

Cite this: *Chem. Sci.*, 2017, 8, 7119

# Photocatalytic oxidation of benzene to phenol using dioxygen as an oxygen source and water as an electron source in the presence of a cobalt catalyst†

Ji Won Han,<sup>‡a</sup> Jieun Jung,<sup>‡ab</sup> Yong-Min Lee,<sup>ID a</sup> Wonwoo Nam<sup>ID \*a</sup>  
and Shunichi Fukuzumi<sup>ID \*ac</sup>

Photocatalytic hydroxylation of benzene to phenol by dioxygen (O<sub>2</sub>) occurs under visible light irradiation of an O<sub>2</sub>-saturated acetonitrile solution containing [Ru<sup>II</sup>(Me<sub>2</sub>phen)<sub>3</sub>]<sup>2+</sup> as a photocatalyst, [Co<sup>III</sup>(Cp\*)(bpy)(H<sub>2</sub>O)]<sup>2+</sup> as an efficient catalyst for both the water oxidation and benzene hydroxylation reactions, and water as an electron source in the presence of Sc(NO<sub>3</sub>)<sub>3</sub>. The present study reports the first example of photocatalytic hydroxylation of benzene with O<sub>2</sub> and H<sub>2</sub>O, both of which are the most green reagents, under visible light irradiation to afford a high turnover number (*e.g.*, >500). Mechanistic studies revealed that the photocatalytic reduction of O<sub>2</sub> to H<sub>2</sub>O<sub>2</sub> is the rate-determining step, followed by efficient catalytic hydroxylation of benzene to phenol with H<sub>2</sub>O<sub>2</sub>, paving a new way for the photocatalytic oxygenation of substrates by O<sub>2</sub> and water.

Received 4th June 2017  
Accepted 21st August 2017

DOI: 10.1039/c7sc02495a

rsc.li/chemical-science

## Introduction

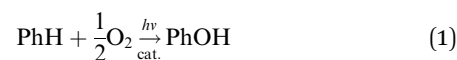
Phenol, which is an important precursor for many chemicals and industrial products (*e.g.*, dyes, polymers, *etc.*), is currently produced from benzene by a three step cumene process.<sup>1,2</sup> Since the efficiency of the cumene process is low (~5% yield of phenol) under severe conditions (*e.g.*, high temperature, high pressure, and strong acidic conditions), it is highly desired to develop a one-step synthesis of phenol from benzene using homogeneous and heterogeneous inorganic catalysts.<sup>3–5</sup> Among various oxidants, hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) is frequently used as a green oxidant, which produces water or dioxygen (O<sub>2</sub>) as products, in catalytic benzene hydroxylation.<sup>6–13</sup> O<sub>2</sub> is a more ideal oxidant than H<sub>2</sub>O<sub>2</sub> because of its abundance in nature, low cost, and environmental benignity.<sup>4,14,15</sup> However, catalytic aerobic oxidation of benzene to phenol has required sacrificial reducing agents such as H<sub>2</sub>, ascorbic acid, and NADH analogs.<sup>16</sup> In addition, the catalytic aerobic oxidation of benzene without sacrificial reducing agents has so far required harsh conditions, such as high temperatures or UV light photoirradiation.<sup>14,15,17,18</sup> Moreover,

although the photocatalytic hydroxylation of benzene to phenol using organic photocatalysts without overoxidation has been demonstrated,<sup>19–22</sup> the turnover numbers (TONs) of the catalytic hydroxylation of benzene to phenol are still low (*e.g.*, 13).<sup>20</sup>

We report herein an efficient photocatalytic hydroxylation of benzene to phenol (PhOH) using O<sub>2</sub> as an oxidant in the presence of Sc(NO<sub>3</sub>)<sub>3</sub> and utilizing [Ru<sup>II</sup>(Me<sub>2</sub>phen)<sub>3</sub>]<sup>2+</sup> (Me<sub>2</sub>phen = 4,7-dimethyl-1,10-phenanthroline) as a photocatalyst, water as an electron source, and [Co<sup>III</sup>(Cp\*)(bpy)(H<sub>2</sub>O)]<sup>2+</sup> (Cp\* = η<sup>5</sup>-pentamethylcyclopentadienyl and bpy = 2,2-bipyridine) as an efficient catalyst for water oxidation as well as benzene hydroxylation to attain a TON of greater than 500 (Scheme 1). Mechanistic aspects of the benzene hydroxylation reaction under the visible light irradiation conditions have been discussed as well.

## Results and discussion

[Co<sup>III</sup>(Cp\*)(bpy)(H<sub>2</sub>O)]<sup>2+</sup> (**1**) and [Ru<sup>II</sup>(Me<sub>2</sub>phen)<sub>3</sub>]<sup>2+</sup> were synthesised according to the literature methods.<sup>23,24</sup> Visible light irradiation of an O<sub>2</sub>-saturated solution of CH<sub>3</sub>CN (MeCN) and H<sub>2</sub>O (v/v = 23 : 2) containing a catalytic amount of **1** (1.0 μM), [Ru<sup>II</sup>(Me<sub>2</sub>phen)<sub>3</sub>]<sup>2+</sup> (1.0 mM), Sc(NO<sub>3</sub>)<sub>3</sub> (100 mM), and benzene (1.0 M) at 298 K resulted in the formation of PhOH [eqn (1)],



as shown in Fig. 1. PhOH is produced *via* the catalytic oxygenation of hydrogen peroxide (H<sub>2</sub>O<sub>2</sub>) with **1**, which acts as not only a benzene oxygenation catalyst but also a water oxidation catalyst

<sup>a</sup>Department of Chemistry and Nano Science, Ewha Womans University, Seoul 03760, Korea. E-mail: fukuzumi@chem.eng.osaka-u.ac.jp; wwnam@ewha.ac.kr

<sup>b</sup>Department of Chemistry, Graduate School of Science, Nagoya University, Chikusa, Nagoya 464-8602, Japan

<sup>c</sup>Faculty of Science and Engineering, Meijo University, SENTAN, Japan Science and Technology Agency (JST), Nagoya, Aichi 468-8502, Japan

† Electronic supplementary information (ESI) available: Table S1 and Fig. S1–S8. See DOI: 10.1039/c7sc02495a

‡ These authors contributed equally to this work.





The photocatalytic hydroxylation reaction proceeds as shown in Scheme 1. Visible light irradiation of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  resulted in the formation of  $\text{H}_2\text{O}_2$  by oxidative quenching of the photoexcited state  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+*}$  (\* denotes the excited state) with  $\text{O}_2$  in the presence of  $\text{Sc}(\text{NO}_3)_3$  (Scheme 1, reaction pathways b and c), as reported for the case of  $[\text{Ru}^{\text{II}}(\text{bpy})_3]^{2+}$ .<sup>26</sup> The time courses of  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  generation and  $\text{H}_2\text{O}_2$  production were obtained to give the stoichiometry of the photochemical reaction, as shown in eqn (3) (Fig. 3).



The lifetime of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+*}$  and emission quenching have been determined to obtain the rate constants ( $k_{\text{et}}$ ) of photoinduced electron transfer in the absence and presence of  $\text{Sc}(\text{NO}_3)_3$  (Fig. S5 and Table S1, ESI†). The  $k_{\text{et}}$  values in both the absence and presence of  $\text{Sc}(\text{NO}_3)_3$  are close to the diffusion-limited value, showing that  $\text{Sc}(\text{NO}_3)_3$  does not affect the oxidative quenching of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+*}$  by  $\text{O}_2$ . The emission spectra of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  in the absence and presence of  $\text{Sc}(\text{NO}_3)_3$  taken in a solvent mixture of MeCN and  $\text{H}_2\text{O}$  ( $v/v = 23 : 2$ ) containing different concentrations of  $\text{O}_2$  indicate that the amount of  $\text{O}_2$  in air is large enough for efficient  $\text{H}_2\text{O}_2$  photogeneration (Fig. S5d, ESI†).

Visible light irradiation of the reaction solution without benzene results in the formation of  $\text{H}_2\text{O}_2$ , as shown in Fig. 4 (Scheme 1, reaction pathways a–c), and the production of  $\text{H}_2\text{O}_2$  is obtained from the photooxidation of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  by  $\text{O}_2$  [eqn (3)] and the catalytic water oxidation by  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  in the presence of **1** [eqn (4)].<sup>23</sup> Thus, the overall reaction



is the photocatalytic oxidation of  $\text{H}_2\text{O}$  by  $\text{O}_2$  to produce  $\text{H}_2\text{O}_2$ , as given by eqn (5).



Fig. 3 Time courses of the concentrations of  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  (red) and  $\text{H}_2\text{O}_2$  (blue) produced from  $\text{H}_2\text{O}$  and  $\text{O}_2$  in the photocatalytic oxidation of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  by  $\text{O}_2$  under photoirradiation (white light) of an air-saturated solvent mixture of MeCN and  $\text{H}_2\text{O}$  ( $v/v = 23 : 2$ ) containing  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  (100  $\mu\text{M}$ ) in the presence of  $\text{Sc}(\text{NO}_3)_3$  (100 mM) at 298 K.

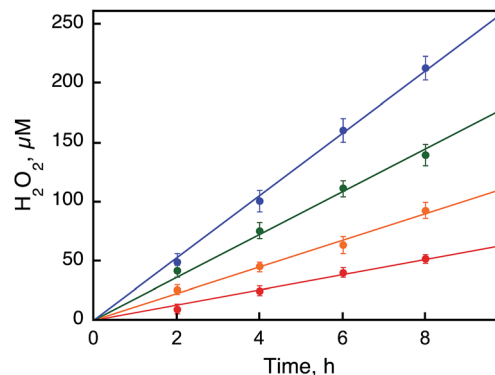
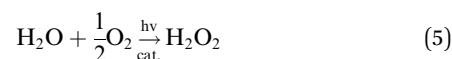
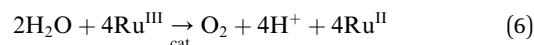


Fig. 4 Time courses of the photocatalytic production of  $\text{H}_2\text{O}_2$  from  $\text{H}_2\text{O}$  and  $\text{O}_2$  with  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  (0.10 mM) in the presence of a catalytic amount of **1** [0.50 mM (red), 1.0 mM (orange), 1.5 mM (green), and 2.0 mM (blue)] in an air-saturated solvent mixture of MeCN and  $\text{H}_2\text{O}$  ( $v/v = 23 : 2$ ) containing  $\text{Sc}(\text{NO}_3)_3$  (100 mM) under photoirradiation (white light) at 298 K.



The rates of photocatalytic oxidation of  $\text{H}_2\text{O}$  by  $\text{O}_2$  to produce  $\text{H}_2\text{O}_2$  in the presence of **1** were also investigated with various concentrations of **1** at 298 K (Fig. S6, ESI†). The zeroth-order rate constants ( $k_{\text{obs}}$ ) were obtained from the initial slopes of the plots of the amount of  $\text{H}_2\text{O}_2$  produced vs. time, being proportional to the concentration of **1** (Fig. 4 and S6, ESI†).

The catalytic water oxidation by  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  with **1** to evolve  $\text{O}_2$  [eqn (6); Scheme 1, reaction pathway a]



was independently confirmed, where  $\text{O}_2$  was evolved, accompanied by the formation of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$ . The evolved  $\text{O}_2$  concentrations were investigated by a Clark oxygen electrode, showing an increase in the amount of  $\text{O}_2$  evolved after addition of  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  (1.0 mM) to a solvent mixture of MeCN and  $\text{H}_2\text{O}$  (1.0 mL;  $v/v = 9 : 1$ ) containing  $\text{Sc}(\text{NO}_3)_3$  (100 mM) and various concentrations of **1** (Fig. 5), where the  $\text{O}_2$  yield increased with an increasing concentration of **1** to approach 100% (1/4 of the initial concentration of  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$ ). The reaction rate was accelerated with an increasing concentration of **1** to account for oxidation of water by  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  with **1** under acidic conditions due to the presence of  $\text{Sc}(\text{NO}_3)_3$ . Formation of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  accompanied by water oxidation was also confirmed by monitoring the UV-vis spectral changes (Fig. S7a, ESI†). The time courses of the oxidation of **1** by  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  were obtained by spectral changes at  $\lambda = 445$  nm due to  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$ , showing that the rate of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  formation from  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  increased with the increase of the concentration of **1** (Fig. S7b, ESI†).

Catalytic oxidation of benzene to phenol with  $\text{H}_2\text{O}_2$  [eqn (7)]

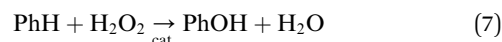




Fig. 5 Time courses of  $\text{O}_2$ -evolution observed in the catalytic oxidation of  $\text{H}_2\text{O}$  by  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  (1.0 mM) with **1** [10  $\mu\text{M}$  (red), 50  $\mu\text{M}$  (orange), 100  $\mu\text{M}$  (green), and 200  $\mu\text{M}$  (blue)] in the presence of  $\text{Sc}(\text{NO}_3)_3$  (100 mM) in a deaerated solvent mixture (1.0 mL) of MeCN and  $\text{H}_2\text{O}$  ( $v/v = 9 : 1$ ) at 298 K. The induction periods were observed due to the reaction time to oxidize **1** by  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  to produce the catalytically active species.

has also been confirmed in a solvent mixture of MeCN and  $\text{H}_2\text{O}$  (23 : 2) containing  $\text{Sc}(\text{NO}_3)_3$  in order to stabilize  $\text{H}_2\text{O}_2$  with the same conditions for the overall photocatalytic reaction as shown in Fig. 6 (Scheme 1, reaction pathway d). The rate of benzene hydroxylation increased with increasing concentrations of **1** (Fig. 6) and  $\text{H}_2\text{O}_2$  (Fig. S8, ESI<sup>†</sup>). A combination of the photocatalytic oxidation of  $\text{H}_2\text{O}$  by  $\text{O}_2$  to produce  $\text{H}_2\text{O}_2$  [eqn (5)] and the catalytic hydroxylation of benzene to phenol by  $\text{H}_2\text{O}_2$  [eqn (7)] affords the overall photocatalytic oxidation of benzene to phenol by  $\text{O}_2$  [eqn (1)]. The photocatalytic reactions of an  $\text{O}_2$ -saturated solution of MeCN and  $\text{H}_2\text{O}$  ( $v/v = 23 : 2$ ) containing **1** (0.50 mM),  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  (0.10 mM), and  $\text{Sc}(\text{NO}_3)_3$  (100 mM) under photoirradiation (white light) were examined in the absence and presence of benzene to compare their reactivities (Fig. 7). The rate of  $\text{H}_2\text{O}_2$  production from  $\text{H}_2\text{O}$  and  $\text{O}_2$  in the absence of benzene was slightly higher than that of PhOH generation in the photocatalytic hydroxylation of benzene, indicating that  $\text{H}_2\text{O}_2$  produced in the photocatalytic oxidation



Fig. 6 Time courses of the catalytic hydroxylation of benzene to PhOH with  $\text{H}_2\text{O}_2$  (1.0 mM) in the presence of **1** [10  $\mu\text{M}$  (black), 50  $\mu\text{M}$  (blue), 100  $\mu\text{M}$  (green), and 500  $\mu\text{M}$  (red)] in a solvent mixture of MeCN and  $\text{H}_2\text{O}$  ( $v/v = 23 : 2$ ) containing benzene (1.0 M) and  $\text{Sc}(\text{NO}_3)_3$  (100 mM) at 298 K.

of  $\text{H}_2\text{O}$  by  $\text{O}_2$  reacted with benzene efficiently in the presence of **1** to produce PhOH.<sup>27</sup> In fact, the rate of PhOH production (59  $\mu\text{M}$  after 1 h) from  $\text{H}_2\text{O}_2$  (1.0 mM) and benzene (1.0 M) with **1** (0.50 mM, red dots in Fig. 6) is much faster than that of PhOH production (estimated to be 7.5  $\mu\text{M}$  after 1 h) in the photocatalytic reaction (Fig. 7), because the concentration of  $\text{H}_2\text{O}_2$  produced in the photocatalytic oxidation of  $\text{H}_2\text{O}$  by  $\text{O}_2$  is much smaller than 1.0 mM. Since the rate of PhOH production from  $\text{H}_2\text{O}_2$  is proportional to the concentration of  $\text{H}_2\text{O}_2$  (Fig. S8, ESI<sup>†</sup>), the rate of PhOH production in the photocatalytic hydroxylation of benzene by  $\text{O}_2$  corresponds to that of PhOH production in the catalytic oxidation of benzene with 127  $\mu\text{M}$   $\text{H}_2\text{O}_2$ , which may be the steady state concentration during the photocatalytic hydroxylation of benzene.

Then,  $^{18}\text{O}_2$ -labeling experiments were performed to confirm that the oxygen atom in the phenol product in eqn (1) derives from  $\text{O}_2$  (Fig. 8). Indeed, Ph $^{18}\text{OH}$  (73%) was formed as the major product together with Ph $^{16}\text{OH}$  (27%) in the  $^{18}\text{O}_2$ -labeling experiments, indicating that the oxygen atom in the PhOH product obtained in the photocatalytic oxidation of benzene by  $\text{H}_2^{18}\text{O}_2$  derived from  $^{18}\text{O}_2$  (98%  $^{18}\text{O}$ -enriched; Fig. 8). The formation of Ph $^{16}\text{OH}$  (27%) indicates that  $^{16}\text{O}_2$  was produced by the oxidation of  $\text{H}_2^{16}\text{O}$  by  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$ , leading to the production of  $\text{H}_2^{16}\text{O}_2$  to oxidize benzene to produce Ph $^{16}\text{OH}$  instead of Ph $^{18}\text{OH}$  (Scheme 1, reaction pathway a). The  $^{18}\text{O}$ -labeling experiments were also performed by replacing  $\text{H}_2^{16}\text{O}$  with  $\text{H}_2^{18}\text{O}$  to support that the oxygen atom in the phenol product derives from  $\text{O}_2$  rather than  $\text{H}_2\text{O}$  (Fig. 9). In this case, Ph $^{16}\text{OH}$  (66%) was the major product together with Ph $^{18}\text{OH}$  (34%), consistent with the  $^{18}\text{O}_2$ -labeling experiments described above.

The photocatalytic mechanism of benzene hydroxylation to phenol by  $\text{O}_2$  is summarized in Scheme 1. The photoexcitation of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  in an  $\text{O}_2$ -saturated solvent mixture of MeCN and  $\text{H}_2\text{O}$  ( $v/v = 23 : 2$ ) containing  $\text{Sc}(\text{NO}_3)_3$  resulted in electron transfer from the triplet excited state of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  to  $\text{O}_2$  to produce  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  and the  $\text{O}_2^{\cdot-}$ - $\text{Sc}^{3+}$  complex.<sup>23,28</sup> The strong binding of  $\text{Sc}^{3+}$  to  $\text{O}_2^{\cdot-}$ , which was

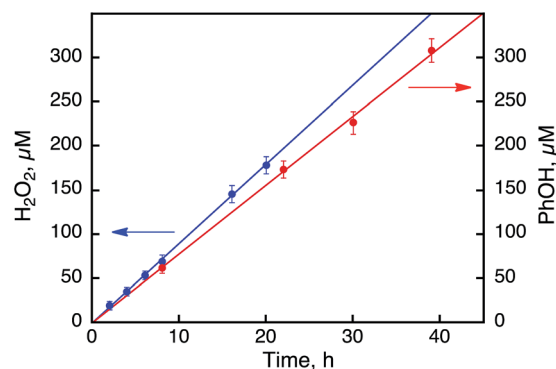


Fig. 7 Time courses of the products from the photoinduced reaction under photoirradiation (white light) in an  $\text{O}_2$ -saturated solvent mixture of MeCN and  $\text{H}_2\text{O}$  ( $v/v = 23 : 2$ ) containing **1** (0.50 mM),  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  (0.10 mM), and  $\text{Sc}(\text{NO}_3)_3$  (100 mM) in the absence (blue) and presence (red) of benzene (1.0 M) at 298 K.



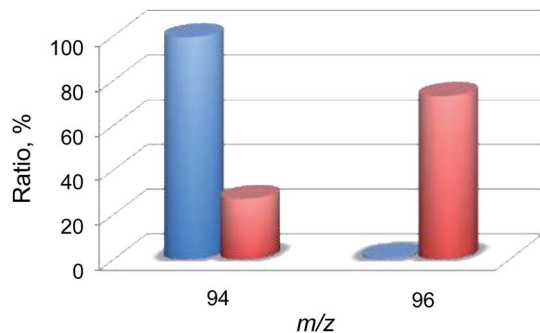


Fig. 8 A comparison of the relative abundances of authentic PhOH (blue) and the PhOH product (red) yielded in the photocatalytic hydroxylation of benzene by **1** (0.10 mM) in the presence of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  (1.0 mM),  $\text{Sc}(\text{NO}_3)_3$  (100 mM), and benzene (2.0 M) under photoirradiation (white light) in an  $^{18}\text{O}_2$ -saturated solvent mixture of MeCN and  $\text{H}_2\text{O}$  (v/v = 23 : 2) at 298 K for 24 h. The peaks at  $m/z = 94$  and 96 correspond to  $\text{Ph}^{16}\text{OH}$  and  $\text{Ph}^{18}\text{OH}$ , respectively.

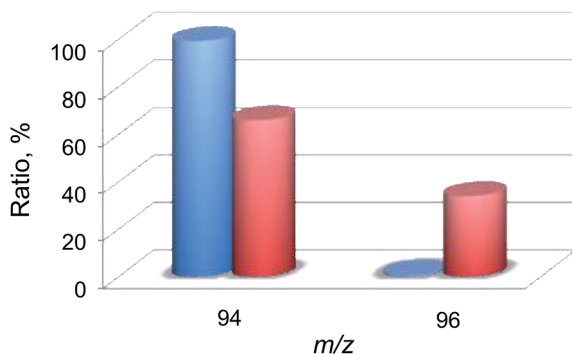


Fig. 9 A comparison of the relative abundances of authentic PhOH (blue) and the produced PhOH (red) in the photocatalytic hydroxylation of benzene by  $\text{O}_2$  with **1** (0.10 mM) in the presence of  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  (1.0 mM),  $\text{Sc}(\text{NO}_3)_3$  (100 mM), and benzene (2.0 M) under photoirradiation (white light) of an  $^{16}\text{O}_2$ -saturated solvent mixture of MeCN and  $\text{H}_2^{18}\text{O}$  (v/v = 23 : 2) at 298 K for 24 h. The peaks at  $m/z = 94$  and 96 correspond to  $\text{Ph}^{16}\text{OH}$  and  $\text{Ph}^{18}\text{OH}$ , respectively.

detected by EPR, inhibits the back electron transfer from the  $\text{O}_2^- - \text{Sc}^{3+}$  complex to  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$ , followed by disproportion with  $\text{H}_2\text{O}$  to produce  $\text{H}_2\text{O}_2$ .<sup>23</sup>  $\text{H}_2\text{O}_2$  has attracted considerable attention as an ideal solar fuel for a one-compartment  $\text{H}_2\text{O}_2$  fuel cell with a theoretical maximum output potential of 1.09 V, which is comparable to that of a hydrogen fuel cell (1.23 V).<sup>23</sup>  $[\text{Ru}^{\text{III}}(\text{Me}_2\text{phen})_3]^{3+}$  can oxidize  $\text{H}_2\text{O}$  to  $\text{O}_2$  with catalysis by **1**. Benzene hydroxylation to PhOH was also catalysed by **1**. As reported previously, the NMR peaks assignable to the bpy and Cp\* of **1** remained the same after the photocatalytic production of  $\text{H}_2\text{O}_2$ , indicating that **1** acted as a homogeneous catalyst during the reaction.<sup>23</sup> However, the catalytically active intermediates such as the Co(IV)-oxo species<sup>29</sup> have yet to be detected in the catalytic benzene hydroxylation with  $\text{H}_2\text{O}_2$ . There was no deuterium kinetic isotope effect (KIE) for benzene (Fig. 2), indicating that C–H bond cleavage is not involved in the rate-determining step of the photocatalytic hydroxylation reaction.

## Conclusions

In conclusion, we have shown that benzene is oxidized to phenol by dioxygen with a high TON (e.g., 500) under visible light irradiation of a reaction solution containing  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$  as a photocatalyst,  $[\text{Co}^{\text{III}}(\text{Cp}^*)(\text{bpy})(\text{H}_2\text{O})]^{2+}$  as an efficient catalyst for both water oxidation and benzene hydroxylation, and water as an electron source in the presence of  $\text{Sc}(\text{NO}_3)_3$ . The combination of the photocatalytic  $\text{H}_2\text{O}_2$  production by  $\text{H}_2\text{O}$  oxidation with  $\text{O}_2$  and the catalytic hydroxylation of benzene to phenol with  $\text{H}_2\text{O}_2$  demonstrated in this study has paved a new road towards direct oxygenation of substrates by  $\text{O}_2$  as the most environmentally benign oxidant as well as oxygen source without any electron source except  $\text{H}_2\text{O}$ , which is also the most environmentally benign hydrogen source.

## Experimental section

### Materials

All solvents and chemicals were of reagent-grade quality, obtained commercially and used without further purification, unless otherwise noted. Ruthenium(III) chloride hydrate, ammonium hexafluorophosphate, *n*-butyllithium solution (2.7 M in heptane), *n*-pentane, and tetrahydrofuran were purchased from Aldrich Chemicals. The chemicals, such as 4,7-dimethyl-1,10-phenanthroline ( $\text{Me}_2\text{phen}$ ), silver sulphate, lead dioxide, and cobalt(II) chloride, were purchased from Alfa Aesar. 2,2'-Bipyridine, 1,2,3,4,5-pentamethylcyclopentadiene, and  $\text{Ti}^{\text{IV}}(\text{O})(\text{tpyp})$  ( $\text{tpyp} = 5,10,15,20\text{-tetra}(4\text{-pyridyl})\text{porphyrinato anion}$ ) were obtained from Tokyo Chemical Industry Co., Ltd.  $\text{Sc}(\text{NO}_3)_3 \cdot 4\text{H}_2\text{O}$  was supplied by Mitsuwa Chemicals Co., Ltd.  $^{18}\text{O}_2$  gas (98%  $^{18}\text{O}$ -enriched) was purchased from ICON Services Inc. (Summit, NJ, USA). The purification of water (18.2 M $\Omega$  cm) was performed with a Milli-Q system (Millipore, Direct-Q 3 UV). Acetonitrile was dried according to published procedures and distilled prior to use.<sup>30</sup> The cobalt(III) starting complex,  $[\text{Co}^{\text{III}}(\text{Cp}^*)(\text{bpy})(\text{H}_2\text{O})]^{2+}$  (**1**, Cp\* =  $\eta^5$ -pentamethylcyclopentadienyl and bpy = 2,2'-bipyridine), and the tris(4,7-dimethyl-1,10-phenanthroline)ruthenium(II) complex,  $[\text{Ru}^{\text{II}}(\text{Me}_2\text{phen})_3]^{2+}$ , were prepared according to the published methods.<sup>25,31</sup>

### Instrumentation

UV-vis spectra were recorded on a Hewlett Packard 8453 diode array spectrophotometer equipped with a UNISOKU Scientific Instruments Cryostat USP-203A. Product analysis for the oxidation reactions was performed with an Agilent Technologies 6890N gas chromatograph (GC) and a Thermo Finnigan (Austin, Texas, U.S.A.) FOCUS DSQ (dual stage quadrupole) mass spectrometer interfaced with a Finnigan FOCUS gas chromatograph (GC-MS). The amount of evolved oxygen was recorded by a Clark-type oxygen electrode made by Hansatech Ltd.



## Product analysis

The products formed in the oxidation of benzene by **1** and O<sub>2</sub> in the presence of [Ru<sup>II</sup>(Me<sub>2</sub>phen)<sub>3</sub>]<sup>2+</sup> and Sc(NO<sub>3</sub>)<sub>3</sub> in a solvent mixture of MeCN and H<sub>2</sub>O (v/v = 23 : 2) at 298 K were identified by GC and GC-MS by a comparison of the mass peaks and retention time of the products with respect to the authentic samples, and the product yields were determined by comparing the responsive peak areas of the sample products against standard curves prepared with known authentic compounds using the internal standard decane. The quantum yield (QY) of the photocatalytic hydroxylation of benzene has been determined under visible light irradiation of monochromatized light using a Compact Xenon Light Source (MAX-302; Asahi Spectra Co., Ltd). The amount of hydrogen peroxide produced was determined by spectroscopic titration with an acidic solution of a [Ti<sup>IV</sup>O(tpypH<sub>4</sub>)]<sup>4+</sup> complex.<sup>31</sup> [Ti<sup>IV</sup>O(tpypH<sub>4</sub>)]<sup>4+</sup> (50 μM) was prepared by dissolving 3.4 mg of the TiO(tpyp) complex into water (100 mL) containing hydrochloric acid (50 mM). A small portion (100 μL) of the photocatalytic reaction solution was taken and diluted up to 1.0 mL with water. To 0.25 mL of the diluted sample, 0.25 mL of perchloric acid (4.8 M) and 0.25 mL of [Ti<sup>IV</sup>O(tpypH<sub>4</sub>)]<sup>4+</sup> (50 μM) were added. The mixed solution was then allowed to stand for 5 min at room temperature. This sample solution was diluted up to 2.5 mL with water and used for the spectroscopic measurement. The absorbance at λ = 434 nm (A<sub>S</sub>) due to [Ti<sup>IV</sup>O(tpypH<sub>4</sub>)]<sup>4+</sup> was measured using a Hewlett Packard 8453 diode array spectrophotometer. In a similar manner, a blank solution was prepared by adding distilled water instead of the sample solution in the same volume with its absorbance designated as A<sub>B</sub>. The difference in absorbance at 434 nm was determined as follows: ΔA<sub>434</sub> = A<sub>B</sub> - A<sub>S</sub>. Based on ΔA<sub>434</sub> and the volume of the solution, the amount of hydrogen peroxide was determined according to the literature.<sup>23</sup> A Clark-type oxygen electrode was used to obtain oxygen evolution data, and calibrated daily using Ar deoxygenated and oxygen saturated atmospheric solutions. Clark electrode experiments were performed by adding 10 μL of [Ru<sup>III</sup>(Me<sub>2</sub>phen)<sub>3</sub>]<sup>3+</sup> (100 mM) to an O<sub>2</sub>-saturated solvent mixture (1.0 mL) of MeCN and H<sub>2</sub>O (v/v = 9 : 1) containing **1** (10–200 μM) and monitoring the O<sub>2</sub> evolved.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was supported by SENTAN projects from the Japan Science and Technology Agency (JST), JSPS KAKENHI (No. 16H02268) to S. F. and by the CRI (NRF-2012R1A3A2048842 to W. N.), GRL (NRF-2010-00353 to W. N.), and Basic Science Research Program (2017R1D1A1B03029982 to Y. M. L. and 2017R1D1A1B03032615 to S. F.) through the NRF of Korea.

## Notes and references

- M. Weber, M. Weber and M. Kleine-Boymann, *Phenol. Ullmann's Encyclopedia of Industrial Chemistry*, Wiley-VCH Verlag GmbH & Co. KGaA, Weinheim, Germany, 2004, vol. 26, p. 503.
- (a) R. Molinari and T. Poerio, *Asia-Pac. J. Chem. Eng.*, 2010, **5**, 191; (b) R. J. Schmidt, *Appl. Catal., A*, 2005, **280**, 89.
- (a) L. Balducci, D. Bianchi, R. Bortolo, R. D'Aloisio, M. Ricci, R. Tassinari and R. Ungarelli, *Angew. Chem., Int. Ed.*, 2003, **42**, 4937; (b) J.-H. Yang, G. Sun, Y. Gao, H. Zhao, P. Tang, J. Tan, A.-H. Lu and D. Ma, *Energy Environ. Sci.*, 2013, **6**, 793.
- S. Fukuzumi and K. Ohkubo, *Asian J. Org. Chem.*, 2015, **4**, 836.
- (a) X. Cai, Q. Wang, Y. Liu, J. Xie, Z. Long, Y. Zhou and J. Wang, *ACS Sustainable Chem. Eng.*, 2016, **4**, 4986; (b) W. Wang, L. Shi, N. Li and Y. Ma, *RSC Adv.*, 2017, **7**, 12738.
- (a) Y. Morimoto, S. Bunno, N. Fujieda, H. Sugimoto and S. Itoh, *J. Am. Chem. Soc.*, 2015, **137**, 5867; (b) Y. Aratani, Y. Yamada and S. Fukuzumi, *Chem. Commun.*, 2015, **51**, 4662.
- (a) D. Wang, M. Wang and Z. Li, *ACS Catal.*, 2015, **5**, 6852; (b) X. Ye, Y. Cui, X. Qiu and X. Wang, *Appl. Catal., B*, 2014, **152–153**, 383; (c) G. Wen, S. Wu, B. Li, C. Dai and D. S. Su, *Angew. Chem., Int. Ed.*, 2015, **54**, 4105.
- (a) M. Yamada, K. D. Karlin and S. Fukuzumi, *Chem. Sci.*, 2016, **7**, 2856; (b) B. Xu, W. Zhong, Z. Wei, H. Wang, J. Liu, L. Wu, Y. Feng and X. Liu, *Dalton Trans.*, 2014, **43**, 15337; (c) L. Wu, W. Zhong, B. Xu, Z. Wei and X. Liu, *Dalton Trans.*, 2015, **44**, 8013.
- (a) W. Ge, Z. Long, X. Cai, Q. Wang, Y. Zhou, Y. Xu and J. Wang, *RSC Adv.*, 2014, **4**, 45816; (b) C. Wang, L. Hu, Y. Hu, Y. Ren, X. Chen, B. Yue and H. He, *Catal. Commun.*, 2015, **68**, 1.
- (a) P. Borah, X. Ma, K. T. Nguyen and Y. Zhao, *Angew. Chem., Int. Ed.*, 2012, **51**, 7756; (b) P. K. Khatri, B. Singh, S. L. Jain, B. Sain and A. K. Sinha, *Chem. Commun.*, 2011, **47**, 1610; (c) Y. Leng, J. Liu, P. Jiang and J. Wang, *Chem. Eng. J.*, 2014, **239**, 1.
- (a) H. Wang, M. Zhao, Q. Zhao, Y. Yang, C. Wang and Y. Wang, *Ind. Eng. Chem. Res.*, 2017, **56**, 2711; (b) S. Verma, R. B. N. Baig, M. N. Nadagouda and R. S. Varma, *ACS Sustainable Chem. Eng.*, 2017, **5**, 3637.
- (a) L. Muniandy, F. Adama, A. R. Mohamed, A. Iqbal and N. R. A. Rahman, *Appl. Surf. Sci.*, 2017, **398**, 43; (b) J. Xu, Q. Jiang, T. Chen, F. Wu and Y.-X. Li, *Catal. Sci. Technol.*, 2015, **5**, 1504; (c) P. R. Makgwane and S. S. Raya, *J. Mol. Catal. A: Chem.*, 2015, **398**, 149.
- (a) L. Carneiro and A. R. Silva, *Catal. Sci. Technol.*, 2016, **6**, 8166; (b) E. Gu, W. Zhong and X. Liu, *RSC Adv.*, 2016, **6**, 98406; (c) E. Gu, W. Zhong, H. Ma, B. Xu, H. Wang and X. Liu, *Inorg. Chim. Acta*, 2016, **444**, 159; (d) B. Xu, W. Zhong, Z. Wei, H. Wang, J. Liu, L. Wu, Y. Feng and X. Liu, *Dalton Trans.*, 2014, **43**, 15337.
- (a) S. Niwa, M. Eswaramoorthy, J. Nair, A. Raj, N. Itoh, H. Shoji, T. Namba and F. Mizukami, *Science*, 2002, **295**,



- 105; (b) K. Sato, T. Hanaoka, S. Niwa, C. Stefan, T. Namba and F. Mizukami, *Catal. Today*, 2005, **104**, 260.
- 15 (a) A. Okemoto, Y. Tsukano, A. Utsunomiya, K. Taniya, Y. Ichihashi and S. Nishiyama, *J. Mol. Catal. A: Chem.*, 2016, **411**, 372; (b) B. B. Sarma, R. Carmieli, A. Collauto, I. Efremenko, J. M. L. Martin and R. Neumann, *ACS Catal.*, 2016, **6**, 6403.
- 16 (a) Z. Long, Y. Zhou, W. Ge, G. Chen, J. Xie, Q. Wang and J. Wang, *ChemPlusChem*, 2014, **79**, 1590; (b) Z. Long, Y. Zhou, G. Chen, P. Zhao and J. Wang, *Chem. Eng. J.*, 2014, **239**, 19; (c) S. Shang, B. Chen, L. Wang, W. Dai, Y. Zhang and S. Gao, *RSC Adv.*, 2015, **5**, 31965.
- 17 Y. Aratani, K. Oyama, T. Suenobu, Y. Yamada and S. Fukuzumi, *Inorg. Chem.*, 2016, **55**, 5780.
- 18 (a) R. Hamada, Y. Shibata, S. Nishiyama and S. Tsuruya, *Phys. Chem. Chem. Phys.*, 2003, **5**, 956; (b) Z. Long, Y. Zhou, G. Chen, W. Ge and J. Wang, *Sci. Rep.*, 2014, **4**, 3651; (c) Z. Long, Y. Liu, P. Zhao, Q. Wang, Y. Zhou and J. Wang, *Catal. Commun.*, 2015, **59**, 1; (d) S. S. Acharyya, S. Ghosh, R. Tiwari, C. Pendem, T. Sasaki and R. Bal, *ACS Catal.*, 2015, **5**, 2850; (e) R. Bal, M. Tada, T. Sasaki and Y. Iwasawa, *Angew. Chem., Int. Ed.*, 2006, **45**, 448.
- 19 (a) T. D. Bui, A. Kimura, S. Ikeda and M. Matsumura, *J. Am. Chem. Soc.*, 2010, **132**, 8453; (b) Y. Ide, M. Torii and T. Sano, *J. Am. Chem. Soc.*, 2013, **135**, 11784; (c) Y. Ide, M. Matsuoka and M. Ogawa, *J. Am. Chem. Soc.*, 2010, **132**, 16762.
- 20 K. Ohkubo, T. Kobayashi and S. Fukuzumi, *Angew. Chem., Int. Ed.*, 2011, **50**, 8652.
- 21 K. Ohkubo, A. Fujimoto and S. Fukuzumi, *J. Am. Chem. Soc.*, 2013, **135**, 5368.
- 22 K. Ohkubo, K. Hirose and S. Fukuzumi, *Chem.–Eur. J.*, 2015, **21**, 2855.
- 23 S. Kato, J. Jung, T. Suenobu and S. Fukuzumi, *Energy Environ. Sci.*, 2013, **6**, 3756.
- 24 (a) Y. Isaka, S. Kato, D. Hong, T. Suenobu, Y. Yamada and S. Fukuzumi, *J. Mater. Chem. A*, 2015, **3**, 12404; (b) Y. Isaka, K. Oyama, Y. Yamada, T. Suenobu and S. Fukuzumi, *Catal. Sci. Technol.*, 2016, **6**, 681.
- 25 (a) A. Das, V. Joshi, D. Kotkar, V. S. Pathak, V. Swayambunathan, P. V. Kamat and P. K. Ghosh, *J. Phys. Chem. A*, 2001, **105**, 6945; (b) D. Hong, J. Jung, J. Park, Y. Yamada, T. Suenobu, Y.-M. Lee, W. Nam and S. Fukuzumi, *Energy Environ. Sci.*, 2012, **5**, 7606.
- 26 It should be noted, however, that the contribution from the catalytic hydroxylation of benzene with  $[\text{Ru}^{\text{III}}(\text{bpy})_3]^{3+}$  in competition with  $\text{H}_2\text{O}$  oxidation cannot be ruled out.
- 27 (a) S. Fukuzumi and K. Ohkubo, *Chem.–Eur. J.*, 2000, **6**, 4532; (b) S. Fukuzumi, M. Patz, T. Suenobu, Y. Kuwahara and S. Itoh, *J. Am. Chem. Soc.*, 1999, **121**, 1605; (c) S. Fukuzumi, H. Ohtsu, K. Ohkubo, S. Itoh and H. Imahori, *Coord. Chem. Rev.*, 2002, **226**, 71.
- 28 W. L. F. Armarego and C. L. L. Chai, in *Purification of Laboratory Chemicals*, Pergamon Press, Oxford, U. K., 6th edn, 2009.
- 29 (a) S. Hong, F. F. Pfaff, E. Kwon, Y. Wang, M.-S. Seo, E. Bill, K. Ray and W. Nam, *Angew. Chem., Int. Ed.*, 2014, **53**, 10403; (b) B. Wang, Y.-M. Lee, W.-Y. Tcho, S. Tussupbayev, S.-T. Kim, Y. Kim, M. S. Seo, K.-B. Cho, Y. Dede, B. C. Keegan, T. Ogura, S. H. Kim, T. Ohta, M.-H. Baik, K. Ray, J. Shearer and W. Nam, *Nat. Commun.*, 2016, **8**, 14839.
- 30 C. Turro, J. M. Zaleski, Y. M. Karabatsos and D. G. Nocera, *J. Am. Chem. Soc.*, 1996, **118**, 6060.
- 31 C. Matsubara, N. Kawamoto and K. Takamura, *Analyst*, 1992, **117**, 1781.

