



IJPPR

INTERNATIONAL JOURNAL OF PHARMACY & PHARMACEUTICAL RESEARCH

An official Publication of Human Journals

ISSN 2349-7203




Human Journals

Research Article


January 2016 Vol.:5, Issue:2

© All rights are reserved by J. Sreeramulu et al.

Synthesis, Characterization and Biological Evaluation of Novel Bio- Active Dopamine Carbamodithiolate Metal Complexes

 **IJPPR**
INTERNATIONAL JOURNAL OF PHARMACY & PHARMACEUTICAL RESEARCH
An official Publication of Human Journals

ISSN 2349-7203



A.Jayaraju, and J. Sreeramulu*

*Department of Chemistry, Srikrishnadevaraya
University, Anantapur, India.*

Submission: 25 December 2015
Accepted: 1 January 2016
Published: 25 January 2016

Keywords: Dopamine, Carbamodithiolate ligand, Elemental Analysis, Copper and Ruthenium Chlorides

ABSTRACT

A series of novel bi dentate carbamodithiolate ligand Dopamine was prepared by new synthetic method. The synthetic sodium salt of Dopamine Carbamodithiolate ligand is followed by the reaction of Copper and Ruthenium Chlorides to get corresponding complexes. The complexes were characterized by Elemental Analysis, IR, ¹HNMR, ESR, TGA-DTA and antimicrobial analysis.



www.ijppr.humanjournals.com

INTRODUCTION

There has been intense interest in the coordination compounds of unsaturated sulphur donor chelating ligands, Carbamodithiolates, and their related molecules from chemists, physicists, biologists and theoreticians alike owing to their interesting chemical properties and possible wide applications¹⁻⁴. Interest in molecular structural investigations and chemical studies of these metal chelates covers a full gamut of areas ranging from general considerations of metal-sulphur bonding and the formation of four-membered chelate rings to the employment of these ligands in inorganic qualitative analysis⁵, their practical application in organic synthesis⁶, medicine⁷, and biology⁸, and their uses as vulcanisation accelerators⁹, floatation agents, "fungicides"¹⁰, pesticides¹¹ radiation protectors,¹² antioxidants¹³ and photostabilisers of polymers¹⁴. Their role in material science has also been quite significant. The interesting low spin, high spin crossover phenomenon was first reported in an iron(III) Carbamodithiolate complex¹⁵. There are several metal Carbamodithiolate complexes with bridging sulphur centres which are known to participate actively in super exchange phenomenon imparting novel magnetic properties to these systems¹⁶. In this article, the interesting ligation characteristics of carbamodithiolates and structural features of their various transition metal complexes in general and the coordination chemistry and stereochemistry of complexes in particular, along with the scope of in this article which covers the relatively unattended aspect of the primary amine derived Carbamodithiolates, synthesis, characterization and antimicrobial activity of Cupper (II) and Ru (II) Complexes of Carbamodithiolate.

MATERIALS AND METHODS

Experimental Section

Copper chloride anhydrous was obtained from Fluka, Dopamine and carbon di sulfide were purchased from Aldrich. Other chemicals used were of analytical reagent or higher purity grade. Solvents used were of reagent grade and purified before use by the standard methods. Conductivity measurement was carried out by a Systronics conductivity bridge 305, using a conductivity cell of cell constant 1.0 double distilled water was used as solvent. Electronic absorption spectra on JAS.CO UV/VIS-7850 recording spectrophotometer. Infrared spectra was recorded on a JAS.Co-460 plus FT-IR spectrophotometer in the range of 4000-400 cm^{-1} in KBr

pellets. Micro chemical analysis of carbon, hydrogen and nitrogen for the complexes were carried out on a Herause CHNO-Rapid elemental analyzer. ^1H NMR spectra were recorded on a Bruker DRX-500 Advance spectrometer at 500MHz in DMSO-discussing tetra methyl silane as internal reference standard. Melting points were measured on a unimelt capillary melting Point apparatus and reported uncorrected.

Preparation of Sodium salt of Carbamodithiolate ligands

0.05 mol of amine was dissolved in 30 ml of absolute alcohol in a clean beaker which was placed in ice bath. To this cold solution add 5 ml of Sodium hydroxide (10N) solution, and then add Pure carbon disulphide (0.05ml) in drop wise with constant stirring. The contents were stirred mechanically for about 30 min, sodium salt of Carbamodithiolate precipitated out. It was dried over and recrystallized from ethanol.

Preparation of Cu (II) and Ru (II) Complexes

Synthesis of $[\text{Cu} ((\text{DCDT}))_2]\text{Cl}_2$

The aqueous solution of 0.05 mol of Cupper Chloride was added with constant stirring to an aqueous solution of 0.01 mol of Sodium salt of Dopamine Carbamodithiolate ligand. The reaction mixture was stirred at room temperature for 2 hours. The colored (yellow) precipitates were obtained. The precipitates were filtered and washed with water and then with methanol and dried over calcium chloride in desiccator's Yield:78% and decomposes at 110 C.

Anal. Calcd. For

C,38.59;H,4.97;N,6.16;Cu,12.70;O,:18.67,S,24.61;Found:C,37..59;H,4.27;N,5.16;Cu,11.70;O,:17.67,S,23.61

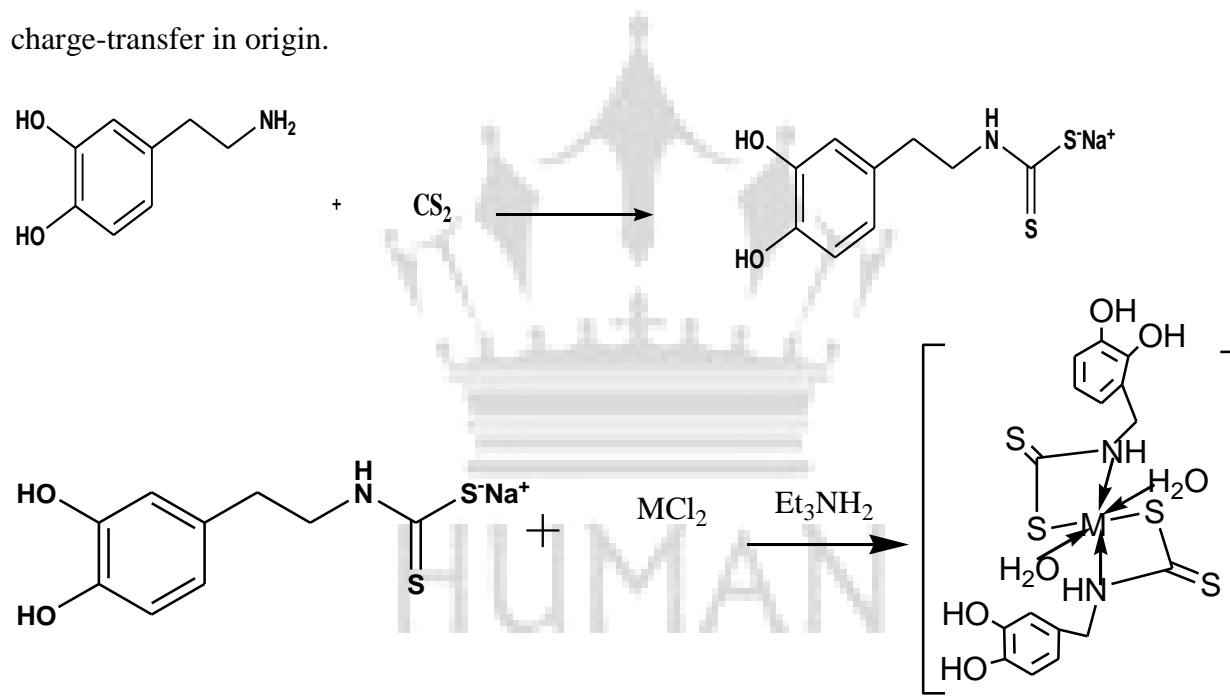
Synthesis of $[\text{Ru} (\text{DCDT})\text{Cl}_2]$

The aqueous solution of 0.05 mol of Manganese Chloride was added with constant stirring to an aqueous solution of 0.01 mol of Sodium salt of Dopamine Carbamodithiolate ligand in the presence of small quantity of triethylamine. The reaction mixture was stirred at room temperature for 2 hours. The colored (gray) precipitates were obtained. The precipitates were filtered and washed with water and then with methanol and dried over calcium chloride in a

desiccator Yield: 80% and decomposes at 110⁰C. Anal. Calcd. For C, 26.43; H, 4.44; N, 6.17; Cl,15.60; Mn,12.09 Found: C:35.16; H:3.9; N:4.82; O:16.53; Ru,17.40; S:22.09

RESULTS AND DISCUSSION

Solid reflectance spectra data for the Cu and Ru of Dopamine metal complexes. The complexes [Cu(DCDT)₂Cl₂] and [Ru(DCDT)₂Cl₂] complexes exhibit magnetic property and has an electronic spectrum which can be assigned to low spin Cu (II) and Ru (II) in an Octahedral Environment. Intra ligand electronic transition in then...C...S...S and S...C...S chromophers of the Carbamodithiolate moiety. Thus the peak at 646 nm and the shoulder at 499 nm arise from 1A_{1g}-1T_{1g} and 1A_{1g}-1T_{2g} transitions, respectively. The other lower peaks are probably charge-transfer in origin.



Infrared Spectrum

Two regions of the IR spectrum of the [Cu (DCDT)₂Cl₂] and [Ru(DCDT)₂Cl₂] complex have proven valuable in arguments concerning the electronic and structural characteristics of this compound. The presence of the thiouride band between 1545-1430 cm⁻¹ suggest a considerable double bond character in the C...N bond vibration of the S₂C-NR₂ group. The band present in the 967 cm⁻¹ range is attributed to the prevailing contribution of (C...S) Vibrations in these ranges have been used defectively in differentiating between monodentate, bidentate

carbamodithiolate ligands. The presence of only one strong band supports bidentate coordination of the dithioligands, whereas a doublet is expected in the case of monodentate coordination. (C...S) and (C..N) Stretching frequencies fall in the 1035 cm^{-1} (1001 cm^{-1} for the free ligand) and 1478 cm^{-1} respectively. The methyl group in the complex, as medium strong bands in the 2960 cm^{-1} range can be related to the asymmetric CH_3 stretching vibration.

H^1 -NMR Spectra

The NMR spectrum of the $[\text{Cu}(\text{DCDT})_2\text{Cl}_2]$ and $[\text{Ru}(\text{DCDT})_2\text{Cl}_2]$ complexes showed at 2.3-2.4 ppm. Which may be assigned to the hydroxyl protons. The peak at 7.9-7.98 attributed to NH protons of thiouraid nitrogens in both complexes. In other signals is also appeared in the region 0.98, 1.5, 3.8 ppm.

Antimicrobial Activity

Antimicrobial test was performed on four bacterias (*Staphylococcus aureus*, *Bacillus subtilis*, *Escherichia coli*, *Pseudomonas aeruginosa*,) and two fungi (*Candida albicans* and *Candida tropicalis*). The media used were prepared by dissolving separately 2g of nutrient broth powder and 38g of the Mueller-Hinton agar powder in 250 ml and 1 L of deionized water, respectively. Two media were sterilized in an autoclave at 121°C for 15 minutes and then stored overnight in a refrigerator after cooling. Cultures of the microorganisms were prepared in sterile nutrient broth and incubated for 24 hours at 37°C for the bacteria and 27°C for the fungi. 0.1 ml of each of the overnight cultures in sterile test tubes with caps were made up to 10 ml with 9.9 ml of sterile deionized water

To give 1:100 or 10⁻² dilution of the microorganisms. The technique used for the study was agar-well diffusion. Solutions of concentration 10 mg/ml of the compounds were made in dimethyl sulphoxide (DMSO). DMSO was also used as the negative control. Positive controls for bacteria and fungi were discs of commercial antibiotics manufactured by Abtek Biological Limited and Fluconazole dissolved in DMSO. The discs were carefully placed on the inoculated media with the aid of sterile forceps. Plates inoculated with bacteria were incubated at 37°C for 24 hours, and those inoculated with fungi were incubated at 27°C for 72 hours. Afterwards, the zones of inhibition of microbial growth that appeared around the wells of the compounds were examined and the diameters measured and recorded in millimeters (mm). Antimicrobial activity

of the Cu (II) and Ru (II) complexes was evaluated *in vitro* against Gram positive bacteria- *Staphylococcus aureus* and *Bacillus subtilis*, Gram negative bacteria- *Escherichia coli*, *Pseudomonas aeruginosa* and fungi, *Candida albicans*, *Candida tropicalis*. The results for the complexes and commercial antibiotics used as positive controls are listed in (Table 1).

Growth inhibition zone in millimeters (mm)						
BACTERIA					FUNGUS	
	Gram +Ve		Gram -Ve			
	<i>S. aur</i>	<i>B. subt</i>	<i>E. coli</i>	<i>P. aerug</i>	<i>C. alb</i>	<i>C. trop</i>
Cu(DCDT) ₂	14	14	15	14	14	15
Ru(DCDT) ₂	13	15	14	114.5	16	14
FLU			----	----	15	15
DMSO	12	14	15	16	----	---

CONCLUSION

Cu (II) and Ru (II) complexes of Dopamine Carbamodithiolate Ligand have been synthesized and characterized. The ligand moiety exhibit a bidentate coordination mode in the Cu (II) and Ru (II) complexes. Solid reflectance spectra and magnetic data indicate that the complexes are Paramagnetic and Octahedral. The complexes show selective activity towards some of the test microorganisms.

ACKNOWLEDGEMENT

The authors are greatly thankful to the Department of Chemistry, Sri Krishnadevaraya University, Anantapur, India is acknowledged for the microanalysis, infrared spectra and thermogravimetric analysis, conductivity and magnetic susceptibility measurements of the compounds.

REFERENCES

- [1] AAM Aly; MM Kamal; MS El-Meligy; ASA Zidan; M El-Shabasy. *Synth. React. Inorg. Met.-Org. Chem.*, **1987**, 17(3), 237–274.
- [2] AAM Aly; MS El-Meligy; ASA Zidan. *Transition Met. Chem.*, **1989**, 14, 366–368.
- [3] AI El-Said; AAM Aly. *Synth. React. Inorg. Met.-Org. Chem.*, **1990**, 1059–1069.

- [4] PA Ajibade; GA Kolawole. *J. Coord. Chem.*, **2008**, 61(21), 3367–3374.
- [5] A Hulanicki. *Talanta*, **1967**, 14, 1371–1392.
- [6] D Coucouvanis. *Prog. Inorg. Chem.*, **1970**, 11, 233–371.
- [7] G Manoussakis; C Bolos; L Ecateriniadou; C Sarris. *Eur. J. Med. Chem.*, **1987**, 22, 421–425.
- [8] L Giovagnini; C Marzano; F Bettio; D Fregona. *J. Inorg. Biochem.*, **2005**, 99, 2139–2150.
- [9] A Manohar; K Ramalingam; R Thiruneelakandan; G Bocelli; L Righi. *Z. Anorg. Allgem. Chem.*, **2006**, 632, 461–464.
- [10] R Pastorek; J Kameníček; J Husárek; V Slovák; M Pavlíček. *J. Coord. Chem.*, **2007**, 60(5), 485–494.
- [11] BA Prakasam; K Ramalingam; R Baskaran; G Bocelli; A Cantoni. *Polyhedron*, **2007**, 26, 1133–1138.
- [12] M Sarwar; S Ahmad; S Ali; SA Awan. *Transition Met. Chem.*, **2007**, 32, 199–203.
- [13] Z Trávníček; R Pastorek; V Slovak. *Polyhedron*, **2008**, 27, 411–419.
- [14] ASA Zidan. *Synth. React. Inorg. Met.-Org. Chem.*, **2001**, 31(3), 457–469.
- [15] V Pawar; S Joshi; V Uma. *J. Chem. Pharm. Res.*, **2011**, 3(1), 169–175.
- [16] AH El-Masry; HH Fahmy; SHA Abdelwahed. *Molecules*, **2000**, 5, 1429–1438.

