SYNTHESIS, CHARACTERIZATION AND ANTIBACTERIAL PROPERTIES OF SOME TRANSITION METAL COMPLEXES OF (1H-PYRROL-2-YL)-ISONICOTINOYLHYDRAZONE

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ABSTRACT. A new ligand, (1H-pyrrol-2-yl)isonicotinoylhydrazone (Pyr-inh) and its metal complexes have been synthesised and characterized by elemental analyses, IR, ¹H and ¹³C-NMR, UV-Vis spectra, molar conductance, and thermogravimetry-differential thermal analysis (TG-DTA). From the results obtained, the structure of the complexes are of general formula [M(Pyr-inh)₂Cl₂]·xH₂O for M = Co(II), and Zn(II) and [M(Pyr-inh)₂(H₂O)₂]Cl₂ for Cu(II). The thermal behaviour of these complexes showed the loss of lattice water in the first step followed by decomposition of the ligand in subsequent steps. The ligand and its transition metal complexes were studied for their in vitro antibacterial activity against ten pathogenic bacterial strains. The data obtained revealed that the ligand was active against Morganella morganii and Escherichia coli, the Co(II) complex active against Klebsiella oxytoca, the Cu(II) complex showed significant activity against five out of the ten bacterial strains and the Zn(II) complex was active against all the pathogenic bacterial strains investigated.

KEY WORDS: (1H-pyrrol-2-yl)isonicotinoylhydrazone, Metal complexes, Schiff base, Synthesis, Thermal analysis, Antibacterial activity

INTRODUCTION

There has been considerable interest in Schiff bases due to the role they played in coordination chemistry as they readily form stable Schiff base complexes with most transition metals [1-3]. Schiff bases and their metal complexes exhibit a broad range of potential applications because of their structural diversity and intrinsic magnetic properties and have been shown to be potential antibacterial [4-15] and anticancer [16-18] agents. The past few decades have seen the introduction of a number of pharmaceutical compounds which contain five, six, and seven-membered rings such as piperazines, piperidines, imidazoles, benzodiazepines, and other heterocycles containing nitrogen, sulfur, and oxygen [1]. The tuberculostatic activity of isonicotinic acid hydrazide and its arylhydrazones containing azomethine nitrogen is attributed to their ability to form stable complexes with d- and f-block metal ions [19-21]. The upsurge interest in hydrazones is due to their antimicrobial, antituberculosis, and antitumor activity [22-24].

The increasing resistance to antibacterial drugs has necessitated the search for new compounds with potential effects against pathogenic bacteria. The incorporation of metals or metal-based systems into molecules of antibacterial or antifungal agents is expected to enhance the bactericidal or fungicidal properties of these drugs and to improve on the activity of compounds with less potentials [4-15]. We have been interested in studying the effect of ligand modification on their physicochemical and biological properties [25, 26].

Isoniazid (INH) is a prodrug of proven therapeutic importance and is used against a broad spectrum of bacterial infections [27]. It is a potent, highly selective antimycobacterial agent

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used against the mycobacterium tuberculosis probably because of its ability to form stable complexes with transition metal ions in living systems. In view of the biological and medicinal properties of this compound, we developed an interest to synthesize a Schiff base derived from isoniazid and pyrrole-2-carboxaldehyde, with the hope of developing metal-based biologically active adducts that could act against resistant bacterial species. We here report on the synthesis, characterisation and biological activities of some transition metal complexes of (1H-pyrrol-2-yl)isonicotinoylhydrazone, Pyr-inh a Schiff base, derived from isoniazid and pyrrole-2-carboxaldehyde.

EXPERIMENTAL

Materials

All chemicals were of analytical AR grade and were used as supplied without further purification. The solvents were doubly distilled according to standard methods. Pyrrole-2-carboxaldehyde was obtained from Acros Organics, isoniazid from BDH Chemicals Ltd, and CuCl₂·2H₂O, ZnCl₂ and CoCl₂·6H₂O from Prolabo Reagents.

Physical measurements

¹H-NMR and ¹³C-NMR spectra were recorded with a 500 MHz Bruker DRX500 spectrophotometer in dimethylsulfoxide (DMSO) as solvent and were reported relative to tetramethylsilane (TMS) as internal standard. The analysis of CHN contents of the ligand and metal complexes were performed on a Euro elemental analyser (Euro vector design). Chloride was estimated by Volhard’s method and metal(II) content by gravimetric method. Conductivity measurements of the metal complexes were determined using a VERNIER LabPro device. Thermogravimetric analyses were carried out using a TGA Perkin-Elmer STA 6000 simultaneous thermal analyzer under a nitrogen atmosphere. IR Spectra were recorded with a Vector 220 FT–IR Spectrophotometer on KBr disc. UV-Visible spectra were recorded on a Shimadzu double-beam spectrophotometer; model UV-150-02 using DMSO as reference.

Synthesis of (1H-pyrrol-2-yl)isonicotinoylhydrazone (Pyr-inh)

Isoniazid (0.288 g, 2.1 mmol) dissolved in 10 mL of methanol was added to pyrrole-2-carboxaldehyde (0.395 g, 2.1 mmol) dissolved in 10 mL of methanol. The mixture was heated under reflux with continuous stirring at about 60 °C for 24 h during which a light brown solution was obtained. The solution was allowed to cool with the formation of light-brown needle-like precipitate. The precipitate was then filtered and dried over anhydrous calcium chloride in a desiccator at room temperature. The formation of the Schiff base was monitored by TLC and its purity was checked by melting point determination and elemental analyses.

Preparation of metal complexes

A general method was used for the synthesis of the metal complexes. These metal complexes were prepared by mixing warm methanolic solutions (0.5 mmol of the metal salts (CoCl₂·6H₂O, CuCl₂·2H₂O and ZnCl₂ dissolved in 10 mL of methanol) and warm methanolic solutions of the ligand (1.0 mmol in 10 mL of methanol). The mixtures were heated under reflux with continuous stirring for 3-4 h, during which the complexes separated out, filtered and washed three times with methanol and dried over anhydrous CaCl₂ in a desiccator.
Biological activity

The ligand and its metal complexes were screened for in vitro antimicrobial activity in DMSO against ten bacterial strains (Staphylococcus aureus, Salmonella typhi, Proteus vulgaris, Enterobacter cloacae, Morganella morganii, Escherichia coli, Citrobacter freundii, Shigella flexneri, Klebsiella pneumoniae and Klebsiella oxytoca). All the bacterial strains were obtained from the Medical Bacteriology Laboratory of Centre Pasteur du Cameroun, Yaoundé Cameroon. The microbial isolates were maintained on an agar slant at 4 °C and the strains were sub-cultured on an appropriate fresh agar plate, 24 h prior antibacterial tests. Samples of the ligand, its metal complexes and the metal salts were diluted in DMSO and 1 mg/mL of each was placed on sterilized filter paper discs and allowed to dry. The reference antibiotic, gentamycin, was also prepared in the same manner in DMSO prior to testing. The antimicrobial diffusion tests were carried out as described by Berghe and Vlietink [27] using a cell suspension of about 1.5 x 10⁶ CFU/mL obtained from the McFarland turbidity Standard No 0.5. The suspension was standardized by adjusting the optical density to 0.1 at 600 nm (Shimadzu UV-120-01 spectrophotometer). The suspension was used for inoculation by floating the surface of the CAM plates. When the excess liquid was allowed to dry under an aspirator hood, holes of 6-mm diameter were made in the medium and filled with 60 µL of test samples.

RESULTS AND DISCUSSION

The condensation of isoniazid with pyrrole-2-carboxaldehyde yielded a new Schiff base (Scheme 1), which was used to prepare the transition metal complexes whose analytical data are presented in Table 1. All the complexes are coloured powders with high melting points. The complexes are non-hygroscopic and are stable in air, insoluble in methyl alcohol, ethyl alcohol, chloroform and acetone but very soluble in DMF and DMSO. The analytical data suggest a ratio of 1:2 (metal:ligand) for all the complexes. The room temperature molar conductivity measurements in DMSO indicated that the complexes are 2:1 electrolytes [28]. The elemental analytical data are consistent with the proposed formula.

![Scheme 1. Synthesis of (1H-pyrrol-2-yl) isonicotinoylhydrazone.](image)

Table 1. Analytical and physical data of the ligand and its complexes.

<table>
<thead>
<tr>
<th>Compound</th>
<th>Colour</th>
<th>Yield %</th>
<th>λ/Ω1 cm⁻¹ mol⁻¹</th>
<th>Elemental analysis</th>
<th>Found (calcd.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>C₃H₁₁NO</td>
<td>Light brown</td>
<td>87.0</td>
<td>-</td>
<td>%C %N %H %M %Cl</td>
<td></td>
</tr>
<tr>
<td>C₂₁H₁₈N₂O₂Cl</td>
<td>Dark green</td>
<td>72.0</td>
<td>46.93 (46.10)</td>
<td>19.91 (18.98)</td>
<td>3.56 (3.87)</td>
</tr>
<tr>
<td>[Zn(C₁₁H₁₀N₄O₂Cl)₂][H₂O]</td>
<td>Light yellow</td>
<td>66.6</td>
<td>46.78 (47.20)</td>
<td>19.84 (18.78)</td>
<td>3.54 (4.32)</td>
</tr>
<tr>
<td>[Co(C₁₁H₁₀N₄O₂Cl)₂][H₂O]</td>
<td>Brown</td>
<td>21.2</td>
<td>47.32 (47.64)</td>
<td>20.08 (21.50)</td>
<td>3.58 (3.78)</td>
</tr>
</tbody>
</table>
IR spectra

A comparative study of the IR spectra of the complexes with that of the ligand indicated that bands due to $\nu$(C=O) of the ligand at 1700 cm$^{-1}$ and $\nu$(NH) at 3240 cm$^{-1}$ were absent in the spectra of the respective complexes (Table 2). This is presumably due to amide-imidol tautomerism (Scheme 2) and their subsequent coordination through the imidol oxygen [29]. Bands at 1600 cm$^{-1}$ due to $\nu$(C=N) of the ligand shifted to lower wave number in the complexes 1565-1575 cm$^{-1}$ indicating the coordination of the azomethine nitrogen. Non ligand bands at 420-427 cm$^{-1}$ and 510-520 cm$^{-1}$ were assigned to $\nu$(M-O) and $\nu$(M-N), respectively. In the ligand, the band at 1581 cm$^{-1}$ due to the pyridine ring nitrogen remained unchanged on complexation indicating the non involvement of the ring nitrogen in the complexation [30]. The overall IR spectral evidence suggests that the ligand is bidentate in nature, coordinating through amide-oxygen and azomethine nitrogen with the formation of a five membered chelate ring. The spectra of all complexes exhibited intense broad bands at 3400-3450 cm$^{-1}$ due to $\nu$(OH) of the lattice water molecule(s) [27, 31]. The presence of the lattice water molecules was also confirmed by elemental and thermogravimetric analysis.

Table 2. Selected infrared absorption bands (cm$^{-1}$) of ligand and its complexes.

<table>
<thead>
<tr>
<th>Compound</th>
<th>$\nu$(OH)</th>
<th>$\nu$(C=N)</th>
<th>$\nu$(C=O)</th>
<th>$\nu$(N-H)</th>
<th>$\nu$(Py-ring)</th>
<th>$\nu$(M-O)</th>
<th>$\nu$(M-N)</th>
</tr>
</thead>
<tbody>
<tr>
<td>C$<em>{11}$H$</em>{10}$N$_4$O$_2$</td>
<td>-</td>
<td>1600</td>
<td>1700</td>
<td>3240</td>
<td>1581</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>[Cu(C$<em>{11}$H$</em>{10}$N$_4$O$_2$)(H$_2$O)$_2$]Cl$_2$</td>
<td>3400</td>
<td>1570</td>
<td>1650</td>
<td>3200</td>
<td>1581</td>
<td>520</td>
<td>420</td>
</tr>
<tr>
<td>[Zn(C$<em>{11}$H$</em>{10}$N$_4$O$_2$)Cl$_2$.2H$_2$O]</td>
<td>3450</td>
<td>1565</td>
<td>1680</td>
<td>3214</td>
<td>1580</td>
<td>518</td>
<td>427</td>
</tr>
<tr>
<td>[Co(C$<em>{11}$H$</em>{10}$N$_4$O$_2$)Cl$_2$.2H$_2$O]</td>
<td>3400</td>
<td>1575</td>
<td>1650</td>
<td>3205</td>
<td>1580</td>
<td>510</td>
<td>424</td>
</tr>
</tbody>
</table>

Scheme 2. Tautomeric forms of the ligand showing (1) amide form and (2) imidol form.

NMR spectra

The $^1$H NMR spectral data of the ligand and the corresponding metal complexes were recorded in DMSO-$d_6$ with TMS as an internal standard. A comparative study of the $^1$H NMR spectra of the ligand and its metal complexes indicated the disappearance of the NH proton signal at $\delta$ 11.62 in the complexes which is considered as an additional evidence of enolization of the ligand during complexation [32]. The free ligand exhibited a signal at $\delta$ 8.51 which was assigned to $\alpha$-protons of the pyridine ring and multiplet signals observed between $\delta$ 6.97 and $\delta$ 8.27 and assigned to $\beta$-protons of the pyridine ring and the aromatic ring protons [33].

The $^{13}$C NMR data of the ligand indicated signals due to carbon attached to the azomethine nitrogen and enolic oxygen at $\delta$ 141.3 and $\delta$ 161.6, respectively. Thus, the $^1$H-NMR and $^{13}$C-NMR spectra confirmed the monobasic bidentate nature of the ligand, already suggested by the IR spectral studies.
**Electronic spectra**

The electronic absorption spectra of the complexes were recorded in DMF solution. The electronic spectrum of the Cu(II) complex showed large bands at 26667 cm$^{-1}$ and 38462 cm$^{-1}$, while the electronic spectrum of Zn(II) complex exhibited two bands at 28571 cm$^{-1}$ and 38462 cm$^{-1}$. Furthermore the electronic spectrum of the Co(II) complex exhibited two bands at 28571 cm$^{-1}$ and 38462 cm$^{-1}$. The absorption bands observed at 26667 cm$^{-1}$ and 28571 cm$^{-1}$ are assignable to a combination of d-d bands and metal-ligand charge transfer (M→LCT) [34, 35]. The electronic spectra measurements is indicative of octahedral geometry for the Cu(II) complex, a tetrahedral geometry for the Zn(II) complex and a square planar geometry for the Co(II) complex.

**Thermal studies**

Thermal analysis has proved to be useful in determining the crystal water content in complexes and their thermal stability and decomposition mode under a controlled heating rate. Thermogravimetric (TG) and differential thermal analyses (DTA) were used to describe the thermal behaviour of the prepared complexes. The typical TG-DTA analysis of the complexes was performed from 20 to 320 °C at a heating rate of 5 K min$^{-1}$ under dynamic air condition. The decomposition of Cu(II) complex takes place in two stages as shown in Figure 1.

The TG/DTA curve of the Cu(II) complex show mass loss between 40-145 °C with DTA endothermic peak at 84 °C due to the dehydration with loss of coordinated water molecules. The mass loss observed in this step is 9.12%. The second stage between 145-300 °C with DTA endothermic peak at 185 °C is due to decomposition of the ligand molecule. The degradation of the molecule was further confirmed by two exothermic peaks at 226 °C and 296 °C with 27.7% mass loss.

The TG/DTA curve for the Zn(II) complex is shown in Figure 2. Thermal decomposition of the Zn(II) complex occurs in three stages and takes place between 40-130 °C with mass loss of 9.22%, attributed to the loss of two moles of lattice water. Two endothermic peaks at 40 °C and 100 °C are observed in the DTA curve. The second stage (from 130-228 °C) is attributed to the decomposition of the ligand moiety. A sharp endothermic peak at 200 °C is noticed with an observed mass loss of 10.39%. The third stage occurred between 228-304 °C with mass loss of 12.31% which may be attributed to the total decomposition of the ligand molecule.

The TG/DTA curves of the Co(II) complex are represented in Figure 3. The Co(II) complex shows almost three stage process as with Zn(II) complex. The first decomposition stage occurs in the temperature range 40-104 °C with a mass loss of 16.34% attributed to the dehydration and loss of lattice water corresponding to an endothermic effect at 60 °C. The second decomposition stage occurred between 104-176 °C with a mass of loss of 7.43% attributed to the decomposition of the ligand moiety. A broad exothermic peak at 94 °C was also observed. The third decomposition occurred between 176-304 °C with mass loss of 28.8% having an endothermic peak at 236 °C and a broad exothermic peak at 270 °C and attributed to the total decomposition of the ligand.

Based on the above analytical and spectral data together with the thermal decomposition studies, the complexes have the structural formula as depicted in Figure 4.
Figure 1. TGA and DTA thermograms of Cu(II) complex.

Figure 2. TGA and DTA thermograms of Zn(II) complex.
Some transition metal complexes of (1H-pyrrol-2-yl)-isonicotinoylhydrazone


Figure 3. TGA and DTA thermograms of Co(II) complex.

Figure 4. Proposed structure of the Cu(II) complexes.

**Biological activity**

Antibacterial activity of the ligand and its metal complexes were tested *in vitro* by the disc-diffusion method [36] against ten bacterial stains (*Salmonella typhi, Staphylococcus aureus, Proteus vulgaris, Enterobacter cloacae, Morganella morganii, Escherichia coli, Citrobacter freundii, Shigella flexneri, Klebsiella pneumoniae* and *Klebsiella oxytoca*). Holes with methanol were used as negative control. The plates were inoculated at 37 °C for 24 h. Antimicrobial activity was evaluated by measuring the diameter of the inhibition zone (IZ) around the hole. The assay was repeated thrice and results recorded as mean ± standard deviation of triplicated experiments. Compounds were considered as active when the IZ was greater than 6 mm.

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The diameter of the susceptibility zones were measured in mm and the results are presented in Table 3. The ligand, INH-PYR, was found to be active against *Morganella morganii* and *Escherichia coli*. The Cu(II) complex showed significant activity against two out of the five bacterial strains (*Klebsiella pneumoniae* and *Klebsiella oxytoca*). The Co(II) complex showed significant activity against *Klebsiella oxytoca*. Isoniazid, INH one of the reagents for the synthesis of the Schiff base is an antibiotic in clinical use. It showed moderate activity against *Klebsiella pneumoniae* while pyrrole-2-carboxaldehyde, another reagent was inactive to all the bacterial strains.

### Table 3. Antibacterial screening data for the ligand and its complexes.

<table>
<thead>
<tr>
<th>Compound</th>
<th>Diameter of inhibition zone (in mm)</th>
<th>S. typhi</th>
<th>M. morganii</th>
<th>E. coli</th>
<th>K. pneumoniae</th>
<th>K. oxytoca</th>
</tr>
</thead>
<tbody>
<tr>
<td>C_{11}H_{10}N_{4}O</td>
<td>13</td>
<td>12</td>
<td>12</td>
<td>12</td>
<td>12</td>
<td>12</td>
</tr>
<tr>
<td>[Cu(C_{11}H_{10}N_{4}O)(H_{2}O)_{2}]Cl</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>19</td>
<td>17</td>
</tr>
<tr>
<td>[Zn(C_{11}H_{10}N_{4}O)Cl_{2}].2H_{2}O</td>
<td>13</td>
<td>8</td>
<td>16</td>
<td>14</td>
<td>12</td>
<td></td>
</tr>
<tr>
<td>[Co(C_{11}H_{10}N_{4}O)Cl_{2}].2H_{2}O</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>RA (Gentamycin)</td>
<td>26</td>
<td>26</td>
<td>25</td>
<td>11</td>
<td>22</td>
<td></td>
</tr>
</tbody>
</table>

The Zn(II) complex showed antibacterial activity against all the five strains of microbes. In this case, it was moderately active against *Morganella morganii* and significantly active against the rest of the bacterial strains. The Zn(II) complex was significantly more active against *Klebsiella pneumoniae* than the reference antibiotic, gentamycin. All metal salts showed activity against all the strains of bacteria.

It is known that chelation tends to improve on the activity of the ligand thus rendering it a more powerful and potent antibactericidal agent than the parent organic ligand (Schiff base) [36]. A possible explanation is that, in the chelated complex, the positive charge of the metal is partially shared with the donor atoms in the ligand and there is \( \pi \)-electron delocalisation over the whole chelating ring. This, in turn, increases the lipophilic character of the metal chelate and favours its permeation through the lipid layers of the membranes of the micro-organism. Apart from this, other factors such as solubility, conductivity and dipole moment influenced by the presence of the metal ions may also be reasons for the increased activity.

**CONCLUSION**

We have been able to synthesise a new Schiff base ligand, (1H-pyrrol-2-yl) isonicotinoylhydrazone and reacted with Co(II), Cu(II) and Zn(II) ions to yield the corresponding metal complexes. This is a neutral ligand, with tridentate NNO donors derived from the the azomethine nitrogen atom, the pyrrole nitrogen atom and the carbonyl oxygen atom. Antibacterial activity of the ligand and its metal complexes reveal that the ligand and the Zn(II) complex exhibited remarkable antibacterial activities against some of the tested bacterial strains. The significant biological activity of the Zn(II) complex clearly demonstrated the effect of chelation on the biological activity of the ligand.

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