INTRODUCTION

Glimepiride 1-(p-(2-(3-ethyle-4-methyl-2-oxo-3-pyroline-1-carboxamido)ethyl)phenyl sulfonyl)-3-(trans-4-methylcyclohexyl) Urea is a third generation hypoglycemic sulfonylurea, which is useful in the treatment of non-insulin dependent diabetes mellitus (NIDDM)\(^1\). Glimepiride is a white crystalline powder, relatively insoluble in water. It exhibits slow gastrointestinal absorption rate and inter individual variation of its bioavailability\(^3\).

The slow absorption rate of drug usually originates from either poor dissolution of drug from the formulation or poor permeability of drug across gastrointestinal membrane. For poorly water soluble and highly permeable drugs the rate of oral absorption is often controlled by the dissolution rate in the gastrentestinal tract\(^4\). Complexation of sulfonylurea with lighter transition metal has been studied in detail by Yoshinaga and Yamamotto (1966)\(^5\), Qureshi and Iqbal (1985)\(^6\). A persual of available literature shows that systemic study on complexation of zinc with sulphonyl ureas is relatively scanty\(^7\).\(^10\). The study of chemistry and chemical reaction of structure co-ordination compound helps in establishing structure activity relationship. It has been reported that in biological activity metal complex is more potent and less toxic as compared to the free ligand\(^11\). In view of the above and in continuation of our work, it is interesting to have an insight into the synthesis of zinc complex with glimepiride and to diagnose various structural aspects of the isolated complex.
Ligand-metal ratio

To confirm the ligand metal ratio, conductometric titrations using monovariation method were carried out at 27±1°C 0.005M solution of glimepiride drug was prepared in 80:20 mixture of DMF and water. Similarly solution of metal salt ZnCl₂ was prepared in the same solvent of 0.01M concentration. 20mL of ligand was diluted to 200ml with the same solvent. The ligand was titrated against metal salt solution using monovariation method. Conductance was recorded after each addition, Graph is plotted between corrected conductance and volume of metal salt added (Fig II). From the equivalence point in the graph it has been concluded that the complex formation has taken place in the ratio of 2:1 (L:M). Stability constant and free energy changes were also calculated using Job's method of continuous variation modified by Turner and Anderson (Fig 3).

Conductometric titration monovariation method
Glimepiride with zinc chloride

Fig. 1: Structure of glimepiride

Fig. 2: Mole Metal Ligand Ratio

Fig. 3: Mole Metal Ligand Ratio
EXPERIMENTAL

All chemicals used were of analytical grade. Pure sample of Glimepiride (Molecular formula C_{24}H_{34}N_{4}O_{5}S and mol.wt 490.62) was obtained from Ipca laboratories Ltd, Ratlam in powdered form m.p 207°C.

Metal salt ZnCl₂ was of merck chemical. The solvent used were distilled water and DMF. Metal-ligand ratio was calculated using Systronics Digital Conductivity meter. Melting point was determined by Parkin Elmer melting point apparatus and are uncorrected pH values determined on LabIndia pH Analyser.

IR spectra of ligand and complex were recorded with perkin Elmer spectrometer in the range of 4000-450cm⁻¹ (CDRI Lucknow). The "^{1}H"NMR spectrum of the ligand glimepiride and their isolated complex zinc-glimepiride, were scanned through Bruker DRX-300 NMR Spectrometer from CDRI Lucknow using deuterated acetone as a solvent. Mass spectral analysis of pure ligand as well as metal complex were obtained from CDRI, Lucknow. X-ray diffraction studies were carried out by X-ray Diffractometer model with 45kV rotating anode and Cuka (1W = 1.54060A⁰) radiation (Panjab University).

Synthesis

Complex was synthesized by mixing the solution (80% DMF) metal salt solutions with that of ligand in 1:2 molar ratios; respectively at room temperature maintaining the pH between (6.5-8) by the addition of dilute NaOH solution. On refluxing the mixture content for 3hrs at 80°C and on cooling the off white coloured crystals were obtained. The complex was washed with 80% DMF or alcohol and weighed (yield-52%).

Table 1: Synthesis and Physicochemical characteristics of Glimepiride-Zinc complex

<table>
<thead>
<tr>
<th>Ligand/Complex</th>
<th>Ligand Metal Ratio</th>
<th>Colour</th>
<th>% Yield</th>
<th>Stability Constant LogK (L/mole)</th>
<th>Free Energy Change (ΔF) KCal/mole</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glimepiride</td>
<td>-</td>
<td>White</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Glimepiride-Zinc Complex</td>
<td>2:1</td>
<td>Off white</td>
<td>52%</td>
<td>11.01</td>
<td>15.13</td>
</tr>
</tbody>
</table>
Analysis of Complex

The resulting complex so formed was characterized by its elemental analysis, physical characteristics, IR, NMR Mass spectral and X-ray diffraction studies.

RESULTS AND DISCUSSION

The synthesized complex is offwhite and stable, being soluble in DMSO, acetone and insoluble in water, ethanol etc. Analytical data (table 2) and conductometric studies suggest 2:1 (L:M) ratio. Measured conductance values of these complex are too low to account for their electrolytic behaviour.

Structure Determination

IR absorption studies

The IR spectrum\(^{18,21}\) of the ligand and the isolated complex were recorded in the range 4000-450 cm\(^{-1}\) and the probable assignments are given in (table 3). The proposed structure for the isolated

<table>
<thead>
<tr>
<th>Ligand Complex</th>
<th>Elemental analysis</th>
<th>Found (calc)</th>
<th>m.p(^{0})C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>C</td>
<td>H</td>
<td>N</td>
</tr>
<tr>
<td>C(<em>{24})H(</em>{34})N(<em>{4})O(</em>{5})S</td>
<td>58.77</td>
<td>6.93</td>
<td>11.92</td>
</tr>
<tr>
<td></td>
<td>(58.50)</td>
<td>(6.95)</td>
<td>(11.94)</td>
</tr>
<tr>
<td>(C(<em>{24})H(</em>{34})N(<em>{4})O(</em>{5})S)(_{2}).Zn</td>
<td>52.22</td>
<td>5.21</td>
<td>7.85</td>
</tr>
<tr>
<td></td>
<td>(52.42)</td>
<td>(5.31)</td>
<td>(7.97)</td>
</tr>
</tbody>
</table>

Table 2: Analytical data of Complex

IR Spectra of Pure Drug Glimepiride

IR Spectra of Glimepiride-Zinc complex
Table 3: IR Absorption data of the complex in cm⁻¹

<table>
<thead>
<tr>
<th>Ligand/Complex</th>
<th>ν(NH)</th>
<th>ν(C=O)</th>
<th>ν(S=O)</th>
<th>ν(C-O)</th>
<th>ν(C=N)</th>
<th>ν(SO₂N)</th>
</tr>
</thead>
<tbody>
<tr>
<td>C₂₄H₃₄N₄O₅S</td>
<td>3681</td>
<td>3681</td>
<td>3681</td>
<td>-</td>
<td>-</td>
<td>3020</td>
</tr>
</tbody>
</table>

Table 6

1H NMR spectral data of pure drug
8.44 (S, 1H NHCO), 7.94 (d, benzene J = 0.97 Hz), 7.53 (d, benzene J = 1H), 6.23 (S, SO₂NH J = 0.39 Hz), 4.1 (Se, CH₂N J = 0.994), 3.61 (S, Pyrollidine), 3.33 (t, CH₂ attached with benzene J = 0.60), 3.02 (q, CH₂ attached with carbonyl, J = 21.83), 2.21 (p, CH₃ attached with methyl J = 1.43 Hz), 1.65 (t, CH₂ attached with cyclohexane J = 1.36 Hz), 1.04 (t, CH₃ group, J = 2.82 Hz).

s = singlet,  d = doublet,  t = triplet,  q = quatrate
Table 7: ¹HNMR spectral data of Glimepiride-Zn complex.

8.43 (S, 1H NHCO), 7.94 (t, benzene J = 2.81 Hₜ), 7.45 (q, benzene, J = 3.88 Hₜ), 6.28 (S, SO₂NH, J = 0.80 H₂), 4.1 (Se, CH₂N. J = 3.45 H₂), 3.72 (S, NHCO-Zn, J = 0.82 H₂), 3.61 (S, pyrrolidine), 3.32 (m, CH₂ attached with benzene, J = 1.37 H₂), 3.06 (q, CH₂ attached carbonyl J = 4.33 H₂), 2.21 (p, CH₂ attached with methyl J = 3.93 H₂), 1.64 (S, CH₂ attached with cyclohexane, J = 2.76 H₂) 1.04 (t, CH₃ group j = 7.99 H₂).

s = singlet     d = doublet   t = triplet  q = quaratr

Structure of glimepiride zinc complex
complex is also supported by IR absorption bands and characterized by the absorption of carbonyl (C=O) and sulphonyl urea group at 1701 cm\(^{-1}\) and 1216 cm\(^{-1}\) respectively. The NH group observed at 3681 cm\(^{-1}\) in the ligand (glimepiride) was shifted to 3752 cm\(^{-1}\) in Zinc glimepiride complex. The next IR band of structural significance of the ligand appears at 1656 cm\(^{-1}\) which may be assigned to \(\nu (\text{C-O})\), which was absent in pure ligand and the considerable frequency of \(\nu (\text{C=N})\) was obtained at 1542 cm\(^{-1}\) in metal complex while absent in pure ligand were indicates that these specific IR absorptions are appeared due to complexation. The linkage through amide-O and sulphone –O- atom was further supported by the appearance of aband in the far IR region at 670cm\(^{-1}\) in the complex that may be assignable to M-O frequency (Fig IV a&b).

\[1^1H\text{NMR spectral analysis} \]

\[1^1H\text{ NMR spectra of Glimepiride and its Zinc For pure ligand Glimepiride.} \]

(300 MHz, Acetone)

The \(^1^H\text{NMR spectrum of the ligands Glimepiride and its isolated complex zinc were scanned through bruker DRX-300 NMR spectrometer from DCRI Lucknow using deuterated acetone as a solvent (table 4). In the spectra of glimepiride-zinc complex (table 5) exhibit a broad singlet signal at 8043 ppm, shows a down field shift, Dd for NH in NHCO group (8.44 ppm) in pure ligand.\]^22-24

CONCLUSION

The differences in melting point of metal-ligand complex as compared to Glimepiride suggested that a new product was formed. The shifts of peaks in IR region as well as new signals around at X-ray diffractogram in X-ray studies further confirmed the drug metal complexation. The tentative structure of the complex are further supported by Mass spectral analysis. The overall studies indicate the glimepiride metal complex is non-ionic and have tetrahedral geometry.

REFERENCES

20. Chandran, A., Varghese, H.T., Yohannan, C.,


