Preparation of TiO$_2$ Sols with High Solid Contents by Sol Gel Method

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Abstract: Titanium oxide has been extensively used as a photocatalytic material because of its excellent UV light-responsive effect, high refractive index, and high chemical stability. TiO$_2$ in anatase phase for applications in optical or electronic devices and photocatalysis has generally been used in the form of a thin film. The aim of this study was to prepare a stable TiO$_2$ sol with high solid content. Therefore one can use less coating time to have a mono layer of TiO$_2$ film on the substrate. In this study, TiCl$_4$ was slowly added to the distilled water in an ice bath. Aqueous solution of sodium hydroxide was added to adjust the pH of the solution to between 8 and 12 to form titanium hydroxide gel. After aging for a period of time, the Ti(OH)$_4$ gel was filtered and sufficiently washed. The filtered cake was repulped in water. Hydrochloric acid, as a catalyst for polycondensation, was slowly added to the solution with stirring. After poly-condensation reaction and crystallization, a transparent suspended TiO$_2$ sol was formed. XRD results show that the crystalline phase was anatase. The suspended TiO$_2$ particles were rhombus primary particles with the major axis ca. 20 nm and the minor axis ca. 5 nm. The sample prepared at pH 8 had the largest BET surface area of 141 cm^3/g among all samples. The best preparation condition to have the smallest TiO$_2$ particles are Ti: H = 1: 1 (atomic ratio), 10% solid content of TiO$_2$, and hydroxypropyl cellulose with viscosity of 150-400 cps as a surfactant. The thin film was obtained by dip-coating the glass in TiO$_2$ sol. Since the sol had high solid content, the coating time became less. The dip-coating on glass was less than three times to have one monolayer TiO$_2$. The transparent TiO$_2$ thin film had super-hydrophilicity after illumination by UV light.

Keywords: Titania, Photocatalysis, Sol-gel, Coating, Anatase

1. Introduction

Titanium dioxide has been used in a wide range of applications. [1-12] It is low cost, chemically stable, non-toxic and bio-compatible. Titania has been used as a photocatalyst, implant material in dental, orthopedic and osteosynthesis applications. TiO$_2$ thin films have a high refractive index that makes TiO$_2$ suitable for optical applications. TiO$_2$ is widely used for photocatalysis to destruct dyes, pollutant, and bacteria under UV light illumination.

TiO$_2$ sol has been used to prepare TiO$_2$ film on substrates by various coating techniques.[1-23] Various preparation methods to prepare TiO$_2$ sols have been reported in the literature. [5-9, 19-21] Most of researchers used titanim alkoxide as the precursor. [2-4, 14, 25] However, titanium alkoxide is expensive and not stable in air. Some researchers used TiCl$_4$ as the precursor. [14, 15] However, most of sols have low solid content. [22-31] It would need many times coating to have one monolayer TiO$_2$ thin film.

The aim of this study was to develop a method to prepare TiO$_2$ sol which has high solid content. In this study, TiCl$_4$ was used as the starting material. It was converted to Ti(OH)$_4$ gel with NaOH. The gel was then converted to TiO$_2$ sol by adding HCl as the catalyst for polycondensation and crysallization, In this study, with carefully control pH value and amount of NaOH and HCl, we have successfully prepare TiO$_2$ sol with 10% solid content.

2. Experimental

2.1. Chemicals

Titanium tetrachloride (> 99.9%), sodium hydroxide

(>96%), and hydrochloric acid (35%) were from Showa Chemicals Co. Two types of hydroxypropyl cellulose (HPC, viscosity = 6-10 cps and 150-400 cps, 2% in water at 20°C) were from Merck.

2.2. Synthesis of TiO₂ Sol

$TiCl_4$ was used as a precursor. It was slowly added to the distilled water in an ice bath at 5°C. When $TiCl_4$ dissolved in distilled water, the heat of the exothermic reaction explosively generated the formation of orthotitanic acid. $Ti(OH)_4$. Because the formation of this species disturbed homogeneous precipitation, 5 ml of hydrochloric acid (35% HCl) per 100 ml of distilled water was added to prevent formation of orthotitanic acid. Aqueous solution of sodium hydroxide solution was slowly added to adjust the pH of the system to 8-12. After aging for a period of time, the peptizate sol was filtered and sufficiently washed to remove chlorine ion from the cake. The cake was repulped in water. Hydrochloric acid was slowly added to the solution with stirring. After condensation reaction and crystallization, the transparent suspended TiO₂ sol was obtained.
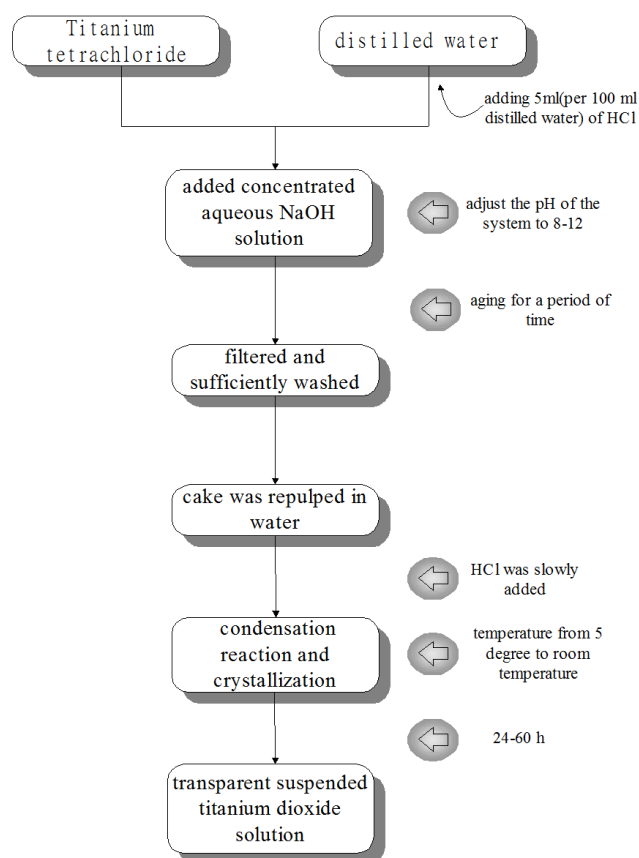

Figure 1. *Preparation method of transparent suspended TiO₂ sol.*

2.3. Characterization

2.3.1. X-ray Diffraction (XRD)
The crystalline structures of the products were analyzed by X-ray powder diffraction using a Siemens D500 automatic powder diffractometer system. Nickel-filtered Cu K_α radiation

(λ= 0.15418 nm) was used with a generator voltage of 40 kV and a current of 29 mA. The Bragg-Brentano focusing geometry was employed with an automatic divergence slit (irradiated sample length was 12.5 nm), a receiving slit of 0.1 nm, a fixed slit of 4° and a proportional counter as a detector. It was operated in the step scan mode, at scanning speeds of 0.05° 2θ/s and 1 second step time from the range 20° to 80°. Scherrer's equation was used to calculate the crystalllite size of TiO₂ crystal:

$$D = \frac{k\lambda}{\beta \cos\theta} \qquad (1)$$

where D is crystallite particle size, κ is a constant of 0.94, λ is X-ray wavelength (0.154 nm), β is half maximum line breadth, and θ is Bragg angle.

2.3.2. N₂ Sorption
The measurement of the surface areas of the samples were achieved by Brunauer-Emmett-Teller (BET) method, and the pore size distributions were determined by Barrett-Joyner-Halenda (BJH) method. The N₂ sorption was carried out in a ASAP 2100 instrument (Micromeritics Co.).

2.3.3. Thermogravimetric Analysis (TGA)
The TGA was carried out at a Perkin Elmer TGA-7 with a heating rate from 50 to 600°C. Those samples were heated in nitrogen at a heating rate of 10°C /min.

2.3.4. Differential Scanning Calorimetery (DSC)
DSC analysis was measure by using Perkin Elmer DSC-7 with a heating rate of 10°C /min from 50 to 450°C.

2.3.5. Scanning Electron Microscopy (SEM)
Sample was dried at 35°C, then placed on a stage for SEM. The sample was coated with Au for 4 min before the experiment and was observed using a Hitachi S-800. It was operated at 200 kV.

2.3.6. Transmission Electron Microscopy (TEM)
The suspended TiO₂ was deposited on a grid with a holey carbon copper film and placed several days at room temperature. It was observed in a JEM-1200 EX II electron microscope operating at 160 kV. The magnification was calibrated in pixel/nm on the camera.

2.3.7. Dynamic Light Scattering (DLS)
The particle size of the sample was measured by DLS using Zetasizer 3000. Dynamic light scattering measures Brownian motion and relates it to size. Random movement of particles due to bombardment by the solvent molecules that surround them, and the larger the particle is, the slower the Brownian motion is. One can calculate the hydrodynamic diameter d(H) by using the Stokes-Einstein equation:

$$d(H) = \frac{\kappa\lambda}{3\pi\eta D} \qquad (2)$$

where d(H) is hydrodynamic diameter, κ is Boltzmann's

constant, T is absolute temperature, η is viscosity, and D is diffusion coefficient.

2.3.8. Ultravillet /Visible Absorption (UV-vis)

The diffuse reflectance UV-vis absorption measurement was carried out with a Jasco UV V-500 spectrophotometer. The sample was loaded in a quartz cell with suprasil windows, and spectra were collected in the range from 300 nm to 800 nm against quartz standard.

2.3.9. Contact Angle Meter

The contact angle analysis was measure by using Model CA-D (Kyowa Interface Science Co., Ltd. Japan). Glass slide was used as a substrate. TiO_2 was coated on the glass substrate several times, then small amount of deionized water was drop on it. The contact angle is based on the equation as follow:

$$\gamma = \frac{rh\rho g}{2\cos\theta} \qquad (3)$$

where γ is surface tension of a capillary, r is radius of a capillary, h is the height of liquid, ρ is the density of liquid, g is gravity, and θ is contact angle.

3. Results and Discussion

This research was devoted to prepare a stable and suspended TiO_2 sol with high solid content, small particles size, and anatase crystalline phase. Five preparation parameters were studied, i.e., pH value, $[H^+]/[Ti]$ ratio, HCl concentration, stored temperature, and surfactant.

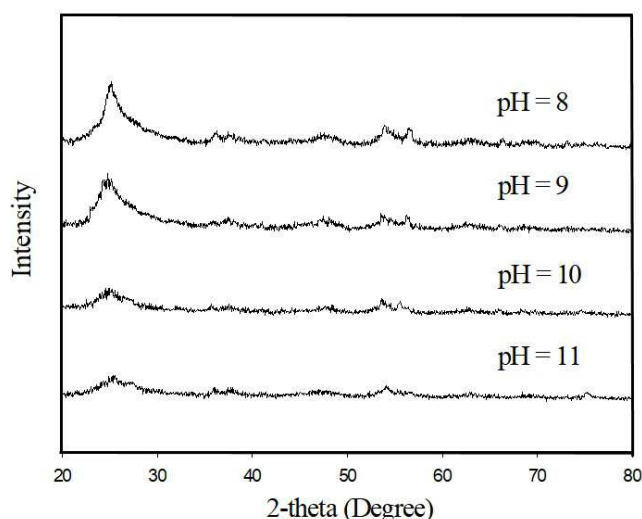*Figure 2. XRD patterns of TiO_2 prepared at different pH values.*

3.1. Particle Size Distribution

Effects of pH value

After dissolving $TiCl_4$ in water, NaOH was added to adjust the pH value of the solution. Based on the literature [32] the isoelectric point (IEP) of TiO_2 is at pH 4.5-6.8. If the pH value is adjusted to this region, there is no repulsive force between particles so as to maintain the dispersion of the particles. The

particles collide in each other, and the particles will aggregate and precipitate due to vander Waals interaction. Due to the drawback described above, the pH value was adjusted to greater than 8 or to value lower than 3 to have stable TiO_2 suspension.

As the TiO_2 particle was small, the XRD peak should appear, but not intense. As shown in Figure 2, all the samples showed anatase XRD peaks with low intensity. The stable anatase TiO_2 was obtained at pH 8-11. The samples prepared at pH 8 and 9 had more intense XRD peaks than those prepared at pH 10 and 11. It should be noted that there was only one major peak in the XRD patterns. It indicates the formation of anatase, but the crystallite size is small. Scherrer's equation was used to calculate the mean diameter of the crystallites. The sample prepared at pH 8 had the smallest crystallites among all samples (Table 1).

Table 1. Mean diameters of TiO_2 crystallites prepared at various pH values (Mean diameter was calculated from XRD result).

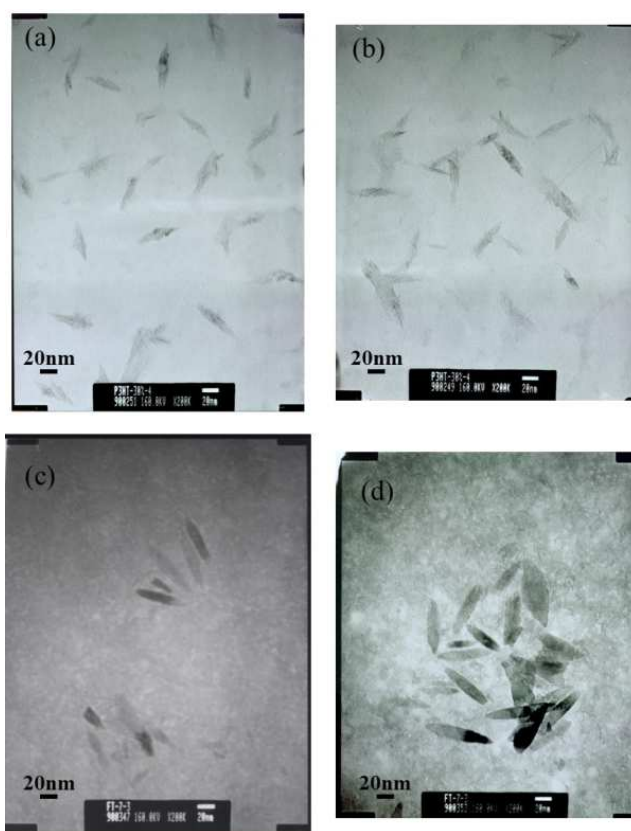| pH value	8	9	10	11	12
Mean diameter (nm) of TiO_2	7.8	7.9	8.1	8.3	8.7

Figure 3. TEM image of TiO_2 prepared at different pH values. (a) pH= 8; (b) pH= 9; (c) pH= 10; (d) pH= 11 (store temperature = 30°C).

Figure 3 shows the TEM images of the samples prepared at various pH values. The samples prepared pH 8 and 9 had small particle sizes than those prepared at pH 10 and 11. As shown in the TEM images (Figure 3), the primary particles of the sample prepared at pH 8 were rhombus shape with major axis of 20 nm and minor axis of 5 nm. At pH 11 and 12, the major axis of the rhombus form was 40-50 nm and the minor

axis was 10 nm. One can conclude that at pH of 8, well-crystallized anatase TiO_2 particles with nano size was obtained. The DLS results show that the particles distributed in 5-10 nm size were 92%, and only 8% of particles aggregated into secondary particles. One can obtain the particles size from XRD, TEM and DLS. The particle sizes of TiO_2 from these three methods are listed in Table 2. It shows that the particle sizes measured from 3 methods are consistent. All of the results show that the particle of TiO_2 were in nano size.

Table 2. The particles size distribution of TiO₂.

Instrument	XRD	TEM	DLS
Mean diameter (nm)	7.8 nm	Major axis of 20 nm Minor axis of 5 nm	Main in 5-10 nm Minor in 30 nm

Table 3 shows the effect of calcinations temperature on the pore structure of TiO_2 powder. The sample calcined at 200°C had the highest BET surface area among all samples. The average pore diameter increased with an increase of calcinations temperature.

Table 3. Surface area and pore structure of the sample from N₂ sorption.

Sample	BET surface area (m²/g)	BJH pore volume (cm³/g)	BJH pore diameter (A°)	H-K Pore diameter (A°)
pH= 8 dried at 80 °C	141	0.13	8.9	5.1
Calcined at 100°C	187	0.15	12.7	5.4
Calcined at 200°C	191	0.15	15.1	5.6
Calcined at 300°C	109	0.10	25.3	6.1

3.2. Effects of [H⁺]/[Ti] Ratio

In general, while using inorganic salt to produce TiO_2, most researchers used strong acid, such as HNO_3 or HCl, as a catalyst to promote poly-condensation reaction of the gel, and the molar ratio of $[H^+]/[Ti]$ was ~1. In this study, $[H^+]/[Ti]$ ratios of 0.6, 0.8, 1, and 1.5, respectively, were used to investigate the effect of $[H^+]/[Ti]$ ratio on the morphology of TiO_2. When $[H^+]/[Ti]$ ratio was at about 0.6 and 0.8, longer time was needed to undergo condensation reaction and crystal growth, the TiO_2 was not well-crystallized and the sol was not transparent. There was some non-dissociate orthotitanic acid that formed TiO_2. At $[H^+]/[Ti]$ ratio of 1 and 1.5, the effects of acid amount on condensation reaction and crystal growth time were observed. As $[H^+]/[Ti]$ ratio of 1.5, the time needed for condensation reaction and crystal growth was about 24 h, while at $[H^+]/[Ti]$ ratio of 1, the time needed was 48 h. As shown in Figure 5, the difference in the crystallities of the samples at $[H^+]/[Ti]$ ratio between 1.5 and 1 was lower than those of 1 and 0.6, 0.8. It should be noted that the intense XRD peak means large crystallite size. The objective of this study was to synthesize small crystals. In other word, the XRD peaks should be detectable and not intense.
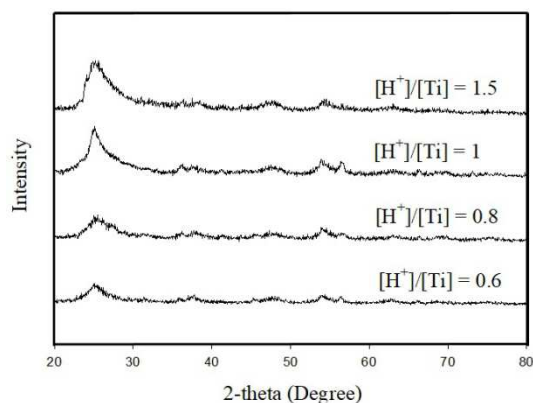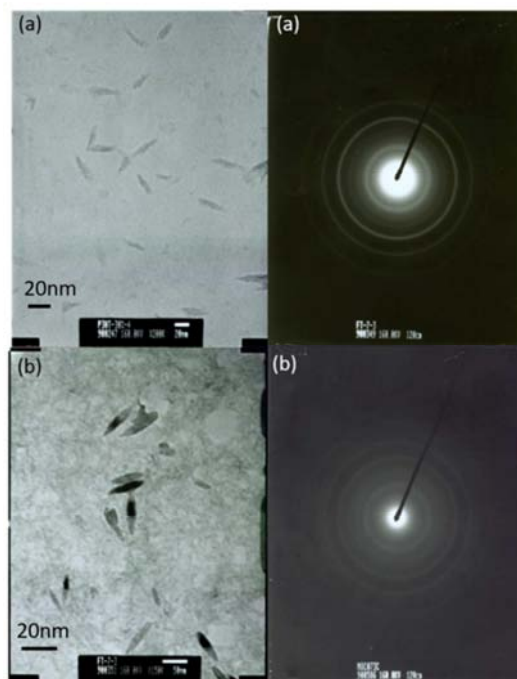

Figure 5. TEM image of TiO₂ prepared at pH=8 and (a) [H⁺]/[Ti] = 1, and (b) TEM image of TiO₂ prepared at pH=8 and [H⁺]/[Ti] = 1.5.

3.3. Effects of HCl Concentration

Figure 4. XRD patterns of TiO₂ prepared at different [H⁺]/[Ti] ratios.
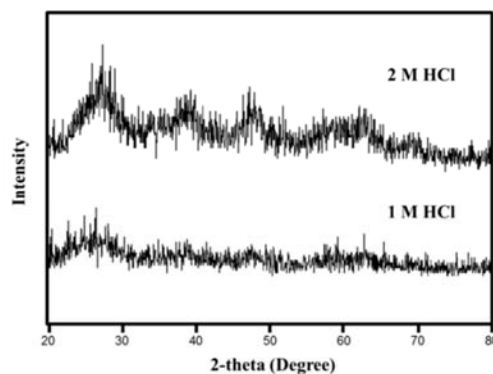

Figure 6. XRD patterns of TiO₂ prepared at different concentrations of HCl (pH =9; [H⁺]/[Ti] = 1).
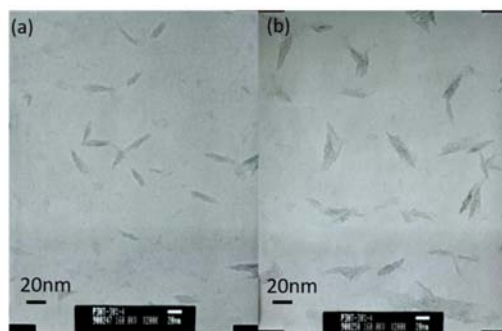
Figure 7. TEM image of TiO₂ prepared at different HCl concentration. (a) pH =8, 1M HCl; (b) pH=8, 2M HCl.

From XRD patterns (Figure 6), one can see that the crystallinity of the sample prepared by 2 M HCl was higher than that prepared by 1 M HCl. Besides, the time needed for condensation reaction and crystal growth by 2 M HCl was 36 h, which was shorter than that needed for the sample prepared by 1 M HCl. Figure 7 shows the TEM images of the samples prepared by 1 M HCl and 2 M HCl. It clears show that the sample prepared by 1 M HCl was smaller than that by 2 M HCl. Based on the above results, one can conclude that the compositions of acid, Ti, and water could affect the formation of TiO₂ nanocrystal. The higher the concentration of acid is, the higher the ratio of $[H^+]/[Ti]$ ratio is, and the shorter the time is needed for condensation reaction and crystal growth. The best crystallized nano-sized TiO₂ was obtained from the short time of crystallization, strong acid, high $[H^+]/[Ti]$ ratio. The zeta potential would change and affected the preservation of the suspension. This will be discussed in later section. If the crystalline particles with small diameter and high crystallinity is needed, one can immediately add basic solution (such as NH₄OH or NaOH aqueous solution) after condensation reaction and crystal growth to change the original pH range of acidic TiO₂ crystal particles suspension solution to their isoelectric point. After precipitation, TiO₂ particles become smaller, and its crystallinity is still high.

3.4. Effects of Stored Temperature on Particle Diameter

In this study, the temperature of preparation system was maintained at 5°C. After pH adjustment and aging, smaller orthotitanic acid particle could undergo condensation reaction and crystal growth, resulting in small diameter of TiO₂ particle. Besides, in filtering and washing steps, the diameter of orthotitanic acid decreased with the number of filtering and washing times, especially when 5°C distilled water was used in washing.

The as-synthesized TiO₂ sol was divided into two parts, one was stored at 30°C and the other at 5°C. TEM images (Figure 8) show that only a small part of the primary particles sample at 5°C aggregated to secondary particles and/or tertiary particles, especially the sol with high TiO₂ solid content. This was due to smaller kinetic energy of particles at low temperature, resulting in fewer collisions, hence double layer (Stern layer and diffuse layer) was larger than van der Waals

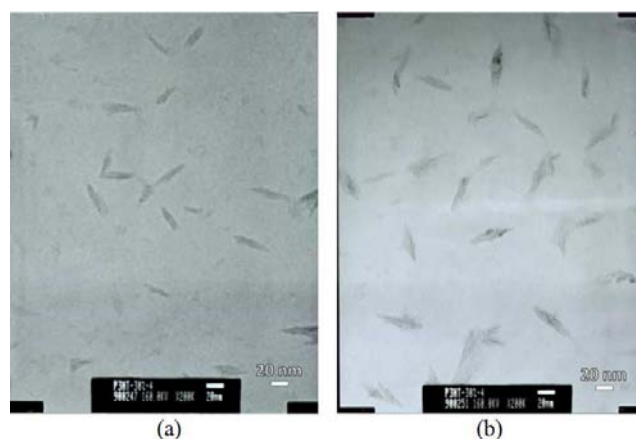force between particles, and stable suspension was obtained.

Figure 8. TEM image of TiO₂ at different stored temperatures, (a) pH= 8, 5°C; (b) pH= 8, 30°C.

3.5. Effects of HPC Surfactant

Many researchers [33] reported that adding surfactant could maintain stable suspension of particles. The surfactants play steric dispersing agent roles, which can keep particles in small diameters. [33] The surfactant can adsorb on TiO₂ surface, change its zeta potential to provide stability of the sol, and avoid the aggregation of particles. Two kinds of HPC were used as surfactants in this study, one with viscosity of 6-10 cps, and the other with viscosity of 150-400 cps. They were added to TiO₂ sol, respectively, to examine their effects on particle dispersion. After TiO₂ sol was formed, 0.005 g/ml HPC was added to suspended TiO₂ sol to disperse particle. It is found that the HPC with viscosity of 150-400 cps was better than that of HPC with viscosity of 6-10 cps. For the sample of TiO₂ sol with solid content greater than 7%, HPC with viscosity of 150-400 cps was added to increase its stored time (Figure 9). Table 4 shows that adding HPC raised the suspension stability of 7% TiO₂, and the stored time for the sample with 10% solid content increased from 14 h to 30 h. This confirms that HPC changed the zeta potential of TiO₂ particle and increased its suspension stability (Table 4).
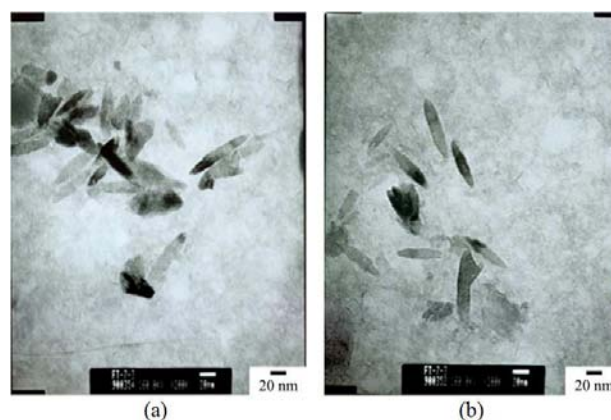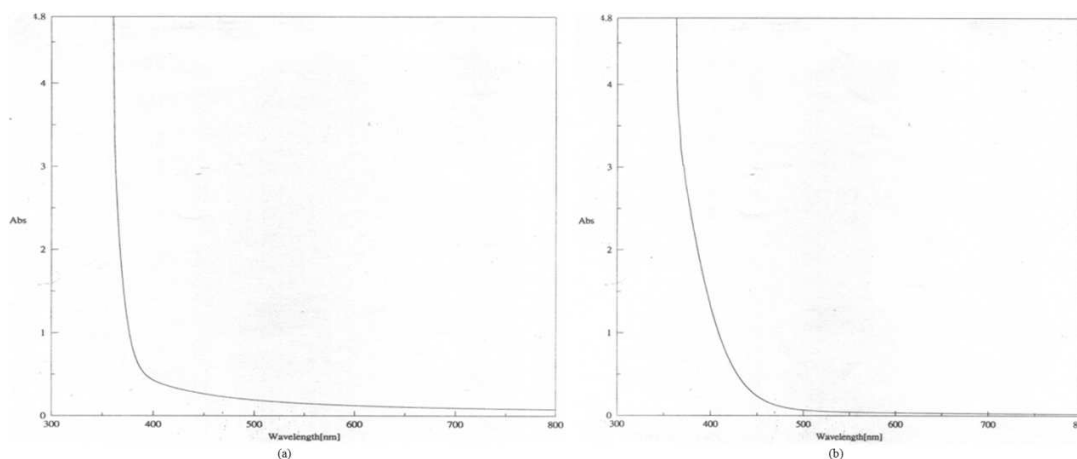

Figure 9. TEM image of TiO₂ with various solid contents by adding HPC surfactant. (a) 10%; (b) 7%.

Table 4. *Efffects of surfactantt.*

	2.5% TiO$_2$	5% TiO$_2$	7% TiO$_2$	10% TiO$_2$ [#]
Stored temp. (30°C)	Stable suspension for 3 months	Stable suspension for 3 months	Stable suspension for 14 h	Stable suspension for 13 h
Stored temp.(5°C)	Stable suspension for 3 months	Stable suspension for 3 months	Stable suspension for 26 h	Stable suspension for 14 h
Adding HPC (30°C)	Stable suspension for 3 months	Stable suspension for 3 months	Stable suspension with a little white color	Stable suspension for 26 h
Adding HPC (5°C)	Stable suspension for 3 months	Stable suspension for 3 months	Stable suspension with a little white color	Stable suspension for 30 h

3.6. UV-vis

The samples prepared in this study showed strong UV-vis absorption bands at 361 nm (Figure 10).

Figure 10. *UV-visible spectra, (a) TiO$_2$ (dried at 80 °C), (b) TiO$_2$ calcined at 150°C.*

Contact angle

TiO$_2$ was coated on glass substrate once or several times, then 40 µl or 140 µl deionized water was dropped on it. The results of contact angle show that if the glass was not washed thoroughly, it had dirt or oil substances, which would cause large degree of the contact angle. The clean glass without TiO$_2$ film attracted water molecules with its ˙OH radicals on slide glasses. The contact angle was less than 10 degrees. The transparent adherent substrates would have smaller contact angle as coating time increases. After coating to a certain amounts, the degree of contact angle would not decrease. The amount of deionized water and illuminating time of UV would also affect the contact angle. The larger amount of the deionized water is, the larger the degree the contact angle is. This study used 40 µl and 140 µl deionized water under 20 min UV light illumination (15W, wavelength 360 nm) for contact angle measurements. The results are shown in Tables 5 and 6. The transparent TiO$_2$ thin film substrates prepared in this study had super-hydrophilic surface after illumination by UV light.

Table 5. *5% TiO$_2$ coating sol on glass.*

Coating time	Contact angle (degree)	
	40 µl	140 µl
1	3.12	5.13
2	2.67	3.44
3	2.19	1.98
4	<1	1.91
5	<1	<1
I	11.23	16.53
II	22.81	28.14

I: slide glasses washed but not coating thin film.
II: slide glasses not washed and not coating thin film.

Table 6. *7% TiO$_2$ coating sol.*

Coating time	Contact angle (degree)	
	40µl	140µl
1	2.37	4.08
2	1.97	2.27
3	<1	<1
4	<1	<1
5	<1	<1

I: slide glasses washed but not coating thin film.
II: slide glasses not washed and not coating thin film.

4. Conclusion

The nanosized TiO$_2$ sol with anatase phase was successfully synthesized by hydrolysis of Ti(OH)$_4$ using HCl as the catalyst for polycondensation. The primary particles were rhombus with major axis of 20 nm and minor axis of 5 nm. It was anatase. The concentrations of acid, Ti precursor, and water could affect the formation of TiO$_2$ crystal. The higher the concentration of acid is, the higher the ratio of [H$^+$]/[Ti] ratio is, the shorter the time needed for condensation reaction and crystal growth is. The small anatase TiO$_2$ crystal could be obtained at concentration of HCl of 1 M and [H$^+$]/[Ti] ratio of 1.

Only a small part of the primary particles sample in 5°C aggregated into secondary particles. This is due to small kinetic energy of particles in low temperature resulting in few collisions; hence, double electric repulsive force (Stern layer and diffuse layer) is larger than van der Waals force between particles. When TiO$_2$ solid content was greater than 7% in the solution, HPC with viscosity of 150-400 cps was added to increase its stability. HPC could adsorb on TiO$_2$

particle surface, changed its zeta potential, and avoid aggregation of particles. The TiO_2 thin film can be heated to 150°C to increase its surface area. The high photocatalytic activity was anticipated, because it had a large surface area of 187 cm^3/g and it is microporous. The transparent TiO_2 thin film on substrate has super-hydrophilic property under UV light irradiation.

Acknowledgements

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