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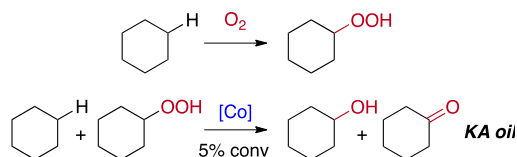
Iron Triflate Salts as Highly Active Catalysts for the Solvent-Free Oxidation of Cyclohexane

Pierre-Adrien Payard,^[a,b] Yu-Ting Zheng,^[a] Wen-Juan Zhou,^{[a]*} Lhoussain Khrouz,^[c] Laurent Bonneviot,^[c] Raphael Wischert,^[a] Laurence Grimaud,^[b] Marc Pera-Titus^{[a]*}

Abstract: Among a series of iron salts, iron triflates revealed as highly active catalysts for the oxidation of cyclohexane by *t*-butyl hydroperoxide into cyclohexanol and cyclohexanone with initial turnover frequencies higher than 10,000 h⁻¹. The structure of the iron complexes under reaction conditions was studied by combining electron paramagnetic resonance (EPR) spectroscopy and DFT calculations. The coordination of the catalytic iron center readily evolved in the presence of the reaction products, leading ultimately to its deactivation. Iron and organic superoxo intermediates were identified as plausible active species allowing to rationalize the high activity of iron ligated by highly-delocalized counter-anions.

Introduction

The oxidation of cyclohexane (CyH) to cyclohexanol (CyOH) and cyclohexanone (CyONE), also known as KA oil, is a well-known industrial process for the production of adipic acid (AA).^[1] This process comprises two consecutive oxidation steps (Scheme 1): (i) non-catalytic autoxidation of CyH by O₂ yielding cyclohexyl hydroperoxide (CyHP), and (ii) deperoxidation of CyHP to KA oil catalyzed by a transition metal, typically a homogeneous cobalt catalyst.^[2] The process is conducted at very low CyH conversion (~5%) with a CyHP selectivity typically of 60-70% in the first step.^[3] The industrial challenge is to increase the catalytic activity for KA formation while keeping the CyHP concentration as low as possible using a clean and safe route. In this view, a fast and selective catalytic pathway using a non-toxic metal would be desirable.



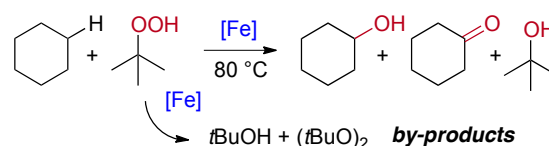
Scheme 1. Industrial process for the oxidation of CyH to KA oil.

High-valent iron-oxo complexes (e.g., cytochrome P450) are known to efficiently catalyze oxidation reactions in living systems.^[4] Iron-oxo systems have been extended to non-heme bio-inspired catalysts based on iron centers,^[5] and to mononuclear iron complexes.^[6] The most studied and simplest non-heme iron system relies on the so-called Fenton reagent consisting of an aqueous solution of hydrogen peroxide and an iron salt.^[7] This reagent has been widely used in the total oxidation of organic pollutants in liquid waste treatment, as it generates very active and non-selective OH[•] radicals.^[8] The search for high selectivity in partial oxidation reactions has led to the development of alternative catalytic systems comprising highly oxidized iron-oxo intermediates (IV or V).^[9] To stabilize high-valent iron intermediates, a ligand is generally used.^[10]

A well-known application of iron complexes for CyH oxidation relies on the pioneering work of Barton, using an iron catalyst combined with metallic zinc and oxygen as oxidant.^[11] This chemistry was improved by replacing Fe(II)/Zn⁰/O₂ by Fe(II)/superoxide. Using an analogous system based on Fe(III)/TBHP (TBHP = *t*-butyl hydroperoxide) based on tris(trimethylacetate)iron(III) [Fe(tma)₃] and tris(1,1,1,2,2,3,3-heptafluoro-7,7-dimethyl-4,6-octanodionate)iron(III) [Fe(fod)₃], Schuchardt *et al* reported a selectivity of 91% at 11% conversion for CyH oxidation.^[12] However, the product (AA) deactivated the active iron centers.

Recently, iron triflates, i.e. Fe^{II}(OTf)₂ and Fe^{III}(OTf)₃, have been reported as active catalysts in low-coordinating solvents for a variety of Lewis-acid catalyzed reactions.^[13] Typical examples include imination of sulfoxides,^[14] aziridination of enol silyl ethers,^[15] hydroaddition to unsaturated C-C bonds,^[16] etherification and transesterification of alcohols,^[17] and glycosylation of sugar derivatives.^[18] Despite the potential of iron triflates for C-H activation in C-N,^[19] C-C,^[20] and C-X (X = F, Cl, I)^[21] coupling reactions, few reports are available on oxidation reactions.^[5b,22]

Herein we report an unprecedented high catalytic activity of Fe^{III}(OTf)₃ for the oxidation of neat CyH towards CyOH and CyONE (KA oil) using TBHP as oxidant (Scheme 2). Fe^{III}(OTf)₃ behaved as precursor of the active iron species for CyH oxidation. We provide detailed insight into the structure of the catalyst under reaction conditions by combining electron paramagnetic resonance (EPR) and DFT calculations.



Scheme 2. Oxidation of CyH to KA oil catalyzed by Fe catalysts.

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Table 1. CyH and TBHP conversion, KA selectivity and yield, A/K molar ratio and TOFs as a function of the catalyst at 0.14 mM Fe^{III}(OTf)₃ (10 ppm Fe)^[a]

Complex ^[b]	CyH conv (%)	TBHP conv (%)	KA sel (%)	KA yield (%)	A/K	TON
Fe ^{III} (OTf) ₃	1.9	73	35	25	6.4	1175
Fe ^{III} (OTf) ₂	1.2	74	21	16	3.3	765
Fe ^{III} (NTf ₂) ₃	1.0	69	23	16	3.1	651
Cl-Fe ^{III} TPP	0.9	69	17	12	2.8	574
Fe ^{III} (OTs) ₃	0.7	57	19	11	3.3	455
Fe ^{III} Cl ₃	0.4	30	16	5	5.5	231

^[a] Reaction conditions: 80 °C, CyH (4.0 g) containing TBHP (7.5 wt%, 3.3 mmol), 0.14 mmol catalyst (10 ppm Fe), 60 min. ^[b] OTf: SO₂CF₃, NTf₂: N(SO₂CF₃)₂, TPP: tetraphenylporphyrine, OTs: OSO₂Tol

Table 2. CyH and TBHP conversion, KA selectivity and yield, A/K molar ratio and TON as a function of the Fe^{III}(OTf)₃ concentration^[a]

C _{Fe} (mM)	CyH conv (%)	TBHP conv (%)	KA sel (%)	KA yield (%)	A/K	TON
0.014	0.43	6.6	86	5.7	82	2658
0.070	1.4	52	34	18	8.6	1670
0.14	1.8	81	29	24	4.0	1140
0.56	2.1	88	30	27	2.3	310
1.95	2.3	96	32	30	2.1	101

^[a] Reaction conditions: 80 °C, CyH (4.0 g) containing TBHP (7.5 wt%), 60 min

Table 3. CyH and TBHP conversion, KA selectivity and yield, A/K molar ratio and TON as a function of the temperature^[a]

T (°C)	CyH conv (%)	TBHP conv (%)	KA sel (%)	KA yield (%)	A/K	TON
40	0.4	30	16	5	5.5	231
45	0.7	57	19	11	3.3	455
50	0.9	69	17	12	2.8	574
55	1.0	69	23	16	3.1	651
60	1.2	74	21	16	3.3	765
65	1.3	78	25	19	3.6	811
70	1.5	80	28	22	3.6	922
80	1.8	81	29	24	4.0	1140

^[a] Reaction conditions: 80 °C, CyH (4.0 g) containing TBHP (7.5 wt%), 60 min, 0.14 mM Fe^{III}(OTf)₃ (10 ppm Fe), 60 min

Results and Discussion

Definition of the catalyst performance metrics

The catalytic performance of the different Fe complexes was primarily characterized in terms of the TBHP conversion, KA selectivity and KA yield with respect to TBHP, as well as by the turnover frequency at time $t = 0$ (TOF₀) and the turnover number (TON) at 60 min.^[23] More details on the reaction metrics can be found in the SI (Experimental section, section 1.3).

CyH oxidation using TBHP as oxidizing agent

Screening of Fe complexes

The catalytic activity of a series of Fe complexes was first investigated using CyH as reactant and TBHP as an oxidant (7.5 wt% or 0.65 mM in CyH, see SI for preparation, section 1.2) at 80 °C under reflux. For comparison, a porphyrin based Fe salt (Cl-Fe-TPP) was chosen as reference catalyst, and its reactivity was compared to that of Fe(II) and Fe(III) salts, i.e. Fe^{II}(OTf)₂, Fe^{III}(OTf)₃, Fe^{III}(NTf₂)₃, Fe^{III}(OTs)₃ and Fe^{III}Cl₃, featuring in all cases a highly delocalized, poorly coordinating, counter anion.

The products and reactants were analyzed by GC (see SI, section 1.3 for detailed method) after 60 min (Table 1). Among the different Fe complexes, the least soluble in CyH is Fe^{III}(OTf)₃ with a solubility of 0.59 mM and 0.24 mM at 20 °C and 80 °C, respectively (see SI, section 1.4 for solubility measurements). In light of these results, a Fe concentration of 0.14 mM (10 ppm Fe) was chosen to compare the catalytic activity of the complexes. The most active complexes for CyH oxidation also appear to be the most selective towards KA oil (entries 1–3) by minimizing undesired TBHP disproportionation to *t*-butyl alcohol and O₂, and the formation of ^tBuO-O^tBu (Scheme 2). Note that ^tBuO-OCy was not detected, opposing previous studies encompassing CyH oxidation by TBHP using solvents (e.g., acetonitrile) under dilute conditions.^[24]

Among the different Fe complexes, Fe^{III}(OTf)₃ exhibits the highest TBHP conversion and KA yield with values of 73% and 25%, respectively. Also, the TON after 60 min is the highest with a value of ca. 1200. Surprisingly, both triflate and triflimide Fe salts perform much better than the porphyrin based Fe salt. The TON decreases in the order: Fe^{III}(OTf)₃ > Fe^{III}(OTf)₂ > Fe^{III}(NTf₂)₃ > Cl-Fe^{III}TPP > Fe^{III}(OTs)₃ > Fe^{III}Cl₃. Noteworthy, at the reaction conditions tested, Fe^{II}(OTf)₂ is slightly less active than Fe^{III}(OTf)₃. However, Fe^{III}(OTf)₃ shows a higher alcohol-to-ketone molar ratio (A/K) (6.4 vs. 3.3).

Effect of the Fe^{III}(OTf)₃ concentration

The catalytic properties of Fe^{III}(OTf)₃ were then explored at variable catalyst concentration in the range 0.014–1.95 mM (1–140 ppm Fe) (Table 2). Increasing the Fe concentration from 0.014 mM (1 ppm Fe) to 0.56 mM (40 ppm Fe) clearly promotes the CyH and TBHP conversion, but at the expense of the TON. This observation suggests that a high catalyst concentration (especially >0.14 mM or 10 ppm Fe) does not enhance the catalytic activity. The highest TON at 60 min (about 2600) is achieved at very low KA yield (5.7%) and at the lowest Fe^{III}(OTf)₃ concentration (0.014 mM). Besides, the A/K molar ratio declines at higher Fe concentration from 82 at 0.014 mM to 4.0 at 0.14 mM, suggesting that CyONE is a secondary oxidation product of CyH.

Effect of the reaction temperature

The effect of the temperature was studied in the range 40–100 °C using 0.14 mM Fe^{III}(OTf)₃ (10 ppm Fe) (Table 3). CyH oxidation is enhanced at higher temperature at the expense of disproportionation into *t*-butyl alcohol and O₂, with KA oil being enriched in CyOH. In parallel, the A/K molar ratio decreases with the temperature from an initial value of 5.5 at 40 °C to 2.8–4.0 in the temperature range 45–80 °C.

Reactant and product evolution during the reaction

The reaction kinetics was measured at different temperatures during the first 20 min of reaction using Fe^{III}(OTf)₃ and Fe^{III}(NTf₂)₃ as catalysts (Figure 1). For both catalysts, KA oil formation proceeds fast during the first 15 min with a decrease of the A/K molar ratio, but stabilizes after 10–20 min for temperatures higher than 50 °C. At first sight, this observation

suggests catalyst deactivation, which could be caused either by the reaction products, or from by-products. Indeed, traces of AA were detected, which combined with water are known to deactivate Fe centers.^[5c,12] To gain more insight into the underlying mechanism of Fe deactivation, CyOH, CyONE and AA were added to the reaction media in the beginning of the reaction. While CyOH or CyONE only exert a slight effect, AA strongly lowers the KA yield, thus confirming the deactivating role of AA (Table 4).

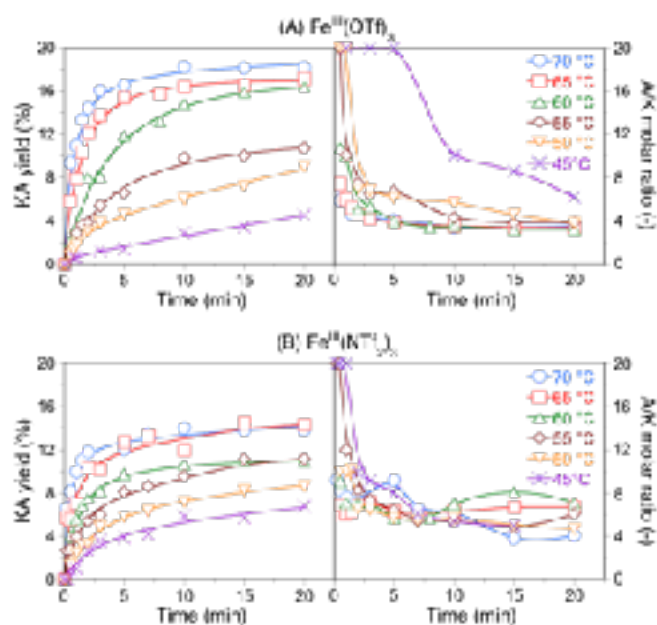


Figure 1. Time-evolution of KA yield and A/K molar ratio in CyH oxidation catalyzed by (A) $\text{Fe}^{\text{III}}(\text{OTf})_3$ and (B) $\text{Fe}^{\text{III}}(\text{NTf}_2)_3$ with TBHP as oxidant. Reaction conditions: 45–70 °C, 0.14 mM catalyst (10 ppm Fe), 4 g of CyH containing 7.5 wt% (3.3 mmol) TBHP, 0–20 min.

Table 4. Evolution of the KA yield upon addition of CyOH, CyONE and AA before the reaction.^[a] The concentration of products was adjusted to magnify the effect on the KA yield.

Entry	Additive	KA Yield %	Δ KA Yield %
0	-	25	0
1	CyOH (0.212 mM)	28	+3
2	CyONE (0.084 mM)	21	-4
3	AA (0.095 mM)	16	-9
4	CyOH (0.198 mM) + CyONE (0.045 mM)	22	-2
5	CyOH (0.220 mM) + AA (0.080 mM)	12	-13
6	CyONE (0.059 mM) + AA (0.091 mM)	9	-16
7	CyOH (0.195 mM) + CyONE (0.049 mM) + AA (0.078 mM)	8	-17

^[a] Reaction conditions: 80 °C, 10 ppm $\text{Fe}^{\text{III}}(\text{OTf})_3$, 4 g of CyH containing 7.5 wt% TBHP (3.3 mmol), 60 min.

TOF and activation energy for CyH oxidation

The reaction rate (i.e. TOF_0) was measured at time $t = 0$ in the temperature range 45–70 °C for $\text{Fe}^{\text{III}}(\text{OTf})_3$ and $\text{Fe}^{\text{III}}(\text{NTf}_2)_3$. In both cases, very high TOFs of 14,000 h^{-1} and 31,000 h^{-1} , respectively, at 60 °C, are observed. The activation energy measured from the Arrhenius plots of $\ln(\text{TOF}_0)$ vs. $1/T$ is 145 ± 12 kJ/mol for $\text{Fe}^{\text{III}}(\text{OTf})_3$ and 115 ± 6 kJ/mol for $\text{Fe}^{\text{III}}(\text{NTf}_2)_3$

(Figure 2). At all temperatures, the TOF_0 is higher for $\text{Fe}^{\text{III}}(\text{NTf}_2)_3$. However, higher KA yields and TONs at 60 min are obtained with $\text{Fe}^{\text{III}}(\text{OTf})_3$ (Table 1, entries 1 and 3). This observation points out a faster deactivation of $\text{Fe}^{\text{III}}(\text{NTf}_2)_3$ during the reaction despite its higher initial reaction rate.

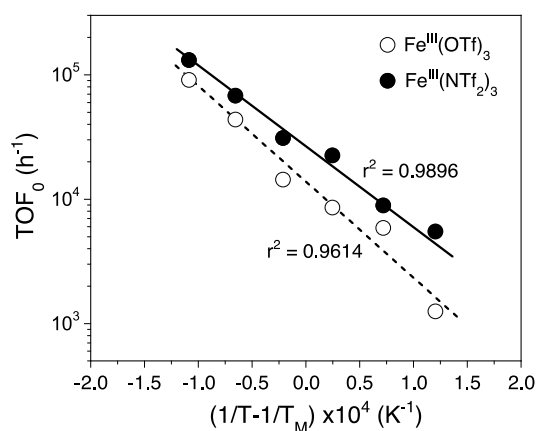


Figure 2. Arrhenius plots of TOF_0 for $\text{Fe}^{\text{III}}(\text{OTf})_3$ and $\text{Fe}^{\text{III}}(\text{NTf}_2)_3$. TM refers to the mean temperature of the series ($T_M = 330.7$ K). Reaction conditions as in Figure 1.

EPR investigation of Fe(III) coordination

EPR is a sensitive technique for the analysis of Fe(III) species with unpaired electrons at very low concentration. In frozen solution, the g -tensor component of Fe(III) can be measured at low- and high-spin states providing information on the symmetry and coordination sphere of the species and their evolution during the reaction. In the case of high-spin Fe(III), one should also consider the contribution of the zero-field-splitting (ZFS), which can be assessed by the so-called the D tensor. This tensor includes two components, i.e. D and E , accounting for the intensity and anisotropy of the interaction.^[25] $\text{Fe}(\text{II})$ ions, which are also paramagnetic are silent in conventional X-band (9.6 GHz) spectrometers.

Coordination sphere of Fe(III)

A CyH solution of $\text{Fe}^{\text{III}}(\text{OTf})_3$ exhibits an EPR signal at $g = 2.02$ (Figure 3), which is also observed on the solid salt (Figure S1). Since the triflate ion is usually described as a weak ligand, one could reasonably expect it to be in high-spin state ($S = 5/2$). Indeed, $\text{Fe}^{\text{III}}(\text{OTf})_3$ and $\text{Fe}^{\text{III}}(\text{NTf}_2)_3$ have been described as high-spin complexes, as also confirmed by DFT calculations (see SI, Table S1).^[26] In our case, the fit of the EPR spectrum of $\text{Fe}^{\text{III}}(\text{OTf})_3$ using the Matlab[®] routine Easyspin is consistent with a single weakly distorted high-spin Fe center ($g = 2.012$, $D = 0.01$ cm^{-1} , $E/D = 0.1$, see fitted spectrum in SI).^[27] DFT calculations (see SI for Computational Details) yield a structure for $\text{Fe}^{\text{III}}(\text{OTf})_3$ featuring three bidentate triflate anions interacting with the Fe center through two oxygen atoms (Figure 3). The metal-ligand interaction in $\text{Fe}^{\text{III}}(\text{OTf})_3$ is mainly ionic as confirmed by NBO analysis: both Fe and O are strongly charged (+1.7 on Fe and -0.9 on O), and the spin is located mainly on Fe (spin density on Fe: 4.0, on O: 0.1). The values $g_x = g_y = g_z = 2.01$ and $D = 0.09$ cm^{-1} were obtained using the EPR calculation module of ORCA, in good agreement with the experimental spectrum.

Coordination sphere of Fe(III) in the presence of CyOH and CyONE

As pointed out above, CyONE and CyOH are potential ligands to Fe(III), but showing a contrasting effect on the catalytic activity:

CyONE seems to promote deactivation, whereas CyOH enhances the activity. To gain more insight into the coordination ability of CyONE and CyOH on the Fe(III) centers, a dedicated EPR study was performed.

The addition of CyONE to a CyH solution of $\text{Fe}^{\text{III}}(\text{OTf})_3$ (10 mM) does not alter the EPR signal (Figure 4, Figure S2). This result is rather unexpected, since ketones should interact with Fe(III), generating a Lewis adduct. Indeed, the Gibbs free energies of coordination computed by DFT strongly support this idea (Table 5): the most stable adduct between $\text{Fe}^{\text{III}}(\text{OTf})_3$ and CyONE, i.e. *mer*- $\text{Fe}^{\text{III}}(\text{OTf})_3(\text{CyONE})_3$, is favored by -134 kJ mol^{-1} . However, the electronic structure of *mer*- $\text{Fe}^{\text{III}}(\text{OTf})_3(\text{CyONE})_3$ is very similar to that of $\text{Fe}^{\text{III}}(\text{OTf})_3$, resulting in almost the same EPR spectrum (Figure 4, blue and red curves). A high-spin, highly charged Fe(III) center (+1.8) is observed with a spin mainly located on Fe (spin density = 4.0). This center is prone to interact with negatively charged oxygen atoms (O of OTf -1.0, O of CyONE -0.6). Since the partial charge is slightly lower on the CyONE oxygen atom, the Fe-O bond is slightly longer compared to the Fe-O bond generated with the O atoms of triflates (2.11 Å vs 1.94 Å). Accordingly, the adduct geometry still approaches a slightly distorted octahedron. As the coordination of CyONE has little impact on the electronic density of Fe, no relevant difference is expected in the EPR spectrum. Indeed, the parameters calculated by DFT are close to the previous ones: $g = 2.01$ and $D = 0.06 \text{ cm}^{-1}$.

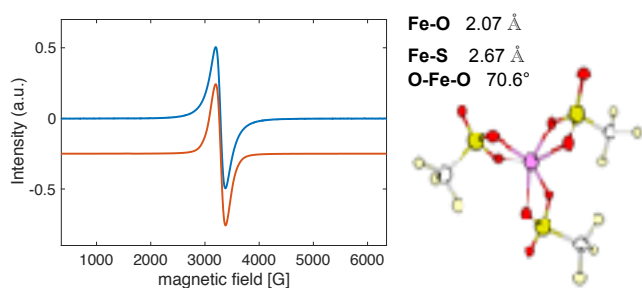


Figure 3. Experimental (blue) and fitted (orange) EPR spectra at 110 K of $\text{Fe}^{\text{III}}(\text{OTf})_3$ in CyH (10 mM).

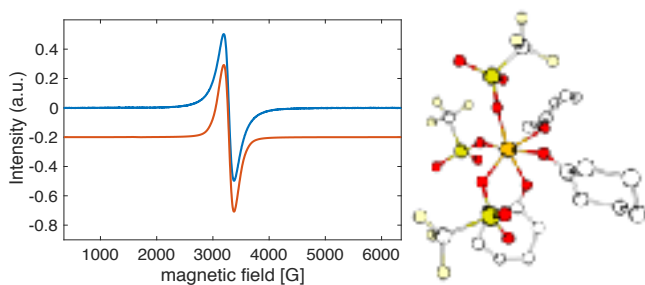


Figure 4. Experimental (blue) and fitted (red) EPR spectra at 110 K of $\text{Fe}^{\text{III}}(\text{OTf})_3$ (10 mM) in CyH in the presence of CyONE (10 equiv).

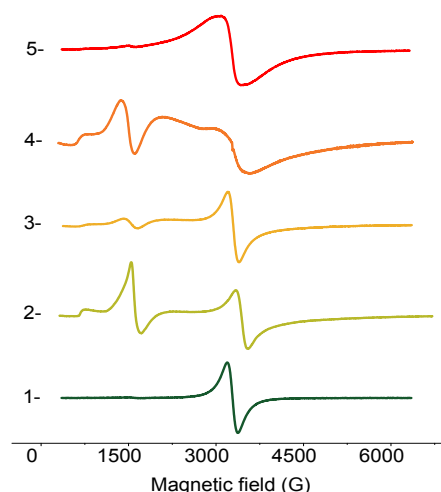


Figure 5. EPR spectra at 110 K of $\text{Fe}^{\text{III}}(\text{OTf})_3$ (10 mM) in CyH in the presence of: (1) CyOH (3 equiv), (2) CyOH (10 equiv), (3) CyONa (1 equiv), (4) CyOH (10 equiv, after 1 night), and (5) CyONa (10 equiv).

Table 5. DFT-calculated enthalpy (ΔH , kJ mol^{-1}) and Gibbs free energy at 25 °C (ΔG , kJ mol^{-1}) of high spin complexes resulting from the coordination of CyOH and CyONE to $\text{Fe}^{\text{III}}(\text{OTf})_3$.

Species	$\Delta G^{[a]}$ (kJ mol^{-1})	$\Delta H^{[a]}$ (kJ mol^{-1})	g	$ D $	E/D
$\text{Fe}(\text{OTf})_3$ (1)	0	0	2.0_1	0.087	0.049
$\text{Fe}(\text{OTf})_3(\text{CyONE})$ (2)	-47.6	-104.4	2.0_1	0.085	0.181
$\text{Fe}(\text{OTf})_3(\text{CyONE})_2$ (3)	-90.8	-214.5	2.0_1	0.078	0.113
<i>mer</i> - $\text{Fe}(\text{OTf})_3(\text{CyONE})_3$ (4)	-133.3	-307.4	2.0_1	0.065	0.152
<i>fac</i> - $\text{Fe}(\text{OTf})_3(\text{CyONE})_3$ (5)	-98.9	-276.0	2.0_1	0.050	0.291
$\text{Fe}(\text{OTf})_3(\text{CyOH})$ (6)	-22.6	-89.4	2.0_1	0.099	0.320
$\text{Fe}(\text{OTf})_3(\text{CyOH})_2$ (7)	-99.3	-224.7	2.0_1	0.078	0.300
<i>mer</i> - $\text{Fe}(\text{OTf})_3(\text{CyOH})_3$ (8)	-155.8	-338.0	2.0_1	0.027	0.305
<i>fac</i> - $\text{Fe}(\text{OTf})_3(\text{CyOH})_3$ (9)	-93.7	-284.4	2.0_1	0.066	0.095
$\text{Fe}(\text{OTf})_2(\text{CyO})$ (10)	-19.8	-67.1	2.0_1	0.403	0.212
$\text{Fe}(\text{OTf})_2(\text{CyOH})(\text{CyO})$ (11)	-44.8	-151.7	2.0_1	0.377	0.065
$\text{Fe}(\text{OTf})_2(\text{CyOH})_2(\text{CyO})$ (12)	-85.3	-247.7	2.0_1	0.417	0.147

^[a] The enthalpy and Gibbs free energies of complex formation were calculated with respect to free $\text{Fe}^{\text{III}}(\text{OTf})_3$, CyOH and CyONE. In the case of complexes **10** to **12**, the formation of CyOH_2OTf ion pair was considered ($\Delta G = -6.8 \text{ kJ mol}^{-1}$, $\Delta H = -61.6 \text{ kJ mol}^{-1}$), since triflic acid is less favorable.

Likewise, the addition of CyOH to $\text{Fe}(\text{OTf})_3$ does not modify the EPR spectrum upon addition up to 3 equiv (Figure 5-1). As for CyONE, DFT calculations confirm that simple coordination of CyOH is not expected to modify the signal to an important extent (Table 5, complexes **3-5**). Conversely, in the presence of an excess of CyOH, a new signal appears at ca. 1500 G (Figure 5-2, Figure S3, S4), while the solubility of the salt increases significantly. The fitting of the spectrum reveals the presence of at least three different high-spin species. The first species corresponds to that described in Figure 5-1, whereas the remaining two species possess a slightly expanded g -tensor

[2.008 2.0005 2.000] and a higher zero field splitting interaction ($D = 0.5 \text{ cm}^{-1}$; $E/D = 0.33$ and $D = 0.7 \text{ cm}^{-1}$; $E/D = 0.3$), being responsible for the low-field signal. From these observations, we could assume that both signals are ascribed to Fe-alkoxy complexes. Indeed, the simulated D and E parameters drastically increase when considering an alkoxy as ligand (Table 5, complexes **10-12**).

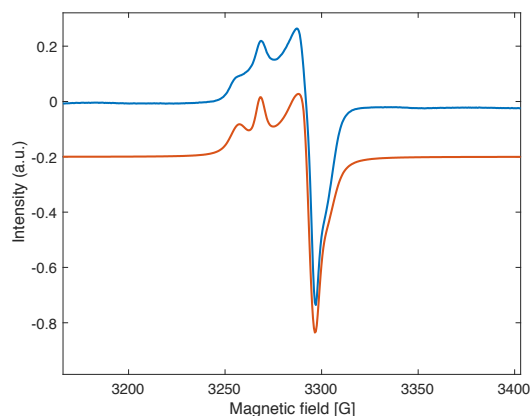


Figure 6. Experimental (blue) and fitted (red) EPR spectra at 110 K of $\text{Fe}^{\text{III}}(\text{OTf})_3$ (10 mM) in a solution of TBHP in CyH (7.5 wt%).

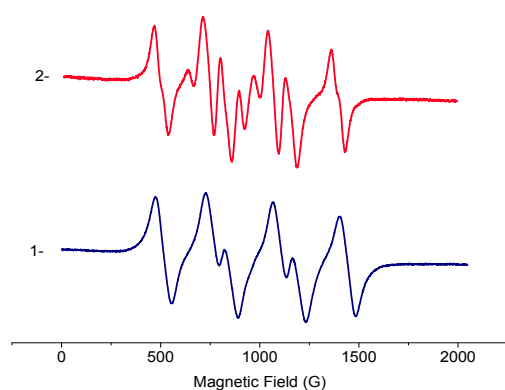


Figure 7. EPR spectra at room temperature of $\text{Fe}^{\text{III}}(\text{OTf})_3$ (10 mM) in a solution of TBHP in CyH (7.5 wt%) in the presence of: (1) DMPO (5 μL), (2) DMPO (5 μL) and CyONE (10 equiv).

The formation of monoalkoxy Fe species, i.e. $(\text{Fe}(\text{OTf})_3 + (n + 2) \text{CyOH} = \text{Fe}(\text{OTf})_2(\text{CyO})(\text{CyOH})_n + \text{CyOH}_2.\text{OTf}$, with $n = 0, 1$ and 2 , complexes **10-12**), turns out to be thermodynamically favorable, while species featuring multiple alkoxy ligands such as $\text{Fe}(\text{OTf})(\text{CyO})_2(\text{CyOH})_n$ and $\text{Fe}(\text{CyO})_3(\text{CyOH})_n$, with $n = 0, 1, 2$ and 3 (Table S2), are less favorable.^[28] The latter complexes are all predicted to be high-spin species (see SI). The formation of iron-alkoxy species was experimentally confirmed by adding variable amounts of sodium cyclohexanate (CyONa) to $\text{Fe}^{\text{III}}(\text{OTf})_3$ (Figure 5-3, Figure S5). Indeed, after addition of 1 equiv of CyONa, the low-field signal immediately emerges. However, under CyONa excess, only a single broad signal appears at around $g = 2$ (Figure 5-5). Interestingly, a similar evolution occurs when heating at $40 \text{ }^\circ\text{C}$ overnight a solution of $\text{Fe}^{\text{III}}(\text{OTf})_3$ with CyOH (10 equiv) in CyH (Figure 54). Since the formation of low-spin complexes seems unlikely with only alkoxy ligands, which is confirmed by DFT (Table S1), this observation might be explained by the formation of Fe(III) clusters featuring CyO^- as bridging ligand. In this view, the weak dipolar interaction between localized spin momenta would broaden the signal due to unresolved coupling.^[29] Indeed, these signals were

successfully fitted using an increasing proportion of Gaussian part in the function (Figure S5, S6).

Catalyst ageing

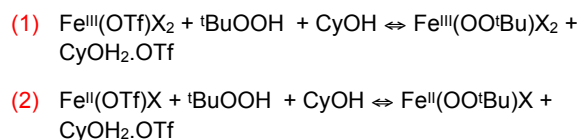
The signal attributed to Fe clusters is also observed in the EPR spectrum measured after the reaction, when the catalyst is supposed to be deactivated (Figure S7). Besides the above mentioned broad signal,^[30] the spectrum can be decomposed into a mixture of high-spin complexes with high ZFS interaction ($D = 0.6 \text{ cm}^{-1}$ $E/D = 0.33$, and $D = 0.7 \text{ cm}^{-1}$, $E/D = 0.21$). All the signals may be attributed to alkoxy or hydroxy Fe complexes. Overall, CyOH is expected to solubilize the salt, but in excess may inhibit the Fe centers.

Reaction intermediates and proposed mechanism

When TBHP is added to a solution of $\text{Fe}^{\text{III}}(\text{OTf})_3$ in CyH, the signal ascribed to Fe(III) vanishes and only sharp signals at $g = [2.024 \ 2.009 \ 2.007]$ and $g = [2.031 \ 2.011 \ 2.003]$ are visible (Figure 6, Figure S8). These signals are characteristic of the superoxide radical $\text{O}_2^{\cdot-}$, either free or stabilized on an EPR-silent Fe center ($\text{Fe}^{\text{II}}-\text{OO}^\cdot$).^[31] Surprisingly, no signal ascribed to $\text{Fe}^{\text{III}}(\text{OO}^t\text{Bu})$ is observed.^[5c] Radical trapping experiments were also carried out using 5,5-dimethyl-1-pyrroline N-oxide (DMPO) (Figure 7).^[32] Under standard reaction conditions, a single radical is observed (Figure 7-1, $S = 1/2$, $g = 2.004$, $a_N = 13.25 \text{ G}$, $a_H = 9.98, 1.45 \text{ G}$), corresponding to the adduct formed between DMPO and $^t\text{BuO}^\cdot$ (or $^t\text{BOO}^\cdot$).^[33] No traces of OH^\cdot , typical of Fenton-type systems, or $^t\text{Bu}^\cdot$ trapped by DMPO, are observed. The addition of CyONE (10 equiv) results in a similar signal, but less intense, while a second species appears (Figure 7-2, $S = 1/2$, $g = 2.006$, $a_N = 7 \text{ G}$, $a_H = 3.5 \text{ G}$, see also Figure S9, S10) due to DMPO oxidation.^[34] This experiment reinforces the idea of Fe poisoning by CyONE.

With these results in hand and supported by the literature, we provide in the following lines a tentative list of reactions participating during CyH oxidation with the detected species highlighted in bold. For a critical review on Fe-peroxo catalytic mechanisms, see Barbusinsky et al.^[35]

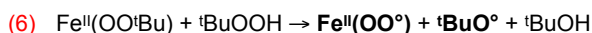
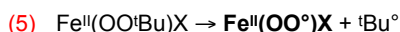
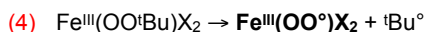
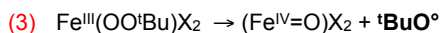
The first reaction involves the deprotonation of TBHP to yield Fe-peroxo complexes (Eq 1-2) ($X = \text{OTf}, \text{CyO}^-, \text{OH}^-$ or $^t\text{BuOO}^-$). Note that the coordination sphere may be completed by additional neutral ligands such as CyOH, CyONE and H_2O , which, for the sake of clarity, have been omitted.



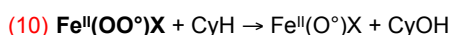
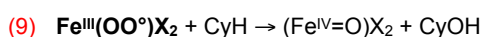
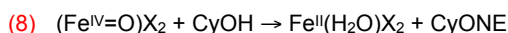
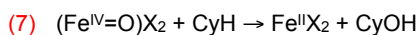
These reactions (in equilibria) should be favored under TBHP excess, since one TfOH molecule is generated upon addition of CyOH to $\text{Fe}^{\text{III}}(\text{OTf})_3$. However, the release of free TfOH proved to be thermodynamically unfavorable, as inferred by DFT. Hence, we propose instead the participation of a $\text{CyOH}_2.\text{OTf}$ adduct ($X = \text{OTf}$, $n = 0$, $\Delta G_1 = -11 \text{ kJ mol}^{-1}$).

The $\text{Fe}^{\text{III}}(\text{OO}^t\text{Bu})\text{X}_2$ complex may further evolve by oxidation of Fe(III) to Fe(IV) generating $^t\text{BuO}^\cdot$ radicals (Eq 3),^[36] matching the observation by spin trapping (*vide supra*). Homolytic cleavage of the C-O bond to form $^t\text{Bu}^\cdot$ (Eq 4-5) is energetically unfavorable ($X = \text{OTf}$, $n = 0$, $\Delta G_4 = +336 \text{ kJ mol}^{-1}$).^[31b] and this radical was not experimentally detected. Accordingly, the formation of superoxo radicals by this pathway is unlikely. As an

alternative, TBHP-assisted C-O cleavage could be driven by Fe^{III}(OO^tBu)X₂ complexes according to Eq 6^[31b]



The as-generated ferryl-oxo species [i.e. (Fe^{IV}=O)X₂] (Eq 3) should be able to activate the C-H bond in CyH (Eq 7) and CyOH (Eq 8).^[37] Besides, FeOO[°] could also oxidize CyH (Eq 9 and Eq 10).^[31b]



We were curious to figure out why Fe^{III}(OTf)₃ and Fe^{III}(NTf₂)₃ salts are highly efficient for CyH oxidation. Assuming that C-H activation by a Fe(IV)=O moiety is the rate-limiting step of the reaction, it may be interesting to question the very high activity of these salts. In a recent study, Sautet and coworkers reported that the activation energy for C-H activation and the energy level of the acceptor frontier orbital of the initial Fe(IV)=O intermediate (σ* and π* for quintuplet and triplet state complexes, respectively) are linearly correlated.^[39] Relying on this work, we computed the orbital energies for some plausible Fe(IV)=O intermediates featuring triflate ligands and for a porphyrin-ligated Fe(IV)=O as a reference (Table 6).

Irrespective of the ligand, the quintuplet state is favored for triflate-containing complexes, except in the case of CyO⁻, for which both the quintuplet and triplet states are very close in energy (<1 kJ mol⁻¹). In contrast, the triplet state is favored for the porphyrin-ligated Fe(IV)=O. Poorly ligated triflate Fe(IV)=O is the most electrophilic species with a very low σ* orbital at -5.4 eV. CyOH- and CyONE-ligated triflate Fe(IV)=O are predicted to be a few less active (-4.2 eV and -3.9 eV). This lower electrophilicity upon coordination may account for the rapid decrease of TON along the reaction (*vide infra*). Coordination of the CyO⁻ ligand is expected to strongly increase the σ* level (-3.5 eV), which is in line with a potential catalyst deactivation. Finally, porphyrin-ligated Fe(IV)=O exhibits the highest frontier orbitals (π*: -5.6 eV, σ* = -3.3 eV), in agreement with the lower KA yield of Cl-Fe^{III}TPP compared to that of Fe^{III}(OTf)₃ (Table 1). Thus, the remarkable activity of Fe^{III}(OTf)₃ for CyH oxidation may be ascribed to the higher electrophilicity of triflate-ligated Fe(IV)=O species, allowing a relatively low C-H activation barrier compared to other Fe-ligated catalysts.

Fe^{II}X₂ may be re-oxidized to Fe(III) or Fe(IV) by O₂ or TBHP according to Eq 11 and Eq 12

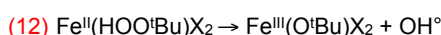
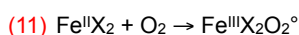


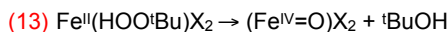
Table 6. Energy level of frontier orbitals (in eV) of S=1 and S=2 spin Fe(IV)=O complexes^a

Species	Spin state	σ* (eV), S = 2	π* (eV), S = 1
Fe(IV)O(OTf) ₂	2	-5.4	-9.1
Fe(IV)O(OTf) ₂ (CyOH) ₃	2	-4.2	-7.9
Fe(IV)O(OTf) ₂ (CyONE) ₃	2	-3.9	-7.3

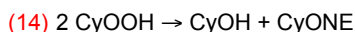
Fe(IV)O(CyO) ₂ (CyOH) ₂	2 or 1	-3.5	-6.6
Fe(IV)O(TPP)	1	-3.3	-5.6

^a Fe(IV) oxo complexes can be on singlet, triplet or quintuplet states. Both triplet and quintuplet states were optimized for each complex. The energy of the acceptor orbitals (relevant to estimate the activation energy of the H-atom transfer) are indicated in eV. The energy of the π* and σ* orbitals are reported for the triplet (S = 1) and quintuplet state (S = 2), respectively, in agreement with the conclusions in ref [38]

However, the above reactions seem unlikely, since no traces of OH[°] or Fe^{III}X₂O₂[°] were experimentally detected.^[39] On the contrary, Eq 13 would allow the regeneration of (Fe^{IV}=O)X₂



Finally, mechanisms involving the formation of CyOOH and its disproportionation to CyOH and CyONE are unlikely, since the A/K molar ratio evolves during the reaction, excluding a termination reaction according to Eq 14.^[5c]



Conclusion

Highly electrophilic high-valent iron triflate salts proved to be of high interest for the C-H activation of cyclohexane. These salts selectively and efficiently catalyze the oxidation of cyclohexane to cyclohexanol and cyclohexone (KA oil) with very high turnover frequencies. Combined EPR spectroscopy with DFT calculations helped us to propose structures intermediates, as well as to find out possible sources of catalyst deactivation.

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Keywords: Iron Triflate • Oxidation • KA oil • C-H activation • EPR • DFT • Cyclohexane • Homogeneous Catalysis

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