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A non-heme diiron complex for (electro)catalytic reduction of dioxygen: tuning the selectivity through electron-delivery

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Abstract

In the Oxygen Reduction Reaction (ORR) domain, the investigation of new homogeneous catalysts is a crucial step towards the full comprehension of the key structural and/or electronic factors that control catalytic efficiency and selectivity. Herein, we report a unique non-heme diiron complex that can act as a homogeneous ORR catalyst in acetonitrile solution. This iron(II)-thiolate dinuclear complex, $[\text{Fe}^{\text{II}}_2(\text{LS})(\text{LSH})]$ ($[\text{Fe}_2^{\text{SH}}]^+$) ($\text{LS}^{2-} = 2,2'-(2,2'\text{-bipyridine-6,6'-diyl})\text{bis}(1,1\text{-diphenylethanethiolate})$) contains a thiol group in the metal coordination sphere. $[\text{Fe}_2^{\text{SH}}]^+$ is an efficient ORR catalyst both in the presence of a one-electron reducing agent as well as under electrochemically assisted conditions. However, its selectivity is dependent on the electron delivery pathway, in particular, the process is selective for H_2O_2 production under chemical conditions (up to ~95%), whereas H_2O is the main product during electrocatalysis (less than ~10% H_2O_2). Based on computational work alongside the experimental data, a mechanistic proposal is discussed that rationalizes the selective and tunable reduction of dioxygen.

Introduction

The development of efficient, stable and noble metal-free catalysts for the O_2 reduction reaction (ORR) with selective production of either water or hydrogen peroxide remains one of the central challenges in the energy conversion domain.¹⁻² Fuel cell design is focused on the production of electricity based on the four-electron O_2 reduction (to H_2O) at the cathode, associated with the oxidation of a fuel (such as H_2 or $MeOH$) at the anode.³⁻⁵ On the other hand, the two-electron O_2 reduction is an interesting strategy to produce H_2O_2 , which represents a valuable commodity chemical and a versatile and clean oxidizing agent.⁶⁻⁸ Heterogeneous ORR catalysts have been mainly developed towards implementation into exploitable devices for such applications.⁹⁻¹⁰ Among them, the most promising noble metal-free materials are nitrogen-doped nanocarbon materials, containing molecularly-defined active sites based on metal porphyrinic moieties.¹¹⁻¹² Regarding molecular homogeneous catalysts, their study generally allows for a deeper comprehension of the ORR mechanism by identifying the rate limiting steps for the reaction and the key structural and electronic factors that control their activity and selectivity (two proton/two electron *vs* four proton/four electron O_2 reduction).^{1, 13} Such investigation should eventually lead to enhanced efficiency through rational ligand design (e.g. by introduction of proton relays or Lewis acidic sites) for guiding the bifurcation processes into the wanted reaction channel.

Regarding homogeneous ORR catalysts, the most common and well-studied ones are based on iron or cobalt complexes embedded in macrocyclic heme-like ligands (such as porphyrins, phthalocyanines, or corroles).¹⁴⁻²¹ Detailed electrochemical studies of a series of such complexes evidenced how Fe-based catalysts are generally more selective for 4-electron O_2 reduction than their Co analogues.^{1, 13, 22} This difference in reactivity is proposed to be a consequence of the generation of high-valent metal-oxo species that are easier to generate for hemes than for porphyrinic complexes based on other metal ions. Catalysts with non-heme ligands have been much less explored, although some studies on copper-based complexes²³⁻³⁰ and, to a lesser extent, manganese-based systems have been reported.³¹⁻³⁴ Even though in several cases the nature of the acid and/or the presence of a proton relay were investigated on such systems, the selectivity towards $4H^+/4e^-$ *vs* $2H^+/2e^-$ O_2 reduction catalysis is still difficult to predict.

For homogeneous catalysts, generally two ways to provide electrons have been adopted, namely through the use of an appropriate reducing agent or an electrode (with application of a suitable electrochemical potential). Both approaches have been widely tested but rarely with the same catalyst. Using an Fe-based porphyrin system, Mayer et al. observed the same

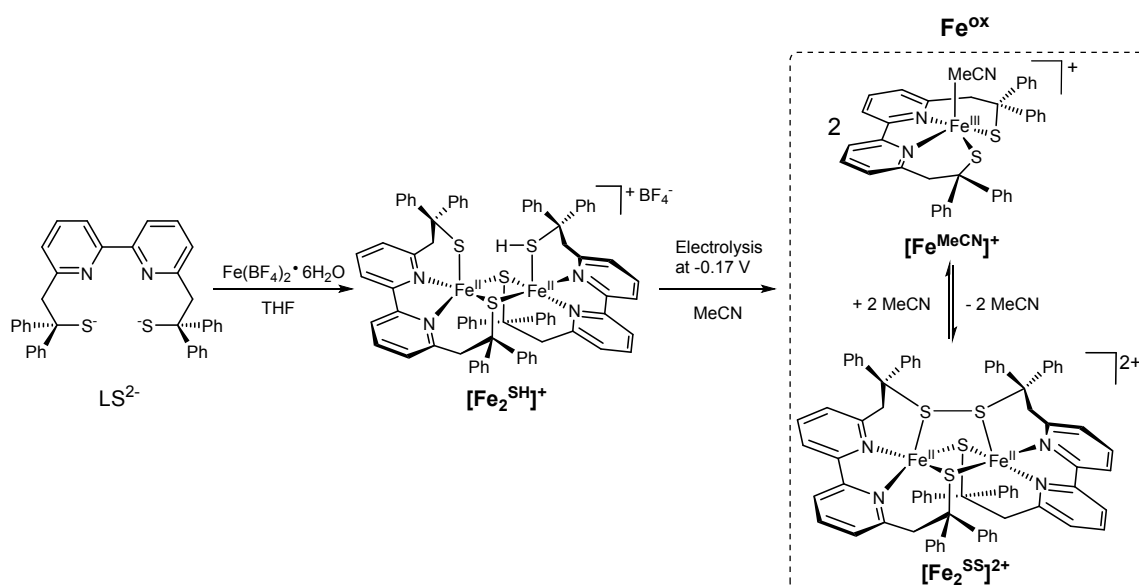
efficiency and comparable selectivities for H₂O₂ production for both the chemically- and electrochemically-driven processes (10(±3)% vs 10-20%).³⁵ A mononuclear Cu complex supported by a tetrapodal nitrogen-donor ligand was also reported and found to be selective for 4H⁺/4e⁻ O₂ reduction catalysis in both cases. However, in that example electrocatalysis was performed in protic solvent,³⁶ while chemical reduction was carried out in acetone and hence the environmental conditions affected the catalysis and selectivity.²³

In the present work, we report the synthesis, characterization and reactivity of a dinuclear non-heme iron complex, [Fe^{II}₂(LS)(LSH)] (LS²⁻ = 2,2'-(2,2'-bipyridine-6,6'-diyl)bis(1,1-diphenylethanethiolate)),³⁷ that acts as a homogeneous O₂ reduction catalyst and displays a change in selectivity under chemical versus electrochemical conditions. The process is selective for H₂O₂ production when octamethylferrocene (Me₈Fc) is used as reducing agent, whereas H₂O is dominant when catalysis is driven electrochemically, while the other main experimental parameters (solvent, catalyst concentration, nature and concentration of the acid) remained unchanged. With the aim of understanding how the selectivity could be fine-tuned as a function of the different electron sources, a mechanistic investigation supported by theoretical calculations has been performed. Surprisingly, even though the iron-oxygen chemistry has been extensively investigated,³⁸⁻⁴⁴ our study reports the first non-heme iron-based ORR catalyst. In addition, it provides insights on how to control the ORR selectivity in separate pathways for 2-electron versus 4-electron reduction. Finally, comparison of the data gained in this work with those previously reported for the isostructural Mn complex^{32,45} allowed us to provide insights into the role of the transition metal in ORR catalysis mediated by such dimetallic complexes.

Results

Synthesis and solid-state properties of the dimercapto-bridged Fe^{II} dinuclear complex.

Under an inert atmosphere the LS²⁻ ligand³⁷ reacts with Fe(BF₄)₂•6H₂O (1.2 equiv.) in THF to yield a brown precipitate that was characterized as [Fe^{II}₂SH]⁺ or [Fe^{II}(LS)(LSH)]⁺ (Scheme 1). Single crystal X-ray diffraction data collected on the isolated product reveal a dimercapto-bridged dinuclear Fe^{II} complex [Fe^{II}₂(LS)(LSH)]BF₄ ([Fe₂SH]⁺), isostructural to the parent Mn^{II} compound ([Mn^{II}₂(LS)(LSH)]BF₄, [Mn₂SH]⁺ reported previously.³²



Scheme 1. Synthetic routes for the different iron-thiolate complexes described in this work.

Similarly to $[Mn_2SH]^+$, the structure of the $[Fe_2SH]^+$ cation (shown in Figure 1a, selected bond distances and angles are reported in the Supporting Information) displays: (i) a planar $\{Fe_2S_2\}$ diamond core with deviations from the $Fe1S2Fe2S3$ plane of ~ 0.010 Å, (ii) two distinguishable metal sites, each Fe^{II} center being pentacoordinated and surrounded by an N_2S_3 donor set in a distorted trigonal bipyramidal environment ($\tau_5 = 0.59$ for Fe1 and $\tau_5 = 0.68$ for Fe2),⁴⁶ (iii) no direct metal-metal interaction, with a metal...metal distance of $3.1107(7)$ Å, and most importantly (iv) the unusual presence of one metal-bound thiol.⁴⁷⁻⁵⁰ The protonation of one thiolate is attested by the difference between the $Fe1-S1$ and $Fe2-S4$ bond lengths ($2.3566(11)$ and $2.5387(11)$ Å, respectively) that gives evidence of the protonation at S4. A similar ~ 0.2 Å discrepancy has been previously observed between the two $Mn-S_{terminal}$ distances in $[Mn_2SH]^+$ ($Mn1-S1 = 2.4399(10)$ Å and $Mn2-S51 = 2.6462(10)$ Å). Thiolate protonation in $[Fe_2SH]^+$ is further confirmed by significant electron density assigned to a hydrogen atom (H4) found on the difference Fourier map near S4 ($S4-H4 = 1.171(17)$ Å). The resulting $S1 \cdots S4$ distance ($3.5940(10)$ Å) is compatible with the presence of an intramolecular $S4-H4 \cdots S1$ hydrogen-bond, while in the parent Mn complex the $S \cdots S$ distance is too long for such interaction ($3.772(3)$ Å).

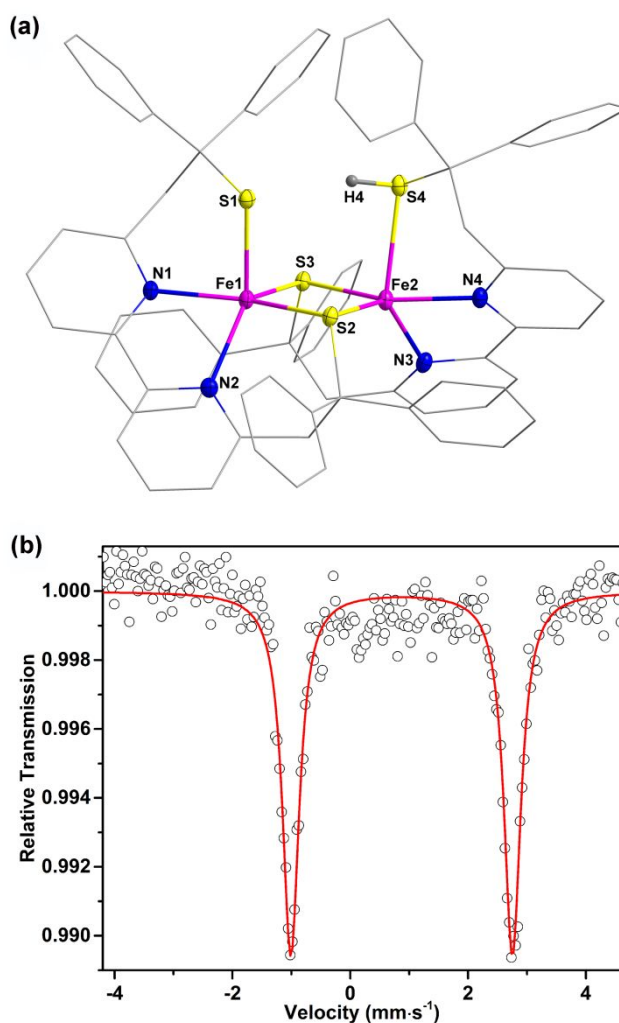


Figure 1. (a) Molecular structure of $[\text{Fe}_2^{\text{SH}}]\text{ClO}_4 \cdot 1.5\text{MeCN} \cdot 0.5\text{Et}_2\text{O}$ determined by X-ray crystallography with the thermal ellipsoids of the metal core drawn at 30% probability level. All hydrogen atoms except H4, all anions and solvent molecules are omitted for clarity. (b) Solid-state zero-field Mössbauer spectrum of $[\text{Fe}_2^{\text{SH}}]^+$ recorded at 80 K.

The +2 oxidation state of both Fe ions is confirmed by Mössbauer spectroscopy (Figure 1b): the zero-field spectrum of $[\text{Fe}_2^{\text{SH}}]^+$ recorded on a solid sample at 80 K displays a single doublet ($\delta = 0.87 \text{ mm}\cdot\text{s}^{-1}$ and $\Delta E_Q = 3.76 \text{ mm}\cdot\text{s}^{-1}$) characteristic of two high-spin ($S = 2$) Fe^{II} ions. Taken together with the charge balance of the complex (+1), these data are consistent with the presence of one Fe-bound thiol in $[\text{Fe}_2^{\text{SH}}]^+$. The two Fe centers are indistinguishable by Mössbauer spectroscopy, reflecting the only small differences in their coordination spheres, and suggesting that elongation of one Fe-S bond does not significantly alter the metal ion's electronic structure.

Solution and redox properties. In MeCN solution, $[\text{Fe}_2^{\text{SH}}]^+$ remains dinuclear, as indicated by (i) ESI-mass spectrometry, with a major peak at 1269.3 m/z (with isotope pattern corresponding to $[\text{Fe}^{\text{II}}_2(\text{LS})(\text{LSH})]^+$; Figure S1) and the absence of any peak originating from mononuclear species, and (ii) the paramagnetic ^1H NMR spectrum of $[\text{Fe}_2^{\text{SH}}]^+$, displays no large paramagnetic shift as observed in the spectrum of the parent mononuclear complex $[\text{Fe}^{\text{II}}(\text{LS})\text{Cl}]^{51}$ (Figure S2), consistent with an antiferromagnetically coupled dinuclear high spin Fe^{II} complex.

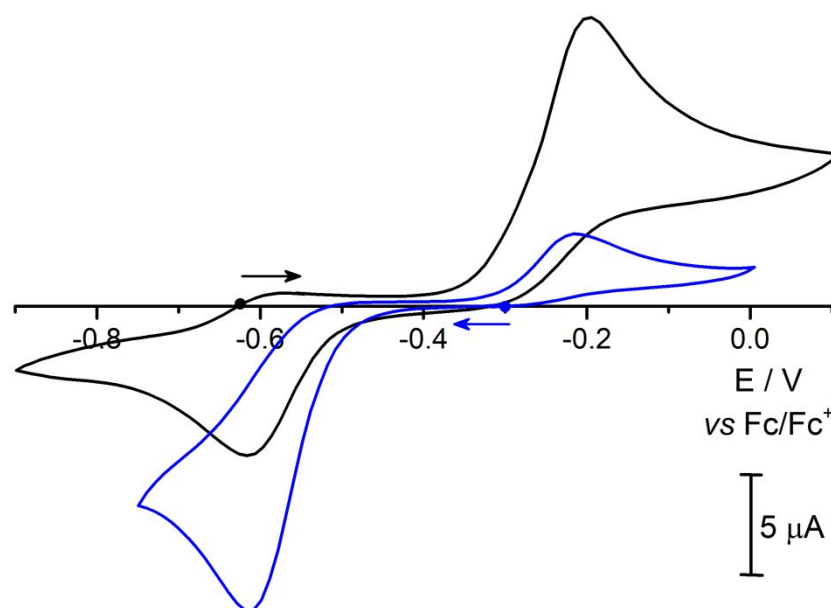


Figure 2. Cyclic voltammograms of $[\text{Fe}_2^{\text{SH}}]^+$ (~1.0 mM) before (black line) and after (blue line) bulk electrolysis at -0.17 V vs Fc^+/Fc in MeCN under argon atmosphere. Supporting electrolyte: 0.1 M Bu_4NClO_4 , GC working electrode ($\varnothing = 3$ mm), scan rate: 100 $\text{mV}\cdot\text{s}^{-1}$.

The cyclic voltammogram (CV, Figure 2) of $[\text{Fe}_2^{\text{SH}}]^+$ displays an electrochemically irreversible oxidation step ($E_{\text{pa}} = -0.19$ V vs Fc^+/Fc and $E_{\text{pc}} = -0.63$ V), which is partially reversible chemically ($i_{\text{pc}}/i_{\text{pa}} \sim 0.6$). On the basis of coulometric and spectroscopic studies (vide infra), this redox system has been assigned to the two-electron metal-based oxidation of the dinuclear complex ($\text{Fe}^{\text{II}}\text{Fe}^{\text{II}} \rightarrow \text{Fe}^{\text{III}}\text{Fe}^{\text{III}}$) followed by chemical rearrangements. A mixture of two different oxidized species is generated by bulk electrolysis carried out at -0.17 V vs Fc^+/Fc of a MeCN solution of $[\text{Fe}_2^{\text{SH}}]^+$ (6.65 C of charge passed for 0.0368 mmol of initial complex, corresponding to 0.94 electrons per Fe atom). By different spectroscopic techniques (including UV-vis, ^1H -NMR, and cw X-band EPR), they have been identified⁵¹ as: (i) a mononuclear Fe^{III} -thiolate complex, $[\text{Fe}^{\text{III}}(\text{LS})(\text{MeCN})]^+$ ($[\text{Fe}^{\text{MeCN}}]^+$), and (ii) a dinuclear Fe^{II} -

disulfide complex, $[\text{Fe}^{\text{II}}_2(\text{LSSL})]^{2+}$ ($[\text{Fe}_2^{\text{SS}}]^{2+}$, LSSL^{2-} = disulfide form of the LS^{2-} ligand, Scheme 1). This assignment relies on our previous investigation showing that an identical mixture is obtained when $[\text{Fe}_2^{\text{SS}}]^{2+}$ is dissolved in MeCN. Note that in solid state, Fe^{ox} can be isolated in the dimeric form, $[\text{Fe}_2^{\text{SS}}]^{2+}$.⁵¹ For simplicity, in the following paragraphs the $[\text{Fe}^{\text{MeCN}}]^+ + [\text{Fe}_2^{\text{SS}}]^{2+}$ mixture ($\sim 2:1$ ratio)⁵¹ is designated Fe^{ox} . Taken together, these results imply that fast deprotonation accompanies the double electron transfer before the Fe^{ox} complex is formed.

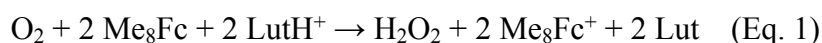
A similar redox behavior is observed for the parent Mn-based dinuclear complex $[\text{Mn}_2^{\text{SH}}]^+$, which displays a ~ 180 mV anodic shift for the $\text{M}_2^{\text{II}}/\text{M}_2^{\text{III}}$ redox system with respect to the Fe complex ($E_{\text{p}_a} = -0.19$ V for Fe vs $E_{\text{p}_a} = -0.01$ V for Mn).^{32, 45} This shift indicates that $[\text{Fe}_2^{\text{SH}}]^+$ is expected to be more easily oxidized by dioxygen with respect to the corresponding Mn complex. For a previously described series of mononuclear M^{II} -thiolate complexes, an even larger cathodic shift (290 mV) was observed between the Fe and Mn derivatives.⁵²

Homogeneous catalysis and electrocatalysis for dioxygen reduction. The capability of Fe^{ox} and $[\text{Fe}_2^{\text{SH}}]^+$ to catalyze ORR was investigated in MeCN solution, in the presence of protons and electrons. The 2,6-lutidinium tetrafluoroborate acid (LutHBF_4) was employed as proton source, whereas electrons were delivered by a chemical reducing agent (octamethylferrocene, Me_8Fc , $E_{1/2} = -0.41$ V vs Fc^+/Fc , Figure S11) or alternatively by applying an adequate electrochemical potential to a glassy carbon electrode. In the presence of LutH^+ , $[\text{Fe}_2^{\text{SH}}]^+$ reacts with O_2 to generate Fe^{ox} (Figure S3). Note that this complex can also be prepared by bulk electrolysis of $[\text{Fe}_2^{\text{SH}}]^+$ (see above), or by dissolving the $[\text{Fe}_2^{\text{SS}}](\text{ClO}_4)_2$ solid in MeCN.⁵¹ Despite the fact that the redox potential of the $\text{Me}_8\text{Fc}^+/\text{Me}_8\text{Fc}$ couple is too high to reduce Fe^{ox} ($E_{\text{p}_c} = -0.63$ V, see above), Fe^{ox} is efficiently reduced by Me_8Fc in the presence of a large excess (50 and 150 equiv.) of LutH^+ to generate the oxygen reactive $[\text{Fe}_2^{\text{SH}}]^+$ species (see the UV-vis experiment in Figure S4). As expected, in the absence of protons, no reaction is observed by mixing Fe^{ox} and Me_8Fc in MeCN solution under Ar atmosphere. The protonation step that occurs after the reduction of Fe^{ox} to generate $[\text{Fe}_2^{\text{SH}}]^+$ is therefore proposed to trigger the electron transfer process. Accordingly, in the CV of Fe^{ox} the cathodic wave is 60 mV shifted to a less negative potential ($E_{\text{p}_c} = -0.57$ V) after the addition of 150 equiv. of LutH^+ (Figure S12).

As revealed by UV-vis absorption monitoring (Figure 3a, Table S3), the addition of a catalytic amount of Fe^{ox} (100 μM) to an air-saturated (~ 1.6 $\text{mM}\cdot\text{atm}^{-1}$ O_2) MeCN solution of Me_8Fc (2 mM, 20 equiv.) and LutH^+ (15 mM, 150 equiv.) results in the progressive oxidation

of Me₈Fc to Me₈Fc⁺ ($\lambda_{\text{max}} = 750 \text{ nm}$, $\epsilon = 420 \text{ M}^{-1}\text{cm}^{-1}$), up to a $\sim 75(\pm 10)\%$ amount. A similar behaviour is observed when [Fe₂^{SH}]⁺ is employed as ORR catalyst (Figure S5), indicating that both species are involved in the same catalytic cycle. In the absence of catalyst (and on the timescale of catalysis) Me₈Fc is not oxidized by O₂. The absence of catalysis when the Fe complexes [Fe(H₂O)₆](BF₄)₂ or [Fe(bpy)₃](PF₆)₂ are employed, highlights the remarkable beneficial effect of the coordination sphere provided by the thiolate-rich LS²⁻ ligand. The non-quantitative Me₈Fc oxidation in the presence of Fe^{ox} or [Fe₂^{SH}]⁺ can be rationalized in term of partial catalyst deactivation during the ORR, most likely because of sulfur oxygenation in the presence of *in-situ* generated H₂O₂ (Figure S9).

The amount of hydrogen peroxide produced in the reaction solution during the ORR catalysis was quantified from a spectrophotometric assay using the Ti-TPyP reagent for the titration (see Table S3).⁵³⁻⁵⁴ In the presence of Fe^{ox} as catalyst, under the typical conditions of Figure 3, a 0.67(±0.03) mM concentration of H₂O₂ is detected after 10 min. If we take into account the overall reaction stoichiometry (Eq. 1), this implies a 94(±4)% selectivity of the catalytic process for H₂O₂ production.



Under these conditions, the ORR selectivity using Fe^{ox} is thus comparable to that measured with the parent Mn catalyst (82 (±2)%).

The kinetic profiles for the generation of Me₈Fc⁺ catalyzed by Fe^{ox} and the parent Mn catalyst ([Mn₂^{SH}]⁺) (Figure S6) indicate one order of magnitude faster apparent kinetics for the iron system (TOF_i Fe^{ox} = 8 (±1) 10³ h⁻¹) compared to the manganese system (TOF_i [Mn₂^{SH}]⁺ ≈ 4.0 (±0.4) 10² h⁻¹, TOF_i = initial turnover frequency).

In the case of Fe^{ox}, the TOF_i is neither significantly sensitive to the relative concentration of acid (from 5 to 15 mM) nor of reductant (from 2 to 4 mM), but the selectivity for H₂O₂ drops when the concentration of reductant increases (46(±4)% with 4 mM of Me₈Fc, Table S3).

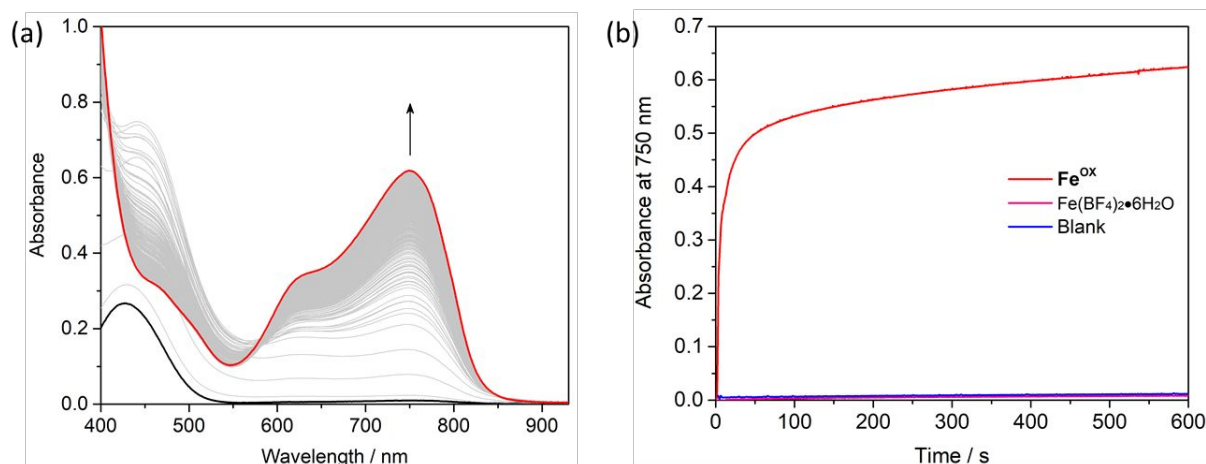


Figure 3. (a) UV-vis spectral changes observed during ORR catalyzed by Fe^{ox} in the presence of LutH^+ and Me_8Fc in MeCN at 293 K: 2.0 mM Me_8Fc , 15 mM LutHBF_4 , 0.1 mM (dimer) Fe^{ox} , air-saturated solution, 1 cm path length, $\Delta t = 0.5$ s. (b) Representative kinetic profiles for the formation of Me_8Fc^+ at 750 nm in the presence of Fe^{ox} ($[\text{Fe}_2^{\text{SH}}]^+$ in Figure S5), $\text{Fe}(\text{BF}_4)_2 \cdot 6\text{H}_2\text{O}$, or in the absence of catalyst (blank).

To test the ability of the air-stable Fe^{ox} catalyst to serve as ORR electrocatalyst, CV experiments of Fe^{ox} (0.1 mM) were performed in the presence of a buffered LutH^+/Lut MeCN solution (15 mM, 150 equiv.) under saturated argon (Figure S14a) or oxygen atmosphere (Figure 4a). Under O_2 saturated atmosphere, an irreversible oxygen-reduction peak appears at $E_{\text{cat}/2} = -0.50$ V vs Fc^+/Fc only in the presence of Fe^{ox} . The ORR peak onset is only slightly shifted (to less negative potentials) with respect to the cathodic peak of Fe^{ox} under Ar atmosphere ($E_{\text{pc}} = -0.60$ V), but is ~ 5 -fold more intense under the same experimental conditions (see Figure 4a vs Figure S14a). These data unequivocally demonstrate the electrocatalytic activity of Fe^{ox} towards O_2 reduction in MeCN solution. A rinse test confirms that the activity arises from the iron catalyst in solution and is not due to any electrode-adsorbed material (see Figure S13).

The standard potentials for $2\text{H}^+/2\text{e}^-$ and $4\text{H}^+/4\text{e}^-$ O_2 reduction can be estimated in the buffered LutH^+/Lut MeCN solution (lutidinium, $\text{pK}_{\text{a}} = 13.92$ in MeCN)⁵⁵: $E^0(\text{O}_2, 4\text{LutH}^+/2\text{H}_2\text{O}) = 1.21 - 0.0592\text{pK}_{\text{a}} \approx 0.39$ V vs Fc^+/Fc , by using the method reported by Mayer et al.,^{1, 22} or $E^0(\text{O}_2, 4\text{LutH}^+/2\text{H}_2\text{O}) = 2.038 - 0.0592\text{pK}_{\text{a}} \approx 1.21$ V vs NHE, corresponding to 0.68 V vs Fc^+/Fc ,⁵⁶ by using the method reported by Nocera et al.⁵⁷; $E^0(\text{O}_2, 2\text{LutH}^+/\text{H}_2\text{O}_2) = 0.68 - 0.0592\text{pK}_{\text{a}} \approx -0.14$ V vs Fc^+/Fc (Mayer method).⁵⁵ On the basis of the experimental mid-wave potential of the catalytic wave ($E_{\text{cat}/2} = -0.50$ V vs Fc^+/Fc), the $4\text{H}^+/4\text{e}^-$ O_2 reduction overpotential $\eta(\text{O}_2, 4\text{LutH}^+/2\text{H}_2\text{O})$ ($\eta = E_{\text{cat}/2} - E^0$) is estimated in the range 0.89-1.18 V (while the $2\text{H}^+/2\text{e}^-$ O_2 reduction overpotential $\eta(\text{O}_2, 2\text{LutH}^+/\text{H}_2\text{O}_2)$ is determined to 0.36 V). The

estimation of $\eta(\text{O}_2, 4\text{LutH}^+/2\text{H}_2\text{O})$ is typical for homogeneous metal catalysts (mainly Fe- and Co-based) for $4\text{H}^+/4\text{e}^- \text{O}_2$ reduction (0.52-0.96 V reported range).⁵⁸

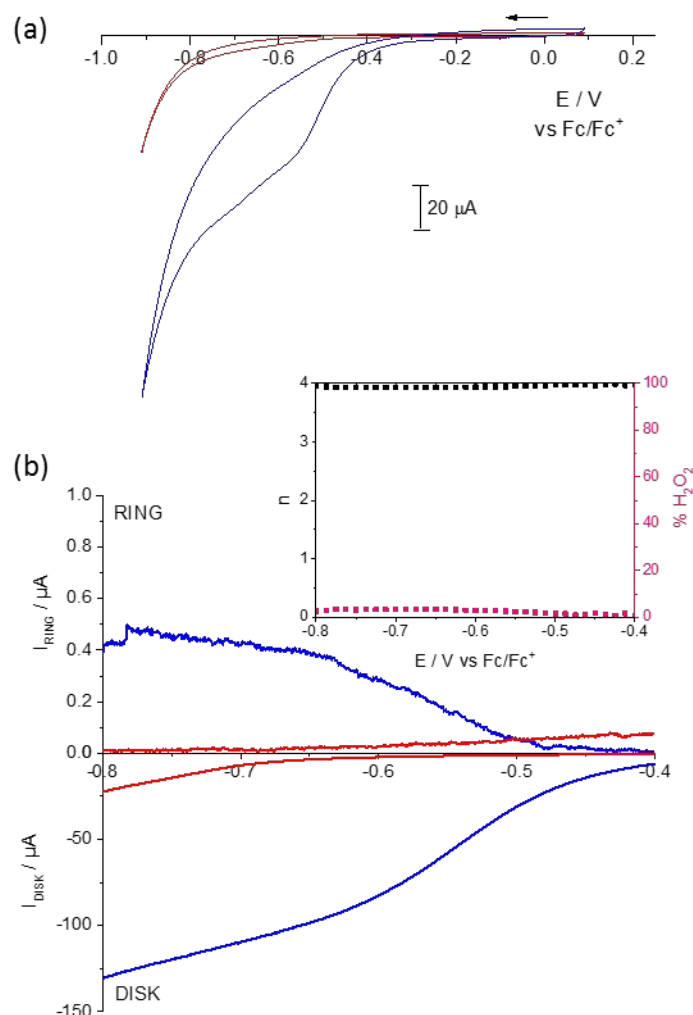


Figure 4. (a) Cyclic voltammograms (CV) on GC disk and (b) rotating ring-disk electrode voltammograms (RRDEV) under oxygen-saturated atmosphere of a 15.0 mM LutH^+ + 15.0 mM Lut solution in the absence (red lines) or presence (blue lines) of 0.1 mM (dimer) Fe^{ox} . MeCN 0.1 M Bu_4NClO_4 , GC ($\varnothing = 5 \text{ mm}$); scan rates: $100 \text{ mV}\cdot\text{s}^{-1}$ (CV), $5 \text{ mV}\cdot\text{s}^{-1}$ (RRDEV, the response does not change within the range 2-10 $\text{mV}\cdot\text{s}^{-1}$); for RRDEV: rotation rate 1500 rpm; ring current recorded with the Pt ring held at 0.2 V vs Fc^+/Fc (for the blue curves, $i_{\text{ring}}^{0.2\text{V}}$ has been corrected by subtracting $i_{\text{ring}}^{0\text{V}}$, see Figure S16). The inset of (b) displays the RRDEV-determined % of produced H_2O_2 and the number of electrons (n) consumed for reduction of one molecule of O_2 vs applied potential.

Rotating ring-disk electrode voltammetry (RRDEV) experiments were carried out for estimating $\text{H}_2\text{O}_2/\text{H}_2\text{O}$ selectivity and quantifying the production of H_2O_2 under electrochemical conditions (Figures 4b, S14-S15). The potential at the GC disk was scanned through the catalytic region (from -0.4 V to -0.8 V vs Fc^+/Fc), and the potential at the Pt ring (E_{ring}) was fixed at 0.2 (Figure 4b) or 0.43 V (Figure S15) vs Fc^+/Fc ⁵⁷ in order to detect the oxidation of the ORR-generated hydrogen peroxide.

Under argon atmosphere, the ring current observed in the presence of Fe^{ox} and LutH^+/Lut is mainly attributed to the re-oxidation of $[\text{Fe}_2^{\text{SH}}]^+$ formed on the disk, to regenerate Fe^{ox} (Figure S14). The ring current at 0.2 V vs Fc/Fc^+ (blue trace in Figure 4b) in the presence of Fe^{ox} and LutH^+/Lut under dioxygen atmosphere can be thus attributed to the oxidation of H_2O_2 generated in the catalytic ORR process. Under these conditions, the Faradaic efficiency for H_2O_2 production is $\leq 3.5\%$ in the explored potential window, corresponding to ≥ 3.9 electrons (n) exchanged during the ORR process (see inset of Figure 4b). Similar results have been found at $E_{\text{ring}} = 0.43$ V vs Fc/Fc^+ : $\leq 12.5\%$ H_2O_2 , $n \geq 3.7$ (Figure S15). These RRDE studies, therefore, give evidence that the Fe^{ox} catalyst is selective for water production, and thus catalyzes almost exclusively $4\text{H}^+/4\text{e}^-$ O_2 reduction under electrochemical conditions. A bulk electrolysis carried out at -0.43 V vs Fc^+/Fc on an O_2 -saturated acidic solution of Fe^{ox} in MeCN (0.1 mM complex, 15 mM LutH^+), followed by Ti-TPyP reagent analysis, evidences no formation of H_2O_2 in agreement with RRDE experiments.

Mechanistic investigations of the O_2 reduction process

In support of the experimental observations, a series of density functional theory (DFT) calculations were performed to gain insight into the mechanistic details for catalytic reduction of dioxygen by $[\text{Fe}_2^{\text{SH}}]^+$ (or Fe^{ox}). We calculated individual structures along the proposed reaction cycle of Scheme 2 (using two different models that give comparable results, see Supporting Information, including Figures S17 – S22) and investigated reaction (free) energies of dioxygen binding and activation.

The catalytic process is initiated by the reduction of Fe^{ox} into $[\text{Fe}_2^{\text{SH}}]^+$, the complex that does interact with O_2 . The DFT-optimized structure of $[\text{Fe}_2^{\text{SH}}]^+$ fully matches the X-ray structure, and the high spin state of the Fe^{II} ions is well predicted. Specifically, the calculated distances of Fe1-S1 and Fe2-S4 of 2.39 and 2.61 Å match the crystal structure bond lengths excellently and indeed show elongation of the Fe-S bond for the thiol-ligated system. Note as well the predicted hydrogen bonding interaction between the terminally bound thiolate and the thiol ($\text{S-S} = 3.582$ Å) in the optimized structure of $[\text{Fe}_2^{\text{SH}}]^+$ as observed experimentally. An almost

thermoneutral dioxygen binding step with $\Delta G = 5.6 \text{ kcal mol}^{-1}$ ($\Delta E + ZPE = -2.2 \text{ kcal mol}^{-1}$) leads to the formation of the *trans*-1,2- μ -peroxo dinuclear Fe^{III} complex, $[\text{Fe}_2^{\text{OO/S}}]^+$ (via a transient high-energy superoxo species, $[\text{Fe}_2^{\text{OO/S}}]^+$, Figure S18 and Table S5). In contrast to the corresponding DFT-calculated structure of the dinuclear Mn^{III} -peroxo complex,³⁰ the diiron complex $[\text{Fe}_2^{\text{OO/S}}]^+$ is much less symmetric with an almost linear Fe–O–Fe angle of 173° , while the second oxygen atom is coordinated to one iron only (Figure 5). As such, the structure of $[\text{Fe}_2^{\text{OO/S}}]^+$ resembles a side-on Fe^{III} -peroxo linked to a second mononuclear Fe^{III} unit. Such $\mu\text{-}\eta^1\text{:}\eta^2$ -peroxo coordination mode has been already observed in heme-peroxo copper models of the cytochrome *c* oxidase.^{59–61}

Attempts to optimize either a more symmetrical structure for the *trans*-1,2- μ -peroxo diiron complex or a *cis*- μ -1,2-peroxo coordination mode failed and converged back to the geometry shown in Figure 5. The *cis*- μ -1,2-peroxo mode is the exclusive coordination mode found in non-heme Fe^{III} peroxo-bridged complexes,^{62–64} in which one or more additional bridges (e.g. carboxylate, phenolato) are often present to stabilize such species. Conversely, complexes with unsupported peroxo-bridged dimers as in Fe^{III} -tetraphenylporphyrins display a *trans*- μ -1,2 coordination mode as in the present case.⁶⁵

We thus evaluated the capacity of the peroxo $[\text{Fe}_2^{\text{OO/S}}]^+$ complex to evolve either via an internal proton transfer to form a hydroperoxo complex or via the breaking of the O–O bond to generate a high-valent Fe-oxo species.

Concerning the intramolecular proton transfer pathway in $[\text{Fe}_2^{\text{OO/S}}]^+$ (Figure 5), a kinetic barrier of $12.8 \text{ kcal mol}^{-1}$ has been calculated to generate the diiron(III)-hydroperoxo complex $[\text{Fe}_2^{\text{OOH}}]^+$ (Figure S22). Regarding the cleavage of the O–O bond, even if two different pathways were located (see SI for further details), the most favorable one corresponds to the generation of dinuclear Fe^{IV} di-oxo intermediate $[\text{Fe}_2(\mu\text{-O})_2^{\text{SH}}]^+$ (Figure 5). However, the energy barrier of $15.9 \text{ kcal mol}^{-1}$ for this reaction is larger than the barrier predicted to generate the protonated $[\text{Fe}_2^{\text{OOH}}]^+$ peroxo complex. These calculations are thus consistent with the fact that H_2O_2 can be selectively produced (in the presence of Me_8Fc), through the reaction of $[\text{Fe}_2^{\text{OOH}}]^+$ with an additional proton and the formation of Fe_2^{ox} (Scheme 2, pathway B).

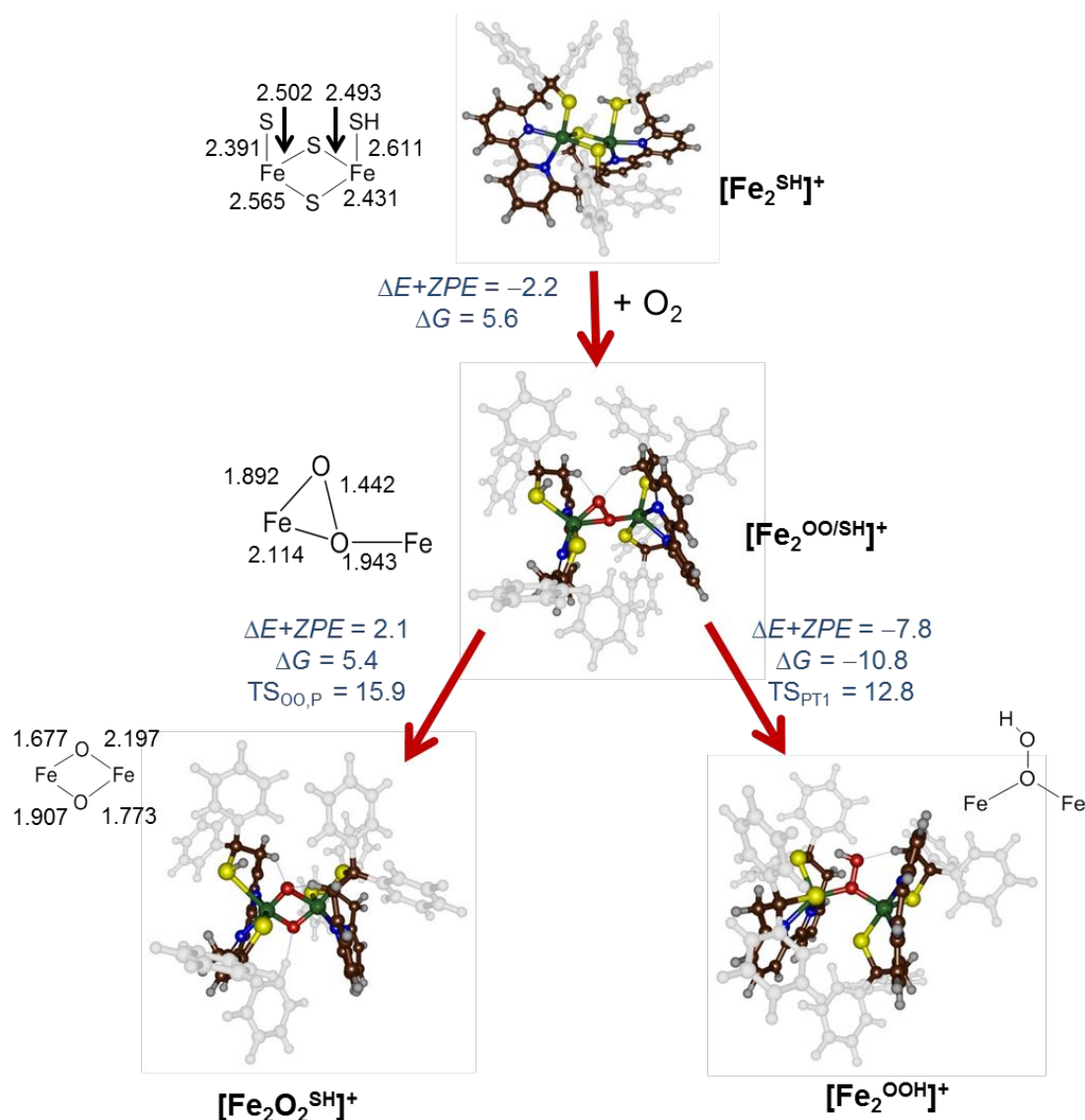
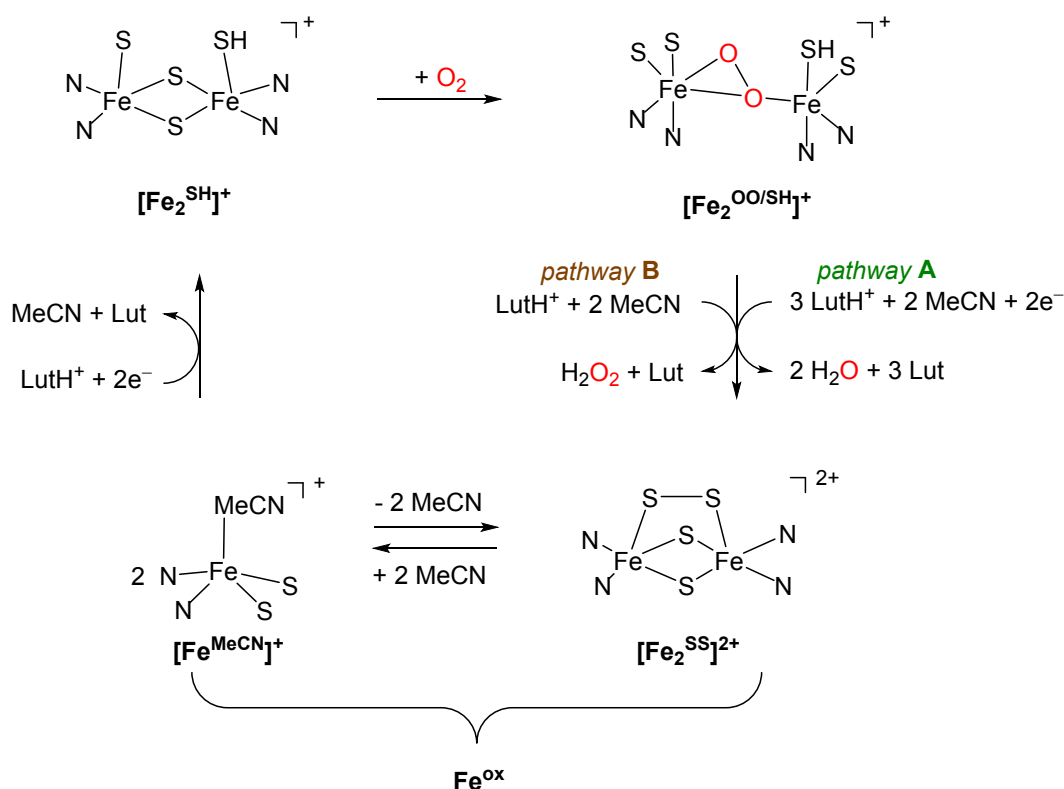


Figure 5. Optimized geometries of [Fe₂SH]⁺, [Fe₂OO/SH]⁺, [Fe₂(μ-O)₂SH]⁺ and [Fe₂OOH]⁺ in the antiferromagnetically coupled singlet spin state with bond lengths in angstroms. For the sake of simplicity, the peroxo intermediate is considered only in its protonated form. Reaction energies ($\Delta E+ZPE$) and reaction free energies (ΔG) at 298 K are given in kcal mol⁻¹ for the large model structures. Transition state energies (in kcal mol⁻¹) are reported at UB3LYP/BS1//UB3LYP/BS2 and contain zero-point and solvent corrections.

When electrons are delivered electrochemically, catalysis mainly occurs through a 4H⁺/4e⁻ O₂ reduction mechanism. In this case we propose that O₂ activation involves the formation of either [Fe₂OO/SH]⁺ (or [Fe₂OOH]⁺), which is then immediately reduced at the electrode before it can react with an additional proton. Indeed in Scheme 2, pathway **A** requires the use of three protons and two electrons to convert [Fe₂OO/SH]⁺ into Fe^{ox} with production of water. The full prediction of this complicated mechanism involving four protons and four electrons in

sequential and/or coupled manner to produce H₂O through an O-O bond cleavage is out of the scope of the present study.

Pathways **A** and **B** are both exergonic, even if route **B** has a much lower driving force ($\Delta G^{\text{B}} = -9.2 \text{ kcal mol}^{-1}$ vs $\Delta G^{\text{A}} = -39.2 \text{ kcal mol}^{-1}$). The fact that $[\text{Fe}_2^{\text{SH}}]^+$ has a proton located in the metal coordination sphere means that during the catalytic cycle there is competition between dioxygen bond cleavage and proton transfer, whereby a possible pathway to H₂O₂ formation becomes possible.



Scheme 2. Proposed 4H⁺/4e[−] (electrochemically driven) vs 2H⁺/2e[−] (chemical reducing agent) O₂ reduction catalysis by $[\text{Fe}_2^{\text{SH}}]^+$ (or Fe^{ox}). For the sake of simplicity, only the peroxo $[\text{Fe}_2^{\text{OO/S}}]^+$ intermediate is depicted (and not the hydroperoxo $[\text{Fe}_2^{\text{OOH}}]^+$ species).

Discussion and Conclusion

In this study, we report the reactivity of a unique dinuclear iron-thiolate complex, $[\text{Fe}_2^{\text{SH}}]^+$ (and of its oxidized form, Fe^{ox}), as an efficient ORR (electro)catalyst. Thiolate ligation has been previously proposed to promote the activation of molecular oxygen,⁶⁶ via (i) stabilization of metal–peroxo intermediates,^{67–72} (ii) decrease of the activation barrier for O₂ binding,^{52, 73} (iii) decrease of the metal redox potential,⁷⁴ (iv) increase of the basicity of bound substrates⁷⁵ or (v) labilization of the trans sites.⁷⁴ Our study confirms and further generalizes

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3 this aptitude: the thiolate-rich N₂S₂-donor LS²⁻ ligand, in the presence of Fe^{II} (this study) or
4 Mn^{II} (previously reported work)^{32, 45} promotes O₂ activation at the coordinated transition
5 metal ion. Although the continuous regeneration of the initial complex [Fe₂^{SH}]⁺ during
6 catalytic turnover entails that sulfur-oxygenation is limited in the present system during the
7 ORR catalysis, this is not the case for other mononuclear Fe-thiolate complexes mainly
8 reported by the Limberg and Goldberg groups.⁷⁶⁻⁸⁰ Clearly, the thiolate groups of the LS²⁻
9 ligand system are much more difficult to oxidize than the cysteinate group used in the
10 Limberg and Goldberg systems. Indeed, our calculations indicate little or no spin density on
11 the thiolate groups.

12
13 A peculiar feature of these [M₂^{SH}]⁺ catalysts (M = Fe, Mn) is the presence of one thiol group
14 coordinated to the metal centers that can act as a proton relay during catalysis. This is a
15 unique example of a sulfur-based pendant acid in ORR catalysis, because most of them are O-
16 and/or N-based (carboxylic acids, ammonium, urea, etc.).^{1, 81-82} The nature of the proton relay
17 can still play a role as potential tunable agent of the selectivity; however no general trend can
18 be drawn up now. Interestingly, while Kojima et al observed an enhancement for 4H⁺/4e⁻ O₂
19 reduction catalysis in the presence of pendant pyridine close to a Cu site,²⁸ Machan et al.
20 described a Schiff-based Mn catalyst with a pendant phenol that displays selective 2H⁺/2e⁻ O₂
21 reduction activity.³³

22
23 The investigation of isostructural iron- and manganese-based catalysts evidences that even if
24 both display similar selectivity in the presence of a chemical reducing agent under similar
25 experimental conditions, [Fe₂^{SH}]⁺ is ~ 6 times more reactive than [Mn₂^{SH}]⁺, (on the basis of
26 the TOF_i values determined in the presence of Me₈Fc). This difference in efficiency can be
27 related to the reduction potential of the catalysts, [Fe₂^{SH}]⁺ being more easily oxidized than
28 [Mn₂^{SH}]⁺ (see above).

29
30 The most attractive properties of [Fe₂^{SH}]⁺ as an ORR catalyst are that: (i) this is the first
31 homogeneous ORR catalyst based on non-heme iron and (ii) its selectivity can be tuned as a
32 function on the mode of electron delivery (chemical vs electrochemical catalysis).

33
34 Although iron porphyrin systems have been largely investigated,^{1, 13-15, 83} this is not the case
35 for non-heme iron complexes. To our knowledge, only one family of Fe complexes supported
36 by non-heme ligands has been previously reported in the ORR domain.⁸⁴ However, the
37 described tetrapyrrolyl mononuclear complexes only react with O₂ stoichiometrically, and not
38 catalytically (the initial complex can be regenerated only in a two-step process). We believe
39 that the discovery of such non-heme iron catalyst for ORR catalysis will pave the way to new
40 structure-reactivity correlation studies encompassing both heme and non-heme compounds.
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As hydrogen peroxide is generated in both cases (predominantly and up to ~95% or in less than ~10% in chemical and electrochemical catalysis, respectively), it can be proposed that a common intermediate, that is, the calculated iron-peroxo complex $[\text{Fe}_2^{\text{OO/S}}]^+$, is generated during catalysis (Scheme 2). The reactivity of such a peroxo intermediate has been proposed to be central for rationalizing the selectivity of homogeneous ORR catalysts. A study on a series of Cu^{II} catalysts with tri- and tetrapodal nitrogen-donor ligands has evidenced that the electronic and structural properties of the generated peroxo intermediates are critical for tuning the selectivity of the ORR process.²³

The tunable selectivity of the $[\text{Fe}_2^{\text{SH}}]^+$ catalyst as a function of the electron delivering manner implies that the peroxo $[\text{Fe}_2^{\text{OO/S}}]^+$ (or $[\text{Fe}_2^{\text{OOH}}]^+$) intermediate can either (i) react with one proton to produce H_2O_2 (with Me_8Fc as reducing agent), or (ii) be reduced to generate H_2O as the main product after successive reduction and protonation steps (under electro-assisted catalysis). These two pathways are supported by DFT calculations, which predict that both are thermodynamically feasible. The production of H_2O_2 is further supported by the fact that the formation of the hydroperoxo intermediate is thermodynamically and kinetically favored over the generation of high valent Fe^{IV} di-oxo species arising from “direct” O-O bond rupture (see Figure 5). Concerning H_2O production, the O-O bond cleavage should thus occur after further reduction of the (hydro)peroxo complex. The tuning of the selectivity can be interpreted in terms of a competition between acid-base and redox processes with the $[\text{Fe}_2^{\text{OO/S}}]^+$ (or $[\text{Fe}_2^{\text{OOH}}]^+$) peroxo as the key intermediate. Indeed, when the concentration of the chemical reductant is increased less H_2O_2 is generated (see Table S3), suggesting that the peroxo intermediate needs to be in a more reducing environment to be reduced (and then to break the O-O bond to generate H_2O) before it gets protonated (to generate H_2O_2). When electrochemically driven, the electron transfer kinetics for its reduction is likely faster because it occurs in the reaction-diffusion layer at the immediate vicinity of the electrode, compared to bulk solution.

The present study thus represents a substantial contribution towards the understanding of the factors controlling catalytic efficiency and selectivity of homogeneous ORR catalysis, in terms of ligand design, nature of transition metal(s), proton and electron delivery.

Supporting Information. Additional experimental data, crystallographic data, spectroscopic characterizations, redox data and DFT calculations. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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