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Sodium hexatitanate photocatalysts prepared by a flux method for reduction of carbon dioxide with water

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Abstract

Flux method was applied to prepare sodium hexatitanate ($\text{Na}_2\text{Ti}_6\text{O}_{13}$) photocatalysts for reduction of carbon dioxide with water. The preparation conditions were examined, such as the starting material, the flux, the composition in the molten mixture, the holding temperature, and the cooling rate of the molten mixture, so as to obtain several sodium hexatitanate samples consisting of the crystals with various morphologies and sizes. These samples were loaded with 0.1 wt% of Ag cocatalyst nanoparticles ($\text{Ag}/\text{Na}_2\text{Ti}_6\text{O}_{13}$) and examined for photocatalytic reduction of carbon dioxide with water, where carbon monoxide, hydrogen and oxygen were formed as products. It was found that the photocatalytic production rate of carbon monoxide varied with the morphology and the size of the sodium hexatitanate crystals. Among the prepared samples, thick rod-like crystals exhibited high photocatalytic activity for the reduction of carbon dioxide.

Keywords: Heterogeneous photocatalyst; CO_2 reduction; Artificial photosynthesis; Flux method.

1. Introduction

An increasing concentration of atmospheric carbon dioxide (CO₂), which should originate from large consumption of fossil fuels, has been considered as the most responsible reason for the current climate change [1]. Since the activity of the photosynthesis in leaf for CO₂ fixation on the earth is limited, artificial photosynthesis producing valuable compounds from CO₂ by utilizing solar energy has been much attractive in view of reducing the CO₂ emission, recycling CO₂ as a carbon source and converting solar energy to the chemical potential of the products [2–8].

Heterogeneous photocatalysis has been studied for the reduction of CO₂ with water to produce formate, carbon monoxide (CO), methanol and methane [9–15]. Especially, in recent years, many kinds of Cu or Ag-modified photocatalysts to produce CO, hydrogen (H₂) and oxygen (O₂) have been reported, such as Cu/ZrO₂ [16], Ag/BaLa₄Ti₄O₁₅ [17], Ag/Ga₂O₃ [18–21], Ag/Zn-Ga₂O₃ [22], Ag/KCaSrTa₅O₁₅ [23], Ag/La₂Ti₂O₇ [24], and Ag/CaTiO₃ [25, 26]. In these systems, two reactions competitively proceed, i.e., photocatalytic reduction of CO₂ to form CO and O₂ (eq. 1) and photocatalytic water splitting into H₂ and O₂ (eq. 2).

The reaction selectivity between the two reactions varies with the employed photocatalyst. On the surface of the photocatalyst, both reduction of CO₂ and proton by the excited electrons (eq. 3 and 4) and oxidation of water into O₂ and proton (eq. 5) can occur.

In these photocatalytic systems, however, the enough production rates have not been achieved yet for the practical use. Thus, a clear guidance for catalyst design has been required for further development.

In some systems, it is reported that the morphological structure of the photocatalyst affects the photocatalytic activity for the CO₂ reduction. For example, on the Ag/BaLa₄Ti₄O₁₅ photocatalyst of anisotropic plate-like structure [17], the reduction of CO₂ to produce CO can take place on the Ag cocatalyst at the edges of the plates while the oxidation can progress on the BaLa₄Ti₄O₁₅ surfaces. In other words, it can be recognized that the separated reaction fields for reduction and oxidation would contribute to the high photocatalytic performance. In our previous study [25], we prepared several Ag/CaTiO₃ photocatalyst samples, where a clear relationship was revealed between the particle size of polyhedral CaTiO₃ crystals in micro meter scale and the photocatalytic production rate of CO, i.e., the larger size of crystal provided higher activity for the production of CO.

Flux method employed in the previous and the present studies can provide fine single crystals or poly crystals and thus has been applied to fabricate various micro or nano-sized photocatalysts in recent years. For example, many kinds of titanate photocatalysts were reported such as Na₂Ti₆O₁₃ [27], Na₂Ti₃O₇ [28], K₂Ti₆O₁₃ [29,30], CaTiO₃ [25,26], SrTiO₃ [31], La₂Ti₂O₇ [32,33], PbTiO₃ [34], CaZrTi₂O₇ [35] and AgLi_{1/3}Ti_{2/3}O₂ [36].

In the present study, we prepared several Ag/Na₂Ti₆O₁₃ photocatalyst samples by using a flux method. The Na₂Ti₆O₁₃ semiconductor has a unique tunnel structure [27] and has a bandgap (3.5 eV) [37] larger than that of anatase TiO₂ (3.2 eV) and rather closed to that of CaTiO₃ (3.6 eV) [25,38]. Thus, it is expected to be also active for the photocatalytic reduction of CO₂. Actually in the present study, it was found that the Ag/Na₂Ti₆O₁₃ photocatalyst can promote the photocatalytic reduction of CO₂ to CO. In addition, a good relationship was revealed between the structure and the photocatalytic activity.

2. Experimental

2.1. Sample preparation

Most of the Na₂Ti₆O₁₃ samples in the present study were synthesized by a flux method in the same manner described our previous works [25,26,30] from Na₂CO₃ (Rare metallic, 99.9%, melting point is 1124 K) and TiO₂ (rutile, Kojundo, 99.9%, 2 μm, melting point is 2116 K) as start materials by using some chlorides such as LiCl, NaCl, KCl and CaCl₂ as a flux (the melting points are 878, 1074, 1043 and 1045 K, respectively). The molar ratio of Na₂CO₃ to TiO₂ was 1:6, and various ratios of the substrate (*x* mol% as Na₂Ti₆O₁₃) in the mixture with the flux were examined, where *x* was defined as: $x \text{ [mol\%]} = 100 \times (\text{amount of Na}_2\text{Ti}_6\text{O}_{13})$

[mol]) / (amount of $\text{Na}_2\text{Ti}_6\text{O}_{13}$ [mol] + amount of a flux [mol]). The mixture of starting materials was put into a platinum crucible, which was loosely covered by a lid, heated at a rate of 200 K h^{-1} to 973, 1073 or 1273 K (standard: 1273 K) in an electric furnace, held at this temperature for 10 h, and then slowly cooled at a rate of -100 , -10 or -5 K h^{-1} (standard: -100 K h^{-1}) to 773 K, followed by being naturally cooled to room temperature in the furnace. It is considered that the decarbonation of Na_2CO_3 would take place to form Na_2O species during heating, the mixture of the starting materials would be molten, and then both melted Na_2O and melted TiO_2 would react with each other to form $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystallites in the molten salt during the cooling step. The obtained products were well washed with hot water (353 K) three times to remove the flux. These samples are referred to as NTO(flux, x , *note if any*), *e.g.*, NTO(NaCl ,50), NTO(NaCl ,5,1073K), NTO(NaCl ,50, -10Kh^{-1}) and so on.

One sample was prepared by a solid state reaction method (SS method). The same starting materials, Na_2CO_3 (0.03 mol) and TiO_2 (rutile, 0.18 mol), were dried at 373 K, mechanically mixed by a wet ball-milling method with 150 g of alumina balls and 80 mL of acetone in a plastic bottle (300 mL) for 24 h at 120 rpm, dried in an oven at 353 K overnight, and heated in air atmosphere at 1273 K for 10 h in an alumina crucible in the furnace. The sample is referred to as NTO(SS). Another sample was prepared by the flux method with anatase TiO_2 supplied by Catalysis Society of Japan, JRC-TIO-8 ($338 \text{ m}^2\text{g}^{-1}$) as the start material, *i.e.*, NTO(NaCl ,50,anatase).

Ag cocatalyst was loaded on the surface of the $\text{Na}_2\text{Ti}_6\text{O}_{13}$ samples by a photodeposition method, where AgNO_3 and methanol (25 vol.%) was used as the Ag source and the reductant, respectively. The loading amount of Ag as metal was 0.1 wt%. The samples are referred to as Ag/NTO, *e.g.*, Ag/NTO(NaCl ,50).

2.2. Characterizations

Several kinds of analyses were employed to characterize the prepared samples, such as XRD, XRF, ICP, SEM, TEM, DR UV-vis, BET, DTA, pH_{ZPC} , and XAFS, the details of which are mentioned in the Electric Supplementary Information (ESI).

2.3. Photocatalytic reaction tests

Photocatalytic reaction tests were carried out under ambient temperature and pressure in the same manner as the previous study [25]. The details are described in the ESI. Detected products were CO , H_2 , and O_2 .

Other products were not observed in this system even in the liquid phase after the reaction, although the possibility of the formation of undetected byproducts was not denied especially in the liquid phase. In the present study, the selectivity to CO in the reduced products was defined as follows: $S_{CO} (\%) = 100 \times (\text{the production rate of CO}) / (\text{sum of the production rates of CO and H}_2)$.

3. Results and discussion

3.1. Preparation with various fluxes

Since molten salt as a flux can regulate the crystal growth to provide unique morphology of the crystals in many cases, some kinds of the salts such as LiCl, NaCl, KCl, and CaCl₂ were examined for the preparation of Na₂Ti₆O₁₃ samples. Fig. 1 shows XRD patterns of the obtained samples with that from database [39] as a reference and that of the rutile TiO₂ sample used as the start material. Among them, the NTO(NaCl,50) sample, the NTO(KCl,50) sample and the NTO(SS) sample (Fig. 1b, 1c, and 1e) exhibited almost the same pattern to that of the reference data (Fig. 1f), indicating that Na₂Ti₆O₁₃ crystallites of monoclinic structure were obtained. It is noted that the distribution of the diffraction line intensity of the NTO samples prepared by the flux method are different from that of the NTO(SS) sample and the reference, for example, the intensity of the line at 24.5° corresponding to (110) plane was relatively low for the former patterns and high for the latter patterns, implying that the Na₂Ti₆O₁₃ crystals prepared by the flux method have an anisotropic morphology. Although the NTO(NaCl,50) sample and the NTO(KCl,50) sample exhibited almost similar diffraction patterns, the latter showed a small shift to lower angle. XRF measurement revealed that the NTO(KCl,50) sample contained small amount of potassium and the composition of the sample was estimated to be Na_{1.46}K_{0.54}Ti₆O₁₃, where potassium cation having the larger size than sodium cation would enlarge the lattice constant.

The NTO(LiCl,50) sample showed a quite different pattern than others (Fig. 1a). By using XRF and ICP, the composition of this sample was estimated to be Li_{1.17}Na_{1.40}Ti₆O_{14.2}, indicating that pure phase of sodium hexatitanate (Na₂Ti₆O₁₃) was not obtained. It is also revealed that the NTO(CaCl₂,50) sample was a mixture of TiO₂ (major) and CaTiO₃ (minor) (Fig. 1d). These facts indicate that both LiCl and CaCl₂ salts are not suitable for the preparation of Na₂Ti₆O₁₃ crystals.

The XRF measurement revealed that these samples did not contain a detectable amount of Cl anions, meaning that the removal of the flux was almost completed by washing with hot water.

SEM images elucidated various morphologies of these NTO samples prepared with various fluxes in the flux method and without any flux in the SS method (Fig. 2). The NTO(LiCl,50) sample consisted of large particles (Fig. 2a), the NTO(NaCl,50) and NTO(KCl,50) samples showed rod-like morphology (Fig. 2b and 2c), the NTO(CaCl₂,50) sample exhibited particles covered with small fragments (Fig. 2d), and the NTO(SS) sample showed ununiformed particles (Fig. 2e). In some cases, various facets could be observed on the surfaces of the particles prepared by the flux method (Fig. 2, a–c). It was clarified that both the use of flux and the property of the flux much affected the shape of the particles. This confirms that the molten salt can regulate the crystal growth of each facet to form the unique morphology and this effect depends on the property of the cation such as Li⁺, Na⁺, K⁺ and Ca²⁺. Among them, it is remarkable that the two Na₂Ti₆O₁₃ samples prepared by the flux method, i.e., the NTO(NaCl,50) sample and the NTO(KCl,50) sample, exhibited rod-like crystals. They were similar to the reported ones in the literatures for Na₂Ti₆O₁₃ [27] and K₂Ti₆O₁₃ [30], where the cross-section of the rod-like structure was typically hexagonal. In the present samples some steps were also observed. On the other hand, the NTO(SS) sample of the Na₂Ti₆O₁₃ structure prepared by the SS method showed granular polyhedral particles connected to each other.

As a conclusion in this section, it is noted that the NaCl flux is suitable for the preparation of the pure phase of the hexagonal rod-like Na₂Ti₆O₁₃ crystals.

3.2. Preparation with NaCl flux with various conditions

Some samples were prepared by using the NaCl flux with various solute concentrations in the standard condition. Fig. 3 shows their SEM images. It was elucidated that these samples prepared by the flux method consisted of similar hexagonal rod-like crystals and both the size and the aspect ratio varied with the solute concentration x in the molten mixture. The lowest solute concentration (5%) provided the clear shape of the fine rod-like crystals (Fig. 3a), while the highest (70%) gave somewhat irregular shape (Fig. 3c). The sizes of the crystals were evaluated from the SEM images and listed in Table 1, entries 1–4. The NTO(NaCl,5) sample consisted of the longest rod-like crystals of 6.4 nm in length on the average. The average length clearly decreased with an increase of the solute concentration x in the molten mixture among the samples prepared by the flux method. On the other hand, the average width of them once increased with an increase

of x to the maximum, i.e., 1.1 nm at $x=50$, then decreased. The aspect ratio of the crystals in these samples decreased from 8.7 to 2.6 with an increase of x , and for the sample prepared by the SS method it was 1.3, almost close to unity (Table 1, entry 5).

Fig. 4 shows a representative TEM image of the NTO(NaCl,50) sample with an electron diffraction pattern. A rod-like particle was obviously found. It is noticed that more transparent perimeter can be observed in the rectangular image of the rod-like crystal, supporting the hexagonal cross-section. When the electron diffraction pattern was monitored at several points of the rod-like crystal, the clear and regular pattern of bright spots was observed for each location (Fig. 4, inset). This fact evidenced that the rod-like crystal was a single crystal.

Fig. 5 shows representative DR UV-visible spectra, which give information for the electronic band structure. The NTO(NaCl,50) sample showed a wide absorption band less than 370 nm in wavelength, indicating that the estimated bandgap was 3.58 eV. This value was very slightly smaller than 3.62 eV for the NTO(NaCl,5) sample and 3.61 eV for the NTO(SS) sample. This fact might be related to the average width of the rod-like crystals, i.e., the thick crystals gave the smaller bandgap. However, no drastic difference on the electronic band structure was observed from each other. The NTO(SS) sample exhibited an additional small band at 360–400 nm, suggesting the presence of very small amount of unreacted titanium oxide that could not be detected by XRD in this sample or some kinds of crystal defects.

The NTO(NaCl,5,973K) sample and the NTO(NaCl,5,1073K) sample were prepared with lower holding temperatures, 973 and 1073 K, respectively. These temperatures are lower than or almost equal to the melting point of pure NaCl (1074 K). XRD profile (not shown) of the former indicated that the holding at 973 K gave a mixture of rutile TiO_2 and $\text{Na}_2\text{Ti}_6\text{O}_{13}$ phases. The SEM image of this sample (Fig. S1a) also shows the mixture of roundish particles and needle-like crystals, assignable to TiO_2 and $\text{Na}_2\text{Ti}_6\text{O}_{13}$ phases, respectively. The sizes of the needle-like crystals were very small as listed in Table 1, entry 6. On the other hand, the holding at 1073 K was sufficient for providing a pure $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystals (XRD was not shown). The SEM images shown in Fig. S1b clarified that the rod-like crystals were formed. The average length and width of the rod-like crystals in this sample were 6.2 and 0.33 nm, respectively as listed in Table 1, entry 7, which were smaller than those for the NTO(NaCl,5) sample prepared at 1273 K (Table 1, entry 1). The variation coefficients for the length and width were 42% and 28% for the sample prepared at 1073 K and 46% and 40% for the sample prepared at 1273 K. These facts indicate that the lower temperature provided

smaller rod-like crystals with narrower size distribution. It is considered that the higher holding temperature would provide enough time for crystal growth in the molten mixture at the longer cooling step.

The cooling rate was also one of the most important parameters controlling the structure of the rod-like $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystals. Fig. 6 shows SEM images of the $\text{NTO}(\text{NaCl}, 50, -y\text{K h}^{-1})$ samples prepared with various cooling rate, -100 , -10 and -5 K h^{-1} , from 1273 K to 773 K . As shown, the slow cooling rate such as -10 and -5 K h^{-1} provided the thicker rod-like crystals than the fast cooling rate as -100 K h^{-1} . The sizes estimated from several SEM images were listed in Table 1 entries 3, 8 and 9. The slow cooling rate would enhance the crystal growth according to Ostwald ripening. The size distribution of the $\text{NTO}(\text{NaCl}, 50, -5\text{K h}^{-1})$ sample seems to become larger with the slow cooling (Fig. 6c), which was supported by the high variation coefficients, 62% and 57%, for the length and width of the thick rod-like crystals, respectively.

The structure of the start material also influenced the morphology. When anatase TiO_2 powder with high specific surface area ($338\text{ m}^2\text{g}^{-1}$) was employed as the start material instead of rutile powder with low specific surface area ($2.5\text{ m}^2\text{g}^{-1}$), the obtained $\text{NTO}(\text{NaCl}, 50, \text{anatase})$ sample consisted of smaller size of rod-like crystals (Fig. S2a, and Table 1, entry 10). In the present study, however, it was not clarified which factor contributed to the smaller size, the anatase structure or the high surface area of the start TiO_2 material. The detailed discussion was described in the ESI.

In this study, it was also confirmed that the composition of sodium titanate can be determined by the molar ratio of the two starting materials, Na_2CO_3 and TiO_2 . In other words, the NaCl flux usually could not react with TiO_2 to form sodium titanate in the present condition. For example, when a mixture of Na_2CO_3 and TiO_2 with the ratio of 1:3 was melted in the NaCl flux and followed by cooling in the same manner, a sample of $\text{Na}_2\text{Ti}_3\text{O}_7$ phase was obtained. When a mixture of only the NaCl flux and rutile TiO_2 powder was examined for the preparation, most of TiO_2 remained and some kinds of sodium titanates were formed as minor products, which was confirmed by the XRD (not shown). In this case, a part of NaCl would be oxidized with the atmospheric O_2 to form Na_2O species in the molten mixture and it would react with TiO_2 to form sodium titanates.

In this section, it was confirmed that well controlled thick hexagonal rod-like $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystals were obtained when the sample was prepared from rutile TiO_2 by using moderate amount of NaCl flux through high holding temperature at 1273 K and the moderately slow cooling rate of -10 K h^{-1} .

3.3. Deposition of the Ag cocatalyst nanoparticles.

Silver nanoparticles were deposited as a cocatalyst on these prepared $\text{Na}_2\text{Ti}_6\text{O}_{13}$ samples by the photodeposition method. Ag loading amount was 0.1 wt% for each sample. Fig. 7 shows DR UV-visible spectra of the representative Ag loaded samples. The Characteristic bands in the visible light region assignable to the surface plasmon resonance (SPR) of the Ag nanoparticles confirmed the deposition of the Ag cocatalyst. The peak position of the SPR band for the Ag/NTO(NaCl,5) and Ag/NTO(NaCl,50) sample was at 500 nm in wavelength while the Ag/NTO(NaCl,70) sample exhibited it at 530 nm. This suggests that the Ag nanoparticles deposited on the Ag/NTO(NaCl,70) sample would be larger than those on the other samples. In the photodeposition method, it is considered that adsorbed Ag ions on the surface of the $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystals are reduced by the photoexcited electron generated in the $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystals. Thus, the structure of the $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystals would affect the photocatalytic property and vary the size of the deposited Ag nanoparticles.

3.4. Photocatalytic activity

The prepared Ag/ $\text{Na}_2\text{Ti}_6\text{O}_{13}$ samples were examined for the photocatalytic reduction of CO_2 by water. The products observed in the present study were carbon monoxide (CO), hydrogen (H_2), and oxygen (O_2), which consisted with the results in our previous study [25,26] and other reports [16–24]. In these systems including the present one, the reduction of CO_2 and water splitting would take place competitively as mentioned above.

In the representative time course of the production rate in the photocatalytic reaction with the Ag/NTO(NaCl,50) sample (Fig. S3), the production rates for H_2 and CO were initially high, but gradually decreased, and finally became almost constant after a long time passing around 24 h.

Although O_2 was not observed initially, it was actually observed later. The stoichiometric ratio of the reduced products (H_2 and CO) and oxidized products (O_2) did not confirmed in the present study, which might be due to the small amount of production and large experimental error to determine the amount of O_2 . However, as discussed in the ESI, there are further some possible reasons for this phenomenon, such as photoadsorption of O_2 on the photocatalyst surface [40–44], the competitive oxidation of water to hydrogen peroxide [45], and the competitive oxidation of Cl^- anion, which is a possible residual from the flux, to form

chlorine (Cl_2) and then hypochlorite species (ClO^-) [46]. To clarify this phenomenon, further study is necessary.

Some kinds of blank tests confirmed that the photocatalyst, light, water, CO_2 gas were absolutely necessary to yield CO. Although CO was produced from CO_2 or NaHCO_3 by photochemical reaction with the irradiated light even in the absence of the photocatalyst or water, the amount of CO was very small. In addition, it was also confirmed that the Ag cocatalyst well accelerated the CO_2 reduction, which is believed to be the active sites for CO production [17–26]. Even if the number of all Ag atoms in the reactor was used as the number of the active sites, the estimated turnover numbers for CO production for 24 h ($\text{TON}_{24\text{h}}$) were larger than 5 over the Ag/NTO(NaCl ,50,–10 Kh^{-1}) sample, which exhibited the highest CO production in the present study. In addition, the reaction took place continuously even after 25 h at the constant rate. These results show that the reduction of CO_2 in the present system proceeds photocatalytically. However, the reaction rate was still too low to estimate the quantum yield exactly in this condition.

The production rates of CO and H_2 after the reaction for 6 and 24 h over the Ag/ $\text{Na}_2\text{Ti}_6\text{O}_{13}$ photocatalysts are listed in Table 2. As mentioned above, although the production of O_2 was really observed, the production rates are not listed in Table 2 since the evaluation would be often imprecise.

The results in Table 2, entries 1–4 show the effect of the fluxes. Among the four samples, the Ag/NTO(NaCl ,50) sample exhibited the highest CO production rate (Table 2, entry 2). The Ag/NTO(KCl ,50) sample (actually, Ag/ $\text{Na}_{1.46}\text{K}_{0.54}\text{Ti}_6\text{O}_{13}$) with similar crystal structure to the Ag/NTO(NaCl ,50) sample exhibited higher CO production rate than the others, i.e., the Ag/NTO(LiCl ,50) sample and the Ag/NTO(CaCl_2 ,50) sample (Table 2, entries 1, 3, and 4). This result suggests that the sample of the $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystal structure exhibited higher photocatalytic activity.

The results in Table 2, entries 2 and 5–8 show the effect of the flux method and the solute concentrations in the molten mixture. The Ag/NTO(SS) sample prepared by the solid state reaction method (Table 2, entry 8) gave a lower CO production rate than the Ag/NTO(NaCl , x) samples prepared by the flux method (Table 2, entry 2 and 5–7), indicating that the present flux method would be more suitable for the preparation of the $\text{Na}_2\text{Ti}_6\text{O}_{13}$ photocatalyst for the CO_2 reduction by water. Further, the photocatalytic activity of the Ag/NTO(NaCl , x) samples also varied with the solute concentration. Fig. 8A shows the variation of the CO production rate on these samples with the solute concentration, x , where the result on the Ag/NTO(SS) sample (Table 2 entry 8) was also plotted at $x=100$. The CO production rate increased with an increase of

the solute concentration to the maximum at 50 mol% and then decreased, which were so-called volcano-shaped plots. On the other hand, Fig. 8B shows the variation of the average length and width of the rod-like crystals (Table 1, entries 1–5) with the solute concentration. Although the length of the rod-like crystals simply decreased with increasing the solute concentration, the width increased then decreased, where the maximum was found at 50 mol%. It is interesting that both CO production rate and the width of the rod-like crystals similarly varied with the solute concentration to show the volcano-shaped plots, suggesting a relationship between them. Such a volcano-shaped plot was not observed for the BET specific surface area listed in Table 1.

The holding temperature of the molten mixture also affected the photocatalytic activity (Table 2, entries 5, 9 and 10). The Ag/NTO(NaCl,5,973K) sample, which did not consist of pure $\text{Na}_2\text{Ti}_6\text{O}_{13}$ phase, exhibited very low activity for the CO production (Table 2, entry 9). The Ag/NTO(NaCl,5,1073K) sample exhibited higher activity for the CO_2 reduction at 24 h than the Ag/NTO(NaCl,5) sample prepared by the high holding temperature at 1273 K (Table 2, entries 5 and 10). The higher surface area of this sample treated lower temperature at 1073 K might contribute the high photocatalytic activity.

The Ag/NTO(NaCl,50, -10K h^{-1}) sample and the Ag/NTO(NaCl,50, -5K h^{-1}) sample, prepared with the low cooling rate (Table 2 entries 11 and 12), exhibited much higher CO production rate than the Ag/NTO(NaCl,50) sample prepared with the high cooling rate, -100 K h^{-1} (Table 2 entry 2), i.e., the sample prepared with the slower cooling rate, consisting of thicker rod-like $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystals, exhibited higher CO production rate. The Ag/NTO(NaCl,50, -10K h^{-1}) sample exhibited 8 times higher CO production rate than the Ag/NTO(SS) sample after the reaction for 24 h.

The Ag/NTO(NaCl,50,anatase) sample prepared from anatase TiO_2 powder with high specific surface area, consisting of the fine rod-like crystals, showed a lower CO production rate than the corresponding sample prepared from rutile, i.e., the Ag/NTO(NaCl,50) sample (Table 2, entry 2 and 13). But it showed a higher rate than the Ag/NTO(SS) sample (Table 2, entry 5).

As a conclusion of this section, the efficient preparation conditions are summarized: The high activity for CO production was obtained by the thick rod-like $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystals prepared in the flux method from rutile TiO_2 by using moderate amount of NaCl flux through high holding temperature at 1073 K or 1273 K and the moderately slow cooling rate of -10 K h^{-1} . This is the relation between the preparation conditions and the

photocatalytic activity. However, fundamentally it is more important to know the relationship between the structure and the photocatalytic activity, which will be discussed further in the next section.

3.5. Relation between the structure and the photocatalytic activity

Fig. 9 shows the relationship between the average width of the rod-like crystals or the particle size in the various $\text{Na}_2\text{Ti}_6\text{O}_{13}$ samples and the photocatalytic activity for CO_2 reduction over the corresponding Ag loaded samples (Table 2, entries 2, 5–8, and 11–13). These $\text{Na}_2\text{Ti}_6\text{O}_{13}$ samples were prepared by the flux method by using NaCl flux with various solute concentrations of 5–70% in the molten mixture, with various cooling rates (-100 , -10 or -5 K h^{-1}) and from rutile or anatase as the starting material. The sample prepared by the solid-state reaction method is also included. All these samples were prepared through heating at 1273 K , and the same amount of Ag ($0.1 \text{ wt\%}$) was loaded on them.

As clearly shown, a good relationship was obtained between them, i.e., the thick rod-like crystal exhibited higher photocatalytic activity for the CO_2 reduction to produce CO molecule, typically larger than $0.7 \mu\text{m}$ in width. Similar correlation was also found in our previous study for the photocatalytic CO_2 reduction with Ag/ CaTiO_3 photocatalysts similarly prepared by the flux method [25]. Especially the $\text{Na}_2\text{Ti}_6\text{O}_{13}$ samples prepared with the slow cooling rate exhibited higher activity, which would originate from the large width of the $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystals such as $1.3 \mu\text{m}$. The aspect ratio of 3.2–3.9 of the thick rod-like structure of these samples (Table 1, entries 8 and 9) might provide a good balance for both reductive and oxidative reactions. It is noted that the clear relationship was found with the CO formation rate but not with the H_2 production rate, the thick structure might be responsible for the state of the Ag nanoparticles as the cocatalyst for the CO production.

3.6. Variation of the Ag cocatalyst nanoparticles

Here, the state of the Ag nanoparticles will be discussed. Fig. 10 shows SEM images of the representative Ag/ $\text{Na}_2\text{Ti}_6\text{O}_{13}$ photocatalysts before and after the photocatalytic reaction test. Since these SEM images were recorded in the reflection mode, the bright dots indicate the Ag nanoparticles. Before the reaction test, only a few and very small white dots could be observed on the Ag/NTO(NaCl,5) sample and the Ag/NTO(NaCl,50) sample, suggesting that the most of the Ag nanoparticles would be small and well dispersed on the crystals (Fig. 10, a and b). The images recorded after using in the reaction tests showed

some larger Ag nanoparticles (Fig. 10, d and e). It is thus proposed that the Ag nanoparticles could aggregate during the photocatalytic reaction. The aggregation was observed also on the Ag/NTO(SS) sample (Fig. 10, c and f). This would be the reasons why the photocatalytic production rates varied with time in the initial period.

As for the variation of the state of the Ag nanoparticles, two possibilities are considered as follows: (i) The Ag nanoparticles might migrate on the surface of the $\text{Na}_2\text{Ti}_6\text{O}_{13}$ crystals under photoirradiation and aggregate to become larger particles. (ii) On the oxidative facets, where the photocatalytic oxidation by positive holes preferably occur, Ag atoms in the Ag nanoparticles could be oxidized and dissolved into aqueous solution as Ag^+ cations, and then they would be deposited and reduced on other Ag nanoparticles located on the reductive facets, where the photocatalytic reduction preferably occurs by the photoexcited electron, as illustrated in Fig. 11.

In the previous study of the Ag/ CaTiO_3 photocatalysts [25], the CaTiO_3 crystals were similarly prepared by a flux method and facets were observed on the crystals. After the photocatalytic reaction test, the Ag nanoparticles were found only on some selected facets of the CaTiO_3 crystals, where the reductive reaction with excited electrons should take place preferentially. In other words, the place stabilizing the Ag nanoparticles would be the reductive sites in the photocatalytic reaction. The Ag cocatalyst nanoparticles stabilized on the reductive sites would be very effective for the photocatalytic CO_2 reduction since the Ag nanoparticles were considered as the receiver of the excited electrons and the cocatalyst to promote the CO_2 reduction to CO.

In the present study, it was notable that the Ag nanoparticles lined along the rod-like crystals in the direction of the long axis on the Ag/NTO(NaCl ,50) sample after the photocatalytic reaction test (Fig. 10e). It seems that the particles are supported by two neighboring facets of the aggregated rod-like crystals, which correspond to the faces vertical to (010) plane such as (001) plane [27]. Since the Ag nanoparticles were stabilized on the long facets, these neighboring facets of the rod-like crystals would be the reductive sites. It is proposed that the neighboring two reductive facets would stabilize the Ag nanoparticles on them from the two directions and more efficiently supply the photoexcited electrons for the two electron step for CO formation (eq. 3), which should contribute to the high activity for the CO production in the photocatalytic CO_2 reduction (Fig. 12). The wide reductive surface would be suitable for holding the large numbers of the

Ag cocatalyst, which would be one of the reasons why the photocatalytic CO₂ reduction was enhanced with the thick rod-like crystals.

The structural variation of the Ag nanoparticles on the Na₂Ti₆O₁₃ samples under the photocatalytic reaction condition was observed in the DR UV-vis spectra of the Ag/NTO(NaCl,50) sample (Fig. S4). The SPR band around 500 nm originated from metallic Ag nanoparticles (Fig. S4a) changed to a broad and weak one (Fig. S4b), indicating that the nanoparticles would change in the size or the oxidation state, supporting the dynamic variation of the Ag nanoparticles as mentioned above. From Ag K-edge XANES spectra of the Ag/NTO(NaCl,50) sample before and after the photocatalytic reaction test (Fig. S5, b and c), it was suggested that the Ag/NTO(NaCl,50) samples before and after the reaction would consist of metallic and oxidized Ag species, suggesting that the Ag cocatalyst nanoparticles would be easily oxidized and reduced during the photocatalytic reaction. This supports the proposed mechanism shown in Fig. 11.

When an aqueous solution of Na₂CO₃ (pH=11) was used for the photocatalytic reaction test with the Ag/NTO(NaCl,30) sample (Table 2, entry 14), the CO production rate and the CO selectivity were higher than that in the solution of NaHCO₃ (pH=8.3–9.2) (Table 2, entry 6). Although the different reaction condition might change the concentration of dissolved CO₂ species in the solution and thus it might increase the reaction rate, the variation of the Ag nanoparticles during the photocatalytic reaction is discussed in the present study. The zero point of charge for the Na₂Ti₆O₁₃ crystals measured by the titration method using the NTO(NaCl,50) sample was pH_{ZPC}=9.3, meaning that the surface of the Na₂Ti₆O₁₃ crystal would be positively charged or close to neutral in the NaHCO₃ solution and negatively charged in the Na₂CO₃ solution. The DR UV-visible spectra after the reaction in this condition (Fig. S4c) showed that the Ag nanoparticles on the negatively charged surface still exhibited the surface plasmon resonance and shifted to shorter wavelength, meaning that the Ag nanoparticles would become more dispersed during the reaction. It is proposed that the dissolved Ag⁺ cation would be preferentially deposited on the negative Na₂Ti₆O₁₃ crystal surface rather than the Ag nanoparticles, which would tend to enhance the dispersion of the Ag nanoparticles on the reductive surface and contribute to the enhancement of the CO production rate. In this condition the highest CO selectivity such as 60% was achieved (Table 2, entry 14), which might be contributed by the highly dispersed Ag nanoparticles as the cocatalyst for the CO production. On the other hand, the Ag nanoparticles on the positive or neutral surface lost the SPR feature of nanoparticle during the photocatalytic reaction test (Fig. S4b) and the CO formation rate decreased drastically (Table 2, entry 6).

compared to the negatively charged surface (Table 2, entry 14), which would be due to the aggregation of the Ag nanoparticles. The control of the Ag species would be one of the key factors to realize the high photocatalytic performance.

4. Conclusions

In the present study, several kinds of Ag loaded $\text{Na}_2\text{Ti}_6\text{O}_{13}$ photocatalysts were prepared with the flux method and the photodeposition method, and the samples were examined for the photocatalytic CO_2 reduction by water in the aqueous solution of NaHCO_3 , which produced CO, H_2 and O_2 . The highest CO production rate in the current reaction condition was $0.89 \mu\text{mol h}^{-1}$ at the steady state (24 h later) over the Ag/NTO(NaCl ,50, -10K h^{-1}) sample, although this value was one third lower than a Ag/ CaTiO_3 photocatalyst of the same Ag content reported in our previous study [25]. It should be mentioned that the production rate of H_2 was much higher than the previous one, implying that the optimization of the loading amount, the structure of the Ag cocatalyst and so on, may contribute to the further improvement of the CO production rate.

The highest CO rate achieved with the $\text{Na}_2\text{Ti}_6\text{O}_{13}$ photocatalyst consisting of thick rod-like crystals, which was obtained by the flux method by using the NaCl flux with 50% of solute concentration in the molten mixture at high holding temperature (1273 K) and followed by the slow cooling rate such as -10 K h^{-1} . The large side facets such as (001) plane are the reductive facets, which can stabilize the Ag nanoparticles on them and supply the photoexcited electrons to the Ag cocatalyst for the CO production. The thick rod-like structure with suitable aspect ratio such as 3.2 would realize a good balance for both reductive and oxidative reactions, realizing high photocatalytic performance for CO production as a result.

On the other hand, the highest selectivity for the CO production in the present study was 60% when the photocatalytic reaction was carried out in the basic aqueous solution of Na_2CO_3 , where the highly dispersed Ag cocatalyst nanoparticles would be formed more. It is expected that further optimization of the reaction condition would also improve the photocatalytic activity for both the CO production rate and the CO selectivity. This will be examined in the next study.

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Appendix A. Supplementary data

Electronic supplementary information (ESI) is available on the web.

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Tables.

Table 1 The average sizes of the rod-like crystals in the NTO samples.

Entry	Sample	average size of crystals ^a /μm			aspect ratio	S _{BET} ^b /m ² g ⁻¹
		length	width			
<i>Various solute concentration, x</i>						
1 ^c	NTO(NaCl,5)	6.4	0.74	8.7		2.0
2 ^c	NTO(NaCl,30)	4.3	0.81	5.3		
3 ^c	NTO(NaCl,50)	4.1	1.1	3.9		1.8
4 ^c	NTO(NaCl,70)	2.4	0.91	2.6		
5	NTO(SS)	0.88	0.68	1.3		1.6
<i>Various hold temperatures</i>						
6 ^d	NTO(NaCl,5,973K)	1.2	0.049	25		
7	NTO(NaCl,5,1073K)	6.2	0.33	18		3.6
<i>Various cooling rate</i>						
8	NTO(NaCl,50,−10Kh ^{−1})	4.1	1.3	3.2		
9	NTO(NaCl,50,−5Kh ^{−1})	4.7	1.3	3.8		
<i>Using anatase TiO₂ as a start material</i>						
10	NTO(NaCl,50,anatase)	1.7	0.24	7.2		

^a The variation coefficient ranged between 28–62%. ^b BET specific surface area.

^c Samples were heated at 1273 K for 10 h, and cooled at a rate of −100 K h^{−1}. ^d the data for the rod-like crystals in the mixture.

Table 2 Results of the reaction tests for the photocatalytic reduction of CO₂ with H₂O with the Ag/NTO samples.

Entry	Sample ^a	Production rate (5 h later) / $\mu\text{mol h}^{-1}$		Production rate (24 h later) / $\mu\text{mol h}^{-1}$		$S_{CO}(\%)^b$
		CO	H ₂	CO	H ₂	
<i>Various fluxes</i>						
1	Ag/NTO(LiCl,50)	0.054	0.708	— ^c	— ^c	—
2	Ag/NTO(NaCl,50)	0.155	0.081	0.042	0.115	27
3	Ag/NTO(KCl,50)	0.109	0.298	0.031	0.089	26
4	Ag/NTO(CaCl ₂ ,50)	0.085	0.380	0.011	0.085	11
<i>Various solute concentration, x</i>						
5	Ag/NTO(NaCl,5)	0.088	0.184	0.019	0.246	7.1
6	Ag/NTO(NaCl,30)	0.126	0.155	0.030	0.079	28
7	Ag/NTO(NaCl,70)	0.079	0.097	0.030	0.103	22
8	Ag/NTO(SS)	0.022	0.042	0.011	0.026	30
<i>Various hold temperatures</i>						
9	Ag/NTO(NaCl,5,973K)	0.017	0.168	— ^c	— ^c	—
10	Ag/NTO(NaCl,5,1073K)	0.085	0.347	0.034	0.630	5.7
<i>Various cooling rate</i>						
11	Ag/NTO(NaCl,50,−10Kh ^{−1})	0.349	1.76	0.089	0.941	8.6
12	Ag/NTO(NaCl,50,−5Kh ^{−1})	0.279	2.30	0.070	3.14	2.2
<i>Others</i>						
13	Ag/NTO(NaCl,50,anatase)	0.086	0.636	0.017	0.214	7.4
14 ^d	Ag/NTO(NaCl,30)	0.140	0.539	0.059	0.040	60

^a Loading amount of Ag cocatalyst was 0.1 wt% and the used photocatalyst was 0.2 g. ^b Selectivity to CO after the reaction at 24 h later, see text. ^c Not measured due to very low production rate. ^d The photocatalytic reaction was carried out in an aqueous solution of Na₂CO₃ (pH=11) instead of the solution of NaHCO₃ (pH=8.3).

Figures.

Fig. 1 XRD patterns of the prepared $\text{Na}_2\text{Ti}_6\text{O}_{13}$ samples; the NTO(LiCl,50) sample (a), the NTO(NaCl,50) sample (b), the NTO(KCl,50) sample (c), the NTO(CaCl_2 ,50) sample (d), the NTO(SS) sample (e), a calculated one from a database (ICSD#182965) (f), and the rutile TiO_2 powder as the starting material (g). The NTO samples were prepared in the standard condition (hold temperature was 1273 K, the cooling rate was -100 K h^{-1}).

Fig. 2 SEM images of the prepared NTO samples; the NTO(LiCl,50) sample (a), the NTO(NaCl,50) sample (b), the NTO(KCl,50) sample (c), the NTO(CaCl₂,50) sample (d), and the NTO(SS) sample (e). The samples were prepared in the standard condition (hold temperature was 1273 K, the cooling rate was -100 K h^{-1}).

Fig. 3 SEM images of the prepared $\text{Na}_2\text{Ti}_6\text{O}_{13}$ samples by using NaCl flux with various concentrations of the solute in the molten mixture; the NTO(NaCl,5) sample (a), the NTO(NaCl,30) sample (b), and the NTO(NaCl,70) sample (c). The samples were prepared in the standard condition (hold temperature was 1273 K, the cooling rate was -100 K h^{-1}). See Fig. 2b for the NTO(NaCl,50) sample.

Fig. 4 TEM image and electron diffraction pattern (inset) of the NTO(NaCl,50) sample prepared by the flux method in the standard condition (hold temperature was 1273 K, the cooling rate was -100 K h^{-1}).

Fig. 5 DR UV-visible spectra of the prepared $\text{Na}_2\text{Ti}_6\text{O}_{13}$ samples; the NTO(NaCl,5) sample (a) and the NTO(NaCl,50) sample (b), and the NTO(SS) sample (c). The samples were prepared in the standard condition (hold temperature was 1273 K, the cooling rate was -100 K h^{-1}).

Fig. 6 SEM images of the NTO(NaCl,50,-100Kh⁻¹) sample (a), the NTO(NaCl,50,-10Kh⁻¹) sample (b), and the NTO(NaCl,50,-5Kh⁻¹) sample (c). Image (a) is shown again for comparison, which is the same as Fig. 2c.

Fig. 7 DR UV-visible spectra of the Ag/NTO(NaCl,5) sample (a), the Ag/NTO(NaCl,50) sample (b) and the Ag/NTO(NaCl,70) sample (c), and the Ag/NTO(SS) sample (d). The samples were prepared in the standard condition (hold temperature was 1273 K, the cooling rate was -100 K h⁻¹).

Fig. 8 (A) Variation of the photocatalytic production rate of CO with the solute concentration x in the preparation of the $\text{Ag}/\text{NTO}(\text{NaCl}, x)$ samples by the flux method in the standard condition, where $x=100$ means the $\text{Ag}/\text{NTO}(\text{SS})$ sample prepared by the SS method. (B) variation of the average size, length and width, of the rod-like crystals of the $\text{NTO}(\text{NaCl}, x)$ samples with the solute concentration x . Values are taken from Tables 1 and 2.

Fig. 9 Effect of the average width of the rod-like crystals or the particle size in the various $\text{Na}_2\text{Ti}_6\text{O}_{13}$ samples prepared by the flux method by using NaCl flux with various solute concentrations (open circles), with various cooling rates (open squares), and from anatase as the starting material (closed triangle), and prepared by the solid-state reaction method (closed circle) on the photocatalytic CO production rate over the corresponding 0.1 wt% $\text{Ag}/\text{Na}_2\text{Ti}_6\text{O}_{13}$ samples at 24 h later from the start of the reaction test. Values are taken from Tables 1 and 2.

Fig. 10 SEM images of the Ag/NTO(NaCl,5) sample (a, d), the Ag/NTO(NaCl,50) sample (b, e) and the Ag/NTO(SS) sample (c, f) before (a–c) and after (d–f) the use in the photocatalytic CO₂ reduction with water. The bright dots indicated by arrows are assignable for the Ag cocatalyst nanoparticles.

Fig. 11 Proposed mechanism for the growth of the Ag nanoparticles on the photocatalyst during the photocatalytic reaction.

Fig. 12 Photocatalytic reduction of CO_2 on the Ag nanoparticles stabilized on the neighboring reductive facets of the $\text{Na}_2\text{Ti}_6\text{O}_{13}$ photocatalyst crystals prepared by the flux method.