Metal–Ligand Exchange Coupling Promotes Iron-Catalyzed Electrochemical CO₂ Reduction at Low Overpotentials

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ABSTRACT: Biological and heterogenous catalysts for electrochemical CO$_2$ reduction often exhibit a high degree of electronic delocalization that serves to minimize overpotential and maximize selectivity over hydrogen evolution. Here, we report a molecular iron(II) complex that achieves a similar feat as a result of strong metal–ligand exchange coupling. This interaction promotes an open-shell singlet electronic structure that drives the electrochemical reduction of CO$_2$ to CO with over 90% selectivity and turnover frequencies of 100,000 s$^{-1}$ at low overpotentials, with no degradation over 20 hours. The decrease in the thermodynamic barrier engendered by this strong metal–ligand exchange coupling enables homogeneous CO$_2$ reduction catalysis in water without compromising reaction selectivity.

INTRODUCTION

The electrochemical carbon dioxide reduction reaction provides opportunities to synthesize value-added products in a sustainable manner.$^{1,2}$ Efficient catalysts for this reaction are required to selectively drive CO$_2$ reduction over the thermodynamic and kinetically competitive hydrogen evolution reaction.$^3$ Biological$^4$, heterogenous$^5$, and molecular$^3$ catalysts have been studied for CO$_2$ reduction, and molecular systems in particular offer potential advantages such as their small size, relative to enzymes, and the fact that they can be tuned with a level of atomic precision that is inaccessible with heterogeneous materials. However, despite advances in the development of molecular catalysts with highly specialized ligand scaffolds, including cyclams$^{6,7}$, porphyrins and phthalocyanines$^{8-10}$, and tricarbonyl-bipyridines$^{11,12}$, it remains a challenge to design molecules that exhibit low overpotentials, high turnover numbers, and compatibility with aqueous electrolytes. In this regard, one key advantage of biological and heterogeneous catalysts is their ability to minimize overpotential and maximize selectivity for CO$_2$ reduction via electronic delocalization, either through electron-tunneling pathways or a high local density of states,
respectively. Such systems can thereby achieve organization and separation of reducing equivalents in a way that is challenging to design in molecular systems.

Previous work from our laboratories has explored the chemistry of the PY5Me2 ligand (2,6-bis(1,1-bis(2-pyridyl)ethyl)pyridine) and its molybdenum13–15 and cobalt15,16 derivatives as catalysts for the hydrogen evolution reaction. We reasoned that by incorporating a redox-active pendant such as terpyridine within this ligand scaffold, it would be possible to increase metal–ligand orbital mixing and delocalization of electron density away from the metal center to favor CO2 reduction (Figure 1a). Indeed, the suppression of metal-centered reduction can limit the formation of off-pathway intermediates necessary for hydrogen evolution (e.g., metal hydrides)17,18 and thereby favor CO2 reduction catalysis. Our synthetic approach was inspired by mononuclear iron hydrogenase enzymes, which catalyze the reduction of methenyl-H4MPT+ to methylene-H4MPT in methanogenic archaea19. In these systems, heterolysis of H2 and hydride shuttling is enabled by a single redox-innocent iron(II) site that synergistically interacts with a redox non-innocent guanyl pyridinol cofactor in the primary coordination sphere to provide reducing equivalents (Figure 1a).

Here we report an iron polypyridine complex [Fe(tpyPY2Me)(CH3CN)]2+ (tpyPY2Me = 6-(1,1-di(pyridin-2-yl)ethyl)-2,2′:6′,2″-terpyridine), that is the first example of a molecule to demonstrate this principle of electronic delocalization. In particular, strong exchange coupling between the redox-active tpyPY2Me ligand and iron(II) center promotes electrochemical reduction of CO2 to CO at low overpotentials with greater than 90% selectivity and high turnover frequencies. Synthetic reduction of the complex enables access to the catalytically active species [Fe(tpyPY2Me)]0, which exhibits an unusual open-shell singlet ground state comprised of an intermediate-spin iron(II) (SFe = 1) antiferromagnetically coupled to a doubly-reduced triplet
terpyridine ligand ($S_{tpy} = 1$). Significantly, this unique electronic structure also enables this catalyst to operate in water with a high selectivity for CO$_2$ reduction over the hydrogen evolution reaction.

RESULTS AND DISCUSSION

The tpyPY2Me ligand and corresponding iron(II) complex $[\text{Fe(tpyPY2Me)(CH}_3\text{CN)}]^2^+$ ([Fe1]$^{2^+}$) were prepared following modifications of previously reported procedures for PY5Me$_2$ and its complexes (see supplementary information). In order to understand the role of the iron center in catalysis, we also prepared the diamagnetic zinc(II) analog $[\text{Zn(tpyPY2Me)}]^2^+$ ([Zn1]$^{2^+}$). Finally, we prepared the complexes $[\text{Fe(PY5Me}_2\text{(CH}_3\text{CN)}]^2^+^{(20)}$ and $[\text{Zn(PY5Me}_2\text{(CH}_3\text{CN)}]^2^+^{(16)}$ to aid in identifying the role of the redox-active terpyridine fragment of the polypyridyl scaffold. A comparison of the solid-state structures of [Fe1]$^{2^+}$ and $[\text{Fe(PY5Me}_2\text{(CH}_3\text{CN)}]^2^+^{(20)}$ obtained via single-crystal X-ray diffraction revealed that the enhanced rigidity of the tpyPY2Me ligand enforces a more distorted pseudo-octahedral ($O_h$) geometry in [Fe1]$^{2^+}$ relative to $[\text{Fe(PY5Me}_2\text{(CH}_3\text{CN)}]^2^+$, which adopts an almost ideal $O_h$ geometry (Figure 1b and Figure S1; Tables S1–S4). In particular, occupation of the iron equatorial plane by the three terpyridine nitrogen atoms and coordination of the pyridine arms to axial and equatorial positions induces severe axial distortions in [Fe1]$^{2^+}$ that are absent in $[\text{Fe(PY5Me}_2\text{(CH}_3\text{CN)}]^2^+^{(20)}$ (Table S2). This distortion also significantly compresses the axial bond angle in [Fe1]$^{2^+}$ ($173.64(6)^\circ$) whereas the corresponding angle in $[\text{Fe(PY5Me}_2\text{(CH}_3\text{CN)}]^2^+$ is nearly linear ($178.53(7)^\circ$) (Figure 1C).

The structure of [Fe1]$^{2^+}$ remains more distorted than that of $[\text{Fe(PY5Me}_2\text{(CH}_3\text{CN)}]^2^+^{(20)}$ in solution, as verified by $^1\text{H}$ NMR spectra collected for both complexes in CD$_3$CN at 295 K. The spectrum of $[\text{Fe(PY5Me}_2\text{(CH}_3\text{CN)}]^2^+^{(20)}$ features five sharp signals consistent with a $C_{2v}$ symmetric complex with low-spin Fe$^{2^+}$ ($S = 0$), whereas the $^1\text{H}$ NMR spectrum of [Fe1]$^{2^+}$ features slightly more broadened resonances with no well-defined symmetry (Figure S2). Using the Evans method,
we measured a room temperature magnetic moment of 1.93 $\mu_B$ for [Fe1]$^{2+}$, which is indicative of an intermediate-spin Fe$^{2+}$ ($S = 1$) ground state (Figure S3). Because the [Fe1]$^{2+}$ bond lengths measured at 100 K are in agreement with those of other low-spin Fe$^{2+}$ polypyridyl complexes$^{20}$, the collective data suggest that [Fe1]$^{2+}$ undergoes a spin-state transition between 100 K and room temperature. In line with this interpretation, at ~300 K the $^1$H NMR spectrum of [Fe1]$^{2+}$ displays severe line broadening consistent with paramagnetic iron(II), whereas cooling to 233 K results in a spectrum with much shaper peaks, indicative of an increase in the population of low-spin iron(II) (Figure S4). Zero-field $^{57}$Fe Mössbauer spectra collected for [Fe1]$^{2+}$ at 5 and 295 K further support such a spin state transition (Figures S5–6 and Tables S5–6). At 5 K, the spectrum features a
symmetric Lorentzian doublet with isomer shift ($\delta$) and quadrupole splitting ($|\Delta E_Q|$) values of 0.3562(4) and 1.0203(8) mm/s, respectively, consistent with a low-spin iron(II) as corroborated by density functional theory (DFT) calculations (Tables S5–6)\textsuperscript{21,22}. In contrast, the Mössbauer spectrum measured at 295 K features an asymmetric quadrupole doublet that can be fit to two iron sites. The parameters of the major site (84(1)%) are nearly identical to the 5 K spectrum when accounting for second-order doppler effects ($\delta = 0.285(1)$ mm/s, $|\Delta E_Q| = 1.056(2)$ mm/s), while the isomer shift of the minor site (16(1)%) is 0.62(1) mm/s, indicative of a higher spin-state iron(II) species. The distinct iron(II) spin states in $[\text{Fe1}]^{2+}$ and $[\text{Fe(PY5Me2)(CH3CN)}]^{2+}$ can be understood from the simplified Walsh diagram presented in Figure 1c. Axial distortions engendered by the rigid tpyPY2Me ligand lead to a decrease in symmetry from idealized $O_h$ to $C_{2v}$, and this distortion significantly stabilizes the $d_{z^2}$ orbital and destabilizes the $d_{xz}$ through $z$-component mixing of the $t_{1u}$ and $t_{2g}$ orbitals. The stabilization of the $d_{z^2}$ orbital and disruption of $t_{2g}$ degeneracy facilitates the formation of intermediate-spin iron(II) in $[\text{Fe1}]^{2+}$ near ambient temperature. These marked differences in the ground-state electronic structures of $[\text{Fe1}]^{2+}$ and $\text{Fe(PY5Me2)(CH3CN)}^{2+}$ contribute to their distinct electrochemical and catalytic behavior (\textit{vide infra}).

The presence of a redox-active terpyridine moiety in tpyPY2Me also gives rise to unique electrochemical behavior for $[\text{Fe1}]^{2+}$ and $[\text{Zn1}]^{2+}$ relative to their PY5Me2 analogues. The room-temperature cyclic voltammogram (CV) of $[\text{Fe1}]^{2+}$ collected in CH\textsubscript{3}CN under Ar exhibits two reversible features: a one-electron reduction at $E_{1/2} = 0.66$ V (versus Fc/Fc\textsuperscript{+}) assigned to the iron(III/II) couple and a two-electron redox process assigned to tpyPY2Me\textsuperscript{0/2−} at −1.43 V (Figure 2a). No additional features were observed when scanning to more negative potentials. Under the same conditions, the cyclic voltammogram of $[\text{Zn1}]^{2+}$ exhibits two reversible, one-electron ligand-centered reductions at $E_{1/2} = −1.56$ and −2.07 V and an additional irreversible redox event
(tpyPY2Me^{2-/3-}) at −2.41 V (Figure 2a), which indicates a degree of communication between the reduced tpyPY2Me ligand and iron that is not possible in the isostructural zinc complex. In contrast, ligand-based reduction in [Fe(PY5Me2)(CH3CN)]^{2+} and [Zn(PY5Me2)(CH3CN)]^{2+} is irreversible and occurs at potentials negative of −2.0 V, highlighting the crucial role of the terpyridyl fragment in achieving electrochemically reversible reduction events at relatively positive potentials.

Under an atmosphere of CO2, CV data collected for [Fe1]^{2+} exhibit only a subtle increase in current relative to scans under Ar; however, when excess phenol (3.5 M) is added as a proton source, two large catalytic waves appear at ca. −1.50 and −2.0 V versus Fc/Fc+ (Figure 2c). [Fe(PY5Me2)(CH3CN)]^{2+} also displays catalytic currents after addition of CO2 and phenol, but the onset current for catalysis is >500 mV more negative than that observed for [Fe1]^{2+} (Figure 2d). Phenol titrations conducted with solutions of [Fe1]^{2+} under a CO2 atmosphere revealed a dose-
dependence on phenol concentration (Figure S7). The electrochemical response is mitigated when control phenol titrations are collected under an Ar atmosphere, suggesting that $[\text{Fe}1]^{2+}$ selectively catalyzes electrochemical $\text{CO}_2$ reduction over the hydrogen evolution reaction, even at high concentrations of organic acid (Figure S7). Indeed, preparative-scale controlled potential electrolysis (CPE) experiments were conducted in $\text{CO}_2$-saturated acetonitrile with 3.5 M phenol across a range of overpotentials ($\eta = 0.006$–0.636 V), and quantification of products by gas chromatography revealed that CO is the major product under all conditions (Figures S8–S10). The thermodynamic potential for $\text{CO}_2$ reduction to CO was determined in acetonitrile using the $pK_a$ for carbonic acid$^{23}$. $[\text{Fe}1]^{2+}$ is highly selective for $\text{CO}_2$ reduction over hydrogen evolution across the entire potential window examined, with current densities for CO production ($j_{\text{CO}}$) reaching as high as 3.6 mA/cm$^2$ (Figure 2e and Figure S11). The Faradaic efficiency for CO production ($\text{FE}_{\text{CO}}$) reaches 94% at an overpotential of 116 mV and increases to nearly 100% with the application of overpotentials between 246 and 636 mV (Figures 2e and S11). Significantly, only small amounts of $\text{H}_2$ ($\text{FE}_{\text{H}_2} = 44\%$) were detected in analogous CPE experiments with $[\text{Fe(PY5Me}_2)(\text{CH}_3\text{CN})]^{2+}$ at much larger overpotentials (Figure S12). Control experiments strongly support the homogenous molecular nature of the catalysis with $[\text{Fe}1]^{2+}$ and its high selectivity for $\text{CO}_2$ reduction. First, $\text{CO}_2$ reduction was not observed in the absence of $[\text{Fe}1]^{2+}$ (Figure S13). Furthermore, cyclic voltammetry and UV-vis data collected before and after 1-hour CPE experiments are indistinguishable, attesting to the bulk stability of $[\text{Fe}1]^{2+}$ under electrolysis conditions (Figure S14). Finally, the possible formation of an electrode-adsorbed active catalyst was excluded through a series of control experiments. Peak currents for $[\text{Fe}1]^{2+}$ and $[\text{Zn1}]^{2+}$ scale linearly with the square root of the scan rate, indicative of freely diffusing species in solution (Figures S15–16). In addition, CPE rinse tests, which probe for electrode deposition, give complete inversion of
product selectivity, with H₂ as the predominant product and negligible generation of CO (Figure S17).

Encouraged by the high selectivity of [Fe1]²⁺ for CO₂ reduction over the hydrogen evolution reaction, we sought to evaluate its kinetic performance. Cyclic voltammograms acquired for [Fe1]²⁺ under CO₂-saturated electrolyte in the presence of 3.5 M phenol (Figure S18) exhibit scan rate dependence and noncanonical peak-shaped waves, indicative of competitive, nonideal processes (e.g., substrate consumption, catalyst deactivation, etc.). As a result, \( k_{\text{obs}} \) (TOF\(_{\text{max}}\)) obtained from catalytic plateau current analysis underestimates rates of CO₂ reduction\(^2\). We instead determined kinetic parameters directly from our variable-potential CPE experiments with direct product detection (Figure 2f)\(^2\) and in parallel applied foot-of-the-wave analysis as pioneered by Savéant\(^2\) to extract kinetic information from our CV data (Figure S19). The latter analysis estimates the idealized catalytic reactivity by considering the initial portion of the catalytic wave where the effects of side phenomena are expected to be minimal. Reasonable agreement between the two methods was found, and a TOF of 100,000 s\(^{-1}\) was measured at an applied overpotential of 116 mV by direct CPE product analysis. Slight discrepancies between the two methods are attributed to the limited linear foot-of-the-wave region (Figure S19). Nevertheless, catalytic Tafel plots constructed from either the variable-potential CPE data (Figure 2f) or foot-of-the-wave analysis (Figure S19) identify [Fe1]²⁺ as one of the most active molecular electrocatalysts reported to date, achieving TOFs >500,000 s\(^{-1}\) at overpotentials below 500 mV (Figure S20). Notably, this rapid, homogeneous CO₂ reduction catalysis is achieved without any modifications to the secondary coordination sphere to introduce hydrogen bond/proton-relay functionalities to break electronic scaling relationships\(^23,26–31\). Finally, to assess the catalytic stability of [Fe1]²⁺, four consecutive five-hour CPE experiments were conducted at −1.98 V (versus Fc/Fc⁺), the highest
potential investigated, without addition of fresh $[\text{Fe1}]^{2+}$ between each experiment (Figure S21). $[\text{Fe1}]^{2+}$ is remarkably stable with over 200 C of charge passed over the 20-hour window without any loss in selectivity for CO (average $\text{FE}_{\text{CO}} = 87\%$).

We reasoned that the observed electrocatalytic efficiency for $[\text{Fe1}]^{2+}$ for CO$_2$ reduction stems from a synergistic effect between the iron center and the redox-active tpyPY2Me ligand. To establish a molecular-level understanding of such cooperativity, we sought to isolate and characterize the catalytically relevant, two-electron reduced complex $[\text{Fe(tpyPY2Me)}]^0 ([\text{Fe1}]^0)$.

Chemical reduction of $[\text{Fe1}]^{2+}$ with 2.2 equivalents of decamethylcobaltocene (Figure 3a) produced $[\text{Fe1}]^0$ as a dark purple solid. Interestingly, room temperature $^1\text{H}$ NMR analysis of $[\text{Fe1}]^0$ in CD$_3$CN revealed that the compound is diamagnetic with the aryl and aliphatic signals shifted...
upfield (Figure 3b). A nearly identical NMR spectrum was obtained for the product \([\text{Zn1}]^0\) generated from the chemical reduction of \([\text{Zn1}]^{2+}\) with potassium graphite (see Figure S22 and details in the supplementary materials). These data are consistent with our assignment of the reduction features for both \([\text{Fe1}]^{2+}\) and \([\text{Zn}]^{2+}\) as ligand-centered events. Given the unique diamagnetic ground state of \([\text{Fe1}]^0\), two electronic structures are possible. The first possibility is a low-spin iron(II) center \((S_{\text{Fe}} = 0)\) coordinated to a reduced closed-shell singlet ligand \((S_{\text{tpy}} = 0)\) generated by both electrons occupying the same \(\pi^*\) orbital. The other possibility is an intermediate-spin iron(II) center \((S_{\text{Fe}} = 1)\) antiferromagnetically coupled to a reduced ligand with an open-shell triplet configuration \((S_{\text{tpy}} = 1)\) through half occupation of both low-lying \(\pi^*\) terpyridine orbitals, leading to an overall open-shell singlet electronic structure. The large anodic shift (> 600 mV) in the ligand-based reductions of \([\text{Fe1}]^{2+}\) relative to \([\text{Zn1}]^{2+}\) strongly supports the latter electronic configuration. Mössbauer data for \([\text{Fe1}]^{2+}\) and \([\text{Fe1}]^0\) further revealed that reduction causes a significant decrease in the isomer shift and quadrupole splitting (from \(\delta = 0.3562(4)\) mm/s and \(|\Delta E_0| = 1.0203(8)\) mm/s to \(\delta = 0.2608(2)\) mm/s and \(|\Delta E_0| = 0.7637(4)\) mm/s, respectively), consistent with the formation of intermediate-spin iron(II) (Figure 3c).\(^{32}\) Mössbauer parameters predicted from DFT also support this assignment (Table S6).\(^{21,22}\) The antiferromagnetic coupling between the reduced biradical ligand (tpyPY2Me\(^2^-\)) and intermediate-spin Fe\(^{2+}\) is expected to be quite strong, given that identical Mössbauer spectra are obtained for \([\text{Fe1}]^0\) at 295 and 5 K (after accounting for second-order Doppler effects) (Figure S23) and a diamagnetic \(^1\)H NMR spectrum is observed at room temperature. Indeed, DFT calculations predict a large coupling constant of \(-1023\) cm\(^{-1}\), giving rise to a singlet-quintet gap of 4400 cm\(^{-1}\) (see the supplementary materials).

The solid-state structure of \([\text{Fe1}]^0\) also supports the presence of an open-shell singlet electronic structure promoted through strong metal–ligand exchange coupling. Single crystals of \([\text{Fe1}]^0\) were
obtained by slow evaporation of a saturated acetonitrile solution, and X-ray diffraction analysis revealed a distorted square pyramidal geometry (Figure 3d) resulting from the loss of the axial acetonitrile ligand in \( \text{[FeI]}^{2+} \). The terpyridine bond metrics enable us to directly distinguish between the two possible electronic configurations of \( \text{[FeI]}^0 \), as a result of the unique bonding interactions of the terpyridine LUMOs. The experimental bond lengths agree most closely with those from the optimized structure of the antiferromagnetically coupled open-shell singlet (Figure 3d; Table S9). Furthermore, a comparison of the terpyridine bond lengths to those reported for reduced, homoleptic bis(terpyridine) complexes \( ([\text{M(tpy)}_2]^{n+}; \text{M} = \text{Cr, Fe})^\text{34,35} \) strongly suggests that both \( \pi^* \) orbitals in \( \text{[FeI]}^0 \) are half-occupied (Table S10). Taken together, the electrochemical, NMR, Mössbauer, and X-ray diffraction data establish that the catalytically relevant \( \text{[FeI]}^0 \) species possesses a unique, open-shell singlet ground state composed of an intermediate-spin iron(II) center \( (S_{\text{Fe}} = 1) \) that is antiferromagnetically coupled to a doubly-reduced tpyPY2Me \( (S_{\text{tpy}} = 1) \). This strong exchange coupling shifts the tpyPY2Me-based reductions of \( \text{[FeI]}^{2+} \) to more positive potentials and ultimately promotes selective and efficient \( \text{CO}_2 \) reduction catalysis.

We carried out DFT calculations using the \( \omega \text{B97X-D} \) functional to generate a more accurate molecular orbital picture of our \( \text{[FeI]}^{2+} \) system. Optimized structures and predicted redox potentials for \( \text{[FeI]}^{2+} \) and \( \text{[ZnI]}^{2+} \) are in good agreement with experimental data (Tables S11–12). Interestingly, although the DFT data indicate the metal center in \( \text{[FeI]}^{2+} \) is best described as low-spin iron(II), the calculated singlet-quintet gap is small, explaining the observed thermal population of a higher spin-state at room temperature (Tables S13–14). The first reduction of \( \text{[FeI]}^{2+} \) to \( \text{[FeI]}^+ \) is predicted to generate a ground state doublet composed of a low-spin iron(II) and a tpyPY2Me-based radical with a predicted redox potential of \(-1.46 \text{ V} \) (Figure 4a–c). The spin density plot of this \( \text{[FeI]}^+ \) species indicates ligand-based reduction with almost no excess spin on
the iron center (Figure 4b). The second reduction populates a second low lying $\pi^*$ orbital of tpyPY2Me and induces loss of the axial acetonitrile molecule, providing access to an intermediate-spin iron(II) state (Figure 4a,d,e). The $S_{Fe} = 1$ iron center is strongly antiferromagnetically coupled to the two electrons in the tpyPY2Me $\pi^*$ orbitals (Figure 4e). Localized orbital bonding analysis confirmed that iron remains in the 2+ oxidation state during the second reduction process, which has a predicted reduction potential of −1.51 V. This broken symmetry solution is significantly lower in energy (12 kcal/mol) than the other feasible singlet consisting of low-spin iron(II) ($S_{Fe} = 0$) and doubly reduced ligand ($S_{tpy} = 0$) (Tables S13–14). To further probe the entanglement of the
two ligand-based orbitals and two metal d-orbitals, we employed multiconfigurational wave-function methods (see the supplementary information), which confirmed the antiferromagnetically coupled ground state. In particular, the axial distortion away from octahedral symmetry allows the $d_{z^2}$ and $d_{xy}$ metal orbitals to interact with the two tpyPY2Me $\pi^*$ orbitals in a weak $\pi$-type interaction (Figure S24). Interestingly, the calculated potential difference for the second reduction event in $[\text{Zn1}]^{2+}$ and $[\text{Fe1}]^{2+}$ is approximately 0.5 V. This shift agrees with the energy difference between the two $[\text{Fe1}]^0$ singlet states, further indicating that significant stabilization is achieved through this unique open-shell singlet conformation.

![Figure 5. Aqueous CO\textsubscript{2} reduction performance. (A) Faradaic efficiencies for CO (blue) and H\textsubscript{2} (red) formation at varying applied potentials for 60 min (averages of three experiments). The average FE\textsubscript{CO} is indicated by the black dashed line. (B) Catalytic Tafel plot for $[\text{Fe1}]^{2+}$ obtained from CPE experiments in 0.10 M NaHCO\textsubscript{3}. The dashed blue line shows data extrapolation with an approximate TOF\textsubscript{max} of 50,000 s\textsuperscript{-1}.](image)

Given the low overpotentials required for $[\text{Fe1}]^{2+}$ to catalyze homogeneous electrochemical CO\textsubscript{2} reduction in organic solvent, we sought to evaluate its activity in water (Figure 5). For this
purpose, we prepared the more water-soluble nitrate analog of \([\text{Fe1}]^{2+}\) (see the supplementary materials), and X-ray diffraction analysis revealed that anion exchange does not alter the catalyst structure (Figure S25; Tables S15–16). As a cheap, abundant, and benign solvent and proton source, water is an attractive medium for CO\(_2\) reduction. However, molecular CO\(_2\) reduction catalysts typically require precious metals or mercury working electrodes to operate effectively in aqueous electrolytes, because their onset potentials are too negative to be compatible with more desirable carbon-based electrodes that preferentially catalyze competing water reduction\(^{36-40}\).

Cyclic voltammograms collected for \([\text{Fe1}]^{2+}\) in 0.10 M NaHCO\(_3\) with a carbon paste working electrode show the formation of a catalytic wave when saturated with CO\(_2\) (Figure S26). Notably, variable potential CPE experiments revealed that \([\text{Fe1}]^{2+}\) maintains its activity and selectivity for CO\(_2\) reduction to CO in water (average \(\text{FE}_{\text{CO}} = 92\%\) and \(\text{FE}_{\text{H}_2} < 1\%\)) with maximum turnover frequencies reaching \(>50,000\) s\(^{-1}\) at overpotentials of 500 mV (Figure 5). Importantly, control rinse tests passed negligible current during one-hour CPE experiments, and the selectivity was inverted such that \(\text{H}_2\) is the major product (Figures S27–32). Electrochemical stripping behavior is also observed, which may contribute to slower catalysis in water relative to organic solvent (Figure S33). Nevertheless, these data show that \([\text{Fe1}]^{2+}\) is an effective molecular catalyst for homogeneous CO\(_2\) reduction in water. Crucially, given its molecular nature, the activity, stability, and overpotential of this catalyst can in principle be tuned through ligand modifications accessible by synthetic chemistry. Further, the study of this catalyst and related systems has the potential to shed important insights on the behavior of materials catalysts bearing atomically dispersed iron sites\(^{41}\).

CONCLUDING REMARKS
The ability to create molecular compounds with strong exchange coupling between synergistic, redox non-innocent ligands and metal centers has broad implications for catalyst design, including enabling control over the density and arrangement of functional active sites. Indeed, redox balancing through electronic delocalization is pivotal for biological and materials catalysts to achieve complex chemical transformations at or near thermodynamic potentials. However, discrete molecular compounds offer a level of tunability via coordination chemistry that remains unparalleled in protein design and nanostructure engineering. Thus, inorganic molecules are uniquely positioned to open up access to a range of new properties and offer a desirable alternative strategy for lowering energy barriers for redox catalysis.

ASSOCIATED CONTENT

Supporting Information. Experimental and computational details, compound characterization, supplemental electrochemical and spectroscopic data, and DFT geometry optimized atomic xyz coordinates is available free of charge.

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**REFERENCES**


TOC Graphic:

- Metal-Ligand Exchange Coupling
- $\eta < 100$ mV
- $\text{IE}_{\text{red}} > 90\%$
- TOF $> 100,000$ s$^{-1}$
- Robust
- Water Compatible