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# Leveraging ligand-based proton and electron transfer for aerobic reactivity and catalysis

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While O<sub>2</sub> is an abundant, benign, and thermodynamically potent oxidant, it is also kinetically inert. This frequently limits its use in synthetic transformations. Correspondingly, direct aerobic reactivity with O<sub>2</sub> often requires comparatively harsh or forcing conditions to overcome this kinetic barrier. Forcing conditions limit product selectivity and can lead to over oxidation. Alternatively, O<sub>2</sub> can be activated by a catalyst to facilitate oxidative reactivity, and there are a variety of sophisticated examples where transition metal catalysts facilitate aerobic reactivity. Many efforts have focused on using metal–ligand cooperativity to facilitate the movement of protons and electrons for O<sub>2</sub> activation. This approach is inspired by enzyme active sites, which frequently use the secondary sphere to facilitate both the activation of O<sub>2</sub> and the oxidation of substrates. However, there has only recently been a focus on harnessing metal–ligand cooperativity for aerobic reactivity and, especially, catalysis. This perspective will discuss recent efforts to channel metal–ligand cooperativity for the activation of O<sub>2</sub>, the generation and stabilization of reactive metal–oxygen intermediates, and oxidative reactivity and catalysis. While significant progress has been made in this area, there are still challenges to overcome and opportunities for the development of efficient catalysts which leverage this biomimetic strategy.

## 1 Introduction

O<sub>2</sub> is an abundant, inexpensive, and thermodynamically powerful oxidant. This thermodynamic potency is perhaps most simply illustrated by the formal electrochemical potential of 1.23 V vs. SHE for O<sub>2</sub> reduction.<sup>1</sup> Importantly, this net reduction of O<sub>2</sub> results in water as a benign byproduct. Despite these advantages, O<sub>2</sub> is comparatively kinetically inert due to its triplet ground spin state. This triplet ground state leads to inhibited reactivity with many compounds, such as most organic substrates that typically have singlet spin states.<sup>2</sup> This kinetic inertness is also illustrated by the bond dissociation free energy (BDFE) of ~51 kcal mol<sup>−1</sup> in solvent for the first proton/electron in the reduction of O<sub>2</sub> to water.<sup>3</sup> To overcome these spin-forbidden processes, and to participate in oxidative reactivity, O<sub>2</sub> must enter the higher energy singlet spin state.<sup>2</sup> This means that direct aerobic reactivity often requires forcing conditions, such as the high temperatures used for combustion, or activation *via* a catalyst. Transition metal complexes feature prominently in aerobic catalysis, which has led to a wide array of elegant examples in which O<sub>2</sub> is activated for catalytic transformations. The Wacker oxidation, which was developed in 1956, is one classic example that uses co-catalytic Pd and Cu

to perform the oxidation of ethylene to acetaldehyde with O<sub>2</sub> as a terminal oxidant.<sup>4–6</sup> Cu is essential for this process as it helps with mediating efficient redox exchange between Pd and O<sub>2</sub>. Alternatively, a supporting ligand can be similarly used to facilitate proton/electron transfer and subsequent O<sub>2</sub> activation. For example, Stahl and coworkers have investigated the utility of organic cocatalysts such as aminoxyl or quinone compounds that work in conjunction with metal salts to facilitate aerobic oxidation chemistry.<sup>7–16</sup> There has also been extensive research into metal-free aerobic reactivity, primarily using aminoxyl compounds.<sup>17–25</sup>

While there have been many successes in this area, it is noteworthy that Nature activates O<sub>2</sub> through elaborate active sites that coordinate transition metal centers with a carefully positioned organic secondary coordination sphere. Secondary coordination spheres in active sites for aerobic catalysis nearly always feature themes such as hydrogen bonding motifs, redox-active cofactors for electron transfer, and proton shuttling machinery, all of which aid in O<sub>2</sub> activation.<sup>26–33</sup> In many cases this activation can lead to more thermodynamically potent oxidants, such as terminal oxo complexes, through the controlled addition of H-atom equivalents. Thus, the elaborate “ligand” of enzymatic secondary coordination spheres is critical for the activation of O<sub>2</sub> and subsequently efficient and selective aerobic oxidations.

Despite the biological ubiquity of coupling transition metals with tailored secondary coordination spheres to facilitate aerobic reactivity, there has only recently been an allied focus

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on leveraging these same influences in synthetic complexes.<sup>13,34</sup> Major advances include hydrogen bonding ligands,<sup>35–38</sup> proton shuttling ligands,<sup>39–46</sup> and redox-active ligands for cooperative electron transfer.<sup>47–60</sup> While these approaches are now broadly used in the design of new synthetic catalysts, their application for O<sub>2</sub> activation, particularly aerobic catalysis, is still nascent. Supporting ligand scaffolds that can cooperatively transfer both protons and electrons to activate O<sub>2</sub> for aerobic reactivity in a biomimetic manner are even more rare.

This perspective will cover recent advances in leveraging metal–ligand cooperativity for O<sub>2</sub> activation and aerobic reactivity (Fig. 1). We will particularly focus on systems that can shuttle both protons and electrons to O<sub>2</sub>, beginning with examples where this cooperativity enables the characterization of well-defined intermediates which feature O<sub>2</sub> binding or reduction. We will then focus on examples which display stoichiometric or catalytic oxidations. Last, we will summarize prospects and challenges, laying out key directions for development and investigation as this field develops. One clear conclusion is that this area holds promise for developing sustainable oxidation reactions, with exciting developments likely coming in the near future.

## 2 Metal–ligand cooperativity enabling the activation or reduction of O<sub>2</sub>

Metal–ligand cooperativity involving ligand-stored protons and electrons has been shown to facilitate the formation and subsequent reduction of various metal–oxygen species such as metal-oxos and metal hydroperoxos from O<sub>2</sub>. Again, this is a common mode of reactivity and O<sub>2</sub> activation in biology, but it has also become a useful approach in synthetic systems. Quinoidal motifs are some of the classic building blocks used in redox non-innocent ligands, and ligands with incorporated

quinones/quinols can further provide an accessible proton and electron source for O<sub>2</sub> reduction. Agapie and coworkers effectively used such a quinone based ligand scaffold with Pd to facilitate O<sub>2</sub> reduction to water (Fig. 2A). The diphosphine hydroquinone ligand is able to donate 2 electrons and 2 protons towards O<sub>2</sub> reduction. When a Pd atom is coordinated to the ligand, O<sub>2</sub> reduction is found to proceed *via* a side-on  $\eta^2$ -peroxo complex, the formation of which can be observed by low temperature ultraviolet-visible (UV-vis) spectroscopy. Low temperature <sup>31</sup>P nuclear magnetic resonance (NMR) spectroscopy of this species supports the formation of an asymmetric complex, and low temperature infra-red (IR) spectroscopy shows no indication of a C=O double bond stretch, indicating that the side on Pd  $\eta^2$ -peroxo complex forms prior to the involvement of the backbone quinone. While this first intermediate observed by UV-vis spectroscopy is long-lived at low temperature, the second intermediate generated upon warming decays more rapidly. Spectroscopic data of this second species suggests a Pd<sup>2+</sup> complex coordinated to a semiquinone ligand, but concrete assignment of the partially hydrogenated oxygen ligand was not obtained due to the transient nature of this intermediate. The reduction of O<sub>2</sub> to H<sub>2</sub>O as a final product was confirmed by <sup>1</sup>H NMR spectroscopic analysis of the reaction volatiles after transferring to a J. Young tube.<sup>61</sup>

Agapie and coworkers have also illustrated the utility of a related isomeric quinone-based ligand scaffold on Mo complexes for a similar reduction of O<sub>2</sub> to water (Fig. 2A). In this case it was found that activation of O<sub>2</sub> led to initial reduction to H<sub>2</sub>O<sub>2</sub> rather than H<sub>2</sub>O. H<sub>2</sub>O<sub>2</sub> then putatively interacts with a second equivalent of the Mo complex for the net 4e<sup>−</sup> reduction to H<sub>2</sub>O to occur. This illustrates how both the ligand and the metal can influence the selectivity of reductions, even in closely related systems. As a final note, while this perspective focuses on the reduction of O<sub>2</sub> with protons and electrons, Mo complexes supported by this general ligand framework, but with Si- or B-functionalized catecholate motifs, make (R<sub>2</sub>SiO)<sub>n</sub> and (R<sub>2</sub>BO)<sub>n</sub> byproducts rather than H<sub>2</sub>O as the final reaction product.<sup>62</sup> Regardless of the terminal O-accepting reagent, all of these examples illustrate how redox-active quinone-based ligands can facilitate selective reduction of O<sub>2</sub> to H<sub>2</sub>O.

While the above examples rely on the incorporation of quinone/quinol moieties directly into the ligand backbone, the use of a pendent quinol on a ligand arm has also been shown to facilitate the reduction and/or activation of O<sub>2</sub>. In particular, Goldsmith and coworkers have shown that appending such a quinol group to a chelating amine/pyridine scaffold facilitates superoxide dismutase (SOD) like reactivity with either Mn<sup>2+</sup> or Zn<sup>2+</sup>.<sup>63,64</sup> Mn<sup>2+</sup> complexes with a ligand featuring one pendent phenol mediate SOD reactivity *via* an outer-sphere mechanism. However, when the ligand features one or two pendent quinols, the reaction accelerates *via* an inner-sphere mechanism.<sup>63</sup> This quinol assisted inner-sphere pathway can also be leveraged to render a redox-inactive metal ion, Zn<sup>2+</sup>, active for SOD catalysis.<sup>64</sup> Beyond this example, Goldsmith and coworkers have also investigated the electrochemical reduction of O<sub>2</sub> using a Co<sup>2+</sup> metal center bound to the same ligand framework (Fig. 2B). They found that when a pendent quinol rather than a pendent

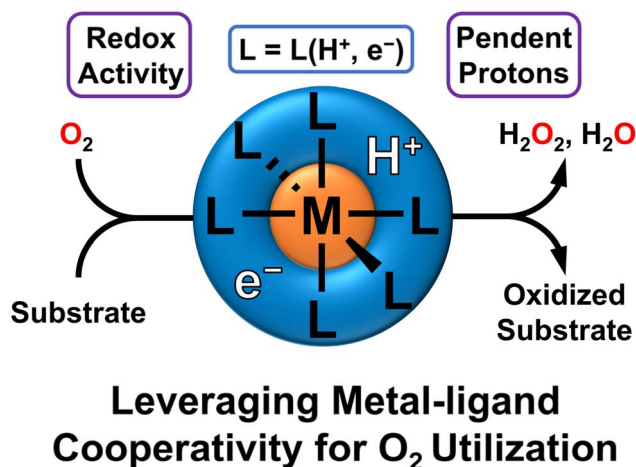


Fig. 1 Metal–ligand cooperativity that incorporates redox-active ligands and pendent H-equivalents in the secondary sphere can activate O<sub>2</sub> for oxidative reactivity with substrates. This enables the ligand to serve as a proton and electron source.



Fig. 2 Metal–ligand cooperative transfer of protons and electrons enabling the activation of  $O_2$ . (A and B) Quinone-based non-innocent ligands that can donate protons and electrons to reduce  $O_2$ . (C) A non-innocent ligand stabilizes metal–oxygen intermediates and acts as a proton shuttle to reduce  $O_2$  to water in the presence of external reductant. (D) Protons and electrons from a non-innocent ligand produce a spectroscopically characterizable Fe–hydroperoxo complex.

phenol was incorporated into the ligand scaffold, the product selectivity and mechanism shifts from a 2 electron, 2 proton reduction of  $O_2$  to form  $H_2O_2$  to a 4 electron, 2 proton reduction of  $O_2$  to form  $H_2O$ .<sup>65</sup> This switch demonstrates that inclusion of groups that donate both electrons and protons not only enables catalytic reactivity, but also provides determinative control over selectivity in some processes. This conclusion meshes with other observations in biological and synthetic systems. As one example, Machan and coworkers and Stahl and coworkers have both shown the utility of 1,4-hydroquinone/benzoquinone as a cocatalyst in facilitating the reduction of  $O_2$  to  $H_2O$  electrocatalytically.<sup>9,66</sup> While the 1,4-hydroquinone/benzoquinone group is not directly incorporated into the ligand backbone in these cases, the electrocatalytic reaction mechanism requires the close association of 1,4-hydroquinone/benzoquinone to produce  $H_2O$  as the final product. As with the Co example above, omission of 1,4-hydroquinone as a cocatalyst in this example from Stahl and coworkers<sup>9</sup> instead results in the preferential reduction of  $O_2$  to  $H_2O_2$ .

An alternate strategy for metal–ligand cooperativity is the use of the ligand to create a secondary sphere that stabilizes reactive oxygenated intermediates through hydrogen bonding interactions. The use of H-bonding or proton shuttling ligands has been of great recent interest.<sup>67</sup> While these scaffolds do not strictly shuttle electrons, there are several ligand scaffolds that feature conjugated arms that may potentially also be redox-active. As such, some select cases are also highlighted here. As one example among many,<sup>35–38,68,69</sup> Borovik and coworkers demonstrated how designing a secondary sphere to stabilize intermediates and shuttle proton equivalents enables the

catalytic reduction of  $O_2$ . They were able to synthesize a side on Mn-peroxo species and Mn-oxo species stabilized by hydrogen bonding interactions (Fig. 2C). Characterization of these intermediates proved crucial to the development of a catalytic cycle converting  $O_2$  to  $H_2O$  at room temperature.<sup>70</sup> Catalysis was only observed with the addition of hydrazine or diphenylhydrazine as a terminal reductant. Fout and coworkers utilized a similar strategy to stabilize  $Fe^{3+}$ -oxo or  $Fe^{3+}$ -hydroxo complexes formed *via* reactivity between an Fe complex and  $O_2$ .<sup>35</sup> Both complexes are stabilized by hydrogen bonding interactions from the ligand, and the ligand shuttles protons to the  $O_2$  substrate.<sup>67</sup> Szymczak and coworkers have also used hydrogen bonding motifs to stabilize a Zn-peroxo dimer formed from reactivity with  $O_2$  in the presence of cobaltocene. However, we note that neither protons nor electrons are shuttled by the ligand for  $O_2$  reduction in this case.<sup>69</sup> The fact that no H-equivalents are transferred to  $O_2$  makes the formation of the Zn-peroxo dimer reversible with added oxidant, such as ceric ammonium nitrate or iodobenzene dichloride. Another related example comes from Garcia-Bosch and coworkers. They recently reported a Cu complex that uses hydrogen bonding interactions from the ligand to stabilize and reduce  $O_2$  to an axial hydroperoxo ligand. However, in this case, the terminal H-atom on the hydroperoxo ligand is abstracted from the ligand of a second Cu complex rather than the oxidized Cu complex itself.<sup>71</sup> In all examples, hydrogen bonding interactions help to stabilize reactive intermediates, with some examples also invoking H-atom abstraction and proton shuttling for catalytic reactivity. While definitive evidence for ligand-based redox reactivity is less obvious in most of these cases, it is possible, if not likely, that



the conjugated arms in these or related scaffolds might also be non-innocent under the right conditions or with appropriate synthetic modifications.

It is noteworthy that while the quinone/quinol systems discussed above formally transfer H-atoms, the H-bonding examples feature N-based ligands which do not as explicitly exhibit net H-atom reactivity. However, there are other ligands with N-H's that more clearly shuttle reducing equivalents to  $O_2$ . David Goldberg and coworkers found that they could use a macrocyclic tetraamine ligand combined with electronic effects from a trans thiolate ligand to stabilize an  $Fe^{3+}$ -hydroperoxo complex (Fig. 2D).<sup>72</sup> Here, the electron and proton equivalents came from the ligand scaffold *via* desaturation to form an imine. This complex can be generated at  $-78\text{ }^\circ\text{C}$ , and was characterized by UV-visible spectroscopy, electron paramagnetic resonance (EPR) spectroscopy, and electrospray ionization mass spectrometry (EI-MS). Using thiolate ligands for stabilizing Fe-oxygen intermediates has been a topic of some interest due to the presence of this motif in enzyme active sites such as superoxide reductase (SOR) or cytochrome P450.<sup>73,74</sup> However, these biomimetic  $H_nO_y$  complexes are rarely generated directly from  $O_2$  since their formation requires the movement of both protons and electrons. Instead, Fe-hydroperoxo complexes are often generated from exogenous reducing and acid equivalents.<sup>75–84</sup> The reactivity observed in this case would be impossible without metal-ligand cooperativity arising from desaturation of the macrocyclic ligand that facilitates the transfer of both electrons and protons.

Similar in structure to quinols, pendent phenol functionalities on the ligand backbone can act as a proton and electron source for the hydrogenation of  $O_2$ . Karlin and coworkers used a Cu system with an aliphatic amine pincer ligand and a pendent phenol to capture  $O_2$  as a bis( $\mu$ -oxo) $Cu_2^{2+}$  dimer at  $-135\text{ }^\circ\text{C}$ .<sup>85</sup> The pendent phenol group on each ligand then donates one proton and one electron to each O atom, resulting in a bis( $\mu$ -hydroxo) $Cu_2^{2+}$  dimer where each Cu center features a ligand-based radical on the O atom of the pendent phenol. These ligand-based radicals can be detected with EPR spectroscopy by a characteristic signal at  $g = 2.00$ . Upon warming in the presence of excess 1-hydroxy-2,2,6,6-tetramethylpiperidine (TEMPOH) as an H-atom source, this dimer releases two equivalents of  $H_2O$  and forms a  $Cu^{2+}$  complex with an overall +1 charge, where the Cu center is now coordinated to the NNN amine pincer ligand as well as the O in the pendent phenol (Fig. 3). This reactivity mimics the reactivity observed in the active site of oxy-tyrosinase, which catalyses the regioselective hydroxylation of monophenols with a *para* substitution to catechols.<sup>86,87</sup>

In each of the examples presented in this section, metal-ligand cooperativity facilitates the shuttling of proton and electron equivalents to  $O_2$  resulting in its reduction. Despite this general reactivity paradigm, the products of these reactions are variable depending on the system, with  $H_2O$ ,  $H_2O_2$ , or stabilized metal-oxygen complexes (*i.e.* hydroperoxo species) that can be spectroscopically characterized all being commonly formed. While these examples demonstrate the utility of incorporating redox active ligands with pendent protons for the

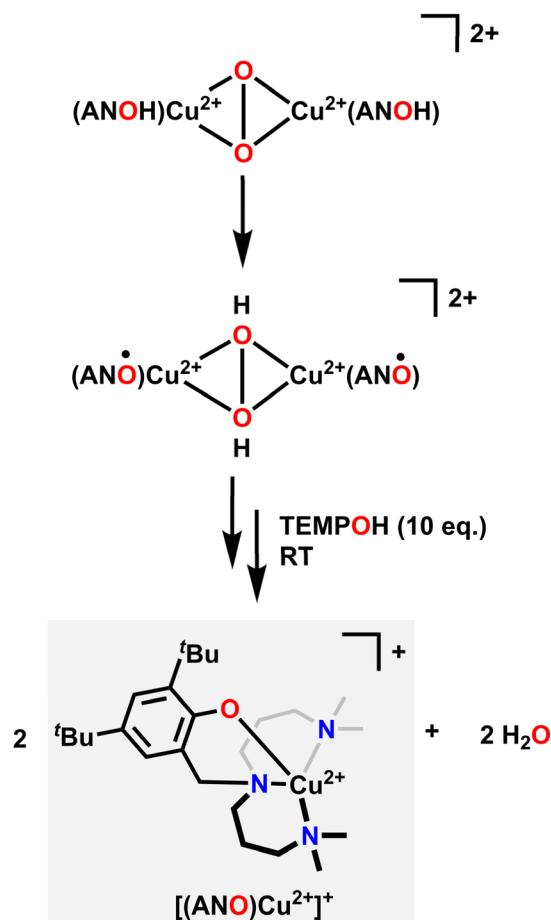


Fig. 3 Mechanism of  $O_2$  capture, hydrogenation, then release of water upon reacting with TEMPOH.

hydrogenation of  $O_2$ , there is still a lack of clarity on what exact design features facilitate the formation of different products. Simple topics of future investigation might reasonably include examining how the positioning of H-donors dictates product selectivity, or how the thermodynamic properties of these donors (BDFE, acidity, *etc.*) can be tuned for desired reactivity. Beyond these fundamental studies, there is also great potential in harnessing the thermodynamic potential of  $O_2$  for subsequent substrate oxidations after activation. Indeed, the next sections of this perspective focus on recent advances in these areas.

### 3 Metal–ligand cooperativity enabling the activation of $O_2$ for alcohol oxidation

Alcohol oxidation is a widely used transformation in synthetic chemistry. Metal-ligand cooperativity can facilitate alcohol oxidation reactions by either forming an oxygenated intermediate that serves as the active oxidant in the key bond forming/breaking steps or as an inexpensive and benign terminal oxidant for the regeneration of an oxidized transition metal intermediate that instead directly oxidizes the substrate, as seen

in Wacker chemistry.<sup>4,5</sup> In both instances, O<sub>2</sub> acts as a formal H-atom acceptor, but there are key mechanistic differences whether an *in situ* oxygenated intermediate acts as the oxidant or whether O<sub>2</sub> simply acts as a regenerating terminal oxidant.

A good example where O<sub>2</sub> generates an active oxygenated transition metal intermediate that performs alcohol oxidation comes from Garcia-Bosch and coworkers (Fig. 4).<sup>88</sup> They use a ureanyl functionalized 1,2-phenylenediamido Cu<sup>2+</sup> platform where the diureanyl ligand can formally store both protons and electrons as well as provide H-bonding interactions. This platform enables the stabilization of a superoxo intermediate *via* H-bonding interactions from the ligand. This transiently stabilized Cu-superoxo then serves as the active oxidant for the oxidation of primary alcohols to aldehydes. The ligand further facilitates the reaction by mediating the movement of proton and electron equivalents during the dehydrogenation of the alcohol and subsequent H-transfer to the coordinated O<sub>2</sub> molecule resulting in both aldehyde and H<sub>2</sub>O<sub>2</sub> as the terminal reaction products. Throughout the course of this reaction mechanism, Cu remains in a 2+ state. This means that the Cu<sup>2+</sup> center does not participate in the alcohol oxidation reaction as an electron source, but rather as a docking site for the primary alcohol/O<sub>2</sub> to coordinate such that the ligand secondary sphere can mediate the resulting oxidation and reduction respectively. Interestingly, this suggests that transition metal centers, which are frequently considered as redox mediators in many catalytic cycles, can instead be tuned/utilized as simple Lewis acid sites for coordination if a suitably redox-active supporting ligand is used.<sup>64,89–91</sup> This is similar to how classic reactions in organic chemistry, such as the Sharpless epoxidation, proceed through use of a catalyst that acts as a Lewis acid.<sup>92</sup> Additionally, the reduction of O<sub>2</sub> to H<sub>2</sub>O<sub>2</sub> allows for the regeneration of the

starting complex and thus allows for a catalytic process. This work is in contrast to the ping-pong mechanism seen in galactose oxidase (GAO), where a primary alcohol is first oxidized to an aldehyde, then O<sub>2</sub> is reduced to H<sub>2</sub>O<sub>2</sub> as the terminal regenerating oxidant to reform the active catalyst. The amino acids within the GAO active site act as proton acceptors, whereas Cu plays a more active role in the catalytic cycle of GAO, by acting as an electron acceptor/donor. This ping-pong mechanism has been more closely reproduced by Stack and coworkers (Fig. 5A).<sup>93</sup> Here the authors were able to synthesize a Cu complex and related intermediates in the catalytic oxidation of primary alcohols and demonstrate pathways that match those proposed for GAO. In their Cu system, alcohol oxidation occurs first with the ligand scaffold acting as an H-atom acceptor, O<sub>2</sub> reduction to H<sub>2</sub>O<sub>2</sub> occurs second, and the Cu<sup>2+</sup> center is reversibly reduced to Cu<sup>1+</sup> during the course of the catalytic cycle.<sup>93</sup> As such, this example from Stack serves as a more faithful biomimetic Cu system for the mechanism observed in GAO where O<sub>2</sub> serves as a regenerating oxidant.

Pioneering work from Wieghardt and coworkers demonstrated that different mononuclear Cu and Zn complexes featuring an ONNO ligand structurally similar the ligand used by Stack and coworkers were also able to facilitate alcohol oxidation of primary and benzylic alcohols using O<sub>2</sub> as an oxidant to regenerate the starting metal complex catalytically and release H<sub>2</sub>O<sub>2</sub> (Fig. 5B).<sup>94,95</sup> Mechanistic and kinetic studies of their system showed evidence for H-atom abstraction by the ligand from the coordinated alcohol, providing evidence for ligand noninnocence and metal–ligand cooperativity during the alcohol oxidation reaction.

Another early example of a Cu complex facilitating alcohol oxidation in conjunction with O<sub>2</sub> reduction comes from Urch,



Fig. 4 Mechanism of alcohol oxidation where O<sub>2</sub> participates in the mechanism in conjunction with ligand non-innocence. O<sub>2</sub> generates the Cu-superoxo that enables alcohol oxidation before being reduced *via* metal–ligand cooperativity.





Fig. 5 (A) Proposed mechanism of alcohol oxidation reported by Stack and coworkers closely mimicking the ping-pong mechanism observed in GAO. O<sub>2</sub> participates in the mechanism in conjunction with ligand non-innocence. O<sub>2</sub> generates a Cu-hydroperoxo and displaces the oxidized substrate, before being reduced via metal-ligand cooperativity to H<sub>2</sub>O<sub>2</sub>. (B) Proposed mechanism of primary alcohol oxidation reported by Wieghardt and coworkers. In this case O<sub>2</sub> serves as a terminal oxidant, allowing the starting metal catalyst to regenerate.

Brown, and coworkers.<sup>96,97</sup> In their system, CuCl was mixed with phenanthroline (phen), a substituted azo compound, such as di-*tert*-butylazodicarboxylate (DBAD), an alcohol, O<sub>2</sub>, and K<sub>2</sub>CO<sub>3</sub>, to form a Cu<sup>1+</sup> complex that facilitates alcohol oxidation *via* hydrogen transfer from the coordinated alcohol to the coordinated azo ligand. Upon release of the aldehyde or ketone, O<sub>2</sub> then coordinates to form a Cu<sup>2+</sup> dimer with a bridging peroxo ligand. The peroxo ligand is then homolytically cleaved when heated to putatively form a Cu<sup>2+</sup>-oxyl complex. This then abstracts an H-atom from both the substituted hydrazine ligand and the alcohol substrate to release H<sub>2</sub>O and restart the catalytic cycle. Interestingly, although this is one of the earliest examples invoking metal-ligand cooperativity for alcohol oxidation with a Cu complex where O<sub>2</sub> is used as a terminal oxidant, it is also one of the examples with the widest substrate scope. Urch, Brown, and coworkers found that this system is able to catalytically oxidize primary, allylic, and benzylic alcohols.<sup>97</sup>

While there are entropic and orientational advantages to having a tethered ligand for H-shuttling, these scaffolds can frequently be difficult to synthesize or tune, which limits their application in organic methodology. An alternative strategy that has been employed to great effect is the use of metal complexes in conjunction with an exogenous organic cocatalyst that acts as a proton/electron reservoir. Stahl and coworkers have been particularly active in this area with extensive applications of Cu bipyridine (bpy) complexes with (2,2,6,6-tetramethylpiperidin-1-yl)oxidanyl (TEMPO) as a cocatalyst for alcohol oxidation

(Fig. 6A).<sup>98</sup> They find that half an equivalent of O<sub>2</sub> and one equivalent of TEMPO are necessary to generate a Cu<sup>2+</sup> hydroxide complex which can then bind alcohol. TEMPO then serves as an H-atom acceptor in tandem with the Cu<sup>2+</sup> center serving as a one e<sup>-</sup> oxidant to regenerate the starting catalyst and generate the aldehyde product. Other aminoxyl co-catalysts are also competent for this reactivity; when TEMPO is switched out for 9-azabicyclo[3.3.1]nonane N-oxyl (ABNO), the oxidation of both primary and secondary alcohols becomes accessible.<sup>99</sup> The transition metal used can also be changed; Stahl and coworkers have also had success using Fe(NO<sub>x</sub>) catalysts instead of Cu(bpy) in conjunction with aminoxyl compounds.<sup>100</sup>

While both Cu and Fe are frequently used for oxidative chemistry with O<sub>2</sub>, particularly in biological or biomimetic contexts, other first row transition metals with redox active ligands and pendent protons can also use O<sub>2</sub> for alcohol oxidation. Many of these examples again leverage metal-ligand cooperativity. As one example, the Paul group used 2-(4-chlorophenylazo)-1,10-phenanthroline ligand with Co<sup>2+</sup> to catalyze alcohol oxidation with benzyl alcohols (Fig. 6B).<sup>101</sup> The proposed reaction mechanism begins by coordinating the alcohol and releasing HCl. The catalytic cycle then proceeds *via* β-hydride elimination from the bound alcohol to release an aldehyde or a ketone while simultaneously transferring the hydride onto Co<sup>2+</sup> to form a transient Co hydride intermediate. The hydride ligand can then transfer onto a diazene fragment of the supporting ligand before a new alcohol equivalent coordinates to Co<sup>2+</sup> as an alkoxide, with the ligand

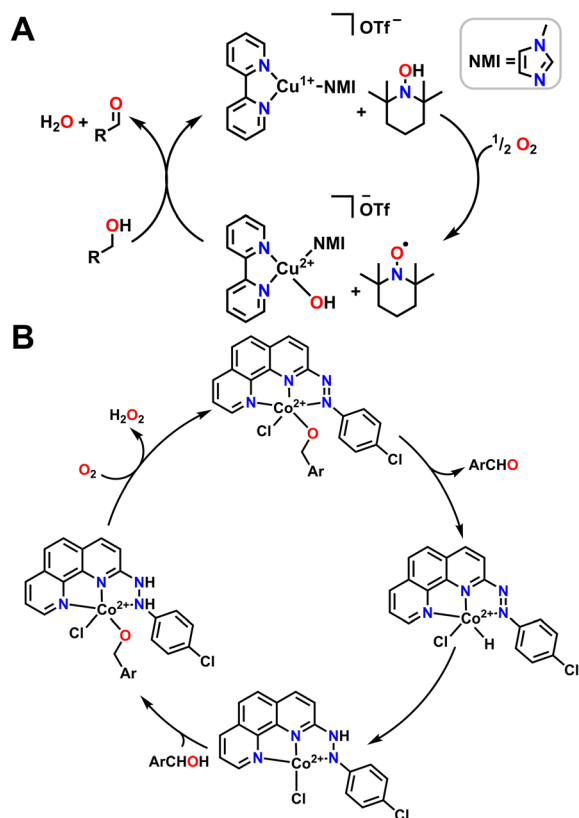


Fig. 6 Mechanisms of alcohol oxidation where O<sub>2</sub> participates in the mechanism in conjunction with ligand non-innocence. (A) O<sub>2</sub> is reduced with a Cu complex and external organic compound that serves as a proton and electron source to regenerate the active molecule for alcohol oxidation. NMI coordinates through the less sterically hindered N atom. (B) O<sub>2</sub> regenerates starting metal complex.

accepting a proton. An O<sub>2</sub> molecule then oxidizes the ligand, removing a formal H<sub>2</sub> equivalent and releasing H<sub>2</sub>O<sub>2</sub> to reform the starting complex in the catalytic cycle (Fig. 6B).

The examples discussed in this section offer some general mechanistic paradigms for catalytic systems featuring both metal–ligand cooperativity and O<sub>2</sub>. Generally, O<sub>2</sub> is either used in the generation of the active oxidant, or it is used to regenerate the active catalyst. In both cases, hydrogenation of O<sub>2</sub> occurs in addition to the targeted alcohol oxidation as part of the catalytic cycle. The production of H<sub>2</sub>O or H<sub>2</sub>O<sub>2</sub> can result in its own challenges since these molecules may participate in side-reactions or competitive coordination, particularly as their relative concentration increases. This is a challenge that will need to be addressed in any future large scale catalytic process, particularly in examples which generate H<sub>2</sub>O<sub>2</sub>.

## 4 Metal–ligand cooperativity enabling the activation of O<sub>2</sub> for O-atom transfer and other dehydrogenation reactions

Beyond using O<sub>2</sub> as a reagent to drive overall oxidative reactivity, metal–ligand cooperativity can also facilitate using O<sub>2</sub> as an O-

atom transfer reagent. This typically occurs by first forming an oxygenated metal complex such as metal-superoxo, -hydroperoxo, or -oxo species which then acts as the active O-atom transfer reagent. Several examples where O<sub>2</sub> can be used for net O-atom transfer have come from our laboratory using a dihydrazonopyrrole (DHP) based ligand scaffold and various first row transition metals. This ligand scaffold has the ability to reversibly donate or accept two protons and two electrons, while additional electron equivalents can be accessed from the metal center itself.<sup>102,103</sup> This unique redox flexibility directly aids in the activation of O<sub>2</sub>, in many cases through spectroscopically observable intermediates. Metalation with Fe<sup>2+</sup> enables the isolation and characterization of a reduced and protonated DHP complex with a formal H<sub>2</sub> equivalent stored on the ligand backbone. Assignment of this complex is supported by suite of spectroscopic techniques, including single crystal X-ray diffraction (SXRD) and IR spectroscopy, both of which support the presence of ligand-based N–H protons. In the presence of O<sub>2</sub>, this complex reacts to form a transiently stable intermediate which can be assigned as an Fe<sup>3+</sup>-hydroperoxo complex with a ligand based DHP radical (Fig. 7A).<sup>104</sup> This reaction can be followed by UV-vis spectroscopy at –40 °C. Isotopic labelling studies with <sup>18</sup>O<sub>2</sub> show an O–O stretch by Raman spectroscopy that provides good support for the Fe<sup>3+</sup>-hydroperoxo assignment. Additionally, IR spectroscopy of deuterium labelled N–H complex followed by reaction with O<sub>2</sub> shows that the signals associated with the N–H and O–H stretches in the Fe<sup>3+</sup>-hydroperoxo complex originate from the ligand scaffold and hydroperoxo moiety respectively. This Fe complex can then release H<sub>2</sub>O<sub>2</sub> upon warming. Alternatively, if this complex is instead warmed in the presence of triphenylphosphine (PPh<sub>3</sub>) one equivalent of triphenylphosphine oxide (OPPh<sub>3</sub>) can form. It was also found that the Fe<sup>3+</sup>-hydroperoxo complex will perform a net H-atom abstraction followed by O-atom transfer with the toluene solvent to form benzaldehyde. While only 0.131 equivalents of benzaldehyde could be detected by gas chromatography-mass spectrometry (GC-MS), this reaction demonstrates how the combination of redox-active ligands and pendent protons can enable the reductive activation of O<sub>2</sub> for both C–H activation as well as O-atom transfer.

A second example of O-atom transfer *via* metal–ligand cooperativity with these DHP systems features a Ni<sup>2+</sup> T-shaped complex with a ligand radical (Fig. 7B).<sup>105</sup> Exposure of this complex to O<sub>2</sub> results in the formation of a Ni<sup>2+</sup>-superoxo complex with an oxidized DHP ligand. In this case, oxidation occurs at the ligand rather than at the metal center upon O<sub>2</sub> binding, paralleling the reactivity proposed in quercetin dioxygenase.<sup>106,107</sup> Unlike the above-mentioned Fe<sup>3+</sup>-hydroperoxo complex with a DHP ligand radical, the Ni<sup>2+</sup>-superoxo complex utilized ligand-based electron transfer only, instead of both proton and electron transfer. This superoxo intermediate is moderately stable at room temperature. While this Ni<sup>2+</sup>-superoxo complex can perform stoichiometric O-atom transfer to PPh<sub>3</sub>, it must be heated to elicit H-atom abstraction from a C–H bond and subsequent O-atom transfer to toluene. The Ni<sup>2+</sup>-superoxo complex can also perform alcohol oxidation of benzyl alcohol, but once again requires heat for reactivity to occur. This



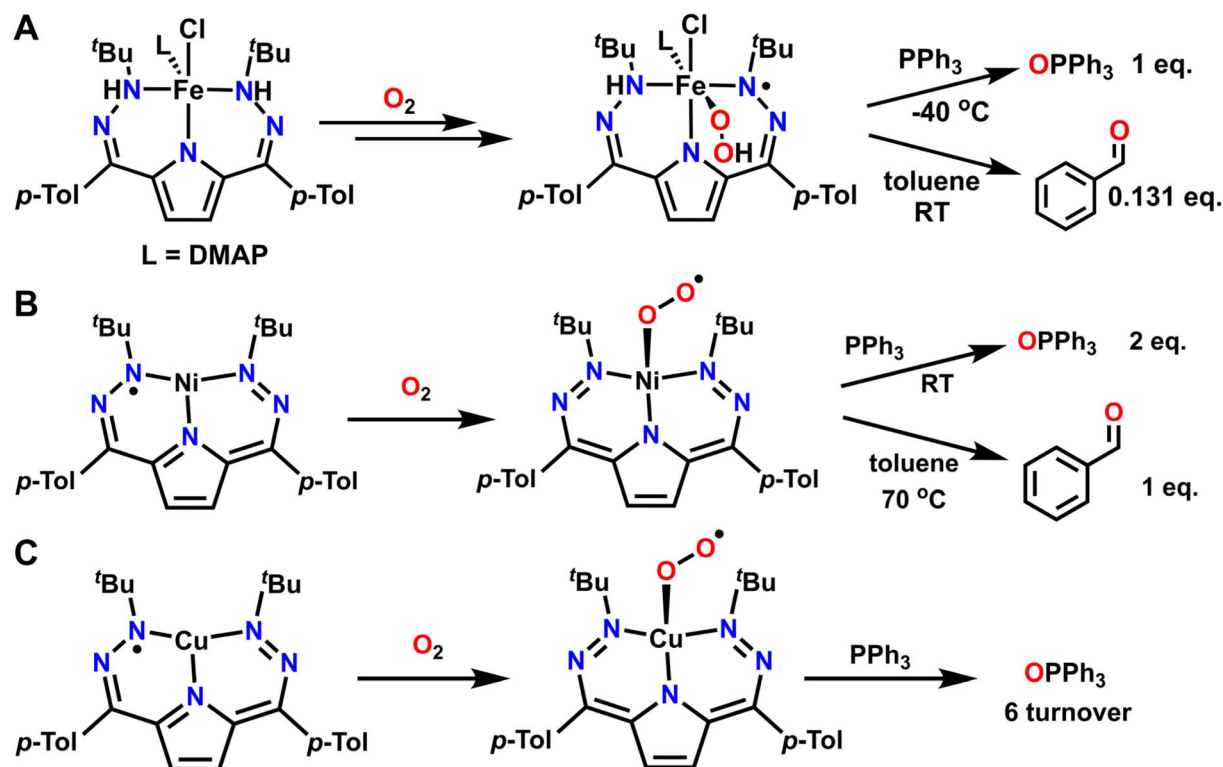


Fig. 7 Exposure to  $\text{O}_2$  forms metal–oxygen complexes that can then facilitate O-atom transfer reactions to phosphines and C–H bonds via metal ligand cooperativity. (A) Reactivity of a fully reduced and hydrogenated Fe(DHP)– $\text{H}_2$  complex. (B) Formation and reactivity of a Ni(DHP)–superoxo complex. (C) Formation and reactivity of a Cu(DHP)–superoxo complex.

is a stark contrast to the previously discussed Fe complex which could perform these reactions at low temperature or room temperature respectively. Unfortunately, the exact mechanism of oxidation for both the Fe and the Ni complexes is difficult to conclusively determine due to decomposition of the metal complex during reactivity.

The above examples with the DHP scaffold on Fe and Ni resulted in only stoichiometric reactivity. However, moving late in the first-row to Cu results in aerobic catalysis. A related  $\text{Cu}^{2+}$ -superoxo complex with an oxidized and closed-shell DHP ligand can be synthesized and characterized with a suite of spectroscopic methods (Fig. 7C). This superoxo complex engages in a variety of catalytic aerobic reactivity including deformylation of 2-phenylpropionaldehyde, dehydrogenation of N–H and O–H bonds, and O-atom transfer to  $\text{PPh}_3$ .<sup>108</sup> As such, this Cu complex displays the most robust and versatile oxidative reactivity among these DHP ligated complexes. All three examples demonstrate how metal–ligand cooperativity can stabilize metal–oxygen intermediates that serve as potent oxidants. However, only the Fe(DHP) complex has provided direct evidence, *via* spectroscopy, of how ligand-based hydrogen equivalents can be transferred to  $\text{O}_2$ . Other first row transition metals featuring the DHP ligand, including Ni and Co, have been shown to participate in hydrogenation where a more direct role for the storage of H-equivalents on the ligand is invoked.<sup>103,109</sup> An interesting hypothesis is that the DHP ligand plays an active role in accepting protons or H-atoms during the

C–H, N–H, or O–H activation step of various oxidations, but additional studies will be required to test this proposal. Regardless, a key conclusion from this series is how different oxidative reactivity is realized with different metal centers all on the same DHP ligand. This highlights how cooperativity between the metal and the ligand is essential beyond the fundamental design of the ligand scaffold itself.

While O-atom transfer to an outside substrate can be extremely challenging, metal–ligand cooperativity can more commonly facilitate intramolecular O-atom transfer, for instance in degradation processes. O-atom transfer to the ligand is often more accessible as the ligand as a substrate is held in close proximity to the activated O-atom equivalents. Unfortunately, it also provides a parasitic pathway that frequently precludes the possibility of performing catalytic oxidations of external substrates. These examples still provide insights into how metal–ligand cooperativity enables the activation of  $\text{O}_2$  for oxidative chemistry. As such, a few examples of ligand oxidation are described here. As a final note, while initial efforts in this area have focused on designing ligands for controlling the storage/flow of electrons and protons, realizing scaffolds which are oxidatively robust is an increasingly important hurdle for broader catalytic applications (as discussed below).

A few examples of intramolecular O-atom transfer have been demonstrated by Karen Goldberg and coworkers. Here, the authors have explored the reactivity of Pt complexes featuring



redox active ligands with  $O_2$  and they have observed oxidation of the ligand *via* C–O bond formation to one O-atom and coordination to the Pt center through the second O-atom to form a 6-member metallocycle ring (Fig. 8A). This Pt–peroxide complex then undergoes O–O bond cleavage to form a 5-membered metallocycle ring with one O-atom and insertion of the other O-atom into a ligand C–H bond. As such, the ligand of this complex stabilizes the coordination of  $O_2$  and subsequently engages in O-atom transfer to the ligand *via* insertion into a C–H bond.<sup>110,111</sup> Net C–H activation of the ligand has been previously observed when reactive metal–oxygen complexes are generated.<sup>112,113</sup> However, we will not discuss these other examples in detail since the reactive complex in question was not generated directly from  $O_2$ .

Another example of  $O_2$  activation and subsequent reactivity with the ligand comes from Milstein and coworkers. Reaction of an  $Fe^{2+}$  hydride complex supported by a PCP pincer ligand results in hydrogenation of one oxygen equivalent of  $O_2$  to  $H_2O$  and insertion of the second oxygen into the Fe–C ligand bond (Fig. 8B). A proton from the ligand backbone O–H and the hydride ligand are the hydrogen sources for this net hydrogenation reaction to form water. This results in an overall oxidized PCP ligand and one equivalent of  $H_2O$  from the net reaction with  $O_2$ .<sup>114</sup> This reactivity is reminiscent of extradiol-cleaving catechol dioxygenases, which use the substrate as the ligand when activating  $O_2$ . In particular, the mechanism for these dioxygenases invokes an Fe–oxygen–ligand metallocycle prior to O-atom insertion into the ligand and the subsequent reduction of the second O-atom to water.<sup>115</sup> A similar O-atom insertion

into a ligand C–H bond and oxidation of the auxiliary ligand, in this case trimethylphosphine ( $PMe_3$ ) to trimethylphosphine oxide ( $OPMe_3$ ), is also observed with a DHP Ni complex related to those discussed above.<sup>116</sup> Abakumov and coworkers have also reported an  $Sb^{5+}$  complex that can coordinate  $O_2$ , then undergo insertion of the terminal O-atom into the ligand to form a metallocycle and dearomatize the ligand scaffold. This reaction is reversible, and under moderate heating will reform the original  $Sb^{5+}$  complex and  $O_2$ .<sup>117,118</sup>

In both the Pt and Fe examples,  $O_2$  is first coordinated to the metal center before proceeding through O-atom transfer to the ligand and hydrogenation. However, metal–ligand cooperativity can facilitate other  $O_2$  binding events, particularly through interactions with ligand-based radicals. A good example of this interaction can be observed with a  $Ru^{2+}$  complex reported by Lahiri and coworkers. Here,  $O_2$  initially reacts with a ligand-based radical to form a C–O–O bond, where a radical primarily resides on the terminal oxygen and effectively dearomatizes the ligand (Fig. 8C). This complex then undergoes a net 1,3-H-atom shift to rearomatize the ligand and provide a hydroperoxo intermediate. Finally, O–O bond homolysis results in the release of hydroxyl radical and overall O-atom transfer to the ligand. Importantly, this net reactivity is again enabled by metal ligand cooperativity between the  $Ru^{2+}$  center and the pincer supporting ligand.<sup>119</sup>

## 5 Opportunities and challenges

Metal–ligand cooperativity, particularly with a scaffold capable of storing both electron and proton equivalents, is a powerful



Fig. 8 Examples where exposure to  $O_2$  results in O-atom transfer to the ligand *via* metal–ligand cooperativity where the ligand donates protons and electrons for the reduction of  $O_2$ . (A), ref. 110; (B), ref. 114; (C), ref. 119.



Alternatively, future studies may find success in utilizing more robust and modular “cooperative” ligands that can donate and accept both protons and electrons to enable flexibility and broad scope that are needed for useful synthetic methodologies.

#### 5.4 Building in sufficient oxidative stability to enable efficient turnover

Some of the most challenging aerobic oxidations also involve substrates which are similar to ligand scaffolds (*i.e.* aliphatic C–H groups). Thus, designing appropriate flanking groups on ligands is important to preclude intramolecular decomposition. Examples of this intramolecular decomposition were presented in the final section of this perspective. While those examples provided important evidence for how metal–ligand cooperativity can enable O-atom transfer to C–H bonds, they also demonstrated how intramolecular oxidation of the ligand scaffold precludes the possibility of catalysis. One possible avenue of investigation on this front includes replacing C–H bonds in the ligand framework with more oxidatively robust groups such as C–F bonds or oxygenated groups (*i.e.* sulfonates, carbonyls, *etc.*). However, maintaining optimized cooperative reactivity with such oxidatively robust flanking groups is a notable synthetic challenge.

## 6 Conclusions

Significant progress has been made using metal–ligand cooperativity and O<sub>2</sub> for oxidative chemistry, but there is still a great deal of space for improvement in this area. In particular, there is a need to build ligand scaffolds that balance stabilizing the active oxidant and providing sufficient activation of the active oxidant to enable catalytic reactivity. Furthermore, a balance between building a ligand scaffold that minimizes intramolecular oxidation while also providing coordination and reaction space for a wide range of substrates to bind and undergo oxidation must be found. Currently, much of the research in this area functions as fundamental chemistry research, which demonstrates the viability of the approach but lacks applicability to a wide substrate scope. Future work should focus on bridging metal–ligand cooperativity in conjunction with O<sub>2</sub> with the field of organic methodology to make transition metal complexes incorporating metal–ligand cooperativity an essential part of a synthetic chemist's toolkit. While this will require further fundamental studies on the principles that govern cooperative activation of O<sub>2</sub>, more applied investigations into tuning these systems for catalysis is also needed.

## Author contributions

Both authors contributed equally to this work.

## Conflicts of interest

There are no conflicts to declare.

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