.** Methylene blue (MB) aqueous solution absorption spectra after treatment using (a) Ag-AgCl-TNT under dark conditions, (b) bulk Ag-AgCl particles under visible light, and (c) Ag-AgCl-TNT under visible light for different durations. (d-e) Adsorption and visible light degradation performance of different silver/silver halide based photocatalysts/conditions for MB removal.<sup>108</sup> (f) Absorption spectra of MB solution ( $20 \text{ mg L}^{-1}$ ,  $50 \text{ mL}$ ) in the presence of sodium titanate nanoflowers ( $20 \text{ mg}$ ) under dark conditions. Inset represents the photograph of MB solutions at varying times. (g) Adsorption of MB ( $20 \text{ mg L}^{-1}$ ,  $50 \text{ mL}$ ) by sodium titanate nanoflowers, sodium titanate nanobelts and P25.  $C_0$  is the initial concentration of the MB solution, and  $C$  is the concentration of that at different intervals during the adsorption. Error bars show standard deviations from two independent experiments, while each experiment was performed in triplicate and the average of concentrations was taken.<sup>107</sup>

#### 4.2 Photocatalytic application

The mechanism of photocatalytic degradation (Figure 24) of organic compounds is similar to the  $\text{TiO}_2$ -based photocatalysts. When the ultraviolet (UV) radiation photons from sunlight or illuminated light source are absorbed by the photocatalyst, it will produce electron-hole pairs. The electrons initially rested in the valence band of

titanium dioxide are excited by the photon energy, and are promoted into the vacant conduction band of titanium dioxide. The resultant positive holes in the conduction band of titanium dioxide can then attack water molecule to generate hydroxyl radical ( $\text{OH}\cdot$ ), while the negative-electrons in the conduction band can attack oxygen molecule to generate peroxide anion. The generation of surface  $\text{OH}^-$  groups and peroxide anion are required for photocatalytic degradation of organic compounds.



**Figure 24.** Principles of oxidative decomposition of photocatalysts.<sup>112</sup>

Titanate materials are semiconductors with a bandgap of  $3.4 \sim 3.7 \text{ eV}$ . The area of photocatalysis is one important application that these titanate materials are suited for, but the literature tackling this aspect of application has been limited. Uncertainties exist over the photocatalytic properties of the titanate materials. For example, a few researchers insisted that enough potentials are possessed by the as-prepared sodium- or hydrogen-titanates nanotubes towards the degradation of the organic pollutants.<sup>113-117</sup> Recent work indicated that the photocatalytic activity of as-prepared titanate materials (nanotubes, nanowires, nanosheet) was not as good as the commercial standard P25 catalyst when they were tested for oxidation of  $\text{NH}_3$ <sup>48</sup> and organic dyes<sup>118, 119</sup> in aqueous suspensions. Either sodium impurities in as-prepared titanate or their moderate crystallinity have been proposed to be the cause of the observed low activity. It is known that recombination centered created by  $\text{Na}^+$  impurities can lower the photocatalytic activity of  $\text{TiO}_2$ .<sup>120</sup> After alkaline hydrothermal reaction, the amount of sodium ions being intercalated into the titanate crystal is extremely high; the photocatalytic activity of the as-prepared titanates is significantly affected. The negative correlation has been confirmed recently between sodium content in the titanate and their photocatalytic dye oxidation activity.<sup>121, 122</sup> Luminescence quenching, which removes sodium ions in the structure, was observed after the protonation of titanate nanotubes.<sup>123, 124</sup> On the other hand, some researchers hold the opinion that neither sodium nor hydrogen titanate nanotubes are efficient photocatalyst due to their intrinsic inertness for photocatalytic reactions.<sup>125-128</sup> Researches have shown that no improvement in photocatalytic activity was observed when acid was used to ion-exchange the as-prepared titanate samples. However, calcination was found to have led to apparent increase in photocatalytic activity for these titanate nanotubes. Further complication arises due to the scatter on the reported photocatalytic

activity comparison experiment. For example, a 3 times better photocatalytic activity than commercial Degussa P25 was reported by Yu et al.<sup>127</sup> on the calcined samples. However, Zhang et al.<sup>122, 126</sup> claimed that, despite the large range of calcination temperatures they have tried, the resultant photocatalytic activity of the nanotubes samples are still inferior to the commercially available powders. The inconsistency among existing reports may be due to the different preparation and test conditions from different groups.

Titanium dioxides, derived from nano-structured layered titanates, have attracted great attention during the past decade. Titanate materials provide a good platform for delicate phase- and morphological tailoring, by annealing<sup>37, 129</sup> or wet chemical reaction<sup>115, 130</sup>, to obtain various TiO<sub>2</sub> or controlled mixed-phase nanostructures. Lots of effort has been made to improve the performance of TiO<sub>2</sub>-based materials during the past four decades, with the hope to bring it into industrial scale application. Some modest gains have been achieved so far in spite of the complexity and difficulty of the task. The main approaches could be sub-classified into three groups (as illustrated next), although more than one method might be used concurrently to modify the pristine titanium dioxide.

#### 4.2.1 Development of pristine TiO<sub>2</sub> based active photocatalysts

##### Optimizing the crystallinity, surface area and exposed facets:

The photocatalytic activity of four titanium dioxide polymorphs has been reported before, namely anatase, rutile, brookite and TiO<sub>2</sub>(B). Their crystal structures are discussed earlier. The information, together with the band gap and density are summarized in Table 2.<sup>131, 132</sup>

**Table 2.** Bulk properties of TiO<sub>2</sub> in its different phases<sup>18, 131, 132</sup>

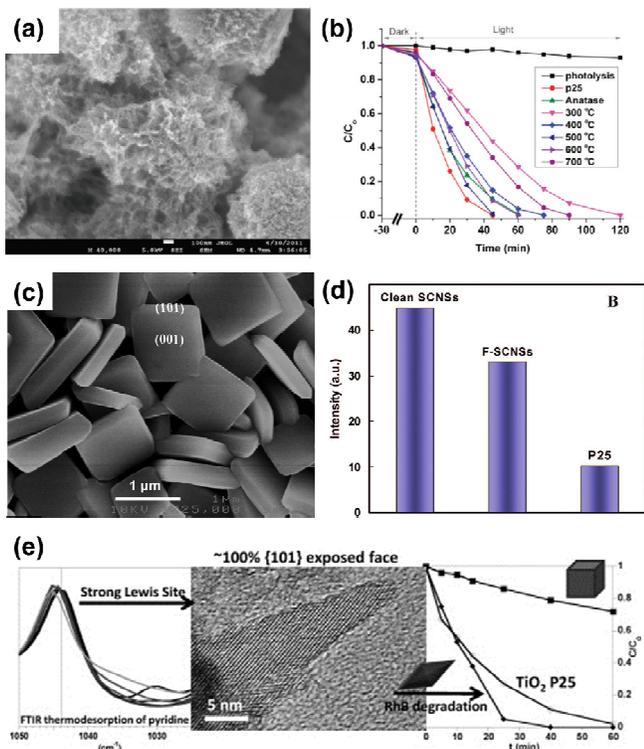
Crystal Structure	Crystal System	Bandgap (eV)	Density (g cm <sup>-3</sup> )
Rutile	Tetragonal	3.00	4.13
Brookite	Orthorhombic	3.30	3.99
Anatase	Tetragonal	3.20	3.79
TiO <sub>2</sub> (B)	Monoclinic	3.20	3.64

Anatase and rutile TiO<sub>2</sub> have been intensively studied for photocatalysis, in part due to their ease of preparation. Of the two common forms, anatase has a conduction band minimum sufficiently negative to reduce protons. Its photoactivity is also generally superior to that of rutile. TiO<sub>2</sub>(B), as a relative new phase, was first synthesized by Marchand et al. in 1980,<sup>133</sup> and have since attracted lots of attention as an alternative photocatalyst due to its relative open structure, although its photoactivity is generally lower than that of anatase TiO<sub>2</sub>. Furthermore, a few groups<sup>18, 134, 135</sup> reported impressive photocatalytic activity of brookite TiO<sub>2</sub>, even better than known active anatase TiO<sub>2</sub> for hydrogen production.<sup>18, 135</sup>

As a special type of heterogeneous catalyst, in principle photocatalysts also benefit from similar textural effects, e.g., dispersion into smaller particles increases the specific surface area, thereby promoting faster and stronger adsorption of reactants at the interface, ultimately leading to higher rates of reaction. Problems of

recombination of photo-generated charges can be mitigated by enhancing the crystallinity of the photocatalyst. Some improvements have been achieved by fabricating it in higher surface area form while maintaining a high degree of crystallization on the nano-scale with different morphologies, such as nanoparticle, nanotube, nanowire, nanoflake, etc.<sup>136-139</sup> A hybrid TiO<sub>2</sub> nanoflake/nanoparticle hierarchical structure formed at 500 °C was found to exhibit a higher photocatalytic activity for the degradation of organic compounds (Figure 25a-b).<sup>138</sup> Unfortunately, this approach is generally difficult because the above attributes are mutually antagonistic and often self-cancelling. A process that most readily increases the crystallinity through thermal treatment inexorably leads to surface area loss due to coarsening, thus dictating a practical compromise between these two factors. More recently, some success was achieved by synthesizing titanium dioxide with a high percentage of active facets.<sup>140-148</sup> For anatase TiO<sub>2</sub>, both the theoretical prediction and experimental results show that its (001) facets are more active than the (101) facets, however, the latter normally dominates the surfaces in the equilibrium state. By fluorine-termination of anatase TiO<sub>2</sub>, the desired (001) facets can comprise up to 50% of the surface.<sup>140</sup> Figure 25c shows the representative scanning electron microscope image of the highly truncated bipyramidal anatase TiO<sub>2</sub> single-crystal nanosheets. Statistically, the synthesized single-crystal nanosheets (SCNSs) have 64% the (001) facets and display superior photoreactivity (more than 5 times) compared to P25 as a benchmarking material (Figure 25d).<sup>147</sup> Also, the coexistence of low-index facets with a highly photoactive (001) facet in anatase TiO<sub>2</sub> nanocrystals has been found beneficial to enhance the photocatalytic performance of TiO<sub>2</sub> via a synergistic effect. Dufour et al.<sup>149</sup> fabricated four different morphologies of pure anatase nanoparticles with exposed surfaces and studied the impact of the exposed surfaces on the photocatalytic efficiency. They found that the most efficient photocatalyst for rhodamine B degradation is the (101) facets TiO<sub>2</sub>, which possesses the stronger acidic surface sites. Recently, our group found that this synergistic effect can be extended from a single crystal to interconnected nanocrystals with dominating (001) and (010) facets in close contact with each other in a hierarchically structured porous particle form.<sup>150</sup> The two faceted building blocks both belong to anatase TiO<sub>2</sub>, but they display different band gap values and band potential positions due to the dominating crystal planes. As a result, photo-generated electrons and holes can transfer from one to the other, giving rise to effective charge separation as commonly observed in a two-semiconductor heterojunction. The particles synthesized at the optimal condition showed outstanding photocatalytic hydrogen production of 364.2 μmol g<sup>-1</sup> h<sup>-1</sup>, which is about four times as much as that of commercial P25 (96.5 μmol g<sup>-1</sup> h<sup>-1</sup>).<sup>150</sup> Although the promising properties of anatase TiO<sub>2</sub> with exposing specific surfaces have been widely investigated, a clear assessment of the role of the crystal faces in photocatalytic processes is still under debate. Recently, D'Arienzo et al.<sup>151</sup> revealed that the concentration of trapped holes (O<sup>-</sup> centers) increases with increasing the (001) surface area and photoactivity, while the amount of Ti<sup>3+</sup> centers increases with the specific surface area of the (101) facets in vacuum conditions, and the highest value occurs for the sample with the worst photooxidative efficacy. These results suggest that (001) surfaces can be considered essentially as oxidation sites with a key role in the photooxidation, while the (101) surfaces provide reductive sites which do not directly assist the oxidative processes. Based on this work, they also demonstrated that the correlation of the reactivity on the type of crystal facets and photogenerated defects.<sup>152</sup> The electron spin resonance spectra reveal that the amount of Ti<sup>3+</sup> (electron traps) is parallel to the H<sub>2</sub> evolution rate and becomes a maximum for the nearly rectangular TiO<sub>2</sub> nanocrystals, which

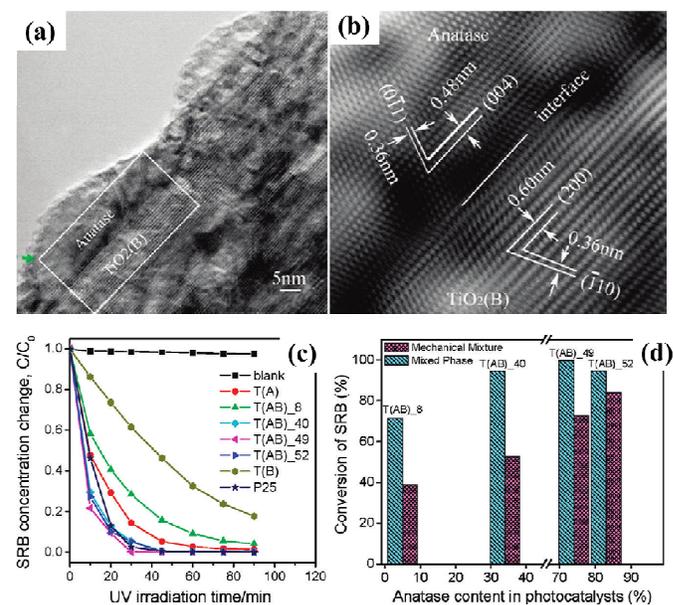
display the highest area of the (010) surfaces and lowest (010) area but also involve a significant fraction of the (010) facets.



**Figure 25.** (a) SEM images of hybrid nanoflake/nanoparticle hierarchical structure and (b) its photocatalytic activity;<sup>138</sup> (c) Typical SEM images of the anatase TiO<sub>2</sub> single-crystal nanosheets (SCNSs) and (d) normalized fluorescence intensity per unit surface area with different photocatalysts: clean SCNSs, TiO<sub>2</sub> SCNSs capped by F atoms (denoted as F-SCNSs), and Degussa P25 TiO<sub>2</sub>.<sup>147</sup> (e) A correlation between the nature of the exposed surfaces, the corresponding surface acidic behavior. Photocatalytic degradation curves of the RhB solution by the anatase samples with (101) exposed facet and the P25 reference.<sup>149</sup>

**Development of TiO<sub>2</sub> based mixed-phase photocatalyst:** Although the relative photocatalytic efficiency of anatase and rutile TiO<sub>2</sub> is still under debate, it is now well-established that mixed-phase photocatalysts show enhanced performance as compared to either single phase. The success of Degussa P25 TiO<sub>2</sub>, the benchmark widely used for comparison of other photocatalysts, is a prime example though it is not purposely designed for such benefit. Commercial Degussa P25 is made by the “aerosil process”, yielding individual particles consisting of roughly 20% rutile and 80% anatase.<sup>153</sup> Its high performance is generally attributed to the abundance of “rectifying” phase junctions that facilitate unidirectional (and opposite) flow of free carriers, thereby facilitating charge separation.<sup>154</sup> However, the nature of the atomic level contact between anatase and rutile is still not well understood,<sup>155</sup> and it has been demonstrated that the distribution of phases can be further improved for photocatalytic action by controlled annealing.<sup>156</sup> Based on the above principle, anatase and TiO<sub>2</sub>(B) core-shell structured nanofibers have been synthesized by hydrothermal treatment of titanate nanobelts, and their efficiencies were found to be comparable to that of commercial Degussa P25 (Figure 26).<sup>129, 157</sup> Highly crystalline pure brookite and two-phase anatase/brookite TiO<sub>2</sub> nanostructures were also successfully synthesized via a simple hydrothermal method with titanium sulfide

as the precursors in sodium hydroxide solutions, and the similar synergy of two phase system has been confirmed.<sup>18</sup>



**Figure 26.** (a-b) HRTEM images of the joint of anatase nanocrystal and TiO<sub>2</sub>(B) core in a single TiO<sub>2</sub> fiber; (c) Photocatalytic decomposition of sulphorhodamine dye with different fibril TiO<sub>2</sub> photocatalysts under UV irradiation; (d) Comparison of conversion rate for the decomposition of SRB with mechanically mixed TiO<sub>2</sub> photocatalysts.<sup>129</sup>

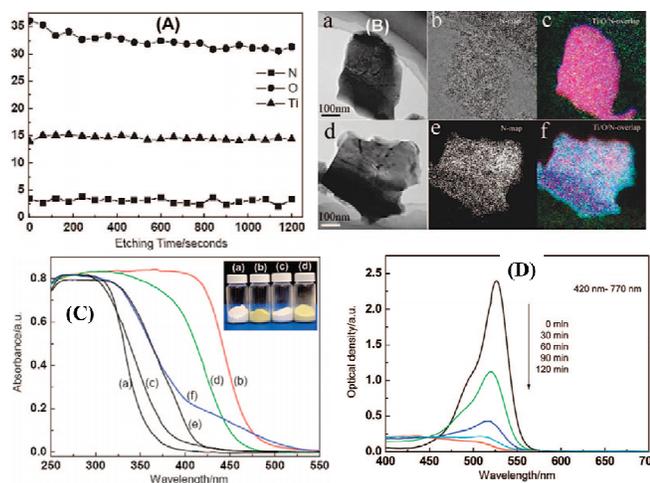
#### 4.2.2 Development of visible light active photocatalysts

While TiO<sub>2</sub> is an excellent and by far the most widely used photocatalyst, it suffers from an intrinsic drawback due to its large band gap (3.2 eV for anatase TiO<sub>2</sub>). Such a wide band gap only allows utilization of 4~5% solar irradiation in the ultraviolet region (UV light).<sup>158-160</sup> For better use of solar energy, a key topic in photocatalysis research now is to confer visible absorption, either by shifting the absorption edge suitably, or by sensitizing TiO<sub>2</sub> for visible photo-activity with surface additives. Several commonly adopted visible light sensitization strategies are discussed next.

**Bulk doping of TiO<sub>2</sub>:** The optical response of any material is determined by its intrinsic electronic structure, which is a strong function of chemical composition, crystal structure, and, to some extent, physical dimension (quantum size effects). The chemical composition of TiO<sub>2</sub> can be altered either by replacing the cation (titanium) or the anion (oxygen), referred to here as metal and non-metal doping, respectively.<sup>161-163</sup> Historically, the most popular approach has been explored by doping with color centers, such as transition metal cations, but with mixed results. When TiO<sub>2</sub> is doped with multivalent transition metal ions, such as Co<sup>II</sup>, Ni<sup>II</sup>, Cu<sup>II</sup>, Fe<sup>II/III</sup>, Cr<sup>III</sup>, V<sup>III</sup>, Mn<sup>IV</sup>, Nb<sup>V</sup>, Mo<sup>V/VI</sup>, an electron occupied level will be formed and the electrons are localized around each dopant.<sup>131</sup> As the dopant concentration increases, the localized level shifts to lower energy and can even reach the top of valence band for Co<sup>II</sup>.<sup>162</sup> Electrons can be excited from this defect state to the conduction band by photons with a lower energy, i.e., in the visible range. Band gap narrowing of TiO<sub>2</sub> can also be achieved by non-metal doping, using anions such as N, C, B, F, P, S and I.<sup>131, 163-168</sup> N and C doping are the two most investigated cases. In the pioneering work by Asahi et al.,<sup>163</sup> substitutional doping of N (for O) was proposed as the most

effective band gap narrowing due to mixing of p states between N and O. However, accumulating evidence suggests that N-doping introduces mid-gap levels slightly above the O 2p valence band.<sup>131</sup> For C-TiO<sub>2</sub>, the dopant introduces deep states in the gap.<sup>166, 167</sup> The intrinsic defects of TiO<sub>2</sub>, such as reduced Ti species and oxygen vacancies, also lead to the formation of localized states which may merge to form sub-band gap level, thus creating a lower energy excitation pathway.<sup>131, 169</sup>

Titanate, as an alternative starting material with relative open structure, is ideal for high concentration homogenous doping. In contrast with most reported nitrogen-doped titanium dioxide photocatalysts with some localized states in the intrinsic band gap and small visible light absorption shoulders induced by inhomogeneous nitrogen doping near the particle surface, the homogeneous substitution of O by N in all particles of layered titanates exhibited extraordinary band-to-band excitation in visible-light ranging up to blue light (Figure 27).<sup>161</sup> The actual photocatalytic efficiency of doped TiO<sub>2</sub>-based material is largely dependent on dopant concentration and homogeneity, which are highly dependent on the preparation method. Although doping with either metal or nonmetal elements definitely confers visible light absorption and associated photo-activity, the overall performance under UV-Vis irradiation may even fall below that of pristine TiO<sub>2</sub> due to deterioration of crystallinity and/or creation of particular defect sites that tend to promote charge recombination.

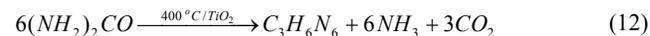
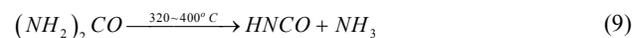


**Figure 27.** (A) XPS depth profiles of the elements of Ti, O and N in the H<sub>0.68</sub>Ti<sub>1.83</sub>O<sub>4-x</sub>N<sub>x</sub> upon Ar<sup>+</sup> sputtering. (B) Energy-filtered TEM images of (a, d) general morphology, (b, e) N map, and (c, f) Ti/O/N overlapped maps of (a-c) Cs<sub>0.68</sub>Ti<sub>1.83</sub>O<sub>4</sub> and (d-f) Cs<sub>0.68</sub>Ti<sub>1.83</sub>O<sub>4-x</sub>N<sub>x</sub>, respectively; (C) UV-visible light absorption spectra, (D) Comparison of photocatalytic activity for different photocatalysts.<sup>161</sup>

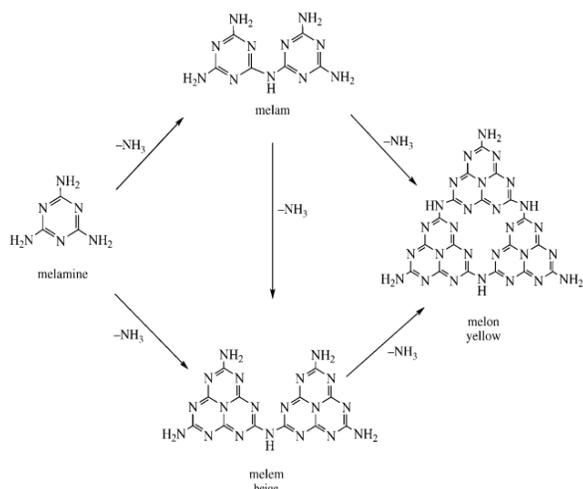
**Surface modification with melon and carbon nitride:** While the evidence has been steadily accumulated that visible light activity in TiO<sub>2</sub> results from introducing N, bulk-doping is a complex process, and the coloration due to the sole presence of N is not easy to prove because oxygen vacancies are created simultaneously to maintain charge balance. Most of the preparation methods use an organic compound as N-source in a wet chemical process, usually followed by calcination at mild temperatures. The choice of nitrogen source, as well as the preparation conditions, have a large influence on photocatalytic activity in the resulting product, possibly due to the presence of diverse nitrogen species, such as oxidic (NO<sub>x</sub>), nitridic or amidic (NH<sub>2</sub>). Recent works focusing on cheaper wet-chemical

methods, typically based on urea, triethylamine, etc., as nitrogen source, have seen some success even after mild calcination of the impregnated material, where substantial bulk (ionic) diffusion is unlikely. Consequently, some researchers even doubt that doped- or surface states of N exist as such<sup>170, 171</sup> and the term “N-modification” is generally preferred. There is now strong evidence that “melon-like” surface layers are obtained<sup>170, 171</sup> when starting from urea. The most commonly used nitrogen sources are urea and its derivatives. A careful study by Mitoraj and Kisch<sup>170-172</sup> based on urea has clearly proved that a condensed polymeric layer of melon is formed on the TiO<sub>2</sub> surface, and it is the chromophore that is responsible for visible light activity. As this surface-overlayer-based system is quite distinct from the (bulk) doped type, they state that it is best classified as “melon-modified” TiO<sub>2</sub>.

The formation of melon from urea is a condensation process involving the surface hydroxyl groups of the substrate. It can be conveniently divided into three stages:<sup>170, 171</sup> (1) thermal decomposition of urea into isocyanic acid and ammonia at 320–400 °C (Eqn. 9); (2) cyanamide, and eventually melamine, formation by the reaction of isocyanic acid with surface OH groups of TiO<sub>2</sub> (Eqns. 10, 11). The balance of these reactions is melamine formation from urea in the presence of TiO<sub>2</sub> at 400 °C (Eqn. 12); (3) melamine then undergoes polycondensation to form melem, melam and finally melon (Figure 28). The (yellow) melon is believed to be the principal structure conferring visible light sensitization.

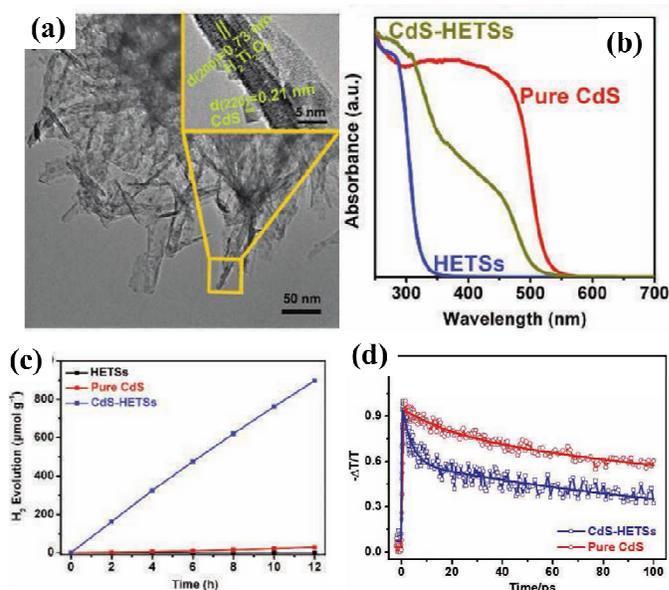


As compared to titanium dioxide, the layered titanate nanotubes/nanobelts offer a practical advantage by promoting the build-up of melon from urea. This is believed to be linked to the higher density coverage of titanates by surface OH groups and their Brønsted acidic properties, which promote polymerization.<sup>173</sup> A dual-phase material composed of hydrated titanate and anatase (TiO<sub>2</sub>) synthesized through a low-temperature one-pot process in the presence of trimethylamine (TEA) as the N-source also exhibited visible light photocatalytic activity that can be attributed to a polyunsaturated hydrocarbonaceous overlayer.<sup>174</sup> Similar observations were discovered for the so-called “C-doped” titanium dioxide. Kisch et al.<sup>175</sup> demonstrated that some organic species containing carboxylic groups can be extracted from the surface of “C-doped” samples by heating them in the alkaline solution, thus the visible light activity of many previously reported so-called “C-doped” titania powder probably does not originate from the presence of lattice carbon atoms or oxygen defects, but from the surface species.



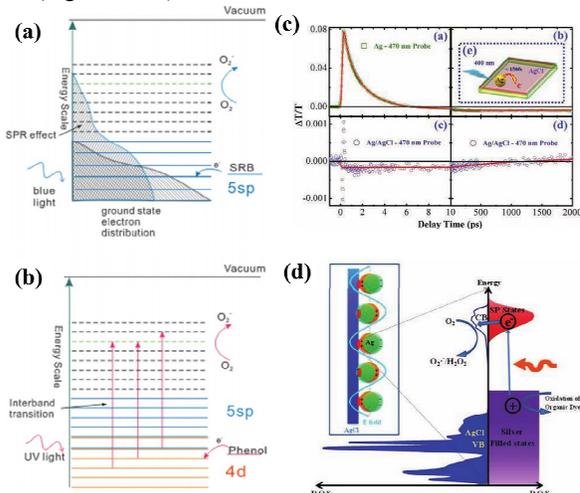
**Figure 28.** Scheme illustrating polycondensation of melamine to melam, melon and melon yellow.<sup>171</sup>

**Metal and narrow band gap semiconductor sensitization:** Another alternative approach to extend the light absorption to visible range is to create  $\text{TiO}_2$ -based composites with colored (narrow band gap) semiconductors such as CdS, CdSe, etc.,<sup>176-179</sup> where a synergy is evident due to directional inter-phase charge transfer analogous to mixed-phase  $\text{TiO}_2$ . For examples, Zhang et al.<sup>176</sup> demonstrated that CdS-titanate spheres, termed as hierarchical echinus-like titanate spheres (HETSs), showed effective photocatalytic activity for visible-light-driven  $\text{H}_2$  (Figure 29). This work confirms that there is fast electron injection ( $\sim 0.32$  ns) between these two kinds of semiconductors (CdS and titanate). This process where the photoinduced electrons on the CdS quantum dot rapidly localizes to the titanate material significantly enhanced the charge separation, therefore promoting the  $\text{H}_2$  yielding compared to the pure CdS sample. On the other hand, noble metal nanoparticles have been widely used as surface modifiers for  $\text{TiO}_2$ -based photocatalysts. These enhance charge separation by acting as “electron sinks”,<sup>180,181</sup> thereby promoting efficient photo-reduction. Noble metal nanoparticles, especially Au and Ag, have the unique property of being optically excited in the visible region due to the so-called surface plasmonic resonance (SPR) effect.<sup>182-185</sup> These materials are now under intensive investigation as potential visible sensitizers.<sup>186</sup> In well-defined nanostructures, localized surface plasmon resonance (LSPR) can occur when confined free electrons oscillate with the incident light irradiation at the same frequency. Under LSPR excitation, plasmon can “excite” the electrons in the conduction band to produce highly energetic “hot electrons”. These “hot electrons” can escape from the plasmonic nanostructures and transfer to a semiconductor in contact, thereby forming a metal–semiconductor Schottky junction.<sup>182, 183, 187</sup> The semiconductor in contact with plasmonic metal nanostructures has a great effect on the charge injection efficiency. Good alignment of the Fermi level of the plasmonic metal nanostructures with the semiconductor band favors efficient photo-generated charge carrier injection.<sup>188</sup>



**Figure 29.** (a) TEM image of CdS-hydrogen titanate spheres; (b) UV-Vis diffused reflectance spectra of pure CdS, HETSs and CdS-HETSs samples; (c)  $\text{H}_2$  evolution of pure CdS, HETSs and CdS-HETSs; (d) Transient absorption (TA) decay profiles (over the first 100 ps) for the pure CdS and CdS-HETSs samples obtained with 400 nm pump/550 nm probe pulses.<sup>176</sup>

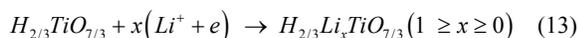
Although gold or silver nanoparticles supported on insulating oxides are reported to have photocatalytic activity under visible light,<sup>189-192</sup> this has been attributed to artifacts, e.g., enhanced thermal catalysis caused by equilibrated local-heating effects<sup>189</sup> and/or instantaneous heat-assisted photo-excitation of Ag electrons to higher energy states (Figure 30a-b).<sup>190</sup>  $\text{TiO}_2$  is by far the most frequently used semiconductor material. A series of papers have reported that Au nanoparticles supported on  $\text{TiO}_2$  surface has much more pronounced activity under visible light irradiation compared with that supported on insulator materials, and, more importantly, the action spectrum for photo-activity resembles the SPR absorption spectrum implying that injection of the plasmonic “hot electrons” from the metal nanostructures to the semiconductor in contact is indeed possible.<sup>184, 193, 194</sup> Other semiconductors, such as silver halides (AgCl, AgBr, AgI), can also serve as electron acceptors,<sup>195-204</sup> and the electron generated from the Ag nanoparticles due to the surface SPR effect is proven (Figure 30c-d).<sup>203</sup>



**Figure 30.** (a-b) The diagram of the band structures of the supported silver NPs and the proposed photocatalysis mechanism;<sup>190</sup> (c) comparisons of 470 nm transient absorption kinetics of Ag nanoparticles and Ag nanoparticles coupled with AgCl nanoparticles after 400 nm excitation; (d) The proposed mechanism of photocatalytic organic dye degradation for the Ag/AgCl hybrid structures.<sup>203</sup>

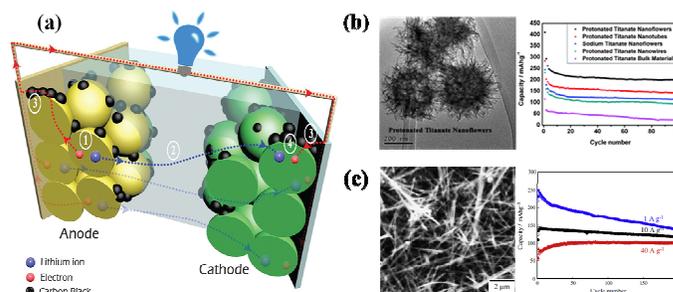
### 4.3 Lithium-ion batteries applications

Rechargeable lithium-ion batteries (LIBs) have attracted intense attentions in recent years for their potential applications in electric vehicles.<sup>205-209</sup> As shown in Figure 31a, the basic operating principle of LIBs is based on  $\text{Li}^+$  shuttling reaction between the cathode and the anode. During cell discharging process,  $\text{Li}^+$  intercalates the positive materials through the electrolyte, while electrons flow through external circuits to provide electricity. During cell charging, the process is reversed. Therefore, the properties of the active Li storage materials greatly influence the performance of rechargeable LIBs. In general, electrode materials that can accommodate large number of lithium will lead to high specific capacity of the battery, while electrodes materials with open structure and short diffusion length allowing for fast ionic/electronic transfer can lead to high current density, and consequently high power density. Conventional graphite electrodes suffer from poor safety characteristics, short cycle life and poor low temperature performances. In searching for better substituents, layered titanate materials emerge as attractive alternatives owing to their open, mesoporous structures with intrinsic high safety. As discussed before, the interlayer distance of titanate nanotubes/nanowires is typically around 0.8 nm, while  $\text{H}^+/\text{Na}^+$  ion are loosely distributed in the interlayer space. This makes them good candidates to provide fast diffusion channels for reversible lithium ion intercalation and de-intercalation, allowing lithium ions to be readily transported and effectively ion-exchanged. The overall lithiation reaction for titanate materials is listed in equation 13. Factors including the number of available lithium reduction sites and the surface area determine their capacity and power characteristics collectively.



So far, high specific capacity and high rate capacity, accompanied by long-term stability and high safety in operation have been achieved in the literature using titanate materials.<sup>14, 210</sup> In terms of specific capacity, earlier work by Li et al. suggested that an initial discharge capacities of 296.6 and 282 mA h  $\text{g}^{-1}$  at current densities of 0.3 and 0.24 A  $\text{g}^{-1}$  can be achieved by hydrogen titanate nanowires and nanotubes electrodes, respectively.<sup>211, 212</sup> Potassium titanate nanowires electrode displayed an higher initial discharge capacity of around 305 mA h  $\text{g}^{-1}$  at a current density of 0.05 mA  $\text{cm}^{-2}$ .<sup>213</sup> Hierarchical titanate microspheres synthesized by Xu's group exhibits even larger initial discharge capacity of 390 mA h  $\text{g}^{-1}$  at current densities of 0.01 A  $\text{g}^{-1}$ , which is superior to that of the hydrogen titanate nanotube, which is 352 mA h  $\text{g}^{-1}$  at the same rate.<sup>214</sup> Later, Zhang et al.<sup>215</sup> also demonstrated that hierarchical titanate spheres (Figure 31b) with large surface area (406.41  $\text{m}^2 \text{g}^{-1}$ ) possess a higher reversible capacity and better cycling stability compared to 1D titanate nanostructures (tubes and wires). Specifically, an initial discharge/charge capacity of 419 and 302 mA h  $\text{g}^{-1}$  is achieved at current rate of 50 mA  $\text{g}^{-1}$ , even after 100 cycles, a high reversible capacity of 200 mA h  $\text{g}^{-1}$  can be obtained, which is superior to other titanate nanomaterials. In terms of rate capacity, Wang et al. synthesized hydrogen titanate nanowires with  $\text{Ti}(\text{OC}_4\text{H}_9)_4$  as the

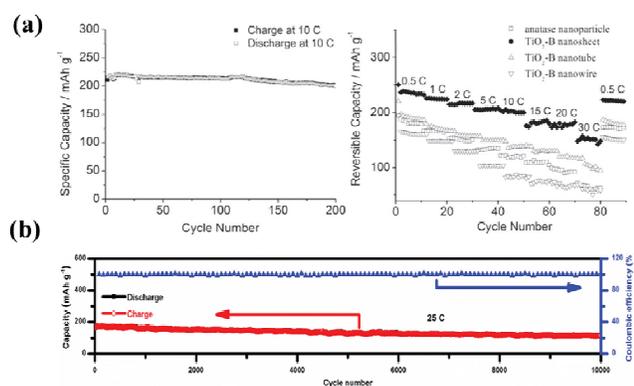
precursor, which delivers a high capacity of 282 and 194 mA h  $\text{g}^{-1}$  at current rate of 1 and 2 A  $\text{g}^{-1}$ .<sup>216</sup> The best rate performances of the hydrogen titanate based materials are reported by Zhou's group (Figure 31c),<sup>217</sup> where TiAl was used as the precursor and emerged into the concentrated NaOH solution for hydrothermal treatment, the obtained hydrogen nanowires ( $\text{H}_2\text{Ti}_3\text{O}_7$ ) exhibit excellent rate capacities. In particular, at current rate of 1 A  $\text{g}^{-1}$ , a capacity of 140 mA h  $\text{g}^{-1}$  can be obtained after 200 cycles, even at extremely high rate of 10 and 40 A  $\text{g}^{-1}$ , a stable capacity of 120 and 100 mA h  $\text{g}^{-1}$  are delivered.



**Figure 31.** (a) Schematic on the operating principle of traditional rechargeable LIBs;<sup>205</sup> (b) Hierarchical titanate nanostructures for lithium ion battery application.<sup>215</sup> (c) hydrogen titanate nanowires as high rate performance electrodes.<sup>217</sup>

Although good capacity and rate capacity have been achieved with titanate materials, the as-obtained hydrogen titanate usually contains structural water, which will react with the organic electrolyte and result in low columbic efficiency in the first cycle and decreased cycle stability. By thermal treatment of the pristine protonated titanate materials, titanate materials will lose the structural water and transform into  $\text{TiO}_2(\text{B})$  nanostructures. Compared to titanate, the nanotubular morphology and open channel structure are retained in the newly formed  $\text{TiO}_2(\text{B})$  materials with improved stability and cycle capacity. Of the most studied polymorphs,<sup>218</sup> nanostructured  $\text{TiO}_2(\text{B})$  has the highest capacity with promising high rate capabilities.  $\text{TiO}_2(\text{B})$  is able to accommodate one  $\text{Li}^+$  per Ti, giving a capacity of 335 mA h/g for nanotubular and nanoparticulate  $\text{TiO}_2(\text{B})$ . Pioneering works regarding the lithium-ion battery performances of the  $\text{TiO}_2(\text{B})$  electrodes are done by Bruce et al.<sup>219, 220</sup> The fabricated  $\text{TiO}_2(\text{B})$  nanowires exhibit a first cycle discharge capacity of 305 mA h  $\text{g}^{-1}$  at a current density of 0.5 A  $\text{g}^{-1}$ , corresponding to  $x = 0.91$  for  $\text{Li}_x\text{TiO}_2(\text{B})$ . Full cell based on  $\text{TiO}_2(\text{B})$  nanowire anode coupled with  $\text{LiFePO}_4$  and  $\text{LiNi}_{0.5}\text{Mn}_{1.5}\text{O}_4$  cathode also demonstrated good cycling stability and rate capability.<sup>221</sup> Similarly,  $\text{TiO}_2(\text{B})$  nanotubes also offer good capacity and cycle stability.<sup>222, 223</sup> The original method in Bruce's work involves static hydrothermal process developed by Kasuga<sup>20</sup> to synthesize  $\text{TiO}_2(\text{B})$  materials, which limits the length of  $\text{TiO}_2(\text{B})$  nanotubes to be less than 1.0  $\mu\text{m}$ . Recently, hierarchical porous  $\text{TiO}_2(\text{B})$  with thin nanosheets, which combines the superiorities of the  $\text{TiO}_2(\text{B})$  polymorph with porous structure as well as thin nanosheets, demonstrated faster insertion and extraction of Li-ion than its nanotube and nanowire counterparts (Figure 32a).<sup>224</sup> Tang et al. synthesized ultralong  $\text{TiO}_2$  nanotubes via a novel stirring hydrothermal technique; the electrode demonstrates excellent rate capacities and cycle life owing to intrinsic high aspect ratio and high conductivity of the long nanotubes. In particular, at rate of 25 C, a capacity of 114 mA h  $\text{g}^{-1}$  can be delivered after 10,000 cycles, maintaining 100% efficiency (Figure 32b).<sup>10</sup> In addition, benefited from the highly viscous nature of the nanotubes, additive-free  $\text{TiO}_2$  electrode could be fabricated and the correlation between aspect-ratio and electrochemical performances of the  $\text{TiO}_2$  materials is

unveiled.<sup>54</sup> In particular, high aspect-ratio TiO<sub>2</sub> nanotubes deliver superior high rate capacities and exceptional long cycle life.



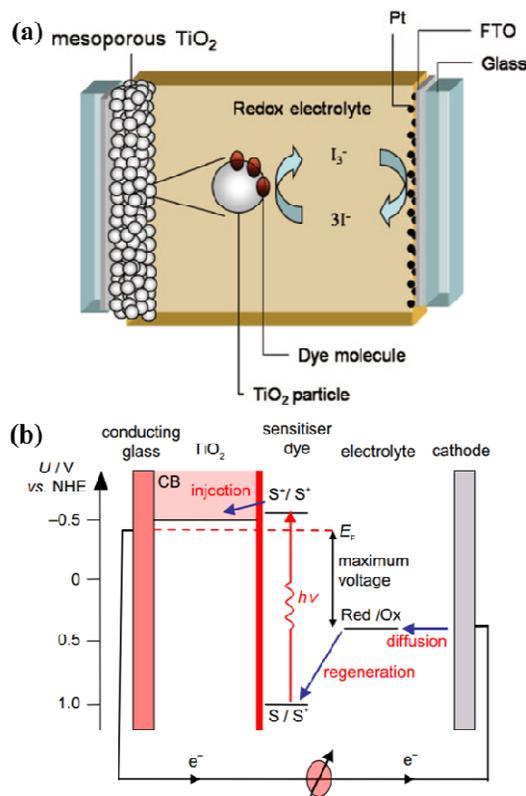
**Figure 32.** (a) Cycling performance of anatase nanoparticles, TiO<sub>2</sub>-B nanosheets, TiO<sub>2</sub>-B nanotubes, and TiO<sub>2</sub>-B nanowires at different charge-discharge rates;<sup>224</sup> (b) Long-term cycling performance of elongated TiO<sub>2</sub>(B) nanotube electrodes at a high current density of 25 C.<sup>10</sup>

These results show the potential of titanate and TiO<sub>2</sub>(B) electrodes in replacing the commercial carbon negative electrodes to offer excellent rate capacity and ultralong cycle life with enhanced safety benefited from the high lithiation potential. The problems related to lithium electroplating and charge loss arising from the formation of a solid-electrolyte interface layer are also avoided.

#### 4.4 Photovoltaic cells and electrochromic devices

##### 4.4.1 Photovoltaic cells application

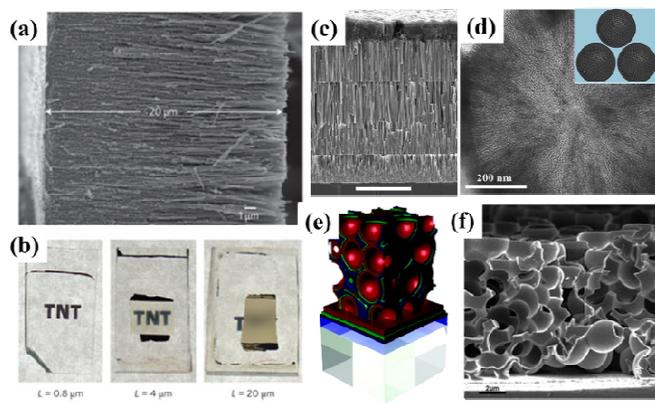
The dye-sensitized solar cell (DSSC), developed by Michael Grätzel,<sup>225</sup> is widely studied for achieving the goal of high efficiency and low cost in the utilization of solar energy.<sup>225, 226</sup> The working principle is as followed.<sup>227-229</sup> The cell is composed of a layer of nanocrystalline semiconductor (TiO<sub>2</sub> is most widely used) particles on a conducting substrate, a platinum counter electrode, and an adsorbed Ru-dye as a sensitizer and an electrolyte. Under irradiation, excited electrons are injected from the lowest unoccupied molecular orbital (LUMO) of the dye into the conduction band of the TiO<sub>2</sub> electrode (Figure 33) through a metal-to-ligand charge transfer pathway. Then the oxidized dye is reduced to its original state by electron donation from the redox couple in the electrolyte. Usually the electrolyte is an organic solvent or ionic liquid containing an I<sup>-</sup>/I<sup>3-</sup> redox system. After the I<sup>3-</sup> ion is oxidized to I, it regenerates the dye. The resulting I species can diffuse to the Pt counter electrode where they are regenerated to I<sup>3-</sup> by reduction.<sup>230</sup>



**Figure 33.** (a) Schematic overview of a dye-sensitized solar cell (DSSC) and (b) its energy level diagram. The potentials for a DSSC based on the N3 dye, TiO<sub>2</sub>, and I<sup>-</sup>/I<sup>3-</sup> redox couple are shown.<sup>231, 232</sup>

At early stage of the development, porous TiO<sub>2</sub> nanoparticle network was used as the photoanode of the DSSC,<sup>225</sup> and promising photo-conversion efficiency was achieved. For example, Tebby et al.<sup>233</sup> reported a UV-irradiation route towards low-temperature processing of nanoporous thin films by hydrothermal approach, and an overall energy conversion efficiency of 2.4–2.5% was obtained on 1–3 μm-thick films modified with Dyesol N3 dye. Magne et al.<sup>234</sup> investigated the effect of TiO<sub>2</sub> polymorphism (anatase, rutile and brookite) on DSSC properties. Their results revealed that the conductivity and diffusion coefficients followed the order of rutile < brookite < anatase, and anatase particles showed the best DSSC performance (ca. 8%). However, for TiO<sub>2</sub> nanoparticles, the overall efficiency is limited due to electron loss during percolation through the nanoparticle network and photo-induced carrier recombination. In order to increase the overall conversion efficiency, different TiO<sub>2</sub> structures, surface modification and morphology design have been explored to enhance the electron transfer ability. Since many excellent reviews have summarized the recent development on the DSSC with electrode structure, organic dye and redox electrolyte design and development, we only focus on the typical examples on the TiO<sub>2</sub> nanostructure for the photoanode of DSSC applications in the current review. 1D nanostructure is an active research area for DSSC. For example, long, vertically aligned titania nanotubes<sup>235</sup> were prepared on transparent conducting oxide glass (Figure 34a-b) with lengths between 0.3 and 33.0 μm using a novel electrochemistry approach. Dye sensitized solar cells containing these arrays yielded a power conversion efficiency of 6.9%. Moreover, a 50 μm thick layer of TiO<sub>2</sub>-coated ZnO nanowire array coating<sup>236</sup> (Figure 34c) was developed for solid-state DSSC cell, which yielded an average power conversion efficiency of 5.65%.

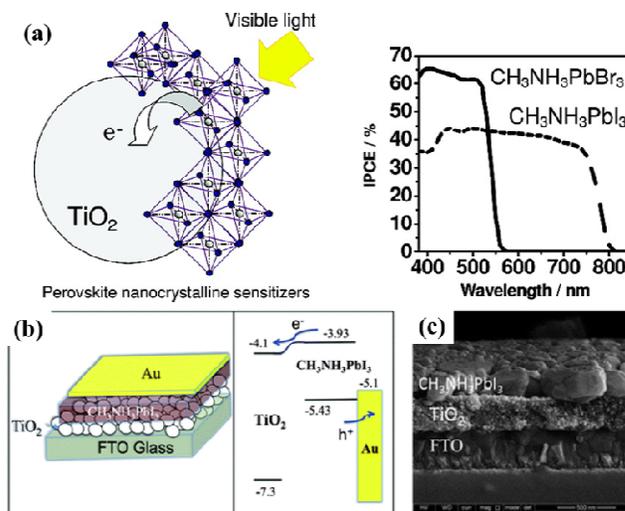
Unfortunately, these 1D arrays suffer from a greatly reduced surface area per unit volume due to their collapse and lack of nanoporosity associated with the roughness of nanometre-sized crystals.<sup>237</sup> Much effort was then drawn towards the fabrication of hierarchical structures with high surface area. More recently, Zhao et al.<sup>238</sup> reported a simple evaporation-driven oriented assembly method to synthesize three-dimensional open mesoporous TiO<sub>2</sub> microspheres (Figure 34d), which has a photoelectric conversion efficiency of up to 12.1%. Tétreault et al.<sup>239</sup> presented a bottom-up synthesis of a high electron mobility and highly light scattering macroporous photoanode (Figure 34e-f) for dye-sensitized solar cells, which reduces recombination and a large surface-area mesoporous anatase guest for high dye loading.



**Figure 34.** (a) FESEM image of a 20-mm-long nanotube array and (b) Digital images of transparent films of different thickness.<sup>235</sup> (c) SEM image of a four-layer TiO<sub>2</sub>-coated ZnO nanowire array by the multistep process.<sup>236</sup> Scale bar, 20 μm. (d) TEM images of a single ultramicrotomed, mesoporous TiO<sub>2</sub> microsphere (inset is the model).<sup>238</sup> (e) Schematic representation of the synthesis method for a 3D host-guest DSSC and (f) SEM of a self-assembled 3D Al/ZnO [TiO<sub>2</sub>] host-guest photoanode.<sup>239</sup>

Stemming from DSSC, perovskite solar cells based on organometal halides represent an emerging photovoltaic technology, which is selected as one of the biggest scientific breakthroughs of 2013.<sup>240, 241</sup> In 2009, a power conversion efficiency (PCE) of around 3–4% was achieved on methylammonium lead halide (CH<sub>3</sub>NH<sub>3</sub>PbI<sub>3</sub>) perovskite / nanocrystalline TiO<sub>2</sub> based on a liquid-based DSSC structure (Figure 35a).<sup>242</sup> However, the stability issues (e.g., dissolution of the perovskite) of liquid-based perovskite solar cell limit their practical applications. In 2012, a long-term stable, and high efficiency (10%) perovskite solar cell was developed by substituting the solid hole conductor with a liquid electrolyte.<sup>243</sup> Moreover, a CH<sub>3</sub>NH<sub>3</sub>PbI<sub>3</sub> / TiO<sub>2</sub> heterojunction solar cell (Figure 35b-c) with a p-n junction structure was reported in 2013.<sup>244</sup> The CH<sub>3</sub>NH<sub>3</sub>PbI<sub>3</sub> was used as a p-type semiconductor. This perovskite solar cell initially showed a PCE of 5.5% with a 500 nm-thick nanosheet TiO<sub>2</sub> film as the n-type layer, and the higher performance approaching 11% was achieved based on this concept.<sup>241</sup> In 2013, the PCE efficiency exceeding 15% was reported using organolead halide perovskite.<sup>245</sup> In 2015, Seok's team<sup>246</sup> showed that the incorporation of methylammonium lead halide perovskite (MAPbBr<sub>3</sub>) into formamidinium lead iodide (FAPbI<sub>3</sub>) stabilizes the perovskite phase of FAPbI<sub>3</sub> and improves the power conversion efficiency of the solar cell to more than 18 per cent. Most recently, a new record certified nonstabilized efficiency of 20.1% was achieved by KRIC.<sup>247</sup> The main advantage of perovskite solar cells is their ability to perform all three basic tasks required for solar cell operation: light harvesting, charge generation

and transport. In addition, their absorption coefficients can be as high as one order of magnitude greater than ruthenium complexes, and they feature long-range charge transport properties.<sup>248</sup> Since PCE values over 20% are realistically anticipated with the use of cheap organometal halide perovskite materials, perovskite solar cells are a promising photovoltaic technology. The recent progresses on perovskite materials and perovskite solar cells are summarized in the recent reviews.<sup>241, 248</sup>



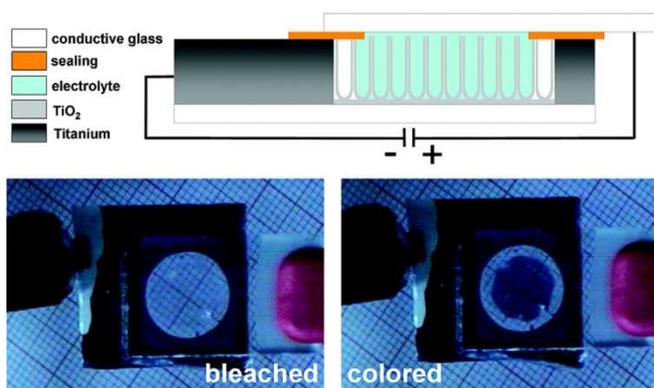
**Figure 35.** (a) Two organolead halide perovskite nanocrystals, CH<sub>3</sub>NH<sub>3</sub>PbBr<sub>3</sub> and CH<sub>3</sub>NH<sub>3</sub>PbI<sub>3</sub>, were found to efficiently sensitize TiO<sub>2</sub> for visible-light conversion in photoelectrochemical cells.<sup>242</sup> (b) Schematic illustration of the CH<sub>3</sub>NH<sub>3</sub>PbI<sub>3</sub>/TiO<sub>2</sub> heterojunction solar cell. (b) Energy level diagram of the discussed solar cell; (c) SEM cross section of the the heterojunction solar cell.<sup>244</sup>

#### 4.4.2 Electrochromic devices application

Similar to some active transition metal oxides (WO<sub>3</sub>, MnO<sub>2</sub>, Nb<sub>2</sub>O<sub>5</sub>, etc.), TiO<sub>2</sub>-based nanostructure is also an excellent electrochromic material for reversible intercalation of extra small ions (e.g., H<sup>+</sup> and Li<sup>+</sup>) into interstitial positions upon applying and releasing an electric field.<sup>249, 250</sup> The effect of ion intercalation into TiO<sub>2</sub> host, accompanied with a reduction of Ti<sup>4+</sup> to Ti<sup>3+</sup>, leads to the apparent change of visible light absorption, which is ascribed to the electronic bandgap changing from the UV (E<sub>g</sub> ≈ 3.0-3.2 eV) to the visible (E<sub>g</sub> ≈ 2.2-2.5 eV) range. In principle, the small ions insertion/extraction switching process (reduction and oxidation of lattice ions) is reversible. Therefore, this visible electrochromism effect is actively utilized for responsively optical devices such as smart windows and smart displays.

Because the specific surface area of the host materials (ion intercalation capacitance) and the defined supply of intercalating ions (electrolyte diffusion path) have great effect on the electrochromic reversibility, aligned TiO<sub>2</sub>-based nanowires and nanotubes constructed by hydrothermal and electrochemical anodizing strategies are particularly interesting in view of electrochromic devices assembling. Campet et al.<sup>251</sup> demonstrated that TiO<sub>2</sub> nanoparticle films constructed by a combination of hydrothermal and doctor-blade process exhibited high electrochromic responses and coloration efficiencies in a stable and environmental friendly ionic liquid electrolyte containing a lithium salt. Schmuki et al.<sup>252</sup> reported that vertical aligned TiO<sub>2</sub> nanotubes films formed by electrochemical anodizing directly on Ti substrate

also displayed better electrochromic abilities regarding  $\text{Li}^+$  and  $\text{H}^+$  ions intercalation than nanoparticle coatings or compact oxide films. To achieve a superior optical contrast and cycle stability, researchers have also developed various strategies to create  $\text{TiO}_2$  nanotube films on transparent conductive substrate, such as ITO or FTO glass (Figure 36).<sup>253-255</sup> Furthermore, modification of  $\text{TiO}_2$  nanostructured films include decoration of electrochromically more active nanoparticles (e.g.,  $\text{WO}_3$ ,  $\text{NiO}$ ,  $\text{V}_2\text{O}_5$ ,  $\text{MoO}_3$ ), single mixed oxide or multilayered metal oxide grown from some transition metal alloys (e.g., Ti-W, Ti-Nb, or Ti-V) were demonstrated to lead to a positive effect on the threshold responsive potentials, higher transmittance contrast, and/or longer cycling stability.<sup>250, 256</sup>



**Figure 36.** Schematic illustration and optical images of the electrochromic device based on transparent  $\text{TiO}_2$  nanotube film on conducting glass substrate in the different coloration (bleached or colored) states.<sup>255</sup>

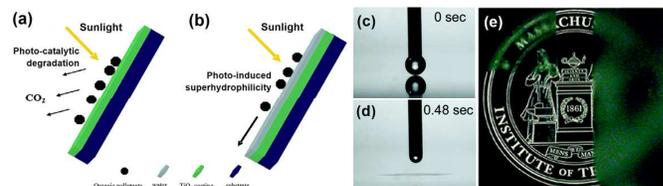
#### 4.5 Self-cleaning, oil-water separation and biomedical application

##### 4.5.1 Self-cleaning

In nature, there exist many kinds of natural self-cleaning surfaces from land to ocean after hundreds of millions of years' evolution. Bio-inspired artificial surface for self-cleaning has been developed since late 20<sup>th</sup> century, and some achievements have led to practical applications. The applications of self-cleaning are very broad and vary from window glasses to solar cell panels. As is known, wettability is closely related to functional surfaces with self-cleaning ability, which provides important clues for antifouling. Three types of surfaces, including superhydrophilic, superhydrophobic, underwater superoleophobic and even superamphiphobic offer effective way to keep a surface clean.

Since the report of  $\text{TiO}_2$  with photocatalytic abilities under UV light illumination and photo-induced superhydrophilic properties, numerous promising applications have been focused based on these materials. Organic pollutants would be photocatalytically degraded and brought away by water layer on superhydrophilic  $\text{TiO}_2$  substrates (Figure 37a,b).<sup>257</sup> Utilizing this strategy, a wide variety of superhydrophilic  $\text{TiO}_2$ -based materials have been constructed, exhibiting environmental friendly applications for self-cleaning, anti-fogging, underwater superoleophobicity, etc.. Lee et al. constructed multi-layer composite coatings with superhydrophilic, self-cleaning and anti-fogging/reflection abilities with a layer-by-layer deposition of  $\text{TiO}_2$  and  $\text{SiO}_2$  nanoparticles.<sup>258</sup> The presence of nanoporous  $\text{TiO}_2$ -based layers in composite coatings resulted in quick wicking of water into the highly hydrophilic network,

exhibiting a long term superhydrophilic and anti-fogging properties (Figure 37c-e).



**Figure 37.** (a) Schematic depicting the self-cleaning via a photocatalytic degradation process on a superhydrophobic  $\text{TiO}_2$  substrate covered with organic pollutants. (b) Schematic diagram of the moving of organic pollutants or dusts when water completely spreading on a superhydrophilic  $\text{TiO}_2$  substrate.<sup>257</sup> Images of a water droplet before (c) and after (d) contacting with the superhydrophilic  $\text{TiO}_2$ - $\text{SiO}_2$  composite coating, and image demonstrating the anti-fogging abilities of  $\text{TiO}_2$ -based multi-layers in composite coatings (e).<sup>258</sup>

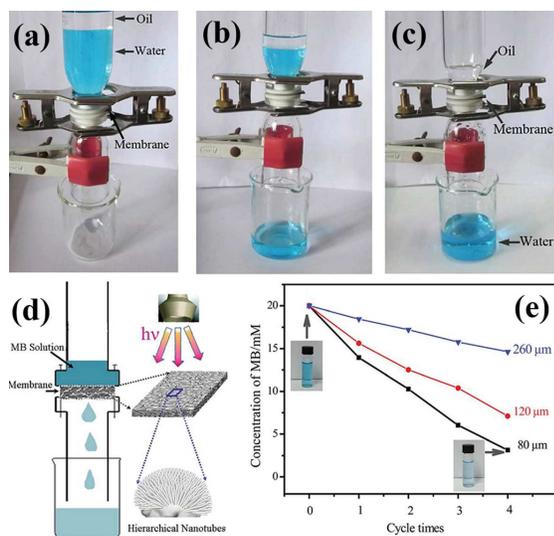
Surface self-cleaning effect is also a significant application of superhydrophobic surface. For superhydrophobic  $\text{TiO}_2$  surfaces, the solid/water contact area is minimized. Water forms spherical droplets that easily slide off the surface and collect the dusts and photocatalytically degraded products on the way. Therefore, the superhydrophobic rough surfaces with low adhesion always exhibit a very low degree of contamination.<sup>259, 260</sup> Zhou et al. reported a long-term superhydrophobic self-cleaning coating prepared by simply blending ambient-cured fluorinated polysiloxane binder with  $\text{TiO}_2$  nanoparticles. The obtained coating exhibited excellent anti-wetting stability in various environments and photocatalytic self-cleaning property for practical applications.<sup>261</sup> Kamegawa et al. reported superhydrophobic surfaces with photocatalytic self-cleaning properties through coating a nanocomposite  $\text{TiO}_2$  photocatalyst and hydrophobic polytetrafluoroethylene onto a structured substrate by applying a co-deposition technique.<sup>262</sup> Recently, Parkin et al. developed a practical approach to construct robust superhydrophobic self-cleaning surfaces by taking advantage of commercial adhesive to bond the hydrophobic  $\text{TiO}_2$  nanoparticles to various substrates such as clothes, paper, glass, and steel.<sup>263</sup> They also reported that the oil (hexadecane) contaminated surface still maintained self-cleaning ability under various environments including underwater, in air or even oil. Moreover, these treated substrates kept the excellent anti-wetting ability after finger-wipe, knife-scratch, or up to 40 sandpaper abrasion cycles.

In addition to superhydrophobicity and superhydrophilicity in air, underwater superoleophobicity is also a promising characteristic for self-cleaning surfaces, which can be easily achieved with the combination of suitable structure and high energy chemical component or constructed  $\text{TiO}_2$ -based surfaces after light irradiation. In nature, fish scale and shark skin are known to be able to prevent oil contamination due to a thin water layer on their surface, exhibiting excellent underwater self-cleaning and bio-antifouling. In general, anti-fouling, low drag, slippery surfaces, and photocatalytic activities play a vital role in the underwater self-cleaning effect.<sup>264, 265</sup> Recently, Sawai et al. reported the preparation of smooth  $\text{TiO}_2$  film for sustainable underwater self-cleaning activity attributed to the photo-induced hydrophilic surface for thin water film formation to effectively prevent oils from contacting the  $\text{TiO}_2$  film.<sup>265</sup> The research on underwater superoleophobicity is still in its infancy, and the characterization techniques for self-cleaning have to be further developed.

Besides pristine TiO<sub>2</sub>, other TiO<sub>2</sub>-based materials such as titanates have also been attracted attention for self-cleaning applications. For example, highly hydrophobic titanate nanobelt (TNB) particle can be easily prepared by a hydrogen-bond-driven assembly of pre-hydrolysed fluoroalkylsilane (FAS).<sup>266-268</sup> Using this kind of TNB/FAS powder, a facile spinning, spraying or dipping process was applied to coat a various substrates with cross-aligned superhydrophobic film.<sup>268</sup> The as-prepared coating displayed good self-cleaning ability and high chemical stability in a wide range of pH solutions.

#### 4.5.2 Oil-water separation

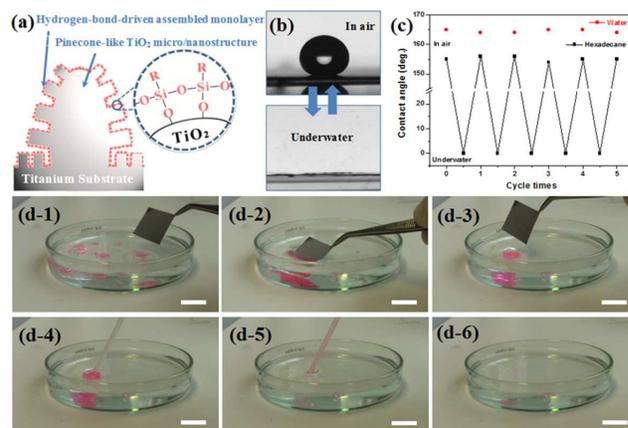
Oil/water separation technology is very important for a wide range of environmental, biomedical, agricultural, and industrial applications.<sup>269-274</sup> Separation of liquids, especially oil-water separation is a worldwide challenging problem arising from the increasing industrial oily wastewater and the frequent oil spill accidents. The conventional strategy for oil-water separation depends on the special wetting membranes with simultaneous superhydrophobicity and superoleophilicity for the separation of oil from the oil-water mixture.<sup>275, 276</sup> For example, Li *et al.* reported a facile electrochemical anodizing method for fabricating porous TiO<sub>2</sub> membrane with special wettability.<sup>275</sup> Such a multifunctional membrane can be successfully used for the separation of insoluble oil from water due to its high wetting contrast (Figure 38a-c), as well as the degradation of soluble pollutants in water under UV light because of the photocatalytic abilities of TiO<sub>2</sub> (Figure 38d,e), making it a promising candidate for water purification. The integration of the inherent photo-induced hydrophilicity, chemical stability, and photocatalytic activity of TiO<sub>2</sub> material enables the convenient removal of the contaminants by ultraviolet irradiation, and allows facile recovery of the separation ability, making it promising for sustainable and highly efficient oil-water separation applications.<sup>277</sup>



**Figure 38.** Proof-of-concept oil-water separation device and process (a-c) using anodized porous TiO<sub>2</sub> membrane. The pre-wetted membrane was fixed between two glass vessels. A mixture of petroleum ether and water was poured into the upper vessel (a). Water permeated through the anodized porous membrane (b), while oil stayed in the upper glass vessel (c), which let to oil-water separation. (d) Schematic diagram for the flow-through photocatalysis device with anodized porous TiO<sub>2</sub> membrane for the photocatalytic degradation of methylene blue (MB). (e) The

concentration variation of MB solution after the photocatalytic degradation with various anodized porous membrane photocatalyst and circulated for different times. Inset: the photographs of MB solution before and after 4-cycles flow-through photocatalytic reaction with a maximal pore size of 80 μm.<sup>275</sup>

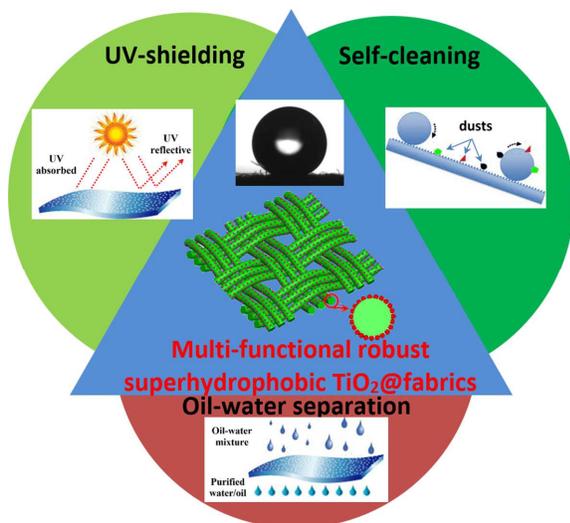
Recently, inspired by living creatures with special wettability and their precise arrangement of topographical structure and chemical component, we described a facile one-step approach to large scale construct superamphiphobic TiO<sub>2</sub> particles film.<sup>278</sup> The as-prepared TiO<sub>2</sub> films exhibit excellent superamphiphilic property in air, changes to superamphiphobicity with good dynamical stability after silane modification (Figure 39a). Moreover, such superamphiphobic surface reversibly can switch from superoleophobic to superoleophilic in air and underwater, respectively (Figure 39b,c). The 3D functional surface would be a versatile platform in a wide range of applications, e.g., self-cleaning, friction reduction, and repeatable oil-water separation. In a proof-of-concept study, we applied the fluoroalkylsilane-modified pinecone-like TiO<sub>2</sub> particles samples with both superamphiphobic in air and superoleophilic underwater as “oil capture hands” to gather oil droplets in water (Figure 39d, process 1-3). The aggregated oils were then extracted to achieve effective oil-water separation (process 4-6). More importantly, it is worth noting that the sample maintain self-cleaning without any oil residue because of the excellent superamphiphobic property in air. This is different from the conventional superhydrophobic surface that is completely oil contaminated after oil absorption in water, which makes it difficult for repeated use. This study opens up a new avenue for the treatment of oily waste water and points out a promising application in industrial fields.



**Figure 39.** (a) The hydrogen-bonding-driven self-assembled process on the hydroxylated TiO<sub>2</sub> particles surface. (b) The oil droplet image on the modified TiO<sub>2</sub> surface in air or underwater environment. (c) The reversible wettability change of oil droplet on the TiO<sub>2</sub> surface by alternating the environment. (d) The oil capture and collection process with a superoleophobic plate and underwater superoleophilic property. Process 1-3: a superoleophobic plate touches, captures and collects the sprayed oil drops in bottom of a glass container. Process 4-6: the gathered oil droplets are collected from the water. The oil was colored pink for convenient observation through the capture and collection process.<sup>278</sup>

As discussed above, multifunctional TiO<sub>2</sub> surfaces with special wettability and adhesion have attracted great amount of attention partially owing to their promising applications in anti-bioadhesion, self-cleaning, oil-water separation, and photocatalysis. Recently, Huang *et al.*<sup>279</sup> constructed a TiO<sub>2</sub>@fabric hierarchical composite via

a facile strategy for loading TiO<sub>2</sub> nanoparticles through a one-pot hydrothermal reaction (Figure 40). By taking advantage of the excellent superhydrophobic property with UV-resistance, low droplet sliding angle and high contrast of wettability for oil and water, the as-constructed UV-durable TiO<sub>2</sub>@fabrics pave the way for their practical applications in various fields such as UV-shielding, self-cleaning and oil-water separation.



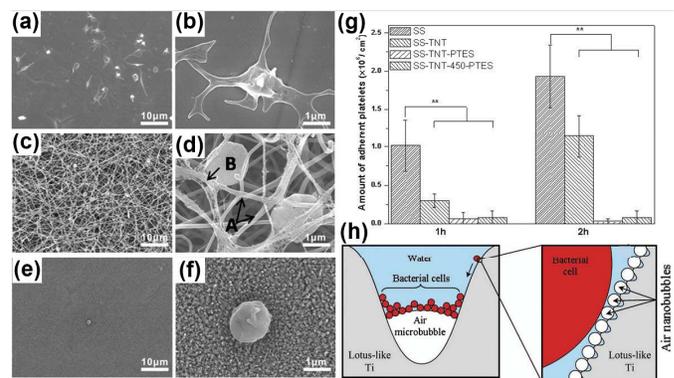
**Figure 40.** Schematic representation of robust superhydrophobic TiO<sub>2</sub>@fabrics prepared via a facile one-pot hydrothermal strategy for hierarchical TiO<sub>2</sub> particles construction and fluoroalkylsilane modification as a multi-functional platform for UV shielding, self-cleaning, and oil-water separation.<sup>279</sup>

#### 4.5.3 Anti-bioadhesion & biomedical application

Beyond the environmental and energy applications, TiO<sub>2</sub>-based functional materials have shown positive effects on biomedical engineering such as bioactive scaffolds, implants, vascular stents, biosensors and drug delivery systems.<sup>280-285</sup> Many factors, such as composition, topography, charge and wettability, are vital in the wettability of TiO<sub>2</sub>-base surface and resultant practical applications in biological science related fields.<sup>286-290</sup> Surface with unique wettability is one of the most important factors influencing protein adsorption, platelet adhesion, bacteria adhesion and cell culture.<sup>291-294</sup> Wall adsorption on superhydrophilic or even hydrophilic surfaces is a common problem in microfluidic devices, particularly when proteins are involved. Superhydrophobic surface with low solid-liquid adhesion was verified to be effective to suppress the attachment and activation of bacteria and cell. In general, the grafting of low energy materials on superhydrophilic surface with rough dual-scale structures is an effective strategy to construct superhydrophobic film. For example, we realized the construction of superhydrophobic surface via a specific condensation reaction between the pre-hydrolysis fluoroalkylsilanol groups of -SiOH and the surface hydroxyl groups of -TiOH for the self-assembling of low energy fluorocarbon groups on hydrophilic or even superhydrophilic TiO<sub>2</sub>-based surface.<sup>245</sup> Recently, Levanen and co-workers

showed that superhydrophobic surfaces can be used to reduce bacterial adhesion as well as to kill bacterial by highly reactive species created by the photocatalysis of TiO<sub>2</sub>.<sup>295</sup>

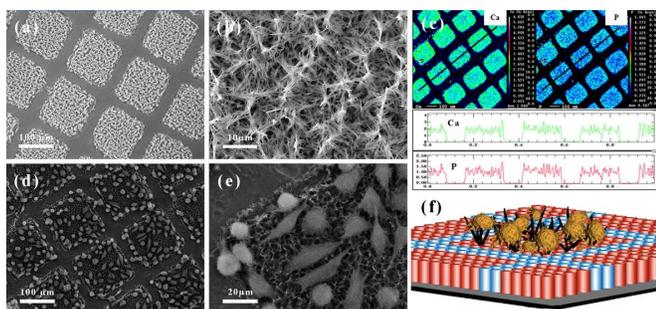
Platelet adhesion and activation on biomedical material surfaces may result in blood coagulation and thrombosis. Therefore, regulating the wettability to increase the blood-compatibility is an efficient way for platelet anti-adhesion. Recently, Yang and Huang et al. combined superhydrophobicity with photocatalysis by coating a TiO<sub>2</sub> surface with fluoroalkylsilane to induce anti-platelet effect.<sup>296, 297</sup> The *in vitro* blood compatibility experiment indicates that the superhydrophobic TiO<sub>2</sub> nanotube layers exhibit a remarkable ability in reducing platelets adhesion. Compared to the abundant platelets adhered on smooth stainless steel surface (Figure 41a,b), relatively low amount of platelets entrapped into fibrin network on hydrophilic rough TiO<sub>2</sub> nanotube arrays upon stainless steel substrate, high TiO<sub>2</sub> layers (Figure 41c,d), and only very few platelets were found to adhere on the superhydrophobic TiO<sub>2</sub> nanotube layers (Figure 41e,f).<sup>298</sup> Moreover, even though some platelets were occasionally attached on the superhydrophobic surface, they looked smooth without any growth of pseudopods, implying that the platelets adhered on the superhydrophobic TiO<sub>2</sub> nanotube surface remain inactive and hardly grow or spread out. From the *in vitro* evaluation, the superhydrophobic TiO<sub>2</sub> layers exhibit excellent blood compatibility and remarkable performance in preventing platelets or bacteria from adhering to the implant surface due to the creation of nanobubbles or microbubbles so that the direct attachment of cells on substrate is reduced (Figure 41g-h). Tang *et al.* also observed similar bacterial adhesion behaviour on TiO<sub>2</sub> structures with various surface wettability.<sup>299</sup> Compared to ordinary hydrophilic or hydrophobic film, the as-prepared super-antiwetting film can greatly decrease the adhesion and activation of *Staphylococcus aureus*. This indicates that the special topography and wettability play critical role in blood-compatibility. Attempts for developing hybrid anti-bacterial materials by combining superhydrophobicity with anti-bacterial metals or with photocatalysis have been verified to enhance the anti-bacterial ability while maintaining the primary function of preventing bacterial adhesion. The effort of loading anti-bacterial metallic ions (e.g., Ag, Cu and Mo) with superhydrophobicity has been more successful than combining photocatalysis with superhydrophobicity. In another effort, Lai et al. demonstrated that the combined super-antiwetting and photocatalysis on TiO<sub>2</sub> surface, together with loading of Ag particles would largely enhance the antibacterial activity.<sup>300</sup>



**Figure 41.** SEM images of adherent platelets on the surface of stainless steel (a, b), superhydrophilic TiO<sub>2</sub> nanotube film on stainless steel (c, d), and superhydrophobic modified TiO<sub>2</sub> nanotube film on stainless steel (e, f). (g) Number of adherent platelets on different samples.<sup>297</sup> (f) Proposed mechanism by which bacterial

cells accumulate at the tri-phase interface on immersed superhydrophobic Ti surfaces.<sup>298</sup>

On the basis of photocatalytic lithography patterning technique on TiO<sub>2</sub> structures, two dimensional (2D) superhydrophilic-superhydrophobic pattern has been prepared to guide the selective deposition of bioactive calcium phosphate (CaP) layer (Figure 42a-c), and further induce the cell attachment onto the CaP/TiO<sub>2</sub> nanotube film (Figure 42d-f).<sup>300-302</sup> It can be clearly seen that CaP nanobelts are preferred to be deposited within the superhydrophilic regions to form CaP pattern while the superhydrophobic areas are kept intact. In the case of the bioadhesion, cells preferentially attached and activated on CaP/TiO<sub>2</sub> regions to generate 3D pattern. Following this principle, arrays of TiO<sub>2</sub>, CdS, ZnO, Ag and molecular probes<sup>303-305</sup> were assembled directly from solution onto Ti substrates at the desired locations precisely. These patterned structures can then be fabricated into arrays of photodetector, gas sensor or matrix devices for large-area microelectronic applications or bio-compatible coatings where drugs can be encapsulated in specific areas of the coating.<sup>306</sup> This patterning strategy will be helpful to develop various micropatterned functional nanocomposite materials. However, a great challenge for wide-spread practical applications still remains to be the development of high-throughput, low-cost, and easily-controllable techniques to achieve desired orientation and order of the nano-blocks.



**Figure 42.** (a) Typical SEM images of the 3D spatially patterned CaP micro-nanostructure films on TiO<sub>2</sub> film. (b) Magnified SEM of CaP crystals on superhydrophilic region. (c) Element distribution maps and corresponding line-scan signal intensity profiles of Ca and P elements across the square pattern (black line direction). (d,e) SEM images of the MG-63 osteosarcoma cells adhering on the CaP patterning surface. (f) Schematic diagram of the site-selective cell immobilization on 3D rough CaP@TiO<sub>2</sub> scaffold.<sup>300</sup>

## 5. Conclusions and perspectives

In this review, we present the state-of-the-art of the development of TiO<sub>2</sub>-based materials, namely titanate and titania, for environmental and energy applications. The titanate and titania materials can be easily transformed to each other via simple processes. For example, titanate materials can be transformed to titania via thermal annealing or hydrothermal treatment of hydrogen titanate, and the titania material can react with the alkaline solutions to form titanate materials. A clear advantage of synthesizing titania from titanate is to maintain or make use of the various nanostructures easily obtainable for titanate materials. Guided by the basic formation principles of bulk materials by solid-state reaction, the preparation strategies of the formation of different dimensional titanate materials via liquid- or vapor- phase hydrothermal reaction, external and internal modified hydrothermal reaction, molten-salt reaction,

high temperature oxidation, and electrochemical spark discharge spallation (ESDS) process are summarized in this review. Among them, hydrothermal route is most popular for the synthesis of 1D, 2D and 3D titanate materials due to the easy operation and rational control of the nanostructures.

Due to the unique crystal structure and physical-chemical properties, titanate materials show great potential for applications in adsorption, lithium ion batteries, biomedical applications, self-cleaning, and oil-water separation. Since titanate materials possess large interlayer spacing and adjustable lattice parameters, they can be widely used as adsorbents due to their unique ion-exchange property. To be a good adsorbent, the adsorption capacity and removal rate are important considerations. The adsorption performance depends on the surface area and the number of exchangeable cations in the titanate materials.

For the photocatalytic application, due to their wide band gap (3.4~3.7 eV) and the crystal defects, the photocatalytic performance of titanate materials is not as good as TiO<sub>2</sub>. Thus, it is essential to obtain the titania materials with required crystal structure and designed nano-construction from titanate by rational control of the morphology of the starting material. An interesting strategy is to combine the excellent adsorption ability of titanates and high photocatalytic activity of titania to promote faster photodegradation of organic pollutants in wastewater. Also, the doping technology and hybrid structures are also introduced to enhance the visible light activity.

For the energy storage application, titanate and titania materials show good potential in replacing the commercial carbon negative electrodes, offering excellent rate capacities and ultralong cycle life with enhanced safety benefited from the high lithiation potential, and absence of the lithium electroplating and the charge loss problem arising from the formation of a solid-electrolyte interface layer. As an important photoanode for the DSSC and perovskite solar cells, the engineering of TiO<sub>2</sub> nanostructures towards prolonging the charge carrier lifetime, and bandgap matching with dye or (organometal halides) are critical to achieve the goal of high efficiency and low cost in the utilization of solar energy. By making use of the reversible transition from Ti<sup>4+</sup> to Ti<sup>3+</sup> into TiO<sub>2</sub> host by ion intercalation/de-intercalation, the visible electrochromism effect can be achieved on the TiO<sub>2</sub> materials.

Lastly, multifunctional TiO<sub>2</sub>-based surfaces with special wettability and adhesion have attracted great attention owing to their promising applications in anti-bioadhesion, self-cleaning, oil/water separation.

Overall, substantial work has been carried out in the synthesis of titanate and titania nanostructures and their applications. Excellent proof-of-concept work has been realized at laboratory scale. Future effort should be focused on the improvement of its functionalities through materials and structural optimization, eventually leading to practical environmental, energy, and other related applications. Practical application of titanate and titania materials is realistically achievable as these materials are intrinsically stable in a wide range of chemical media and under UV illumination, earth-abundant, and low cost in production. For the absorbent application, the mass production on titanate materials with high surface area is an important future effort towards low-cost and effective pollutant removal, especially for the toxic metal ions and non-degradable organic dye. Little attention has been paid so far on the disposal or recovery of the

titanate materials (together with some metal ions of high value) after adsorption. More research effort should be made in this aspect the future.

As a well-studied photocatalyst system in the past decades, the photocatalytic efficiency of TiO<sub>2</sub> under full solar energy spectrum is still low. New photoconversion concept based on rational materials design or bandgap engineering is urgently needed to achieve high solar-to-fuel conversion efficiency. For other energy related applications, much effort is needed to commercialize these TiO<sub>2</sub>-based materials for rechargeable lithium ion battery and DSSC applications. In particular, the exploration on the TiO<sub>2</sub> materials towards solid state, wearable and printable energy devices will have a promising future for the development of efficient and flexible optoelectronics. For the wetting ability control on TiO<sub>2</sub> nanostructure, the rate of wetting property change, stability under UV, and mechanical strength should be carefully studied in the future.

### Acknowledgements

The authors thank the Environment and Water Industry Programme Office (EWI) under the National Research Foundation (NRF) of Singapore (grant MEWR651/06/160), National Natural Science Foundation of China (grant 21501127; 51502185), Natural Science Foundation of Jiangsu Province of China (grant BK20130313), Priority Academic Program Development of Jiangsu Higher Education Institutions (PAPD) for financial support of this work.

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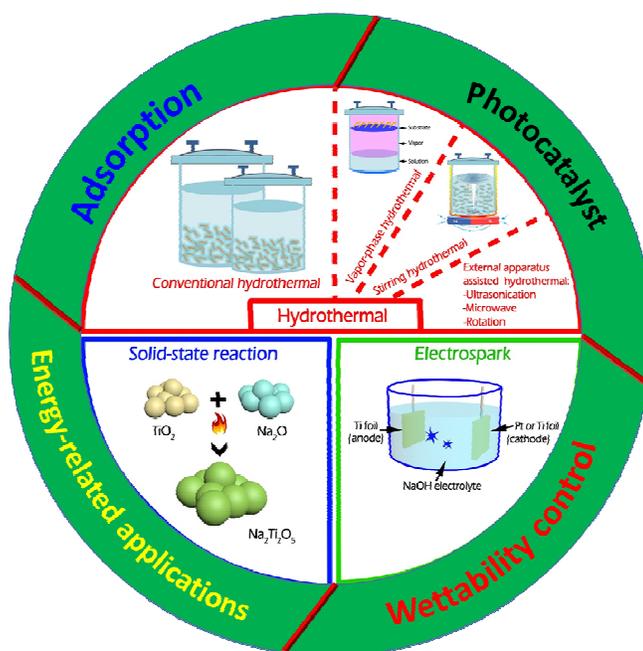
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TOC images



In this review, we present the state-of-the-art development of fabrication strategies of titanate/titania nanostructures and the corresponding applications including adsorbent, photocatalysis, lithium-ion batteries, photovoltaics, electrochromic device, biomedical application, self-cleaning and oil-water separation.