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# Comparison of High-Spin and Low-Spin Nonheme Fe<sup>III</sup>–OOH Complexes in O–O Bond Homolysis and H-Atom Abstraction Reactivities

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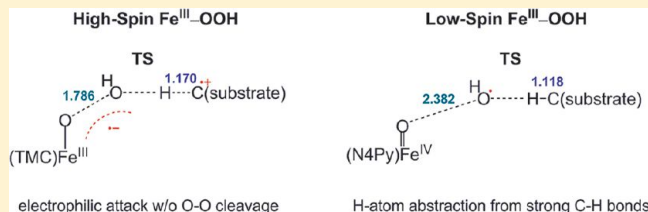
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## S Supporting Information

**ABSTRACT:** The geometric and electronic structures and reactivity of an  $S = 5/2$  (HS) mononuclear nonheme (TMC)Fe<sup>III</sup>–OOH complex are studied by spectroscopies, calculations, and kinetics and compared with the results of previous studies of  $S = 1/2$  (LS) Fe<sup>III</sup>–OOH complexes to understand parallels and differences in mechanisms of O–O bond homolysis and electrophilic H-atom abstraction reactions. The homolysis reaction of the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex is found to involve axial ligand coordination and a crossing to the LS surface for O–O bond homolysis. Both HS and LS Fe<sup>III</sup>–OOH complexes are found to perform direct H-atom abstraction reactions but with very different reaction coordinates. For the LS Fe<sup>III</sup>–OOH, the transition state is late in O–O and early in C–H coordinates. However, for the HS Fe<sup>III</sup>–OOH, the transition state is early in O–O and further along in the C–H coordinate. In addition, there is a significant amount of electron transfer from the substrate to the HS Fe<sup>III</sup>–OOH at transition state, but that does not occur in the LS transition state. Thus, in contrast to the behavior of LS Fe<sup>III</sup>–OOH, the H-atom abstraction reactivity of HS Fe<sup>III</sup>–OOH is found to be highly dependent on both the ionization potential and the C–H bond strength of the substrate. LS Fe<sup>III</sup>–OOH is found to be more effective in H-atom abstraction for strong C–H bonds, while the higher reduction potential of HS Fe<sup>III</sup>–OOH allows it to be active in electrophilic reactions without the requirement of O–O bond cleavage. This is relevant to the Rieske dioxygenases, which are proposed to use a HS Fe<sup>III</sup>–OOH to catalyze *cis*-dihydroxylation of a wide range of aromatic compounds.



## 1. INTRODUCTION

Mononuclear nonheme iron enzymes catalyze a wide range of chemical reactions that are important in medical, pharmaceutical, and environmental applications.<sup>1–5</sup> Within this broad class, bleomycin, extradiol dioxygenases, pterin-dependent hydroxylases,  $\alpha$ -KG-dependent dioxygenases, and Rieske dioxygenases utilize an Fe<sup>II</sup> site to activate O<sub>2</sub> for attack on substrates. Among these enzymes, Fe<sup>III</sup>–hydroperoxo species are observed and proposed to be key intermediates in the catalytic cycles of bleomycin and Rieske dioxygenases.<sup>6–11</sup>

Bleomycins are a family of glycopeptide antibiotics that exhibit high intrinsic anticancer cytotoxicity, which is due to their ability to effect single- and double-strand cleavage of DNA.<sup>12–16</sup> DNA strand scission is initiated by abstraction of the C-4' hydrogen atom from the backbone deoxyribose sugar by an Fe<sup>III</sup>–hydroperoxo intermediate termed activated bleomycin (ABLM).<sup>17,18</sup> From nuclear resonance vibrational spectroscopy (NRVS), ABLM is a low-spin (LS)  $S = 1/2$  Fe<sup>III</sup> species with an end-on hydroperoxo ligand.<sup>8</sup> Recent real-time

kinetic studies using circular dichroism determined that DNA accelerates the decay of ABLM, supporting a mechanism that involves direct H-atom abstraction by ABLM.<sup>6</sup> DFT calculations of the reaction of ABLM with DNA show that direct H-atom abstraction by LS Fe<sup>III</sup>–OOH is thermodynamically and kinetically favored over other proposed reaction pathways (e.g., heterolytic O–O cleavage to form a P450 compound I-like species).<sup>7,8</sup>

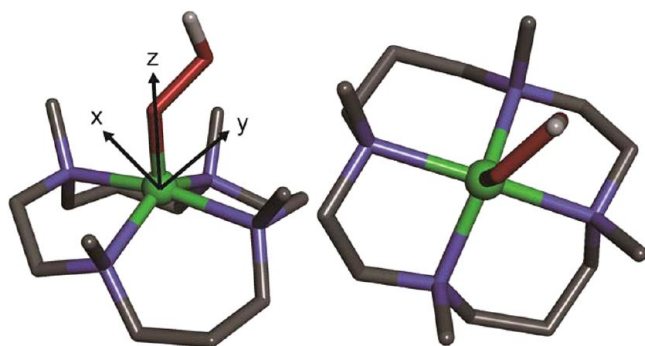
The Rieske dioxygenases catalyze stereo- and regiospecific electrophilic *cis*-dihydroxylation of aromatic compounds as the first step in the metabolism of aromatic compounds by bacteria.<sup>19</sup> Crystal structures of two Fe<sup>III</sup>–(hydro)peroxo intermediates have been reported.<sup>9,11</sup> A third one has been characterized by EPR and Mössbauer spectroscopy coupled with DFT calculations that favor a Fe<sup>III</sup>–hydroperoxo species.<sup>10</sup> The open question is whether this HS Fe<sup>III</sup>–hydroperoxo

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intermediate performs *cis*-dihydroxylation directly or whether the O–O bond first cleaves to generate a high-valent  $\text{HO}^\bullet\text{--Fe}^{\text{IV}}\text{=O}$  or  $\text{HO--Fe}^{\text{V}}\text{=O}$  species that performs the *cis*-dihydroxylation.<sup>10,20–22</sup>

Recently, we trapped a HS  $S = 5/2$   $\text{Fe}^{\text{III}}$ –hydroperoxo species  $[(\text{TMC})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  (TMC = 1,4,8,11-tetramethyl-1,4,8,11-tetraazacyclotetradecane).<sup>23</sup> Combination of resonance Raman, Fe K-edge XAS, and DFT calculations defined this to be a five-coordinate species with the hydroperoxo ligand syn to the methyl groups of the TMC ligand (see Figure 1).<sup>23</sup> In the



**Figure 1.** DFT-optimized structure of the  $[(\text{TMC})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  complex (left, side view; right, top view). Fe atom is in green, N atoms are in blue, C atoms are in black, O atoms are in red, H atom is in white. Only the H atom on the OOH ligand is shown, whereas other H atoms are omitted for clarity. Coordinate system is shown in the left panel, where *z* axis is defined as along the Fe–O bond, *y* axis and *x* axis are in and perpendicular to the Fe–O–O plane, respectively.

present study, we performed low-temperature Abs, magnetic circular dichroism (MCD), variable-temperature variable-field MCD (VTVH MCD), and resonance Raman profiling experiments to understand the electronic structure of this complex and calibrate electronic structure calculations. Second, this HS  $[(\text{TMC})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  species was found to undergo O–O bond homolysis, which however appears inconsistent with the results that O–O bond homolysis of a HS  $\text{Fe}^{\text{III}}\text{--OOH}$  system has an extra  $\sim 10$  kcal/mol barrier relative to LS  $\text{Fe}^{\text{III}}\text{--OOH}$  systems which do undergo O–O bond homolysis.<sup>24</sup> We therefore conducted temperature-dependent kinetic studies of O–O bond homolysis of the HS  $[(\text{TMC})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  complex and obtained analogous data of a well-defined LS  $\text{Fe}^{\text{III}}\text{--hydroperoxo}$  species,<sup>25,26</sup>  $[(\text{N4Py})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  (N4Py = *N,N*-bis(2-pyridylmethyl)-*N*-bis(2-pyridyl)methylamine) for quantitative comparison. These data were coupled with the experimentally calibrated DFT calculations to elucidate the O–O bond homolysis mechanism of the HS  $[(\text{TMC})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  complex and its relation to that of the LS  $\text{Fe}^{\text{III}}\text{--OOH}$  species. Finally, the HS  $[(\text{TMC})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  species was shown to abstract H atoms from substrates with weak C–H bonds, such as xanthene (75.5 kcal/mol) and 9,10-dihydroanthracene (77 kcal/mol).<sup>23</sup> Quantitative comparison of the direct H-atom abstraction reactivities of the HS and LS  $\text{Fe}^{\text{III}}\text{--OOH}$  species was therefore performed and correlated to the calibrated DFT calculations to elucidate their different reactivities in H-atom abstraction reactions and their dependence on the properties of substrates. These studies provide new insights into the electrophilic reactivities of LS and HS  $\text{Fe}^{\text{III}}\text{--OOH}$  species relevant to bleomycin and Rieske dioxygenases.

## 2. MATERIALS AND METHODS

**2.1. Synthesis of Materials.** All chemicals were obtained from Aldrich Chemical Co. as the highest purity available and used without further purification unless otherwise indicated. Solvents were dried and distilled under argon prior to use.<sup>27</sup> TMC ligand was purchased from Aldrich Chemical Co. Deuterated substrate, xanthene-*d*<sub>2</sub>, was prepared by taking xanthene (0.16 g, 0.87 mmol) in DMSO-*d*<sub>6</sub> (5 mL) along with NaH (0.1 g, 4.2 mmol) under an inert atmosphere.<sup>28</sup> The reaction solution was stirred for 8 h and quenched with D<sub>2</sub>O (5 mL). Crude product was filtered and washed with copious amounts of H<sub>2</sub>O. <sup>1</sup>H NMR confirmed >99% deuterated. The nonheme ferrous complexes,  $[\text{Fe}^{\text{II}}(\text{TMC})(\text{CH}_3\text{CN})_2](\text{CF}_3\text{SO}_3)_2$  and  $[\text{Fe}^{\text{II}}(\text{N4Py})(\text{CH}_3\text{CN})](\text{ClO}_4)_2$ , were prepared according to literature methods.<sup>24,29–32</sup> The HS nonheme ferric complex,  $[(\text{TMC})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$ , was prepared by following the literature method.<sup>21</sup> The LS nonheme ferric complex,  $[(\text{N4Py})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  (536 nm,  $\epsilon = 1000 \text{ M}^{-1} \text{ cm}^{-1}$ ), was generated by adding H<sub>2</sub>O<sub>2</sub> (25 mM) into a solution containing  $[\text{Fe}^{\text{II}}(\text{N4Py})(\text{CH}_3\text{CN})](\text{ClO}_4)_2$  (0.5 mM) in a solvent mixture of acetone and trifluoroethanol (3:1 v/v) at 288 K.<sup>24</sup> The rate of natural decay was determined to be  $5.4 \times 10^{-4} \text{ s}^{-1}$ . **CAUTION:** Perchlorate salts of metal complexes with organic ligands are potentially explosive and should be handled with care!

**2.2. Kinetic Studies.** All reactions were followed by monitoring UV–vis spectral changes of reaction solutions with a Hewlett-Packard 8453 spectrophotometer equipped with a circulating bath or UNISOKU cryostat system (USP-203; UNISOKU, Japan). Temperature-dependent kinetic experiments for O–O bond homolysis were performed on HS  $[(\text{TMC})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  (1 mM) and LS  $[(\text{N4Py})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  (0.5 mM) complexes by monitoring the decay of absorption bands at 526 and 536 nm, respectively, in a solvent mixture of acetone and trifluoroethanol (3:1 v/v) at the given temperatures. Temperatures were varied from 233 to 308 K. Rates of H-atom abstraction reactions by HS and LS nonheme  $\text{Fe}^{\text{III}}\text{--hydroperoxo}$  complexes,  $[(\text{TMC})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  (1 mM) and  $[(\text{N4Py})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  (0.5 mM), were examined with xanthene (5–40 mM) by monitoring spectral changes at 526 and 536 nm, respectively, in a solvent mixture of acetone and trifluoroethanol (3:1 v/v) at given temperatures. Pseudo-first-order fitting of the kinetic data allowed us to determine  $k_2$  values for the H-atom abstraction reactions of xanthene at given temperatures. Primary kinetic isotope effect values of the reaction between  $[(\text{TMC})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  or  $[(\text{N4Py})\text{Fe}^{\text{III}}\text{--OOH}]^{2+}$  and xanthene were calculated by comparing  $k_2$  values obtained from H-atom abstraction reactions of xanthene and that of xanthene-*d*<sub>2</sub> in a solvent mixture of acetone and trifluoroethanol (3:1 v/v). Reactions were run at least in triplicate, and data reported represent the average of these reactions.

**2.3. Spectroscopic Methods.** X-band EPR spectra were collected on a Bruker EMX spectrometer with Bruker ER 041XG microwave bridges and ER 4102ST/ER 5106QT cavities. Spectra were collected at temperatures between 3.8 and 50 K using an Oxford ITC503 temperature controller with an SR 900 continuous flow cryostat. Low-temperature UV–vis Abs spectra were taken on a Cary 500 spectrometer equipped with a Janis Research Super Vari-temp helium cryogenic dewar at 7 K. MCD spectra were taken on a Jasco J-810D spectropolarimeter equipped with an extended S20 photomultiplier tube and a SM-4000-7T superconducting magnet. VTVH MCD data were collected using a calibrated Cernox resistor (Lakeshore Cryogenics, calibrated 1.5–300 K) inserted into the sample cell to accurately measure the sample temperature. Resonance Raman spectra were obtained using a triple monochromator (Spex 1877 CP) with 1200, 1800, and 2400 grooves/mm holographic spectrograph gratings and Andor iDus and Newton CCD detectors cooled to  $-80^\circ\text{C}$ . Excitation was provided by Kr (Coherent 190C–K) ion, Ar (Innova Sabre 25/7) ion, and Dye (Coherent 599 equipped with Rhodamine 6G) lasers with incident powers of  $\sim 20$  mW using an  $\sim 135^\circ$  backscattering configuration. Samples were prepared in *d*<sub>6</sub>-acetone with 5 mM Fe concentration in NMR tubes cooled to 77 K in a liquid nitrogen finger dewar (Wilmad). <sup>18</sup>O samples were used for resonance

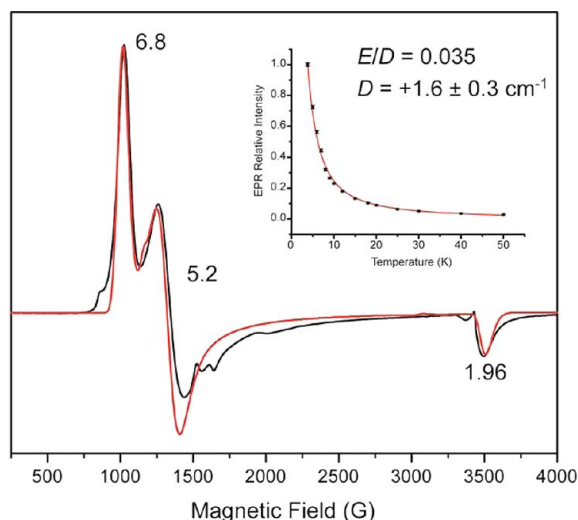
Raman profiling to avoid overlapping with solvent bands, and solvent peaks were used for internal standards.

**2.4. Density Functional Calculations.** Spin-unrestricted DFT calculations were performed with the Gaussian 09 package.<sup>33</sup> Geometry optimizations including transition state searches were performed using the hybrid density functional B3LYP with the 6-311G\* basis set for Fe and the 6-311G basis set for all other atoms. Frequency calculations were done at the same level of theory to ensure true local minima with no negative eigenvalues and that transition states had one and only one negative eigenvalue. Single-point and time-dependent DFT calculations were done at the B3LYP/6-311+G\*\* level of theory. Solvation effects were accounted for using the polarized continuum model (PCM) with acetone as the solvent. Optimization of the transition state in solvent yields similar energies and geometric/electronic structures. For the O–O bond homolysis and H-atom abstraction reaction calculations of the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex, acetonitrile was used as the axial ligand, since it is observed experimentally to bind to the [(TMC)Fe<sup>IV</sup>=O]<sup>2+</sup> product in acetonitrile solvent.<sup>23</sup> Orbital components were analyzed with QMForge.<sup>34</sup>

### 3. RESULTS AND ANALYSIS

**3.1. Spectroscopic.** **3.1.1. EPR.** Our previous EPR study of the [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex showed a signature  $S = 5/2$  spectrum with effective  $g$  values of 6.8, 5.2, and 1.96 (reproduced in Figure 2) described by the spin Hamiltonian

$$\hat{H}_{\text{spin}} = \beta \vec{B} g_0 \vec{S} + D \left[ S_z^2 - \frac{1}{3} S(S+1) \right] + E [S_x^2 - S_y^2] \quad (1)$$



**Figure 2.** X-band (3.8 K, 9.64 GHz) 2 mW EPR spectrum of the [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex (black) and simulation (red).<sup>37</sup> (Insert) Boltzmann fit (red curve) to the Curie law of the relative EPR intensities (black points) of the temperature dependence of the  $g = 6.8$  signal.

where  $\beta$  is the Bohr magneton,  $g_0$  is the free electronic  $g$  value,  $\vec{B}$  is the magnetic flux density,  $\vec{S}$  is the ground state total spin, and  $D$  and  $E$  are the axial and rhombic zero-field splitting parameters.<sup>23</sup> Simulation of the EPR spectrum gives a close to axial  $E/D$  value of 0.035, which is distinct from that of the rhombic HS Fe<sup>III</sup>–alkylperoxo complex, which was the first HS nonheme Fe<sup>III</sup>–OOR species that had been previously reported.<sup>24,35</sup> In order to obtain the sign and magnitude of  $D$ , EPR data of the [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex were collected between 3.8 and 50 K under nonsaturating

conditions. The temperature dependence of the relative intensity of the  $g = 6.8$  signal is shown in the insert in Figure 2. A Boltzmann fit of the data to the Curie law (eq 2,

$$A = \frac{C}{T} \left[ \frac{1}{1 + e^{-2D/kT} + e^{-6D/kT}} \right] \quad (2)$$

where  $A$  is the relative intensity,  $C$  is the Curie constant,  $T$  is the absolute temperature, and  $k$  is the Boltzmann constant) gives a value of  $D = +1.6 \pm 0.3 \text{ cm}^{-1}$ . The  $E/D$  and  $D$  values are similar to the  $E/D$  of 0.097 and  $D$  of  $+2.5 \text{ cm}^{-1}$  obtained from Mössbauer spectroscopy for the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex prepared in acetonitrile.<sup>36</sup>

**3.1.2. Abs, MCD, VTVH MCD, and Resonance Raman Profiles.** The low-temperature UV–vis absorption spectrum of the [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex is given in Figure 3A. A broad band at  $\lambda_{\text{max}}$  of  $19\,120 \text{ cm}^{-1}$  ( $523 \text{ nm}$ ,  $\epsilon \approx 2100 \text{ M}^{-1} \text{ cm}^{-1}$ ) is observed. The absorption intensity increases to higher energy with no distinct feature. This spectrum is similar to that of the HS Fe<sup>III</sup>–alkylperoxo complex, which had a broad band at  $\lambda_{\text{max}}$  of  $17\,900 \text{ cm}^{-1}$  ( $560 \text{ nm}$ ,  $\epsilon = 2000 \text{ M}^{-1} \text{ cm}^{-1}$ ).<sup>24</sup> The [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex has an additional weak band on the low-energy tail of the broad band (at  $\sim 13\,000 \text{ cm}^{-1}$ ,  $\epsilon = 280 \text{ M}^{-1} \text{ cm}^{-1}$ ). No equivalent low-intensity band was observed in the HS Fe<sup>III</sup>–alkylperoxo complex. If an equivalent low-energy band is present in the Fe<sup>III</sup>–alkylperoxo complex, it would have to be closer in energy to its intense broad band and thus not resolved.

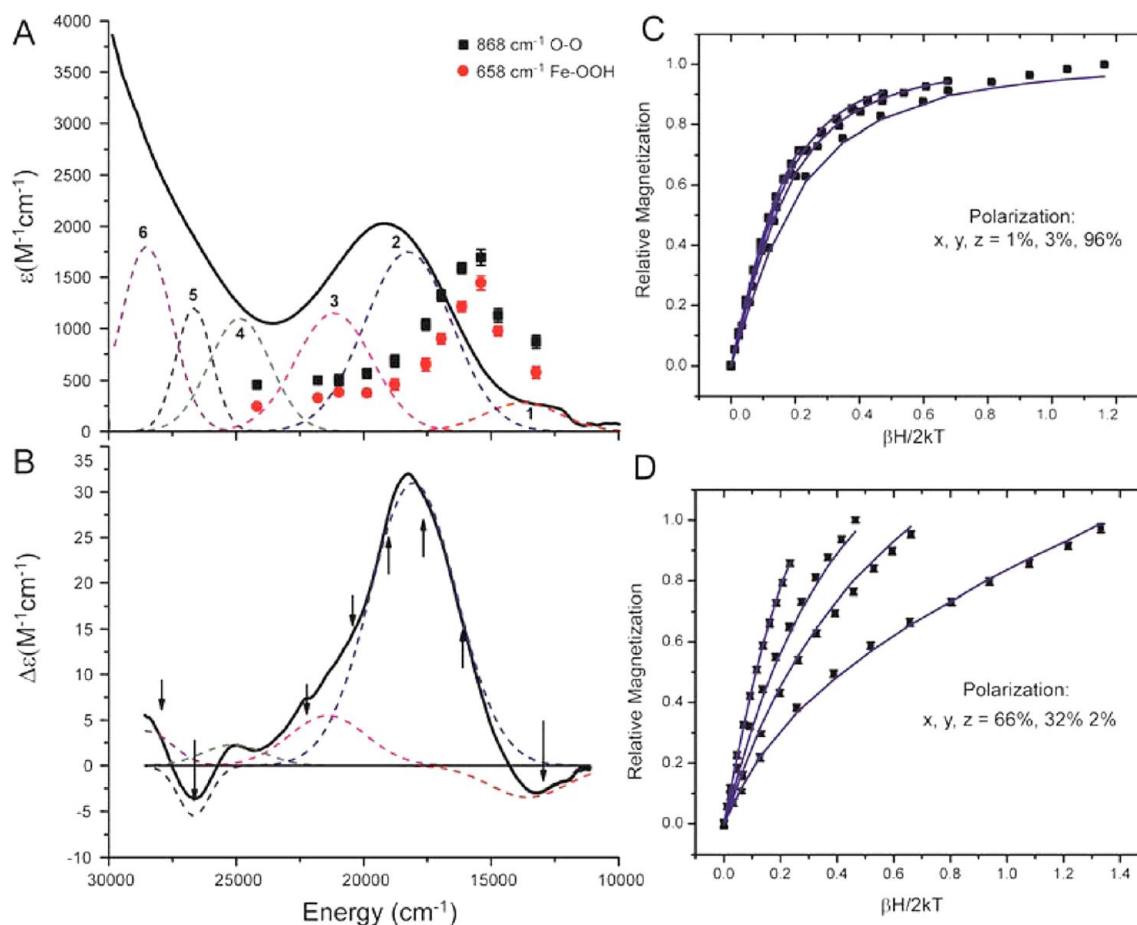
The low-temperature MCD spectrum of the [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex is shown in Figure 3B. The main feature is the broad, intense band centered at  $\sim 18\,000 \text{ cm}^{-1}$ . There is also a positive shoulder at  $\sim 22\,000 \text{ cm}^{-1}$ . The weak  $\sim 13\,000 \text{ cm}^{-1}$  feature in Abs is clearly resolved in the MCD spectrum as a negative band. Three other weak features are also resolved in the energy region above  $\sim 25\,000 \text{ cm}^{-1}$ . Simultaneous Gaussian fit of the Abs and MCD spectra resolves 6 bands, numbered in Figure 3A and summarized in Table 1.

Variable-temperature variable-field (VTVH) MCD data were collected at the energies indicated by the arrows shown in Figure 3B to determine the polarizations of these bands. VTVH MCD isotherms are recorded across the  $18\,000 \text{ cm}^{-1}$  band (band 2) at  $16\,000$ ,  $17\,795$ ,  $19\,085$ , and  $20\,490 \text{ cm}^{-1}$ . The VTVH MCD isotherms collected at  $17\,794$  and  $19\,084 \text{ cm}^{-1}$  are equivalent and show nesting behavior (Figure 3C), while the VTVH isotherms collected at  $16\,000$  and  $20\,492 \text{ cm}^{-1}$  show contributions from overlap with bands 1 and 3, respectively. The isotherms recorded at  $22\,124 \text{ cm}^{-1}$  for band 3 (Figure 3D) show a much more nested behavior and indicate a different polarization. VTVH MCD data collected at  $13\,000$ ,  $26\,455$ , and  $28\,090 \text{ cm}^{-1}$  were too noisy to analyze due to low intensity and overlap with the bands. VTVH MCD data were modeled using eq 3<sup>38</sup>

$$\frac{\Delta\epsilon}{E} = \frac{\gamma}{4\pi S} \int_0^\pi \int_0^{2\pi} \sum_i N_i (l_x \langle S_z \rangle_i M_{xy}^{\text{eff}} + l_y \langle S_y \rangle_i M_{xz}^{\text{eff}} + l_x \langle S_x \rangle_i M_{yz}^{\text{eff}}) \sin \theta \, d\theta \, d\phi \quad (3)$$

where  $N_i$  is the temperature-dependent Boltzmann population,  $x$ ,  $y$ , and  $z$  are the principle axes of the ZFS tensor,  $l_x$ ,  $l_y$ , and  $l_z$  are the direction cosines for the magnetic field relative to the molecular coordinate system,  $\langle S_x \rangle_i$ ,  $\langle S_y \rangle_i$ , and  $\langle S_z \rangle_i$  are the spin expectation values for the  $i$ th ground sublevel in the defined coordinate,  $M_{ij}^{\text{eff}}$  are the products of the  $i$ ,  $j$  polarizations of





**Figure 3.** (A and B) UV-vis absorption (A, 10 K) and MCD (B, 5 K and 7 T) spectra of the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex with simultaneous best Gaussian fit (bands 1–6). Arrows in the MCD spectrum indicate energies where VTVH MCD data were collected. Resonance Raman profiles of  $\nu(\text{O}-\text{O})$  ( $868\text{ cm}^{-1}$ , black rectangles) and  $\nu(\text{Fe}-\text{OOH})$  ( $658\text{ cm}^{-1}$ , red circles). (C and D) VTVH MCD isotherms (black rectangles) and best fits to the data (blue lines). Data were collected at 1.8, 3.5, 5, and 10 K. VTVH MCD taken at  $17\,794\text{ cm}^{-1}$  (C) and  $22\,124\text{ cm}^{-1}$  (D).

**Table 1. Results of the Simultaneous Gaussian Fit of the UV-Vis Absorption and MCD Spectra**

band	$\nu_{\text{max}}$ ( $\text{cm}^{-1}$ )	$\epsilon_{\text{max}}$ ( $\text{M}^{-1}\text{cm}^{-1}$ )	$\Delta\epsilon_{\text{max}}$ ( $\text{M}^{-1}\text{cm}^{-1}$ )	$f_{\text{osc}}^a$	$ \Delta\epsilon/\epsilon $
1	13 563	280	−3.5	0.0045	0.0125
2	18 050	1750	31.0	0.0321	0.0177
3	21 404	1150	5.5	0.0186	0.0048
4	25 109	1100	2.3	0.0142	0.0021
5	26 657	1200	−5.5	0.00905	0.0046
6	28 514	1800	3.8	0.0176	0.0021

<sup>a</sup>Oscillator strength,  $f_{\text{osc}} = 4.3 \times 10^{-9} \times \text{area under the absorption band}$ .

electronic transitions, and  $\gamma$  is a collection of constants. A fit of the VTVH MCD intensity to eq 3 using the spin Hamiltonian parameters  $D$  and  $E/D$  obtained from the EPR data, insert in Figure 2, allows effective transition moments  $M_{ij}^{\text{eff}}$  to be obtained. These transition moments can then be used to obtain the polarization using eq 4; % $y$  and % $z$  are obtained from cyclic permutations of the indices.<sup>38</sup>

$$\%x = 100 \times \left[ \frac{(M_{xy}^{\text{eff}} M_{xz}^{\text{eff}})^2}{(M_{xy}^{\text{eff}} M_{xz}^{\text{eff}})^2 + (M_{xy}^{\text{eff}} M_{yz}^{\text{eff}})^2 + (M_{yz}^{\text{eff}} M_{xz}^{\text{eff}})^2} \right] \quad (4)$$

From analysis of the VTVH MCD data taken at  $17\,790\text{ cm}^{-1}$ , band 2 is dominantly  $z$  polarized (96%  $z$  and 4%  $x,y$  polarized), which is the same as band 1 of the HS  $\text{Fe}^{\text{III}}$ -alkylperoxo complex.<sup>24</sup> From analysis of VTVH MCD data taken at  $22\,124\text{ cm}^{-1}$ , band 3 is dominantly  $x,y$  polarized (98%  $x,y$  and 2%  $z$  polarized).

Figure 3A also contains the resonance Raman profiles for the  $\nu(\text{O}-\text{O})$  stretch at  $868\text{ cm}^{-1}$  and  $\nu(\text{Fe}-\text{O})$  stretch at  $658\text{ cm}^{-1}$ . These two modes are resonance enhanced by both bands 1 and 2 and have maximum enhancement between these two bands. Thus, the excited states of these electronic transitions distort mostly along the Fe–O and O–O bonds relative to the ground state. Therefore, bands 1 and 2 correspond to hydroperoxo to iron charge transfer transitions. The HS  $\text{Fe}^{\text{III}}$ -alkylperoxo complex has its resonance Raman profiles corresponding only to the intense broad band, consistent with its lack of the lower energy band.<sup>24</sup>

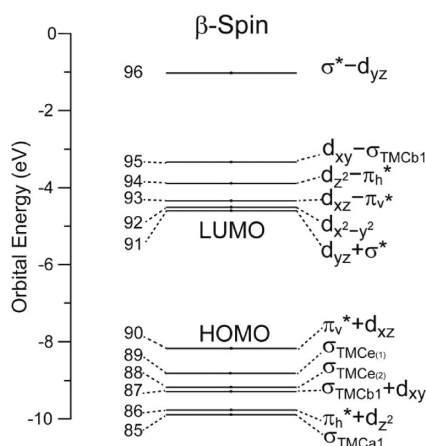
Since band 2 is  $z$  polarized and corresponds to the hydroperoxo to iron charge transfer, which is polarized along the Fe–O bond, the  $z$  axis of the ZFS tensor must be aligned with the Fe–OOH bond. This allows the coordinate system to be defined in Figure 1.

### 3.2. DFT Calculations: Correlation to Spectroscopy.

**3.2.1. Geometric Structure.** Top and side views of the DFT-optimized geometry of the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex are shown in Figure 1. As described in the Introduction, the

complex does not have an axial ligand trans to the OOH ligand. From DFT calculations,  $\angle\text{Fe}-\text{O}-\text{O}$  is  $120^\circ$  and the  $\text{Fe}-\text{O}-\text{O}$  plane bisects cis  $\text{N}-\text{Fe}$  bonds. The optimized  $\text{O}-\text{O}$  bond is 1.48 Å, and the predicted  $\text{Fe}-\text{O}$  bond and averaged  $\text{Fe}-\text{N}$  bonds are 1.84 and 2.19 Å, which agree well with EXAFS results of 1.85 and 2.16 Å, respectively.<sup>23</sup>

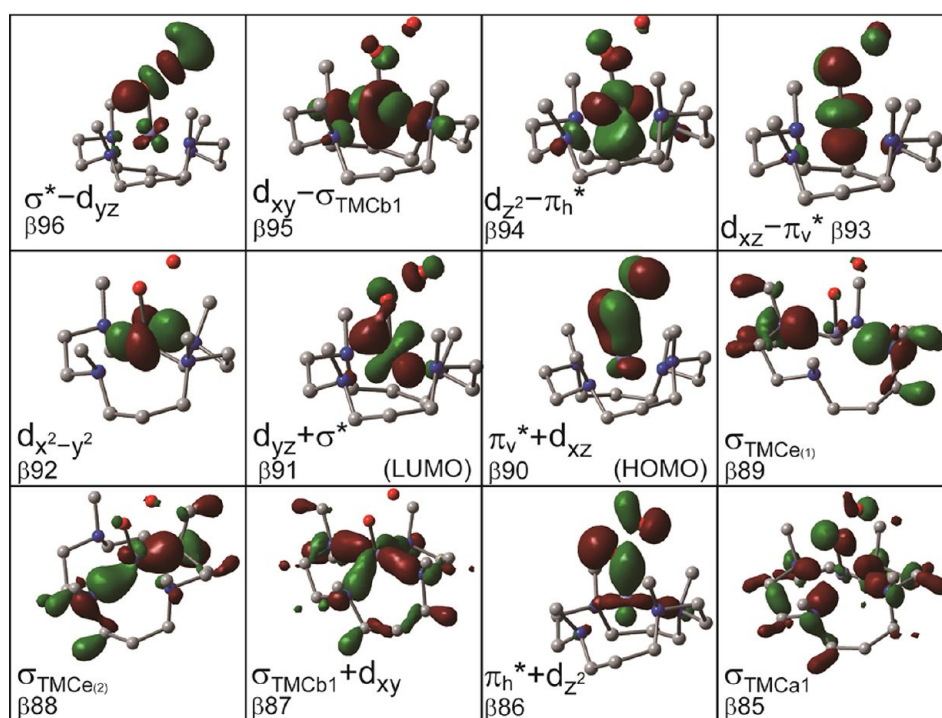
**3.2.2. Electronic Structure.** Since the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex has a ground state of  $S = 5/2$ , all five  $\alpha$ -spin d orbitals are occupied and all five  $\beta$ -spin d orbitals are unoccupied. As with the alkylperoxo ligand in ref 24, the hydroperoxo only acts as a donor ligand; therefore, only the  $\beta$ -spin d orbitals can contribute to net bonding. These are given in the spin-unrestricted molecular orbital (MO) diagram in Figure 4 with boundary surface plots in Figure 5. The



**Figure 4.**  $\beta$ -Spin molecular orbital diagram of the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex in the coordinate system defined in Figure 1. “a+b” and “a–b” notations mean bonding and antibonding molecular orbitals, respectively, between orbitals a and b, and a is the major component of the resulting molecular orbital.

coordinate system used is given in Figure 1 ( $z$  axis along the  $\text{Fe}-\text{O}$  bond, and  $y$  and  $x$  are in and perpendicular to the  $\text{Fe}-\text{O}-\text{O}$  plane, respectively, which bisects the equatorial  $\text{L}-\text{M}$  bonds).

We previously presented a detailed analysis of the electronic structure of a HS  $\text{Fe}^{\text{III}}$ -alkylperoxo complex.<sup>24</sup> Since the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex is similar, we only present a brief description here. The hydroperoxo ligand has two HOMO  $\pi^*$  orbitals that have donor interactions with Fe d orbitals and contribute to the  $\text{Fe}-\text{OOH}$  bonding. The out-of-plane  $\pi_v^*$  orbital of the hydroperoxo interacts with the  $d_{xz}$  orbital of the Fe in a  $\pi$ -type fashion. The resulting bonding orbital,  $\pi_v^*+d_{xz}$  ( $\beta 90$ ), is the  $\beta$ -spin HOMO. Similar to the HS  $\text{Fe}^{\text{III}}$ -alkylperoxo complex, the  $\pi_v^*+d_{xz}$  ( $\beta 90$ ) orbital has 25% iron and 73% hydroperoxo character, indicating a very covalent  $\pi$ -donor bond.<sup>24</sup> The corresponding antibonding orbital,  $d_{xz}-\pi_v^*$  ( $\beta 93$ ) (71% iron and 22% hydroperoxo), is unoccupied and the highest energy  $d\pi$  orbital. The second  $\pi^*$  orbital of the hydroperoxo (the in-plane  $\pi_h^*$ ) interacts with the Fe  $d_z^2$  to form a  $\sigma$ -type bonding molecular orbital  $\pi_h^*+d_z^2$  ( $\beta 86$ ) (25% iron and 73% hydroperoxo character) and a corresponding antibonding orbital  $d_z^2-\pi_h^*$  ( $\beta 94$ ) (73% iron and 10% hydroperoxo). Note that the  $d\sigma$   $d_z^2-\pi_h^*$  is only 0.46 eV higher than the  $d\pi$   $d_{xz}-\pi_v^*$  orbital, which is small relative to that of the HS  $\text{Fe}^{\text{III}}$ -alkylperoxo complex of 1.6 eV.<sup>24</sup> This reflects the lack of an axial ligand trans to the hydroperoxo ligand in the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex, which results in a lower  $d_z^2$  orbital energy. There are four TMC  $\sigma$ -donor orbitals from combination of the four amine  $\sigma$ -donor orbitals with  $a_1$ ,  $b_1$ , and  $e$  symmetries in approximate  $C_{4v}$ . The  $\sigma_{\text{TMCb1}}$  orbital interacts with the  $d_{xy}$  orbital of Fe in a  $\sigma$ -donor fashion and forms the  $\sigma_{\text{TMCb1}}+d_{xy}$  ( $\beta 87$ ) bonding and  $d_{xy}-\sigma_{\text{TMCb1}}$  ( $\beta 95$ ) antibonding orbitals, the latter being at higher energy than the other  $d\sigma$  orbital ( $d_z^2-\pi_h^*$  ( $\beta 94$ )). The  $\beta$ -spin LUMO is a  $d\pi$ -type orbital,  $d_{yz}+\sigma^*$  ( $\beta 91$ ) (85% iron and 5% hydroperoxo). It is the

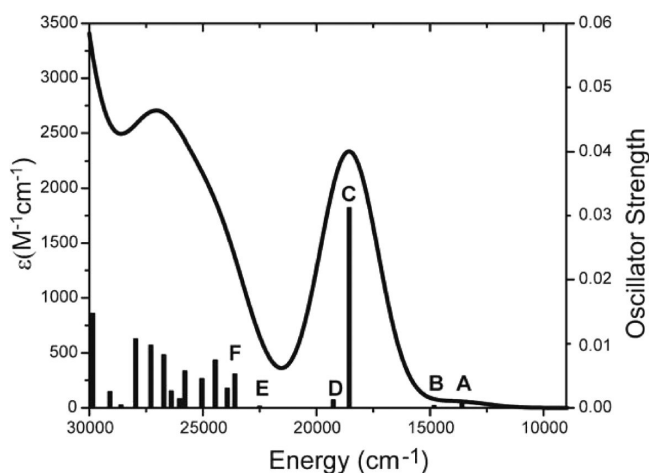


**Figure 5.** Boundary surface plots of the important  $\beta$ -spin MOs of the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex.

bonding combination of iron  $d_{yz}$  and the  $\sigma^*$  LUMO of the hydroperoxo ligand; the corresponding antibonding combination is the  $\sigma^*-d_{yz}$  ( $\beta 96$ ) (9% iron and 90% hydroperoxo) orbital, located  $\sim 3$  eV above the d manifold. Since both are unoccupied, there is no net contribution to bonding, but this shifts the  $d_{yz}+\sigma^*$  ( $\beta 91$ ) orbital to lower energy than the nonbonding  $d_{x^2-y^2}$  ( $\beta 92$ ) orbital. This will be significant with respect to the spectroscopy and reactivity.

### 3.2.3. TD DFT Calculations: Spectroscopic Assignments.

From Figure 3A, the absorption spectrum of the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex features an intense broad band at  $\sim 19\,120\text{ cm}^{-1}$  and a weak band on its low energy side at  $\sim 13\,000\text{ cm}^{-1}$ . These two features are well reproduced by TD DFT calculations, Figure 6. The weak band is predicted to arise



**Figure 6.** TD DFT predicted absorption spectrum of the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex. Bars represent the values of calculated oscillator strength (right axis) of the transitions at the calculated energies indicated. First six excited states are labeled as A–F, respectively.

from excitations A and B at  $13\,595$  and  $14\,830\text{ cm}^{-1}$ , respectively. Transitions A and B correspond to the  $\pi_v^*+d_{xz} \rightarrow d_{yz}+\sigma^*$  (92% pure, HOMO to LUMO) and  $\pi_v^*+d_{xz} \rightarrow d_{x^2-y^2}$  (96% pure) charge transfer (CT) excitations in Figure 4, with low predicted oscillator strengths of  $0.0006$  and  $0.0003$ , respectively. This is consistent with the resonance Raman enhancement of the Fe–O and O–O stretches observed for band 1 in Figure 3A. The low intensity of these transitions is due to the small coefficient of mixing between the donor (mostly hydroperoxide  $\pi_v^*$ ) and the acceptor ( $d_{yz}+\sigma^*$  and  $d_{x^2-y^2}$ ) MOs. As mentioned in section 3.1.2, no equivalent low-energy low-intensity band was observed in the HS  $\text{Fe}^{\text{III}}$ -alkylperoxo species in ref 24. This is because the acceptor  $d_{yz}+\sigma^*$  orbital is closer to the acceptor  $d_{xz}-\pi_v^*$  orbital in the HS  $\text{Fe}^{\text{III}}$ -alkylperoxo than in the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex ( $0.112$  vs  $0.252\text{ eV}$ ). Transitions C and D in Figure 6 at  $18\,560$  and  $19\,260\text{ cm}^{-1}$  give rise to the intense broad Abs band that reproduces the energy position and intensity of the  $19\,120\text{ cm}^{-1}$  band 2 in the absorption spectrum in Figure 3A. Excitations C and D correspond to the intense  $\pi_v^*+d_{xz} \rightarrow d_{xz}-\pi_v^*$  (81% pure) and weak  $\pi_v^*+d_{xz} \rightarrow d_{z^2}-\pi_h^*$  (86% pure) CT transitions, with predicted oscillator strengths of  $0.0312$  and  $0.0012$ , respectively. The high intensity of the  $z$ -polarized  $\pi_v^*+d_{xz} \rightarrow d_{xz}-\pi_v^*$  CT transition reflects the high covalency of the  $\pi$  bond between the  $d_{xz}$  and the hydroperoxo  $\pi_v^*$  orbitals. Coupled to the VTVH MCD data, band 2 of the Abs spectrum

(Figure 3A) is assigned to be the  $\pi_v^*+d_{xz} \rightarrow d_{xz}-\pi_v^*$  CT transition. Transition D corresponds to a CT to the  $d_{z^2}$  orbital and is close in energy to C (CT to a  $d\pi^*$ -type orbital) due to the lower energy of the  $d_{z^2}$  orbital associated with the lack of axial ligand for this complex. Excitations E and F are two TMC to iron CT transitions,  $\sigma_{\text{TMCb1}}+d_{xy} \rightarrow d_{x^2-y^2}$  (82% pure) (a  $\sigma$  bonding  $\rightarrow$  nonbonding) and  $\sigma_{\text{TMCe(1)}} \rightarrow d_{x^2-y^2}$  (49% mixed character) at  $22\,500$  and  $23\,600\text{ cm}^{-1}$ , with predicted oscillator strengths of  $0.0002$  and  $0.0052$ , respectively. These transitions are  $xy$  polarized consistent with VTVH MCD on band 3 in Figure 3D. At higher energy are other states corresponding to TMC and hydroperoxo  $\pi_h^*+d_{z^2}$  to iron CT transitions, which are summarized in Supporting Information Table S1, consistent with the additional bands in Figure 3A.

In summary, the spectral data combined with TD DFT calculations on the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex show that the  $\pi_v^*$  orbital of the hydroperoxo ligand forms a strong covalent  $\pi$ -donor bond with the  $d_{xz}$  orbital of  $\text{Fe}^{\text{III}}$  and gives rise to the dominant intense  $z$ -polarized CT transition band 2. The  $xy$ -polarized band 3 arises from TMC to  $\text{Fe}^{\text{III}}$  CT transitions. It is important to note that the low-energy band 1 present in the HS  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex reflects a particularly low-energy CT to the  $d_{yz}+\sigma^*$  orbital and indicates the presence of a configuration interaction with the higher energy hydroperoxo  $\sigma^*$  orbital which will provide significant insight into reactivity (vide infra). The density functional and basis set (geometric optimization by B3LYP/6-311g\* for Fe, 6-311g for other atoms; single point by B3LYP/6-311+g\*\* with solvent correction) used in these calculations reproduce well the geometric and electronic structure of the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex and are used for reaction coordinate calculations below.

**3.3. O–O Bond Homolysis.** **3.3.1. Kinetics.** In our recent study, the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex was found to undergo O–O bond homolysis.<sup>23</sup> This is the first HS ferric hydroperoxide species reported to undergo homolysis. This observation however does not appear to be consistent with the results in ref 24 that the O–O bond homolysis of the HS system has an extra  $\sim 10\text{ kcal/mol}$  barrier relative to the LS  $\text{Fe}^{\text{III}}-\text{OOH}$  system which does undergo O–O bond homolysis.<sup>24</sup> The additional O–O bond homolysis barrier of HS systems is due to an allowed orbital crossing that originates from the noninteracting nature of the redox-active orbitals on the HS  $\text{Fe}^{\text{III}}$  and the hydroperoxide. Alternatively, O–O bond homolytic cleavage of the LS system involves a forbidden orbital crossing that originates from the strong configuration interaction of the redox-active orbitals and thus has no additional barrier for this endergonic reaction.<sup>24</sup> It is important to note, however, that the product of  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  homolysis is a 6C LS  $S = 1\text{ Fe}^{\text{IV}}=\text{O}$  species, and thus, a spin surface crossing must occur at some point in the reaction coordinate.

In order to quantitatively compare the reactivities of HS and LS  $\text{Fe}^{\text{III}}-\text{OOH}$  species in O–O bond homolysis, temperature-dependent kinetic experiments were performed on HS  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  and on the LS  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex. The LS 6C  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex undergoes O–O homolysis to also form a 6C LS  $S = 1\text{ Fe}^{\text{IV}}=\text{O}$  species.<sup>39</sup> Kinetic parameters obtained are summarized in Table 2. Arrhenius plots (Figure S1, Supporting Information) of the temperature-dependent kinetic data show that the activation energy  $E_a$  of the O–O bond homolysis of the HS complex is slightly higher than that of the LS complex. The Eyring plots



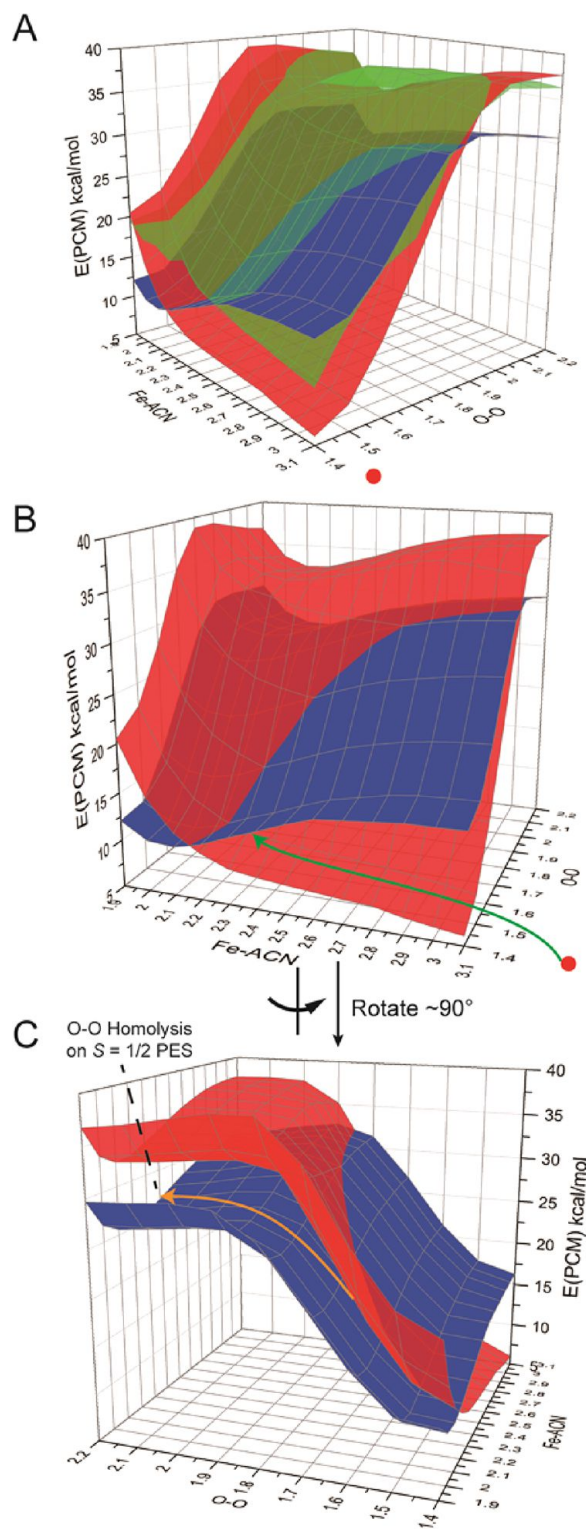
**Table 2. Kinetic Parameters of HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> and LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> O–O Homolytic Cleavages**

	[(TMC)Fe <sup>III</sup> –OOH] <sup>2+</sup>	[(N4Py)Fe <sup>III</sup> –OOH] <sup>2+</sup>
$E_a$ (kcal/mol)	13.9(5)	12.9(2)
$\Delta H^\ddagger$ (kcal/mol)	13.4(5)	12.6(3)
$\Delta S^\ddagger$ (cal/mol·K)	–17.9(4)	–28.9(2)
$\Delta G^\ddagger$	18.7(5) (298 K)	20.9(3) (298 K)

(Figure S2, Supporting Information) of the temperature-dependent kinetic data yield a slightly higher  $\Delta H^\ddagger$  and a less negative  $\Delta S^\ddagger$  for the HS complex than those of the LS complex. At 298 K,  $\Delta G^\ddagger$  of the HS complex is 2.2 kcal/mol lower than that of the LS complex.<sup>24</sup> Thus, the experimental O–O bond homolysis energetics of the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> and the LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> complexes are very similar, which does not appear to be consistent with the previous prediction that a HS hydroperoxo system will have an additional barrier for O–O bond homolysis.<sup>24</sup>

**3.3.2. DFT Calculations of O–O Bond Homolysis.** Since the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex changes coordination number and spin state along the reaction coordination, a potential energy surface crossing must occur.<sup>23</sup> In order to evaluate this process, we constructed high-spin (HS,  $S = 5/2$ ), intermediate-spin (IS,  $S = 3/2$ ), and low-spin (LS,  $S = 1/2$ ) two-dimensional potential energy surfaces (PES), where the O–O bond is scanned from 1.4 to 2.2 Å with 0.1 Å increments and the Fe–acetonitrile bond (the solvent which is capable of coordinating to Fe<sup>III</sup>) is scanned from 1.9 to 3.1 Å with 0.1 Å increments. The overlay of these three PESs is shown in Figure 7A. Note that the freely optimized HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex has an O–O bond of 1.48 Å and an Fe–acetonitrile distance of 4.37 Å. Its approximate position on the HS PES (red, transparent) is indicated as a red solid dot in Figure 7A. In general, the IS PES (green, transparent) is at higher energy than the HS PES (red, transparent) over most of the coordinates surveyed, Figure 7A. Although the HS PES is lower in energy than the LS PES (blue, opaque) at long Fe–acetonitrile (Fe–ACN) and short O–O bond lengths, the LS PES is lower in energy than the HS PES at short Fe–ACN and long O–O bond lengths. This is because the [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex energetically favors binding the axial ligand in the  $S = 1/2$  spin state and disfavors axial ligand binding in the  $S = 5/2$  spin state. Thus, the HS and LS PESs cross and result in an isoenergetic two-dimensional seam, Figure 7B and 7C.

All three PESs have their O–O bond homolysis saddle points in the regions around an Fe–ACN bond length of ~2.1 Å and an O–O bond length of ~1.9 Å. The LS PES has the lowest barrier of ~26 kcal/mol, the HS PES has a barrier of ~34 kcal/mol, and the IS PES has a barrier of ~36 kcal/mol. The energy difference for O–O bond homolysis barriers on the HS and LS PESs is thus consistent with the previous prediction in ref 24 that the barrier on the HS PES is ~10 kcal/mol higher than that of the LS PES, and, in fact, O–O bond homolysis does not occur on the HS PES.<sup>24</sup> The PESs in Figure 7, combined with the experimental results that the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex undergoes O–O bond homolysis to form a LS 6C  $S = 1$  Fe(IV)=O species and that the O–O bond homolysis of the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> and the LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> complexes have similar activation energies, indicate that the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex first crosses to the LS PES and then undergoes O–O bond homolysis on the LS PES. In Figure 7B, the green arrow



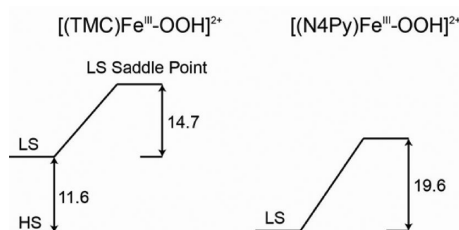
**Figure 7.** 2D potential energy surfaces of  $S = 5/2$  (red, transparent),  $3/2$  (green, transparent), and  $1/2$  (blue, opaque) spin states of O–O bond homolysis of the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex with acetonitrile (ACN) as a trans axial ligand to the hydroperoxo group. Two coordinates scanned are O–O and Fe–ACN (in Ångströms). Energies are related to the freely optimized HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex with an O–O bond length of 1.48 Å and an Fe–ACN distance of 4.37 Å. Its approximate position on the  $S = 5/2$  PES is indicated as a red solid dot: (A) overlay of  $S = 5/2$ ,  $3/2$ , and  $1/2$  PESs; (B) overlay of  $S = 5/2$  and  $1/2$  PESs; (C) 90° rotation of B.



indicates a reasonable pathway for O–O bond homolysis of the HS complex, where the Fe–ACN coordinate first contracts without O–O bond elongation. As the Fe–ACN distance decreases, the HS PES increases in energy and the LS PES decreases. These become isoenergetic at the crossing point on the seam of the HS and LS PES. At an Fe–ACN of  $\sim 2.2$  Å and an O–O of  $\sim 1.5$  Å, the energy is  $\sim 10$  kcal/mol higher than that of the freely optimized HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex (Fe–ACN distance = 4.37 Å). In this region of the 2D surfaces, the IS PES is only  $\sim 7$  kcal/mol higher in energy and would provide a mechanism for second-order spin–orbit coupling between the HS and the LS states, which allows their crossing. After the HS to LS cross over, the system proceeds along the minimum energy path (mostly along the O–O coordinate, orange arrow in Figure 7C) with an Fe–ACN distance of  $\sim 2.1$  Å (Figure 7C) and undergoes O–O bond homolysis to produce the  $S = 1$  [(TMC)Fe<sup>IV</sup>=O]<sup>2+</sup> acetonitrile-bound product and a hydroxyl radical.

Focusing on  $\Delta H^\ddagger$  to avoid added issues associated with entropy at the transition state, the DFT-calculated  $\Delta H^\ddagger$  of the O–O bond homolysis of the LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex is 19.6 kcal/mol, in Scheme 1 (right). Compared to

**Scheme 1.** DFT-Calculated  $\Delta H^\ddagger$  (298 K, in kcal/mol) of the O–O Bond Homolysis Saddle Point on the LS PES (right) Compared with Freely Optimized HS and LS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> without and with Acetonitrile Bound as an Axial Ligand, Respectively (left)



the experimental results, the DFT calculations overestimate this  $\Delta H^\ddagger$  value by 7.4 kcal/mol, Table 2. The DFT-calculated  $\Delta H^\ddagger$  of the O–O bond homolysis of the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex which occurs on the LS PES is 26.3 kcal/mol, which is 12.9 kcal/mol higher than the experimental  $\Delta H^\ddagger$  value. The DFT-calculated  $\Delta H^\ddagger$  value for [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> can be decomposed into two contributions, the energy difference (11.6 kcal/mol) between LS and HS states and the energy difference (14.7 kcal/mol) between the LS PES saddle point and LS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup>. Taking into account the 7.4 kcal/mol that the DFT overestimates the O–O bond homolysis of the LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex, the spin state energy difference in the B3LYP calculations would be off by 5.5 kcal/mol. This spin state splitting is known to be very sensitive to functional. When the Hartree–Fock exchange of 20% of the B3LYP hybrid functional is decreased to 10% and 0%, this HS and LS energy splitting does in fact decrease from 11.6 kcal/mol in B3LYP to  $-1.7$  and  $-11.6$  kcal/mol (0% HF exchange).

In summary, the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex first crosses to the LS PES (via second-order spin–orbit coupling) through axial ligand binding at a short O–O length and then undergoes O–O bond homolysis on the LS PES.

### 3.4. H-Atom Abstraction Reactivities. 3.4.1. Kinetics.

In an earlier study, several LS Fe<sup>III</sup>–OOH model complexes, including [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup>, were found to be incapable

of oxidizing cyclohexene (C–H bond of 83.9 kcal/mol).<sup>40</sup> This is in contrast to activated bleomycin, a LS Fe<sup>III</sup>–OOH species, which performs direct H-atom abstraction reaction for the strong C–H bond of sugar (92 kcal/mol).<sup>6–8</sup> In our recent study, we provided direct experimental evidence that the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> is capable of abstracting an H atom from weak C–H bonds (xanthene (75.5 kcal/mol) and 9,10-dihydroanthracene (77 kcal/mol)), showing that the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> is as reactive as the LS [(TMC)Fe<sup>IV</sup>=O]<sup>2+</sup> species in H-atom abstraction.<sup>23</sup> In order to compare the electrophilic reactivities of HS and LS Fe<sup>III</sup>–OOH species, we performed analogous kinetic experiments on HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> and LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> with the same substrate (e.g., xanthene) in the same solvent system (e.g., a solvent mixture of acetone and trifluoroethanol (3:1 v/v)).

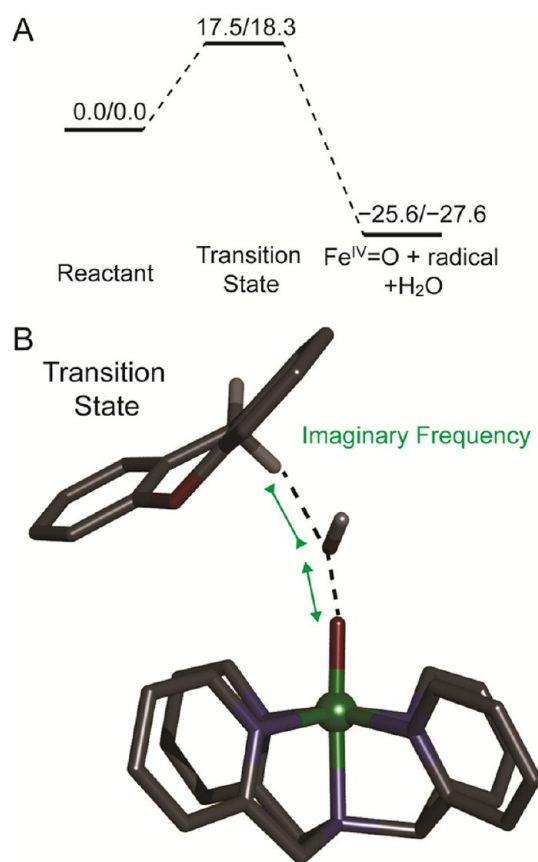
A second-order rate constant of  $6.4 \times 10^{-2} \text{ M}^{-1} \text{ s}^{-1}$  was determined for the C–H bond activation of xanthene by the LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex at 288 K, which is slower than the second-order rate constant ( $8.1 \times 10^{-1} \text{ M}^{-1} \text{ s}^{-1}$ ) for this reaction by HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> at 253 K. Temperature-dependent kinetic data are summarized in Table 3. The

**Table 3.** Kinetic Parameters of HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> and LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> Reactions with Xanthene

	[(TMC)Fe <sup>III</sup> –OOH] <sup>2+</sup>	[(N4Py)Fe <sup>III</sup> –OOH] <sup>2+</sup>
$E_a$ (kcal/mol)	11.7(3)	14.4(2)
$\Delta H^\ddagger$ (kcal/mol)	11.2(3)	13.9(3)
$\Delta S^\ddagger$ (cal/mol·K)	–21.5(2)	–15.8(2)
$\Delta G^\ddagger$ (kcal/mol)	16.7(2) (253 K), 17.4(3) (288 K)	18.4(3) (288 K)
$K_{\text{H}_2\text{O}}/K_{\text{D}_2\text{O}}$ (H <sub>2</sub> O vs D <sub>2</sub> O) (kcal/mol)	1.2(2)	1.3(2)
$K_{\text{H}}/K_{\text{D}}$ (xanthene vs <i>d</i> <sub>2</sub> -xanthene) (kcal/mol)	5.0(2)	3.0(2)

Arrhenius plots (Figure S3, Supporting Information) of the temperature-dependent kinetic data show that the activation energy  $E_a$  of the H-atom abstraction of xanthene by the HS complex is 2.7 kcal/mol lower than that of the LS complex. From the Eyring plots (Figure S4, Supporting Information) of the temperature-dependent kinetic data,  $\Delta G^\ddagger$  of the HS complex is 1 kcal/mol lower than that of the LS complex at 288 K, which is consistent with the fact that the HS complex has a higher second-order rate constant than that of the LS complex. Thus, the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex is slightly more reactive than the LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex in H-atom abstraction from xanthene.

**3.4.2. DFT-Calculated H-Atom Abstraction Reaction Coordinate of LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup>.** The DFT-predicted reaction coordinate with xanthene as the substrate on the  $S = 1/2$  PES is shown in Figure 8A. The transition state is 17.5/18.3 kcal/mol ( $\Delta H^\ddagger/\Delta G^\ddagger$ ) higher in energy than the reactant complex, which is in reasonable agreement with the experimental result (13.9/18.4 kcal/mol), Table 3. Products are predicted to be an  $S = 1$  [(N4Py)Fe<sup>IV</sup>=O]<sup>2+</sup> species, a H<sub>2</sub>O, and an  $S = 1/2$  radical; the reaction is exothermic at  $-25.6/-27.6$  kcal/mol ( $\Delta H/\Delta G$ ). The calculated transition state has only one imaginary frequency and a vibrational motion associated with O–O bond cleavage and O–H bond formation (see Figure 8B). Important structural and electronic properties of the reactant, transition state, and product are



**Figure 8.** (A)  $S = 1/2$  PES ( $\Delta H/\Delta G$  at 288 K, with solvent correction, in kcal/mol) for reaction of the LS  $[(N4Py)Fe^{III}-OOH]^{2+}$  complex with xanthene. (B) Transition state structure of the direct H-atom abstraction reaction between the LS  $[(N4Py)Fe^{III}-OOH]^{2+}$  complex and xanthene. Fe atom is in green, N atoms are in blue, C atoms are in black, O atoms are in red, H atom is in white. Only important H atoms are shown; others are omitted for clarity.

summarized in Table 4. At the transition state, the O–O bond length is 2.382 Å compared to 1.507 Å in reactant, which

**Table 4. Structural and Electronic Parameters Changes of the H-Atom Abstraction Reaction with Xanthene by the LS  $[(N4Py)Fe^{III}-OOH]^{2+}$  Complex**

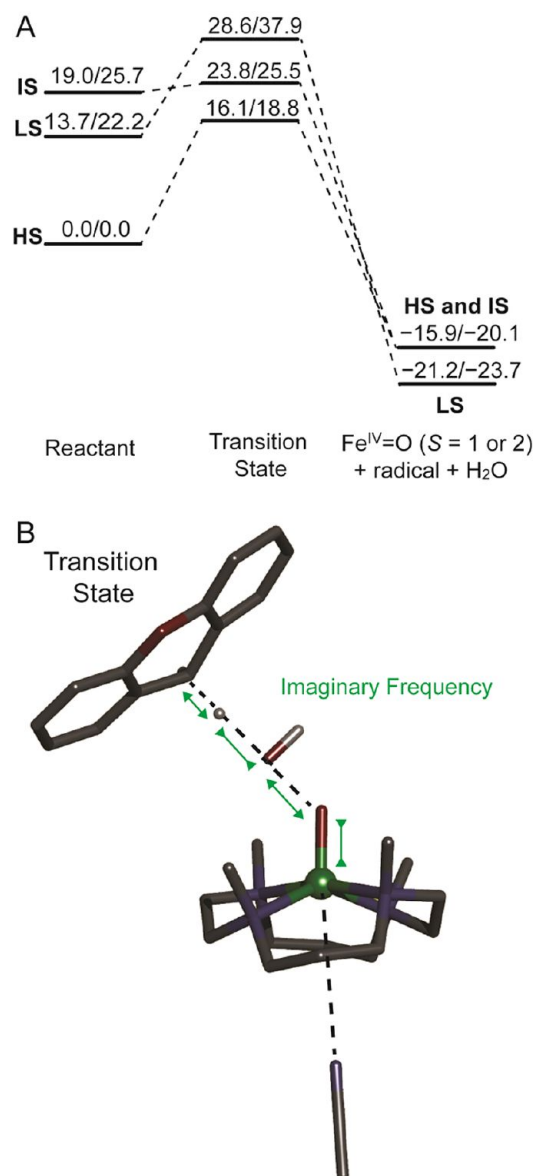
	reactant	transition state	product
structure			
$r(Fe-O)$	1.779 Å	1.635 Å	1.645 Å
$r(O-O)$	1.507 Å	2.382 Å	
$r(O_{dist}-H)$	2.947 Å	1.979 Å	0.971 Å
$r(H-C)$	1.098 Å	1.118 Å	
spin densities			
Fe	0.83	1.22	1.21
$O_{prox}$	0.21	0.74	0.84
$O_{dist}H$	0.02	−0.80	0.00
xanthene–H	0.00	−0.10	−1.00
H atom	0.00	0.00	0.01

indicates that the O–O bond is essentially broken; the C–H bond of xanthene is only slightly elongated from 1.098 to 1.118 Å; the O–H distance is still fairly long at 1.979 Å. These bond lengths indicate the transition state occurs late in O–O cleavage coordinate and early in H-atom transfer coordinate.

The  $O_{dist}H$  unit has a Mulliken spin density of −0.80 at the transition state, which indicates the  $O_{dist}H$  unit is essentially a hydroxyl radical resulting from O–O bond homolysis. The fact that the Fe–O bond is 1.635 Å compared to 1.779 Å in the reactant and the Mulliken spin density on the Fe–O moiety is  $\sim 2$  compared to  $\sim 1$  in the reactant indicates that Fe–O is essentially an  $S = 1$   $Fe^{IV}=O$  at transition state, resulting from O–O bond homolysis. This is supported by molecular orbital occupation analysis. These structural and electronic properties are very similar to the transition state for the direct H-atom abstraction reaction from the C4'–H of DNA by ABLM (i.e., a LS  $Fe^{III}-OOH$  species).<sup>7</sup> The similarities of these transition states indicate that the LS  $[(N4Py)Fe^{III}-OOH]^{2+}$  complex and ABLM perform direct H-atom abstraction using the equivalent reaction coordinates.

**3.4.3. DFT-Calculated H-Atom Abstraction Reaction Coordinate of HS  $[(TMC)Fe^{III}-OOH]^{2+}$ : Comparison to LS  $[(N4Py)Fe^{III}-OOH]^{2+}$ .** Since the HS  $[(TMC)Fe^{III}-OOH]^{2+}$  O–O bond homolysis undergoes a HS to LS cross over before O–O bond cleavage, we evaluated the reaction coordinate of  $[(TMC)Fe^{III}-OOH]^{2+}$  for H-atom abstraction on the  $S = 5/2$ ,  $3/2$ , and  $1/2$  (HS, IS, and LS) PESs, as shown in Figure 9A. The ground state of the reactant is HS, and the  $S = 5/2$  transition state is lower in energy than the other two ( $S = 1/2$  and  $3/2$ ) transition states. Therefore, in contrast to the O–O bond homolysis process, the H-atom abstraction reaction on the HS PES proceeds through the HS  $S = 5/2$  transition state. For xanthene as a substrate, the  $S = 5/2$  transition state is calculated to be 16.1/18.8 kcal/mol ( $\Delta H^\ddagger/\Delta G^\ddagger$ ) higher in energy than the reactant complex. The DFT-predicted  $\Delta H^\ddagger$  value is  $\sim 5$  kcal/mol higher than the experimental value (11.2 kcal/mol, Table 3). However, it is lower than the DFT-predicted  $\Delta H^\ddagger$  (17.5 kcal/mol) of the LS  $[(TMC)Fe^{III}-OOH]^{2+}$  H-atom abstraction reaction and thus reproduces the experimental trend. The calculated  $S = 5/2$  transition state has only one imaginary frequency and vibrational motion associated with O–O bond cleavage, O–H bond formation, and Fe–O shortening (see Figure 9B).

The geometric and electronic structures of the  $S = 5/2$  HS  $[(TMC)Fe^{III}-OOH]^{2+}$  transition state (HS TMC TS) are very different from that of the  $S = 1/2$  LS  $[(N4Py)Fe^{III}-OOH]^{2+}$  transition state (LS N4Py TS) (see Tables 4 and 5). The HS TMC TS has an O–O bond of 1.786 Å, which is much shorter than that of the LS N4Py TS (2.382 Å). According to the Mayer bond order,<sup>41</sup> the HS TMC O–O bond is  $\sim 52\%$  broken at the transition state, whereas the LS N4Py TS O–O bond is  $\sim 83\%$  broken. The C–H bond of xanthene in HS TMC TS is elongated from 1.098 to 1.170 Å, which is longer than that of the LS N4Py TS (1.118 Å). Again, according to Mayer bond order analysis,<sup>41</sup> the HS TMC TS C–H bond is  $\sim 23\%$  broken whereas the LS N4Py TS C–H bond is only  $\sim 5\%$  broken. The longer C–H bond in the HS TMC TS is consistent with the primary KIE values of 5.0 and 3.0 for HS and LS  $Fe^{III}-OOH$  species, respectively (see Table 3). These bond lengths indicate the HS TMC TS occurs early in both the O–O and the H–C cleavage coordinates. The close to zero Mulliken spin density on the  $O_{dist}H$  unit of HS TMC TS indicates that in contrast to the Mulliken spin density of −0.80 of the  $O_{dist}H$  unit of the LS N4Py TS the  $O_{dist}H$  unit is not a hydroxyl radical. It is important to note that there is 0.33  $e^-$  transfer from xanthene to HS  $[(TMC)Fe^{III}-OOH]^{2+}$  at the TS; however, almost no electron (0.04  $e^-$ ) transfer occurs from xanthene to the LS  $[(N4Py)Fe^{III}-OOH]^{2+}$  at the TS. The electronic structure

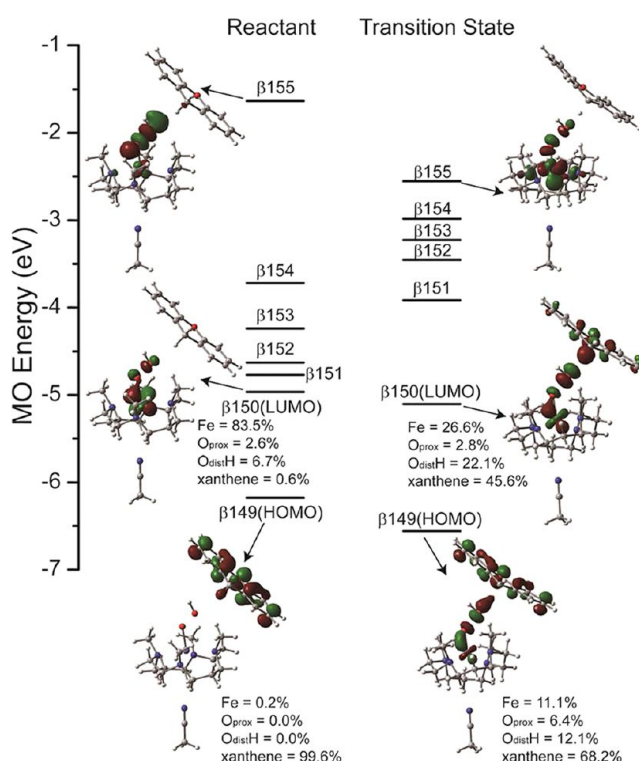


**Figure 9.** (A)  $S = 1/2$ ,  $3/2$ , and  $5/2$  (LS, IS, and HS) PES ( $\Delta H/\Delta G$  at 253 K, with solvent correction, in kcal/mol) for reaction of the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex with xanthene. Transition states energies were calculated as the energy difference between the transition state and reactant components calculated in one box. Product energies were calculated as the energy difference between product and reactant components calculated in separate boxes to prevent electron transfer from product radical to  $[(\text{TMC})\text{Fe}^{\text{IV}}=\text{O}]^{2+}$ . (B)  $S = 5/2$  transition state structure for direct H-atom abstraction reaction between the HS  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex and xanthene. Fe atom is in green, N atoms are in blue, C atoms are in black, O atoms are in red, H atom is in white. Only important H atoms are shown; others are omitted for clarity. Note that the axial ligand, modeled as acetonitrile for the reason mentioned in section 2.4, is not coordinated in the TS (4.39 Å) on the HS PES.

calculations (Figure 10) reveal that the  $0.33 e^-$  comes solely from the xanthene-based  $\beta$ -spin HOMO ( $\beta 149$ , 99.6% xanthene) and is transferred into the  $\text{Fe}^{\text{III}}-\text{OOH}$ -based  $\beta$ -spin LUMO ( $\beta 150$ , 83.5% Fe and 9.3%  $\text{OOH}^-$ , which is the  $d_{yz}+\sigma^*$  orbital). This  $\beta$ -spin LUMO is responsible for the low-energy absorption band of the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex in Figure 3A. In summary, the geometric and electronic

**Table 5. Structural and Electronic Parameters Changes from Reactant to Transition of the H-Atom Abstraction Reaction with Xanthene by the HS  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  Complex on  $S = 5/2$  PES**

	reactant	transition state
	structure	
$r(\text{Fe}-\text{O})$	1.862 Å	1.749 Å
$r(\text{O}-\text{O})$	1.512 Å	1.786 Å
$r(\text{O}_{\text{dist}}-\text{H})$	2.016 Å	1.606 Å
$r(\text{H}-\text{C})$	1.098 Å	1.170 Å
spin densities		
Fe	4.02	3.85
$\text{O}_{\text{prox}}$	0.35	0.26
$\text{O}_{\text{dist}}\text{H}$	0.06	0.05
xanthene-H	0.00	0.32
H atom	0.00	0.01



**Figure 10.** Energies and boundary surface plots of the important  $\beta$ -spin MOs of the reactant and transition state of the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  reaction with xanthene on the  $S = 5/2$  PES (using acetonitrile as the potential axial ligand, see section 2.4). Mulliken population analyses are given for the  $\beta$ -spin HOMO and LUMO of the reactant and transition state.

structures of the transition states of HS TMC and LS N4Py are very different and imply different reaction coordinates, which will be discussed below.

The energies of the experimentally defined H-atom abstraction reaction products (e.g.,  $[(\text{TMC})\text{Fe}^{\text{IV}}=\text{O}]^{2+}$ ,  $\text{H}_2\text{O}$ , and xanthene-H radical) were obtained by calculating the individual components. The  $S = 1$   $[(\text{TMC})\text{Fe}^{\text{IV}}=\text{O}]^{2+}$  product is predicted to be the ground state, which is in agreement with experiment.<sup>23</sup> Thus, the system needs to cross from the  $S = 5/2$  to the  $S = 1/2$  PES, Figure 9A. Since after the transition state  $\text{Fe}^{\text{IV}}=\text{O}$  has an  $S = 2$  electronic structure, its crossing to the  $S = 1$  PES was evaluated. PES scans along the Fe–O coordinate



(Figures S7A and S7B, Supporting Information) of  $S = 2$  and 1  $[(\text{TMC})\text{Fe}^{\text{IV}}=\text{O}]^{2+}$  reveal that these PESs do not cross, independent of whether the axial acetonitrile binds to Fe. Since the  $S = 1$   $[(\text{TMC})\text{Fe}^{\text{IV}}=\text{O}]^{2+}$  has shorter Fe–equatorial ligand bonds than those of  $S = 2$   $[(\text{TMC})\text{Fe}^{\text{IV}}=\text{O}]^{2+}$  (average Fe–equatorial bond lengths of 2.09 vs 2.16 Å, respectively), we further surveyed the PESs for contraction along these coordinates, Figure S7C, Supporting Information, which demonstrates that the energies of the  $S = 2$  and 1  $[(\text{TMC})\text{Fe}^{\text{IV}}=\text{O}]^{2+}$  species are indeed sensitive to one pair of Fe–equatorial ligand bonds, defined as Fe–eqB. Therefore, 2D PESs along both the Fe–O and the Fe–eqB coordinates were evaluated, Figure S7D, Supporting Information, and a reasonable reaction pathway is indicated by the black arrows. Starting from the transition state, the Fe–O bond contracts on the  $S = 2$  PES. At short Fe–O and long Fe–eqB bond length, the system crosses from the  $S = 2$  to the  $S = 1$  PES via an allowed spin–orbit coupling and then proceeds to shorten the Fe–eqB coordinate on the  $S = 1$  PES to form the LS product observed experimentally.<sup>23</sup>

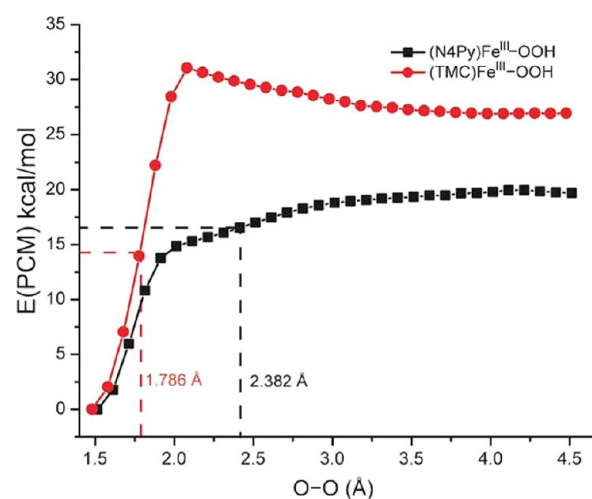
It is of interest to note that when the HS system is propagated past its TS, a HS  $S = 5/2$   $\text{Fe}^{\text{III}}=\text{O}$  product is observed that is at  $\sim -45$  kcal/mol (Figure S6, Supporting Information), which represents an effective hydride abstraction reaction that is not observed experimentally.<sup>23</sup> The electronic structure of the HS TMC TS described above clearly shows an H-atom abstraction process. Therefore, a second electron transfer from the substrate follows an initial H-atom abstraction. A recent computational study of H-atom abstraction by HS  $S = 2$   $\text{Fe}^{\text{IV}}=\text{O}$  species observed an analogous second electron transfer.<sup>42</sup> For the HS  $\text{Fe}^{\text{III}}-\text{OOH}$  system, the second electron transfer, which does not occur experimentally, can be attributed to the self-interaction error inherent in DFT when the system is positively charged.<sup>43–45</sup>

In summary, the H-atom abstraction reaction by the HS  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex occurs on the  $S = 5/2$  PES with a different reaction coordinate compared to that of the LS  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex. For the HS system, the transition state features a shorter O–O and longer C–H bonds than those of the LS transition state.

#### 4. DISCUSSION

In agreement with ref 24, O–O bond homolysis of HS  $\text{Fe}^{\text{III}}-\text{OOH}$  complexes should have a  $\sim 10$  kcal/mol higher cleavage barrier relative to LS  $\text{Fe}^{\text{III}}-\text{OOH}$  complexes as shown in Figure 11. The O–O bond homolysis observed in the five-coordinate  $S = 5/2$  (HS)  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex in fact appears to occur by first crossing to the LS PES through axial ligand binding at short O–O lengths and then undergoing O–O bond homolysis on the LS PES to produce the six-coordinate  $S = 1$   $[(\text{TMC})\text{Fe}^{\text{IV}}=\text{O}]^{2+}$  species that is observed in experiment.

With respect to H-atom abstraction from xanthene, the LS  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  features a transition state late in the O–O coordinate and early in the C–H coordinate. This transition state essentially reflects O–O bond homolysis, generating a hydroxyl radical, which then performs H-atom abstraction from the substrate. This is consistent with the fact that the O–O bond homolysis energy at the O–O bond length associated with the transition state for H-atom abstraction (2.382 Å) is  $\sim 20$  kcal/mol (Figure 11). To evaluate the contribution of O–O homolysis energy to the transition state energies of the HS and LS  $\text{Fe}^{\text{III}}-\text{OOH}$  H-atom abstraction reactions, we decomposed the transition state energies into



**Figure 11.** One-dimensional PESs of HS  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  (red dots) and LS  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  (black rectangles) along the O–O bond.

three contributions: the O–O distortion energy of the  $\text{Fe}^{\text{III}}-\text{OOH}$  at the transition state, the C–H distortion energy of the substrate at the transition state, and the interaction energy of  $\text{Fe}^{\text{III}}-\text{OOH}$ /substrate at the transition state. As shown in Table 6, summation of these contributions is in good agreement with the directly calculated TS energies, which validates this decomposition scheme. For reaction of the LS  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  with xanthene, the substrate distortion energy is only 0.6 kcal/mol, the O–O distortion energy is 19.6 kcal/mol, and the TS energy is 20.0 kcal/mol. Therefore, the O–O distortion energy is the dominant contribution to the activation energy for H-atom abstraction by the LS  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex.

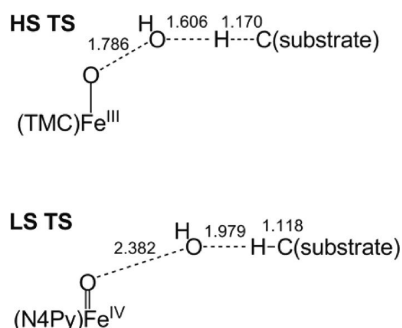
The transition state for H-atom abstraction from xanthene by  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  is very similar to that of the direct H-atom abstraction reaction from DNA by ABLM.<sup>7</sup> Interestingly, the ABLM reaction is even more reactive ( $E_a = 4.7$  kcal/mol) against a stronger C–H bond (92 kcal/mol of sugar vs 75.5 kcal/mol of xanthene). This is likely due to the equatorial ligand difference between  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  and ABLM. ABLM has one negatively charged deprotonated amide ligand, which dominates the Fe  $d\pi$  hole orientation of the LS  $\text{Fe}^{\text{III}}-\text{OOH}$  complex, and this reduces the overlap between the  $d\pi$  hole and the hydroperoxo ligand.<sup>46</sup> This would destabilize the reactant. This ligand would also stabilize the transition state due to its electrostatic stabilization of the  $\text{Fe}^{\text{IV}}=\text{O}$  formed. Both would contribute to a lower  $E_a$  for ABLM relative to  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$ , enabling its reaction with the strong C–H bond of DNA.

With respect to the H-atom abstraction reaction by the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  species, in contrast to the O–O homolysis, this reaction occurs on the  $S = 5/2$  PES. The HS TS for the H-atom abstraction reaction is strikingly different from the LS TS in its geometric structure, Scheme 2. In contrast to the LS TS, the HS TS is early in the O–O coordinate (TS O–O bond length is 1.786 Å for HS compared to 2.382 Å for LS). From Figure 11, for O–O homolysis on the HS surface, the PES is very steep between 1.5 and 2.1 Å, which means that a slight increase of the O–O bond length corresponds to a large increase of energy. Therefore, the barrier would be too high for HS  $\text{Fe}^{\text{III}}-\text{OOH}$  to undergo O–O bond homolysis to produce a hydroxyl radical for the H-atom abstraction. Also, the HS TS is further along in the C–H

**Table 6.** Energy Decomposition of H-Atom Abstraction Reaction Transition States of HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> and LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> with the Series of Substrates Defined in Figure S8, Supporting Information<sup>a</sup>

substrates	xanthene 0F	1F	2F	3F	4F	6F	dioxine	dioxine 3F	dioxine 6F
HS [(TMC)Fe <sup>III</sup> –OOH] <sup>2+</sup>									
TS energy	18.4	19.2	19.7	22.6	24.3	25.2	25.3	27.0	28.5
Fe <sup>III</sup> –OOH/substrate interaction energy	–6.3	–5.0	–5.5	–6.1	–6.8	–6.3	–12.1	–11.1	–8.5
Fe <sup>III</sup> –OOH distortion energy	22.1	20.9	20.3	22.2	24.4	24.9	31.4	31.9	30.1
substrate distortion energy	2.8	2.7	2.8	5.1	7.7	7.2	7.4	7.8	8.5
summation	18.6	18.6	17.7	21.2	25.3	25.7	26.8	28.6	30.1
LS [(N4Py)Fe <sup>III</sup> –OOH] <sup>2+</sup>									
TS energy	20.0	19.9	19.7	19.8	20.3	20.4	20.8	21.3	20.9
Fe <sup>III</sup> –OOH/substrate interaction energy	0.2	0.0	0.0	–0.2	–0.9	–0.9	–3.1	–3.3	–3.4
Fe <sup>III</sup> –OOH distortion energy	19.3	19.3	19.4	20.1	20.0	19.9	21.2	21.0	20.6
substrate distortion energy	0.6	0.5	0.3	1.3	2.0	1.7	3.6	4.2	5.2
summation	20.1	19.9	19.8	21.3	21.1	20.9	21.7	21.9	22.4

<sup>a</sup>Electronic energies are given with solvation correction. Important geometric parameters of these transition states are shown in Table S3, Supporting Information. The interaction energy is obtained as the difference between the energy of Fe<sup>III</sup>–OOH complex and the substrate at the transition state geometries but kept 10 Å away and the energy of the transition state.

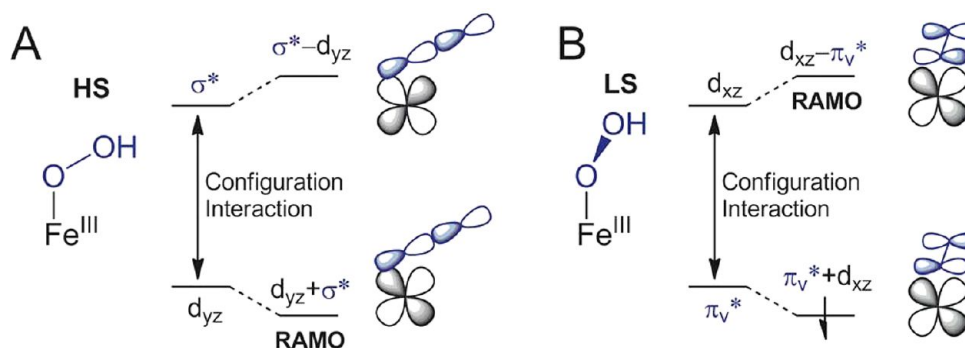
**Scheme 2.** Schematics of the Transition States (TS) of HS and LS Fe<sup>III</sup>–OOH in the H-Atom Abstraction Reaction<sup>a</sup>

<sup>a</sup>HS TS features shorter O–O and longer C–H bonds than those of the LS TS. Bond lengths are in Angstroms.

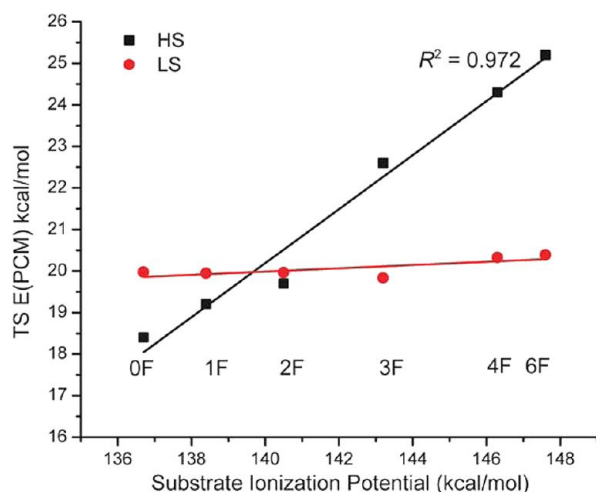
coordinate than the LS TS (1.170 vs 1.118 Å), which is in agreement with the experimental difference in primary KIE effect (e.g., HS:LS = 5:3). The HS TS is also distinctly different from the LS TS in electronic structure. From section 3.4.3, 0.33 e<sup>–</sup> transfers from the substrate to the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> at the transition state; however, almost no electron (0.04 e<sup>–</sup>) is transferred from substrate to the LS complex at its transition state. The HS *S* = 5/2 complex might be expected to be harder to reduce due to the greater loss of exchange

stabilization. However, DFT calculations indicate that the reduction potential of HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> is in fact 810 mV higher than that of the LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup>. This is consistent with experiment for iron complexes with similar ligands and the same charge but different spin state.<sup>47</sup> The longer Fe–ligand bonds of the HS state result in less charge donation and thus a higher *Z*<sub>eff</sub> for Fe<sup>III</sup>. Also, the redox-active MO (RAMO) of the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> is the β-spin LUMO (*d*<sub>yz</sub>+σ\* orbital in Figure 5), which undergoes configuration interaction with the higher energy hydroperoxo σ\* orbital that further lowers its energy and also contributes to the higher reduction potential, Scheme 3A. Alternatively, the RAMO of the LS [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> complex is a strong antibonding Fe *d*π orbital (with the hydroperoxo π<sub>v</sub>\*, Scheme 3B) and therefore higher in energy.<sup>25</sup> Together, these favor electron transfer from substrate to the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup>.

Since there is significant e<sup>–</sup> transfer from the substrate to Fe<sup>III</sup>–OOH in the HS TMC TS but not in the LS N4Py TS and since the substrate C–H bond contributes more in the HS TMC TS than the LS N4Py TS, the H-atom abstraction reactivity of the HS Fe<sup>III</sup>–OOH should be more dependent on substrate than that of the LS Fe<sup>III</sup>–OOH. In order to evaluate this idea, we carried out DFT calculations to compare the dependence of the energies of the transition states for H-atom abstraction of the HS [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> and the LS

**Scheme 3.** Configuration Interaction between (A) *d*<sub>yz</sub> and Hydroperoxo σ\* Orbitals of [(TMC)Fe<sup>III</sup>–OOH]<sup>2+</sup> Lowers the Energy of the *d*<sub>yz</sub>+σ\* Orbital as the RAMO and Hydroperoxo π<sub>v</sub>\* and *d*<sub>xz</sub> Orbitals of [(N4Py)Fe<sup>III</sup>–OOH]<sup>2+</sup> Increases the Energy of *d*<sub>xz</sub>–π<sub>v</sub>\* Orbital as the RAMO

$[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  on the ionization potential and C–H bond strength of substrates. Xanthene and fluorinated xanthenes (abbreviated as xanthene 0F, 1F, 2F, 3F, 4F, and 6F; structures and full formulas are given in Figure S8, Supporting Information) were chosen to test the substrate ionization potential dependence. These compounds are calculated to have very different ionization potentials (varying from 136.7 kcal/mol for xanthene to 148.1 kcal/mol for hexafluoroxanthene, Table S2, Supporting Information) but with very similar C–H bond strengths (within 1.6 kcal/mol, Table S2, Supporting Information). As shown in Figure 12, the



**Figure 12.** Dependence of the activation energies of HS  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  (black rectangles) and LS  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  (red dots) H-atom abstraction reactions with xanthene and fluorinated xanthenes as indicated in the figure. Full names of these compounds are shown in Figure S8, Supporting Information.

HS  $\text{Fe}^{\text{III}}-\text{OOH}$  is more reactive (i.e., lower transition state energy) than the LS  $\text{Fe}^{\text{III}}-\text{OOH}$  toward xanthene 0F, 1F, and 2F (substrates with low ionization potentials) but less reactive toward 3F, 4F, and 6F (substrates with high ionization potentials). Thus, the activation energies for H-atom abstraction by HS  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  (black rectangles) are clearly correlated with the substrate ionization potentials ( $R^2 = 0.972$ ). Alternatively, from Figure 12, the activation energies for H-atom abstraction by LS  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  (red dots) are independent of substrate ionization potential. These results are consistent with the model that  $e^-$  transfer occurs in the HS TMC TS but not in the LS TMC TS. For xanthene as the substrate, its  $e^-$  transfer to the  $\text{Fe}^{\text{III}}-\text{OOH}$  has about a 6 kcal/mol stabilization effect on the HS TS, Table 6.

Alternatively, to examine the dependence of reactivity on the strength of C–H bonds, the transition states of H-atom abstraction from xanthene and a dioxine derivative (4b,10a-dihydrobenzo[*b*]benzo[3,4]cyclobuta[1,2-*e*][1,4]dioxine, abbreviated as dioxine, Figure S8, Supporting Information) were evaluated. These compounds have very similar ionization potentials (1.3 kcal/mol from experiment<sup>48</sup> and 0.3 kcal/mol from calculation, Table S2, Supporting Information) but are calculated to have very different C–H bond strengths (dioxine is 21 kcal/mol stronger than xanthene, Table S2, Supporting Information). As shown in Table 6, the activation energies of HS  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  with xanthene and dioxine reactions are 18.4 and 25.3 kcal/mol, respectively, clearly dependent on C–H bond strength. However, for H-atom

abstraction reactions of LS  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$ , the activation energy is only slightly higher (0.8 kcal/mol) even though this is a much stronger C–H bond. This is consistent with the small variation of activation energy for H-atom abstraction by the hydroxyl radical from methane and halomethanes which have large variations of C–H bond strength.<sup>49,50</sup> In conclusion, the reactivity of HS  $\text{Fe}^{\text{III}}-\text{OOH}$  is dependent on substrate ionization potential and C–H bond strength. In contrast, the reactivity of LS  $\text{Fe}^{\text{III}}-\text{OOH}$  is independent of substrate ionization potential and only slightly dependent on substrate C–H bond strength.

## 5. CONCLUSION

LS  $\text{Fe}^{\text{III}}-\text{OOH}$  complexes should be more reactive toward substrates with strong C–H bonds as in ABLM, whereas HS  $\text{Fe}^{\text{III}}-\text{OOH}$  complexes should be more reactive toward substrates with low ionization potentials and weak C–H bonds. The higher reduction potentials of the HS  $\text{Fe}^{\text{III}}-\text{OOH}$  complexes also allow these to be active in electrophilic reactions without the requirement of O–O cleavage. This is important for the reaction coordinate in Rieske dioxygenases, which catalyze *cis*-dihydroxylation of a wide range of aromatic compounds. As mentioned in the Introduction, a HS  $\text{Fe}^{\text{III}}-\text{OOH}$  intermediate is believed to be involved in the catalytic cycles of these enzymes. The open question is whether this intermediate is able to directly carry out *cis*-dihydroxylation with aromatic substrates or whether the O–O bond first cleaves to generate a high-valent  $\text{HO}^\bullet-\text{Fe}^{\text{IV}}=\text{O}$  or  $\text{HO}-\text{Fe}^{\text{V}}=\text{O}$  species for the *cis*-dihydroxylation.<sup>10,20–22</sup> While the former mechanism has been favored by calculations,<sup>51</sup> the latter mechanism has been favored based on a small amount of  $^{18}\text{O}$  incorporation into product from solvent.<sup>52</sup> This study suggests that the former reaction coordinate is a plausible mechanism.

## ■ ASSOCIATED CONTENT

### Supporting Information

Detailed assignments for the TD DFT predicted absorption spectrum of the  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  complex; descriptions of the substrates used in Discussion, DFT predicted ionization potentials and C–H bond strengths of these substrates, and the geometric parameters of the transition states of their reactions with  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  and  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$ ; Arrhenius and Eyring plots of the unimolecular decay and xanthene reactions of  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  and  $[(\text{N4Py})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$ ; geometric and electronic parameters of the transition states for the reactions of  $[(\text{TMC})\text{Fe}^{\text{III}}-\text{OOH}]^{2+}$  with xanthene on the  $S = 3/2$  and  $1/2$  PESs; 2D PESs for the spin crossover of the  $S = 2$  and 1 spin states of  $[(\text{TMC})\text{Fe}^{\text{IV}}=\text{O}]^{2+}$ . This material is available free of charge via the Internet at <http://pubs.acs.org>.

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### Notes

The authors declare no competing financial interest.

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