SYNTHESIS, STUDIES THEIR QUANTUM MECHANICAL PROPERTIES AND BIOLOGICAL ACTIVITY OF NEWER ANTIPYRIN DERIVATIVE AND ITS (CU2+) COMPLEX

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ABSTRACT

The synthesis of (C16H14N3O2) ligand and (Cu2C32H28N6O12S2) complex are prepared in this work . The newly prepared compounds were characterized by IR spectroscopy and elemental analysis . The determination of the antimicrobial activity of ligand and its complex was carried out on sample of Escherichia coli , Klebsiella pneumonia , Staphylococcus aureus and Bacillus cereus . The quantitative antimicrobial activity test results proved that all the prepared compounds are very active against Bacillus cereus and that complex very active against Escherichia coli. Quantum mechanical calculation confirmed the assumption of stability of prepared compounds which agreement with antimicrobial activity results .

INTRODUCTION

Antipyrine and its derivatives have been widely used in different field of medicine over the last few years , the pharmacological and biological activities(1-4) of antipyrine derivatives as well as the antitumour(5,6) , antimicrobial activities(7) are also expected . Therefore with the aim of producing biologically active drugs , the newly synthesized compound (C16H14N3O2) and its complex (Cu2C32H28N6O12S2) Were established based on the elemental and spectral analyses which are in good agreement with these structure , Quantum . chemical parameters help to study the geometrical optimization and heat of formation of these compound . The antimicrobial activity of the antipyrine complexes have been discussed previously(8-15).

Complex Combinations of copper are used in the treatment of a variety of disease, including inflammatory processes, cancer, ulcers, nervous system and heart diseases further more we have tested the antimicrobial activity of the prepared compound and its complex using strains of Escherichia coli, klebsiella pneumonia staphylococcus aurous and Bacillus cereus.

Experimental Part :

1 – Instruments used :

Infra – red spectrophotometer (pye – Unicam model sp- 3 - 300 S) CHN analysis (Carlo Erba EA 1108 elemental analyzer)

2 - Synthesis of (C16H14N3O2) Compound(15):

A solution of 4 – amino antipyrin (1.25m mol) in methanol (15 ml) was added to a solution of terephthalic aldehyde (0.5m mol)in methanol(15ml) the resulting yellow solution was refluxed for (30 minutes) and then left at room temperature for(6 hrs). The intense yellow precipitate formed was filtered, washed with methanol and dried.

3 - Synthesis of (Cu2 C32 H28 N6 O12 S2) Complex(13): -

A DMF solution (15 ml) (C16 H14 N3 O2) (0.5 m mol) was added to aqueous copper sulphate (1 m mol) dissolved in distiller water (15ml) the resulting green solution was refluxed for (2 hours). this solution was left at room temperature for (4 days). A brown precipitate was formed which was filtered, washed with ethanol and dried.

4 - Quantum - mechanical calculations :

For quantum-chemical calculations we have used two important programs Alchemy 2000 and Hyperchem (MNDO semiempirical method) wich were used for geometrical optimization and heat of formation calculation of investigated compounds.

5 – Biological assay : -

The fresh cultures obtained from clinical isolates were suspended in distiller water and adjusted to a standard density of (0.5 macfarland).

The microbial suspensions were plated on solid Mueller hinton medium

and solutions of the test compounds (10 μ L) prepared in DMF(1 mg/ml)

were added on filter paper disks concomitantly, these disks were impregnated with the same concentration of gentamycin, which was used as reference standard for reporting the antibiotic sensitivity. The plates were incubated at (37° C) for (24 hrs). The quantitative antimicrobial activity assays were performed by the two - fold serial microdilution method in liquid solution of test compound medium (nutrient broth)(8).

Serial tow – fold dilutions of a solution of test compound in DMF from (1000 to 62.5μ g/ml) in a total volume of (200 μ l)medium and standard microbial suspension (50 ml) was added in each well. After (18-24hours) The plates are examined visually for evidence of bacterial growth .

RESULTS AND DISCUSSION

1 – infra – red spectra

IR spectra were recorded by using KBr disc were shown in table (1). IR

Spectra of (C16H14N3O2)compound was observed that the frequency of the specific band of the (Vc=N) bond (1654 cm-1) is moved towards lower wavenumbers by approx. (15 cm-1) in the spectra of the complex (Cu2 C32 H28 N6 O12 S2) which confirms the coordination of the nitrogen atom to the metallic ion. The specific (V>c=o) bond (1595cm-1) of ligand moves towards lower wave number (1565 cm-1) of complex, suggesting the coordination of the ligand to the metallic ion via the (>c=o) group. In complex a characteristic band corresponding to the bidentate coordination of the (So4²-) ion appears. Thus the (V1)and(V2) frequencies specific to a (Td) arrangement appear as medium intensity bands, (V3) and (V4) frequencies each split in to three bands, which suggest a low symmetry(16) in complex the (Cu²+) ion has a deformed tetrahedral geometry (17).

2 – Elemental analysis :

The percentage of carbon, hydrogen and nitrogen of prepared compounds are shown in table (2), the elemental analysis results of prepared compounds are in a good agreement with the calculated values

Tabl (1): IR spectral data for the prepared ligand and con	plexe
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				١	Vso42-	
compound	Vc=N	V>C=O	V1	V2	V 3	V4
C16H14N3O2	1654	1595				
					1050	574
Cu2 C32 H28 N6 O12 S2	1639	1565	995	462	1105	610
					1170	641

Table (2): (CHN) analysis of ligand and complex

compound	Practical values				Theoretical values			
	C%	Н%	N%	Cu%	C%	H%	N%	Cu%
C16H14N3O2	70.58	6.28	16.2 1		68.5 7	5.00	15.0 0	
Cu2 C32 H28 N6 O12 S2	46.12	3.31	9.32	14.09	43.6 7	3.18	9.55	14.44

3- Antimicrobial activity assays : -

The antimicrobial activity assays of the (C16 H14 N3 O2) and (Cu2 C32 H28 N6 O12 S2) were performed by quantitative methods based on serial two–fold dilutions of the test compounds in order to establish the corresponding minimal inhibitory concentrations(MIC)at the highest dilution(lowest concentration) of the tested compound

that completely inhibited microbial growth four bacterial strains(Escherichia coli , Klebsiella pneumonia , staphylococcus aureus and Bacillus cereus) were cultivated on solid media and incubated at (37°C) for (24 hrs) prior to testing .The quantitative assay results (Figure -1-) showed that the tested compounds exhibited variable (MICs) and selective antimicrobial activity depending on the microbial strains . All tested compounds proved to be active on Bacillus cereus well known for its high constitutive and acquired resistance rates and(Cu2 C32 H28 N6 O12 S2) complex showed good activity against E.coli proving their potential usefulness as broad spectrum antimicrobial agents.

4 – Quantum mechanical results :-

Quantum mechanical calculation of heat of formation (Hf) for ligand and its complex which were prepared in this work has been shown in table (3)

Compound	Hf (kcal / mol)
C16H14N3O2	197.45
Cu2C32H28N6O12S2	175.06

Table (3): Heat of formation (Hf) of ligand and complex

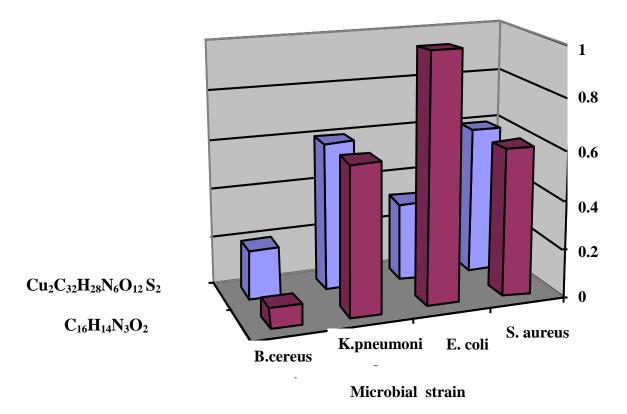


Figure 1:The graphic representation of the MIC values (mg / ml) of the tested compounds towards different bacterial strains

CONCLUSION

- 1 The IR spectral and (CHN) analysis results indicate that the structure of prepared compounds are as expected.
- 2 The quantitive anti microbial activity test results proved that both the ligand and their complex have specific anti microbial activity depending on the microbial species tested.
- 3 Heat of formation results proved that the ligand stability increases when its coordinated with (Cu²+) ion therefore its has a great covalency and prevent it from decomposition when it penetrate the cell well then these are agreement with anti – microbial activity test results which its increases when ligand compound are coordinated with (Cu²+) ion.

تحضير ودراسة الخواص الميكانيكية والفعالية البايلوجية لمشتقات جديدة للانتي بايرون ابتهال حنظل التميمي ، رواء محمد كلية الصيدلة – جامعة البصرة – البصرة – العراق

الخلاصة

أهـ تمت الدراسـة ف_ي هـذا البحـث تحض ير الليكاند (C16H14N3O2) والمـعقد (O12S2) Cu2C32H28N6) وقد شخصت بطريقة تحليل العناصر والمطيافيه وعند دراسة الفعالية البايلوجية ضد أربعة أنواع من البكتريا أظهرت النتائج أن جميع المركبات المحضرة لها فعالية ضد بكتريا B.cereus وان المعقد الناتج من اتحاد الليكاند مع (+Cu2) أظهر فعاليه ضد بكتريا (E.coli) ايضا وان المعالجة الكمية للمركبات المحضرة في هذه الدراسة بينت إن أستقر ارية المعقد أعلى من الليكاند و هذا يتوافق مع نتائج دراسة الفعالية البايلوجية ضد البكتريا لهذه المركبات

REFERENCES

- 1 Harb. A.A., Elagamey .A.A. and Khodeir.M.N.: Imidazolene derivatives of antipyrin.
 Egypt. J. pharm. Sci. 29. 279- 288 (1988).
- 2 Eid. A. I., Mikhael. A.N. and Rashad. S: Synthesis of some pyrazolone derivatives of expected pharmacological activity. Egypt. J. pharm Sci. 29. 381- 391 (1988).
- 3 Elagamey. A.A., Harb. A.A. Khodeir. M.N. and Sawellim.S.Z., Synthesis of midazole derivatives containing biologically active units Arch. pharm. Res., 10. 153-157 (1987).
- 4 Dahlobom. R. and Ekstrand. T.: 10 –Aminoacctylphenothiaziene derivatives. Acta. Chem., 6. 1285 (1952).
- 5 Ebeid. M. Y., El Moghazy. S. M., Eissa. A.A.H. and Osman. A. M.: synthesis and antitumour poiential of isosteres of acridine antineop lastic agents. Egypt. J. pharm. Sci., 31. 427 – 436 (1990).
- 6 El Moghazy. S. M. and Safwat. H.M.: Synthesis and antiumour activity of some acridonanil derivatives. Egypt. J. pharm. Sci.31 505 513 (1990).
- 7 Abu Shady. H. A., Ragab. F. A. and Ali. E. L. Hydrazones, thiosemicarbazide. tiosemicarbazones and 1,3-thiazolidin-4-ones derived from acridine, synthesis, cyeotoxic and antimicrobial activities. Egypt. J. pharm. Sci., 31. 285 – 300 (1990)

- 8– Raman,N., Kulandaisamy,A., Shunmugasundaram, A., Jeyasubramanian,K., synthesis,Spectral,redox and antimicrobial activities of Schiff base complexes derived from1–phenyl–2-3–dimethyl-4-aminopyrazol - 5 – one and acetoacetanilide. Transit. Metal chem.,26.131-135, (2001).
- 9 Raman,N., Kulandaisamy,A., Jeyasubramanian, K. Synthesis,Spectral, redox and antimicrobial activites of Schiff base transition metal (II) complexes derived from 4aminoantipyrine and benzil Synth. React. Inorg. Met-Org. Nano– Met. Chem.,32.1583 – 1610, (2002).
- 10 Singh, L., Sharma, D.K., Singh,U., Kumar, A. synthesis and spectral studies of Cu(II) coordination compounds of 4[N-(cinnamalidene) amino] antipyrine semicarbazone. Asian J.Chem., 16, 577–580, (2004).
- 11 Raman, N., Thangaraja, C., Johnsonraja, S. Synthesis, spectral characterization, redox and antimicrobial activity of Schiff base transition metal (II) complexes derived from 4-aminoantipyrine and 3-salicylideneacetylacetone.Centr. Eur. J.chem., 3, 537-555, (2005).
- 12 Pandey,O. P., Sengupta,S.K.,Dwivedi,A. Organophosphorus derivatives containing antipyrine ring as chemotherapeutics against fungal pathogens of sugarcane. Electron. J. Environ. Agr.Food chem., 4, 886 891, (2005).
- 13 Agarwal,R.K.,Singh,L., Sharma,D.K. synthesis. spectral and biological properties of copper (II) complexes of thiosemicarbazones of schiff bases derived from 4aminoantipyrine and aromatic aldehydes. Bioinorg. chem.Appl. article ID 59509, (2006).
- 14 Agarwal, R.K., Gargb,R.K., sindhub, S.K. synthesis and magneto- Zpectral investigations of some six and nine coordinated complexes of lanthanides (III) derived from 4 [N-(2-hydroxyl -p- napthalidene) amino] antipyrine thiosemicarbazone, J. Iran.Chem.Soc.,2, 203 211 (2005).
- 15 Agarwal R. K., Prasad. S.,Synthesis and spectral investigations of some platinum metals ions coordination compounds of 4[N-(furan-2- carboxalidene) amino] antipyrine thiosemicarbozone and 4 [N (3,4,5-trimethoxybenzalidene) amino]antipyrine thiosemicarbazone. Turk. J.Chem.,29,289-29,(2005).

- 16 Nakamoto,K.,Infrared Spectra of Inorganic and Coordination compounds Wiley and Sons,New york, 248 249, (1986).
- 17 Lever, A. B. P. Inorganic Electronic Spectroscopy. 2nd Edn, Elsevier science, New york, 560 – 571, (1984).