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towards bifunctional catalysts of tunable basicity

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Selective catalytic 2e⁻-oxidation of organic substrates by an FeII complex having an *N*-heterocyclic carbene ligand in water

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An FeII complex, 1, having a pentadentate ligand with an NHC moiety catalyzes substrate oxidation to afford 2e⁻-oxidized products with high selectivity by suppression of overoxidation in water. A Bell-Evance-Polanyi plot for the substrate oxidation catalyzed by 1 exhibited an inflection point around 86 kcal mol⁻¹, indicating strong C-H abstraction ability of the reactive species derived from 1.

High-valent Fe-oxo complexes have been well explored as important reactive species in oxidation reactions performed by metalloenzymes and synthetic oxidation reactions.[1–3] For example, nonheme iron enzymes such as α-ketoglutarate-dependent taurine dioxygenase (TauD)[4] generate an FeIV-oxo intermediate as the reactive species to hydroxylate taurine. The activation of C-H bonds by the FeIV-oxo species is proposed to occur through hydrogen-atom abstraction mechanism.[5,6] Oxidation enzymes such as TauD are able to oxidize substrates selectively to the corresponding 2e⁻-oxidized products.[4] A lot of synthetic FeIV-oxo complexes have been prepared and characterised to explore the reactivity of those complexes and to gain mechanistic insights into C-H oxidation.[7–9] However, these synthetic nonheme FeIV-oxo complexes do not perform selective 2e⁻ oxidation in organic solvents and aqueous media.[9] For example, an FeIV-oxo complex has been reported to oxidize ethylbenzene, toluene and cyclohexane to 1-phenylethanol (58% yield), benzaldehyde (58%) and cyclohexanone (57%), respectively, in CH₃CN.[9] Therefore, catalysts that can oxidize substrates to selectively afford 2e⁻-oxidation products in aqueous media are required to establish a sustainable oxidation strategy.

In general, a relationship between the activation energies for substrate oxidation and bond dissociation energies (BDEs) of the substrates can be analysed in light of the Bell-Evans-Polanyi (BEP) equation[10] as given bellow:

$$E_a = \alpha\Delta H + C \qquad (1)$$

In most cases, oxidation reactions of C-H bonds by metal-oxo complexes obeyed the BEP relation.[11,12] However, the improvement of product selectivity in substrate oxidation have never been examined on the basis of the difference in BDEs between substrates and the corresponding oxidation products.

On the other hand, Long and co-workers have reported selective 2e⁻-oxidation by low-spin FeIV-oxo complexes with a pentadentate pyridyl ligands in aqueous media.[13] Furthermore, they also have reported that an FeII-acetonitrile complex having a pentadentate ligand with an *N*-heterocyclic carbene (NHC) moiety at the axial position.[14] In this case, the NHC moiety of the ligand exerts strong σ-donation and *trans*-influence as represented by a longer coordination bond between the FeII centre and the CH₃CN ligand.[14] Influence of an NHC ligand located at the *trans* position to an oxo ligand on the reactivity should be of interest in terms of the regulation of the reactivity of a high-valent Ru-oxo complex.[15–17] For such a FeII-NHC complex,[14] however, the reactivity in substrate oxidation has yet to be explored.

Herein, we report a novel FeII-aqua complex having a pentadentate ligand with an NHC moiety, [Fe(PY₄Cl₂BIm)(OH₂)](NO₃)₂ (**1**·(NO₃)₂) (PY₄Cl₂BIm: 5,6-dichloro-1,3-dihydro-1,3-bis(2-pyridyl)-methyl-2*H*-benz-imidazol-2-ylidene) as an efficient catalyst for substrate oxidation in water. In this study, we have focused on the catalytic oxidation by using complex **1**·(NO₃)₂ as a catalyst, Na₂S₂O₈ as a sacrificial electron-transfer (ET) oxidant in water to observe selective 2e⁻-oxidation of substrates. Synthesis of **1**·(NO₃)₂ was performed as described in Scheme 1 and the details are given in the ESI†. Characterization of the low-spin FeII complex was made by ¹H NMR (Fig. S1) and UV-vis absorption
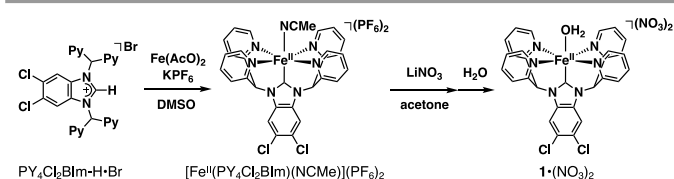
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† Electronic Supplementary Information (ESI) available: Further experimental details, supporting figures and tables. See DOI: 10.1039/x0xx00000x
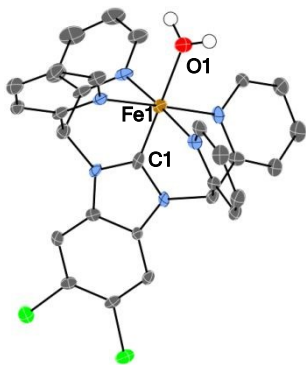
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Scheme 1. Synthesis of an Fe^{II}(NHC)-aqua complex, **1**·(NO$_3$)$_2$.

Fig. 1   An ORTEP drawing of the cation moiety of **1**·(ClO$_4$)$_2$ with 50% probability thermal ellipsoids. Hydrogen atoms except for the aqua ligand and two ClO$_4^-$ ions are omitted for clarity. Grey, carbon; blue, nitrogen; red, oxygen; light green, chlorine.

spectroscopies, ESI-TOF-MS spectrometry (Fig. S2), X-ray crystallography, and elemental analysis. The crystal structure of **1**·(ClO$_4$)$_2$ (for preparation, see ESI) in place of **1**·(NO$_3$)$_2$ indicates that the NHC moiety of the PY$_4$Cl$_2$BIm ligand binds at the *trans*-position of the aqua-ligand as shown in Fig. 1. The Fe-O bond length of **1**·(ClO$_4$)$_2$ is 2.012(5) Å[18] and longer than that of the pyridyl counterpart (1.9907(13) Å),[13] indicating that the NHC ligand exerts strong *trans*-influence.

Electrochemical studies of **1**·(NO$_3$)$_2$ were performed in water in the presence of 0.1 M KNO$_3$ as an electrolyte to observe a reversible redox wave of **1**·(NO$_3$)$_2$ at +0.38 V (*vs.* SCE), indicating that the Fe^{II} centre was oxidized to the Fe^{III} state (Fig. S3). The Fe^{III} centre was confirmed to be in a low-spin ($S = 1/2$) state by ESR spectroscopy at 5 K after a reaction of **1**·(NO$_3$)$_2$ with (NH$_4$)$_2$-[Ce^{IV}(NO$_3$)$_6$] (CAN) in water (Fig. S4). In the Pourbaix diagram of **1**, the Fe^{II}/Fe^{III} redox potential of **1** in water showed no dependence on solution pH values (Fig. S5). Since the acidity of the aqua ligand in **1**·(NO$_3$)$_2$ decreased due to the strong *trans*-influence of the NHC ligand, the pK_a value of the aqua ligand could not be determined in the pH range from 1.0 to 12 by UV-vis spectroscopic titration (Fig. S6). Thus, 1e$^-$ oxidation of **1**·(NO$_3$)$_2$ forms the corresponding Fe^{III}-OH$_2$ complex in the pH range. Note that, in CH$_2$Cl$_2$, two oxidation waves were observed in a square-wave voltammogram; the first one at +0.60 V (*vs.* SCE) should be due to the Fe^{II}/Fe^{III} process and the second one at +1.92 V should be due to oxidation of the Fe^{III} complex to an Fe^{IV} species (Fig. S7).

Catalytic oxidation was performed in a glass tube containing D$_2$O (1.0 mL, pD was not adjusted), **1**·(NO$_3$)$_2$ (0.1 mM), oxidant (50 mM) and a substrate (10 mM) at 323 K under Ar atmosphere. Catalytic turnover numbers (TONs) after stirring for 3 h were determined by ^{1}H NMR spectroscopy using sodium trimethyl-silylpropanesulfonate (DSS) as an internal standard. Here, sodium ethylbenzenesulfonate (EtPhS), sodium cumene-sulfonate (CumS) and sodium styrenesulfonate (StyS) were used as water-soluble substrates. At first, we explored the reactivity of various oxidants in the presence of **1**·(NO$_3$)$_2$ and consequently Na$_2$S$_2$O$_8$ was found to be the best oxidant among them (Table S1). In addition, the ESI-TOF-MS measurement of the mixture after the catalytic reaction indicates that the Fe^{III} species

remaining in the solution is an Fe(PY$_4$Cl$_2$BIm) complex having the substrate as a ligand (Fig. S8). Thus, complex **1** still holds the PY$_4$Cl$_2$BIm ligand even after the reaction. The TONs obtained for three substrates with Na$_2$S$_2$O$_8$ are listed in Table 1. Formation of CO$_2$ and O$_2$ was not observed in GC-MS measurements for the reaction mixtures in all cases. In the case of oxidation of EtPhS and CumS, the 2e$^-$ oxidation proceeded to afford the corresponding benzyl alcohol derivatives in TONs of 42 and 26, respectively.[19] In the EtPhS oxidation, CH$_3$CH(OH)PhS was obtained in 86% selectivity; however, catalytic oxidation of EtPh by a nonheme Fe^{IV}-oxo complex has been reported to afford CH$_3$CH(OH)Ph in *ca.* 50% selectivity.[9,20] Oxidation of StyS selectively afforded a diol derivative, which should be derived from hydrolysis of the corresponding epoxide as a 2e$^-$-oxidized product, in full conversion. On the other hand, 2e$^-$-oxidation products were not observed in the absence of catalyst **1**·(NO$_3$)$_2$ (Table S2). These results indicate that **1**·(NO$_3$)$_2$ selectively catalyses 2e$^-$-oxidation of substrates by suppression of over-oxidation.

Table 1   Catalytic substrate oxidation by **1**·(NO$_3$)$_2$ in water

Substrate	Product	TONs^a	Alcohol/ketone	Conversion, %
EtPhS	(OH product)	42	6.0	49
	(O product)	7		
CumS	(OH product)	26	–	26
StyS	(diol product)	93	23	97
	(CHO product)	2		
	HCOOH	2		

^{a} TONs = [Product]/[catalyst]. Conditions: [**1**·(NO$_3$)$_2$] = 0.1 mM, [Na$_2$S$_2$O$_8$] = 50 mM, [Substrate] = 10 mM, T = 323 K, and reaction time = 3 h. TONs were determined by ^{1}H NMR spectroscopy using DSS as an internal standard. Any side-reaction products except those listed in the table were not observed.

To confirm the origin of the oxygen atoms of CH$_3$CH(OH)PhS obtained from the oxidation reaction of EtPhS, ^{18}O-labeled water (H$_2$^{18}O) was used as a solvent. In the ESI-TOF-MS spectrum of the reaction mixture in H$_2$^{18}O, a peak cluster was observed at m/z = 203.17 for CH$_3$CH(^{18}OH)PhS, not at m/z = 201.10 for CH$_3$CH(^{16}OH)PhS (Fig. S9). This indicates that Na$_2$S$_2$O$_8$ works simply as an ET oxidant, not as an oxygen-atom-transfer reagent in the reaction.[21] Thus, it has been confirmed that an oxygen atom in the active species derived from water as a solvent is transferred to a substrate, affording an oxygenated product.

To gain mechanistic insights into the oxidation process of EtPhS, kinetic studies were performed. At first, catalytic oxidation reactions of EtPhS were conducted in D$_2$O (pD was not adjusted) in the presence of Na$_2$S$_2$O$_8$ (50 mM) and various concentration of **1**·(NO$_3$)$_2$ (0.01–0.1 mM) at 323 K and the progress of the reaction was monitored by ^{1}H NMR spectroscopy. The initial rates, v_0, were determined with the

slope of time-course of the CH$_3$CH(OH)PhS formation (Fig. S11). On the basis of the dependence of v_0 on the concentration of **1**·(NO$_3$)$_2$ obeyed first-order kinetics (Fig. 2a). Thus, one molecule of the catalyst participates in the catalytic oxidation of EtPhS. Based on this dependence, the rate constants, k_{cat}, were determined. The k_{cat} values were also determined for various initial concentrations of EtPhS (Fig. 2b). As the results, saturation of k_{cat} values against the concentration of EtPhS was observed, indicating that the existence of a pre-equilibrium process between an oxidatively active species and EtPhS prior to the oxidation. Curve fitting for the plots of k_{cat} against the concentration of EtPhS with eq (2)[22,23] afforded the equilibrium constant, K, as $(1.3 \pm 0.2) \times 10^2$ M^{-1} for the pre-equilibrium process and the first-order rate constant, k, as $(7.9 \pm 0.4) \times 10^{-3}$ s^{-1} for the oxidation reaction (Table S3).

$$k_{cat} = kK[\text{EtPhS}]/(1 + K[\text{EtPhS}]) \tag{2}$$
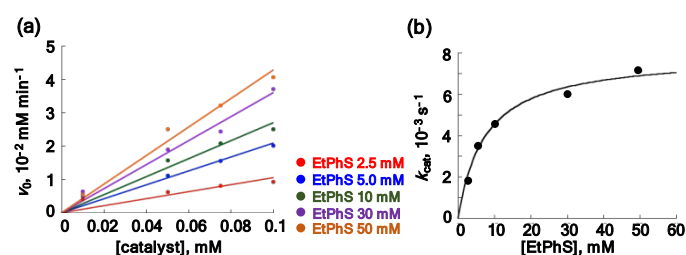

Fig. 2   (a) Dependence of the initial rates on the catalyst concentration, and (b) dependence of the rate constants, k_{cat}, on the initial concentration of EtPhS for the oxidation of EtPhS by **1**·(NO$_3$)$_2$ as the catalyst (0.1 mM) and Na$_2$S$_2$O$_8$ as an oxidant (50 mM) in D$_2$O (pD was not adjusted) at 323 K.

To obtain further information on the oxidation process by **1**·(NO$_3$)$_2$, the kinetic isotope effects (KIE) were also studied using sodium triphenylmethanesulfonate (TPMS) and its mono-deuterated derivative (TPMS-d_1) in a D$_2$O solution at 323 K and the reactions were monitored by ^{1}H NMR spectroscopy. The k_{cat} values for the TPMS and TPMS-d_1 oxidation also displayed saturation behaviour as in the case of EtPhS oxidation described above (Fig. 3). The k and K values were also determined for TPMS ($k^H = (4.4 \pm 0.1) \times 10^{-3}$ s^{-1}, $K^H = (1.9 \pm 0.3) \times 10^2$ M^{-1}) and TPMS-d_1 ($k^D = (1.2 \pm 0.5) \times 10^{-3}$ s^{-1}, $K^D = 46 \pm 5$ M^{-1}) based on eq 2 (Table S3). The oxidation of TPMS-d_1 was clearly slower than that of TPMS. The KIE value was determined to be 3.7 as the ratio of the rate constants (k^H/k^D) for the oxidation reactions of
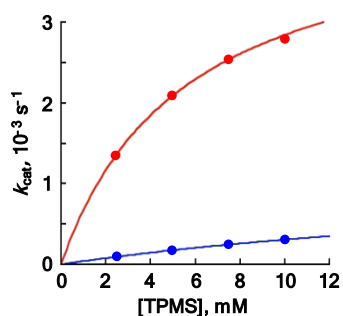
TPMS and TPMS-d_1. The KIE value indicates that the hydrogen-atom abstraction from the methine C-H bond is involved in the rate-determining step (RDS).

To investigate the dependence of the k values on BDEs of the C-H bonds for substrates, we determined the k values for oxidation of six substrates, TPMS (BDE = 79.0 kcal mol^{-1}),[24] sodium diphenylmethane-sulfonate (DPMS; 82.0),[24] CumS (83.5),[24] EtPhS (85.4),[24] benzyl alcohol (BnOH; 87.5),[24] 2-propanol (2-PrOH; 90.0),[24] by **1**·(NO$_3$)$_2$ as the catalyst and Na$_2$S$_2$O$_8$ as the sacrificial oxidant at 323 K (Table S3). The log (k/n) values (n: the number of equivalent C-H hydrogen atoms) were plotted against the BDE values of the C-H bonds to be cleaved in the substrates (Fig. 4). In contrast to other Fe catalysts,[11,12] the rate constants for the oxidation reactions by **1**·(NO$_3$)$_2$ showed an inflection point: For substrates having BDE values greater than *ca.* 86 kcal mol^{-1}, the log (k/n) values decreases with increasing BDE as expected on the basis of the eq (1) with $\alpha = -0.5$. In sharp contrast, for substrates having BDEs less than *ca.* 86 kcal mol^{-1}, log (k/n) values showed no dependence on the BDE values.

Tanko and co-workers have reported on oxidation reactions of organic substrates by ^{t}BuO$^•$ as the oxidant and discussed causes of the insensitivity of the rate constants to the BDEs of the C-H bonds in substrates.[25] It has been known that the BDE of the O-H bond in ^{t}BuOH is 105 kcal mol^{-1}, indicating that ^{t}BuO$^•$ has high reactivity in hydrogen-atom abstraction. Furthermore, their BEP plot for oxidation of various substrates by ^{t}BuO$^•$ exhibited an inflection point at *ca.* 92 kcal mol^{-1}, which is about 13 kcal mol^{-1} lower than that of the O-H bond of ^{t}BuOH. Therefore, our results suggest that the O-H bond formed after C-H oxidation in a putative Fe-oxo reactive species derived from **1**·(NO$_3$)$_2$ should have a large BDE value (*ca.* 100 kcal mol^{-1}), in light of the inflection point at *ca.* 86 kcal mol^{-1}.

The relatively high 2e$^-$-oxidation selectivity by suppression of overoxidation in the present system should be derived from a strongly BDE-controlled reaction mechanism as demonstrated by the BEP plot (Fig. 4): C-H abstraction from the 1-phenyl-ethanol derivative (BDE: 88.3 kcal mol^{-1})[24] as the 2e$^-$-oxidized product of EtPhS (85.4 kcal mol^{-1}) should be strongly retarded, in light of the BEP plot. On the basis of the product analysis and kinetic studies described above, a plausible mechanism for the oxidation reaction of EtPhS catalysed by **1**·(NO$_3$)$_2$ is proposed as shown in Fig. 5. As the first step, **1** is oxidized to the corres-

Fig. 3   Pseudo-first-order kinetic analysis for oxidation reactions of TPMS (red line) and TPMS-d_1 (blue line) by **1**·(NO$_3$)$_2$ as the catalyst (0.1 mM) and Na$_2$S$_2$O$_8$ as oxidant (50 mM) in D$_2$O (pD was not adjusted) at 323 K.
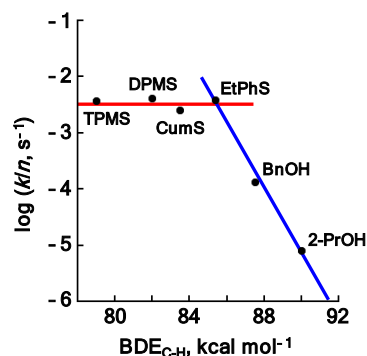

Fig. 4   A plot of the log (k/n) at 323 K for the oxidation of five substrates (TPMS, DPMS, CumS, EtPhS, BnOH, and 2-PrOH) by **1** against BDEs of C-H bonds to be cleaved.

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ponding Fe^{III}-OH_2 complex through ET oxidation by $S_2O_8^{2-}$, as confirmed by ESR spectroscopy after the reaction of **1**·$(NO_3)_2$ with $Na_2S_2O_8$ (step I in Fig. 5; see Fig. S12). The Fe^{III}-OH_2 complex is further oxidized by $SO_4^{•-}$ to form an Fe^{IV}=O active species, which is known to exhibit extremely strong oxidizing power,[26] enough to oxidize the Fe^{III}-OH_2 complex to the Fe^{IV}=O complex; E(Fe^{III}/Fe^{IV}) = +1.92 V *vs.* SCE in CH_2Cl_2 (see above, step II). To clarify the characteristics of the reactive species formed by $2e^-$-oxidation of **1**, DFT calculations were performed on a Fe-O complex including two H_2O molecules interacted with the oxygen ligand at the B3LYP/6-311+G** level of theory (Fig. S13). The results suggest that the triplet Fe^{IV}=O state is the most stable (Table S4). The Fe=O distance in the optimized structure is 1.668 Å and the bond order is 1.83. After the pre-equilibrium process to form an adduct between the active species and EtPhS (step III, see Fig. 2), EtPhS is oxidized through the C-H bond cleavage as the RDS to afford the $2e^-$-oxidized product (step IV, KIE shown in Fig. 3). In the final step V, the oxidized product coordinated to Fe centre is substituted by a water molecule to complete the catalytic cycle (see Fig. S9).

In conclusion, we have demonstrated catalytic oxidation reactions by an NHC-ligated Fe-complex, **1**·$(NO_3)_2$ as a catalyst to afford $2e^-$-oxidized products with high selectivity by suppression of overoxidation in water. Although details of the reactive species in the reactions have yet to be clarified, the reactive species derived from **1** should have a strong oxidative activity due to the strong *trans*-influence from the NHC moiety.
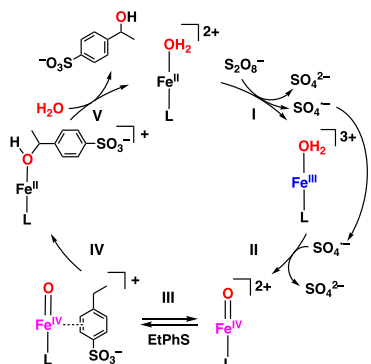

Fig. 5   A proposed mechanism of the oxidation of EtPhS by **1** and $S_2O_8^{2-}$.

Conflicts of interest

There are no conflicts to declare.

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Strong trans-influence

86% selectivity

Selective 2e⁻-oxidation

$Na_2S_2O_8$ in H_2O