Chemical **Science**

EDGE ARTICLE

Check for updates

Cite this: Chem. Sci., 2020, 11, 11877

d All publication charges for this article have been paid for by the Royal Society of Chemistry

Received 1st July 2020 Accepted 23rd September 2020

Introduction

Electrochemical oxidation presents an attractive alternative to traditional chemical reagents for large-scale applications, mostly owing to the generation of less toxic waste than that produced by current chemical processes.1-4 However, electrochemical oxidation of unactivated C-H bonds, the most ubiguitous structural motifs in complex natural products, remains a significant challenge. The difficulty in such oxidation is the high redox potentials required to oxidize such moieties by direct electrolysis (frequently above 3.0 V vs. SCE, which is significantly higher than the thermodynamic potential).^{5,6} A solution to this problem was envisaged by the use of "electron-proton transfer mediator" (EPTM) molecules7 such as N-hydroxyphthalimide (NHPI), quinuclidine, and TEMPO, which have allowed the oxidation of compounds with C-H, including hydrocarbons^{8,9} and alcohols.^{3,10,11} In the electrochemical oxidation of compounds with unactivated C-H bonds using an EPTM, the first step involves the H-atom abstraction from the C-H bond by oxidized N-containing tertiary amines (especially quinuclidine) via proton-coupled electron transfer (PCET).8 Subsequently, the radical formed reacts with dioxygen to form an oxygenated product via a radical process, which significantly reduces the regioselectivity of these oxidation reactions.12-15 In

Oxoiron(v) mediated selective electrochemical oxygenation of unactivated C-H and C=C bonds using water as the oxygen source[†]

Bittu Chandra, 🕑 Hellan K. M., Santanu Pattanayak and Sayam Sen Gupta ២ *

An efficient electrochemical method for the selective oxidation of C-H bonds of unactivated alkanes (BDE \leq 97 kcal mol⁻¹) and C=C bonds of alkenes using a biomimetic iron complex, [(bTAML)Fe^{III}-OH₂]⁻, as the redox mediator in an undivided electrochemical cell with inexpensive carbon and nickel electrodes is reported. The O-atom of water remains the source of O-incorporation in the product formed after oxidation. The products formed upon oxidation of C-H bonds display very high regioselectivity (75:1, 3° : 2° for adamantane) and stereo-retention (RC ~99% for cyclohexane derivatives). The substrate scope includes natural products such as cedryl acetate and ambroxide. For alkenes, epoxides were obtained as the sole product. Mechanistic studies show the involvement of a high-valent oxoiron(v) species, [(bTAML)Fe^V(O)]⁻ formed via PCET (overall 2H⁺/2e⁻) from [(bTAML)Fe^{III}-OH₂]⁻ in CPE at 0.80 V (vs. Aq/AqNO₃). Moreover, electrokinetic studies for the oxidation of C-H bonds indicate a second-order reaction with the C-H abstraction by oxoiron(v) being the rate-determining step.

> Nature, the selectivity in oxidation is achieved by the catalytic generation of high-valent oxometal species, especially highvalent oxoiron intermediates.16-21 These reactive intermediates abstract H-atoms from C-H bonds via PCET and subsequently hydroxylate via a "rebound mechanism, a process that avoids the formation of long-lived radical intermediates that can escape into the bulk". These high-valent oxometal intermediates are typically formed upon activation of the enzyme-bound metal ion with chemical oxidants such as O_2 (together with a biological e^{-}/H^{+} donor) or with H_2O_2 . Alternatively, highvalent oxometal intermediates are also generated via a PCET process from metal ions ligated to aquo ligands (oxidation of [M $- OH_2$ ⁿ⁺ to [M(O)]ⁿ⁺²), which is postulated to occur in the active site of photosystem II.22,23 Such a PCET process has been used to generate oxometal complexes from their corresponding aquabound metal precursors in synthetic complexes of first and second-row transition metals.24-29 For example, Ru^{VI}/Ru^V-oxo species are synthesized electrochemically via PCET from corresponding Ru^{II}–OH₂ complexes and explored for the oxidation of C-H bonds (e.g. benzyl alcohol, ethylbenzene, and 3° C-H bonds of amine derived substrates).^{30–32} Recently, a [Fe^{III}-TAML] (TAML = tetra-amido macrocyclic ligand)³³ complex has been shown to oxidize activated benzylic C-H bonds via electrochemical formation of a dimeric µ-O-Fe^{IV}₂ intermediate in a divided cell.34 The advantage of such electrochemical oxidation is that it eliminates the necessity of terminal oxidants like O₂, H₂O₂, or mCPBA. In all these examples, the O-atom of water remains the source of the O-incorporation in the products. Recently, heterogenized iron porphyrins have been used to catalytically oxidize hydrocarbons, including unactivated C-H

View Article Online

View Journal | View Issue

Department of Chemical Sciences, Indian Institute of Science Education and Research Kolkata Mohanpur, West Bengal, India-741246. E-mail: sayam.sengupta@iiserkol.ac. in

[†] Electronic supplementary information (ESI) available: Materials, general instrumentation and additional figures. See DOI: 10.1039/d0sc03616a

bonds using molecular O2 at cathodic potentials.³⁵ However, the selective homogeneous electrochemical oxidation of unactivated C-H and C=C bonds in organic substrates using water as the O-atom source via the formation of high-valent oxoiron intermediates has been elusive. We have earlier demonstrated that the 5th generation biuret modified TAML, [Et₄N]₂[FebTAML(Cl)], catalyzes the oxidation of 3° C-H bonds with unprecedented selectivity over 2° C-H bonds using NaOCl or mCPBA as the terminal oxidant.^{36,37} Epoxidation of C=C bonds has also been demonstrated with both NaOCl and O2.38,39 The active intermediate involved in this reaction was identified to be a room temperature stable oxoiron(v) intermediate, [(bTAML) $Fe^{V}(O)$]⁻ [in short, oxoiron(v) or $Fe^{V}(O)$] which has been extensively characterized spectroscopically.36,40 However, this reaction utilizes stoichiometric chemical oxidants, and concurrently produces toxic waste such as m-chlorobenzoic acid at the end of the reaction. Recently, we have demonstrated that this oxoiron(v) intermediate can also be generated electrochemically both in water and in a mixed CH₃CN: H₂O solvent.⁴¹ Since this generation of oxoiron(v) uses only water and electricity, coupling this to the catalytic oxidation of C-H bonds would constitute a clean synthetic process. In this work, we demonstrate the selective electrochemical alkane hydroxylation and epoxidation by employing [(bTAML)Fe^{III}(OH₂)]⁻ as the catalyst and water as the oxygen atom source. The substrate included cedryl acetate, a natural product, where selective hydroxylation of an unactivated 3° C-H bond was demonstrated. The electrochemical oxidations were carried out in a simple undivided cell using inexpensive electrodes made from carbon and nickel. To the best of our knowledge, this represents the first example of the use of an iron-complex to catalyze electrochemically selective oxidation of strong C-H bonds (with BDE \leq 97 kcal mol⁻¹) and C=C bonds using water as the O-atom donor (Scheme 1).

Results and discussion

The electrochemical properties of the bTAML complexes 1a and 1b were investigated in 0.1 M Bu₄NPF₆ acetonitrile solution using glassy carbon (GC) as the working electrode, Pt wire as a counter electrode, and Ag/AgNO3 (0.01 M) as the reference electrode. Complex 1a with an axial chloro (Cl) ligand was converted to its corresponding axially agua (H_2O) complex (1b,PPh₄ salt) by a method published before for similar complexes.⁴¹ X-ray diffraction quality single crystals of 1b were grown from a concentrated water solution over a week (Fig. S1[†]). For complex **1a**, a reversible one-electron wave at $E_{1/2} = 0.13$ V with a near Nernstian peak-to-peak separation of $\Delta E_{\rm p} = 66 \text{ mV}$ was observed by cyclic voltammetry (CV). A second quasireversible one-electron wave at $E_{1/2} = 0.72$ V with an almost similar peak-to-peak separation of $\Delta E_{\rm p} = 66$ mV was also seen. The peak-to-peak separation of 66 mV of the first and second oxidation process indicates that the processes at 0.18 V and 0.72 V are both 1e⁻ redox processes. We assigned these redox processes to the Fe^{IV}/Fe^{III} and Fe^V/Fe^{IV} redox couple, respectively (Fig. 1A). Variation of the scan rates shows that the peak currents (i_p) associated with both of the redox events vary linearly with the square root of the scan rates (Fig. S4[†]) and follow Randles-Sevcik diffusion eqn (1).

$$i_{\rm p} = 0.446 n_{\rm p} F A[{\rm Fe}] \sqrt{\frac{n_{\rm p} F v D}{RT}}$$
(1)

where i_p is the current at peak potential for the anodic and cathodic peak ($E_{p,a}$ and $E_{p,c}$), n_p is the number of electrons transferred in the absence of the substrate, D is the diffusion coefficient, F is the Faraday constant, A is the electrode area, R is the universal gas constant, T denotes temperature, [Fe] is the catalyst concentration, and ν is the scan rate. Using eqn (1), the electrochemical diffusion coefficient (D) of **1a** is determined to



Scheme 1 (A) Various approaches of forming high-valent oxometal complexes. (B) Electrochemical approach of forming an oxoiron(v) complex from 1a/1b followed by oxidation of alkanes and alkenes using water as the O-atom source.



Fig. 1 (A) CV of 1a (0.2 mM) in CH₃CN, scan rate = 100 mV s⁻¹. (B) CV of 1b (0.2 mM) in CH₃CN (95%)–H₂O (5%), scan rate = 100 mV s⁻¹. (C) CV of 1b (0.2 mM) with adamantane (30 mM) in CH₃CN (90%)–H₂O (10%) with a scan rate of 300 mV s⁻¹ and (D) CV of 1b (0.2 mM) with styrene (4 mM) in CH₃CN (90%)–H₂O (10%) with a scan rate of 300 mV s⁻¹.

be 6.0×10^{-6} cm² s⁻¹ and 1.46×10^{-5} cm² s⁻¹ for first and second oxidation processes, respectively (Fig. S4†).

In the CV of 1b, which contains an axial H₂O ligand instead, the corresponding Fe^{IV}/Fe^{III} was shifted to 0.41 V vs. Ag/AgNO₃ in CH_3CN (95%)- H_2O (5%). The peak shift to a higher potential for **1b** is reflective of the change in the overall charge of the Fe^{III}b-TAML complex from -2 to -1, upon replacement of the axial anionic Cl ligand with the neutral OH2, in line with similar observations reported earlier.42 Besides, the greater solvation of anionic species due to an increase in the solvent polarity as a result of water addition could also lead to this shift.43 The one-electron redox event at 0.41 V was found to be pH-dependent when CVs and DPVs were recorded at different pH (CH₃CN, 80% - 5 mM phosphate buffer, 20%; Fig. S9[†]). In comparison to that of CH₃CN (100%), the Fe^{V}/Fe^{IV} peak of the catalyst (1a) in CH₃CN (90%)-H₂O (10%) slightly shifted anodically to an $E_{1/2}$ of 0.74 V vs. Ag/AgNO₃ (Fig. 1B). This change observed might be due to bias voltage change because of the presence of a mixed solvent. However, in contrast to the Fe^{IV}/Fe^{III} peak, the Fe^V/Fe^{IV} peak is pH-independent (Fig. S9[†]). Additionally, the same shifts were observed when the DPV of 1a was performed in CH₃CN (90%)-H₂O (10%) (Fig. S7†), indicating that the axial chloride ligand was replaced by water under these conditions, as has been reported earlier.41,42 For 1b, the peak currents (i_p) also vary linearly with the square root of scan rates (0.01 to 1 V s^{-1}) for both the redox processes (Fig. S5[†]).

The controlled potential electrolysis (CPE) of a dilute solution (60 μ M) of **1b** in CH₃CN (90%)–H₂O (10%) at 0.80 V vs. Ag/AgNO₃ for 60 s led to the immediate formation of a light green solution (Fig. S2†). This solution displayed UV-Vis peaks (445 nm and 613 nm), and HR-MS spectra that are characteristic of the well-characterized [(bTAML)Fe^V(O)]⁻ (3) that had been electrochemically synthesized and reported by our group earlier.⁴¹ When the same CPE was carried out at higher concentrations (>0.2 mM of **1b**) in CH₃CN (90%)–H₂O (10%), the formation of a violet species (UV-Vis peaks at 421 nm and 750–1050 nm) characteristic of μ -O-Fe^{IV}₂ (2) was observed (Fig. S3†).

All these studies indicate that for **1b**, the second peak in the CV at $E_{1/2} = 0.74$ V can be assigned to a $[(bTAML)Fe^{V}(O)]^{-/}$ $[(bTAML)Fe^{IV}(O)]^{2-}$ couple (*viz.* spectroelectrochemistry and pH-independent CV). Since the variation of i_p with the square root of the scan rate indicated a diffusion-controlled $1e^{-}$ redox process, the possibility of an EC mechanism, during which the electro-generated oxoiron(v) underwent very fast comproportionation with bulk **1b** to form the dimeric μ -O-Fe^{IV}₂ complex³⁸ on the CV time scale was excluded. The formation of the μ -O-Fe^{IV}₂ complex, observed in CPE experiments at higher concentrations of **1b**, was a result of electro-generated oxoiron(v) diffusing out of the reaction layer and undergoing a comproportionation reaction with **1b** in bulk (Fig. S3†).

The electrochemical generation of oxoiron(v) at 0.80 V on the CV time scale prompted us to explore the possibility of electrochemical oxidation of substrates containing unactivated C-H and C=C bonds. Such substrates, which have C-H bonds with BDE \leq 97 kcal mol⁻¹, have been shown to react with the chemically synthesized oxoiron(v) with high selectivity and fast reaction rates.^{36,37} The CVs of **1b** recorded in 0.1 M Bu₄NPF₆ CH₃CN (90%)–H₂O (10%) medium in the presence of either

styrene or adamantane show an almost unperturbed first oxidation wave at $E_{p,a} = 0.40$ V, and a slight increase in current at an E_{p.a} of 0.80 V (Fig. 1C, D and S10[†]). The enhancement of the current in the second oxidation wave was indicative of a chemical reaction that was coupled to the oxidation of 1b. Further, the peak currents (i_p) did not vary linearly with the square root of scan rates (Fig. S6[†]). The possibility of the chemical reaction was ascertained by performing CPE at 0.80 V (vs. $Ag/AgNO_3$) in the presence of either adamantane or styrene for 6 hours. GC-MS analysis of the reaction medium showed the formation of styrene oxide when styrene was added during CPE. For adamantane, predominantly 1-adamantanol was observed as the product with \sim 3% 2-adamantanone, which indicated a selectivity of 75:1 for 3° C-H bonds over 2° C-H bonds in adamantane. The anodic CV waveform observed in the presence of the strong C-H bond (adamantane) was indicative of low substrate consumption at a high scan rate of 300 mV s^{-1} (Fig. 1C) as compared to a low scan rate of 50 mV s⁻¹ (Fig. 2C). In both the CV traces, the cathode waveform displays a hybrid peak-plateau shape, where the initial peak shape observed due to depletion of oxoiron(v) at the reaction layer is controlled by a plateau resulting from the slow subsequent catalytic oxidation reaction.44,45 The formation of styrene oxide and 1-adamantanol during the CPE experiments prompted us to perform bulk electrochemical oxidation reactions with a variety of substrates containing C-H bonds with BDE ≤ 97 kcal mol⁻¹ and C=C bonds. All the electrochemical oxidation reactions were performed under constant potential electrolysis (CPE) conditions in an IKA ElectraSyn instrument using a simple 3-electrode undivided electrochemical cell, consisting of a working electrode (RVC), a counter electrode (Ni foam) and a reference electrode (Ag/AgNO₃). A reaction mixture of 1a/1b (5 mol%), substrate, and Bu_4NPF_6 (0.1 M) as the supporting electrolyte in (4:1) CH₃CN-H₂O was maintained at a constant potential of 0.80 V (vs. Ag/AgNO₃) at ambient temperature and pressure. No product formation was observed when the reaction was carried out either in the absence of the catalyst (1a/1b) or water or at constant potential less than 0.80 V (Table S1[†]). Further, no changes in the yield and the composition of the products were observed for the reaction performed in the presence or absence of O2. Finally, optimization of the reaction conditions showed that the addition of pH 8 phosphate buffer (5 mM) accelerated the catalytic reaction but the yield of the reaction remained unchanged (Table S1[†]). Phosphate promoted acceleration of reaction rates has been reported by us earlier for the oxidation of C-H bonds using Fe-bTAML and chemical oxidants.37 Since the second redox process at 0.80 V was unaffected by the presence of phosphate buffer, the catalytic reactions were carried out using pH 8 phosphate buffer (5 mM) (Fig. S8[†] and S9[†]).



View Article Online Edge Article

With the optimized reaction conditions, various organic substrates containing 3° C-H bonds with BDE \leq 97 kcal mol⁻¹ and activated 2° C-H bonds were explored using this electrochemical method (Table 1 & Fig. S17-S24[†]). Oxidation at the 3° position occurred preferentially in the presence of the statistically more abundant 2° C-H bond, with excellent conversion, yield, and retention of configuration (RC). For example, oxidation of adamantane, possessing twelve 2° C-H (BDE = 98.5 kcal mol⁻¹)⁴⁶ and four 3° C–H bonds (BDE = 96.5 kcal mol⁻¹),⁴⁶ afforded very high yields of 1-adamantanol $(\sim 84\%)$ with very high regioselectivity (*i.e.*, 75 : 1) of 3° C–H over 2° C-H bonds (Table 1; entry 1). Moreover, a moderate value of faradaic efficiency (\sim 42%) was observed (Table S2^{\dagger}), which suggests a parallel competitive reaction involving the reaction of electro-generated oxoiron(v) with 1b, resulting in the formation of the μ -O-Fe^{IV}₂ dimer. *cis*-1,2-Dimethylcyclohexane (3° C-H BDE = 93.9 kcal mol⁻¹),⁴⁶ trans-1,2-dimethylcyclohexane $(3^{\circ} \text{ C-H BDE} = 97.4 \text{ kcal mol}^{-1})$,⁴⁶ and *cis*-decalin $(3^{\circ} \text{ C-H BDE})$ = 93.5 kcal mol⁻¹)⁴⁶ (Table 1; entries 2, 3 and 4) are the cyclohexane derivatives where the stereochemical orientation of the 3° C-H bonds (axial or equatorial) governs the regioselectivity outcome of the reaction. For cis-dimethylcyclohexane and cisdecalin, mainly 3° hydroxylated products were obtained in 85% and 76% yields, respectively (along with \sim 99% RC). On the other hand, electrochemical oxidation of trans-dimethylcyclohexane, afforded moderate yields (62%). Besides, trans-dimeregioselectivity, thylcyclohexane exhibits compromised resulting in the formation of both the alcohol and ketone in a ratio $(3^\circ : 2^\circ)$ of 97.8 : 2.2. The difference in reactivity between the cis-isomer (cis-dimethylcyclohexane and cis-decalin) and trans-isomer (trans-dimethylcyclohexane) can be attributed to the strain release in the transition state for the *cis*-isomers.⁴⁷ The results obtained in the electrochemical oxidation of these hydrocarbons mirror those obtained in chemical oxidations with complex 1a and mCPBA/NaOCl as the oxidant.^{36,37} This supports the likely involvement of a high-valent oxoiron intermediate during the electrochemical oxidation reactions with 1a/ 1b.

The protocol for oxidation of 3° C-H bonds in simple hydrocarbons was then extended to natural product derivatives. For example, cedryl acetate, a natural product derivative of cedrol, a sesquiterpene alcohol, found in essential oil and used in the chemistry of aroma compounds, having a rigid structure with five 3° C-H bonds, affords a single hydroxylated product in 52% yield (Table 1, entry 5). Ambroxide, a naturally occurring terpenoid used in perfumery, undergoes oxidation at the alpha ethereal C-H bond predominantly among many other electronically and sterically accessible secondary and tertiary C-H bonds. This results in the formation of the ketone product, sclareolide in 75% yield (Table 1, entry 6). The electrochemical oxidation of ambroxide was also attempted at a 1 mmol scale using the same electrochemical setup, and sclareolide was formed in 80% yield (based on conversion). We believe that this methodology can be further scaled up by suitable modifications of the electrodes and the electrochemical cell. Substrates having activated benzylic C-H bonds such as ethylbenzene and diphenylmethane were also explored. Primarily ketone products

(acetophenone and benzophenone) were formed with high conversion and yield for ethylbenzene and diphenylmethane, respectively (Table 1, entries 7 and 8). The formation of a ketone as the main product was attributed to the over oxidation of the hydroxylated product, which was first formed during the oxygenation reaction. Over oxidation of alcohols to ketones has been shown by us earlier for the oxidation of cyclohexane (oxidation of cyclohexanol with 3 is ~350 times faster than cyclohexane oxidation).⁴⁸

Next, electrochemical oxidation of alkenes was performed at constant potential (0.80 V vs. Ag/AgNO₃) using complex 1a/1b as the mediator and water as the oxygen atom source (Table 2 & Fig. S25-S29[†]). Analysis of the products indicated the predominant formation of alkene oxides in moderate to high yields with only a trace amount of the side-product (mainly aldehydes, \sim 5– 7% with respect to epoxides). Styrene afforded styrene oxide as the predominant product in 86% yield (Table 2; entry 1). Subsequently, different para-substituted styrene derivatives such as 4-chlorostyrene and 4-methoxystyrene afforded their corresponding epoxides with yields of up to 92% (Table 2; entry 2 and 3). The higher yields obtained for the electron-rich styrene derivative in comparison to the electron-deficient styrene derivative support the likely involvement of an electrophilic high-valent oxoiron intermediate. Oxidation of cis-stilbene, a substrate that contains a sterically constrained double bond resulted in 64% yield (72% conversion) with a cis/trans product ratio of 8:1 (Table 2; entry 4). cis-Cyclooctene was oxidized to cis-cyclooctene oxide in 75% yield (Table 2; entries 5), which also shows the selectivity for the oxidation of the C=C bond over the weaker allylic C-H bond.

Electrochemical epoxidation using iron and manganese complexes and nanoparticles has been reported earlier. For example, Groves and coworkers49 reported the electrochemical generation of a model compound II analog (oxoiron(IV)) from the corresponding porphyrin-Fe^{III}-OH. Although this intermediate was competent in oxidizing styrene derivatives to their epoxides, the selectivity and efficiency were much lower than those of desired model compound I (which is isoelectronic to oxoiron(v)). Collman and coworkers50 and Tanaka and coworkers⁵¹ reported electrochemical epoxidation using Mn complexes (porphyrins and salen), but the mechanism involved the electrochemical generation of the chemical oxidant (H_2O_2) and OCl⁻), which would subsequently activate the metal complex in a process similar to chemical oxidation. Murray and coworkers⁵² and recently Dey and coworkers³⁵ showed electrochemical reductive activation of O2 bound to metal porphyrin complexes, which would subsequently form high-valent oxometal intermediates to catalyze oxidation reactions. The only example, which demonstrates the use of water as the O-atom source for epoxidation reactions has been recently reported by Manthiram and coworkers.53 They used manganese oxide nanoparticles to catalyze cyclooctene epoxidation using water as an oxygen source with a faradaic efficiency of \sim 30%.

A study of the mechanism of electrochemical oxidation of 3° C–H bonds with BDE ≤ 97 kcal mol⁻¹ was then attempted with the substrate adamantane. We hypothesized that for the electrochemical reaction, the starting Fe(III)-OH₂ (**1b**) is converted to

Table 1	Electrochemical	oxidation of	of alkanes	using	Fe-bTAML
---------	-----------------	--------------	------------	-------	----------

Entry	Substrate (% conversion)	Products (% yield)	Selectivity
1	(90%)	(84%)	$3^{\circ}: 2^{\circ} = 75: 1$
2	(90%)	(85%)	<i>cis</i> : <i>trans</i> = 99 : 1, 3° : 2° = 99 : 1
3	(70%)	(62%)	<i>trans</i> : $cis = 99 : 1, 3^{\circ} : 2^{\circ} = 87.8 : 22.2$
4	H (80%)	(76%)	$cis: trans = 99: 1, 3^{\circ}: 2^{\circ} = 98.3: 1.7$
5	(68%)	о (52%)	Only alcohol
6		(75%)	Only ketone
7	(99%)	(92%)	Only ketone
8	(98%)	(95%)	Only ketone

^{*a*} Reaction conditions: **1***a*/**1***b* (0.75 mM) and substrate (15 mM) in acetonitrile aqueous phosphate buffer (4 : 1 v/v, 5 mM, pH \sim 8) at a constant potential of 0.80 V (*vs.* Ag/AgNO₃) at room temperature for 10 h. Yields and conversions were estimated by GC–MS.

oxoiron(v), (3) through PCET followed by another electron transfer, which then reacts with adamantane to give rise to the product 1-adamantanol, in turn generating **1b** back. The adamantane consumption is small in the reaction layer, and the plateau current is governed by the slow chemical reaction. This reaction can be approximated as proceeding through the E_rC_i mechanism (see ref. 56 for discussion).^{44,45} To understand the influence of the varying substrate concentration and catalyst concentration on the catalytic current we assume that the maximum catalytic current ($i_{cat,max}$) would change according to eqn (2), where [Ada] is the concentration of the substrate adamantane, and n_p and n_{cat} are the number of electrons transferred in the absence of the substrate and in the catalytic reaction respectively.

$$i_{\text{cat, max}} = n_{\text{cat}} FA[\text{Fe}] \sqrt{Dk_{\text{cat}}} = n_{\text{cat}} FA[\text{Fe}] \sqrt{Dk_2[\text{Ada}]}$$
 (2)

The background-corrected i_{cat} increases with increasing concentrations of the catalyst **1b** in CH₃CN (90%)–H₂O (10%)

(Fig. 2A) and shows a linear dependence with the catalyst concentration (Fig. 2B). Hence, the catalytic activity is first-order with the concentration of **1b**. The catalytic current also varies linearly with the square-root of [Ada], suggesting a first-order kinetic dependence of adamantane (eqn (2), Fig. 2D). When combined, the rate of adamantane oxidation can be expressed as rate = $k_2 \times$ [Fe] × [Ada]. Similar second-order dependence on the substrate has been found for the electrochemical oxidation of alcohols, which involves high-valent oxoruthenium intermediates.^{30,31} Finally, the same rate expression was also obtained earlier for hydrocarbon oxidation with chemically synthesized oxoiron(v) under single turnover conditions.³⁶

It's important to note that the CV shape observed for varying catalyst and substrate concentrations is not precisely the S-shaped curves obtained in zone KS (see ref. 57 for discussion), which can be generally achieved at high substrate concentrations or high scan rates.^{44,45} We were unable to reach a perfect KS zone due to the limited solubility of adamantane in CH_3CN (90%)– H_2O (10%), and an increasing probability of

Table 2 Electrochemical oxidation of alkenes using Fe-bTAML^a



^a Reaction conditions: 1a/1b (0.75 mM) and substrate (15 mM) in acetonitrile aqueous phosphate buffer (4:1 v/v, 5 mM, pH ${\sim}8)$ at a constant potential of 0.80 V (vs. Ag/AgNO₃) at room temperature for 4-8 h. Yields and conversions were estimated by GC-MS.

dimerization at low scan rates. As the CVs show a hybrid peak and plateau at varying substrate concentrations (Fig. 2C & S11[†]), the plateau current analysis according to eqn (2) to determine kinetic parameters would result in an approximated maximum rate value ($k_{2,\text{max}}$).⁴⁴ The $k_{2,\text{max}}$ value of 0.23 \pm 0.02 M⁻¹ s⁻¹ determined from the slope (Fig. 2D and S11[†]) using eqn (2) was around 6-fold higher than that determined from the reaction of adamantane with chemically generated oxoiron(v) in CH₃CN.³⁶ The second-order rate constant $(k_{2,\max})$ was determined from the catalytic response obtained at various substrate concentrations at a fixed scan rate (Fig. 2C), using values of D estimated earlier. The presence of 10% water and higher ionic strength during the electrochemical oxidation could have afforded enhanced rates during the oxidation reactions, as has been reported earlier.54 Moreover, an approximate kinetic isotope effect (KIE) value of 4 was obtained from comparing the catalytic current for adamantane and adamantane-d₁₆ (Fig. S12†). This KIE value is comparable with our previously reported KIE value (~ 5) of adamantane during chemical oxidation.⁵⁵

However, the CVs recorded in the presence of a substrate having C-H bonds with low BDE (xanthene; BDE = 75.5 kcal mol^{-1}) were indicative of a no substrate consumption zone (Fig. S13[†]).^{44,45} Catalytic current increased linearly with varying catalyst concentration and with the square-root of the xanthene concentration (Fig. S14 and S15[†]). Hence, a similar



Fig. 2 (A) CV of 1b in the presence of adamantane (5 mM) in CH₃CN (90%)-H₂O (10%) with varying concentration of the catalyst (scan rate = 300 mV s⁻¹). (B) Plot of i_{cat} vs. [Fe-bTAML]. (C) CV of **1b** (0.08 mM) in CH₃CN (90%)-H₂O (10%) with varying concentration of adamantane (scan rate = 50 mV s⁻¹). (D) Plot of i_{cat} vs. [Ada]^{0.5}

second-order rate for xanthene oxidation can be expressed as rate = $k_2 \times [Fe] \times [xanthene]$. From the slope of linear variation of i_{cat} vs. $v^{-0.5}$, the second rate constant $(k_{2,\text{max}})$ for xanthene oxidation was determined to be $182 \pm 40 \text{ M}^{-1} \text{ s}^{-1}$ (Fig. S16[†]). This observation does indicate that the rate of the chemical reaction following the electron transfer has an impact on the CV shape.

We proposed a catalytic cycle for the electrochemical oxidation of alkenes and alkanes based on the product analysis, cyclic voltammetry, controlled potential electrolysis, UV-Vis spectroscopy, and previously reported studies. The electrocatalytic cycle begins at the anode (RVC) immediately after applying a constant potential (0.80 V) where complex 1b undergoes an overall $2H^{+}/2e^{-}$ (PCET) transfer reaction to form oxoiron(v) species, $[(bTAML)Fe^{V}(O)]^{-}$ (3). For the hydroxylation reaction, the high-valent oxoiron(v) species (3) hydroxylates the C-H bond in substrates to form the corresponding alcohol by the typical "rebound mechanism" as has been reported earlier.^{29,36,37} The high regioselectivity observed in adamantane $(3^\circ: 2^\circ \text{ of } 75: 1)$ and stereo-retention found for cis-1,2-dimethylcyclohexane $(\sim 99\%)$ show that the alkyl radical formed during the oxidation does not escape the reaction layer to react with O2 present in the solvent to generate unwanted oxygenated products. Such "nonrebound" pathways, if operational, lead to much reduced regioselectivity and stereo-retention. In fact, when catalytic oxidations of adamantane with 1a and NaOCl/mCPBA were carried out in the presence of O_2 , the regioselectivity of $3^\circ : 2^\circ$ hydroxylation dropped to 30:1 (compared to 75:1, when O2 was rigorously excluded).55 This had indicated that for catalytic oxidation reactions using chemical oxidants, both the "rebound" and "non-rebound" pathways were operational, although oxoiron(v) was shown to be the active intermediate in these reactions. However, during the electrochemical oxidation reactions, the presence or absence of O2 did not affect the selectivity of the reaction, indicating that the radicals formed in

the reaction layer at the applied potential exclusively formed the corresponding alcohol via the rebound process. Such a mechanism would regenerate the starting Fe^{III}-OH₂ complex (1b), which is expected to be seen in the bulk. However, spectroelectrochemical studies indicate the formation of the µ-O- $Fe^{IV}_{2}(2)$ dimer in bulk during the reaction. Such dimeric species are typically formed by the comproportionation reaction of 3 with 1b. The formation of 2 during the electrochemical catalytic reaction is expected since 3 generated in the anode does not quantitatively react with the substrate due to the inherently slow rates of the reaction. This fact is corroborated in the CV studies of 1b (in the presence of the substrate, Fig. 1C), where the peak due to the reduction of oxoiron(v) to oxoiron(v) is discernible. Hence a part of 3 generated in the anode diffuses to the bulk solution and is readily converted to the dimeric species µ-O- Fe_{2}^{IV} (2) via a comproportionation reaction with bulk 1b. However, 2 is not stable in the reaction medium and reduces back to the parent 1b. In fact, upon removal of the applied potential during the reaction, spectroelectrochemical studies indicate that the conversion of 2 into the starting Fe^{III} precursor (1b) occurs within minutes. We have reported that the chemically synthesized dimeric µ-O-Fe^{IV}₂ complex in CH₃CN reduces back to the starting Fe^{III} precursor over time, and the rate of reduction increases with an increase in the water concentration. A plausible mechanism for the reduction involves the disproportionation of the dimeric species to corresponding Fe(v)-oxo and Fe(III) complexes in acetonitrile-aqueous phosphate buffer solution. The aqueous phosphate buffer is known to axially bind with the catalyst and can accelerate the disproportionation reaction. Fe(v)-oxo reacts with the substrate and eventually leads to the formation of Fe(III). As there is no applied potential, Fe(v)oxo is not formed again, and the complete regeneration of Fe(III) is observed. Thus, during the catalytic reaction, the dimeric µ-O-Fe^{IV}₂ species formed spontaneously converts to Fe^{III} in bulk, which is then re-oxidized in the anode to continue the catalytic cycle. This competitive reaction involving the comproportionation of 3 with 1b and the subsequent self-reduction to the parent Fe^{III} complex is reflected in the low faradaic efficiency of the reaction. On the other hand, the mechanism of the epoxidation reaction probably involves the electrophilic attack of oxoiron(v) onto the alkene followed by a fast ring-closing step to form the epoxide.38 However, additional studies are required to understand the mechanism in more detail.

Finally, there are two things that are important to discuss. First, it is worth comparing the electrochemical oxidation of substrates having 3° C–H bonds with the corresponding chemical oxidation methods using *m*CPBA as the terminal oxidant. During catalytic chemical oxidation reactions for adamantane with complex **1a**, the yield of the product 1-adamantanol obtained was less than 5%.⁵⁵ Studies have indicated that the excess oxidant and its by-product present in the reaction mixture promoted the demetallation of **1a**, resulting in complete loss of its activity. This oxidant-induced demetallation process was completely averted during electrochemical oxidations, which allowed them to be used for reactions with moderate efficiency. The second is the comparison of the prototype Fe-TAML complex with our biuret-substituted FebTAML complex. The prototype Fe-TAML complex has been recently shown to catalyze the electrochemical oxidation of alcohols and substrates containing benzylic C-H bonds, viz. substrates having C-H bonds with BDE less than 90 kcal mol⁻¹.³⁴ In this electrocatalytic approach, the breadth of reactivity demonstrated with the Fe-bTAML complex is not achieved due to the oxidative self-decomposition of the prototype Fe-TAML catalyst under the bulk electrolysis conditions, which limits the catalyst lifetime. In addition, the instability of oxoiron(v) of the prototype Fe-TAML (decomposition at temperatures $> -40 \degree \text{C}$ ³³ leads to its conversion to the corresponding μ -O-Fe^{IV}₂ dimer, which is a much weaker oxidant. This limits the reactivity to only weak C-H bonds. In contrast, the increased stability of the oxoiron(v) intermediate in our FebTAML (stable at RT) allows this highly reactive intermediate [BDE (Fe^{IV}O-H) = 99 kcal mol⁻¹] to catalyze the oxidation of C-H bonds with a BDE of up to 97 kcal mol^{-1} with very high selectivity.36

Conclusion

In conclusion, we have demonstrated an electrochemical method for selective oxidation reactions of alkanes having sp³ C-H bonds with BDE ≤ 97 kcal mol⁻¹ using water as the oxygen source. Typically such electrochemical oxidations require very "high overpotential" which can also oxidize other functional groups present in the substrate. However, the use of the Fe^{III} complexes, 1a/1b, allows such oxidations to be achieved at much lower overpotentials and with high selectivity. Spectroelectrochemical analysis shows the involvement of a highvalent oxoiron(v) species, [(bTAML)FeV(O)]-, formed via PCET from the iron(III) complex, [(bTAML)Fe^{III}-OH₂]⁻. Determination of the rate law from cyclic voltammetry experiments and product analysis allows us to conclude that the mechanism of hydroxylation involves abstraction of the H-atom by oxoiron(v) followed by the "rebound" mechanism to form the hydroxylated product. This electrochemical oxidation was performed with a series of substrates including natural products such as cedryl acetate and ambroxide to afford oxygenated products with high conversion and yield. The oxidation of ambroxide was also scaled up to a 1 mmol scale. The electrochemical oxidation was also extended to the epoxidation of alkenes. The main attractiveness of this method includes the use of (i) inexpensive carbon (RVC) and nickel electrodes in a simple undivided cell, (ii) an earth-abundant iron complex as a redox mediator, and (iii) water as the O-atom source, (iv) under an open atmosphere. To the best of our knowledge, our report with 1a/1b demonstrates the first example of homogeneous electrochemical hydroxylation and epoxidation involving the high-valent oxoiron(V) intermediate and using the O-atom of water as the oxygen source.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

S. S. G. acknowledges SERB, New Delhi (Grant no EMR/2017/ 003258), and IISER-Kolkata ("Start-up Grant") for funding. B. Chandra acknowledges UGC-New Delhi for his fellowship.

Notes and references

- 1 M. Yan, Y. Kawamata and P. S. Baran, *Chem. Rev.*, 2017, **11**7, 13230–13319.
- 2 M. D. Kärkäs, Chem. Soc. Rev., 2018, 47, 5786-5865.
- 3 F. Wang and S. S. Stahl, Acc. Chem. Res., 2020, 53, 561-574.
- 4 K. J. Jiao, Y. K. Xing, Q. L. Yang, H. Qiu and T. S. Mei, Acc. Chem. Res., 2020, 53(2), 300-310.
- 5 H. G. Roth, N. A. Romero and D. A. Nicewicz, *Synlett*, 2016, 27, 714–723.
- 6 O. R. Luca, J. L. Gustafson, S. M. Maddox, A. Q. Fenwick and D. C. Smith, *Org. Chem. Front.*, 2015, **2**, 823–848.
- 7 R. Francke and R. D. Little, *Chem. Soc. Rev.*, 2014, **43**, 2492–2521.
- 8 Y. Kawamata, M. Yan, Z. Liu, D. H. Bao, J. Chen, J. T. Starr and P. S. Baran, *J. Am. Chem. Soc.*, 2017, **139**, 7448–7451.
- 9 E. J. Horn, B. R. Rosen, Y. Chen, J. Tang, K. Chen, M. D. Eastgate and P. S. Baran, *Nature*, 2016, 533, 77–81.
- 10 A. Badalyan and S. S. Stahl, Nature, 2016, 535, 406-410.
- 11 C. Costentin, ACS Catal., 2020, 10, 6716-6725.
- 12 H. Sterckx, B. Morel and B. U. W. Maes, *Angew. Chem., Int. Ed.*, 2019, **58**, 7946–7970.
- 13 G. B. Shul'pin, Alkane Funct., 2018, 2, 47-72.
- 14 J. E. Lyons, P. E. Ellis and H. K. Myers, *J. Catal.*, 1995, **155**, 59–73.
- 15 B. Chandra, P. De and S. Sen Gupta, *Chem. Commun.*, 2020, 56, 8484–8487.
- 16 L. Que and W. B. Tolman, Nature, 2008, 455, 333-340.
- 17 M. Costas, M. P. Mehn, M. P. Jensen and L. Que, *Chem. Rev.*, 2004, **104**, 939–986.
- 18 P. R. Ortiz De Montellano, Chem. Rev., 2010, 110, 932-948.
- 19 B. Meunier, S. P. de Visser and S. Shaik, *Chem. Rev.*, 2004, 104, 3947–3980.
- 20 X. Huang and J. T. Groves, Chem. Rev., 2018, 118, 2491-2553.
- 21 M. Sono, M. P. Roach, E. D. Coulter and J. H. Dawson, *Chem. Rev.*, 1996, **96**, 2841–2887.
- 22 J. P. McEvoy and G. W. Brudvig, *Chem. Rev.*, 2006, **106**, 4455–4483.
- 23 R. Gupta, T. Taguchi, B. Lassalle-Kaiser, E. L. Bominaar, J. Yano, M. P. Hendrich and A. S. Borovik, *Proc. Natl. Acad. Sci. U. S. A.*, 2015, **112**, 5319–5324.
- 24 M. K. Coggins, M. T. Zhang, A. K. Vannucci, C. J. Dares and T. J. Meyer, J. Am. Chem. Soc., 2014, 136, 5531–5534.
- 25 M. J. Collins, K. Ray and L. Que, *Inorg. Chem.*, 2006, **45**, 8009–8011.
- 26 W. A. Lee, T. S. Calderwood and T. C. Bruice, *Proc. Natl. Acad. Sci. U. S. A.*, 1985, 82, 4301–4305.
- 27 H. Kotani, T. Suenobu, Y. M. Lee, W. Nam and S. Fukuzumi, *J. Am. Chem. Soc.*, 2011, **133**, 3249–3251.
- 28 W. C. Ellis, N. D. McDaniel, S. Bernhard and T. J. Collins, J. Am. Chem. Soc., 2010, 132, 10990–10991.

- 29 B. Chandra, K. K. Singh and S. Sen Gupta, *Chem. Sci.*, 2017, **8**, 7545–7551.
- 30 A. Paul, J. F. Hull, M. R. Norris, Z. Chen, D. H. Ess, J. J. Concepcion and T. J. Meyer, *Inorg. Chem.*, 2011, 50, 1167–1169.
- 31 A. K. Vannucci, Z. Chen, J. J. Concepcion and T. J. Meyer, *ACS Catal.*, 2012, **2**, 716–719.
- 32 S. G. Robinson, J. B. C. Mack, S. N. Alektiar, J. Du Bois and M. S. Sigman, Org. Lett., 2020, 22(18), 7060–7063.
- 33 (a) T. J. Collins and A. D. Ryabov, *Chem. Rev.*, 2017, 117, 9140–9162; (b) F. T. de Oliveira, A. Chanda, D. Banerjee, X. Shan, S. Mondal, L. Que Jr, E. L. Bominaar, E. Münck and T. J. Collins, *Science*, 2007, 315, 835–838.
- 34 A. Das, J. E. Nutting and S. S. Stahl, *Chem. Sci.*, 2019, **10**, 7542–7548.
- 35 M. Mukherjee and A. Dey, ACS Cent. Sci., 2019, 5, 671-682.
- 36 M. Ghosh, K. K. Singh, C. Panda, A. Weitz, M. P. Hendrich, T. J. Collins, B. B. Dhar and S. Sen Gupta, *J. Am. Chem. Soc.*, 2014, **136**, 9524–9527.
- 37 S. Jana, M. Ghosh, M. Ambule and S. Sen Gupta, *Org. Lett.*, 2017, **19**, 746–749.
- 38 K. K. Singh, M. K. Tiwari, B. B. Dhar, K. Vanka and S. Sen Gupta, *Inorg. Chem.*, 2015, **54**, 6112–6121.
- 39 K. K. Singh and S. Sen Gupta, *Chem. Commun.*, 2017, 53, 5914–5917.
- 40 S. Pattanayak, A. J. Jasniewski, A. Rana, A. Draksharapu, K. K. Singh, A. Weitz, M. Hendrich, L. Que, A. Dey and S. Sen Gupta, *Inorg. Chem.*, 2017, 56, 6352–6361.
- 41 S. Pattanayak, D. R. Chowdhury, B. Garai, K. K. Singh,
 A. Paul, B. B. Dhar and S. Sen Gupta, *Chem. Eur. J.*, 2017,
 23, 3414–3424.
- 42 S. Groni, C. Hureau, R. Guillot, G. Blondine, G. Blain and E. Anxolabéhère-Mallart, *Inorg. Chem.*, 2008, **47**, 11783– 11797.
- 43 A slight change in the CV shape observed in the second redox process in CH_3CN-H_2O (9 : 1) could be the result of axial ligation of water molecules to the oxoiron(v) formed. Since chemically synthesized oxoiron(v) undergoes ¹⁶O to ¹⁸O scrambling upon addition of H_2O ,¹⁸ such change in the coordination environment after formation of oxoiron(v) is possible.
- 44 (a) E. S. Rountree, B. D. McCarthy, T. T. Eisenhart and J. L. Dempsey, *Inorg. Chem.*, 2014, 53, 9983–10002; (b)
 K. J. Lee, N. Elgrishi, B. Kandemir and J. L. Dempsey, *Nat. Rev. Chem.*, 2017, 1, 0039.
- 45 (a) J.-M. Sav'eant, Elements of Molecular and Biomolecular Electrochemistry, John Wiley & Sons, Inc., Hoboken NJ, 2006; (b) A. J. Bard and L. R. Faulkner, Electrochemical Methods, John Wiley & Sons, Inc., Hoboken NJ, 2nd edn, 2001.
- 46 Y.-R. Luo, Handbook of Bond Dissociation Energies in Organic Compounds, CRC Press, 2004, vol. 126.
- 47 (a) K. Chen, A. Eschenmoser and P. S. Baran, *Angew. Chem., Int. Ed.*, 2009, 48, 9705–9708; (b) M. S. Chen and M. C. White, *Science*, 2010, 327, 566–571.
- 48 M. Ghosh, Y. L. K. Nikhil, B. B. Dhar and S. Sen Gupta, *Inorg. Chem.*, 2015, **54**, 11792–11798.

- 49 J. T. Groves, Z. Gross and M. K. Stern, *Inorg. Chem.*, 1994, 33, 5065–5072.
- 50 H. Nishihara, K. Pressprich, R. W. Murray and J. P. Collman, *Inorg. Chem.*, 1990, **29**, 1000–1006.
- 51 H. Tanaka, M. Kuroboshi, H. Takeda, H. Kanda and S. Torii, *J. Electroanal. Chem.*, 2001, **507**, 75–81.
- 52 S. E. Creager, S. A. Raybuck and R. W. Murray, *J. Am. Chem. Soc.*, 1986, **108**, 4225–4227.
- 53 K. Jin, J. H. Maalouf, N. Lazouski, N. Corbin, D. Yang and K. Manthiram, *J. Am. Chem. Soc.*, 2019, **141**, 6413–6418.
- 54 K. K. Singh, M. K. Tiwari, M. Ghosh, C. Panda, A. Weitz, M. P. Hendrich, B. B. Dhar, K. Vanka and S. Sen Gupta, *Inorg. Chem.*, 2015, 54, 1535–1542.
- 55 M. Ghosh, S. Pattanayak, B. B. Dhar, K. K. Singh, C. Panda and S. Sen Gupta, *Inorg. Chem.*, 2017, **56**, 10852–10860.
- 56 The " E_rC_i' mechanism" associates a reversible electrode electron transfer with a first-order (or pseudo-first-order) follow-up irreversible homogeneous chemical reaction. It is one of the simplest reaction schemes where a heterogeneous Nernstian electron transfer is coupled to an irreversible chemical reaction that takes place in the adjacent solution.^{44,45}
- 57 A "KS zone" is a zone in a kinetic zone diagram representing the various regimes of competition between diffusion and the follow-up reaction.^{44,45} In the KS zone of the EC zone diagram for homogeneous catalysis an S-shaped (where the forward and reverse scans trace each other exactly) CV response is expected.