



Cite this: *RSC Adv.*, 2017, 7, 50713

Synthesis and reactivity of a 4His enzyme model complex†

Jia Li,^a Atanu Banerjee,^a Timothy A. Hasse,^a Reza Loloee,^b Shannon M. Biros,^c Richard J. Staples^c and Ferman A. Chavez^{id}*^a

A new iron(II) complex has been prepared and characterized. $[\text{Fe}(\text{TrIm})_4(\text{OTf})_2]$ (**1**, TrIm = 1-tritylimidazole). The solid state structure of **1** has been determined by X-ray crystallography. Compound **1** crystallizes in triclinic space group $P\bar{1}$, with $a = 13.342(7)$ Å, $b = 13.5131(7)$ Å and $c = 13.7025(7)$ Å. The iron center resides in distorted octahedral geometry coordinated to four equatorial imidazole groups and two axial triflate oxygens groups. The complex is high spin between 20 K and 300 K as indicated by variable field variable temperature magnetic measurements. A fit of the magnetic data yielded $g = 2.24$ and $D = -0.80$ cm⁻¹. A large HOMO–LUMO gap energy (3.89 eV) exists for **1** indicating high stability. Addition of H₂O₂ or ^tBuOOH to **1** results in formation of an oxygenated intermediate which upon decomposition results in oxidation of the trityl substituent on the imidazole ligand.

Received 25th August 2017
 Accepted 9th October 2017

DOI: 10.1039/c7ra09456f

rsc.li/rsc-advances

Introduction

Carotenoids are isoprene-containing compounds possessing conjugated π systems.¹ In plants, carotenoids carry out a variety of critical functions. They operate as accessory light-harvesting components (antenna complex)^{2,3} of photosynthetic systems, light induced antioxidants (brought about by the low-lying triplet state poised to quench triplet chlorophyll and singlet oxygen⁴), membrane fluidity regulators, pigments (yellow to red-orange; important in attracting pollinators)² for leaves, vegetables, and fruits.⁵ Along with carotenoids being essential in plants, carotenoid-derived products serve as important sources of bioactive compounds called apocarotenoids.⁶ Apocarotenoids serve as pigments, hormones (abscisic acid and strigolactones),⁷ aroma and scent compounds, regulatory compounds and molecules with yet to be determined functions. Carotenoids are also present in various fungi and heterotrophic bacteria as well as all photosynthetic organisms.⁶ In animals, carotenoids yield essential precursors necessary for vision, embryonic development, cellular homeostasis, and immunity (vitamin A).^{2,8,9} Carotenoid Cleavage Dioxygenases (CCDs) are enzymes possessing a nonheme site and are found in all life forms. CCD processes carotenoids and related substrates.^{10,11} CCDs generally catalyze the cleavage of carotenoid carbon–carbon double bonds to yield

aldehydes and/or ketones. CCD must have appeared early in evolution since one of the products (retinal) is critical for photoreception by the ancient type 1 opsin protein family, a process critical in phototactic and phototrophic organisms.¹²

Synthesis of abscisic acid (ABA), occurs during the oxidative cleavage of the 11,12 carbon–carbon double bond of a 9-*cis*-epoxycarotenoid (9-*cis*-violaxanthin).¹³ This reaction is carried out by the enzyme viviparous14 (VP14) in maize (*Zea mays*). The X-ray structure of VP14 has been determined (PDB: 3NPE) and reveals an active site consisting of a nonheme iron coordinated to four histidine side chain residues. The remaining sites are occupied by water (or hydroxide) and dioxygen.¹³ The putative bound oxygen molecule is coordinated to iron in an end-on fashion. This type of iron coordination is also seen in photosystem II, superoxide reductase, and fumurate reductase.^{14–16} Apocarotenoid cleavage oxygenase (ACO), a CCD from cyanobacteria catalyzes the formation of 3-hydroxyretinal from 3-hydroxy-8'-apo- β -carotenol. The X-ray structure of ACO from *Synechocystis* was determined and also possesses iron ligated by four histidines.¹⁷ Retinal and its derivatives are involved in several mammalian biological processes and in particular the immune system and vision.¹⁸ CCDs which form retinal include β -carotene-15,15'-oxygenase (BCO1) generating retinal from symmetrical cleavage β -carotene. Retinal becomes reduced to retinol and is converted to all-*trans* retinyl esters by lecithin retinol acyltransferase.¹⁹

In our efforts to model the 4His active site of CCD, we have employed a bulky imidazole ligand (1-tritylimidazole, TrIm) to synthesize a 4His nonheme site ($[\text{Fe}(\text{TrIm})_4(\text{OTf})_2]$ (**1**), Scheme 1). The synthesis, structure, and characterization (UV-vis and FTIR spectroscopy, and SQUID magnetometry) as well as theoretical studies (DFT) are presented. Reactivity studies are also presented.

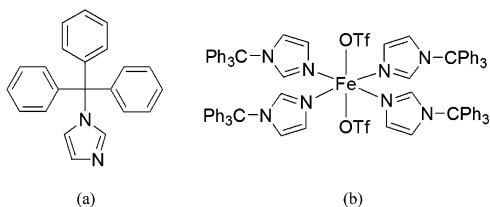
^aDepartment of Chemistry, Oakland University, Rochester, MI 48309, USA. E-mail: chavez@oakland.edu; Tel: +1 248 370-4092

^bDepartment of Physics and Astronomy, Michigan State University, East Lansing, MI 48824, USA

^cDepartment of Chemistry, Michigan State University, East Lansing, MI 48824, USA

† Electronic supplementary information (ESI) available: CCDC 1548010. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c7ra09456f





Scheme 1 (a) Structure of 1-tritylimidazole (TrIm) and (b) its iron(II) complex $[\text{Fe}(\text{TrIm})_4(\text{OTf})_2]$ (**1**).

Experimental details

Materials

1-Tritylimidazole and ^tBuOOH (TBHP, 70% dried over anhydrous magnesium sulfate) were purchased from Sigma-Aldrich Chemical Co. H₂O₂ (30% dried over anhydrous magnesium sulfate to afford 78% as determined by density) was purchased from EMD Millipore. Fe(OTf)₂·2CH₃CN was synthesized according to a literature method.²⁰ Pure dry solvents were afforded using an Innovative Technologies, Inc. Solvent Purification System. Elemental analysis was performed on pulverized crystalline samples that were heated under vacuum and sealed in a glass ampule prior to submission (Atlantic Microlabs, Inc., Norcross, GA).

Synthesis

[Fe(TrImA)₄(OTf)₂]·2Et₂O (1·2Et₂O). To a stirring suspension of 1-tritylimidazole (158.9 mg, 0.512 mmol) in dichloromethane (2 mL) was added a solution of Fe(OTf)₂·2CH₃CN (55.8 mg, 0.128 mmol) in dichloromethane (2 mL) under dry nitrogen atmosphere. After 1 h of stirring, the solution was filtered and placed in an ether diffusion chamber. After 24 h, colorless crystals were deposited. The crystals were collected and washed with ether. Yield: 155.7 mg (76%). Selected IR bands (KBr pellet, cm⁻¹): 1489 (s), 1444 (s), 1308 (s), 1235 (s), 1216 (s), 1167 (s), 1112 (s), 1078 (s), 1025 (s), 932 (s), 845 (m), 747 (s), 663 (s), 633 (s) and 516 (m). UV-vis (CH₃CN; λ_{max}, nm (ε, M⁻¹ cm⁻¹): 255 (8202, sh), 322 (2040, sh); (CH₂Cl₂; λ_{max}, nm (ε, M⁻¹ cm⁻¹): 256 (6775, sh), 294 (2320, sh). Anal. calcd for C₉₀H₇₂N₈O₆F₆S₂Fe (**1**): C, 67.75; H, 4.55; N, 7.02. Found: C, 67.96; H, 4.99; N, 6.63.

Physical characterization

A Varian 3100 Excalibur Series was used to collect the FTIR spectra. Optical spectra were collected using a Cary 50 UV-vis spectrophotometer. Variable temperature magnetic susceptibilities were measured using a Quantum Design MPMS3 SQUID magnetometer calibrated with a 765-Palladium standard purchased from NIST (formally NSB). A powdered sample of **1** was placed in a plastic bag. The sample was measured in the temperature range 2–300 K with $H = 0.6$ T. The sample was placed in a plastic drinking straw for the measurement. The magnetic contribution of the bag was determined between 2–300 K as well and subtracted from the sample. The molar magnetic

susceptibility was corrected for the diamagnetism of the complex using tabulated values of Pascal's constants to obtain a corrected molar susceptibility. The program julX written by E. Bill was used for the simulation and analysis of magnetic susceptibility data.²¹ The Hamiltonian operator (H) was

$$g\beta\hat{S}\cdot\hat{B} + D\left[\hat{S}_z^2 - 1/3S(S+1) + E/D(\hat{S}_x^2 - \hat{S}_y^2)\right] \quad (1)$$

where g is the average electronic g value, D the axial zero-field splitting parameter, and E/D is the rhombicity parameter. Magnetic moments were obtained from numerically generated derivatives of the eigenvalues of eqn (1), and summed up over 16 field orientations along a 16-point Lebedev grid to account for the powder distribution of the sample. Intermolecular interactions were considered by using a Weiss temperature, θ_w , as perturbation of the temperature scale, $kT' = k(T - \theta_w)$. GCMS experiments were performed on an HP 6890/5973 gas chromatograph. The products were identified and quantified by comparing the retention times along with mass spectra and peak areas with authentic standards.

Computational details

Quantum chemical calculations providing energy minimized molecular geometries, molecular orbitals (HOMO–LUMO), and vibrational spectra for compound **1** were carried out using density functional theory (DFT) as implemented in the GAUSSIAN09 (Rev. C.01) program package.²² We employed the hybrid functional PBE0.²³ The basis set used was 6-31G(d).²⁴ Full ground state geometry optimization was performed without any symmetry constraints. Only the default convergence criteria were used during the geometry optimizations. Initial geometry coordinates were taken from the crystal structure in the quintet state. Optimized structures were confirmed to be local minima (no imaginary frequencies for both cases). Selected theoretically derived metric parameters are shown in Table 2. Molecular orbitals were generated using Avogadro²⁵ (an open-source molecular builder and visualization tool, Version 1.1.0. <http://avogadro.openmolecules.net/>).

X-ray crystallography

Compound **1** was crystallized from a dichloromethane/ether solution. A colorless prism crystal with dimensions 0.224 × 0.185 × 0.064 mm was mounted on a Nylon loop using a very small amount of paratone oil. Data were collected using a Bruker CCD based diffractometer equipped with an Oxford Cryostream low-temperature apparatus operating at 173 K. Data were measured using omega and phi scans of 0.5° per frame for 10 s. The total number of images was based on results from the program COSMO²⁶ where redundancy was expected to be 4.0 and completeness of 100% out to 0.83 Å. Cell parameters were retrieved using APEX II software²⁷ and refined using SAINT on all observed reflections. Data reduction was performed using the SAINT software,²⁸ which corrects for Lp. Scaling and absorption corrections were applied using SADABS²⁹ multi-scan technique, supplied by George Sheldrick. The structures are solved by the direct method using the SHELXS-97 program and



refined by least squares method on F^2 , SHELXL-2014,³⁰ which are incorporated in OLEX2.³¹ All non-hydrogen atoms were refined anisotropically. Hydrogen atoms were calculated by geometrical methods and refined as a riding model. The crystal used for the diffraction study showed no decomposition during data collection. All drawings are done at 50% ellipsoids.

Results and discussion

Synthesis

[Fe(TrIm)₄(OTf)₂] (**1**) was synthesized in good yield at room temperature under dry nitrogen conditions by reacting 4 equiv. 1-tritylimidazole (TrIm) with Fe(OTf)₂·2CH₃CN in dichloromethane. Suitable crystals of **1** for X-ray studies were grown from dichloromethane/ether.

Crystallography

The crystal structure of **1** was determined. Crystallographic parameters are shown in Table 1 while selected bond distances and angles are given in Table 2. The structure of **1** was solved in the $P\bar{1}$ space group. The structure reveals iron(II) in distorted octahedral geometry with ligands derived from two triflate oxygens and four 3-imidazole nitrogens (Fig. 1). The Fe–N_{TrIm} distances are 2.1823(15) Å (Fe1–N1) and 2.1807(15) Å (Fe1–N1A) which is in the range of other known high spin Fe^{II}–N_{TrIm} compounds (2.103–2.272 Å).^{32–38} The iron triflate Fe1–O1 distance is 2.1843(13) Å which is in the range of other known high spin Fe^{II}–O_{triflate} bond distances (2.025–2.211 Å).^{20,39–50} Hydrogen bonding (D⋯A: 2.995 Å) is observed between an imidazole hydrogen located on C1A and a coordinated triflate oxygen (O1). Close contacts between a triflate fluorines and phenyl carbons (F3⋯C7': 3.320 Å; F3⋯C8': 3.322 Å; F1⋯C19' 3.133 Å; F1⋯C16': 3.295 Å; F1⋯C18':

3.120 Å) are observed. Each iron is well isolated from other irons with the closest distance between iron centers being 13.234 Å. No π – π stacking is observed among aromatic groups and all phenyl rings are approximately orthogonal within each trityl group.

Magnetic susceptibility

A powdered sample of **1** was characterized using variable temperature magnetic susceptibility (χ_M). The measurements were taken between 2–300 K with an applied field of 0.6 T. At 300 K $\chi_M T$ has a value of 3.73 cm³ mol^{−1} K (Fig. 2) consistent with high spin iron(II) ($S = 2$), 3.25–4.06 cm³ mol^{−1} K.⁵¹ As the temperature is lowered, the $\chi_M T$ value begins to decrease at 50 K quickly towards 1.17 cm³ mol^{−1} K at 2 K. This trend is likely due to zero-field splitting of high spin iron(II). This behavior is consistent with low symmetry at the iron center.^{52–54} The data was simulated using the program julX²¹ which gave a very good fit using $g = 2.24$ and $D = -0.80$ cm^{−1} which is typical for distorted octahedral geometry.⁵⁵

DFT studies

Theoretical calculation (DFT) were performed on **1** in the gas phase using X-ray coordinates as the starting point to optimize

Table 2 Selected bond lengths (Å) and angles (°) for [Fe(TrIm)₄(OTf)₂]·2Et₂O (1·2Et₂O). Calculated values are in brackets

Distances			
Fe1–O1	2.1843(13) [2.159]	N1–C2	1.374(2) [1.372]
Fe1–N1	2.1823(15) [2.173]	N2–C1	1.355(2) [1.358]
Fe1–N1A	2.1807(15) [2.178]	N2–C3	1.375(2) [1.381]
N1–C1	1.318(2) [1.319]	N2–C4	1.503(2) [1.487]
Angles			
N1–Fe1–N1A	93.13(6) [92.87]	N1–Fe1–O1'	86.93(5) [91.56]
N1–Fe1–N1'	180.00(8) [180.00]	N1A–Fe1–O1	95.24(6) [83.88]
N1–Fe1–N1A'	86.87(6) [87.13]	N1A–Fe1–O1'	84.76(6) [86.12]
N1A–Fe1–N1'	86.87(6) [87.15]	N1'–Fe1–O1	86.93(5) [91.56]
N1A–Fe1–N1A'	180.00(9) [180.00]	N1'–Fe1–O1'	93.07(5) [88.44]
N1'–Fe1–N1A'	93.13(6) [92.85]	N1A'–Fe1–O1	84.76(6) [86.12]
N1–Fe1–O1	93.07(5) [88.44]	N1A'–Fe1–O1'	95.24(6) [83.88]

Table 1 Crystallographic parameters for [Fe(TrIm)₄(OTf)₂]·2Et₂O (1·2Et₂O)

1·2Et ₂ O	
Empirical formula	C ₉₈ H ₉₂ N ₈ O ₈ F ₆ S ₂ Fe
Formula weight	1743.76
Temperature/K	173.15
Wavelength (Å)	0.7103
Crystal system	Triclinic
Space group	$P\bar{1}$
$a/\text{Å}$	13.2342(7)
$b/\text{Å}$	13.5131(7)
$c/\text{Å}$	13.7025(7)
$\alpha/^\circ$	98.0060(10)
$\beta/^\circ$	116.1990(10)
$\gamma/^\circ$	93.3490(10)
Volume/Å ³	2157.39(19)
Z	1
$D_{\text{calc}}/\text{Mg m}^{-3}$	1.342
μ/mm^{-1}	0.300
Goodness-of-fit on F^2	1.035
Final R indexes [$I > 2\sigma(I)$]	$R_1 = 0.0384$, $wR_2 = 0.0948$
Final R indexes [all data]	$R_1 = 0.0508$, $wR_2 = 0.1032$

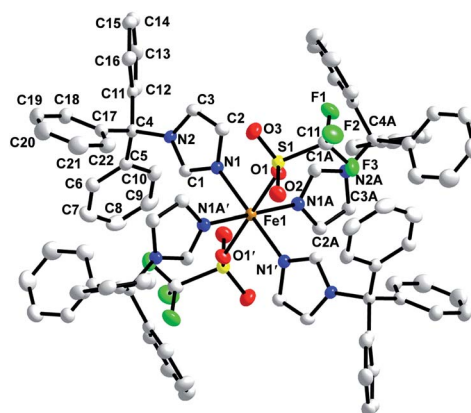


Fig. 1 X-ray crystal structure of [Fe(TrIm)₄(OTf)₂] (**1**) with thermal ellipsoids drawn at the 50% probability level.



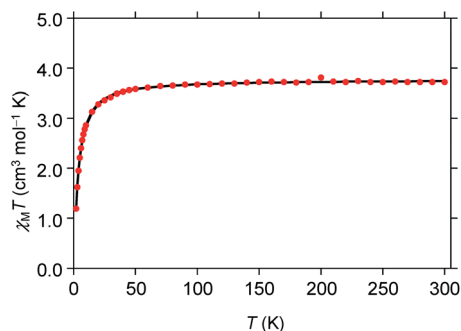


Fig. 2 Temperature-dependent molar magnetic susceptibility ($\chi_M T$) for $[\text{Fe}(\text{TrIm})_4(\text{OTf})_2]$ (**1**) at $H = 0.6$ T.

the ground state structure. The energies were obtained using PBE0/6-31G(d). The calculated metric parameters for the optimized structure were compared to those empirically determined (Table 2). The largest difference between the calculated and experimental bond distances is 0.025 Å (Fe1–O1), while the bond angles are in good agreement. Despite slight differences, the calculated structure is very close to the experimental structure and therefore the electronic properties can be confidently surmised. The calculated Mulliken charge value on the iron(II) of **1** is +1.35 consistent with high spin iron(II).⁵⁶ Fig. 3 illustrates selected molecular orbitals: HOMO, HOMO–1, LUMO, and LUMO+1 for **1**. It is found that the highest occupied molecular orbital (HOMO, β -412) is largely distributed over the d_{xz}/d_{yz} orbital while the HOMO–1 (α -416) is associated with the $d_{x^2-y^2}$

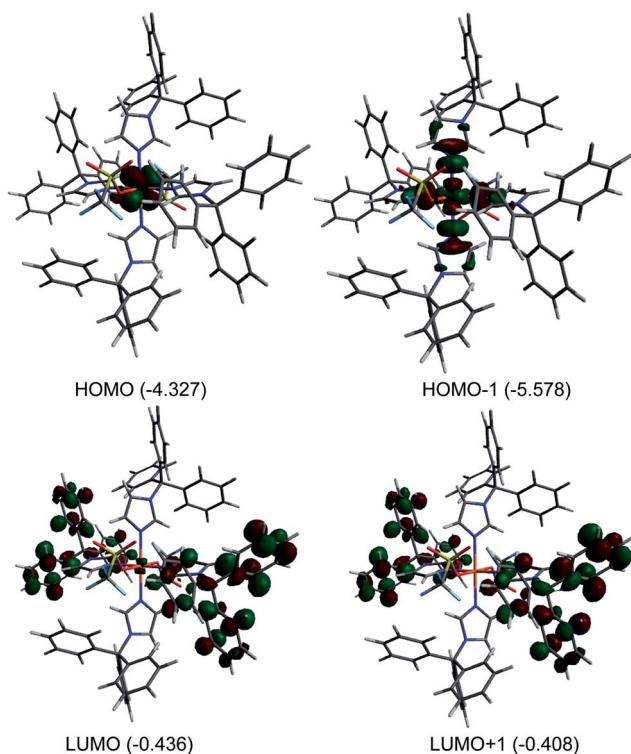


Fig. 3 Plots of molecular orbitals: HOMO–1, HOMO, LUMO, and LUMO+1 for $[\text{Fe}(\text{TrIm})_4(\text{OTf})_2]$ (**1**). Orbital energies (eV) are indicated.

and nitrogen sp^2 orbital in an antibonding fashion. The lowest unoccupied molecular orbital (LUMO, β -413) is primarily distributed over ring unhybridized p orbitals and to a lesser extent the d_{xz}/d_{yz} orbital while the LUMO+1 (β -414) consists of unhybridized p orbitals. The HOMO–LUMO gap for **1** is large (3.891 eV) indicating high stability.^{57,58}

Reactivity studies

In our efforts to mimic CCD reactivity we reacted **1** with 1 equiv. *cis*-stilbene in the presence of dioxygen. It was anticipated that formation of benzaldehyde would be observed should biomimetic activity be present. The initial colorless solution immediately turned pale yellow when dioxygen was bubbled into the solution (Fig. 4(a)). No temperature sensitive intermediates were observed during oxygenation. Analysis of the reaction mixture (after addition of acid and extraction with CH_2Cl_2) by GCMS, however, indicated that the *cis*-stilbene was unaffected and that no other oxygenated products were detected. We then decided to use H_2O_2 , a stronger oxidizing agent, for this reaction. Addition of 20 equiv. hydrogen peroxide (78%) to **1** in dichloromethane in the presence of *cis*-stilbene at 25 °C yielded

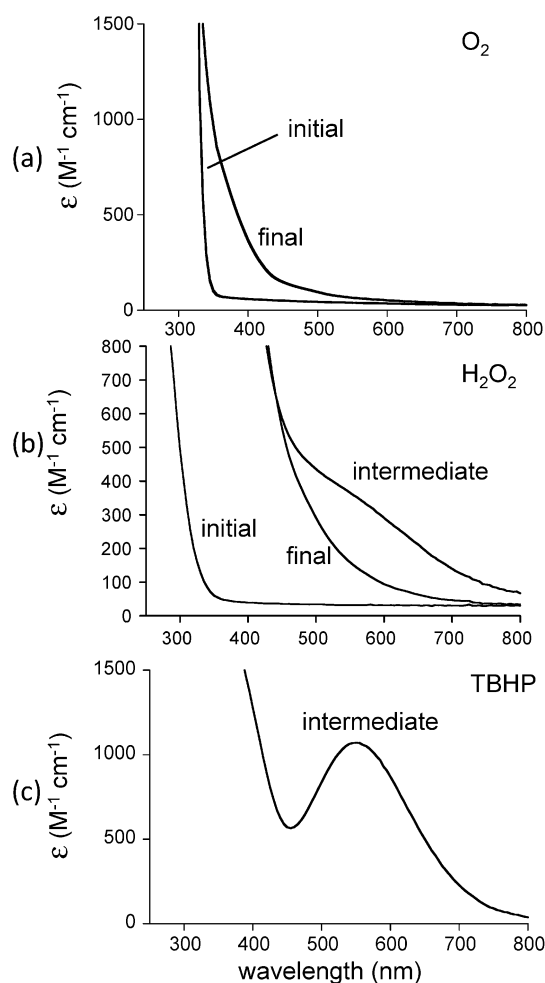
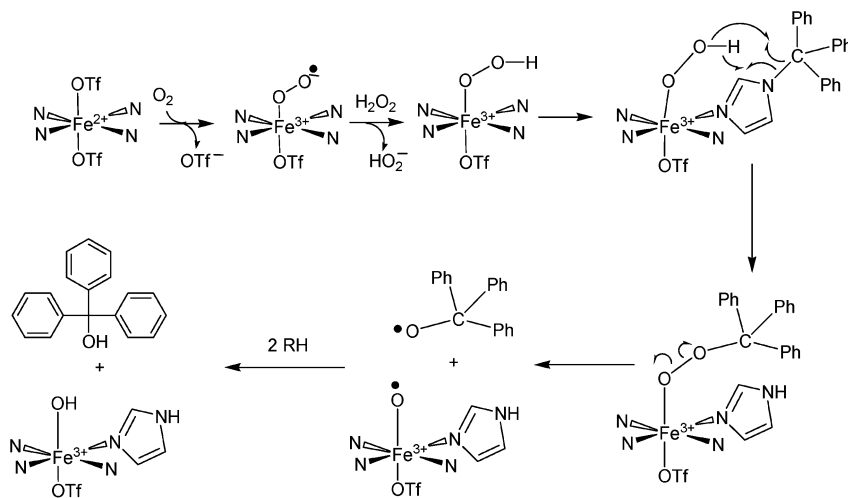


Fig. 4 Reaction of $[\text{Fe}(\text{TrIm})_4(\text{OTf})_2]$ (**1**) in CH_2Cl_2 with (a) O_2 in the presence of *cis*-stilbene at 25 °C, (b) 20 equiv. H_2O_2 at –30 °C, and (c) $t\text{BuOOH}$ (TBHP) at –30 °C.





Scheme 2 Proposed mechanism for oxidation of the trityl group attached to $[\text{Fe}(\text{TrIm})_4(\text{OTf})_2]$ (**1**) when H_2O_2 is added.

a dark purple-brown solution which gradually turned orange. Analysis of the reaction mixture by GCMS in this case indicated that triphenylmethanol (0.5 equiv.) had formed with no *cis*-stilbene-derived products. A control reaction (lacking iron) resulted in no oxidation products. This result indicates the preference for ligand oxidation over *cis*-stilbene oxidation which is likely due to the close juxtapositioning of the ligand to the oxygenated intermediate compared to *cis*-stilbene. In our attempts to stabilize the purple-brown intermediate, we conducted a low temperature UV-vis experiment. A solution of **1** in dichloromethane was first cooled to -30°C then 20 equiv. H_2O_2 was introduced to generate the intermediate species with a shoulder at $\sim 590\text{ nm}$ (Fig. 4(b)). The absorption intensity of shoulder began to decrease almost immediately and was gone after several minutes.

To explain these results, we have proposed a reaction mechanism which is shown in Scheme 2. In this mechanism, the iron(II) center in **1** first becomes oxidized to Fe(III) (due to the presence of ambient oxygen and possibly forming an $\text{LFe}^{\text{III}}-\text{O}_2^-$ species⁵⁹) and reacts with the hydrogen peroxide to form what is believed, based on spectroscopic characteristics, to be a $\text{LFe}^{\text{III}}-\text{OOH}$ species.^{60–62} The protonated superoxide species would be displaced. Since triphenylmethyl is known to form the stable triphenylmethyl radical ($\text{Ph}_3\text{C}^\bullet$),⁶³ we hypothesize that this species exchanges with the hydroperoxo group and forms $\text{LFe}^{\text{III}}-\text{OOCPh}_3$ which along with $\text{LFe}^{\text{III}}-\text{OOH}$ could account for the dark brown-purple intermediate species. Homolytic scission of the O–O bond in $\text{LFe}^{\text{III}}-\text{OOCPh}_3$ could produce $\text{LFe}^{\text{III}}-\text{O}^\bullet$ and $^\bullet\text{OOCPh}_3$ radicals. The $^\bullet\text{OOCPh}_3$ species could abstract an H atom from an RH species (solvent or ligand) to form triphenylmethanol while the $\text{LFe}^{\text{III}}-\text{O}^\bullet$ species could additionally abstract an H atom to form $\text{LFe}^{\text{III}}-\text{OH}$. It should also be noted that a long-lived purple species ($\lambda_{\text{max}} = 550\text{ nm}$, $\epsilon = 1070\text{ M}^{-1}\text{ cm}^{-1}$) is generated by reacting **1** with $^t\text{BuOOH}$ at -30°C (Fig. 4(c)). This intermediate is expected to be $\text{Fe}-\text{OO}^t\text{Bu}$.^{59–61} Upon decomposition, Ph_3COCH_3 is the primary product detected presumably formed by combination of $^\bullet\text{OOCPh}$ and $^\bullet\text{CH}_3$ (derived from decomposition of $^t\text{BuO}^\bullet$).⁶⁴

Conclusion

In this study we have utilized 1-tritylimidazole (TrIm) to model enzymatic 4His nonheme sites. Reaction of 4 equiv. TrIm with $\text{Fe}(\text{OTf})_2 \cdot 2\text{MeCN}$ yields $[\text{Fe}(\text{TrIm})_4(\text{OTf})_2]$ (**1**) in good yield. The four imidazole ligands are bonded in a square plane while triflate oxygens are in the apical positions. Variable temperature magnetic susceptibility measurements revealed **1** to be high spin between 20 K and 300 K. Simulations of the data yielded $g = 2.24$ and $D = -0.80\text{ cm}^{-1}$. DFT studies reproduced metric parameters for **1** and the HOMO–LUMO gap energy (3.89 eV) indicated a high degree of stability. Reactivity studies of **1** with hydrogen peroxide or $^t\text{BuOOH}$ indicate that an oxygenated intermediate is formed which subsequently decomposes to yield oxidized trityl substituent products.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

FAC acknowledges the receipt of an OU-REF grant. JL acknowledges a graduate fellowship from OU. TAH acknowledges an OU Provost Award. We thank Prof. M. M. Szczyński for assistance with the DFT calculations. NIH Grant No. R15GM112395 and NSF Grant No. CHE-0748607 and CHE-0821487 are gratefully acknowledged.

Notes and references

- X. W. Sui, M. Golczak, J. Y. Zhang, K. A. Kleinberg, J. von Lintig, K. Palczewski and P. D. Kiser, *J. Biol. Chem.*, 2015, **290**, 30212–30223.
- B. Zhang, C. Liu, Y. Q. Wang, X. Yao, F. Wang, J. S. Wu, G. J. King and K. D. Liu, *New Phytol.*, 2015, **206**, 1513–1526.
- M. H. Walter and D. Strack, *Nat. Prod. Rep.*, 2011, **28**, 663–692.



- 4 S. A. Baba, D. Jain, N. Abbas and N. Ashraf, *J. Plant Physiol.*, 2015, **189**, 114–125.
- 5 V. A. Tu, A. Kaga, K. H. Gericke, N. Watanabe, T. Narumi, M. Toda, B. Brueckner, S. Baldermann and N. Mase, *J. Org. Chem.*, 2014, **79**, 6808–6815.
- 6 M. Bruno, P. Beyer and S. Al-Babili, *Arch. Biochem. Biophys.*, 2015, **572**, 126–133.
- 7 L. H. Liu, Z. Y. Shao, M. Zhang and Q. M. Wang, *Mol. Plant*, 2015, **8**, 28–39.
- 8 J. Amengual, M. A. K. Widjaja-Adhi, S. Rodriguez-Santiago, S. Hessel, M. Golczak, K. Palczewski and J. von Lintig, *J. Biol. Chem.*, 2013, **288**, 34081–34096.
- 9 S. Raghuvanshi, V. Reed, W. S. Blaner and E. H. Harrison, *Arch. Biochem. Biophys.*, 2015, **572**, 19–27.
- 10 M. Yahyaa, A. Berim, T. Isaacson, S. Marzouk, E. Bar, R. Davidoyich-Rikanati, E. Lewinsohn and M. Ibdah, *J. Agric. Food Chem.*, 2015, **63**, 8275–8282.
- 11 S. Holger, R. Kurtzer, W. Eisenreich and W. Schwab, *J. Biol. Chem.*, 2006, **281**, 9845–9851.
- 12 O. P. Ernst, D. T. Lodowski, M. Elstner, P. Hegemann, L. S. Brown and H. Kandori, *Chem. Rev.*, 2014, **114**, 126–163.
- 13 S. A. J. Messing, S. B. Gabelli, I. Echeverria, J. T. Vogel, J. C. Guan, B. C. Tan, H. J. Klee, D. R. McCarty and L. M. Amzel, *Plant Cell*, 2010, **22**, 2970–2980.
- 14 C. R. D. Lancaster, A. Kroger, M. Auer and H. Michel, *Nature*, 1999, **402**, 377–385.
- 15 G. Katona, P. Carpentier, V. Niviere, P. Amara, V. Adam, J. Ohana, N. Tsanov and D. Bourgeois, *Science*, 2007, **316**, 449–453.
- 16 J. P. McEvoy and G. W. Brudvig, *Biochemistry*, 2008, **47**, 13394–13403.
- 17 X. W. Sui, P. D. Kiser, T. Che, P. R. Carey, M. Golczak, W. X. Shi, J. von Lintig and K. Palczewski, *J. Biol. Chem.*, 2014, **289**, 12286–12299.
- 18 D. P. Kloer, S. Ruch, S. Al-Babili, P. Beyer and G. E. Schulz, *Science*, 2005, **308**, 267–269.
- 19 D. Babino, M. Golczak, P. D. Kiser, A. Wyss, K. Palczewski and J. von Lintig, *ACS Chem. Biol.*, 2016, **11**, 1049–1057.
- 20 K. S. Hagen, *Inorg. Chem.*, 2000, **39**, 5867–5869.
- 21 julX, http://www.mpiibac.mpg.de/bac/index_en.php/logins/bill/julX_en.php.
- 22 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. A. Montgomery Jr, J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, T. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, J. M. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, O. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski and D. J. Fox, *Gaussian09, Revision C.01*, Gaussian Inc., Wallingford CT, 2010.
- 23 C. Adamo and V. Barone, *J. Chem. Phys.*, 1999, **110**, 6158–6170.
- 24 G. A. Petersson and M. A. Al-Laham, *J. Chem. Phys.*, 1991, **94**, 6081–6090.
- 25 M. Hanwell, D. Curtis, D. Lonie, T. Vandermeersch, E. Zurek and G. Hutchison, *J. Cheminf.*, 2012, **4**, 1–17.
- 26 *COSMO V1.61, Software for the CCD Detector Systems for Determining Data Collection Parameters*, Bruker Analytical X-ray Systems, Madison, WI, 2009.
- 27 *APEX2 V2010.11-3. Software for the CCD Detector System*, Bruker Analytical X-ray Systems, Madison, WI, 2010.
- 28 *SAINT V 7.68A Software for the Integration of CCD Detector System*, Bruker Analytical X-ray Systems, Madison, WI, 2010.
- 29 SADABS V2008/2 Program for absorption corrections using Bruker-AXS CCD based on the method of Robert Blessing; R. H. Blessing, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 1995, **51**, 33–38.
- 30 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2008, **64**, 112–122.
- 31 O. V. Dolomanov, L. J. Bourhis, R. J. Gildea, J. A. K. Howard and H. Puschmann, *J. Appl. Crystallogr.*, 2009, **42**, 339–341.
- 32 N. Brefuel, C. Duhayon, S. Shova and J. P. Tuchagues, *Chem. Commun.*, 2007, 5223–5225.
- 33 S. W. Jin, W. Z. Chen and H. Y. Qiu, *Cryst. Growth Des.*, 2007, **7**, 2071–2079.
- 34 I. Morgenstern-Badarau, F. Lambert, A. Deroche, M. Cesario, J. Guilhem, B. Keita and L. Nadjo, *Inorg. Chim. Acta*, 1998, **276**, 234–241.
- 35 A. K. Patra, K. S. Dube, G. C. Papaefthymiou, J. Conradi, A. Ghosh and T. C. Harrop, *Inorg. Chem.*, 2010, **49**, 2032–2034.
- 36 U. Ray, D. Banerjee, J. C. Liou, C. N. Lin, T. H. Lu and C. Sinha, *Inorg. Chim. Acta*, 2005, **358**, 1019–1026.
- 37 W. Liu, G. Zhang, X. Li, B. L. Wu and H. Y. Zhang, *Acta Crystallogr., Sect. E: Struct. Rep. Online*, 2009, **65**, M938–U1085.
- 38 E. Ohshima, K. Yoshida, K. Sugiyama and H. Uekusa, *Acta Crystallogr., Sect. E: Struct. Rep. Online*, 2012, **68**, 1093–1094.
- 39 J. Li, A. Banerjee, P. L. Pawlak, W. W. Brennessel and F. A. Chavez, *Inorg. Chem.*, 2014, **53**, 5414–5416.
- 40 L. Benhamou, A. Thibon, L. BreLOT, M. Lachkar and D. Mandon, *Dalton Trans.*, 2012, **41**, 14369–14380.
- 41 M. Grau, A. Kyriacou, F. C. Martinez, I. M. de Wispelaere, A. J. P. White and G. J. P. Britovsek, *Dalton Trans.*, 2014, **43**, 17108–17119.
- 42 G. Panchbhai, W. M. Singh, B. Das, R. T. Jane and A. Thapper, *Eur. J. Inorg. Chem.*, 2016, 3262–3268.
- 43 K. Umehara, S. Kuwata and T. Ikariya, *Inorg. Chim. Acta*, 2014, **413**, 136–142.
- 44 J. England, R. Gondhia, L. Bigorra-Lopez, A. R. Petersen, A. J. P. White and G. J. P. Britovsek, *Dalton Trans.*, 2009, 5319–5334.
- 45 S. Gosiewska, J. J. L. M. Cornelissen, M. Lutz, A. L. Spek, G. van Koten and R. J. M. K. Gebbink, *Inorg. Chem.*, 2006, **45**, 4214–4227.



- 46 G. J. P. Britovsek, J. England and A. J. P. White, *Dalton Trans.*, 2006, 1399–1408.
- 47 N. J. Hardman, X. G. Fang, B. L. Scott, R. J. Wright, R. L. Martin and G. J. Kubas, *Inorg. Chem.*, 2005, **44**, 8306–8316.
- 48 G. J. P. Britovsek, J. England and A. J. P. White, *Inorg. Chem.*, 2005, **44**, 8125–8134.
- 49 D. W. Blakesley, S. C. Payne and K. S. Hagen, *Inorg. Chem.*, 2000, **39**, 1979–1989.
- 50 J. Y. Chen and L. K. Woo, *Polyhedron*, 1999, **18**, 825–830.
- 51 C. J. O'Connor, *Prog. Inorg. Chem.*, 1982, **29**, 203–283.
- 52 B. N. Figgis and M. A. Hitchman, *Ligand Field Theory and Its Applications*, Wiley-VCH, New York, 2000.
- 53 T. Fujinami, K. Nishi, N. Matsumoto, S. Iijima, M. A. Halcrow, Y. Sunatsuki and M. Kojima, *Dalton Trans.*, 2011, **40**, 1230–12309.
- 54 Y. Sunatsuki, R. Kawamoto, K. Fujita, H. Maruyama, T. Suzuki, H. Ishida, M. Kojima, S. Iijima and N. Matsumoto, *Inorg. Chem.*, 2009, **48**, 8784–8795.
- 55 R. Boca, *Coord. Chem. Rev.*, 2004, **248**, 757–815.
- 56 G. J. Long, S. Tanase, F. Remacle, G. Periyasamy and F. Grandjean, *Inorg. Chem.*, 2009, **48**, 8173–8179.
- 57 N. D. Sousa, R. B. de Lima, A. L. P. Silva, A. A. Tanaka, A. B. F. da Silva and J. D. G. Varela, *Comput. Theor. Chem.*, 2015, **1054**, 93–99.
- 58 A. L. P. Silva, L. F. de Almeida, A. L. B. Marques, H. R. Costa, A. A. Tanaka, A. B. F. da Silva and J. D. G. Varela, *J. Mol. Model.*, 2014, **20**, 2131.
- 59 M. Costas, M. P. Mehn, M. P. Jensen and L. Que Jr, *Chem. Rev.*, 2004, **104**, 939–986.
- 60 F. Namuswe, G. D. Kasper, A. A. Sarjeant, T. Hayashi, C. M. Krest, M. T. Green, P. Moenne-Loccoz and D. P. Goldberg, *J. Am. Chem. Soc.*, 2008, **130**, 14189–14200.
- 61 M. P. Jensen, M. Costas, R. Y. Ho, J. Kaizer, A. Mairata i Payeras, E. Munck, L. Que Jr, J. U. Rohde and A. Stubna, *J. Am. Chem. Soc.*, 2005, **127**, 10512–10525.
- 62 G. Roelfes, V. Vrajmasu, K. Chen, R. Y. Ho, J. U. Rohde, C. Zondervan, R. M. La Crois, E. P. Schudde, M. Lutz, A. L. Spek, R. Hage, B. L. Feringa, E. Munck and L. Que Jr, *Inorg. Chem.*, 2003, **42**, 2639–2653.
- 63 W. P. Nuemann, W. Uzick and A. K. Zarkadis, *J. Am. Chem. Soc.*, 1986, **108**, 3762–3770.
- 64 M. Finn, R. Friedline, N. K. Suleman, C. J. Wohl and J. M. Tanko, *J. Am. Chem. Soc.*, 2004, **126**, 7578–7584.

