Ammonia photosynthesis via an association pathway using a plasmonic photoanode and a zirconium cathode

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Most conventional photoelectrochemical-based methods for synthesizing NH3 show low selectivity due to the generation of H2 as a by-product. In principle, two types of reaction mechanisms can occur in the reduction of N2 to NH3. One is an associative pathway in which N2 molecules on the catalyst are hydrogenated. The other is a dissociative pathway in which nitrogen and hydrogen react after the cleavage of the strong N2 triple bond. Understanding the mechanism of NH3 formation on the electrode will facilitate the development of selective and efficient NH3 synthesis techniques. In this study, we constructed a two-electrode system composed of a strontium titanate photocatalytic anode in which the plasmon effect is expressed by plasmonic gold nanoparticles and a zirconium cathode, which was connected to the external circuit to investigate the reaction by electrochemical analysis in addition to analysis of the product. The bias and pH dependences of the reaction were then systematically investigated, and the plasmon-induced synthesis of NH3 on Zr was proposed to proceed via an associative pathway.

Introduction

Ammonia (NH3) has attracted considerable attention as a hydrogen carrier and combustion fuel in addition to being a key precursor of synthetic fertilizer.1-3 NH3 is industrially produced by the Haber-Bosch process, which involves the heterogeneous reaction of N2 and H2 produced from fossil fuels under harsh conditions at high pressure (20 MPa) and temperature (400 °C).4-6 Although researchers have attempted to synthesize NH3 under milder conditions by improving the thermochemical catalyst,7, 8 the cleavage of the N2 triple bond still requires high energy. Therefore, another approach that produces NH3 with low energy consumption, such as an enzymatic method, is needed if NH3 is to be used as an energy carrier.

Previously, numerous efforts have been devoted towards developing electrochemical9-12 and photocatalytic13-17 methods for synthesizing NH3 under ambient conditions. However, most of these synthetic methods show low selectivity due to the formation of H2 as a by-product. Additionally, in particular, it is difficult that a semiconductor photocatalyst utilizes a broadband solar spectrum is difficult because of the band structure limitations. Plasmon-related photoreactions are expected to be a novel way to utilize the visible and near-infrared light in the solar spectrum because of the light harvesting properties.18-23 Recently, we reported that NH3 was selectively synthesized through the reduction of nitrogen using water as the electron source by using strontium titanate (SrTiO3) loaded with gold nanoparticles (Au-NPs) on one side and zirconium (Zr) on the other side.24, 25 The reaction was driven via plasmon-induced charge separation on the Au-NP-loaded SrTiO3 photoanode, and Zr served as the cathode and co-catalyst. The proposed mechanism of plasmon-induced water oxidation is described below.26 The hot-electron in the plasmonic metal nanoparticles is excited through the decay of the localized surface plasmon resonance (LSPR). A hole with a high oxidation ability remained at the surface states of the semiconductor near the metallic nanoparticle, and this hole has the potential to photocatalytically oxidize water.26, 27 However, the mechanism of nitrogen reduction, such as electron flow, that is active in NH3 production on the cathode is still unclear because the conventional plasmonic NH3 synthesis system, which integrates both the anode and cathode into one substrate, did not provide electrochemical information including the photocurrent flow and potential differences between the anode and cathode.

In this study, we constructed a two-electrode system connected to an external circuit to investigate the reaction by electrochemical analysis in addition to product analysis. We also investigated the effect of the bias and pH on the plasmon-induced NH3 synthesis on Zr.

Results and Discussion

Construction of plasmon-induced NH3 synthesis device

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For the measurement of evolved H₂, a sealed cell in which N₂ flow. The two-electrode system used a similar circuit with the SrTiO₃ electrode loaded with Au-NPs on one side and Zr on the other side except an ion exchange membrane served as an ion exchange film and N₂ was continuously bubbled through the cathodic chamber during the reaction. Although the cathodic chamber was not sealed, a contamination of ambient air was prevented by the continuous supply of N₂ gas.

**Photoelectrochemical measurements on plasmon-induced NH₃ synthesis**

Figure 2a depicts the current density and voltage (J-V) characteristics of the two-electrode system using a SrTiO₃ photoanode loaded with Au-NPs and a Zr coil cathode (Au-NPs/SrTiO₃|Zr coil). A good rectification property derived from the SrTiO₃ was observed in dark conditions. Additionally, a stable photocurrent was observed under visible light irradiation over a wide range of applied biases. The photocurrent is derived from hot-electron-induced charge separation due to LSPR excitation because the action spectrum of the incident photon-to-current efficiency (IPCE) corresponds to the plasmon resonance spectrum, as shown in Figure 2b, and the SrTiO₃ does not absorb visible light longer than 390 nm.

Figure 2c shows the amount of NH₃ evolved in the cathodic chamber as a function of irradiation time with visible light (410-810 nm) with an applied bias of 1.0 V. The amount of NH₃ formed with (c) the two-electrode system using a Au-NPs/SrTiO3|Zr coil. The red and black lines indicate the current under irradiation and dark conditions, respectively. The applied biases were 0 and 1.0 V for the IPCE and NH₃ measurements, respectively. The irradiation wavelength was 410-810 nm for the J-V and NH₃ measurements.

**Associative Pathway**

\[
\text{N}_2 \rightarrow \text{N}_2^* \rightarrow \text{H} + \text{N} + \text{H} \rightarrow \text{NH}_3
\]

**Dissociative Pathway**

\[
\text{N}_2 \rightarrow \text{N}_2^* \rightarrow \text{H} + \text{N} + \text{H} \rightarrow \text{NH}_3
\]

**Figure 3. Generic mechanisms for N₂ reduction to NH₃ on heterogeneous catalysts.**

In principle, two types of reaction mechanisms are possible in ammonia synthesis systems. The larger reaction rate of NH₃ formation was 10 nmol h⁻¹ cm⁻². This rate is approximately 10 times larger than the reported with plasmonic ammonia synthesis systems. The larger reaction rate of NH₃ photosynthesis was achieved due to the efficient ion transport path, the greater cathode surface area, and the continuous supply of N₂ gas.

**Bias and pH dependence of plasmon-induced NH₃ synthesis**

In principle, two types of reaction mechanisms are possible in the reduction of N₂ to NH₃, as shown in Figure 3. One is an associative pathway in which N₂ molecules on the catalyst are...
hydrogenated. The other is a dissociative pathway in which nitrogen and hydrogen react after the cleavage of the strong N₂-H₂ triple bond. Enzymes such as nitrogenase follow the associative pathway as the activation energy is lower. However, the Haber-Bosch method uses the dissociative pathway. Understanding the reaction mechanism of NH₃ formation on Zr will facilitate the development of selective and efficient NH₃ synthesis techniques. Therefore, the effects of the applied potential and concentration of reactants on the reaction are important. Figure 4a shows the faradaic efficiency (FE) of NH₃ formation calculated from the amount of NH₃ formed and the photocurrent as a function of the applied bias. Hydrazine (N₂H₄) was also measured by the colorimetric method because N₂H₄ is a possible intermediate in NH₃ production through the associative pathway.

When the applied bias is small, no NH₃ was detected. A small amount of NH₃ was observed with an applied bias of 0.7 V. The FE of NH₃ increased as the applied bias increased, reaching approximately 80% when the applied bias was 1.0 V. Additionally, N₂H₄ was observed when the applied bias was from 0.5 to 0.9 V. To understand the applied potential on the cathode in well-defined condition, the three-electrode electrochemical reaction was performed using Zr as the working electrode. The current-voltage characteristics of Zr under N₂ (h₂) and Ar (Iₐ) bubbling conditions and the current enhancement by N₂ calculated as (Iₐ/h₂-Iₐ)/Iₐ was larger than 1/0 over a wide range of applied potentials, and the current enhancement showed a maximum at ~0.3 V vs. Ag/AgCl of the applied bias.

Figure 4b depicts the FE of N₂H₄ as a function of pH under an applied bias of 1.0 V. It was confirmed that the pH was not increased during the reaction (Figure S3). The FE of NH₃ was very poor when the pH was larger than 4.0. However, the FE of N₂H₄ increased dramatically as the pH decreased below 4. In contrast, N₂H₄ was observed when the pH was between 3.5 and 4.0. This result indicates that the proton concentration directly affects NH₃ production.

Notably, both the bias and pH dependencies indicate that intermediate values of bias and pH produce N₂H₄. These results suggest that reaction acceleration is needed for NH₃ production, and N₂H₄ may be an intermediate in NH₃ production. However, the reduction potential from N₂ to N₂H₄ (~0.33 V vs. the reversible hydrogen electrode (RHE)) is negative relative to that from N₂ to NH₃ (0.09 V vs. RHE). Therefore, plasmon-induced NH₃ formation via a six-electron and six-proton transfer process on Zr proceeded kinetically rather than thermodynamically. Also, there is a possibility that the produced N₂H₄ was consumed as a redox reagent due to the high reduction ability. Additionally, the sum of the FE of NH₃ and N₂H₄ does not equal 100% in the intermediate region. One of the other electron-consuming processes is the self-reduction of the Zr surface, and the other process is H₂ production via proton reduction. H₂ production was evaluated by using a sealed reaction cell in which N₂ was encapsulated in the chamber. As a result, approximately 50% of the photocurrent was used for proton reduction under the applied bias of 0.7 V (Figure S1b). Also, the FE of H₂ was less than detection limit under the applied bias of 1.0 V.

**Proposed reaction mechanism**

In previous reports using isotopic reactants, water oxidation on the plasmonic photoanode and NH₃ evolution by N₂ reduction on the Zr cathode proceeded via plasmon-induced NH₃ synthesis. In particular, it was predicted that N₂ is reduced by proton addition in the aqueous solution rather than the addition of a hydrogen adatom.
Based on the results described above, we propose the plasmon-induced NH₃ synthesis on Zr proceeds via an associative pathway in which N₂ is hydrogenated by protons (Figure 5). Initially, the N₂ molecule is preferentially adsorbed on the Zr surface (step i). Subsequently, the reduction of N₂ proceeds via proton addition (step ii) because the hypothesis that protons attach to the adsorbed N₂ is supported by the dependence of the NH₃ production on the proton concentration, as shown in Figure 3b. When the pH was large and applied potential was low, H₂ was also produced as a by-product (step ii', iii'). Additionally, when the applied potential and proton concentration are moderate (pH 3.5-4.0 at applied bias of 1.0 V, or applied bias 0.5-0.9 V at pH 3.0), N₂H₄ is produced as the major product via a four-electron transfer process (step iii-iv). If the reaction rate is high enough with a large applied bias and a high proton concentration, NH₃ is produced as the main product via a six-electron transfer process (step v). We conclude that the reason for the high selectivity in N₂ conversion on Zr is due to the association process without direct cleavage of the strong N₂ triple bond.

Experimental

Preparation of plasmonic photoanode

Au-NPs/SrTiO₃ photoanode was fabricated according to the previous reports. The Au-NPs/SrTiO₃ photoelectrode was fabricated by Helicon sputtering (ULVAC, MPS-4000C1/HC1) of 3 nm Au film on a single crystal of 0.05wt% Nb-doped SrTiO₃ (10 × 10 × 0.5 mm³, Furuuchi Chemical) with a (110) surface, and subsequent annealing at 800°C in N₂. The morphology of Au-NPs on the SrTiO₃ was observed using field-emission scanning electron microscopy (FE-SEM, JSM-6700FT, JEOL); the maximum resolution attainable at an electron acceleration voltage of 15 kV was 1 nm. The cross-sectional structure of the Au-NPs/SrTiO₃ was observed by STEM (Hitachi HD-2000 operated at 200 kV). The cross-sectional STEM sample was prepared using the focused ion beam technique (JEOL JIB-4600F/HKD) with Ga ions.

Photoelectrochemical reaction

The N₂ fixation device comprised reaction cells with two reaction chambers separated by the Nafion® 117 proton exchange membrane (Aldrich). An InGa alloy (2:1 weight ratio) film was pasted onto the backside of the plasmonic photoanode and then connected to an electrochemical analyzer (ALS/CH Instruments 760DH (ALS)) as a photoanode with a copper lead wire. A Zr coil was used as a cathode. The surface area of the Zr coil was 0.628 cm². An H₂SO₄ aqueous solution (pH3) was filled in both chambers as the supporting electrolyte solution without any specific electron donor, and the N₂ was bubbled in the cathodic chamber with 50 sccm during the reaction. When the pH of the solution was less than 4, the concentration of the electrolyte was kept at 0.05 mol dm⁻³ by the addition of Na₂SO₄. The photoanode was irradiated in an area of φ6 mm (0.283 cm²) by a xenon lamp (800 W, Newport) using an arbitrary light intensity and wavelength, and the amount of NH₃ formed per area was calculated by dividing the value by the irradiation area. For the IPCE calculation, the photocurrent was measured at 0 V between two-electrode. The IPCE was calculated by the following equation:

\[
\text{IPCE} = \frac{1240 \cdot J(A/cm^2)}{\lambda (nm) \cdot P(W/cm^2)} \times 100\%
\]

where J is the photocurrent density generated by monochromatic light with a bandwidth of less than 15 nm, λ is the wavelength of the monochromatic incident light, and P is the light intensity. Bandpass filters with a bandwidth of less than 15 nm at the full-width at half-maximum were used.

The faradaic constant, 96485 C/mol, n is the number of electrons involved in the reaction, Q is the total charge passed through the whole reaction.

Assays

The quantity of NH₃ formed and evolved H₂ were determined using the procedure using sodium salicylate, pyrazole, sodium hypochlorite, and sodium hydroxide. We passed 0.5 mL of deionized water through the cathodic chamber to fully dissolve the NH₄Cl (aqueous solution) formed. After the solution was neutralized to pH 7 using 1.5 mol dm⁻³ sodium hydroxide aqueous solution, the following materials were added successively: 0.08 mL of an aqueous solution of ethylenediaminetetraacetic acid tetrasodium salt hydrate (1.3 mol dm⁻³); 0.16 mL of an aqueous solution of sodium salicylate (1.46 mol dm⁻³) and pyrazole (0.24 mol dm⁻³); 0.52 mL of an aqueous solution of sodium hydroxide (1.25 mol dm⁻³) and sodium hypochlorite (0.25 mol dm⁻³). The sodium salicylate formed a dimer in the presence of NH₃ with absorption at λmax=650 nm. Ammonium chloride was purchased (Wako chemical, LTD) and used for calibration. The absorption spectra were monitored using a UV/vis spectrometer (UV-3100, Shimadzu). The value for NH₃ formation at 0 hours was subtracted from all of the data as the background.

The quantity of formed N₂H₄ was determined by a colorimetric method using organic probe 3-cyano-7-hydroxycoumarin levulinate (CHCL). CHCL was synthesized according to the literature as shown in Scheme S1. Oxalyl chloride (0.82 mL, 8.6 mmol) and DMF (10 μL) were added to a suspension of levulinic acid (502 mg, 4.3 mmol) in dichloromethane (50 mL). The reaction mixture was stirred at room temperature for 4 h, and the volatiles were evaporated under reduced pressure and subsequently dried with vacuum pumping. The residue was dissolved in a small amount of dry dichloromethane. The solution was slowly added to the dispersed dichloromethane solution (50 mL) containing 3-cyano-7-hydroxycoumarin (CHC, 284.6 mg, 1.5 mmol) and triethylamine (0.55 mL, 3.9 mmol) at 0°C. After stirring for 12 h
The utilization of NH₃ as an energy carrier.

The chromogenic signalling behaviour of CHCL was investigated in a 30% aqueous acetonitrile solution at pH 6.8 (phosphate buffer, 10 mM). CHCL revealed moderate UV-vis absorption at 336 nm. Upon the interaction of CHCL with hydrazine, a prominent absorption band centered at 416 nm developed which is a characteristic of CHC. The changes in absorption bands by the hydrazine-induced deprotection process were used as a ratiometric analysis for the transformation of probe CHCL to CHC.

The calibration curves for NH₃ and N₂H₄ were shown in Figure S4.

Conclusions

In summary, a two-electrode plasmon-induced NH₃ synthesis system that enables electrical and chemical analysis was fabricated. The bias and pH dependences of the reaction were systematically investigated, and the mechanism of the plasmon-induced NH₃ synthesis on Zr was proposed follow an associative pathway. The results presented herein provide crucial insight into the design of catalysts and electrodes for plasmon-induced NH₃ synthesis. As an aspect of the plasmonic photoanode, higher absorptivity and charge separation efficiency are required to produce a larger number of hot-electrons.

Future studies are aimed at increasing the reaction yield while maintaining high selectivity. The innovative development of the plasmonic photoanode, such as the coupling between plasmons and the other photonic mode, offers enhanced charge separation efficiency. Time-resolved measurements of the near and far fields are also important for understanding the plasmon decay and the generation and transportation of the photocarriers. Additionally, the development of oxidation co-catalysts is indispensable for quickly consuming the photogenerated holes. The development of high-performance cathodes and co-catalysts for nitrogen reduction is also indispensable. Although further improvement of the reaction activity is needed, this robust plasmonic NH₃ synthesis system under visible light irradiation could facilitate the utilization of NH₃ as an energy carrier.

Conflicts of interest

The authors declare no conflict of interest.

Acknowledgements

We acknowledge financial support from JSPS KAKENHI (Grant Nos. JP18H05205, JP17H01041, JP19H02737, JP19H04667, and JP18K05053), the Nanotechnology Platform (Hokkaido University), and the Dynamic Alliance for Open Innovation Bridging Human, Environment and Materials (Five-Star Alliance) of MEXT.

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