

Development of a Highly Efficient Hydrogen Generation System by Plasmon-Induced Charge Separation Using Sunlight as an Energy Source

太陽光をエネルギー源としたプラズモン誘起電荷分離による高効率水素発生システムの開発

TAKAHASHI Yukina

高橋 幸奈

Although the total amount of sunlight is large, the energy density per unit area is small and the supply is unstable. Therefore, it is important in practical use to devise some means to store and increase the energy density. The localized surface plasmon resonance (LSPR) of metal nanoparticles increases the density of solar energy, and the phenomenon of plasmon-induced charge separation (PICS), which occurs when metal nanoparticles are combined with semiconductors, can be used to convert light energy into electrochemical energy, which can be used to generate hydrogen from solar energy. This paper describes a research plan to develop a highly efficient hydrogen generation system using solar energy.

太陽光は総量が大きいですが、単位面積あたりのエネルギー密度は小さく、供給量が不安定であるため、何らかの手段で貯めたり、エネルギー密度を高める工夫を施すことが、実用において重要である。金属ナノ粒子の局在表面プラズモン共鳴 (localized surface plasmon resonance: LSPR) によって太陽光エネルギーの密度を高めるとともに、金属ナノ粒子と半導体とを組み合わせた時に生じるプラズモン誘起電荷分離(plasmon-induced charge separation: PICS)という現象を利用して光のエネルギーを電気化学エネルギーに変換することで、太陽光による高効率な水素発生システムの開発を目指す研究計画について述べる。

Introduction

A stable supply of renewable energy that does not rely on fossil resources such as oil and coal is essential to the realization of a sustainable society. Among the energy sources such as solar, wind, hydro, geothermal, tidal, and biomass, which can always be supplied from nature, solar is a particularly promising energy source. This is because the energy is obtained from outside the closed system of the earth, where the law of conservation of energy holds. The sun generates enormous amounts of energy through nuclear fusion. Only a small fraction of that energy reaches the Earth's surface. The energy reaching the surface of the earth and the sea is approximately $3 \times 10^{24} \text{ J y}^{-1}$.^[1] This value is approximately 5000 times as large as world energy consumption.^[2] However, although an enormous amount of light energy falls on the earth in total, it has the disadvantage of having a small energy density per unit area. In addition, since it is dependent on nature, the supply is unstable because it is greatly affected by the

seasons and weather, with the exception of some regions such as the Sahara Desert. In other words, it is an important issue in the implementation of the system to store a large amount of energy when it is supplied in large quantities, to downsize the system to produce only the necessary amount in small quantities when needed on site, and to devise ways to increase the energy density. Among them, we are focusing on ways to increase energy density.

Problems with conventional PICS system

However, conventional PICS has several problems due to the use of n-type semiconductors. One is the low stability of metal nanoparticles. Since oxidation reactions occur on the metal nanoparticles, it may cause the metal nanoparticles themselves to dissolve, even silver nanoparticles, which are known to be relatively stable. We have been studied to solve the problem. We succeeded in improving the thermal stability of spherical silver nanoparticles, by introducing an Al_2O_3 nanomask as a template of the nanoparticles, and the chemical stabilities by coating with a thin and dense titanium dioxide film.^[14] In same way, by coating with titanium dioxide film, we improved stabilities of gold nanorods,^[15] and silver nanoplates,^[16] which are effective for the use of near-infrared light due to their shape anisotropy but are easily deformed into a spherical shape. However, the use of cheaper and less unstable metals was desired.

Second is the low efficiency of the charge separation. The photoelectric conversion efficiency (η) of the conventional PICS was approximately 1%.^[10] In titanium dioxide photocatalysts, metal nanoparticles supported on the photocatalyst function as an electron pool that suppresses recombination of excited electrons and holes, as evidenced by their use as reduction sites.^[17] Therefore, electrons transferred to the semiconductor after charge separation by conventional PICS are likely to undergo reverse electron transfer to the metal nanoparticles (recombination in the broad sense). It was supported by the reports that charge separation lifetime improved from picoseconds to microseconds by changing the semiconductor combined with the metal nanoparticles from n-type to p-type.^[18]

Highly efficient hydrogen generation via PICS system with a p-type semiconductor

We therefore propose that the above problem can be solved by using a p-type semiconductor, which reverses the charge transfer, because the reduction reaction can proceed on the metal nanoparticles used in PICS (Figure 3).^[19] The catalytic effect of the metal nanoparticles in the reduction reaction can also be expected. As a result, the selectivity of the reaction products can be improved. For example, copper is known to be able to reduce carbon dioxide to C2 compounds such as ethanol and ethylene when used as an electrode in electrochemical reactions.^[20] Therefore, it is expected that by controlling the type of metal and crystal planes that make up the nanoparticles, arbitrary chemical reactions can be promoted, such as the generation of hydrogen and the generation of specific hydrocarbons through the reduction of carbon dioxide. In same way, as the p-type

semiconductor, for example, iridium oxide is a proven material as an electrode for the electrochemical oxidation of water,^[21] and is expected to water splitting into oxygen and hydrogen. The proposed system is theoretically expected to be dramatically more efficient than conventional systems.

It was noted above that conventional PICS system can be applied to sensing devices. The proposed system can also work as a sensor with higher sensitivity than conventional one. It may be possible to simultaneously detect the hydrogen generation reaction as an electrical signal in the novel system. It will be able to detect not only hydrogen generation reactions, but also the progress of beneficial chemical reactions. This sensing system will be useful in a variety of systems in the future.

Conclusion

We have described the possibility of a novel, highly efficient hydrogen generation system using PICS with a p-type semiconductor. The proposed novel system is expected to realize high-sensitivity sensing at the same time. We believe that the realization of this system, which is expected to be able to effectively harvest and utilize energy of sunlight, will contribute to solving energy problems of the world.

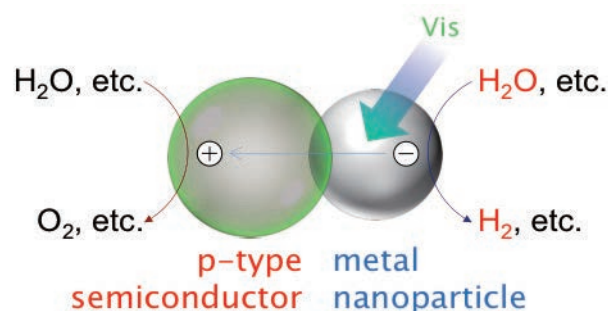


Figure 3 Schematic illustration of the PICS with p-type semiconductor.

References

- [1] Solar Energy Perspectives, OECD/IEA, 2011.
- [2] bp Statistical Review of World Energy 2020.
- [3] M. Fleischman, P. J. Hendra, A. J. McQuillan, *Chem. Phys. Lett.*, **26**, 163 (1974).
- [4] C. Wen, K. Ishikawa, M. Kishima, K. Yamada, *Sol. Energy Mater. Sol. Cells*, **61**, 339 (2000).
- [5] S. D. Standridge, G. C. Schats, T. Hupp, *J. Am. Chem. Soc.*, **131**, 8407 (2009).
- [6] K. R. Catchpole, S. Pillai, *J. Luminescence*, **121**, 315 (2006).
- [7] D. Derkacs, S. H. Lim, P. Matheu, W. Mar, E. T. Yu, *Appl. Phys. Lett.*, **89**, 093103 (2006).
- [8] K. Nakayama, K. Tanabe, H. A. Atwater, *Appl. Phys. Lett.*, **93**, 121904 (2008).
- [9] P. Anger, P. Bharadwaj, L. Novotny, *Phys. Rev. Lett.*, **96**, 113002 (2006).
- [10] Y. Tian, T. Tatsuma, *J. Am. Chem. Soc.*, **127**, 7632 (2005).
- [11] T. Tatsuma, Y. Katagiri, S. Watanabe, K. Akiyoshi, T. Kawawaki, H. Nishi, E. Kazuma, *Chem. Commun.*, **51**, 6100 (2015).
- [12] Y. Takahashi, T. Tatsuma, *Appl. Phys. Lett.*, **99**, 182110 (2011).
- [13] E. Kowalska, R. Abe, B. Ohtani, *Chem. Commun.*, **241** (2009).
- [14] Y. Takahashi, T. Tatsuma, *Nanoscale*, **2**, 1494 (2010).
- [15] Y. Takahashi, N. Miyahara, S. Yamada, *Anal. Sci.*, **29**, 101 (2013).
- [16] Y. Takahashi, K. Suga, T. Ishida, S. Yamada, *Anal. Sci.*, **32**, 275 (2016).
- [17] W. Kubo, T. Tatsuma, *J. Mater. Chem.*, **15**, 3104 (2005).
- [18] Z. Lian, M. Sakamoto, H. Matsunaga, J. J. M. Vequizo, A. Yamakata, M. Haruta, H. Kurata, W. Ota, T. Sato, T. Teranishi, *Nat. Commun.*, **9**, 2314 (2018).
- [19] Y. Takahashi, Y. Yamadori, T. Murayama, S. Shingo, S. Yamada, in preparation.
- [20] Y. Hori, R. Takahashi, Y. Yoshinami, A. Murata, *J. Phys. Chem. B*, **101**, 7075 (1997).
- [21] Y. Takahashi, T. Tatsuma, *Electrochemistry*, **82**, 749 (2014).



Dr. TAKAHASHI Yukina

高橋 幸奈

Associate Professor,
International Institute for Carbon-Neutral Energy
Research (I²CNER),
Kyushu University