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# Electrochemical Water Oxidation Using a Copper Complex

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## Abstract

Herein, we report the application of the mononuclear copper complex **1**, [Cu<sup>II</sup>(L)]<sup>2+</sup>, in the electrochemical water oxidation catalysis (L = 1,3-bis(((1-methyl-1H-imidazol-2-yl)methyl)amino)propan-2-ol). The complex exhibits a N<sub>4</sub> donor set consisting of two amine and two imidazole units and a dangling OH unit in close proximity to the copper ion. **1** exhibits a moderate apparent rate constant  $k_{\text{cat}}$  of 0.12 s<sup>-1</sup> in catalysis and operates at an overpotential of 0.83 V. Detailed investigation allowed us to derive a mechanism for water oxidation. The catalysis proceeds only under basic conditions, where [Cu<sup>II</sup>(L)(OH)]<sup>+</sup>, 1H<sub>-1</sub>, is the main solution species, which indicates that a negatively charged ligand is necessary to drive catalysis. Initial oxidation of 1H<sub>-1</sub> is coupled to proton loss forming a copper(III) species and further oxidation initiates oxygen evolution. Initial oxidation of **1** under neutral, i.e. non-catalytic, conditions is pH independent, highlighting the importance of PCET steps during catalysis. We collected reasonable evidence that catalysis proceeds *via* a water nucleophilic attack mechanism. The electrolyte presumably acts as proton acceptor in catalysis as the onset potential depends of the buffer employed.

## Introduction

The production of sustainable fuels such as H<sub>2</sub>, methanol, CO or formate requires the reduction of the oxidised precursors H<sup>+</sup> or CO<sub>2</sub>.<sup>1</sup> Water is considered as a cheap and abundant electron and proton source in this regard. Its oxidation requires a potential of 1.23 V under standard conditions. One of the most efficient catalysts in the electrochemical oxygen evolution catalysis (OEC) is platinum metal.<sup>2</sup> However, platinum's high price makes it rather unattractive and therefore catalysts based on Earth-abundant metals have been developed. Molecular electrocatalysts based on iron, nickel, cobalt, and copper have been identified, which lower the overpotential substantially and are stable under the highly oxidative conditions of the catalysis.<sup>3</sup> Since the initial report of the copper bipyridine system, **A**, by Mayer and co-workers in 2012,<sup>4</sup> copper catalysts attracted considerable attention.<sup>5</sup> Selected molecular copper complexes with N-donor ligands, which have been explored in molecular OEC, are shown in Table 1.

Table 1. Selected mononuclear copper complexes, which have been explored in molecular OEC and key catalytic properties. Structures **A-H** see Figure S 1.

	Donor set L	pH	$\eta$ /V	$k_{\text{cat}}$ / s <sup>-1</sup>	TON
<b>A</b> <sup>d</sup>	N <sub>2</sub>	12.5	0.75 <sup>b</sup>	100	30
<b>B</b> <sup>5a</sup>	N <sub>4</sub> O	11	0.52 <sup>a</sup>	33	13
<b>C</b> <sup>5c</sup>	N <sub>2</sub>	12.4	0.64 <sup>c</sup>	0.4	1
<b>D</b> <sup>5e</sup>	N <sub>5</sub>	8	0.64 <sup>c</sup>	~20	19
<b>E</b> <sup>15f</sup>	N <sub>4</sub>	11.5	0.40	3.6	0.5
<b>E</b> <sup>25f</sup>	N <sub>4</sub>	11.5	0.17	0.16	0.5
<b>F</b> <sup>5k</sup>	N <sub>4</sub>	7	0.88 <sup>b</sup>	7	n.a. <sup>d</sup>
<b>G</b> <sup>5m</sup>	N <sub>2</sub>	12	0.30 <sup>a</sup>	35	n.a. <sup>d</sup>
<b>H</b> <sup>5n</sup>	N <sub>4</sub>	12.5	1.00	33	1.43

$\eta$ :  $E(\text{O}_2|\text{H}_2\text{O})$  relative to <sup>a</sup> half peak potential, <sup>b</sup> onset potential, <sup>c</sup> applied potential during electrolysis; <sup>d</sup> n.a. = not available,

The complexes show moderate to high apparent rate constants  $k_{\text{cat}}$  at overpotentials as low as 0.17 V.<sup>5f</sup> Some of these complexes (**E**, **F**) exhibited an equatorial N donor ligand-set and such complexes proved to be stable under the oxidative conditions of the catalysis. This prompted

us to investigate **1** as catalyst in the water oxidation reaction (Figure 1).<sup>6</sup> **1** exhibits a tetradentate N ligand set and a dangling OH unit, which could allow for efficient proton transfer (PT) pathways during catalysis.<sup>7</sup> The complex also catalyses the electrochemical dihydrogen evolution from seawater – Earth's most abundant water source,<sup>6b</sup> which would potentially allow water splitting catalysis in a one cell compartment with cheap electrode materials.

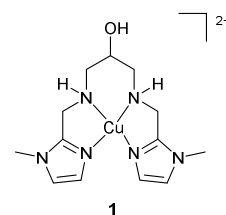


Figure 1. Copper complex that catalyses the electrochemical hydrogen<sup>6b</sup> and oxygen evolution reaction.

## Results and discussion

The synthesis, purification, and characterisation of **1** has been described previously.<sup>6</sup> The purity of the complex was confirmed by elemental analysis. A common three electrode setup employing an SCE reference electrode, a platinum counter electrode and a GC working electrode was used for electrochemical measurements. If not otherwise noted, potentials are referenced vs. NHE. The complex exhibits a reduction process at -0.28 V in neutral water. Under basic conditions an increased current was observed at applied potentials above 1 V. A pre-wave was visible at ~1 V and a catalytic wave with an onset potential of ~1.3 V at solution's pH above 10 (Figure 2, Figure S 2).

CV measurements of **1** in acetonitrile and acetonitrile/water mixtures indicate that the catalytic wave could indeed belong to water oxidation. In dry acetonitrile, no oxidation process of **1** was observed, but the addition of 10 % basic water led to an oxidation



process at a potential of 0.49 V vs.  $\text{Fc}^{+/0}$  and a pronounced wave at a potential of 0.91 V vs.  $\text{Fc}^{+/0}$  (Figure S 3).

We investigated the influence of the working electrode on the catalysis and utilised a boron doped diamond electrode (BDD). However, considerable background activity was visible under basic conditions and therefore the GC electrode was used for all further investigation (Figure S 4).

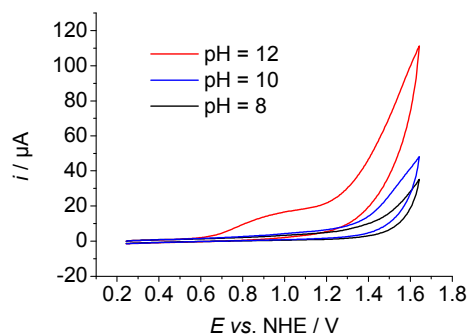


Figure 2. pH dependent CV data of **1** in Millipore water ( $[\mathbf{I}] \approx 1 \text{ mM}$ ,  $I = 0.1 \text{ M PO}_4^{3-}$ ,  $v = 0.1 \text{ mVs}^{-1}$ ).

Controlled potential electrolysis experiment (CPE) at an applied potential of 1.494 V confirmed oxygen evolution under basic conditions, i.e. pH = 12 (Figure S 5). Gas bubbles were visible at the electrode surface and the colour of the solution changed from purple to blue. The amount of evolved  $\text{O}_2$  was analysed by gas chromatography (Figure S 19). Catalysis exhibits a Faraday efficiency of 60 % (average of 6 runs) and a maximum TON of 6 after 30 min of electrolysis, which is rather high in comparison to previous reports employing copper complexes (Table 1). The solution's pH decreased from 12.04 to 11.8 during electrolysis due to proton formation. No hydrogen peroxide was detected after the CPE experiment with a potassium iodide/starch test.

We collected reasonable evidence that the catalyst remains homogeneous during catalysis. Visual inspection of the electrode after CPE did not reveal any deposit. The rinse test was negative, that is a CV and CPE measurement of the electrode after a CPE experiment in freshly prepared buffer solution was similar to the background measurement (Figure S 9). Scanning electron microscopy (SEM) and energy-dispersive X-ray spectroscopy (EDX) of the electrode surface revealed very little deposition of copper containing material (Figures S 23, 24). The electrode surface has a similar roughness and shape as after a CPE experiment without the complex (Figures S 25, 26). In line with this, virtually no current passed and no notable oxygen formation has been detected, when we reused the GC electrode in a second experiment employing complex free buffer solution (Figure S 10). That is to say that the very small amounts of deposited copper containing material are not active in the OER. The concentration of **1** after chemical oxidation and after CPE was the same as determined by visible spectroscopy (Figure S 22). CPE with  $\text{Cu(II)SO}_4$  was used as reference system for copper deposition (Figures S 8, S 27, S 28).<sup>5b</sup>

Subsequently, we investigated the kinetics of the catalysis. Notably, in order to obtain reproducible results, the electrode was polished after each CV scan (Figure S 11). Catalysis proceeds at an overpotential of 0.83 V at pH = 12 as determined from CV data ( $E = 1.35 \text{ V}$  at  $i_{\text{cat}}/2$ ,  $[\mathbf{I}] = 5 \text{ mM}$ ).<sup>8</sup> In order to estimate the apparent rate constant  $k_{\text{cat}}$  of the catalysis, scan rate dependent data were collected (Figure S 12).  $k_{\text{cat}}$  is proportional to  $i_c^2$  as shown in eq. (2).<sup>9</sup>  $i_c$  denotes the catalytic peak current,  $n_{\text{cat}}$  the number of transferred electrons for water oxidation,  $F$  the Faraday constant,  $A$  the electrode surface,  $c_{\text{cat}}$  the catalyst concentration, and  $D$  the diffusion coefficient in phosphate buffer at pH 12.  $D$  was determined from the  $\text{Cu}^{\text{II}}/\text{Cu}^{\text{I}}$  redox pair.

$$i_c = n_{\text{cat}} \cdot F \cdot A \cdot c_{\text{cat}} \cdot D^{\frac{1}{2}} \cdot k_{\text{cat}}^{\frac{1}{2}} \quad (2)$$

The apparent rate constant can be calculated from the  $i_c$  over  $i_p$  vs. the square root of the scan rate plot (eq. (3)).  $i_p$  denotes the current in the no-catalytic event and it has been determined from the  $\text{Cu}^{\text{II}}/\text{Cu}^{\text{I}}$  reduction process of **1** (eq. (S1), Figure S 13).

$$\frac{i_c}{i_p} = \frac{n_p}{0.4463 \cdot n_p} \left( \frac{R \cdot T}{n_p \cdot F} \right)^{\frac{1}{2}} \left( \frac{k_{\text{cat}}}{v} \right)^{\frac{1}{2}} \quad (3)$$

The apparent rate constant at 1.494 V equals  $0.12 \text{ s}^{-1}$ , which is at the lower limit of previously reported catalysts.

A plot of the catalytic current at  $E = 1.494 \text{ V}$  vs. solution's pH indicates a current increase under basic conditions with a rather sharp onset above solution's pH of 10 (Figure 3). The first derivative of the CV traces at different solution's pH revealed two pH-dependent inflection points. The first one is accompanied by a small current increase and the second one by a pronounced one (Figure S 2, Figure S 15).

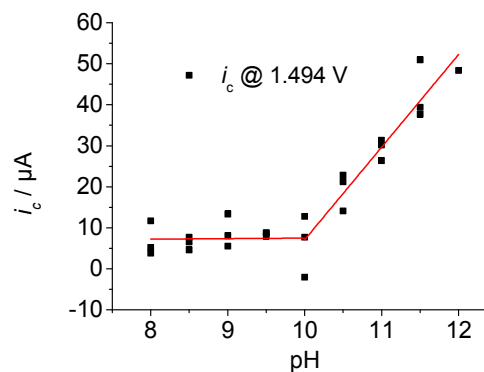


Figure 3. Plot of the current  $i_c$  at  $E = 1.494 \text{ V}$  in dependence of the pH, 3 runs, ( $[\mathbf{I}] \approx 1 \text{ mM}$ ,  $I = 0.1 \text{ M PO}_4^{3-}$ ,  $v = 0.1 \text{ mVs}^{-1}$ ).

We assigned the first oxidation process to an oxidation of the initial copper(II) species to a copper(III) species, as the corresponding zinc complex is redox inactive in this potential region (Figure S 14). No catalysis has been observed at an applied potential of the wave of the first oxidation process ( $E^{\text{app1}} = 0.944 \text{ V}$ , Figure S 5). Thus, further oxidation beyond the copper(III) state initiates



catalysis. Previous speciation studies showed that  $[\text{Cu}(\text{L})(\text{H}_2\text{O})]^{2+}$  is the dominant species below solution's pH 11 and the deprotonated species  $[\text{Cu}(\text{L})(\text{OH})]^+$  above pH of 11.<sup>6b</sup> This indicates, that the doubly charged complex exhibits no noticeable activity and negatively charged ligands are important to enter catalysis. The peak potential of the first oxidation process is pH independent below pH  $\sim 10$  and it shifts by 54 mV per pH above pH  $\sim 10$ , which is characteristic for a  $1e^-/1H^+$  coupled redox process according to the Nernst equation (Figure S 15). Since catalysis only takes place under basic conditions, this fact highlights the importance of PCET steps during oxidation. The second oxidation process shifts by 25 mV per pH indicating proton involvement during oxidation. Indeed, the catalytic wave exhibits a strong kinetic isotopic effect (KIE) (Figure S 16): Changing from  $\text{H}_2\text{O}$  to  $\text{D}_2\text{O}$  leads to a pronounced current decrease of the catalytic wave. A KIE of 5 was calculated according to eq. (4).

$$\text{KIE} = \frac{k_{\text{H}_2\text{O}}}{k_{\text{D}_2\text{O}}} = \left( \frac{i_{c,\text{H}_2\text{O}}}{i_{c,\text{D}_2\text{O}}} \right)^2 = 5.25 \quad (4)$$

Three mechanisms are discussed in literature for O–O bond formation pathways in water oxidation catalysis, water nucleophilic attack at high valent metal species (WNA),<sup>10</sup> radical coupling of two high valent metal species (RC)<sup>11</sup> or redox isomerisation of a dinuclear  $\text{Cu}_2\text{O}_2$  species (RI).<sup>5n</sup> The first type proceeds *via* single sites, while the latter two mechanisms involve binuclear species. Such dimeric species have not been observed previously for single site Cu complexes bearing an equatorial  $N_4$  donor sets and it was also very unlikely for **1**: (i) Previous speciation studies of **1** in solution showed the formation of a mononuclear complex, when a copper to ligand ratio of one is employed<sup>6b</sup> and (ii) the equatorial  $N_4$  coordination mode of the ligand precludes dimerisation as observed for **A**<sup>4</sup> or **H**<sup>5n</sup>. Indeed, concentration dependent measurements showed a first order kinetics in **1** as the current increases linearly with increasing catalyst concentration ( $[\text{1}] = 0.5 - 5.0 \text{ mM}$ , Figure 4).

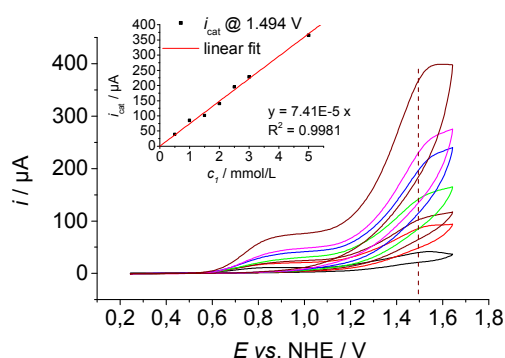
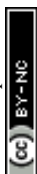


Figure 4. Concentration dependent CV data of **1** ( $[\text{1}] \approx 0.5 - 5.0 \text{ mM}$ ,  $I = 0.1 \text{ M PO}_4^{3-}$ ,  $\text{pH} = 12$ ,  $v = 0.1 \text{ Vs}^{-1}$ ). Inset: Plot of the catalytic current  $i_c$  at  $E = 1.494 \text{ V}$  vs.  $[\text{1}]$ .

This indicates that the catalysis proceeds through a WNA mechanism. In the WNA mechanism, water attacks the highly oxidised copper species forming a  $\text{Cu}^{\text{IV}}\text{OOH}$

species, and further oxidation of the intermediate leads to dioxygen release. Meyer and co-workers proposed a  $\text{Cu}^{\text{IV}}=\text{O}/\text{Cu}^{\text{III}}-\text{O}^{\cdot}$  species as crucial intermediate after initial  $2e^-$  oxidation of **B**, which reacts with water forming a copper(II)peroxide species<sup>5a</sup> while Llobet and co-workers gave evidence for the formation of  $[(\text{L})\text{Cu}^{\text{III}}(\text{OH})]$  after initial  $2e^-$  oxidation of **E**, which binds an additional hydroxide ion only weakly leading to a SET-WNA mechanism.<sup>5f,12</sup> Ligand oxidation was less likely in **1**, as the zinc complex is redox inactive under oxidative conditions (Figure S 14). CV measurements in the presence of various amounts of  $\text{H}_2\text{O}_2$  did not allow to conclude on the formation of a peroxide intermediate, as  $\text{H}_2\text{O}_2$  oxidation occurs at a similar potential (Figure S 17). Addition of  $\text{H}_2\text{O}_2$  leads to a colour change from purple to blue, and Vis spectroscopy showed a blue-shift of the d-d-band of **1** from 580 nm over 660 nm (10 eq.) to 675 nm (100 eq., Figure S 20). The colour change and the blue-shift of the d-d-bands was similar to the shift observed after CPE (Figure S 21):  $\lambda_{\text{max}}$  shifted to 655 nm after CPE at 0.944 V and to 675 nm after CPE at 1.494 V.

Finally, the influence of the electrolyte on catalysis was evaluated, as the redox potential shift per pH and the large H/D isotope effect indicates PCET during catalysis and the buffer could act as proton acceptor. The first oxidation wave, which we assigned to the  $\text{Cu}^{\text{II}}/\text{Cu}^{\text{III}}$  oxidation, has a similar potential in all measurements independently of the electrolyte (Figure 5, phosphate, borate, acetate and carbonate salt). We also observe a catalytic wave in all four solutions indicating that the activity is an inherent property of **1** and not due to an impurity in the electrolytes (Figure S 18). By far the highest catalytic current is observed with borate buffer. For phosphate, carbonate, and acetate, the catalytic current is very similar, indicating similar catalytic activity in all three cases (Table 2). The onset of the catalytic wave depends on the electrolyte and therefore we assume, that the anion is involved in the catalysis as proton acceptor, which was discussed previously.<sup>5a, e</sup> Notably, we observe a trace crossing of the forward and reverse scan, when carbonate or acetate is employed as electrolyte. Such trace crossing could indicate the formation of an oxidised species, that reacts in a chemical reaction to a second species, which has a less positive standard potential with respect to the initial catalyst.<sup>13</sup> Such scenario is consistent with the formation of a peroxide intermediate, which is easier to reduce than the initial catalyst.<sup>5b</sup>



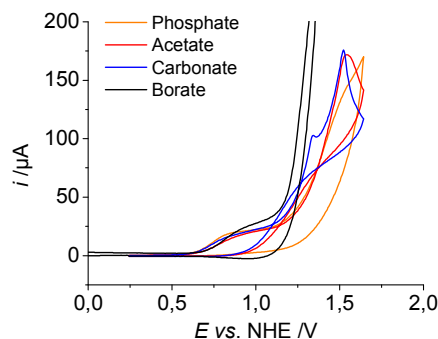
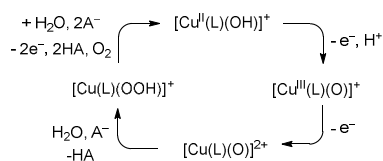


Figure 5. CV data of **1** with various electrolytes, ( $[I] \approx 1$  mM,  $I = 0.1$  M,  $pH = 12$ ,  $\nu = 0.1$  Vs<sup>-1</sup>).

Table 2.  $i_c$  over  $i_p$  ratios of **1** in various different buffer solutions,  $i_c$  at 1.494 V vs. NHE,  $i_p$  of the Cu<sup>II</sup>/Cu<sup>I</sup> couple ( $[I] \approx 1$  mM,  $I = 0.1$  M,  $pH = 12$ ,  $\nu = 0.1$  Vs<sup>-1</sup>).

Electrolyte	$E^{\text{onset}}$ i @ 50 $\mu$ A	$E$ @ $i_c/2$	$i_c/i_p$
B(OH) <sub>3</sub>	1.18 V	1.32 V	40
Na <sub>2</sub> CO <sub>3</sub>	1.24 V	1.39 V	9
NaOAc	1.30 V	1.36 V	10
K <sub>2</sub> HPO <sub>4</sub>	1.29 V	1.35 V	7

Taking all experimental observations together, we derived a minimum mechanism for water oxidation employing **1** in phosphate buffer (Scheme 1). The initial oxidation of [Cu<sup>II</sup>(L)(OH)]<sup>+</sup> forming [Cu<sup>III</sup>(L)(O)]<sup>+</sup> is metal based and coupled to proton loss. Further oxidation of [Cu<sup>III</sup>(L)(O)]<sup>+</sup> initiates catalysis. This step is also coupled to PT events and we have reasonable evidence, that the buffer acts as proton acceptor, since the onset potential of the catalysis depends on electrolyte. Subsequently, a peroxide intermediate is formed *via* WNA at the high valent copper species, which is oxidised at a lower standard potential than the initial catalyst and this oxidation leads to oxygen evolution.



Scheme 1. Proposed mechanism for the electrochemical water oxidation catalysed by **1** under basic conditions.

### Conclusion

**1** proved to be a stable, homogeneous catalyst in the electrochemical-driven water oxidation catalysis. In comparison to previously reported copper catalysts, the complex exhibits a moderate apparent rate constant and a reasonable overpotential. The dangling OH unit has no impact on catalysis. The complex is active under basic conditions, highlighting the need of a negatively charged ligand at the metal centre and PCET steps during oxidation. Under neutral conditions, the complex exhibits neutral ligands and its initial oxidation is not coupled to PT, while 1H<sub>-1</sub> is the main solution species under basic conditions, and its initial oxidation is accompanied by a

proton loss. Further oxidation is also coupled to PT events and initiates catalysis under basic conditions. The catalysis proceeds presumably *via* a single site mechanism involving the formation of a peroxide intermediate through nucleophilic attack of water at and high valent copper-oxido species.

### Experimental

**Synthesis.** The synthesis and purification of **1** has been described previously.<sup>6</sup>

**Electrochemistry.** A common three electrode setup was used with a glassy carbon electrode (CH Instruments,  $A = 7.1$  mm<sup>2</sup>) or boron doped diamond (Windsor Scientific,  $A = 7.1$  mm<sup>2</sup>) as working electrode, a platinum wire as counter electrode (99.999 %, 1 mm diameter) and a SCE as reference electrode (ALS). All electrochemical data were referenced against NHE by adding 0.244 V to the measured potentials. Before *each scan* the GC working electrode was pre-treated by rinsing with millipore water, 30 seconds polishing with an alox-slurry (0.05  $\mu$ m), rinsing with millipore water, 3 minutes sonification in millipore water, rinsing with millipore water and drying. Electrolyte solution (0.1 M) were prepared by titrating a 0.1 M K<sub>2</sub>HPO<sub>4</sub> solution with KOH. Unless otherwise noted, the electrolyte solution was stirred over Chelex® (100-200 mesh particle size, sodium form) for 20 minutes and the filtrate was used to improve the reproducibility and to remove traces of other metals, if present.

For bulk electrolysis the solutions were degassed by “pump, freeze, thaw” cycles. The solutions were frozen, evacuated and then the solutions were allowed to defreeze. This was repeated 4 times. A glassy carbon rod (7 mm diameter) was used as working electrode and a platinum spiral as counter electrode. Bulk electrolysis was performed in a custom made air tight two compartment cell. Working and counter electrode were separated by a glass frit. For better reproducible results the working electrode was cycled 30 times between 1.84 V and -0.76 V with 500 mVs<sup>-1</sup> in electrolyte solution before electrolysis. For the surface analysis GC plates (1x1 cm) were used in the same air tight two compartment cells as for the bulk electrolysis and pre-treated in the same way as the working electrode for bulk electrolysis. After electrolysis the plates were carefully rinsed with Millipore water and dried.

### Acknowledgements

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### Conflicts of interest

There are no conflicts to declare.

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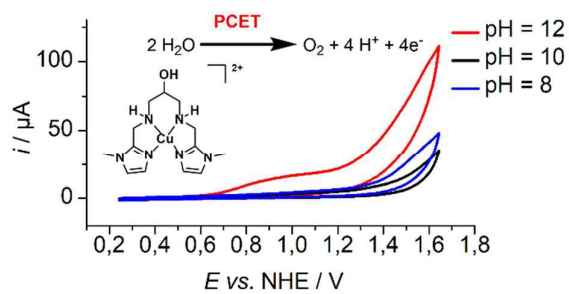
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## TOC



The study highlights the importance of proton coupled electron transfer (PCET) during electrochemical-driven water oxidation catalysis employing a copper complex.

