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Catalytic Oxidation with a Non-Heme Iron Complex That Generates a Low-Spin FeIIIOOH Intermediate

Gerard Roelfes,[a] Marcel Lubben,[a] Ronald Hage,[b] Lawrence Que, Jr.,*[c] and Ben L. Feringa*[a]

Abstract: The antitumor drug bleomycin (BLM) is proposed to act via a low-spin iron(III) hydroperoxide intermediate called “activated bleomycin”. To gain more insight into the mechanistic aspects of catalytic oxidation by these intermediates we have studied the reactivity of $\{N4Py\}Fe(CH,CN)\{(ClO4)2$ (I) $\{N4Py\}=N4,N-bis(2$-pyrldylmethyl)-N-bis(2-pyridyl)methylamine) with excess H2O2. Under these conditions a transient purple species is generated, $\{(N4Py)FeOOH\}^{2+}$ (2), which has spectroscopic features and reactivity strongly reminiscent of activated bleomycin. The catalytic oxidation of alkanes such as cyclohexane, cyclooctane, and adamantane by I with H2O2 gave the corresponding alcohols and ketones in up to 31% yield. It was concluded, from the O2 sensitivity of the oxidation reactions, the formation of brominated products in the presence of methylene bromide, and the nonstereospecificity of the oxidation of cis- or trans-dimethylycyclohexane, that long-lived alkyl radicals were formed during the oxidations. Oxidation of alkenes did not afford the corresponding epoxides in good yields but resulted instead in allylic oxidation products in the case of cyclohexene, and cleavage of the double bond in the case of styrene. Addition of hydroxyl radical traps, such as benzene and acetone, led to only partial quenching of the reactivity. The kinetic isotope effects for cyclohexanol formation, ranging from 1.5 in acetone to 2.7 in acetone with slow addition of H2O2, suggested the involvement of a more selective oxidizing species in addition to hydroxyl radicals. Monitoring the UV/Vis absorption of 2 during the catalytic reaction showed that 2 was the precursor for the active species. On the basis of these results it is proposed that 2 reacts through homolysis of the O–O bond to afford two reactive radical species: $\{(N4Py)Fe^{\cdot}O\}^{\cdot}$ and ‘OH. The comparable reactivity of I and Fe – BLM raises the possibility that they react through similar mechanistic pathways.

Keywords: homogeneous catalysis • iron • N ligands • O–O activation • oxidations

Introduction

Iron–peroxo species are invoked in the mechanisms of several iron-requiring biological oxidation catalysts,[1] Such intermediates have been observed for non-heme diiron enzymes such as methane monooxygenase,[2] ribonucleotide reductase,[3] and stearoyl acyl carrier protein Δ9-desaturase.[4] These peroxo intermediates serve as precursors for high-valent iron–oxy species that effect substrate oxidation.[5] In the case of cytochrome P450, an iron(III)–peroxo species is assumed to be formed when the key second electron is injected into oxy-P450 (FeIII–O2–). Due to the apparently low oxidative reactivity of FeIII–μ2–O2 species, it is proposed that this moiety, if formed in the P450 active site, must be protonated to form an FeIIIOOH species before its conversion to a high-valent iron–oxy species.[6]

A low-spin FeIIIOOH species has been characterized for “activated BLM”,[7] the active form of the antitumor drug bleomycin (BLM), which is a metalloglycopeptide.[8] Activated BLM is formed by the reaction of the FeIIb form, O2, and a one-electron reductant to form a metastable FeIIIOOH species, of which the formulation has been established by electrospays ionization mass spectrometry.[9] The decomposition of this intermediate is thought to be responsible for the drug’s ability to cleave DNA by an oxidative mechanism.[9] The accepted mechanism for bleomycin action[9, 10] involves hydrogen abstraction by activated BLM at the deoxyribose unit of a nucleotide to form a C4’ carbon radical whose fate is...
determined by two subsequent pathways, one that requires additional O₂ and another that does not. Besides DNA cleavage, activated BLM is also capable of the epoxidation of styrene,[10] the oxidation of stilbene to give the corresponding epoxide and other oxidation products such as benzoin and benzaldehyde,[11] the hydroxylation of naphthalene and anisole,[12] and the demethylation of N,N-dimethylaniline (DMA).[12]

How the low-spin FeIIIOOH moiety of activated BLM is involved in its oxidative reactions has been the subject of considerable debate. Three pathways can be considered. First, following the heme enzyme precedent, the iron hydroperoxide intermediate may undergo O–O bond heterolysis to give rise to a (formally) FeV=O species,[8] analogous to heme peroxidase (Scheme 1). This is supported by the observation of olefin epoxidation activity of FeII–BLM with oxygen atom donors such as iodosylbenzene,[12] but it is weakened by the fact that iodosylbenzene can also be activated by redox-inactive Lewis-acidic metal centers.[13] A second pathway is the homolytic scission of the O–O bond to give an FeV=O species and ‘OH, which would generate a highly reactive oxidant with low discrimination. Thirdly, the FeIIIOOH intermediate itself could be involved in substrate oxidation. This option has the advantage of involving an oxidant with more moderate reactivity and thus greater selectivity, and is considered to rationalize the reactivity of certain heme enzymes.[16]

There has been much effort to improve understanding of the chemistry of Fe–BLM through the use of model complexes. To date, the 2-(2,5-diazapentyl)-5-bromopyridine-6-carboxylic acid N-[2-(4'−imidazolyl)ethyl]amide (pma) ligand designed by Mascharak and co-workers and the N-[6-[[[(5'-2-amino-2-carboxamidophenyl)amino]methyl]pyridine-2-carboxyl]-1-histidine (pyml) ligand most closely reproduce the iron coordination environment of BLM. Indeed, like FeII–BLM, FeIII–pma and FeII–pmyl react with O₂ to give rise to intermediates with EPR parameters similar to those of activated BLM.[15] However, the FeIIIOOH formulation has not been established in these cases. More recently, a number of synthetic iron complexes have been found to react with H₂O₂ or alkyl hydroperoxides to give low-spin FeIIIIOOH[18] and FeIIIOR intermediates[19] analogous to activated BLM. Several of these complexes have been found to be good catalysts for hydrocarbon oxidation. From a detailed mechanistic study, it has been established that the [Fe(tpa)(H₂O)₂][ClO₄]₂/BuOOH system (tpa = tris(2-pyridylmethyl)amine) generates a low-spin FeIIIOOR intermediate which reacts by O–O bond homolysis giving FeIV=O and ‘OH.[20] The ‘OH radical reacts with the substrate to give a radical substrate which can then be trapped by the FeV=O moiety to give alcohol or undergo a radical chain autoxidation reaction with O₂ to generate equimolar amounts of alcohol and ketone. On the other hand, [Fe(tpa)(CH₃CN)₂][ClO₄]₂ and [Fe(bpmen)(CH₃CN)₂][ClO₄]₂ (bpmen = N,N-dimethyl-N,N-bis(2-pyridylmethyl)ethylene-1,2-diamine) are capable of stereospecific oxidation of alkanes and alkenes with H₂O₂, results that are inconsistent with free radical chemistry.[18d, 21]

We previously reported the synthesis and characterization of iron complexes of the ligand N,N-bis(2-pyridylmethyl)-N-bis(2-pyridylmethyl)amine (N₄Py).[18e, 22] Like BLM, N₄Py can act as a pentadentate ligand, leaving one available coordination site in the corresponding iron complex. We have demonstrated by X-ray crystallography that this site can be occupied by CH₃CN or Cl⁻ in FeIII complexes and by CH₂O– or O₂⁻ in FeIIII complexes. Reaction of [Fe(N₄Py)(CH₃CN)](ClO₄)₂ (I) with H₂O₂ resulted in the formation of a transparent purple species which has been characterized spectroscopically (UV/Vis, EPR, resonance Raman[23] and ESI-MS) as [FeIII(N₄Py)OOH]²⁺ (2) (Scheme 2). [18d]

$$\text{[N₄PyFe(CH₃CN)]}^{2+} + \text{H}_2\text{O}_2 \rightarrow \text{[N₄PyFeOOH]}^{2+}$$

Scheme 2. Reaction of the N₄Py−iron(II) complex I with H₂O₂ to generate the FeIIIOOH intermediate 2 and (inset) the ligand N₄Py.

Here we report on a detailed study of the reaction of the [Fe(N₄Py)(CH₃CN)](ClO₄)₂/H₂O₂ system in catalytic oxidations, giving a mechanistic interpretation of the results and discussing the relevance of these results to an understanding of the chemistry of Fe–BLM.

**Results and Discussion**

Catalytic oxidations with 1 were examined, focusing on: i) parameters that affect the oxidation pathway; and ii) the reactivity of the key peroxide complex 2. To determine whether 2 reacts by homolysis or heterolysis, catalytic oxidations in acetoni trile were investigated. Studies with hydroxyl radical traps such as acetone and benzene and mechanistic probes such as kinetic isotope effects, tertiary/secondary (3/2') ratios in adamantane oxidation, and stereo-selectivity in the oxidation of cis- and trans-1,2-dimethylcyclohexane led to a proposed mechanism entailing homolysis of the O–O bond of 2, affording two active oxidizing species: [(N₄Py)FeIV=O]²⁻ and ‘OH.

Scheme 3 summarizes the reactions catalyzed by 1. They include oxidation of alkanes, alkenes, alcohols, benzene, and N,N-dimethylaniline (DMA). All reactions were carried out under an argon atmosphere, unless noted otherwise, at 25 °C. The reaction was started by adding 100 equivalents of H₂O₂ to a solution containing the catalyst and 1000 equivalents of substrate. Acetonitrile and acetone were each used as solvent for the catalytic oxidation reactions; samples for GC analysis were taken after 30 min (acetone) or 90 min (acetonitrile). We
have shown previously that 2 is formed in either acetone or acetonitrile, although the intermediate is formed quantitatively only in acetone.\cite{22}

**Oxidations in acetonitrile:** Under the conditions mentioned above, the oxidation of alkanes gave considerable yields of the corresponding alcohols and ketones. In the case of cyclohexane, cyclohexanol and cyclohexanone were formed in 31% combined yield (based on hydrogen peroxide) in acetonitrile (Table 1, entry 1). The large yields observed make 1 among the most reactive and efficient non-heme iron oxidation catalysts. Cyclohexanol was the main product, with an alcohol/ketone (A/K) ratio of 1.4, when the reaction was carried out under Ar (Table 1, entry 1). A/K decreased to nearly 1 when the reaction was in air (Table 1, entry 2) and increased to 1.9 upon syringe-pump addition of H$_2$O$_2$ under argon (entry 3), suggesting the involvement of O$_2$ that propagates a radical chain autoxidation process.\cite{20}

Experiments were performed under a vigorous argon purge in an attempt to remove all traces of dioxygen and block the radical chain reaction. Since cyclohexane is too volatile for this purpose, cyclooctane was used as substrate. Under the standard reaction conditions in acetonitrile both cyclooctanol and cyclooctanone were produced (Table 2, entry 1), but in this case the ketone proved to be the main product (A/K = 0.3). A similar shift in alcohol/ketone selectivity on going from cyclohexane to cyclooctane as substrate was observed.

### Table 1. Catalytic oxidation of cyclohexane to cyclohexanol and cyclohexanone.

<table>
<thead>
<tr>
<th>Entry</th>
<th>Additive</th>
<th>Alcohol[$^\text{[d]}$]</th>
<th>Ketone[$^\text{[d]}$]</th>
<th>A/K</th>
<th>$k_{\text{al}}/k_{\text{ket}}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>acetonitrile</td>
<td>18.5</td>
<td>12.9</td>
<td>1.4</td>
<td>1.5</td>
</tr>
<tr>
<td>2</td>
<td>air</td>
<td>14.9</td>
<td>15.6</td>
<td>0.96</td>
<td>-</td>
</tr>
<tr>
<td>3</td>
<td>syringe pump (200 min)</td>
<td>17.5</td>
<td>9.1</td>
<td>1.9</td>
<td>-</td>
</tr>
<tr>
<td>4</td>
<td>CH$_4$Br$_2$ (2500 equiv)</td>
<td>13.4</td>
<td>9.3</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>5</td>
<td>benzene (10% v/v)</td>
<td>16.7</td>
<td>8.4</td>
<td>2.0</td>
<td>-</td>
</tr>
<tr>
<td>6</td>
<td>benzene (50% v/v)</td>
<td>10.5</td>
<td>4.0</td>
<td>2.6</td>
<td>1.7</td>
</tr>
<tr>
<td>7</td>
<td>acetone</td>
<td>16.9</td>
<td>6.6</td>
<td>2.6</td>
<td>2.3</td>
</tr>
<tr>
<td>8</td>
<td>air</td>
<td>13.4</td>
<td>14.9</td>
<td>0.9</td>
<td>-</td>
</tr>
<tr>
<td>9</td>
<td>syringe pump (70 min)</td>
<td>17.7</td>
<td>4.8</td>
<td>3.7</td>
<td>-</td>
</tr>
<tr>
<td>10</td>
<td>syringe pump (200 min)</td>
<td>12.4</td>
<td>2.4</td>
<td>5.2</td>
<td>2.7</td>
</tr>
<tr>
<td>11</td>
<td>CH$_4$Br$_2$ (2500 equiv)</td>
<td>11.4</td>
<td>5.3</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>12</td>
<td>[D$_3$]acetone</td>
<td>7.9 (RBr)</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

[a] Turnover number (TON) = mol product per mol catalyst.

### Table 2. Catalytic oxidation of other substrates.

<table>
<thead>
<tr>
<th>Entry</th>
<th>Substrate</th>
<th>Products</th>
<th>TON in CH$_3$CN</th>
<th>TON in acetone</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>1[$^\text{[c]}$]</td>
<td>cyclooctane</td>
<td>cyclooctanol</td>
<td>2.7</td>
<td>-</td>
<td>A/K = 0.3</td>
</tr>
<tr>
<td>2[$^\text{[b]}$]</td>
<td>cyclooctane</td>
<td>cyclooctanone</td>
<td>9.0</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>3</td>
<td>cis-1,2-dimethylcyclohexane</td>
<td>cis-1,2-dimethylcyclohexan</td>
<td>1.4</td>
<td>2.3</td>
<td>cis/trans 1.8 (CH$_3$CN)</td>
</tr>
<tr>
<td>4</td>
<td>trans-1,2-dimethylcyclohexane</td>
<td>trans-1,2-dimethylcyclohexan</td>
<td>1.1</td>
<td>1.7</td>
<td>cis/trans 1.4 (CH$_3$CN)</td>
</tr>
<tr>
<td>5</td>
<td>adamantane[$^\text{[c]}$]</td>
<td>1-adamantan</td>
<td>8.3</td>
<td>-</td>
<td>3/2* = 3.1</td>
</tr>
<tr>
<td>6[$^\text{[c]}$]</td>
<td>adamantane[$^\text{[c]}$]</td>
<td>1-adamantan</td>
<td>8.9</td>
<td>-</td>
<td>3/2* = 3.3</td>
</tr>
<tr>
<td>7</td>
<td>cyclohexene</td>
<td>cyclohexen</td>
<td>27.8</td>
<td>23.1</td>
<td>-</td>
</tr>
<tr>
<td>8</td>
<td>cyclooctane</td>
<td>many products</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>9</td>
<td>styrene</td>
<td>benzaldehyde</td>
<td>21.3</td>
<td>25.6</td>
<td>-</td>
</tr>
<tr>
<td>10</td>
<td>benzene</td>
<td>phenol</td>
<td>16.6</td>
<td>2.4(3.4)[$^\text{[c]}$]</td>
<td>-</td>
</tr>
<tr>
<td>11</td>
<td>cyclohexan</td>
<td>cyclohexanen</td>
<td>13.6</td>
<td>10.6</td>
<td>-</td>
</tr>
<tr>
<td>12</td>
<td>benzyl alcohol</td>
<td>benzaldehyde</td>
<td>54.9</td>
<td>64.4</td>
<td>-</td>
</tr>
<tr>
<td>13</td>
<td>N,N-dimethylaniline</td>
<td>N-methyl aniline</td>
<td>-</td>
<td>15.3</td>
<td>-</td>
</tr>
</tbody>
</table>

[a] 50 equivalents of H$_2$O$_2$ used. [b] Ar purge. [c] 100 equivalents of substrate were suspended in the solvent. [d] Under air. [e] In [D$_3$]acetone.
with the \([\text{Fe}_2\text{O}(\text{bpy})_3(\text{H}_2\text{O})_2]\) \((\text{ClO}_4)^-/\text{H}_2\text{O}\) system (bpy = 2,2'-bipyridine).\(^{34}\) Under argon purge, the yields of both alcohol and ketone were strongly decreased (Table 2, entry 2), proving that trace amounts of \(O_2\) play an important role during the reaction.

The behavior of \(1/\text{H}_2\text{O}_2\) towards alkane substrates can be compared with several groups of non-heme iron catalysts: a) the Gy family of catalysts, which afford mainly ketone products,\(^{25,26}\) b) catalysts with \(A/K \approx 1\) such as \([\text{Fe}_2\text{O}(`\text{OAc})_2(\text{bpy})_2]\text{Cl}_2\), \([\text{Fe}_2\text{O}(\text{OAc})(\text{tmia})_2]\) \((\text{ClO}_4)_2\) (tmia = tris(1-methylimidazol-2-yl)methylamine),\(^{20}\) \([\text{Fe}(`\text{pm}a)](\text{ClO}_4)\),\(^{27}\) and \([\text{Fe}_2\text{O}(\text{bpy})(\text{H}_2\text{O})_2]\) \((\text{ClO}_4)_2\) and c) catalysts with large \(A/K\) ratios such as \([\text{Fe}(`\text{bpm}en)(\text{CH}_3\text{CN})]_2(\text{ClO}_4)_2\) (A/K = 6.3)\(^{21}\) and \([\text{Fe}(`\text{tpa})(\text{CH}_3\text{CN})]_2(\text{ClO}_4)_2\) (A/K = 4.3).\(^{18}\) Thus the A/K ratio found for \(1/\text{H}_2\text{O}_2\) corresponds most closely to those associated with the catalysts in group b. An A/K ratio of approximately 1 is symptomatic of the presence of free alkyl radical intermediates, which react rapidly with \(O_2\) to initiate a radical chain autoxidation.\(^{28,29}\) In support, significant amounts of cyclohexyl bromide were formed in the presence of excess methylene bromide (Table 1, entries 4 and 11), which serves as an excellent trap for free alkyl radicals.\(^{30}\) These radicals can then be trapped by dioxygen to form alkylperoxy radicals that afford equimolar amounts of ketone and alcohol in a Russell termination reaction\(^{28,29}\) or by Fe\(^{4+}\)O species to give alcohols.\(^{28a}\)

The behavior of \(1/\text{H}_2\text{O}_2\) with olefins supports mechanistic conclusions derived from the alkane oxidation experiments. Styrene was converted to styrene oxide and benzaldehyde (Table 2, entry 9), whereas cyclohexene oxidation afforded the corresponding allylic alcohol and ketone, and very little epoxide (Table 2, entry 7). With cyclooctene, which is less susceptible to allylic oxidation,\(^{31}\) many different oxidation products were found, each accounting for less than one turnover (Table 2, entry 8). These observations contrast with those for \([\text{Fe}(`\text{tpa})(\text{CH}_3\text{CN})]_2(\text{ClO}_4)_2\) and \([\text{Fe}(\text{cyclam})]_2(\text{CF}_3\text{SO}_3)_2\) (cyclam = 1,4,8,11-tetraazaacyclotetradecane),\(^{30}\) non-heme iron complexes which catalyze the stereospecific epoxidation of olefins with \(\text{H}_2\text{O}_2\) as oxidant. The observations that the \([\text{Fe}(\text{N}^4\text{Py})(\text{CH}_3\text{CN})]_2(\text{ClO}_4)_2/\text{H}_2\text{O}_2\) oxidation of cyclohexene gives mainly allylic oxidation products and that the oxidation of styrene gives benzaldehyde as the major product point to the involvement of a radical oxidant. The formation of small amounts of styrene oxide might implicate a two-electron oxidant, but the absence of significant amounts of epoxide in the oxidation of cyclohexene and cyclooctene suggests that the styrene oxide observed is more likely to be the result of a radical addition to the double bond followed by ring closure, as proposed for the epoxidation of styrene by \([\text{tmp}]\text{Fe}^{4+}\text{O}(`\text{tmp} = \text{tetrasmethylporphyrin dianion})\).\(^{31}\)

The \(1/\text{H}_2\text{O}_2\) system also catalyzed efficient oxidation of cyclohexanol to cyclohexanone and benzal alcohol to benzaldehyde (Table 2, entries 11 and 12) as well as the \(N\)-demethylation of \(N,N\)-dimethylamine (Table 2, entry 13). However, since such oxidations are not unique for a particular oxidizing agent, little mechanistic insight can be drawn from these results.

**Effect of hydroxyl radical traps on alkane oxidation:** The alkane hydroxylation mechanism was further elucidated with benzene and acetonitrile as hydroxyl radical traps. The use of benzene as a substrate under standard conditions resulted in the formation of phenol (Table 2, entry 10), and no diphenyl was detected. Although this oxidation may occur with a metal-based oxidant, the pronounced effect of the solvent strongly suggests the attack of OH radicals.\(^{32}\) In acetonitrile, 17 turnovers of benzene to phenol were found, but in acetone only 2.6 turnovers were obtained, a decrease in activity of more than sixfold. These observations may be rationalized by the fact that acetonitrile is a good trap for OH radicals as well,\(^{33}\) thus reducing the turnover number (TON) of benzene to phenol in that solvent. Since a \(k_{\text{B}}/k_{\text{A}}\) ratio of approximately 2 has been reported for hydrogen abstraction from acetone by OH,\(^{33}\) the use of \([\text{D}_3]\) acetone as solvent should lead to increased yields of oxidized products; indeed the turnover number for phenol increased to 3.4. The presence of benzene during cyclohexene oxidation decreased the yields of the oxidation products. With 10% (v/v) benzene, the turnover numbers for cyclohexanol and cyclohexanone were 16.7 and 8.4, respectively (Table 1, entry 5); they decreased further, to 10.5 and 4.0, respectively, in 50% (v/v) benzene (Table 1, entry 6), presumably because of the trapping of hydroxyl radicals. Furthermore, the A/K ratios observed increased steadily from 1.4 in pure acetonitrile to 2.0 in 10% (v/v) benzene and 2.6 in 50% (v/v) benzene. These results strongly suggest the involvement of other oxidation mechanisms.

Alkane hydroxylation experiments in acetone as solvent support the possible participation of other mechanisms of cyclohexene oxidation. The oxidation of cyclohexane in acetone afforded 16 turnovers of alcohol and 6 of ketone (Table 1, entry 7), a 35% decrease in turnover number relative to that in acetonitrile, consistent with the trapping of hydroxyl radicals. In support, the turnover numbers in \([\text{D}_3]\) acetone increased to 20.0 and 9.4 for cyclohexanol and cyclohexanone (Table 1, entry 12), respectively; these are comparable with those found in acetonitrile. As in the case of benzene, the A/K ratio increased from 1.4 in acetonitrile to 2.1 in \([\text{D}_3]\) acetone and 2.6 in acetone, but the latter value decreased to 0.9 for the reaction in air (Table 1, entry 8), demonstrating that free alkyl radicals were formed in the reaction and trapped by \(O_2\). However the A/K ratios increased to 3.7 and 5.2 when the \(\text{H}_2\text{O}_2\) was introduced by syringe pump over 70 and 200 min periods, respectively (Table 1, entries 9 and 10). These results strongly suggest that OH radicals do play a role in the oxidation reaction catalyzed by 1, particularly in acetonitrile. However, trapping of OH radicals by acetone or benzene leads to only partial quenching of cyclohexene oxidation. In the presence of these traps, the A/K ratio can jump from 1.3 to as much as 5.2, suggesting that the radical chain autoxidation pathway can be suppressed. These observations raise the possibility that a second oxidizing species may be involved.

**Mechanistic probes of alkane hydroxylation:** The \(k_{\text{B}}/k_{\text{A}}\) ratios for the formation of cyclohexanol were determined in competition experiments between cyclohexane and \([\text{D}_3]\) cyclohexane. In acetonitrile the value of 1.5 (Table 1, entry 1) approached that associated with hydroxyl radicals.\(^{30}\) As the reaction conditions were modified to trap hydroxyl radicals,
the $k_{ij}/k_0$ ratio rose (Table 1, entries 6, 7, and 12), and it could be further increased to 2.7 by syringe-pump addition of $\text{H}_2\text{O}_2$ in acetone over a 200 min period (Table 1, entry 10). Values of 1–2, generally associated with radical chain autoxidations,[20] have been reported for many systems, for example, $\text{Fe(Clo}_3\text{)}_2$·$\text{H}_2\text{O}$ (1.5),[20] $\text{Fe}_3\text{O}_4$($\text{OAc}$)$_2$(bpy)$_2$Cl$_2$ (1.4),[20] and $\text{Fe}_3\text{O}_4$(bpy)$_2$(H$_2$O)$_2$][ClO$_4$] (2.1).[20] Larger $k_{ij}/k_0$ ratios have been found for hydrogen abstraction by sterosepecific alkane hydroxylation catalysts such as $\text{[Fe(tpa)(CH}_3\text{CN})]_2}$[ClO$_4$]$_2$ (3.5)[24] and iron porphyrins (10–24).[33] The increase in $k_{ij}/k_0$ under conditions that diminish the effect of hydroxyl radicals supports the deduction from the radical trap experiments that an oxidant more selective than the hydroxyl radical is also involved in the oxidation.

With adamantane as substrate, oxidation occurred at both secondary and tertiary carbon centers (Table 2, entries 5 and 6), but there was selectivity for oxidation at the tertiary position (3/2: 3.1–3.3, normalized on a per-hydrogen basis). For comparison: 3/2 ratios of 2.7, on average, have been found for Gif-type oxidations,[33] and the selectivity for the oxidation of adamantane by $\cdot OH$,[36] 3.5 for oxidation with $\text{Fe}_3\text{O}_4$($\text{OAc}$)$_2$·(bpy)$_2$Cl$_2$H$_2$O;[20] 9.5–10 for oxidation with $\text{Fe}_3\text{O}_4$(bpy)$_2$·(H$_2$O)$_2$][ClO$_4$] or $\text{BuOOH}$ or $\text{FeCl}_3$(tpa)Cl][ClO$_4$]($\text{BuOOH}$,[36] and 11–48 for oxidations with PhIO catalyzed by P450 mimics.[39] Thus the oxidant involved in $\text{[Fe(N4Py)(CH}_3\text{CN})]_2}$[ClO$_4$]$_2$/H$_2$O is not as reactive as hydroxyl radicals, but only slightly less so.

The stereoselectivity of the alkane hydroxylation reaction was examined with cis- and trans-1,2-dimethylcyclohexane as substrates (Table 2, entries 3 and 4). Both isomeric 1,2-dimethylcyclohexanols were formed, with cis/trans ratios of 1.4–1.9. These ratios were in sharp contrast to the stereospecificity found for $\text{[Fe(tpa)(CH}_3\text{CN})_2]}$[ClO$_4$]$_2$/H$_2$O;[10] $\text{[Fe(bpmien)(CH}_3\text{CN})_2]}$[ClO$_4$]$_2$/H$_2$O, and cytochrome P450 models,[89] but were more in the range of the cis/trans ratios found for catalytic autoxidation reactions (1.1–1.3).[41] Again, this indicates the formation of alkyl radicals with a lifetime sufficient to allow epimerization at the radical site.

Nature of the key oxidizing species: The fact that 1 reacts with $\text{H}_2\text{O}_2$ to form the $\text{Fe}^{II}$OOH intermediate 2 raises the question of its involvement in the oxidation reaction, either as the oxidant or as its precursor by cleavage of the O–O bond. The 530 nm absorbance of 2 in acetone was monitored concomitantly with the oxidation of cyclohexane to cyclohexanol and cyclohexanone. Figure 1 shows that the reaction is essentially complete after 15 min, coincident with the disappearance of the characteristic visible absorption of the intermediate, and demonstrates that the intermediate is indeed involved in the catalytic oxidation. However, the lifetime of the intermediate was not affected when $\text{[D}_2\text{]}$cyclohexane was used as substrate, despite a kinetic isotope effect (KIE) of 2.3 for cyclohexane oxidation. It thus appears likely that the $\text{Fe}^{II}$OOH intermediate itself is not the active oxidant, but serves as the precursor for the active species.

Decay of the $\text{Fe}^{III}$OOH intermediate to form the active oxidant can occur through O–O bond homolysis or heterolysis. From the results described above it is clear that the reactivity of the $\text{I}/\text{H}_2\text{O}_2$ system can be explained only in terms of one-electron oxidation and not by the involvement of a two-electron oxidant such as the (formally) Fe$^0$ species proposed for cytochrome P450. The arguments for a radical-type oxidation are:

i) The oxygen sensitivity of the oxidation of cyclohexane and cyclooctane and the formation of cyclohexyl bromide in the presence of methylene bromide reveals the involvement of free alkyl radicals, which is further supported by the lack of stereoselectivity in the oxidation of dimethyl-cyclohexane. This indicates hydrogen abstraction by a radical species and not by an Fe$^0$ species followed by oxygen rebound as proposed for cytochrome P450 and P450 model compounds.[6–13] Furthermore, addition of radical scavengers resulted in partial quenching of the reactivity towards cyclohexane.

ii) The KIE values of approximately 2 are in the range for radical-type oxidations. Oxidation by two-electron oxidants typically results in much higher KIE values.

iii) The observed C3/C2 ratio of approximately 3 in adamantane oxidation is comparable with that found for radical-type oxidations, whereas cytochrome P450 mimics give higher selectivity for tertiary positions.

iv) Complex 1 is unable to catalyze epoxidation, a typical two-electron oxidation process. Although the oxidation of styrene yielded the oxide in significant amounts, this result can also be explained in terms of radical addition to the double bond followed by ring closure. Furthermore, the formation of large amounts of benzaldehyde during styrene oxidation indicates a radical-type oxidation. Hence it can be concluded that the $\text{Fe}^{II}$OOH intermediate does not react through heterolysis of the O–O bond to give a (formally) Fe$^0$ species, which would be a two-electron oxidant. All the evidence points to one-electron oxidants and strongly suggests that 2 reacts by homolysis to give two radical species: $\text{[N4Py][Fe}^{III}$O$]$ and $\cdot OH$ (Scheme 4).

The involvement of $\cdot OH$ radicals is evident from the results with acetonitrile as solvent: a low A/K ratio in the oxidation of cyclohexane, a low KIE for cyclohexanol formation and significant hydroxylation of benzene to phenol. Furthermore, the turnover numbers for the last of these reactions decreases drastically in the presence of a radical trap such as acetone. These observations lead to the conclusion that at least part of the observed reactivity results from free radical chemistry initiated by hydroxyl radicals.
Whether the [(N4Py)FeIV(O)]3+ species itself is (re)active is more difficult to solve. In heme chemistry oxidative transformations by the FeIV(O) moiety, such as the oxygenation of triphenylphosphine to triphenylphosphine oxide,[42] and the nonstereospecific epoxidation of olefins,[31, 43] have been reported. Recently, the intramolecular hydroxylation of an aromatic ring has been observed for a non-heme FeIV(O) moiety also.[44] No examples of hydrogen abstraction from saturated alkanes by FeIV(O) moieties have been reported, to our knowledge. However, an important difference between heme systems and 1 is that the porphyrin ligands have a double negative charge and N4Py is a neutral ligand. Therefore, a more electron-negative FeIV(O) species will be formed in the homolysis of 2, making hydrogen abstraction from saturated alkanes more likely. Several observations support the idea that [(N4Py)FeIV(O)]3+ may be involved in oxidations catalyzed by 1. First, trapping of ‘OH radicals by added radical traps such as acetone leads to only a minor reduction in turnover numbers, strongly suggesting that the ‘OH radical is not the sole species responsible for oxidation. Indeed, the only catalytic reaction that is significantly affected by the use of acetone as solvent is the hydroxylation of benzene, which is typically carried out by ‘OH radicals. Furthermore, the use of ‘OH radical traps leads to an increase in the A/K ratio in cyclohexane oxidation, a result inconsistent with radical chain autoxidation chemistry. Most convincing, however, are the KIE values obtained for cyclohexane oxidation. The low value in acetonitrile indicates large contribution of ‘OH radicals. Trapping of ‘OH radicals by benzene or acetone leads to an increase in k_OH/k_H2O to 2.3. With slow addition of H2O2, to decrease the effect of radical chain autoxidations, k_OH/k_H2O increases further to 2.7, which is significantly beyond the range normally observed for ‘OH oxidations. This strongly suggests the involvement of a more selective oxidizing species, for example metal-based. On the basis of these observations we propose that the iron hydroperoxide intermediate 2 reacts through homolysis of the O-O bond and that the resulting species, [(N4Py)FeIV(O)]3+ and ‘OH, are both capable of effecting hydrogen bond abstraction of organic substrates.

Comparison with other systems: Comparison of the chemistry of the low-spin FeIII(OOH) intermediates involved in the present system with that of those in oxidations catalyzed by [Fe(tpa)(CH3CN)]2(CIO4)2 and Fe–BLM reveals marked differences, which appear to be related to the number of coordinating N atoms in the ligand. The systems (1 and Fe–BLM) with pentadentate ligands show similar reactivity to a number of substrates. In both systems epoxidation of styrene is accompanied by formation of benzaldehyde, hydroxylation of aromatic compounds, and demethylation of DMA. Furthermore, the observation that 1/H2O2 reacts with alkanes to form long-lived alkyl radicals which react mainly with O2 to give the products corresponds with the proposed mechanism for DNA degradation by Fe–BLM,[9] according to which hydrogen abstraction occurs at the C4’ position of the deoxyribose ring to generate a long-lived alkyl radical that can trap dioxygen to form a peroxy radical intermediate that degrades further to cause DNA strand cleavage (the oxygen-dependent route). These observations suggest that Fe–BLM and 1 react through similar mechanistic pathways, through homolysis of the O–O bond of the FeIII(OOH) intermediate. The alternative mechanism, which involves a (formally) FeO species derived from O–O bond heterolysis, has been considered for Fe–BLM, mainly on the basis of the observed reactivity of Fe–BLM with iodosylobenzene. As discussed in the literature,[13] the interpretation of these results is complicated by the fact that iodosylobenzene can also be activated by redox-inactive Lewis acids, so the involvement of an FeO moiety is not established. Therefore our analysis of the cumulative data presented here on the reactivity of 2, an intermediate strongly reminiscent of activated bleomycin, provides strong support for a mechanism for Fe–BLM involving homolytic scission of the O–O bond in activated bleomycin, giving [BLM–FeIV(O)] and ‘OH.

The oxidation chemistry of the [Fe(tpa)(CH3CN)]2(CIO4)2 system[10] differs from those observed for 1 and Fe–BLM, even though an FeIV(OOH) intermediate very similar to 2 and activated BLM is observed.[7] The tpa-based catalysts are capable of stereospecific epoxidation of olefins and hydroxylation of alkanes with H2O2. This could suggest heterolysis of the O–O bond of the [Fe(tpa)OOH]3+ (3) intermediate to give (formally) [(tpa)FeO]3+. Evidence in support of this hypothesis was provided recently for the Fe[(bpmen)-(CH3CN)]2(CIO4)2 system.[21] Incorporation of 18O in the product in the presence of H218O suggested the involvement of an oxidant that could undergo solvent exchange.

Since resonance Raman spectroscopy showed that the O–O bonds in 2 and 3 were of comparable strength (v(O–O) 790 and 789 cm−1, respectively)[28] it is difficult to imagine why 2 would react through homolysis of the O–O bond and 3 through heterolysis. Therefore there appears to be a more important factor controlling the decomposition pathway of the FeIII(OOH) intermediate.

The explanation may be that tpa is a tetradentate ligand, leaving two “open” coordination sites in the iron complex, in contrast to the pentadentate N4Py (and BLM). Whereas 2 reacts through homolysis of the O–O bond of the η1-coordinated hydroperoxide as described above, [Fe(tpa)(η1-OOH)]3+ (3) may react through a transition state in which the hydroperoxide is bound in an η2 fashion (Figure 2), analogously to proposed mechanisms for other peroxide-utilizing transition metal catalysts such as the Sharpless epoxidation catalyst[40] and MeReO3.[46] The FeIII–(η2–OOH) complex could then react through heterolysis of the O–O bond to give a (formally) FeO species that elicits stereospecific oxidation of a substrate. Whether this difference in peroxide coordina-
Conclusion

We have shown that the N4Py-Fe system is one of the most reactive non-heme iron catalysts known to date, capable of oxidizing a wide range of organic substrates including alkanes, alkenes, alcohols, benzene, and DMA. Complex 1 reacts with H2O2 to give the well characterized [(N4Py)Fe(II)OOH]2− intermediate 2, with properties strongly reminiscent of activated BLM. Intermediate 2 reacts through homolysis of the O−O bond, affording two species: [(N4Py)Fe(IV)O]2+ and OH. Although OH radicals are involved in substrate oxidation, the results obtained cannot be explained solely by the action of OH radicals. Therefore we propose that the Fe(IV)O species also plays a prominent role in oxidation of the substrate. The involvement of this species and the nature of the oxidizing complex need to be elucidated further. The formation of a low-spin iron(III) intermediate by both 1 and Fe–BLM, and their similar oxidation chemistry, lead to the attractive hypothesis that 1 and Fe–BLM react through the same mechanistic pathways. In view of the results described above, the chemistry of Fe–BLM should be considered in terms of homolysis of the O−O bond in activated bleomycins.

Experimental Section

Instrumentation and materials: UV/Vis spectra were recorded on a Hewlett Packard 8453 UV–Visible Spectrophotometer. GC analyses were performed on a Hewlett Packard 6890 Gas Chromatograph using an HP-1 dimethyl polysiloxane column, an HP-5 5% phenyl methyl siloxane column, or a CP-wax 52 CB column. Retention times of oxidation products were compared with commercial or independently prepared samples. Complex 1 was prepared according to published procedures.14a, 47 Caution: Perchlorate salts are potentially explosive and should be handled with care.

Catalytic oxidations: All experiments were carried out under argon, unless noted otherwise, in a water bath thermostatted at 25 °C.

In a typical procedure, cyclohexane (0.38 mL, 1000 equiv) was added to a solution of 1 (8.75 × 10−3 mol, 4 mL) and a known amount of bromobenzene (internal standard) in acetone. The reaction was started by addition of 30% H2O2 (35 μL). After 30 min an aliquot (1 mL) was taken from the reaction and filtered over a small silica column. The silica was washed thoroughly with diethyl ether or diethyl ether/10% methanol. The sample was concentrated to 2 mL by passing an argon stream over the solution, then analyzed by GC.

Kinetic isotope effect determination: In essentially the procedure described above, a cyclohexane/[D2]2-cyclohexane (1:1) mixture was used. The KIE was determined by comparing the turnover numbers for cyclohexanol and [D2]2-cyclohexanol (determined by GC with the CP-wax 52 CB column) and corrected for the relative concentrations of cyclohexane and [D2]2-cyclohexane.

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