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Oxygen-atom transfer photochemistry of a molecular copper bromate complex†

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We report the synthesis and oxygen-atom transfer (OAT) photochemistry of $[\text{Cu}(\text{tpa})\text{BrO}_3]\text{ClO}_4$. *In situ* spectroscopy and *in crystallo* experiments indicate OAT proceeds from a Cu–O fragment generated by sequential Cu–O bond cleavage and OAT from BrO_x to $[\text{Cu}(\text{tpa})]^+$. These results highlight synthetic opportunities in M–O photochemistry and demonstrate the utility of *in crystallo* experiments to evaluating photochemical reaction mechanisms.

Reactive ligand-supported metal–oxygen (M–O) species are at the heart of oxygen-atom transfer catalysis in synthesis and biology and have featured prominently in the development of modern theories of metal–ligand (M–L) bonding. As a result, the synthesis and characterization of reactive metal–ligand fragments have garnered significant interest. In the context of Cu–O chemistry, which are responsible for methane hydroxylation in both particulate monooxygenases (pMMOs)^{1–5} and copper-doped porous materials, enormous synthetic effort has been directed towards the synthesis of molecular site mimics.^{6–11} These studies have resulted in the isolation of copper hydroxides, peroxides, hydroperoxides, and superoxides, but the reactivity of terminal Cu–O species has thus far precluded isolation of this structural motif.²

Synthetic photochemistry can enable cryogenic synthesis and thus can enable observation and characterization of species that are otherwise short-lived. To this end, photolysis of metal oxanion complexes has been explored as a potential source of reactive metal oxygen fragments. In 2015, Zhang and coworkers reported that photolysis of Fe(III) bromate complexes afforded either Fe(IV) or Fe(V) oxo complexes depending on the structure of the supporting porphyrin ligand (Fig. 1a).¹² In related studies of matrix isolated Fe(III) oxanions complexes,

Suslick and coworkers demonstrated that oxanion photochemistry was initiated by Fe–O bond homolysis and suggested that subsequent OAT from the extruded radical to the reduced metal center was responsible for oxo complexes observed in steady-state, solution-phase experiments (Fig. 1b).^{13–15} In the context of Cu oxanion chemistry, in 2020 Roithová *et al.* reported the observation of $\text{Cu}(\text{tpa})\text{O}$ during collision-induced dissociation mass spectrometry (CID-MS) analysis of $[\text{Cu}(\text{tpa})\text{OClO}_2]^+$ (Fig. 1c).¹⁶ While this experiment enabled observation of a transient Cu–O species, no spectroscopic or structural data were available.

Here we describe the synthesis, characterization, and OAT photochemistry of copper bromate complex $[\text{Cu}(\text{tpa})\text{BrO}_3]\text{ClO}_4$ ($[\mathbf{8}]\text{ClO}_4$). Steady-state photolysis of $[\mathbf{8}]\text{ClO}_4$ results in OAT to carbon monoxide as well as olefinic, benzylic, and aliphatic substrates. Mass spectrometry, *in situ* spectroscopy, and

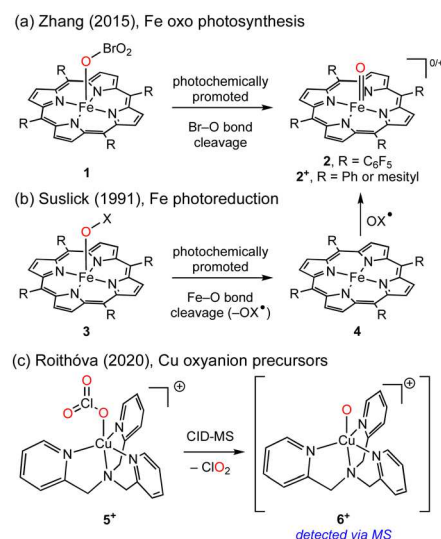


Fig. 1 (a) Ligand-dependent Fe oxo photochemistry of Fe(III) porphyrin bromate complexes. (b) Sequential Fe–O homolysis followed by OAT to the reduced metal fragment could generate M–O species. (c) Collision-induced dissociation mass spectrometry (CID-MS) observation of Cu–O fragment $[\mathbf{6}]^+$.

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Fig. 2 Synthesis of Cu bromate **[8]ClO₄**. Displacement ellipsoid plot of **[8]ClO₄**. H-atoms and counter anions have been omitted for clarity. Ellipsoids drawn at 50% probability. Selected bond lengths (Å): Cu–O = 1.955(3), Cu–N_{pyridine} = 2.072(4), and Cu–N_{amine} = 2.019(4).

in crystallo photochemical experiments are consistent with initial Cu–O bond cleavage to generate $[\text{Cu}(\text{tpa})]^+$ and $\text{BrO}_3^\bullet$. Subsequent OAT from $\text{BrO}_3^\bullet$ (or $\text{BrO}^\bullet$ obtained by decomposition of $\text{BrO}_3^\bullet$ to $\text{BrO}^\bullet$ and O_2) to Cu generates a transient Cu–O fragment that is responsible for substrate functionalization. These results provide new entry into reactive Cu–O species and highlight the power of *in crystallo* photochemistry to probe the reactions of molecular inorganic species.

Treatment of $[\text{Cu}(\text{tpa})\text{Br}]\text{ClO}_4$ (**[7]ClO₄**) with AgBrO_3 affords $[\text{Cu}(\text{tpa})\text{BrO}_3]\text{ClO}_4$ (**[8]ClO₄**) in 65% yield (Fig. 2). The ^1H NMR spectrum of **[8]ClO₄** features two paramagnetically shifted peaks at 10 and 29 ppm (Fig. S1, ESI†). The UV-vis spectrum of **[8]ClO₄** in acetone displays a weak absorbance centered at 426 ($\epsilon = 2.3 \times 10^4 \text{ M}^{-1} \text{ cm}^{-1}$) and stronger absorbances at 710 ($\epsilon = 3.0 \times 10^2 \text{ M}^{-1} \text{ cm}^{-1}$), and 920 nm ($\epsilon = 5.4 \times 10^2 \text{ M}^{-1} \text{ cm}^{-1}$) (Fig. S2, ESI†). The IR spectrum of **[8]ClO₄** displays spectral features at 831, 842, and 852 cm^{-1} , which are attributed to stretching modes of a Cu-bound bromate ligand (Fig. S3, ESI†).¹⁷ Electrospray ionization mass spectrometry (ESI-MS) analysis of **[8]ClO₄** displays a signal at $m/z = 479.98$ with the isotope distribution expected for 8^+ . Consistent with an $S = 1/2$ ground state, the X-band EPR spectrum of **[8]ClO₄** measured at 4 K features an isotropic signal at $g = 2.126$ (Fig. S4, for X-band EPR spectrum of **[7]ClO₄**, see Fig. S5, ESI†).

The solid-state structure of **[8]ClO₄** is illustrated in Fig. 2 (see Fig. S6 and Table S1 for refinement details, ESI†). The Cu(II) ion in **[8]ClO₄** is five-coordinate and exhibits a distorted trigonal bipyramidal geometry ($\tau_5 = 0.77$),¹⁸ with the three pyridine donors from the tpa ligand occupying the equatorial plane. The apical sites are coordinated by the tertiary amine donor of the tpa ligand and an O-bound bromate ligand. The Cu–N distances in **[8]ClO₄** (Cu–N_{pyridine} = 2.072(4) Å, Cu–N_{amine} = 2.019(4) Å) are similar to other crystallographically characterized Cu(II)(tpa) complexes and the Cu–O distance (1.955(3) Å) is well-matched to crystallographically characterized Cu(II)(tpa) oxyanion complexes (Table S2, ESI†).

Copper(II) bromate **[8]ClO₄** participates in photochemically promoted oxygen-atom transfer (OAT) and hydrogen-atom abstraction (HAA) reactions, which are characteristic of the reactivity patterns of reactive M–O species (Fig. 3). Photolysis ($\lambda > 335 \text{ nm}$) of a CO-saturated CH_3CN solution of complex **[8]ClO₄** afforded CO_2 (detected by GC analysis of the reaction headspace, Fig. S7, ESI†). Similar OAT to olefinic substrates was observed: Photolysis of **[8]ClO₄** in the presence of styrene or 1-octene afforded epoxides **9** and **10** in 91% and 36%

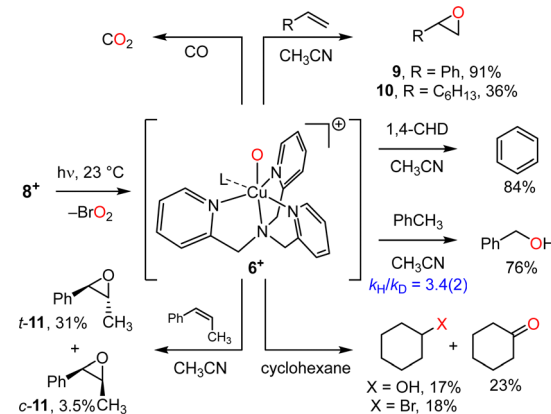


Fig. 3 Summary of photochemically promoted substrate functionalization chemistry. Both OAT and HAA reactivity could arise from a transiently generated reactive CuO fragment (*i.e.*, $[\text{Cu}(\text{tpa})(\text{L})\text{O}]^+$ (**6⁺**); L = solvent or vacant coordination site).

yields, respectively (Fig. S8 and S9, ESI†). Photolysis in the presence of 1,4-cyclohexadiene (bond dissociation energy (BDE) = $76.0 \text{ kcal mol}^{-1}$)¹⁹ yielded benzene (84% yield), the product of HAA (Fig. S10, ESI†). Photolysis of **[8]ClO₄** in toluene (BDE = $89.7 \text{ kcal mol}^{-1}$)²⁰ afforded benzyl alcohol in 76% yield (Fig. S11, ESI†)²¹ and photolysis in the presence of cyclohexane (BDE = $99.5 \text{ kcal mol}^{-1}$)²² afforded a mixture of cyclohexanone (23% yield), cyclohexanol (17% yield), and bromocyclohexane (18% yield) (Fig. S12, ESI†). Photolysis of a CH_3CN solution of **[8]ClO₄** in the presence of 1 : 1 H_8 -toluene/ D_8 -toluene provided a kinetic isotope effect (KIE; $k_{\text{H}}/k_{\text{D}}$) of 3.4(2) for benzylic hydroxylation, which is consistent with significant C–H cleavage in the rate-determining transition state (Fig. S13, ESI†). Photolysis of **[8]ClO₄** in the presence of pentane (BDE = 97.5)²³ or benzene (BDE = 112.9)²⁰ resulted in no observed substrate oxidation. In no case were products of OAT or HAA observed without photolysis (*i.e.*, there are no background reactions with any of the substrates).

To gain additional insight into the mechanism of OAT and the potential intermediacy of $[\text{CuO}]$ species (*e.g.* **6⁺**), we scrutinized the photochemical epoxidation of stereochemically defined 1,2-disubstituted olefins. Epoxidation of *cis*- β -methylstyrene is not stereospecific: Photolysis of **[8]ClO₄** in the presence of *cis*- β -methylstyrene afforded a 10 : 1 mixture of *trans*:*cis*-**11** in 35% yield (Fig. S14, ESI†). For comparison, we also carried out the epoxidation of *cis*- β -methylstyrene with $[\text{Cu}(\text{tpa})]\text{BF}_4$ (**[12]BF₄**) with PhIO and obtained a 10 : 1 mixture of *trans*:*cis*-**11** (Fig. S15, ESI†). The identical stereochemical outcome suggests that both photo-promoted and Cu(I) mediated processes proceed *via* a common intermediate, *e.g.* a CuO fragment. Taken with the primary KIE observed for benzylic hydroxylation of toluene, these data are consistent with stepwise substrate activation *via* carbon-centered radicals (*i.e.*, alkyl radicals generated by either radical addition to olefins or *via* H-atom abstraction (HAA) from alkanes).

In analogy to the oxyanion photochemistry described in Fig. 1, we envisioned that the formation of a $[\text{CuO}]$ intermediate could occur *via* direct photooxidation, *i.e.*, cleavage of the O–Br

bond to generate $[\text{Cu}(\text{tpa})\text{O}]^+$ (6^+) and BrO_2 , or *via* a photo-reduction/OAT sequence, *i.e.*, initial cleavage of the Cu–O bond to generate a $[\text{Cu}(\text{tpa})]^+$ fragment (12^+) and $\text{BrO}_3^\bullet$ following by OAT from $\text{BrO}_3^\bullet$ (or BrO derived thereof) to 12^+ . A combination of MALDI-MS, *in situ* spectroscopy data, and *in crystallo* photochemistry suggest the latter mechanism is operative under steady-state photolysis of $[\text{8}]\text{ClO}_4$.

Mass spectrometry data provides evidence for the ions expected from sequential photoreduction and subsequent OAT from $\text{BrO}_3^\bullet$ to 12^+ : $[\text{Cu}(\text{tpa})]^+$ is the base ion in the MALDI-MS of $[\text{8}]\text{ClO}_4$ (indicated by the set of peaks at 353.0 and 355.0 m/z with the expected isotopic distribution) and $\text{BrO}_3^\bullet$ is detected as BrO_3^- *via* negative-mode MALDI-MS at 126.9 and 128.9 m/z (Fig. 4a inset). OAT from extruded $\text{BrO}_3^\bullet$ to $[\text{Cu}(\text{tpa})]^+$ would be accompanied by the evolution of BrO_2 ; negative-mode MALDI-MS features at 110.9 and 112.9 m/z correspond to expected isotopic distribution for BrO_2 . Finally, CID-MS data indicates the formation of Cu–O fragment 6^+ (Fig. 4). In addition to mass spectrometry, the formation of BrO_2 is also evidence by *in situ* infrared (IR) spectroscopy: The IR spectrum of Cu bromate $[\text{8}]\text{ClO}_4$ in a KBr pellet displays characteristic bromate stretches at 831, 842, and 852 cm^{-1} . Photolysis of this KBr pellet results in the disappearance of

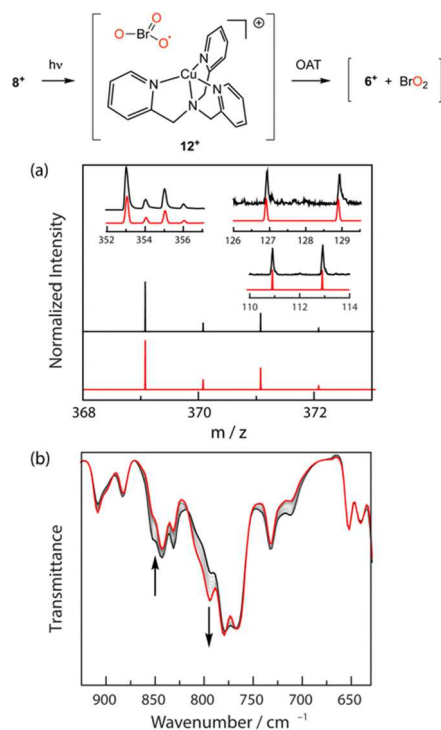


Fig. 4 Photoreduction of 8^+ by Cu–OBrO₂ homolysis followed by OAT from $\text{BrO}_3^\bullet$ to 12^+ would generate 6^+ and BrO_2 . (a) CID-MS analysis shows peaks at 369.1 and 371.1 m/z which correspond to $[6]^+$; MALDI-MS analysis shows peaks at 110.9 and 112.9 m/z which correspond to BrO_2 (black —), simulation (red —) (inset); peaks at 126.9 and 128.9 m/z which correspond to photoextruded BrO_3 (black —), simulation (red —) (inset); and peaks at 353.0 and 355.0 m/z which correspond to $[\text{Cu}(\text{tpa})]^+$ (black —), simulation (red —) (inset). (b) IR spectra collected during the photolysis of a KBr pellet of $[\text{8}]\text{ClO}_4$ shows a new peak at 795 cm^{-1} , which is attributed to BrO_2 .

those spectral features and the evolution of a new peak at 795 cm^{-1} , which is well-matched to a reported stretch for BrO_2 (Fig. 4b).²³ The other stretching mode expected BrO_2 (*i.e.*, 845 cm^{-1}) overlaps with a stretching mode of the $[\text{Cu}(\text{tpa})]^+$ fragment. Diffuse reflectance and X-band EPR spectra obtained after solid-state photolysis are depicted in Fig. S16 and S17 (ESI[†]).

Attempts to characterize the photochemistry of $[\text{8}]\text{ClO}_4$ by low-temperature solution-phase spectroscopy have been stymied by a combination of insolubility in, or reaction with, common glassy solvents. To avoid these challenges and to build on emerging *in crystallo* photochemical strategies to characterize reactive species relevant to C–H functionalization, we were attracted to the potential to apply *in crystallo* photochemistry to directly visualize the primary photochemical events relevant to the OAT chemistry described above. We reasoned that photo-oxidation *via* BrO_2 loss would be differentiable from initial photoreduction *via* $\text{BrO}_3^\bullet$ elimination. To these ends, X-ray diffraction data was collected during 365 nm irradiation of a single crystal of $[\text{8}]\text{ClO}_4$ at 100 K using 30 keV synchrotron radiation. Solid-state reaction progress was monitored by free refinement of the Cu–OBrO₂ fragment.

Refinement of the X-ray diffraction data obtained following *in crystallo* photolysis of $[\text{8}]\text{ClO}_4$ are consistent with initial activation of the Cu–O bond (Fig. 5, refinement details collected in Table S3, ESI[†]). The most significant structural changes induced by photolysis are Cu–O elongation from 1.955(3) Å to 2.22(1) Å and Br–O contraction from 1.657(4) Å (avg.) to 1.53(1) Å (avg.). Accompanying Cu–O elongation, the Cu1–O1–Br1 angle opens from 124.28(2)° to 118.9(5)°. The average O–Br–O angles change minimally from 105.4(2)° to 103.6(8)°. In contrast to the significant elongation of the Cu–O vector, the Cu–N bonds are not significantly different following photolysis (Fig. 5a). Based on comparison to the metrics of $\text{Cu}(\text{tpa})\text{Cl}$,²⁴ *in crystallo* photoreduction of $[\text{8}]\text{ClO}_4$ to generate $\text{BrO}_3^\bullet$ and Cu(I) would be anticipated to elongate the Cu–N(4) vector significantly (0.4 Å in $[\text{Cu}(\text{tpa})\text{Cl}]^+$). Furthermore, the optimized geometry of $[\text{Cu}(\text{tpa})]^+$, featuring no fifth ligand as would be expected for *in crystallo* photoreduction, features a Cu–N(4) bond length of 2.2 Å (Table S4, ESI[†]). As such, the metrics expected for Cu(I) are inconsistent with those observed

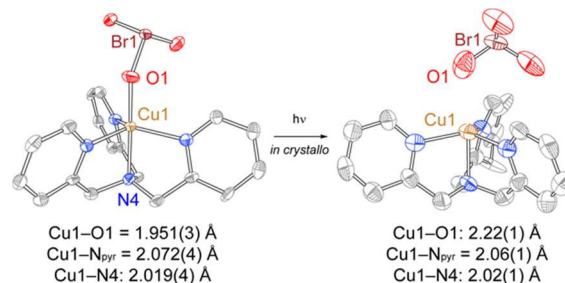


Fig. 5 Displacement ellipsoid plots generated by *in crystallo* photochemistry of $[\text{8}]\text{ClO}_4$. Ellipsoids of $[\text{8}]\text{ClO}_4$ are drawn at 50% probability; following photolysis ellipsoids are drawn at 30% probability. H atoms and counter anions are removed for clarity.

following *in crystallo* photolysis. Instead, the observed structural parameters are more consistent with assignment of the *in crystallo* product of Cu(II)^{2+} and BrO_3^- , which could arise from Cu–O homolysis followed by electron transfer (ET) from the initially formed Cu(I) to generate Cu(II). Rapid *in crystallo* ET may be driven by the dense crystal packing of $[\text{8}]\text{ClO}_4$. Unlike previous *in crystallo* photoreactions in which gaseous, closed-shell small molecule leaving groups (e.g. N_2 and CO) are extruded from the molecular photoprecursors,^{25–29} *in crystallo* Cu–O cleavage would generate a solid, reactive leaving group ($\text{BrO}_3^\bullet$) that is held in proximity to the Cu(I) by virtue of crystal packing (Fig. S18, ESI†).

Finally, we examined the potential that photogenerated $\text{BrO}_3^\bullet$ was the active oxidant in substrate functionalization chemistry (*i.e.* reactions pictured in Fig. 3), we evaluated the photochemistry of a mixture of KBrO_3 and $(\text{NH}_4)_2\text{S}_2\text{O}_8$ in the presence of toluene. Photolysis of a mixture of KBrO_3 and $(\text{NH}_4)_2\text{S}_2\text{O}_8$ has previously been reported to afford $\text{BrO}_3^\bullet$, which was detected by TA spectroscopy.³⁰ In our experiment, we irradiated a mixture of KBrO_3 and $(\text{NH}_4)_2\text{S}_2\text{O}_8$ in acetonitrile with 10 equivalents of toluene added. This reaction afforded a mixture of benzyl alcohol, benzaldehyde, benzoate (Fig. S19, ESI†), which contrasts the selective generation of benzyl alcohol during the photolysis of $[\text{8}]\text{ClO}_4$. $\text{BrO}_3^\bullet$ is known³⁰ to undergo rapid decomposition to BrO and O_2 and the control experiment described here does not differentiate between whether OAT to $[\text{Cu}(\text{tpa})]^+$ is accomplished by $\text{BrO}_3^\bullet$ or reactive oxidants derived thereof. To verify that the observed reactivity is $\text{BrO}_3^\bullet$ based, and not derived from transient intermediates of $(\text{NH}_4)_2\text{S}_2\text{O}_8$ photochemistry, we photolyzed a toluene solution of $(\text{NH}_4)_2\text{S}_2\text{O}_8$ and observed no oxidation products. Control experiments with cyclohexane are described in Fig. S20 (ESI†).

In summary, we report the synthesis and OAT photochemistry of a molecular Cu bromate complex. Photoactivation results in net OAT to CO, olefinic substrates, and C–H bonds to afford CO_2 , epoxides, and alcoholic products. *In crystallo* photochemistry is consistent with initial Cu–O bond cleavage and complementary mass spectrometry and *in situ* spectroscopy experimental suggest that initial Cu–O homolysis (*i.e.*, photoreduction) precedes the formation of a reactive Cu–O species that is responsible for substrate functionalization. These studies add to a growing body of literature on the prevalence of one-electron photochemistry of metal oxanion complexes³¹ and demonstrate new tools to elucidate the atomistic details of molecular photoreactions.

G. P. V. T., D. S., and D. C. P. conceived of the project. G. P. V. T., D. S., and R. R. T. carried out experimental work. All authors participated in data analysis, manuscript writing, and editing.

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Conflicts of interest

There are no conflicts to declare.

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