Modelling the *cis*-oxo-labile binding site motif of non-heme iron oxygenases. Water exchange and remarkable oxidation reactivity of a novel non-heme iron(IV)-oxo compound bearing a tripodal tetradentate ligand

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Abstract

The spectroscopic and chemical characterization of a new synthetic non-heme iron(IV)-oxo species $[Fe^{IV}(O)(^{Me,H}Pytacn)(S)]^{2+}$ (**2**, ^{Me,H}Pytacn = 1-(2′-pyridylmethyl)-4,7-dimethyl-1,4,7-triazacyclononane, S = CH_3CN or H_2O) is described. **2** has been prepared by reaction of $[Fe^{II}(CF_3SO_3)_2(^{Me,H}Pytacn)]$ (**1**) with peracetic acid. Complex **2** bears a tetradentate N_4 ligand that leaves two *cis*- sites available for binding an oxo group and a second external ligand but, unlike related iron(IV)-oxo of tetradentate ligands, it is remarkably stable at room temperature ($t_{1/2} > 2h$ at 288 K). Its ability to exchange the oxygen atom of the oxo ligand with water has been analyzed in detail by means of kinetic studies, and a mechanism has been proposed on the basis of DFT calculations. Hydrogen-atom abstraction from C-H bonds and oxygen atom transfer to sulfides by **2** have also been studied. Despite its thermal stability, **2** proves to be a very powerful oxidant that is capable of breaking the strong C-H bond of cyclohexane (BDE = 99.3 kcal·mol^{-1}).

Keywords

Bioinorganic Chemistry; High Valent; Reaction Mechanisms; Iron; Nonheme Oxigenases

Introduction

Iron(IV)-oxo compounds are proposed as the active species in the catalytic cycles of several O_2-activating non-heme iron enzymes such as isopenicillin N-synthase, pterin-dependent hydroxylases and 2-oxoglutarate-dependent oxygenases.[1–3] Quite recently, high spin (S = 2) iron(IV)-oxo intermediates have been trapped and spectroscopically characterized in five

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different non-heme iron enzymes: namely taurine:α-ketoglutarate dioxygenase (TauD),[4,5] halogenase CytC3,[6] propyl-4-hydroxylase[7], tyrosine hydroxylase,[8] and aliphatic halogenase, SyrB2.[9] Parallel synthetic efforts have succeeded in the preparation of octahedral S = 2 $[Fe^{IV}(O)(H_2O)_5]^{2+}$,[10] and octahedral S = 1 $Fe^{IV}=O$ species using tetradentate N_4 ligands with available *trans*-binding sites,[11,12] tetradentate N_4 ligands with available *cis*-binding sites,[13–17], pentadentate N_5 ligands.[18–22] and a pentadentate N_4S ligand.[23] Very recently, two examples of trigonal bipyramidal S = 2 iron(IV)-oxo complexes, which bears structural and electronic resemblance to the enzymatic non-heme intermediates, have been described.[24–26]

These metastable and highly oxidizing complexes have been characterized by various spectroscopic techniques including, in selected cases, X-ray crystallography.[11,22,25–27] The ability of synthetic non-heme $Fe^{IV}(O)$ species to act as oxidants has been proved[28,29] and several studies demonstrate that they are capable of performing several oxidation reactions including oxygen-atom transfer to sulfides and triphenylphosphine,[16,30,31] epoxidation[13,19,21] and possible dihydroxylation[21,32] of alkenes, alcohol oxidation, [33] hydrogen atom abstraction,[18,34,35] aliphatic and aromatic hydroxylation,[36,37] oxo-transfer to other Fe^{II} complexes,[38] and glutathione and peptide oxidation.[39,40] On the other hand, by analogy to the better studied heme systems,[41] it is widely postulated that non-heme iron-oxo species can exchange their oxygen atom with water *via* a oxo-hydroxo tautomerism before attack to a substrate (Scheme 1). Thus indirect evidence for the participation of these compounds in the catalytic cycle of enzymes[42] and/or synthetic models[43] comes from the incorporation of oxygen from water into oxidized products. Recently, this has been experimentally proved in two synthetic non-heme ion(IV)-oxo species; well-defined $[Fe^{IV}(O)(tmc)(CH_3CN)]^{2+}$ and $[Fe^{IV}(O)(N4Py)]^{2+}$ (Scheme 2) have been shown to undergo exchange with water (Scheme 1).[44]

Because the vast majority of non-heme iron oxygenases contain *cis*-labile binding sites, synthetic iron(IV)-oxo complexes with tripodal tetradentate ligands are especially relevant from a biological point of view. However, they have proved to be less stable than their pentadentate counterparts,[28] rendering their study more difficult. For this reason most of the studies have been performed using pentadentate N_5 ligands (N4Py) or equatorial tetradentate N_4 ligands with two labile sites in a *trans* configuration (tmc). In this work we address this issue and we describe the preparation and spectroscopic and chemical characterization of a new $Fe^{IV}(O)$ species bearing the tripodal tetradentate ^{Me,H}Pytacn ligand (^{Me,H}Pytacn = 1-(2′-pyridylmethyl)-4,7-dimethyl-1,4,7-triazacyclononane). This species shows an unprecedented high stability at room temperature allowing us to evaluate several biologically relevant aspects of its reactivity. These studies include the evaluation of its ability to perform oxygen-atom transfer to sulfides and hydrogen-atom abstraction reactions. Moreover, the exchange of its oxo group with the oxygen atom from $H_2^{18}O$ has been studied by means of kinetic and computational methods.

Results and Discussion

Preparation and characterization of the $Fe^{IV}(O)$ species

Reaction of $[Fe^{II}(CF_3SO_3)_2(^{Me,H}Pytacn)]$ (**1**) (Scheme 3)[45] with 2 equivalents of peracetic acid (CH_3CO_3H) in CH_3CN at 15°C produces within 20 minutes a novel species (**2**) with a UV-vis spectrum showing a band at λ_{max} = 750 nm (ε = 200 $M^{-1}cm^{-1}$) and a weaker shoulder around 900 nm ($\varepsilon \sim$ 100 $M^{-1}cm^{-1}$) (Figure 1). **2** is relatively stable in solution, with a half-life time ($t_{1/2}$) of 2.4 hours at 15°C. The nature of the transient species **2** has been determined by a combination of Mössbauer, ^{1}H-NMR spectroscopy, resonance Raman, ESI-MS spectrometry and DFT calculations. We have studied the Mössbauer spectra of **2** in acetonitrile between 4.2 K and 100 K in parallel applied magnetic fields up to 8.0 T. Two

representative spectra are shown in Figure 2. In zero applied magnetic field (Figure 2A), **2** exhibits at 4.2 K a quadrupole doublet, representing 85(5)% of the total iron, with quadrupole splitting $\Delta E_Q = 0.73$ mms^{-1} and isomer shift $\delta = 0.05$ mm.s^{-1}. These parameters, together with the fact that **2** has integer electron spin (as witnessed by the observation that **2** displays a doublet rather than magnetically split spectra in zero field), show that **2** is an Fe^{IV} complex. Spectra recorded in applied magnetic fields, e. g. Figure 2B, have features very similar to those reported for $S = 1$ Fe^{IV}(O) complexes; $S = 2$ Fe^{IV}(O) complexes display quite different spectra.[6,8,10,24] We have analyzed the spectra of **2** with the commonly used $S = 1$ spin Hamiltonian[23]; the parameters obtained are very similar to those obtained for other $S = 1$ Fe^{IV}(O) complexes and are listed in the caption of Figure 2. The spectrum of Figure 2A indicates the presence of an $S = 0$ diferric contaminant with $\Delta E_Q = 1.60$ mm.s^{-1} and $\delta = 0.45$ mm·s^{-1}, representing ca. $\approx 15\%$ of Fe. For the present chemistry such complexes are decomposition products that frequently represent the thermodynamic sink.[13]

High-resolution ESI-MS definitively proves the nature of **2** (Figure 1, inset) as it reveals a clean spectrum with a major peak with a mass value at m/z = 469.08 and an isotopic pattern fully consistent with the ion {[Fe^{IV}(O)(CF$_3$SO$_3$)(^{Me,H}Pytacn)]}$^+$. Evidence for the presence of the oxo ligand was provided by vibrational spectroscopy. Resonance Raman analysis on frozen acetonitrile solutions of **2** with laser excitation at $\lambda_{exc} = 407.9$ nm shows a resonance enhanced feature at 831 cm^{-1} that downshifts to 788 cm^{-1} when **2** is allowed to react with H$_2^{18}$O. The energy of the band falls in the range of reported synthetic[11,15,22,24,46] and enzymatic[47] nonheme Fe=O stretching vibrations, and the –43 cm^{-1} shift is in reasonable agreement with the shift of –37 cm^{-1} predicted by Hooke's law for a diatomic Fe-O stretch and –40 cm^{-1} predicted by DFT calculations.[48,49] In conclusion, the spectroscopic data allow us to formulate **2** as a new iron(IV)-oxo species [Fe^{IV}(O)(^{Me,H}Pytacn)(S)]$^{2+}$, where S = CH$_3$CN or H$_2$O. The nature of the sixth ligand could not be experimentally determined, but insight into its nature can be obtained from DFT calculations described in the next section. Acetonitrile binding was previously documented in the crystallographic characterization of [Fe^{IV}(O)(tmc)(CH$_3$CN)]$^{2+}$.[11] However, in the present case, the presence of a water ligand is proposed on the basis of DFT calculations (*vide infra*), which indicated that these species are thermodynamically more stable than the corresponding analogue containing an acetonitrile ligand.[13,14]

Because of the inequivalence of the two binding positions not occupied by ^{Me,H}Pytacn, two isomeric forms are possible for **2** (**2a** and **2b**, Scheme 3). These two isomers differ not only in the group disposed *trans* to the oxo ligand but also in the relative orientation of the pyridine ring with respect to the Fe-O axis. However, the ^{1}H-NMR spectrum of **2** (Figure 3) showed a unique set of sharp signals that could be assigned to the pyridine β (42 ppm), γ (10 ppm) and β′ (–13 ppm) protons. The shift pattern for these signals is reminiscent of that found in the ^{1}H-NMR spectrum of [Fe^{IV}(O)(N4Py)]$^{2+}$ in which all four pyridine rings are disposed parallel to the Fe-O axis, and different from the pattern associated with pyridine rings oriented perpendicular to the Fe=O axis.[27] Thus, in the present case, the NMR spectrum indicates that only isomer **2a** is present in solution, with the pyridine ring disposed parallel to the Fe-O axis.

Despite the inherently low stability of high valent oxo-iron(IV) species, the ^{1}H-NMR spectrum of **2** lacks signals corresponding to **1**[50] and **3** (the dinuclear Fe(III) decomposition product, *vide infra*, Sup Info), suggesting that **2** is prepared with good purity, in accord with the Mössbauer analysis (*vide supra*).

DFT analysis of the structures of the two isomers of 2

DFT calculations were employed to help clarifying the molecular and electronic structure of **2**.[48,49] The nature of the sixth ligand that completes the octahedral coordination sphere of the iron center was first studied by comparing the relative energies of the corresponding $[Fe^{IV}(O)(^{Me,H}Pytacn)(S)]^{2+}$ (S = CH_3CN and H_2O) complexes. Computational analysis for S = H_2O indicates that hydrogen bonding between the water ligand and a second, external water molecule is very strong because of the dicationic nature of **2**. This interaction stabilizes $[Fe^{IV}(O)(^{Me,H}Pytacn)(H_2O)]^{2+}$ relative to $[Fe^{IV}(O)(^{Me,H}Pytacn)(CH_3CN)]^{2+}$ by $\Delta G = 16.1$ kcal·mol^{-1}. Acetonitrile binding has been previously demonstrated in the X-ray structure of $[Fe^{IV}(O)(tmc)(CH_3CN)]^{2+}$,[11] for which the hydrophobic pocket created by the N-Me groups of tmc isolates the acetonitrile ligand. This is likely to prevent intermolecular interactions with additional water molecules. In contrast, the tripodal tetradentate $^{Me,H}Pytacn$ ligand lacks such a sterically restricted hydrophobic pocket, which may constitute the key structural feature that favors water binding over acetonitrile in **2**.

Optimized molecular structures of $[Fe^{IV}(O)(^{Me,H}Pytacn)(H_2O)]^{2+}$ (**2a** and **2b**) obtained by DFT calculations are shown in Figure 4, with selected structural parameters given in the figure caption.[51] The relative DFT energies of the two isomeric forms of **2** indicate that **2a** is energetically favored by 2.1 kcal·mol^{-1}, thus supporting our structural assignment based on NMR spectroscopy. Energies corresponding to singlet (S = 0) and quintet states (S = 2) of **2a** and **2b** were also computed (Table S7). The substantially higher energies of these states, when compared with the triplet (S = 1) state further support the Mössbauer-based spin state assignment of **2**. Structural parameters for **2a** and **2b** are very similar. The computed Fe-O distances in **2a** and **2b** are 1.62 Å and 1.63 Å, respectively, in good agreement with structurally characterized examples of synthetic[11,17,22,24–27,52] and enzymatic[53,54] non-heme iron(IV)-oxo species. The oxo ligand exerts a *trans* effect that is indicated by a ~ 0.1 Å lengthening of the corresponding *trans* Fe-N distance (compare $d_{Fe1-N4} = 2.19$ Å in **2b** with 2.06 Å in **2a**, and $d_{Fe1-N3} = 1.99$ Å in **2b** with 2.11 Å in **2a**). The Fe-N$_{py}$ distance is 2.03 Å in **2a** and 2.02 Å in **2b**, values that appear somewhat intermediate between the short d_{Fe-Npy}~ 1.95 Å of the pyridine rings *cis* to the oxo ligand in $[Fe^{IV}(O)(N4Py)](ClO_4)_2$[27] and the longer $d_{Fe-Npy} = 2.118(3)$ Å of $[Fe^{IV}(O)(Pytmc)](CF_3SO_3)_2$[22] for which the pyridine is bound *trans* to the oxo ligand.

Relative thermal stability and decay of 2

$[Fe^{IV}(O)(^{Me,H}Pytacn)(S)]^{2+}$ (**2**) joins the growing family of iron(IV)-oxo species formed with tetradentate ligands tmc,[11] tpa[16], bqen[17], bpd[35] and bpmcn[14] (Scheme 2). In the case of tmc, the ligand topology gives rise to an iron center with two available *trans* sites, one of which is occupied by an oxo ligand in the Fe^{IV}-oxo compound. Instead, ligands like $^{Me,H}Pytacn$, tpa, bpd, bqen and bpmcn wrap around the metal ion, so that two sites in a *cis* relative disposition are ready for interaction with exogenous ligands. Thus, in the corresponding iron(IV)-oxo compounds, the oxo group binds to one of these positions and the other site is presumably occupied by a neutral labile molecule, such as an acetonitrile or water (Scheme 2). Despite the similar coordination spheres around the metal center, the resulting iron(IV)-oxo compounds exhibit remarkably different stabilities. $[Fe^{IV}(O)(bpmcn)(S)]^{2+}$, $[Fe^{IV}(O)(tpa)(S)]^{2+}$ and $[Fe^{IV}(O)(bpd)(S)]^{2+}$ (S = CH_3CN or H_2O) can only be prepared in good yields at -40 °C because they easily decompose at higher temperatures. [13,14,35] $[Fe^{IV}(O)(bqen)(S)]^{2+}$ has a lifetime of $t_{1/2}$~ 30 min at 0 °C.[17] In contrast, **2** can be generated and remains relatively stable at 15 °C.

Thermal decay of **2** leads to a new species **3** that has been identified as $[Fe^{III}_2(\mu\text{-}O)(\mu\text{-}CH_3CO_2)(^{Me,H}Pytacn)_2]^{3+}$ by means of UV-vis spectroscopy, ESI-MS and comparison with an independently prepared sample characterized by X-ray crystallography (Figure 5).[55]

The relatively poor stability observed for iron(IV)-oxo species with tripodal tetradentate ligands, in comparison with those that contain pentadentate or tetradentate ligands with *trans*- topologies, can be understood by considering that only the latter two offer effective steric protection of the oxo-ligand against decomposition into a diiron complex. In addition, these "stable" iron(IV) compounds lack a *cis*-labile site adjacent to the reactive oxo ligand, precluding pre-binding of a putative substrate. Based on these arguments, we suggest that the surprising thermal stability of **2** derives from a combination of the oxidatively robust nature of the ^{Me,H}Pytacn ligand and some steric hindrance imposed by the N-Me groups of the ligand that project into the space surrounding the Fe=O unit. Consistent with this notion, in the absence of acetate ion, which facilitates dimerization, the oxidation of **1** with H_2O_2 in an aqueous:acetonitrile solvent mixture does not result in the common rapid formation of oxo-bridged ferric dimers,[56] but instead in the formation of the mononuclear species formulated as $\{[Fe^{III}(OH)(^{Me,H}Pytacn)](CF_3SO_3)\}^+$ on the basis of its ESI-MS spectra. The remarkable stability of **2** has allowed us to study some aspects of its biologically relevant reactivity which are detailed below.

Oxygen-atom exchange with water

Because of the biological relevance of water-exchange reactions at metal centers,[57] the ability of **2** to exchange its oxo ligand with $H_2^{18}O$ was studied by taking advantage of ESI-MS spectrometry. **2** was generated by reaction of **1** with 2 equivalents of CH_3CO_3H in acetonitrile. Subsequently, 600 equivalents of $H_2^{18}O$ were added to the reaction solution of **2**, and several ESI-MS spectra were taken over time. The initial single peak at m/z = 469.1 associated with $\{[Fe^{IV}(^{16}O)(CF_3SO_3)(^{Me,H}Pytacn)]\}^+$ decreased gradually, while a new peak at m/z = 471.1 corresponding to $\{[Fe^{IV}(^{18}O)(CF_3SO_3)(^{Me,H}Pytacn)]\}^+$ appeared. Exchange of the oxygen atom with labeled water was complete in approximately 3 minutes at room temperature. Addition of pyridine-N-oxide, a strong σ-donating ligand, inhibited the water exchange exhibited by **2** suggesting that the available coordination site in a *cis* configuration with respect to the oxo group plays an important role in facilitating the water-exchange process.

Kinetic analyses of water exchange in **2** were performed by monitoring ^{18}O incorporation into **2** directly *via* ESI-MS, and indirectly by quenching of a solution of **2** and $H_2^{18}O$ with thioanisole at different reaction times and analyzing the percentage of labeled sulfoxide generated by GC-MS. Within error, the two methods afforded identical reaction rates, supporting the validity of the second approach. In any case, exchange rate constants were obtained by non-linear regression fitting of the data to a single exponential function.

The rate of water incorporation into **2** (k_{obs}) was found to be linearly dependent on $[H_2^{18}O]$, and independent of [**2**]. The data are consistent with the rate law in equation 1, where the use of a relative large $[H_2^{18}O]$ allows for its analysis as a pseudo-first order reaction;

$$\frac{d[Fe^{IV}={}^{18}O]}{dt} = k_{exc}[Fe^{IV}={}^{16}O][H_2^{18}O] = k_{obs}[Fe^{IV}={}^{16}O]$$

A bimolecular rate constant of $k_{exc} = 0.028 \pm 0.003$ M^{-1} s^{-1} at 295K was extracted from this analysis. Activation parameters for water exchange were also determined by measuring exchange rates between 275K and 305K, affording $\Delta H^\ddagger = 10.2 \pm 0.8$ kcal·mol^{-1} and $\Delta S^\ddagger = -32 \pm 3$ cal·K^{-1}·mol^{-1} (Figure 6). The negative activation entropy is consistent with an associative event in the transition state, most likely a bimolecular collision between the $Fe^{IV}(O)$ species and a water molecule. Second-order rate constants for water exchange at $[Fe^{IV}(O)(tpa)(S)]^{2+}$ were also measured (Table 1). The accumulated data indicate that water

exchange rate for **2** is substantially faster than those for $[Fe^{IV}(O)(tmc)(CH_3CN)]^{2+}$ and $[Fe^{IV}(O)(N4Py)]^{2+}$, which lack *cis*-available sites to the oxo ligand, and also somewhat faster than $[Fe^{IV}(O)(tpa)(S)]^{2+}$. Nevertheless, considering the pentadentate nature of N4Py, the higher reaction rate observed for **2** represents a rather modest enhancement. Furthermore, the smaller reaction rate measured for $[Fe^{IV}(O)(tpa)(S)]^{2+}$ suggests that reaction rates are not solely determined by the presence of a *cis*-labile site. Analysis of the enthalpy and entropy activation parameters indicates that the origin of the faster exchange rate in **2** has an entropic origin, which compensates for its higher activation enthalpy.

Computational analysis of the water exchange mechanism

An obvious goal arising from these studies is to establish the fundamental chemical steps by which **2** can exchange its oxygen atom with water. The answers to this question may be of fundamental interest due to the structural similarity between **2** and biologically relevant non-heme oxo-iron(IV) species, in which the oxo group binds in a *cis* configuration with respect to one or more available coordination sites. In the heme paradigm, the incorporation of oxygen from water is explained by the so-called oxo-hydroxo tautomerism proposed by Meunier and co-workers,[41] which entails the binding of labeled water *trans* to the oxo group and then tautomerization of this species to a symmetric *trans*-dihydroxoiron(IV) intermediate that scrambles the label (Scheme 1). This pathway clearly cannot be operative for **2**, since there is no binding site *trans* to the oxo group available for coordination with water, and an alternative mechanism must apply in the present case. The mechanism depicted in Scheme 4 starts with the iron(IV)-oxo complex **2a**, with a coordinated water molecule in a *cis* position with respect to the oxo group. In the DFT study[49,58] of the water exchange mechanism, the solvent effects and the dispersion correction have been taken into account in the single point calculations. Solvent effects for acetonitrile were computed using the Polarizable Continuum Model (PCM)[59] as implemented in Gaussian09.[50] The dispersion energy has been computed using the DFT-D3 method and program of S. Grimme *et al.*[60] According to our theoretical calculations,[58] a key aspect of the reaction is the presence of an exogenous water molecule at close distance of the oxo and the water ligands. In the first step the exogenous water assists the transfer of one H^+ from the bound H_2O to the oxo ligand, via a transition state (**TS1**), with a barrier of $\Delta G = 9.3$ kcal·mol^{-1}. After the first proton transfer event, an $Fe^{IV}(OH)_2$ species (**I1**) is formed which is energetically favored compared to the starting complex **2a** by 2.2 kcal·mol^{-1}. Our spectroscopic data do not provide evidence for a significant accumulation of **I1** in solution, but a dihydroxomanganese(IV) species has recently been described for a related macrocyclic N$_4$-ligand,[61] and analogous LFe^{IV}(OH)$_2$ species have been proposed on the basis of DFT methods for a bispidine-based complex (Scheme 2).[32,62] It is possible that the energy of **I1** is underestimated in our calculations, and that it is slightly higher in energy than the corresponding oxo-aqua isomeric species (see also below for a discussion based on computed Mössbauer parameters). In **I1**, the exogenous water molecule has a H-bond interaction with the O5H hydroxide ligand (oxo group in **2a**). In the next step, the exogenous water changes its interaction from the O5H hydroxo group to the second hydroxo ligand (O6H, aqua in **2a**) and the intermediate **I2** is formed. This step is endothermic with a free energy cost of 4.2 kcal·mol^{-1}. When the free energy corrections are included **TS2** and **I2** are essentially isoenergetic. Close inspection reveals that **TS2** and **I2** differ in the orientation of the O5-H bond, which is not involved in hydrogen bonding with the external water molecule. **I2** evolves *via* a second H^+ transfer, passing through **TS3** and forming **2b,** where the oxo ligand is *cis* to the **NCH$_2$py** ligand. Considering that **TS2** and **I2** are nearly isoenergetic, the larger barrier along the overall process is **TS2** + **TS3** with $\Delta G = 13.0$ kcal·mol^{-1}, and corresponds to the second proton transfer. Proton transfer not assisted by the second water molecule has an activation enthalpy barrier about three times higher. Isomer **2b** could not be experimentally observed because it is 2.11 kcal·mol^{-1} higher in energy than

2a, and because the activation barriers associated to each of the elemental steps that connect the backwards conversion of **2a** to **2b** are small. In this scenario, water ligand exchange at **2b** and microscopic microreversibility leads to the formation of **2a**, where the oxygen atom of the oxo group (*trans* to a NCH$_2$Py unit) comes from water. Interestingly, the mechanism for water exchange at **2** bears resemblance to the DFT-computed conversion of [Fe^{IV}(OH)$_2$L]$^{2+}$ (where L stands for a tetradentate bispidine ligand related to bpd, Scheme 2) to [Fe^{IV}(O)(OH$_2$)L]$^{2+}$.[32,62] Calculations indicate that this formal tautomerization takes place with a small energy barrier of 8.3 kcal·mol^{-1}. Unlike in **2**, the reaction involves a spin-crossover from the $S = 1$ Fe^{IV}(OH)$_2$ to the thermodynamically more stable (4.4 kcal.mol^{-1}) $S = 2$ Fe^{IV}(O)(OH$_2$) tautomer, without assistance of a second water molecule. In contrast, all iron(IV) species in our calculations remain on the S = 1 surface.

Taking this computed mechanism into consideration, the first order kinetic dependence on water concentration is attributed to the initial equilibrium step involving the binding of a labeled water molecule, which replaces the initially bound unlabeled water. Nam, Que and coworkers have proposed that a seven-coordinate *cis*-oxo-aqua intermediate is formed in the first step of the reaction between [Fe(O)(N4Py)]$^{2+}$ and [Fe(O)(tmc)(CH$_3$CN)]$^{2+}$ with water (Scheme 1).[34] Such mechanism may be also possible for **2** and [Fe(O)(tpa)(S)]$^{2+}$, but owing to the presence of the labile water/acetonitrile ligand, a six-coordinate intermediate **2b** formed via a ligand exchange reaction is favored. The breaking of the Fe–OH$_2$ bond is likely to be the origin of the higher activation enthalpy which however, is entropically compensated because of the dissociative nature of the event.

An obvious consequence of the present mechanism is that, owing to the asymmetry of the two *cis*-labile sites, oxygen incorporation from water into the oxo ligand requires two oxo-hydroxo tautomerism reactions. That constitutes a fundamental mechanistic difference from oxygen atom incorporation from water in metalloporphyrins.[41]

DFT computation of the Mössbauer parameters of the species implicated in the water exchange mechanism

Considering that the DFT-computed water-exchange mechanism predicts that tautomeric *bis*-hydroxo-iron(IV) (**I1** and **I2**) and oxo-iron(IV) (**2a** and **2b**) are close in energy, the corresponding Mössbauer parameters were calculated with DFT methods and they are collected in Table 2. (See experimental section for details on the DFT analysis). The calculated isomer shifts for **2a** and **2b** are a bit too high relative to the experimental values. The Pittsburgh group, using Gaussian with the B3LYP functional (calibration of Vrajmasu et al.[14a]), has observed that the calculated δ values for TMC come out quite well while those of some most Fe^{IV}=O complexes with pyridine and amine ligands are generally too high, by as much as 0.09 mm/s. For the question at hand it is significant that the ΔE_Q values of **I1** and **I**2 are much larger than those of **2a** and **2b**. In agreement, large ΔE_Q values have been reported for [Fe^{IV}(OH)(OO^{t}Bu)(bpmcn)]$^{2+}$,[14b] $\Delta E_Q = 1.75$ mm·s^{-1} and the Fe^{IV}-OH site of [(OH)(L$_{OMe}$)Fe^{IV}(μ-O)Fe^{IV}(O)(L$_{OMe}$)]$^{3+}$, L = *tris*((4-methoxy-3,5-dimethylpyridin-2-yl)d$_2$-methyl)amine),$\Delta E_Q = 1.96$ mm·s-1.[14c] We conclude that, despite **I1** being the lower energy species in the DFT-calculated mechanism, the Mössbauer parameters favor the oxo-iron(IV) formulation, in agreement with the resonance Raman evidence for an Fe=O unit.

Oxidation of sulfides: oxygen atom transfer

The oxygen atom transfer ability of **2** was evaluated for the oxidation of sulfides. Complex **2** reacted rapidly at 273 K in CH$_3$CN with 10 equiv of thioanisole affording the corresponding sulfoxide (methylphenylsulfoxide) in 98% yield. The reaction was monitored with UV-vis spectroscopy by following the decrease of the band at 750 nm characteristic of the iron(IV)-

oxo complex **2**. Under conditions of excess substrate (5 – 50 equiv with respect to **2**) the reactions showed pseudo first-order behavior so that the observed reaction rates (k_{obs}) were linearly dependent on substrate concentration. From this analysis, a second-order rate constant (k_H) of 1.0 ± 0.1 M^{-1}s^{-1} was obtained for the oxidation of thioanisole (see Supp. Info). Additional information about the mechanism of this transformation was gained through the initial observation that the reaction rates were highly dependent on the *para* substituent of the aromatic ring. Thus, we measured the second-order rate constants (k_X) for a series of *para*-substituted methyl phenyl sulfides, *p*-X-thioanisoles (X = CN, Cl, CH$_3$, OCH$_3$). Plotting of $\log(k_X/k_H)$ for the different substrates against the corresponding Hammett parameters (σ_p) afforded a good linear correlation, and a Hammett value of $\rho = -1.5$ was obtained (Figure 7). The negative ρ value reflects the electrophilicity of the oxo group in **2**. On the other hand, a plot of the $\log(k_X)$ against the one-electron oxidation potentials of each *p*-X-thioanisoles (Eo_{ox}) afforded a linear correlation with a slope of -3.0 (Supporting Information). As slope values around 10 would have been expected for a process initiated by 1-electron transfer,[63, 64] the observed slope of 3 indicates that the oxidation of sulfides by **2** does not occur by such a process but rather *via* a direct oxygen atom transfer mechanism. Further evidence for a direct oxygen atom transfer between the terminal oxo ligand and the sulfide came from labeling of the oxo group with ^{18}O. This was achieved by oxygen atom exchange of **2** with H$_2$^{18}O and subsequent reaction of the resulting [Fe^{IV}(^{18}O)(^{Me,H}Pytacn)(H$_2$O)]$^{2+}$ with thioanisole. Analysis of the resulting sulfoxide by GC-MS indicated that 80% of the product was ^{18}O-labeled.

Despite the similarities between the oxygen atom transfer process in **2** and in other synthetically prepared iron(IV)-oxo species, reactions rates are highly dependent on the specific structure of the ligand. In particular, the oxidation of thioanisole by [Fe^{IV}(O)(N4Py)]$^{2+}$ and [Fe^{IV}(O)(Bn-tpen)]$^{2+}$ (Scheme 2)afforded second-order rate constants of 0.065 M^{-1}s^{-1} (273 K) and 0.075 M^{-1}s^{-1} (253 K), respectively, and this rate was even smaller for [Fe^{IV}(O)(tmc)(CH$_3$CN)]$^{2+}$ for which a value of 0.029 M^{-1}s^{-1} at 35°C was obtained.[31] Comparison with the reactivity of [Fe^{IV}(O)(tpa)(S)]$^{2+}$ is hampered due to its inherent instability which prevents proper measurement of the second-order rate constant at relatively high temperatures (a value of 0.44 M^{-1}s^{-1} corresponding to the reaction with thioanisole at 228 K has been reported).[31] Nevertheless, from these studies it can be concluded that **2** and [Fe^{IV}(O)(tpa)(S)]$^{2+}$, both of which bear tripodal tetradentate based ligands, are much more reactive in the oxygen atom transfer reaction than the corresponding complexes bearing tmc (tetradentate planar) or pentadentate (N4Py and Bn-tpen) ligands.

Oxidation of activated C-H bonds: hydrogen atom abstraction

The reaction of **2** with a series of substrates for H-atom abstraction was studied. **2** reacted with 5 equiv 9,10-dihydroanthracene (DHA) at 258 K affording anthracene as the only detected product by GC-MS. When the reaction was run under N$_2$, 0.45 mmol anthracene/mmol of **2** was measured. The UV-vis spectra at the end of the reactions did not show formation of **1**. In addition, ESI-MS spectrum of the resulting solution was dominated by ions corresponding to mononuclear Fe^{III} species. Therefore, considering that Mössbauer spectra indicate a 85(5)% Fe^{IV} content in our preparations, on the basis of stoichiometric considerations we conclude that in this reaction **2** acts as a 1e$^-$ oxidant (equation 2), and that the yield of the reaction with DHA is nearly quantitative.

$$(2)$$

The kinetics of reactions of **2** with a series of substrates were studied by UV-vis following the decay of the band at 750 nm characteristic of the iron(IV)-oxo species **2**. The decay of **2** followed good first-order kinetics under conditions of excess substrate and k_{obs} values were linearly dependent on substrate concentration in all cases, thus allowing us to determine the corresponding second order rate constants, k_2 (see supporting information). These values are collected in Table 3. Reaction of **2** with alkylaromatic substrates (toluene and ethylbenzene) caused rapid formation of brown-dark blue intense chromophores, indicative of alkylphenolate-bound iron(III) species. Considering the high bond dissociation energy (BDE) of the C-H bond in aromatic substrates, such reactions could not occur via H-atom abstraction but instead, they originate from electrophilic attack of the aromatic ring by **2**.[37] For this reason, these substrates were not further addressed in the present study.

2 proved to be a very powerful oxidant in C-H oxidation reactions (Tables 3 and 4). Compared to related dicationic iron(IV)-oxo complexes, under analogous conditions, reaction rates obtained with **2** are somewhat higher than those obtained with $[Fe^{IV}(O)(N4Py)]^{2+}$ and nearly two orders of magnitude faster than with $[Fe^{IV}(O)(tmc)(CH_3CN)]^{2+}$ [24,34] but somewhat lower than that found for $[Fe^{IV}(O)(bpd)(CH_3CN)]^{+2}$ (Scheme 2).[35] Furthermore, the second order rate constant for the oxidation of DHA with **2** appears to be several orders of magnitude higher than those reported for low-spin $Fe^{III}Fe^{IV}$ and $Fe^{IV}Fe^{IV}$ oxo-dimers, and approaches the remarkably high rate ($k_2 = 28$ M^{-1}s^{-1} at 193 K) recently measured for the high-spin $[(HO)(L_{OMe})Fe^{III}-O-Fe^{IV}(O)(L_{OMe})]^{2+}$, L_{OMe} = tris((4-methoxy-3,5-dimethyl-pyridin-2-yl)methyl)amine (Table 4).[65] The high reactivity of the latter has been rationalized on the basis of the high spin (S = 2) state of the iron ions, and the compound has limited stability even at −80 °C. On the other hand, as already discussed, the metal ion found in **2** has a S = 1 spin state, and it is moderately stable at room temperature. As expected from its higher oxidation state, reaction rates with **2** are 2–3 orders of magnitude faster than those obtained with the lipoxygenase model $[Fe^{III}(OMe)(Py5)]^{2+}$ (Py5 = 2,6-bis(bis(2-pyridyl)methoxymethane)pyridine).[66] Indeed, **2** proved to be competent for performing the oxidation of strong C-H bonds such as THF (BDE = 93 kcal·mol^{-1}) with a reaction rate $k_2 = (2.3 \pm 0.3) \times 10^{-3}$ M^{-1} s^{-1} at 258 K.[67] Most remarkably, **2** is also competent to perform the oxidation of cyclohexane (BDE = 99.3 kcal·mol^{-1}) at room temperature with a second order rate constant, after correction for the number of C-H bonds, of $k_2' = 3.6 \times 10^{-5}$ M^{-1} s^{-1}. GC analysis of the reactions reveals the formation of 0.12 mol cyclohexanone/mol of **2** and 0.28 mols cyclohexanol/mol of **2**. Nevertheless, this reaction rate is relatively very slow when compared with the fast oxidation observed in catalytic **1**/H$_2$O$_2$ cyclohexane oxidations, where a $[Fe^{V}(O)(OH)(^{Me,H}Pytacn)]^{2+}$ species has been inferred from isotopic labeling experiments and DFT analysis.[45,50]

Comparative reactivity profiles of the reaction of **2** against different substrates show that the rate constants decreased with the increase of the C-H BDE and more interestingly the log(k_2') values correlated linearly with the C-H BDE values of the substrates, giving a slope of approximately −0.2 (Table 3, Figure 8). The slope is comparable to those obtained for hydrogen abstraction reactions mediated by $[Fe^{IV}(O)(N4Py)]^{2+}$[16] and $[Fe^{III}(OMe)(Py5)]^{2+}$,[66] and intermediate between the −0.4 and −0.1 observed for $[Ru^{IV}(O)(bpy)_2(py)]^{2+}$ and $[Mn^{III}(OMe)(Py5)]^{2+}$,[69,70] respectively. Such a correlation strongly suggests that reactions take place via a H-atom abstraction mechanism, [69,71–73] as early established for related Fe^{IV} compounds.[18,34,74]

Parallel reactions with deuterated 9,10-dihydroanthracene (d_4-DHA) yielded a kinetic isotope effect (KIE) of 27, a value which is consistent with a C-H bond cleavage being the rate-determining step.[68] This large KIE value is well above the semi-classical limit of 7, suggesting a hydrogen atom transfer mechanism dominated by quantum mechanical tunneling.[74,77,78] For comparison, KIE values around 17 were obtained for the oxidation

of DHA by $[Fe^{IV}(O)(tmc)(X)]^{n+}$ (X = CH$_3$CN, CF$_3$COO$^-$ or N$_3^-$)[34] and even larger values were reported in the oxidation of benzyl alcohol by $[Fe^{IV}(O)(N4Py)]^{+2}$ (KIE = 48) and $[Fe^{IV}(O)(tpa)(S)]^{+2}$ (KIE = 58).[33] Moreover, such large isotope effects have also been observed in hydrogen atom abstraction reactions performed by the iron(IV)-oxo intermediate of TauD (KIE ~ 37)[79], the iron(III)-hydroxo active species of lypoxygenase (KIE~ 50)[80] and compound Q (Fe^{IV}Fe^{IV}) of the diiron enzyme soluble methane monooxygenase (sMMO) (KIE > 50).[81]

A particular case that deserves some comment is the reaction with the NADH analogue 10-methyl-9,10-dihydroacridine (AcrH$_2$), which is considered a hydride donor.[82] The reaction of AcrH$_2$ with **2** results in rapid formation (k_2 = 86 ± 19 M^{-1} s^{-1}) of the acridinium cation AcrH$^+$, as evidenced by UV-vis spectroscopy. UV-vis quantification based on the characteristic spectral features of the cation accounts for 1 mmol AcrH$^+$/mmol **2**, indicating that **2** acts as a 2e$^-$ oxidant, and that the reaction is best described as a formal hydride transfer to the iron(IV)-oxo moiety. Interestingly, the corresponding oxidation rate constant ($k_2{'}$) appears to be only slightly faster than expected from the log($k{'}_2$) *vs* BDE graph correlation (Figure 8) suggesting that a common rate determining step is operative for all the series of substrates plotted. In addition, the KIE evaluated from the reaction of **2** with AcrD$_2$ at 258 K is 4.7 (Figure S11), which is intermediate between the values observed for $[Fe^{IV}(O)(N4Py)]^{+2}$ (KIE = 13.5),[82] and those obtained recently with $[Fe^{IV}(O)(bpd)(CH_3CN)]^{2+}$, (KIE = 2.3) (Scheme 2).[35] On this basis, we propose that the reaction takes place via a rate determining hydrogen atom abstraction (HAT) that can be regarded as a proton coupled electron transfer (PCET), followed by a fast electron-transfer.[68,76,83,84]

Considering that C-H bond oxidation by **2** occurs *via* a hydrogen atom abstraction mechanism,[73,85,86] the balance between the strength of the cleaved C-H bond and of the formed Fe^{III}-OH bond must determine the observed reaction rates. Along this interpretation, we notice that reactions mediated by **2** are comparable to those associated with the most reactive Fe^{IV}(O) species containing nitrogen based pentadentate ligands such as $[Fe^{IV}(O)(N4Py)]^{2+}$, which constitute one of the very rare examples of well defined metal-oxo species that are capable of breaking the strong C-H bond of cyclohexane at room temperature.[18,87–89] The similar reaction rates observed for **2** and $[Fe^{IV}(O)(N4Py)]^{2+}$ should be interpreted as both of them forming a Fe^{III}O-H bond of approximately the same energy. Previous literature reports on the estimation of the Fe^{III}O-H bond in $[Fe^{IV}(O)(N4Py)]^{2+}$ have provided somewhat different values. Two independent theoretical studies yielded Fe^{III}O–H BDE's of 84[90,91] and 97 kcal·mol^{-1},[92] and some of us have experimentally estimated a more modest BDE = 78 kcal·mol^{-1} from its experimentally observed redox potential in water.[75] We can take the latter value as a lower limit estimate of the corresponding Fe^{III}O–H bond formed by H-abstraction in **2**. Thus, as previously noted for $[Fe^{IV}(O)(N4Py)]^{2+}$,[90] despite its rather surprising thermal stability, **2** is a very powerful oxidant. Besides the high oxidation state, one of the reasons at the origin of such a high reactivity is likely to be the positive charge of the $[LFe^{IV}(O)]^{2+}$ unit when L = neutral ligand (L = N4Py or ^{Me,H}Pytacn), which results in a very electrophilic oxidant.

The remarkably high oxidative reactivity exhibited by **2** both in O-atom transfer and in H-abstraction reactions is unprecedented and deserves some comment. The factors that determine the reactivity of oxo-iron(IV) species have been the focus of intense debate. Accessibility of multiple spin states modulated by the nature of the ligand *trans* to the oxo group have been invoked to explain inverted tendencies in oxygen atom transfer *vs* H-abstraction reaction by series of Fe^{IV} compounds.[34] Basic anionic ligands X in a series of $[Fe^{IV}(O)(X)(tmc)]^+$ complexes decrease their electrophilicity and O-atom transfer reactivity, but enhance their H-abstraction reactivity by populating a reactive quintet state. Population of an excited, highly reactive S = 2 state has also been proposed to account for the

exceptional ability of $[Fe^{IV}(O)(N4Py)]^{2+}$ to cleave strong C-H bonds via a H-atom abstraction mechanism.[90] The nature of the ligands *cis* to the oxo group is considered to have a more modest effect on the oxidative reactivity.[93] Computational studies have thus suggested that high-spin $S = 2$ $Fe^{IV}(O)$ species could be much more reactive than their corresponding low spin $S = 1$ analogues. However, the recent preparation and characterization of the first synthetic $S = 2$ $Fe^{IV}(O)$ species raises some questions about this prediction, because it exhibits reactivity only comparable to that of $[Fe^{IV}(O)(N4Py)]^{2+}$. [24,25] Steric effects were invoked by the authors to account for these observations. **2** contains a $S = 1$ spin state, and DFT computations show that it is well separated in energy from the $S = 2$ spin state (See table S7 in sup. Info). Therefore a two-state reactivity (TSR) scenario appears highly unlikely. In addition, high reactions rates are measured for both O-atom transfer and hydrogen-atom abstraction reactions. Its high oxidative reactivity may then add to the rationale of substrate accessibility to the reactive oxo ligand as a major factor dictating reactivity in these species. The high reactivity of **2** in C-H oxidation reactions finds a nice precedent in that recently described for $[Fe^{IV}(O)(bpd)(CH_3CN)]^{+2}$ (Scheme 2).[35] Both compounds have a N_4-based ligand that enforces a *cis*-oxo-labile site. However, the bispidine is a weaker field ligand[32,62] than $^{Me,H}Pytacn$,[50] and the TSR scenario is more likely. In this scenario, we propose that **2** is more reactive than most low spin iron(IV)-oxo species bearing pentadentate and/or planar tetradentate ligands because the oxo ligand in **2** is more exposed, and susceptible to interaction with small substrate molecules. However, compared with previously reported tetradentate ligands, $^{Me,H}Pytacn$ affords a very significant degree of steric protection against bimetallic dimerization reactions.

Conclusion

In this work we have reported the preparation of a new S = 1 iron(IV)-oxo species (**2**) with a tripodal tetradentate ligand that has remarkable thermal stability, especially when compared with structurally related complexes like $[Fe^{IV}(O)(bpmcn)(S)]^{2+}$, $[Fe^{IV}(O)(tpa)(S)]^{2+}$, and $[Fe^{IV}(O)(bpd)(CH_3CN)]^{+2}$. **2** has a labile site *cis* to the oxo group, which constitutes a very common structural feature of non-heme iron oxygenases. **2** rapidly exchanges its oxygen atom with $H_2{}^{18}O$ following a mechanism in which an exogenous water molecule assists the hydrogen transfer from the coordinated water molecule to the oxo group. Despite its remarkable thermal stability, **2** is a very good oxidant both with respect to oxygen atom transfer to sulfides and hydrogen-atom abstraction of alkane C-H bonds, demonstrating a highly electrophilic character arising from the Fe^{IV} oxidation state in a neutral N-based ligand environment. The higher oxidative reactivity of **2** when compared with complexes containing pentadentate ligands is attributed to the tetradentate nature of the $^{Me,H}Pytacn$ ligand, which does not provide steric encumbrance of the oxo ligand in **2**, thus providing the basis of its remarkable oxidative reactivity.

Experimental Section

Materials and methods

Reagents and solvents used were of commercially available reagent quality unless otherwise stated. Preparation and handling of air-sensitive materials were done under an inert atmosphere either on a Schlenk line or in a glove box. Acetonitrile was purchased from Scharlau. $H_2{}^{18}O$ (95% ^{18}O-enriched) was received from ICON Isotopes.

Synthesis of complexes

The starting complex$[Fe(CF_3SO_3)_2(^{Me,H}Pytacn)]$ (**1**) was prepared following the previously reported experimental procedure.[45]

Preparation of iron(IV)-oxo complex (**2**) was performed as follows: to a 1 mM solution of **1** in acetonitrile (2 mL total volume) 2 equiv of peracetic acid solution (100 µL of a 40 mM solution in CH$_3$CN obtained by dilution of commercially available 32% w/w peracetic acid solution in acetic acid) were added at 288 K. The formation of **2** was followed by UV-vis spectroscopy showing the appearance of a band at 750 nm (λ_{max} = 200 M^{-1}cm^{-1}) with a shoulder at 900 nm which developed in about 10 minutes. The resulting solution of **2** (1 mM) was directly used for subsequent reactivity studies.

Direct preparation of [Fe^{III}$_2$(µ-O)(µ-CH$_3$COO)(^{Me,H}Pytacn)$_2$](ClO$_4$)$_3$·CH$_3$CN (3·CH$_3$CN)

54.6 mg of compound 1 (90 µmols) were dissolved in MeCN (2 mL) under an inert atmosphere. 200 µL of an acetonitrile solution 0.30 M in CH$_3$COOH and Et$_3$N (60 µmols CH$_3$COOH and Et$_3$N) were added at once, which caused an immediate color change from dark pink to bright yellow. Immediately a balloon filled with O$_2$ was connected into the reaction vessel and the solution became red-brown in a few seconds. After stirring for 3 h, the solvent from the resulting solution was removed under reduced pressure which afforded a brown oil. The resulting product was redissolved in acetonitrile and 19 mg NaClO$_4$·H$_2$O (135 µmols) were added. The solution was stirred for about 2 hours, filtered through Celite and diethyl ether was slowly diffused. 37 mg of brown crystals of **3·CH$_3$CN** suitable for X-ray diffraction were obtained (38 µmols, 84%). ^{1}H-NMR (200 MHz, CDCl$_3$, 300K) δ = 26.43, 18.27, 17.41, 14.47, 6.57 ppm. FT-IR (ATR) ν = 1611, 1523 (CH$_3$COO), 1447 (C=C$_{ar}$), 1073, 621 cm^{-1} (ClO$_4$).UV/Vis (CH$_3$CN): λ_{max} (ε/Fe) = 427 (660), 462 (690), 513 (500), 692 nm (58 mol^{-1}·cm^{-1}). ESI-MS: m/z: 227.6 [M-3ClO$_4$]$^{+3}$, 880.9 [M-ClO$_4$]$^+$.

Oxygen atom exchange with water

Kinetic studies on oxygen atom exchange of **2** with H$_2$^{18}O were performed by quenching aliquots of the reaction mixture with thioanisole at different times, and analyzing the percentage of ^{18}O-labeled methylphenyl sulfoxide generated by GC-MS. pH dependence of the reaction rates was not studied. In a typical experiment, to 2.5 mL of a stirred solution of **2** in acetonitrile (1 mM), the appropriate amount of H$_2$^{18}O (from 6 to 60 µL) was added at once. At different reaction times, an aliquot of the reaction mixture (350 µL) was directly poured into a solution containing 7 µL of thioanisole in 300 µL of CH$_3$CN. The resulting yellow solution was stirred for 30 minutes at room temperature, filtered through a short path of basic alumina and washed with 2 mL of ethyl acetate. The sample was analysed by GC-MS. The percentage of ^{18}O-labeled sulfoxide was determined by the isotopic pattern showed by the peaks at m/z = 125 and 140. In a typical experiment, a total of 7 aliquots were taken out from the reaction mixture. The temperature of the reaction mixture (between 0 and 30 °C) was controlled by means of a cryostat or a thermostatized water bath. The percentage of ^{18}O-labeled sulfoxide is directly related to the percentage of iron(IV)-oxo species (**2**) that has exchanged its oxygen atom with H$_2$^{18}O. The percentage of ^{18}O-labeled sulfoxide over time could be fitted to a pseudo-first-order kinetics that allowed us to measure the rate constants (k_{obs}) corresponding to oxygen-atom exchange with ^{18}O-labeled water in **2**.

Oxidation of activated C-H bonds and sulfides

The appropriate amounts of substrates (diluted in acetonitrile) were added to a solution of **2** (1 mM) and the subsequent decay of the spectral changes corresponding to the iron(IV)-oxo were directly monitored by UV-vis spectroscopy. The kinetic studies were performed at specific temperatures: 273 K for the oxidation of sulfides and 258 K for the oxidation of activated C-H bonds. Unless specifically stated, reactions were run under air. Rate constants, k_{obs}, were determined by pseudo-first-order fitting of the decrease of the absorption band at 750 nm. Product analyses for the oxidation of thioanisole and DHA were performed by gas chromatography. Prior to injection, an internal standard (biphenyl) was added to the solution

which was further filtered through basic alumina and washed with ethyl acetate. Calibration curves with authentic products were generated to obtain quantitative determination of the oxidized products.

Details on the DFT calculations of the Mossbauer parameters

Details on the DFT calculations are collected in refs The Fe^{IV} DFT Mössbauer parameters were calculated at OPBE/TZP[94] level within the conductor-like screening (COSMO) solvation model[95] (with dielectric constant ε =37.5, Acetonitrile) using the Amsterdam Density Functional (ADF) suite of program.[96] The isomer shift (δ), which is proportional to the electron density [ρ(0)] difference at the Fe nuclei between the studied system and a reference system (normaly α-Fe at 300K), were calculated according to the procedure described in Refs.[97] with utility programs provided by Han and Noodleman. The isomer shift is given by:

$$\delta = \alpha(\rho(0) - A) + C$$

where ρ(0) is obtained using ADF and Han and Noodleman code; A is a constant chosen close to the electron density at the Fe nucleus in the reference state; and α and C were determined by linear regression between the calculated ρ(0) and experimental δ, Han and Noodleman for their $Fe^{2.5+,3+,3.5+,4+}$ training set obtained $\alpha = -0.312 \pm 0.022$ and $C = 0.373 \pm 0.014$ mm s^{-1} for OPBE. The quadrupole splitting, which is proportional to the electric field gradient (EFG) at the Fe nucleus, is directly given by ADF.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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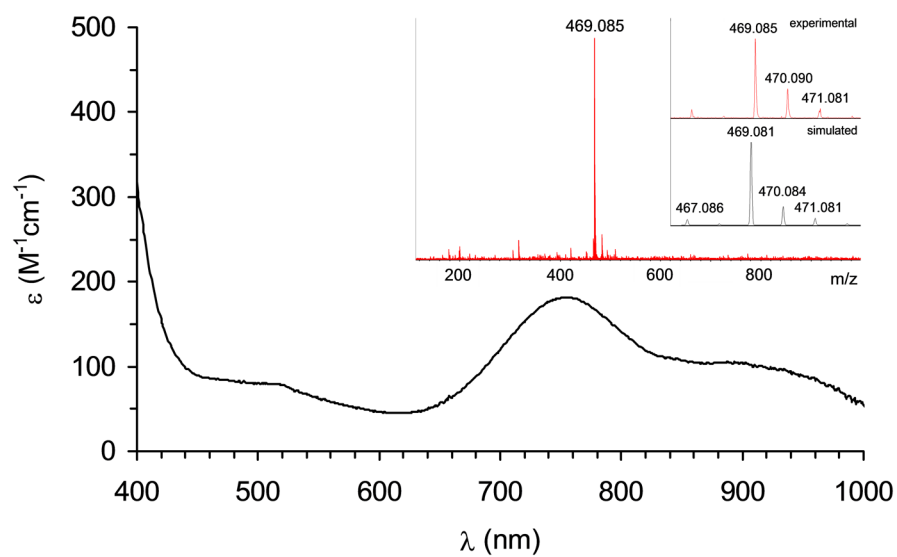**Figure 1.**
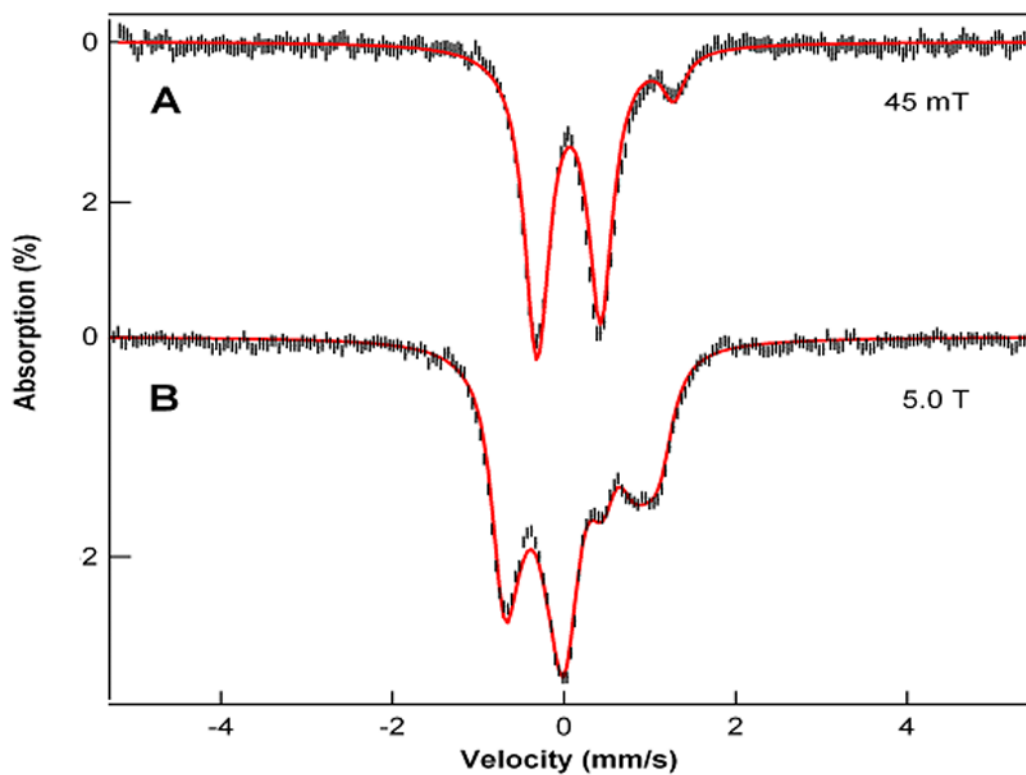UV-vis spectrum of **2** along with its ESI-MS spectrum.

Figure 2.
Mössbauer spectrum of **2** in acetonitrile, recorded at 4.2 K in parallel field as indicated. Solid lines are spectral simulations based on an S = 1 spin Hamiltonian[23] using the zero-field splitting parameters D = +27 cm^{-1}, E/D = 0, ^{57}Fe magnetic hyperfine tensor $A_{x,y,z}/g_n\beta$ = (−23.7,−20.5, −4.5) T, quadrupole parameters ΔE_Q = 0.73 mm·s^{-1}, η = 1, and isomer shift δ = 0.05 mm·s^{-1}.
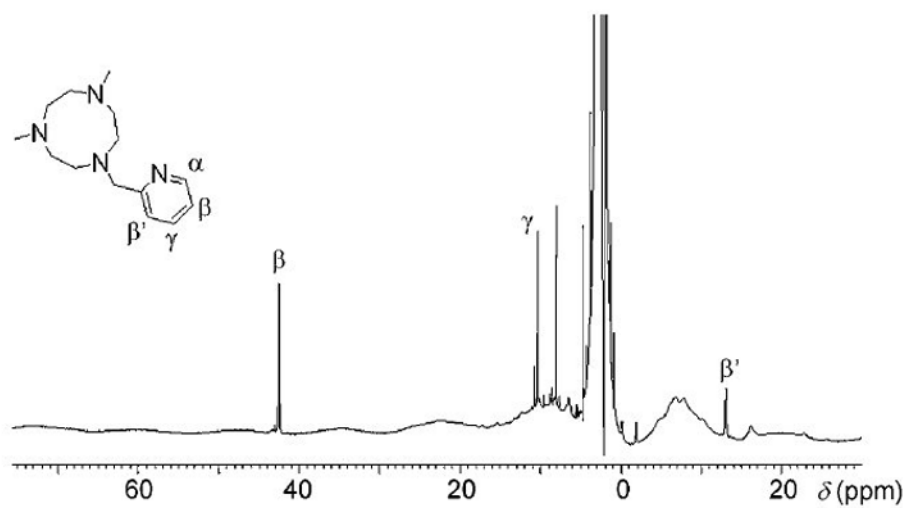
Figure 3.
^{1}H NMR spectra of **2** at 298 K in CD$_3$CN, along with selected signal identifications.
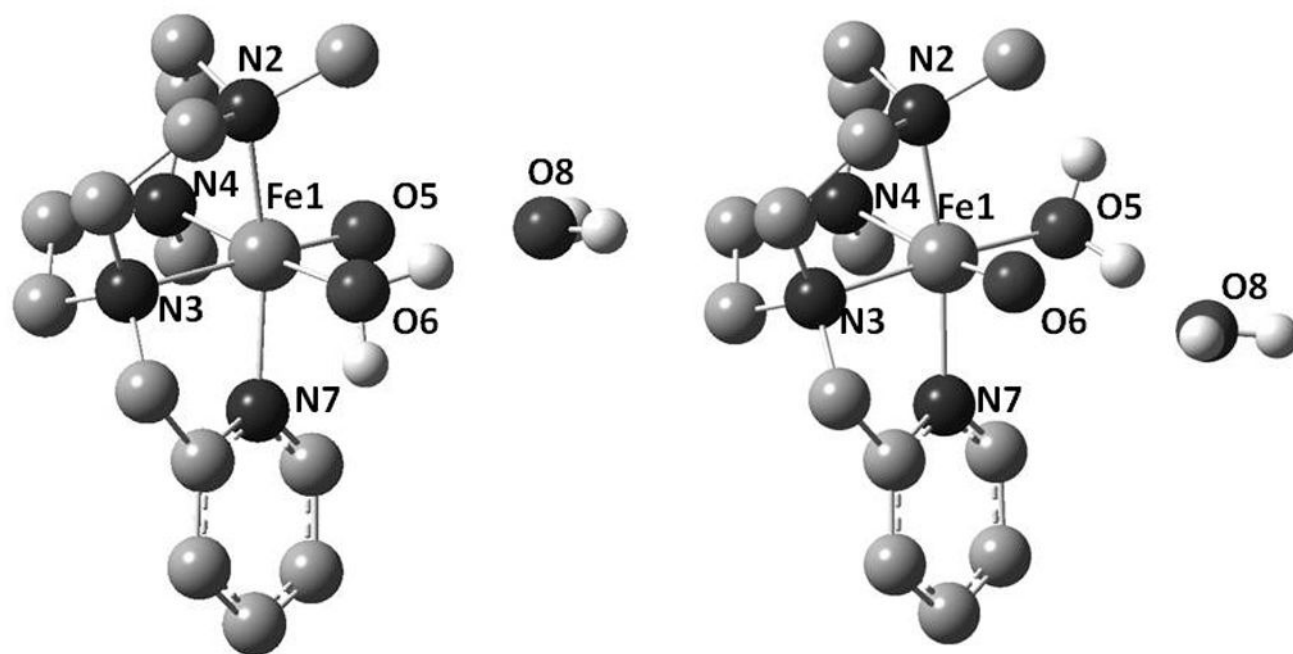
Figure 4.
DFT-calculated structures of **2a** (left) and **2b** (right). Selected distances (Å) for **2a**; Fe1-N2 2.05, Fe1-N3 2.11, Fe1-N4 2.06, Fe1-O5 2.06, Fe1-O6 1.62, Fe1-N7 2.03 and **2b**; Fe1-N2 2.07, Fe1-N3 1.99, Fe1-N4 2.19, Fe1-O5 1.63, Fe1-O6 2.03, Fe1-N7 2.02.
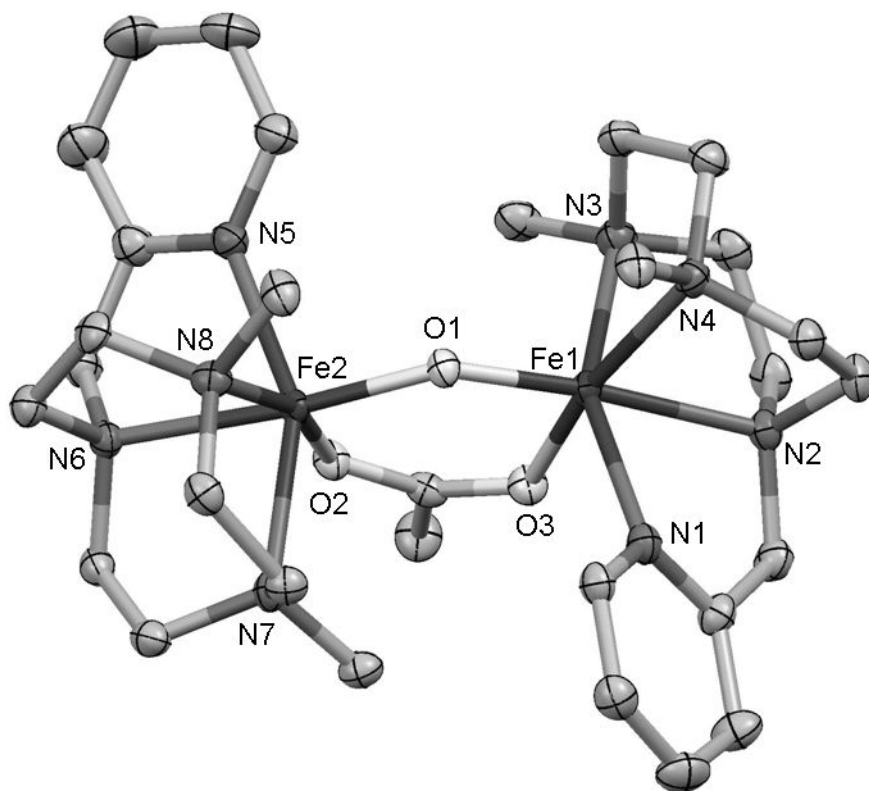
Figure 5.
ORTEP plot (50% probability ellipsoids) of [Fe$^{III}_2$(μ-O)(μ-CH$_3$CO$_2$)(^{Me,H}Pytacn)$_2$] (ClO$_4$)$_3$·CH$_3$CN, (**3**·CH$_3$CN). Selected distances in Å: Fe1-O1 1.801(3), Fe1-O3 2.062(4), Fe1-N1 2.167(4), Fe1-N3 2.180(4), Fe1-N4 2.197(4), Fe1-N2 2.225(4), O1-Fe2 1.805(3), Fe2-O2 2.017(3), Fe2-N5 2.163(4), Fe2-N8 2.184(4), Fe2-N7 2.196(4), Fe2-N6 2.247(4). ∠ O1-Fe1-O3 95.42(15)o, Fe2-O1-Fe1 135.2(2)o.
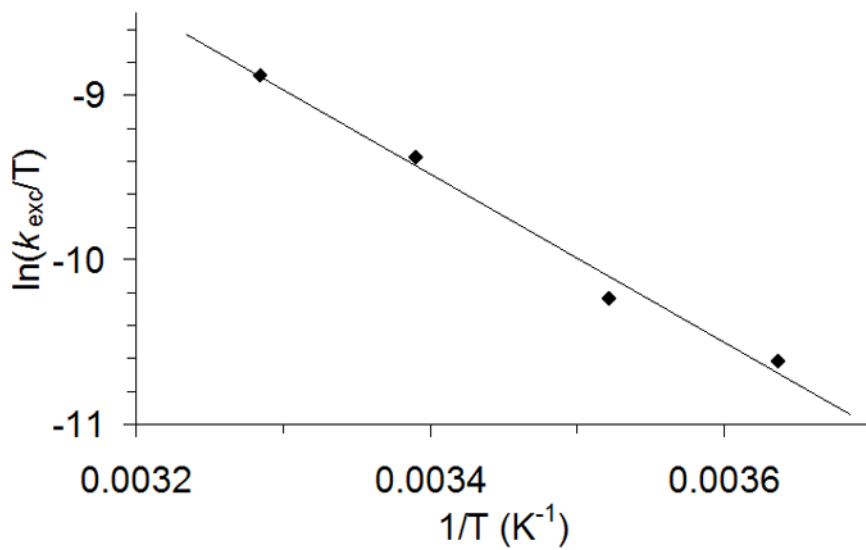
Figure 6.
Eyring plot for the determination of the activation parameters of the oxygen atom exchange of **2** with $H_2^{18}O$ ($[H_2^{18}O] = 0.2$ M).
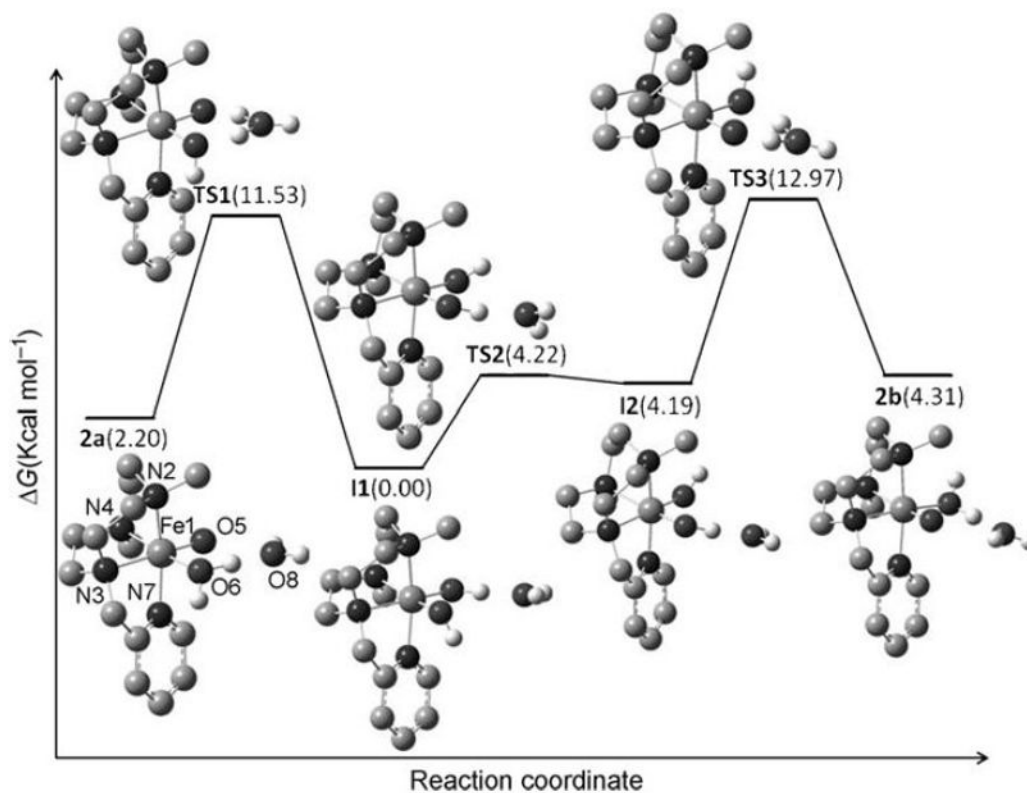
Figure 7.
Hammett plot representing log(k_X/k_H) against the Hammett parameter (σ_p) for the reaction of **2** with *p*-X-thioanisoles at 273 K.
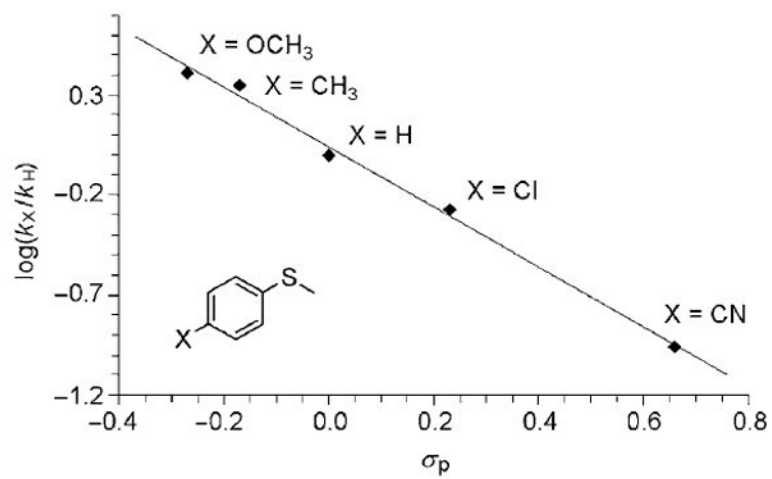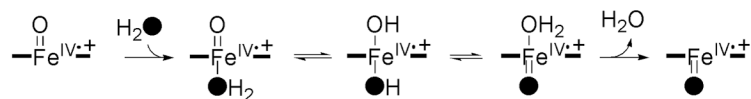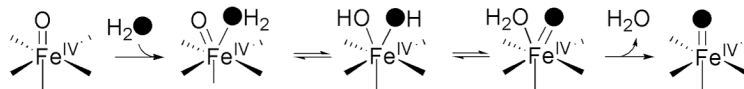
Figure 8.
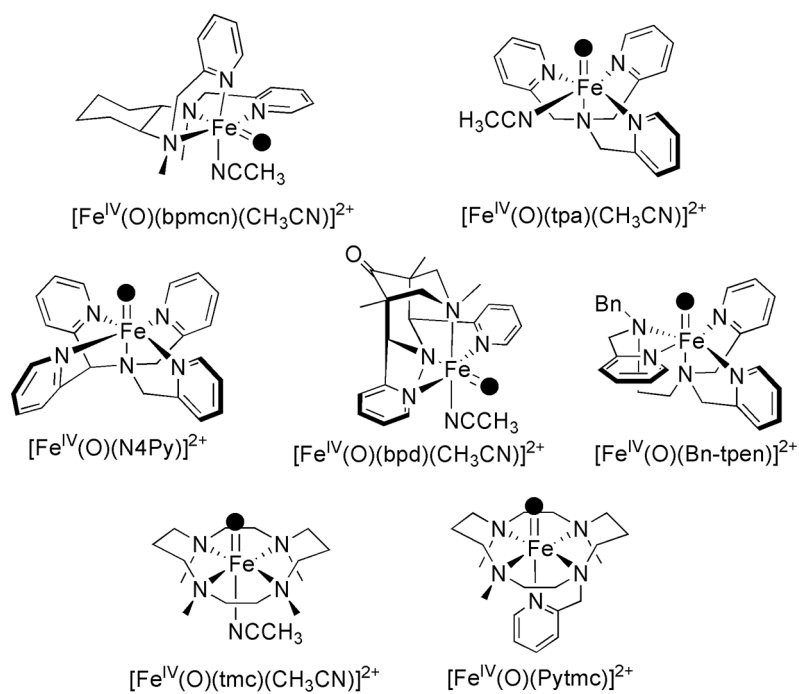Plot of $\log(k'_2)$ (determined at $-15°C$) against C-H BDE of different substrates.

heme systems

non-heme iron models: tmc and N4Py

Scheme 1.
Top: Proposed oxo-hydroxo tautomerism in heme systems. Bottom: Proposed oxo-hydroxo tautomerism in $[Fe^{IV}(O)(tmc)(CH_3CN)]^{2+}$ and $[Fe^{IV}(O)(N4Py)]^{2+}$.

[Fe^{IV}(O)(bpmcn)(CH₃CN)]²⁺

[Fe^{IV}(O)(tpa)(CH₃CN)]²⁺

[Fe^{IV}(O)(N4Py)]²⁺

[Fe^{IV}(O)(bpd)(CH₃CN)]²⁺

[Fe^{IV}(O)(Bn-tpen)]²⁺

[Fe^{IV}(O)(tmc)(CH₃CN)]²⁺

[Fe^{IV}(O)(Pytmc)]²⁺

Scheme 2.
Schematic representation of synthetic $Fe^{IV}(O)$ complexes relevant to this work.
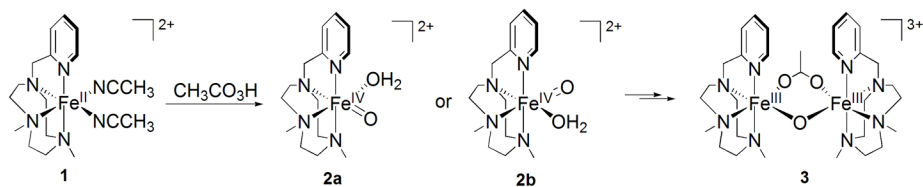
Scheme 3.
Schematic view of formation and decay of **2**, along with the proposed chemical structures of **1**, **2** and **3**.
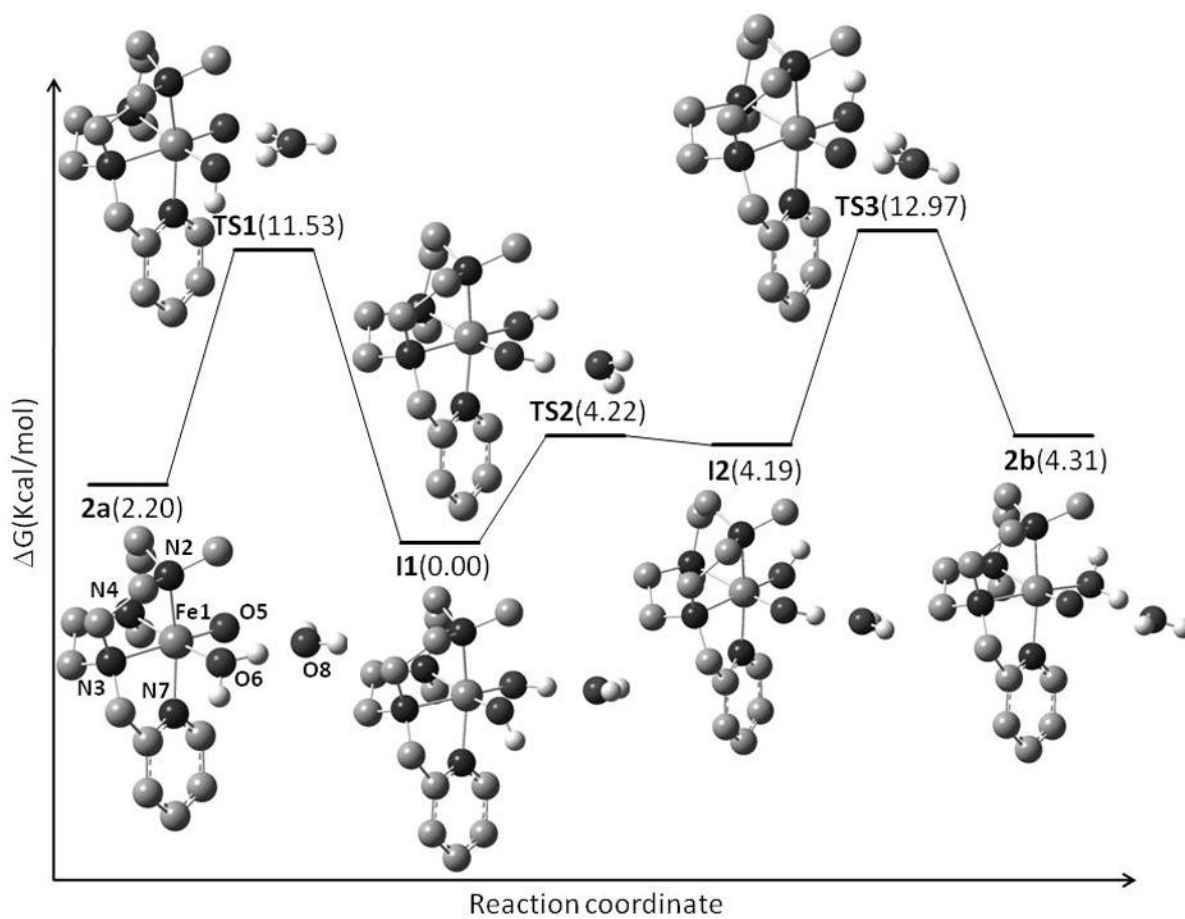
Scheme 4.
DFT mechanism for water exchange at **2**.

Table 1

Kinetic parameters for the water exchange reaction at selected oxo-iron(IV) complexes

Compound	$^a k_{exc} \times 10^{-3} M^{-1} s^{-1}$	ΔH kcal mol^{-1}	ΔS cal K^{-1} mol^{-1}	Ref
2	28±3	10.2±0.8	−32±3	-
[Fe(O)(tmc)(CH$_3$CN)]$^{2+}$	2.0^b	4.1±0.6	−57±8	[44]
[Fe(O)(N4Py)]$^{2+}$	2.4^b	4.0±0.6	−57±8	[44]
[Fe(O)(tpa)(S)]$^{2+}$	8±2	c	c	-

^{a}At 295K

^{b}Calculated from experimental data in ref. 44.

^{c}Eyring analysis could not be performed because of limited thermal stability.

Table 2

DFT-Calculated Mössbauer parameters[a] of oxo-iron(IV) (**2a** and **2b**) and *bis*-hydroxo-iron(IV) (**I1** and **I2**) tautomeric species.

Compound	ΔE_Q (mm·s^{-1})[a]	δ (mm·s^{-1})
2a	0.57	0.18
I1	2.28	0.12
I2	2.44	0.18
2b	0.83	0.17
Exp	0.73	0.05

[a]See experimental section for details on the DFT analysis.

Table 3

C-H Bond dissociation energies and reaction rates for the reaction of **2** with different substrates.[a]

Substrate	BDE (kcal·mol^{-1})[67],[68]	k_2 (M^{-1}s^{-1})
10-methyl-9,10-	73.7	86 ± 19
dihydroacridine (AcrH$_2$) xanthene	75.5	8.1 ± 0.8
9,10-dihydroanthracene (DHA)	77	5.7 ± 0.9
1,4-cyclohexadiene (CHD)	78	4.2 ± 0.3
fluorene	80	1.2 ± 0.2
2,3-dimethyl-2-butene	84	$(1.6 \pm 0.2) \times 10^{-2}$
tetrahydrofuran (THF)	93	$(2.3 \pm 0.3) \times 10^{-3}$
cyclohexane[b]	99.3	$(4 \pm 1) \times 10^{-4}$

[a]Reactions were run in acetonitrile at 258 K, using **2** generated in situ from the reaction of **1** (1 mM) with 2 equiv of peracetic acid at 15°C.

[b]298 K

Table 4

DHA oxidation rates of various iron-oxo complexes.

Complex	k_2 $(M^{-1}s^{-1})$	T (K)	Ref
2	5.7 ± 0.9	258	
$[Fe^{IV}(O)(N4Py)]^{+2}$	2.8	258	[75]
$[Fe^{IV}(O)(bpd)(CH_3CN)]^{+2}$	8.0	238	[35]
$[Fe^{IV}(O)(tmc)(CH_3CN)]^{+2}$	0.016	243	[24]
$[Fe^{III}(OMe)(Py5)]^{+2}$	5.0×10^{-3}	298	[66]
$[(OH)(L_{OMe})Fe^{III}(\mu-O)Fe^{IV}(O)(L_{OMe})]^{+2}$	28	193	[65]
$[Fe^{III}Fe^{IV}(\mu-O)_2(L_{OMe})_2]^{+3}$	10^{-5}	193	[65]
$[Fe^{IV}_2(\mu-O)_2(L_{OMe})_2]^{+4}$	10^{-4}	193	[65]
$[(OH)(L_{OMe})Fe^{IV}(\mu-O)Fe^{IV}(O)(L_{OMe})]^{+3}$	0.027	193	[65]
$[Fe^{IV}(O)(tmp)]$	2.7	258	[76]
$[Fe^{IV}(O)(TMG_3tren)]^{+2}$	0.090	243	[24]

L_{OMe} = *tris*((4-methoxy-3,5-dimethylpyridin-2-yl)d$_2$-methyl)amine, tmp = tetramesitylporphinate, TMG$_3$tren = 1,1,1-*tris*{2-[N^2-(1,1,3,3-tetramethylguanidino)]ethyl}amine.