

Short communication

Novel porous TiO₂ glass-ceramics with highly photocatalytic abilityTetsuo Yazawa^{*}, Fumiko Machida, Kaori Oki,
Atsushi Mineshige, Masafumi Kobune*Graduate School of Engineering, University of Hyogo, 2167 Shosha, Himeji-City, Hyogo 671-2201, Japan*

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Abstract

A porous TiO₂ glass-ceramics with high photo-oxidative activity was successfully obtained from the SiO₂–Al₂O₃–B₂O₃–CaO–TiO₂ glass system. Rutile-type TiO₂ was observed in the crystallization temperature range of 973–1173 K. The band gap of the glass-ceramics coincided approximately with that of rutile-type TiO₂. The photocatalytic activity of this glass-ceramics was about four times larger than that of a TiO₂-coated photocatalyst fabricated by the sol–gel process. Furthermore, as this porous TiO₂ glass-ceramics contained TiO₂ in composition form, it could prevent peeling of the TiO₂ from the substrate. As well, this glass-ceramics can be easily shaped into sheets, tubes, rods, etc.

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1. Introduction

Photocatalyst such as TiO₂ is widely used in various products such as air purifiers, deodorants, and for sterilization and anti-fouling [1]. However, the photocatalysts are used in dispersed form and are very difficult to collect after use. To overcome this shortcoming, photocatalysts have recently been developed in which TiO₂ was coated on appropriate substrates, such as a ceramic, glass or metal plate, or glass tube, by the sol–gel process [2–5]. However, in these applications, the coated TiO₂ easily peels off from the substrate.

One of significant characteristics of glass-ceramics is the excellent formability [6]. In the present study, a novel porous TiO₂ glass-ceramics with highly photocatalytic ability was prepared in the SiO₂–Al₂O₃–B₂O₃–CaO–TiO₂ glass system by the crystallized glass process. As this porous TiO₂ glass-ceramics contains TiO₂ in composition form, it prevents peeling of the TiO₂ from the substrate. And further, this glass-ceramics can be easily shaped into sheets, tubes, rods, etc. Consequently, this glass-ceramics is highly suitable for incorporation into a photocatalytic reactor.

2. Experimental**2.1. Glass melting**

The glass was prepared by mixing the desired amounts of the analytical-grade reagents TiO₂, CaCO₃, Al(OH)₃, SiO₂, MgHPO₄, H₃BO₃, ZrO₂ and Mg(NO₃)₂. 50 g of the powdered mixture was melted at 1773 K for 3 h in a platinum crucible in an electric furnace, and then cast on a carbon plate.

2.2. Crystallization and acid leaching

Crystallization was performed in the temperature range of 773–1173 K over a period of 15 h. A porous TiO₂ glass-ceramics was obtained by acid leaching with an aqueous solution of 3-N HNO₃ at 371 K for the desired time [7,8].

2.3. Characterization of the glass

Crystals in the glass were analyzed by X-ray diffraction (XRD) measurement with Cu K α radiation (Mini Flex Rigaku Co.). The TiO₂ content was analyzed by X-ray fluorescence analysis (SEA2001 Seiko Instruments Inc.). The measurement of pore characteristics was performed by N₂ adsorption method

^{*} Corresponding author. Tel.: +81 79 267 4896; fax: +81 79 267 4896.

E-mail address: yazawa@eng.u-hyogo.ac.jp (T. Yazawa).

Table 1
Composition of initial glasses.

	A	B	C	D	E	F
TiO ₂	18	20	26	22	20	20
SiO ₂	36	29	26	32	25	29
Al ₂ O ₃	12	13	10	13	13	13
CaO	25	28	28	27	28	28
MgO	4	5	5	1.5	9	5
P ₂ O ₅	2	2		3.5		
B ₂ O ₃	2	2	5		5	2
ZrO ₂	1	1		1		1
Na ₂ O						2

(BELSORP-mini BELL JAPAN INC.). The diffusive reflectance spectra were determined by using a photometer (V-650 JASCO Co.) equipped with an integrating sphere accessory.

2.4. Evaluation of photocatalytic activity

Photocatalytic activity was evaluated by measuring the photo-degradation of a 200 mdm³ aqueous methylene blue solution with an initial concentration of 56 $\mu\text{mol/dm}^3$ [9,10]. A quartz cell was used as the reaction vessel under the irradiation of a 10-W low-pressure mercury lamp (Model UVL10DL-12, Sen Tokusyu Kogen Ltd.). The weight of the glass-ceramics used in each measurement was 1.0 g. The concentration of methylene blue was determined by a spectrophotometer (U-1100 HITACHI Co.) at 664 nm.

To avoid the influence of adsorption on the porous glass-ceramics surface, the samples were soaked in an aqueous methylene blue solution with the same concentration for 24 h under a reduced pressure of 0.1 abs. MPa in order to attain adsorption equilibrium prior to photocatalytic activity measurement. To attain adsorption equilibrium rapidly, the samples

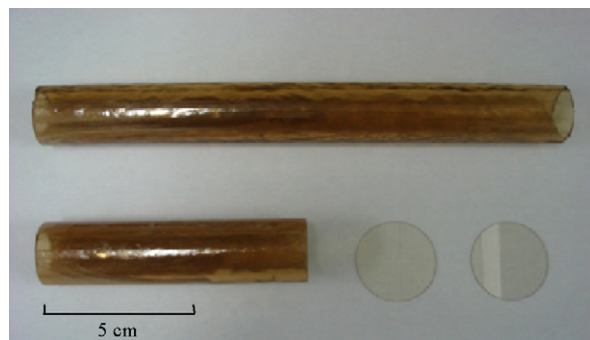


Fig. 1. Photograph of glass shaped into a tube and disk.

were acid leached for 15 min in order to obtain a thin porous layer on the sample surface.

3. Results and discussion

3.1. Glass

Table 1 tabulates the compositions of the initial glasses used. Composition F was the best glass-forming, consequently the best formability judging from its transparency and XRD measurement. This may be attributed to the addition effect of Na₂O. The second and third were A and B, respectively. A and B were slightly brownish, clear glasses. After crystallization, F showed no TiO₂ crystal peak, as revealed by XRD, while A showed a very clear TiO₂ peak, as described in Section 3.2. The addition of Na₂O might play some role on the inhibition of TiO₂ crystallization by decreasing glass transition temperature [11]. The larger the content of TiO₂ crystals in glass-ceramics, the larger the photocatalytic activity. In this meaning, C is expected of high photocatalytic activity. However, the larger the content of TiO₂ in glass-ceramics, the smaller the formability by its

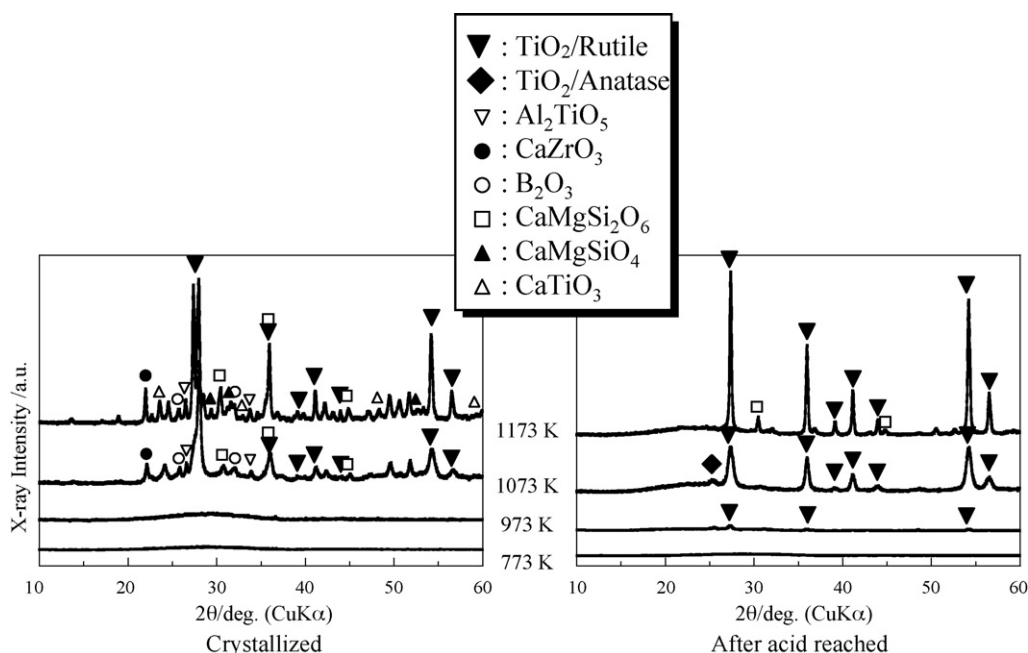


Fig. 2. XRD patterns of the glass-ceramics for various crystallization temperatures.

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