<table>
<thead>
<tr>
<th>Title</th>
<th>Photocatalytic activity of amorphous−anatase mixture of titanium(IV) oxide particles suspended in aqueous solutions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Author(s)</td>
<td>Ohtani, Bunsho; Ogawa, Yoshimasa; Nishimoto, Seiichi</td>
</tr>
<tr>
<td>Citation</td>
<td>Journal of physical chemistry. B, 101(19): 3746-3752</td>
</tr>
<tr>
<td>Issue Date</td>
<td>1997-05-08</td>
</tr>
<tr>
<td>Doc URL</td>
<td><a href="http://hdl.handle.net/2115/52791">http://hdl.handle.net/2115/52791</a></td>
</tr>
<tr>
<td>Type</td>
<td>article</td>
</tr>
<tr>
<td>File Information</td>
<td>jp962702+.pdf</td>
</tr>
</tbody>
</table>

Hokkaido University Collection of Scholarly and Academic Papers: HUSCAP
Photocatalytic Activity of Amorphous–Anatase Mixture of Titanium(IV) Oxide Particles Suspended in Aqueous Solutions

Bunsho Ohtani,* Yoshimasa Ogawa, and Sei-ichi Nishimoto
Department of Energy and Hydrocarbon Chemistry, Graduate School of Engineering, Kyoto University, Kyoto 606-01, Japan

Received: August 30, 1996; In Final Form: January 30, 1997

Titanium(IV) oxide (TiO<sub>2</sub>) powders of various amorphous–anatase compositions were prepared by heat treatment (573–1073 K) of amorphous TiO<sub>2</sub> in air and characterized by differential scanning calorimetry (DSC), powder X-ray diffraction (XRD), and BET surface area measurements. An exothermic peak at ca. 723 K in the DSC pattern was assigned to the crystallization of amorphous phase to anatase, and its heat was used to evaluate the weight fraction of amorphous phase. The fraction of anatase crystallinity (f(anatase)) was calculated as the remainder after the amorphous phase and contaminated water or organic residue. The XRD data showed that the size of anatase crystallites was slightly decreased but almost constant along with the increase in f(anatase), being consistent with the small change in the BET surface area. These results suggest that each amorphous particle transforms into an anatase crystal of similar size without sintering or crystal growth. The particles of mixture of amorphous and anatase were used for three kinds of photocatalytic reactions in deaerated aqueous suspensions: silver metal deposition with oxidation of 2-propanol by bare powders, and dehydrogenation of 2-propanol by ex situ and in situ platinized powders. For all the reactions, the photocatalytic activity of amorphous TiO<sub>2</sub> was negligible, increased almost linearly with f(anatase), and further improved by calcination of completely crystallized powder. The negligible activity of amorphous TiO<sub>2</sub> was attributed to recombination of photoexcited electron and positive hole at defects located on the surface and in the bulk of particles.

I. Introduction

Semiconductor photocatalytic reaction has been investigated extensively, with applications focusing on solar energy conversion and storage, reductive fixation of carbon dioxide, or organic synthesis. Among semiconductors, titanium(IV) oxide (TiO<sub>2</sub>) is one of the most popular and promising materials as a photocatalyst because it is stable in various solvents under photoirradiation, available commercially, and easy to prepare in the laboratory and has potential ability to induce various types of redox reactions. However, it has been also clarified that its photocatalytic activity strongly depends on its physical properties; a certain TiO<sub>2</sub> powder showed no appreciable activity but another powder was quite active. There have been many articles reporting the correlation of photocatalytic activity with the physical properties of TiO<sub>2</sub> powders, such as crystal structure, surface area, particle size, surface hydroxyls, and so on. It is believed by most researchers in this field that the crystal structure (anatase, rutile, or brookite) is the most primitive and essential property to predict the photocatalytic activity, e.g., anatase, but not rutile, crystallites have relatively high photocatalytic activity. On the other hand, few papers have revealed the photocatalytic activity of amorphous TiO<sub>2</sub>: to the best of our knowledge, only one paper reported the negligible activity of amorphous TiO<sub>2</sub> (same as that used in this study) by comparison with commercial crystalline TiO<sub>2</sub> catalysts, but no interpretation for such behavior was given.

Although a few reports described the photocatalytic activity of less (or negligibly) crystallized TiO<sub>2</sub> particles, as amorphous TiO<sub>2</sub>, prepared by hydrolytic precipitation from titanium(IV) alkoxides or halides, the particles may consist of, presumably, titanium(IV) hydroxide (Ti(OH)<sub>2</sub>) or oxyhydroxide (TiO(OH)<sub>2</sub>) or, in another expression, hydrated TiO<sub>2</sub> (TiO<sub>2</sub>·2H<sub>2</sub>O and TiO<sub>2</sub>·H<sub>2</sub>O).

In a series of studies on TiO<sub>2</sub> photocatalytic reactions, we have shown that the overall kinetics depend on both amount of substrate(s) adsorbed on the surface to be reduced or oxidized by photoexcited electron (e<sup>-</sup>) or positive hole (h<sup>+</sup>), respectively, and rate of geminate recombination of e<sup>-</sup> and h<sup>+</sup>. To obtain highly active TiO<sub>2</sub> photocatalyst, therefore, it is necessary to give simultaneously a couple of properties, generally of contradiction, to the powders; large surface area to adsorb substrates and high crystallinity (or lesser surface and bulk defects) to diminish the e<sup>-</sup>−h<sup>+</sup> recombination. This hypothesis has been supported by the ultra-highly active TiO<sub>2</sub> photocatalyst prepared by a newly developed method, hydrothermal crystallization in organic media (HyCOM) and thermal decomposition of titanium(IV) alkoxide (TD). The design and development of polycrystalline TiO<sub>2</sub> photocatalysts are in progress along this line. Our research interest, then, focused on photocatalytic activity of amorphous, not crystalline, semiconductor particles. In this paper, the results of investigation on mixture of amorphous and anatase TiO<sub>2</sub> powders are described and discussed.

II. Experimental Section

Materials. Amorphous TiO<sub>2</sub> (type UF-IT-S) was supplied from Idemitsu Kosan Co., Ltd., as a flocculent white powder. A 5–10 g portion of this powder was calcined in air at various temperature (573–1073 K) and duration (1–3 h) in an electric furnace equipped with a time-programmed temperature controller. A commercial anatase TiO<sub>2</sub> sample was supplied from Merck and used without pretreatment. Ex situ loading of platinum (Pt; 2 wt %) on the TiO<sub>2</sub> powders was achieved as...
follows. To an aqueous suspension of the powder was added dropwise an aqueous solution of chloroplatinic acid; the mixture was stirred thoroughly and then evaporated to dryness. The resulting pale yellow powder was heat-treated under hydrogen (H₂) stream at 453 K for 4 h in an electric furnace to obtain gray or grayish white powder of TiO₂·Pt.

The other chemicals used in this study were purchased from Wako Pure Chemical or Nacarai Tesque in the highest available grades. Ion-exchanged water prepared with a Corning Mega-Pure System MP-190 (>16 MΩ cm) was used to make the aqueous solutions throughout this study.

**Characterization of TiO₂ Powders.** Differential scanning calorimetry was performed with a Shimadzu DSC-50 calorimeter under nitrogen (N₂) atmosphere. An ca. 2 mg portion of sample powder was packed in an aluminum cell and set in the calorimeter. The calorimeter measured the heat flux from the sample with a unit of mJ s⁻¹ mg⁻¹. The rate of heating was 20 K s⁻¹. Therefore, the vertical axis of the results shown in Figure 1 had a dimension of mJ mg⁻¹ K⁻¹ and the heat of crystallization could be calculated from the peak area in the unit of mJ mg⁻¹. X-ray diffraction (XRD) pattern was recorded on a Rigaku Geigerflex 2013 diffractometer (Cu Kα, Ni filter, 35 kV, 20 mA) at a scanning rate of 1 deg min⁻¹. To prepare the XRD calibration curve (Figure 3), a commercial anatase TiO₂ (Merck) was mixed with the original amorphous TiO₂ with different weight ratio and CaCO₃ corresponding to 20 wt % of the mixture was added. The details of XRD measurement were reported previously.²⁰ The specific surface area of TiO₂ samples was determined from N₂ adsorption at 77 K on the basis of the BET equation by using a glass apparatus for constant-volume adsorption measurement equipped with an oil diffusion pump and an MKS Baratron type 122A capacitance manometer. To minimize the change in physical properties of TiO₂ samples, a drying process prior to this surface-area measurement was operated at 383 K under reduced pressure. Reflection spectra of powders in ultraviolet and visible region were recorded on a Hitachi U-4000 spectrophotometer. The powder samples are loaded in a cell equipped with a quartz window and reflectance was measured with a reference spectrum with barium sulfate. Reflectance (R/R₀) was normalized at 500 nm to be 1.

**Photocatalytic Activity of Amorphous—Anatase Mixture of TiO₂**

**Figure 1.** Representative DSC pattern of amorphous—anatase mixture of TiO₂ powders: (a) as-received amorphous, (b) mixture (fanatase) ca. 0.5% prepared from a by heat treatment (573 K, 2 h), and (c) remeasuring of a. Patterns were translated for clarity along the vertical axis.

was suspended in an aqueous 2-propanol solution and irradiated without H₂PtCl₆ under Ar at 298 K for 3 h. To the photoirradiated reaction mixture was added a deaerated aqueous solution of MV²⁺ (0.5 mmol, 0.5 cm³). The resulting pale blue suspension containing reduced form of MV²⁺, MV⁺, was filtered and the supernatant solution was loaded in a sealable quartz cell (light path length 1 mm) in an N₂-purged vacuum glovebox. From the absorbance of solution at 602 nm (a Shimadzu MP-1000 spectrophotometer) and reported extinction coefficient of MV⁺ (13 700 mol⁻¹ dm⁻³ cm⁻¹ at 602 nm), the molar amount of Ti³⁺ was estimated on the assumption that equimolar amounts of Ti³⁺ and MV²⁺ react to produce MV⁺.

**III. Results and Discussion**

**Characterization of Amorphous—Anatase Mixture.** Figure 1 is a representative DSC pattern of TiO₂ samples. A characteristic exothermic peak was observed at ca. 723 K (curve a). An endothermic peak below ca. 550 K was attributed to liberation or decomposition of water and/or organic residue, since it disappeared in the pattern of heat-treated (573 K, 2 h) sample (curve b). The as-received powder (giving no XRD peak of anatase crystallite, as described later) gave almost constant heat at ca. 723 K in several repeated measurements. This exothermic peak is attributable to crystallization of amorphous TiO₂ to anatase crystals since similar exothermic peak at 743 K has been also observed in differential thermal analysis under atmospheric conditions.⁴⁶ The fact that decrease in the sample weight was seen at elevating temperature up to 573 K and thereafter it negligibly changed suggests that the exothermic peak is independent of the dehydration or decomposition of organic residue which must be accompanied by the change in weight. The heat of the crystallization was determined from the average of peak area in the several measurements to be 217 J g⁻¹ (4.1 kcal mol⁻¹), which is in the same order of magnitude of transformation of anatase into rutile,⁴⁷ 392 J g⁻¹ (7.5 kcal mol⁻¹), or amorphous germanium to its crystal,⁴⁸ 158–178 J g⁻¹ (2.8–3.1 kcal mol⁻¹). This heat of crystallization was used as a standard to determine the weight fraction of amorphous TiO₂ (f(amorphous)), and that of anatase (f(anatase)) was estimated by an equation

\[
f(\text{anatase}) = 1 - f(\text{amorphous}) - f(\text{water})\]  

where f(water) is weight fraction of water or organic residue in the sample and evaluated from the decrease in weight after DSC measurements. Actually, f(water) was smaller than 0.04 for heat-treated samples and 0.08 for the as-received amorphous powder.
Along with the decrease in amorphous fraction in the TiO₂ samples, the XRD peak at 25.4° assigned to anatase crystallite, increased (Figure 2). Before the calcination, no characteristic peak could be seen, showing that the as-received powder consisted of amorphous TiO₂. Under the present calcination conditions (temperature < 1073 K), formation of rutile or brookite crystallites could not be observed. The content of anatase crystallites could be also evaluated from the ratio of peak area between the peaks of anatase and an internal standard, calcium carbonate (CaCO₃; Wako, 20 wt %). A calibration peak area between the peaks of anatase and an internal standard, anatase crystallites could be also evaluated from the ratio of brookite crystallites could not be observed. The content of anatase crystallites could be also evaluated from the ratio of peak area between the peaks of anatase and an internal standard, calcium carbonate (CaCO₃; Wako, 20 wt %). A calibration line was drawn with the mixture of a commercial anatase TiO₂ (Merck) and CaCO₃; calculated weight fraction of anatase (anatase) evaluated by two methods, DSC and XRD

\[ f(\text{anatase}) = 1.063 R(\text{anatase}/\text{CaCO}_3) - 0.091 \]  

In the previous report, similar linear correlation was obtained for the mixture of anatase and rutile. However, by assuming a linear relation between the XRD peak intensity and fraction of given crystallites, the intercept should be zero. The discrepancy might be caused by the marked difference in the size of anatase and amorphous standard samples, though we could not obtain an anatase standard sample (without amorphous or hydrated TiO₂) of crystallite size similar to amorphous TiO₂ to confirm this. Anyway, practically the fraction of anatase in the samples could be evaluated from the XRD measurements. As shown in Figure 4, \( f(\text{anatase}) \) evaluated by two methods, DSC and XRD

\begin{align*}
\text{Figure 2. Representative XRD pattern of amorphous-anatase mixture of TiO}_2\text{ powders: (a) as-received amorphous, (b) mixture (f(anatase)} \\
c\text{ca. 0.3), and (c) completely crystallized anatase. A broad peak at 2} \\
\theta = 19° \text{ and small peaks at 21}° \text{ and 23.5}° \text{ are caused by vaseline used to} \\
\text{fix the powders on an XRD sample holder. Sample b contained 20 wt} \\
\% \text{ of CaCO}_3 \text{ as an internal standard.}
\end{align*}
Photocatalytic Activity of Amorphous—Anatase Mixture of TiO₂

smaller along with the decrease in crystallite size. These facts teach us the difficulty of measuring content of anatase crystallites of size smaller than 30 nm from XRD if the calibration line is drawn with larger crystallites of standard anatase sample (in this case the size of anatase crystallites of standard was >200 nm). This problem is clarified first, to our knowledge, in this study. The absorption edge of amorphous TiO₂ particles of almost the same size (anatase) range from 0 to 0.98. This indicates that both amorphous and anatase crystal–mesostructures. Under the acidic conditions, where surface hydroxyls are protonated to be positively charged, cations such as Ag⁺⁺ are inhibited to be adsorbed and therefore they do not participate in the reaction system A (Figure 8 upper), the molar amount of deposited Ag was approximately twice that of acetone as an oxidation product from 2-propanol, satisfying a stoichiometry of

\[ 2\text{Ag}^+ + (\text{CH}_3)_2\text{CHOH} = 2\text{Ag} + (\text{CH}_3)_2\text{CO} + 2\text{H}^+ \quad (3) \]

In the previous paper, we have shown that under such reaction conditions of low pH, the rate of the photocatalytic reaction is governed by the reaction of a photogenerated positive hole (h⁺) with 2-propanol, i.e., the overall rate (r) is proportional to the amount of surface-adsorbed 2-propanol, but not Ag⁺. The following rate expression could be reasonably fit to the dependence of rate (r) on the 2-propanol concentration ([2-propanol]ₐd).

\[ r = Iφk_a[2\text{-propanol}]ₐd/k_r \quad (4) \]

where I and φ are flux of photons absorbed by the photocatalyst and efficiency of production of e⁻ and h⁺, and \( k_a \) and \( k_r \) are rate constants of reaction by h⁺ and recombination of e⁻ and h⁺, respectively. This equation is based on the idea that in a primary stage of photocatalytic reaction the photogenerated e⁻−h⁺ pairs undergo recombination to give appreciable chemical reaction unless they react with the surface-adsorbed substrates. Under the acidic conditions, where surface hydroxyls are protonated to be positively charged, cations such as Ag⁺⁺ are inhibited to be adsorbed and therefore they do not participate

Photocatalytic Activities of the Amorphous—Anatase Mixture. Figure 8 shows the rate of photocatalytic reaction in three different systems evaluated from the slope of the almost linear time-course curves. In all the systems, the rate increased linearly with f(anatase) up to unity and markedly enhanced at f(anatase) = 1. This fact suggests that only anatase crystallites, but not amorphous, induce the photocatalytic reactions and that the activity of anatase crystallites is improved further by the heat treatment. The latter finding will be discussed later.

In the reaction system A (Figure 8 upper), the molar amount of deposited Ag was approximately twice that of acetone as an oxidation product from 2-propanol, satisfying a stoichiometry of

\[ 2\text{Ag}^+ + (\text{CH}_3)_2\text{CHOH} = 2\text{Ag} + (\text{CH}_3)_2\text{CO} + 2\text{H}^+ \quad (3) \]

In the previous paper, we have shown that under such reaction conditions of low pH, the rate of the photocatalytic reaction is governed by the reaction of a photogenerated positive hole (h⁺) with 2-propanol, i.e., the overall rate (r) is proportional to the amount of surface-adsorbed 2-propanol, but not Ag⁺. The following rate expression could be reasonably fit to the dependence of rate (r) on the 2-propanol concentration ([2-propanol]ₐd).

\[ r = Iφk_a[2\text{-propanol}]ₐd/k_r \quad (4) \]

where I and φ are flux of photons absorbed by the photocatalyst and efficiency of production of e⁻ and h⁺, and \( k_a \) and \( k_r \) are rate constants of reaction by h⁺ and recombination of e⁻ and h⁺, respectively. This equation is based on the idea that in a primary stage of photocatalytic reaction the photogenerated e⁻−h⁺ pairs undergo recombination to give appreciable chemical reaction unless they react with the surface-adsorbed substrates. Under the acidic conditions, where surface hydroxyls are protonated to be positively charged, cations such as Ag⁺⁺ are inhibited to be adsorbed and therefore they do not participate

Photocatalytic Activities of the Amorphous—Anatase Mixture. Figure 8 shows the rate of photocatalytic reaction in three different systems evaluated from the slope of the almost linear time-course curves. In all the systems, the rate increased

Photocatalytic Activities of the Amorphous—Anatase Mixture. Figure 8 shows the rate of photocatalytic reaction in three different systems evaluated from the slope of the almost linear time-course curves. In all the systems, the rate increased
in the primary stage. The equation could reasonably imply the experimental results for polycrystalline TiO$_2$ photocatalysts. Assuming that $k_0$ is constant for the present amorphous–anatase samples (it seems to be reasonable to see their similar reflection spectra shown in Figure 7) and [2-propanol]$_{ad}$ is proportional to the BET surface area, we can attribute the rate dependence in Figure 8 predominantly to the difference in the ratio of $S_{BET}$ $k_h/k_r$ between amorphous and anatase particles (therefore, we would regard the ratio, estimated from the ratio of practical reaction rate to the surface area ($rS_{BET}$) which was sometimes used in the studies of photocatalysis, as an intrinsic photocatalytic activity). Since, in the region of $f$(anatase)$<1$, the BET surface area was almost constant or rather a little decreased, the increasing rate with $f$(anatase) should be attributed to increasing ratio of $k_h/k_r$. The fact that the rate was negligible for amorphous TiO$_2$ ($f$(anatase) = 0) suggests the negligible $k_h/k_r$ ratio for amorphous particles.

Similar rate dependence was seen for the reaction systems B and C in which dehydrogenation of 2-propanol proceeds to yield equimolar amount of H$_2$ and acetone (acetone was liberated in reaction C, but was not measured qualitatively).

$$(\text{CH}_3)_2\text{CHOH} = (\text{CH}_3)_2\text{CO} + \text{H}_2 \quad (5)$$

It has been also clarified that the rate of this 2-propanol dehydrogenation is expressed by an equation similar to eq 4 for the platinized commercial TiO$_2$ powders. Hence, similar interpretation that amorphous TiO$_2$ is inactive also for the 2-propanol dehydrogenation may be plausible. It should be noted that the slope of linear relation for system B (Figure 8, middle) was almost equal to that for system A (Figure 8, upper); i.e., the limiting rate at $f$(anatase) = 1 was almost the same in these systems. This is consistent with eq 4 which does not include the terms relating to reduction by $e^-$. Under the similar reaction conditions of same pH, [2-propanol]$_{ad}$ should be the same and the common reaction of 2-propanol with h$^+$ should predominate the overall reaction rates regardless of the different reduction processes.

The loaded platinum deposit acts as a site to reduce proton (or water) into H$_2$. It has been proved that for the photocatalytic 2-propanol dehydrogenation by small TiO$_2$ particles (of a few dozens of nm) at least one Pt deposit makes each TiO$_2$ particle photoactive and the activity is constant or rather a little diminished by further loadings; i.e., the rate is almost constant regardless of the number of Pt deposits on particles so long as they have at least one Pt deposit. On the basis of these results, the photocatalytic activity of different kind of TiO$_2$ powders should be compared with consideration of platinization, i.e., number of Pt deposits on each TiO$_2$ particles. The above-mentioned fact that the rate of 2-propanol oxidation was almost the same regardless of the different reduction processes suggests that all the TiO$_2$ (anatase) particles work as a photocatalyst; the impregnation followed by $H_2$ reduction may produce Pt deposits which is enough for uniform platinization of the TiO$_2$ particles.

On the other hand, the rate for system C was ca. 3 times faster than that of system B though the same 2-propanol dehydrogenation proceeded, maybe due to the difference in [2-propanol]$_{ad}$ in eq 4 since system C was neutral while system B was acidic. The actual [2-propanol]$_{ad}$ could not be measured because the change in 2-propanol concentration by the adsorption in the dark was below the detectable limit. Anyway, the rate dependences on $f$(anatase) in systems B and C are similar, showing that the different method of platinization gives no influence on the tendency of photocatalytic activity and, thereby, the rate shown in Figure 8 reflects the intrinsic photocatalytic activity of TiO$_2$ particles.

### Negligible Photocatalytic Activity of Amorphous TiO$_2$

As described above, the present results are reasonably interpreted by assuming that the amorphous particle exhibits negligible photocatalytic activity for the reactions where 2-propanol is oxidized by h$^+$ and Ag$^+$ or H$^+$ is reduced by e$^-$. Judging from the rate expression (eq 4), the negligible activity is accounted for by the negligible ratio of rate constants, $k_h/k_r$. We have, at present, no evidence to distinguish which rate constant governs the activity. However, we consider that large $k_r$, i.e., facilitated e$^-$–h$^+$ recombination in amorphous TiO$_2$, cancels the photocatalytic activity, since it is probable that $k_h$ reflects the structure of surface and/or adsorbed 2-propanol while $k_r$ depends strongly on the bulk structure; the difference in effect of bulk structure to $k_r$ between amorphous and anatase particles might be more extensive than that of surface structure to $k_h$. Amorphous TiO$_2$ must contain many imperfections, i.e., impurities, dangling bonds, or microvoids, which lead to electronic states in the band gap, and they behave as a recombination center for e$^-$–h$^+$. On the other hand, the surface structure depends on a short-range order of bulk structure, which should be not so different between anatase and amorphous TiO$_2$ particles; at least we could not expect the significant $k_h$ difference which makes the $k_h/k_r$ ratio to be negligible.

Bickle and co-workers have reported that a commercial TiO$_2$ (Degussa P-25) is a mixture of anatase crystallites with amorphous and rutile phases mainly on the basis of microscopic investigations. They attributed the unusually high photocatalytic activity of the TiO$_2$ particles to their complex structure and suggested that localized Anderson states of amorphous TiO$_2$ may increase the lifetimes of e$^-–h^+$ and thereby enhance the photocatalytic activity. This is in contrast with the present results, but the discrepancy may arise from the difference in the structure of TiO$_2$ particles; the P-25 powder comprised coagulated anatase–amorphous and anatase–rutile particles, while in the present study the samples presumably consist of independent anatase and amorphous particles, as described in the preceding section.

Figure 9 shows the molar amount of Ti$^{3+}$ formed during the photoirradiation in the presence of 2-propanol but in the absence of loaded Pt deposits. Since neither appropriate electron acceptor nor catalyst (such as Pt for H$_2$ evolution) is present in the reaction system, TiO$_2$ itself is reduced by e$^-$; the suspension turned gray or blue gray by the photoirradiation. This Ti$^{3+}$ formation is caused by the trapping of e$^-$ at the defects along with the oxidation of 2-propanol with h$^+$ and its amount seems to be an indication of oxygen anion (O$^{2-}$) vacancy or O$^{2-}$ weakly bonded to Ti$^{4+}$. Since both Ti$^{3+}$ and MV$^{+}$ are highly reactive toward O$_2$ in the atmosphere and are oxidized quickly,
i.e., they are readily quenched by small amount of O₂ contaminant during the measurements, large experimental error should be included in the results shown in Figure 9. However, a tendency is seen that appreciable amount of Ti³⁺ was formed in partly crystallized samples (f(anatase) < 1) but was negligible when completely crystallized (f(anatase) = 1). It seems that the Ti³⁺ amount is roughly proportional to the BET surface area depicted in Figure 5, and therefore Ti³⁺ species might be located on the surface of both amorphous and anatase particles; i.e., the Ti³⁺ amount should be a measure of the common e⁻–h⁺ recombination site on the surface. The fact that even the TiO₂ particles consisting of predominantly anatase (high f(anatase) but not unity) which showed relatively high activity gave appreciable amount of Ti³⁺ suggests that not only the surface defect, i.e., recombination center, detected as Ti³⁺, but also bulk defect(s) accelerates the recombination leading to the negligible activity of amorphous TiO₂.

Photocatalytic Activity of Anatase Crystallites. As depicted in Figure 8, a large difference in the photocatalytic activity of anatase TiO₂ powders (f(anatase) = 1) was seen. This is in accord with the fact that many anatase TiO₂ powders available commercially or prepared in laboratories show extensively different photocatalytic activity; i.e., the crystal phase is not a sole decisive factor of activity. The rate for system C is plotted in Figure 10 as a function of anatase crystallite size. It seems that the rate of H₂ production, i.e., photocatalytic activity, in system C increased with the growth of anatase particle size of ca. 30 nm, and thereby we could conclude that amorphous TiO₂ has negligible photocatalytic activity. The referee also suggested the difficulty in estimation of photoabsorption characteristics. Previous reports have demonstrated the more sophisticated procedure, e.g., Palmsano, L.; Augugliaro, V.; Compostrini, R.; Shiavello, M. J. Catal. 1993, 143, 149. In this paper, however, simple procedures were employed for convenience to evaluate the photocatalytic activity.

Acknowledgment. The authors thank Dr. Yukihiro Tsuchida (Idemitsu Kosan Co., Ltd.) for the supply of amorphous TiO₂ samples. Professor Yoshihiko Kera, Dr. Hiroshi Kominami, and Mr. Jun-ichi Kato (Kinki University) are gratefully acknowledged for the permission to use the reflection spectrometer and for the help in measurements. The authors are indebted to Messrs. Satoshi Kusakabe and Takayuki Yako (Kyoto University) for their help in Ti³⁺ measurements and data processing, respectively. Financial support by Foundation of Space Science Promotion and Kansai Research Foundation for Technology Promotion is acknowledged. This investigation was partly supported by a Grant-in-Aid from the Ministry of Education, Science, Sports, and Culture of Japan.

References and Notes


(33) Davidson, R. S.; Morrison, C. L.; Abraham, J. *J. Photochem.* 1984, 24, 27.


