

Mechanistic Dichotomy in Proton-Coupled Electron-Transfer Reactions of Phenols with a Copper Superoxide Complex

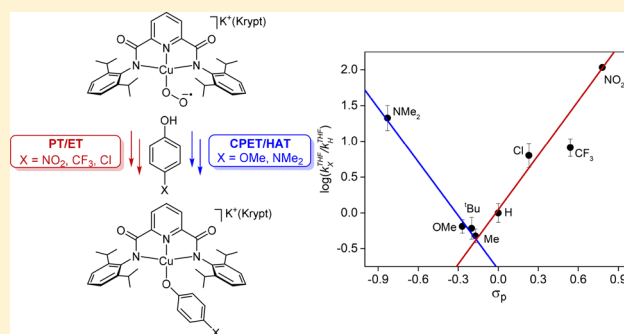
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Supporting Information

ABSTRACT: The kinetics and mechanism(s) of the reactions of $[K(\text{Krypt})][\text{LCuO}_2]$ (Krypt = 4,7,13,16,21,24-hexaoxa-1,10-diazabicyclo[8.8.8]hexacosane, L = a bis(arylcarboxamido)pyridine ligand) with 2,2,6,6-tetramethylpiperidine-*N*-hydroxide (TEMPOH) and the para-substituted phenols $^X\text{ArOH}$ (X = para substituent NO₂, CF₃, Cl, H, Me, ^tBu, OMe, or NMe₂) at low temperatures were studied. The reaction with TEMPOH occurs rapidly ($k = 35.4 \pm 0.3 \text{ M}^{-1} \text{ s}^{-1}$) by second-order kinetics to yield TEMPO[•] and $[\text{LCuOOH}]^-$ on the basis of electron paramagnetic resonance spectroscopy, the production of H₂O₂ upon treatment with protic acid, and independent preparation from reaction of $[\text{NBu}_4][\text{LCuOH}]$ with H₂O₂ ($K_{\text{eq}} = 0.022 \pm 0.007$ for the reverse reaction). The reactions with $^X\text{ArOH}$ also follow second-order kinetics, and analysis of the variation of the k values as a function of phenol properties (Hammett σ parameter, O–H bond dissociation free energy, $\text{p}K_{\text{a}}$, $E_{1/2}$) revealed a change in mechanism across the series, from proton transfer/electron transfer for X = NO₂, CF₃, Cl to concerted-proton/electron transfer (or hydrogen-atom transfer) for X = OMe, NMe₂ (data for X = H, Me, ^tBu are intermediate between the extremes). Thermodynamic analysis and comparisons to previous results for LCuOH, a different copper–oxygen intermediate with the same supporting ligand, and literature for other $[\text{CuO}_2]^+$ complexes reveal significant differences in proton-coupled electron-transfer mechanisms that have implications for understanding oxidation catalysis by copper-containing enzymes and abiological catalysts.



INTRODUCTION

A key to understanding enzymes¹ and synthetic catalysts² that use single copper sites to perform challenging oxidative transformations is discerning the nature of the involved copper–oxygen intermediates. Postulates of such intermediates are shown in Figure 1^{3–5} in which supporting ligands are omitted and each species is shown in a form emphasizing either proposed formal oxidation states (blue) or overall charge for the core (black, in brackets). Synthetic complexes that incorporate many of these cores have been characterized and their reactivity studied, shedding important light on their viability in catalysis.^{4–10} Yet, compounds incorporating many of the indicated cores have not been unequivocally isolated to date, and much is still unknown about the fundamental chemistry of those that have been prepared, particularly with respect to their interconversions (potentially involving reactions with organic substrates) by the indicated proton and/or electron transfers (PT, vertical arrows; ET, horizontal arrows; PCET, proton-coupled electron transfer, red diagonal arrows). An overarching research goal, and one which underlies the work reported herein, is to discern the relative propensities for these reaction paths (kinetics and thermodynamics) and how these are influenced by

the conditions (supporting ligand environment, secondary coordination sphere, temperature, solvent, etc.). Such research ultimately aims to develop rules for oxidase and/or oxygenase catalyst design.

Deep insights into the nature of proton-coupled electron-transfer mechanisms^{11,12} such as those depicted in Figure 1 have been attained through examinations of the reactivity of metal–oxygen species with phenols (important substrates in biological proton-coupled electron-transfer processes) that have tunable properties because of the presence of variable para substituents.^{13–16} By examination of the course and kinetics of the reactions with a series of para-substituted phenols, trends can be ascertained that, in concert with thermodynamic analysis of relevant square schemes, enable stepwise (PT/ET or ET/PT) and concerted proton-coupled electron transfer (CPET) mechanisms to be distinguished. In a recent application of this approach, we examined the kinetics of the reactions of LCuOH comprising the $[\text{CuOH}]^{2+}$ core supported by strongly electron-donating bis(arylcarboxamido)pyridine ligands (L^{2-} ; Figure 2)

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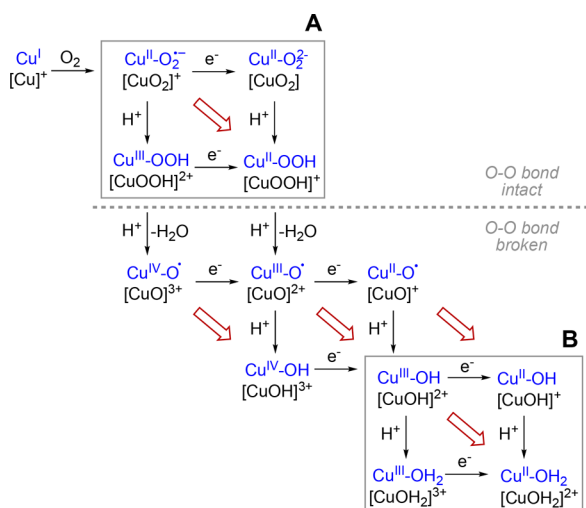


Figure 1. Intermediates proposed for O_2 activation at a single copper site, emphasizing proton transfers (PT, vertical arrows), electron transfers (ET, horizontal arrows), and proton-coupled electron transfers (PCET, diagonal red arrows) that connect them. Arrows are drawn for simplicity as unidirectional, although the processes are equilibria characterized by parameters $\text{p}K_a$ (PT), $E_{1/2}$ (ET), or BDFE (PCET), respectively, where BDFE = bond dissociation free energy. Gray boxes indicate the square schemes A and B emphasized in this work, and each intermediate is written in two forms: indicating possible formal oxidation states (blue) or overall charge (black). Figure inspired by and adapted from one published in ref 3a.

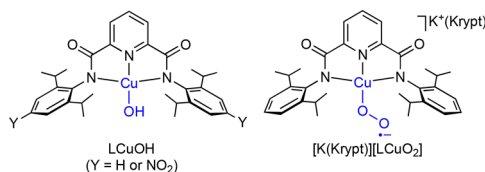


Figure 2. Copper complexes comprising $[\text{CuOH}]^{2+}$ and $[\text{CuO}_2]^+$ cores supported by bis(arylcarboxamido)pyridine ligands. Krypt = Kryptofix222.

with para-substituted phenols $^X\text{ArOH}$ (X = para-substituent NO_2 , CF_3 , Cl , H , Me , OMe , NMe_2) using low-temperature stopped-flow methods and reference to the square scheme B (Figure 1).¹⁶ A key finding was that a CPET (or hydrogen-atom transfer, HAT) pathway was followed for all X , but with anomalies in the cases of $X = \text{NO}_2$ and CF_3 that were interpreted to indicate contributions from a PT/ET route for acidic phenol substrates.

At the other end of the cascade of possible copper–oxygen intermediates (Figure 1) is the initially formed 1:1 Cu/O_2 adduct with the $[\text{CuO}_2]^+$ core (typically formally $\text{Cu}^{\text{II}}-\text{O}_2^{\bullet-}$) that must be involved in any system that uses a single copper ion to bind and activate O_2 . A number of synthetic complexes with this core have been characterized,^{4–8} including one ($[\text{K}(\text{Krypt})][\text{LCuO}_2]$)¹⁷ with the same supporting ligand as in LCuOH (Figure 2). We envisioned that unique insights would be attained by comparing the reactivity of these two different monocopper–oxygen cores supported by the same ligand. Moreover, such comparisons would address an important fundamental question with parallels in many other metal–oxygen systems:^{18,19} How do the reactivities and mechanisms traversed differ for species that contain an intact O–O bond (like $[\text{CuO}_2]^+$) versus those that do not (such as $[\text{CuOH}]^{2+}$)?

Herein, we describe the results of studies aimed at answering this general question in a specific case by examining the kinetics of the reactions of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ with the same series of phenols $^X\text{ArOH}$ used in previous studies¹⁶ of LCuOH . We also studied the reactivity of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ with a substrate having a particularly weak O–H bond, 2,2,6,6-tetramethylpiperidine-*N*-hydroxide (TEMPOH). Analysis via reference to square scheme A and comparisons to the previous results for LCuOH and literature for other $[\text{CuO}_2]^+$ complexes reveal significant differences in proton-coupled electron-transfer mechanisms. These differences have implications for understanding the reactivity of copper–oxygen species proposed in enzymatic and other catalytic systems.

RESULTS

Detailed studies of the reactivity of $[\text{LCuO}_2]^-$ were enabled by the discovery (reported in detail elsewhere^{17b}) that stable solutions ($t_{1/2} \sim 12$ h at -60 °C) in predominantly tetrahydrofuran (THF) with minimal CH_3CN (THF/ CH_3CN ratios 10–40:1) could be obtained by using 4,7,13,16,21,24-hexaoxa-1,10-diazabicyclo[8.8.8]-hexacosane (Kryptofix222, abbreviated further here as “Krypt”) to solubilize the reactant KO_2 . The resulting solutions of the complex $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ contain a characteristic UV–vis feature at 628 nm ($\epsilon \approx 1400 \text{ M}^{-1} \text{ cm}^{-1}$) that was monitored in reactions with external substrates.

Reactivity of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ with TEMPOH. Addition of a solution of TEMPOH in THF (40–200 mM) to a freshly prepared solution of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ (0.65 mM) in THF/ CH_3CN (v/v 30:1) at -80 °C resulted in the rapid decay of the UV–vis feature at 628 nm to yield a final spectrum with absorptions at 485 nm ($\epsilon = 386 \text{ M}^{-1} \text{ cm}^{-1}$) and 655 nm ($238 \text{ M}^{-1} \text{ cm}^{-1}$), with a shoulder at 513 nm (Figure 3). The final

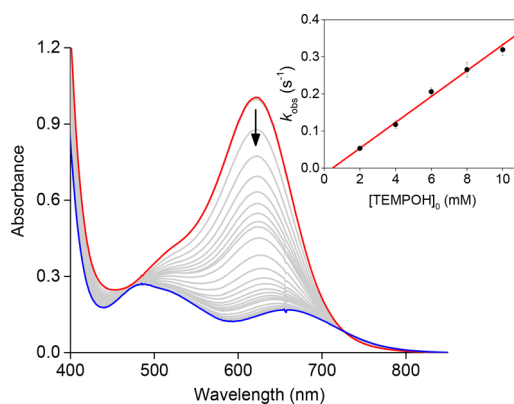


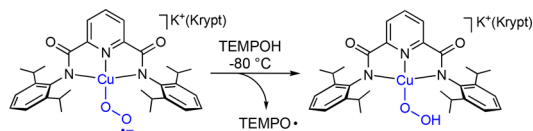
Figure 3. UV–vis spectra as a function of time observed upon reaction of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ (red) with TEMPOH. Conditions: $[\text{Cu}] = 0.7$ mM, $[\text{TEMPOH}]_0 = 1.4$ mM, THF:MeCN 10:1, -80 °C. Inset shows linear fit in plot of k_{obs} versus $[\text{TEMPOH}]_0$ (slope = 34.9 ± 1.5 , intercept = -0.016 ± 0.005 , $R^2 = 0.993$).

product spectroscopic features remained unchanged for long periods up to -20 °C, but decomposed at room temperature. Single-wavelength monitoring at 628 nm showed the reactions with excess TEMPOH to obey first-order kinetics on the basis of fits to simple exponential decays (Figure S3). A plot of the observed pseudo-first-order rate constants (k_{obs}) versus $[\text{TEMPOH}]_0$ was fit to a straight line with an effectively zero intercept within error, consistent with an overall second-order reaction with slope = k of $34.9 \pm 1.5 \text{ M}^{-1} \text{ s}^{-1}$ (inset to Figure 3).

This result was corroborated by an alternative analysis using multiwavelength global fitting of the UV–vis spectra that gave $k = 35.4 \pm 0.3 \text{ M}^{-1} \text{ s}^{-1}$ (Figure S4).

We propose on the basis of evidence described below that the products of the reactions with TEMPOH are the TEMPO• radical and the new complex $[\text{K}(\text{Krypt})][\text{LCuOOH}]$ (Scheme 1). The X-band electron paramagnetic resonance (EPR)

Scheme 1. Reaction of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ with TEMPOH



spectrum of the product mixture from the reaction of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ and 10 equiv of TEMPOH displayed a characteristic signal for a LCu^{II} species overlaid with that for free TEMPO• radical ($g = 2.005$; Figure S5). Spin integration compared to either a Cu(II) or TEMPO• standard indicated two $S = 1/2$ systems in a 1:1 ratio; no excess TEMPO• was formed, indicating that the Cu(II) reaction product does not react further with TEMPOH. The copper hyperfine splitting constant ($A_z = 578 \text{ MHz}$) and g_z value (2.185) for the Cu(II) signal were found to be comparable to those reported previously for other LCu^{II} complexes (Table S3).

The nature of the Cu(II) product of the reactions of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ with TEMPOH was further probed by adding triflic acid (HOTf) to the product mixture and assaying it for H_2O_2 . In the experiment, addition of HOTf in Et_2O (20 mM) to a THF solution of $[\text{K}(\text{Krypt})][\text{LCuOOH}]$ (made from addition of 2 equiv of TEMPOH to $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ at -60°C) resulted in the immediate appearance of bands at 406 ($2450 \text{ M}^{-1} \text{ cm}^{-1}$) and 642 nm ($370 \text{ M}^{-1} \text{ cm}^{-1}$) (Figure S6). Titration of the product solution into saturated NaI in MeCN revealed 73% yield of H_2O_2 relative to copper,²⁰ consistent with protonolysis of the proposed hydroperoxide species $[\text{K}(\text{Krypt})][\text{LCuOOH}]$. Further corroborating the formulation of the latter complex, addition of a solution of H_2O_2 in Et_2O (predried with MgSO_4 to remove most H_2O) to a solution of $[\text{NBu}_4][\text{LCuOH}]$ in THF yielded a UV–vis spectrum closely analogous to that of the solution resulting from reaction of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ with TEMPOH (Figure S7). The reaction with H_2O_2 was reversed upon addition of H_2O , consistent with the equilibrium eq 1, for which K_{eq} was estimated to be 0.022 ± 0.007 (Figure S8, Supporting Information). Taken together, the data support formation of $[\text{LCuOOH}]^-$ via both routes. Unfortunately, attempts to identify the $[\text{CuOOH}]^+$ core more directly by resonance Raman spectroscopy on solution state samples prepared by addition of H_2O_2 in ether (1 equiv) to $[\text{NBu}_4][\text{LCuOH}]$ (10 mM in THF) at -20°C (90° backscattering geometry) were unsuccessful; discernible peaks derived from the sample were not observed using any of the available excitation wavelengths (406, 457, 515, 561, and 660 nm).



Finally, in an attempt to determine the $E_{1/2}$ for the $[\text{CuOOH}]^{2+/+}$ couple, we measured the cyclic voltammogram of the solution obtained upon reaction of $[\text{NBu}_4][\text{LCuOH}]$ (2 mM) with anhydrous H_2O_2 in Et_2O (1 equiv) in 1,2-difluorobenzene at 0°C (0.2 M NBu_4PF_6 , Figure S10). An

irreversible oxidation wave was observed with $E_{\text{p,a}} = -0.177 \text{ V}$ versus Fc^+/Fc at a scan rate of 100 mV/s; no return wave was observed even at high scan rates (2000 mV/s). The data suggest that oxidation to the $[\text{CuOOH}]^{2+}$ core occurs, but that this core is unstable at 0°C . The irreversible voltammograms were batch-simulated using the Digisim 3.03b software package from BASi (Figure S11). Data collected at scan rates of 50, 100, 200, 750, 1500, and 2000 mV/s were simultaneously modeled using a two-step mechanism: initial heterogeneous electron transfer (eq S6) followed by a fast, homogeneous chemical event which quenches the oxidized copper species (eq S7). From this model, a theoretical redox potential (that is necessarily approximate) for the $[\text{CuOOH}]^{2+/+}$ couple was derived with a value of $E_{1/2} = -0.215 \text{ V}$ (versus Fc^+/Fc) (Figure S11).

Reactivity of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ with para-Substituted Phenols. To solutions of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ in THF/ CH_3CN (18:1) were added solutions of the substituted phenols $^{\text{X}}\text{ArOH}$ (20 equiv, X = para-substituent NO_2 , CF_3 , Cl, H, Me, ^tBu , OMe, or NMe_2) in THF at -60°C (Figure 4). Monitoring

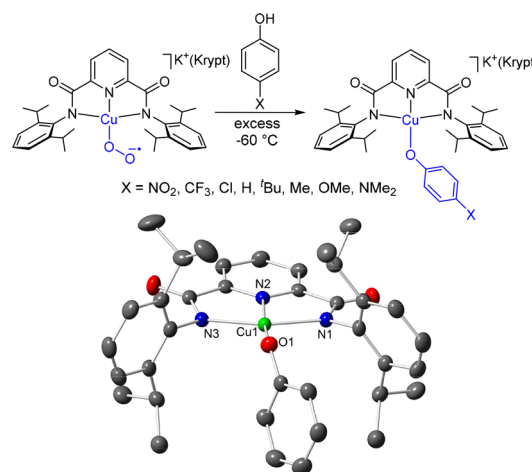


Figure 4. Reactions of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ with phenols $^{\text{X}}\text{ArOH}$, with a representation of the X-ray crystal structure for the case X = H, showing all nonhydrogen atoms as 50% thermal ellipsoids (hydrogen atoms and NEt_4 counterion omitted). Selected interatomic distances (Å) and angles (deg): Cu1–O1: 1.8767(13), Cu1–N2: 1.9296(14), Cu1–N1: 2.0064(15), Cu1–N3: 2.0220(15), O1–Cu1–N2: 172.98(6), O1–Cu1–N1: 96.94(6), O1–Cu1–N3: 102.46(6) N1–Cu1–N3: 160.44(6).

the reactions by UV–vis spectroscopy showed that the features due to $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ decayed, ultimately affording spectra with weak bands $>400 \text{ nm}$ associated with Cu(II) products (Figure S12). Those products were identified as salts of the phenolate complexes $[\text{LCuOAr}^{\text{X}}]^-$ on the basis of comparison to data obtained for independently synthesized and characterized complexes for several cases (X = NO_2 , H, OMe, NMe_2 ; the formulations for X = CF_3 , ^tBu , Me, Cl were made by analogy). Thus, the complexes $[\text{NR}_4][\text{LCuOAr}^{\text{X}}]$ were prepared either by reaction of (a) $\text{LCu}(\text{CH}_3\text{CN})$ with $[\text{NR}_4][\text{OAr}^{\text{X}}]$ (R = Bu, X = NO_2 ,¹⁶ R = Et, X = H or OMe) or (b) $[\text{NBu}_4][\text{LCuOH}]$ with $^{\text{X}}\text{ArOH}$ (X = NMe_2). The products were isolated and characterized by UV–vis and EPR spectroscopy (Figures S14 and S1, respectively), CHN analysis or high-resolution mass spectrometry (X = NMe_2), and X-ray crystallography (illustrative structure for X = H in Figure 4; others shown in Figure S2). Importantly, the close similarity of the UV–vis spectroscopic features for the isolated complexes

and the final solutions resulting from the reactions of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ with ${}^X\text{ArOH}$ indicated that the final products of the latter were the respective complexes $[\text{LCuOAr}^X]^-$.

The UV–vis spectra as a function of time for the reactions of $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ with ${}^X\text{ArOH}$ were analyzed by multiwavelength global analysis (Figures S12 and S13). In all cases, the initial exponential decays of $[\text{LCuO}_2]^-$ were fit to a simple second-order model, yielding second-order rate constants (k) listed in Table 1. For the cases $X = \text{NMe}_2$ and OMe, data

Table 1. Second-Order Rate Constants (k) for the Reactions between $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ and ${}^X\text{ArOH}$ at $-60\text{ }^\circ\text{C}$

X	k ($\text{M}^{-1} \text{s}^{-1}$)	$\log(k)$	$\log(k^{\text{DMSO}})$
NMe ₂	30(13)	1.4(2)	0.1(2)
OMe	0.7(1)	-0.16(9)	-1.45(9)
^t Bu	0.8(3)	-0.2(2)	-1.4(2)
Me	0.5(1)	-0.3(1)	-1.6(1)
H	1.2(3)	0.0(1)	-1.3(1)
Cl	8(3)	0.8(2)	-0.7(2)
CF ₃	10(3)	0.9(1)	-0.7(1)
NO ₂	114(6)	2.06(2)	0.29(2)

acquired over longer time periods indicated a more elaborate rate law (see below), but the k values in Table 1 were found to be reasonable for comparative purposes. To evaluate the data using thermodynamic parameters reported in dimethyl sulfoxide (DMSO),^{21,22} these rate constants were converted to values in that solvent (k^{DMSO}) according to a previously described protocol¹⁶ specific for HAT/CPET reactions.²³ Plots of $\log(k^{\text{DMSO}})$ versus phenol σ values (Hammett), pK_a 's, and $E_{1/2}$ values (for the ${}^X\text{ArOH}^{0/+}$ couples) are shown in Figure 5. In each plot, the data point for $X = \text{NMe}_2$ is clearly an outlier and a distinct break in continuity is apparent by the indicated linear fits, which occurs approximately at $X = \text{Me}$, with maximum rates at the extremes $X = \text{NO}_2$ and NMe_2 . Such a break in these plots points to a change in mechanism across the series. We hypothesize that two mechanisms are operative, one being rate-determining PT (line through point for $X = \text{NO}_2$, rate increases as pK_a decreases) and the other being rate-determining CPET/HAT (line through point for $X = \text{NMe}_2$, which has lowest $E_{1/2}$ and highest pK_a).

Further evaluation of the kinetic UV–vis spectroscopic data over longer reaction times revealed additional evidence in support of the above mechanistic hypotheses. For the reactions with ${}^X\text{ArOH}$ ($X = \text{NMe}_2$, OMe, and to a lesser extent ^tBu, and Me), distinct losses of isosbestic points in the UV–vis traces over long time periods are apparent (Figure 6, Figure S16) and an intermediate spectrum (purple, “Int.”) slowly converts to that of the final $[\text{LCu}(\text{OAr}^X)]^-$ product (blue). The two different time courses are illustrated for select wavelengths in the inset to Figure 6. Additional kinetics runs with ${}^X\text{ArOH}$ ($X = \text{OMe}$) were performed using variable concentrations of substrate (5–25 equiv based on Cu), followed over 1 h at $-60\text{ }^\circ\text{C}$ by UV–vis spectroscopy. In all cases, multiwavelength global analysis of the full kinetic traces were fit to a two-phase reaction mechanism (eq 2), with average rate constants of $k_1 = 0.44(9)\text{ M}^{-1} \text{s}^{-1}$ and $k_2 = 0.12(2)\text{ M}^{-1} \text{s}^{-1}$ (Figure S17).²⁴ Multiwavelength global analysis was also used to fit the reaction of $[\text{LCuO}_2]^-$ and NMe_2ArOH (20 equiv) to a two-phase reaction mechanism, with $k_1 = 13\text{ M}^{-1} \text{s}^{-1}$ and $k_2 = 1.1\text{ M}^{-1} \text{s}^{-1}$ (Figures S16 and S17, Scheme S1). Furthermore, included in the model was the

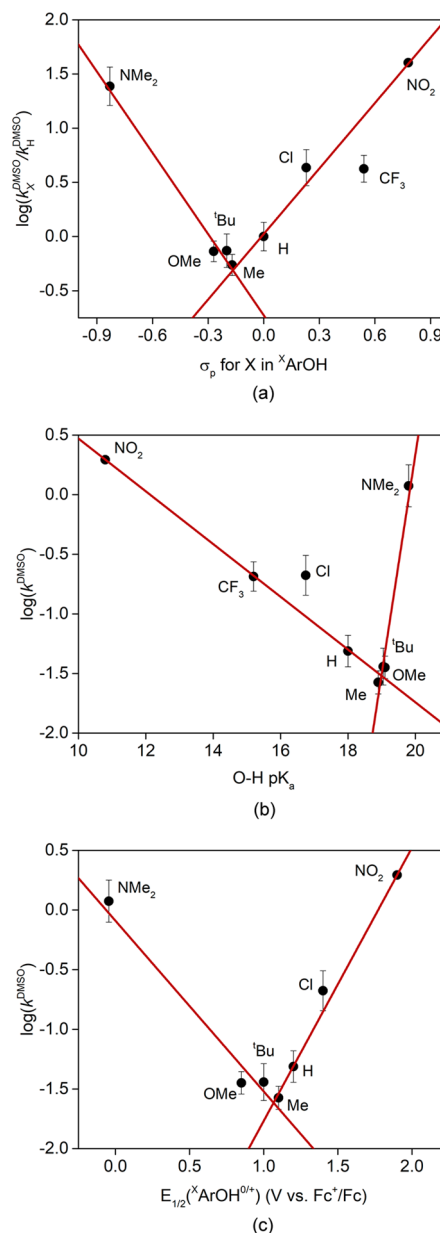


Figure 5. (a) Hammett plot of $\log(k_x^{\text{DMSO}}/k_{\text{H}}^{\text{DMSO}})$ versus ${}^X\text{ArOH}$ σ_p , with X labeled. Separate linear fits are shown to data for $X = \text{NMe}_2$, OMe, ^tBu, and Me (slope $\rho = -2.5$, $R^2 = 0.97$) and for $X = \text{Me}$, H, Cl, CF₃, and NO₂ (slope $\rho = +2.0$, $R^2 = 0.96$). (b) Plot of $\log(k^{\text{DMSO}})$ versus ${}^X\text{ArOH}$ pK_a . Separate linear fits are shown to data for $X = \text{NMe}_2$, OMe, ^tBu, and Me (slope $= +1.84$, $R^2 = 0.924$) and for $X = \text{Me}$, H, Cl, CF₃, and NO₂ (slope $= -0.22$, $R^2 = 0.998$). (c) Plot of $\log(k^{\text{DMSO}})$ versus ${}^X\text{ArOH}^{0/+}$ $E_{1/2}$. Separate linear fits are shown to data for $X = \text{NMe}_2$, OMe, ^tBu, and Me (slope $= -1.43$, $R^2 = 0.92$) and for $X = \text{Me}$, H, Cl, and NO₂ (slope $= +2.29$, $R^2 = 0.996$).

appearance of the transient phenoxyl radical, $\text{NMe}_2\text{ArO}^\bullet$, with a fitted spectrum closely matching that reported previously ($\lambda_{\text{max}} = 495\text{ nm}$).²⁵ The appearance of this radical is consistent with a CPET/HAT mechanism for $X = \text{NMe}_2$. The spectrum of the initially formed intermediate in both reactions ($X = \text{OMe}$ and NMe_2 ; “Int.” in eq 2) is indicative of LCu(II) species, and is not consistent with formally Cu(III) complexes that are characterized by intense ligand to metal charge transfer (LMCT) absorptions.^{26–28} We hypothesize that the intermediate is $[\text{LCuOOH}]^-$ derived from CPET/HAT, which is favorable for

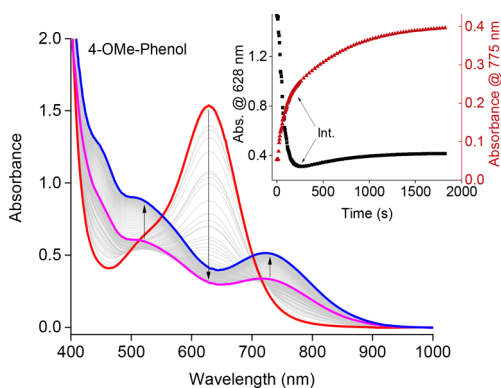


Figure 6. UV-vis spectral traces for the reaction of [K(Krypt)][LCuO₂] (red) with ^XArOH (X = OMe, 20 equiv) at -60 °C, 19:1 THF:CH₃CN, over ~30 min. Inset shows the decay trace over time following the signal at 628 (black squares) and 775 nm (red triangles). “Int.” is defined as the initially formed Cu(II) intermediate and is highlighted in the UV-vis spectrum in purple.

substrates with relatively weak O–H bonds (especially X = NMe₂, BDFE = 79 kcal/mol; OMe, BDFE = 83 kcal/mol; Figure 7).²¹ The intermediate is then consumed in a protonolysis event

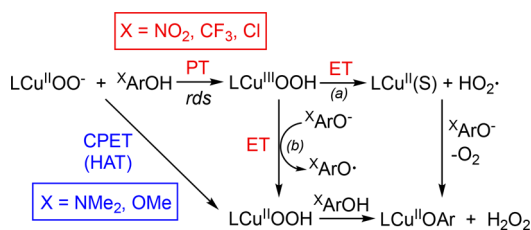
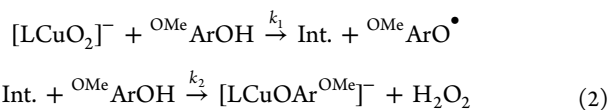


Figure 7. Proposed PT/ET (red) and CPET (blue) mechanisms for the reactions of [K(Krypt)][LCuO₂] with ^XArOH. The data are ambiguous for X = H, Me, ^tBu, suggesting that in these cases both pathways may be operative (not shown).

by excess phenol (k_2). In support of this hypothesis, reaction of the independently synthesized [NBu₄][LCuOOH] with NMe₂ArOH (20 equiv) resulted in a UV-vis spectrum matching that of [LCuOAr^{NMe2}]⁻ and a calculated second-order rate constant of $k = 0.54 \text{ M}^{-1} \text{ s}^{-1}$ (Figure S18). We surmise that because these phenols are weak acids (X = NMe₂, pK_a = 19.8; OMe, pK_a = 19.1), the protonolysis is relatively slow, thus enabling observation of the intermediate.



For substrates that are more acidic (especially X = NO₂, pK_a = 10.8; CF₃, pK_a = 15.2)^{21,22} the reactions exhibit clean isosbestic points and no intermediate is observed. We propose that the rate-limiting step in these cases is an initial PT event that yields a highly reactive and unstable (formally Cu(III)) species LCuOOH. This species could decay quickly via ET from the phenoxide anion and subsequent protonolysis, or by some other decay pathway (such as Cu–O bond homolysis)^{29,30} followed simply by phenoxide coordination (Figure 7). Because of the larger $E_{1/2}$ values for the ^XArO[•]/^XArO⁻ couple (X = NO₂ and CF₃; $E_{1/2} = 0.314$ and 0.079 V versus Fc⁺/Fc)^{21,22} compared to the approximated [CuOOH]^{2+/+} couple ($E_{1/2} = -0.215$ V versus

Fc⁺/Fc), pathway (b) is unlikely from a thermodynamic standpoint. Furthermore, reactions with [LCuO₂]⁻ and only 1 equiv of ^XArOH (X = NO₂, CF₃; Figure S20) yield the same phenoxide product in the same amount (calculated by extinction of final UV-vis spectrum) as those performed with an excess of phenol substrate, favoring pathway (a) where excess ^XArOH is not needed (Figure 7). Only ^{Cl}ArO⁻ ($E_{1/2} = -0.232$ V versus Fc⁺/Fc)²¹ could undergo a thermodynamically favorable ET to [LCuOOH] by pathway (b). Formulation of the mechanisms for X = H, Me, ^tBu are less clear because of their position in the plots (Figure 5) at the intersection of the linear fits, and we suspect that contributions from both pathways may occur in these intermediate cases. Further aspects of these mechanistic hypotheses are described below (see Discussion section).

Thermodynamic Analysis. To obtain a deeper understanding of the reactivity of [K(Krypt)][LCuO₂] with the phenols ^XArOH, we attempted to evaluate the thermodynamics of PCET via the square scheme A (Figure 1). Complicating the analysis is the instability of all the species in the scheme (except [K(Krypt)][LCuO₂]). Nonetheless, we were able to obtain a rough estimate of key pK_a and redox potential values, enabling estimation of the O–H BDE for the [CuOOH]⁺ core. As noted above, cyclic voltammograms measured on solutions of [LCuOOH]⁻ prepared by reaction of [LCuOH]⁻ with H₂O₂ contained an irreversible wave ascribed to oxidation of the [CuOOH]⁺ core with $E_{p,a} = -0.177$ V versus Fc⁺/Fc. By simulating the half-wave at varying scan rates, a value for $E_{1/2}$ for the [CuOOH]^{2+/+} couple was estimated at -0.215 V versus Fc⁺/Fc. To estimate the pK_a value of the [CuOOH]²⁺ complex, we make the crude assumption that at the mechanistic crossover point in the plots shown in Figure 5 (about at X = Me) the thermodynamic driving forces for PT and CPET are approximately the same ($\Delta G_{\text{PT}} \sim \Delta G_{\text{CPET}}$).³¹ The driving force for the CPET reaction (ΔG_{CPET}) is estimated by the difference between the BDFE of ^{Me}ArOH in DMSO and the BDE of [LCuOO–H]⁻ in THF (eq 3). The [LCuOO–H]⁻ BDE is represented by eq 4, where pK_a is the acidity of LCu^{III}OO–H, E° is approximated by $E_{1/2}$ of [LCuOOH]⁻ (-0.215 V), and a value of 66 kcal mol⁻¹ is used for the constant C_H (in THF).³² With use of the reported value of 87.1 kcal mol⁻¹ for the BDFE of ^{Me}ArOH,²¹ and the incorporation of eq 4, eq 3 may be simplified to eq 5, which relates ΔG_{CPET} to the unknown pK_a of LCu^{III}OOH. The driving force for proton transfer (ΔG_{PT}) is estimated from the difference in pK_a of ^{Me}ArOH and LCu^{III}OOH, the latter being the unknown variable (eq 6, where R is the ideal gas constant and T is the temperature). With use of the known pK_a of ^{Me}ArOH (19.1, DMSO),²¹ eq 6 simplifies to eq 7. Working under the assumption that $\Delta G_{\text{PT}} \sim \Delta G_{\text{CPET}}$, eq 5 and eq 7 may be set to be equal to one another, enabling calculation of the pK_a of LCu^{III}OOH to be ~19 (THF). Inserting this value back into eq 4 enables calculation of the BDE of [LCu^{II}OOH]⁻ as ~87 kcal mol⁻¹. We hypothesize that this approximated value represents an upper estimate; on the basis of the proposed CPET mechanism followed in the reaction of [LCuO₂]⁻ and ^{OMe}ArOH (BDFE = 83 kcal/mol),²¹ it is likely that the true BDE of [LCuOO–H]⁻ falls in the range 83–87 kcal/mol.

$$\Delta G_{\text{CPET}} = \text{BDFE}_{\text{MeArO-H}} - \text{BDE}_{\text{CuOO-H}} \quad (3)$$

$$\text{BDE}_{\text{CuOO-H}} = 1.37\text{pK}_a + 23.06E^\circ + C_H \quad (4)$$

$$\Delta G_{\text{CPET}} = 25.18 - 1.37\text{pK}_a\text{CuOO-H} \quad (5)$$

$$\Delta G_{\text{PT}} = 2.303RT(\text{p}K_{\text{aMeArO-H}} - \text{p}K_{\text{aCuOO-H}}) \quad (6)$$

$$\Delta G_{\text{PT}} = 18.43 - 0.975\text{p}K_{\text{aCuOO-H}} \quad (7)$$

DISCUSSION

Comparisons to other previously reported complexes with $[\text{CuO}_2]^+$ cores for which reactivity with TEMPOH (or derivatives) and phenols have been studied are useful for placing our findings in perspective (Figure 8).^{4,14,33–39} All are

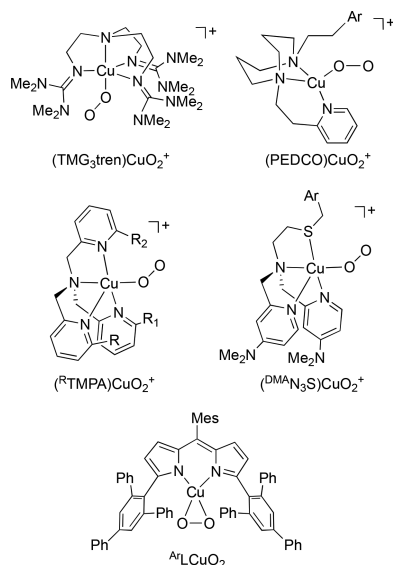


Figure 8. Complexes with $[\text{CuO}_2]^+$ cores for which reactivity studies with TEMPOH (or derivatives) and phenols have been reported. ^{DMA}TMPA: R = NMe₂, R₁ = R₂ = H; ^{DMM}TMPA: R = OMe, R₁ = R₂ = H; ^{PV}TMPA: R = NH(CO)^tBu, R₁ = R₂ = H; ^{NEO}TMPA: R = R₁ = NH(neopentyl), R₂ = H. Ar = aryl, Mes = mesityl. Note: The complex ^{Ar}LCuO₂ is proposed to equilibrate with an end-on (η^1) isomer (ref 39).

postulated to exhibit end-on η^1 binding of superoxide, except ^{Ar}LCuO₂; however, this η^2 complex was proposed to exist in equilibrium with an end-on isomer.³⁹ The complex [K(Krypt)]-[LCuO₂] differs from the complexes shown in Figure 8 with respect to overall charge and supporting ligand geometry and donor set. These structural differences are accompanied by disparities in their reactivity with TEMPOH (or derivatives) and phenols, as discussed below. We also discuss reactivity variations between [K(Krypt)]-[LCuO₂] and LCuOH, which contains a different monocopper–oxygen core supported by the same ligand.

Reaction with TEMPOH. Reaction of [K(Krypt)]-[LCuO₂] with TEMPOH (Scheme 1) yields TEMPO• and a Cu(II) product that we formulate as [LCuOOH][−] on the basis of EPR spectroscopy, the production of H₂O₂ upon treatment with protic acid, independent preparation from reaction of [NBu₄]-[LCuOH] with H₂O₂ (eq 1, $K_{\text{eq}} = 0.022 \pm 0.007$), and analogy to previously reported and more stable complexes [LCuOOR][−] (R = cumyl, ^tBu).²⁸ The rapid CPET/HAT reaction followed the second-order rate law: rate = $k[\text{complex}][\text{TEMPOH}]$, with $k = 34.9 \pm 1.5 \text{ M}^{-1} \text{ s}^{-1}$ at -80°C . Similar conversions were reported previously for $[\text{CuO}_2]^+$ complexes supported by TMG₃tren,³³ ^{NEO}TMPA,³⁸ PEDCO,¹⁴ and ^{Ar}L[−]³⁹ (Figure 8). It would thus appear that, for the substrate TEMPOH having a quite weak O–H bond (BDFE ~67.5 kcal/mol),⁴⁰ a common reaction course involving apparent HAT is followed for the

$[\text{CuO}_2]^+$ core irrespective of the overall charge and supporting ligand in the complexes. Nonetheless, in the one other report of reaction kinetics having been evaluated ($(\text{PEDCO})\text{CuO}_2^+$),¹⁴ the second-order rate constant ($k = 2.4 \pm 0.05 \text{ M}^{-1} \text{ s}^{-1}$ at -85°C) is about an order of magnitude smaller than that we observed for [K(Krypt)]-[LCuO₂]. The complex LCuOH also abstracts an H atom from TEMPOH at -80°C , but at a rate so fast that we have not been able to monitor it with much precision or accuracy. This greater PCET reactivity for LCuOH is reflected in the phenol reaction kinetics as well.

Reactions with Phenols. The reactions of [K(Krypt)]-[LCuO₂] with phenols ^XArOH (X = para-substituent NO₂, CF₃, Cl, H, Me, ^tBu, OMe, or NMe₂) yields the complexes [LCuOAr^X][−]. These products were identified by spectroscopic comparison to independently synthesized samples, several of which were characterized by X-ray crystallography (X = NO₂, H, OMe, NMe₂; Figure 4 and Figure S2); similar formulations for X = CF₃, ^tBu, Me, Cl were made by analogy. Kinetics studies of the reactions monitored by UV–vis spectroscopy showed that the decay of the features due to [K(Krypt)]-[LCuO₂] followed second-order kinetics (k and k^{DMSO} values listed in Table 1). The initial formation of an intermediate that slowly transformed to the final [LCuOAr^X][−] product was indicated for X = NMe₂ and OMe (also ^tBu and Me, although the data were less clear). We postulate that, for these cases, the intermediate is the same product seen in the TEMPOH reactions, [LCuOOH][−], and that the mechanism traversed differs from those for the phenols with other X substituents.

The plots of $\log(k^{\text{DMSO}})$ versus phenol σ values (Hammett), $\text{p}K_{\text{a}}$'s, and $E_{1/2}$ values (for the ^XArOH^{0/+} couples) shown in Figure 5 corroborate a change in mechanism across the series of X groups.²³ For the series X = NO₂, CF₃, Cl, H, Me the rate constant increases with increasing electron-withdrawing properties ($\rho = +2.0$, Figure 5a), increasing acidity (lower $\text{p}K_{\text{a}}$, Figure 5b), and increasing redox potential for the phenol substrates (Figure 5c). These results are consistent with a mechanism involving rate-determining proton transfer (PT) from the phenol to [K(Krypt)]-[LCuO₂] to yield the formally Cu(III) complex LCuOOH (Figure 7, top). Supporting the feasibility of such a complex, related species LCuOOR (R = ^tBu, cumyl) have been identified by spectroscopy.²⁸ The failure to observe LCuOOH as an intermediate suggests that it reacts rapidly, with one possible pathway involving reduction by the phenolate ion to yield [LCuOOH][−] that then undergoes fast protonolysis by the excess (acidic) phenol (Figure 7, path (b)). A similar PT process but an alternative subsequent decay pathway was proposed to explain the results of reactions of $(\text{PEDCO})\text{CuO}_2^+$ with a series of para-substituted phenols ^XArOH (X = OPh, Me, H, F, Cl)¹⁴ and ^{Ar}LCuO₂ with ^XArOH (X = H, NO₂) and 2-amino-4-*tert*-butylphenol.³⁹ These reactions also yielded phenolate complexes, and mechanisms were proposed involving initial PT, but followed by Cu–O homolysis and loss of HO₂ radical that then disproportionates to yield H₂O₂ and O₂ (Figure 7, path (a)). We cannot unequivocally distinguish between paths (a) and (b) with the available evidence, although the observation of the same yield of final product in reactions using only 1 equiv of substrate phenol argues against path (b). Importantly, both paths involve ET, such that the overall reaction is a PT/ET process.

The lines through the data for X = NMe₂, OMe, ^tBu, and Me in Figure 5 paint a different picture. The rate constant decreases with increasing electron-withdrawing properties ($\rho = -2.5$, Figure 5a), increasing acidity (lower $\text{p}K_{\text{a}}$, Figure 5b), and

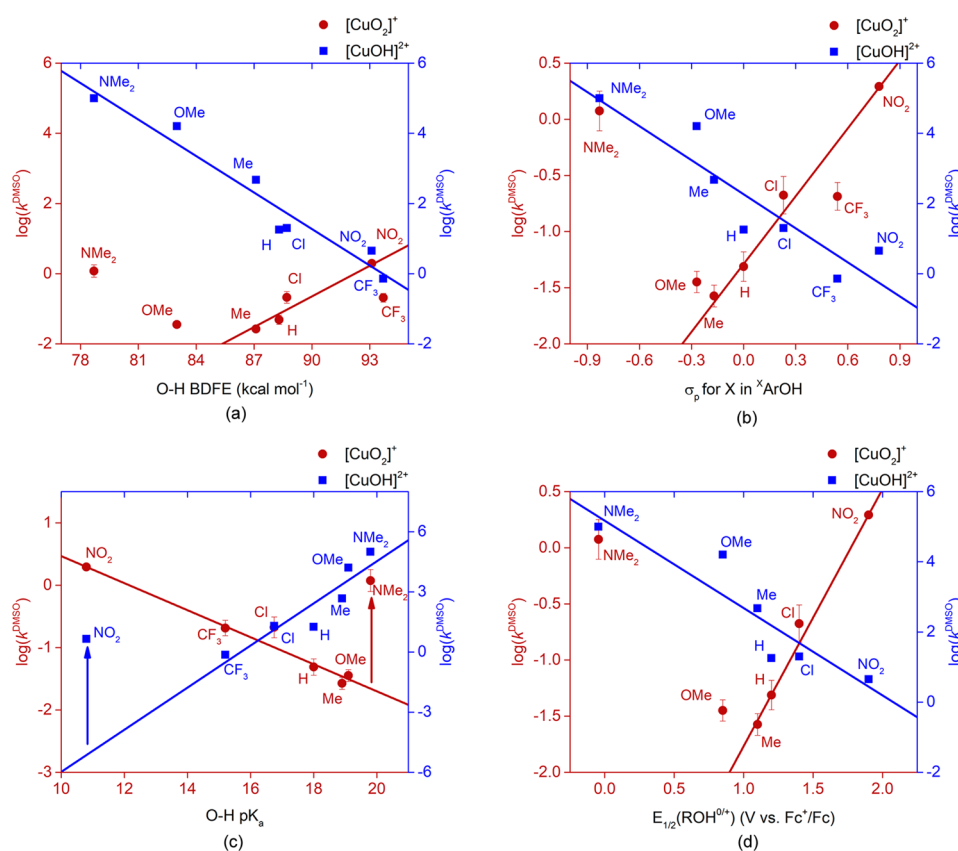


Figure 9. (a) $\log(k^{\text{DMSO}})$ versus $^X\text{ArOH}$ BDFE overlay plot for the reaction of $[\text{CuO}_2]^+$ (red circles, -60°C) and $[\text{CuOH}]^{2+}$ (blue squares, -80°C) cores with $^X\text{ArOH}$ ($X = \text{NMe}_2, \text{OMe}, \text{Me}, \text{H}, \text{Cl}, \text{CF}_3, \text{NO}_2$). The red line depicts an arbitrary linear correlation for the data points for $X = \text{Me}, \text{H}, \text{Cl}, \text{CF}_3$, and NO_2 ($R^2 = 0.779$, slope = 0.29). The blue line is a fit to the data for all X ($R^2 = 0.932$, slope = -0.35). (b) $\log(k^{\text{DMSO}})$ versus $^X\text{ArOH}$ σ_p overlay plot for the $[\text{CuO}_2]^+$ (red circles, -60°C) and $[\text{CuOH}]^{2+}$ (blue squares, -80°C) cores. The red line depicts a fit to the data for $X = \text{Me}, \text{H}, \text{Cl}, \text{CF}_3$, and NO_2 ($R^2 = 0.959$, slope = 2.0). The blue line depicts a fit to the data for all X ($R^2 = 0.806$, slope = -3.2). (c) $\log(k^{\text{DMSO}})$ versus $^X\text{ArOH}$ pK_a overlay plot for the $[\text{CuO}_2]^+$ (red circles, -60°C) and $[\text{CuOH}]^{2+}$ (blue squares, -80°C) cores. The red and blue lines are fits to the data for all X except NMe_2 (red, $R^2 = 0.992$, slope = -0.22) or except NO_2 (blue, $R^2 = 0.811$, slope = 1.1). (d) $\log(k^{\text{DMSO}})$ versus $^X\text{ArOH}^{0/+}$ $E_{1/2}$ overlay plot for the $[\text{CuO}_2]^+$ (red circles, -60°C) and $[\text{CuOH}]^{2+}$ (blue squares, -80°C) cores ($E_{1/2}$ values referenced to the Fc^+/Fc redox couple). The red line is a fit for $X = \text{Me}, \text{H}, \text{Cl}$, and NO_2 ($R^2 = 0.996$, slope = 2.3). The blue line is a fit to the data for all X ($R^2 = 0.798$, slope = -2.5).

increasing redox potential for the phenol substrates (Figure 5c). We interpret these results, as well as the observation in these cases of an intermediate formulated as $[\text{LCuOOH}]^-$, to indicate that a CPET/HAT process is followed, akin to that seen in the reaction with TEMPOH (Figure 7, blue). This contrasting reactivity is analogous to that reported for $^{\text{Ar}}\text{LCuO}_2$, which performs HAT with 2-hydroxy-2-azaadamantane but PT/ET with some phenols.³⁹ Yet, to our knowledge, the observation of a changeover in reaction pathway for one type of substrate that we observe is unique in copper–oxygen chemistry.

Deeper insights may be drawn by comparing the data for the reactions of $^X\text{ArOH}$ with $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ to those we reported previously for LCuOH ,¹⁶ which features a different $[\text{CuOH}]^{2+}$ core but with the same supporting ligand (Figure 9). Overall, the rate constants for the reactions with LCuOH measured at -80°C are significantly larger ($\log k^{\text{DMSO}}$ range from ca. 0 to +5) relative to those for $[\text{K}(\text{Krypt})][\text{LCuO}_2]$ ($\log k^{\text{DMSO}}$ range from ca. -1.6 to +0.4) measured at -60°C .⁴¹ Clearly, at parity of the supporting ligand, the $[\text{CuOH}]^{2+}$ core is more reactive than the $[\text{CuO}_2]^+$ unit. Even more striking are the essentially opposite slopes to the linear fits (blue versus red, Figure 9). In the $\log(k^{\text{DMSO}})$ versus O–H BDFE plot (a), for $[\text{CuOH}]^{2+}$ (blue) the rate constant increases as the bond strength decreases, as expected for a predominate CPET/HAT

process. The data for the $[\text{CuO}_2]^+$ complex is much more scattered, with a fit to the data for $X = \text{Me}, \text{H}, \text{Cl}, \text{CF}_3$, and NO_2 , indicating a poor correlation ($R^2 = 0.779$) and an opposite trend (faster rate for substrates with stronger O–H bonds). Notably, similar rate constants are observed for $X = \text{NO}_2$ and NMe_2 that have drastically different O–H BDFE values (93 versus 79 kcal/mol, respectively). These data argue against a predominant CPET/HAT process and support a mechanism change across the series. Similarly, the Hammett plot (b) and the $\log(k^{\text{DMSO}})$ versus $E_{1/2}$ plot (d) show divergent slopes for the $[\text{CuOH}]^{2+}$ (blue) and $[\text{CuO}_2]^+$ (red) cores, with the data for $X = \text{NMe}_2$ for the $[\text{CuO}_2]^+$ case being a clear outlier. The $\log(k^{\text{DMSO}})$ versus pK_a plot (c) is particularly revealing. For the case of the $[\text{CuOH}]^{2+}$ (blue) complex, the rate decreases as the acidity increases, but with $X = \text{NO}_2$ being an outlier. In contrast, for the $[\text{CuO}_2]^+$ (red) complex, the rate increases as the acidity increases, but with $X = \text{NMe}_2$ being an outlier. We rationalize these opposite behaviors by (a) a predominantly CPET/HAT pathway for the $[\text{CuOH}]^{2+}$ core, with a PT/ET contributor for the most acidic ($X = \text{NO}_2$) phenol (indicated by arrow), and (b) a predominantly PT/ET pathway for the $[\text{CuO}_2]^+$ core that shifts to CPET/HAT for the least acidic ($X = \text{NMe}_2$) phenol (indicated by arrow). The observation of anomalously high rates for the substrate with $X = \text{NMe}_2$ that has the weakest O–H

bond, most electron-donating substituent, and lowest reduction potential also is consistent with a CPET/HAT mechanism for its reaction with the $[\text{CuO}_2]^+$ unit.

A key question concerns why the two cores exhibit such different behaviors. Comparison of the thermodynamics as reflected in the square schemes A and B (Figures 1 and 10) is

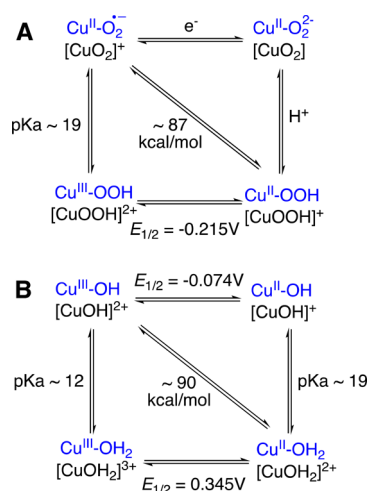


Figure 10. Square schemes A and B (from Figure 1), but with measured or estimated thermodynamic parameters from this work (A) or reported previously (B)⁴² indicated.

revealing. The measured/estimated pK_a , $E_{1/2}$, and BDE values for each core supported by the same ligand L^{2-} are shown in Figure 10.⁴² It is notable that the O–H BDE value for the $[\text{CuOOH}]^+$ core is at least ~ 3 kcal/mol lower than that for the $[\text{CuOH}_2]^{2+}$ unit, consistent with the generally faster rates for PCET by $[\text{CuOH}]^{2+}$ than for $[\text{CuO}_2]^+$. Another notable difference is the basicity of these latter cores; the ~ 7 pK_a unit difference between the pK_a values for the $[\text{CuOOH}]^{2+}$ and $[\text{CuOH}_2]^{3+}$ cores is significant. Also significant is the difference in the $E_{1/2}$ values for the $[\text{CuOOH}]^{2+/+}$ and $[\text{CuOH}_2]^{2+/3+}$ couples (~ 0.5 V higher for the latter). Both of these large differences (pK_a and $E_{1/2}$) are consistent with the one unit differences in charge between the species being compared; the higher charge for the $[\text{CuOH}_2]^{3+}$ core results in a lower pK_a and higher $E_{1/2}$ relative to those of the $[\text{CuOOH}]^{2+}$ unit.

The aforementioned differences in acidity and redox potential translate into key differences in thermodynamics that we postulate underlie the differences in observed reactivity. As described previously, for $[\text{CuOH}]^{2+}$ (B),¹⁶ the difference between the free energy for CPET/HAT (defined as the difference in the BDE values for the phenols and the $[\text{CuOH}_2]^{2+}$ O–H bonds) and the free energy for PT (estimated from the difference in pK_a values for the phenols in DMSO and that of the $[\text{CuOH}_2]^{3+}$ unit in THF) is large for most of the phenols, thus rendering PT unlikely and favoring CPET/HAT (Figure 11a). The exceptions are for the most acidic phenols ($X = \text{CF}_3$ or NO_2), where PT becomes thermodynamically feasible (dashed box). In contrast (Figure 11b), for $[\text{LCuO}_2]^-$ the overall thermodynamic driving forces are smaller, the pK_a for the $[\text{CuOOH}]^{2+}$ core is ~ 7 units higher than that for $[\text{CuOH}_2]^{3+}$, and the enhanced basicity of the $[\text{CuO}_2]^+$ core thus renders PT much more favorable such that only for the least acidic phenols is CPET/HAT preferred. Put another way, as graphically shown in Figure 11, the dashed box (where PT/ET and CPET/HAT paths have similar driving forces) is positioned differently along

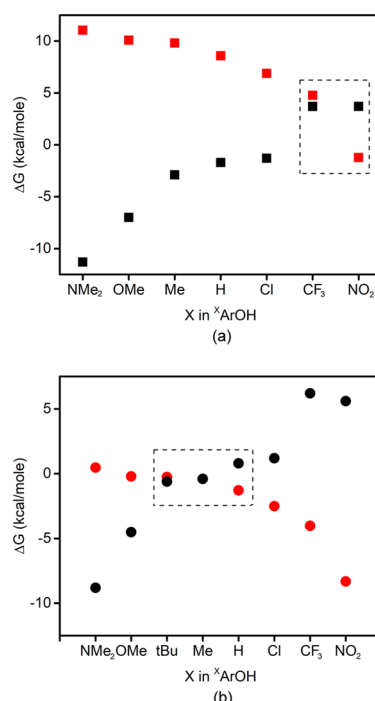


Figure 11. Plots of ΔG for the corresponding PT (red) or CPET (black) reactions with LCuOH (top, squares, adapted from ref 17) and $[\text{LCuO}_2]^-$ (bottom, circles). The dashed regions highlight where there is thermodynamic potential for both PT and CPET mechanisms to occur.

the phenol Hammett σ axis for the two cores, predominantly because of significant differences in the PT thermodynamics (more favorable for $[\text{LCuO}_2]^-$).

Notably relevant to our work are the results of reactions of the $[\text{CuO}_2]^+$ complexes shown in Figure 8 with the weak O–H bonds of 2,2,6,6-tetramethylpiperidine-*N*-hydroxide (TEMPOH) or phenols.^{4,14,33–39} All of the complexes react with TEMPOH to give TEMPO \cdot and compounds with the $[\text{CuOOH}]^+$ core, consistent with hydrogen-atom abstraction (HAT) from the O–H bond having a BDFE ~ 67.5 kcal/mol. Differences arise in reactions with phenols. The $[\text{CuO}_2]^+$ complexes supported by TMG_3tren ,³³ DMA-TMPA ,³⁴ DMM-TMPA ,³⁵ and P^VTMPA ³⁷ yield products derived from phenoxyl radicals and $[\text{CuOOH}]^+$ cores upon reaction with various substituted phenols, again consistent with HAT pathways. Such a pathway for $[(\text{DMM-TMPA})\text{CuO}_2]^+$ was further supported by a detailed mechanistic study and thermodynamic analysis, which notably led to a PT/ET pathway being ruled out.³⁵ In the context of our findings, we hypothesize that for all of these cationic complexes supported by TMG_3tren and R^nTMPA derivatives, HAT is favored because the basicity of the $[\text{CuO}_2]^+$ core is insufficient to make initial PT feasible. Consistent with this notion, even a strong acid like trifluoroacetic acid does not protonate $[(\text{TMG}_3\text{tren})\text{CuO}_2]^+$; instead, an adduct forms that is then susceptible to reduction by octamethyl- or decamethylferrocene.⁴³

On the other hand, reactions of $[(\text{PEDCO})\text{CuO}_2]^+$ with a series of para-substituted phenols $^X\text{ArOH}$ ($X = \text{O}^i\text{Ph}$, Me, H, F, Cl) yielded copper(II)–phenolate complexes $[(\text{PEDCO})\text{-CuOAr}^X]^+$, and a pathway was proposed involving proton transfer to the superoxide (PT) followed by loss of HO_2 radical that then disproportionates to yield H_2O_2 and O_2 (essentially, a PT/ET pathway).¹⁴ Consistent with initial PT, a plot of $\log k$

versus pK_a for the phenols showed an increase in rate as the acidity increased (slope of linear fit = -0.18 , similar to the value of -0.22 that we observed; Figure 9c), although in the study of $[(\text{PEDCO})\text{CuO}_2]^+$ the span in pK_a 's measured was small (2 units) and there was significant deviation in the data. In studies of reactions of $^{\text{Ar}}\text{LCuO}_2$ with phenol and *tert*-butyl-substituted derivatives, PT/ET was implicated for the former and HAT was found for the latter.³⁹ The basis for the apparently similar tendency of the $[\text{CuO}_2]^+$ cores supported by PEDCO and L^{2-} to react with phenols via initial PT, and for both cores supported by $^{\text{Ar}}\text{L}^-$ and L^{2-} to follow either PT/ET or HAT depending on the nature of the phenol, is unclear. It would be useful to have more extensive thermodynamic data for the systems supported by PEDCO and $^{\text{Ar}}\text{L}^-$ in needed efforts to better understand how supporting ligands influence the mechanisms of reactions of $[\text{CuO}_2]^+$ cores.

Mechanistic dichotomies between PT/ET, CPET, or ET/PT within the same system have been observed previously in high valent metal-oxo species. For example, mechanistic changes between HAT and ET/PT were seen for a Mn^{IV} -oxo complex with changing temperature,⁴⁴ and for an Fe^{IV} -oxo complex on addition of scandium.⁴⁵ A changeover between PCET and PT/ET has also been observed by tuning the ligand basicity in Fe^{IV} -oxo species.⁴⁶ The degree to which proton and electron transfer are coupled in CPET processes has recently been highlighted for an array of Fe^{IV} -oxo compounds through the introduction of the asynchronicity factor, η ,⁴⁷ and has been implemented experimentally to describe C–H bond activation by a terminal Co^{III} -oxo complex.⁴⁸ In the latter system, reaction rates were found to be inversely related to the substrate pK_a (as observed in our work for acidic phenols). However, experimental evidence did not support a stepwise process (PT/ET), but instead supported basic asynchronous CPET behavior, where a more asynchronous process resulted in higher rates. This example has important ramifications for other metal-oxo systems, where the relative reactivity with R–H substrates has been linked to the basicity of the oxo ligand,⁴⁹ as well as to our work on systems supported by L^{2-} that we report herein. Indeed, through an analysis akin to that reported previously involving determination of η and assessment of a $\Delta E_{\text{eff}}^{\circ}$ and $\Delta pK_{a,\text{eff}}$ correlation plot (Figure S21),^{47,48} we confirmed that the reaction of LCuOH with $^{\text{X}}\text{ArOH}$ via CPET is exergonic for phenols with $\text{X} = \text{NMe}_2$, OMe , Me , H , Cl , but endergonic for $\text{X} = \text{NO}_2$. Moreover, none of the phenols react through a true synchronous CPET mechanism; instead, asynchronicity in CPET favoring ET ($\text{X} = \text{NMe}_2$) or favoring PT ($\text{X} = \text{OMe}$, Me , H , Cl) is proposed, while a stepwise PT/ET mechanism is proposed for $\text{X} = \text{NO}_2$ (CPET endergonic). Unfortunately, because of our current inability to measure all thermodynamic parameters for $[\text{LCuO}_2]^-$ (Figure 10 (A)), we were unable to perform a corresponding analysis of asynchronicity for this complex.

CONCLUSIONS

Studies of the reactions with the O–H bonds in TEMPOH and the series of phenols $^{\text{X}}\text{ArOH}$ ($\text{X} = \text{para}$ -substituent NO_2 , CF_3 , Cl , H , Me , $^{\text{t}}\text{Bu}$, OMe , or NMe_2) have shown $[\text{K}(\text{Krypt})]^-$ $[\text{LCuO}_2]$ to be capable of divergent proton-coupled electron-transfer mechanisms spanning PT/ET and CPET/HAT processes, depending on the nature of the substrate. Proton transfer (PT) is notably favorable with respect to kinetics and thermodynamics, such that CPET (or HAT) prevails for substrates that have relatively low acidities, low redox potentials, and weak O–H bonds. We surmise that the dominance of PT/

ET paths is due in large part to the anionic nature of the complex, a unique attribute arising from the dianionic supporting ligand L^{2-} relative to other known species with $[\text{CuO}_2]^+$ cores. The reactivity is notably unlike that of LCuOH , a particularly interesting comparison to draw because the complex contains a different $[\text{CuOH}]^{2+}$ core supported by the same ligand L^{2-} . Reactions of LCuOH with $^{\text{X}}\text{ArOH}$ are significantly faster, and CPET/HAT paths are favored, such that PT/ET is apparent only for the substrate $^{\text{NO}_2}\text{ArOH}$ with the highest acidity, highest redox potential, and strongest O–H bond. The significantly higher oxidizing power, lower basicity, and stronger O–H bond in the product $[\text{CuOH}_2]^{2+}$ core all contribute to the divergent reactivity of LCuOH relative to $[\text{LCuO}_2]^-$. An implication of these findings is that at parity of supporting ligand donors, one would expect similarly divergent reactivity of $[\text{CuO}_2]^+$ and $[\text{CuOH}]^{2+}$ cores in enzymes or other catalysts. In addition, one would anticipate that for the $[\text{CuO}_2]^+$ unit, or others in the upper left quadrant of the cascade in Figure 1 (with intact O–O bonds), to exhibit reactivity competitive with the $[\text{CuOH}]^{2+}$ core, or others in the lower right quadrant of Figure 1 (lacking O–O bonds), perturbations of the supporting ligand and/or local environment would be needed.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/jacs.9b00466.

Experimental details, including procedures, syntheses, and spectroscopic and kinetic data (PDF)

X-ray crystallographic data (CIF)

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The authors declare no competing financial interest.

X-ray data has been deposited in the CSD (1880915, 1880916, and 1880917) and may be accessed at <https://www.ccdc.cam.ac.uk/>.

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(24) In the two-phase reaction, k_1 ($0.44(9) \text{ M}^{-1} \text{ s}^{-1}$) for ^{OMe}ArOH is statistically equal (3σ) to the second-order rate constant, k ($0.7(1) \text{ M}^{-1} \text{ s}^{-1}$), derived from the single exponential decay phase. For continuity with other ^XArOH substrates, all arguments invoking rate data use k from the single exponential decay phase.

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