# Microbial Screening of Cobalt (II) and Nickel (II) Complexes of Acetone Thiosemicarbazone, Synthesis and Characterisation

#### V. Sudha<sup>1</sup>, G. Gohulavani<sup>2</sup>

<sup>1</sup>Assistant Professor, Department of Chemistry, SNMV College of Arts and Science, Coimbatore-50

<sup>2</sup>Assistant Professor, Department of Chemistry, Ethiraj College for Women, Chennai-08

**Abstract:** The present investigation involves the preparation and characterization of Nickel (II) and Cobalt (II) nitrate complexes using acetone thio semicarbazone as ligand. The complexes were characterized by conductivity, spectral and biological studies.

Keywords: Acetone thio semicarbazone, conductivity, spectral studies, biological studies

#### 1. Introduction

Thiosemicarbazone complexes of transition metals have attracted special attention in the field of biology, due to their effective biological commotion against virus, protozoa and certain kinds of tumor and also antitubercular activity <sup>(1-3)</sup>. The most active research area in the field of inorganic chemistry is Co- ordination chemistry. Co-ordination compounds play an essential role in industry and in Biology and its significance depends on elements, particularly metals in various oxidation states which exhibit the capacity of binding donor molecules. The Co-ordination complexes formed by Cobalt (II) are found to be brown or brownish black crystalline solids <sup>(4-5).</sup> The co-ordination complexes formed by Nickel (II) are found to be Octahedral or distorted Octahedral having light pink or blue or green crystalline solids <sup>(6-9).</sup>

#### 2. Experimental Work

#### 2.1 Preparation of Ligands

#### Preparation of acetone thiosemicarbazone

The following general method was adopted for the preparation of Acetone thio semicarbazone

Ethanolic solution of thiosemicarbazide and acetone were mixed in the molar ratio of 1: 1. The mixture was refluxed for about 6 hours in a water bath. After the completion of reaction, excess of the solvent was removed by filtration and the white coloured product obtained was dried. It was further purified by recrystallisation using the same solvent.

#### 2.2 Preparation of complexes

#### i) <u>Preparation of Nickel (II) nitrate complexes of the</u> <u>ligands acetone thiosemicarbazone:</u>

Ethanolic solution of nickel (II) nitrate and acetone thiosemicarbazone were mixed in the molar ratio of 1 :3. The mixture was heated in a water bath for 2 hours. The excess solvent was evaporated and dried. Washing with acetone and ether for several times purified this. The complex was pink colour solid. Decomposition temperature was found to be above  $190^{\circ}$  C.

## ii) <u>Preparation of Cobalt (II) nitrate complex of the ligands acetone thiosemicarbazone</u>:

Ethanolic solutions of cobalt (II) nitrate and acetone thiosemicarbazone were mixed in the molar ratio1: 3 and the mixture was heated on a water bath for 2 hours. Then excess alcohol was evaporated and the residue was washed with acetone and ether several times. The complex was found to undergo decomposition above190°C. The complex was soluble in water and sparingly soluble in alcohol.

#### 3. Results and Discussion

The present investigation involves the preparation, characterization and biological investigation of Nickel (II) and Cobalt (II) nitrate complex using acetone thio semicarbazone as a ligand. The complexes were characterized by conductivity, spectral and biological studies.

The Nickel (II) nitrate complex was prepared using the ligand acetone thio semicarbazone. The Nickel(II) nitrate complex was found to be pink coloured amorphous solid. It was soluble in water and sparingly soluble in alcohol but not in ether, acetone and chloroform. The Cobalt (II) nitrate complex was prepared using the ligand acetone thio semicarbazone. The complex thus obtained was found to be brown colored amorphous solid. It was soluble in water and sparingly soluble in alcohol but not in ether, acetone and chloroform.

#### **Physical and Conductivity study**

The melting point of the ligand and the yield of Nickel (II) and Cobalt (II) complexes and the conductance data of these two complexes are given in the table 1.

**Table 1** Physical Data of the Ligand and Ni(II) and Co(II)

 Complexes

Name	% Yield	Melting Point	Conductance
Ligand	82%	196	56
Nickel(II) Complexes	86%	128	65
Cobalt(II) Complexes	85%	136	87

## Volume 7 Issue 1, January 2018

#### Licensed Under Creative Commons Attribution CC BY

#### **Infrared spectra**

The IR spectra of acetone thio semicarbazone, Nickel (II), Cobalt (II) nitrate complex were taken and analyzed mainly for certain specific frequency which are directly involved in complex formation and the values are given below the table 2.The spectra of the respective ligand and complexes are shown in the figure 1,2,3.

Considerable changes in I-R spectra of the Nickel (II) complex is the appearance of v(C=N) band in the region of 1604.77as compared to 1652.32cm<sup>-1</sup> in the ligand thus showing a slit towards lower wave number. Coordination of azomethane is confirmed by the presence of a new band at 421-492 cm<sup>-1</sup> which can be assigned to v(M-N)for Nickel(II) and Cobalt(II)complexes.

The IR Spectra of the ligand exhibit a medium intensity band at1268.51 cm-<sup>1</sup>due to C=S stretching vibrations. The corresponding band in the spectra of the Nickel (II), Cobalt (II) nitrate complex appear in the region of 1235.9&1230.0 respectively. This lower shift may due to the participation of thiosemicarbonyl group in complexation. This is also supported by the appearance of new band in the region of 420 due to v (M-S) stretching vibrations. The broad band in the region 3400-2900 cm<sup>-1</sup> indicates the presence of water molecules. These regions appear due to OH stretching vibrations merged with NH absorption bands. A sharp peak at 1635.44 is due to H-O-H bending vibrations.

Generally the IR spectrum of the nitrate ion v2 (out-of plane deformations), v3 (double degenerate) v4 (double degenerate in the plane bending). The IR spectra of coordinated nitrate has been reported in literature survey.v1 (symmetric) is normally inactive but sometimes becomes weakly active through crystal; interactions. The magnitude of v4-v1 decides the nature of the coordinating nitrate group. A difference if nearly 100 cm<sup>-1</sup>shows the presence if monodentate nitrate groups and for bidentate nitrate groups this difference is greater than 150 cm<sup>1</sup>. Both the Nickel (II), Cobalt (II) nitrate complexes exhibit absorption at 1384 cm<sup>-1</sup> characteristic of the ionic nitrate groups.

Table 2: Im	portant IR Absor	ption Bands (in o	cm) Of Ligands.	Ni(II) and C	Co(II) Ni	trate Coplexes
	portune 110 1000	phon Dunus (m	only of Eigunds,	i ii(ii) uiia (	20(11) 1 11	unde copieses

1 1		· /	U I		~ /		1	
Name of the Sample	ν	ν	ν	ν	ν	ν	ν	ν
	N-H	C-H	C=N	C=S	CO-H	H-OH	N-N	NO <sub>3</sub> Ionic
LIGAND- ACETONE THIOSEMICARBAZONE	-	2997.9	-	1268.5	3371.8	-	-	-
NI(II)NITRATE COMPLEX	-	2954.8	1604.8	1235.9	3350.4	1635.4	434.1	1384.1
CO(II)NITRATE COMPLEX	-	082.8	-	1230.0	-	1629.3	-	1384.2

#### **Biological studies**

Thiosemicarbazone possess a wide spectrum of medicinal properties. The ligand and the metal complexes were screened for antibacterial activities against some of the pathogenic bacteria like Pseudomonas putdita and Pseudomonas aures by using cup method. The diameter (mm) of the inhibition zone around each cup was measured after 24hours and results are listed in table 3.

The results clearly illustrate that the compounds have antibacterial activity against all the organisms tested. The ligand acetone thiosemicarbazone have no effect on the growth of bacterial compared to metal complexes. The antibacterial activity of nickel (II) complexes against Pseudomonas putdita and Pseudomonas aures were much higher compared to other bacterias used. Further it was confirmed all the complexes have higher activity at higher concentration (1000 $\mu$ g) than lower concentration (500  $\mu$ g).

The work assumes importance as many metal thiosemicarbazone are found to have significant antibacterial activities. The results reveal that all the metal complexes are more toxic than the ligand.

Tabl	e 3	5	Antiba	cterial	Activ	ity	of	the	Liga	nd	and	Metal
Com	plex	xe	s Evalı	lated E	Bymini	imu	m I	nhib	itory	Co	ncen	ration

Ligand/Comlexes	Pseudomonas	Pseudomona	
Engund/Connexes	auras	putdita	
S1- CO(II) Thiosemicarbazone complex	+ + + +	++++	
S2- Ligand	+	+	
S3- NI(II) Thiosemicarbazone complex	+ + + +	++++	

**Inhibition diameter in MM (% inhibition);** +, 6-10 (27-45%); ++ 10-14 (45-64%); ++ +: 14-18 (64-82%); ++ ++, 18-22 (82-100%). Per anti inhibition values are relative to inhibition (22mm) with 100% inhibition.

The ligand and the metal complexes were screened for antifungal activities against some of the pathogenic fungal like Aspergillus Fumigates and Trichoderma Veridi using cup method. The diameter (mm) of the inhibition zone around each cup was measured after 24hours and results are listed in table 4

Table	4:	Antifungal	Activity	of	the	Ligand	and	Metal
Comp	lexe	s Evaluated l	by Minim	um	Inhit	oitory Co	ncent	ration

Ligand/Complexes	Aspergillus	Frichoderma
	Fumigatus	Veridi
S1- Co(II) thiosemicarbazone complex	+ + + +	+ + + +
S2- Ligand	+	+
S3 - Ni(II) thiosemicarbazone complex	+ + + +	+ + + +

Inhibition diameter in MM (% inhibition) ; +, 6-10 (27-45%) ; +1 10-14 (45-64%) ; + + + :14-18 (64-82%) ; + + + +,18-22 (82-100%). Per anti inhibition values are relative to inhibition (22mm) with 100% inhibition.

## 4. Conclusion

Acetone thio semicarbazone and their complexes were found to have significant role in antimicrobial activities. The yield obtained was good and the biological studies also resulted well.

Volume 7 Issue 1, January 2018 www.ijsr.net

Licensed Under Creative Commons Attribution CC BY

#### References

- [1] Negi Parul, Nandy Subhangkar, Mahato Arun, IRJP 2012.3(5).
- [2] T. Priya Devi\*and R. K. Hemakumar Singh, RASAYAN j.chem, Vol.3, No.2 (2010), 266-270
- [3] Hossain Mohammad Zakir, Mele Jesmin and Shaikh Mohammad Mohsin Ali, Asian J. Med. Pharm. Res. 6(4). 32-40, Dec 25, 2016.
- [4] Mehdi Amirnasr etal, Polyhedron 21 (2002) 2733\_/2742
- [5] M.K Gupta etal, AsianJ.Chem, Vol.19, No.1(2007), 5-9
- [6] Hitesh dahyabhai Patel,etal,Indian J,Chem, Vol.52B, April 2013,pp 535-545
- [7] Tripti Kamalpuria, Megha Chourey, Nameeta Bende, Anish Ghuraiya, Arpan Bhardwaj, Int. J. Curr. Res. Chem. Pharm. Sci. (2016). 3(5): 30-35
- [8] Valbona Bashari\*, Danielle Rinke\*, and Floyd Beckford, Journal of Undergraduate Chemistry Research, 2006, 2, 99
- [9] Mei-Hsiu Shih \*, Yu-Yuan Xu, Yu-Sheng Yang and Tzu-Ting Lin, *Molecules* 2015, 20, 5184-5201