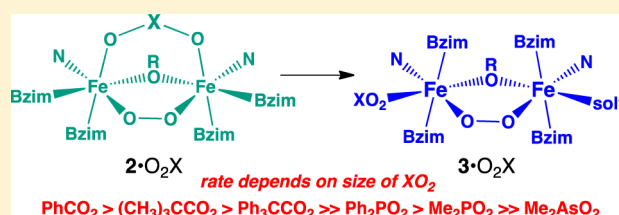


Factors Affecting the Carboxylate Shift Upon Formation of Nonheme Diiron-O₂ AdductsJonathan R. Frisch,[†] Ryan McDonnell,[‡] Elena V. Rybak-Akimova,^{*,‡} and Lawrence Que, Jr.^{*,†}[†]Department of Chemistry and Center for Metals in Biocatalysis, University of Minnesota, 207 Pleasant St. S.E., Minneapolis, Minnesota 55455, United States[‡]Department of Chemistry, Tufts University, 62 Talbot Ave., Medford, Massachusetts 02155, United States

S Supporting Information

ABSTRACT: Several [Fe^{II}₂(N-EtHPTB)(μ-O₂X)]²⁺ complexes (1·O₂X) have been synthesized, where N-EtHPTB is the anion of *N,N,N',N'*-tetrakis(2-benzimidazolylmethyl)-2-hydroxy-1,3-diaminopropane and O₂X is an oxyanion bridge. Crystal structures reveal five-coordinate (μ-alkoxo)diiron(II) cores. These diiron(II) complexes react with O₂ at low temperatures in CH₂Cl₂ (−90 °C) to form blue-green O₂ adducts that are best described as triply bridged (μ-η¹:η¹-peroxo)diiron(III) species (2·O₂X). With one exception, all 2·O₂X intermediates convert irreversibly to doubly bridged, blue (μ-η¹:η¹-peroxo)diiron(III) species (3·O₂X). Where possible, 2·O₂X and 3·O₂X intermediates were characterized using resonance Raman spectroscopy, showing respective ν_{O−O} values of ~850 and ~900 cm^{−1}. How the steric and electronic properties of O₂X affect conversion of 2·O₂X to 3·O₂X was examined. Stopped-flow analysis reveals that oxygenation kinetics of 1·O₂X is unaffected by the nature of O₂X, and for the first time, the benzoate analog of 2·O₂X (2·O₂CPh) is observed.



INTRODUCTION

Nonheme diiron enzymes have attracted great interest because they perform a wide variety of reactions despite having very similar active sites.^{1–8} For example, soluble methane monooxygenase inserts an oxygen atom into an alkane C–H bond; toluene and *o*-xylene monooxygenases, as well as phenol hydroxylase, insert oxygen atoms into aromatic C–H bonds; the R2 subunit of class I ribonucleotide reductases extracts a hydrogen atom, producing a stable radical; and, as its name implies, stearoyl acyl carrier protein Δ⁹-desaturase dehydrogenates a fatty acid hydrocarbon chain. In addition to having similar active site structures, these enzymes all share one other property. The putative catalytic cycles of these enzymes all contain peroxide-bridged diiron(III) complexes formed upon reduction of molecular oxygen. In some cases, these intermediates are stable enough to be trapped and characterized.^{9–17} While peroxo-diiron(III) species are endemic in these enzymes, their precise role in catalysis is poorly understood, even though they have been well-studied. Synthetic attempts to study peroxo-diiron(III) intermediates have resulted in a variety of biomimetic complexes.^{18–27} These compounds often form peroxide-bridged diiron(III) intermediates, especially when stabilized by carboxylate bridges. They are commonly reported as having a (μ-η¹:η¹-peroxo)diiron(III) motif.

In a previous publication,²⁸ we discussed how replacing the carboxylate bridges commonly employed in synthesis of these biomimetic complexes affected the behavior of the peroxo intermediates formed. Using the dinucleating ligand N-EtHPTB [anion of *N,N,N',N'*-tetrakis(2-benzimidazolylmethyl)-2-hydroxy-1,3-diaminopropane] combined with 2 equiv of

iron(II) and 1 equiv of either benzoate, diphenylphosphinate, or dimethylarsinate,²⁹ we demonstrated that two different (μ-η¹:η¹-peroxo)diiron(III) species can be formed upon reduction of dioxygen. A doubly bridged (μ-alkoxo)(μ-1,2-peroxo) intermediate was obtained from the benzoate-bridged diiron(II) precursor at −40 °C, in contrast to the triply bridged (μ-alkoxo)(μ-1,2-peroxo)(μ-1,3-dimethylarsinato) intermediate produced by the dimethylarsinate-bridged diiron(II) precursor. The diphenylphosphinate-bridged diiron(II) precursor was unique in that it formed a triply bridged peroxo intermediate as the initial kinetic product, which converted to a metastable doubly bridged peroxo intermediate by a shift of the phosphinate from a bridging mode to a terminal monodentate binding mode before peroxo decomposition, analogous to the “carboxylate shift” notion described by Lippard 20 years ago.³⁰ We concluded that the nature of the oxyanion bridge (O₂X) strongly influenced the nature and stability of any (μ-η¹:η¹-peroxo)diiron(III) species formed, but we felt further work was required before the effects of bridge differences could be clearly understood. This work is intended to address that issue.

To this end, we focused on three effects: (i) those produced by O₂X O···O bite distance differences, (ii) those produced by O₂X electronic differences, and (iii) those produced by O₂X steric differences. Examination required synthesis of several new diiron(II) complexes (1·O₂X) using N-EtHPTB and various O₂X ligands. We found that all of these new species reacted

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with O₂ in solution to produce blue-green (μ - η^1 : η^1 -peroxo)-diiron(III) intermediates ($2\cdot\text{O}_2\text{X}$) at -90°C . With one exception, they then converted to deep-blue (μ - η^1 : η^1 -peroxo)-diiron(III) species ($3\cdot\text{O}_2\text{X}$) before decaying to yellow products ($4\cdot\text{O}_2\text{X}$) at higher temperature. In most cases, addition of OPPh₃ to solutions of $2\cdot\text{O}_2\text{X}$ led to conversion to purple-blue species ($3'\cdot\text{O}_2\text{X}$) similar to $3\cdot\text{O}_2\text{X}$. In this paper, we report crystallographic details of three diiron(II) complexes, kinetic data on their reactions with dioxygen, and spectroscopic characterization of a variety of (μ - η^1 : η^1 -peroxo)diiron(III) intermediates. Implications of O₂X bite distances as well as electronic and steric differences are discussed.

EXPERIMENTAL SECTION

Materials and Syntheses. All reagents and solvents were purchased from commercial sources and were used as received, unless noted otherwise. The ligand N-EtHPTB was synthesized using a published procedure.³¹ Solvents were dried according to published procedures and distilled under Ar prior to use.³² The ¹⁸O₂ (97%) used in resonance Raman experiments was purchased from Cambridge Isotope Laboratories, Inc., Andover, MA. Preparation and handling of air sensitive materials were carried out under an inert atmosphere by using either standard Schlenk and vacuum line techniques or a glovebox. Elemental analyses were performed by Atlantic Microlab, Inc., Norcross, GA.

1-O₂CPh, 1-O₂PPh₂, and 1-O₂AsMe₂ were synthesized using published procedures.²⁸

1-O₂PMe₂. N-EtHPTB (157 mg, 0.217 mmol) was dissolved in MeOH (~10 mL) along with Et₃N (0.19 mL, 1.4 mmol). Dimethylphosphinic acid (21.6 mg, 0.230 mmol) was added and allowed to dissolve. Fe(OTf)₂·2MeCN³³ (189 mg, 0.434 mmol) was added, producing a yellow solution. After 5 min, NaBPh₄ (149 mg, 0.435 mmol) was added, resulting in immediate precipitation of a white powder. The solid was filtered and dried in vacuo. Recrystallization from MeCN and Et₂O produced colorless crystals, some suitable for X-ray diffraction structural analysis. Yield: 153 mg (80%). Anal. Calcd for [Fe₂(N-EtHPTB)(O₂PMe₂)](BPh₄)(OTf) (C₇₀H₇₃BF₃Fe₂N₁₀O₆PS): C, 60.27; H, 5.42; N, 10.04. Found: C, 59.82; H, 5.49; N, 10.38.

1-O₂P(OPh)₂. N-EtHPTB (176 mg, 0.243 mmol) was dissolved in MeOH (~10 mL) along with Et₃N (0.19 mL, 1.4 mmol). Diphenylphosphoric acid (67.7 mg, 0.271 mmol) was added and allowed to dissolve. Fe(OTf)₂·2MeCN (222 mg, 0.509 mmol) was added, producing a yellow solution. After 5 min, NaBPh₄ (173 mg, 0.506 mmol) was added, resulting in immediate precipitation of a white powder. The solid was filtered and dried in vacuo. Recrystallization from MeCN and Et₂O produced milky crystals, some suitable for X-ray diffraction structural analysis. Yield: 331 mg (79%). Anal. Calcd for [Fe₂(N-EtHPTB)(O₂P(OPh)₂)](BPh₄)₂ (C₁₀₃H₉₉B₂Fe₂N₁₀O₅P): C, 71.87; H, 5.80; N, 8.14. Found: C, 71.77; H, 5.94; N, 8.41.

1-O₂CCPh₃. N-EtHPTB (208 mg, 0.288 mmol) was dissolved in MeOH (~10 mL) along with Et₃N (0.19 mL, 1.4 mmol). Triphenylacetic acid (83.3 mg, 0.289 mmol) was added and allowed to dissolve. Fe(OTf)₂·2MeCN (257 mg, 0.589 mmol) was added, producing a yellow solution. After 5 min, NaBPh₄ (205 mg, 0.598 mmol) was added, resulting in immediate precipitation of a white powder. The solid was filtered and dried in vacuo. Recrystallization from MeCN and Et₂O produced colorless crystals. Yield: 370 mg (73%). Anal. Calcd for [Fe₂(N-EtHPTB)(O₂CCPh₃)](BPh₄)₂ (C₁₁₁H₁₀₄B₂Fe₂N₁₀O₃): C, 75.78; H, 5.96; N, 7.96. Found: C, 75.46; H, 5.95; N, 7.96.

1-O₂CCMe₃. N-EtHPTB (153 mg, 0.212 mmol) was dissolved in MeOH (~10 mL) along with Et₃N (0.19 mL, 1.4 mmol). Trimethylacetic acid (21.7 mg, 0.212 mmol) was added and allowed to dissolve. Fe(OTf)₂·2MeCN (186 mg, 0.426 mmol) was added, producing a yellow solution. After 5 min, NaBPh₄ (152 mg, 0.443 mmol) was added, resulting in immediate precipitation of a white

powder. The solid was filtered and dried in vacuo. Recrystallization from MeCN and Et₂O produced milky crystals. Yield: 130 mg (50%). Anal. Calcd for [Fe₂(N-EtHPTB)(O₂CCMe₃)](OTf)₂ (C₅₀H₅₈F₆Fe₂N₁₀O₉S₂): C, 48.71; H, 4.74; N, 11.36. Found: C, 48.91; H, 4.77; N, 11.39.

1-O₂CC₆H₂-3,4,5-(OMe)₃. N-EtHPTB (72.9 mg, 0.101 mmol) was dissolved in MeOH (~10 mL) along with Et₃N (0.077 mL, 0.56 mmol). 3,4,5-Trimethoxybenzoic acid (21.4 mg, 0.101 mmol) was added and allowed to dissolve. Fe(OTf)₂·2MeCN (94.2 mg, 0.216 mmol) was added, producing a yellow solution. After 5 min, NaBPh₄ (87.2 mg, 0.255 mmol) was added, resulting in immediate precipitation of a pale yellow powder. The solid was filtered and dried in vacuo. Recrystallization from MeCN and Et₂O produced yellow crystals, some suitable for X-ray diffraction structural analysis. Yield: 116 mg (74%). Anal. Calcd for [Fe₂(N-EtHPTB)(O₂CCH₂-3,4,5-(OMe)₃)](BPh₄)(OTf) (C₇₈H₈₀BF₃Fe₂N₁₀O₉S): C, 61.92; H, 5.33; N, 9.26. Found: C, 62.14; H, 5.55; N, 8.94.

1-O₂CC₆H₃-3,4-(OMe)₂. N-EtHPTB (100.0 mg, 0.138 mmol) was dissolved in MeOH (~10 mL) along with Et₃N (0.097 mL, 0.69 mmol). 3,4-Dimethoxybenzoic acid (25.1 mg, 0.138 mmol) was added and allowed to dissolve. Fe(OTf)₂·2MeCN (120.3 mg, 0.276 mmol) was added, producing a yellow solution. After 5 min, NaBPh₄ (196.5 mg, 0.574 mmol) was added, resulting in immediate precipitation of a pale yellow powder. The solid was filtered and dried in vacuo. Recrystallization from MeCN and Et₂O produced milky crystals. Yield: 162 mg (89%). Anal. Calcd for [Fe₂(N-EtHPTB)(O₂CC₆H₃-3,4-(OMe)₂)](OTf)₂ (C₅₄H₅₈F₆Fe₂N₁₀O₁₁S₂): C, 49.40; H, 4.45; N, 10.67. Found: C, 49.65; H, 4.44; N, 10.39.

1-O₂CC₆H₃-3,5-(OMe)₂. N-EtHPTB (146.1 mg, 0.202 mmol) was dissolved in MeOH (~10 mL) along with Et₃N (0.142 mL, 1.02 mmol). 3,5-Dimethoxybenzoic acid (37.0 mg, 0.203 mmol) was added and allowed to dissolve. Fe(OTf)₂·2MeCN (187.8 mg, 0.431 mmol) was added, producing a yellow solution. After 5 min, NaBPh₄ (190.6 mg, 0.557 mmol) was added, resulting in immediate precipitation of a pale green-yellow powder. The solid was filtered and dried in vacuo. Recrystallization from MeCN and Et₂O produced pale yellow crystals. Yield: 238.5 mg (80%). Anal. Calcd for [Fe₂(N-EtHPTB)(O₂CC₆H₃-3,5-(OMe)₂)](BPh₄)(OTf)₂ (C₇₇H₇₈BF₃Fe₂N₁₀O₈S): C, 62.36; H, 5.30; N, 9.44. Found: C, 62.56; H, 5.11; N, 9.33.

1-O₂CC₆H₄-4-OMe. N-EtHPTB (149.2 mg, 0.206 mmol) was dissolved in MeOH (~10 mL) along with Et₃N (0.144 mL, 1.03 mmol). 4-Methoxybenzoic acid (31.3 mg, 0.206 mmol) was added and allowed to dissolve. Fe(OTf)₂·2MeCN (189.2 mg, 0.433 mmol) was added, producing a yellow solution. After 5 min, NaBPh₄ (205.4 mg, 0.600 mmol) was added, resulting in immediate precipitation of a pale green-yellow powder. The solid was filtered and dried in vacuo. Recrystallization from MeCN and Et₂O produced milky crystals. Yield: 245.4 mg (93%). Anal. Calcd for [Fe₂(N-EtHPTB)(O₂CC₆H₄-4-OMe)](OTf)₂ (C₅₃H₅₆F₆Fe₂N₁₀O₁₀S₂): C, 49.62; H, 4.40; N, 10.92. Found: C, 49.41; H, 4.51; N, 11.08.

Physical Methods. UV-vis spectra were recorded on a Hewlett-Packard 8453 diode array spectrophotometer (2-nm resolution) equipped with an Unisoku Scientific Instruments cryostat (Osaka, Japan). Resonance Raman spectra were collected on an ACTON AM-506M3 monochromator with a Princeton LN/CCD data collection system using a Spectra-Physics Model 2060 krypton laser. Low-temperature spectra of the peroxo intermediates in CH₂Cl₂ and MeCN were obtained at 77 K using a 135° backscattering geometry. Samples were frozen onto a gold-plated copper coldfinger in thermal contact with a Dewar flask containing liquid nitrogen. Raman frequencies were referenced to the features of indene. Slits were set for a band-pass of 4 cm⁻¹ for all spectra.

Time-resolved spectra of rapid oxygenation reactions were acquired with a Hi-Tech Scientific (Salisbury, Wiltshire, UK) SF-43 multimixing anaerobic cryogenic stopped-flow instrument combined with either a monochromator (low intensity light irradiation of the sample) or a diode array rapid scanning unit (strong UV-vis irradiation of the sample), or with a TgK Scientific (formerly HiTech Scientific, Salisbury, Wiltshire, UK) SF-61DX2 cryogenic stopped-flow system equipped with a J&M diode array (Spectralytics). All manipulations

Table 1. Crystal Data and Structure Refinement for 1·O₂PMe₂(BPh₄)(OTf)·MeCN, 1·O₂P(OPh)₂(BPh₄)₂·MeCN, and (1·O₂CC₆H₂-3,4,5-(OMe)₃)₂(BPh₄)₂(OTf)₂·2MeCN

	[1·O ₂ PMe ₂](BPh ₄)(OTf)·MeCN	[1·O ₂ P(OPh) ₂](BPh ₄) ₂ ·MeCN	[1·O ₂ CC ₆ H ₂ -3,4,5-(OMe) ₃] ₂ (BPh ₄) ₂ (OTf) ₂ ·2MeCN
empirical formula	C ₇₂ H ₇₈ BF ₃ Fe ₂ N ₁₁ O ₆ PS	C ₁₀₅ H ₁₀₂ B ₂ Fe ₂ N ₁₁ O ₅ P	C ₁₆₀ H ₁₆₆ B ₂ F ₆ Fe ₄ N ₂₂ O ₁₈ S ₂
fw	1435.99	1762.27	3108.29
T (K)	173(2)	173(2)	173(2)
Mo Kα λ, Å	0.71073	0.71073	0.71073
space group	P2 ₁ /c	P2 ₁ /n	P $\bar{1}$
a (Å)	16.0709(9)	23.063(4)	12.576(4)
b (Å)	15.6360(9)	18.942(4)	19.153(7)
c (Å)	28.4703(15)	23.888(4)	36.053(12)
α (deg)	90	90	96.743(5)
β (deg)	103.5800(10)	99.205(3)	92.147(5)
γ (deg)	90	90	103.100(5)
V (Å ³)	6954.1(7)	10301(3)	8381(5)
Z	4	4	2
ρ (calc), Mg/m ³	1.372	1.136	1.232
abs coeff (mm ⁻¹)	0.539	0.352	0.437
R1 ^a	0.0398	0.0446	0.0713
wR2 ^b	0.0984	0.1271	0.1955

$$^a R1 = \sum ||F_o| - |F_c|| / \sum |F_o|, \quad ^b wR2 = [\sum [w(F_o^2 - F_c^2)^2] / \sum [w(F_o^2)^2]]^{1/2}.$$

Table 2. Selected Interatom Distances and Bond Angles for [Fe₂(N-EtHPTB)(O₂X)]²⁺

	1·O ₂ AsMe ₂ ^a	1·O ₂ PPh ₂ ^a	1·O ₂ PMe ₂	1·O ₂ P(OPh) ₂	1·O ₂ CC ₆ H ₂ -3,4,5-(OMe) ₃ ^b	1·O ₂ CPh ^a
	0.81	0.77	0.82	0.91	0.89	0.93
	τ_{ave}					
	Interatom Distances (Å)					
Fe1–O1	2.0145(13)	2.004(2)	2.0184(18)	2.0057(16)	1.967(3)	1.976(5)
Fe2–O1	2.0037(14)	1.992(2)	1.9995(18)	2.0031(15)	1.970(3)	1.964(5)
Fe1–N1	2.3366(16)	2.294(3)	2.323(2)	2.3141(18)	2.328(4)	2.316(6)
Fe2–N2	2.3892(16)	2.352(3)	2.372(2)	2.309(2)	2.314(4)	2.280(7)
Fe1–O2	1.9825(14)	2.008(2)	2.0037(19)	2.0562(16)	2.036(3)	2.057(5)
Fe2–O3	1.9863(14)	2.021(2)	2.0123(18)	2.0506(16)	2.025(3)	2.019(6)
Fe1–N3	2.1372(17)	2.086(3)	2.105(2)	2.0668(19)	2.045(4)	2.064(6)
Fe1–N5	2.1023(18)	2.115(3)	2.127(2)	2.0732(19)	2.046(4)	2.069(6)
Fe2–N7	2.1088(17)	2.075(3)	2.098(2)	2.0551(19)	2.063(4)	2.080(6)
Fe2–N9	2.0759(17)	2.098(3)	2.080(2)	2.075(2)	2.044(4)	2.064(6)
As1/P1/C44–O2	1.6740(15)	1.514(2)	1.509(2)	1.4762(17)	1.254(5)	1.264(9)
As1/P1/C44–O3	1.6776(14)	1.512(2)	1.517(2)	1.4784(18)	1.273(5)	1.253(9)
Fe1...Fe2	3.5357(5)	3.5405(10)	3.5364(6)	3.6211(7)	3.4879(13)	3.4749(31)
O2...O3	2.7991(21)	2.5600(32)	2.5545(28)	2.5520(24)	2.2253(46)	2.2251(74)
	Bond Angles (deg)					
Fe1–O1–Fe2	123.27(6)	124.76(11)	123.32(9)	129.19(8)	124.73(15)	123.8(2)
O1–Fe1–O2	105.77(6)	105.55(9)	102.50(8)	98.99(6)	98.27(13)	98.6(2)
O1–Fe2–O3	107.92(6)	99.44(9)	103.98(8)	98.93(7)	100.51(13)	101.5(2)
Fe1–O2–As1/P1/C44	127.24(8)	132.01(14)	131.47(12)	133.92(11)	138.1(3)	136.4(5)
Fe2–O3–As1/P1/C44	128.15(8)	138.88(15)	132.54(12)	133.24(10)	134.7(3)	133.9(5)
O2–As1/P1/C44–O3	113.26(7)	115.59(13)	115.17(12)	119.47(10)	123.4(4)	124.2(7)

^aData from ref 28. ^bThe unit cell of this compound contains virtual twins of 1·O₂CC₆H₂-3,4,5-(OMe)₃ produced through a pseudoinversion center. As there is no substantive difference between the two cations, values listed here are taken from a single cation.

with diiron(II) complexes and their solutions were done using an argon atmosphere glovebox, airtight syringes, and the anaerobic stopped-flow instrument to avoid contamination with air. Saturated solutions of O₂ in CH₂Cl₂ and CH₃CN were prepared by bubbling the dry O₂ gas for 20 min in a septum-closed cylinder with the solvent at a constant temperature (20 or 25 °C). The solubility of O₂ was accepted to be 5.8 mM in dichloromethane at 20 °C and 8.1 mM in acetonitrile at 25 °C.³⁴ Solutions of O₂ with smaller concentrations were prepared by diluting the O₂-saturated solution with argon-saturated solvent using graduated gastight syringes equipped with three-way valves. For the kinetic experiments, dioxygen was always taken in large excess so

that its concentration did not change significantly during the reaction with 1·O₂X. The solutions of 1·O₂X and O₂ were cooled to a preset temperature (±0.1 °C) in the stopped-flow instrument before mixing. Data analysis was performed with the IS-2 Rapid Kinetics Software (Hi-Tech Scientific) for kinetic traces at a single wavelength.

X-ray Crystallography. X-ray diffraction data were collected on a Bruker SMART platform CCD diffractometer at 173(2) K.³⁵ Preliminary sets of cell constants were calculated from reflections harvested from three sets of 20 frames. These initial sets of frames were oriented such that orthogonal wedges of reciprocal space were surveyed. The data collection was carried out using Mo Kα radiation

(graphite monochromator). Randomly oriented regions of reciprocal space were surveyed to the extent of one sphere and to a resolution of 0.84 Å. The intensity data were corrected for absorption and decay using SADABS.³⁶ Final cell constants were calculated after integration with SAINT.³⁷ The structures were solved and refined using SHELXL-97.³⁸ The space groups $P2_1/c$, $P2_1/n$, and $P\bar{1}$ were determined on the basis of systematic absences and intensity statistics. Direct-methods solutions were calculated, which provided most non-hydrogen atoms from the E -map. Full-matrix least-squares/difference Fourier cycles were performed, which located the remaining non-hydrogen atoms. All non-hydrogen atoms were refined with anisotropic displacement parameters. All hydrogen atoms were placed in ideal positions and refined as riding atoms with relative isotropic displacement parameters. The SQUEEZE function of the program PLATON³⁹ was used to remove 143 effective electrons in diffuse scattering from a volume of 1555.2 Å³ per cell of $1 \cdot \text{O}_2\text{P}(\text{O}^i\text{Ph})_2(\text{BPh}_4)_2 \cdot \text{MeCN}$ and 120 effective electrons in diffuse scattering from a volume of 1097.4 Å³ per cell of $(1 \cdot \text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3)_2(\text{BPh}_4)_2(\text{OTf})_2 \cdot 2\text{MeCN}$. Brief crystal data and intensity collection parameters for the crystalline complexes are shown in Table 1.

RESULTS

X-ray Crystallography. For this work, we synthesized diiron(II) complexes using a variety of O_2X ligands, namely O_2PMe_2 , $\text{O}_2\text{P}(\text{O}^i\text{Ph})_2$, and several aliphatic and aromatic carboxylates, to augment a list that includes O_2AsMe_2 , O_2PPh_2 , and benzoate. We were able to obtain crystal structures of three of these $1 \cdot \text{O}_2\text{X}$ compounds [$\text{O}_2\text{X} = \text{O}_2\text{PMe}_2$, $\text{O}_2\text{P}(\text{O}^i\text{Ph})_2$, and $\text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$] (Table 2). Like those of previously published complexes,²⁸ the new complexes possess two distorted trigonal bipyramidal iron(II) centers, with N-EtHPTB amine nitrogen atoms and oxygen atoms from the bridging O_2X moiety occupying the axial positions. The equatorial sites are occupied by two benzimidazole nitrogen atoms and the alkoxide oxygen atom (Figure 1). With few exceptions, the respective atoms in the first coordination sphere of each complex reflect approximately equal interatomic distances and angles. Differences of note include average τ values,⁴⁰ O_2X bite distances, O-X-O angles, and inter-iron distances. The τ values can be grouped into two sets with $\tau_{\text{ave}} \sim 0.8$ ($\text{O}_2\text{X} = \text{O}_2\text{AsMe}_2$, O_2PPh_2 , and O_2PMe_2) and $\tau_{\text{ave}} \sim 0.9$ [$\text{O}_2\text{X} = \text{O}_2\text{P}(\text{O}^i\text{Ph})_2$, $\text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$, and O_2CPh]. Comparing bite distances of all complexes, we see $\text{O} \cdots \text{O}$ distances of ~ 2.55 Å ($\text{X} = \text{P}$) and ~ 2.23 Å ($\text{X} = \text{C}$). The lone $\text{X} = \text{As}$ complex is in a class of its own with a bite distance of ~ 2.80 Å. It is also apparent that the O-X-O angle varies with the identity of the X atom ($\text{X} = \text{As}$, 113.26° ; $\text{X} = \text{P}$, 115.17° – 119.47° ; $\text{X} = \text{C}$, 123.4° – 124.2°). The O-X-O angle measurement correlates inversely with O_2X bite distance; the species with the longest $\text{O} \cdots \text{O}$ distance ($1 \cdot \text{O}_2\text{AsMe}_2$) has the most acute O-X-O angle, while the $\text{X} = \text{C}$ complexes have the shortest bite distances and largest O-X-O angles, with the $\text{X} = \text{P}$ compounds falling in between these extremes. This inverse correlation reflects variations in X-O bond lengths ($\text{As-O}_{\text{ave}} = 1.676$ Å, $\text{P-O}_{\text{ave}} = 1.501$ Å, $\text{C-O}_{\text{ave}} = 1.261$ Å). Even though the O-X-O angle decreases, concomitant lengthening of the X-O bonds results in a greater $\text{O} \cdots \text{O}$ distance.

In four of the six diferrous complexes crystallographically characterized, increased bite distance correlates to increased $\text{Fe} \cdots \text{Fe}$ distance. With the other two complexes, we observed that (i) $1 \cdot \text{O}_2\text{AsMe}_2$ has an $\text{O} \cdots \text{O}$ distance ~ 0.24 Å longer than any other complex, although its inter-iron distance is within 0.01 Å of those found in both $1 \cdot \text{O}_2\text{PPh}_2$ and $1 \cdot \text{O}_2\text{PMe}_2$. (ii) $1 \cdot \text{O}_2\text{P}(\text{O}^i\text{Ph})_2$ has an $\text{O} \cdots \text{O}$ distance on par with $1 \cdot \text{O}_2\text{PPh}_2$ and $1 \cdot \text{O}_2\text{PMe}_2$, yet its inter-iron distance (3.6211 Å) is the greatest

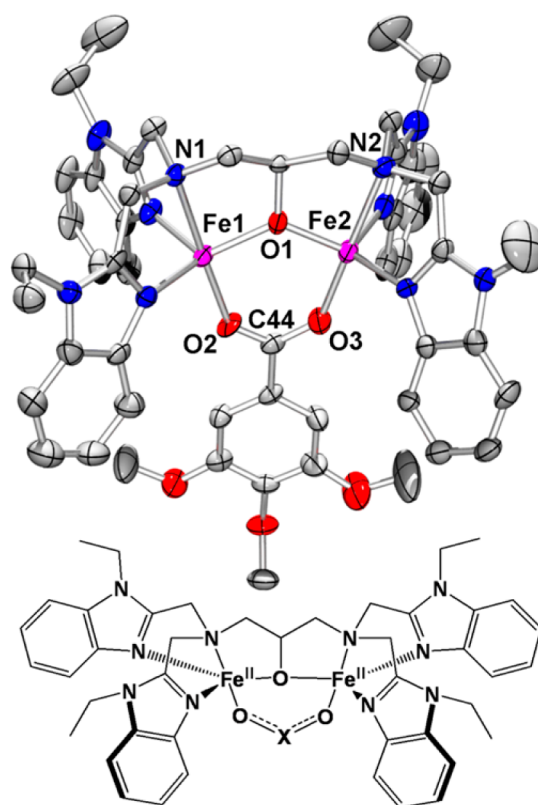


Figure 1. Crystal structure of the cation $1 \cdot \text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$ (50% ellipsoids) with hydrogen atoms removed. Generic cartoon of $1 \cdot \text{O}_2\text{X}$: $\text{O}_2\text{X} = \text{O}_2\text{AsMe}_2$ ($1 \cdot \text{O}_2\text{AsMe}_2$), O_2PPh_2 ($1 \cdot \text{O}_2\text{PPh}_2$), O_2PMe_2 ($1 \cdot \text{O}_2\text{PMe}_2$), $\text{O}_2\text{P}(\text{O}^i\text{Ph})_2$ ($1 \cdot \text{O}_2\text{P}(\text{O}^i\text{Ph})_2$), $\text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$ ($1 \cdot \text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$), and O_2CPh ($1 \cdot \text{O}_2\text{CPh}$).

of all the compounds, exceeding the distances found in $1 \cdot \text{O}_2\text{AsMe}_2$ (3.5357 Å) and the two other $\text{X} = \text{P}$ species (3.5405 and 3.5364 Å) by ~ 0.08 Å. This is greater than the ~ 0.06 Å difference observed between the $\text{Fe} \cdots \text{Fe}$ distances of the $\text{X} = \text{C}$ complexes (3.4879 and 3.4749 Å) and those with $\text{Fe} \cdots \text{Fe}$ distances of ~ 3.54 Å. The unusually long $\text{Fe} \cdots \text{Fe}$ distance in $1 \cdot \text{O}_2\text{P}(\text{O}^i\text{Ph})_2$ is most likely not due to packing effects, because a similar inter-iron distance of 3.649 Å was reported when $1 \cdot \text{O}_2\text{P}(\text{O}^i\text{Ph})_2$ crystallized in a different space group ($P\bar{1}$) with different anions (ClO_4).⁴¹

There are two crystallographically interesting observations regarding the compounds we report here and those examined in our previous work.²⁸ First, we note that $[\text{Fe}_2(\text{N-EtHPTB})(\text{O}_2\text{PMe}_2)](\text{BPh}_4)(\text{OTf})(\text{MeCN})$ and $[\text{Fe}_2(\text{N-EtHPTB})(\text{O}_2\text{AsMe}_2)](\text{BPh}_4)(\text{OTf})(\text{MeCN})$ are crystallographically isostructural. Both fall into the space group $P2_1/c$ with $Z = 4$ and share virtually identical unit cells with respective values for a , b , and c of 16.0709(9) vs 16.1151(12) Å, 15.6360(9) vs 15.7370(12) Å, and 28.4703(15) vs 28.473(2) Å. The α and γ angles are exactly 90° and the β angles are respectively $103.5800(10)^\circ$ and $103.4900(10)^\circ$. The respective cations, anions, and solvent molecules in each unit cell lie in essentially the same positions with the same orientations. The minor variations in unit cell parameters and atom positions arise from differences around the X atom of the O_2X bridging moieties. Small angle changes around X as well as X-C and X-O bond length variations produce minor differences in the cations that slightly alter the position of every other atom in the unit cell due to packing effects. The second observation is that the unit

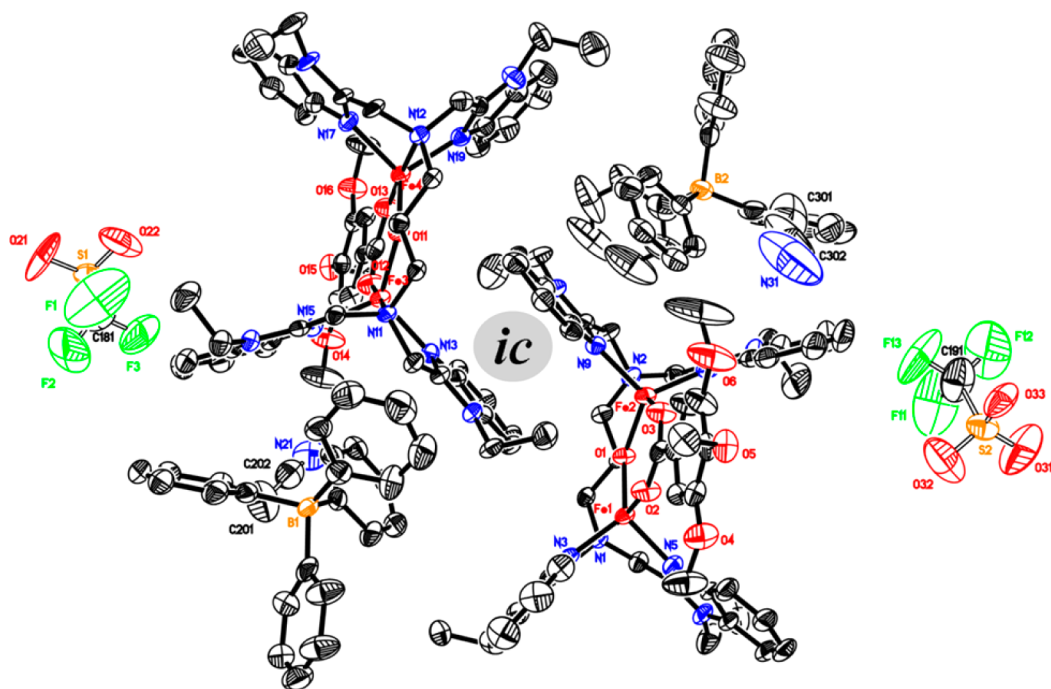


Figure 2. ORTEP diagram (50% ellipsoids) of the unit cell containing two molecules of $[\text{Fe}_2(\text{N-EtHPTB})(\text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3)]^{2+}$ and accompanying anions and solvent molecules (hydrogen atoms removed for clarity). There is a pseudoinversion center (*ic*) between the cations. Comparing the cations, we find that the “inversion” is almost true, whereas comparing the BPh_4 ions reveals differences in thermal ellipsoid size and minor differences in atom locations, indicating that the inversion center is not real.

cell containing the cation $[\text{Fe}_2(\text{N-EtHPTB})(\text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3)]^{2+}$ contains a pseudoinversion center lying between cations (Figure 2). This effectively doubles the number of unique elements per unit cell, thus doubling the size of the unit cell. The cation “inversion” is reasonably true, but the anions are farther away from the pseudoinversion center, resulting in greater distortion during “inversion”. Because the differences between the cations are not significant, we chose to focus on one cation rather than redundantly discussing what essentially amounts to duplicate cations.

Observation of Two Peroxo Intermediates. Upon reaction with O_2 , the virtually colorless solutions of $1\cdot\text{O}_2\text{X}$ produce blue-green O_2 adducts ($2\cdot\text{O}_2\text{X}$), which in all cases except two convert to deep blue species ($3\cdot\text{O}_2\text{X}$) before decaying to yellow final products ($4\cdot\text{O}_2\text{X}$). Previously, we reported the formation of $2\cdot\text{O}_2\text{PPh}_2$ and its conversion to $3\cdot\text{O}_2\text{PPh}_2$ at -30°C .²⁸ Corresponding studies with $\text{O}_2\text{X} = \text{benzoate}$ afforded only evidence for the formation of $3\cdot\text{O}_2\text{CPh}$ under these conditions.^{29,42,43} However, visible spectral evidence was obtained for the formation of both $2\cdot\text{O}_2\text{CR}$ and $3\cdot\text{O}_2\text{CR}$ by going to -90°C in CH_2Cl_2 , as shown in Figure 3 for $\text{O}_2\text{X} = \text{O}_2\text{CCPh}_3$ with respective λ_{max} values at 708 and 630 nm. In general, the visible chromophores of $2\cdot\text{O}_2\text{X}$ and $3\cdot\text{O}_2\text{X}$ fall in the ranges of 630–710 and 580–620 nm, respectively (Table 3), which have been previously assigned to peroxo-to-iron(III) charge transfer transitions for $\text{O}_2\text{X} = \text{benzoate}$, Ph_2PO_2 , and Me_2AsO_2 by resonance Raman spectroscopy.^{28,42}

Resonance Raman spectra for $2\cdot\text{O}_2\text{CCPh}_3$, $3\cdot\text{O}_2\text{CCPh}_3$, $2\cdot\text{O}_2\text{P(OPh)}_2$, and $3\cdot\text{O}_2\text{P(OPh)}_2$ shown in Figure 4 are representative of the species observed in this study. $^{16}\text{O}_2$ and $^{18}\text{O}_2$ labeling experiments reveal O–O stretching frequencies (Table 4) that by comparison to those previously published^{28,42} allow us to sort them into two clusters. The $2\cdot\text{O}_2\text{X}$ cluster exhibits $\nu_{\text{O-O}}$ values ranging from 839 to 851 cm^{-1} that

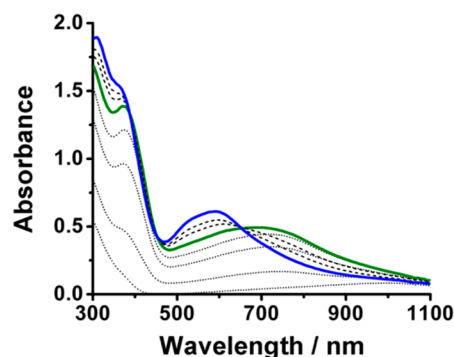
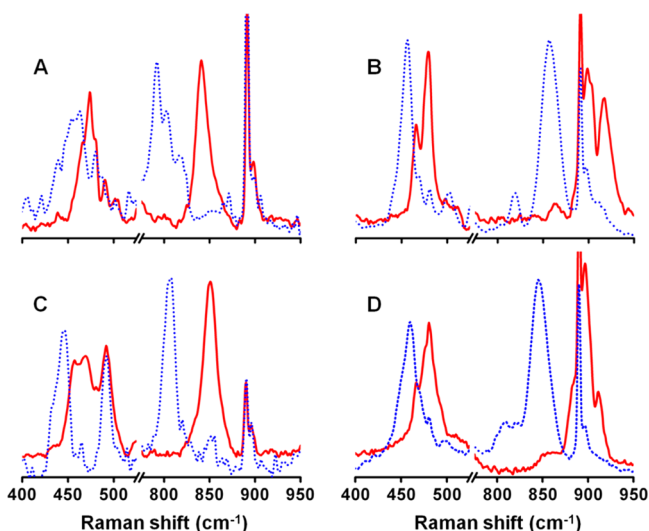


Figure 3. Selected UV-vis spectra obtained in the reaction of O_2 and $1\cdot\text{O}_2\text{CCPh}_3$ in CH_2Cl_2 at -90°C showing initial formation of $2\cdot\text{O}_2\text{CCPh}_3$ (solid green line) and subsequent conversion to $3\cdot\text{O}_2\text{CCPh}_3$ (solid blue line). Dotted lines correspond to spectra leading to the appearance of $2\cdot\text{O}_2\text{CCPh}_3$ within the first 40 s, while dashed lines correspond to spectra associated with the conversion of $2\cdot\text{O}_2\text{CCPh}_3$ to $3\cdot\text{O}_2\text{CCPh}_3$ over the course of the following hour.

downshift to 791–807 cm^{-1} upon ^{18}O substitution, while the $3\cdot\text{O}_2\text{X}$ cluster has $\nu_{\text{O-O}}$ values ranging from 897 to $\sim 910\text{ cm}^{-1}$ that downshift to 845–857 cm^{-1} with ^{18}O incorporation. On the other hand, no systematic difference is observed for the $\nu_{\text{Fe-O}}$ values of the $2\cdot\text{O}_2\text{X}$ and $3\cdot\text{O}_2\text{X}$ complexes, all of which range from 457 to 479 cm^{-1} (Table 4). These features can be assigned to the symmetric Fe–O stretch of the Fe– O_2 –Fe moiety and often appear as Fermi doublets. The difference in the $\nu_{\text{O-O}}$ values of $2\cdot\text{O}_2\text{X}$ and $3\cdot\text{O}_2\text{X}$ complexes has previously been attributed to a change in the Fe–Fe distance in the O_2 adducts on the basis of a systematic study of 1,2-peroxo-bridged diiron(III) complexes by Fiedler et al.²⁷ For the $2\cdot\text{O}_2\text{X}$ and $3\cdot\text{O}_2\text{X}$ complexes, the change in the Fe–Fe distance indicated by the difference in the $\nu_{\text{O-O}}$ values is postulated to derive from

Table 3. UV–Vis Properties of $2\cdot\text{O}_2\text{X}$, $3\cdot\text{O}_2\text{X}$ and $3'\cdot\text{O}_2\text{X}$ in CH_2Cl_2

O_2X	λ_{max} (nm), $[\epsilon]$ ($\text{M}^{-1} \text{cm}^{-1}$)		
	$2\cdot\text{O}_2\text{X}$	$3\cdot\text{O}_2\text{X}$	$3'\cdot\text{O}_2\text{X}$
O_2AsMe_2	632 [2100]	— ^a	— ^b
O_2PMe_2	674 [1800]	— ^a	616 [2400]
O_2PPh_2	678 [2100]	621 [1800]	592 [2600]
$\text{O}_2\text{P(OPh)}_2$	680 [2800]	576 [3100]	594 [3600]
O_2CCPh_3	708 [2500]	630 [2200]	595 [3300]
O_2CCMe_3	706 [—] ^c	580 [3100]	577 [3800]
O_2CPh	704 [—] ^c	588 [1500] ^d	592 [2300]
$\text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$	707 [2600]	590 [3200]	592 [4300]
$\text{O}_2\text{CC}_6\text{H}_3\text{-3,4-(OMe)}_2$	704 [—] ^c	578 [6200]	590 [5300]
$\text{O}_2\text{CC}_6\text{H}_3\text{-3,5-(OMe)}_2$	706 [—] ^c	588 [5400]	592 [5600]
$\text{O}_2\text{CC}_6\text{H}_4\text{-4-OMe}$	705 [—] ^c	580 [3000]	590 [4000]

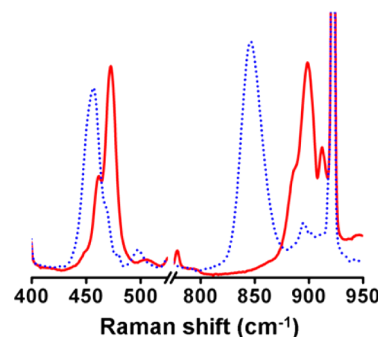
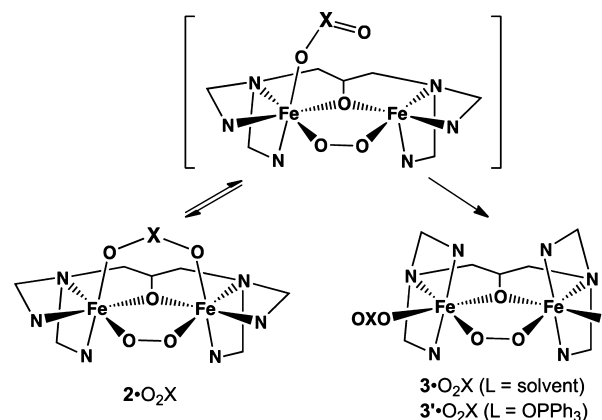
^aConversion to $3\cdot\text{O}_2\text{X}$ did not take place at any temperature.^bConversion to $3'\cdot\text{O}_2\text{X}$ did not take place at any temperature, even with addition of 100 equiv of OPPh_3 . ^cConversion of $2\cdot\text{O}_2\text{X}$ to $3\cdot\text{O}_2\text{X}$ began before complete formation of $2\cdot\text{O}_2\text{X}$, so ϵ was not determined.^dValues from ref 42.Figure 4. Resonance Raman spectra of $2\cdot\text{O}_2\text{CCPh}_3$ (A), $3\cdot\text{O}_2\text{CCPh}_3$ (B), $2\cdot\text{O}_2\text{P(OPh)}_2$ (C), and $3\cdot\text{O}_2\text{P(OPh)}_2$ (D). Solid red lines ($^{16}\text{O}_2$) and dotted blue lines ($^{18}\text{O}_2$); $T = 77 \text{ K}$, $\lambda_{\text{ex}} = 647 \text{ nm}$.

the conversion of the O_2X moiety from a bidentate bridging ligand in $2\cdot\text{O}_2\text{X}$ to a terminal monodentate ligand in $3\cdot\text{O}_2\text{X}$ (Scheme 1).

Attempts to obtain resonance Raman spectra for either $2\cdot\text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$ or $2\cdot\text{O}_2\text{CC}_6\text{H}_3\text{-3,4-(OMe)}_2$ were unsuccessful, even though they exhibited a longer half-life than $2\cdot\text{O}_2\text{CCPh}_3$. A color change was observed during the freezing process. Their respective Raman spectra were comparable to those of $3\cdot\text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$ and $3\cdot\text{O}_2\text{CC}_6\text{H}_3\text{-3,4-(OMe)}_2$, indicating that conversion took place during the phase change (Figure S1, Supporting Information). On the other hand, the longer lifetimes of the $3\cdot\text{O}_2\text{X}$ intermediates allowed us to collect spectra of every one (Figures 4 and S1, Supporting Information), with the exception of $3\cdot\text{O}_2\text{PMe}_2$, which did not form in quantities sufficient for characterization. The resonance Raman spectrum of $3'\cdot\text{O}_2\text{PPh}_2$, formed by treatment of $2\cdot\text{O}_2\text{PPh}_2$ with an excess of OPPh_3 (Figure 5), was also acquired. As it was nearly identical to the previously reported spectrum of $3'\cdot\text{O}_2\text{CPh}$,²⁷ we did not carry

Table 4. Fe–O and O–O Stretching Frequencies of $2\cdot\text{O}_2\text{X}$ and $3\cdot\text{O}_2\text{X}$ Complexes Determined Using Resonance Raman Spectroscopy

O_2X	$2\cdot\text{O}_2\text{X}$		$3\cdot\text{O}_2\text{X}$	
	$\nu_{\text{Fe-O}} (\text{cm}^{-1})$ [^{18}O]	$\nu_{\text{O-O}} (\text{cm}^{-1})$ [$^{18}\text{O}_2$]	$\nu_{\text{Fe-O}} (\text{cm}^{-1})$ [^{18}O]	$\nu_{\text{O-O}} (\text{cm}^{-1})$ [$^{18}\text{O}_2$]
O_2AsMe_2	464 [433]	845 [796]	—	—
O_2PMe_2	467, 479 [449, 460]	839 [791]	—	—
O_2PPh_2	465, 476 [455]	845, 853 [807]	477 [458]	897 [848]
$\text{O}_2\text{P(OPh)}_2$	457, 469 [436, 446]	851 [806]	479 [460]	897 [845]
O_2CCPh_3	466, 474 [447, 454]	841 [792]	466, 479 [448, 456]	903, 917 [857]
O_2CCMe_3	—	—	466, 475 [447, 457]	897, 912 [846]
O_2CPh^a	—	—	476 [460]	900 [850]
$\text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$	—	—	465, 476 [449, 457]	900, 914 [853]
$\text{O}_2\text{CC}_6\text{H}_3\text{-3,4-(OMe)}_2$	—	—	466, 474 [449, 456]	898, 912 [847]
$\text{O}_2\text{CC}_6\text{H}_3\text{-3,5-(OMe)}_2$	—	—	465, 476 [449, 458]	899, 913 [848]
$\text{O}_2\text{CC}_6\text{H}_4\text{-4-OMe}$	—	—	466, 474 [448, 457]	899, 913 [847]

^aValues from ref 42.Scheme 1. Conversion of $2\cdot\text{O}_2\text{X}$ to $3\cdot\text{O}_2\text{X}$ or $3'\cdot\text{O}_2\text{X}$ Figure 5. Resonance Raman spectra of $3'\cdot\text{O}_2\text{PPh}_2$ ($^{16}\text{O}_2$ = solid red line, $^{18}\text{O}_2$ = dotted blue line).

out the corresponding experiments with other $3'\cdot\text{O}_2\text{X}$ intermediates.

Conversion of $2\cdot\text{O}_2\text{X}$ to $3\cdot\text{O}_2\text{X}$. The spectral differences between $2\cdot\text{O}_2\text{X}$ and $3\cdot\text{O}_2\text{X}$ make UV–vis absorption spectrosc-

copy an excellent tool for following the transition from the first peroxo intermediate to the second. On the basis of the data shown in Figure 3, analysis of the spectral changes observed upon oxygenation of $1\text{-O}_2\text{CCPh}_3$ at -90°C afforded a first-order rate constant of $1.7 \times 10^{-3} \text{ s}^{-1}$ for the conversion of $2\text{-O}_2\text{CCPh}_3$ to $3\text{-O}_2\text{CCPh}_3$. This value is comparable to that previously reported for the conversion of $2\text{-O}_2\text{PPh}_2$ to $3\text{-O}_2\text{PPh}_2$ at -40°C , thus implicating a more facile conversion for the carboxylate-bridged species. Figure 6 provides evidence

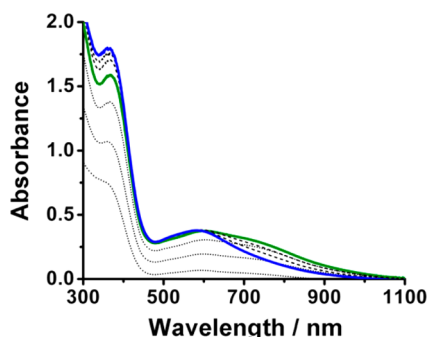


Figure 6. Selected UV-vis spectra obtained in the reaction of O_2 with $1\text{-O}_2\text{CPh}$ at -90°C showing evidence for the formation of two peroxo intermediates. The solid green line represents the spectrum with the largest fraction of $2\text{-O}_2\text{CPh}$ formed, while the solid blue line corresponds to the subsequently formed $3\text{-O}_2\text{CPh}$. Dotted lines correspond to spectra leading to the maximum amount of $2\text{-O}_2\text{CPh}$ formed within the first 90 s after oxygenation, while dashed lines correspond to spectra associated with the conversion of $2\text{-O}_2\text{CPh}$ to $3\text{-O}_2\text{CPh}$ over the course of the following 30 min.

for the formation of two peroxo species at -90°C even for the well-studied complex $1\text{-O}_2\text{CPh}$,^{29,42,44} although they are not as spectroscopically distinct as those of $2\text{-O}_2\text{CCPh}_3$ and $3\text{-O}_2\text{CCPh}_3$. Stopped-flow kinetic analysis at -80°C revealed the conversion from $2\text{-O}_2\text{CPh}$ to $3\text{-O}_2\text{CPh}$ with a k_{obs} of $5.3 \times 10^{-2} \text{ s}^{-1}$, much faster than that observed for $2\text{-O}_2\text{CCPh}_3$ to $3\text{-O}_2\text{CCPh}_3$ at -90°C . This comparison provides a rationale for why $2\text{-O}_2\text{CPh}$ was not observed previously.^{29,42,44} The rate of conversion of $2\text{-O}_2\text{X}$ to $3\text{-O}_2\text{X}$ decreased as electron-donating substituents were introduced onto the benzoate ring or as the steric bulk of the carboxylate bridge was increased (Table 5). The rate of conversion was further decreased by replacing the trigonal carboxylate moiety with tetrahedral anions such as $(\text{PhO})_2\text{PO}_2^-$, Ph_2PO_2^- , Me_2PO_2^- , and $\text{Me}_2\text{AsO}_2^-$ (Table 5).

In our previous publication,²⁸ we postulated that the conversion of $2\text{-O}_2\text{X}$ to $3\text{-O}_2\text{X}$ in acetonitrile involves a change in O_2X binding mode from bridging to terminal and coordination of a solvent molecule in position L (Scheme 1). For this work we added OPPh_3 to solutions of $2\text{-O}_2\text{X}$ in hopes of substituting it into the L position. Upon OPPh_3 addition, purple-blue species ($3'\text{-O}_2\text{X}$) indeed formed with λ_{max} values at wavelengths near each respective $3\text{-O}_2\text{X}$, although the extinction coefficients of the new intermediates are higher than those observed for each respective $3\text{-O}_2\text{X}$ (Table 3), with the exception of the $\text{O}_2\text{CC}_6\text{H}_3\text{-3,4-(OMe)}_2$ -based species. While facile conversion to $3'\text{-O}_2\text{X}$ from most $2\text{-O}_2\text{X}$ intermediates was accomplished by adding 20 equiv of OPPh_3 at -90°C , $2\text{-O}_2\text{PMe}_2$ and $2\text{-O}_2\text{PPh}_2$ had to be warmed to -40°C to attain reasonable rates of conversion. In contrast,

Table 5. $\text{p}K_{\text{a}}$ Values of HO_2X and First-Order k_{obs} Values for Conversion of $2\text{-O}_2\text{X}$ to $3\text{-O}_2\text{X}$ and $2\text{-O}_2\text{X}$ to $3'\text{-O}_2\text{X}$

O_2X	$\text{p}K_{\text{a}}$ of HO_2X	k_{obs} (s^{-1})	
		$2 \rightarrow 3^a$	$2 \rightarrow 3'^b$
O_2AsMe_2	6.27	— ^c	— ^d
O_2PMe_2	3.08 ^e	— ^c	$1.8(4) \times 10^{-4}$ (-40°C)
O_2PPh_2	2.32 ^e	3.1×10^{-4f} (-40°C)	$4.3(4) \times 10^{-3}$ (-40°C)
$\text{O}_2\text{P}(\text{OPh})_2$	1.85 ^e	$1.2(1) \times 10^{-4}$	$3.1(1) \times 10^{-2}$
$\text{O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$	4.24	$4.2(6) \times 10^{-4}$	0.13(5)
$\text{O}_2\text{CC}_6\text{H}_3\text{-3,4-(OMe)}_2$	4.36	$6.6(9) \times 10^{-4}$	0.12(1)
O_2CCPh_3	3.96	$1.7(2) \times 10^{-3}$	$2.6(3) \times 10^{-2}$
$\text{O}_2\text{CC}_6\text{H}_3\text{-3,5-(OMe)}_2$	3.97	$4.8(2) \times 10^{-3}$	0.20(1)
$\text{O}_2\text{CC}_6\text{H}_4\text{-4-OMe}$	4.50	$4.7(4) \times 10^{-3}$	0.12(1)
O_2CCMe_3	5.03	$1.8(1) \times 10^{-2}$	$8.0(3) \times 10^{-2}$
O_2CPh	4.19	$5.3(1) \times 10^{-2g}$ (-80°C)	— ^h

^aAll rates measured at -90°C , except where noted. ^bAll rates measured after addition of 20 equiv of OPPh_3 at -90°C , except where noted. ^cConversion to observable concentrations of $3\text{-O}_2\text{X}$ did not take place at any temperature. ^dConversion to observable concentrations of $3'\text{-O}_2\text{X}$ did not take place at any temperature, even with addition of 100 equiv of OPPh_3 . ^eMeasured in 7% EtOH (ref 53). ^fAt -40°C in CH_2Cl_2 , $3\text{-O}_2\text{PPh}_2$ starts to decay before complete conversion from $2\text{-O}_2\text{PPh}_2$ occurs. For this reason, k_{obs} was calculated from the y-intercept of the OPPh_3 concentration dependence plot for the conversion of $2\text{-O}_2\text{PPh}_2$ to $3'\text{-O}_2\text{PPh}_2$ (Figure S2, Supporting Information). ^gRate measured at -80°C using stopped-flow techniques. ^hBy the time enough $2\text{-O}_2\text{CPh}$ had formed to allow for addition of OPPh_3 , significant conversion to $3\text{-O}_2\text{CPh}$ had occurred, preventing accurate rate determination for the conversion of $2\text{-O}_2\text{CPh}$ to $3'\text{-O}_2\text{CPh}$.

$2\text{-O}_2\text{AsMe}_2$ appears to be unaffected by addition of up to 100 equiv of OPPh_3 , even at temperatures as high as 20°C .

After finding evidence indicating the formation of $2\text{-O}_2\text{CPh}$, we opted to further examine the oxygenation kinetics of both $1\text{-O}_2\text{PPh}_2$ and $1\text{-O}_2\text{CPh}$ using stopped-flow techniques. $1\text{-O}_2\text{PPh}_2$ offers an excellent opportunity to characterize the initial oxygen coordination with the formation of $2\text{-O}_2\text{PPh}_2$, since the subsequent rearrangement of $2\text{-O}_2\text{PPh}_2$ to $3\text{-O}_2\text{PPh}_2$ is relatively slow. The reaction between $1\text{-O}_2\text{PPh}_2$ and O_2 in either MeCN or CH_2Cl_2 , which is accompanied by rapid growth of the absorption band with $\lambda_{\text{max}} \approx 680 \text{ nm}$, was studied. At low temperatures (down to -80°C in CH_2Cl_2), the initially formed diiron- O_2 intermediate $2\text{-O}_2\text{PPh}_2$ is stable for at least 1 h. However, noticeable decay of this intermediate was observed under intense illumination with polychromatic, UV-rich light of the arc lamp. In order to avoid complications from undesirable photodecomposition, quantitative kinetic measurements were performed in a single-wavelength mode, with low-intensity monochromatic light and a sensitive photomultiplier detector. Under these conditions, photobleaching was not observed. In order to determine the rate law, kinetic traces were acquired under a large excess of O_2 , showing single-exponential growth of $2\text{-O}_2\text{PPh}_2$ and yielding the values of observed pseudo-first-order rate constants, k_{obs} . The observed rate constants did not depend on the initial concentration of $1\text{-O}_2\text{PPh}_2$, in agreement with a rate law that is first-order in the diiron precursor. The observed rate constants increased linearly

with an increase in the concentration of O_2 (Figure S3, Supporting Information), indicating that the reaction was first-order in dioxygen. It can be concluded that the formation of $2\cdot O_2PPh_2$ is a second-order process (first-order in diiron complex and first-order in O_2): $\nu = k_2[1\cdot O_2PPh_2][O_2]$

As the temperature was raised, the oxygenation rate of $1\cdot O_2PPh_2$ increased modestly, and the conversion of $2\cdot O_2PPh_2$ to $3\cdot O_2PPh_2$ became pronounced. The Eyring plot (Figure 7)

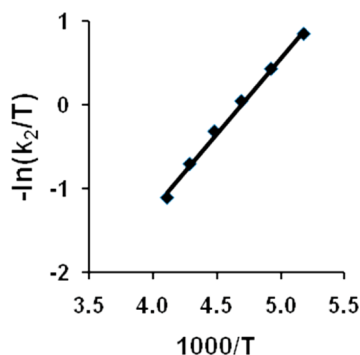


Figure 7. Eyring plot for O_2 binding to $1\cdot O_2PPh_2$ in CH_2Cl_2 .

for the oxygen binding step was linear over a broad temperature range (from -80 to -20 °C in CH_2Cl_2), yielding the activation parameters that are summarized in Table 6. The kinetics of oxygen binding to $1\cdot O_2PPh_2$ in MeCN was also examined, and very similar activation parameters were extracted (Table 6). Subsequent disappearance of $2\cdot O_2PPh_2$ was readily observed in MeCN at temperatures above -40 °C (Figure S4, Supporting Information). The spectral changes agreed well with the conversion of $2\cdot O_2PPh_2$ into $3\cdot O_2PPh_2$ described in detail by Frisch et al.²⁸ Low activation enthalpies and large negative activation entropies for the oxygenation of $1\cdot O_2PPh_2$ leading to $2\cdot O_2PPh_2$ are typical of associative O_2 coordination at vacant or labile iron sites,^{34,45} which are present in the starting diiron(II) complex. The rates and activation parameters for the oxygenation of $1\cdot O_2PPh_2$ in CH_2Cl_2 or in acetonitrile are very similar to previously published kinetic parameters for the oxygenation of $1\cdot O_2CPh$ in propionitrile.⁴³ The kinetics of oxygenation of five-coordinate diiron(II) complexes with a related dinucleating ligand HPTP is also very similar (Table 6).⁴⁶

A closer examination of dioxygen binding to $1\cdot O_2CPh$ identified the individual reaction steps of the overall reaction. Stopped-flow observation of the reaction of $1\cdot O_2CPh$ with dioxygen (2.9 mM) in CH_2Cl_2 at -80 °C reveals a two-step process (Figure S5, Supporting Information), in agreement with sequential formation of intermediates $2\cdot O_2CPh$ and $3\cdot O_2CPh$. At longer wavelengths (greater than ca. 650 nm), the initial absorbance increase takes place within 15 s and is followed by decay that is complete within 100 s. At 640 nm, the initial absorbance increase was also seen, but after 15 s the

kinetic trace becomes flat. At shorter wavelengths, a biexponential absorbance increase was seen. Two-exponential fit of the variable-wavelengths data gave the following value of the observed rate constant: $k_1 = 0.22\text{ s}^{-1}$ at -80 °C ($76\text{ M}^{-1}\text{ s}^{-1}$ under the assumption of a second-order process). This rate constant corresponds to the first reaction step, formation of $2\cdot O_2CPh$. Interestingly, it is in excellent agreement with the reported value of the rate constant for the reaction of $1\cdot O_2CPh$ with O_2 ($135\text{ M}^{-1}\text{ s}^{-1}$ at -75 °C in propionitrile, with the activation parameters of $\Delta H^\ddagger = 15.4\text{ kJ/mol}$, $\Delta S^\ddagger = -121\text{ J K}^{-1}\text{ mol}^{-1}$).⁴³ In that work, $3\cdot O_2CPh$ was characterized spectroscopically, and it was assumed that the kinetics of oxygenation corresponded to the formation of $3\cdot O_2CPh$. On the basis of our studies, the kinetic parameters reported by Feig et al.⁴³ correspond to the first oxygenation step, the formation of $2\cdot O_2CPh$; the next step, conversion of $2\cdot O_2CPh$ to $3\cdot O_2CPh$, is very rapid at most temperatures.

DISCUSSION

One step in the catalytic activation of dioxygen by biological diiron(II) systems often produces $(\mu-\eta^1:\eta^1\text{-peroxo})$ diiron(III) moieties. Some of these intermediates exhibit a level of stability that allows for them to be trapped and characterized.^{9–16} The same is true for many synthetic $(\mu-\eta^1:\eta^1\text{-peroxo})$ diiron(III) complexes,^{18,25,27,41,47} some of which are so stable they have been crystallographically characterized.^{19–22} In previously published work,²⁸ we used the dinucleating ligand N-EtHPTB and different oxanions (O_2X) to synthesize a set of three dioxygen binding diiron(II) complexes ($1\cdot O_2X$). Our efforts revealed that the $(\mu-\eta^1:\eta^1\text{-peroxo})$ diiron(III) intermediates produced upon oxygenation of the diiron(II) precursors come in two forms, green-blue $2\cdot O_2X$ and deep-blue $3\cdot O_2X$ (Scheme 1). In the initial form, the oxanion acts as a three-atom bridge between the iron centers and the pendant benzimidazoles are cis to each other on each iron. In many cases, this intermediate converts to a second form, wherein the O_2X ligand has moved to a terminal position, allowing the pendant benzimidazoles of the N-EtHPTB ligand to rearrange from a cis to a trans disposition on each iron. We found that the stability of $2\cdot O_2X$ is influenced by the identity of O_2X ($O_2X = O_2AsMe_2$, O_2PPh_2 , and O_2CPh) and concluded that the dominant factor governing $2\cdot O_2X$ stability is the bite distance ($O\cdots O$) of the O_2X moiety in $1\cdot O_2X$ as determined by X-ray crystallography. Anions with greater bite distances are better able to accommodate the $>3\text{ \AA}$ $Fe\cdots Fe$ distance in $2\cdot O_2X$ and gave rise to more stable $2\cdot O_2X$ intermediates; for $X = O_2AsMe_2$, the $2\cdot O_2X$ intermediate was so stable that no observable conversion to $3\cdot O_2X$ was observed prior to decomposition.

For this work, we synthesized several more $1\cdot O_2X$ complexes to assess additional factors that may affect the stability of $2\cdot O_2X$. Specifically, we were interested in examining effects produced by electronic and steric changes in O_2X and how those results relate to effects produced by differences in O_2X

Table 6. Kinetic Parameters for Reactions of Diiron(II) Complexes with Dioxygen

complex	solvent	T range (°C)	k (−40 °C) ($M^{-1}\text{ s}^{-1}$)	$\Delta H^\ddagger$ (kJ/mol)	$\Delta S^\ddagger$ (J/K·mol)	ref
$1\cdot O_2PPh_2$	CH_2Cl_2	−80 to −20	4.7×10^2	15.7(4)	−124(6)	this work
$1\cdot O_2PPh_2$	MeCN	−40 to 0	2.55×10^3	13.7(4)	−119(10)	this work
$1\cdot O_2CPh$	EtCN	−75 to −15	$1.03(12) \times 10^2$	15.4(6)	−121(3)	43
$[Fe_2(HPTP)(O_2CPh)]^{2+}$	MeCN	−40 to 0	7.3×10^3	15.8(4)	−101(10)	46
$[Fe_2(HPTP)(O_2CPh)]^{2+}$	CH_2Cl_2	−80 to 0	67	16.7(2)	−132(8)	46

bite distances. We also investigated how OPPh_3 could be used to destabilize some $2\text{-O}_2\text{X}$ intermediates. In addition, we used low-temperature stopped-flow techniques to demonstrate that the oxygenation of $1\text{-O}_2\text{CPh}$ was in fact a two-step process like those we have described for the other $1\text{-O}_2\text{X}$ complexes. Indeed, $2\text{-O}_2\text{CPh}$ is short-lived, even at -90°C , so the adduct we originally observed at -40°C and assigned to be $2\text{-O}_2\text{CPh}$ in 1990^{29,42} is in actuality the more stable $3\text{-O}_2\text{CPh}$ isomer. Finally, we determined the activation parameters for the oxygenation of $1\text{-O}_2\text{PPh}_2$ in CH_2Cl_2 and in MeCN and found them to be very similar to those reported earlier for $1\text{-O}_2\text{CPh}$ in EtCN (Table 6),⁴³ strongly suggesting a common rate-determining step for these reactions, corresponding to the formation of $2\text{-O}_2\text{X}$.

The focus of the experiments reported in this paper has been to gain further insight into the factors that affect the conversion of $2\text{-O}_2\text{X}$ to $3\text{-O}_2\text{X}$, in which irreversible conversion to $3\text{-O}_2\text{X}$ is preceded by movement of the O_2X moiety from a bridging to a terminal position (Scheme 1). We postulate that there is a rapid equilibrium between bridging and terminal coordination modes of the O_2X ligand in $2\text{-O}_2\text{X}$ and only the isomer with a terminal O_2X ligand can undergo conversion to $3\text{-O}_2\text{X}$ during which the benzimidazole arms of the N-EtHPTB ligand shift from a cis relationship to each other to a trans configuration on both iron centers. Thus, the initial preequilibrium should be affected by the basicity of O_2X ligands, with the more basic ligand favoring the bridging mode and thereby decreasing the fraction of monodentate O_2X isomer available to undergo conversion to $3\text{-O}_2\text{X}$. On the other hand, as the transformation from $2\text{-O}_2\text{X}$ to $3\text{-O}_2\text{X}$ entails a significant rearrangement of the coordination spheres about each iron center, the conversion should be slowed down by an increase in the steric bulk of O_2X .

The trend in the stability of the $2\text{-O}_2\text{X}$ complexes is best discerned by an examination of the kinetic data for the large subset of complexes with carboxylate bridges. A perusal of Table 5 suggests that, while it may be possible to observe an effect of ligand basicity in comparisons of select pairs (e.g., O_2CPh vs $\text{O}_2\text{CC}_6\text{H}_4\text{-4-OMe}$, and O_2CPh vs O_2CCMe_3), for the most part steric considerations supersede ligand basicity arguments. For example, the stability of $2\text{-O}_2\text{CR}$ is enhanced 30-fold in the series, $\text{R} = \text{Ph} < \text{CMe}_3 < \text{CPh}_3$, commensurate with the increase in steric bulk on the carbon atom adjacent to the carboxylate function. Although the higher $\text{p}K_{\text{a}}$ (5.03) of HO_2CCMe_3 ⁴⁸ relative to benzoic acid ($\text{p}K_{\text{a}} = 4.19$)⁴⁸ could be used to rationalize the 3-fold longer lifetime of $2\text{-O}_2\text{CCMe}_3$, the 30-fold greater stability of $2\text{-O}_2\text{CCPh}_3$ than $2\text{-O}_2\text{CPh}$ cannot be explained by the slightly lower $\text{p}K_{\text{a}}$ of HO_2CCPh_3 (3.96)⁴⁹ but can easily be rationalized by the much greater bulk of the triphenylmethyl group relative to phenyl.

Complexes with methoxy-substituted benzoate bridges support the above arguments. The conversion of $2\text{-O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$ to its $3\text{-O}_2\text{X}$ form is 100-fold slower than for $2\text{-O}_2\text{CPh}$, despite the fact that the two carboxylic acids have essentially identical $\text{p}K_{\text{a}}$ values (4.24 and 4.19, respectively).^{48,50} Therefore, the considerable difference in the stabilities of $2\text{-O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$ and $2\text{-O}_2\text{CPh}$ (Table 5) must arise from steric considerations. Other methoxy-substituted benzoate complexes exhibit intermediate rates of conversion; $2\text{-O}_2\text{CC}_6\text{H}_3\text{-3,4-(OMe)}_2$ converts just slightly faster than $2\text{-O}_2\text{CC}_6\text{H}_2\text{-3,4,5-(OMe)}_3$, while $2\text{-O}_2\text{CC}_6\text{H}_3\text{-3,5-(OMe)}_2$ and $2\text{-O}_2\text{CC}_6\text{H}_4\text{-4-(OMe)}$ are 10-fold faster. From these results, it seems clear that additional steric bulk in the

meta and/or para positions of benzoate-based oxyanions is the primary factor that stabilizes the resultant $2\text{-O}_2\text{X}$ complexes.

The lifetimes of $2\text{-O}_2\text{CR}$ complexes are shortened upon addition of OPPh_3 . We previously found that OPPh_3 had a good binding affinity for the parent $\text{Fe}^{\text{III}}_2(\text{N-Et-HPTB})$ -peroxo complex and exerted such a significant stabilizing effect that its presence led to the crystallization and structural characterization of $[\text{Fe}^{\text{III}}_2(\mu\text{-1,2-O}_2)(\text{N-Et-HPTB})(\text{OPPh}_3)_2]^{3+}$.²⁰ Addition of 20 equiv of OPPh_3 to $2\text{-O}_2\text{CR}$ at -90°C in fact accelerated the conversion to $3'\text{-O}_2\text{CR}$ with rates of 0.03–0.2 s^{-1} , representing a much smaller range of values than for the conversions in the absence of OPPh_3 (Table 5). These results suggest that the binding of OPPh_3 to the diiron center facilitates the ligand rearrangement required to convert from $2\text{-O}_2\text{X}$ to $3\text{-O}_2\text{X}$ (Scheme 1).

Tetrahedral O_2X bridges also increase the stability of $2\text{-O}_2\text{X}$. In this series, we compared complexes with O_2AsMe_2 , O_2PMe_2 , O_2PPh_2 , and $\text{O}_2\text{P(OPh)}_2$ bridges that differ in steric properties and basicity ($\text{p}K_{\text{a}} = 6.27, 3.08, 2.32$, and 1.85 , respectively).^{51–53} The order of stability in this subset is $\text{O}_2\text{AsMe}_2 > \text{O}_2\text{PMe}_2 > \text{O}_2\text{PPh}_2 > \text{O}_2\text{P(OPh)}_2$ (Table 4), following the trend of decreasing $\text{p}K_{\text{a}}$ values. Only $2\text{-O}_2\text{P(OPh)}_2$ undergoes conversion to the corresponding $3\text{-O}_2\text{X}$ form at -90°C , with $k_{\text{obs}} = 1.2(1) \times 10^{-4} \text{ s}^{-1}$, a rate that is slower than observed for five of the six $2\text{-O}_2\text{CR}$ complexes at -90°C (Table 5). While $2\text{-O}_2\text{PPh}_2$ is indefinitely stable at -90°C , it converts to $3\text{-O}_2\text{PPh}_2$ when warmed to -40°C , at a rate comparable to that of $2\text{-O}_2\text{P(OPh)}_2$ at -90°C . Neither $2\text{-O}_2\text{PMe}_2$ nor $2\text{-O}_2\text{AsMe}_2$ generate an observable $3\text{-O}_2\text{X}$ form; instead they appear to decay directly to $4\text{-O}_2\text{X}$. However, $2\text{-O}_2\text{PMe}_2$ can be converted to $3'\text{-O}_2\text{PMe}_2$ at -40°C by addition of OPPh_3 , whereas $2\text{-O}_2\text{AsMe}_2$ does not convert to $3'\text{-O}_2\text{AsMe}_2$, even when reacted with 100 equiv of OPPh_3 at room temperature. The increasing stability of $2\text{-O}_2\text{X}$ with the bite distance of the oxyanion bridge ($\text{X} = \text{C}, 2.23 \text{ \AA}$; $\text{X} = \text{P}, 2.55 \text{ \AA}$; $\text{X} = \text{As}, 2.80 \text{ \AA}$) reflects the ability of the O_2X bridge to span the $\text{Fe}\cdots\text{Fe}$ distance required by the dinucleating HPTB ligand framework without imposing a strain on the six-member ring formed by the three atoms of the oxyanion bridge, the two iron atoms, and the alkoxide oxygen on the N-EtHPTB ligand. The O_2AsMe_2 complex, having the longest $\text{O}\cdots\text{O}$ bite distance, represents the most stable of the $2\text{-O}_2\text{X}$ intermediates.

In this paper, we have investigated the various factors that influence the conversion of $2\text{-O}_2\text{X}$ O_2 adducts to corresponding $3\text{-O}_2\text{X}$ species. This conversion entails the shift of the O_2X ligand from a bidentate bridging mode to a terminal monodentate mode, providing another example of a diiron complex involved in a mechanistically important “carboxylate shift” first recognized by Lippard and co-workers 20 years ago.³⁰ An analogous carboxylate shift has been suggested by Do et al.⁵⁴ to occur upon protonation of O_2 adducts of $[\text{Fe}^{\text{II}}_2(\text{N-EtHPTB})(\mu\text{-O}_2\text{CR})]^{2+}$ complexes ($\text{R} = \text{Ph}$ or C_6F_5) at -30°C . However, although unequivocal evidence was provided for the protonation of the carboxylate ligand, it is clear from a comparison with the spectroscopic data we have presented in this paper that the O_2 adducts studied by Do et al.⁵⁴ must be assigned structures we now associate with $3\text{-O}_2\text{CR}$ complexes, which have terminal monodentate carboxylates. On the other hand, the corresponding $2\text{-O}_2\text{CR}$ complexes (with bridging carboxylates) are in fact formed upon oxygenation of the diiron(II) precursors (as previously assumed^{29,42,54}) but can be observed only at -90°C , as they readily undergo the “carboxylate shift” at that temperature to isomerize to

corresponding $3\text{-O}_2\text{CR}$ species. However, the key take-home message of the study by Do et al.⁵⁴ is that protonation of a carboxylate ligand on a diiron complex can occur and give rise to a complex with a carboxylic acid ligand. Indeed, a carboxylate ligand may serve as a convenient conduit for delivering a proton to a bound peroxide to facilitate O–O bond cleavage, for example, in the conversion of intermediate **P** to **Q** in the case of sMMO, for which a proton is clearly implicated.^{13,55}

■ ASSOCIATED CONTENT

■ Supporting Information

Figures S1–S5. 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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