



LIGANDS OF TRANSITION METAL COMPLEXES INCLUDING SCHIFF BASE: AN OVERVIEW

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ABSTRACT :

The purpose of the research and synthesis of new metal complexes for therapeutic, diagnostic and catalytic applications and other biological and chemical applications is to introduce the Coordination Chemistry Base, the Coumarine Moieties and the Quinoline Aldehyde.

KEYWORDS : metal complexes, Schiff base, The Ligands,

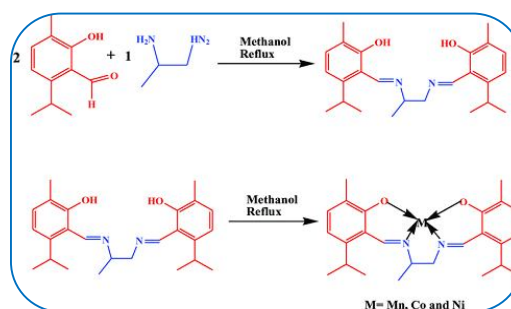
INTRODUCTION :

In 1893, Alfred Werner, considered the father of coordination chemistry in 1913, proposed the first theory on coordination chemistry. Coordination compounds are called compounds whose dissolution does not create ions, but rather complex ions. These compounds show different properties depending on the metal ion, nature of the metal and the form of ligand. During these times multiple colours were divided, later called the typical coordination compounds. the increase of coordinating chemistry goes back to the French Revolution. The work of Jorgenson and Werner began with the modern study of coordination compounds. The reason for this complex work is the formulation of the electronic metal ion structure in coordination compounds such as the theory of the crystal field (CFT), molecular orbital theory (MOT) and the theory of the Ligand Field (LFT).

APPLICATIONS OF COORDINATION COMPOUNDS

Many uses have been identified, including the treatment, management and diagnosis of diseases using medicinal products, according to a literature survey. Some metal complexes used for medicine, catalysis and so forth. For example: Examples:

- The highly efficient DNA and RNA objective 24 cleavage catalysis of amino glycoside is shown to be highly efficient.
- Ziegler-Natta catalysts are used for the low-druck polymerization of polyethylene articles, which are an aluminium and titanium complex. Even the alkaline, alkaline and noble metals have been used in treating disease, such as gold compounds for arthritis and platinum compounds for some cancers, and calcium and metal ions in the magnesium play a key role in enzyme activation, nucleic acid nerve impulsion complexation, muscle contraction, etc.

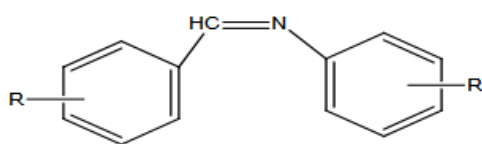
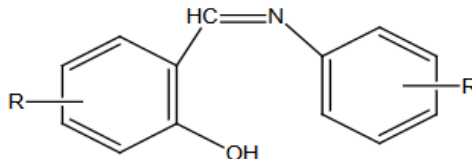
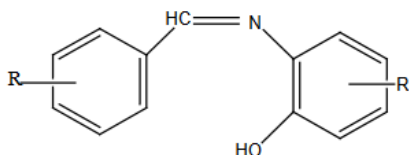
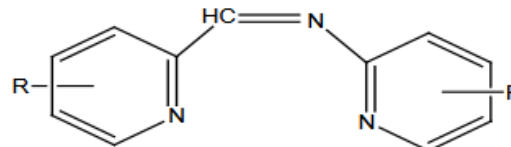
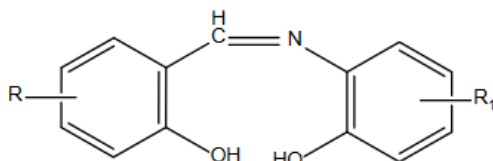


COORDINATION BEHAVIOR OF THE LIGANDS:

In the preparation of metal complexes with specific spectroscopic and magnetic properties, macrocyclic ligands have also been used. Azomethines can be called ligands with C = N structure, and these are commonly known as the Schiff's base, hydrazones, halving carbazones, thiosemicarbazones and oximes.

SCHIFF BASES

The co-ordination chemists recently focussed on Schiff's bases as a multifunctional spacer due to their preparedness and structural variant. They are condensation products of carbonyl compounds of primary amines. These compounds are named after him and contain an aryl or alkyl group with a double carbon nitrogen bond with a nitrogen atom. These compounds are called azomethine imine and. Compounds of this kind also have a variable nomenclature. These products are provided by groups, imines and Schiff's base in chemical abstracts. The Schiff Basis synthesis is based on many reaction routes. Amine and aldehyde or ketone condensation reactions in different solvents under different conditions are the most frequently induced in acid catalysed. The acid-catalyzed reaction is done by acetroping the carbonyl compound and amine. Schiff's base can, as follows by typical examples, be divided into subgroups;

**Figure 1: Neutral bidentate.****Figure 2: Monovalent bidentate****Figure 3: Neutral tridentate.****Figure 4: Monovalent tridentate.****Figure 5: Bivalent tridentate.**

In the following complexes, the neutral tridentate was isolated and chemically analysed, the octahedral complexes with transition metals were shown, where the metal ion is covered with two ligands in an octahedral range.

a) Monodentate Schiff's base

The fundamental strength of the C = N group is not enough to achieve stable complexes by coordinating the imino nitrogen atom with a metal ion. Therefore, it is necessary to preserve a metal nitrogen bond in at least one additional group. The ligands are normally stabilised by resonance by aryl group attached to either O or N. Monodentate Schiff are well known in the literature as a foundation such as N-benzylideneaniline derivative (Figure: 6).

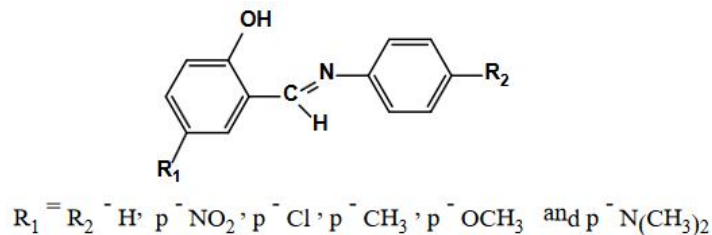


Figure: 6

b) Bidentate Schiff's base

In preparing metal complexes, the foundation of Bidentate Schiff is commonly used.

In N, O donor atom set Large bidentate Schiff base groups are used as ligands for the preparation of the N and O donor set metal complexes. The group OH also includes oxygen. The Schiff ligands of these Schiff's are typically chelated monoamines. A variety of example examples of possible bidentate Schiff's ligands with N, O donor sets are derived from 2-isopropylaniline (Figure 7), 2-Amino-3-acetophenone (Figure 8) (Figure 9).

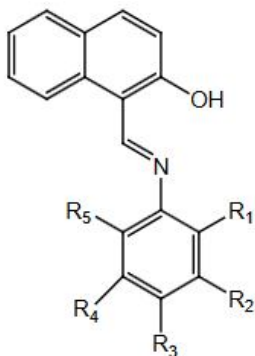


Figure: 7

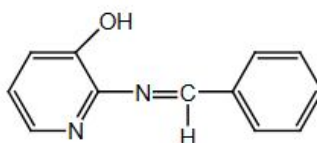


Figure: 8

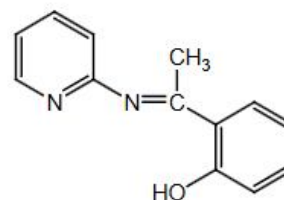


Figure: 9

Patel et al reporter have identified Mononuclear Cu(II), Ni(II), Zn(II), Fe(III), Mn(II) and Cd(II) Bidentate Schiff Base Complexes (Figure 10) characterised by elementary analysis, magnetic measurements, infrared spectra, electronic spectra and thermographic analysis

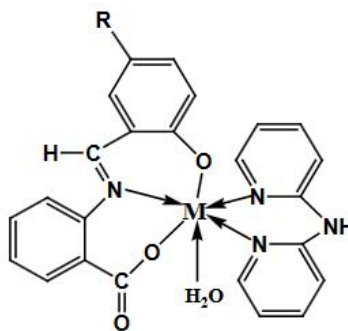


Figure: 10 R = H and Br

Naveer et al. study Co(II) and Ni(II) complexes of the bidentate ligand (figure 11) where they are coordinated in ligands in the enolized form (4-oxo-4H-1-benzopyrene 3-carboxyaldehyde-4-chloromethyl

benzyl hydrazone). The four donor atoms N2 and O2 are square foundations in this complex and two water molecules are on either side of the plane which create octahedral geometry around central metal atom.

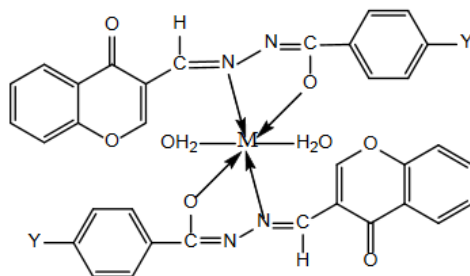


Figure : 11 M = Co(II) and Ni(II), Y= Cl and CH₃

ii) N, N donor set The base of Schiff with 2 donor-nitrogen atoms can be derived by either dialdehyde condensation or diketones with two imine molecules, or by diamine- and ketone-based reactions. Pyridine-2-carboxaldehyde with 1,3-diaminopropane ligand (Figure 12), 2-[2-(9-pyridylamino)]pyridine ligand (Figure 13), may be possible examples. Application is given in Figure 13. These ligands have been extensively studied for their structure and coordination components.

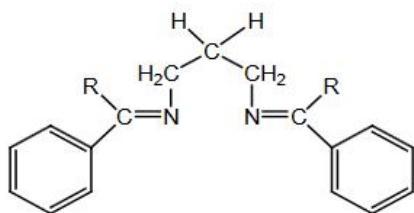


Figure : 12

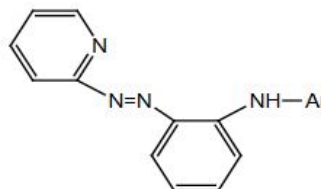
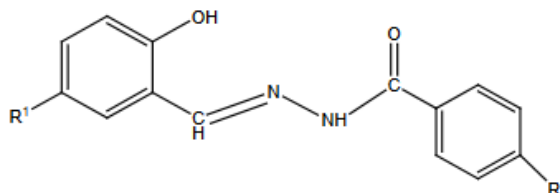


Figure: 13

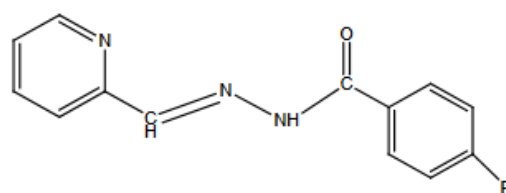
c) Tridentate Schiff's base

The tridentate Schiff's foundation, which includes donors from N2O, N2S, NO2 and NSO. The inclusion of another donor group may usually be regarded as derived from the bidentate analogous. It may be that the base ligands of such Schiff may also serve as a bridge between two metal centres, which gives a number of polynuclear complexes of the base ligands of some of the tridentate Schiff (Fig. 14), (Fig. 15) and (Fig. 16) below.



R = R¹ = H, Cl, Br, CH₃

Figure: 14



R = H, Cl, Br, CH₃

Figure: 15

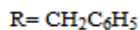
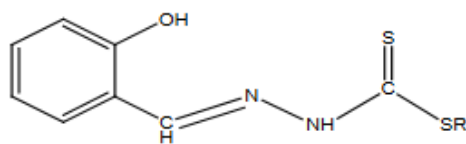


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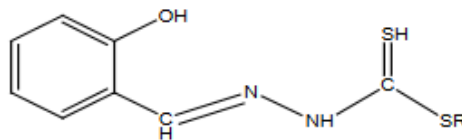


Figure: 17

d) Tetradentate Schiff's base

Tetradentate Schiff's N₂O₂ and N₂S₂ donor bases were studied for their coordination capability with metal ions. Electronic nature and conformativity of the ligands are the properties of the complexes derived from the base of their shipping ligands.³⁹Tetradentate Ethylene diamine foundation Schiff can often be categorised as I acen (Fig.18), II salen (Fig.19) and III moxen (Fig.20).

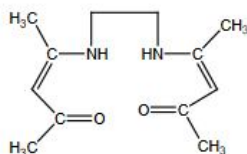


Figure: 18

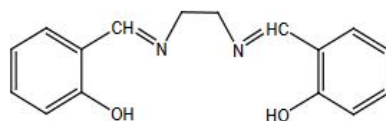


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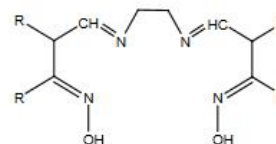


Figure:20

Because of the complicated existence of the IR spectra, Berger et al. carried out some simple structural details. Then Mossbauer tin-spectroscopy was shown to provide knowledge of this kind on organotines, and these inorganic tin adducts with Mossbauer techniques (Fig.21) and (Fig.22) were thought to be worth studying.

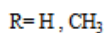
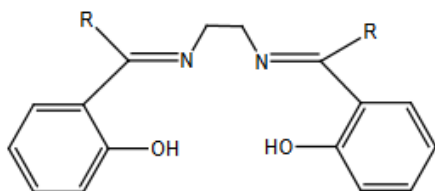


Figure: 21

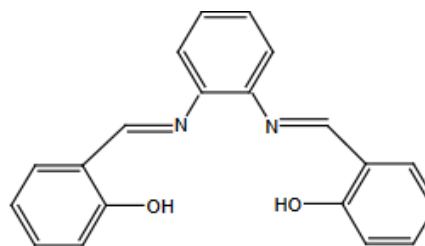


Figure: 22

P. G.Cozzi et al.,⁴² performed a Cu(II) complex (bis)(mercaptobenzylidene)diamine (Fig. 23) electron spin resonance spectral test and arrived at Cu(II) ion configuration, namely squareplanar (Fig.23). The computer simulated programmes have determined different parameters.

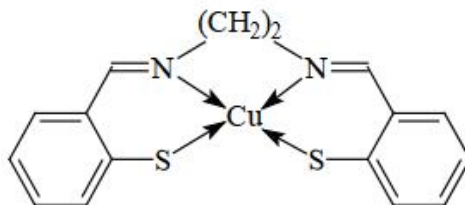


Figure: 23

As a result of expansion, transition metals form multi-coordinate complexes, especially Co(II), Ni(II) and Cu(II) complexes are formed. Such a case also happens with square planar compounds by taking the penta and hexa-co-ordinated complexes of one or two Schiff-base ligands. Different solvents are also correlated with the colour variations in some Co(II) and Ni(II) complexes, which result in complex higher coordination numbers.

In case of studying the implications of comprehensive conjugation of coordinates of boundary and structural imines, the magnetic and spectral studies show the intense conjugation of the C = N group does not promote the attack of imines, which are induced by Natarajan / Paleniandavar 2-hydroxychalconic imines with their Co(II), Ni(II) and Cu(II) complexes. The square-planar configurations of all complexes. Other forms of complexes of chalconium are also mentioned in the literature and are also considered to have square planar geometry, Cu (II) complexes with 2,2'-dihydroxychalcone imines.

Schiff bases have a broad range of uses in fruit, dye, analytical chemicals, catalyses, fungicides, agrochemical and biological activities. With the rise in the incidence of deep mycoses, the screening of new and more effective antimicrobials with low toxicity has been increasingly emphasised. Based on their ready accessibility and their structural variety, Schiff base complexes are considered as one of the most important stereochemical models in main group and transition metal coordination chemistry.

CONCLUSION:

Adipichydrazides with quinoline aldehyde, coumarinhydrazide with substituted acetophenones, and 2-amino-5-iodobenzoic acid hydrazide with quinoline aldehyde are used for the present study to synthesise the Schiff base ligands and their metal complexes.

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