Z-scheme photocatalyst systems employing Rh- and Ir-doped metal oxide materials for water splitting under visible light irradiation†

Akihiko Kudo,†a Shunya Yoshino,a Taichi Tsuchiya,a Yuhei Udagawa,a Yukihiro Takahashi,a Masaharu Yamaguchi,a Ikue Ogasawara,a Hiroe Matsumotoa and Akihide Iwaseab

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Various types of Z-scheme systems for water splitting under visible light irradiation were successfully developed by employing Rh- and Ir-doped metal oxide powdered materials with relatively narrow energy gaps (EG): BaTa2O6:Ir,La (EG: 1.9–2.0 eV), NaTaO3:Ir,La (EG: 2.1–2.3 eV), SrTiO3:Ir (EG: 1.6–1.8 eV), NaNbO3:Rh,Ba (EG: 2.5 eV) and TiO2:Rh,Sb (EG: 2.1 eV), with conventional SrTiO3:Rh (an H2-evolving photocatalyst) or BiVO4 (an O2-evolving photocatalyst), and suitable electron mediators. The Z-scheme systems were classified into three groups depending on the combination of H2- and O2-evolving photocatalysts and electron mediator. The Z-scheme systems combining BaTa2O6:Ir,La with BiVO4, and NaTaO3:Ir,La with BiVO4 were active when a [Co(bpy)3]3+/2+ redox couple was used rather than an Fe3+/2+ one. The combination of SrTiO3:Ir with SrTiO3:Rh gave an activity when the [Co(bpy)3]3+/2+ and Fe3+/2+ redox couple ionic mediators were used. The Z-scheme systems combining NaNbO3:Rh,Ba and TiO2:Rh,Sb with SrTiO3:Rh showed activities by using the [Co(bpy)3]3+/2+ and Fe3+/2+ redox couples and also via interparticle electron transfer by just contact with/without reduced graphene oxide (RGO). These suitable combinations can be explained based on the impurity levels of doped Rh3+ and Ir3+ toward the redox potentials of the ionic mediators for the Z-scheme systems employing ionic mediators, and p-/n-type and onset potentials of the photocurrent in the photoelectrochemical properties of those photocatalyst materials for the Z-scheme systems working via interparticle electron transfer.

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Introduction

Artificial photosynthesis has attracted attention from the view point of solar energy conversion to storable chemical energy. Because solar water splitting is one of the representative reactions, photocatalytic water splitting has extensively been studied. \(^1\)–\(^7\) Powder-based metal oxide materials are attractive for photocatalytic water splitting because the cost will be low \(^8\)–\(^10\) and stability will be high compared with other materials such as chalcogenides. Although the efficiency of powder-based photocatalyst systems is behind that of systems of photovoltaic + electrolysis at the present stage, there are some advantages to powder-based photocatalyst systems.

It is crucial to demonstrate a solar water splitting system employing powder-based oxide photocatalysts. The present stage of research of artificial synthesis is along this topic. A reactor system including a gas separation system of evolved hydrogen from oxygen has been studied in addition to the development of photocatalyst materials aimed towards the practical use of solar water splitting. \(^10\) The separation system can be achieved by the use of a suitable separation membrane. Moreover, the safety issue of the co-evolved H\(_2\) and O\(_2\) has also been examined via the use of a suitable gas transportation tube. However, even if an excellent reactor system with a gas separation membrane system is established, which photocatalyst is employed for it is still a key issue as high efficient photocatalysts for real solar water splitting into H\(_2\) and O\(_2\) without any sacrificial electron donors and acceptors have not yet been developed. It is essential to develop an efficient photocatalyst for demonstrating a solar water splitting system taking gas separation and safety issues into account.

High efficiency of a photocatalyst can be brought about by a high quantum yield and a response to light with a long wavelength. From this viewpoint, it is vital to develop photocatalysts that can utilize up to 600–700 nm of the solar spectrum. Although the use of noble metals might prevent the photocatalyst materials from practical use, the usage would be allowed if the amount is small and the materials can be recycled.

There are single particulate and Z-scheme systems in powder-based photocatalysts. \(^2\)–\(^11\) Z-scheme photocatalyst systems have the advantage that photocatalysts active for either H\(_2\) or O\(_2\) evolution can be employed. This means that a Z-scheme system can be constructed with various combinations of H\(_2\)- and O\(_2\)-evolving photocatalysts. \(^12\)–\(^45\) SrTiO\(_3\):Rh (an H\(_2\)-evolving photocatalyst) and BiVO\(_4\) (an O\(_2\)-evolving photocatalyst) are representative metal oxide materials for the Z-scheme system. \(^1\)–\(^2\),\(^16\) It should be stressed that photocatalyst sheets prepared by a particle transfer method using SrTiO\(_3\):Rh,La and BiVO\(_4\):Mo powders demonstrate a quantum efficiency of 30% at 420 nm and a solar to hydrogen energy conversion efficiency of 1%. \(^17\)–\(^19\) This result suggests that the Z-scheme system is a promising photocatalyst system for practical solar water splitting. H\(_2\) evolution separated from O\(_2\) evolution is possible in the Z-scheme system if a suitable reactor is designed. \(^8\) However, this photocatalyst sheet consisting of the SrTiO\(_3\):Rh,La and BiVO\(_4\):Mo powders responds up to 520 nm because the energy gap (EG) of SrTiO\(_3\):Rh,La and the band gap (BG) of BiVO\(_4\):Mo are 2.3 eV and 2.4 eV, respectively. So, it is a key issue to develop Z-scheme photocatalyst systems consisting of metal oxide photocatalysts with a response at longer wavelengths than 520 nm.
Transition metal doping is one strategy to make wide band gap photocatalysts responsive to visible light.\textsuperscript{1,2} Rh and Ir are effective dopants in this strategy. We have reported that SrTiO$_3$:Rh (EG: 2.3 eV),\textsuperscript{20} SrTiO$_3$:Rh,Sb (EG: 2.2–2.4 eV),\textsuperscript{21} SrTiO$_3$:Ir (EG: 1.6–1.8 eV),\textsuperscript{20,22} BaTa$_2$O$_6$:Ir,La (EG: 1.9–2.0 eV),\textsuperscript{23} and NaTaO$_3$:Ir,La (EG: 2.1–2.3 eV)\textsuperscript{24} are active for sacrificial H$_2$ evolution in the presence of electron donors such as methanol, while SrTiO$_3$:Rh,Sb,\textsuperscript{21} SrTiO$_3$:Ir\textsuperscript{20} and TiO$_2$:Rh,Sb (EG: 2.1 eV)\textsuperscript{25} are active for sacrificial O$_2$ evolution in the presence of electron acceptors such as Ag$^+$. In these photocatalyst materials, the visible light responses are due to electronic transition from the impurity levels formed by the dopants to the conduction bands of the host materials. These energy and band gaps are close to, or narrower than, those of SrTiO$_3$:Rh (EG: 2.3 eV)\textsuperscript{20} and BiVO$_4$ (BG: 2.4 eV).\textsuperscript{26,27} So, it is attractive to utilize these Rh- and Ir-doped metal oxide photocatalysts for the construction of Z-scheme systems.

There are several types of Z-scheme system employing different electron mediators such as Fe$^{3+/2+}$ and [Co(bpy)$_3$]$^{3+/2+}$ redox couples,\textsuperscript{16,28,29} and reduced graphene oxide (RGO)\textsuperscript{30–32} as shown in Fig. 1(a) and (b). Moreover, some Z-scheme systems work even without electron mediators, as shown in Fig. 1(c).\textsuperscript{33,34} In this case, electron transfer proceeds through an interface contacted between the particles of the H$_2$- and O$_2$-evolving photocatalysts. So, it is important to examine the electron mediator in the Z-scheme system.

In the present paper, sacrificial H$_2$ and O$_2$ evolutions over Rh- and Ir-doped metal oxide photocatalysts under visible light irradiation were examined first to see the relationship between the photocatalytic properties and their band structures. Then, these H$_2$- and O$_2$-evolving photocatalysts were employed for various types of Z-scheme system for water splitting into H$_2$ and O$_2$ in stoichiometric amounts under visible light irradiation, without any sacrificial reagents. The photocatalytic performances for the sacrificial H$_2$ and O$_2$ evolutions and the Z-schematic water splitting were discussed based on the band structures and

![Fig. 1 Various types of powder material-based Z-scheme systems for water splitting.](image-url)
photoelectrochemical properties of the photocatalyst materials such as p-/n-types and the onset potentials of photocurrents.

**Experimental**

**Preparation of Rh- or Ir-doped metal oxide photocatalysts**

SrTiO$_3$:Rh(1%),$^{29}$ SrTiO$_3$:Ir(0.2%),$^{22}$ TiO$_2$:Rh($x\%$),Sb($2x\%$) ($x = 0.5$ or $1.3$),$^{25}$ NaTaO$_3$:Ir(1%),La(2%),$^{24}$ BaTa$_2$O$_6$:Ir(1%),La(2%)$^{23}$ and BiVO$_4$,$^{26,27}$ were prepared by a solid-state reaction, a borate-flux method, and a liquid–solid reaction according to previous reports. In addition to them, NaNbO$_3$:Rh($x\%$),Ba($y\%$) ($x, y = (1.2, 1.44$) or $(1.0, 2.0$) was newly prepared by a solid-state reaction. The starting materials, Na$_2$CO$_3$ (Kanto Chemical; 99.5 or 99.8%), Nb$_2$O$_5$ (Kanto Chemical; 99.99% or Kojundo Chemical; 99.99%), Rh$_2$O$_3$ (Wako Chemical; 98%), and BaCO$_3$ (Kanto Chemical; 99%), were mixed at a molar ratio of Na/Nb/Rh/Ba = $1.05 - 1.05y : 1 - x : x : y$. An excess of sodium was added in the starting materials to compensate for volatilization. The starting materials mixture was calcined at 1173 K for 1 h, and then 1423–1473 K for 10 h once or twice. The excess sodium was washed out with water after the calcination. The obtained powders had nonspecific shapes with aggregations, judging from the SEM images (Jeol; JSM-6700F) (Fig. S1†). The obtained samples were identified using X-ray diffraction (Rigaku; MiniFlex, Cu K$_a$). Diffuse reflectance spectra were obtained by a UV-vis-NIR spectrometer (JASCO, V-570) equipped with an integrator sphere and were converted to absorbance measurements via the Kubelka–Munk method.

**Preparation of an RGO–metal oxide composite**

An RGO-incorporated O$_2$-evolving photocatalyst was prepared by photocatalytic reduction of graphene oxide (GO) on the photocatalyst according to previous reports.$^{30–32}$ GO prepared by the Hummers’ method$^{35}$ and the O$_2$-evolving photocatalyst were dispersed in an aqueous methanol (Kanto Chemical; 99.8%) solution (50 vol%). The suspension was irradiated with visible light from a 300 W Xe lamp (PerkinElmer; CERMAX PE300BF) with a long pass filter (HOYA; L42) under a N$_2$ atmosphere with a pressure of 1 atm to obtain the RGO–photocatalyst composite. The methanol was carefully removed by washing with water. The RGO–photocatalyst composite was collected by filtration and was dried at room temperature in air.

**Sacificial H$_2$ and O$_2$ evolutions (half reactions of water splitting)**

H$_2$ and O$_2$ evolutions from aqueous solutions containing the sacrificial reagents CH$_3$OH (Kanto Chemical; 99.8%) and AgNO$_3$ (Kojima Chemical; 99.9% or Toyo Chemical; 99.9%) that were half reactions of water splitting were examined using a top-irradiation reaction cell with a Pyrex window and a 300 W Xe lamp (PerkinElmer; CERMAX PE300BF). NaTaO$_3$:Ir,La, BaTa$_2$O$_6$:Ir,La, NaNbO$_3$:Rh,Ba, and TiO$_2$:Rh,Sb were used as prepared, whereas SrTiO$_3$:Ir without a cocatalyst was reduced under 1 atm of H$_2$ at 473 K for 2 h as a pretreatment for sacrificial O$_2$ evolution. The photocatalyst powders (0.1–0.3 g) were suspended in aqueous solutions (120–150 mL) and irradiated with visible light. For the H$_2$ evolution, Pt (0.3 wt%) cocatalyst, working as an H$_2$ evolution site, was loaded on photocatalysts by photodeposition from an aqueous methanol solution containing
H$_2$PtCl$_6$ (Tanaka Kikinzoku; 37.55% as Pt). The wavelength of the irradiation light was controlled to visible light using long-pass filters (HOYA; L42 and Y44). The amounts of evolved H$_2$ and O$_2$ were determined using an online gas chromatograph (Shimadzu; GC-8A, MS-5A column, TCD, Ar carrier).

**Z-schematic water splitting**

Z-schematic water splitting was conducted using a gas-closed system with a top-irradiation cell with a Pyrex window. H$_2$-evolving photocatalyst and O$_2$-evolving photocatalyst powders (0.05 or 0.1 g, respectively) were suspended in 120 mL of water. For the interparticle Z-scheme systems without an electron mediator and with an RGO solid-state electron mediator, water not containing any ionic mediators was used. For the Z-scheme system with ionic mediator, an aqueous solution containing [Co(bpy)$_3$]SO$_4$ or FeCl$_3$ as a mediator was used. The pH was adjusted with H$_2$SO$_4$ in each of the Z-scheme systems with and without electron mediator, if necessary. Ru (0.7 wt%) cocatalyst, functioning as an H$_2$ evolution site, was loaded on SrTiO$_3$:Rh (an H$_2$-evolving photocatalyst) by photodeposition from an aqueous methanol solution containing RuCl$_3$·nH$_2$O (Tanaka Kikinzoku; 36% as Ru in RuCl$_3$·nH$_2$O). Pt (0.3–1 wt%) was loaded on the NaTaO$_3$:Ir,La and BaTa$_2$O$_6$:Ir,La (H$_2$-evolving photocatalysts), and SrTiO$_3$:Ir (an O$_2$-evolving photocatalyst) by an impregnation method. The photocatalyst powders and an aqueous H$_2$PtCl$_6$ solution were placed in a porcelain crucible and dried on a hot plate. The H$_2$PtCl$_6$-impregnated NaTaO$_3$:Ir,La and BaTa$_2$O$_6$:Ir,La powders were calcined at 673 K for 2 h in air, whereas the SrTiO$_3$:Ir was not. The Pt-loaded NaTaO$_3$:Ir,La and Pt-loaded BaTa$_2$O$_6$:Ir,La were subsequently reduced at 673 K for 1 h under 1 atm of H$_2$ as a pretreatment, while Pt-loaded SrTiO$_3$:Ir was reduced at 573 K for 1 h. The light source and GC setup were the same as those for the sacrificial H$_2$ and O$_2$ evolutions.

**Photoelectrochemical measurements**

A squeegee method was used to prepare the SrTiO$_3$:Rh(1%) photoelectrode and a drop-casting method was used for the NaTaO$_3$:Ir(1%),La(2%) and BaTa$_2$O$_6$:Ir(1%),La(2%) with and without H$_2$-reduction; SrTiO$_3$:Ir(0.2%) with H$_2$-reduction; NaNbO$_3$:Rh(1%),Ba(2%) and TiO$_2$:Rh(0.5%),Sb(1%) without H$_2$-reduction, and BiVO$_4$ photoelectrodes using powdered photocatalyst materials. For the SrTiO$_3$:Rh(1%) photoelectrode, a paste consisting of 20 mg of SrTiO$_3$:Rh(1%) photocatalyst powder, 20 µL of acetylacetone (Kanto Chemical; 99.5%) and 40 µL of distilled water was coated on an indium tin oxide transparent electrode (ITO). For the other photoelectrodes, the photocatalyst powders were dispersed in ethanol (1–2 mg mL$^{-1}$) by sonication. The suspensions were drop-cast onto a fluorine-doped tin oxide transparent electrode (FTO) to obtain 1–2 mg cm$^{-2}$ of photocatalyst on the FTO. The H$_2$-reduced SrTiO$_3$:Ir-loaded FTO substrate was not calcined, whereas the other photocatalyst-loaded ITO and FTO substrates were calcined at 573–673 K for 2 h in air. The photoelectrochemical properties were evaluated with a three-electrode system consisting of working, Ag/AgCl reference, and Pt counter electrodes with a potentiostat (Hokuto Denko; HZ-series or HSV-110) using a conventional H-type cell with a Nafion membrane. The electrolyte was 0.1 mol L$^{-1}$ K$_2$SO$_4$, 0.025 mol L$^{-1}$ KH$_2$PO$_4$ + 0.025 mol L$^{-1}$ Na$_2$HPO$_4$ pH buffer.
was added, if necessary. A 300 W Xe lamp (PerkinElmer; CERMAX PE300BF) with a long-pass filter (HOYA; L42) was employed as a light source.

**Results**

**Photocatalytic activities for sacrificial H₂ and O₂ evolutions over Rh- and Ir-doped metal oxide materials and their band structures**

Sacrificial H₂ and O₂ evolutions of half reactions were carried out as test reactions of water splitting over Rh- and Ir-doped metal oxide photocatalysts using a sacrificial electron donor and acceptor to see the ability for photocatalytic H₂ or O₂ evolution, as shown in Table 1, prior to conducting water splitting.

The energy gaps were determined from the diffuse reflectance spectra and wavelength dependence of the photocatalytic activities as shown in Fig. 2 and 3. In general, trivalent and tetravalent Rh and Ir species are doped at the Ti⁴⁺, Nb⁵⁺ and Ta⁵⁺ sites in metal oxide materials.²⁰–²⁵,³⁷,³⁸ Among the species, Rh³⁺ and Ir³⁺ contribute to the visible light response for metal oxide photocatalysts.²⁰,²⁴,³⁷,³⁸ Sb⁵⁺ was codoped with Rh³⁺ at the Ti⁴⁺ and Nb⁵⁺ sites for charge compensation to enhance the formation of Rh³⁺ and Ir³⁺ and suppress the formation of Rh⁴⁺ and Ir⁴⁺ as efficient recombination centers between photogenerated electrons and holes, while Ba²⁺ and La³⁺ were replaced at the alkali and alkaline earth metal sites for the same purpose. SrTiO₃:Ir was reduced with H₂ at 473 K to form the dopant Ir³⁺.

In the case of the Ir-doped photocatalysts, SrTiO₃:Ir was active for both the sacrificial H₂ and O₂ evolutions,²⁰,²² whereas BaTa₂O₆:Ir,La and NaTaO₃:Ir,La were active only for the sacrificial H₂ evolution.²³,²⁴ BaTa₂O₆:Ir,La and NaTaO₃:Ir,La were not active for sacrificial O₂ evolution even if they were reduced with H₂ at 673 K, as SrTiO₃:Ir was, to reduce the Ir⁴⁺ species. The Rh-doped photocatalysts

### Table 1  Sacrificial H₂ and O₂ evolutions under visible light irradiation over Rh- or Ir-doped metal oxide photocatalysts

<table>
<thead>
<tr>
<th>Photocatalyst</th>
<th>Energy gap/eV</th>
<th>Incident light/nm</th>
<th>Activity/μmol h⁻¹</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pt/NaTaO₃:Ir(1%),La(2%)</td>
<td>2.1–2.3</td>
<td>λ &gt; 420</td>
<td>H₂</td>
<td>3.2</td>
</tr>
<tr>
<td>NaTaO₃:Ir(1%),La(2%)</td>
<td>2.1–2.3</td>
<td>λ &gt; 420</td>
<td>O₂</td>
<td>—</td>
</tr>
<tr>
<td>Pt/BaTa₂O₆:Ir(1%),La(2%)</td>
<td>1.9–2.0</td>
<td>λ &gt; 420</td>
<td>H₂</td>
<td>4.6</td>
</tr>
<tr>
<td>BaTa₂O₆:Ir(1%),La(2%)</td>
<td>1.9–2.0</td>
<td>λ &gt; 420</td>
<td>O₂</td>
<td>—</td>
</tr>
<tr>
<td>Pt/SrTiO₃:Ir(0.2%)</td>
<td>1.6–1.8</td>
<td>λ &gt; 440</td>
<td>H₂</td>
<td>8.6</td>
</tr>
<tr>
<td>SrTiO₃:Ir(0.2%)</td>
<td>1.6–1.8</td>
<td>λ &gt; 420</td>
<td>O₂</td>
<td>4.3</td>
</tr>
<tr>
<td>Pt/NaNbO₃:Rh(1%),Ba(2%)</td>
<td>2.5</td>
<td>λ &gt; 420</td>
<td>H₂</td>
<td>0.2</td>
</tr>
<tr>
<td>NaNbO₃:Rh(1%),Ba(2%)</td>
<td>2.5</td>
<td>λ &gt; 420</td>
<td>O₂</td>
<td>5.3</td>
</tr>
<tr>
<td>Pt/TiO₂:Rb(0.5%),Sb(1%)</td>
<td>2.1</td>
<td>λ &gt; 440</td>
<td>H₂</td>
<td>0</td>
</tr>
<tr>
<td>TiO₂:Rb(0.5%),Sb(1%)</td>
<td>2.1</td>
<td>λ &gt; 440</td>
<td>O₂</td>
<td>7.5</td>
</tr>
</tbody>
</table>

*Photocatalyst: 0.1–0.3 g; light source: 300 W Xe lamp with long-pass filters (λ > 420 nm or λ > 440 nm); reaction cell: top-irradiation cell with a Pyrex window.¹ Cocatalyst: Pt (0.3 wt%, photodeposition); reactant solution: 10 vol% aqueous methanol solution. Cocatalyst: none; reactant solution: 0.02–0.05 mol L⁻¹ aqueous AgNO₃ solution (120–150 mL). Treatment: H₂-reduction at 473 K for 2 h.*
TiO$_2$:Rh,Sb$^{25}$ and NaNbO$_3$:Rh,Ba were active for sacrificial O$_2$ evolution using Ag$^+$ as an electron acceptor. NaNbO$_3$:Rh,Ba showed very low activity for the sacrificial H$_2$ evolution. These properties will be discussed based on the band structure in the Discussion section. These results of the sacrificial H$_2$ and O$_2$ evolutions suggest that NaTaO$_3$:Ir,La and BaTa$_2$O$_6$:Ir,La can be used as H$_2$-evolving photocatalysts.

**Fig. 2** The wavelength dependence of H$_2$ evolution from a 10 vol% aqueous methanol solution (closed circles) and diffuse reflectance spectra (solid line) of (a) NaTaO$_3$:Ir(La(1%),La(2%)) and (b) BaTa$_2$O$_6$:Ir(1%),La(2%). Photocatalyst: 0.1 g, cocatalyst: Pt (photodeposition) for H$_2$ evolution, reactant solution: 120 mL, light source: 300 W Xe lamp with long-pass filters, and reaction cell: top-irradiation cell with a Pyrex window. Samples of diffuse reflectance spectra were reduced at 473 K.

TiO$_2$:Rh,Sb$^{25}$ and NaNbO$_3$:Rh,Ba were active for sacrificial O$_2$ evolution using Ag$^+$ as an electron acceptor. NaNbO$_3$:Rh,Ba showed very low activity for the sacrificial H$_2$ evolution. These properties will be discussed based on the band structure in the Discussion section. These results of the sacrificial H$_2$ and O$_2$ evolutions suggest that NaTaO$_3$:Ir,La and BaTa$_2$O$_6$:Ir,La can be used as H$_2$-evolving photocatalysts.

**Fig. 3** The wavelength dependence of O$_2$ evolution from a 0.05 mol L$^{-1}$ aqueous silver nitrate solution (closed circles) and diffuse reflectance spectra (solid line) of (a) TiO$_2$:Rh(1.3%),Sb(2.6%), (b) NaNbO$_3$:Rh(1.2%),Ba(1.44%) and (c) SrTiO$_3$:Ir(0.2%) with H$_2$ reduction at 473 K. Photocatalyst: 0.1–0.3 g, reactant solution: 150 mL, light source: 300 W Xe lamp with long-pass filters, and reaction cell: top-irradiation cell with a Pyrex window.
photocatalysts for the construction of Z-scheme systems, while SrTiO$_3$:Ir, NaNbO$_3$:Rh,Ba and TiO$_2$:Rh,Sb are expected to be employed as O$_2$-evolving photocatalysts.

Fig. 2 and 3 show the diffuse reflectance spectra and wavelength dependence of the photocatalytic H$_2$ and O$_2$ evolutions of the Rh- and Ir-doped metal oxide photocatalysts in the presence of sacrificial reagents. The wavelengths were controlled with long-pass filters. It is vital to see the wavelength dependency of the photocatalytic activity because it is not guaranteed that photocatalysts with visible light absorption bands always give activities under visible light irradiation. The onsets of the wavelength dependence agreed with those of the diffuse reflection spectra. The onset wavelengths for the H$_2$ evolutions were 640 and 600 nm for BaTa$_2$O$_6$:Ir,La and NaTaO$_3$:Ir,La, respectively. These onset wavelengths were longer than the 540 nm of SrTiO$_3$:Rh (a conventional H$_2$-evolving photocatalyst). The onset wavelengths for O$_2$ evolution were 600, 500 and 700 nm for TiO$_2$:Rh,Sb, NaNbO$_3$:Rh,Ba and SrTiO$_3$:Ir, respectively. It is noteworthy that TiO$_2$:Rh,Sb and SrTiO$_3$:Ir responded at longer wavelengths than the BiVO$_4$ (BG: 2.4 eV) (a conventional O$_2$-evolving photocatalyst).

Fig. 4 shows the band structures of the Rh- and Ir-doped metal oxide photocatalysts. The impurity levels of Rh$^{3+}$ and Ir$^{3+}$ were estimated from the energy gaps determined by diffuse reflection spectra supposing that the valence band consisting of O 2$p$ located at +3 V vs. NHE at a pH of 0.$^9$ The absorption bands in the visible light region shown in Fig. 2 and 3 are due to electronic transition from the impurity levels consisting of Rh$^{3+}$ and Ir$^{3+}$ to the conduction bands of the host materials. The impurity levels formed with electron-filled orbitals of Ir$^{3+}$ were around 1.0–1.2 V for NaTaO$_3$ and BaTa$_2$O$_6$, while Ir$^{3+}$ in SrTiO$_3$ formed an impurity level around 1.4–1.6 V that was deeper than those in the cases of the tantalates. The energy levels formed with electron-filled orbitals of Rh$^{3+}$ located around 2.0–2.1 V that were similar to those of SrTiO$_3$:Rh and SrTiO$_3$:Rh,Sb.$^{20,21}$ The reason Rh$^{3+}$ forms a deeper impurity level than Ir$^{3+}$ is due to Ir$^{4+}$ being more stable than Rh$^{4+}$ in metal oxides. Therefore, electronic transition from the Ir$^{3+}$ impurity level to a conduction band is easier than that from Rh$^{3+}$, resulting in the formation of the shallow impurity level by Ir$^{3+}$.

![Fig. 4](image-url)  
**Fig. 4** The band structures of Rh- or Ir-doped metal oxide photocatalysts at pH 0.
Z-schematic systems for photocatalytic water splitting employing Rh- and Ir-doped metal oxide materials

The combination of SrTiO₃:Rh and BiVO₄ photocatalysts can be a benchmark of a Z-scheme photocatalyst system. The Z-schematic water splitting proceeds using Fe³⁺/²⁺ and [Co(bpy)$_3$]³⁺/²⁺ redox couples as ionic mediators (Fig. 1(a)) and also via interparticle electron transfer between SrTiO₃:Rh and BiVO₄ particles with and without RGO (Fig. 1(b) and (c)). In general, water splitting via Z-schematic interparticle electron transfer by contact between the particles of the H₂- and O₂-evolving photocatalysts with and without RGO can be achieved when the H₂- and O₂-evolving photocatalysts satisfy the following two requirements; (i) H₂- and O₂-evolving photocatalysts possess p- and n-type semiconductor properties, respectively, (ii) there is a certain electrode potential at which the cathodic photocurrent of the p-type semiconductor overlaps with the anodic photocurrent of the n-type semiconductor, being similar to water splitting using a photoelectrochemical cell working with no applied external bias as shown in Fig. 1(d). Moreover, H₂- and O₂-evolving photocatalysts must have contact with each other for interparticle electron transfer. In contrast to this, a Z-scheme system employing an ionic electron mediator could work regardless of the p- or n-type properties of the H₂- or O₂-evolving photocatalysts, if the photocatalysts have potentials for the reduction or oxidation of redox couple ionic mediators and the adsorption abilities for the redox couples. The SrTiO₃:Rh and BiVO₄ photocatalysts satisfy these factors resulting in all of the Z-scheme systems showing activities for water splitting into H₂ and O₂ in stoichiometric amounts without any sacrificial reagents (Fig. 1(a)–(d)).

Various types of Z-scheme photocatalyst systems employing Ir- and Rh-doped photocatalysts for water splitting into H₂ and O₂ under visible light irradiation are shown in Table 2. Here, the absolute evaluation of the performance of photocatalyst systems for water splitting is how much H₂ and O₂ is obtained under certain experimental conditions. In this sense, the activities of the different Z-scheme systems in Table 2 are comparable with each other, but not with those of Table 1, because of the almost identical experimental conditions for the water splitting, even if the kinetics would be different among the systems.

For the construction of Z-scheme systems, BiVO₄ (an O₂-evolving photocatalyst) was combined with BaTa₂O₆:Ir,La and NaTaO₃:Ir,La (H₂-evolving photocatalysts), while SrTiO₃:Rh (an H₂-evolving photocatalyst) was combined with TiO₂:Rh,Sb, NaNbO₃:Rh,Ba and SrTiO₃:Ir (O₂-evolving photocatalysts) as suggested by their H₂ and O₂ evolution abilities. In addition to the suspension system, the photoelectrochemical properties of photocatalyst powders immobilized on a conducting substrate as shown in Fig. 1(d) were examined using a Pt counter electrode without any sacrificial reagents to see the cathodic or anodic photocurrent and the onset potentials of semiconductor properties, not as a water splitting device, as shown in Fig. 5, in order to consider the potential overlap for Z-schematic water splitting via interparticle electron transfer with and without RGO. It has been reported that SrTiO₃:Rh and BiVO₄ function as a photocathode and photoanode, respectively. NaTaO₃:Ir,La showed only an anodic photocurrent. Although BaTa₂O₆:Ir,La and SrTiO₃:Ir showed cathodic and anodic photocurrents, the photocurrents were poor and the onset potentials shifted with the sweeping direction of the CV curves. This result implies that
Table 2  Z-schematic water splitting under visible light irradiation using Rh- or Ir-doped metal oxide photocatalysts

<table>
<thead>
<tr>
<th>H₂-photocat.</th>
<th>O₂-photocat.</th>
<th>Mediator</th>
<th>Initial pH</th>
<th>Activity/µmol h⁻¹</th>
<th>H₂</th>
<th>O₂</th>
<th>Ref.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pt/NaTaO₃:Ir(1%),La(2%)</td>
<td>BiVO₄</td>
<td>None</td>
<td>4.2</td>
<td>Trace</td>
<td>Trace</td>
<td>Trace</td>
<td>This work</td>
</tr>
<tr>
<td>Pt/NaTaO₃:Ir(1%),La(2%)</td>
<td>BiVO₄</td>
<td>RGO</td>
<td>4.2</td>
<td>Trace</td>
<td>Trace</td>
<td>Trace</td>
<td>This work</td>
</tr>
<tr>
<td>Pt/NaTaO₃:Ir(1%),La(2%)</td>
<td>BiVO₄</td>
<td>Fe⁺³⁺/²⁺f</td>
<td>2.4</td>
<td>0.8</td>
<td>0.3</td>
<td>This work</td>
<td></td>
</tr>
<tr>
<td>Pt/NaTaO₃:Ir(1%),La(2%)</td>
<td>BiVO₄</td>
<td>[Co(bpy)₃]³⁺/²⁺g</td>
<td>4.2</td>
<td>0.9</td>
<td>0.4</td>
<td>This work</td>
<td></td>
</tr>
<tr>
<td>Pt/BaTa₂O₆:Ir(1%),La(2%)</td>
<td>BiVO₄</td>
<td>None</td>
<td>4.2</td>
<td>0.9</td>
<td>0.4</td>
<td>This work</td>
<td></td>
</tr>
<tr>
<td>Pt/BaTa₂O₆:Ir(1%),La(2%)</td>
<td>BiVO₄</td>
<td>RGO</td>
<td>4.2</td>
<td>0.1</td>
<td>0.4</td>
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<tr>
<td>Pt/BaTa₂O₆:Ir(1%),La(2%)</td>
<td>BiVO₄</td>
<td>Fe⁺³⁺/²⁺f</td>
<td>2.4</td>
<td>5.9</td>
<td>2.1</td>
<td>22</td>
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<tr>
<td>Ru/SrTiO₃:Rh(1%)</td>
<td>Pt/SrTiO₃:Ir(0.2%)</td>
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<td>3.5</td>
<td>0.4</td>
<td>This work</td>
<td></td>
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</tr>
<tr>
<td>Ru/SrTiO₃:Rh(1%)</td>
<td>Pt/SrTiO₃:Ir(0.2%)</td>
<td>RGO</td>
<td>3.5</td>
<td>0.8</td>
<td>This work</td>
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<tr>
<td>Ru/SrTiO₃:Rh(1%)</td>
<td>Pt/SrTiO₃:Ir(0.2%)</td>
<td>Fe⁺³⁺/²⁺f</td>
<td>2.4</td>
<td>2.5</td>
<td>1.0</td>
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<tr>
<td>Ru/SrTiO₃:Rh(1%)</td>
<td>Pt/SrTiO₃:Ir(0.2%)</td>
<td>[Co(bpy)₃]³⁺/²⁺h</td>
<td>7.8</td>
<td>5.1</td>
<td>2.4</td>
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<td></td>
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<tr>
<td>Ru/SrTiO₃:Rh(1%)</td>
<td>NaNbO₂:Rh(1%),Ba(2%)</td>
<td>None</td>
<td>2.4</td>
<td>2.1</td>
<td>1.1</td>
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<tr>
<td>Ru/SrTiO₃:Rh(1%)</td>
<td>NaNbO₂:Rh(1%),Ba(2%)</td>
<td>Fe⁺³⁺/²⁺f</td>
<td>2.4</td>
<td>5.7</td>
<td>2.3</td>
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<tr>
<td>Ru/SrTiO₃:Rh(1%)</td>
<td>NaNbO₂:Rh(1%),Ba(2%)</td>
<td>[Co(bpy)₃]³⁺/²⁺h</td>
<td>3.8</td>
<td>4.1</td>
<td>1.7</td>
<td>This work</td>
<td></td>
</tr>
<tr>
<td>Ru/SrTiO₃:Rh(1%)</td>
<td>TiO₂:Rh(0.5%),Sb(1%)</td>
<td>None</td>
<td>3.5</td>
<td>5.4</td>
<td>2.4</td>
<td>This work</td>
<td></td>
</tr>
<tr>
<td>Ru/SrTiO₃:Rh(1%)</td>
<td>TiO₂:Rh(0.5%),Sb(1%)</td>
<td>Fe⁺³⁺/²⁺f</td>
<td>2.4</td>
<td>12</td>
<td>5.3</td>
<td>This work</td>
<td></td>
</tr>
<tr>
<td>Ru/SrTiO₃:Rh(1%)</td>
<td>TiO₂:Rh(0.5%),Sb(1%)</td>
<td>[Co(bpy)₃]³⁺/²⁺h</td>
<td>3.5</td>
<td>28</td>
<td>13</td>
<td>This work</td>
<td></td>
</tr>
</tbody>
</table>

Photocatalyst: 0.05 or 0.1 g each; light source: 300 W Xe lamp with a long-pass filter (λ > 420 nm); reaction cell: top-irradiation cell with a Pyrex window. Cocatalyst: Pt (0.3 wt%, impregnation at 673 K for 2 h and subsequent H₂-reduction at 673 K for 1 h). Cocatalyst: Ru (0.7 wt%, photodeposition). Cocatalyst: Pt (1 wt%, impregnation without calcination and subsequent H₂-reduction at 573 K for 1 h). Reactant solution: H₂SO₄ solution. Reactant solution: 2 mmol L⁻¹ FeCl₃ solution. Reactant solution: 0.02 mmol L⁻¹ [Co(bpy)₃]SO₄ solution (120 mL).
those photocurrents might not be due to H₂ and O₂ evolutions, but possibly due to redox reactions of the doped Ir species. In contrast, TiO₂:Rh,Sb, and NaNbO₃:Rh,Ba gave clear anodic photocurrents indicating an n-type semiconductor character. The onset potential of NaNbO₃:Rh,Ba was more negative than that of TiO₂:Rh,Sb, whereas the anodic photocurrent of NaNbO₃:Rh,Ba was much smaller than that of TiO₂:Rh,Sb. These anodic photocurrents overlapped with the cathodic photocurrent of SrTiO₃:Rh at a certain electrode potential.

The Z-scheme systems were classified into three groups depending on the combination of H₂- and O₂-evolving photocatalysts and an electron mediator;
being active for only a [Co(bpy)₃]³⁺/²⁺ redox couple, active for [Co(bpy)₃]³⁺/²⁺ and Fe³⁺/²⁺ redox couple ionic mediators, active not only for [Co(bpy)₃]³⁺/²⁺ and Fe³⁺/²⁺ redox couples but also via interparticle electron transfer with and without RGO. NaTaO₃:Ir,La + BiVO₄ and BaTa₂O₆:Ir,La + BiVO₄ were active when not Fe³⁺/²⁺ but a [Co(bpy)₃]³⁺/²⁺ redox couple was used. Although the BaTa₂O₆:Ir,La + BiVO₄ showed activities via interparticle electron transfer at a pH of 4.2 with and without RGO, the activities were smaller than that with the ionic electron mediator [Co(bpy)₃]³⁺/²⁺. SrTiO₃:Rh + SrTiO₃:Ir was active when the [Co(bpy)₃]³⁺/²⁺ and Fe³⁺/²⁺ redox couple ionic mediators were used, whereas it was not active via interparticle electron transfer with and without RGO at pH 3.5. SrTiO₃:Rh + NaNbO₃:Rh,Ba and SrTiO₃:Rh + TiO₂:Rh,Sb were active for the [Co(bpy)₃]³⁺/²⁺ and Fe³⁺/²⁺ redox couples and via interparticle electron transfer without RGO. These Z-scheme systems with RGO using NaNbO₃:Rh,Ba and TiO₂:Rh,Sb were not examined in detail, because the GOs were not suitably photoreduced on the NaNbO₃:Rh,Ba and TiO₂:Rh,Sb due to their poor reducing activities, as expected from their poor H₂ evolution activities as shown in Table 1. SrTiO₃:Rh + TiO₂-Rh,Sb showed the best performances among the Z-scheme systems in Table 2.

Discussion

H₂ and O₂ evolution abilities (shown in Table 1) can be considered based on the band structure, as shown in Fig. 4. Photogenerated holes in the Ir³⁺ levels in NaTaO₃:Ir,La and BaTa₂O₆:Ir,La have no potentials for water oxidation. In contrast, the Ir³⁺ level of SrTiO₃:Ir possesses the water oxidation potential, though the driving force is not so large. Conduction bands consisting of Ta 5d orbitals in BaTa₂O₆:Ir,La and NaTaO₃:Ir,La possess thermodynamically enough potentials for water reduction to form H₂. On the other hand, the photogenerated holes in the Rh³⁺ levels in NaNbO₃:Rh,Ba and TiO₂:Rh,Sb have enough potential for water oxidation to form O₂. These energy levels were consistent with the H₂ and O₂ evolution abilities, as shown in Table 1. Of course, other kinetic factors of the active sites would exist for the H₂ and O₂ evolution abilities in addition to the energy levels of the thermodynamic factor.

Let us consider the reason why the Z-scheme systems shown in Table 2 are classified into three groups, based on the band structure and photoelectrochemical properties. Fig. 6 shows energy diagrams for the Z-schemes employing Fe³⁺/²⁺ and [Co(bpy)₃]³⁺/²⁺ redox couples at pH 2.4 and 4.0, respectively. It is assumed that the band levels shift with –0.059 V pH⁻¹ because those materials are metal oxides.

NaTaO₃:Ir,La + BiVO₄ and BaTa₂O₆:Ir,La + BiVO₄ were active with a [Co(bpy)₃]³⁺/²⁺ redox couple at pH 4.2, but not with Fe³⁺/²⁺ at pH 2.4. Photogenerated holes in the Ir³⁺ levels in NaTaO₃:Ir,La and BaTa₂O₆:Ir,La (an H₂-evolving photocatalyst) possess small driving forces for oxidation of Fe²⁺ (an electron mediator) at pH 2.4 as shown in Fig. 6(a), whereas they possess enough potential for oxidation of [Co(bpy)₃]²⁺ at pH 4.2 as shown in Fig. 6(b). The activities from the interparticle electron transfer were very small or negligible because of poor photoresponse and poor overlaps of the potentials giving cathodic photocurrents of NaTaO₃:Ir,La and BaTa₂O₆:Ir,La and an anodic photocurrent of BiVO₄ as shown in Fig. 5, and also may be due to poor contact between NaTaO₃:Ir,La or BaTa₂O₆:Ir,La (an H₂-evolving photocatalyst) and BiVO₄ (an O₂-
evolving photocatalyst) in the suspension at pH 4.2. In contrast to them, SrTiO$_3$:Rh + SrTiO$_3$:Ir was active when the \([\text{Co(bpy)}_3]^{3+/2+}\) and \(\text{Fe}^{3+/2+}\) redox couple ionic mediators were used, because the conduction band of SrTiO$_3$:Ir (an O$_2$-evolving photocatalyst) has enough potential for the reduction of Fe$^3+$ and \([\text{Co(bpy)}_3]^{3+}\) (Fig. 6(c) and (d)). However, because the anodic photocurrent of SrTiO$_3$:Ir hardly overlapped with the cathodic photocurrent of SrTiO$_3$:Rh, the activities by interparticle electron transfer with and without RGO were negligible.

The conduction bands of TiO$_2$:Rh,Sb and NaNbO$_3$:Rh,Ba (O$_2$-evolving photocatalysts) also possessed enough potential for the reduction of Fe$^3+$ and \([\text{Co(bpy)}_3]^{3+}\) as well as SrTiO$_3$:Ir (Fig. 6(c) and (d)). Moreover, in these cases, the anodic photocurrents of the TiO$_2$:Rh,Sb and NaNbO$_3$:Rh,Ba photoelectrodes overlapped enough with the cathodic photocurrent of SrTiO$_3$:Rh. So, it is reasonable that SrTiO$_3$:Rh + NaNbO$_3$:Rh,Ba and SrTiO$_3$:Rh + TiO$_2$:Rh,Sb were active not only for the \([\text{Co(bpy)}_3]^{3+/2+}\) and Fe$^{3+/2+}$ redox couples but also via interparticle electron transfer with and without RGO. The activity of SrTiO$_3$:Rh + TiO$_2$:Rh,Sb was higher than that of SrTiO$_3$:Rh + NaNbO$_3$:Rh,Ba for all of the types of Z-scheme system. In the cases with the use of ionic electron mediators, it is probably due to the higher activity for O$_2$ evolution and the narrower energy gaps of TiO$_2$:Rh,Sb than those of NaNbO$_3$:Rh,Ba, as shown in Table 1. The reason why SrTiO$_3$:Rh + TiO$_2$:Rh,Sb showed a higher activity than SrTiO$_3$:Rh + NaNbO$_3$:Rh,Ba via interparticle electron transfer is that TiO$_2$:Rh,Sb gave much larger anodic photocurrents than NaNbO$_3$:Rh,Ba, as shown in Fig. 5.

---

Fig. 6 The band structures of Rh- or Ir-doped metal oxide photocatalysts, and redox potentials of Fe and Co-complex ionic electron mediators at pH 2.4 and 4.0. (a) and (b) represent Z-scheme systems of Ir-doped metal oxides as H$_2$-evolving photocatalysts combined with BIVO$_4$. (c) and (d) represent Z-scheme systems of Rh-doped metal oxides and SrTiO$_3$:Ir as O$_2$-evolving photocatalysts combined with SrTiO$_3$:Rh.
Conclusions

We have successfully developed Z-scheme photocatalyst systems for water splitting under visible light irradiation employing Rh- and Ir-doped metal oxide photocatalysts with longer wavelength responses than conventional SrTiO$_3$:Rh and BiVO$_4$. The impurity levels of doped Ir$^{3+}$ and Rh$^{3+}$ that contributed to the visible light responses for the photocatalytic reactions were determined from diffuse reflectance spectra and supposing +3.0 V vs. NHE of a valence band of the oxide photocatalyst materials. The impurity levels of Ir$^{3+}$ in NaTaO$_3$:Ir,La and BaTa$_2$O$_6$:Ir,La have sufficient potentials for the oxidation of the electron mediator [Co(bpy)$_3$]$^{2+}$, but insufficient potentials for water oxidation to form O$_2$ and oxidation of the electron mediator Fe$^{2+}$. The conduction bands of the NaTaO$_3$:Ir,La and BaTa$_2$O$_6$:Ir,La have enough potential for water reduction to form H$_2$. Therefore, the NaTaO$_3$:Ir,La and BaTa$_2$O$_6$:Ir,La could be used as H$_2$-evolving photocatalysts only when a [Co(bpy)$_3$]$^{3+/2+}$ redox couple was used. The impurity levels of Ir$^{3+}$ in SrTiO$_3$:Ir and Rh$^{3+}$ in TiO$_2$:Rh,Sb and NaNbO$_3$:Rh,Ba have potentials for water oxidation and their conduction bands possess the potentials for reduction of the electron mediators Fe$^{3+}$ and [Co(bpy)$_3$]$^{3+}$. This property means that SrTiO$_3$:Ir, TiO$_2$:Rh,Sb and NaNbO$_3$:Rh,Ba could be employed as O$_2$-evolving photocatalysts for the construction of Z-scheme systems employing ionic electron mediators. Moreover, photoelectrochemical measurements using photocatalyst powders immobilized on conducting substrate revealed that the n-type characters and relatively negative onset potentials of the TiO$_2$:Rh,Sb and NaNbO$_3$:Rh,Ba photoanodes enabled the Z-scheme systems to work by interparticle electron transfer. Although the efficiencies of the present Z-scheme systems are low at the present stage, these will be improved by interfacial controls from a kinetic point of view in basic research. These results and discussion will contribute to the design of a highly active photocatalyst system for water splitting into H$_2$ and O$_2$, aiming for the demonstration of actual solar water splitting using a suitable reactor.

Conflicts of interest

There are no conflicts to declare.

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References


