Preparation and Reaction of Titania Particles Encapsulated in Hollow Silica Shells as an Efficient Photocatalyst for Stereoselective Synthesis of Pipecolinic Acid

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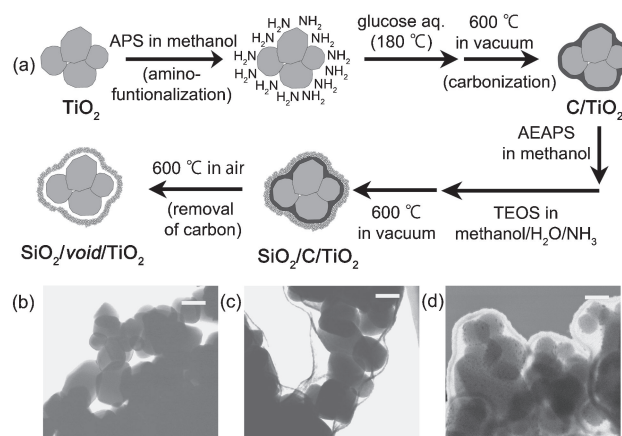
Hollow core–shell particles of titania core and silica shell were synthesized by multistep process, and the core–shell particles showed improved stereoselectivity in the photocatalytic redox-combined synthesis of L-pipecolinic acid from L-lysine in an aqueous suspension without reducing the original activity of the bare titania core.

Photocatalytic reactions occurring on the surface of photo-irradiated titania (TiO_2) have garnered a wide interest due to their potential environmental applications.[1,2] An example is photoinduced removal of chemical contaminants under atmospheric conditions, being attributed to the ability of TiO_2 photocatalyst to cleave chemical bonds nonselectively, i.e., mineralization. However, selective reactions of targeted chemicals are also possible. One of the most useful approaches for selective photocatalytic reaction is operation of a photocatalytic reaction under deaerated conditions where undesirable excessive oxidation through the radical chain reaction with oxygen (O_2) is prohibited, and thereby an alternative electron acceptor should be used.[3,4] Another approach for the selective organic synthesis is utilization of photocatalysts of or in defined microstructures; TiO_2 particles or isolated titanium oxide species are distributed onto or into inorganic supports.[5,6] According to previous works,[7,8] one of the most widely used methods to provide the selectivity is encapsulation of TiO_2 particles into porous substances. However, the surface coverage of these substances causes decrease in intrinsic photocatalytic activity of the medial TiO_2.
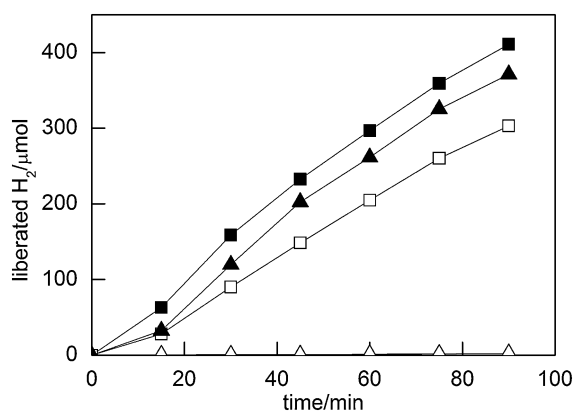
We have reported[9,10] fabrication of a novel core–shell composite photocatalyst which consisted of commercially available TiO_2 particles incorporated in a hollow silica shell (SiO_2/void/TiO_2). The composite possesses size-selective properties in the photodecomposition of organic compounds; SiO_2/void/TiO_2 showed photocatalytic activity for decomposition of small substrates retaining the activity of original bare TiO_2, while negligible activity for polymers was observed, i.e., SiO_2/void/TiO_2 exhibits molecular size selectivity. Recently, we have attempted to use the SiO_2/void/TiO_2 photocatalyst for the synthesis of L-pipecolinic acid (L-PCA), a useful intermediate material for various fine chemicals,[11] and found another function of silica shell to improve stereoselectivity, instead of molecular-size selectivity.

Preparation of SiO_2/void/TiO_2 was performed by coating of TiO_2 with a carbon layer and a silica layer followed by heat treatment to remove the carbon layer,[9] as shown schematically in Figure 1a (For details, see SI[20]). An SEM image of source TiO_2 core (Ishihara Sangyo ST-41) is shown in Figure 1b. An angular morphology of the sample was distinctly observed. The particle

Figure 1. (a) Schematic representation of the procedure for preparation of SiO_2/void/TiO_2, SEM image taken in transmission mode for (b) TiO_2, (c) $SiO_2(0.5)$/void/TiO_2, and (d) $SiO_2(0.5)$/void/TiO_2 after deposition of Pt particles. Scale bar corresponds to 100 nm.

size was in the range of 100–300 nm. The TiO_2 powder was treated with 3-aminopropyltrimethoxysilane (APS), and the APS-modified TiO_2 was then subjected to hydrothermal reaction in aqueous glucose at 180 °C for 6 h. The resulting polysaccharide (PS)-covered particles were recovered and heated at 600 °C under vacuum for 2 h. This resulted in the encapsulation of the particle aggregates with a thick uniform layer of carbon (C/TiO_2). The thickness of layer was 30–80 nm. Then, C/TiO_2 was treated with N-(2-aminoethyl)-3-aminopropyltrimethoxysilane (AEAPS) and then with tetraethyl orthosilicate (TEOS) followed by heat treatment under vacuum at 600 °C to obtain TiO_2 particles covered with a carbon layer and a silica layer (SiO_2(silylation time/h)/C/TiO_2). Finally, the carbon layer was removed by calcinations at 600 °C for 2 h in air, thus successfully yielding TiO_2 encapsulated in a hollow silica shell (SiO_2/void/TiO_2). SEM image in transmission mode (Figure 1c) showed the presence of void space of 3–10 nm in width between shell of around 9–10 nm in thickness and core TiO_2 particles for $SiO_2(0.5)$/void/TiO_2. The presence of void space was also supported by the fact that specific surface area (BET method) of SiO_2/void/TiO_2 ($29 m^2 g^{-1}$) was more than twice that of the original TiO_2 ($13 m^2 g^{-1}$). As a reference, TiO_2 mechanically mixed with silica (mec-SiO_2 + TiO_2) and TiO_2 directly coated with silica (dir-SiO_2/TiO_2) were also prepared, the latter of which was prepared according to the procedures by Graf et al.[12] with slight modification (For details, see SI[20]).

Figure 2. Time-course curves of H_2 liberated from aqueous methanol solutions by TiO_2 (filled squares), $SiO_2(0.5)/void/TiO_2$ (filled triangles), $mec\text{-}SiO_2 + TiO_2$ (open squares), and $dir\text{-}SiO_2/TiO_2$ (open triangles) preirradiated in aqueous $H_2[PtCl_6]$ solutions.

Since platinum (Pt) deposits on the TiO_2 surface are required for the photocatalytic synthesis of L-PCA,[13] all samples were platinized (2 wt %) using two-step photodeposition. First, a sample was suspended in water containing the required amount of hydrogen hexachloroplatinate(IV) ($H_2[PtCl_6]\cdot6H_2O$), irradiated by a 400-W mercury arc (Eiko-sha 400; ca. 25 mW cm^{-2} at 300–400 nm) for 1.5 h, and then irradiated for an additional 1.5 h in the presence of 50 vol % methanol.

Figure 2 shows the time-course curves of hydrogen (H_2) liberation from aqueous methanol solutions in the second step of the platinization. Almost linear increase in the amount of H_2 was observed after some induction period for all the samples except for $dir\text{-}SiO_2/TiO_2$, suggesting that reduction of platinum complex to metallic state, to induce methanol dehydrogenation, required 5–10 min irradiation. As shown in this figure, $dir\text{-}SiO_2/TiO_2$ was almost inactive with negligible H_2 liberation possibly due to retardation of adsorption of substrates, methanol, and $H_2[PtCl_6]$, participating in the reaction onto the TiO_2 surface by the covering silica layer to result in practically no Pt deposition. The activity of $SiO_2/void/TiO_2$ was almost the same as that of bare TiO_2 despite the presence of silica shell and even higher than that of $mec\text{-}SiO_2 + TiO_2$. SEM observation of the sample after the platinization shown in Figure 1d clearly indicates the deposition of fine Pt particles onto TiO_2 without any collapse of the silica shells. A similar finding was observed in our previous research, and this can be attributed to the presence of pores in silica shell and void spaces between the shell and core TiO_2 particles.[10] These structures led to efficient mass transfers through a silica shell to supply substrates that participate in this reaction to the naked active surface of the TiO_2 core.

For the photocatalytic reaction of redox-combined stereoselective synthesis of L-PCA from L-lysine (L-Lys), a Pt-loaded photocatalyst (0.05 g as TiO_2) was suspended in an aqueous solution (5.0 cm^3) containing L-Lys (100 μmol) and photoirradiated with a high-pressure mercury arc (Eiko-sha, 400 W) under argon (Ar) under magnetic stirring (1000 rpm). The photoirradiation was performed through a cylindrical Pyrex glass filter and a glass reaction tube (18 mm in diameter and 180 mm in length) so that light of wavelength >290 nm reached the suspension. The temperature of the suspension during photo-
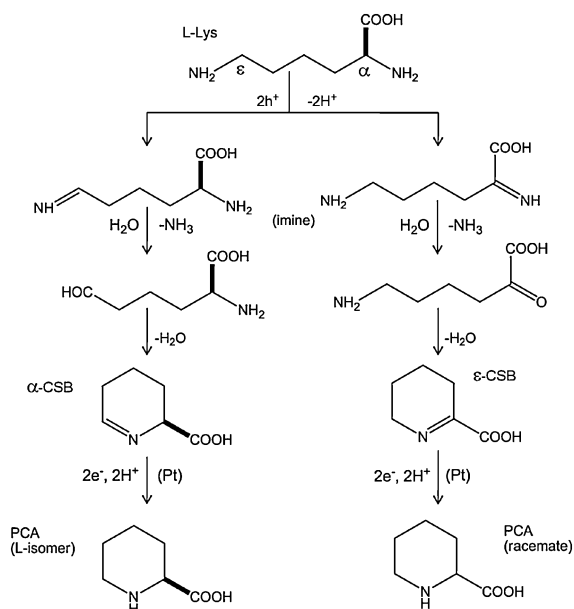
Table 1. Synthesis of PCA from L-Lys using various platinized TiO_2 photocatalysts

Photocatalyst	Conversion /%	S_{PCA}[a] /%	OP_{PCA}[b] /%	R_{PCA}[c]	Y_{H2}[d] /μmol
TiO_2	100	51	57	27	75
$mec\text{-}SiO_2 + TiO_2$	100	52	59	27	63
$dir\text{-}SiO_2/TiO_2$[e]	14	26	—[f]	0.2	2
$SiO_2(0.5)/void/TiO_2$	98	43	70	22	72
$SiO_2(1.5)/void/TiO_2$	96	50	70	25	50
$SiO_2(3.0)/void/TiO_2$	96	46	70	23	57

[a]Selectivity of PCA production based on amount of consumed L-Lys. [b]Optical purity of L-PCA. [c]Rate of PCA formation in the unit of μmol h^{-1}. [d]Yield of H_2. [e]Platinization via photodeposition was unsuccessful (see text). [f]Not determined.

irradiation was maintained at $25 \pm 0.5\,°C$ by the use of a thermostated water bath. After irradiation for 2 h, a portion (0.2 cm^3) of the gas phase of the sample was withdrawn with a syringe and subjected to gas chromatographic analysis (GC, Shimadzu GC-8A with an MS-5A column and a TCD detector) for H_2. The yield of enantiomers of PCA, as well as the amount of unreacted L-Lys, was measured by HPLC (Shimadzu LC-6A equipped with a Daicel Chiral-Pak MA(+) column and an ultraviolet absorption detector).

Table 1 summarizes the results for the synthesis of L-PCA from L-Lys by 2-h photoirradiation using various platinized TiO_2 photocatalysts. Photoirradiation of the TiO_2 photocatalysts suspended in an aqueous solution of L-Lys under Ar led to the formation of PCA, as reported previously.[14,15] Complete consumption of L-Lys was achieved using TiO_2 and also $mec\text{-}SiO_2 + TiO_2$. These photocatalysts showed very similar results in terms of selectivity (S_{PCA}), optical purity (OP_{PCA}), and the rate of PCA formation (R_{PCA}), suggesting that the mechanical mixing of silica with TiO_2 does not give any effect on this reaction as only the TiO_2 part was responsible for the photocatalytic reaction. As expected, $dir\text{-}SiO_2/TiO_2$ showed poor photocatalytic activity to convert only 14% of L-Lys, thus proving that direct coverage of the TiO_2 surface with silica hinders the activity of the TiO_2 by prohibiting Pt deposition as well as L-Lys adsorption onto the bare TiO_2 surface. The $SiO_2(0.5)/void/TiO_2$ particles prepared with 0.5 h of silylation period showed the performance almost the same as that of bare TiO_2. Although the selectivity was slightly lower than that of bare TiO_2, $SiO_2/void/TiO_2$ exhibited the highest OP_{PCA}, 13% more than that of platinized bare TiO_2, among all the samples. In order to further prove the effectiveness of the hollow core–shell structure, $SiO_2/void/TiO_2$ with a thicker layer of silica shell was also prepared, by extending the silylation period (1.5 and 3.0 h). The thickness of the silica layer was increased to 14–32 at 1.5 h and 28–45 nm at 3.0 h from 9 to 10 nm for $SiO_2(0.5)/void/TiO_2$. While $SiO_2(1.5)/void/TiO_2$ exhibited the best performance among the tested samples, it seemed that the photocatalytic performance (conversion, S_{PCA}, OP_{PCA}, and R_{PCA}) was almost independent of the silica shell thickness. This suggests that the silica shell behaves as highly porous optically transparent penetration-free layer which surrounds the TiO_2 core and that this swollen sponge-like silica layer controls the stereoselectivity of the reaction.

Scheme 1. Proposed mechanism of the photocatalytic N-cyclization of L-Lys on platinized TiO₂ photocatalysts.

It has been proposed that PCA formation from L-Lys proceeds through redox-combined mechanism shown in Scheme 1:[16] one of the amino groups in L-Lys is oxidized by positive holes (h^+) to imines, which are then hydrolyzed to an aldehyde or keto acid by ε- or α-amino group oxidation, and then cyclic Schiff base (CSB) intermediates formed by intramolecular condensation are reduced by photoexcited electrons (e^-) to yield PCA. According to this mechanism, OP_{PCA} is regulated by (1) selectivity in the position in the first oxidation process and (2) difference in efficiency in the following second process of conversion from imine into PCA between ε- and α-routes; S_{PCA} corresponds to the average efficiency of the second process. On the assumption of the same efficiency in the second process for α- and ε-routes, OP_{PCA} shows proportion of the ε-route, since ε- and α-routes yield L- and racemic PCA, respectively. A possible reason for improved OP_{PCA}, with almost the same S_{PCA}, by the use of SiO₂/$void$/TiO₂ is increase in the proportion of ε-route, presumably due to the acidity of silica.[17] It has been observed that operation of the reaction at lower pH improved OP_{PCA} and decreased R_{PCA} when platinized (bare) TiO₂ particles were used as a photocatalyst.[18] Since ε-amino group is protonated to be an ammonium group ($-NH_3^+$), compensating negative charge of carboxylate and leaving α-amino group in neutral form under the conditions employed in this study,[19] preferential oxidation of ε-amino group cannot be expected with ordinary photocatalyst particles. Possible acidic microenvironment of the core TiO₂ surface induced by silica shell might lead to protonation of α-amino group to result in the retardation of α-route due to higher (more anodic) oxidation potential of ammonium form of amino groups. As reported previously,[15] blocking of ε-amino group by carbamoyl derivatization not to be protonated and thereby preferential protonation of α-amino group in Lys led to the production of optically pure L-PCA. Preliminary study on acid properties of samples by ammonia TPD (Belsorp, Bel Japan)

suggested the presence of a small amount of weak acid sites (desorption at 150–300 °C; ca. 1 µmol in total) presumably due to the SiO₂ layer. Measurements of the L-Lys adsorption and acid properties through ammonia TPD and ζ-potential analysis are under study.

In conclusion, the present hollow core–shell structured photocatalyst provides the improved OP_{PCA} keeping the S_{PCA} and R_{PCA} without addition of any chemicals, such as an acid or a buffer solution, which must be separated in the post reaction procedure. Though improved OP_{PCA} was still not so high (70%), this enables purification of L-PCA by recrystallization.[15] It is expected that modification of silica with more acidic functional groups and/or choice of appropriate thickness of void space between core and shell improve the performance of photo-catalysts for stereoselective synthesis of L-PCA, and study along this line is now in progress.

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19　L-Lysine hydrochloride was used and equimolar amount of sodium hydrochloride was added to neutralize the acid.
20　Supporting Information is available electronically on the CSJ-Journal Web site, http://www.csj.jp/journals/chem-lett/index.html.