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1901Tuning the stereoelectronic factors of iron(II)-
2-aminophenolate complexes for the reaction
with dioxygen: oxygenolytic C–C bond cleavage
vs. oxidation of complex†Sayanti Chatterjee, ^{‡a} Sridhar Banerjee,^a Rahul Dev Jana,^a
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Oxidative C–C bond cleavage of 2-aminophenols mediated by transition metals and dioxygen is a topic of great interest. While the oxygenolytic C–C bond cleavage reaction relies on the inherent redox non-innocent property of 2-aminophenols, the metal complexes of 2-aminophenolates often undergo 1e[−]/2e[−] oxidation events (metal or ligand oxidation), instead of the direct addition of O₂ for subsequent C–C bond cleavage. In this work, we report the isolation, characterization and dioxygen reactivity of a series of ternary iron(II)-2-aminophenolate complexes [(Tp^{Ph,Me})Fe^{II}(X)], where X = 2-amino-4-*tert*-butylphenolate (4-*t*Bu-HAP) (**1**); X = 2-amino-4,6-di-*tert*-butylphenolate (4,6-di-*t*Bu-HAP) (**2**); X = 2-amino-4-nitrophenolate (4-NO₂-HAP) (**3**); and X = 2-anilino-4,6-di-*tert*-butylphenolate (NH-Ph-4,6-di-*t*Bu-HAP) (**4**) supported by a facial tridentate nitrogen donor ligand (Tp^{Ph,Me} = hydrotris(3-phenyl-5-methylpyrazol-1-yl)borate). Another facial N₃ ligand (Tp^{Ph2} = hydrotris(3,5-diphenyl-pyrazol-1-yl)borate) has been used to isolate an iron(II)-2-anilino-4,6-di-*tert*-butylphenolate complex (**5**) for comparison. Both [(Tp^{Ph,Me})Fe^{II}(4-*t*Bu-HAP)] (**1**) and [(Tp^{Ph,Me})Fe^{II}(4,6-di-*t*Bu-HAP)] (**2**) undergo regioselective oxidative aromatic ring fission reaction of the coordinated 2-aminophenols to the corresponding 2-picolinic acids in the reaction with dioxygen. In contrast, complex [(Tp^{Ph,Me})Fe^{II}(4-NO₂-HAP)] (**3**) displays metal based oxidation to form an iron(III)-2-amidophenolate complex. Complexes [(Tp^{Ph,Me})Fe^{II}(NH-Ph-4,6-di-*t*Bu-HAP)] (**4**) and [(Tp^{Ph2})Fe^{II}(NH-Ph-4,6-di-*t*Bu-HAP)] (**5**) react with dioxygen to undergo 2e[−] oxidation with the formation of the corresponding iron(III)-2-iminobenzosemiquinonato radical species implicating the importance of the –NH₂ group in directing the C–C bond cleavage reactivity of 2-aminophenols. The systematic study presented in this work unravels the effect of the electronic and structural properties of the redox non-innocent 2-aminophenolate ring and the supporting ligand on the C–C bond cleavage reactivity vs. the metal/ligand oxidation of the complexes. The study further reveals that proper modulation of the stereoelectronic factors enables us to design a well synchronised proton transfer (PT) and dioxygen binding events for complexes **1** and **2** that mimic the structure and function of the nonheme enzyme 2-aminophenol-1,6-dioxygenase (APD).

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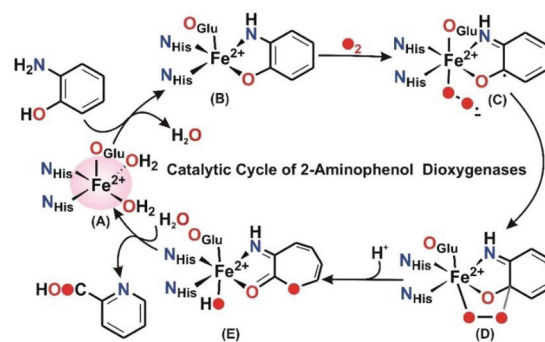
Introduction

Oxidative C–C bond cleavage reaction is a key metabolic step in the biodegradation of toxic and xenobiotic compounds.^{1–3} The resulting ring cleavage products are used by soil bacteria as the only source of energy and carbon in nature, which in turn is critical to maintaining the global carbon cycle.¹ Ring cleaving dioxygenases constitute a large family of mononuclear nonheme enzymes which are involved in the aerobic degradation of aromatic compounds, such as *ortho*-hydroxy phenols (catechols) or *o*-aminophenols or gentisic acid to aliphatic products in the soil with the incorporation of both atoms of dioxy-

gen into the cleavage products.^{1,4,5} In general, for the aromatic diols (catechols), depending upon the position of the C–C bond cleavage, these enzymes are classified into two categories: intradiol catechol dioxygenases and extradiol catechol dioxygenases.^{6,7} 2-Aminophenol-1,6-dioxygenase (APD), isolated and purified from *Pseudomonas pseudoalcaligenes* JS45, is a ring cleaving enzyme in a small group of nonheme extradiol type dioxygenases and is responsible for the C–C bond cleavage of 2-aminophenols (Scheme 1).⁸

In the catalytic cycle of the C–C bond cleavage pathway of the enzyme 2-aminophenol-1,6-dioxygenase (APD), the C1–C6 bond adjacent to the –OH group is cleaved^{9,10} in the presence of dioxygen.^{11,12} The C–C cleavage product, 2-aminomuconic acid semialdehyde, spontaneously loses a water molecule to form 2-picolinic acid (Scheme 1).¹³ A related enzyme, 3-hydroxyanthranilate-3,4-dioxygenase (HAD), isolated from *Saccharomyces cerevisiae*, catalyzes the ring opening of 3-hydroxyanthranilate to quinolinic acid in the tryptophan catabolism pathway.⁹ Structural studies reveal that the active sites of HAD^{14–16} and APD¹⁷ contain an iron(II) center coordinated by the “2-His-1-Glu facial triad” motif.^{6,18} 2-Aminophenol/3-hydroxyanthranilic acid substrates bind to the iron(II) center in a bidentate mode to initiate the aromatic ring cleavage reaction by activating dioxygen. On the basis of the structural and biochemical studies on HAD and APD, a mechanistic proposal similar to that of extradiol catechol dioxygenases has been proposed.⁹ The substrate binds to the ferrous ion (A) and the enzyme–substrate complex (B) activates dioxygen to form initially an iron(III)-superoxide intermediate that receives an electron from the redox non-innocent 2-aminophenolate unit to form an iron(II)-superoxide diradical species (C) (Scheme 2). An iron(II)-peroxide intermediate (D) is subsequently generated by the attack of the iron(II)-superoxo on the ligand based radical. The peroxide moiety then undergoes O–O bond heterolysis to form a lactone intermediate (E). One oxygen atom from the peroxide unit is incorporated into the lactone ring. Hydrolysis of the lactone affords the C–C bond cleavage product, 2-amino muconic acid semialdehyde, which is then converted to 2-picolinic acid through a nonenzymatic pathway (Scheme 2).¹⁹

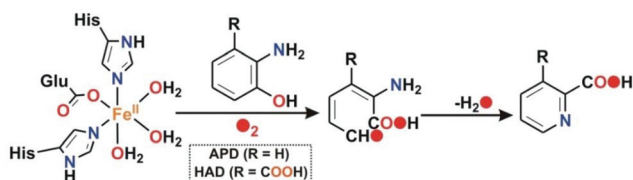
The oxidative transformation of 2-aminophenol to 2-picolinic acid by molecular oxygen has attracted the attention of biomimetic chemists to develop synthetic models of 2-aminophenol dioxygenase. Moreover, similar to catechols, *o*-aminophenols are well-known “redox non-innocent” ligands^{20–23} and



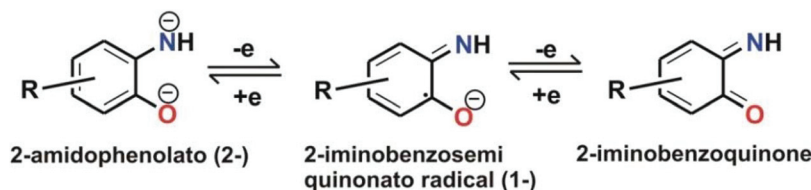
Scheme 2 Proposed mechanism for the catalytic cycle of APD.

are stabilized in different redox states upon binding with suitable metal ions (Scheme 3).

Reports of several iron(II) complexes coordinated by the non-innocent *o*-aminophenolate ligands in different redox states have been published by Wieghardt and coworkers.^{21,22,24–28} In biomimetic chemistry, dioxygen reactivity of model iron(II)-2-aminophenolate is less explored compared to the catecholate complexes.⁶ Paine's group reported the O₂-dependent C–C bond cleavage reactivity of iron(II)-2-aminophenolate complexes supported by neutral N₄ donor ligands.^{29,30} The complexes have been shown to react with dioxygen under ambient conditions to afford substituted 2-picolinic acids mimicking the function of 2-aminophenol dioxygenases (APD and HAD).^{29,30} These model complexes provide insights into the mechanism of aromatic ring cleavage of 2-aminophenols. Subsequently, the Paine group developed catalytic models of 2-aminophenol dioxygenases using neutral tridentate³¹ and tetradentate ligands.³² The role of the second sphere residue³³ as well as the pH of the reaction solution in controlling the catalytic reactivity and selectivity has been proposed. However, the reported catalytic model of 2-aminophenol dioxygenase with a tridentate ligand was not structurally characterized.³¹ In fact, there is no report of structurally characterized iron(II)-2-aminophenolate complex that closely mimics the active site structure of APD and at the same time displays C–C cleavage activity. Monoanionic tris(pyrazolyl)borates are often used as bioinspired N₃ donor ligands to mimic the ‘2-His-1-carboxylate facial triad’ found in the active sites of many dioxygen activating nonheme mono-iron enzymes.^{34–43} Since APD and HAD belong to the superfamily of nonheme iron enzymes with the ‘2-His-1-carboxylate’ facial triad, tris(pyrazolyl)borates could be appropriate ligands to develop biomimetic models of these enzymes. Fiedler's group reported the synthesis and characterization of iron(II)-2-aminophenolate complexes with the monoanionic Tp^{Ph2} ligand (Tp^{Ph2} = hydrotris(3,5-diphenyl-pyrazol-1-yl)borate) and neutral Ph²TIP tris(4,5-diphenyl-1-methylimidazol-2-yl)phosphine ligand.^{44–46} The iron(II)-2-aminophenolate complex, supported by the Tp^{Ph2} ligand, when oxidized chemically using one electron oxidant led to the isolation of an iron(II)-2-iminobenzosemiquinone radical intermediate with *S* = 3/2 spin state.⁴⁴ The



Scheme 1 Reactions catalyzed by nonheme enzyme APD and HAD.



Scheme 3 Redox non-innocence behavior of 2-aminophenols.

radical species has relevance to an intermediate proposed in the catalytic cycle of APD. Although the dioxygen reactivity of the Tp^{Ph_2} supported iron(II)-2-aminophenolate complex was not reported, the electronic structure of the intermediate provided mechanistic information on the catalytic cycle of the enzyme.⁴⁴ On the other hand, the dioxygen reactivity of the neutral Ph_2TIP ligand led to the $2e^-$ oxidation process to form $[(\text{Ph}_2\text{TIP})\text{Fe}^{\text{III}}(4,6\text{-di-}t\text{Bu-ISQ})]^+$, where ISQ is 2-iminobenzosemiquinone. Similar $2e^-$ oxidation events of 2-aminophenol in iron complexes of monoanionic N3C donor ligands have been reported by Paine and co-workers. The low-spin complexes do not exhibit aromatic C–C bond cleavage reactions.^{47,48} These studies reveal that the difference in the electronic structure of the biomimetic complexes with a change in ligand backbones apparently controls reactivity, whether a simple $1e^-/2e^-$ oxidation event or the direct addition of O_2 to form the C–C bond cleavage reactions occurs in the biomimetic iron(II)-2-aminophenolate complexes.^{30,44,46} Recently, Fiedler and coworkers have reported the dioxygen reactivity of pentacoordinate cobalt(II)-2-aminophenolate complexes, $[\text{Co}(\text{Tp}^{\text{Me}_2})(4,6\text{-di-}t\text{Bu-HAP})]$ and $[\text{Co}(\text{Tp}^{\text{Ph}_2})(4,6\text{-di-}t\text{Bu-HAP})]$.⁴⁹ Both the $\text{Co}(\text{II})$ -4,6-di-*t*Bu-HAP complexes exhibit hydrogen atom transfer reactivity (HAT) at room temperature upon exposure to dioxygen to form the corresponding $(\text{Tp}^{\text{R}_2})\text{Co}^{\text{II}}$ -2-iminobenzosemiquinone species. However, at low temperature (-70°C) the $[\text{Co}(\text{Tp}^{\text{R}_2})(4,6\text{-di-}t\text{Bu-HAP})]$ complex supported by the Tp^{Me_2} ligand backbone reacts with dioxygen to form an oxygen adduct which was identified as the Co^{III} -superoxo species. The Tp^{Ph_2} supported Co^{II} -4,6-di-*t*Bu-HAP complex on the other hand is unreactive towards dioxygen at lower temperatures suggesting that sterics can modulate the energetics of oxygen binding. Interestingly, the cobalt(II)-2-iminobenzosemiquinone intermediate supported by the Tp^{Me_2} ligand further reacts with dioxygen at reduced temperature to form a Co^{III} -alkylperoxo intermediate bound to the substituted iminobenzosemiquinone ligand. The observed Co^{III} -superoxo as well as the Co^{III} -alkylperoxo species mimic the proposed intermediates in the enzymatic cycle.⁹ In fact the crystal structure of Co^{III} -alkylperoxo resembles the iron-alkylperoxo intermediate reported by Lipscomb in a crystal structure of catechol dioxygenase HPCD.⁵⁰ Unfortunately, none of these synthetic oxygen intermediates led to the C–C bond cleavage of substituted 2-aminophenol, in contrast to the enzymatic system. Thus tuning of stereoelectronic properties of the supporting ligand and 2-aminophenol substrates is one of the key parameters that may be given due consideration while developing models of APD.

With this background, we have explored in this work the dioxygen reactivity of a series of iron(II)-2-aminophenolate complexes $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(\text{X})]$ (X = 2-amino-4-*tert*-butylphenolate (4-*t*Bu-HAP) (1); X = 2-amino-4,6-di-*tert*-butylphenolate (4,6-di-*t*Bu-HAP) (2); X = 2-amino-4-nitrophenolate (4- NO_2 -HAP) (3); and X = 2-anilino-4,6-di-*tert*-butylphenolate (NH-Ph-4,6-di-*t*Bu-HAP) (4) of the facial tridentate nitrogen ligand ($\text{Tp}^{\text{Ph,Me}}$ = hydrotris(3-phenyl-5-methylpyrazol-1-yl)borate) (Chart 1). Another facial N_3 ligand (Tp^{Ph_2} = hydrotris(3,5-diphenylpyrazol-1-yl)borate) (Chart 1) has also been utilized to isolate the corresponding iron(II)-2-anilino-4,6-di-*tert*-butylphenolate complex (5). We have envisioned that the dioxygen activation and subsequent aromatic ring cleavage activity of substituted 2-aminophenols could be controlled *via* fine-tuning of the supporting ligand as well as controlling the electron density on the aromatic ring of 2-aminophenols. The structural tuning of the monoanionic tris(pyrazolyl)borate ligand allows us not only to design close structural mimics but also reveal the functional activity of 2-aminophenol dioxygenases. By using the $\text{Tp}^{\text{Ph,Me}}$ = hydrotris(3-phenyl-5-methylpyrazol-1-yl)borate, which is sterically less congested and at the same time has proven superior activity compared to the Tp^{Ph_2} backbone in biomimetic chemistry,⁵¹ we expected a different reactivity pattern of the corresponding iron(II)-2-aminophenolate complexes toward dioxygen. A comparative study of the dioxygen reactivity of the iron(II)-2-aminophenolate complexes is presented in this work. The results of this study shed light on the effect of the supporting ligand and of the substituted 2-aminophenolate in directing the course of reactivity towards oxygenolytic C–C bond cleavage reactivity *vs.* oxidation of the iron(II)-2-

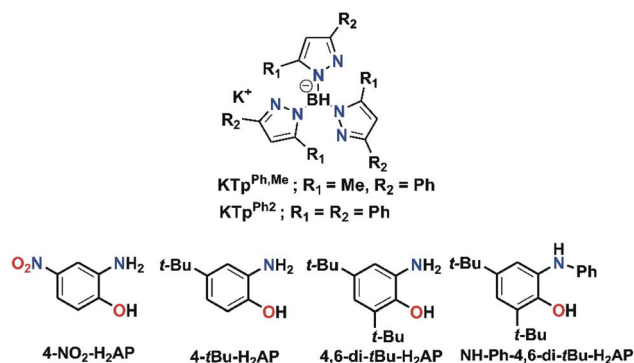


Chart 1 Facial N_3 ligands and substituted 2-aminophenols used in this work.



aminophenolate complexes (1 or 2 electron metal and/or ligand oxidation).

Results and discussion

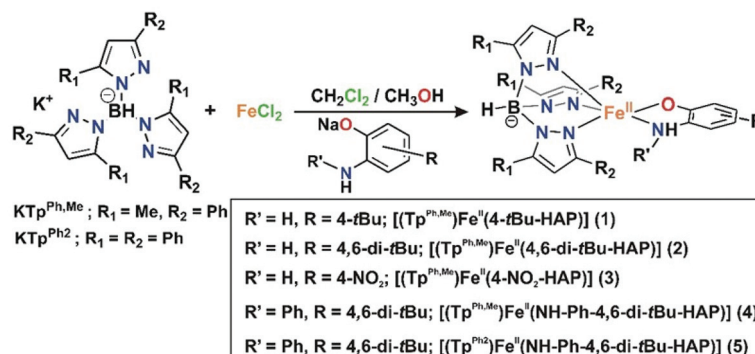
The iron(II)-2-aminophenolate complexes $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(4\text{-}t\text{Bu-HAP})]$ (**1**), $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(4,6\text{-di-}t\text{Bu-HAP})]$ (**2**) and $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(4\text{-NO}_2\text{-HAP})]$ (**3**) were synthesized by mixing equimolar amounts of the ligand ($\text{KTp}^{\text{Ph,Me}}$), iron(II) chloride, and the sodium salt of the respective 2-aminophenol in a dichloromethane-methanol solvent mixture at room temperature under a nitrogen atmosphere (Scheme 4). The complexes $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(\text{NH-Ph-4,6-di-}t\text{Bu-HAP})]$ (**4**) and $[(\text{Tp}^{\text{Ph}_2})\text{Fe}^{\text{II}}(\text{NH-Ph-4,6-di-}t\text{Bu-HAP})]$ (**5**) were isolated from the reaction of the respective ligand ($\text{Tp}^{\text{Ph,Me}}$ for **4** and Tp^{Ph_2} for **5**) with iron(II) salt (iron chloride for **4** and iron perchlorate for **5**) and 2-anilino-4,6-di-*tert*-butylphenol in the presence of triethylamine as a base. Complexes **4** and **5** can also be synthesized by the reaction of the respective ligand ($\text{Tp}^{\text{Ph,Me}}$ for **4** and Tp^{Ph_2} for **5**) with iron(II)-chloride and the sodium salt of 2-anilino-4,6-di-*tert*-butylphenol in a dichloromethane-methanol solvent mixture at room temperature under a nitrogen atmosphere (Scheme 4). Room temperature magnetic moment values ranging between 4.89 and 4.86 μ_{B} indicate the high-spin nature of the complexes (ESI^\dagger). The ^1H NMR spectra of the complexes show relatively sharp, well-resolved, and paramagnetically shifted resonances comparable to other high-spin iron(II) complexes of $\text{Tp}^{\text{R,R'}}$ ligands (ESI^\dagger).^{40,43,51–53}

Crystal structures of **1**, **2**, and **5**

Complexes **1**, **2** and **5** were characterized by single-crystal X-ray diffraction studies. The X-ray structures reveal that each of the neutral iron complexes (**1** and **2**) contains a five-coordinate center coordinated by one $\text{Tp}^{\text{Ph,Me}}$ ligand and one monoanionic 2-aminophenolate. Three nitrogen donors from the ligand ($\text{Tp}^{\text{Ph,Me}}$) coordinate facially to the iron center and the remaining two sites are occupied by the bidentate 2-aminophenolates (Fig. 1). The average Fe–N distance of all the three complexes

is comparable to that of the reported high-spin iron(II) complex tris(pyrazolyl borate) ligands (Tables S3 and S4, ESI^\dagger).^{43,44,51,54,55} In the five-coordinate complexes, two pyrazole nitrogens from the supporting ligand and the phenolate oxygen from 2-aminophenolate unit form the equatorial plane while the amine nitrogen of 2-aminophenol and one pyrazole nitrogen of the ligand occupies the axial positions giving rise to distorted trigonal bipyramidal ($\tau = 0.63$ for **1** and $\tau = 0.69$ for **2**) coordination geometry at the iron center (Fig. 1). The C–C bond distances of the coordinated 2-aminophenolate ring in both **1** and **2** are similar to those of $\text{Tp}^{\text{Ph}_2}\text{Fe}^{\text{II}}(4,6\text{-di-}t\text{Bu-aminophenolate})$ reported by Fiedler and coworkers⁴⁴ supporting the aromatic nature of the 2-aminophenolate ring (Table S3, ESI^\dagger).

In the case of complex **5** (Fig. 2), a five-coordinate iron(II) center coordinated by one Tp^{Ph_2} ligand and one monoanionic NHPh-2-aminophenolate unit is observed. The 2-aminophenolate oxygen O1 binds to the iron center *trans* to the pyrazole nitrogen N3 with the O1–Fe1–N3 angle of $131.46(5)^\circ$ (Table S4, ESI^\dagger). The nitrogen atom N7 of the 2-aminophenolate unit occupies an axial position and is *trans* to the pyrazole nitrogen N5 with an N7–Fe1–N5 angle of $161.55(5)^\circ$. The oxygen atom O1 from 2-aminophenolate and two nitrogen donors (N3 and N1) from the ligand form the equatorial plane of the distorted trigonal bipyramidal complex ($\tau = 0.50$).⁵⁵ Compared to complexes **1** and **2**, the distortion in **5** toward trigonal bipyramid is less. The C–C bond distances in the coordinated 2-aminophenolate moiety indicate the retention of the aromatic character without any quinonoid distortion (Table S4, ESI^\dagger). Although X-ray diffraction quality single crystals could not be isolated for $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(\text{NH-Ph-4,6-di-}t\text{Bu-HAP})]$ (**4**), the structure and binding motif of **4** is expected to be very similar to that of $[(\text{Tp}^{\text{Ph}_2})\text{Fe}^{\text{II}}(\text{NH-Ph-4,6-di-}t\text{Bu-HAP})]$ (**5**) having a coordinated 2-anilino-4,6-di-*tert*-butylphenolate. The spectroscopic and analytical data of **3** suggest the complex to be mononuclear with five-coordinate geometry at the iron(II) center similar to that in **1** and **2**. The binding mode of the 2-aminophenolate complexes (**1**) and (**2**) at the iron(II) centre bears similarity with the reported crystal structure of APD from *Comamonas* sp. strain CNB-1.^{14–17} However, unlike the enzymatic system which has an anionic NNO facial environment around the metal



Scheme 4 Syntheses of iron(II)-2-aminophenolate complexes.



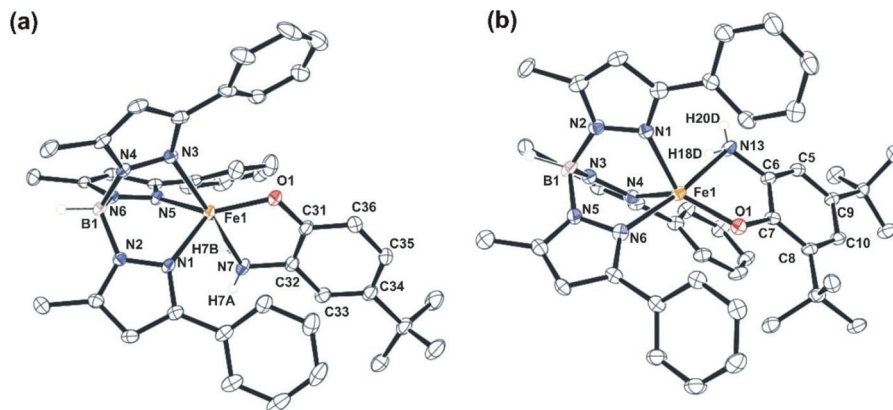


Fig. 1 Molecular structures of the neutral complexes (a) $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(4\text{-tBu-HAP})]$ (1) (All hydrogen atoms except those on N7 and B1 have been omitted for clarity) and (b) $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(4,6\text{-di-tBu-HAP})]$ (2). (All hydrogen atoms except those on N13 and B1 have been omitted for clarity).

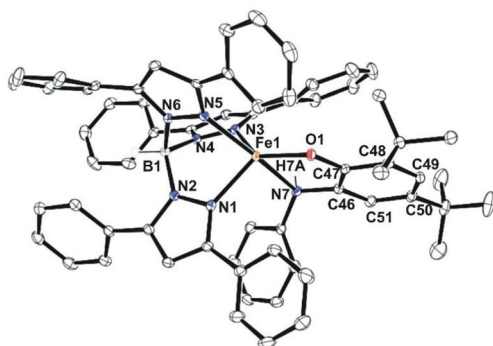


Fig. 2 Molecular structures of $[(\text{Tp}^{\text{Ph}_2})\text{Fe}^{\text{II}}(\text{NH-Ph-4,6-di-tBu-HAP})]$ (5). All hydrogen atoms except those on N7 and B1 have been omitted for clarity.

centre, the complexes (1 and 2) reported here have anionic NNN facial environment and thus serve as close structural mimics of 2-aminophenol dioxygenase.

Reactions of iron(II)-2-aminophenolate complexes with O_2

Dioxygen reactivity of 1 and 2

The iron(II)-2-aminophenolate complex $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(4\text{-tBu-HAP})]$ (1) reacts with O_2 at room temperature, with the formation of two charge-transfer bands at 618 nm and 807 nm within 1 min (Fig. 3). The ESI-mass spectrum of the solution shows a molecular ion peak at $m/z = 702.3$ with the isotope distribution patterns calculated for $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}(4\text{-tBu-AP})]^+$ (Fig. 3, inset). The X-band EPR spectrum of the reaction solution at this point exhibits a rhombic signal at $g = 4.3$, typical of a high-spin iron(III) complex (Fig. S6, ESI†).^{29,30} In the presence of dioxygen, complex 1 is oxidized to its iron(III)-2-amidophenolate form (1^{ox}). In the next step of the reaction, the band at 618 nm shifts to 660 nm and the band at 807 nm shifts to

lower energy to generate a broad band at 900 nm within 2 min (Fig. 3b). The time-dependent EPR spectrum suggests the formation of a ligand-based radical ($g = 1.99$) along with a rhombic signal at $g = 4.3$ (Fig. S6, ESI†).^{29,30} In the final step, the bands at 660 nm and 900 nm slowly decay over a period of 2 h to form a brown-green solution (Fig. 3b). The final EPR solution exhibits no radical signature, but the rhombic signal at $g = 4.3$ remains (Fig. S6, ESI†). The ESI-mass spectrum of the final oxidized solution shows an ion signal at $m/z = 718.3$ with the isotope distribution patterns calculated for $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}(4\text{-tert-butyl-2-picolinate}) + \text{H}]^+$ (Fig. 3b, inset) which shifts by two mass units in the presence of $^{18}\text{O}_2$ (Fig. S9, ESI†).

Exposure of a dichloromethane solution of $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(4,6\text{-di-tBu-HAP})]$ 2 to dioxygen at room temperature immediately results in the formation of a greenish-blue solution. The colour change associated with the reaction of 2 with dioxygen can be monitored by optical spectral change, which indicated a multistep reaction. At first, the greenish blue solution exhibits a broad band at 650 nm typical of iron(III)-2-amidophenolate complexes (Fig. 4a).^{30,47} The X-band EPR spectrum of 2 at 77 K in a dichloromethane-toluene glass shows a rhombic signal at $g = 4.3$ typical of high-spin iron(III) complexes (Fig. 5, right).³⁰ Thus, complex 2 directly converts to its iron(III)-2-amidophenolate (2^{ox}) form unlike the iron(II)-2-iminobenzosemiquinonate intermediate formed with a Tp^{Ph_2} supporting ligand.⁴⁴ However in a subsequent step, a new band appears near 898 nm which decays over a period of 2 h (Fig. 4b) and consequently a sharp signal at $g = 1.99$ is observed presumably due to the formation of a ligand-based radical species in solution (Fig. 5, right).³⁰ Notably, the intensity of the EPR signal at $g = 1.99$ slowly diminishes as the reaction proceeds further. The final step is a slow process where the initial charge-transfer band at 650 nm slowly decays over a period of 5 h to form a greenish brown solution (Fig. 5, left). The final solution displays a rhombic EPR signal at $g = 4.3$ due to the formation of a high-spin iron(III) species (Fig. 5, right).

Interestingly, simulation of the EPR spectra of the time dependent dioxygen reactivity study for both complexes 1 and



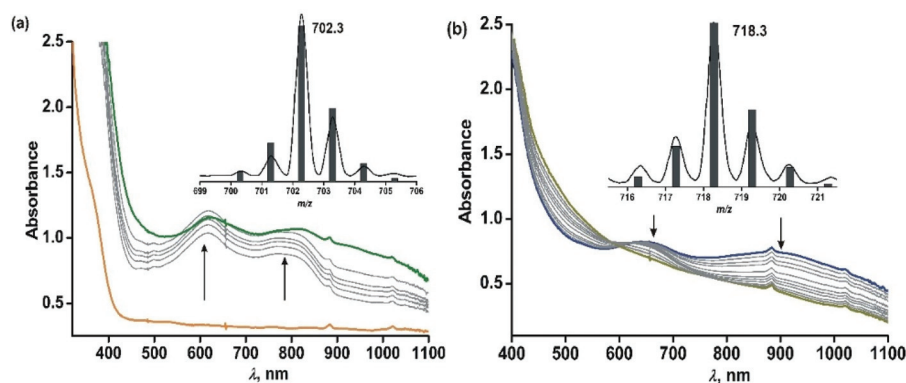


Fig. 3 Time-dependent optical spectral changes of **1** (0.5 mM in dichloromethane, 298 K) during the reaction with dioxygen (a) within 1 min. Inset: ESI-MS of the solution formed after 1 min (b) during the reaction with dioxygen after 1 min. Inset: ESI-MS of the final oxidized solution.

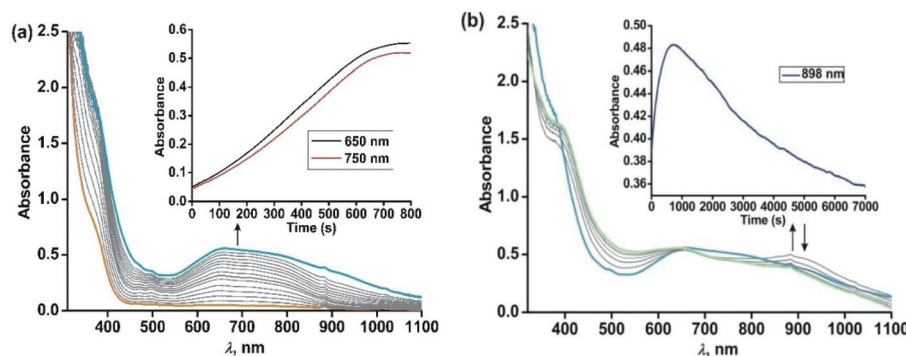


Fig. 4 Time-dependent optical spectral changes of **2** (0.5 mM in dichloromethane) during the reaction with dioxygen at room temperature. (a) Step 1 and (b) step 2. Inset: Plot of absorbance vs. time(s).

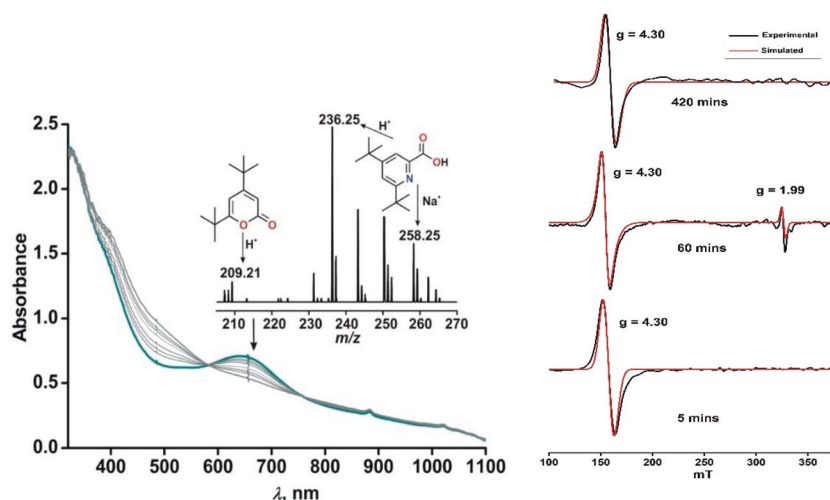


Fig. 5 (Left) Time dependent optical spectral changes of **2** (0.5 mM in dichloromethane) during the reaction with dioxygen at room temperature (step 3). Inset: ESI-MS of the final oxidized solution of **2** after reaction with dioxygen. (Right) X-band EPR spectral changes with time during the reaction of **2** with dioxygen in dichloromethane–toluene glass at 77 K. The red traces show the simulated EPR spectra. For detailed simulation parameters see Table S1.†

2 revealed a significant change in the zero-field splitting ($|D|$) (see the ESI, Table S1†). For paramagnetic systems with $S \geq 1$, a zero-field splitting (ZFS) can often be observed which arises

from dipolar and spin-orbit couplings, causing differences among energy levels in the absence of an external magnetic field. There is still a limited understanding of how ZFS para-



meters relate to the geometric and electronic structures of transition metals compounds, including how metal–ligand bonding affects ZFS.^{56,57} However studies from Neese and co-workers have revealed a correlation between the increase of D with the spectrochemical series showing a decrease of the ligand field strength.⁵⁷ Looking at our zero-field splitting values for the $S = 5/2$ spin system, from the simulated EPR spectra it may be qualitatively predicted that compared to the iron(III)-2-aminophenolate species (1^{ox} and 2^{ox}), the binding of dioxygen changes the coordination geometry and decreases the $|D|$ value. In the final stage of the reaction, complexes show a significant increase in the $|D|$ value, in line with the fact that compared to strongly bound 2-aminophenolate ligand, the C–C bond cleavage results in weakly coordinating iron(III)-2-picolinate complexes, thereby increasing the zero-field splitting.

Analyses of organic products from 1 and 2 and labelling experiments

The final oxidised products obtained from the reaction of 1 with dioxygen was analysed by ^1H NMR spectrum after acidic work-up, maintaining the pH and extraction in diethyl ether and quantification using 1,4-benzoquinone as the internal standard, following the literature procedure.²⁹ A quantitative transformation of 2-amino-4-*tert*-butylphenol to 4-*tert*-butyl-2-picolinic acid was observed in 2.5 h (Scheme 5 and Fig. S10, see the ESI† for details).²⁹ No starting material, 2-amino-4-*tert*-butylphenol, was detected. The ESI-MS of the reaction solution after work up shows the peak at $m/z = 180.21$ which corresponds to the distribution pattern calculated for $[4\text{-}t\text{-butyl-2-picolinic acid} + \text{H}^+]$ (Fig. S7, ESI†). The GC-mass spectrum of the organic product (2-picolinic acid was esterified with diazomethane, see ESI†) shows a distinct ion peak at $m/z = 193$ with the expected fragmentation patterns for methyl-4-*tert*-butyl-2-picolinate (Fig. S8a, ESI†). The ion peak at $m/z = 193$ is shifted to 195 when the reaction is performed with $^{18}\text{O}_2$ supporting the origin of oxygen atoms from dioxygen (Fig. S8b, ESI†).

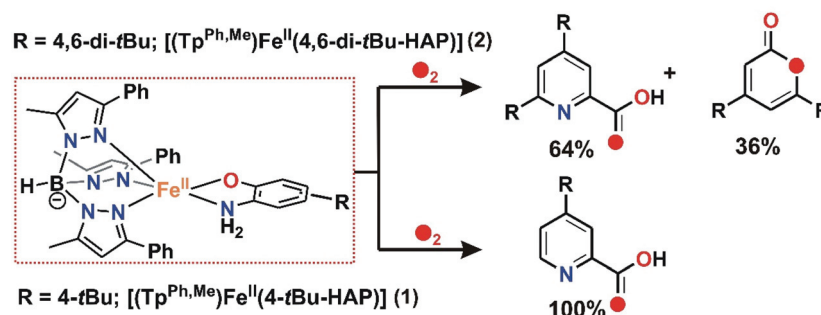
The oxidized products derived from 2 were similarly extracted into an organic solvent after acid treatment, following the literature procedure,^{30,31} and the organic products were analyzed by ESI-MS and GC-MS and quantified by ^1H NMR spectroscopy using 1,4-benzoquinone as the internal

standard (see the ESI†). The ^1H NMR spectrum derived from the dioxygen reactivity of 2 reveals that 2-amino-4,6-di-*tert*-butylphenol quantitatively transforms to a mixture of C–C bond cleavage products, 4,6-di-*tert*-butyl-2H-pyran-2-one (36%) and 4,6-di-*tert*-butyl-2-picolinic acid (64%) (Scheme 5 and Fig. S13†).^{30,31} The ESI-MS shows distinct ion peaks at $m/z = 209.21$ which correspond to the distribution pattern calculated for $[4,6\text{-di-}t\text{-butyl-2H-pyran-2-one} + \text{H}^+]$ and at $m/z = 236.35$ and 258.25 for $[4,6\text{-di-}t\text{-butyl-2-picolinate} + \text{H}^+]$ and $[4,6\text{-di-}t\text{-butyl-2-picolinate} + \text{Na}^+]$, respectively (Fig. S11, ESI†). The GC-mass spectrum of organic products (2-picolinic acid was esterified with diazomethane) shows two distinct ion peaks at $m/z = 208$ and 249 with the expected fragmentation patterns of 4,6-di-*tert*-butyl-2H-pyran-2-one and methyl-4,6-di-*tert*-butyl-2-picolinate (Fig. S12a, ESI†). When the reaction is carried out with $^{18}\text{O}_2$, the ion peak at $m/z = 208$ is shifted two mass units higher to $m/z = 210$ and the peak at $m/z = 249$ is shifted to $m/z = 251$ (Fig. S12b, ESI†). The labeling experiment confirms the incorporation of one ^{18}O atom into each of the cleavage products.

Thus, both the substituted iron(II)-2-aminophenolate complexes (1 and 2) react with dioxygen to quantitatively form the C–C bond cleavage product of the corresponding 2-aminophenols. For iron(II)-2-amino-4-*tert*-butylphenolate complex (1), 4-*tert*-butyl-2-picolinic acid is observed as the sole C–C bond cleavage product after the reaction with molecular oxygen. On the other hand, the reaction of iron(II)-2-amino-4,6-di-*tert*-butylphenolate complex (2) with oxygen results in a mixture of 4,6-di-*tert*-butyl-2H-pyran-2-imine and 4,6-di-*tert*-butyl-2-picolinic acid. However, in both the cases, the products are found to be regioselective and result from the C1–C6 bond cleavage of the respective 2-aminophenols upon dioxygen activation (Scheme 5).

Dioxygen reactivity of 3

For the $-\text{NO}_2$ substituted 2-aminophenol, complex $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(4\text{-NO}_2\text{-HAP})]$ (3), the light yellow solution in dichloromethane rapidly transforms to a greenish-blue species (3a) upon exposure to dioxygen. During the reaction, two broad bands at 530 nm and 720 nm, characteristic of the LMCT transitions of iron(III)-2-aminophenolate species (Fig. 6), are generated over a period of 2 h. The X-band EPR spectrum of the



Scheme 5 Aromatic ring cleavage products of substituted 2-aminophenolates during the reaction of 1 and 2 with dioxygen.



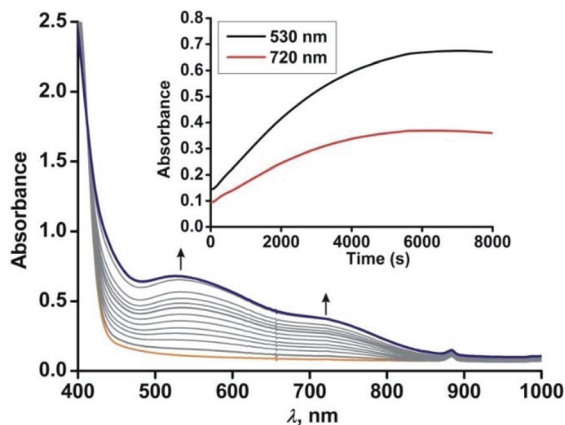


Fig. 6 Optical spectral changes with time during the reaction of **3** (0.5 mM solution in dichloromethane) with dioxygen at 298 K. Inset: Plot of absorbance vs. time.

final oxidized solution exhibits an intense rhombic signal at $g = 4.3$ (Fig. S14, ESI†). Similar to the iron(II)-catecholate complexes,^{38,55} iron(II)-2-aminophenolate complexes are converted to the corresponding iron(III)-2-amidophenolate species (Scheme 6). The intensity and position of the ligand-to-metal charge-transfer bands of **3a** remain unchanged even after several hours under an oxygen atmosphere. Analyses of the organic product by ^1H NMR after acidic workup of the bluish-green solution reveal that 4- NO_2 -aminophenol is quantitatively recovered without the formation of any C–C cleavage product.

Role of $-\text{NH}_2$ group in the C–C bond cleavage reactivity of iron(II)-2-amino-phenolate complexes

Both the iron(II)-2-aminophenolate complexes **4** and **5** with $-\text{NH}-\text{Ph}$ groups instead of $-\text{NH}_2$ are sensitive towards dioxygen in solution. The light yellow solution of $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(\text{NH}-\text{Ph}-4,6\text{-di-}t\text{Bu-HAP})]$ (**4**) in dichloromethane rapidly transforms into greenish-blue species (**4a**) upon exposure to dioxygen. Initially, two broad peaks at 680 nm and 800 nm are generated rapidly whose feature and absorbance match with the reported iron(II)-2-iminobenzosemiquinonate species.⁴⁴ However, the two broad peaks merge extremely fast and during the sub-

sequent course of reaction of **4** with dioxygen, two prominent bands at 440 nm and 750 nm (Fig. 7) generate over a period of 4 h. The time dependent EPR spectrum reveals the formation of an intense rhombic signal at $g = 4.3$ for a high-spin iron(III) species after 5 min. However, with time, the intensity of the $g = 4.3$ signal decreases and the final oxidized solution after 4 h becomes EPR silent (Fig. S15, ESI†). The intensity and position of the optical transition of **4a** remain unchanged even after several hours under an oxygen atmosphere. No oxidative C–C bond cleavage of 2-anilino-4,6-di-*tert*-butylphenol is observed upon the reaction of **4** with dioxygen. Thus, in the presence of dioxygen, complex **4** undergoes two electron oxidation on the redox non-innocent 2-aminophenol to form an EPR silent iron (III)-2-iminobenzosemiquinonate species (Scheme 7 and Fig. S15, ESI†).^{32,46–48} However, attempts to crystallize the final oxidized solution were unsuccessful.

Similar to complex **4**, complex $[(\text{Tp}^{\text{Ph}_2})\text{Fe}^{\text{II}}(\text{NH}-\text{Ph}-4,6\text{-di-}t\text{Bu-HAP})]$ (**5**) is reactive towards dioxygen. However, changing the ligand to Tp^{Ph_2} , from $\text{Tp}^{\text{Ph,Me}}$, enables to monitor three distinct oxidation steps, as evident from the optical spectrum. Upon exposure to dioxygen, the yellow solution of **5** immediately turns green. Three charge transfer transitions at 510 nm, 680 nm and 850 nm arise within 3 min (Fig. 8). The X-band EPR spectrum of the oxidized solution at this point shows an intense rhombic signal at $g = 4.3$ (Fig. S16, ESI†). The spectral data suggest the formation of an iron(III)-2-amidophenolate species. During the next steps of the reaction under an oxygen atmosphere, the charge-transfer transitions at 680 nm and 850 nm decay over a period of 3 h to form a single broad band

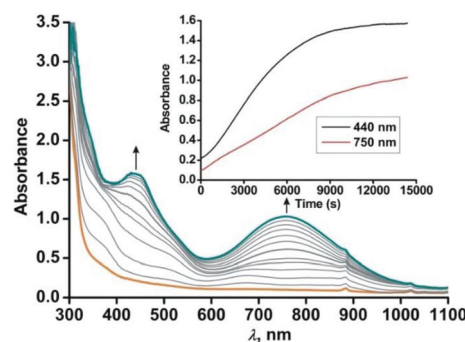
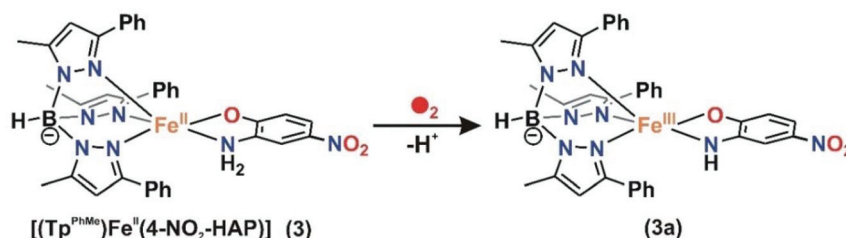
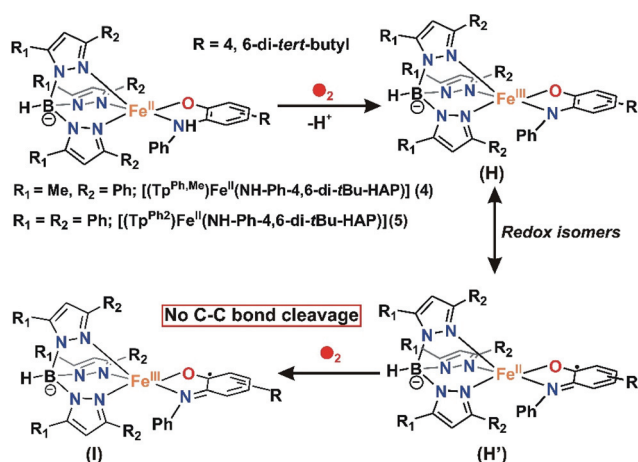


Fig. 7 Optical spectral changes of **4** (0.5 mM in dichloromethane) during the reaction with dioxygen at 298 K. Inset: Plot of absorbance vs. time.



Scheme 6 Reaction of $[(\text{Tp}^{\text{Ph,Me}})\text{Fe}^{\text{II}}(4\text{-NO}_2\text{-HAP})]$ (**3**) with dioxygen.





Scheme 7 Proposed mechanism for the reaction of 4 and 5 with dioxygen.

at around 800 nm (Fig. 8). The molar extinction coefficient and position of the optical transition after 3 h resemble that of the optical spectrum of the chemically one electron oxidized form of $[\text{Tp}^{\text{Ph2}}\text{Fe}^{\text{II}}(2\text{-amino-4,6-di-}tert\text{-butylphenol})]$, i.e. an iron(II)-2-iminobenzosemiquinonate species reported by Fiedler *et al.*⁴⁴

However, the X-band EPR spectrum at 77 K reveals signals at $g = 1.99$ possibly due to the presence of a ligand-based radical species in solution along with a small peak at $g = 4.3$ (Fig. S16, ESI†).³⁰ No peak corresponding to the $S = 3/2$ spin state was observed at 77 K. Attempts to isolate the species formed at this point, lead to the isolation of greenish blue crystals, which were of extremely poor quality for exact structure determination. Finally, the broad band at 800 nm undergoes a blue-shift to 750 nm and slowly increases in intensity over a period of 7 h (Fig. 9) and remains stable for several days. The final oxidized solution is EPR silent.^{32,46–48} Such a stable UV-vis peak at the higher wavelength region of around 750 nm with a molar absorption coefficient of approx. $4000 \text{ M}^{-1} \text{ cm}^{-1}$ is characteristic for the literature reported iron(III)-2-iminobenzosemiquinone species.^{44,46} Reports from Paine and coworkers as well as Fiedler and coworkers have revealed such iron(III)-iminobenzosemiquinone species to be EPR silent likely due to

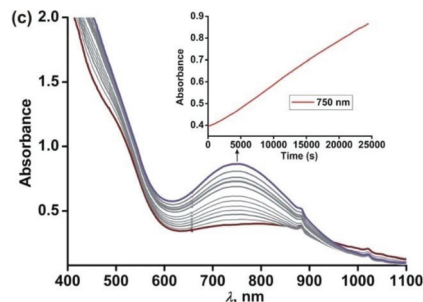


Fig. 9 Optical spectral changes of 5 (0.5 mM in dichloromethane) during the reaction with dioxygen at 298 K over a period of 7 h (step 3). Inset: Plot of absorbance vs. time.

the coupling between iron(III) and iminobenzosemiquinone ligand based radicals, making it overall an integer spin EPR silent species.^{32,46–48} The iron(II)-2-anilino-4,6-di-*tert*-butylphenolate complex, thus undergoes two electron oxidation *via* its iron(II)-2-iminobenzosemiquinonate form. No oxidative C–C bond cleavage of 2-anilino-4,6-di-*tert*-butylphenol was observed in the reaction of 5 with dioxygen.

Despite sufficient electron density on the aromatic ring for 2-anilino-4,6-di-*tert*-butylphenol in complexes 4 and 5, they undergo two electron oxidation of the redox non-innocent ligand,^{44,47} instead of showing C–C bond cleavage reactivity. This demonstrates the importance of the $-\text{NH}_2$ group in substituted 2-aminophenols in displaying C–C bond cleavage reactions. The Fe(II)-2-iminobenzosemiquinonate species (H') reacts with dioxygen and the lack of the $-\text{NH}_2$ group directs the course of reactivity towards further oxidation on the metal center to form Fe(III)-2-iminobenzosemiquinonate complex (I)^{44,46} and no further reaction with dioxygen occurs (Scheme 7). However, Mukherjee and coworkers have shown an example of an iron complex of 2-aminophenol-appended supporting ligand where reaction with dioxygen resulted in a dimeric iron(III)₂ oxo-bridged complex coordinated by a furan derivative in each metal unit, which was proposed to be formed as a result of the oxidative aromatic C–C bond cleavage product of 2-aminophenol.⁵⁸

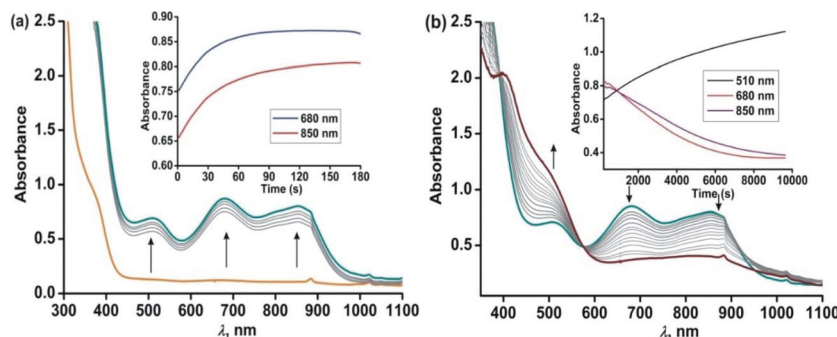


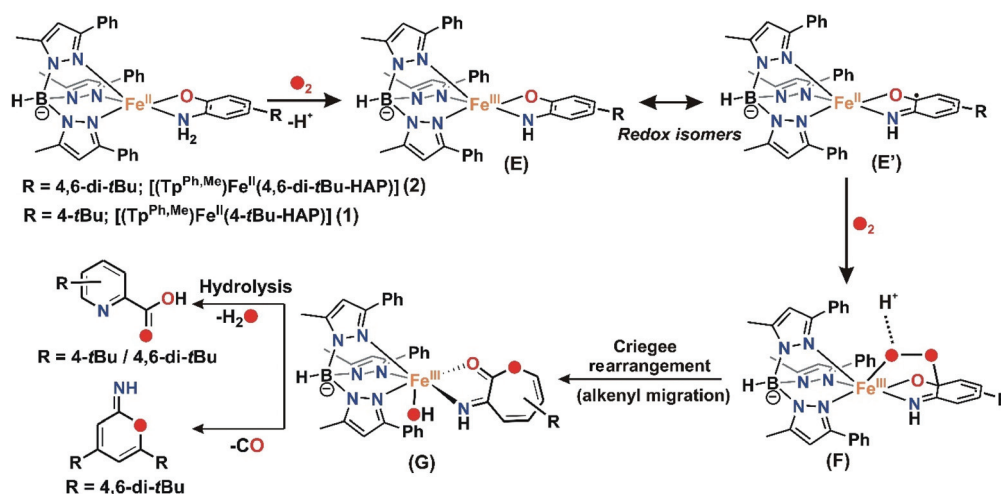
Fig. 8 Optical spectral changes of 5 (0.5 mM in dichloromethane) during the reaction with dioxygen at room temperature over a period of (a) 2 min (step 1) and (b) over a period of 3 h (step 2). Inset: Plot of absorbance vs. time.

Discussion and proposed C–C bond cleavage mechanism of iron(II)-2-aminophenolate complexes

For iron(II)-catecholate model complexes, an outer sphere one electron oxidation of iron(II)-catecholate to iron(III)-catecholate species have been reported by Que and Moro-oka.^{6,38} The activation of dioxygen is initiated at the iron(II)-semiquinone intermediate (a redox isomer of iron(III)-catecholate) to form an iron(III)-alkylperoxo intermediate for the C–C bond cleavage of catechol. Paine and coworkers reported the first functional model of 2-aminophenol dioxygenases supported by a tetradentate tripodal ligand system that undergoes C–C bond cleavage *via* an iron(III)-2-amidophenolate species.^{29,30} The reactivity (and stability) of the intermediate, which directs the course of the aromatic ring cleavage reaction, was reported to be dependent on the nature of ring substituent on 2-aminophenols.³⁰ Fiedler and co-workers have isolated and structurally characterized an iron(II)-2-aminophenolate complex,^{44–46} $[(\text{Tp}^{\text{Ph}_2})\text{Fe}^{\text{II}}(4,6\text{-di-}t\text{-butylaminophenol})]$, which was converted to an iron(II)-2-iminobenzosemiquinonato radical species upon one-electron chemical oxidation. On the basis of our experimental observations and literature precedents, it is proposed that complexes **1**, **2** and **3** with $\text{Tp}^{\text{Ph},\text{Me}}$ as the supporting ligand react with dioxygen to undergo one-electron oxidation to form an iron(III)-2-amidophenolate species in the first step (Schemes 6 and 8). The presence of an electron withdrawing nitro group at the 4-position of the 2-aminophenolate ring in **3** makes it unreactive towards dioxygen. Only electron transfer *via* an outer sphere mechanism operates and so complex **3** becomes stable in its iron(III)-2-amidophenolate (**3a**) form and does not react further with dioxygen (Scheme 6).³⁰ However, the incorporation of electron donating groups on the aromatic ring of 2-aminophenol (*tert*-butyl) for complex **1** and **2** results in the oxygenolytic aromatic ring cleavage reaction.

The proper steric geometry provided by the $\text{Tp}^{\text{Ph},\text{Me}}$ ligand along with appropriate electron density on the aromatic ring of 2-aminophenols triggers the iron(II)-iminobenzosemiquinonato species (**E**), a redox isomer of iron(III)-2-amidophenolate species (**E'**) from **1** and **2** to react with dioxygen and generate an iron(III)-alkyl peroxo intermediate (**F**). The iron(III)-alkyl peroxo intermediate undergoes an alkenyl Criegee rearrangement, ultimately leading to the regioselective C–C bond cleavage of substituted 2-aminophenols (Scheme 8). However, the isolation and stabilization of putative species like iron(II)-2-iminobenzosemiquinonato under an oxygen environment (unlike chemical oxidation) is extremely challenging. The reported models (**1** and **2**) undergo regioselective C–C bond cleavage of substituted 2-aminophenols at a rate much faster than those reported for the tetradentate N_4 system by Paine and coworkers.^{29,30} Whereas complex **1** takes only 2.5 h to undergo the C–C bond cleavage of 4-*t*Bu-HAP upon reaction with dioxygen, the corresponding iron(II)-2-aminophenolate complex of the tetradentate ligand 6Me₃TPA takes around 6 h for the reaction. Similarly, there is a three-fold increase in the rate of reaction for C–C bond cleavage of 4,6-di-*t*Bu-HAP by complex **2** (7 h) compared to the iron(II)-4,6-di-*t*Bu-HAP complex supported by the 6Me₃TPA ligand (22 h). Though we were unable to isolate any reactive intermediate, it might perhaps not be hyperbolic to term the complexes (**1** and **2**) as first models simultaneously resembling the structural motif as well as emulating the functional reactivity of 2-aminophenol dioxygenases.

So far from this study and previous literature reports (see Table S5† for a comparative analysis of dioxygen reactivity of reported 2-aminophenol model complexes), it is evident that unlike catechol cleaving dioxygenases, the development of the functional models of 2-aminophenol dioxygenase is more challenging. It is instructive from both the experimental and theoretical studies that differences in the denticity of the supporting ligand backbone, electronic factors of both the supporting ligands and the substituted 2-aminophenols as well as



Scheme 8 Proposed mechanism for the C–C bond cleavage of 2-aminophenolates by biomimetic iron(II) complexes **1** and **2** using O₂ as the oxidant.



the proton shuttle in the reaction medium and probable second sphere residues exhibit a remarkable effect on the ring cleaving activity of model 2-aminophenol complexes.^{30–32,59} Whereas $[(\text{Tp}^{\text{Ph}_2})\text{Fe}^{\text{II}}4,6\text{-di-}t\text{Bu-HAP}]$ or $[(\text{TIP}^{\text{Ph}_2})\text{Fe}^{\text{II}}4,6\text{-di-}t\text{Bu-HAP}]$ undergoes chemical one-electron oxidation or reaction with dioxygen and undergo the HAT reaction to form the corresponding $\text{Fe}^{\text{II}}\text{-2-iminosemibenzoquinone}$ species, these complexes do not show the C–C bond cleavage reactivity of substituted 2-aminophenols, likely due to the steric and/or electronic factors of the supporting ligands.⁴⁹ For the $\text{Co}^{\text{II}}\text{-2-iminosemibenzoquinone}$ species formed by HAT upon exposure to the dioxygen of $\text{Tp}^{\text{R}_2}\text{Co}^{\text{II}}4,6\text{-di-}t\text{Bu-HAP}$ ($\text{R}_2 = \text{Me}_2$ or $\text{R}_2 = \text{Ph}_2$), dioxygen adduct formation was hindered for the sterically bulky Ph_2 substituted ligand compared to its Me_2 congener, highlighting once again the importance of steric effect in modulating the energetics of oxygen binding. Thus, in this work, the success of the C–C bond cleavage of substituted 2-aminophenols was governed by both the steric effect and electronics of the supporting ligand $\text{Tp}^{\text{Ph,Me}}$ and also the electron density on the 2-aminophenolate ring (reactivity of $4\text{-}t\text{Bu-H}_2\text{AP} > 4,6\text{-di-}t\text{Bu-H}_2\text{AP} > 4\text{-NO}_2\text{-H}_2\text{AP}$). Furthermore, for iron(II) complexes **4** and **5**, despite having similar facial N_3 coordination motif by the supporting ligands (Tp^{Ph_2} and $\text{Tp}^{\text{Ph,Me}}$) around the metal centre, the lack of a $-\text{NH}_2$ group in coordinated 2-anilino-4,6-di-*tert*-butylphenolate directs the course of reactivity towards the two-electron oxidised $\text{Fe}(\text{III})\text{-2-iminobenzosemiquinonate}$ complex instead of undergoing C–C bond cleavage. Thus a synchronized way of both proton transfer (PT) and O_2 binding along with the proper control of stereoelectronic factors are perhaps the keys for the oxygenolytic scission of C–C bond of substituted 2-aminophenols. The efficient management of these events is well executed in the enzymatic system in harmony with the active site and the conserved second sphere residues.

Conclusion

In conclusion, we have synthesized a series of iron(II)-2-aminophenolate complexes supported by tridentate monoanionic facial ligands which are all reactive towards dioxygen. The works presented herein show how the electronic and structural properties of 2-aminophenolate rings as well as the supporting ligand influence the C–C bond cleavage of 2-aminophenols and also highlight the role of $-\text{NH}_2$ group in controlling the oxidative C–C bond cleavage reactivity. This work represents the first closest structural mimic of 2-aminophenol dioxygenases which exhibits excellent functional selectivity of the enzyme to regioselectively cleave the C1–C6 bond of substituted 2-aminophenols. Future directions would aim to explore detailed spectroscopic and theoretical calculations to evaluate the subtle changes in the geometric and electronic factors that modulate the dioxygen reactivity pathway from one or two electron oxidation events to regioselective C–C bond cleavage of 2-aminophenols in model complexes.

Experimental section

See ESI.†

Conflicts of interest

There are no conflicts to declare.

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