



Vol. 6 | No.2 | 161-163 | April-June | 2013

ISSN: 0974-1496 | e-ISSN: 0976-0083 | CODEN: RJCABP http://www.rasayanjournal.com

http://www.rasayanjournal.co.in

# pH- METRIC STUDIES OF TERNERY COMPLEXES OF SOME TRANSITION METAL(II) IONS WITH ASPARTIC ACID AND GLUTAMIC ACID AS PRIMARY LIGANDS AND NICOTINIC ACID AND ASCOBIC ACID AS SECONDARY LIGANDS

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#### **ABSTRACT**

The interaction of Mn (II), Co (II), Ni (II), Cu (II) and Zn (II) metal ions with nicotinic acid (NA) and ascorbic acid (AA) have been studied by pH-metric technique at 0.1 M (KNO<sub>3</sub>) ionic strength at  $302 \pm 0.5 \text{ K}$  in aqueous medium. The data obtained were used to evaluate the values of proton-ligand and metal-ligand stability constants using Irving-Rossotti titration technique. Mixed ligand complex studies of these metal ions using aspartic acid (ASP) and glutamic acid (GLU) as primary ligands and NA and AA as secondary ligands also have been carried out pH-metrically at the same conditions.

**Keywords:** Ternary complexes, Transition metal ions, nicotinic acid, ascorbic acid, aspartic acid and glutamic acid.

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#### INTRODUCTION

Recently, there has been considerable interest in the study of binary, ternary and quaternary complexes by pH-metric method<sup>1-3</sup>. The ligand nicotinic acid<sup>4</sup> (NA) and ascorbic acid<sup>5, 6</sup> (AA) are well known vitamins. NA is chiefly used in the treatment of pellagra, disease due to diet deficiency and AA is necessary for the prevention and cure of the deficiency disease scurvy.

Ternary complexes of Ni (II) and Cu (II) with nicotinic acid as primary ligand and imidazoles, benzimidazole, histamine and L-histidine as secondary ligands have been studied by Nair and Neekantan<sup>7</sup>. Patil<sup>8</sup> has studied the stability constants of Mn (II), Co (II), Ni (II), Cu (II) and Zn (II) with aspartic acid and glutamic acid as primary ligands and pyridoxine hydrochloride and ethambutol hydrochloride as secondary ligands potentiometrically. Nigam and coworkers<sup>9</sup> have studied the ternary complexes of Mn (II), Co (II), Ni (II), Cu (II) and Zn (II) using ASP as primary ligand and thymine as secondary ligand potentiometrically. Ternary complexes of Cu(II) using ASP and GLU as primary ligands have been reported potentiometrically by Pandeya and Patel<sup>10</sup>.Kalshetti and coworkers<sup>11</sup> have studies binary and ternary complexes of Co (II), Ni (II), Cu (II) and Zn(II). Mixed ligand complexes of transition metal(II) ions with N-(2-hydroxybenzylidene)-2,3-dimethylaniline as primary ligand and N-(2-hydroxy-1-naphthylidene)-4-chloroaniline as secondary ligand has been studied by Mapari and Mangaonkar<sup>12</sup>. Solution equilibria of ternary systems involving transition metal ions, hydroxamic acids, and bioligands have been studied by Khalil and Mahmoud<sup>13</sup>. Gazala Mohamed<sup>14</sup> studied the ternary complexes of Co (II) involving nitrilotriacetic acid and some biologically active ligands.

In this paper the stability constants of ternary complexes of Mn (II), Co (II), Ni (II), Cu (II) and Zn (II) ions with ASP and GLU as primary ligands and AMP and CEP as secondary ligands at  $302 \pm 0.5$  K and at fixed ionic strength,  $\mu = 0.1$ M KNO<sub>3</sub> using modified form of Irving-Rossotti pH-metric technique<sup>15</sup> in aqueous medium have been studied.

#### **EXPERIMENTAL**

The ligand NA and AA are obtained from LOBA Chemie. These ligands were used as such. Carbonate free sodium hydroxide solution was prepared by standard method<sup>16</sup>. All other solutions were prepared in doubly distilled water.

The pH-metric measurements were carried out by using Elico digital pH-meter model L-120 with combined glass-calomel electrode with an accuracy of  $\pm$  0.01 of pH unit at 302  $\pm$  0.5 K. The pH-meter was standardized against 0.05 M potassium hydrogen phthalate solution in acid medium and 0.01M borax solution in alkaline medium.

For determination of proton-ligand stability constant of the secondary ligand and the metal-ligand stability constants of ternary complexes, the following set of solutions were prepared and titrated against standard alkali solution.

#### **Ternary Systems**

- i. 9.6 X 10<sup>-3</sup>M HNO<sub>3</sub>
- ii.  $9.6 \times 10^{-3} \text{M HNO}_3 + 1.0 \times 10^{-3} \text{M secondary ligand}$
- iii.  $9.6 \times 10^{-3} \text{M HNO}_3 + 1.0 \times 10^{-3} \text{M primary ligand} + 1.0 \times 10^{-3} \text{M metal ion}$
- iv.  $9.6 \times 10^{-3} \text{M HNO}_3 + 1.0 \times 10^{-3} \text{M primary ligand} + 1.0 \times 10^{-3} \text{M metal ion} + 1.0 \times 10^{-3} \text{M}$  secondary ligand

The ionic strength was maintained constant (0.1M) by adding required volume of 1M KNO<sub>3</sub>. The ratio of metal (M): primary ligand (A): secondary ligand (L) was maintained at 1: 1: 1 in each of the ternary system.

#### RESULTS AND DISCUSSION

## **Proton-Ligand Stability Constants**

The plots of volume of alkali (NaOH) against pH-meter readings were used to evaluate the proton-ligand stability constants of NA and AA. The deviation between free acid titration curve and secondary ligand titration curve was used to evaluate the formation functions  $\overline{\eta_A}$ . The proton-ligand formation curves were then obtained by plotting the values of  $\overline{\eta_A}$  versus pH-meter readings. From the graphs, the values of

 $logK^H$  were evaluated by half integral method (method A) and point wise calculation method (method B) and presented in Table-1.

Table-1: Proton-Ligand Stability Constants of NA and AA

Ligand	logK <sub>1</sub> <sup>H</sup>		log	$K_2^H$	logβ <sup>H</sup>		
	Method(A)	Method(B)	Method(A)	Method(B)	Method(A)	Method(B)	
NA	4.73	4.74	-	-	4.73	4.74	
AA	11.45	11.50	4.10	4.12	15.55	15.62	

The formation curves for the proton-ligand system of NA extended between 0 to 1 in the  $\overline{\mathbf{I}_{\mathbf{A}}}$  scale at fixed ionic strength and temperature indicating that the dissociation of HNA as follows:

$$HNA \rightleftharpoons NA^- + H^+$$

The formation curves for the proton-ligand system of AA extended between 0 to 2 in the scale at fixed ionic strength and temperature indicating that the dissociation of HNA as follows:

$$H_2AA \rightleftharpoons HAA^- + H^+ \rightleftharpoons AA^{2-} + H^+$$

#### **Metal-Ligand Stability Constants of Ternary Complexes**

The metal-ligand stability constants of the ternary complexes were evaluated assuming that the formation of hydrolyzed products, polynuclear complexes, hydrogen and hydrogen bearing complexes were absent. An examination of titration curves indicates that complex formation has taken place in the solution on the following grounds:

The ternary complex titration curves show the displacement with primary complex titration curves.
 The horizontal distance was measured between acid curve and the secondary ligand curve (V<sub>2</sub> - V<sub>1</sub>) and subtracted through the horizontal distance between ternary complex curves and primary complex titration curves (V<sub>4</sub> - V<sub>3</sub>) show the positive difference which proves the earlier released of protons in the formation of ternary complexes.

2. The hydrolysis of the metal ions was suppressed and precipitation did not result.

The values of  $\overline{\eta}$  vary from 0 to 1.0, thus confirming the formation of 1:1:1 mixed ligand complexes. The values of  $\log K_{MAL}^{ASP}$  and  $\log K_{MAL}^{GLU}$  have been evaluated from the formation curves ( $\overline{\eta}$  vs. pL). At  $\overline{\eta} = 0.5$  in the formation curve, pL =  $\log$  K. The metal-ligand stability constant of NA and AAas secondary ligands and ASP and GLU as primary ligands are presented in Table-2.

System	$\log K_{MAL}^{MA}$									
	Mn (II)		Co (II)		Ni (II)		Cu (II)		Zn (II)	
	A	В	A	В	Α	В	A	В	A	В
[M (II) - (ASP) (NA)]	4.52	4.51	4.59	4.58	4.77	4.75	4.90	4.91	4.55	4.54
[M (II) - (GLU) (AA]	4.53	4.53	4.57	4.56	4.66	4.65	4.69	4.68	4.63	4.62
[M (II) - (ASP) (NA)]	5.87	5.90	7.26	7.27	7.44	7.45	8.28	8.28	7.32	7.36
[M (II) - (GLU) (AA)]	5.32	5.33	6.39	6.38	7.40	7.42	8.28	8.27	6.67	6.68

Table-2: Metal - Ligand Stability Constants of Ternary Complexes

The Irving-Williams order  $^{17, 18}$  of stability constants was observed in ternary complexes which is Mn (II) < Co (II) < Ni (II) < Cu (II) > Zn (II)

This sequence of stability of complexes with respect to metal ion is due to deceasing atomic radius and increasing the second ionization potential.

#### **ACKNOWLEDGEMENTS**

The authors are thankful to Principal, P. N, College, Pusad for providing necessary facilities.

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[RJC-1038/2013]