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Nanofabrication of Metal Oxide Nanostructures in Aqueous Solutions

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1. Introduction

Metal oxides have been widely used in electro devices, optical devices, etc. Recently, liquid phase syntheses of them attract much attention as future technology and novel academic field. Especially, liquid phase syntheses of anisotropic particles or films are expected for next generation metal oxide devices. This section describes liquid phase morphology control of anisotropic metal oxide. They were realized by precise control of nucleation and crystal growth. They showed high performance of solution systems for future metal oxide devices. Liquid phase morphology control of anisotropic metal oxide particles and films would contribute to development of metal oxide science and technology.

2. Morphology control of acicular BaTiO₃ particles¹

Acicular BaTiO₃ particles were developed using solution systems. The morphology of BaC₂O₄ · 0.5H₂O was controlled to acicular shape. Its phase transition to BaTiO₃ was realized by introducing Ti ions from the coprecipitated amorphous phase. Acicular BaTiO₃ particles have an aspect ratio as high as 18 and the particle size can be controlled by varying the growth period of BaC₂O₄ · 0.5H₂O which governs the size of BaC₂O₄ · 0.5H₂O particles. Acicular particles of crystalline BaTiO₃ can be used for ultra-thin multilayer ceramic capacitors.

Multilayer ceramic capacitors (MLCC) are indispensable electronic components for advanced electronic technology²⁻¹², but larger capacity and smaller size are needed for future electronic devices. To meet these needs, BaTiO₃ particles were downsized, but ferroelectric ceramics lose their ferroelectricity when their particle size is decreased and lose ferroelectricity entirely at a critical size²⁻¹¹. This is known as the size effect and it impedes the progress of MLCC, so a novel solution has been eagerly anticipated.

Here, we propose MLCC using acicular BaTiO₃ particles¹³. An ultra-thin ferroelectric layer and high capacity can be realized by acicular particles having a high aspect ratio. The short side provides an ultra-thin ferroelectric layer and the large volume caused by the long side avoids the loss of ferroelectricity at the critical size. Anisotropic BaTiO₃ particles are thus a candidate for MLCC. BaTiO₃ has, however, an isotropic cubic or tetragonal structure, and its morphology is extremely difficult to control due to its isotropic crystal faces. We focused on triclinic BaC₂O₄ · 0.5H₂O which has an anisotropic crystal structure and controlled the

morphology of these particles by precisely controlling crystal growth. We also achieved phase transition of $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ to crystalline BaTiO_3 by introducing Ti ions from the coprecipitated amorphous phase. Having developed several key technologies, we were successfully able to produce anisotropic acicular BaTiO_3 particles.

Morphology control of BaTiO_3 to rod-shape was reported previously. Additionally, metal oxalates (MC_2O_4) have been used for synthesis of rod-shaped oxides or hydroxides. Y. Hayashi *et al.* reported preparation of rod-shaped BaTiO_3 from rod-shaped $\text{TiO}_2 \cdot n\text{H}_2\text{O}$ and BaCO_3 in molten chloride at high temperature¹⁴. Li *et al.* reported preparation of nanoflakes and nanorods of $\text{Ni}(\text{OH})_2$, $\text{Co}(\text{OH})_2$ and Fe_3O_4 by hydrothermal conversion at 160 °C for 12h from $\text{MC}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$ in NaOH solutions¹⁵. Sun *et al.* prepared flowerlike SnC_2O_4 submicrotubes in ethanol solutions containing SnCl_2 and oxalic acid. They were annealed at 500 °C for 2 h in ambient atmosphere to obtain flowerlike SnO_2 submicrotubes¹⁶.

Oxalic acid (252 mg) was dissolved into isopropyl alcohol (4 ml) (Fig. 1)¹. Butyl titanate monomer (0.122 ml) was mixed with the oxalic acid solution, and the solution was then mixed with distilled water (100 ml). The pH of the solution was increased to pH = 7 by adding NaOH (1 M) and distilled water, while the volume of the solution was adjusted to 150 ml by these additions. The aqueous solution (50 ml) with barium acetate (39.3 mg) was mixed with the oxalic acid solution. The mixed solution containing barium acetate (0.77 mM), butyl titanate monomer (2 mM) and oxalic acid (10 mM) was kept at room temperature for several hours with no stirring, and the solution gradually became cloudy. Stirring causes the collision of homogeneously nucleated particles and destruction of large grown particles, and so was avoided in this process. The size of the precipitate was easily controlled from nanometer order to micrometer order by changing the growth period. Large particles were grown by immersion for several hours to evaluate the morphology and crystallinity in detail.

Fig. 1. Conceptual process for fabricating acicular BaTiO_3 particles. Morphology control of $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ particles and phase transition to BaTiO_3 . Reprinted with permission from Ref.¹, Masuda, Y., Yamada, T. and Koumoto, K., 2008, *Cryst. Growth Des.*, 8, 169. Copyright ©American Chemical Society

Oxalate ions ($\text{C}_2\text{O}_4^{2-}$) react with barium ions (Ba^{2+}) to form barium oxalate ($\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$). $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ is dissolved in weak acetate acid provided by barium acetate ($(\text{CH}_3\text{COO})_2\text{Ba}$), however, it can be deposited at pH 7 which is adjusted by adding NaOH. $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ was thus successfully precipitated from the solution.

Acicular particles were homogeneously nucleated and precipitated from the solution. They were on average 23 μm (ranging from 19 to 27 μm) in width and 167 μm (ranging from 144 to 189 μm) in length, giving a high aspect ratio of 7.2 (Fig. 2). They had sharp edges and

clear crystal faces, indicating high crystallinity. A gel-like solid was also coprecipitated from the solution as a second phase.

Fig. 2. SEM micrograph and XRD diffraction pattern of acicular $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ particles precipitated from an aqueous solution at $\text{pH} = 7$. XRD diffraction measurement data (first step), XRD pattern calculated from crystal structure data¹⁶ (second step) and XRD pattern of JCPDS No. 20-134 (third step) are shown for triclinic $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$. Reprinted with permission from Ref.¹, Masuda, Y., Yamada, T. and Koumoto, K., 2008, *Cryst. Growth Des.*, 8, 169. Copyright ©American Chemical Society

XRD diffraction patterns for the mixture of acicular particles and gel-like solid showed sharp diffraction peaks of crystalline $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ with no additional phase. Acicular particles were crystalline $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ and the gel-like solid would be an amorphous phase.

Fortunately, $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ has a triclinic crystal structure as shown by the model calculated from structure data¹⁷ (Fig. 2 XRD first step) and thus anisotropic crystal growth was allowed to proceed to produce an acicular shape. Each crystal face has a different surface energy and surface nature such as zeta potential and surface groups. Anisotropic crystal growth is induced by minimizing the total surface energy in ideal crystal growth. Additionally, site-selective adsorption of ions or molecules on specific crystal faces suppresses crystal growth perpendicular to the faces and so induces anisotropic crystal growth. These factors would cause anisotropic crystal growth of $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ and hence allow us to control morphology and fabricate acicular $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ particles. The positions of diffraction peaks corresponded with that of JCPDS No. 20-0134 (Fig. 2 XRD third step) and that calculated from crystal structure data¹⁷ (Fig. 2 XRD second step), however, several diffraction peaks, especially 320 and 201, were enhanced strongly compared to their relative intensity. The enhancement of diffraction intensity from specific crystal faces would be related to anisotropic crystal growth; a large crystal size in a specific crystal orientation increases the x-ray diffraction intensity for the crystal face perpendicular to the crystal orientation.

EDX elemental analysis indicated the chemical ratio of the precipitate, which included acicular particles and gel-like solid, to be about $\text{Ba} / \text{Ti} = 1$ to 1.5. The chemical ratio indicated that the coprecipitated amorphous gel contained Ti ions. Additional Ba ions can be

transformed into BaCO_3 by annealing and removed by HCl treatment in the next step. The ratio was thus controlled to slightly above $\text{Ba} / \text{Ti} = 1$ by adjusting the volume ratio of acicular particles and gel-like solid. Consequently, acicular particles of crystalline $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ with Ti-containing gel-like solid were successfully fabricated in an aqueous solution process.

In comparison, isotropic particles of barium titanyl oxalate ($\text{BaTiO}(\text{C}_2\text{O}_4)_2 \cdot 4\text{H}_2\text{O}$) were precipitated at pH 2. TiOC_2O_4 was formed by the following reaction in which the reaction of oxalic acid ($\text{H}_2\text{C}_2\text{O}_4 \cdot 2\text{H}_2\text{O}$) with butyl titanate monomer ($(\text{C}_4\text{H}_9\text{O})_4\text{Ti}$) and hydrolysis can take place simultaneously¹⁸.

$\text{TiO}(\text{C}_2\text{O}_4)$ was then converted to oxalotitanic acid ($\text{H}_2\text{TiO}(\text{C}_2\text{O}_4)_2$) by the reaction:

Alcoholic solution containing oxalotitanic acid ($\text{H}_2\text{TiO}(\text{C}_2\text{O}_4)_2$) formed by reaction (b) was subjected to the following cation exchange reaction by rapidly adding an aqueous solution of barium acetate at room temperature:

$\text{BaTiO}(\text{C}_2\text{O}_4)_2$ isotropic particles were formed by reaction (c).

On the other hand, neither $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ nor $\text{BaTiO}(\text{C}_2\text{O}_4)_2$ was precipitated at pH 3 to pH 6. Gel-like solid was formed in the solution and their XRD spectra showed no diffraction peaks. The amorphous gel that precipitated at pH = 3 to 6 would be the same as the amorphous gel coprecipitated at pH 7.

These comparisons show that the crystal growth and morphology control of $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ are sensitive to the solution conditions.

The precipitate was annealed at 750 °C for 5 h in air. Acicular $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ particles were reacted with Ti-containing amorphous gel to introduce Ti ions to transform into crystalline BaTiO_3 . X-ray diffraction of the annealed precipitate showed crystalline BaTiO_3 and an additional barium carbonate phase (BaCO_3). Excess precipitation of $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ caused the generation of barium carbonate phase (BaCO_3) as expected.

The annealed precipitate was further immersed in HCl solution (1 M) to dissolve barium carbonate (BaCO_3). Acicular particles of crystalline BaTiO_3 were successfully fabricated with no additional phase. Particles showed acicular shape with $2.8 \times 10 \times 50 \text{ } \mu\text{m}$ and x-ray diffraction of single-phase crystalline BaTiO_3 (Fig. 3). The high aspect ratio of the particles ($17.8 = 50 / 2.8$) would be provided by that of $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ particles. The particle size of acicular BaTiO_3 can be easily controlled by the growth period and solution concentration for $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ precipitation which decides the particle size of $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$.

BaTiO_3 has a cubic crystal structure at high temperature above phase transition and has a tetragonal crystal structure at room temperature. The cubic crystal structure is completely isotropic and the tetragonal crystal structure results from stretching a cubic lattice along one of its lattice vectors. For both of the crystal structures it is difficult to control anisotropic crystal growth, however, with our newly developed process we could successfully control the morphology and fabricate acicular particles. This was achieved by controlling the morphology of triclinic $\text{BaC}_2\text{O}_4 \cdot 0.5\text{H}_2\text{O}$ to acicular shape and the phase transition to BaTiO_3 .

by introducing Ti ions from the coprecipitated amorphous phase. The novel concept can be applied to a wide variety of morphology control and crystal growth control for advanced electronic devices composed of crystalline materials.

Fig. 3. SEM micrograph and XRD diffraction pattern of acicular BaTiO₃ particles after annealing at 750 °C for 5 h and HCl treatment. XRD diffraction measurement data (first step) and XRD pattern of JCPDS No. 05-0626 (second step) are shown for tetragonal BaTiO₃. Reprinted with permission from Ref.¹, Masuda, Y., Yamada, T. and Koumoto, K., 2008, *Cryst. Growth Des.*, 8, 169. Copyright ©American Chemical Society

3. Morphology control of ZnO particles^{19,20}

Ethylenediamine (H₂N-CH₂CH₂-NH₂, 15–45 mM, Sigma-Aldrich) was added to the zinc acetate aqueous solution (Zn(CH₃COO)₂, 15 mM, Kishida Chemical Co., Ltd.) to promote deposition of ZnO¹⁹. Zinc chelate (Zn(H₂N-CH₂CH₂-NH₂)₂²⁺) was formed from zinc acetate and ethylenediamine in reaction (a). ZnO was crystallized from zinc chelate and hydroxide ion (OH⁻) in reaction (c).

The solution became turbid shortly after adding ethylenediamine. The molar ratio of ethylenediamine to Zn was [ethylenediamine] / [Zn] = (a) 1 : 1, (b) 2 : 1 or (c) 3 : 1. pH of the solutions were (a) pH=7.3, (b) pH=8.0 or (c) pH=8.7, respectively. Crystal growth rate and deposition of ZnO were attempted to control to change particle morphology. Si substrate (Newwingo Co., Ltd.) was immersed to evaluate deposited ZnO particles and particulate films. The solution in a glass beaker was kept at 60 °C for 3 h using a water bath. The silicon substrate was cleaned before immersion as described in references. The substrate was rinsed with distilled water after immersion.

ZnO particles having hexagonal cylinder shape were homogeneously nucleated and deposited in the aqueous solution containing 15 mM ethylenediamine ([ethylenediamine] / [Zn] = (a) 1 : 1) (Fig. 4a). X-ray diffraction patterns showed the deposition to be well crystallized ZnO (Fig. 5a). The relative intensity of (10-10) and (0002) is similar to that of randomly deposited ZnO particles, indicating the random orientation of deposited ZnO hexagonal cylinders, which is consistent with SEM observations. Crystals showed hexagonal facets of about 100–200 nm in diameter and about 500 nm in length. ZnO has a hexagonal crystal structure and thus hexagonal cylinders can be obtained by sufficiently slow crystal growth. A low crystal growth rate allows enough ions to diffuse to form a complete crystal structure.

Fig. 4. SEM micrographs of (a) ZnO hexagonal cylinder particles, (b) ZnO long ellipse particles, (c1) ZnO hexagonal symmetry radial whiskers and (c2) magnified area of (c1). Reprinted with permission from Ref.¹⁹, Y. Masuda, N. Kinoshita and K. Koumoto, *J. Nanosci. Nanotechnol.*, in press. Copyright ©American Scientific Publisher

Fig. 5. XRD patterns of (a) ZnO hexagonal cylinder particles, (b) ZnO long ellipse particles, (c) ZnO hexagonal symmetry radial whiskers and ZnO (JCPDS No. 36-1451). Reprinted with permission from Ref.¹⁹, Y. Masuda, N. Kinoshita and K. Koumoto, *J. Nanosci. Nanotechnol.*, in press. Copyright ©American Scientific Publisher

ZnO having hexagonal cylinder shape showed strong photoluminescence intensity in the UV region at about 370–400 nm and weak intensity in the visible light region at about 530–550 nm by 350 nm excitation light which appears in spectra (Fig. 6a). ZnO crystals were reported to show UV luminescence (around 390 nm) attributed to band-edge luminescence and visible-light luminescence caused from oxygen vacancy (450–600 nm)^{21,22}. Oxygen vacancies would be generated in ZnO during crystallization to show visible-light luminescence.

Concentration of ethylenediamine was increased twice to [ethylenediamine] / [Zn] = (b) 2 : 1. ZnO particles with long ellipse shape were deposited homogeneously from the solution (Fig. 4b). ZnO particles were about 100–200 nm in diameter and about 500 nm in length, and were similar to those of hexagonal cylinder shape. XRD showed the deposition to be well crystallized ZnO (Fig. 5b). Relative intensity of (10-10) diffraction is much stronger than that of (0002), indicating that mainly ZnO particles with long ellipse shape were laid on the silicon substrate. This was also observed in SEM micrographs (Fig. 4b). Deposition speed of ZnO with long ellipse shape was slightly faster than that of ZnO with hexagonal cylinder shape because of the high concentration of ethylenediamine. Ethylenediamine accelerates the crystallization of ZnO. In other words, supersaturation degree of the solution was increased by increasing the ethylenediamine concentration. As a result, ZnO grew slightly faster and formed not sharp hexagonal facets but rounded hexagonal cylinders, i.e., long ellipse shape. The photoluminescence spectrum of ZnO with long ellipse shape (Fig. 6b) was similar to that of ZnO with hexagonal cylinder shape (Fig. 6a). The luminescence property

was clearly shown to be stable and not influenced by the synthesis conditions in the range of [ethylenediamine] / [Zn] = 1 : 1 to 2 : 1. High repeatability and stability of the photoluminescence property without being affected by the deposition conditions are major advantages of this system for large-scale production.

Fig. 6. Photoluminescence spectra of (a) ZnO hexagonal cylinder particles, (b) ZnO long ellipse particles and (c) ZnO hexagonal symmetry radial whiskers. Reprinted with permission from Ref.¹⁹, Y. Masuda, N. Kinoshita and K. Koumoto, *J. Nanosci. Nanotechnol.*, in press. Copyright ©American Scientific Publisher

Further control the morphology was attempted by increasing the supersaturation degree. Ethylenediamine was added to the solution to be [ethylenediamine] / [Zn] = (c) 3 : 1 in the deposition process. ZnO whiskers were successfully grown and deposited from the solution. The morphology was drastically changed by the precise investigation and control of solution conditions. The whiskers were about 10–100 nm in diameter and about 1000–2000 nm in length (Fig. 4c1). A high aspect ratio was realized by high crystal growth rate of end faces to elongate the whiskers. Details of morphology were further evaluated. They were dispersed in water and dropped on a silicon substrate. Whiskers were connected to form hexagonal symmetry radial whiskers (Fig. 4c2). Six whiskers connected to form one particle. They had hexagonal symmetry. Tips of whiskers were finer than that at center of the particles. XRD showed the whiskers to be well crystallized ZnO (Fig. 5c). The relative intensity of (10-10) diffraction is much stronger than that of (0002), showing that mainly ZnO whiskers were laid on the substrate. This was also observed in SEM micrographs (Fig 4

c). Deposition of ZnO whiskers having high aspect ratio on the substrate would provide a ZnO network film having a high specific surface area. These whisker films can be applied to gas sensors^{23,24} or solar cells²⁵ which require high specific surface area. The whisker films also have high conductivity per unit volume²⁶ compared to conventional particulate films or mesoporous materials because the whiskers carry an electric current for a long distance without grain boundaries. Photoluminescence intensity in the visible light region was quite different from that of ZnO with hexagonal cylinder shape or long ellipse shape (Fig. 5c). The large change of morphology indicates that the crystal growth mechanism differs greatly between ZnO particles having hexagonal cylinder or long ellipse shape and ZnO whiskers. Basically, ethylenediamine increased crystal growth rate to generate oxygen vacancies. The oxygen vacancies increased photoluminescence intensity in visible light region. However, hexagonal symmetry radial whiskers prepared in the solution of [ethylenediamine] / [Zn] = (c) 3 : 1 showed very weak photoluminescence intensity in visible light region compared to hexagonal cylinders or long ellipse particles. Excess ethylenediamine would decrease photoluminescence intensity in visible light region. Ion concentration of the solution of [ethylenediamine] / [Zn] = (c) 3 : 1 would drastically change at the initial stage. The solutions of [ethylenediamine] / [Zn] = (a) 1 : 1 and (b) 2 : 1 became turbid shortly after adding ethylenediamine and they changed to transparent gradually. The solutions of [ethylenediamine] / [Zn] = (c) 3 : 1 also became turbid shortly after adding ethylenediamine, however, they changed to transparent rapidly compared to that of [ethylenediamine] / [Zn] = (a) 1 : 1 and (b) 2 : 1. It indicated that ions were consumed to form particles rapidly in the solution of [ethylenediamine] / [Zn] = (c) 3 : 1 and ion concentration would decrease drastically with color change. It can be assumed that high concentration of ethylenediamine increased crystal growth rate to form ZnO particles at the first stage. The particles generated in the first stage would not be whiskers but small ZnO particles. Ion concentration decreased rapidly to make solution transparent by formation of small particles because Zn ions were consumed in formation of the particles. ZnO whiskers would grow slowly on the small particles in dilute solutions at second stage. Consequently, hexagonal symmetry radial whiskers were formed in the solution of [ethylenediamine] / [Zn] = (c) 3 : 1. This phenomenon was consistent with reported ZnO whiskers which had high crystallinity, high photoluminescence intensity in UV region and low photoluminescence intensity in visible light region. Novel properties such as unique morphology, high specific surface area, high conductivity per unit volume, low photoluminescence intensity in visible light region and high photoluminescence intensity in UV region may pave the way to a new age of ZnO devices. Furthermore, they can be fabricated on low heat resistant materials such as polymers, paper or organic materials for flexible devices.

4. Morphology control of multi-needle ZnO particles and their particulate films²⁷

ZnO has attracted much attention as a next-generation gas sensor for CO²⁸⁻³⁰, NH₃³¹, NO₂³², H₂S³³, H₂^{29,34}, ethanol^{34,35}, SF₆³⁴, C₄H₁₀³⁴ or gasoline³⁴ and dye-sensitized solar cells^{26,36-39}. Sensitivity directly depends on the specific surface area of the sensing material. ZnO particles, particulate films or mesoporous material having high specific surface area were thus strongly required.

ZnO has been crystallized to a hexagonal cylinder shape for gas sensors or solar cells in many studies²⁶ by using the hexagonal crystal structure of ZnO at low supersaturation

degree. However, strategic morphology design and precise morphology control for high specific surface area should be developed to improve the properties. ZnO particles should be controlled to have multi-needles or high surface asperity to increase the specific surface area by the crystallization at high supersaturation degree.

Recently, morphology control^{20,40-42} and nano/micro manufacturing⁴³⁻⁴⁶ of oxide materials were proposed in solution systems. Solution systems have the advantage of adjustment of supersaturation degree and high uniformity in the system for particle morphology control. However, many factors affect the system compared to gas phase systems or solid state reactions. Solution chemistry for oxide materials is therefore being developed and many areas remain to be explored.

Morphology control of ZnO particles to hexagonal cylinder shape, ellipse shape and multi-needle shape was recently developed²⁰. Photoluminescence property was improved by changing the morphology and oxygen vacancy volume in this system. Morphology control of ZnO has also been proposed based on control of crystal growth⁴⁷⁻⁵⁰. W. Q. Peng et al. reported flower-like bunches synthesized on indium-doped tin oxide glass substrates through a chemical bath deposition process⁴⁷. Y. R. Lin et al. fabricated nanowires on a ZnO-buffered silicon substrate by a hydrothermal method⁴⁸. H. Zhang et al. prepared flowerlike, disklike, and dumbbell-like ZnO microcrystals by a capping-molecule-assisted hydrothermal process⁴⁹. K. H. Liu et al. reported a hierarchical polygon prismatic Zn-ZnO core-shell structure grown on silicon by combining liquid-solution colloids together with the vapor-gas growth process⁵⁰. These studies showed high morphology controllability of ZnO, however, morphology should be optimized to have high specific surface area to apply to solar cells or gas sensors.

In this study, the morphology design of ZnO particles was proposed for solar cells or gas sensors in which high specific surface area, high electrical conductivity and high mechanical strength are required. Multi-needle ZnO particles having ultrafine surface relief structure, as well as particulate films constructed from multi-needle particles and thin sheets were fabricated²⁷. Morphology control was realized based on a new idea inspired from the morphology change in our former study²⁰. High supersaturation degree of the solution was used for fast crystal growth which induces the formation of multi-needle particles and low super saturation was used for the formation of ZnO thin sheets.

Zinc nitrate hexahydrate ($\text{Zn}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, > 99.0%, MW 297.49, Kanto Chemical Co., Inc.) and ethylenediamine ($\text{H}_2\text{NCH}_2\text{CH}_2\text{NH}_2$, > 99.0%, MW 60.10, Kanto Chemical Co., Inc.) were used as received. Glass (S-1225, Matsunami Glass Ind., Ltd.) was used as a substrate. Zinc nitrate hexahydrate (15 mM) was dissolved in distilled water at 60 °C and ethylenediamine (15 mM) was added to the solution to induce the formation of ZnO. Glass substrate was immersed in the middle of the solution at an angle and the solution was kept at 60 °C using a water bath for 80 min with no stirring. The solution became clouded shortly after the addition of ethylenediamine. Ethylenediamine plays an essential role in the formation of crystalline ZnO. ZnO was homogeneously nucleated and grown to form a large amount of particles to make the solution clouded. ZnO particles were gradually deposited and further grown on a substrate. Homogeneously nucleated particles precipitated gradually and the solution became light white after 80 min. The supersaturation degree of the solution was high at the initial stage of the reaction for the first 1 h and decreased as the color of the solution changed.

ZnO particulate films constructed from ZnO particles and thin sheets were fabricated by immersion for 48 h. The glass substrate was immersed in the middle of the solution at an

angle and the solution was kept at 60 °C using a water bath for 6 h with no stirring. The solution was then left to cool for 42 h in the bath. The solution became clouded shortly after the addition of ethylenediamine and clear after 6 h. The bottom of the solution was covered with white precipitate after 6 h. The supersaturation degree of the solution was high at the initial stage of the reaction for the first 1 h and then decreased as the color of the solution changed.

Morphology of ZnO particles and particulate films was observed by a field emission scanning electron microscope (FE-SEM; JSM-6335FM, JEOL Ltd.) after heating at 150 °C for 30 min in vacuum for drying of carbon paste (Vacuum oven, VOS-201SD, EYELA, Tokyo Rikakikai. Co., Ltd.) and Pt coating for 3 nm (Quick cool coater, SC-701MCY, Sanyu Electronic Company). Crystal phases were evaluated by an X-ray diffractometer (XRD; RINT-2100V, Rigaku) with CuK α radiation (40 kV, 40 mA).

Morphology design of ZnO particles and particulate films. The morphology of ZnO particles was designed to increase the specific surface area, electrical conductivity and mechanical strength of the base material of solar cells and sensors (Fig. 7). Typical ZnO particles grown at low supersaturation degree are shown in Fig. 7a²⁰. The particles show edged hexagonal faces and elongate parallel to the *c*-axis. ZnO particles grow to have a hexagonal cylinder shape by slow crystal growth due to the hexagonal crystal structure of ZnO. The morphology of ZnO particles was controlled to have a multi-needle shape in an aqueous solution (Fig. 7b)²⁰. Multi-needle particles have a high specific surface area compared to hexagonal cylinder particles, but particulate films constructed from small particles have many grain boundaries which reduce the electrical conductivity (Fig. 7c). Particles should thus have large grain size to decrease grain boundaries and increase electrical conductivity (Fig. 7d). Furthermore, the specific surface area of ZnO particles should be increased to improve the sensing performance of sensors or generating efficiency of solar cells (Fig. 7e). An ultrafine surface relief structure on ZnO particles is a candidate

Fig. 7. Design and morphology control of ZnO particles and particulate films for high specific surface area, high electrical conductivity and high mechanical strength. Reprinted with permission from Ref.²⁷, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 2633. Copyright ©American Chemical Society

morphology for increasing the specific surface area. Additionally, particles should be connected to each other and with a substrate by a combined member such as ZnO thin sheets (Fig. 7f). The thin sheets increase the mechanical strength of particulate films and help raise the electrical conductivity and specific surface area. Morphology control of ZnO particles was attempted based on the strategic morphology design for application to sensors.

Morphology control of ZnO particles. After having been immersed in the solution for 80 min, the substrate covered with ZnO particles was evaluated by SEM and XRD. ZnO particles were shown to be multi-needle shape in which many needles were grown from the center of the particles (Fig. 8). The particles have more needles compared with the particles previously reported which were constructed from two large needles and several small needles²⁰. The size of particles was in the range from 1 – 5 μm which is larger than the particles prepared previously²⁰ (Fig. 7b). Needles were constructed from an assembly of narrow acicular crystals and thus the side surfaces of needles were covered with arrays of pleats. The tips of the needles were rounded V-shape with many asperities. Edged hexagonal shapes were observed at the tips of needles, thus clearly showing high crystallinity and the direction of the *c*-axis. The *c*-axis would be the long direction of multi-needles and narrow acicular crystals. Elongation of the *c*-axis observed by SEM is consistent with high diffraction intensity of 0002 (Fig. 9). The 0002 diffraction intensity of multi-needle ZnO particles was much stronger than 1010 or 1011 peaks though 0002 diffraction is weaker than 1010 or 1011 diffractions in randomly orientated ZnO particles (JCPDS No. 36-1451). High diffraction intensity from (0002) planes which are perpendicular to the *c*-axis would be caused from the crystalline ZnO particles which grew to elongate the *c*-axis. The particles have more stacks of (0002) crystal planes compared to that of (1010) planes which are parallel to the *c*-axis or (1011) planes and hence the intensity from (0002) planes was stronger than that from (1010) or (1011) planes.

Fig. 8. SEM micrograph (a-d) of multi-needle ZnO particles having ultrafine surface relief structure. Reprinted with permission from Ref.²⁷, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 2633. Copyright ©American Chemical Society

ZnO grows to a hexagonal cylinder shape at low supersaturation degree because of its hexagonal crystal structure, however, ZnO grows to a multi-needle shape at high supersaturation degree which induces fast crystal growth. ZnO was thus grown to a multi-needle shape in our solution in spite of its hexagonal crystal structure. The growth of ZnO was halted by a rapid decrease of supersaturation degree and removal of particles from the solution to obtain ZnO multi-needle particles having an ultrafine surface relief structure. The morphology of the ZnO particles was controlled by the fast crystal growth due to high supersaturation degree and by the suppression of crystal growth due to the rapid decrease of supersaturation degree and removal of particles from the solution.

Fig. 9. XRD diffraction pattern of multi-needle ZnO particles having ultrafine surface relief structure. Reprinted with permission from Ref.²⁷, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 2633. Copyright ©American Chemical Society

Morphology control of ZnO particulate films. ZnO particulate films showed a multi-needle shape and were connected to each other by thin sheets (Fig. 10a-c). The morphology of the particles was similar to that of the particles prepared by immersion for 80 min to have a high specific surface area. Thin sheets had a thickness of 10 – 50 nm and width of 1 – 10 μm and were connected to particles closely with no clearance. The particulate films had continuous open pores ranging from several nm to 10 μm in diameter. The particulate films showed x-ray diffraction patterns of ZnO crystal with no additional phase (Fig. 11). Diffraction peaks were very sharp, showing high crystallinity of the particulate films. The high intensity of 0002 would be caused by elongation of multi-needle particles in the *c*-axis direction which increases the stacks of (0002) crystal planes.

ZnO multi-needle particles having an ultrafine surface relief structure were prepared at 60 $^{\circ}\text{C}$ in the white solution during the initial 80 min (Fig. 7e). The supersaturation degree was high at the initial stage of the reaction due to the high concentration of ions. ZnO particles were then precipitated, making the bottom of the solution white and the solution itself clear. Ions were consumed to form ZnO particles and thus the ion concentration of the solution decreased rapidly. Thin sheets were formed at 25 $^{\circ}\text{C}$ in the clear solution after the formation of multi-needle particles (Fig. 7f). Solution temperature and supersaturation degree would

influence on precipitates. Consequently, the particulate films constructed from multi-needle particles and thin sheets were successfully fabricated by the two-step growth (Fig. 7f).

Fig. 10. SEM micrograph (a-d) of ZnO particulate films constructed from ZnO multi-needle particles and thin sheets. Reprinted with permission from Ref.²⁷, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 2633. Copyright ©American Chemical Society

Fig. 11. XRD diffraction pattern of ZnO particulate films constructed from ZnO multi-needle particles and thin sheets. Reprinted with permission from Ref.²⁷, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 2633. Copyright ©American Chemical Society

For comparison, assemblies of thin sheets were prepared at air-liquid interfaces of the same solution we used in this study⁵¹. XRD patterns of the sheets were assigned to ZnO. The sheets had *c*-axis orientation parallel to the sheets, i.e., in-plane *c*-axis orientation. TEM observations showed the sheets were dense polycrystals consisted of nano-sized ZnO crystals. Electron diffraction pattern showed strong isotropic diffraction ring from (0002) planes. It suggested in-plane *c*-axis orientation of ZnO crystals which was consistent with XRD evaluations. Mechanical strength and electrical property would be affected by crystal

orientation and microstructures. The sheets would have stronger mechanical strength compared to porous sheets because of their dense structure. They would have different electrical properties from randomly oriented sheets due to in-plane *c*-axis orientation, because ZnO has anisotropic electrical properties caused from anisotropic hexagonal crystal structure. The sheets prepared at air-liquid interfaces⁵¹ would be similar to that prepared in the solutions (Fig. 10) because both of them were prepared from the same solution and showed XRD patterns assigned to ZnO⁵¹ (Fig. 11). The sheets prepared in this study (Fig. 4, 5) would have similar mechanical and electrical properties to the sheets prepared at air-liquid interfaces⁵¹.

Thin sheets were transformed to particles and porous particulate films by annealing at 500 °C for 1 h in air. The sheets did not maintain their thin sheet shape due to high slimness and/or phase transformation. Thin sheets would be inorganic films containing Zn ions such as crystalline ZnO, amorphous ZnO or zinc hydroxide and were transformed to porous ZnO particulate films by the annealing. Further investigation of the thin sheets would contribute to more precise morphology control of ZnO structure and further improvement of specific surface area. Additionally, precise evaluation of mechanical strength and electrical properties should be performed to clarify the potential of ZnO particulate films for sensors or solar cells and to produce guidelines for improving their properties.

Multi-needle ZnO particles having an ultrafine surface relief structure were successfully fabricated by the precise control of crystal growth in an aqueous solution. The morphology of ZnO was further controlled for ZnO particulate films constructed from ZnO multi-needle particles and thin sheets. The thin sheets connected particles to each other and with a substrate. The morphology design and morphology control described here will facilitate the progress of crystal science for developing future advanced materials and devices.

5. Morphology control of high *c*-axis oriented stand-alone ZnO self-assembled film⁵¹

Recently, crystalline ZnO^{20,26,47-50,52-59} have been synthesized to utilize the high potential of the solution process for future devices and to realize green chemistry for a sustainable society. For instance, ZnO nanowire arrays have been synthesized using seed layers in aqueous solutions for dye-sensitized solar cell^{26,52}. Full sun efficiency of 1.5% is demonstrated in this study. O'Brien et al. prepared specular ZnO films consisting of clumps of elongated triangular crystals⁵³, small ZnO spherical clumps consisting of particles of ca. 100 nm⁵³, ZnO films consisting of randomly rod-shaped particles of up to 1000 nm in length⁵³, ZnO films consisting of very thin random rod-shaped particles of ca. 1000 nm⁵³ and specular films consisting of flowers with well-formed triangle features⁵⁹ in aqueous solutions.

However, ZnO films have usually been prepared on substrates^{54,57,60-76}, and in particular crystalline ZnO films having high *c*-axis orientation require expensive substrates such as single crystals or highly-functional substrates. A simple and low-cost process for self-supporting crystalline ZnO films is expected to be used for a wide range of applications such as windows of optical devices or low-value-added products. Self-supporting crystalline ZnO films can also be applied by being pasting on a desired substrate such as low heat-resistant polymer films, glasses, metals or papers.

In this section, high *c*-axis oriented stand-alone ZnO self-assembled films was fabricated using an air-liquid interface⁵¹. ZnO was crystallized from an aqueous solution without heat

treatment or catalyst. The ZnO film was fabricated at the air-liquid interface without using ammonia vapor.

Zinc nitrate hexahydrate ($\text{Zn}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$, > 99.0%, MW 297.49, Kanto Chemical Co., Inc.) and ethylenediamine ($\text{H}_2\text{NCH}_2\text{CH}_2\text{NH}_2$, > 99.0%, MW 60.10, Kanto Chemical Co., Inc.) were used as received. Zinc nitrate hexahydrate (15 mM) was dissolved in distilled water at 60°C and ethylenediamine (15 mM) was added to the solution to induce the formation of ZnO. The solution was kept at 60°C using a water bath for 6 h with no stirring. The solution was then left to cool for 42 h in the bath. Polyethylene terephthalate (PET) film, glass (S-1225, Matsunami Glass Ind., Ltd.) and an Si wafer (p-type Si [100], NK Platz Co., Ltd.) were used as substrates.

Morphology of ZnO film was observed by a field emission scanning electron microscope (FE-SEM; JSM-6335FM, JEOL Ltd.) and a transmission electron microscope (TEM; H-9000UHR, 300 kV, Hitachi). Crystal phase was evaluated by an X-ray diffractometer (XRD; RINT-2100V, Rigaku) with $\text{CuK}\alpha$ radiation (40 kV, 40 mA). Si wafer was used as a substrate for XRD evaluation. The crystal structure model and diffraction pattern of ZnO were calculated from ICSD (Inorganic Crystal Structure Database) data No. 26170 (FIZ Karlsruhe, Germany and NIST, USA) using FindIt and ATOMS (Hulinks Inc.).

The solution became clouded shortly after the addition of ethylenediamine by the homogeneous nucleation and growth of ZnO particles. ZnO particles were gradually deposited to cover the bottom of the vessel, and the solution became light white after 1 h and clear after 6 h. The supersaturation degree of the solution was high at the initial stage of the reaction for the first 1 h and decreased as the color of the solution changed.

White films were formed at the air-liquid interface and they grew to large films. The films had sufficiently high strength to be obtained as stand-alone films. Additionally, a film was scooped to past onto a desired substrate such PET film, Si wafer, glass plate or paper, and the pasted ZnO film was then dried to bond it to the substrate. Both sides of the film can be pasted on substrate. The film physically adhered to the substrate. The film maintained its adhesion during immersion in lightly ultrasonicated water, however, it can be easily peeled off again by strong ultrasonication. The film can be handled easily from substrate to other substrate. It also can be attached strongly to substrate by annealing or addition of chemical reagents such as silane coupling agent to form chemical bonds between the film and the substrate

The film grew to a thickness of about 5 μm after 48 h, i.e., 60°C for 6 h, and was left to cool for 42 h.

The air side of the stand-alone film had a smooth surface over a wide area due to the flat air-liquid interface (Fig. 12-a1), whereas the liquid side of the film had a rough surface (Fig. 12-b1). The films consisted of ZnO nano-sheets were clearly observed from the liquid side (Fig. 12-b2) and the fracture edge-on profile of the film (Fig. 12-c1, 12-c2). The nano-sheets had a thickness of 5-10 nm and were 1-5 μm in size. They mainly grew forward to the bottom of the solution, i.e., perpendicular to the air-liquid interface, such that the sheets stood perpendicular to the air-liquid interface. Thus, the liquid side of the film had many ultra-fine spaces surrounded by nano-sheet and had a high specific surface area. The air side of the film, on the other hand, had a flat surface that followed the flat shape of the air-liquid interface. The air-liquid interface was thus effectively utilized to form the flat surface of the film. This flatness would contribute to the strong adhesion strength to substrates for pasting of the film. The air-side surface prepared for 48 h had holes of 100-500 nm in diameter (Fig. 12-a2), and were hexagonal, rounded hexagonal or round in shape. The air-side surface prepared for 6 h, in contrast, had no holes on the surface. The air-side surface was well

crystallized to form a dense surface and ZnO crystals would partially grow to a hexagonal shape because of the hexagonal crystal structure. Well-crystallized ZnO hexagons were then etched to form holes on the surface by decrease in pH. The growth face of the film would be liquid side. ZnO nano-sheets would grow to form a large ZnO film by Zn ion supply from the aqueous solution. Further investigation of the formation mechanism would contribute to the development of crystallography in the solution system and the creation of novel ZnO fine structures.

Fig. 12. SEM micrographs of high c -axis oriented stand-alone ZnO self-assembled film. (a1) Air-side surface of ZnO film. (a2) Magnified area of (a1). (b1) Liquid-side surface of ZnO film. (b2) Magnified area of (b1). (c1) Fracture cross section of ZnO film from air side. (c2) Magnified area of (c1). Reprinted with permission from Ref.⁵¹, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 275. Copyright ©American Chemical Society

The film showed a very strong 0002 x-ray diffraction peak of hexagonal ZnO at $2\theta = 34.04^\circ$ and weak 0004 diffraction peak at $2\theta = 72.16^\circ$ with no other diffractions of ZnO (Fig. 13). (0002) planes and (0004) planes were perpendicular to the c -axis, and the diffraction peak only from (0002) and (0004) planes indicates high c -axis orientation of ZnO film. The inset figure shows that the crystal structure of hexagonal ZnO stands on a substrate to make the c -axis perpendicular to the substrate. Crystallite size parallel to (0002) planes was estimated from the half-maximum full-width of the 0002 peak to 43 nm. This is similar to the threshold limit value of our XRD equipment and thus the crystallite size parallel to (0002) planes is estimated to be greater than or equal to 43 nm. Diffraction peaks from a silicon substrate were observed at $2\theta = 68.9^\circ$ and $2\theta = 32.43^\circ$. Weak diffractions at $2\theta = 12.5^\circ$, 24.0° , 27.6° , 30.5° , 32.4° and 57.6° were assigned to co-precipitated zinc carbonate hydroxide ($\text{Zn}_5(\text{CO}_3)_2(\text{OH})_6$, JCPDS No. 19-1458).

Fig. 13. XRD diffraction pattern of high c -axis oriented stand-alone ZnO self-assembled film. Reprinted with permission from Ref.⁵¹, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 275. Copyright ©American Chemical Society

Stand-alone ZnO film was further evaluated by TEM and electron diffraction. The film was crushed to sheets and dispersed in an acetone. The sheets at the air-liquid interface were skimmed by a copper grid with a carbon supporting film. The sheets were shown to have uniform thickness (Fig. 14a). They were dense polycrystalline films constructed of ZnO nanoparticles (Fig. 14b). Lattice image was clearly observed to show high crystallinity of the particles. The film was shown to be single phase of ZnO by electron diffraction pattern. These observations were consistent with XRD and SEM evaluations.

The film pasted on a silicon wafer was annealed at 500°C for 1 h in air to evaluate the details of the films. ZnO film maintained its structure during the annealing (Fig. 15). The air side of the film showed a smooth surface (Fig. 15-a1) and the liquid side showed a relief structure having a high specific surface area (Fig. 15-b1, 15-b2). The air side showed the film consisted of dense packing of small ZnO nanosheets and the size of sheets increased toward the liquid-side surface (Fig. 15-a2). ZnO sheets would grow from the air side to the liquid side, i.e., the sheets would nucleate at the liquid-air interface and grow down toward the bottom of the solution by the supply of Zn ions from the solution. Annealed film showed X-ray diffractions of ZnO and Si substrate with no additional phases. As-deposited ZnO nanosheets were shown to be crystalline ZnO because the sheets maintained their fine structure during the annealing without any phase transition. High c -axis orientation was also maintained during the annealing, showing a very strong 0002 diffraction peak.

Fig. 14. (a) TEM micrograph of ZnO nano-sheets. (b) Magnified area of (a). (Insertion) Electron diffraction pattern of ZnO. Reprinted with permission from Ref.⁵¹, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 275. Copyright @American Chemical Society

Fig. 15. SEM micrographs of high *c*-axis oriented stand-alone ZnO self-assembled film annealed at 500°C for 1 h in air. (a1) Fracture edge-on profile of ZnO film from air side. (a2) Cross-section profile of ZnO film from air side. (b1) Fracture edge-on profile of ZnO film from liquid side. (b2) Cross-section profile of ZnO film from liquid side. Reprinted with permission from Ref.⁵¹, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 275. Copyright @American Chemical Society

The solution was further kept at 25°C for 1 month to evaluate the details of the deposition mechanism. The film prepared at the air-liquid interface for 1 month was not hexagonal ZnO. The film showed strong X-ray diffractions of zinc carbonate hydroxide single phase. ZnO would be dissolved by decrease in pH. ZnO would be crystallized at the initial reaction stage for the first 48 h. ZnO was then gradually etched and dissolved by nitric acid and zinc carbonate hydroxide was crystallized using Zn ions which were supplied by the dissolution of crystalline ZnO.

High *c*-axis oriented stand-alone ZnO self-assembled film was fabricated using a simple solution process. The film consisted of ZnO nanosheets was crystallized at air-liquid interface. The nanosheets grew perpendicular to the film. The film had high *c*-axis orientation and showed a strong 0002 diffraction peak and weak 0004 peak. The air side of the film had a flat surface, whereas the liquid side had a rough surface having many ultra-fine spaces surrounded by ZnO nano-sheets. The rough surface of the liquid side can be utilized for sensors or dye-sensitized solar cells. The ZnO film was also pasted on a desired substrate such as PET films, Si substrate or glass plates. The surface of low heat-resistant flexible polymer film was modified with high *c*-axis oriented crystalline ZnO film without heat treatment. This low-cost, low-temperature technique can be used for a wide range of applications including sensors, solar cells, electrical devices and optical devices using the various properties of high *c*-axis oriented crystalline ZnO.

6. Morphology control of nanocrystal assembled TiO₂ particles⁷⁷

Anatase TiO₂ particles, 100–200 nm in diameter, were developed in aqueous solution at 50°C. The particles were assemblies of nano TiO₂ crystals covered with nanorelief surface structures. The crystals grew anisotropically along the *c*-axis to form acicular crystals. The particles showed *c*-axis orientation due to high-intensity X-ray diffraction from the (004) planes. The particles had a BET specific surface area of 270 m²/g. BJH and DFT/Monte-Carlo analysis of adsorption isotherm indicated the existence of pores ~3 nm and ~1 nm in diameter. Crystallization and self-assembly of nano TiO₂ were effectively utilized to fabricate nanocrystal assembled TiO₂ particles having high surface area and nanorelief surface structure.

Nanoporous TiO₂ architecture with micropores (<2 nm), mesopores (2–50 nm) and/or macropores (>50 nm) is of considerable interest for both scientific and technical applications. The latter include cosmetics, catalysts⁷⁸, photocatalysts^{79–82}, gas sensors^{83,84}, lithium batteries^{85–87}, biomolecular sensors⁸⁸ and dye-sensitized solar cells^{89,90}. Crystalline anatase generally exhibits higher properties than rutile in photocatalysts, biomolecular sensors and dye-sensitized solar cells. Electrons are obtained from dyes adsorbed on TiO₂ electrodes in sensors and solar cells. Photoelectric conversion efficiency strongly depends on dye adsorption volume and surface area of TiO₂. High surface area is required to achieve high efficiency and sensitivity of the devices. Additionally, the surface of TiO₂ should be covered with nano/microrelief structures to adsorb large amounts of dye, molecules and DNA for biomolecular sensors and dye-sensitized solar cells.

TiO₂ nanoparticles have been prepared by flame synthesis^{91,92}, ultrasonic irradiation^{93,94}, chemical vapor synthesis⁹⁵, sol-gel methods^{79,96–100}, sonochemical method¹⁰¹ and liquid phase deposition of amorphous TiO₂^{102–105}. High temperature in the treatment processes, however, causes aggregation of nanoparticles and decreased surface area. Formation of nanorelief structures on the surface of the particles is difficult to achieve in these processes.

Highly porous materials have been prepared via template-based methods, including soft templates (surfactants, chelating agents, block polymers, etc.)¹⁰⁶⁻¹⁰⁹ and hard templates (porous anionic alumina, porous silica, polystyrene spheres, carbon nanotubes, etc.)^{110,111}. However, the nanostructures of these materials usually change due to amorphous-phase crystallization to anatase TiO₂ during annealing. This decreases the surface area and damages the surface nanostructures.

In this section, porous anatase TiO₂ particles were developed in aqueous solution. Nano TiO₂ was crystallized in the solution to assemble into particles 100–200 nm in diameter. The surface of the particles was covered with nanorelief structures. The particles showed *c*-axis orientation due to anisotropic crystal growth of TiO₂ along the *c*-axis. BET surface area of the particles was estimated to be 270 m²/g¹¹². BJH and DFT/Monte-Carlo analysis of adsorption isotherm indicated the existence of pores ~2.8 nm and ~3.6 nm in diameter, respectively. The existence of micropores ~1 nm was also indicated. Crystallization and self-assembly of acicular TiO₂ were effectively utilized to fabricate nanocrystal assembled TiO₂ particles having high surface area and nanorelief surface structure.

Ammonium hexafluorotitanate ([NH₄]₂TiF₆) (Morita Chemical Industries Co., Ltd., FW: 197.95, purity 96.0%) and boric acid (H₃BO₃) (Kishida Chemical Co., Ltd., FW: 61.83, purity 99.5%) were used as received. Ammonium hexafluorotitanate (12.372 g) and boric acid (11.1852 g) were separately dissolved in deionized water (600 mL) at 50°C. Boric acid solution was added to ammonium hexafluorotitanate solution at a concentration of 0.15 and 0.05 M, respectively.

The solution was kept at 50°C for 30 min using a water bath with no stirring, after which it was centrifuged at 4000 rpm for 10 min (Model 8920, Kubota Corp.). Preparation for centrifugation, centrifugation at 4000 rpm, deceleration from 4000 to 0 rpm and preparation for removal of supernatant solution took 4 min, 10 min, 10 min and 8 min, respectively. The particles contacted with the solution, the temperature of which was gradually lowered for 32 min after maintaining 50°C for 30 min. Precipitates were dried at 60°C for 12 h after removal of supernatant solution.

The crystal phase of the particles was evaluated by X-ray diffractometer (XRD; RINT-2100V, Rigaku) with CuK α radiation (40 kV, 30 mA). Diffraction patterns were evaluated using JCPDS, ICSD (Inorganic Crystal Structure Database) data (FIZ Karlsruhe, Germany and NIST, USA) and FindIt. Morphology of TiO₂ was observed by transmission electron microscopy (TEM; JEM2010, 200 kV, JEOL). Zeta potential and particle size distribution were measured by electrophoretic light-scattering spectrophotometer (ELS-Z2, Otsuka Electronics Co., Ltd.) with automatic pH titrator. Samples of 0.01 g were dispersed in distilled water (100 g) and ultrasonicated for 30 min prior to measurement. The pH of colloidal solutions was controlled by the addition of HCl (0.1 M) or NaOH (0.1 M). Zeta potential and particle size distribution were evaluated at 25°C and integrated 5 and 70 times, respectively.

Nitrogen adsorption-desorption isotherms were obtained using Autosorb-1 (Quantachrome Instruments) and samples of 0.137 g were outgassed at 110°C under 10⁻² mmHg for 6 h prior to measurement. Specific surface area was calculated by BET (Brunauer-Emmett-Teller) method using adsorption isotherms. Pore size distribution was calculated by BJH (Barrett-Joyner-Halenda) method using adsorption isotherms because an artificial peak was observed from BJH size distribution calculated from desorption branches. Pore size distribution was further calculated by DFT/Monte-Carlo method (N₂ at 77 K on silica (cylinder/sphere, pore, NLDFT ads. model), adsorbent: oxygen) using adsorption branches.

Total pore volume (V) and average pore diameter ($4V/A$) were estimated from pores smaller than 230 nm (diameter) at $P/P_o = 0.99$ through the use of BET surface area (A). However, adsorption volume increased drastically at high relative pressure in isotherms, indicating a large data error at high relative pressure. Therefore, total pore volume and average pore diameter estimated from pores smaller than 11 nm at $P/P_o = 0.80$ would provide useful information. Relative pressure $P/P_o = 0.80$ was selected for estimations because it was lower than the drastic adsorption increase and higher than adsorption hysteresis. The data can be compared to that of TiO_2 having similar morphology with small errors. Total pore volume (cumulative pore volume) was also estimated by BJH method from pores smaller than 154 nm. This usually has a small error compared to that estimated from isotherm data including high relative pressure such as $P/P_o = 0.99$ because estimation by BJH method is not effected by adsorption volume errors at high relative pressure.

Liquid phase crystal deposition of anatase TiO_2 . The solution became clouded about 10 min after mixing ammonium hexafluorotitanate solution and boric acid solution. The particles were homogeneously nucleated in the solution, turning the solution white.

Deposition of anatase TiO_2 proceeds by the following mechanisms^{43,113,114}:

Equation (a) is described in detail by the following two equations:

Fluorinated titanium complex ions gradually change into titanium hydroxide complex ions in an aqueous solution, as shown in Eq. (c). The increase of F^- concentration displaces Eqs. (a) and (c) to the left; however, the produced F^- can be scavenged by H_3BO_3 (BO_3^{3-}) as shown in Eq. (b) to displace Eqs. (a) and (c) to the right. Anatase TiO_2 is formed from titanium hydroxide complex ions ($\text{Ti}(\text{OH})_6^{2-}$) in Eq. (d).

Crystal phase of TiO_2 particles. X-ray diffraction peaks for the particles were observed at $2\theta = 25.1, 37.9, 47.6, 54.2, 62.4, 69.3, 75.1, 82.5$ and 94.0° after evaluation of N_2 adsorption. They were assigned to the 101, 004, 200, 105 + 211, 204, 116 + 220, 215, 303 + 224 + 312 and 305 + 321 diffraction peaks of anatase TiO_2 (JCPDS No. 21-1272, ICSD No. 9852) (Fig. 16).

The 004 diffraction intensity of randomly oriented particles is usually 0.2 times the 101 diffraction intensity as shown in JCPDS data (No. 21-1272). However, the 004 diffraction intensity of the particles deposited in our process was 0.36 times the 101 diffraction intensity. Additionally, the integral intensity of the 004 diffraction was 0.18 times the 101 diffraction intensity, indicating the c -axis orientation of the particles. Particles were not oriented on the glass holder for XRD measurement. Therefore, TiO_2 crystals would be an anisotropic shape in which the crystals were elongated along the c -axis. The crystals would have a large number of stacks of c planes such as (001) planes compared to stacks of (101) planes. The diffraction intensity from the (004) planes would be enhanced compared to that from the (101) planes.

Crystallite size perpendicular to the (101) or (004) planes was estimated from the full-width half-maximum of the 101 or 004 peak to be 3.9 nm or 6.3 nm, respectively. Elongation of crystals in the c -axis direction was also suggested by the difference in crystallite size.

Fig. 16. XRD diffraction pattern of anatase TiO_2 particles. Reprinted with permission from Ref.⁷⁷, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 3213. Copyright @American Chemical Society

TEM observation of TiO_2 particles. The particles were shown to be assemblies of nano TiO_2 crystals (Fig. 17a). Particle diameter was estimated to be 100–200 nm. Relief structures had formed on the surfaces and open pores had formed inside because the particles were porous assemblies of nanocrystals.

Nanocrystals were shown to have acicular shapes (Fig. 17b). They were about 5–10 nm in length. The longer direction of acicular TiO_2 is indicated by the black arrow. The inserted FFT image shows the 101 and 004 diffractions of anatase TiO_2 . Nanocrystals are assigned to the single phase of anatase TiO_2 . It is notable that the diffraction from the (101) planes has a ring shape due to random orientation but that from the (004) planes was observed only in the upper right region and lower left region in the FFT image. Anisotropic 004 diffractions indicated the direction of the c -axis, which was perpendicular to the (004) planes, as shown by the white arrow. It was roughly parallel to the longer direction of acicular TiO_2 . These results suggest that acicular TiO_2 grew along the c -axis to enhance the diffraction intensity from the (004) planes. Crystal growth of anatase TiO_2 along the c -axis was previously observed in TiO_2 films¹¹³. Anisotropic crystal growth is one of the features of liquid phase crystal deposition.

Acicular nanocrystals showed lattice images of anatase TiO_2 (Fig. 17c). They were constructed of anatase TiO_2 crystals without amorphous or additional phases. Anatase crystals were not covered with amorphous or additional phases even at the tips. Bare anatase crystal with nanosized structure is important to achieve high performance for catalysts and devices.

Crystallization of TiO_2 was effectively utilized to form assemblies of acicular nanocrystals in the process. Open pores and surface relief structures were successfully formed on the particles.

Fig. 17. (a): TEM micrograph of anatase TiO_2 particles. (b): Magnified area of (a) showing morphology of acicular crystals. Insertion in (b): FFT image of (b) anatase TiO_2 . (c): Magnified area of (a) showing lattice images of anatase TiO_2 . Reprinted with permission from Ref.⁷⁷, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 3213. Copyright @American Chemical Society

Zeta potential and particle size distribution. The dried particles were dispersed in water to evaluate zeta potential and particle size distribution after evaluation of N_2 adsorption. The particles had positive zeta potential of 30.2 mV at pH 3.1, which decreased to 5.0, -0.6, -11.3 and -36.3 mV at pH 5.0, 7.0, 9.0 and 11.1, respectively (Fig. 18). The isoelectric point was estimated to be pH 6.7, slightly higher than that of anatase TiO_2 (pH 2.7–6.0)¹¹⁵. Zeta potential is very sensitive to the particle surface conditions, ions adsorbed on the particle surfaces, and the kind and concentration of ions in the solution. The variations in zeta potential were likely caused by the difference in the surface conditions of TiO_2 particles, affected by the interaction between particles and ions in the solution.

Fig. 18. Zeta potential of anatase TiO_2 particles as a function of pH. Reprinted with permission from Ref.⁷⁷, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 3213. Copyright @American Chemical Society

Mean particle size was estimated to be ~550 nm in diameter with a standard deviation (STD) of 220 nm at pH 3.1 (Fig. 19a). This was larger than that observed by TEM. Slight aggregation occurred at pH 3 because the particles were dried completely prior to measurement. Particle size increased with pH and showed a maximum of near the isoelectric point (550 nm at pH 3.1, 3150 nm at pH 5, 4300 nm at pH 7, 5500 nm at pH 9 or 2400 nm at pH 11.1) (Fig. 19b). Strong aggregation resulted from the lack of repulsion force between particles near the isoelectric point.

The particles were generated in the solution at pH 3.8 in this study. It would be suitable to obtain repulsion force between particles for crystallization without strong aggregation.

Fig. 19. (a): Particle size distribution of anatase TiO₂ particles at pH 3.1. (b): Particle size of anatase TiO₂ particles as a function of pH. Reprinted with permission from Ref.⁷⁷, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 3213. Copyright ©American Chemical Society

N₂ adsorption characteristics of TiO₂ particles. TiO₂ particles exhibited N₂ adsorption-desorption isotherms of Type IV (Fig. 20a). The desorption isotherm differed from adsorption isotherm in the relative pressure (P/P_0) range from 0.4 to 0.7, showing mesopores in the particles. BET surface area of the particles was estimated to be 270 m²/g (Fig. 20b). This is higher than that of TiO₂ nanoparticles such as Aerioxide P25 (BET 50 m²/g, 21 nm in diameter, anatase 80% + rutile 20%, Degussa), Aerioxide P90 (BET 90–100 m²/g, 14 nm in diameter, anatase 90% + rutile 10%, Degussa), MT-01 (BET 60 m²/g, 10 nm in diameter, rutile, Tayca Corp.) and Altair TiNano (BET 50 m²/g, 30–50 nm in diameter, Altair Nanotechnologies Inc.)¹¹⁶. A high BET surface area cannot be obtained from particles having a smooth surface even if the particle size is less than 100 nm. A high BET surface area would be realized by the unique morphology of TiO₂ particles constructed of nanocrystal assemblies.

Total pore volume and average pore diameter were estimated from pores smaller than 230 nm at $P/P_0 = 0.99$ – 0.431 cc/g and 6.4 nm, respectively. They were estimated to be 0.212 cc/g and 3.1 nm, respectively, from pores smaller than 11 nm at $P/P_0 = 0.80$. Total pore volume was also estimated by the BJH method from pores smaller than 154 nm to be 0.428 cc/g. Average pore diameter was estimated to be 6.3 nm using BET surface area.

Fig. 20. (a): N₂ adsorption-desorption isotherm of anatase TiO₂ particles. (b): BET surface area of anatase TiO₂ particles. (c): Pore size distribution calculated from N₂ adsorption data of anatase TiO₂ particles using BJH equation. (d): N₂ adsorption-desorption isotherm and DFT/Monte-Carlo fitting curve of anatase TiO₂ particles. (e): Pore size distribution calculated from N₂ adsorption data of anatase TiO₂ particles using DFT/Monte-Carlo equation. Reprinted with permission from Ref.⁷⁷, Masuda, Y. and Kato, K., 2008, *Cryst. Growth Des.*, 8, 3213. Copyright ©American Chemical Society

Pore size distribution was calculated by the BJH method using adsorption isotherms (Fig. 20c). It showed a pore size distribution curve having a peak at ~ 2.8 nm and pores larger than 10 nm. TiO_2 particles would have mesopores of ~ 2.8 nm surrounded by nanocrystals. Pores larger than 10 nm are considered to be interparticle spaces. The pore size distribution also suggested the existence of micropores smaller than 1 nm.

Pore size distribution was further calculated by the DFT/Monte-Carlo method. The model was in fair agreement with adsorption isotherms (Fig. 20d). Pore size distribution showed a peak at ~ 3.6 nm that indicated the existence of mesopores of ~ 3.6 nm (Fig. 20e). The pore size calculated by the DFT/Monte-Carlo method was slightly larger than that calculated from the BJH method because the latter method is considered to have produced an underestimation¹¹⁷⁻¹¹⁹. The pore size distribution also suggested the existence of micropores of ~ 1 nm, probably resulting from microspaces surrounded by nanocrystals and the uneven surface structure of nanocrystals.

The particles were shown to have a large surface area as well as micropores of ~ 1 nm, mesopores of ~ 2.8 – 3.6 nm and pores larger than 10 nm, by N_2 adsorption characteristics. Assembly of acicular nanocrystals resulted in unique features and high surface area.

TiO_2 particles were generated in the solutions at 90°C for 1h using an oil bath with no stirring for comparison. The solutions became clouded after the addition of boric acid solutions into ammonium hexafluorotitanate solutions. High temperature accelerated crystal growth of TiO_2 . Hydrogen chloride of 0.6 ml was added into the solutions of 200ml to decrease crystallization speed of TiO_2 . The pH of the solutions was 2.4 one hour after mixing the solutions. BET surface area of the particles was estimated to $18 \text{ m}^2/\text{g}$. This is much lower than that of the particles prepared at 50°C and slightly lower than that prepared at 90°C for 8 min in our previous work ($44 \text{ m}^2/\text{g}$)¹²⁰. Formation of TiO_2 was accelerated at high temperature and it decreased surface area. The particles grew in the solutions to decrease surface area as function of time. Crystallization of TiO_2 was shown to be strongly affected by growth conditions such as solution temperature and growth time.

Anatase TiO_2 particles, 100–200 nm in diameter, were successfully fabricated in aqueous solution. They were assemblies of nanocrystals 5–10 nm that grew anisotropically along the *c*-axis to form acicular shapes. The particles thus had nanorelief surface structures constructed of acicular crystals. They showed *c*-axis orientation due to high-intensity X-ray diffraction from the (004) crystal planes. The particles had a high BET surface area of $270 \text{ m}^2/\text{g}$. Total pore volume and average pore diameter were estimated from pores smaller than 230 nm at $P/P_0 = 0.99$ – 0.43 cc/g and 6.4 nm, respectively. They were also estimated from pores smaller than 11 nm at $P/P_0 = 0.80$ – 0.21 cc/g and 3.1 nm, respectively. BJH and DFT/Monte-Carlo analysis of adsorption isotherm indicated the existence of pores ~ 2.8 and ~ 3.6 nm, respectively. Additionally, the analyses suggested the existence of micropores of ~ 1 nm. Crystallization and self-assembly of nano TiO_2 were effectively utilized to fabricate nanocrystal assembled TiO_2 particles having high surface area and nanorelief surface structure.

7. Morphology control of multi-needle TiO_2 particles¹²¹

Flower-like multi-needle anatase TiO_2 particles were developed in aqueous solutions. They were pure anatase TiO_2 crystals containing no cores, organic binders or solvents. Furthermore, micro-structured silicon wafers were covered with the TiO_2 particles uniformly in the solutions. Their unique crystals growth and physicochemical profiles were precisely evaluated and discussed.

Micro-structured silicon wafers and their surface modification to super-hydrophilic surfaces

Silicon wafers were modified to have micro-structures on the surfaces with cutting work. They were cut using a precise diamond cutter under running water. Width and height of salient lines were 200 μm and 150 μm , respectively. They were formed at 500 μm intervals. They were blown by air to remove dust and were exposed to vacuum-ultraviolet light (VUV light, low-pressure mercury lamp PL16-110, air flow, 100 V, 200 W, SEN Lights Co., 14 mW/cm² for 184.9 nm at a distance of 10 mm from the lamp, 18 mW/cm² for 253.7 nm at a distance of 10 mm from the lamp) for 10 min in air. Bare silicon surfaces were covered with small amount of surface contamination. The VUV irradiation modified them to clean surfaces that showed super hydrophilic surfaces of water contact angle about 0-1 °.

Morphology control of anatase TiO₂

Ammonium hexafluorotitanate (206.20 mg) and boric acid (186.42 mg) were separately dissolved in deionized hot water (100 mL) at 50°C. Boric acid solution was added to ammonium hexafluorotitanate solution at concentrations of 15 mM and 5 mM, respectively. The silicon wafers having patterned surfaces were immersed in the middle of the solutions with the bottom up at an angle. They were tilted at 15 degrees to the upright. The solutions were kept at 50°C for 19 hours or 7 days using a drying oven (Yamato Scientific Co., Ltd., DKN402) with no stirring. The substrates were washed with running water and dried by air blow. The solutions were centrifuged at 4000 rpm for 10 min (Model 8920, Kubota Corp.). Precipitated particles were dried at 60°C for 12 h after removal of supernatant solutions. The particles were dispersed in distilled water. They were centrifuged and dried again for purification.

1. Morphology control of multi-needle TiO₂ particles

TEM observation of multi-needle TiO₂ particles

Flower-like multi-needle TiO₂ particles were successfully formed in aqueous solutions (Fig. 21a, b). Needle shaped crystals grew from the center of the particles. Especially, the needles grew parallel to TEM observation direction from the center of the particles as shown in red circles. They clearly showed that needles radiated in all directions to form flower-like morphology. Each particle had about 6-10 taper needles. Width and length of them were about 200 nm and 100 nm, respectively. Aspect ratio was about 2 (200 nm / 100 nm). Width of the needles became smaller as growth direction to have tips. The particles had no core or pore at the center of their bodies. Nucleation and crystal growth were well controlled to have flower-like multi-needle morphology.

Electron diffraction pattern showed that the particles were single phase of anatase TiO₂ crystals (Fig. 21c). Interplanar spacing of (004), (200), and (204) planes were estimated to 0.243 nm, 0.201 nm and 0.151 nm. Diffractions from (004), (200), and (204) planes were clear single spots (Fig. 21c). Needle shaped crystal was shown to single crystal of anatase TiO₂. Long direction of the needle shaped crystals was perpendicular to (004) crystal faces (Fig. 21c). It indicated that anatase TiO₂ crystals grew along *c*-axis to form needle shape morphology. They were thus surrounded by *a*-faces of anatase TiO₂ crystals.

Surfaces of the needle crystals were observed carefully (Fig. 21c, d). There were no amorphous layers or second phase layers on the surfaces. The particles had pure and bare anatase TiO₂ surfaces.

Fig. 21. Transmission electron microscope images of the multi-needle TiO_2 particles. (a, b) Low magnification images of the particles. (c) High resolution image of the needles grew from the center of the particle showing taper shape of the needle. (d) High resolution image of surface of the needles showing bare TiO_2 surface. (c) and (d) are magnified area of (a) and (b). Insert in (c) is electron diffraction pattern indicating crystal phase and interplanar spacing. Red circles indicate typical images of the multi-needle TiO_2 particles. Reprinted with permission from Ref.¹²¹, Masuda, Y., Ohji, T. and Kato, K., 2010, *Cryst. Growth Des.*, 10, 913. Copyright @American Chemical Society

XRD analysis of multi-needle TiO_2 particles

XRD analysis showed that the particles were single phase of anatase TiO_2 . X-ray diffraction peaks were observed at $2\theta = 25.12, 36.8, 37.7, 47.7, 53.7, 54.7, 62.4, 68.5, 69.8, 74.7, 82.2, 93.7$ and 94.4° . Pure anatase TiO_2 particles were obtained from the aqueous solutions in this study. Anatase phase have been prepared using high temperature annealing in many reports. It caused deformation of nano/micro-structures, aggregation of the particles and decrease of surface area. However, crystallization of anatase TiO_2 was realized at 50°C in this study to avoid degradation of the properties.

Crystallite size perpendicular to the (101), (004) or (200) planes was estimated from the full-width half-maximum of the 101, 004 or 200 peak to be 18.2, 29.8 or 19.2 nm, respectively. Crystallite size perpendicular to (004) was larger than that of others. Difference in crystallite size indicated anisotropic crystal growth along *c*-axis.

For comparison, nanocrystal assembled TiO₂ particles were prepared in aqueous solutions at 50°C for 30 min in previous report⁷⁷. Crystallite size perpendicular to the (101), (004) or (200) planes was estimated to be 3.9, 6.3 or 4.9 nm, respectively. The crystalline degree of the multi-needle TiO₂ nanostructures was much improved compared with that of the previous nanocrystal⁷⁷. The key to achieve highly crystalline degree even at low-temperature, i.e., 50°C, was slow and long-term growth. It was achieved by low super saturation degree of the solutions. Concentration of ammonium hexafluorotitanate and boric acid were one-tenth of the previous nanocrystal⁷⁷. Synthesis parameters of multi-needle TiO₂ nanostructures were developed based on previous reports as follows. Acicular nanocrystals were homogeneously formed immediately after mixing of two solutions in previous report⁷⁷. They aggregated into nanocrystal assembled particles. The particles were removed from the solutions 30 minutes after the mixing to prevent further crystal growth. The crystallite size was thus small and it contributed to high specific surface area. On the other hand, TiO₂ films were prepared in the same solutions at 50°C for several hours^{114,122,123}. The films were consisted of two layers. Under layer was consisted of small nanocrystals. They were formed at an early stage of immersion period in high ion concentration solutions. Ions were consumed gradually to form the crystals. Upper layer of acicular crystal assembly was then formed. They grew in the solutions with low ion concentrations. It indicated that low ion concentration and low super saturation degree realized formation of acicular TiO₂ crystals. Additionally, size of acicular crystals was much larger than initially deposited nanocrystals. We tried to form multi-needle particles consisted of large acicular crystals on the basis of these results. Crystal growth in the solution with low ion concentration and low super saturation degree was utilized for anisotropic crystal growth. They were grown to large crystals with long period such as 7 days. Slow growth rate caused formation of euhedral crystals that were affected by crystal structure of tetragonal anatase.

Raman spectroscopy of multi-needle TiO₂ particles

The particles showed Raman peaks at 157 cm⁻¹, 412 cm⁻¹, 506 cm⁻¹ and 628 cm⁻¹. It was typical of anatase TiO₂ phase. They were assigned to Eg (v6) mode (157 cm⁻¹), B1g mode (412 cm⁻¹), doublet of the A1g and B1g modes (506 cm⁻¹), and Eg (v1) mode (628 cm⁻¹), respectively. Additional peaks indicating rutile TiO₂ or other phases were not observed. It was notable that Eg (v6) mode shifted with respect to those of bulk crystals or sintered powders. For the single crystal, Ohsaka et al. determined following allowed bands: 144 cm⁻¹ (Eg), 197 cm⁻¹ (Eg), 399 cm⁻¹ (B1g), 513 cm⁻¹ (A1g), 519 cm⁻¹ (B1g) and 639 cm⁻¹ (Eg)¹²⁴. TiO₂ particles (P25, Degussa) showed peaks at 143.3 cm⁻¹ (Eg), 196 cm⁻¹ (Eg), 396 cm⁻¹ (B1g), 516 cm⁻¹ (A1g+B1g) and 638 cm⁻¹ (Eg). Raman peak of Eg (v6) mode (144 cm⁻¹) has been reported to shift due to several factors such as effect of crystalline size (quantum size confinement effect)^{125,126}, temperature¹²⁵ or pressure¹²⁷. Additionally, nitrogen doped anatase TiO₂ particles of 8.40-9.80 nm in diameter were reported to have Eg (v6) mode at 151 cm⁻¹¹²⁶. The multi-needle particles showed large peak shift of Eg (v6) mode from 144 cm⁻¹ to 157 cm⁻¹ because of fluorine doping effect and size effect. The large shift of Eg (v6) mode was one of the characteristics of the multi-needle TiO₂ particles.

FT-IR absorbance of multi-needle TiO₂ particles

The particles showed absorption spectra in infrared light region. The absorption bands related to TiO₂ were observed at 907 cm⁻¹ and 773 cm⁻¹. They were assigned to stretching vibrations of Ti=O and -Ti-O-Ti-, respectively. They were consistent with TEM, XRD and Raman analyses. The bands observed in the range of 1400-1750 cm⁻¹ were attributed to bending vibrations of O-H¹²⁸. Absorption band in frequency range 600-450 cm⁻¹ were reported to be attributed to stretching vibrations of Ti-F bonds in TiO₂ lattice^{129,130}. The bands were observed at 513, 532, 540, 558 or 568 cm⁻¹ from xTiOF₂ · yBaF₂ · zMnF₂ glasses¹²⁹. They indicated that F ions were partially replaced to O ions in the lattice¹²⁹. Actually, absorption bands were observed at 455, 488, 505, 520, 552 and 567 cm⁻¹ in the spectra of the multi-needle TiO₂ particles. Additionally, absorption band at 1080 cm⁻¹ was reported to be assigned to surface fluorinated Ti-F species¹³¹. Absorption band was observed at about 1058 cm⁻¹ from the multi-needle TiO₂ particles. These analyses suggested that F ions were partially doped into TiO₂ crystals to replace O ions.

UV-Vis spectroscopy and band gap of multi-needle TiO₂ particles

Optical property of the particles was evaluated with UV-Vis spectroscopy. Transparency gradually decreased as decrease of wavelength. The bulk band gap structures are known as direct-transition for anatase and indirect-transition for rutile in titania, respectively. Band gap of the particles was estimated to 3.20 eV (388 nm) by assuming the direct-transition. It was similar to that of anatase TiO₂ nanostructures such as anatase TiO₂ nanorods (3.2 eV (388 nm))¹³², anatase TiO₂ nanowalls (3.2 eV (388 nm))¹³², anatase TiO₂ nanotubes (3.2 eV (388 nm))¹³³, single-layered TiO₂ nano-sheet with a thickness less than 1 nm (3.15 eV (394 nm))¹³⁴, stacked TiO₂ nano-sheets (3.15 eV (394 nm))¹³⁴ and anatase TiO₂ films (3.2 eV (388 nm))¹³⁵. It was higher than that of rutile TiO₂ films (2.9 eV (428 nm))¹³⁵, rutile TiO₂ nanorods (3.0 eV (414 nm))¹³², rutile TiO₂ single crystal (3.0 eV (414 nm), SHINKOSHA Co., Ltd.), and lower than that of amorphous TiO₂ films (3.5 eV (355 nm))¹³⁵.

N₂ adsorption characteristics of multi-needle TiO₂ particles

BET surface area of the particles was estimated to be 178 m²/g from an adsorption branch in the range of P/Po = 0.1-0.29. Average pore diameter was estimated to be 11.1 nm using BET surface area. They were estimated to 166 m²/g and 11.9 nm, 151 m²/g and 17.6 nm from P/Po = 0.05-0.1 or P/Po = 0.02-0.07, respectively. BET surface area was higher than that of TiO₂ nanoparticles such as Aerioxide P25 (BET 50 m²/g, 21 nm in diameter, anatase 80% + rutile 20%, Degussa), Aerioxide P90 (BET 90-100 m²/g, 14 nm in diameter, anatase 90% + rutile 10%, Degussa), MT-01 (BET 60 m²/g, 10 nm in diameter, rutile, Tayca Corp.) and Altair TiNano (BET 50 m²/g, 30-50 nm in diameter, Altair Nanotechnologies Inc.)¹¹⁶. Pore size distribution calculated by BJH method indicated that spaces of 2-3 nm were existed in the particles. DFT/Monte-Carlo analysis showed several types of mesopores in the range of 3-10 nm (3.5 nm, 4.9 nm, 6.3 nm and 10 nm) and micropores of ~0.8 nm. Total pore volume and average pore diameter were estimated from pores smaller than 241 nm at P/Po = 0.99 to be 0.493 cc/g and 13 nm, respectively. They were estimated to be 0.158 cc/g and 4.2 nm, respectively, from pores smaller than 11 nm at P/Po = 0.81. Total pore volume was also estimated by the BJH method from pores smaller than 156 nm to be 0.505 cc/g using desorption isotherm. Total pore volume was also estimated by the BJH method from pores smaller than 160 nm to be 0.628 cc/g using adsorption isotherm.

2. Surface coating of micro-structured substrates with multi-needle TiO₂ particles

SEM observation of the surface coatings

Micro-structured silicon wafers were immersed in the aqueous solutions at 50°C for 7 days. The substrates were successfully covered with multi-needle TiO₂ particles (Fig. 22-a). Both of salient regions and concave regions were modified with the particles uniformly (Fig. 22-b1, c1). The original micro-structure of silicon wafers were well maintained because the TiO₂ surface coatings were uniform thin layers of about 200-600 nm in thickness (Fig. 22-b2, c2). The coating layers were consisted of flower-like multi-needle TiO₂ particles of about 200-400 nm in diameter (Fig. 22-b3, c3, red circles). The particles had several needles which grew from the center of the particles. The needles had taper shape along growth direction. The particle in red circle of Fig. 22-b3 clearly indicated that tetragonal crystal phase of anatase TiO₂ caused fourfold symmetry crystal growth of four needles parallel to the substrate. The *a*-axis and *c*-axis of the particle were parallel and perpendicular to the substrate, respectively. This shape was basic morphology of the particles in this system. Other needles also grew from the center to form multi-needle shape as shown in red circles in Fig. 22-c3. Salient regions were slightly trapezoidal geometry (Fig. 22-a, b1). TiO₂ particles were deposited on walls of them. The area between the salient and concave regions was thus white in SEM images.

For comparison, micro-structured silicon wafers were immersed in the aqueous solutions at 50°C for 19 hours. The multi-needle TiO₂ particles were formed on both of salient regions and concave regions (Fig. 23-a). The micro-structures of the substrates were maintained because the surface coatings were uniform thin layers (Fig. 23-b1, c1). However, coverage of the surfaces was different from that immersed for 7 d. The TiO₂ particles covered approximately half of the salient regions and one third of concave regions (Fig. 23-b2, c2). Coverage of the salient regions was higher than that of concave regions because concave regions set back far from the salient region surfaces. The particles and ions were not well supplied to concave regions. The particles had several needles grown from the center (Fig. 23-b3, c3). The morphology of them was similar to that immersed for 7d, in contrast, their size was slightly smaller to be about 100-300 nm. These observations indicated growth mechanism as follows. The particles were homogeneously nucleated from ions in the solutions. They were then adhered on the substrates to form surface coatings. The particles further grew to increase number and size of the needles.

XPS analysis of the surface coatings

Surfaces of the TiO₂ coatings were analyzed with XPS. Titanium, oxygen, carbon, silicon and fluorine were observed from the surface before (Fig. 24-1a) and after (Fig. 24-1b) Ar⁺ sputtering. Ti 2p_{3/2} spectrum was observed at 458.6 eV (Fig. 24-2a). The binding energy was higher than that of Ti metal (454.0 eV), TiC (454.6 eV), TiO (455.0 eV), TiN (455.7 eV) and Ti₂O₃ (456.7 eV), and similar to that of TiO₂ (458.4-458.7 eV)¹³⁶⁻¹³⁸. This suggested that the titanium atoms in the particles were positively charged relative to that of titanium metal by formation of direct bonds with oxygen. Ti 2p spectrum changed its shape by the sputtering due to the decrease of cation valence (Fig. 24-2b). The phenomenon was often observed in sputtering of titanium oxide¹³⁶. The spectrum after sputtering was not suitable for estimation of chemical ratio because the ratio was slightly changed by decrease of titanium cation valence. The chemical ratio of Ti to O was thus estimated from the spectra before the sputtering.

Fig. 22. (a) SEM micrograph of a micro-structured substrate coated with the multi-needle TiO_2 particles deposited for 7 d. (b1) Salient region coated with the multi-needle TiO_2 particles. (b2) Magnified area of (b1) showing uniform surface coating. (b3) Magnified area of (b2) showing morphology of the multi-needle TiO_2 particles. (c1) Concave region coated with the multi-needle TiO_2 particles. (c2) Magnified area of (c1) showing uniform surface coating. (c3) Magnified area of (c2) showing morphology of the multi-needle TiO_2 particles. Black arrows show salient regions. White arrows show concave regions. Red circles indicate typical images of the multi-needle TiO_2 particles. Reprinted with permission from Ref.¹²¹, Masuda, Y., Ohji, T. and Kato, K., 2010, *Cryst. Growth Des.*, 10, 913. Copyright ©American Chemical Society

Fig. 23. (a) SEM micrograph of a micro-structured substrate coated with the multi-needle TiO_2 particles deposited for 19 h. (b1) Salient region coated with the multi-needle TiO_2 particles. (b2) Magnified area of (b1) showing uniform surface coating. (b3) Magnified area of (b2) showing morphology of the multi-needle TiO_2 particles. (c1) Concave region coated with the multi-needle TiO_2 particles. (c2) Magnified area of (c1) showing uniform surface coating. (c3) Magnified area of (c2) showing morphology of the multi-needle TiO_2 particles. Black arrows show salient regions. White arrows show concave regions. Red circles indicate typical images of the multi-needle TiO_2 particles. Reprinted with permission from Ref.¹²¹, Masuda, Y., Ohji, T. and Kato, K., 2010, *Cryst. Growth Des.*, 10, 913. Copyright ©American Chemical Society

O 1s spectrum was decomposed into two Gaussian curves after removal of background (Fig. 24-3a). The integral intensity ratio of O 1s peak at 532.12 eV to O 1s peak at 529.85 eV was 0.662 : 0.338. The binding energy of O 1s peak at 529.85 eV was similar to that of TiO₂ (529.9 eV¹³⁸, 530.1 eV^{136,137}) showing that oxygen was negatively charged compared to neutral oxygen molecules (531.0 eV) through the formation of direct bonds with Ti. High binding energy component at 532.12 eV was assigned to oxygen atoms combined to carbon atoms as C-O or C=O (532.8 eV). It was decreased by the sputtering for 10 sec (Fig. 24-3b). It was included in surface contaminations. Drastic decreasing of C 1s spectrum by the sputtering supported this ascription (Fig. 24-4a, 4b). Chemical ratio of Ti to O was estimated to 1 : 1.76 using Ti 2p at 458.6 eV (Fig. 24-2a) and O 1s at 529.85 eV (Fig. 24-3a). It was slightly smaller than that expected from TiO₂. Oxygen vacancy and doping of fluorine ions would decrease oxygen volume. Fluorine was, in fact, observed from the surfaces at 684.5 eV (Fig. 24-5a). Chemical ratio was estimated to Ti : O : F = 1 : 1.76 : 0.18. Sum of oxygen and fluorine was 1.94 which was similar to 2 expected from TiO₂. Fluorine spectrum intensity was not decreased by the sputtering to be Ti : F = 1 : 0.16 (Fig. 24-5b). These indicated that fluorine

Fig. 24. XPS spectra of (1) wide scan, (2) Ti 2p, (3) O 1s, (4) C 1s, (5) F 1s and (6) Si 2p for a micro-structured substrate coated with the multi-needle TiO₂ particles deposited for 7d (a) before and (b) after Ar⁺ sputtering. (3a) O1s spectra decomposed to two peaks after removal of background. Reprinted with permission from Ref.¹²¹, Masuda, Y., Ohji, T. and Kato, K., 2010, *Cryst. Growth Des.*, 10, 913. Copyright ©American Chemical Society

would not be included in surface contaminations but in the TiO_2 particles. Fluorine has been reported to improve properties of TiO_2 . Fluorination of TiO_2 increased surface acidity due to the strongest electronegativity of fluorine¹³⁹. Fluorine doping into TiO_2 increased the surface OH radicals that were suitable for photocatalytic reactions and improved photocorrosion resistance¹⁴⁰⁻¹⁴⁵. These properties expand application area of TiO_2 .

Si $2p_{3/2}$ was observed at 98.6 eV and 102.7 eV (Fig. 24-6a). It was detected from silicon regions uncovered with TiO_2 . The binding energies were similar to those in silicon wafers (99.6 eV) and SiO_2 (103.4 eV). The latter component was decreased by the sputtering because surface native oxide layer of amorphous SiO_2 was removed (Fig. 24-6b).

In this section, Multi-needle TiO_2 nanostructures having high surface area, fluorine doping and novel physicochemical characteristics were successfully fabricated. Unique aqueous synthesis realized their distinct morphologies and properties. The needle crystals grew along *c*-axis from the center of the particles. They were surrounded by *a*-faces of anatase TiO_2 crystals. Width and length of needles were about 200 nm and 100 nm. Aspect ratio was about 2 (200 nm / 100 nm). Diffraction patterns showed that the particles were single phase of anatase TiO_2 . Interparticle spaces and micro/meso pores of 1-10 nm allowed us to realize high surface area of 178 m^2/g . Large Raman peak shift of E_g (ν_6) mode suggested fluorine doping and size effect. Band gap was estimated to 3.20 eV (388 nm) with UV-Vis. These were characteristics of the multi-needle TiO_2 particles. Furthermore, self-assembly surface coating of micro-structured substrates was successfully realized. The coating layers were consisted of multi-needle TiO_2 particles of about 200-400 nm in diameter. XPS analyses indicated chemical bonds between Ti and O. Chemical ratio was estimated to Ti : O : F = 1 : 1.76 : 0.18 suggesting fluorine doping in TiO_2 . The multi-needle particles and the surface coating with the particles having nano/micro structures would contribute to future metal oxide devices of solar cells, photo catalysts and high sensitive sensors.

8. Morphology control of nano-sheet assembled tin oxide¹⁴⁶

Tin oxide particles were synthesized in aqueous solutions. They were consisted of nanosheets of tin oxide crystals. The sheets were about 50-100 nm in size and 5-10 nm thickness. X-ray diffraction analysis revealed that the particles were crystals of SnO_2 and SnO . The particles had BET surface area of 85 m^2/g estimated with N_2 adsorption characteristics. BJH analysis indicated that mesopores of 3.9 nm in size contributed to increase surface area.

Tin dioxide (SnO_2), with a rutile-type crystalline structure, is an n-type wide band gap (3.5 eV) semiconductor. It is an oxide of great interest for gas sensors[1-3], optical device¹⁴⁷, lithium batteries¹⁴⁸⁻¹⁵¹, white pigments for conducting coatings, transparent conducting coatings for furnaces and electrodes [9], surge arrestors (varistors)^{152,153}, catalysts^{154,155}, opto-conducting coatings for solar cells¹⁵⁶, dye-sensitized molecular sensors, etc. Transparency, semiconductivity and surfaced properties of tin oxide are suitable for these applications.

Nanoparticles of tin dioxide have been synthesized by several methods such as precipitation^{151,157}, hydrothermal synthesis^{158,159}, sol-gel^{160,161}, hydrolytic¹⁶², carbothermal reduction¹⁶³ and polymeric precursor¹⁶⁴ methods. A variety of SnO_2 nanostructures including nanowires, nanobelts¹⁶⁵, nanotubes^{150,166,167}, nanorods^{168,169}, spirals, nanorings¹⁷⁰, zigzag nanobelts¹⁷¹, grains¹⁷², flakes¹⁷², plates¹⁷², meshes¹⁷³ and columnar thin film¹⁴⁹ were synthesized.

Tin chlorides (SnCl_2 or SnCl_4) were commonly used in many reports. However, chlorine ions were difficult to remove from the systems and it seriously altered superficial and electrical properties. For instance, chlorine ions caused sensitivity degradation of gas sensors¹⁷⁴, aggregation of particles¹⁷⁵ and increasing of sintering temperatures¹⁷⁶. Chloride problems can be avoided through usage of organic tin compounds, such as alkoxides. However, these reagents were costly which makes industrial syntheses implementation hardly attainable. Additionally, sol-gel processes using alkoxides formed amorphous phases. High temperature annealing which increased cost, energy consumption and CO_2 emission was necessary to obtain tin oxide crystals.

Recently, aqueous syntheses of metal oxide crystals including tin oxide were developed^{1,20,177}. Morphology control of metal oxide crystals were realized by anisotropic crystal growth. They were induced by crystal growth control or organic additives. Thermodynamically stable crystal faces, for instance, depend on crystal growth conditions. Organic molecules adsorbed on typical crystal faces suppress crystal growth perpendicular to the faces. Morphology control realized improvement of properties of metal oxide devices. In this study, tin oxide particles were prepared in aqueous solutions¹⁴⁶. They were assembly of nanosheets. Morphology, crystal phases and N_2 adsorption characteristics were evaluated. Specific surface area and size distribution of pores were analyzed using isotherms.

Aqueous synthesis of nano-sheet assembled tin oxide particles. SnF_2 (Wako Pure Chemical Industries, Ltd., No. 202-05485, FW: 156.71, purity 90.0%) was used as received. Distilled water in polypropylene vessels (200 mL) were capped with polymer films and kept at 90°C . SnF_2 (870.6 mg) was dissolved in the distilled water at 90°C to be 5 mM. The solutions were kept at 90°C for 30 min using a drying oven (Yamato Scientific Co., Ltd., DKN402) with no stirring. The solutions became clouded shortly after the addition of SnF_2 . The bottoms of the vessels were covered with white precipitates. The solutions were centrifuged at 4000 rpm for 10 min (Model 8920, Kubota Corp.). Precipitated particles were dried at 60°C for 12 h after removal of supernatant solutions.

The solutions became clouded shortly after the addition of SnF_2 because of the homogeneous nucleation and growth of tin oxide particles. SnO_2 and SnO are formed in the aqueous solutions as follows:

The particles were gradually deposited to cover the bottom of the vessels. The supersaturation degree of the solutions was high at the initial stage and decreased as the color of the solutions changed.

Precipitated particles showed broad X-ray diffraction peaks at $2\theta = 27, 34, 38, 52$ and 57.5 . They were assigned to SnO_2 (JDPDS No. 41-1445). The peaks had large width due to small crystallite size of SnO_2 .

Morphology of nano-sheet assembled tin oxide particles. Nano-sheet assembled tin oxide particles were successfully fabricated. Crystallization of tin oxide nanosheets in aqueous solutions allowed us to obtain unique morphology of tin oxide. The particles with 300-800 nm in diameter were observed with FE-SEM (Fig. 25). They were not dense particles but assemblies of tin oxide nanosheets. The sheets were about 50-100 nm in size and 5-10 nm thickness. Aspect ratio was estimated to about 10. They were randomly aggregated to form the particles. The particles had thus continuous open pores inside. It contributed to increase surface area of the particles.

The nanosheets would be generated in the solutions homogeneously. They were aggregated to form particles which made solutions cloudy. They were then precipitated to cover bottoms of vessels. These were consistent with color change observation of solutions.

Fig. 25. SEM micrographs of (a) tin oxide particles and (b) magnified area of (a). Reprinted with permission from Ref.¹⁴⁶, Masuda, Y. and Kato, K., 2009, *J. Cryst. Growth*, 311, 593.

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N₂ adsorption characteristics of nano-sheet assembled tin oxide particles. The particles exhibited N₂ adsorption-desorption isotherms of Type IV. The desorption isotherm differed from adsorption isotherm in the relative pressure (P/P_0) range from 0.45 to 0.97, showing mesopores in the particles. BET surface area of the particles was estimated to be 85 m²/g. This is higher than that of SnO₂ nanoparticles such as SnO₂ (BET 47.2 m²/g, 18.3nm in diameter, No. 549657-25G, Aldrich), SnO₂ (BET 25.9 m²/g, 34nm in diameter, Yamanaka & Co., Ltd.), SnO₂ (BET 23 m²/g, 26nm in diameter, No. 37314-13, NanoTek, C. I. Kasei Co.,Ltd.) and In₂O₃-SnO₂ (BET 3-6m²/g, 100-300nm in diameter, Sumitomo Chemical Co., Ltd.). The particles were assembly of nanosheets. Unique morphologies of the sheets contributed to increase surface area of the particles.

Total pore volume and average pore diameter were estimated to 0.343 cc/g and 16.1 nm, respectively from pores smaller than 259 nm at $P/P_0 = 0.9925$. They were estimated to be 0.088 cc/g and 4.1 nm, respectively, from pores smaller than 10.6 nm at $P/P_0 = 0.7994$. Total pore volume was also estimated by the BJH method from pores smaller than 174 nm to be 0.354 cc/g.

Pore size distribution was calculated by the BJH method using desorption braches. It showed a pore size distribution curve having a strong peak at ~3.9 nm and a broad peak

from 25 nm to 175 nm. The particles would have a large amount of mesopores with ~ 3.9 nm in diameter and mesopores with 25-175 nm in diameter. The pores were spaces surrounded by nanosheets and interparticle spaces. Micropores smaller than 2 nm were not suggested by BJH pore size distribution.

In this section, nanosheet assembled tin oxide particles were fabricated by aqueous solution synthesis. Anisotropic crystal growth of tin oxide was effectively utilized to form nanosheet assembled structure. They were crystallized at ordinary temperature without annealing. It allowed us to avoid aggregation of the particles and decrease of surface area. The particles were 300-800 nm in diameter and were crystals of SnO_2 and SnO . The sheets were about 50-100 nm in size and 5-10 nm thickness. The particles had BET surface area of $85 \text{ m}^2/\text{g}$ estimated with N_2 adsorption characteristics. BJH analysis indicated that mesopores of 3.9 nm in size contributed to increase surface area. The particles were prepared by environmentally friendly process. The system had advantages on low cost, low energy consumption and low CO_2 emission.

9. Enhancement of surface area of tin oxide¹⁸⁰

Tin oxide nano-crystals with high surface area were firstly synthesized in aqueous solutions at 50°C . BET surface area was successfully reached to $194 \text{ m}^2/\text{g}$ ¹⁸⁰. It was much higher than that of SnO_2 (BET $47.2 \text{ m}^2/\text{g}$, Aldrich), SnO_2 (BET $25.9 \text{ m}^2/\text{g}$, Yamanaka & Co., Ltd.), SnO_2 (BET $23 \text{ m}^2/\text{g}$, C. I. Kasei Co., Ltd.) and $\text{In}_2\text{O}_3\text{-SnO}_2$ (BET $3\text{-}6 \text{ m}^2/\text{g}$, Sumitomo Chemical Co., Ltd.). N_2 adsorption characteristics revealed that they had pores of 1-3 nm which contributed high surface area. TEM, ED and XRD indicated morphology, crystal structure and chemical composition of nano-crystals. Novel process allowed us to avoid sintering and deformation of the crystals and hence, realized high surface area and unique morphology.

In this study, tin oxide nanosheets were formed in the solutions. BET surface area was successfully reached to $194 \text{ m}^2/\text{g}$. Origin of high surface area was discussed with pore size distribution and morphology observations.

Synthesis of Tin Oxide Nanosheets. SnF_2 (870.6 mg) was dissolved in distilled water (200 mL) of 50°C to be 5 mM. The solutions were kept at 50°C for 20 min and then at 28°C for 3 days without stirring. The nanocrystals precipitated to cover bottom of the vessels. For comparison, the solutions were centrifuged at 4000 rpm for 10 min after keeping at 50°C for 20 min. Precipitated particles were dried at 60°C for 12 h after removal of supernatant solutions.

Morphology and crystal phase of tin oxide nanosheets.

The nanosheets synthesized at 50°C for 20 min and at 28°C for 3 days were mixture of SnO_2 main phase and SnO additional phase (Fig. 26a). X-ray diffraction peaks at $2\theta = 26.5, 33, 51.4, 62, 64.5, 80$ and 89 were assigned to 110, 101, 211, etc. of SnO_2 (JDPDS No. 41-1445). 101, 110 and 002 diffraction peaks of SnO (JDPDS No. 06-0395) were overlapped to peaks of SnO_2 . For comparison, diffraction pattern of the nanosheets synthesized at 50°C for 20 min was shown in Fig. 26b. Half maximum full-width of the peaks was smaller than that in Fig. 26a. Lower SnO content resulted sharp peaks of SnO_2 in Fig. 26b. These observations indicated that SnO was mainly formed at 28°C rather than at 50°C .

Fig. 26. XRD patterns of (a) tin oxide nanosheets fabricated at 50°C for 20min and at 28°C for 3 days, and (b) tin oxide nanosheets fabricated at 50°C for 20min. Reprinted with permission from Ref.¹⁸⁰, Masuda, Y., Ohji, T. and Kato, K., 2010, *J. Am. Ceram. Soc.*, 93(8), 2140. Copyright ©Wiley

The nanosheets were well dispersed in ethanol. The nanosheets in supernatant solutions were skimmed with Cu grids for TEM observations. They were 20-50 nm in diameter having uniform thickness (Fig. 27A). Similar structures were observed from many areas. We observed that they curled up during long-term observation. It indicated that they had sheet structure. Electron beams damaged them to transform the structures. Some of them tightly connected each other (Fig. 27B). They had clear interfaces without pores or small grains. The nanosheet showed electron diffractions (Fig. 27B, Insert). Lattice spacing calculated from spots indicated with a red line and a yellow line were 0.283 nm and 0.154 nm, respectively. They can be assigned to SnO₂ and/or SnO. The former one can be assigned to 101 crystal plane of SnO₂ (0.264 nm) or 110 crystal plane of SnO (0.269 nm). The later one can be assigned to 310 crystal plane of SnO₂ (0.149 nm), 221 crystal plane of SnO₂ (0.148 nm), 202 crystal plane of SnO (0.149 nm) or 103 crystal plane of SnO (0.148 nm). Additionally, diffraction spots related to lattice spacing of 0.283 nm were observed at upper part. They were shown with two white circles and a red line in Fig. 27B Insert. The double spots indicated that the area shown in a white circle in Fig. 27B was consisted of two crystals, i.e., stacked two nanosheets. High magnification image was also obtained from other observation area (Fig. 27C). The structure was thinner than that in Fig. 27A and 27B. Clear image was not obtained in low magnification images due to low contrast. However, it showed clear high magnification image and lattice fringe (Fig. 27C). Electron diffraction patterns showed lattice spacing of 0.277 nm (red line), 0.279 nm (green line) and 0.196 nm

(yellow line). They were assigned to 110, 1-10 and 200 crystal planes of SnO, respectively. Lattice spacing and their angles were well matched to that of SnO. Chemical composition was estimated from several points of tightly-packed area. The area included nanosheets and spherical crystals. Chemical composition varied among each positions in the range of Sn : O = 1 : 1.7-2.7 which was similar to that of SnO₂ rather than SnO. These observations indicated that the crystals were mixture of SnO₂ and SnO.

Fig. 27. TEM micrographs and electron diffraction patterns of tin oxide nanosheets. (b): high magnification image of (a). (c): high magnification image of other area. Reprinted with permission from Ref.¹⁸⁰, Masuda, Y., Ohji, T. and Kato, K., 2010, *J. Am. Ceram. Soc.*, 93(8), 2140. Copyright @Wiley

N₂ adsorption characteristics of tin oxide nanosheets.

BET surface area of the nanosheets was estimated from N₂ adsorption isotherm (Fig. 28a). The nanosheets had successfully high surface area of 194 m²/g (Fig. 28b). It was much higher than that of SnO₂ nanoparticles such as SnO₂ (BET 47.2 m²/g, 18.3nm in diameter, No. 549657-25G, Aldrich), SnO₂ (BET 25.9 m²/g, 34nm in diameter, Yamanaka & Co., Ltd., Osaka, Japan), SnO₂ (BET 23 m²/g, 26nm in diameter, No. 37314-13, NanoTek, C. I. Kasei Co.,Ltd., Tokyo, Japan) and In₂O₃-SnO₂ (BET 3-6m²/g, 100-300nm in diameter, Sumitomo

Chemical Co., Ltd.). Additionally, it was more than double higher than that of previous report ($85 \text{ m}^2/\text{g}$)¹⁸¹. Pore size distribution was analyzed with BJH method using adsorption isotherm (Fig. 28c). It indicated that the nanosheets included pores of 1-2 nm. Micropore analysis was performed with DFT/Monte-Carlo Fitting which was completely-consistent with isotherms (Fig. 28d). Pores of 1-3 nm were shown to be in the nanosheets (Fig. 28e). The micro and meso-pores contributed high surface area of $194 \text{ m}^2/\text{g}$ in this system. For comparison, BET surface area of the nanosheets synthesized at 50°C for 20min was estimated to $146 \text{ m}^2/\text{g}$. It was also higher than that of nanoparticles in former studies.

Fig. 28. (a): N_2 adsorption-desorption isotherm of tin oxide nanosheets. (b): BET surface area of tin oxide nanosheets. (c): Pore size distribution calculated from N_2 adsorption data of tin oxide nanosheets using BJH equation. (d): N_2 adsorption-desorption isotherm and DFT/Monte-Carlo fitting curve of tin oxide nanosheets. (e): Pore size distribution calculated from N_2 adsorption data of tin oxide nanosheets using DFT/Monte-Carlo equation. Reprinted with permission from Ref.¹⁸⁰, Masuda, Y., Ohji, T. and Kato, K., 2010, *J. Am. Ceram. Soc.*, 93(8), 2140. Copyright ©Wiley

In this section, nanosheets of tin oxides were fabricated in aqueous solutions at ordinary temperature. BET surface area successfully reached to 194 m²/g. It was much higher than that of nanoparticles in former studies. 2-dimensional sheet structure was one of ideal structures for high surface area per unit weight. Nano sized thickness directly contributed high surface area. Crystalline nanosheets were prepared without high temperature annealing which degraded surface area and nanostructures. High surface area and unique nanostructures of the sheets can be applied to gas sensors, dye-sensitized solar cells and molecular sensors.

10. Tin oxide coating on polytetrafluoroethylene films in aqueous solutions¹⁸²

Tin Polytetrafluoroethylene(PTFE) films were successfully coated with tin oxide in aqueous solutions. Tin oxide was crystallized in the solution and formed nanocrystal coatings on the polymer films. The coatings were consisted of SnO₂ and SnO crystals. They were assemblies of tin oxide nanosheet of about 10 nm to 50 nm in size and about 5 nm in thickness. The nanocrystal films can be exfoliated from the PTFE substrates. Tin oxide nanocrystal films had a rough liquid-surface and a dense substrate-side surface. Transparency of PTFE films coated with tin oxide was same to that of bare PTFE films in the range from 400 nm to 800 nm. Tin oxide decreased transparency about 25 % at 320 nm. The PTFE films coated with tin oxide nanocrystals can be pasted on desired substrates.

Organic-inorganic hybrid materials such as metal oxide electronics on polymer flexible films have received considerable attention in recent years for light-weight flexible sensors, displays, dye-sensitized solar cells, etc. The polymer films offer the advantages of flexibility, light-weight, low-cost and impact resistance.

Tin oxide (SnO₂) is an important semiconductor with a wide band gap of 3.6 eV at room temperature. They have been widely used in gas sensors^{167,183}, optical devices¹⁴⁷, lithium batteries¹⁴⁸⁻¹⁵⁰, etc. A novel type of bio-sensor was proposed to detect environmental toxins such as bisphenol-A or dioxin^{88,184}. Tin oxide is a candidate material for the sensor because of its suitable band gap, surface characteristics, and high transparency.

Metal oxides including tin oxide have been synthesized with high temperature processes at several hundred degrees for many years. Recently, the aqueous syntheses of metal oxides have attracted much attention as a next generation science and technology^{1,51,114,177}. Aqueous systems are environmentally friendly and have advantages of low energy consumption, low cost and an organic solvent free process.

In recent years, metal oxide films and their microstructures were fabricated on organic surfaces—such as polyethylene terephthalate (PET) films^{185,186} or self-assembled monolayers^{20,113,187,188}. However, metal oxide formation on polytetrafluoroethylene (PTFE) was difficult compared to PET films. PTFE films are widely used in electronic applications because of low chemical reactivity, low coefficient of friction, high melting point (327 °C), high-corrosion resistance, a high dielectric strength over many different frequencies, a low dissipation factor, and a high surface resistivity. They were selected as substrates in this study.

In this section, tin oxide nanocrystals were prepared on PTFE films in aqueous solutions^{182,184}. They were crystallized in the solution containing tin ions to form films consisted of nanosheets. The process realized tin oxide film formation without high temperature annealing and unique morphology of tin oxide crystals.

1. Aqueous synthesis of tin oxide nanocrystals on PTFE films

SnF_2 (870.6 mg) was dissolved in the distilled water (200 mL) at 90°C to be 5 mM. PTFE films (thickness: 50 μm , ASF-110, Chukoh) with silicone adhesive (thickness: 30 μm) were pasted on quartz substrates ($25 \times 50 \times 1$ mm). They were immersed in the middle of the solutions with the bottom up at an angle or with the top up at an angle. They were tilted at 15 degrees to the upright. The solutions were kept at 90°C using a drying oven (Yamato Scientific Co., Ltd., DKN402) for 2 h with no stirring. The solutions became lightly clouded after 2 h. The as-deposited nanocrystals on substrates were rinsed under running water and dried by a strong air spray. Additionally, the solutions kept at 90°C for 2 h were centrifuged at 4000 rpm for 10 min (Model 8920, Kubota Corp.). Precipitated particles were dried at 20°C for 12 h after the removal of supernatant solutions.

2. Morphology of tin oxide nanocrystals on PTFE films

Bare PTFE films pasted on quartz substrates had cracks of about 50 nm to 200 nm in length (Fig. 29). Longer directions of cracks were perpendicular to the extensional direction. They were formed during adhesive processes. The quartz substrates and the PTFE films were transparent or slightly white, respectively. The PTFE films can be pasted on desired substrates such as quartz, metals and polymers with silicone adhesive.

Fig. 29. SEM micrograph of a bare PTFE film. Reprinted with permission from Ref.¹⁸², Masuda, Y. and Kato, K., 2010, *Polym. Adv. Technol.*, 21(3), 211. Copyright @Wiley

The surface of PTFE films was completely covered with assemblies of nanosheets (Fig. 30a). Uniform formation of tin oxide coatings is one of the advantages of solution processes. Large sheets were also observed from the surfaces (Fig. 30b). They were about 200 nm to 300 nm in size and about 10 nm in thickness (Fig. 30b). Some of them stood perpendicular to the PTFE films at an angle. They had an angular outline, which was connected by straight lines. They were caused from anisotropic crystal growth of tin oxide reflected in their crystal structure. The large sheets connected in a cross shape were also observed at the lower right (Fig. 30b). Nanosheets were roughly estimated to about 10 nm to 50 nm in size and about 5 nm in thickness (Fig. 30c). They connected to each other to form continuous films on the PTFE surfaces. The tin oxide films were exfoliated from PTFE films by scratch using a metal spatula. Exfoliated tin oxide films were placed on a substrate for SEM observation. Three exfoliated tin oxide films were observed to be partially-overlapping (Fig. 30d). The top film and bottom film showed their liquid-side surfaces. They were similar to that observed in Fig. 30a-c. The middle film showed a substrate-side surface, which was contacted with PTFE film during the immersion period. Substrate-side surface of the middle film was indicated

by a white arrow. It had a dense surface, which consisted of nanocrystals of about 5 nm to 10 nm in size. These observations indicated that dense films consisted of nanocrystals of about 5 nm to 10 nm in size were formed on the PTFE films at first stage of tin oxide film formation. Tin oxide then grew to sheet-shapes to form assemblies of nanosheets. Additionally, large sheet crystals grew on the tin oxide dense films.

Fig. 30. SEM micrographs of (a) tin oxide nanocrystals on a PTFE film and (b, c) magnified area of (a) showing morphology of nanocrystals and (d) exfoliated tin oxide nanocrystal films showing liquid-side surface and substrate-side surface. Substrate-side surface is indicated by a white arrow. Reprinted with permission from Ref.¹⁸², Masuda, Y. and Kato, K., 2010, *Polym. Adv. Technol.*, 21(3), 211. Copyright ©Wiley

3. Crystal phase of tin oxide nanocrystals on PTFE films

X-ray diffraction pattern obtained from a PTFE film coated with tin oxide was similar to that from a bare PTFE film. Diffraction peaks assigned to tin oxide were not observed because of thin film thickness.

Precipitated particles were evaluated after drying. They showed broad X-ray diffraction peaks at $2\theta = 27, 34, 38$ and 52 . They were assigned to 110, 101, 200 or 211 diffraction peaks from SnO_2 (JDPDS No. 41-1445) (Fig. 3c). The peaks were wide in width due to the small crystallite size of SnO_2 . Peaks were also observed $2\theta = 29.5, 32.0, 37.4, 48.0, 50.4$ and 57.4 . They were assigned to 101, 110, 002, 200, 112 or 211 diffraction peaks from SnO (JDPDS No. 06-0395).

XRD analysis indicated that the particles obtained from the solutions consisted of SnO_2 crystals and SnO crystals.

4. XPS analysis of tin oxide nanocrystals on PTFE films

Carbon, fluorine and oxygen were detected from bare PTFE films (Fig. 31b). Chemical ratio of them was estimated to $\text{C} : \text{F} : \text{O} = 1 : 2.05 : 0.02$. It was consistent with chemical composition of polytetrafluoroethylene (PTFE, $\text{C} : \text{F} = 1 : 2$). Small amount of oxygen was detected from surface contamination. PTFE films coated with tin oxide showed spectra of tin, oxygen, fluorine and carbon (Fig. 31a). Spectral peak corresponding to $\text{Sn } 3d_{5/2}$ was observed at 487.2 eV (Fig. 31c1). The binding energy was similar to that of SnO_2 (486.3 eV¹⁸⁹,

486.5 eV¹⁹⁰, 486.6 eV¹⁹¹, 487.3 eV) and higher than that of Sn metal (484.8 eV, 484.85 eV, 484.87 eV, 484.9 eV, 485.0 eV), which suggested tin atoms in surface coatings were positively charged by forming direct bonds with oxygen. Binding energy of O 1s centered at about 531.2 eV corresponds to that of SnO₂ (Fig. 31c2). Chemical ratio of the coatings was estimated to Sn : O : F : C = 1 : 1.88 : 7.25 : 3.82 and C : F = 1 : 1.90. It was indicated that the surface coatings consisted mainly of SnO₂ (Sn : 1 : 2). They were formed on polytetrafluoroethylene films (PTFE, C : F = 1 : 2). Difference and similarity between XPS analyses and XRD analyses suggested deposition mechanism of tin oxides. Crystal phases and chemical compositions of the surface coatings were different from those of the precipitated particles. SnO₂ crystallized on the films to form surface coatings, on the other hand, SnO₂ and SnO homogeneously crystallized to form particles in the solutions. Pure SnO₂ coatings were thus successfully formed on the PTFE films.

Fig. 31. XPS spectra of (a) tin oxide nanocrystals on a PTFE film and (b) a bare PTFE film. (c1) Sn 3d_{5/2} spectrum and (c2) O 1s spectrum of tin oxide nanocrystals on a PTFE film.

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5. Optical property of tin oxide nanocrystals on PTFE films

Quartz substrates had high transparency in the range from 200 nm to 850 nm (Fig. 32, black line). PTFE films pasted on quartz substrates were visually observed to be slight white. Their transparency was lower than that of quartz (Fig. 32, red line). Especially, it decreased as decrease of wavelength below 350 nm. PTFE films coated with tin oxide showed the same transparency as bare PTFE films in the range from 400 nm to 850 nm (Fig. 32, blue line). They decreased transparency about 25 % at 320 nm. Tin oxide particles precipitated from the solution were evaluated for comparison. They had absorption peak centered at 320 nm. This absorption was caused by tin oxide. These analyses indicated that tin oxide was deposited on PTFE films immersed in the solutions and they decrease transparency at 320 nm.

Polytetrafluoroethylene (PTFE) films were successfully coated with tin oxide nanocrystals. Tin oxide was crystallized in aqueous solutions to form nanosheet-assembled films. They were about 10 nm to 50 nm in size and about 5 nm in thickness. Large sheets of about 200 nm to 300 nm in size and about 10 nm in thickness were also crystallized on the surfaces. X-ray diffraction analysis indicated that tin oxide was a mixture of SnO₂ and SnO. PTFE films coated with tin oxide were transparent in the range from 400 nm to 850 nm. Tin oxide on the films had absorption centered at 320 nm. Tin oxide coated PTFE films can be pasted on desired substrates. Hybrid tin oxide-PTFE composites may be useful for increasing the potential application of tin oxide film as flexible electronics.

Fig. 32. UV-Vis spectra of (a) a quartz substrate (black line), (b) a PTFE film pasted on a quartz substrate (red line) and (c) tin oxide nanocrystals on a PTFE film pasted on a quartz substrate (blue line). UV-Vis spectrum of tin oxide particles. Reprinted with permission from Ref.¹⁸², Masuda, Y. and Kato, K., 2010, *Polym. Adv. Technol.*, 21(3), 211. Copyright @Wiley

11. Summary

Liquid phase morphology control of metal oxide was developed by precise control of crystal growth in this study. Various nano/micro-structures of them were fabricated in solution systems. These would contribute to development of future metal oxide devices and develop new academic fields.

12. References

- [1] Masuda, Y.; Yamada, T.; Koumoto, K. *Cryst. Growth Des.* 2008, 8, 169-171.
- [2] Tanaka, K.; Suzuki, K.; Fu, D. S.; Nishizawa, K.; Miki, T.; Kato, K. *Jpn. J. Appl. Phys.* 2004, 43, 6525-6529.
- [3] Aoyagi, S.; Kuroiwa, Y.; Sawada, A.; Kawaji, H.; Atake, T. *J. Therm. Anal. Cal.* 2005, 81, 627-630.
- [4] Luan, W. L.; Gao, L.; Guo, J. K. *Ceram. Inter.* 1999, 25, 727-729.
- [5] Wang, X. H.; Chen, R. Z.; Gui, Z. L.; Li, L. T. *Mater. Sci. Eng. B* 2003, 99, 199-202.
- [6] Sakabe, Y.; Yamashita, Y.; Yamamoto, H. *J. Euro. Ceram. Soc.* 2005, 25, 2739-2742.
- [7] Polotai, A. V.; Ragulya, A. V.; Randall, C. A. *Ferroelectrics* 2003, 288, 93-102.
- [8] Yashima, M.; Hoshina, T.; Ishimura, D.; Kobayashi, S.; Nakamura, W.; Tsurumi, T.; Wada, S. *J. Appl. Phys.* 2005, 98.
- [9] Frey, M. H.; Xu, Z.; Han, P.; Payne, D. A. *Ferroelectrics* 1998, 206, 337-353.
- [10] Frey, M. H.; Payne, D. A. *Phys. Rev. B* 1996, 54, 3158-3168.
- [11] Wada, S.; Suzuki, T.; Noma, T. *J. Ceram. Soc. Japan* 1996, 104, 383-392.
- [12] Masuda, Y.; Koumura, T.; Okawa, T.; Koumoto, K. *J. Colloid Interface Sci.* 2003, 263, 190-195.
- [13] Masuda, Y.; Koumoto, K.; Ueyama, R. Japan, 2007.
- [14] Hayashi, Y.; Kimura, T.; Yamaguchi, T. *J. Mater. Sci.* 1986, 21, 757-762.
- [15] Li, X. L.; Liu, J. F.; Li, Y. D. *Mater. Chem. Phys.* 2003, 80, 222-227.
- [16] Sun, H.; Kang, S. Z.; Mu, J. *Mater. Lett.* 2007, 61, 4121-4123.

- [17] Mutin, J. C.; Dusauroy, Y.; Protas, J. J. *Solid State Chem.* 1981, 36, 356-364.
- [18] Potdar, H. S.; Deshpande, S. B.; Date, S. K. *J. Am. Ceram. Soc.* 1996, 79, 2795-2797.
- [19] Masuda, Y.; Kinoshita, N.; Koumoto, K. *J. Nanosci. Nanotechnol.*, in press.
- [20] Masuda, Y.; Kinoshita, N.; Sato, F.; Koumoto, K. *Cryst. Growth Des.* 2006, 6, 75-78.
- [21] Wu, X. L.; Siu, G. G.; Fu, C. L.; Ong, H. C. *Appl. Phys. Lett.* 2001, 78, 2285-2287.
- [22] Kang, J. S.; Kang, H. S.; Pang, S. S.; Shim, E. S.; Lee, S. Y. *Thin Solid Films* 2003, 443, 5-8.
- [23] Golego, N.; Studenikin, S. A.; Cocivera, M. J. *J. Electrochem. Soc.* 2000, 147, 1592-1594.
- [24] Sberveglieri, G. *Sens. Actuators B: Chem.* 1995, 23, 103-109.
- [25] Pauporte, T.; Lincot, D. *Electrochem. Acta.* 2000, 45, 3345-3353.
- [26] Law, M.; Greene, L. E.; Johnson, J. C.; Saykally, R.; Yang, P. D. *Nature Mater.* 2005, 4, 455-459.
- [27] Masuda, Y.; Kato, K. *Cryst. Growth Des.* 2008, 8, 2633-2637.
- [28] Gong, H.; Hu, J. Q.; Wang, J. H.; Ong, C. H.; Zhu, F. R. *Sens. Actuators B: Chem.* 2006, 115, 247-251.
- [29] Moon, W. J.; Yu, J. H.; Choi, G. M. *Sens. Actuators B: Chem.* 2002, 87, 464-470.
- [30] Chang, J. F.; Kuo, H. H.; Leu, I. C.; Hon, M. H. *Sens. Actuators B: Chem.* 2002, 84, 258-264.
- [31] Tang, H. X.; Yan, M.; Zhang, H.; Li, S. H.; Ma, X. F.; Wang, M.; Yang, D. R. *Sens. Actuators B: Chem.* 2006, 114, 910-915.
- [32] Shishiyuan, S. T.; Shishiyuan, T. S.; Lupan, O. I. *Sens. Actuators B: Chem.* 2005, 107, 379-386.
- [33] Wagh, M. S.; Patil, L. A.; Seth, T.; Amalnerkar, D. P. *Mater. Chem. Phys.* 2004, 84, 228-233.
- [34] Xu, J. Q.; Pan, Q. Y.; Shun, Y. A.; Tian, Z. Z. *Sens. Actuators B: Chem.* 2000, 66, 277-279.
- [35] Paraguay, D. F.; Miki-Yoshida, M.; Morales, J.; Solis, J.; Estrada, L. W. *Thin Solid Films* 2000, 373, 137-140.
- [36] Baxter, J. B.; Aydil, E. S. *Appl. Phys. Lett.* 2005, 86, 53114.
- [37] Katoh, R.; Furube, A.; Hara, K.; Murata, S.; Sugihara, H.; Arakawa, H.; Tachiya, M. *J. Phys. Chem. B* 2002, 106, 12957-12964.
- [38] Karuppuchamy, S.; Nonomura, K.; Yoshida, T.; Sugiura, T.; Minoura, H. *Solid State Ionics* 2002, 151, 19-27.
- [39] Keis, K.; Bauer, C.; Boschloo, G.; Hagfeldt, A.; Westermarck, K.; Rensmo, H.; Siegbahn, H. J. *Photochem. Photobiol. A* 2002, 148, 57-64.
- [40] Kasuga, T.; Hiramatsu, M.; Hoson, A.; Sekino, T.; Niihara, K. *Adv. Mater.* 1999, 11, 1307-+.
- [41] Oaki, Y.; Imai, H. *J. Am. Chem. Soc.* 2004, 126, 9271-9275.
- [42] Xia, Y. N.; Yang, P. D.; Sun, Y. G.; Wu, Y. Y.; Mayers, B.; Gates, B.; Yin, Y. D.; Kim, F.; Yan, Y. Q. *Adv. Mater.* 2003, 15, 353-389.
- [43] Masuda, Y.; Saito, N.; Hoffmann, R.; De Guire, M. R.; Koumoto, K. *Sci. Tech. Adv. Mater.* 2003, 4, 461-467.
- [44] Masuda, Y.; Ieda, S.; Koumoto, K. *Langmuir* 2003, 19, 4415-4419.
- [45] Masuda, Y.; Jinbo, Y.; Yonezawa, T.; Koumoto, K. *Chem. Mater.* 2002, 14, 1236-1241.
- [46] Nakanishi, T.; Masuda, Y.; Koumoto, K. *Chem. Mater.* 2004, 16, 3484-3488.

- [47] Peng, W. Q.; Qu, S. C.; Cong, G. W.; Wang, Z. G. *Cryst. Growth Des.* 2006, 6, 1518-1522.
- [48] Lin, Y. R.; Yang, S. S.; Tsai, S. Y.; Hsu, H. C.; Wu, S. T.; Chen, I. C. *Cryst. Growth Des.* 2006, 6, 1951-1955.
- [49] Zhang, H.; Yang, D. R.; Li, D. S.; Ma, X. Y.; Li, S. Z.; Que, D. L. *Cryst. Growth Des.* 2005, 5, 547-550.
- [50] Liu, K. H.; Lin, C. C.; Chen, S. Y. *Cryst. Growth Des.* 2005, 5, 483-487.
- [51] Masuda, Y.; Kato, K. *Cryst. Growth Des.* 2008, 8, 275-279.
- [52] Greene, L. E.; Law, M.; Tan, D. H.; Montano, M.; Goldberger, J.; Somorjai, G.; Yang, P. D. *Nano Lett.* 2005, 5, 1231-1236.
- [53] O'Brien, P.; Saeed, T.; Knowles, J. J. *Mater. Chem.* 1996, 6, 1135-1139.
- [54] Yamabi, S.; Imai, H. *J. Mater. Chem.* 2002, 12, 3773-3778.
- [55] Vayssieres, L.; Keis, K.; Lindquist, S. E.; Hagfeldt, A. *J. Phys. Chem. B* 2001, 105, 3350-3352.
- [56] Vayssieres, L. *Adv. Mater.* 2003, 15, 464-466.
- [57] Saito, N.; Haneda, H.; Sekiguchi, T.; Ohashi, N.; Sakaguchi, I.; Koumoto, K. *Adv. Mater.* 2002, 14, 418-421.
- [58] Saito, N.; Haneda, H.; Seo, W. S.; Koumoto, K. *Langmuir* 2001, 17, 1461-1469.
- [59] Saeed, T.; O'Brien, P. *Thin Solid Films* 1995, 271, 35-38.
- [60] Kakiuchi, K.; Hosono, E.; Kimura, T.; Imai, H.; Fujihara, S. *J. Sol-Gel Sci. Technol.* 2006, 39, 63-72.
- [61] Schwenzer, B.; Gomm, J. R.; Morse, D. E. *Langmuir* 2006, 22, 9829-9831.
- [62] Yu, H. D.; Zhang, Z. P.; Han, M. Y.; Hao, X. T.; Zhu, F. R. *J. Am. Chem. Soc.* 2005, 127, 2378-2379.
- [63] Choi, K. S.; Lichtenegger, H. C.; Stucky, G. D.; McFarland, E. W. *J. Am. Chem. Soc.* 2002, 124, 12402-12403.
- [64] Yin, M.; Gu, Y.; Kuskovsky, I. L.; Andelman, T.; Zhu, Y.; Neumark, G. F.; O'Brien, S. J. *Am. Chem. Soc.* 2004, 126, 6206-6207.
- [65] Feng, X. J.; Feng, L.; Jin, M. H.; Zhai, J.; Jiang, L.; Zhu, D. B. *J. Am. Chem. Soc.* 2004, 126, 62-63.
- [66] Yoshida, T.; Tochimoto, M.; Schlettwein, D.; Wohrle, D.; Sugiura, T.; Minoura, H. *Chem. Mater.* 1999, 11, 2657-2667.
- [67] Lee, J. Y.; Yin, D. H.; Horiuchi, S. *Chem. Mater.* 2005, 17, 5498-5503.
- [68] Kopalko, K.; Godlewski, M.; Domagala, J. Z.; Lusakowska, E.; Minikayev, R.; Paszkowicz, W.; Szczerbakow, A. *Chem. Mater.* 2004, 16, 1447-1450.
- [69] Turgeman, R.; Gershevitz, O.; Deutsch, M.; Ocko, B. M.; Gedanken, A.; Sukenik, C. N. *Chem. Mater.* 2005, 17, 5048-5056.
- [70] Turgeman, R.; Gershevitz, O.; Palchik, O.; Deutsch, M.; Ocko, B. M.; Gedanken, A.; Sukenik, C. N. *Cryst. Growth Des.* 2004, 4, 169-175.
- [71] Mirica, E.; Kowach, G.; Evans, P.; Du, H. *Cryst. Growth Des.* 2004, 4, 147-156.
- [72] Lin, Y. R.; Tseng, Y. K.; Yang, S. S.; Wu, S. T.; Hsu, C. L.; Chang, S. J. *Cryst. Growth Des.* 2005, 5, 579-583.
- [73] Gao, Y. F.; Nagai, M. *Langmuir* 2006, 22, 3936-3940.
- [74] Wu, X. D.; Zheng, L. J.; Wu, D. *Langmuir* 2005, 21, 2665-2667.
- [75] Peterson, R. B.; Fields, C. L.; Gregg, B. A. *Langmuir* 2004, 20, 5114-5118.

- [76] Liu, T. Y.; Liao, H. C.; Lin, C. C.; Hu, S. H.; Chen, S. Y. *Langmuir* 2006, 22, 5804-5809.
- [77] Masuda, Y.; Kato, K. *Cryst. Growth Des.* 2008, 8, 3213-3218.
- [78] Carlson, T.; Giffin, G. L. *J. Phys. Chem.* 1986, 90, 5896-5900.
- [79] Zhang, Z. B.; Wang, C. C.; Zakaria, R.; Ying, J. Y. *J. Phys. Chem. B* 1998, 102, 10871-10878.
- [80] Wang, R.; Hashimoto, K.; Fujishima, A. *Nature* 1997, 388, 431-432.
- [81] Choi, W. Y.; Termin, A.; Hoffmann, M. R. *J. Phys. Chem.* 1994, 98, 13669-13679.
- [82] Sung, Y. M.; Lee, J. K.; Chae, W. S. *Cryst. Growth Des.* 2006, 6, 805-808.
- [83] Kumazawa, N.; Islam, M. R.; Takeuchi, M. *J. Electroanal. Chem.* 1999, 472, 137-141.
- [84] Ferroni, M.; Carotta, M. C.; Guidi, V.; Martinelli, G.; Ronconi, F.; Sacerdoti, M.; Traversa, E. *Sens. Actuators B: Chem.* 2001, 77, 163-166.
- [85] Wagemaker, M.; Kentgens, A. P. M.; Mulder, F. M. *Nature* 2002, 418, 397-399.
- [86] Arico, A. S.; Bruce, P.; Scrosati, B.; Tarascon, J. M.; Van Schalkwijk, W. *Nature Mater.* 2005, 4, 366-377.
- [87] Guo, Y. G.; Hu, Y. S.; Maier, J. *Chem. Commun.* 2006, 26, 2783-2785.
- [88] Tokudome, H.; Yamada, Y.; Sonezaki, S.; Ishikawa, H.; Bekki, M.; Kanehira, K.; Miyauchi, M. *Appl. Phys. Lett.* 2005, 87, 213901-213903.
- [89] Nazeeruddin, M. K.; De Angelis, F.; Fantacci, S.; Selloni, A.; Viscardi, G.; Liska, P.; Ito, S.; Takeru, B.; Gratzel, M. G. *J. Am. Chem. Soc.* 2005, 127, 16835-16847.
- [90] Wang, P.; Zakeeruddin, S. M.; Moser, J. E.; Humphry-Baker, R.; Comte, P.; Aranyos, V.; Hagfeldt, A.; Nazeeruddin, M. K.; Gratzel, M. *Adv. Mater.* 2004, 16, 1806-1811.
- [91] Morrison, P. W.; Raghavan, R.; Timpone, A. J.; Artelt, C. P.; Pratsinis, S. E. *Chem. Mater.* 1997, 9, 2702-2708.
- [92] Yang, G. X.; Zhuang, H. R.; Biswas, P. *Nanostruct. Mater.* 1996, 7, 675-689.
- [93] Yu, J. C.; Yu, J. G.; Ho, W. K.; Zhang, L. Z. *Chem. Commun.* 2001, 19, 1942-1943.
- [94] Huang, W. P.; Tang, X. H.; Wang, Y. Q.; Koltypin, Y.; Gedanken, A. *Chem. Commun.* 2000, 15, 1415-1416.
- [95] Seifried, S.; Winterer, M.; Hahn, H. *Chem. Vap. Deposition* 2000, 6, 239-244.
- [96] Scolan, E.; Sanchez, C. *Chem. Mater.* 1998, 10, 3217-3223.
- [97] Wang, C. C.; Ying, J. Y. *Chem. Mater.* 1999, 11, 3113-3120.
- [98] Burnside, S. D.; Shklover, V.; Barbe, C.; Comte, P.; Arendse, F.; Brooks, K.; Gratzel, M. *Chem. Mater.* 1998, 10, 2419-2425.
- [99] Zhang, H. Z.; Finnegan, M.; Banfield, J. F. *Nano Lett.* 2001, 1, 81-85.
- [100] Sung, Y. M.; Lee, J. K. *Cryst. Growth Des.* 2004, 4, 737-742.
- [101] Perkas, N.; Pol, V.; Pol, S.; Gedanken, A. *Cryst. Growth Des.* 2006, 6, 293-296.
- [102] Deki, S.; Aoi, Y.; Hiroi, O.; Kajinami, A. *Chem. Lett.* 1996, 6, 433-434.
- [103] Deki, S.; Aoi, Y.; Yanagimoto, H.; Ishii, K.; Akamatsu, K.; Mizuhata, M.; Kajinami, A. *J. Mater. Chem.* 1996, 6, 1879-1882.
- [104] Deki, S.; Aoi, Y.; Asaoka, Y.; Kajinami, A.; Mizuhata, M. *J. Mater. Chem.* 1997, 7, 733-736.
- [105] Kishimoto, H.; Takahama, K.; Hashimoto, N.; Aoi, Y.; Deki, S. *J. Mater. Chem.* 1998, 8, 2019-2024.

- [106] Choi, S. Y.; Mamak, M.; Coombs, N.; Chopra, N.; Ozin, G. A. *Adv. Func. Mater.* 2004, 14, 335-344.
- [107] Antonelli, D. M.; Ying, J. Y. *Angew. Chem., Int. Ed.* 1995, 34, 2014-2017.
- [108] Shibata, H.; Ogura, T.; Mukai, T.; Ohkubo, T.; Sakai, H.; Abe, M. *J. Am. Chem. Soc.* 2005, 127, 16396-16397.
- [109] Shibata, H.; Mihara, H.; Mlikai, T.; Ogura, T.; Kohno, H.; Ohkubo, T.; Sakait, H.; Abe, M. *Chem. Mater.* 2006, 18, 2256-2260.
- [110] Tian, B. Z.; Liu, X. Y.; Yang, H. F.; Xie, S. H.; Yu, C. Z.; Tu, B.; Zhao, D. Y. *Adv. Mater.* 2003, 15, 1370-+.
- [111] Ryoo, R.; Joo, S. H.; Kruk, M.; Jaroniec, M. *Adv. Mater.* 2001, 13, 677-681.
- [112] Masuda, Y.; Kato, K. Japan, 2007.
- [113] Masuda, Y.; Sugiyama, T.; Seo, W. S.; Koumoto, K. *Chem. Mater.* 2003, 15, 2469-2476.
- [114] Masuda, Y.; Kato, K. *Chem. Mater.* 2008, 20, 1057-1063.
- [115] Furlong, D. N.; Parfitt, G. D. *J. Colloid Interface Sci.* 1978, 65, 548-554.
- [116] Wahi, R. K.; Liu, Y. P.; Falkner, J. C.; Colvin, V. L. *J. Colloid Interface Sci.* 2006, 302, 530-536.
- [117] Kruk, M.; Jaroniec, M. *Chem. Mater.* 2001, 13, 3169-3183.
- [118] Ravikovitch, P. I.; Odomhnaill, S. C.; Neimark, A. V.; Schuth, F.; Unger, K. K. *Langmuir* 1995, 11, 4765-4772.
- [119] Lastoskie, C.; Gubbins, K. E.; Quirke, N. *J. Phys. Chem.* 1993, 97, 4786-4796.
- [120] Katagiri, K.; Ohno, K.; Masuda, Y.; Koumoto, K. *J. Ceram. Soc. Japan* 2007, 115, 831-834.
- [121] Masuda, Y.; Ohji, T.; Kato, K. *Crystal Growth & Design* 2010, 10, 913-922.
- [122] Masuda, Y.; Kato, K. *Thin Solid Films* 2008, 516, 2547-2552.
- [123] Masuda, Y.; Kato, K. *J. Cryst. Growth* 2009, 311, 512-517.
- [124] Ohsaka, T.; Izumi, F.; Fujiki, Y. *J. Raman Spectrosc.* 1978, 7, 321-324.
- [125] Gao, K. *Physica Status Solidi B-Basic Solid State Physics* 2007, 244, 2597-2604.
- [126] Cong, Y.; Zhang, J.; Chen, F.; Anpo, M. *J. Phys. Chem. C* 2007, 111, 6976-6982.
- [127] Ohsaka, T.; Yamaoka, S.; Shimomura *Solid State Communications* 1979, 30, 345-347.
- [128] Yu, J. G.; Yu, H. G.; Cheng, B.; Zhao, X. J.; Yu, J. C.; Ho, W. K. *Journal of Physical Chemistry B* 2003, 107, 13871-13879.
- [129] Ignat'eva, L. N.; Polishchuk, S. A.; Antokhina, T. F.; Buznik, V. T. *Glass Physics and Chemistry* 2004, 30, 139-141.
- [130] Nakamoto, K. *Infrared Spectra of Inorganic and Coordination Compounds*; Wiley: New York, 1970.
- [131] Padmanabhan, S. C.; Pillai, S. C.; Colreavy, J.; Balakrishnan, S.; McCormack, D. E.; Perova, T. S.; Gun'ko, Y.; Hinder, S. J.; Kelly, J. M. *Chemistry of Materials* 2007, 19, 4474-4481.
- [132] Wu, J. J.; Yu, C. C. *Journal of Physical Chemistry B* 2004, 108, 3377-3379.
- [133] Perathoner, S.; Passalacqua, R.; Centi, G.; S. Su, D.; Weinberg, G. *Catalysis Today* 2007, 122, 3-13.
- [134] Sato, H.; Ono, K.; Sasaki, T.; Yamagishi, A. *Journal of Physical Chemistry B* 2003, 107, 9824-9828.

- [135] Naik, V. M.; Haddad, D.; Naik, R.; Benci, J.; Auner, G. W. In *Symposium on Solid-State Chemistry of Inorganic Materials IV held at the 2002 MRS Fall Meeting*; AlarioFranco, M. A., Greenblatt, M., Rohrer, G., Whittingham, M. S., Eds.; Materials Research Society: Boston, Ma, 2002, p 413-418.
- [136] Masuda, Y.; Sugiyama, T.; Lin, H.; Seo, W. S.; Koumoto, K. *Thin Solid Films* 2001, 382, 153-157.
- [137] Huang, D.; Xiao, Z.-D.; Gu, J.-H.; Huang, N.-P.; Yuan, C.-W. *Thin Solid Films* 1997, 305, 110-115.
- [138] Zhang, F.; Mao, Y.; Zheng, Z.; Chen, Y.; Liu, X.; Jin, S. *Thin Solid Films* 1997, 310, 29-33.
- [139] Li, D.; Haneda, H.; Hishita, S.; Ohashi, N.; Labhsetwar, N. K. *Journal of Fluorine Chemistry* 2005, 126, 69-77.
- [140] Wang, C. M.; Mallouk, T. E. *Journal of Physical Chemistry* 1990, 94, 423-428.
- [141] Minero, C.; Mariella, G.; Maurino, V.; Pelizzetti, E. *Langmuir* 2000, 16, 2632-2641.
- [142] Minero, C.; Mariella, G.; Maurino, V.; Vione, D.; Pelizzetti, E. *Langmuir* 2000, 16, 8964-8972.
- [143] Vohra, M. S.; Kim, S.; Choi, W. In *International Symposium on Photochemistry at Interfaces*; Elsevier Science Sa: Sapporo, Japan, 2002, p 55-60.
- [144] Fu, H. B.; Zhang, L. W.; Zhang, S. C.; Zhu, Y. F.; Zhao, J. C. *Journal of Physical Chemistry B* 2006, 110, 3061-3065.
- [145] Irie, H.; Watanabe, Y.; Hashimoto, K. *Journal of Physical Chemistry B* 2003, 107, 5483-5486.
- [146] Masuda, Y.; Kato, K. *J. Cryst. Growth* 2009, 311, 593-596.
- [147] Ginley, D. S.; Bright, C. *MRS Bull.* 2000, 25, 15-21.
- [148] Idota, Y.; Kubota, T.; Matsufuji, A.; Maekawa, Y.; Miyasaka, T. *Science* 1997, 276, 1395 - 1397.
- [149] Zhang, Y. L.; Liu, Y.; Liu, M. L. *Chem. Mater.* 2006, 18, 4643-4646.
- [150] Wang, Y.; Lee, J. Y.; Zeng, H. C. *Chem. Mater.* 2005, 17, 3899-3903.
- [151] Bose, A. C.; Kalpana, D.; Thangadurai, P.; Ramasamy, S. *Journal of Power Sources* 2002, 107, 138-141.
- [152] Pianaro, S. A.; Bueno, P. R.; Longo, E.; Varela, J., A. J. *Mat. Sci. Lett.* 1995, 14, 692-694.
- [153] Bueno, P. R.; de Cassia-Santos, M. R.; Leite, E. R.; Longo, E.; Bisquert, J.; Garcia-Belmonte, G.; Fabregat-Santiago, F. *Journal of Applied Physics* 2000, 88, 6545-6548.
- [154] Tagawa, T.; Kataoka, S.; Hattori, T.; Murakami, Y. *Applied Catalysis* 1982, 4, 1-4.
- [155] Park, P. W.; Kung, H. H.; Kim, D. W.; Kung, M. C. *Journal of Catalysis* 1999, 184, 440-454.
- [156] Chopra, K. L.; Major, S.; Pandya, D. K. *Thin Solid Films* 1983, 102, 1-46.
- [157] Sergent, N.; Gelin, P.; Perier-Camby, L.; Praliaud, H.; Thomas, G. *Sensors and Actuators B-Chemical* 2002, 84, 176-188.
- [158] Baik, N. S.; Sakai, G.; Miura, N.; Yamazoe, N. *J. Am. Ceram. Soc.* 2000, 83, 2983-2987.
- [159] Ristic, M.; Ivanda, M.; Popovic, S.; Music, S. *Journal of Non-Crystalline Solids* 2002, 303, 270-280.

- [160] Broussous, L.; Santilli, C. V.; Pulcinelli, S. H.; Craievich, A. F. *Journal of Physical Chemistry B* 2002, 106, 2855-2860.
- [161] Zhang, J. R.; Gao, L. *Journal of Solid State Chemistry* 2004, 177, 1425-1430.
- [162] Deng, Z. X.; Wang, C.; Li, Y. D. *Journal of The American Ceramic Society* 2002, 85, 2837-2839.
- [163] Leite, E. R.; Gomes, J. W.; Oliveira, M. M.; Lee, E. J. H.; Longo, E.; Varela, J. A.; Paskocimas, C. A.; Boschi, T. M.; Lanciotti, F.; Pizani, P. S.; Soares, P. C. *Journal of Nanoscience and Nanotechnology* 2002, 2, 125-128.
- [164] Leite, E. R.; Maciel, A. P.; Weber, I. T.; Lisboa, P. N.; Longo, E.; Paiva-Santos, C. O.; Andrade, A. V. C.; Pakoscimas, C. A.; Maniette, Y.; Schreiner, W. H. *Advanced Materials* 2002, 14, 905-908.
- [165] Dai, Z. R.; Gole, J. L.; Stout, J. D.; Wang, Z. L. *J. Phys. Chem. B* 2002, 106, 1274 -1279.
- [166] Liu, Y.; Liu, M. L. *Adv. Funct. Mater.* 2005, 15, 57 - 62.
- [167] Huang, J.; Matsunaga, N.; Shimano, K.; Yamazoe, N.; Kunitake, T. *Chem. Mater.* 2005, 17, 3513-3518.
- [168] Liu, Y. K.; Zheng, C. L.; Wang, W. Z.; Yin, C. R.; Wang, G. H. *Adv. Mater.* 2001, 13, 1883 - 1887.
- [169] Sun, J. Q.; Wang, J. S.; Wu, X. C.; Zhang, G. S.; Wei, J. Y.; Zhang, S. Q.; Li, H.; Chen, D. R. *Cryst. Growth Des.* 2006, 6, 1584-1587.
- [170] Yang, R. S.; Wang, Z. L. *J. Am. Chem. Soc.* 2006, 128, 1466-1467.
- [171] Duan, J. H.; Yang, S. G.; Liu, H. W.; Gong, J. F.; Huang, H. B.; Zhao, X. N.; Zhang, R.; Du, Y. W. *J. Am. Chem. Soc.* 2005, 127, 6180 -6181.
- [172] Ohgi, H.; Maeda, T.; Hosono, E.; Fujihara, S.; Imai, H. *Cryst. Growth Des.* 2005, 5, 1079-1083.
- [173] Uchiyama, H.; Imai, H. *Cryst. Growth Des.* 2007, 7, 841-843.
- [174] Niesz, D. E.; Bennett, R. B.; Snyder, M. J. *Am. Ceram. Soc. Bull.* 1972, 51, 677-680.
- [175] Roosen, A.; Hausener, H. *Adv. Ceram. Mater.* 1988, 3, 131.
- [176] Vasylyuk, O.; Sakka, Y. *Journal of The American Ceramic Society* 2001, 84, 2489-2494.
- [177] Niesen, T. P.; DeGuire, M. R. *J. Electroceramics* 2001, 6, 169-207.
- [178] Baes, C. F.; Mesiner, R. E. *The Hydrolysis of Cations*; John Wiley & Sons, Inc., Wiley-Interscience: New York, 1976.
- [179] Ararat Ibarguena, C.; Mosquera, A.; Parrab, R.; Castro, M. S.; Rodríguez-Páez, J. E. *Mater. Chem. Phys.* 2007, 101, 433-440.
- [180] Masuda, Y.; Ohji, T.; Kato, K. *Journal of The American Ceramic Society* 2010, 93, 2140-2143.
- [181] Masuda, Y.; Kato, K. *Journal of Crystal Growth* 2009, 311, 593-596.
- [182] Masuda, Y.; Kato, K. *Polymers for Advanced Technologies* 2010, 21, 211-215.
- [183] Shukla, S.; Seal, S.; Ludwig, L.; Parish, C. *Sens. Actuators B: Chem.* 2004, 97, 256-265.
- [184] Masuda, Y.; Kato, K.; Sonezaki, S.; Ajimi, M.; Bekki, M. *Japan, 2008, p P 2008-227389.*
- [185] Xiang, J. H.; Masuda, Y.; Koumoto, K. *Adv. Mater.* 2004, 16, 1461-1464.
- [186] Xiang, J. H.; Zhu, P. X.; Masuda, Y.; Koumoto, K. *Langmuir* 2004, 20, 3278-3283.
- [187] Masuda, Y.; Yamagishi, M.; Koumoto, K. *Chem. Mater.* 2007, 19, 1002-1008.
- [188] Masuda, Y.; Wakamatsu, S.; Koumoto, K. *J. Euro. Ceram. Soc.* 2004, 24, 301-307.

- [189] Themlin, J. M.; Chtaib, M.; Henrard, L.; Lambin, P.; Darville, J.; Gilles, J. M. *Phys. Rev. B* 1992 46, 2460-2466.
- [190] Yan, L.; Pan, J. S.; Ong, C. K. *Mater. Sci. Eng. B* 2006, 128, 34-36.
- [191] Wagner, C. D. *Practical Surface Analysis*; 2 ed.; John Wiley, 1990; Vol. 1.

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We face many challenges in the 21st century, such as sustainably meeting the world's growing demand for energy and consumer goods. I believe that new developments in science and technology will help solve many of these problems. Nanofabrication is one of the keys to the development of novel materials, devices and systems. Precise control of nanomaterials, nanostructures, nanodevices and their performances is essential for future innovations in technology. The book "Nanofabrication" provides the latest research developments in nanofabrication of organic and inorganic materials, biomaterials and hybrid materials. I hope that "Nanofabrication" will contribute to creating a brighter future for the next generation.

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