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Proton Reduction Catalysts

[FeFe]-Hydrogenase Mimic Employing $\kappa^2$-C$_7$N-Pyridine Bridgehead Catalyzes Proton Reduction at Mild Overpotential


Abstract: Two novel $\kappa^2$-C$_7$N-pyridine bridged [FeFe]-H$_2$ase mimics (1 and 2) have been prepared and are shown to function as efficient molecular catalysts for electrocatalytic proton reduction. The elemental and structural composition of the complexes are confirmed by NMR and IR spectroscopy, high-resolution mass spectrometry and single-crystal X-ray diffraction. Electrochemical investigations reveal that the complexes reduce protons at their first reduction potential, resulting in the lowest overpotential (120 mV) ever reported for [FeFe]-H$_2$ase mimics in proton reduction catalysis when mild acid (phenol) is used as proton source.

Introduction

To facilitate large-scale production of cheap renewable energy that can be stored cost-effectively, there is a great demand for catalysts that can efficiently produce dihydrogen from water and are preferably made from earth-abundant transition metals. The [FeFe]-hydrogenase ([FeFe]-H$_2$ase) enzymes catalyze the reversible reduction of protons at ambient conditions with a low overpotential. It is envisioned that synthetic mimics of the active site of the [FeFe]-H$_2$ase enzyme can serve as efficient catalysts for proton reduction in renewable fuel applications. Ever since the structure of the active site of the [FeFe]-H$_2$ase enzyme was elucidated, a large number of synthetic mimics has been developed. It has been shown that it is possible to develop mimics that operate at similar and even higher rates than the natural enzyme. Next to a high rate it is important to develop catalytic systems that operate at a low overpotential, as it is essential to reduce the loss of energy in the overall conversion of electrical energy to chemical energy. Despite intensive investigations, the development of synthetic [FeFe]-H$_2$ase mimics that operate at a mild overpotential remains a challenge to be solved.

In most [FeFe]-H$_2$ase mimics the two iron atoms are connected via a bridging dithiolato fragment and each iron atom is coordinated by three terminal carbonyl ligands. Along these lines, the first class of mimics was based on structures in which a propanedithiolato ($\mu$-pdt) fragment bridged the two iron atoms, and this group of mimics has been studied in detail by several groups including Pickett, Darensbourg and Rauchfuss. The next generation of mimics focused on analogues with the biologically relevant aza-dithiolato ($\mu$-adt) bridge, wherein the basic amine functionality can act as “proton relay”. Among the many other dithiolato bridges, the more rigid benzenedithiolato ($\mu$-bdt) bridge has been studied in some detail and found to give rather active proton reduction catalysis. Most parent hexacarbonyl complexes can undergo substitution of one or more of the six carbonyls by a great variety of ligands, which by now has resulted in a library of hundreds of reported complexes. Typically, these substitutions lead to an increase in the complex’ overall basicity, resulting in higher reduction potentials. Given the inverse relationship between overall basicity and redox potential, such terminal ligand substitutions generally do not lead to catalysts that operate with high rates at a mild overpotential.

In most [FeFe]-H$_2$ase mimics the two iron atoms are connected via a bridging dithiolato fragment and each iron atom is coordinated by three terminal carbonyl ligands. Along these lines, the first class of mimics was based on structures in which a propanedithiolato ($\mu$-pdt) fragment bridged the two iron atoms, and this group of mimics has been studied in detail by several groups including Pickett, Darensbourg and Rauchfuss. The next generation of mimics focused on analogues with the biologically relevant aza-dithiolato ($\mu$-adt) bridge, wherein the basic amine functionality can act as “proton relay”. Among the many other dithiolato bridges, the more rigid benzenedithiolato ($\mu$-bdt) bridge has been studied in some detail and found to give rather active proton reduction catalysis. Most parent hexacarbonyl complexes can undergo substitution of one or more of the six carbonyls by a great variety of ligands, which by now has resulted in a library of hundreds of reported complexes. Typically, these substitutions lead to an increase in the complex’ overall basicity, resulting in higher reduction potentials. Given the inverse relationship between overall basicity and redox potential, such terminal ligand substitutions generally do not lead to catalysts that operate with high rates at a mild overpotential.

Figure 1. Common [FeFe]-H$_2$ase mimics employing different dithiolato-based bridges.

The beneficial effect on the catalytic overpotential when transitioning from a $\mu$-pdt to a $\mu$-adt bridge suggests that modification of the bridging ligand could be key to lowering the overpotential. The class of hydrogenase mimics based on a
bridging pyridine–monothiolato ligand is relatively unexplored. The synthesis of such complexes has been described from the corresponding thioester,[17] and to the best of our knowledge there is only one earlier report on the catalytic activity in the presence of acetic acid.[18]

With the aim to study the effect of changing the dithiolato bridge for a pyridine bridge on the catalytic overpotential, we report novel [FeFe]-H$_2$ase mimics 1 and 2, in which the iron atoms are connected by a pyridine bridge in a $\kappa^2$-CN fashion (Figure 2). The pyridine ring of 1 is substituted with a thioisopropyl group and that of 2 with a dimethylamine group, which may serve as a “proton relay” due to its basic nature. The catalysts are capable of reducing protons from acids that are weaker than acetic acid at their first reduction potential, resulting in a catalytic overpotential that is up to 240 mV lower than that of [Fe$_2$(µ-bdt)(CO)$_6$].

Figure 2. Structures and relative overpotentials of the novel $\kappa^2$-CN pyridine di-iron complexes.

**Results and Discussion**

**Synthesis and Characterization**

Ligand L$_1$ is obtained in one step according to a literature procedure, by reacting 2,3-dichloropyridine with an excess of sodium isopropylthiolate at 85 °C in an $S_2$Ar reaction.[19] Ligand L$_2$ is prepared in two steps starting from 3-amino-2-chloropyridine. First, the free aminopyridine is methylated using formaldehyde and formic acid in an Eschweiler–Clarke reaction. Subsequently, the chloride substituent is displaced by isopropylthiolate in an $S_2$Ar reaction. Complex 1 is obtained in 27 % yield by heating a mixture of L$_1$ and iron precursor Fe$_2$(CO)$_9$ (2 equiv.) in toluene at 100 °C for 15 min under an inert atmosphere (Scheme 1). Complex 2 (17 % yield) is prepared using an identical procedure in the presence of ligand L$_2$ (for the full synthetic protocol of 1 and 2, see Supporting information, Section 2).

Compounds 1 and 2 are fully characterized by IR and NMR spectroscopy, high-resolution mass spectrometry and single-crystal X-ray diffraction (for full characterization, see Supporting information, Section 2). The IR spectra of both complexes display the typical fingerprint of di-iron hexacarbonyl complexes (Table 1 and Supporting information, Figure S5–S7).[20,21] The bands of 2 appear at lower stretching frequencies compared to those of 1, in line with more electron-rich iron centers of 2 due to the electron-donating amino substituent. The $^1$H NMR spectrum of 1 shows two signals with a doublet splitting pattern for the CH$_3$ groups of the bridging isopropyl substituent (Supporting information, Figure S1). The doublets in the $^1$H NMR spectrum indicate that the CH$_3$ groups are diastereotopic.

Figure 3. X-ray crystal structures of 1 (left) and 2 (right) with displacement ellipsoids drawn at 50 % probability. Hydrogen atoms have been omitted for clarity. Color code: C, gray; N, blue; O, red; sulfur, yellow; iron, orange.

X-ray analysis of 1 confirms that it exists as two enantiomers, making the methyl groups diastereotopic (Supporting information, Figure S8). The $^{13}$C NMR spectrum of 1 shows a signal at 193.5 ppm, which is typical for a species with a Fe–C bond (Supporting information, Figure S2).[22] Complex 2 shows similar features in its $^1$H and $^{13}$C NMR spectrum (Supporting information, Figure S3–S4).

Table 1. IR stretching frequencies of 1 and 2 compared to [Fe$_2$(µ-bdt)(CO)$_6$] and [Fe$_2$(µ-pdt)(CO)$_6$].

<table>
<thead>
<tr>
<th>Catalyst</th>
<th>IR stretches of three main bands [cm$^{-1}$]</th>
<th>Solvent</th>
</tr>
</thead>
<tbody>
<tr>
<td>Complex 1</td>
<td>2065, 2024, 1988</td>
<td>Pentane</td>
</tr>
<tr>
<td>Complex 2</td>
<td>2062, 2022, 1986</td>
<td>Pentane</td>
</tr>
<tr>
<td>[Fe$_2$(µ-bdt)(CO)$_6$]$^{[24]}$</td>
<td>2079, 2044, 2004</td>
<td>Hexane</td>
</tr>
</tbody>
</table>

Complex 1 was crystallized by layering a dichloromethane solution with pentane. Single crystals of 2 were obtained by slow evaporation of a pentane solution under argon at 5 °C. The crystal structures are similar to previously reported $\kappa^2$-CN pyridine bridged di-iron compounds (Figure 3).[17,18] The Fe1–Fe2 bond lengths of 1 and 2 are 2.5741(5) and 2.5726(5) Å, respectively. These bonds are slightly longer than the Fe–Fe bond length in [Fe$_2$(µ-bdt)(CO)$_6$] (2.480(2) Å$^{[20]}$) and [Fe$_2$(µ-pdt)(CO)$_6$] (2.5103(11) Å$^{[25]}$). The Fe2–C7 bond length is 1.996(3) Å for 1 and 1.992(2) Å for 2 and this is comparable to the Fe–C bond length in similar $\kappa^2$-CN bridged hydrogenase mimics.[17] The Fe1–N1 bond is 1.984(2) Å for 1 and a little shorter for 2 (1.9744(19) Å).

Scheme 1. Synthesis of novel [FeFe]-H$_2$ase mimics 1 and 2.
Redox Behavior of 1 and 2 in the Absence of Acid

Cyclic voltammetry of 1 in acetonitrile reveals an irreversible reduction wave with a cathodic peak potential of around –1.8 V (vs. Fc/Fc⁺; all potentials are reported against this redox couple), followed by at least one anodically shifted re-oxidation wave at around –0.6 V (Figure 4). Complex 2 displays similar redox behavior as 1, but with a 50 mV cathodic shift in both the reduction and oxidation event. This shift is caused by the more electron-donating nature of the dimethylamine substituent of 2 as compared to the isopropylthiol substituent of 1, and this is in line with the observations in the IR measurements (vide infra).

Both 1 and 2 are more difficult to reduce than known [FeFe]-H₂ase mimics [Fe₂(μ-bdt)(CO)₆] and [Fe₂(μ-pdt)(CO)₆] (Table 2 and Supporting information, Figures S13–S14). For both 1 and 2 the peak current of the reduction wave varies linearly with the square root of the scan rate, indicative of a solution-based redox event (Supporting information, Figures S9–S12).[26]

Spectroelectrochemical studies provided more insight into the structures of the species that are formed upon reduction of 1 and 2. Linear sweep voltammetry of 1, while probing the IR spectrum, reveals bleaching of IR signals associated to the neutral complex, concomitant with the appearance of new red-shifted bands assigned to the reduced species 1⁻ (Figure 5a). The absorption-difference spectra show a small band growing in at 1730 cm⁻¹, which is characteristic for reduced diiron compounds with a bridging carbonyl ligand.[27,28] Complex 2 shows similar bands in its IR spectrum upon reduction (Figure 5b).

The semi-integral convolution plot of 1 in the presence of an equimolar amount of ferrocene suggests a one-electron reduction process, assuming that the diffusion constant of 1 and ferrocene are similar (Supporting information, Figure S15). Controlled potential coulometry (–1.9 V) of a solution of 1 confirms the passage of one electron per molecule (Supporting information, Figure S16). Moreover, the IR spectrum of 1 in the presence of 1.6 equiv. of the reducing agent decamethylcobaltocene (E₁/₂ = –1.9 V in MeCN)[29] shows complete reduction to 1⁻ (Supporting information, Figure S17). On the contrary, the IR spectrum of [Fe₂(μ-bdt)(CO)₆], which is known to undergo disproportionation and therefore has a two-electron reduction at E₁/₂ = –1.32 V, shows a mixture of neutral and [FeFe]²⁻ species in the presence of the same amount of reductant (Supporting information, Figure S18). Based on these experiments we prudently conclude that 1 undergoes a one-electron reduction, and 2 is expected to display the same electrochemical behavior, albeit at slightly different potential.

DFT Calculations and XAS Analysis on the Reduced Species 1⁻

The irreversible redox behavior of 1 suggests that the complex displays follow-up chemistry upon reduction. More detailed insight into the structure of the mono-reduced species comes from DFT calculations (Supporting information, Section 6). Computations were performed on the monoanionic 1⁻ and
anionic 1− species, both with all terminal carbonyl ligands and with one bridging carbonyl ligand. A comparison of the computed IR spectra with the experimental spectra reveals that mono-anion B, which contains a bridging carbonyl ligand, shows the best fit (Figure 6b). This is consistent with the spectroelectrochemical measurements that indicate the presence of a bridging carbonyl ligand upon reduction of 1. The significant difference in the calculated and experimental wavenumber for the bridging carbonyl ligand is likely due to its position being greatly affected by the solvent.[30] The Fe–N bond and one of the Fe–S bonds are broken in the mono-reduced species, allowing for the structural rearrangement into a bridging carbonyl species (Figure 6a). Such a rearrangement accounts for the irreversibility of the reduction wave observed in the voltamogram and is reminiscent of chemistry observed with benzene dithiolate analogs.[15,16]

To investigate the effect of the κ2-C,N-pyridine bridge on the catalytic performance of complexes 1 and 2, cyclic voltammetric studies were undertaken in the presence of various weak acids and the reactivity was compared to that of [Fe2(μ-bdt)(CO)6]. In the presence of one equiv. acetic acid (AcOH; pKₐ = 22.3 in CH3CN) the peak potential of 1 undergoes an anodic shift of around 15 mV, indicating an electrochemical event followed by protonation (Figure 7a, inset).[31,32] A slightly larger potential shift (around 25 mV) in the presence of one equiv. of AcOH is observed for 2, revealing that protonation of 2 is thermodynamically more favorable than protonation of 1 (Figure 7b, inset).[31] The currents of the reduction waves of 1 and 2 increase as a function of the acid concentration, confirming catalytic proton reduction at the first reduction potential for both complexes (Figure 7a, b).

A comparison of the catalytic performance of 1 and 2 to that of [Fe2(μ-bdt)(CO)6] reveals that these complexes behave differently (Figure 7a, b). While 1 and 2 reduce protons from AcOH at their first reduction potential, [Fe2(μ-bdt)(CO)6] displays catalysis at a considerably more negative potential than its first reduction potential. However, [Fe2(μ-bdt)(CO)6] is a faster catalyst, as is evident from its sharper catalytic waves along with a higher current (Figure 7a, b black traces). The catalytic parameters are determined as previously reported[33–37] and summarized in Table 3 (For a detailed description of the determination of the catalytic parameters, see Supporting information, Section 7). Complex 2 operates with a three times higher rate than 1, but it is still 250 times slower than [Fe2(μ-bdt)(CO)6] (Table 3, of EXAFS analysis, see Supporting Information, Section 5). Additionally, the Fe–Fe contribution is fitted with an elongated bond length of 2.69(2) Å compared to the original value of 2.60(1) Å. This analysis confirms the breakage of the Fe–N bond and one Fe–S bond and the formation of a bridged CO ligand, as suggested by the IR analysis and DFT computations.
Table 3. Catalytic parameters of 1, 2 and [Fe₂(μ-bdt)(CO)₆] in the presence of different acids (0.1 M NBu₄PF₆ in CH₃CN).[^a]

<table>
<thead>
<tr>
<th>Entry</th>
<th>Catalyst</th>
<th>Proton source</th>
<th>Catalytic E₁/₂ (V vs. Fc/Fc⁺)</th>
<th>η [V]</th>
<th>kcat [M⁻¹ s⁻¹][^b]</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1</td>
<td>AcOH</td>
<td>-1.77</td>
<td>0.57</td>
<td>136[^b]</td>
</tr>
<tr>
<td>2</td>
<td>1</td>
<td>AcOH</td>
<td>-1.77</td>
<td>0.57</td>
<td>391[^b]</td>
</tr>
<tr>
<td>3</td>
<td>[Fe₂(μ-bdt)(CO)₆]</td>
<td>AcOH</td>
<td>-1.94</td>
<td>0.74</td>
<td>1×10⁵[^b]</td>
</tr>
<tr>
<td>4</td>
<td>1</td>
<td>PhOH</td>
<td>-1.76</td>
<td>0.12</td>
<td>1.4[^b]</td>
</tr>
<tr>
<td>5</td>
<td>2</td>
<td>PhOH</td>
<td>-1.81</td>
<td>0.17</td>
<td>6.9[^b]</td>
</tr>
<tr>
<td>6</td>
<td>1</td>
<td>ClAcOH</td>
<td>-1.81</td>
<td>0.88</td>
<td>566[^b]</td>
</tr>
<tr>
<td>7</td>
<td>2</td>
<td>ClAcOH</td>
<td>-1.79</td>
<td>0.86</td>
<td>659[^b]</td>
</tr>
</tbody>
</table>

[^a] E_H₂/PH₂ = -0.028 – 0.05916 × pKₐ – 1.64 V for PhOH and -0.93 V for CIC₆H₄COOH. For AcOH the effect of homoconjugation has been described[^34] and by taking this into account a value of ~1.2 V is obtained and applied as the thermodynamic potential.[^b] Calculated using Dubois' formula as described in the Supporting information, Section 7.

Entries 1–3). The calculated overpotential (η) for both 1 and 2 is 0.57 V, which is 170 mV lower than that of [Fe₂(μ-bdt)(CO)₆]. This large decrease in overpotential is a result of 1 and 2 performing catalysis at the potential of their first reduction.

The low overpotential of 1 and 2 in the catalytic proton reduction of the weak acid AcOH encouraged further studies with the even weaker acid phenol (PhOH; pKₐ = 27.2 in MeCN). Cyclic voltammetry of 1 or 2 in the presence of 500 equiv. PhOH as the proton source reveals a catalytic wave at the first reduction potential of the catalysts (Figure 8a, b). Complex 2 operates with a five times higher rate than 1 and again is significantly slower than [Fe₂(μ-bdt)(CO)₆], in line with studies performed with AcOH (Figure 8c and Table 3, entries 4–5). Interestingly, the catalytic overpotentials of both 1 and 2 in the reduction of PhOH are remarkably low, with 1 operating at an overpotential of only 120 mV, which is the lowest overpotential ever reported for a [FeFe]-H₂ase mimic, to the best of our knowledge.

To investigate whether the dimethylamine substituent of 2 can be applied as a “proton relay” and thereby also affect the catalytic overpotential, catalytic studies were undertaken using...
the slightly stronger acid, chloroacetic acid (ClCH₂COOH; \( pK_a = 15.3 \)). Cyclic voltammetry of 2 in the presence of ClCH₂COOH reveals that the acid is not strong enough to protonate the complex prior to reduction, as evident from the lack of an anodic shift in the reduction wave (Figure 9b). A significant anodic shift of the reduction wave of the catalyst would be expected if the complex was protonated prior to reduction. Complex 1 shows similar behavior as 2, whereas [Fe₂(μ-bdt)-

Figure 9. Electrocatalytic reduction of ClCH₂COOH. (a) Cyclic voltammetry (0.1 V s⁻¹) of 1.0 mM 1 in CH₃CN containing 0.1 M NBu₄PF₆ and 0–50 equiv. ClCH₂COOH. (b) Cyclic voltammetry (0.1 V s⁻¹) of 1.0 mM 2 in CH₃CN containing 0.1 M NBu₄PF₆ and 0–50 equiv. ClCH₂COOH. (c) Cyclic voltammetry (0.1 V s⁻¹) of 1.0 mM [Fe₂(μ-bdt)(CO)₆] in CH₃CN containing 0.1 M NBu₄PF₆ and 0–50 equiv. ClCH₂COOH.

Figure 10. (a) Tafel plots of 1, 2 and [Fe₂(μ-bdt)(CO)₆] in the presence of AcOH. (b) Tafel plots of 1 and 2 in the presence of PhOH and ClCH₂COOH. The value of TOF_max is extrapolated for a 1 M concentration of protons.
in chemical bonds. Initial is the key challenge to efficient storage of electrical energy. The impact of proton reduction potential to drive catalysis. As a consequence, proton reduction catalysis is demonstrated with the lowest overpotential (120 mV) ever reported for [FeFe]-H2ase mimics. The higher rate. For ClCH2COOH, the potential is similar efficiency (Figure 10b). In the case of PhOH the lower overpotential than [Fe2(μ-bdt)(CO)6]. The higher catalytic rate than [Fe(μ-bdt)(CO)6]. Xinmin in dichloromethane. 1H NMR (400 MHz, CD2Cl2) δ (ppm) = 7.44 (d, J = 5.4 Hz, 1H), 7.10 (d, J = 7.9 Hz, 1H), 6.67 (dd, J = 7.9, 5.4 Hz, 1H), 3.33 (septet, J = 6.6 Hz, 1H), 2.61 (septet, J = 6.4 Hz, 1H), 1.49 (d, J = 6.4 Hz, 3H), 1.48 (d, J = 6.8 Hz, 3H), 1.33 (d, J = 6.5 Hz, 3H), 1.32 (d, J = 6.5 Hz, 3H). 13C NMR (101 MHz, CD2Cl2) δ (ppm) = 213.60, 211.55, 211.15, 193.54, 150.86, 147.29, 132.49, 120.75, 44.49, 37.95, 30.27, 27.31, 26.88, 22.99. FTIR (penta) cm⁻¹ = 2065, 2026, 1994, 1985, 1972, 1970. The thus obtained pure compound was dissolved in 25 mL of pentane after which the volatiles were removed under vacuum to afford complex 1 as an orange powder (27 % yield with respect to L1). Single crystals suitable for X-ray diffraction analysis were obtained by liquid-liquid diffusion of pentane into a solution of 1 in dichloromethane. 1H NMR (400 MHz, CD2Cl2) δ (ppm) = 7.10 (d, J = 5.4 Hz, 1H), 7.02 (d, J = 7.9 Hz, 1H), 6.67 (dd, J = 7.9, 5.4 Hz, 1H), 3.33 (septet, J = 6.6 Hz, 1H), 2.61 (septet, J = 6.4 Hz, 1H), 1.49 (d, J = 6.4 Hz, 3H), 1.48 (d, J = 6.8 Hz, 3H), 1.33 (d, J = 6.5 Hz, 3H), 1.32 (d, J = 6.5 Hz, 3H). 13C NMR (101 MHz, CD2Cl2) δ (ppm) = 213.60, 211.55, 211.15, 193.54, 150.86, 147.29, 132.49, 120.75, 44.49, 37.95, 30.27, 27.31, 26.88, 22.99. FTIR (penta) cm⁻¹ = 2065, 2026, 1994, 1985, 1972, 1970. HRMS (FD) calcld. for [1+] (C17H17Fe2NO4S2)⁺ 506.91961, found 506.92126. Crystallographic details 1: C17H17Fe2NO6S2, Fw = 507.14, orange block, 0.630 × 0.403 × 0.200 mm, monoclinic, P2₁/n (No: 14), a = 13.9060(12), b = 9.3308(8), c = 17.1082(14), β = 93.308(8), V = 20890.3 Å³, Z = 4, D = 1.612 g/cm³, μ = 1.621 mm⁻¹. 21812 Reflections were measured up to a resolution of (sin θ/λ)max = 0.84 Å⁻¹. 3693 Reflections were unique (Rint = 0.0417), of which 3218 were observed (I > 2σ(I)). 257 Parameters were refined with 0 restraints. R = 0.0308/0.1043. R = 0.0390/0.1216. S = 1.026. Residual electron density between -0.483 and 0.711 e Å⁻³. Synthesis of Complex 2 An oven-dried Schlenk flask was charged with Fe2(CO)9 (82 mg, 0.23 mmol) and equipped with a gas bubbler filled with oil via a needle through the septum of the Schlenk flask. The thus obtained pure compound was dissolved in 8 mL of toluene and transferred to the iron precursor and the resulting mixture was heated to 100 °C in a preheated oil bath. The reaction progress was monitored by IR spectroscopy. After 15 minutes the dark red mixture was cooled to room temperature. The volatiles, including the side-product Fe(CO)5, were carefully removed under vacuum. The crude product was purified by column chromatography (silica, eluent: gradient from hexane to hexane/CH2Cl2, 80:20). The thus obtained pure compound was dissolved in 25 mL of pentane after which the volatiles were removed under vacuum to afford complex 1 as an orange powder (27 % yield with respect to L1). Single crystals suitable for X-ray diffraction analysis were obtained by liquid-liquid diffusion of pentane into a solution of 1 in dichloromethane. 1H NMR (400 MHz, CD2Cl2) δ (ppm) = 7.10 (d, J = 5.4 Hz, 1H), 7.02 (d, J = 7.9 Hz, 1H), 6.67 (dd, J = 7.9, 5.4 Hz, 1H), 3.33 (septet, J = 6.6 Hz, 1H), 2.61 (septet, J = 6.4 Hz, 1H), 1.49 (d, J = 6.4 Hz, 3H), 1.48 (d, J = 6.8 Hz, 3H), 1.33 (d, J = 6.5 Hz, 3H), 1.32 (d, J = 6.5 Hz, 3H). 13C NMR (101 MHz, CD2Cl2) δ (ppm) = 213.60, 211.55, 211.15, 193.54, 150.86, 147.29, 132.49, 120.75, 44.49, 37.95, 30.27, 27.31, 26.88, 22.99. FTIR (penta) cm⁻¹ = 2067, 2063, 2024, 2000, 1991, 1984.
Crystallographic details

2: C_{16}H_{16}Fe_{2}N_{2}O_{6}S, Fw = 476.07, dark yellow block, 0.128 x 0.380 x 0.506 mm, monoclinic, P2_{1}/c (No: 14)), a = 14.2199(9), b = 14.2199(9), c = 17.2070(11) Å, β = 109.50(2)°, V = 3959.7(4) Å³, Z = 8, D_{x} = 1.597 g/cm³, µ = 1.604 mm⁻¹. 114346 Reflections were measured up to a resolution of (sin θ/λ)_{max} = 0.84 Å⁻¹. 6953 Reflections were unique (R_{int} = 0.0423), of which 6052 were observed (I > 2σ(I)). 495 Parameters were refined with 0 restraints. R_{1}/wR_{2} [I > 2σ(I)]: 0.0310/0.01046. R_{1}/wR_{2} [all refl.]: 0.00391/ 0.1171. S = 0.989. Residual electron density between −0.374 and 0.332 e Å⁻³.

CCDC 1893259 (for 1), and 1893262 (for 2) contain the supplementary crystallographic data for this paper. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre.

Conflict of interest

The authors declare no conflict of interest.

Acknowledgments

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Keywords: Proton reduction · Hydrogenase mimics · Iron · Electrochemistry · Homogeneous catalysis