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Mechanism of a one-photon two-electron process in photocatalytic hydrogen evolution from ascorbic acid with a cobalt chlorin complex†

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A one-photon two-electron process was made possible in photocatalytic H_2 evolution from ascorbic acid with a cobalt(II) chlorin complex [Co^{II}(Ch)] *via* electron transfer from ascorbate to the excited state of $[Ru(bpy)_3]^{2+}$ followed by electron transfer from $[Ru(bpy)_3]^{+}$ to Co^{II}(Ch) with proton to give the hydride complex, which reacts with proton to produce H_2. [Co^{III}(Ch)]$^+$ was reduced by ascorbate to reproduce Co^{II}(Ch).

Photocatalytic production of hydrogen (H_2) has attracted increasing attention as a clean energy source because of the ever-increasing demand for energy and climate change on our planet.[1] A number of highly efficient hydrogen evolving systems have been developed including homogeneous and heterogeneous photocatalytic systems.[2–13] Two electrons are required to produce H_2 from protons, although one photon generates normally only one electron. A mechanism of photocatalytic production of H_2 was reported to clarify how photoinduced electron transfer of a photosensitiser (a one-electron process) leads to H_2 production (a two-electron process).[14–16] Disproportionation of one-electron reduced species of metal complexes resulted in formation of the two-electron reduced species from which H_2 is formed.[17] Bimolecular reactions of metal(III)–hydride complexes also generate H_2 accompanied by regeneration of metal(II) complexes.[18] In each case, the maximum quantum yield of H_2 production per photon is 50%, because two photons are required to produce two electrons. Thus there has so far been no example for one photon to generate one H_2 molecule.

We report herein photocatalytic H_2 evolution from ascorbic acid ($AscH_2$) with a cobalt(II) chlorin complex [Co^{II}(Ch)] (a chemical structure shown in Scheme 1)[19] in an aqueous acetonitrile solution (H_2O/MeCN), which proceeds *via* a one-photon two-electron process. The photocatalytic mechanism is clarified by nanosecond laser transient absorption spectra and by examining each step in the catalytic cycle independently.

Visible light irradiation of a deaerated (Ar-saturated) H_2O/MeCN solution (1 : 1 v/v) of $[Ru(bpy)_3]^{2+}$ (bpy = 2,2′-bipyridine) containing ascorbic acid ($AscH_2$) and ascorbate ($AscH^-$) (E_{ox} = 0.43 V *vs.* SCE) as an electron donor and Co^{II}(Ch) (E_{red} = −0.96 V *vs.* SCE) (Fig. S1 in the ESI†) as a catalyst resulted in H_2 evolution (Fig. 1, black line). When the ratio of $AscH^-$ to $AscH_2$ was changed as fixed total concentrations of $AscH_2$ and $AscH^-$ ($[AscH_2] + [AscH^-]$ = 1.1 M), the largest H_2 evolution activity was attained with $AscH^-$ (0.30 M) and $AscH_2$ (0.80 M) (Fig. S2 in ESI†). The smaller concentration of $AscH^-$ results in less efficient reductive quenching of the $[Ru(bpy)_3]^{2+*}$ emission (* denotes the excited state).

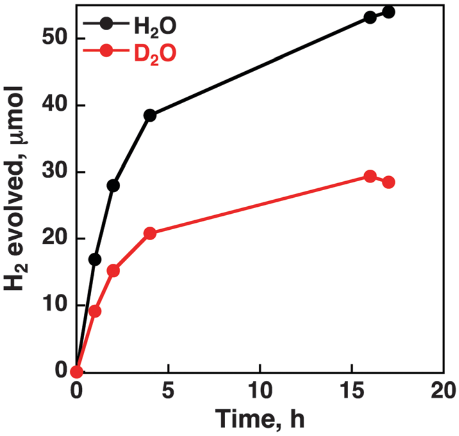

Fig. 1 Time courses of H_2 evolution in the photocatalytic reduction of proton in an Ar-saturated H_2O/MeCN (black) and D_2O/MeCN (red) mixed solution (1 : 1 v/v) containing $[Ru^{II}(bpy)_3]^{2+}$ (2.0 mM), $AscH_2$ (0.80 M), AscHNa (0.30 M) and Co^{II}(Ch) (25 μM) under irradiation of visible light (λ > 420 nm) at 298 K.

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† Electronic supplementary information (ESI) available: Experimental details and cyclic voltammograms (Fig. S1), time courses of H_2 evolution (Fig. S2), emission spectra (Fig. S3 and S5), UV-vis absorption spectra (Fig. S4 and S7) and kinetic data (Fig. S6–S11). See DOI: 10.1039/c5cc05064b

The quenching efficiency of $[Ru(bpy)_3]^{2+*}$ (E_{red} = 0.77 V *vs.* SCE in MeCN)[20] by $AscH^-$ (0.30 M) with $AscH_2$ (0.80 M) was determined to be 95% (Fig. S3 in ESI†). On the other hand, the smaller concentration of $AscH_2$ may retard H_2 production due to decreasing the acidity. When H_2O was replaced by D_2O, D_2 and HD were produced without formation of H_2. Thus, hydrogen was produced from water and ascorbic acid as electron and proton sources. The observed deuterium kinetic isotope effect (KIE) in Fig. 1 (k_H/k_D = 1.8 in the initial stage) suggests that the Co–H bond cleavage of a cobalt hydride intermediate ($[Co^{III}(H)(Ch)]$) by proton may be the rate-determining step for the photocatalytic H_2 evolution (*vide infra*).

The concentration of $Co^{II}(Ch)$ was optimised to be 50 μM for the efficient photocatalytic H_2 evolution. The absorption of $[Ru(bpy)_3]^{2+}$ is blocked by the larger concentration of $Co^{II}(Ch)$ (Fig. S4 in ESI†).[21] The quantum yield of the photocatalytic H_2 evolution was determined to be 12% using a ferric oxalate actinometer (see the Experimental section in ESI†). This value is similar to the highest value reported for photocatalytic H_2 evolution using a cobalt terpyridine complex (Φ = 0.13).[22]

Nanosecond transient absorption spectra of an H_2O/MeCN solution of $[Ru(bpy)_3]^{2+}$ with $AscH_2$ and $AscH^-$ are shown in Fig. 2, where appearance of the absorption band at 500 nm due to $[Ru(bpy)_3]^+$ is observed upon the nanosecond laser excitation. Thus, electron transfer from $AscH^-$ to $[Ru(bpy)_3]^{2+*}$ occurred to produce $AscH^\bullet$ and $[Ru(bpy)_3]^+$. The rate constant of electron transfer from $AscH^-$ to $[Ru(bpy)_3]^{2+*}$ (k_{et}) was determined to be 8.0×10^8 M^{-1} s^{-1} from a slope of Stern–Volmer plot (K_{SV} = 3.5×10^2 M^{-1}) and the lifetime of $[Ru(bpy)_3]^{2+*}$ (0.44 μs in water/MeCN 1 : 1 v/v) (Fig. S5 in ESI†).[23] The decay rate of absorbance at 500 nm due to $[Ru(bpy)_3]^+$ obeyed the second-order kinetics of bimolecular back electron transfer from $[Ru(bpy)_3]^+$ to $AscH^\bullet$. In the presence of $Co^{II}(Ch)$, the decay of absorbance became much faster because of electron transfer from $[Ru(bpy)_3]^+$ to $Co^{II}(Ch)$ as shown in Fig. 2b. The decay rate constant linearly increased with increasing the concentration of $[Co^{II}(Ch)]$ (Fig. S6 in ESI†). The rate constant of electron transfer from $[Ru(bpy)_3]^+$ to $Co^{II}(Ch)$ was determined to be 2.5×10^9 M^{-1} s^{-1} from the slope of dependence of the first-order decay rate constant on concentration of $Co^{II}(Ch)$ (Fig. S6b in ESI†).

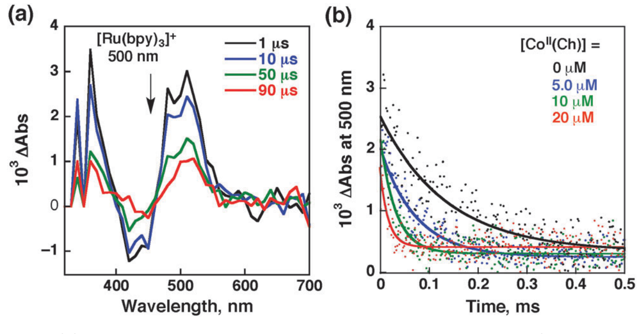


Fig. 2 (a) Transient absorption spectra after laser excitation (λ = 450 nm) of $[Ru^{II}(bpy)_3]^{2+}$ (80 μM) in the presence of $AscH_2$ (0.80 M) and AscHNa (0.30 M) in a deaerated H_2O/MeCN mixed solution (1 : 1 v/v) at 298 K. (b) Time profiles of absorbance at 500 nm due to decay of $[Ru(bpy)_3]^+$ in the presence of various concentrations of $Co^{II}(Ch)$ (0–20 μM) in deaerated H_2O/MeCN mixed solutions (1 : 1 v/v) containing $[Ru^{II}(bpy)_3]^{2+}$ (80 μM), $AscH_2$ (0.80 M), AscHNa (0.30 M).

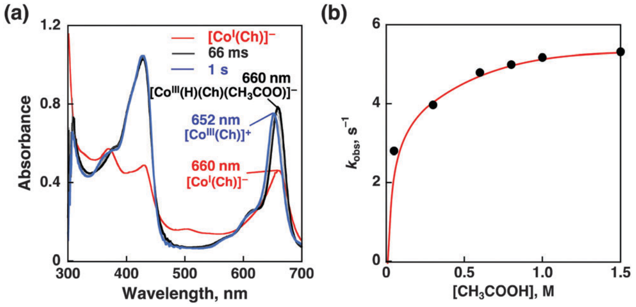
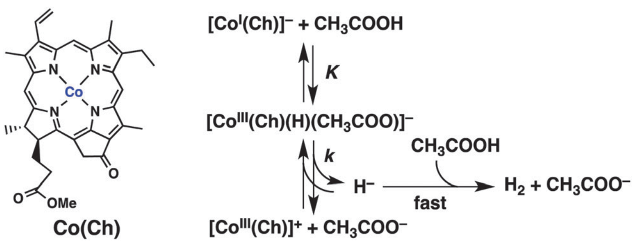


Fig. 3 (a) UV-vis absorption spectral changes of $[Co^I(Ch)]^-$ (30 μM) upon addition of CH_3COOH (0.30 M) in dearated MeCN at 298 K. The black and blue lines show the spectra taken at 66 ms and 1 s after mixing, respectively. The red line shows UV-vis absorption spectrum of $[Co^I(Ch)]^-$ (15 μM) formed by the electron-transfer reduction of $Co^{II}(Ch)$ (15 μM) with $CoCp_2^*$ (300 μM) in dearated MeCN at 298 K. (b) Plot of k_{obs} for the rate of formation of $[Co^{III}(Ch)]^+$ *vs.* $[CH_3COOH]$.

To examine the reaction of $[Co^I(Ch)]^-$ that is produced by electron transfer from $[Ru(bpy)_3]^+$ to $Co^{II}(Ch)$, $[Co^I(Ch)]^-$ was prepared by the one-electron reduction of $Co^{II}(Ch)$ by decamethylcobaltocene $[Co(Cp^*)_2]$ in MeCN (Fig. S7 in ESI†). The UV-vis absorption band of $[Co^I(Ch)]^-$ (red line in Fig. 3a; λ_{max} = 510 nm) decreased with increasing absorption band at 660 nm (black line) at 66 ms after addition of acetic acid (CH_3COOH) (0.30 M). Then, this absorption band was finally blue shifted to λ_{max} = 652 nm, which is due to $[Co^{III}(Ch)]^+$.[24,25] Thus, $[Co^I(Ch)]^-$ may react with CH_3COOH to form the hydride complex ($[Co^{III}(H)(Ch)(CH_3COO)]^-$: λ_{max} = 660 nm), from which H_2 was evolved by the reaction with CH_3COOH to produce $[Co^{III}(Ch)]^+$. The reaction of $[Co^I(Ch)]^-$ with CH_3COOH was monitored by the absorption change at 652 nm due to $[Co^{III}(Ch)]^+$ as shown in Fig. 3, where the rate of the formation of $[Co^{III}(Ch)]^+$ obeyed first-order kinetics (Fig. S8 in ESI†). The first-order rate constant increased with increasing concentration of CH_3COOH to approach a constant value (Fig. 3b). Such a saturation behaviour indicates that CH_3COOH is not involved in the rate-determining step and that the reaction of $[Co^I(Ch)]^-$ with CH_3COOH proceeds *via* formation of the hydride complex ($[Co^{III}(H)(Ch)(CH_3COO)]^-$), followed by the rate-determining heterolytic cleavage of the Co^{III}–H bond. The subsequent reaction of the released hydride ion with CH_3COOH to produce H_2 and $[Co^{III}(Ch)]^+$ may be fast as compared with the back reaction of the Co^{III}–H bond cleavage (Scheme 1). The kinetic equation for the formation of $[Co^{III}(Ch)]^+$ is given by eqn (1),



Scheme 1 Mechanism of hydrogen formation by the reaction of $[Co^I(Ch)]^-$ with CH_3COOH.

$$d[[Co^{III}(Ch)]^+]/dt = k[[Co^{III}(H)(Ch)(CH_3COO)]^-] \quad (1)$$

where k is the rate constant of the hydrogen evolution. From the equilibrium constant (K), the concentration of a complex between $[Co^I(Ch)]^-$ and CH_3COOH is given by eqn (2), where

$$[[Co^{III}(H)(Ch)(CH_3COO)]^-] = K[CH_3COOH]([[Co^I(Ch)]^-]_0 - [[Co^{III}(Ch)]^+])/(1 + K[CH_3COOH]) \quad (2)$$

$[[Co^I(Ch)]^-]_0$ is the initial concentration. Eqn (1) is rewritten by eqn (3).

$$d[[Co^{III}(Ch)]^+]/dt = kK[CH_3COOH]([[Co^I(Ch)]^-]_0 - [[Co^{III}(Ch)]^+])/(1 + K[CH_3COOH]) \quad (3)$$

Under the conditions, the concentration of CH_3COOH is much higher than that of $[Co^I(Ch)]^-$, the k_{obs} value is given by eqn (4). To determine the k value, eqn (4) is rewritten by eqn (5), which predicts

$$k_{obs} = kK[CH_3COOH]/(1 + K[CH_3COOH]) \quad (4)$$

$$k_{obs}{}^{-1} = 1/kK{\cdot}[CH_3COOH]^{-1} + 1/k \quad (5)$$

a linear correlation between $k_{obs}{}^{-1}$ and $[CH_3COOH]^{-1}$ (Fig. S9 in ESI†). The k and K values were determined from the intercept and slope of the linear plot of $k_{obs}{}^{-1}$ *vs.* $[CH_3COOH]^{-1}$ to be 5.9 s^{-1} and 7.1 M^{-1}.

When CH_3COOH was replaced by CH_3COOD, the deuterium kinetic isotope effect (KIE) was observed (Fig. S10 in ESI†),[26] indicating that the cleavage of the Co–H bond of $[Co^{III}(H)(Ch)(CH_3COO)]^-$ or O–H bond of CH_3COOH is involved in the rate-determining step of the reaction of $[Co^I(Ch)]^-$ with CH_3COOH. Because CH_3COOH is not involved in the rate-determining step (*vide infra*), the cleavage of the Co–H bond of $[Co^{III}(H)(Ch)(CH_3COO)]^-$ is the rate-determining step of the reaction of $[Co^I(Ch)]^-$ with CH_3COOH. The KIE value was 1.7 which is virtually the same as observed for the photocatalytic H_2 evolution (KIE = 1.8, Fig. 1), indicating that the heterolytic Co–H bond cleavage of $[Co^{III}(H)(Ch)(CH_3COO)]^-$ is also the rate-determining step in the photocatalytic H_2 evolution.

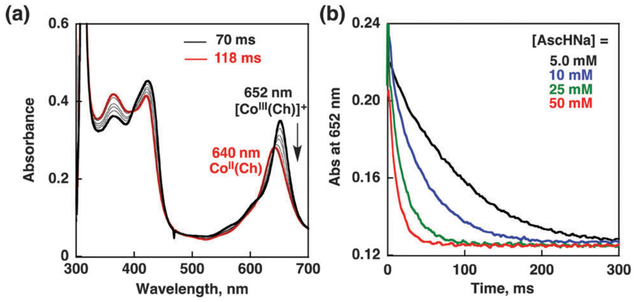
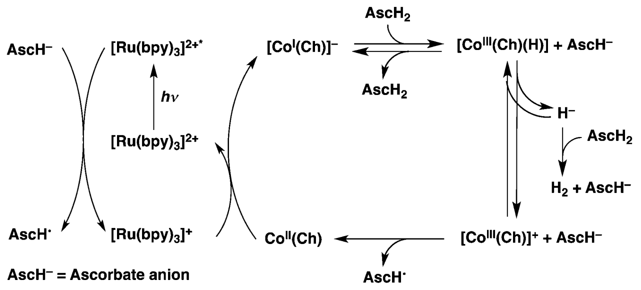


Fig. 4 (a) UV-vis absorption spectral changes in the electron-transfer reduction of $[Co^{III}(Ch)]^+$ (15 μM) with AscHNa (50 mM) in air-saturated H_2O/MeCN mixed solutions (1 : 1 v/v) at 298 K taken at 70 ms and 118 ms after mixing. (b) Decay time profiles of absorbance at 652 nm due to $[Co^{III}(Ch)]^+$ in the presence of various concentrations of AscHNa in air-saturated H_2O/MeCN mixed solutions (1 : 1 v/v) at 298 K.



Scheme 2 Mechanism of photocatalytic hydrogen evolution from $AscH^-$ and $AscH_2$ with $[Ru(bpy)_3]^{2+}$ and $Co^{II}(Ch)$.

$[Co^{III}(Ch)]^+$ produced by the reaction of $[Co^{III}(H)(Ch)(CH_3COO)]^-$ with CH_3COOH is reduced by $AscH^-$ to form $Co^{II}(Ch)$ as shown by stopped-flow measurements in Fig. 4.[27] The rate constant of electron transfer from $AscH^-$ to $[Co^{III}(Ch)]^+$ that was prepared by the one-electron oxidation of $Co^{II}(Ch)$ with $(p\text{-}BrC_6H_4)_3N^{\bullet +}SbCl_6{}^-$ in H_2O/MeCN was determined to be 1.5×10^3 M^{-1} s^{-1} from the linear dependence of the first-order rate constant on concentration of $AscH^-$ (Fig. S11 in ESI†).

The photocatalytic cycle is summarized in Scheme 2. Photoexcitation of $[Ru(bpy)_3]^{2+}$ resulted in electron transfer from $AscH^-$ to $[Ru(bpy)_3]^{2+*}$ to produce $[Ru(bpy)_3]^+$, followed by electron transfer from $[Ru(bpy)_3]^+$ to $Co^{II}(Ch)$ to produce $[Co^I(Ch)]^-$, which reacts with $AscH_2$ to produce $[Co^{III}(H)(Ch)(AscH)]^-$. Hydrogen is generated by the reaction of $[Co^{III}(H)(Ch)(AscH)]^-$ with $AscH_2$ *via* the Co–H bond heterolysis to produce $[Co^{III}(Ch)]^+$,[28,29] which is reduced by $AscH^-$ to regenerate $Co^{II}(Ch)$. In such a case, a one-photon two-electron process is made possible, because one photon is required to produce $[Co^I(Ch)]^-$ for H_2 evolution and another electron is provided thermally by $AscH^-$.

In conclusion, $Co^{II}(Ch)$ acts as an efficient catalyst for photocatalytic H_2 evolution from ascorbic acid with $[Ru(bpy)_3]^{2+}$ as a photocatalyst to attain the high quantum yield *via* a one-photon two-electron process in which the second electron is provided thermally from ascorbic acid.

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24 The spectrum of $[\mathrm{Co^{III}(Ch)}]^+$ obtained by the reaction $[\mathrm{Co^{I}(Ch)}]^-$ with CH_3COOH was identical to that of $[\mathrm{Co^{III}(Ch)}]^+$ prepared by the electron-transfer oxidation of $\mathrm{Co^{II}(Ch)}$ by a one-electron oxidizing reagent of $(p\text{-}\mathrm{BrC_6H_4})_3\mathrm{N}^{\bullet+}\mathrm{SbCl_6}^-$ (E_{red} = 1.05 V *vs.* SCE).[19].

25 $[\mathrm{Co^{III}(Ch)}]^+$ or $\mathrm{Co^{III}(H)(Ch)}$ species is not re-reduced by large excess of $\mathrm{Co(Cp^*)_2}$, under the present reaction conditions because $\mathrm{Co^{II}(Ch)}$ with 20 molar equiv. of $\mathrm{Co(Cp^*)_2}$ is necessary to quantitatively produce $[\mathrm{Co^{I}(Ch)}]^-$ as shown in ESI,† Fig. S7. $\mathrm{Co(Cp^*)_2}$ ($E_{1/2}{}^{+/0}$ = −1.47 V *vs.* SCE) is unstable even in carefully degassed and dehydrated MeCN.

26 The KIE value was determined from the k_{obs} values at $[\mathrm{CH_3COOH}] = [\mathrm{CH_3COOD}]$ = 1.0 M.

27 Neither oxidation of $[\mathrm{Co^{III}(Ch)}]^+$ nor O_2 reduction was observed under the basic reaction conditions.

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