INTRODUCTION

The complexes of 1,4-benzodiazepine with metal ions have also been reported in literature. Alprazolam a benzodiazepine drug is included in the category of tranquilizer drugs. It is used to treat anxiety disorders and as an adjunctive treatment for depression. It has been reported that complexes of 1,4-benzodiazepine also possess anticancer properties. Metal complexes of 1,4-benzodiazepines possessing biological activity may be even more active than free ligand. Copper (II) complex of Alprazolam appears to be quite active, having a rapid onset of action and it also prolongs the duration as compared to that of Alprazolam itself. Different coordination modes have been including coordination with the neutral ligand viz. N(4). However, none of the modes has been ascertained. Neutral complexes of 1,4-benzodiazepines such as Nitrazepam where the ligands act as anions through labilization of proton at N(I) atom giving (imido) complexes, have been reported. The structure of the ligand Alprazolam (AZ) is given in figure (1).

![Fig. 1: Alprazolam (AZ)](image-url)
The proposed work has been taken up to investigate the mode of coordination of the metal ion with (AZ) drug of increased biological activity of the drug on complexation as reported in the literature. The results will throw light on the mode of coordination of the metal ion with the AZ drug as well as on the complex formation of other benzodiazepines in general.

**EXPERIMENTAL**

All the chemical used were of AR grade and their solution were prepared in double distilled water. The complexes were prepared by mixing molar solutions of Cu(NO$_3$)$_2$.3H$_2$O, Co(NO$_3$)$_2$.6H$_2$O (dissolved in double distilled water) and the ligand Alprazolam (dissolved in ethanol). The pH of the mixture was adjusted at 7.5-8.5.

The stoichiometry of the complexes of the drug (0.025M) with Cu(II) and Co(II) (0.025M) metal ions was found by carrying out potentiometric titration against standard (0.1M) NaOH solution in ethanol-water mixture. The pH changes observed during the titration were plotted against moles of alkali(m) added per moles of metal ion and ligand as depicted in figs. 3 and 4. The stoichiometric ratio is confirmed by Job's method. The coloured precipitates were filtered, washed several times with hot water followed by ethanol to free it from the soluble impurities. The complex were finally dried in an oven at 100°C and stored in desicator. The purity of the complexes was checked by TLC. The complexes were dissolved in benzene: Acetic Acid (2:1) and TLC was carried out in ethanol: Benzene (80:20) system. The Retention factor (Rf value) was calculated by following formula.

![Fig. 2:](image)

![Fig. 3: System Cu(II): AZ](image)

![Fig. 4: System Co(II): AZ](image)
The Rf values of the Cu(II) and Co(II) complexes were given in table.

**Experimental Biocidal**

In the present study, metal complexes, metal ion and ligand have been tested for their effect on the growth of microbial cultures to study their interactive role with fungi (*Aspergillus flavous, Aspergillus niger, Penicillium, triticena and Fusarium species*) and bacteria (*Escherichia coli, Salmonella typhi, Bacillus subtilis, Staphylococcus aureus*) using filter paper disc method$^{14}$ and Broth serial dilution method$^{15}$ respectively as detailed below.

The complexes were dissolved in DMF (0.5ml) to get a stock solution of 1000 ppm by adding sterilized distilled water. The dilute solution solution of 750 ppm, 500 ppm and 250 ppm concentration were obtained by further dilution of the stock solution. The activity was studied in all the solutions in the cavity slides by mixing the spore

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### Table 1: The Rf values of Cu(II) and Co(II) complexes of Alprazolam

<table>
<thead>
<tr>
<th>Solvent System</th>
<th>Sample</th>
<th>Rf value</th>
<th>Impurity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ethanol : Benzene</td>
<td>Cu(II)-AZ complexes</td>
<td>0.78</td>
<td>Not visible</td>
</tr>
<tr>
<td>80 : 20</td>
<td>Co(II) – AZ complexes</td>
<td>0.72</td>
<td>Not visible</td>
</tr>
</tbody>
</table>

### Table 2: The physical and analytical data of ligand, Cu(II) and Co(II) complexes

<table>
<thead>
<tr>
<th>Compound</th>
<th>Mol. Wt.</th>
<th>Elemental analysis found (Calculated) %</th>
<th>Colour</th>
<th>M.P.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>C</td>
<td>H</td>
<td>N</td>
<td>Cl</td>
</tr>
<tr>
<td>$[C_{17}H_{13}ClN_4]$</td>
<td>308.8</td>
<td>(66.06)</td>
<td>(4.21)</td>
<td>(18.13)</td>
</tr>
<tr>
<td>AZ Cu(NO$_3$)$_2$</td>
<td>296.34</td>
<td>(41.10)</td>
<td>(2.62)</td>
<td>(16.92)</td>
</tr>
<tr>
<td>AZ Co(NO$_3$)$_2$</td>
<td>491.73</td>
<td>(41.49)</td>
<td>(2.64)</td>
<td>(17.08)</td>
</tr>
</tbody>
</table>

### Table 3: Antibacterial and Antifungal activities of the Drug, Metal nitrates and Complexes

<table>
<thead>
<tr>
<th>Compound</th>
<th>Antibacterial Activity zone of inhibition (in mm.)</th>
<th>Antifungal Activity zone of inhibition (in mm.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Drug (AZ)</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Cu(NO$_3$)$_2$</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Co(NO$_3$)$_2$</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Cu AZ(NO$_3$)$_2$</td>
<td>69.13</td>
<td>47.84</td>
</tr>
<tr>
<td>Co AZ(NO$_3$)$_2$</td>
<td>56.56</td>
<td>64.00</td>
</tr>
<tr>
<td>DMF</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>
Fig. 5: IR Spectrum of AZ Complexes
suspension of the test organism separately. Standard drug was used to check and compare the activity of the complexes. It indicated that metal complexes of the drug had a significant activity at a lower concentration and bacteria were incubated at 24±1°C for 20 hours. The data and results are presented in Table 3.

The IR spectra in KBr matrix were recorded on Perkin-Elemer 842-Spectrometer. Elemental analysis of C, H, N were carried out at CDRI, Lucknow.

RESULTS AND DISCUSSION

The molecular formula of the Cu(II) and Co(II) complexes correspond to \[(C_{17}H_{13}Cl N_4). Cu(NO_3)_2\] and \[(C_{17}H_{13}Cl N_4). Co(NO_3)_2\]. The molecular weights 496.34 and 491.73 respectively for Cu(II), Co(II) complexes were determined on the basis of elemental analysis. The physical and analytical data of ligand Cu(II) and Co(II) complexes are given in Table 2.

The IR spectrum of the ligand exhibits bands in the region 1628 cm\(^{-1}\), 1280 cm\(^{-1}\), 1600 cm\(^{-1}\), 740 cm\(^{-1}\), and 2960 cm\(^{-1}\) which may be assigned to \(\nu(C=N)\), \(\nu(C-N)\), \(\nu(C_6H_5)\), \(\nu(Cl)\), \(\nu(CH_2)\) and \(\nu(-CH_3)\) respectively. The ligand band observed in the range of 1628 cm\(^{-1}\) undergoes the lower shifting to 1612 cm\(^{-1}\) and 1613 cm\(^{-1}\) in Cu(II) and Co(II) complexes respectively, indicating azomethine nitrogen N(4) atom of Benzodiazepine ring in coordination to the metal ion complexes. The Cu(II) and Co(II) complexes the band attributed to the vibration mode \(\nu(C-N)\) appear at 1263 cm\(^{-1}\) and 1265 cm\(^{-1}\) showing lower shifting as compared to the ligand (1280 cm\(^{-1}\)), indicating nitrogen (1) participating in complexation. In IR spectrum of the ligand the bands attributed to vibrational mode \(\nu(-C_6H_5)\), \(\nu(CH_2)\), \(\nu(Cl)\), \(\nu(CH_2)\) appears at 1600 cm\(^{-1}\), 1355 cm\(^{-1}\), 740 cm\(^{-1}\) and 2960 cm\(^{-1}\) showing small positive shift in complexation.

The presence of new bands at 480 cm\(^{-1}\) and 340 cm\(^{-1}\) in Cu (II) complex and 492 cm\(^{-1}\) and 340 cm\(^{-1}\) in Co (II) are attributed to \(\nu(M-N)\) linkage\(^{16}\). Other strong bands at 1383 cm\(^{-1}\) and 861 cm\(^{-1}\) in Cu(II) complexes and 1370 cm\(^{-1}\) and 850 cm\(^{-1}\) in Co (II) complex suggest monodentate nitrate group in complexes\(^{17}\). The representative structure of the complex may be shown as

![Probable structure of metal complexes](image)

Co(II) complexes on bacteria and fungal species are presented in Table 3.

The above data indicate that the zone of inhibition at 500 ppm concentration is best as compared to the standard drug. The results show that the Cu(II) complex of Alprazolam is more effective towards all fungi and bacteria as compared to Co(II) complex of Alprazolam and drug Alprazolam itself.

REFERENCES

5. Cinellue, M.A.; Stoccore, S.; Minghetti, G.