Synthesis, Spectroscopic, Structural Characterization, Conductivity and Electrochemical Studies of a Schiff Base Ligand and Its Copper Complexes

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Abstract

Schiff base ligand (L) derived from glyoxal and 4-aminoantipyrine was synthesized. The ligand (L) has been characterized by IR, NMR, electronic spectral studies and electrochemical studies. Cu(II) complexes of a Schiff base ligand (L) from 4-aminoantipyrine and glyoxal having the composition [CuL₁]X₂ where X = Cl⁻ or NO₃⁻ have been prepared and characterized by elemental analysis, electrical conductivity in non-aqueous solvent, infrared and electronic, as well as cyclic voltammetric studies. L acts as a neutral tetra-dentate ligand coordinating through both the carbonyl oxygen and azo-methine nitrogen. On both the complexes both the anions are not coordinated. A square planar geometry is assigned for complexes. The electrochemical studies of ligand show a typical cyclic voltammogram for an irreversible process. While copper(II) complexes show the typical cyclic voltammograms for quasi reversible process.

Keywords

Schiff Base Ligand, Copper(II) Complexes, Cyclic Voltammetry

1. Introduction

A large number of Schiff bases and their complexes have been studied for their interesting and important properties, e.g; their ability to reversibly bind oxygen [1], catalytic activity in hydrogenation of olefins and transfer of an amino group [2], photochromic properties [3], and complexing ability towards some toxic
metals [4]. The high affinity for the chelation of the Schiff bases towards the transition metal ions is utilized in preparing their solid complexes.

Schiff bases are condensation products of primary amines and carbonyl compounds and they were discovered by a German chemist, Nobel Prize winner, Hugo Schiff in 1864 [5]. In the recent decades, Schiff bases have attracted tremendous interests due to their unique properties and extensive applications in many scientific areas, e.g. anticancer [6] and antibacterial [7], biosensor [8], catalysis [2], analytical chemistry [9] and corrosion prevention [10].

Structurally, Schiff base (also known as imine or azomethine) is an analogue of a ketone or aldehyde in which the carbonyl group (C=O) has been replaced by an imine or azomethine group. Schiff bases are compounds having a formula RR'C = NR'' where R is an aryl group, R' is a hydrogen atom and R'' is either an alkyl or aryl group. However, usually compounds where R'' is an alkyl or aryl group and R’ is an alkyl or aromatic group are also counted as Schiff bases. Schiff base ligands are essential in the field of coordination chemistry, especially in the development of complexes of Schiff bases because these compounds are potentially capable of forming stable complexes with metal ions [11].

During the past two decades, considerable attention has been paid to the chemistry of the metal complexes of Schiff bases containing nitrogen and other donors [12]. This may be attributed to their stability, biological activity and potential applications in many fields such as oxidation catalysis, electrochemistry, etc. In this study, the synthesis of a Schiff base ligand (L) and its Cu(II) complexes were reported. Their spectral properties and electrochemical behavior were investigated.

2. Experimental Part

2.1. Instrumental Measurements

IR spectra were recorded on a PERKIN ELMER SPECTRUM 65 FT-IR spectrometer on KBr pellet in the wave number range of 4000 - 400 cm⁻¹. Electronic spectral studies were conducted on a GENESY’s UV-Visible spectrometer in the wavelength 200 - 800 nm. Nuclear Magnetic Resonance (NMR) analysis was recorded on a Brukeravance 400 MHz spectrometer with tetramethylsilane as internal standard and DMSO-d⁶ as solvent. Molar conductivity of the complexes were measured for 10⁻³ Molar solution in acetonitrile using a SX713 model conductivity meter. All the electrochemical experiments were carried out using a computerized electrochemical analyzer (BAS CV-50 W) with a conventional three electrode system at room temperature with a glassy carbon electrode (diameter = 3 mm) as working electrode, Ag/AgCl/ as reference electrode and a platinum wire as counter electrode in DMF and electrochemical grade Tetra butyl ammonium hexafluoro phosphate (0.01 mol/dm³) as the supporting electrolyte.

2.2. Solvents and Reagents

The metal salts were prepared from Analar BDH copper carbonate and the re-
spective 50% acids (AR) and crystallising of the salts by evaporation of the solutions on a steam bath. All the solvents used in the present study is of analytical Grade and used without any further purification. Tetrabutylammonium hexafluorophosphate (Fluka), Glyoxal (Aldrich Chem. Co. USA), 4-Aminoantipyrene (Sigma Chem. Co. USA) were also used without any further purification.

2.3. Preparation of the Schiff Base

The Schiff base ligand was prepared by the condensation of the 0.44715 g (2.2 mmol) 4-Aminoantipyrene with 0.057 ml (1 mmol) Glyoxal in absolute ethanolic solution (Scheme 1). The resulting mixture was then refluxed for 2 hrs. The yellow precipitate formed was filtered and recrystalized from ethanol to give yellow needles.

2.4. Preparation of the Schiff Base Copper(II) Complexes

Cupric chloride 0.170 g (1 mmole) and copper nitrate 0.18756 g (1 mmole) were dissolved in methanol 5 ml and added in to a refluxing solution of Schiff base 0.4280 g (1 mmole) in ethyl acetate. The reaction mixtures were refluxed for 3 hrs. The complexes were separated as brown solid is cooled, filtered and repeatedly washed with hot ethyl acetate to remove excess ligand if any. It was then dried in a vaccum in a desiccator over P2O5.

2.5. Electrochemical Studies

Electrochemical cyclic voltammetry measurements were performed at room temperature in 3-electrode cell by using a glassy carbon electrode with 0.071 cm² surface area as a working electrode, while a platinum wire served as the counter electrode and a Ag/AgCl quasi reference electrode. A DMF solution of all the ligand and complexes (1 × 10⁻⁴ M) and tetrabutylammonium hexafloro-phosphate (0.01 M) as supporting electrolyte were used in each measurements. Measurements were made over a potential range between 0 V to +1.6 V for Schiff base ligand (L) while 0 V to 0.9 V for the complexes with a scan rate of 0.1 V/s.

3. Results and Discussion

3.1. Proton NMR Spectra of Ligand (L)

The ¹H NMR spectrum (Figure 1) of the free ligand showed a singlet at 9.25 ppm due to the imine protons, multiplet in the range 7.1 - 8.0 ppm due to the aromatic protons, signals appearing at 3.10 - 3.50 δ correspond to methyl protons near to hetero-cyclic atoms and signals at 2.8 - 2.10 δ correspond to methyl protons [13] (Table 1).

The Copper(II) complexes are brown solids (Table 2). They are soluble in acetonitrile, dimethyl formamide, ethanol, and methanol, but insoluble in ethyl acetate.

The molar conductance values of the chloride and nitrate complexes of L₁ (10⁻³ M solution) in acetonitrile, fall in the range expected for 1:2 electrolytes.
Scheme 1. Synthesis of Schiff base ligand (L).

Figure 1. Proton NMR spectra Schiff base ligand L.

Table 1. Chemical shift data of ligand.

<table>
<thead>
<tr>
<th>Compound</th>
<th>Chemical shift (δ, ppm)</th>
<th>Assignment</th>
</tr>
</thead>
<tbody>
<tr>
<td>L</td>
<td>9.25</td>
<td>CH = N imine protons</td>
</tr>
<tr>
<td></td>
<td>7.8 - 7.1</td>
<td>Phenyl ring protons</td>
</tr>
<tr>
<td></td>
<td>3.5 - 3.1</td>
<td>Protons near to hetero-atoms</td>
</tr>
<tr>
<td></td>
<td>2.8 - 2.1</td>
<td>Methyl protons</td>
</tr>
</tbody>
</table>

Table 2. Elemental analyses and electrical conductance data of copper(II) complexes.

<table>
<thead>
<tr>
<th>Compound</th>
<th>M. Wt</th>
<th>Color</th>
<th>% Cu (calcd.)</th>
<th>% Cu (Found)</th>
<th>Molar conductance (Ω⁻¹·Cm²·mol⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>[CuL₁]Cl₂</td>
<td>562.5</td>
<td>Brown</td>
<td>(11.28)</td>
<td>(11.2)</td>
<td>240</td>
</tr>
<tr>
<td><a href="NO%E2%82%83">CuL₁</a>₂</td>
<td>615.5</td>
<td>Dark brown</td>
<td>(10.35)</td>
<td>(10.32)</td>
<td>235</td>
</tr>
</tbody>
</table>

[14]. Thus the complexes may be formulated as [CuL₁]X₂ (where X = NO₃⁻ or Cl⁻).

3.2. Infrared Spectra of Ligand and Its Copper Complexes

The important infrared spectral bands of L and its copper complexes with the tentative assignments are presented in Table 3 and Figures 2-4.
Table 3. Important infrared spectra bands (cm$^{-1}$) of ligand and its copper complexes.

<table>
<thead>
<tr>
<th>Compound</th>
<th>$\nu$(C=O)</th>
<th>$\nu$(C=N)</th>
<th>$\nu$(M-N)</th>
<th>$\nu$(M-O)</th>
<th>$V_2$ (ionic nitrate)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ligand (L)</td>
<td>1659</td>
<td>1563</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>$[\text{Cu(L)}_2]\text{Cl}_2$</td>
<td>1635</td>
<td>1584</td>
<td>502</td>
<td>590</td>
<td></td>
</tr>
<tr>
<td>$[\text{Cu(L)}_2]\text{(NO}_3)_2$</td>
<td>1620</td>
<td>1584</td>
<td>520</td>
<td>573</td>
<td>1374</td>
</tr>
</tbody>
</table>

Figure 2. IR spectrum of ligand.

Figure 3. IR spectrum of chloride complex.

Figure 4. IR spectrum of nitrate complex.
The infrared spectrum of L exhibits two strong bands at 1659 and 1653 cm\(^{-1}\) corresponding to the stretching vibrations of the carbonyl groups of the ligand, also exhibits a strong band at 1563 cm\(^{-1}\) due to the C=N stretching vibration \cite{15}. The infrared band at 1659 cm\(^{-1}\) characteristic of the \(\nu_{\text{C}=\text{O}}\) is shifted to 1635 cm\(^{-1}\) in the chloride complex and 1620 in nitrate complex indicating, the coordination of both carbonyl oxygen to the central metal ion. Also the band at 1563 cm\(^{-1}\) corresponds to C=N stretching is shifted to 1584 cm\(^{-1}\) in chloride and nitrate complex indicating the coordination of both azomethine nitrogens to the central metal ion. A strong peak at 1374 cm\(^{-1}\) which is corresponds to the \(\nu_{\text{out of plane}}\) deformation of the ionic nitrate which is concordance with the conductance data \cite{15}. Further the \(\nu_{\text{Cu-O}}\) and \(\nu_{\text{Cu-N}}\) stretching vibrations are observed at about 520 and 502 cm\(^{-1}\) respectively in both complexes.

3.3. Electronic Spectral Studies

The electronic spectral bands of L and its copper complexes with tentative assignments are presented in Table 4 and Figures 5-7.

The electronic spectra of L shows two band maxima at 26,881 and 39,525 cm\(^{-1}\) corresponding to \(n \rightarrow \pi^*\) and \(\pi \rightarrow \pi^*\) transitions respectively \cite{16}. In copper(II) complexes, both the \(n \rightarrow \pi^*\) and \(\pi \rightarrow \pi^*\) bands are found to be blue shifted and appeared in the region 26,178 - 27,322 cm\(^{-1}\) and 39,840 - 45,248 cm\(^{-1}\) respectively compared to that of L. The spectra of nitrate complex exhibit a band at 21,739 cm\(^{-1}\) corresponding to the d-d transition consistent with square planar copper(II) complexes. The electronic spectra of chloride complex shows three band maxima at 45,248 and 39,840 cm\(^{-1}\) for \(\pi \rightarrow \pi^*\) and 27,322 cm\(^{-1}\) corresponding to \(n \rightarrow \pi^*\) transitions respectively. In chloride complex, the d-d transition cannot be visible because it may be masked by the strong MLCT \cite{16}.

Based on the above studies the following tentative structures can be proposed for the complexes (Figure 8).

3.4. Cyclic Voltammetry

The ligand (L) and copper(II) complexes were subjected to cyclic voltammetric studies with a view to examine its electrochemical behaviour. A glassy carbon electrode was used as working electrode, Ag/AgCl as reference electrode and

<table>
<thead>
<tr>
<th>Compounds</th>
<th>Abs. max. cm(^{-1})</th>
<th>Tentative assignment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ligand (L)</td>
<td>39,525</td>
<td>(n \rightarrow \pi^*)</td>
</tr>
<tr>
<td></td>
<td>26,881</td>
<td>(\pi \rightarrow \pi^*)</td>
</tr>
<tr>
<td>[CuL(_1)]Cl(_2)</td>
<td>45,248</td>
<td>(n \rightarrow \pi^*)</td>
</tr>
<tr>
<td></td>
<td>39,840</td>
<td>(\pi \rightarrow \pi^*)</td>
</tr>
<tr>
<td></td>
<td>27,322</td>
<td>(n \rightarrow \pi^*)</td>
</tr>
<tr>
<td><a href="NO(_3)">CuL(_1)</a>(_2)</td>
<td>40,000</td>
<td>(n \rightarrow \pi^*)</td>
</tr>
<tr>
<td></td>
<td>26,178</td>
<td>(\pi \rightarrow \pi^*)</td>
</tr>
<tr>
<td></td>
<td>21,739</td>
<td>d-d transition</td>
</tr>
</tbody>
</table>
Figure 5. UV-Vis spectrum of ligand.

Figure 6. UV-Vis spectrum of chloride complex.

Figure 7. UV-Vis spectrum of nitrate complex.

Figure 8. Tentative structures of the Cu(II) complexes of L (X = Cl⁻ or NO₃⁻).
platinum wire as auxiliary electrode. The electrochemical studies were conducted with DMF solution of all the ligand and complexes (1 × 10⁻⁴ M) and tetrabutylammonium hexafloro-phosphate (0.01 M) as supporting electrolyte. Measurements were made over a potential range between 0 V to +1.6 V for Schiff base ligand and 0 V to 0.9 V for the corresponding complexes with a scan rate of 0.1 V/s (Table 5).

Table 5. Voltammetric data for ligand and complexes in DMF solution containing 0.01 M Tetrabutylammonium hexafluorophosphate. Scan rate (v) =100 mV/s.

The CV of the Schiff base ligand (L) (Figure 9) which exhibited one irreversible oxidation peak as a result of a large peak separation value (ΔEp = 819 mV) at 0.100 V/s scan rate. A cyclic voltammogram of Cu(II) chloride complex displays a reduction peak at Epc = −326 mV with an associated oxidation peak at Epa = 712 mV at a scan rate of 100 mV/s. The peak separation of this couple (ΔEp) is 386 mV. The ratio of anodic to cathodic peak height was greater than one. However, the peak current increases with the increase of the square root of the scan rates. This establishes the electrode process as diffusion controlled [17]. The copper nitrate complex exhibited two quasi-reversible peaks. The representative CV of copper nitrate complex is shown in Figure 10. A cyclic voltammogram of nitrate (Figure 11) displays two reduction peaks. On the first positive potential sweep, the complex shows an anodic oxidation peaks at Epa(I) = 695 mV and Epa(II) = 1064 mV. The complementary reduction peaks on the negative potential sweep appear at Epc(I) = 134 mV and Epc(II) = 389 mV; corresponding to the Cu(II)/Cu(I). The value of ΔEp is 561 mV and 675 mV for first and second redox couples respectively and increases with scan rate giving evidence for quasi-reversible nature associated with one electron reduction. The ratio of the anodic to cathodic peak current (ipa/ipc) is deviates from one. From these observations it is concluded that the redox process is diffusion controlled.

3.5. Effect of Scan Rate

The effect of scan rate could be shown by recorded the CV at concentration of 1 × 10⁻⁴ M of Schiff base ligand. Schiff base ligand L shows the typical cyclic voltammogram for an irreversible process (Figure 12 & Figure 13). The most obvious indication is the absence of a cathodic reduction signal. Furthermore the oxidation signals significantly shift to more positive potentials with faster scan rates.

Table 5. Voltammetric data of ligand and Cu(II) complexes.

<table>
<thead>
<tr>
<th>Scan rate (mV/s)</th>
<th>Ip (µA)</th>
<th>Ip (µA)</th>
<th>ipa/ipc</th>
<th>Epc (mV)</th>
<th>Epa (mV)</th>
<th>ΔEp (mV)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A 100</td>
<td>0.04</td>
<td>-</td>
<td>819</td>
<td>819</td>
<td>819</td>
<td></td>
</tr>
<tr>
<td>B 100</td>
<td>59.2</td>
<td>86.2</td>
<td>1.456</td>
<td>326</td>
<td>712</td>
<td>386</td>
</tr>
<tr>
<td>C 100</td>
<td>8.9</td>
<td>20.4</td>
<td>2.29</td>
<td>389</td>
<td>1064</td>
<td>675</td>
</tr>
</tbody>
</table>
Figure 9. Cyclic voltammogram of $1 \times 10^{-4}$ M of Schiff base ligand at glassy carbon electrode in DMF solution containing 0.01 M Tetrabutylammonium hexafluorophosphate. Scan rate ($v$) = 100 mV/s.

Figure 10. Cyclic voltammogram of $1 \times 10^{-4}$ M of Cu(II) chloride complex at glassy carbon electrode in DMF solution containing 0.01 M Tetrabutylammonium hexafluorophosphate. Scan rate ($v$) = 100 mV/s.

Figure 11. Cyclic voltammogram of $1 \times 10^{-4}$ M of Cu(II) nitrate complex at glassy carbon electrode in DMF solution containing 0.01 M Tetra butyl ammonium hexafluoro phosphate. Scan rate ($v$) = 100 mV/s.
Cyclic voltammograms of Schiff base ligand on glassy carbon electrodes in 0.01 M tetrabutylammonium hexaflorophosphate at different scan rates (mV/s): 10 - 280 mV/s.

Figure 12. Cyclic voltammograms of Schiff base ligand on glassy carbon electrodes in 0.01 M tetrabutylammonium hexaflorophosphate at different scan rates (mV/s): 10 - 280 mV/s.

Figure 13. Plot of anodic peak current of Schiff base ligand L as a function of square root of scan rate.

A 1 × 10^{-4} M concentrations of copper(II) complexes show the typical cyclic voltammograms for quasi reversible process. The cyclic voltammogram will take longer to record as the scan rate is decreased. At a slow scan rate, the diffusion layer will grow much further from the electrode as compared to a fast scan. This leads to a concentration gradient to the electrode surface that is much lower as compared to a fast scan. The peak heights of the anodic signal and the cathodic signal are not completely equal anymore. However, the most obvious indication that the process is not completely reversible anymore is the separation of anodic and cathodic peak potential. The difference of the peak potentials is significantly bigger than 59 mV and the separation of the two signals increases with faster scan rates [18]. Therefore the reaction of copper(II) complexes at higher scan rates can be considered to be quasireversible (Figure 14 & Figure 15).

For both chloride and nitrate complexes the graph of \( i_{pc} \) and \( i_{pa} \) against \( v^{1/2} \) gave a linear plot with \( R^2 \) greater than 0.98 (Figure 16 & Figure 17). This indicates that the \( i_{pc} \) and \( i_{pa} \) were directly proportional to the square root of the scan rate.
Figure 14. Cyclic voltammograms of chloride complex on glassy carbon electrodes in 0.01 M tetrabutylammonium hexafluorophosphate at different scan rates (mV/s): 10 - 320 mV/s.

Figure 15. Cyclic voltammograms of nitrate complex on glassy carbon electrodes in 0.01 M tetrabutylammonium hexafluorophosphate at different scan rates (mV/s): 10 - 320 mV/s.

Figure 16. Plot of anodic peak current and cathodic peak current of chloride complex as a function of square root of scan rate.
rate. From the graph, it also shows that as the scan rate increases, the peak current also increases.

4. Summary and Conclusion

The Schiff base ligand (L) and its copper(II) complexes were synthesised and characterised by elemental analyses, molar conductance in non-aqueous solvents, infrared and electronic spectra NMR as well as spectra.

The complexes of L have the general formulae \([\text{Cu} (\text{L})_n \text{X}_2]\) (where \(X = \text{Cl}^−, \text{NO}_3^−\)). The ligand L acts a neutral tetradeinate ligand, coordinating through both the carbonyl oxygens and both the azomethine nitrogens in the complexes. Both counter anions remain ionic in the complexes. Electronic spectral studies suggest a square planar geometry around the Cu(II) ion in both the complexes.

The electrochemical studies of ligand show a typical cyclic voltammogram for an irreversible process. The most obvious indication is the absence of a cathodic reduction signal. Furthermore the oxidation signals significantly shift to more positive potentials with faster scan rates.

Copper(II) complexes show the typical cyclic voltammograms for quasi reversible process. The cyclic voltammogram will take longer to record as the scan rate is decreased. At a slow scan rate, the diffusion layer will grow much further from the electrode as compared to a fast scan. This leads to a concentration gradient to the electrode surface that is much lower as compared to a fast scan. The peak heights of the anodic signal and the cathodic signal are not completely equal. However, the most obvious indication that the process is not completely reversible anymore is the separation of anodic and cathodic peak potential. The difference of the peak potentials is significantly bigger than 59 mV and the separation of the two signals increases with faster scan rates. Therefore the reaction of copper(II) complexes at higher scan rates can be considered to be quasi-reversible.
Future research in this area would benefit from the work incorporated in this paper to study catalytic activity, biological activities and even polymer film formation.

**Conflicts of Interest**

The authors declare no conflicts of interest regarding the publication of this paper.

**References**


