Synthesis, Characterisation and Electrochemical Studies of Cu (II) Complex of (E)-4-(2-(4-Methoxyphenyl) Diazenyl) Benzene 1, 3 Diol Including DNA Binding and Antibacterial Activity

Sukdev Maity¹, PK Saha² and Swapan Kumar Bhattacharya¹*

¹Department of Chemistry, Jadavpur University, Kolkata, West Bengal, India
²Department of Textile Technology, GCETTS, Hooghly, West Bengal, India

ABSTRACT

An innermetallic chelated azo-dye complex of second order having the stoichiometry (Cu (L₁)₂, 2H₂O) with ligand L₁,(E)-4-(2-(4-methoxyphenyl) diazenyl) benzene 1, 3 diol, has been synthesised at pH =7.48 and studied in both solid state and aqueous phase using different spectroscopic techniques. The structure of the complex has been carried out from elemental analysis, UV-Vis, IR, and mass spectrophotometry. Thermal decomposition has been studied from TGA-DSC analysis. Cyclic voltammetry of the complex has been carried out to ensure the presence of Cu (II) in the complex (Cu(L₁)₂,2H₂O). Magnetic susceptibility determination of (Cu (L₁)₂, 2H₂O) complex has been taken at a constant magnetic field of 5 KG with powder sample in the temperature range 20-35K. In addition special attention has been given to the studies of ct-DNA binding to the complex (Cu(L₁)₂, 2H₂O) by absorption spectroscopy and cyclic voltammetry measurements under physiological condition (pH=7.4, 25°C) with intrinsic binding constant in the order of 10⁷ M⁻¹. The complex also shows higher antibacterial activity than L₁.

Keywords: Azo-dye Cu (II) complex; Antibacterial activity: Cyclic voltammetry: DNA binding; TGA; UV-visible

INTRODUCTION

A large number of azo-dyes have been synthesized recently and characterized by number of sophisticated spectroscopic methods. In addition, their wide ranges of analytical, industrial and biological applications have been shown earlier [1]. It is worth mentioning that these azo-dyes can function as good N, O donor chelating ligands and are so capable of forming a number of stable transition metal-azodye complexes both in solution phase and solid states (isolable) [2]. The characterizations of such complexes have also been successfully made [3]. But biological activities of such complexes are very limited. In this study we synthesis, characterization and biological activity of Cu (II) complexes have been reported. This paper also describe the structure elucidation of the Cu (II)-L₁ complexes using different techniques, where ligand L₁ is already reported [4,5]. The special attraction in the paper is the study of DNA interaction with Cu (II) azo dye complex [6,7]. Due to presence of azo-group (N=N) which along with some oxidisable functional groups like – OH, -OR, -NH₂ allow such compounds to interact with cells or DNA that may be probed to study such interaction in several biological activity or in medicine [8,9]. As manyazo-dye compounds have been known to have general toxicity andcarcinogenic in characteristics, transition metal azo-dye complexes have been shown to have significantly reduced toxicity as reported in many journals [10,11]. The present paper works with Cu (II) –L₁ complex who’s binding with calf thymus deoxyribonucleic acid (CT DNA) is significant as considered from data collected from cyclic voltammetry and a series of UV-vis spectra of the complex of Cu (II) with L₁. This study has attracted more attention because of their potential use as drugs, regulators of gene expression and tools for molecular biology [12]. These metal complexes are important as they may find their application as
antifungal, antibacterial, anti-convulsion, anti-inflammatory, anti-malarial, analgesic, platelets, anti-tuberculosis, anti-cancer activity. TGA of this complex has also been studied to predict relative thermal stability of the complex [13].

EXPERIMENTAL SECTION

All chemicals and solvents are of the highest purity taken from commercial suppliers such as Merck, BDH etc. and used without further purification. CuSO₄ (99.9% A.R, BDH) is the starting material for preparing (Cu(L₁)₂.2H₂O) complex. Azo-dye ligand L₁ has been synthesized by previous method [4] and then recrystalised to get 99% of purity. Sodium acetate (anhydrous), triple distilled water, all are of analytical grade used for this purpose. For reference electrolyte used in cyclic voltammetry, phosphate buffer (pH=7.4) of A.R. quality has been used. CT DNA purchased from Sigma Chemical Company was dissolved in triple distilled water. Reference electrolyte used in cyclic voltammetry, phosphate buffer (pH=7.4) of A.R. quality has been used. CT DNA purity. Sodium acetate (anhydrous), triple distilled water, all are of analytical grade used for this purpose. For reference electrolyte used in cyclic voltammetry, phosphate buffer (pH=7.4) of A.R. quality has been used. CT DNA purchased from Sigma Chemical Company was dissolved in triple distilled water.

Synthesis of Cu (L₁)₂ Complex

CuSO₄ is collected and dissolved in 100 ml triple distilled water to get a solution of 0.1(M). To this solution is added saturated solution of L₁ in methanol such that Cu (II):L₁ is 1:2 in molar ratio. The mixture is then taken in a well stoppered conical flask which is then agitated by a magnetic stirrer followed by addition of saturated solution of sodium acetate and heating at constant temperature of 50°C for 2 hour. The mixture is then filtered under suction pump. The black brown solid is repeatedly washed by methanol-water mixture till the filtrate is almost colour-less [6]. The deep brown residue is then re-crystalised.

RESULTS AND DISCUSSION

Physical properties and Elemental Analysis

Physical properties and elemental analysis (C,H,N,O) of Cu(II)L₁ complex are recorded in Table 1. A single spot is located in the TLC, which suggests high purity of the complex. It is also supported by a low percent error (with in 5%) in elemental analysis in Table 2 and sharp decomposition temperature determined from TGA-DSC [13-15].

<table>
<thead>
<tr>
<th>Name of complex</th>
<th>Empirical formula</th>
<th>Relative molar mass</th>
<th>Rvalue</th>
<th>pH</th>
<th>Colour</th>
<th>MP °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cu(L₁)₂.2H₂O</td>
<td>Cu(C₁₁H₁₂N₆O₈)₂.2H₂O</td>
<td>583.50(589.26)</td>
<td>0.71</td>
<td>6.52</td>
<td>Black brown</td>
<td>284</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Name of complex</th>
<th>yield %</th>
<th>Carbon %</th>
<th>Hydrogen%</th>
<th>Nitrogen%</th>
<th>Oxygen%</th>
<th>Copper%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cu(L₁)₂.2H₂O</td>
<td>71</td>
<td>53.49(53.28)</td>
<td>3.80(3.76)</td>
<td>11.42(9.56)</td>
<td>21.40(21.52)</td>
<td>10.82(10.86)</td>
</tr>
</tbody>
</table>

Electronic Spectrum of the Complex (UV-Vis)

The UV-vis spectra for (Cu(L₁)₂.2H₂O) has been recorded in (i) aqueous Solution (10⁻⁵M) and (ii) phosphate buffer, pH=7.4 (10⁻³M). From Electronic spectra (UV-Vis) it is speculated that majority of mono nuclear Cu(II) complexes are either square planar or highly distorted octahedral in nature. This ligand being a chelated N,O donor ligand (azo-N and phenolic O) demands that it might form a stable bis- Cu(II) chelate internally compensated with or without axially populated by two H₂O molecules, which might be thought to be a distorted octahedral complexes [16-20]. So (Cu(L₁)₂(H₂O)₂) will show a Cu(N₂O₂) chromophores. This will give rise to UV-Vis spectrum which exhibited n→π* or π→π* charge transfer band at 420 nm. There will be another broad band observed at 381nm and 254 nm which is attributed to d-d transition [21,22]. The experimentally obtained absorption spectrum (UV-Vis) are graphically represented in Figure 1.
The peaks obtained from aqueous solution are (i) ν₁ = 23809 cm⁻¹, corresponds to the transition ²²B₁g → ²²B₂g, (ii) ν₂ = 26246 cm⁻¹, corresponds to the transition ²²B₁g → ²²A₁g, (iii) ν₃ = 39370 cm⁻¹ corresponds to the transition ²²B₁g → ²²E₂g.

All these are d-d bands where the tail of the blue ends shows a shift to reddish brown [11]. The formations of three peaks is attributed to anticipated distorted octahedral ligand field (Jahn-Tellor distortion). The bands appearing in the UV-Vis spectra of Cu(II) complex in phosphate buffer solution is comparable to that recorded in aqueous solution except the missing of the peak at 420 nm in buffer solution and shifting of the peak at 381 nm is shifted to 443 nm that is towards the blue region. At the alkaline pH (phosphate buffer solution 7.4) there is some chances of producing anionic complex or axial H₂O might be replaced by HPO₄²⁻ ion and intra ligand charge transfer band at 420 nm may merge with the band at 443 nm (d-d).

**Infrared Spectra**

IR- spectrum of L₁ has already been reported [4], where strong bands around 1246.75 cm⁻¹, 1177.33 cm⁻¹, 1107 cm⁻¹ and 1025.94 cm⁻¹ have appeared in conformation with the characteristic band for C-O stretching as mixed band of alkyl C-O and aryl C-O stretching [23]. In the IR spectrum of the complex these bands remain unaffected as shown in the Figure 2, which appear around 1249.65 cm⁻¹, 1187.94 cm⁻¹, 1109.83 cm⁻¹ and 1026.91 cm⁻¹ respectively. By comparing with L₁ it is also suggested that free OH band at 3753.64 cm⁻¹ is a weak stretching band and H-bonded OH band at 3456.88 cm⁻¹ is a broad band, are almost missing or appear as very weak bands with high %T, as presented in the Figure 2 in this complex. It means that one phenolic 'O' is attached to Cu(II) which corresponds to stretching at 519.72 cm⁻¹. There is another bonding Cu-N whose stretching occurs at 442.58 cm⁻¹. It is known that the ligand L₁ is an unsymmetrical ether (CH₂-O-Ar). So the two C-O bond coupled to give anti symmetrical and symmetrical C-O stretching absorption for three different IR bands appeared and already mentioned earlier. The bands appearing at 1460.81 cm⁻¹ and 1361.5 cm⁻¹ are distinctly less intense than that obtained in free ligand L₁ in bending frequency of ν₁=ν₁. This lowering in stretching of azo group might correspond to partial drifting of the π-electron cloud towards metal in addition to Cu(II)-N σ-bonding. Consequently N-N bond order decreases and finally it assumes an approximate Cu(II)-N (=N) σ-bonding might go into partial chelation (haptactility). Linear stretching of N=N is absent as it is IR inactive (Figure 2). The band appearing between 1500 cm⁻¹-1600 cm⁻¹ (1595.81 cm⁻¹, 1505.17 cm⁻¹ and %T=30.5732 and 48.8876) obviously indicate C=C (aromatic) bonding which is not appeared in complex [24]. Band offering at 2939.95 cm⁻¹ is very close to Ar-H bonding of theoretically known to be 3030 cm⁻¹.

**Mass Spectral Studies**

It is predicted from C,H,N analysis that molecular formula of Cu(II) L₁ complex is [Cu(L₁)₂·2H₂O] This formula having relative molar mass 585.5 is roughly in agreement with m/z peak at 588.16, a value having 0.6% error might be due to adhering moisture. This complex is primarily considered to an internally compensated with L₁ anions in a square planer ligand field as observed from analysis of number of square planer Cu(II) complex with Cu(N₂O₅) chromophores [11]. Two water molecules may have two different attachments that either two axial sites are occupied by two H₂O molecules or two H₂O molecules may be H-bonded with free phenolic –OH groups. Another (m/z) peak appearing at 610.76 might appear due to knocking out H₂O molecules and attachment of OAc⁻. The impurity OAc⁻ comes from the NaAc soln. which was used during synthesis of the complex. The composition

![Figure 1: UV-Vis of the Cu(II) complex in (a) aqueous Solution and (b) buffer solution](image-url)
(Cu (L) _2(OAc) ) _2 can not be ruled out as Cu^{+2} may have an affinity towards OAc^- and the peak m/z appearing at 668.71 which is the calculated Value of mass as 667.57 the nearby pecks at 662.72, 664.79, 661.69 might be generated by loss H^+ from the complex successively [25,26]. The high peak m/z appearing at 706.26 should indicate the stoichiometry (Cu (L) _2(OAc) ) _2, 2H_2O, due to presence of two H_2O molecules as water of crystallization. Attachment of H_2O to H-bonding with free phenolic OH or axially located H_2O is not unlikely (Figure 3). Loss of one H_2O is possible, which causes another peak characterized by m/z at 688 very close to actually observed peak at 684.29 [27].

**Figure 2:** IR spectrum of Cu(II) complex

**Figure 3:** Mass spectra of sodium salt of Cu(II) complex

**TGA-DSC Studies**

The thermo gravimetric analysis (Figure 4) gives information about the thermal stability of the complex and suggests a general scheme for thermal decomposition of the Cu(II) chelate complex. In the present investigation, heating rates were suitably controlled at 10°C min^-1 under nitrogen atmosphere [12,13,17]. The thermo gram of the Cu (II) complex shows three decomposition steps (Table 3) within the temperature range 25-294°C. The first step involves loss of non-co-ordinated water molecules around 60-106°C with an estimated mass loss 6.53% (calculated mass loss 7.21%), the second step involves loss of two co-ordinated water molecules around 106 - 208°C with an estimated mass loss 12.19% (calculated mass loss 10.32%) and the third step involves loss of one molecule of ligand around 208-294°C with an estimated mass loss 27.46% (calculated mass loss 27.71%). This thermo gram is (Figure 4) accompanied by three exothermic peak at103°C, 190°C, 207°C and 243°C on the DSC curve [18].
Electrochemical Studies of Cu (II) Complex with L₁
Cyclic voltammetry experiments were performed using model number DY2300 series potentiostat, Digi-IVY. The experiments were carried out using the conventional three-electrode system at 25°C. A Pt disc electrode served as the working electrode A calomel electrode Saturated with KCl was used as reference electrode while a platinum wire served as the counter electrode. Electrochemical measurements were performed in a 10 mL electrochemical cell. 5 × 10⁻³(M) metal complex solution was prepared using phosphate buffer (pH=7.4) solution which was used as supporting electrolyte [1,3,14].

The Cu(II) complex of L₁ showed two reduction (cathodic) peaks at +0.30 V (Cu²⁺/Cu⁺) and -0.059V (Cu⁺/Cu) respectively in the Figure 5. For the Cu(II)L₁ complex the first reduction was attributed to reduction of Cu²⁺ + e → Cu⁺ due to single electron gain, while the second reduction was of Cu⁺ + e → Cu due to another single electron gain. In addition to these two reduction (cathode) peaks for the complex, there was one very sharp oxidation (anodic) peak at + 0.135 V which was due to generation of Cu/Cu’ species and another peak at +0.36 V due to conversion of Cu⁺ - e → Cu²⁺. This was supported by a recent finding where an almost identical condition was used for a Cu(I) complex

Figure 5: Cyclic voltammogram of (Cu(L₁)₂2H₂O) complex is recorded using platinum electrode at scan rate 50 mVs⁻¹
Plot of anodic peak current $I_{pa}$ with square root of scan rate ($\sqrt{v}$) being linear, the oxidation for Cu(I) complex (Figure 2) indicate a diffusion controlled process with no adsorption on the electrode surface [11].

**DNA Binding Studies**

**Absorption spectral measurements:**

All experiments involving calf thymus DNA (CT- DNA) were performed in Tris- buffer solution (50 mM NaCl/5 mM Tris–HCl, pH 7.4) at 25 ± 0.2°C. Double distilled water was used to prepare the buffer solution. The concentration of CT- DNA was determined from the intensity of absorbance at 260 nm with a known extinction coefficient value ($\varepsilon_{260}=6000$ M$^{-1}$ cm$^{-1}$) [1]. The ratio of the absorbance of CT DNA at 260 nm and 280 nm was found as 1.85. Therefore, no further purification was attempted [6]. Absorption titration measurements were carried out by varying the concentration of CT DNA from 0 to $10 \times 10^{-6}$ M, while keeping the metal complex concentration constant at $12 \times 10^{-6}$ M. Samples were incubated at 25 ± 0.2°C for 24 hour before recording each spectrum (Figure 6). The intrinsic binding constant ($K_b$) for the interaction of the complex with ctDNA was determined using the following equation [11].

\[
\frac{(\text{DNA})}{(\varepsilon_a - \varepsilon_f)} = \frac{(\text{DNA})}{(\varepsilon_b - \varepsilon_f)} + \frac{1}{K_b (\varepsilon_b - \varepsilon_f)} (1)
\]

where (DNA) is the concentration of CT DNA, the apparent absorption coefficients $\varepsilon_a$, $\varepsilon_f$ and $\varepsilon_b$ correspond to $A_{obsd}/(\text{Cu})$, the extinction coefficient for the free Cu(II) complex and the extinction coefficient for the Cu(II) complex in the fully bound form, respectively. A plot of $(\text{DNA})/(\varepsilon_a - \varepsilon_f)$ vs (DNA) gave a slope of $1/(\varepsilon_b - \varepsilon_f)$ and a Y-intercept, $1/K_b (\varepsilon_b - \varepsilon_f)$, $K_b$ is the ratio of the slope to the Y-intercept (Figure 7). The absorption spectra of the complex in the absence and presence of increasing amounts of CT-DNA concentration are shown in Figure 6. With increasing concentrations of DNA, the complex exhibited hypochromism with slight red shifts of the absorption bands at 443 nm observed in the presence of DNA can be assigned to LMCT transitions in the absorption spectra of all complexes. The changes in the absorbance values with increasing amounts of CTDNA were used to evaluate the intrinsic binding constant $K_b$. Based on the hypochromism exhibited and shifts in absorbance upon addition of CT DNA, non-intercalative interaction probably by an electrostatic interaction between complex ions and negatively charge phosphate groups of the CT DNA can be predicted [6]. However, since DNA possesses several hydrogen bonding sites which are accessible both in the minor and major grooves, a favorable hydrogen bonding may be formed between the coordinated and non-coordinated amine $\text{–NH–}$ groups of the complex with the base pairs in CT DNA. Further, the intrinsic binding constant ($K_b$) of the complex calculated by using Eq. (1) is found to be $1.69 \times 10^4$ M$^{-1}$.

**Figure 6:** Absorption spectra of Cu(II) complex in the absence and in the presence of increasing concentration of CT DNA; the top most spectrum is recorded in the absence of DNA and spectra bellow on successive addition of 1 μl DNA to it.
The data obtained from the spectrophotometric titration of Cu(II) complex with ct DNA was analyzed by Scatchard plot [1]. The intrinsic binding constant ($K'$) and site size ($n_b$) were determined directly using eqn. (5):

$$\frac{r}{c_f} = K' (n-r) \quad (5)$$

$r = C_b/C_D$ where ‘$C_b$’ is the concentration of bound compound and ‘$C_D$’ concentration of ct DNA ‘$C_f$’ refers to the concentration of free compounds. $K'$ is $(1.74 \times 10^4)$ the intrinsic or the overall binding constant of any molecule with a substrate. ‘$n$’ is (3.5) the binding stoichiometry in terms of number of bound compound (Cu(II) complex) per nucleotide while ‘$n_b$’ is (0.28) the reciprocal of ‘$n$’ i.e., the binding site size in terms of number of nucleotide per molecule of the compounds that were used. The overall binding constant $K'$ is also related to $K_{app}$ by $K' = K_{app} \times n_b$ [1]. Therefore knowing $n_b$ and using the $K_{app}$ values the intrinsic binding constant $K'$ could be evaluated for the interaction of the Cu(II) complex with ct DNA. The $K'$ values obtained in this manner for complex was compared with values obtained directly from Scatchard plots (Figure 8). Thus using different modes of evaluation both apparent binding constant ($K_{app}$) and intrinsic binding constant ($K'$) were obtained from where an idea of the strength of interaction of the Cu(II) complex with ct DNA could be obtained.

**Cyclic voltammetric measurements:**

The voltammogram of Cu(II) azo-dye complex showed steady redox peaks in the potential range of +1.0 to -1.0 V in either scan i.e., in forward scan anodic peak and in reverse scan cathodic peak appeared [6]. This cyclic voltammogram of Cu(II) azo-dye complex showed one electron transfer and the reaction is totally reversible electrochemical reaction. By mixing 30, 60, 90, 120 and 150 μM CT-DNA into 2 mM Cu(II) azo-dye complex solution (Figure 9) the change in peak potential and decrease in current $i_{pa}$ was being seen. The decrease in peak current $i_{pa}$ is due to diffusion of Complex into double helix DNA which resulted supramolecular complex formation, due to which transfer of electrons was being reduced as a result number of free molecules was being
decreased. The change in value of formal potential explained the nature of binding between Cu(II) azo-dye complex and DNA. Generally intercalation of small molecules into double helical deoxyribonucleic acid caused positive change in the peak potential, whereas negative change revealed binding of the positively charged molecule with the negatively charged phosphate, \((\text{PO}_4)^{3-}\) moiety present on DNA backbone called the electrostatic interaction [8]. The negative change in peak potential was observed for Cu(II) azo-dye complex by the addition of different concentration of CT-DNA, revealed the electrostatic nature of interaction. The binding constant can be calculated by using following equation: \(1/(\text{DNA}) = K(1-A)/(1-(i/i_0) - K)\), where \(i_0\) and \(i\) are the peak currents in absence and presence of CT-DNA, \(K\) is the binding constant, and \(A\) is the proportionality constant. If we plot a graph between \(1/(\text{DNA})\) and \(1/(1-i/i_0)\), binding constant \((K)\) can be calculated which was \(2.02 \times 10^4\) M\(^{-1}\) (Figures 10 and 11). The changed binding free energy \((-\Delta G = RT \ln K\) at 25°C\) of compound Cu(II) azo-dye complex was calculated to be 30.25 kJ mol\(^{-1}\) exhibited the spontaneity of Cu(II) azo-dye complex -DNA interaction.

The diffusion coefficient of free compounds and compound-DNA adduct was calculated from Randles-Sevcik equation; \(I_{pa} = 2.69 \times 10^5 n^{3/2} A C_0^* D_0^{1/2} v^{1/2}\) where \(I_{pa}\)is referred to anodic peak current in ampere, \(v\) referred as scan rate in V s\(^{-1}\), \(C_0^*\) is concentration in mol cm\(^{-3}\), \(A\) is cross sectional area of electrode in cm\(^2\), \(n\) is number of electrons involved in the reaction, \(D_0\) is diffusion coefficient in cm\(^2\) s\(^{-1}\). The diffusion coefficient of free Cu(II) azo-dye complex was calculated 4.609 \times 10^{-3} whereas diffusion coefficient of Cu(II) azo-dye complex -DNA was found to less i.e., 1.918 \times 10^{-3}. The decreased diffusion coefficient value for Cu(II) azo-dye complex -DNA adduct can be justified as free molecules are easy to diffuse and is of low molecular weight so exhibit more peak current whereas when compound Cu(II) azo-dye complex was interacted with DNA, the quantity of free molecules became less with the obvious a result of decrease in current.

Figure 9: Cyclic voltammograms of 2 mM Cu(II) azo-dye complex in the absence and presence of 30 μM, 60 μM , 90 μM , 120 μM and 150 μM DNA showing a decrease in I from I, and a -ve shift in peak potential indicating electrostatic interactions

Figure 10: The plot of \((i/\Delta-\iota)\)vs \(1/(\text{DNA})\) for determination of binding constant
Antibacterial activity:
The Cu(II) complex was tested against the bacteria Escherichia coli, Bacillus subtilis, Staphylococcus aureus, S. monnellatyphi. Diameter of incubation zone: 15 mm, concentration of bacterial growth 200 μg/mL, in DMSO solution (\*): Inactive. The data in the Table 4, it is observed that the Cu(II) complex is more active against all the test organisms, Bacillus subtilis and Staphylococcus aureus and as well as more active compared to the L1. Such increased activity of the Cu(II) complex can be explained with respect to Overtone’s concept and Tweedy’s chelation theory [8]. According to Overtone’s concept of cell permeability, the lipid membrane that surrounds the cell favours the passage of only the lipid-soluble materials whose liposolubility is an important factor, which controls the antibacterial activity. On chelation, the polarity of the Cu(II) ion is reduced [6,12] to a great extent due to the overlap of the ligand, L1 orbital and partial sharing of the positive charge of the Cu(II) ion with donor groups. On the other hand it increases the delocalization of π electrons over the whole chelate ring and enhances the lipophilicity of the Cu(II) complex. This increased lipophilicity enhances the penetration of the Cu(II) complex into lipid membranes and blocking of the cobalt metal binding sites in the enzymes of micro-organisms. This Cu(II) complex also disturb the respiration process of the cell and thus block the synthesis of proteins, which restricts further growth of the organisms.

Table 4: Antibacterial activity of Cu(II) complex

<table>
<thead>
<tr>
<th>Metal Complex</th>
<th>Rate of oxidation (Antibacterial activity)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cu(L1)2.2H2O</td>
<td>Control</td>
</tr>
<tr>
<td></td>
<td>7</td>
</tr>
</tbody>
</table>

CONCLUSION

The complex of Cu(II) azo-dye has been synthesized and structure of the complex has been established by elemental analysis, IR, electronic, mass spectroscopy, molar conductance. Electronic spectral data suggest that the complex has distorted octahedral geometry. The complex exhibits good redox property. The complex had shown significant binding with CT-DNA by non-intercalative mode i.e., Electrostatic interactions. The complex act as potent bactericidal agent. Further work with analogs is needed.

ACKNOWLEDGEMENT

This work has been funded by Jadavpur University in the form of a “J.U Research Grant” to Prof. Dr. Swapan Kumar Bhattacharya, Department of Chemistry, Jadavpur University. The authors are thankful to the authorities of Chemistry Department of Jadavpur University and Department of Textile Technology, GCETTS, HOOGHLY for allowing them to pursue this research work.
REFERENCES