Set of Fe(II)-3-Hydroxyflavonolate Enzyme–Substrate Model Complexes of Atypically Coordinated Mononuclear Non-Heme Fe(II)-Dependent Quercetin 2,4-Dioxygenase

Ying-Ji Sun, Qian-Qian Huang, and Jian-Jun Zhang

School of Chemistry, Dalian University of Technology, 2 Linggong Road, Dalian 116024, China

ABSTRACT: With the aim of revealing the catalytic role of atypically coordinated (3His-1Glu) active site mononuclear non-heme Fe(II)-dependent quercetin 2,4-dioxygenase (Fe-2,4-QD) and the electronic effects of the model ligands on the reactivity toward dioxygen, a set of p/m-R-substituted carboxylate-containing ligand-supported Fe(II)-3-hydroxyflavonolate complexes, [FeL₅(flav)] (L₅H₂: 2-[(bis(pyridin-2-ylmethyl)amino)methyl]benzoic acid; R: p-OMe (1), p-Br (2), p-NO₂ (5); flav: 3-hydroxyflavonolate), were synthesized and characterized as structural and functional models for the ES (enzyme–substrate) complex of Fe-2,4-QD. [FeL₅(flav)] show relatively high enzyme-type reactivity (dioxygenative ring opening of the coordinated substrate flav, single-turnover reaction) at low temperatures (30–65 °C). The reaction shows a linear Hammett plot (ρ = −1.21), and electron donating groups enhance the reaction rates. The notable difference on the reactivity can be rationalized from the electronic nature of the substituent in the ligands, which could tune the reactivity via tuning Lewis acidity of the Fe(II) ion, electron density, and the redox potential of flav. The properties and the reactivity show approximately linear correlations between λmax or E1/2 of flav and the reaction rate constant k. This work sheds light not only on understanding of electronic effects of the ligands and the property–reactivity relationship but also on the role of the catalytic reaction by Fe-2,4-QD.

INTRODUCTION

Mononuclear non-heme Fe(II)-dependent enzymes (MNHEs) act pivotal roles in the O₂ metabolism, catalyze an amazing variety of oxidative reactions, and have potential medical and pharmaceutical applications. Generally, the Fe(II) ion in the active sites of MNHEs is coordinated by two histidine imidazoles and a carboxylate group of Glu or Asp in a “facial triad” mode. However, atypical coordination modes, such as the three histidine imidazoles 3His mode [as observed in diketone dioxygenase (Dke1)³ and cysteine dioxygenase (CDO)⁴], or three histidine imidazoles and a carboxylate 3His-1Glu mode [as observed in acireductone dioxygenase (ARD)⁵ and quercetin 2,4-dioxygenase (2,4-QD)⁶], were recently reported. The active site structure–function relationship of different active site MNHEs is very interesting and fascinating. Compared with the well-studied typically coordinated MNHEs, biomimetic research of the atypically coordinated MNHEs, especially those bearing the 3His-1Glu mode and the structure–function relationship of different active site MNHEs, remains largely undeveloped.

In the initial step of the quercetin catabolism, 2,4-QD activates dioxygen to catalyze the oxygenative 0-heterocycle ring opening reaction of flavonoid quercetin (3′,4′,5,7-tetrahydroxyflavonol, QUE) to the corresponding phenolic carboxylic acid esters and release carbon monoxide (Scheme 1).⁶

Scheme 1. Reaction of QUE with Dioxygen Catalyzed by 2,4-QD

The fungal 2,4-QD⁷–⁹ is the only known Cu(II)-dependent dioxygenase (Cu-2,4-QD) and has a mononuclear type II Cu(II) reaction center. The active site of 2,4-QD from Aspergillus japonicas comprises two distinct coordination environments. In the first case, the Cu(II) is coordinated by a carboxylate group of Glu73, three histidine imidazoles, and one water molecule to form a distorted trigonal-bipyramidal geometry, whereas another exhibits a distorted tetrahedral geometry without the binding of Glu73. In the enzyme–substrate (ES) complex, the Cu(II) active site shows a distorted square pyramidal geometry, in which a deprotonated substrate 3-hydroxyflavonolate (flav) is bound to Cu(II) via the 3-hydroxy group with the displacement of the water molecule.⑩ The...
mutation of Glu73 induces a loss of enzyme activity, implying that the Glu73 carboxylate group acts an essential catalytic role.

The bacterial 2,4-QD from Bacillus subtilis is an Fe II-dependent dioxygenase (Fe-2,4-QD). It has an atypically coordinated mononuclear non-heme Fe(II) active site, which also exhibits two distinct structures similar to that of Cu-2,4-QD. In the N- or C-terminal cupin motif, the Fe(II) active site shows a distorted trigonal bipyramidal or a square pyramidal geometry involving three histidine imidazoles, a water molecule, and one Glu69 (2.10 Å) or Glu241 (2.44 Å, weakly coordinated).

So far, a lot of Cu-2,4-QD models have been reported, though other metal-based models, especially those containing iron, are still rare. Currently, only five structurally characterized Fe-2,4-QD complexes—[Fe III(4′-MeOfl)2] (Figure 1a), 17 [Fe III(4′-fl)2] (Figure 1b), 17 [Fe III(L1)salen] H2 (L1: 1,6-bis-(2-hydroxyphenyl)-2,5-diaza-hexa-1,5-diene) (Figure 1c), 21a and [Fe III(L4)salen] H2 (L4: 2-[[bis-(pyridin-2-ylmethyl)amino]methyl]benzoic acid) (Figure 1d) 21a—have been reported, though their oxidation states are not the same as that in the ES adduct of the FeII-2,4-QD except [Fe III(L4)salen]. There are some reports about the biomimetic studies that focus on the electronic substituent effects of the substrate fla on the reactivity. Generally, the active site residues of the enzyme and their substituent groups, with the aid from the second coordination sphere, are important factors to stabilize the intermediates, tune the catalytic activity, and the conformation of the enzyme via influencing the coordination geometry of the metal cofactor. There are some reports about the biomimetic studies that focus on the electronic substituent effects of the substrate fla on the reactivity. However, as we know, there are only two reports that focus on the electronic effects of the substituent group of the model ligands of Co(II) and Ni(II) ES model complexes on their reactivities. No report focused on the electronic effects of the substituent group in the model ligands of Fe(II) ES model complexes on their reactivities has been published yet.

With an aim to investigate the electronic substituent effects of the model ligands on the chemical functions of Fe-2,4-QD and the active site structure—function relationship of different active site MNHEs, we synthesized and characterized a set of structural and functional FeII-2,4-QD ES model complexes [Fe III(L4)salen] H2 (R: p-OMe (1), p-Me (2), m-Br (4), and m-NO2 (5) (Figure 1d) (3) has already been reported in our previous paper 21a) of Fe-2,4-QD using p/m-substituted carboxylic-acid-containing ligands. A detailed investigation on their structures, physicochemical properties, electrochemical properties, and dioxygenation reactivity is presented.

RESULTS AND DISCUSSION

Synthesis and Structural Characterization. The ligands L4H (Figure 1d) were synthesized based on the reported methods. The reaction of FeII(OAc)2 with the neutral ligand L4H and flavonol (flaH) under N2 gave the corresponding complexes [FeII(L4)salen] H2 (R: p-OMe (1), p-Me (2), m-Br (4), and m-NO2 (5)). No additional base was added during the reaction, and the acetate ions (AcO−) were proposed to act as proton acceptors. All complexes are relatively stable under air in the solid state, but very sensitive to air in solution (see below).

Thus, their single crystals suitable for X-ray crystallographic analysis cannot be obtained successfully. Only a single crystal of one of the oxidized dioxygenation products, salicylate (sal) complex [FeIII(L4)salen] H2O (1), was obtained during recrystallization of [FeIII(L4)salen] H2O (1) by slow diffusion of ether into the dichloromethane solution of 1 without N2 protection at room temperature. The structure is shown in Figure 2. The crystallographic data and the selected bond lengths and angles are summarized in Tables S1 and S2, respectively.
Figure 2. ORTEP (50% ellipsoid) representation of [FeII(LOMe(sal))]: H2O (1'). For clarity, hydrogen atoms and free solvent molecules are omitted.

1' crystallizes in the monoclinic P2(1)/n space group. It has a mononuclear structure, and the FeIII ion is coordinated by O(4) (carboxylate) and O(6) (2-hydroxylate) of sal, O(1) (benzolate) of LOMe, and three nitrogen atoms of LOMe to form a distorted octahedral geometry. The axial positions are occupied by O(6) and N(3) atoms (amine nitrogen atom).

The bond lengths of Fe−O(1) (benzolate of LOMe) is 1.940(3) Å, which is shorter than those of the resting Fe−2.4-QD (2.10 Å) and the nonsubstituted derivative [FeIII(L)(fla)] (2.0256(18) Å), but close to that of [FeIIi(fl)(L)][ClO4] (1.909(5) Å). The bond lengths of Fe−O(4) and Fe−O(6) (oxygen atoms of salicylate) are 1.962(3) and 1.878(3) Å, 1.940(3) Å, which is shorter than those of the resting FeII-2,4-QD (2.16 Å), which is consistent with its high enzyme−activity. The average Fe−O bond length is 1.93 Å, which is similar to that of [FeIIi(fl)(L)][ClO4] (1.97 Å). The average Fe−N bond length is 2.16 Å, which is also close to those of [FeIIIi(fl)(L)][ClO4] (2.15 Å) and [(6-Ph2TPAFeIIi(fl))2(μ-O)](ClO4)2 (2.21 Å).

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The eventually produced crystals of the enzymes of dioxygenation reaction product sal complex [FeIIIi(OMe(sal))]: H2O (1') during recrystallization of [FeIIIi(OMe(sal))] (1) without O2 protection at room temperature are likely an indication that [FeIIIi(OMe(sal))] (1) very easily reacts with O2 even under air at room temperature, which is consistent with its high enzyme−type reactivity (see below).

Although suitable single crystals of [FeIIi(L)(fla)] cannot be obtained, they are proposed to have a mononuclear structure bearing a distorted octahedral geometry, similar to that of the nonsubstituted derivative [FeIIi(L)(fla)]22 as proved by Fourier-transform infrared spectroscopy (FT-IR), ESI-MS, and electron paramagnetic resonance (EPR) spectroscopic results that will be depicted below.

**Spectroscopic and Redox Properties of the Complexes.**

**Infrared Spectroscopy.** The FT-IR spectra of both solid and ethanol solution state samples of the complexes were measured (Table 1, Supporting Information, Table S3 and Figure S1). As a typical example, the FT-IR spectra of 1 in both states are shown in Figure S1. The spectra between the ethanol solution and solid state samples of each complex are quite similar, suggesting that all complexes keep their mononuclear structure [FeIIi(L)(fla)] in solution, which coincides with ESI-MS and EPR results (see below). The ν(C=O) (C=O stretching vibration) of the bound carbonyl group of fla appears around 1550 cm−1 (ca. 50 cm−1 red-shifted compared with free flavon (1602 cm−1)), which is close to those of reported fla complexes.23−26 In both states, each complex displays the asymmetric νas(COO−) and symmetric νs(COO−) stretching frequencies of the carbonylate group of L1 around 1615 and 1420 cm−1, respectively. The difference Δν between νas(COO−) and νs(COO−) is in the range of 190−200 cm−1, which is a clear evidence to prove that the carbonylate group of ligand in all complexes has a monodentate coordination mode in both states.

**UV-vis Spectroscopy.** All complexes display an intense π−π* transition band of the coordinated fla30 in dimethylformamide (DMF) around 400 nm (ε = 9.59 × 104 M−1 cm−1) for 1, 399 nm (ε = 1.03 × 104 M−1 cm−1) for 2, 396 nm (ε = 1.16 × 104 M−1 cm−1) for 4, and 394 nm (ε = 1.03 × 104 M−1 cm−1) for 5 (Figure 3a and Table 1). The εmax values are comparable with that of [FeIIi(L)(fla)] (3) (398 nm)21a but blue-shifted 2−17 nm than those of models [FeIIi(L)(fla)] (402 nm),32 [FeIIi(MeOfla)] (411 nm),32 [FeIIi(fl)(salen)] (407 nm),19 and [(6-Ph2TPAFeIIi(fl))][ClO4] (415 nm)18, which could be explained by a combination effect of the carbohydrate negative charge in the model ligands and a lower charge of the FeII versus FeIII ion. Compared with free fla (458 nm for Me4Nfla32 and 465 nm for Kfla32), all εmax values are blue-shifted (ca. 60 nm), which are similar to those of the enzymatic25,34,35,36 and other synthetic model systems.37−39

Besides, an order of −OMe (1) > −Me (2) > −H (3)21a > −Br (4) > −NO2 (5) was observed for the εmax values of fla in

| Table 1. Summary of Spectroscopic and CV Results for [FeIIi(L)(fla)] |
|--------------------------|--------------------------|--------------------------|--------------------------|--------------------------|
|                          | [FeIIi(OMe)(fla)] (1)     | [FeIIi(L)(fla)] (2)       | [FeIIi(OMe)(fla)] (4)     | [FeIIi(NO2)(fla)] (5)     |
| FT-IR/cm−1               | ν(C=O)                   | 1545                     | 1560                     | 1550                     | 1547                     |
|                         | νas(CO2)/νs(CO2)         | 1611/1418                | 1620/1420                | 1610/1420                | 1612/1420                |
|                         | Δν/cm−1                  | 193                      | 200                      | 190                      | 192                      |
| UV−vis                  | λ/nm (ε/M−1 cm−1)        | 400 (9.59 × 104)         | 399 (1.03 × 104)         | 396 (1.16 × 104)         | 394 (1.03 × 104)         |
|                         | ν(ν=π)                   | 492 (2.53 × 104)         | 491 (2.54 × 104)         | 485 (2.77 × 104)         | 484 (2.56 × 104)         |
|                         | ν(ν=π)                   | 271 (1.31 × 104)         | 276 (1.23 × 104)         | 287 (1.41 × 104)         | 286 (1.25 × 104)         |
| CV/V                    | Ep/Oε                  | −0.079/−0.149            | −0.050/−0.148            | +0.004/−0.102            | −0.099/−0.049            |
|                         | Ep/ε                  | −0.114/70 mV             | −0.098/98 mV             | −0.049/106 mV            | −0.029/40 mV             |
|                         | tpc/tpa                 | 0.98                     | 0.91                     | 0.91                     | 0.15                     |
|                         | Ep/ε                  | +0.151/0.089             | +0.157/0.096             | +0.250/0.182             | +0.270/0.192             |
|                         | Ep/ε                  | +0.120/62 mV             | +0.126/61 mV             | +0.216/68 mV             | +0.231/78 mV             |
|                         | tpc/tpa                 | 0.88                     | 0.76                     | 0.45                     | 0.47                     |
the complexes, which correlates linearly with the Hammett constants $\sigma$ of the substituent groups of the ligands ($R = 0.99$) (Figure 3b). The stronger electron donating groups induce less blue shift (lower energy) of the $\lambda_{\text{max}}$ values of $f_l$. These results demonstrate that the $\lambda_{\text{max}}$ values of $f_l$ are influenced by the substituent groups in the ligands through the benzoate, Fe(II).
“charge delocalization conduit”, as indicated by wine-red color in Scheme 2. All complexes also show an absorption shoulder around 485 nm, which is probably due to the charge transfer between Fe(II) and ligand and an intense absorption band around 280 nm from the supporting model ligand.

**ESI-MS Spectroscopy.** Because [FeIIOMe(IIa)] was quite sensitive to air in solution, we observed only a peak cluster attributable to the oxidized species [FeIIOMe(IIa)]+ as a main peak at m/z (pos.) = 655.1 for 1, 639.3 for 2, 703.2 for 4, and 670.3 for 5 (Figure 4). Furthermore, no oxo- or fla-bridged dimeric species were observed.

The very good agreement of the m/z values and the isotope distribution patterns of the peak clusters between the experimental and calculated data strongly suggest that [FeIIOMe(IIa)] is easily auto-oxidized to [FeIIOMe(IIa)]+ by air but keep their monomer structures even after auto-oxidation in solution, which are in line with the results of the solution obtained using FT-IR and EPR (see below).

**EPR Spectroscopy.** The X-band EPR spectra of the complexes [FeIIOMe(IIa)] were all silent under N2 at 100 K (Figure 5A, black line spectra), indicating that the valence of the iron ion is +2. The results resemble that of native Fe-2,4-QD.5,11,12

**Cyclic Voltammetry.** The results of cyclic voltammetry (CV) of [FeIIOMe(IIa)] are summarized in Table 1. All redox potentials are reported versus SCE. The cyclic voltammogram of 1 is shown in Figure 6a as a typical example. All complexes display a quasireversible redox couple attributable to the one-electron

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**Figure 5.** EPR spectra of [FeIIOMe(IIa)] (4 mM in 0.5 mL DMF) at 100 K. (a) [FeIIOMe(IIa)] (1), (b) [FeIIOMe(IIa)] (2), (c) [FeIIOMe(IIa)] (4), and (d) [FeIIOMe(IIa)] (5). Line (A): under N2, (B): exposed to air 2 min, and (C): after the reaction with O2.

**Figure 6.** Cyclic voltammograms of [FeIIOMe(IIa)] (1) in DMF at room temperature, (a) under N2, and (b) under a slightly aerobic condition. (c) Plot of $E_{1/2}$ of FeIII/II vs $\sigma$. (d) Plot of $E_{1/2}$ of fla/fla• vs $\sigma$. 

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The redox process between FeII and FeIII with $E_{1/2} = -0.114$ V for [FeIIOMe(fl)] (1), $-0.098$ V for [FeIIOMe(fl)] (2), $-0.049$ V for [FeIIOMe(fl)] (4), and $-0.029$ V for [FeIIOMe(fl)] (5). The $E_{1/2}$ values are comparable with that of [FeIIOMe(fl)] (3) ($E_{1/2} = -0.094$ V). An order of $-\text{OMe (1)} < -\text{Me (2)} < -\text{H (3)} < -\text{Br (4)} < -\text{NO$_2$ (5)}$ was observed for the $E_{1/2}$ values of FeIII/II of the complexes. Besides, the values also correlate linearly with Hammett constants $\sigma$ ($R = 0.98$) (Figure 6c).

Obviously, the order can be explained by stronger electron-donating groups that induce higher electron density on the Fe(II) ion, thus leading to easier oxidation of the Fe(II) ion. Intriguingly, when the solution of [FeIIOMe(fl)] in DMF was made to contact with air slowly, a new quasi-reversible redox couple arose at $E_{1/2} = +0.120$ V for [FeIIOMe(fl)] (1) (Figure 6b), +0.126 V for [FeIIOMe(fl)] (2), +0.216 V for [FeIIOMe(fl)] (4), and +0.231 V for [FeIIOMe(fl)] (5), then vanished slowly. Under anaerobic conditions, this redox couple was not observed. The redox couple emerged during the slow reaction with O$_2$ can be assigned to the one-electron redox process between fla and flavonoxyl radical fla$^\bullet$. The $E_{1/2}$ values of fla/fla$^\bullet$ redox couples are comparable with that of [FeIIOMe(fl)] (3) ($E_{1/2} = +0.130$ V) and exhibit remarkable differences among them. Besides, an order of $-\text{OMe (1)} < -\text{Me (2)} < -\text{H (3)}$ was observed for the $E_{1/2}$ values of fla/fla$^\bullet$ redox couples.

### Table 2. Dioxygenation Reaction Rate Constants and Activation Parameters of [FeIIOMe(fl)] complexes

| complexes         | $T$ (°C) | $10^2 \times k$ (M$^{-1}$ s$^{-1}$) | $\Delta H^\ddagger$ (kJ mol$^{-1}$) | $\Delta S^\ddagger$ (J mol$^{-1}$ K$^{-1}$) | $E_a$ (kJ mol$^{-1}$) | ref |
|-------------------|----------|-----------------------------------|-----------------------------------|---------------------------------|------------------------|-----|
| [FeIIOMe(fl)] (1) | 50       | 150 ± 6                           | 59 ± 4                            | −64 ± 5                        | 61 ± 6                 | this work |
| [FeIIOMe(fl)] (2) | 50       | 59.8 ± 0.7                        | 62 ± 2                            | −59 ± 2                        | 64 ± 1                 | this work |
| [FeIIOMe(fl)] (3) | 50       | 39.8 ± 0.1                        | 65.92                             | −51.81                         | 68.60                  | 21a |
| [FeIIOMe(fl)] (4) | 50       | 11.0 ± 0.4                        | 67 ± 2                            | −57 ± 3                        | 69 ± 3                 | this work |
| [FeIIOMe(fl)] (5) | 50       | 8.71 ± 0.10                       | 67 ± 2                            | −59 ± 3                        | 70 ± 2                 | this work |
| [FeIIOMe(fl)] (1')| 80       | 58.6                              |                                   |                                |                        | 17 |
| [FeIIOMe(fl)] (L')| 80       | 44.8                              |                                   |                                |                        | 17 |
| [FeIIOMe(fl)] (3')| 100      | 80.0                              |                                   |                                |                        | 18 |
| [FeIIOMe(fl)] (salen)| 100 | 2.07                              |                                   |                                |                        | 19 |

Figure 7. Dioxygenation of [FeIIOMe(fl)] in DMF at 50 °C under O$_2$. (a) Time-dependent change of the absorption spectra during the reaction of [FeIIOMe(fl)] (1) (1.0 × 10$^{-4}$ M) with O$_2$. Inset: Time trace of the absorption changes of 1 at 400 nm. Plots of $k$ vs (b) [FeIIOMe(fl)]$_0$ and (c) [O$_2$]$_0$. (d) Hammett plot. Plots of $k$ vs (e) $\lambda_{\text{max}}$ and (f) $E_{1/2}$ of fla/fla$^\bullet$. The redox couple emerged during the slow reaction with O$_2$ can be assigned to the one-electron redox process between fla and flavonoxyl radical fla$^\bullet$. The $E_{1/2}$ values of fla/fla$^\bullet$ redox couples are comparable with that of [FeIIOMe(fl)] (3) ($E_{1/2} = +0.130$ V) and exhibit remarkable differences among them. Besides, an order of $-\text{OMe (1)} < -\text{Me (2)} < -\text{H (3)}$ was observed for the $E_{1/2}$ values of fla/fla$^\bullet$ redox couples.
< −Br (4) < −NO₂ (5) was observed, and the E₁/₂ values are also correlate the linear relationship with Hammett constants σ (R = 0.92) (Figure 6d). The order implies that the bound fla is more readily oxidized in the order of −OME (1) > −Me (2) > −H (3) > −Br (4) > −NO₂ (5), as also confirmed by experiments (see below). The order of E₁/₂ of fla/ffa* can be ascribed to the electronic nature of the substituent in the ligands, which could tune the redox potential of fla via tuning Lewis acidity of the Fe(II) ion and the electron density of fla. The stronger electron donating group, the weaker Lewis acidity of the Fe(II) ion, the weaker interaction between the Fe(II) ion and fla, the larger electron density on the fla moiety, the lower redox potentials of fla, and the easier oxidation of the bound fla.

**Reactivity with Dioxygen (Enzyme-Type Dioxygenation Reactivity).** Aerobic treatment of [FeII(L₆)(fla)] in DMF at 70 °C for 8 h gave the primary products O-benzoylsalicylic acid (HObs) [m/z (pos.): 243.1 (M + H)⁺] (2.4–17%) and CO (over 64% for 1); then, HObs was hydrolyzed to produce salicylic acid [m/z (neg.): 137.1 (M − H)⁻] (81–97%) and benzoic acid [m/z (neg.): 121.1 (M − H)⁻] (67–82%) by a small amount of water presented in the solvent. The benzoic acid was further converted to an amide derivative, N,N-dimethylbenzamide [m/z (pos.): 150.1 (M + H)⁺] (10–16%) (Scheme 2) by the solvent DMF. These products were identified by LC–MS (Figure S2) and ¹H NMR [21a] and summarized in Table S4. Conversions of the complexes are over 99%. As the products of the oxygenating ring opening reaction of the bound fla are similar to that catalyzed by the native enzyme, [FeII(L₆)(fla)] display an enzyme-type reactivity.

The dioxygenation reactions processes of the bound fla of [FeII(L₆)(fla)] were followed by monitoring the absorption change ca. 400 nm band in DMF at 30–65 °C. The kinetic data, activation parameters, and reaction conditions are summarized in Tables 2 and S5. As a typical example, the time-dependent absorption spectral change and time trace during the dioxygenation reaction of [FeII(L₆)(O₃M)] (1) are shown in Figure 7a. The initial reaction rate (v₄°, M s⁻¹) was calculated from the initial straight line part of the plot of ΔA versus time (Figure 7a, inset). Figure 7bc show that the v₄° correlates linearly with the initial concentrations of both [FeII(L₆)(fla)]₀ and [O₂]₀ thus, v₄° = k[FeII(L₆)(fla)][O₂]. The second-order reaction constants k were calculated by dividing v₄° by both [FeII(L₆)(fla)]₀ and [O₂]₀.

Therefore, the apparent second-order reaction constants k were calculated to be 8.71–150 × 10⁻² M⁻¹ s⁻¹ at 50 °C (ΔH° = 59–67 kJ mol⁻¹, ΔS° = −57 to −64 J mol⁻¹ K⁻¹) (Table 2, Supporting Information, Table S5 and Figure S3).

Figure 8. (a) Spectral change of [FeII(L₆)(fla)] (1) (0.5 mM in DMF) in the presence of excess NBT under O₂ (red line) and N₂ (black line) at room temperature. (b) EPR spectrum of 1 in the presence of 2 equiv DMPO under air at room temperature.

Till now, only two examples of Fe(II)–fla complexes [FeII(fl(a)(L₁))₃] and [FeII(L₂)(fla)] (3) (39.8 × 10⁻³ M⁻¹ s⁻¹ at 50 °C and 179 × 10⁻² M⁻¹ s⁻¹ at 70 °C) [21e] showing a dioxygenation reactivity have been reported. [FeII(fl(a)(L₁))] is not structurally characterized and exhibits a relatively lower reactivity (only 58.6 × 10⁻³ M⁻¹ s⁻¹ even at 80 °C) and requires a higher temperature (65–80 °C). Compared to it, our model complexes [FeII(L₆)(fla)] show a higher enzyme-type reactivity at a lower temperature (30–65 °C), which could be rationalized by the electron donation effect of the substituted benzoate group in the ligand. These results further demonstrate that our model complexes are good functional models of Fe-2,4-QD.

It is noteworthy that the dioxygenation reactivity of [FeII(L₆)(fla)] shows a great difference and follows a substituent dependent ranking, −OME (1) > −Me (2) > −H (3) > −Br (4) > −NO₂ (5) (Figure 7, Table 2). The Hammett plot is linear, and the reaction constant is ρ = −1.21 (R = 0.89) (Figure 7d), implying that the electron donating groups quicken the reaction rates remarkably. The notable difference on the dioxygenation reactivity could be rationalized by the electronic nature of the substituent in the ligand, which could tune the reactivity through varying the electron density of fla via the benzoate, Fe(II) ion, and “charge delocalization conduit”. Namely, the stronger electron donation to the Fe(II) ion by the electron donating group via the benzoate group makes Lewis acidity of the Fe(II) ion weaker; thus, the electron density on the fla moiety in the complexes is larger and the redox potential of fla is lower, so the fla moiety can be oxidized easier by O₂.

It should be pointed out that the order of dioxygenation reactivity looks to be in line with the initial concentrations of both the benzoate, Fe(II) ion, and electron-donating group in the ligand, which could tune the reactivity through varying the electron density of fla via the benzoate, Fe(II) ion, and “charge delocalization conduit”. There are some relationships among them. The stronger the electron-donating group in the ligand, the lower the energy of the π*→π transition, the lower the redox potential of fla, and the higher the dioxygenation reactivity. As the first example of a set of structural and functional ES models of atypically coordinated 3His-1Glu active site mononuclear non-heme Fe⁺-dependent Fe-2,4-QD, which focus on the electronic effects of the ligand on the enzyme-type reactivity; this study...
will supply critical insights into the role of Fe-2,4-QD and their property-reactivity relationship.

**Dioxygenation Reaction Mechanism.** As mentioned above, when the ESI-MS spectra of \([\text{Fe}^{III}L^R(\text{fla})]\) were recorded under the air, we only observed a peak cluster assignable to the oxidized species \([\text{Fe}^{III}L^R(\text{fla})]^+\) as a main peak (Table 1 and Figure 4). After the EPR samples of the \([\text{Fe}^{III}L^R(\text{fla})]\) solution were exposed to air for 2 min at room temperature, we observed a typical mononuclear Fe(III) EPR signal around \(g = 4.3\) (Figure 5B, red line spectra). Ca. 10% Fe(II) ions were observed a typical mononuclear Fe(III) signal appeared around \(g = 7.66, 4.27, \text{ and } 2.06\) (Figure 5C, blue line spectra). After the dioxygenation reaction, a peak cluster assignable to \([\text{Fe}^{III}L^R(\text{fla})]_{\text{obs}}^+\) was recorded (Table 1 and Figure 5) when excess NBT2+ was added to a \([\text{Fe}^{III}L^R(\text{fla})]\) solution (4 mM) and 2 equiv DMPO in DMF. Another pathway b: the reaction of \([\text{Fe}^{III}L^R(\text{fla})]_{\text{obs}}^+\) and one \(\text{O}_2\) molecule react quickly to form \([\text{Fe}^{III}L^R(\text{fla})]^+_2\) (A) and \(\text{OO}^\bullet\), then there are two possible pathways from A in the next step. One pathway a: direct reaction of intermediate A with another \(\text{O}_2\) molecule to form an endoperoxide intermediate \([\text{Fe}^{III}L(\text{fla}−\text{O}_2)]^+\) (B) slowly. Another pathway b: the reaction of A with another \(\text{O}_2\) molecule through an electron transfer process from fla to \(\text{O}_2\) to produce flavon ox radical species \([\text{Fe}^{III}L^R(\text{fla})]^{2+}\) (C) and another \(\text{OO}^\bullet\) slowly. There are also two possible pathways from C in the next step. One pathway c: the \(\text{OO}^\bullet\) attacks the flavan moiety in C to generate an alkylperoxide adduct D. Then, an intramolecular cyclization reaction makes place and an endoperoxide intermediate E is formed. Another pathway d: a \(\text{Fe}^{III}\)-superoxide adduct E is formed by the coordination of \(\text{OO}^\bullet\) to \(\text{Fe}^{III}\) in C. Then, a \(\text{Fe}^{III}\)-alkylperoxide intermediate F is produced by the intramolecular radical coupling. From F the same endoperoxide intermediate B can also be generated. Once B is formed, the dioxygenated products complex \([\text{Fe}^{III}L^R(\text{Obs})]^{+}\) (G) is engendered by releasing CO. No direct evidence is available to conclude the accurate reaction pathways, path a or path b and path c or path d at this moment. However, the observed redox couple of fla/fla*, the reduction of NBT2+, and DMPO−OOH EPR signals mentioned above may support the reaction pathway b and pathway d, which involves a flavonox radical species, intermediate C and \(\text{Fe}^{III}_2\text{OO}^\bullet\) species E, respectively. Nevertheless, the possibility of pathway a also could not be excluded. The reaction mechanism of all the complexes are the same and also close to those of the nonsubstituted derivative \([\text{Fe}^{III}L^H(\text{fla})]\) (3)11a and Fe-2,4-QD5,11,12.
CONCLUSIONS

In conclusion, we have successfully synthesized and characterized a set of structural and functional ES models of Fe-2,4-QD, [FeII^fl(ﬂ)] based on p/m-R-substituted carboxylate-containing ligands (R: p-OMe (1), p-Me (2), m-Br (4), and m-NO2 (5)). The model complexes exhibit a relatively high enzyme-type dioxygenation reactivity at low temperature. The dioxygenation reactions display first-order dependence versus both initial concentrations of the complexes and dioxygen. In addition, the reactivity is varied notably with the substituent group R and follows the ranking of −OMe (1) > −Me (2) > −H (3) → −Br (4) → −NO2 (5). The Hammett plot shows a linear relationship (ρ = −1.21), and electron donating groups progress the reaction rates. The electronic nature of the substituent in the ligand tunes the properties and the reactivity of the complexes remarkably, and there are some correlations between k with the λ_{max} of the π → π* transition or the redox potential of the ﬂa moiety in the complexes, respectively. That is to say, the key role of the substituent group is to tune Lewis acidity of Fe(II), the electron density, and the redox potential of ﬂa by electron donation via benzoate, Fe(II) ion, and “charge delocalization conduit”. As far as we know, this is the first example of a set of the ES models of Fe-2,4-QD, which faithfully reproduce the atypically coordinated 3His-1Glu coordination environment of the mononuclear non-heme FeII-dependent Fe-2,4-QD active site structurally and functionally and will provide crucial insights into the role of Fe-2,4-QD and the active site structure–function relationship of different active site MNHES.

EXPERIMENTAL SECTION

General and Physical Methods. The methods are the same as the methods reported previously.23 EPR instrument parameters for DMPO–OOH detection are microwave frequency 9.531841 GHz, modulation frequency 100 kHz, modulation amplitude 3 G, microwave power 1.91 mW, frequency 9.531841 GHz, modulation frequency 100 kHz, on a glass capillary. Data of X-ray diﬀraction were collected using the Bruker Smart APEX II CCD single-crystal diﬀractometer using Mo Kα radiation (λ = 0.71073 Å) to 2θ_{max} of 50.0° at 293 K. The crystallographic calculation was performed by a direct method using SHELXL 2014. The FT-IR data, reaction product analysis data, kinetic analysis methods are the same as the methods reported previously.23

reaction products analysis. The reaction product analysis methods are the same as the methods reported previously.21a Synthesis of the Model Complexes [FeII^fl(ﬂ)]. Ligands L^H (R: p-OMe, p-Me, m-Br, m-NO2) were synthesized based on the reported methods.23 Complexes were synthesized using a similar method. A solution of L^H (0.1 mmol in 1.0 mL dry CH3OH) was added dropwise to a solution of FeII(OAc)2 (17.4 mg, 0.1 mmol, in 2.0 mL dry CH3OH) under a nitrogen atmosphere at room temperature. After stirring for 30 min, a solution of flavonol (23.8 mg, 0.1 mmol, in 1:1 CH3OH/CH2Cl2) was added dropwise to the above solution with strong stirring. The mixture was stirred for 1 day under N2. The [FeII^fl(ﬂ)] was isolated as dark red powder by filtration.

ASSOCIATED CONTENT

Supporting Information

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AUTHOR INFORMATION

Corresponding Author
E-mail: yingjiisdlut.edu.cn (Y.-J.S.)

ORCID
Ying-Ji Sun: 0000-0001-6238-1871

Present Address
Department of Chemistry, XinXiang Medical University, Xinxiang 453003, China (Q.-Q.H.).
Notes
The authors declare no competing financial interest.

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