



Catalytic aspects of a copper(II) complex: biological oxidase to oxygenase activity

BISWAJIT CHOWDHURY^a, MILAN MAJI^c and BHASKAR BISWAS^{a,b,*}

^aDepartment of Chemistry, Raghunathpur College, Purulia, West Bengal 723 133, India

^bDepartment of Chemistry, Surendranath College, 24/2 M G Road, Kolkata, West Bengal 700 009, India

^cDepartment of Chemistry, National Institute of Technology, Mahatma Gandhi Avenue, Durgapur, West Bengal 713 209, India

E-mail: mr.bbiswas@rediffmail.com

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Abstract. A copper(II) complex, $[\text{Cu}(\text{dpa})_2(\text{OAc})](\text{ClO}_4)$ (**1**) [dpa = 2, 2'-dipyridylamine; OAc = acetate], has been synthesized and crystallographically characterized. X-ray structure analysis revealed that this mononuclear Cu(II) complex crystallizes as a rare class of hexa coordination geometry named bicapped square pyramidal geometry with $P2_1/c$ space group. This copper complex displays excellent catalytic efficiency, $k_{\text{cat}}/K_{\text{M}} (\text{h}^{-1}) = 6.17 \times 10^5$ towards the oxidative coupling of 2-aminophenol (2-AP) to aminophenoxazin-3-one. Further, upon stoichiometric addition of copper(II) complex to 3,5-DTBC in presence of molecular oxygen in ethanol medium, the copper complex affords predominantly extradiol cleavage products along with a small amount of benzoquinone and a trace amount of intradiol cleavage products at a rate, $k_{\text{obs}} = 1.09 \times 10^{-3} \text{ min}^{-1}$, which provide substantial evidence for the oxygen activation mechanism. This paper presents a novel addition of a copper(II) complex having the potential to mimic the active site of phenoxazinone synthase and catechol dioxygenase enzymes with significant catalytic efficiency.

Keywords. Copper(II); crystal structure; phenoxazinone synthase activity; catechol dioxygenase; bio-mimetic chemistry.

1. Introduction

Over the past few decades, biochemical oxidation reactions have significantly stimulated the coordination chemists to design various metal complexes as structural and functional mimic of the active site of different metalloproteins and metalloenzymes.^{1–3} Bio-inspired catalysis mediated by different copper proteins and enzymes include primarily dioxygen transport (hemocyanin, Hc),⁴ aromatic ring oxidations (tyrosinase, Tyr,⁵ catechol oxidase,⁶ and quercetin-2,3-dioxygenase,^{7–9} the biogenesis of neurotransmitters and peptide hormones (dopamine- β -monooxygenase, D β M,¹⁰ and peptidylglycine α -amidating monooxygenase, PHM,^{11,12} hydrogen peroxide generation (galactose and glyoxal oxidases,^{13–18} iron homeostasis (ceruloplasmin3 and Fet3p,^{19–21} and methane oxidation (particulate methane monooxygenase, pMMO).^{22–25} A prominent member of

these copper proteins is phenoxazinone synthase which is a copper-containing oxidase that catalyzes the coupling of 2-aminophenols to form the 2-aminophenoxazinone chromophore. This reaction constitutes the final step in the biosynthesis of the potent antineoplastic agent actinomycin.^{26–29} On the other hand, catechol dioxygenases of intradiol class utilizes a non-heme iron(III) cofactor in catalyzing the cleavage of the carbon–carbon bond between the two catechol oxygens; and the extradiol dioxygenases utilize a nonheme iron(II) cofactor in catalyzing the cleavage of the carbon–carbon bond adjacent to the catechol oxygens.^{30,31} In the last decade, Youngme *et al.*, and Choudhury *et al.*, produced same Cu(II)-dipyridyl complex using different reaction methodology.³² Here we introduce a copper(II)-dipyridylamine complex $[\text{Cu}(\text{dpa})_2(\text{OAc})](\text{ClO}_4)$ (**1**) [dpa = 2, 2'-dipyridylamine; OAc = acetate], with potential ability to mimic the functional sites of phenoxazinone synthase enzyme with $k_{\text{cat}} (\text{h}^{-1}) = 1.83 \times 10^3$ and catechol dioxygenase enzyme with the

*For correspondence

decomposition rate, $1.09 \times 10^{-3} \text{ min}^{-1}$, respectively. Coordinative unsaturation of the Cu(II) centre in ethanol medium facilitates the formation of substrate-enzyme adduct and account in favour of such oxidase to oxygenase activity.

2. Experimental

2.1 Materials

High purity 2,2'-dipyridylamine (Aldrich, UK), copper(II) perchlorate hexahydrate (Fluka, Germany), sodium acetate (E. Merck, India), 2-aminophenol (E. Merck, India), 3,5-di-tert-butylcatechol (Sigma Aldrich Corporation, St. Louis, MO, USA) were obtained from commercial sources and used as purchased. All other chemicals and solvents were of analytical grade and were used as received without further purification.

Caution! Perchlorate salts of metal ions are potentially explosive, especially in the presence of organic ligands. Only a small amount of material should be prepared and it should be handled with care.

2.2 Synthesis of $[\text{Cu}(\text{dpa})_2(\text{OAc})](\text{ClO}_4)(\mathbf{1})$

The copper(II) complex was synthesized by addition of aqueous solution of dipyridylamine (0.3420 g, 2 mmol) into a solution of $\text{Cu}(\text{ClO}_4)_2 \cdot 6\text{H}_2\text{O}$ (0.3650 g, 1 mmol) in the same solvent (20 mL) keeping the solution on a magnetic stirrer with stirring. Then solid sodium acetate (0.0820 g, 1 mmol) was added in solid into the blue solution and stirring continued to 30 min more. The blue solutions were turned into green and the supernatant liquids were kept in air for slow evaporation. After 7–10 days the fine microcrystalline compound was separated out and washed with hexane and dried *in vacuo* over silica gel indicator. The spectroscopic measurements and elemental analyses confirm the structural formation of the complex. Yield = $\sim 0.28\text{g}$, ($\sim 77\%$ based on metal salt). Anal. calc. (%) $\text{C}_{22}\text{H}_{22}\text{N}_6\text{ClO}_6\text{Cu}$: C, 46.73; H, 3.92; N, 14.86; Found(%): C, 46.69; H, 3.88; N, 14.89. Selected IR bands (KBr pellet, cm^{-1}): 3321 (m), 1637(s), 1604(s), 1423(m), 1378(s), 1095(s). UV-Vis (λ , nm; 10^{-4}M , 1 cm cell length, abs, ethanol): 298(1.86), 725–745(0.00180) (broad band); ESI-MS (MeCN): m/z , 406.10 ($[\text{Cu}(\text{dpa})_2]\text{-H}^+$); (Calc. 406.09).

2.3 Physical measurements

Infrared spectrum (KBr) was recorded with a FTIR-8400S SHIMADZU spectrophotometer in the range $400\text{--}3600 \text{ cm}^{-1}$. ^1H NMR spectrum in $\text{DMSO-}d^6$ was obtained on a Bruker Avance 300 MHz spectrometer at 25°C and was recorded at 299.948 MHz. Ground-state absorption measurements were made with a Jasco model V-730 UV-Vis spectrophotometer. Elemental analyses were performed on a Perkin Elmer 2400

CHN microanalyser. Electrospray ionization (ESI) mass spectrum was recorded on a Q-TOF MicroTM Mass Spectrometer. The electrochemical studies were carried out using Cyclic voltammograms were recorded in CH_3CN solutions containing 0.1 M TBAP at 25°C using a three-electrode configuration (Pt working electrode, Pt counter electrode, Ag/AgCl reference) and a PC-controlled PAR model 273A electrochemistry system. All the experimental solutions were degassed for 30 min with high-purity argon gas before any cyclic voltammetry of a sample was done. The Electron Paramagnetic Resonance (EPR) spectrum was recorded on a Bruker EMX-X band spectrometer.

2.4 Crystal structure determination and refinement

Single crystal X-ray diffraction data of the copper(II) complex were collected using a Rigaku XtaLABmini diffractometer equipped with Mercury CCD detector. The data were collected with graphite monochromated Mo-K α radiation ($\lambda = 0.71073 \text{ \AA}$) at 293 K using ω scans. The data were reduced using Crystal Clear suite 2.0³³ and the space group determination was done using Olex2. The structure was resolved by direct method and refined by full-matrix least-squares procedures using the SHELXL-2014/7³⁴ software package through the OLEX2 suite.³⁵ The crystallographic bond distance and bond angle are given in Table 1 and Table S1 (in Supplementary Information).

2.5 Catalytic oxidation of 2-aminophenol

In order to examine the penoxazinone synthase activity, $1 \times 10^{-3}\text{M}$ solution of **1** in EtOH was treated with 10 equiv.

Table 1. Crystallographic refinement parameters of $[\text{Cu}(\text{dpa})_2(\text{OAc})](\text{ClO}_4)$ (**1**).

Parameters	Cu(II) compound
Empirical formula	$\text{C}_{22}\text{H}_{21}\text{N}_6\text{O}_6\text{ClCu}$
Formula weight	564.44
Temperature (K)	293
Crystal system	Monoclinic
Space group	$P2_1/c$
a (\AA)	13.885(12)
b (\AA)	7.899(7)
c (\AA)	22.202(18)
Volume (\AA^3)	2582.7(2)
Z	4
ρ (g cm^{-3})	1.541
μ (mm^{-1})	1.058
F (000)	1156
θ ranges ($^\circ$)	3.1° to 27.50°
R_{int}	0.038
R (reflections)	15670
wR2 (reflections)	5578
Final R indices	0.0634, 0.1929
Largest peak and hole (eA^{-3})	0.77, -0.44

of 2-aminophenol (2-AP) under aerobic conditions at room temperature. Absorbance vs. wavelength (wavelength scans) of the solution was recorded at a regular time interval of 15 min for aminophenol oxidation in the wavelength range 300–800 nm. Kinetic experiments were performed spectrophotometrically^{26–28} with Cu(II) complex and 2-AP in EtOH at 25°C for aminophenol oxidation activity. 0.04 mL of the complex solution, with a constant concentration of 1×10^{-3} M, was added to 2 mL of 2-AP of a particular concentration (varying its concentration from 1×10^{-3} M to 1×10^{-2} M) to achieve the ultimate concentration of the complex as 1×10^{-3} M. The conversion of 2-aminophenol to 2-aminophenoxazine-3-one was monitored with time at 433 nm (time scan)²⁸ in EtOH. To determine the dependence of rate on substrate concentration, kinetic analyses were performed in triplicate. Effect of catalyst concentration on the reaction rate for the catalytic aminophenol oxidation in EtOH medium was determined by varying the concentration of copper(II) complex (5×10^{-3} M to 5×10^{-4} M) with a constant concentration (1×10^{-2} M) of each substrate. Here, the kinetic analyses were also performed in triplicate.

2.6 Catechol dioxygenase activity of the copper(II)-dipyridylamine complex

To investigate the catechol dioxygenase activity of the complex, a 10^{-3} M solution of **1** in ethanol (EtOH) solvent was treated with a 10^{-3} M solution of 3,5-di-*tert*-butylcatechol (DTBC) in oxygen saturated ethanol medium at room temperature. Absorbance vs. wavelength (wavelength scans) of the solution was recorded at regular time intervals for 6 h in the wavelength range 200–900 nm.³⁶ It may be noted here that a blank experiment without catalyst did not show the formation of any cleavage products up to 12 h in EtOH. The solvent was equilibrated at the atmospheric pressure of O₂ at 25°C using a known procedure with modifications.³⁷ Investigation of dioxygen reactivity of the *in situ* generated Cu(II)-catecholate adduct was carried out in oxygen saturated ethanol medium at 25°C. Kinetic analyses³⁶ of the catechol cleavage reactions were carried out by time-dependent measurement of the disappearance of the lower energy DBC²⁻-to-Cu(II) LMCT band at 824 nm by exposing to molecular oxygen.

2.7 Determination of 3,5-di-*tert*-butyl catechol cleavage products

0.2260 g (0.04 mmol) of the Cu(II) complex was reacted with 0.088 g (0.04 mmol) of DTBC in an oxygen saturated ethanol (100 mL) at ambient condition and then allowed to stir for 12 h. The reddish brown solution slowly turned to green. The residue was then treated with 15 mL of 3M HCl and the catechol cleavage products were extracted with diethylether (3×10 mL) and dried over sodium sulfate. The catechol cleavage products were analyzed by ESI-MS and were quantified by ¹H NMR spectroscopy. ¹H NMR data for 3,5-di-*tert*-butyl catechol cleavage products (500

MHz, CDCl₃): δ = 3, 5-di-*tert*-butyl-2-pyrone: 6.09 (m, 2H); 4,6-di-*tert*-butyl-2-pyrone: 7.11 (d, 1H), 7.24 (d, 1H); 3,5-di-*tert*-butylbenzoquinone: 6.19 (d, 1H), 6.74 (d, 1H); 3,5-di-*tert*-butyl-5-(carboxymethyl)-2-furanone: 6.84 (s, 1H).

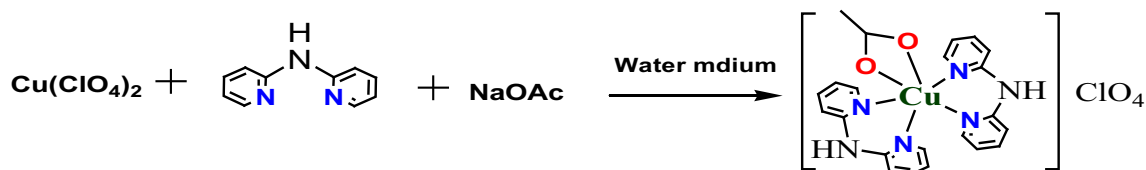
3. Results and Discussion

3.1 Synthesis and formulation of the Cu(II)-dpa complex (**1**)

The copper(II) complex was synthesized by addition of 2, 2'-dipyridylamine to a solution of Cu(ClO₄)₂ · 6H₂O in water followed by the addition of sodium acetate (NaOAc) at room temperature (Scheme 1). The structure was determined by routine spectroscopic techniques including single crystal X-ray diffraction analysis.

3.2 Description of the crystal structure

The X-ray crystal structure analysis of copper(II) complex reveals that it crystallizes in monoclinic crystal system with *P*2₁/*c* space group. An ORTEP of the copper(II) complex is shown in Figure 1. The crystallographic structural parameters are given in Table 1 and bond angles, and bond distances are given in Table S1. Monocationic [Cu(dpa)₂(OAc)]⁺ unit exists in a rare class of six coordination geometry named bicapped square pyramidal geometry and the residual cationic charge is counterbalanced by a perchlorate ion. This copper(II) complex resembles to previously reported bicapped square pyramidal chromophore, [Cu(bpy)₂(O₂NO)](NO₃)³⁸ and this structure is known as a rare class of coordination geometry in the scientific literature. The square plane contains N1, N2 (dpa), N4 (other dpa) atoms and O1 atom of acetate ligand [Cu1-N1, 1.99 Å; Cu1-N2, 2.02 Å; Cu1-N4, 2.01 Å; Cu1-O1, 2.0 Å] while N2 (other dpa) and O2 (acetate) atoms occupy the apical position [Cu1-N3, 2.15 Å; Cu1-O2, 2.70 Å]. Though there are two Cu-O bonds (axial and equatorial) around the Cu centre but Cu1-O2 bond gets axially elongated due to Jahn-Teller distortion.³⁹ In searching for the origin of the existence of this unusual geometry of the copper(II) complex in the solid state, we closely observe the role of H-bonding interaction. In the crystalline state of Cu(II) complex, perchlorate anion has a substantial role to rotate the pyridine rings of the dipyridyl amine. The oxygen atoms of the ClO₄⁻ ion act as donor centres and H-atoms of the pyridine rings as well as secondary amines behave as acceptors [C(10)-H(10)···O5, 2.68 Å; C(17)-H(17)···O6, 2.677 Å; C(17)-H(17)···O4, 2.601 Å; C(8)-H(8)···O3, 2.635 Å; C(14)-H(14)···O3, 2.652 Å; N(6)-H(6)···O3, 2.028 Å;



Scheme 1. Preparative procedure for the Cu(II) complex.

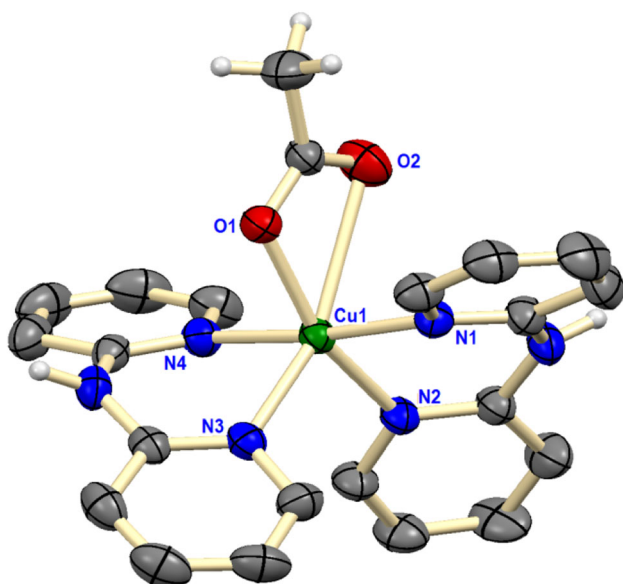


Figure 1. An ORTEP diagram of Cu(II)-dpa complex (30% ellipsoid probability).

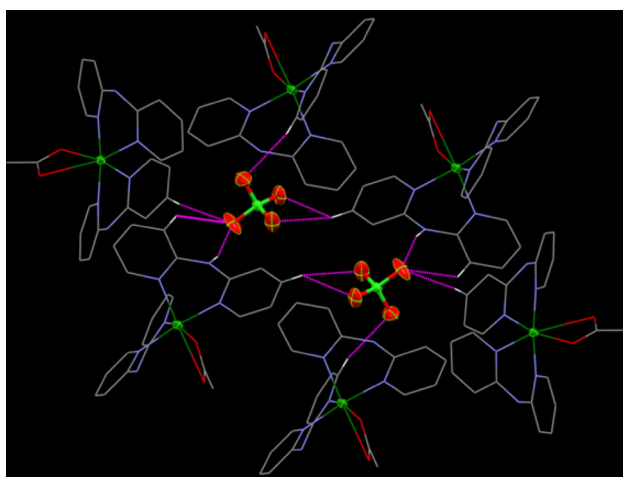


Figure 2. Perchlorate ion mediated 3D crystalline architecture through C/N-H \cdots O interaction in solid state.

Figure 2, Table S2 in Supplementary Information] to form a 3D architecture through C-H \cdots O/N-H \cdots O interactions along *b* axis.

3.3 Electronic, EPR and electrochemical characterization

The solution integrity of the copper complex has been performed through UV-Vis and EPR spectral analysis. The UV-Vis spectrum in ethanol medium at room temperature shows a series of high intensity transitions in the range of 225 to 238 and 295 nm which are originated from the intraligand electronic transitions (Figure S1). A very broad low energy transition centered at 741 nm (Figure S1 in Supplementary Information), which is assigned to the ligand field transition for the copper(II) complex. The X-band EPR spectrum of the copper complex (Figure S2 in SI) at low temperature (LT, 77K) was recorded in frozen MeCN solution (Figure S2 (Left)) and in powder state at 77 K (Figure S2 (Right) in SI) to examine the geometry of copper center in **1**. The bicapped square pyramidal environment was also confirmed from the presence of four lines in the LT spectrum and suggested the existence of a single species (Figure S2).

We have also examined the electrochemical behaviour of this copper(II) complex in ethanol at RT. The reduction and oxidation potentials for Cu(II)-dpa complex was observed, respectively, at -773 and +42 mV (Figure S3 in SI) at the scan rate of 20 mV s⁻¹. The high value of reduction potential suggests that the reduction is quite difficult for central Cu(II) ion.

3.4 Catalytic oxidation of 2-aminophenol

The phenoxazinone synthase activity of the copper(II) complex was studied using 2-aminophenol (2-AP) as a convenient model substrate, in air saturated ethanol solvent at room temperature (25°C). For this purpose, a 1×10^{-3} M solution of the copper(II)-dipyridylamine complex was treated with 1×10^{-2} M (10 equiv.) of 2-AP and the course of the reaction was followed by recording the UV-Vis spectra of the mixture at an interval of 15 min for 3 h.

Spectral bands at 225–238, 295 and 741 nm appeared in the electronic spectrum of the copper complex in methanol, whereas 2-AP showed a single band at 267 nm. As the reaction proceeded, there was a gradual

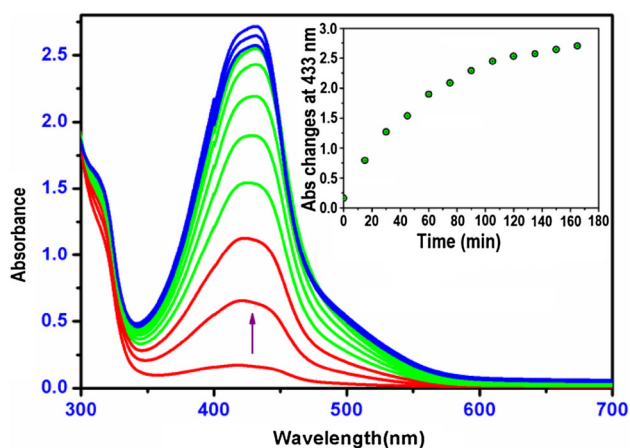


Figure 3. Increase of aminophenoxazinone band at 433 nm after addition of 10 equivalents of 2-AP to a Cu(II) solution in EtOH medium. The spectra were recorded after every 9 min. Inset: Plot of Abs versus time.

decrease in intensity of the band at 267 nm²⁶ and an initial new broad band centered at 433 nm with increasing intensity was formed (Figure 3), which indicated the formation of the respective phenoxazinone species.^{26–29} The phenoxazinone was purified by column chromatography and isolated in high yield (81.6% for **1**) by slow evaporation of the eluant. The product was identified by ¹H NMR spectroscopy. ¹H NMR (CDCl₃, 300 MHz,) δ_{H} : 7.62 (m, 1H), 7.48 (m, 3H), 6.41 (s, 1H), 6.30 (s, 1H).

Kinetic studies were performed to understand the extent of the efficiency. The kinetics of oxidation of 2-AP were determined by the method of initial rates and involved monitoring the growth of the phenoxazinone band at 433 nm as a function of time (Figure S4 in SI).^{28,29} The plot of rate constants versus concentration of the substrate were also analyzed on the basis of the Michaelis-Menten approach of enzymatic kinetics to get the values of the kinetics parameters, $V_{\text{max}} 5.11 \times 10^{-4}$ (Sd. Error. 7.78×10^{-5}), $K_{\text{M}} = 2.98 \times 10^{-3}$ (Sd. Error 9.14×10^{-3}) and $k_{\text{cat}} = 1.83 \times 10^3$. The observed rate constant versus substrate concentration plot for copper(II)-dipyridylamine complex in EtOH is shown in Figure S4 (in SI).

The high catalytic efficiency value ($k_{\text{cat}}/K_{\text{M}} = 2.09 \times 10^5$) for copper(II) complex also indicates its high reactivity towards aminophenol oxidation. We also investigate the effect of catalyst concentration on reaction rate and the rate of catalytic oxidation increase with increasing the concentration of copper(II)-dipyridylamine complex in a linear manner. Reuse of this copper(II)-dipyridylamine catalyst (Figure S9) for aminophenol oxidation was also examined under similar reaction conditions after completion of catalytic

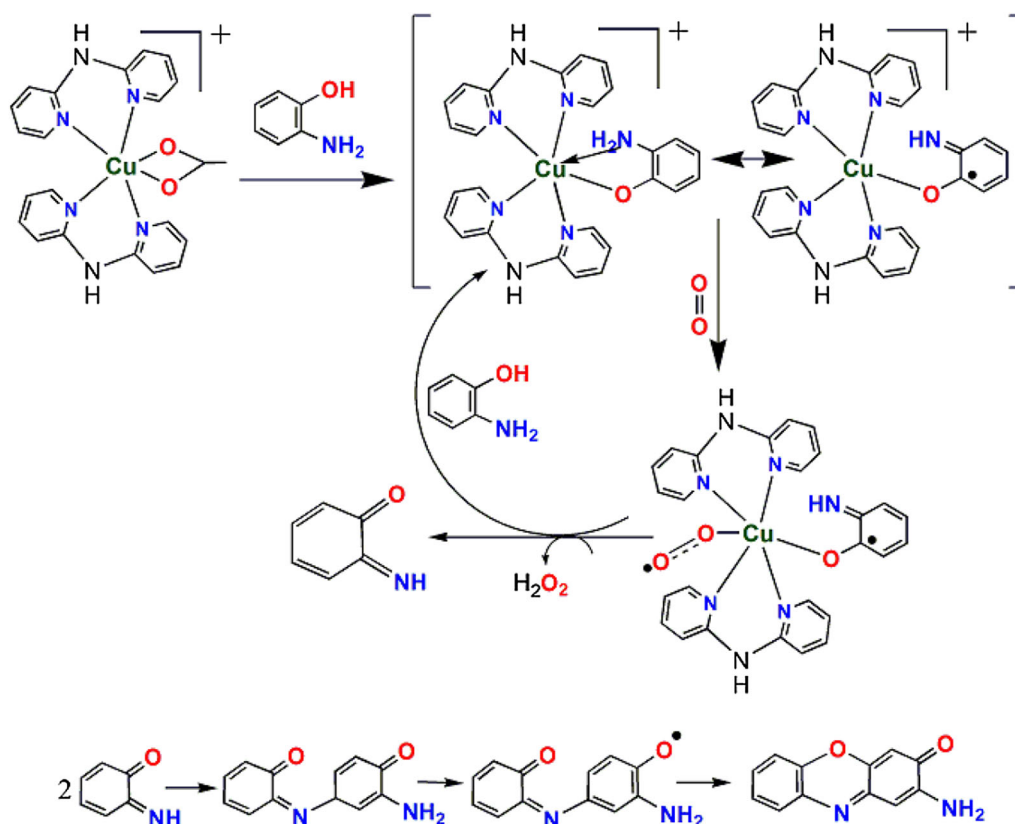
oxidation. The reused copper(II)-dipyridylamine catalyst exhibited no change in reactivity after four cycles also. In order to make a comparison of the phenoxazinone activity between our copper-dpa complex and the reported copper(II) complex, [Cu(bpy)₂(O₂NO)](NO₃) of the same class, we performed phenoxazinone synthase activity for the both complexes under similar reaction conditions. We find that Cu(II)-dpa complex exhibits better catalytic efficiency towards the oxidative coupling of 2-aminophenol than the reported copper complex.³⁸

Though this class of catalytic oxidation reaction under aerobic condition bears special attention, we could not find enough literature reports to make a comparison between phenoxazinone synthase activity of the Cu(II)-dipyridylamine in the present case and previously reported works. Chaudhury *et al.*,^{40a} modelled a tetracopper complex for the catalytic aerial oxidation of 2-aminophenol to 2-amino-phenoxazine-3-one, and proposed an “on-off” mechanism of the radicals together with redox participation of the metal center behind the mimics of six-electron oxidative coupling in the catalytic function of the coppercontaining enzyme phenoxazinone synthase. Begley *et al.*,^{40b} suggested that 2-aminophenoxazinone synthesis proceeds via a sequence of three consecutive 2-electron aminophenol oxidations and that the aminophenol moiety is regenerated during the reaction sequence by facile tautomerization reactions.

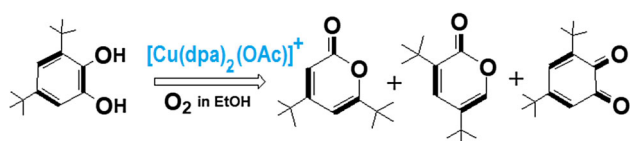
In order to evaluate the mechanistic aspects of the catalytic cycle by copper(II)-dipyridylamine complex (Scheme 2) for the oxidative coupling of phenoxazinone product, mass spectral analysis of the reaction mixture in ethanol medium provides valuable information regarding the reactive species and product formed in the reaction. The mass spectrum of the reaction mixture (Figure S5 in SI) exhibits mainly the characteristic peaks at m/z 213.57 and 557.99 with isotope distribution patterns which further consolidated the corroboration respectively [(2-amino-3*H*-phenoxazine-3-ones)+H⁺] and [[Cu(dpa)₂(2-AP)]+H⁺]. Further, the characteristic peak at $m/z \sim 557.99$ nm indicates the presence of coordinated iminobenzosemiquinonato radical during the generating apx product.

3.5 Catechol dioxygenase activity of the copper(II)-dipyridylamine complex

The oxygenation reactions for the copper(II)-dipyridylamine complex was carried out using 3,5-di-*tert*-butylcatechol (DTBC) as the model substrate in bio-friendly ethanol, and the advantages of using the latter as substrate are the relatively high stability of the main



Scheme 2. Proposed mechanistic routes for aminophenol oxidation by copper(II) complex.



Scheme 3. Oxygenation products of DTBC for Cu(II) complex in ethanol.

cleavage product and the fast reaction of the catecholate complex with dioxygen at room temperature (Scheme 3).

The DTBC²⁻ adducts of Cu(II) complex was generated *in situ* in ethanol solution, and their reactivity toward O₂ was investigated by monitoring the decay of the low energy DTBC²⁻ to Cu(II) LMCT band (Figure 4). The green solution of the *in situ* Cu(II)-catecholate adduct reacts with dioxygen in bio-friendly ethanol medium at ambient conditions over a period of 4 h and two new visible bands with maximum absorption at 516 and 824 nm (Figure 4), were observed for the catecholate adduct of Cu(II) complex. The lower energy visible bands with decreasing in absorbance are attributed to DTBC²⁻ to-copper(II) LMCT transitions involving two different catecholate ligand orbitals.^{30,31} Commonly, it is seen that the energy of the LMCT transition strongly depends on the nature of the ligands

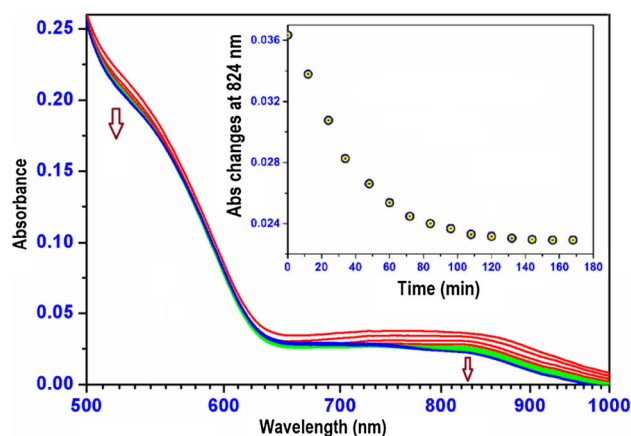


Figure 4. Absorption spectral changes during the reaction of the *in situ* generated adduct [Cu(dpa)₂(2-AP)] with O₂ (The spectra were recorded after every 12 min). Inset: Plot of Abs versus Time.

and Lewis acidity of the metal ion in the complex.^{41,42} The disappearance of the lower-energy catecholate-to-copper(II) LMCT band (Figure 4) on oxygenation exhibits pseudo first-order kinetics, as judged from the linearity of the plot [1 + log(Absorbance)] versus time⁴³ (Figure 5), and the value of k_{obs} was obtained from the

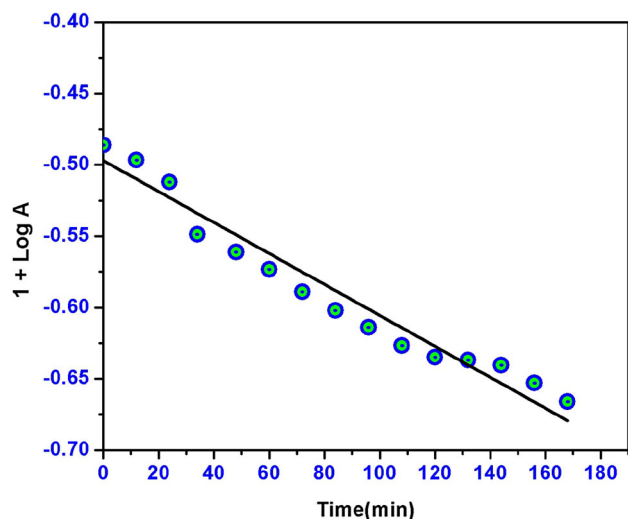
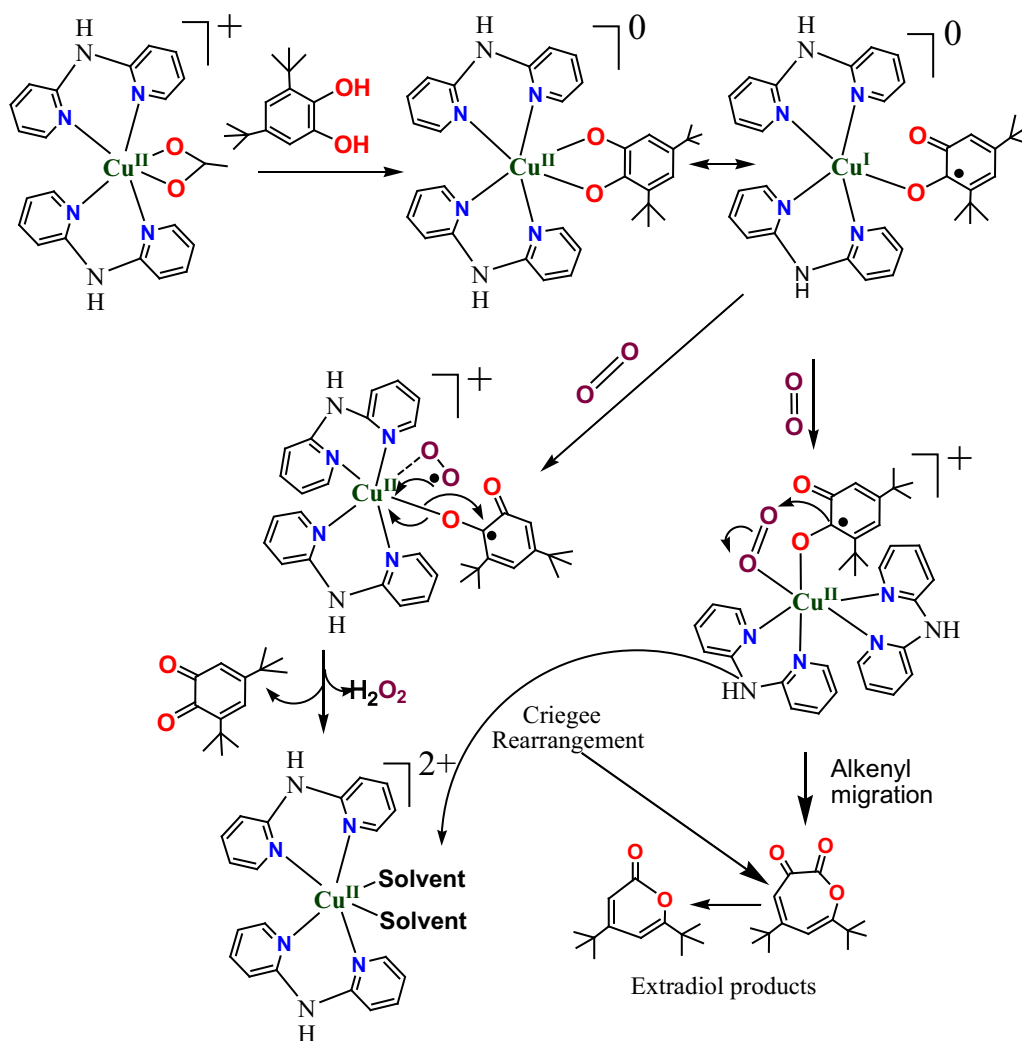


Figure 5. Plot of $[1 + \log(\text{Absorbance})]$ versus time for the reaction of $[\text{Cu}(\text{dpa})_2(\text{DBC})]$ with O_2 at 25°C in EtOH solution.

slope of the plot. The pseudo-first order rate constant was determined as $k_{\text{obs}}: 1.09 \times 10^{-3} \text{ min}^{-1}$.

Further, the cleavage products from catechol bound Cu(II) species were identified and quantified by ^1H NMR spectroscopy (Figure S6 in SI). The distribution of catechol-derived products was found to be principally 4,6-di-*tert*-butyl-2-pyrone and 3,5-di-*tert*-butyl-2-pyrone, as extradiol cleavage products, in major amount and 3,5-di-*tert*-butylbenzoquinone as minor product, while 3,5-di-*tert*-butyl-5-(carboxymethyl)-2-furanone as intradiol cleavage product was found in trace quantity. In the reaction of (Cu complex+DTBC) with O_2 in ethanol, 81% of extradiol cleavage products (42.6%+38.4%) were obtained along with the formation of a trace amount of intradiol cleavage product as a side product (Scheme 4). The percentage of minor oxidation product, 3,5-di-*tert*-butylbenzoquinone, was found to be a small one (16.8%) also. The amount of



Scheme 4. Proposed mechanistic pathway for the formation of cleavage products by **1**.

organic product from catechol cleavage accounts for ~ 95% of 3,5-di-*tert*-butyl catechol. The remaining 5% was accounted as unreacted substrate.

The formation of extradiol cleavage products for the *in situ* catecholate adduct of the Cu(II) complex is expected for having a vacant coordination site on the catecholate adduct for oxygen coordination, which favors dioxygen-activation pathway.^{44–46} From ESI mass spectrometric analysis of the reaction mixture, it is revealed that at the primary stage 3,5-DTBC forms *in situ* adduct but the presence of semiquinone band ~ 516 nm helps to detect DTBSQ radical. The existence of the DTBQ radical in solution during the investigation of dioxygenase activity was confirmed by the EPR signal at $g = 2.075$ (Cu complex with 3,5-DTBC in EtOH) (Figure S7 in SI). But existence of Cu(I)-semiquinone species in solution facilitates dioxygen activation mechanism and provides one electron from Cu(I) to the anti-bonding orbital of dioxygen which generates oxo-species in solution and produces extradiol cleavage products in a major amount.⁴³ A small amount of benzoquinone indicates the natural oxidation of the substrate in solution. Probably, trace amount of intradiol catechol cleavage reaction proceeded by peroxo intermediate that underwent 1,2-Criegee rearrangement⁴ to yield the intradiol catechol cleaved products analogous to the native enzyme. Earlier models have shown that the dioxygenase activity strongly depends on the nature of the ligand set and the coordination mode of the catecholate ligand.^{47–51} Speier *et al.*, reported an aerobic oxidation of a (phenanthrenediolato)copper(II) complex of tmeda, which is in resonance with two valence tautomers of the (phenanthrenediolato) copper(II) and the (phenanthrenesemiquinonato) copper(I) states.⁵² The (phenanthrenesemiquinonato) copper(I) tautomer can bind a dioxygen molecule to form a dioxygen complex, which is decomposed to ring-cleavage products.

4. Conclusions

Herein, we report an isolation of a copper(II)-dipyridylamine complex, $[\text{Cu}(\text{dpa})_2(\text{OAc})](\text{ClO}_4)$ with unusual hexa coordination geometry. X-ray structure shows that this copper complex crystallizes with $P2_1/c$ space group in a monoclinic system. It shows significant catalytic ability, $k_{\text{cat}}/K_{\text{M}}(\text{h}^{-1}) = 6.17 \times 10^5$ for the oxidative coupling of 2-AP to aminophenoxazin-3-one. Upon stoichiometric addition of copper(II) complex to 3,5-DTBC, two catecholate-to-Cu(II) LMCT bands (516 and 824 nm) were observed and the *in situ* generated catecholate intermediate reacts with molecular oxygen at the rate, $k_{\text{obs}} = 1.09 \times 10^{-3} \text{ min}^{-1}$ in ethanol medium

to afford predominantly extradiol cleavage products along with a small amount of benzoquinone, and a trace amount of intradiol cleavage products. The yield of cleavage products is strongly in favour of molecular oxygen activation mechanism. This paper presents a novel addition of a copper(II) complex having the potential to mimic the active site of phenoxazinone synthase and catechol dioxygenase enzyme with significant catalytic efficiency.

Supplementary Information (SI)

CCDC 1513638 contains the supplementary crystallographic data for **1**. These data can be obtained free of charge via <http://www.ccdc.cam.ac.uk/conts/retrieving.html> or from the Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: (+44) 1223-336-033; or e-mail: deposit@ccdc.cam.ac.uk. Experimental data such as IR and UV-Vis spectra, cyclic voltogram, EPR spectra at low temperature, ESI mass spectra, ¹H NMR of cleavage products, rate vs. substrate plot, Lineweaver-Burk plot, bond distance, bond angle & H-bonded interaction parameters are available at www.ias.ac.in/chemsci.

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References

- (a) Kaim W and Schwederski B 1993 *Bioanorganische Chemie*, Teubner, Stuttgart Bioinorganic Catalysis J Reedijk (Ed.) (New York: Marcel Dekker); (b) Biswas B, Patra M, Dutta S, Ganguly M and Kole N 2013 Synthesis, structural characterization and biological activity of a trinuclear zinc(II) complex: DNA interaction study and antimicrobial activity *J. Chem. Sci.* **125** 1445; (c) Dey D, Basu Roy A, Shen C-Y, Tsai H-L, Ranjani A, Gayathri L, Chandaleka S, Dhanasekaran D, Akbarsha M A, Kole N and Biswas B 2015 Synthesis and bio-catalytic activity of isostructural cobalt(III)-phenanthroline complexes *J. Chem. Sci.* **127** 649; (d) Das S, Pasan J, Gayathri L, Saha S, Chandaleka S, Maji M, Dhanasekaran D, Akbarsha M A, Kole N and Biswas B 2016 Recognition of self-assembled water-nitrate cluster in a Co(III)-2,2'-bipyridine host: synthesis, crystal structure, DNA cleavage, molecular docking and anticancer activity *J. Chem. Sci.* **128** 1755

2. (a) Klinman J P 1996 Mechanisms whereby mononuclear copper proteins functionalize organic substrates *Chem. Rev.* **96** 2541; (b) Dey D, Kaur G, Ranjani A, Gyathri L, Chakraborty P, Adhikary J, Pasan J, Dhanasekaran D, Choudhury A R, Akbarsha M A, Kole N and Biswas B 2014 A trinuclear zinc–schiff base complex: biocatalytic activity and cytotoxicity *Eur. J. Inorg. Chem.* **3350**; (c) Chowdhury B, Patra M, Maji M and Biswas B 2015 Ligand centered radical pathway in catechol oxidase activity with a trinuclear zinc-based model: synthesis, structural characterization and luminescence properties *Spectrochim. Acta Part A Mol Biomol. Spect.* **144** 148
3. (a) Solomon E I, Sundaram U and Machonkin T E 1996 Multicopper oxidases and oxygenases *Chem. Rev.* **96** 2563; (b) Biswas B, Al-Hunaiti A, Räisänen M T, Ansalone S, Leskelä M, Repo T, Chen Y-T, Tsai H-L, Naik A D, Railliet A P, Garcia Y, Ghosh R and Kole N 2012 Efficient and selective oxidation of primary and secondary alcohols using an iron(III)/phenanthroline complex: structural studies and catalytic activity *Eur. J. Inorg. Chem.* 4479; (c) Pal A, Biswas B, Mitra M, Purohit C S, Hazra S, Kumar G S, Ghosh R 2013 Synthesis, X-ray structure and DNA binding of a mononuclear iron(II) Schiff base complex *J. Chem. Sci.* **125** 1161
4. Magnus K A, Ton-That H and Carpenter J E 1994 Recent structural work on the oxygen transport protein hemocyanin *Chem. Rev.* **94** 727
5. Sa'nchez-Ferrer A, Rodri'guez-Lo'pez J N, Garcí'a-Ca'novas F and Garcí'aCarmona F 1995 Tyrosinase: a comprehensive review of its mechanism *Biochim. Biophys. Acta* **1** 1247
6. Gerdemann C, Eicken C and Krebs B 2002 The crystal structure of catechol oxidase: new insight into the function of type-3 copper proteins *Acc. Chem. Res.* **35** 183
7. Fusetti F, Schröter K H, Steiner R A, van Noort P I, Pijning T, Rozeboom H J, Kalk K H, Egmond M R and Dijkstra B W 2002 crystal structure of the copper-containing quercetin 2,3-dioxygenase from *Aspergillus japonicus* *Structure* **10** 259
8. Steiner R A, Kooter I M and Dijkstra B W 2002 Functional analysis of the copper-dependent quercetin 2,3-dioxygenase. 1. Ligand-induced coordination changes probed by X-ray crystallography: inhibition, ordering effect, and mechanistic insights *Biochemistry* **41** 7955
9. Kooter I M, Steiner R A, Dijkstra B W, van Noort P I, Egmond M R and Huber M 2002 EPR characterization of the mononuclear Cu-containing *Aspergillus japonicus* quercetin 2,3-dioxygenase reveals dramatic changes upon anaerobic binding of substrates *Eur. J. Biochem.* **269** 2971
10. Stewart L C and Klinman J P 1988 Dopamine beta-hydroxylase of adrenal chromaffin granules: structure and function *Annu. Rev. Biochem.* **57** 551
11. Prigge S T, Mains R E, Eipper B A and Amzel L M 2000 New insights into copper monooxygenases and peptide amidation: structure, mechanism and function *Cell. Mol. Life Sci.* **57** 1236
12. Blackburn N J, Rhames F C, Ralle M and Jaron S 2000 Major changes in copper coordination accompany reduction of peptidylglycine monooxygenase: implications for electron transfer and the catalytic mechanism *J. Biol. Inorg. Chem.* **5** 341
13. Whittaker J W 2003 Free radical catalysis by galactose oxidase *Chem. Rev.* **103** 2347
14. Whittaker J W and Whittaker M M 1998 Radical copper oxidases: one electron at a time *Pure Appl. Chem.* **70** 903
15. Knowles P F and Ito N 1994 In *Perspectives in Bioinorganic Chemistry* (London: Jai Press) Vol. **2** p. 207
16. Halcrow M, Phillips S and Knowles P 2000 In *Subcellular Biochemistry*, **35**, *Enzyme-Catalyzed Electron and Radical Transfer* A Holzenburg and N S Scrutton (Eds.) (New York: Plenum) 183
17. Whittaker M M, Kersten P J, Nakamura N, Sanders-Loehr J, Schweizer E S and Whittaker J W 1996 Glyoxal oxidase from *phanerochaete chrysosporium* is a new radical-copper oxidase *J. W. J. Biol. Chem.* **271** 681
18. Whittaker M M, Kersten P J, Cullen D and Whittaker J W 1999 Identification of catalytic residues in glyoxal oxidase by targeted mutagenesis *J. Biol. Chem.* **274** 36226
19. Kosman D J, Hassett R, Yuan D S and McCracken J 1998 Spectroscopic characterization of the Cu (II) sites in the Fet3 protein, the multinuclear copper oxidase from yeast required for high-affinity iron uptake *J. Am. Chem. Soc.* **120** 4037
20. Blackburn N J, Ralle M, Hassett R and Kosman D 2000 Spectroscopic analysis of the trinuclear cluster in the Fet3 protein from yeast, a multinuclear copper oxidase *J. Biochemistry* **39** 2316
21. Palmer A E, Quintanar L, Severance S, Wang T-P, Kosman D J and Solomon E I 2002 Spectroscopic characterization and O₂ reactivity of the trinuclear Cu cluster of mutants of the multicopper oxidase Fet3p *Biochemistry* **41** 6438
22. Nguyen H-H T, Nakagawa K H, Hedman B, Eliot S J, Lidstrom M E, Hodgson K O and Chan S I 1996 X-ray absorption and EPR studies on the copper ions associated with the particulate methane monooxygenase from *Methylococcus capsulatus* (Bath). Cu(I) ions and their implications *J. Am. Chem. Soc.* **118** 12766
23. Elliott S J, Randall D W, Britt R D and Chan S I 1998 Pulsed EPR studies of particulate methane monooxygenase from *Methylococcus capsulatus* (Bath): evidence for histidine ligation *J. Am. Chem. Soc.* **120** 3247
24. Nguyen H-H T, Elliott S J, Yip J H-K and Chan S I 1998 The particulate methane monooxygenase from *Methylococcus capsulatus* (Bath) is a novel copper-containing three-subunit enzyme isolation & characterization *J. Biol. Chem.* **273** 7957
25. Lieberman RL, Shrestha D B, Doan P E, Hoffman B M, Stemmler T L and Rosenzweig A C 2003 Purified particulate methane monooxygenase from *Methylococcus capsulatus* (Bath) is a dimer with both mononuclear copper and a copper-containing cluster *Proc. Natl. Acad. Sci. U.S.A.* **100** 3820
26. (a) Katz E 1967 Biosynthesis of secondary metabolites: roles of trace metals In *Antibiotics II* D Gottlieb D and P D Shaw (Eds.) p. 276 (New York: Springer); (b) McLain J, Lee J, Groves J T 2000 Biomimetic oxidations catalyzed by transition metal complexes In *Biomimetic Oxidations Catalyzed by Transition Metal Complexes* B Meunier (Ed.) (London: Imperial College

- Press); (c) Simándi T M, Simándi L I, Győr M, Rockenbauer A and Gömöry Á 2004 Kinetics and mechanism of the ferroxime (II)-catalysed biomimetic oxidation of 2-aminophenol by dioxygen. A functional phenoxazinone synthase model *Dalton Trans.* 1056–1060
27. (a) Hollstein U 1974 Actinomycin. Chemistry and mechanism of action *Chem. Rev.* **74** 625; (b) Simándi L I, Németh S and Rumlis N 1987 Study of the oxidation of 2-aminophenol by molecular oxygen catalyzed by cobalt (II) phthalocyaninetetrasodiumsulfonate in water *J. Mol. Catal.* **42** 357
 28. (a) Butenandt A 1957 Über Ommochrome, eine Klasse natürlicher Phenoxazon-Farbstoffe *Angew. Chem.* **69** 16; (b) Simándi TM, Simándi LI, Győr M, Rockenbauer A, Gömöry A, 2004 Kinetics and mechanism of the ferroxime (II)-catalysed biomimetic oxidation of 2-aminophenol by dioxygen. A functional phenoxazinone synthase model *J. Chem. Soc. Dalton Trans.* 1056
 29. (a) Cavill G W K, Clezy P S, Tetaz J R and Werner R L 1959 Synthesis of novel angular diazaphenoxazinone derivatives via palladium catalyzed Buchwald-Hartwig amidation protocols *Tetrahedron* **5** 275; (b) Kaizer J, Csonka R and Speier G 2002 TEMPO-initiated oxidation of 2-aminophenol to 2-aminophenoxazin-3-one *J. Mol. Catal. A: Chem.* **180** 91
 30. (a) Que L, Jr, Lipscomb J D, Münck E and Wood J M 1977 Protocatechuate 3,4-dioxygenase: inhibitor studies and mechanistic implications *Biochim. Biophys. Acta* **485** 60
 31. (a) Ohlendorf D H, Lipscomb J D and Weber P C 1988 Structure and assembly of protocatechuate 3, 4-dioxygenase *Nature* **336** 403; (b) Solomon E I, Sundaram U M, Machonkin T E 1996 Multicopper oxidases and oxygenases *Chem. Rev.* **96** 2563; (c) Ohlendorf D H, Orville A M and Lipscomb J D 1994 Structure of protocatechuate 3, 4-dioxygenase from *Pseudomonas aeruginosa* at 2.15 Å resolution *J. Mol. Biol.* **244** 586; (d) Valley M P, Brown C K, Burk D L, Vetting M W, Ohlendorf D H and Lipscomb J D 2005 Roles of the equatorial tyrosyl iron ligand of protocatechuate 3, 4-dioxygenase in catalysis *Biochemistry* **44** 11024
 32. (a) Youngme S, Phuengphai P, Chaichit N, Mutikainen I, Turpeinen U and Murphy B M 2007 Crystal structures and electronic properties of three fluxional [Cu(di-2-pyridylamine)₂(OXO)]Y complexes *J. Coord. Chem.* **60** 131; (b) Choudhury S R, Chen C-Y, Seth S, Kar T, Lee H M, Colaciu E and Mukhopadhyay S 2009 Anion- π interaction stitching 2-D layers formed by self-assembly of cations of a mononuclear copper(II) complex: synthesis, crystal structure and magnetism of [Cu(OAc)(2,2'-dypam)₂](ClO₄) [HOAc = acetic acid, 2, 2'-dypam = 2,2'-dipyridylamine] *J. Coord. Chem.* **62** 540
 33. *CrystalClear 2.0*; Rigaku Corporation: Tokyo, Japan.
 34. Sheldrick GM, 2008 Crystal structure refinement with SHELXL *Acta Cryst. A* **64** 112
 35. Dolomanov O V, Bourhis L J, Gildea R J, Howard J A K and Puschmann H 2009 OLEX2: a complete structure solution, refinement and analysis program *J. Appl. Cryst.* **42** 339
 36. (a) Paria S, Halder P and Paine T K 2010 A functional model of extradiol-cleaving catechol dioxygenases: mimicking the 2-his-1-carboxylate facial triad *Inorg. Chem.* **49** 4518–4523; (b) De A, Garai M, Yadav H R, Choudhury A R and Biswas B 2017 Catalytic promiscuity of an iron (II)phenanthroline complex *Appl. Organometal. Chem.* **31** 10.1002/aoc.3551
 37. (a) Sawyer D T 1991 *Oxygen Chemistry* (New York: Oxford University Press); (b) Mialane P, Tehertanov L, Banse F, Sainton J and Girerd J 2000 Aminopyridine iron catecholate complexes as models for intradiol catechol dioxygenases. Synthesis, structure, reactivity, and spectroscopic studies *Inorg. Chem.* **39** 2440
 38. Fereday R J, Hodgson P, Tyagi S and Hathaway B J 1981 The crystal structure and electronic properties of bis (2, 2'-bipyridyl)-copper (II) bis (hexafluorophosphate) *J. Inorg. Nucl. Chem. Lett.* **17** 243
 39. De A, Dey D, Yadav H R, Maji M, Rane V, Kadam R M, Choudhury A R and Biswas B 2016 Unprecedented hetero-geometric discrete copper (II) complexes: crystal structure and bio-mimicking of Catecholase activity *J. Chem. Sci.* **128** 1775
 40. (a) Mukherjee C, Weyhermuller T, Bothe E, Rentschler E and Chaudhury P 2007 A Tetracopper (II)-tetraradical cuboidal core and its reactivity as a functional model of phenoxazinone synthase *Inorg. Chem.* **46** 9895; (b) Barry C E, Nayar P G and Begley T G 1989 A fungal metabolite mediates degradation of non-phenolic lignin structures and synthetic lignin by laccase *Biochemistry* **28** 6323
 41. (a) Chatterjee S, Sheet D and Paine T K 2013 An oxido-bridged diiron (II) complex as functional model of catechol dioxygenase *Chem. Commun.* **49** 10251; (b) Balamurugan M, Vadivelu P and Palaniandavar M 2014 Iron (III) complexes of tripodal tetradentate 4N ligands as functional models for catechol dioxygenases: the electronic vs. steric effect on extradiol cleavage *Dalton Trans.* **43** 14653
 42. (a) Ito M and Que L Jr. 1997 *Angew. Chem. Int. Ed. Engl.* **36** 1342; (b) Dey D, De A, Yadav H R, Guin P S, Choudhury A R, Kole N and Biswas B 2016 *ChemistrySelect* **01** 1910; (c) Dey D, Das S and Biswas B 2016 *J. Indian Chem. Soc.* **93** 495
 43. Ito M and Que L Jr. 1997 Biomimetic extradiol cleavage of catechols: insights into the enzyme mechanism *Angew. Chem. Int. Ed. Engl.* **36** 1342
 44. Funabiki T, Mizoguchi A, Sugimoto T, Tada S, Tsuji M, Sakamoto H and Yoshida S 1986 Oxygenase model reactions. 1. Intra- and extradiol oxygenations of 3, 5-di-tert-butylcatechol catalyzed by (bipyridine)(pyridine) iron (III) complex *J. Am. Chem. Soc.* **108** 2921
 45. Die A, Gatteschi D and Pardi L 1993 Synthesis, characterization, and reactivity of catecholato adducts of iron(III) triaza- and tetraazamacrocyclic complexes: chemical evidence of the role of the metal ion in the oxidative cleavage *Inorg. Chem.* **32** 1389
 46. Pascaly M, Duda M, Rompel A, Sift B H, Meyer-Klaucke W and Krebs B 1999 Novel iron (III) complexes with imidazole containing tripodal ligands as model systems for catechol dioxygenases *Inorg. Chim. Acta* **291** 289
 47. Pascaly M, Nazikkol C, Schweppe F, Wiedemann A, Zurlinden K and Krebs B 2000 Structures and properties

- of novel mononuclear iron(III) complexes with benzimidazole containing tripodal tetradentate ligands *Z. Anorg. Allg. Chem.* **626** 50
48. Nishida Y, Shimo H and Kida S 1994 Synthesis, structural characterization, and extradiol oxygenation of iron-catecholato complexes with hydrotris (pyrazolyl) borate ligands *J. Chem. Soc. Chem. Commun.* **1611**
49. Viswanathan R, Palaniandavar M, Balasubramanian T and Mutiah T P 1999 Functional models for catechol 1, 2-dioxygenase. Synthesis, structure, spectra, and catalytic activity of certain tripodal iron (III) complexes *Inorg. Chem.* **37** 2943
50. Pascaly M, Duda M, Schweppe F, Zurlinden K, Müller F K and Krebs B 2001 The systematic influence of tripodal ligands on the catechol cleaving activity of iron (III) containing model compounds for catechol 1, 2-dioxygenases *J. Chem. Soc. Dalton Trans.* **828**
51. Dey D, Das S, Yadav H R, Ranjani A, Gyathri L, Roy S, Guin P S, Dhanasekaran D, Choudhury A R, Akbarsha M A and Biswas B 2016 Design of a mononuclear copper (II)-phenanthroline complex: catechol oxidation, DNA cleavage and antitumor properties *Polyhedron* **106** 106
52. Speier G, Tyeklár Z, Szabo L, Tóth P, Pierpont C G and Hendrickson D N 1993 In *The Activation of Dioxygen and Homogeneous Catalytic Oxidation* D H R Barton, A E Martell and D T Sawyer (Eds.) (New York: Plenum Press) pp. 423–436