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Proceedings of the Nigerian Society of Physical Sciences

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Synthesis, characterization, and antimicrobial activity of Ni (II) and Cu (II) complexes with schiff base derived from pyrrole-2-carboxaldehyde and thiosemicarbazide

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ABSTRACT

A novel Schiff base ligand was synthesized from the condensation of substituted thiosemicarbazide and 1H- 1H-pyrrole-2-carboxaldehyde. The corresponding Ni(II) and Cu(II) complexes were obtained by refluxing the chloride of the metals with the prepared Schiff base in an ethanolic solution. The Schiff base and its complexes were characterized and analyzed using Fourier Transform Infrared (FT-IR), UV-visible, magnetic susceptibility, conductivity measurement, melting point/decomposition temperature, and solubility test. The Infrared spectral data of the Schiff base showed an absorption band at 1585 cm^{-1} , attributed to $\nu(\text{C}=\text{N})$ stretching. However, this band was shifted to a higher frequency of 1590 cm^{-1} and 1596 cm^{-1} indicating the formation of a Ni-N and Cu-N band in the complexes respectively. The UV-Visible studies revealed significant red shifts in the characteristics C=N and C=S bands upon complexation confirming strong ligand-metal coordination. The complexes exhibited enhanced thermal stability (with decomposition temperatures of 217°C for Ni(II), 205°C for Cu(II)). Magnetic measurements indicated a high-spin octahedral geometry for the Ni(II) complex ($\mu_{\text{eff}}=2.9\text{B.M}$) and a distorted octahedral (Jahn-Teller) geometry for the Cu(II) complex ($\mu_{\text{eff}}=1.90\text{B.M}$). The molar conductance value of the Ni(II) complex was observed at $38.20\text{ ohm}^{-1}\text{ cm}^2\text{ mol}^{-1}$ while that of the Cu(II) complex was observed at $24.20\text{ ohm}^{-1}\text{ cm}^2\text{ mol}^{-1}$ suggesting non-electrolytic nature of the complexes. The Schiff base and the complexes were screened for antimicrobial activity. The results demonstrated that metal complexation significantly enhances bioactivity relative to the free ligand, although their activity was lower than that of the standards.

Keywords: Synthesis, Schiff base, Thiosemicarbazide, Pyrrole-2-carboxaldehyde, Antimicrobial activity.

DOI:10.61298/pnspsc.2025.2.176

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1. INTRODUCTION

Schiff base ligands and their metal complexes have garnered significant interest in coordination chemistry due to their versatile bonding capabilities, structural diversity and wide range of ap-

plications in catalysis, electrochemistry and medicine [1]. Schiff bases, characterized by an azomethine ($-\text{C}=\text{N}$) functional group, are typically synthesized via the condensation of primary amines with carbonyl compounds. Their ability to coordinate with transition metals leads to the formation of stable metal complexes with tunable electronic, magnetic, and biological properties [2].

Among Schiff base ligands, thiosemicarbazone derivatives exhibit enhanced coordination flexibility due to the presence of

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multiple donor atoms such as N, S or O stabilizing metal centers in different oxidation states [3]. Additionally, incorporating heterocyclic moiety such as Pyrrole can significantly alter the electronic properties of the ligands leading to enhanced redox activity, biological interactions and unique optical characteristics [4].

Metal-based drugs have gained attention as alternatives to traditional organic molecules, particularly in antimicrobial, anticancer and antioxidant properties due to their ability to interact with biomolecules such as DNA and proteins [6]. Metal-based drugs have played a significant role in medicinal chemistry, particularly in the treatment of cancer and infectious diseases. The incorporation of metal ions into drug molecules has led to improved pharmacokinetics, enhanced biological activity and targeted delivery mechanisms [7].

One of the major advantages of the metal-based drugs is their ability to interact with biomolecules in a manner distinct from purely organic drugs. For instance, platinum-based drugs like cisplatin and carboplatin function by forming DNA adducts leading to apoptosis in cancer cells [8]. Despite these advantages, challenges remain in optimizing the selectivity and minimizing the side effects of metal-based drugs.

Recent efforts in drug design have focused on developing theranostic metal complexes, compounds that combine therapeutic and diagnostic functions to improve patient outcomes and personalized medicine approaches [9]. With increasing research on Schiff base ligands derived from thiosemicarbazide and pyrrole-2-carboxaldehyde, there is growing potential for the development of novel metal complexes with enhanced bioactivity and reduced toxicity [10]. Studies suggest that metal complexes with pyrrole-containing Schiff bases exhibit increased reactive oxygen species (ROS) generation, disrupting bacterial and fungal cell membranes [11]. In particular, Cu (II) and Ni (II) complexes have been reported to show potent biological activity due to their ability to catalyze oxidation reactions in the cellular environment. Thus, the research aimed to synthesize and characterize novel Schiff base ligand and to prepare its corresponding Ni(II) and Cu(II) complexes and to also evaluate the bioactivity of these compounds against bacterial and fungal strains thereby establishing structure-activity relationships for potential therapeutic applications.

2. EXPERIMENTAL

2.1. MATERIALS AND METHODS

All chemicals and solvents used were of analytical grade and were purchased from Sigma-Aldrich. All glass wares used were washed with detergent thoroughly, rinsed with distilled water, and dried in an oven at 110°C. Thiosemicarbazide derivative (T33405) and 2-pyrrolecarboxaldehyde (P73404) were also obtained from Sigma Aldrich and were used without further purification. Weighing was conducted using Metler balance model AB54. Melting point and decomposition temperatures were recorded using the Gallenkamp SMP10 melting point apparatus.

2.1.1. Synthesis of ligand

Ethanol solution of 2-pyrrolecarboxaldehyde (0.952 g, 10 mmol) was added in hot ethanol solution (20 ml) of thiosemicarbazide (0.911 g, mmol) with a few drops of acetic acid with

constant stirring. The mixture was refluxed at 80°C for 2 hours. On cooling, a silver grey colored compound precipitated out. It was then filtered, washed several times with 50% ethanol, dried and recrystallized from methanol [13].

2.1.2. Synthesis of metal complexes

The Schiff base ligand (3.36 g, 20 mmol) was dissolved in ethanol (20ml) under stirring to which a solution of the corresponding metal salts (2.38 g, 10 mmol) for Ni(II) and (1.71g, 10 mmol) for Cu(II) metal salts in ethanol (20ml) was added dropwise. The reaction mixture was refluxed for 4 hours at 80°C until a colored precipitate was formed. The solid was collected, washed with ethanol and dried in a desiccator [13].

2.2. MOLAR CONDUCTIVITY MEASUREMENT OF THE THIOSEMICARBAZONE COMPLEXES IN DMSO

Conductivity was measured using a Jenway conductivity meter (model 4010) to determine the ionic nature of the complexes.

2.3. MAGNETIC SUSCEPTIBILITY MEASUREMENT

Magnetic moments were determined using Sherwood magnetic susceptibility balance to measure the electronic configuration of the metal centers.

2.4. INFRARED SPECTRAL ANALYSIS OF THE THIOSEMICARBAZONE AND ITS METAL (II) COMPLEXES

Infrared spectral analysis was recorded using Fourier Transform Infrared Spectrometer (CARY 630 Agilent technology) in the range 4000-400 cm^{-1} to confirm functional group formation and metal coordination.

2.5. UV-VISIBLE SPECTRAL ANALYSIS OF THE THIOSEMICARBAZONE AND ITS METAL (II) COMPLEXES

The electronic spectra of the ligand and its complexes were recorded between 200-600 nm at room temperature using Perkin-Elmer Lambda 35 to observe d-d transition and ligand-metal charge transfer bands.

2.6. OB'S METHOD OF CONTINUOUS VARIATION

This technique is used to determine the stoichiometry of the metal complexes by mixing equimolar solutions of the Schiff base and the metal salts in varying ratios. The absorbance was measured using UV-Visible spectroscopy and a plot of molar fraction against absorbance was plotted.

2.7. GRAVIMETRIC METHOD FOR CHLORINE DETERMINATION IN THE COMPLEXES

Gravimetric analysis is a quantitative analysis used to determine chlorine content in the complexes. The metal complex was dissolved in an acid solution to release chloride ions which were then precipitated as silver chloride (AgCl) using a standardized AgNO_3 solution. The AgCl precipitate was filtered, washed, dried, and weighed. The mass measured of the AgCl allowed for the calculation and confirmation of the presence of the coordinated chloride in the complexes.

Table 1. Percentage yield and some properties of thiosemicarbazone and its metal (II) complexes.

Compounds	Color	Melting point (°C)	Decomposition Temperature (°C)	Percentage Yield (%)
Ligand, L	Silver grey	179	-	73.10
[NiL ₂ Cl ₂].3H ₂ O	Dark grey	-	217	78.70
[CuL ₂ Cl ₂].5H ₂ O	Dark yellow	-	205	71.36

L=C₆H₈N₄S

Table 2. Conductivity measurement of Ni (II) and Cu (II) complexes in 1×10⁻³ DMSO.

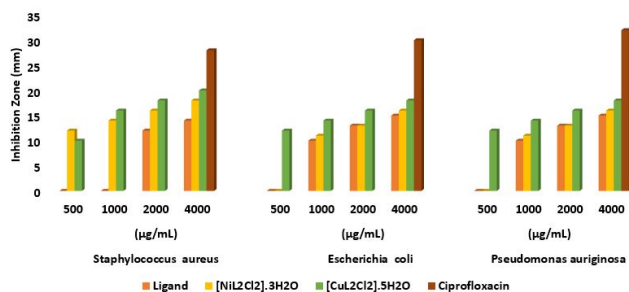
Compounds	Specific conductivity Ω ⁻¹ cm ⁻¹	Molar conductivity Ω ⁻¹ cm ² mol ⁻¹
[NiL ₂ Cl ₂].3H ₂ O	38.20 × 10 ⁻⁶	38.20
[CuL ₂ Cl ₂].5H ₂ O	24.20 × 10 ⁻⁶	24.20

L=C₆H₈N₄S

Table 3. Magnetic susceptibility data of Ni (II) and Cu (II) complexes.

Compounds	X _g (g ⁻¹)	X _m (mol ⁻¹)	μ _{eff} (B.M)	Magnetic property
[NiL ₂ Cl ₂].3H ₂ O	4.6987 × 10 ⁻⁶	2.437 × 10 ⁻³	2.40	Paramagnetic
[CuL ₂ Cl ₂].5H ₂ O	2.0769 × 10 ⁻⁶	1.1623 × 10 ⁻³	1.70	Paramagnetic

L=C₆H₈N₄S

**Figure 1. Bar chart showing antibacterial activity of thiosemicarbazone and its complexes.**

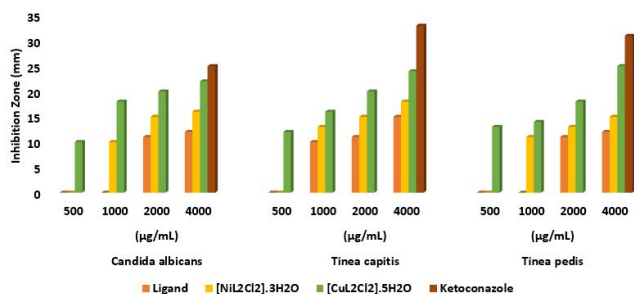
2.8. ANTIMICROBIAL ASSAY

The antibacterial and antifungal activity tests of the synthesized thiosemicarbazone and its respective complexes were carried out against three pathogenic bacterial isolates *Staphylococcus aureus*, *Escherichia coli*, and *Pseudomonas aeruginosa*, and three fungal isolates *Candida albicans*, *Tinea capitis* and *Tinea pedis* using well diffusion methods as reported by Ref. [11].

3. RESULTS AND DISCUSSION

Tables 1 - 6 present the results obtained from the physicochemical analysis, spectral characterization, and *in-vitro* antimicrobial evaluation of the thiosemicarbazones and their complexes.

Table 1 shows the physicochemical properties of the thiosemicarbazone and its metal (II) complexes. The thiosemicarbazone appears silver grey, indicating a neutral electronic environment with minimal conjugation effects. The Ni(II) complex is dark grey while the Cu(II) complex is dark yellow indicating a strong ligand-to-metal charge transfer. The color variation is likely due to metal-ligand interaction, ligand field effect, and hydration contribution [14–17]. The Ni(II) complex decomposed at 217°C compared to the Cu(II) complex (205°C). The higher decomposition temperature indicates a more stable coordination environment [18]. The thiosemicarbazone yield (73.10%) indi-

**Figure 2. Bar chart showing antifungal activity of thiosemicarbazone and its complexes.**

cates an efficient condensation reaction between the pyrrole-2-carboxaldehyde and thiosemicarbazide and also proves the reaction is economically feasible and promising. Both synthesized complexes have good percentage yields 78.70% and 71.36% respectively. These yields are within the range for thiosemicarbazone metal complexes [19, 20].

Table 2 shows the molar conductance measurement of the metal (II) complexes was carried out in DMSO. The values obtained for the respective complexes were 38.20 and 24.20 Ω⁻¹cm²mol⁻¹. These low molar conductance values suggested the non-electrolytic behaviour of the complexes with the chloride ions likely coordinated to the metal center rather than existing freely in solution. This is consistent with literature reports on similar Schiff base metal complexes. Studies have shown that complexes with inner-sphere chloride exhibit molar conductance values, 50 Ω⁻¹cm²mol⁻¹ whereas complexes that dissociate into free ions display significantly higher values (often 70 Ω⁻¹cm²mol⁻¹). Such low conductance values have been reported for Schiff base complexes [6, 21]. The magnetic moment values, as shown in Table 3, suggested that the Ni (II) complex with an observed μ_{eff} value of 2.40 B.M indicated two unpaired electrons in the octahedral high-spin environment which is in agreement with previously reported values for Ni(II) Schiff base complexes with the spin-only value in the range, 2.83-3.3 [19, 22]. For the Cu(II) complex, the measured value of 1.90B.M is characteristic of a Jahn-Teller distorted octahedral geometry where one unpaired electron in the d⁹ system leads to deviation from ideal octahedral symmetry. Reported magnetic moments for similar Cu(II) complexes range between 1.85-2.1 B.M [23, 24].

Table 4 shows the solubility test result of the thiosemicarbazone and its metal (II) complexes in various solvents. The thiosemicarbazone Schiff base was found soluble in DMF and DMSO due to strong dipole interactions [25]. The insolubility of the Schiff base in water, n-hexane and Diethyl ether is probably due to the non-ionic nature and lack of hydrophilic groups, which indicates the non-electrolytic nature of the Schiff

Table 4. Solubility test of thiosemicarbazone and its metal (II) complexes in various solvents.

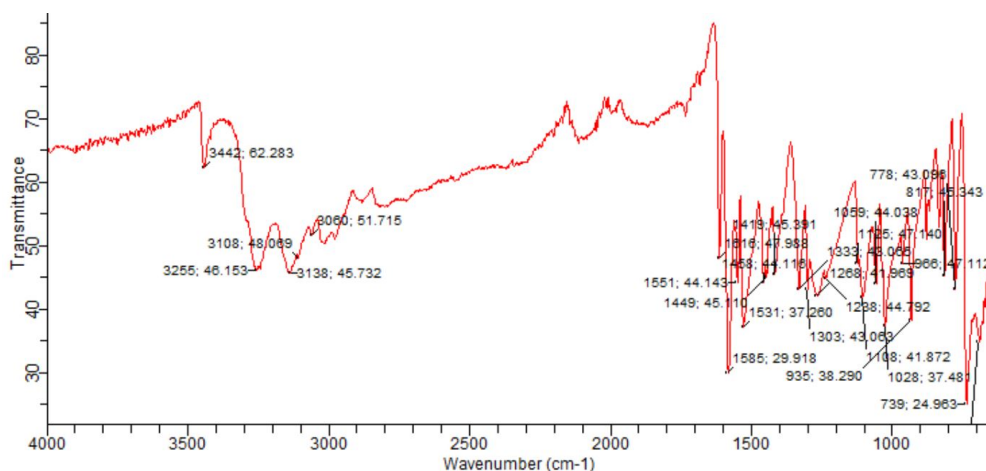
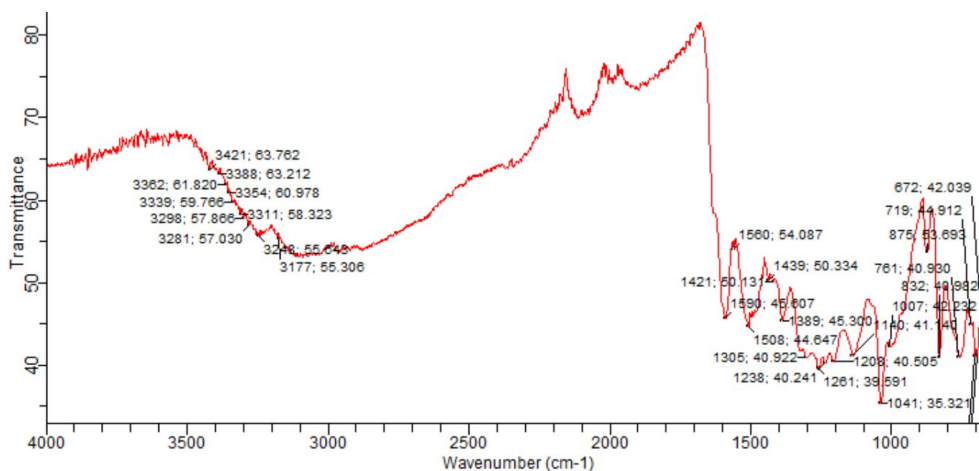
Compounds	Distilled water	Ethanol	DMF	DMSO	n-Hexane	Diethyl ether
Ligand, L	Insoluble	Sparingly soluble	Soluble	Soluble	Sparingly soluble	Sparingly soluble
[NiL ₂ Cl ₂].3H ₂ O	Insoluble	Sparingly soluble	Soluble	Soluble	Insoluble	Insoluble
[CuL ₂ Cl ₂].5H ₂ O	Insoluble	Sparingly soluble	Soluble	Soluble	Insoluble	Insoluble

L=C₆H₈N₄S

Table 5. Infrared Spectral data of thiosemicarbazone and its Metal (II) complexes.

Compounds	$\nu(\text{OH}) \text{ cm}^{-1}$	$\nu(\text{NH})_{\text{sec}} \text{ cm}^{-1}$	$\nu(\text{NH})_{\text{pri}} \text{ cm}^{-1}$	$\nu(\text{C}=\text{N}) \text{ cm}^{-1}$	$\nu(\text{C}=\text{S}) \text{ cm}^{-1}$	$\nu(\text{C}-\text{S}) \text{ cm}^{-1}$	$\nu(\text{M}-\text{N}) \text{ cm}^{-1}$	$\nu(\text{M}-\text{S}) \text{ cm}^{-1}$	$\nu(\text{M}-\text{Cl}) \text{ cm}^{-1}$
Ligand, L	-	3443	3260	1618	1532	817	-	-	-
[NiL ₂ Cl ₂].3H ₂ O	3421	3248	3177	1590	1238	832	513	470	422
[CuL ₂ Cl ₂].5H ₂ O	3414	3248	3146	1596	1261	829	506	482	420

L=C₆H₈N₄S

**Figure 3. FTIR spectrum of thiosemicarbazone.****Figure 4. FTIR spectrum of [NiL₂Cl₂].3H₂O complex.****Table 6. UV-Visible Spectral data of thiosemicarbazone and its Ni(II) and Cu(II) complexes.**

Compounds	$n-\pi^*(\text{nm}) \text{ C}=\text{S}$	$n-\pi^*(\text{nm}) \text{ C}=\text{O}$	$n-\pi^*(\text{nm}) \text{ C}=\text{N}$
Ligand, L	295	327	383
NiL ₂ .3H ₂ O CuL ₂ .5H ₂ O	342 333	348 343	357 350

L=C₆H₈N₄S

base. The metal (II) complexes retain a similar solubility trend. The findings are in agreement with similar report in the literature [26, 27]. Table 5 shows the infrared spectral data of the thiosemicarbazone and its complexes. The $\nu(\text{N}-\text{H})$ stretching bands for secondary and primary amines appear at 3443cm^{-1} and 3260cm^{-1} in the spectra of the thiosemicarbazone as indicated in Figure 3, shifts to 3248cm^{-1} and 3177cm^{-1} in Ni(II) complex and 3248cm^{-1} and 3146cm^{-1} in spectra of the Cu(II)

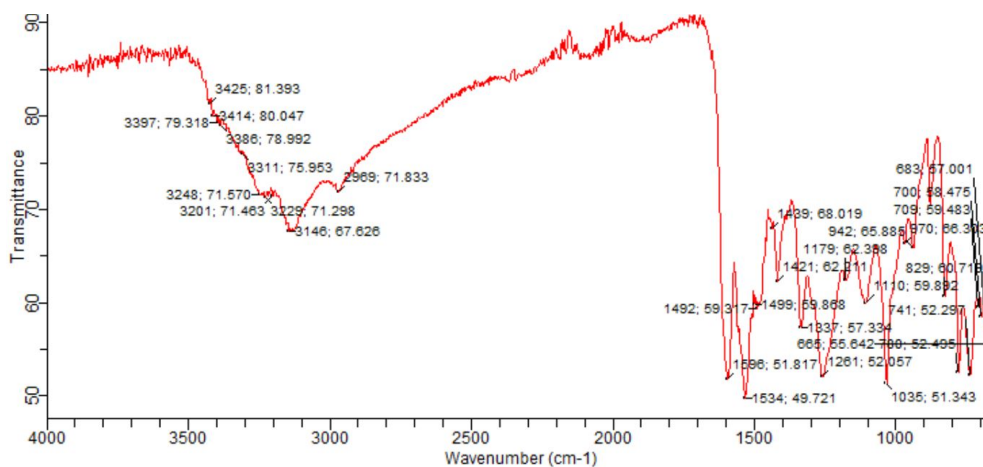


Figure 5. FTIR spectrum of $[\text{CuL}_2\text{Cl}_2]\cdot 5\text{H}_2\text{O}$ complex.

complex which is an indication of coordination through the nitrogen of the azomethine. The $\nu(\text{C}=\text{N})$ stretching vibration at 1618cm^{-1} which indicates the presence of an azomethine group, shifted to a lower frequency of 1590cm^{-1} and 1596cm^{-1} in the spectra of the complexes due to coordination through the nitrogen atom of the azomethine group. Similarly, $\nu(\text{OH})$ stretching at 3421cm^{-1} and 3414cm^{-1} suggests the presence of water molecules. New absorption bands at 513cm^{-1} , 470cm^{-1} , 422cm^{-1} and 506cm^{-1} , 482cm^{-1} , and 420cm^{-1} in the spectra of the complexes were due to M-N, M-S, and M-Cl respectively as shown in Figures 4 and 5. This confirms the coordination of the thiosemicarbazone to the metal ions. This observation agrees with studies on similar Schiff base metal complexes confirming coordination through azomethine nitrogen and thiocarbonyl sulphur atoms [28]. UV-visible spectra of the thiosemicarbazone and that of its complexes were recorded in Table 6. For the thiosemicarbazone absorption bands at $\text{C}=\text{S}(295\text{nm})$, $\text{C}=\text{O}(327\text{nm})$, $\text{C}=\text{N}(383\text{nm})$ corresponds to $n\rightarrow\pi^*$ transitions. The complexes showed high intensities electronic transitions at 342 and $333(n\rightarrow\pi^*, \text{C}=\text{S})$, 348 and $343(n\rightarrow\pi^*, \text{C}=\text{O})$, 357 and 350nm ($n\rightarrow\pi^*, \text{C}=\text{N}$). These shifts in absorption bands to lower λ_{max} (blue shift) or higher λ_{max} (red shift) in the spectra of the complexes are due to coordination of the thiosemicarbazone to the metal ions. These findings are in agreement with reported studies on transition metal Schiff base complexes [29]. Figures 1 and 2 show the *in vitro* antibacterial and antifungal activities of the thiosemicarbazone and its metal (II) complexes. The compounds were tested against three clinically tested pathogenic bacteria (*Staphylococcus aureus*, *Escherichia coli*, and *Pseudomonas aeruginosa*) and three fungal isolates (*Candida albicans*, *Tinea capitis* and *Tinea pedis*). The results indicated that the metal complexes exhibit higher antibacterial and antifungal activities than the free thiosemicarbazone. The incorporation of metal ions into the ligand enhances the activity of the compound thereby increasing the ligand's liposolubility and also is concentration dependant, that is, increases generally with increase in concentrations. But their activities are comparable with that of control as indicated in Figures 1 and 2. Cu(II) complex showed the highest inhibition against all tested strains particularly *Staphylococcus aureus* and *Candida albicans* where it showed activity comparable to the standard drugs.

These findings are in agreement with previous reports that Cu(II) complexes, due to their redox-active nature, exhibit enhanced cell membrane disruption and enzyme inhibition compared to Ni(II) complex [23].

4. CONCLUSION

A Schiff base ligand and its metal (II) complexes were successfully synthesized and characterized using various physicochemical techniques and spectral analysis. Both the Schiff base and the resulting compounds were produced in considerable yields and displayed notable thermal stability. The low conductivity observed in the complexes suggests their non-electrolytic nature, confirming chloride coordination within the inner coordination sphere. Measurements of effective magnetic moments indicate that the complexes are paramagnetic. Infrared spectra exhibited all absorption bands at the anticipated wavenumbers, while electronic spectra revealed $n\rightarrow\pi^*$ transitions at expected wavelengths. Furthermore, results from Job's method and gravimetric analysis indicate a metal-ligand ratio of 1:2, supporting the proposed structural model. The antimicrobial activity showed significant enhancement upon complexation, underscoring the potential pharmaceutical applications of these complexes.

DATA AVAILABILITY

The data will be available on request from the corresponding author.

References

- [1] G. Balachandran, A. Dhamotharan, K. Kaliyamoorthy, K. S. Rajammal, R. Kulandaya & A. Raja, "Synthesis, characterization, and catalytic applications of schiff-base metal complexes", Journal of Engineering Proceedings **61** (2024) 26. <http://dx.doi.org/10.3390/engproc2024061026>.
- [2] M. S. Sinicropi, J. Ceramella, D. Jacopetta, A. Catalano, A. Mariconda, C. Rosano, C. Satumino, H. El-Kashef & P. Longo, "Metal complexes with schiff bases: data collection and recent studies on biological activities", Journal of Molecular Science **23** (2022) 14840. <https://doi.org/10.3390/jms232314840>.
- [3] A. Kanwal, B. Parveen, R. Ashraf, N. Haider & K. G. Ali, "A review on synthesis and applications of some selected schiff bases with their transition metal complexes", Journal of coordination chemistry **75** (2022) 19. <https://doi.org/10.1080/00958972.2022.2138364>.
- [4] A. P. Mishra & S. Monica, "Synthesis, structural, and biological studies of some schiff bases and their metal complexes", Metal-Based Drugs **7** (2008) 875410. <https://doi.org/10.1155/2008/875410>.

- [5] M. A. Lotfi, K. A. Sadeeq, A. A. Laila, M. Sabri, A. M. Jazeem, Y. A. Mohammad & M. A. Wafa, "Synthesis and characterization of different complexes derived from schiff Base and evaluation as a potential anticancer, antimicrobial and insecticide agent", *Journal of Biological Sciences* **30** (2023) 3. <https://doi.org/10.1016/j.sjbs.2023.103598>.
- [6] A. Soroceanu & A. Bargan, "Advanced and biomedical applications of schiff-base ligands and their metal complexes: a review", *Crystals* **12** (2022) 1436. <https://doi.org/10.3390/cryst12101436>.
- [7] N. Sourav, D. Abhijit, D. Arjit & A. Suman, "Metal-based dugs in cancer therapy", *International Journal of Experimental Research and Review* **37** (2024) 159. <https://doi.org/10.52756/ijerr.2024.v37spl.014>.
- [8] A. Hheidari, J. Mohammad, M. Ghodousi, M. Mahmoodi, S. Ebrahimi, E. Pishbin & A. Rahdar, "Metal-based nanoparticles in cancer treatment: lessons, learned and challenges", *Front. Bioeng. Biotechnol.* **12** (2024) 1436297. <https://doi.org/10.1039/D3DT00000X>.
- [9] P. Shreyas, G. Shubhankar, K. Ravi Teja & P. Malay, "Challenges and opportunities in the development of metal-based anticancer theranostic agents", *Biosci Rep* **42** (2022) 5. <https://doi.org/10.1042/bsr20212160>.
- [10] I. Domenico, C. Jessica, C. Alessia, M. Annaluisa, G. Federica, S. Carmela, L. Pasquale & S. S. Maria, "Metal complexes with schiff bases as antimicrobials and catalysts", *Inorganics* **11** (2023) 8. <https://doi.org/10.3390/inorganics11080320>.
- [11] S. S. El-Sayed, T. M. Al-Harbi, A. N. Al-Hakimi & M. M. Abd El-Hady, "Synthesis and characterization of a metal complexes based on aniline derivative schiff base for antimicrobial applications and UV protection of a modified cotton fabric", *Journal of Coatings* **12** (2022) 8. <https://doi.org/10.3390/coatings12081181>.
- [12] C. Takeda, D. Nakane & T. Akitsu, "Recent advances in chiral schiff base compounds in 2023", *Molecules* **28** (2023) 7990. <https://doi.org/10.3390/molecules28247990>.
- [13] M. Chandra, K. Vinod & K. Sarla, "Synthesis, characterization and antibacterial activity of Schiff bases derived from thiosemicarbazide, 2-acetylthiophene and thiophene-2-aldehyde", *Int. Res. J. pharm.* **9** (2018) 153. <http://dx.doi.org/10.7897/2230-8407.097141>.
- [14] L. A. Alfonso-Herrera, S. Rosette-Luna, D. Hernandez-Romero, J. M. Rivera-Villanueva, J. L. Olivares-Romero, J. A. Cruz-Navarro, A. Soto-Contreras, A. Arenaza-Corona, D. Morales-Morales & R. Colorado-Peralta, "Transition metal complexes with tridentate schiff bases (ONO and ONN) derived from salicylaldehyde: an analysis of their potential anticancer activity", *ChemMedChem* **17** (2022) e202200367. <https://doi.org/10.1002/cmdc.202200367>.
- [15] P. S. Saini, V. Kumar, A. K Gupta, G. K. Gupta, "Synthesis, characterization and antibacterial activity of a novel heterocyclic Schiff's base and its metal complexes of first transition series", *Medicinal Chemistry Research* **23** (2014) 690. <https://doi.org/10.1007/s00044-013-0657-6>.
- [16] B. Kenan, T. Nevin, A. Ayhan, C. Naki & O. Ismail, "Synthesis, characterization and catalytic activities of some Schiff base ligands and Pd(II) complexes containing substituted groups", *Journal of Molecular Structure* **1309** (2024) 138185. <https://doi.org/10.1016/j.molstruc.2024.138185>.
- [17] T. J. Saritha & P. Metilda, "Synthesis, spectroscopic characterization and biological applications of some novel Schiff base transition metal(II) complexes derived from curcumin moiety", *Journal of Saudi Chemical Society* **25** (2021) 101245. <https://doi.org/10.1016/j.jscs.2021.101245>.
- [18] I. Tariqul, B. Nur Amin, H. Faruk, A. Ali & Z. Kudrate, "Investigation on biological activities of thiosemicarbazide derived schiff base metal complexes", *Asian Journal of Research in Chemistry* **17** (2024) 6-2. <https://doi.org/10.52711/0974-4150.2024.00002>.
- [19] A. A. Aly, E. M. Abdallah, S. A. Ahmed, M. M. Rabee & S. Brase, "Transition metal complexes of thiosemicarbazides, thiocarbonylhydrazides and their corresponding carbazones with Cu(I), Cu(II), Ni(II), Pd(II), and Ag(I): a Review", *Molecules* **28** (2023) 1808. <https://doi.org/10.3390/molecules28041808>.
- [20] T. Geetha & S. Santhi, "Synthesis, characterization, dft studies and electrochemical biological application studies of a new schiff base on from naphthalene-1,8-diamine and its Co(II), Ni(II), Cu(II) and Zn(II) Complexes", *Oriental Journal of Chemistry* **40** (2024) 6. <https://dx.doi.org/10.13005/ojc/400631>.
- [21] G. Venkatesh, P. Vennila, S. Kaya, S. Ben Ahmed, P. Sumathi, V. Siva, P. Rajendran & C. Kamal, "Synthesis and spectroscopic characterization of schiff base metal complexes, biological activity and molecular docking studies", *ACS Omega* **9** (2024) 8123. <https://doi.org/10.1021/acsomega.3c08526>.
- [22] F. Hoque, N. Nahar, F. S. Annie, A. Rahim, M. K. Hossain, M. N. Abdul Rahman, K. A. A. Alkadi, R. B. Shardin, N. A. Sulong & A. Azad, "Synthesis, characterization and complexation of schiff base ligand p-anisalcefuoxime with cu²⁺ and fe³⁺ ions; antimicrobial and docking analysis with PBP2xt0 study pharmacokinetic parameters", *Mediterr. J. Pharm Sci.* **5** (2025) 48. <https://doi.org/10.5281/zenodo.14647651>.
- [23] S. S. Sadu, S. J. Bhimane & S. Harshavardhan, "Synthesis and characterization of schiff base metal complexes derived from Naphthofuran-2-Carbohydrazide and 4(methylthio)benzaldehyde", *Int. J. of Sci. Res. Archive* **7** (2022) 466. <https://doi.org/10.30574/ijrsra.2022.7.2.0312>.
- [24] A. K. Suleiman, A. H. Sadi, A. Hadiza & D. Aminu, "Synthesis, characterization and antimicrobial evaluation of Schiff base transition metal complexes of (E)-2-methoxybenzylidene amino phenol", *Journal of chemical sciences* **11** (2023) 100. <https://www.unn.edu.ng/wp-content/uploads/2023/06/Synthesis-Characterization-And-Antimicrobial-Evaluation-Of-Schiff-Base-Transition-Metal-Complexes.pdf>.
- [25] R. Giernoth, *Solvents and Solvent Effects in Organic Chemistry*, 4th Edition, Wiley, 2011, pp. 11289-11289. <https://doi.org/10.1002/anie.201105531>.
- [26] N. B. Jitendra & S. R. Shankarsing, "Design, synthesis and biological properties of transition metal complexes of schiff base ligand derived from pyridine derivatives", *J. Adv. Sci. Res.* **13** (2022) 10. <https://scisage.info/index.php/JASR/article/view/1892>.
- [27] I. Waziri, O. O. Wahab, G. A. Mala, S. O. Oselusi, S. A. Egieyeh & H. Nasir, "Zinc (II) complex of (Z)-4-((4-Nitrophenyl)Amino)Pent-3-en-2-one, a potential antimicrobial agent: synthesis, characterization, antimicrobial screening, dft calculation and docking study", *Bull. Hem. Soc. Ethiop.* **37** (2023) 3. <https://dx.doi.org/10.4314/bcse.v37i3.8>.
- [28] A. Sayeed, J. Na'aliya & W. Ethan, "Synthesis, characterization and antimicrobial studies of metal(II) complexes with 4-methoxybenzaldehyde with p-anisidine", *Global Journal of Pure and Applied Chemistry Research* **10** (2022) 1. <https://doi.org/10.37745/gjpacr.2013/vol10no1pp.23-38>.
- [29] S. Karthik, P. Priyab & T. Gomathi, "Synthesis, characterization, antimicrobial, anti-diabetic, anti-inflammatory and anti-cancer studies of Schiff base metal(II) complexes derived from mixed Schiff base ligands", *Indian Journal of Chemistry* **63** (2024) 112. https://www.google.com/url?sa=t&source=web&rct=j&opi=89978449&url=https://or.niscpr.res.in/index.php/IJC/article/view/4556&ved=2ahUKewjgzvQqNKMAxWKWUEAHV0SFDAQFnoECBIQAQ&usq=AOvVaw2LE6mqP8YXOH__MOhEiOhm.