Synthesis and Characterization of Ternary (Tri- Trivalent) Polymetaphosphate Derivatives with Metachromatic studies

Vishnu Kumar Khandelwal*

Department of Chemistry, JECRC University, Jaipur (Rajasthan)-India

ABSTRACT:

Ternary (tri-trivalent) polymetaphosphate derivatives having the composition $[Na_{x}M_{1-x/2a}^{III}PO_{3}]_{n}$ (where $M^{III} \& M'^{III} = Fe(III)$, La(III), Sm(III), Gd(III), x = 2/3, 3/4, a = Valency of metal ion) have been synthesized by fusion technique. Metachromatic studies of ternary polymetaphosphate derivatives are carried out with some thiazene dyes like methylene blue and toludine blue. These metachromatic studies of ternary polymetaphosphate derivatives confirmed their long chain polymeric character

Key words: Polymetaphosphate, Fusion technique, Metachromatic studies, Methylene Blue, Toludine Blue.

INTRODUCTION:

The condensed phosphates form a very important class of phosphorous compounds which exhibit marked stability of polymeric structure in solution. Structurally, condensed phosphates include all those compounds of P-O-P linkage formed by the sharing of PO₄ tetrahedra, in cyclic chain or cross-linked structures. Specifically the polyphosphates consist of PO₄ tetrahedra sharing two of their oxygen atoms with adjoining PO₄ tetrahedra forming long chain polyanions of the formulation $(PO_3)_n$ ⁻ⁿ. These polyanionic chains exhibit pronounced capacity for sequestration of metal ions.

The chief representative of the condense phosphates is Graham's salt, which was prepared by Thomas Graham¹ by thermal dehydration of NaH₂PO₄.2H₂O. Fleitmann and Henneberg² reported that this product was Hexametaphosphate. Another form of difficultly soluble sodium metaphosphate was obtained by tempering the super-cooled melt of [NaPO₃]_n at 550°C. Tammann³ named this derivative as Kurrol's salt. The sequestering ability of Graham's salt towards metal ions was reported by Hall⁴, since then it is used for softening of water for boilers and given trade name 'Calgon'. Wall and Doremus⁵ reported that the exchange of a part of sodium ion in [NaPO₃]_n with strontium, does not affect the properties of the former. Mehrotra and Gupta⁶ also indicated that if a part of the alkali metal in its phosphates is substituted by bivalent metal ions, the derivative so obtained should have properties similar to that of Graham's salt. This assumption was confirmed by the preparation of soluble complex polymetaphosphate derivatives of the composition $[M_x^I M^{II}_{1-x/a}]$ $PO_3]_n$ (where $M^I = Li(I)$, K(I), Cs(I), $M^{II} = Mg(II)$, Ca(II), Ba(II), Zn(II), Cu(II), Ni(II), Pb(II), a = Mg(II), Ca(II), Ca(II), Cu(II), Ni(II), Nivalency of metal ion, x = 2/3, 1/3, 1/2). Mehrotra and Vyas later reported soluble complex derivatives of the composition $[K_x M^{II}_{1-x/a} PO_3]_n$ (where $M^{II} = Mg(II)$, Ba(II), Ca(II), Zn(II), Cu(II), Ni(II),Pb(II), a=valency of metal ion, x=2/3,1/3,1/2). Similarly Mehrotra and Oza⁷ reported soluble complex of derivatives of lithium and cesium polymetaphosphate. All these derivatives are found to be polymeric in nature on the basis of various physico-chemical studies used for characterization of Graham's salt. The complex polymetaphosphates of the composition $[M_x^{I}M^{III}_{1-x/a}PO_3]_n$ where M^{III} = Pr(III), Nd(III) and x = 2/3, 1/2, 1/3, a = valency of metal ion) were synthesized and their polyelectrolytic behaviour and spectral characteristics were studied by Mamta Oza⁸. A few complex bimetallic and trimetallic polymetaphosphate derivatives containing alkaline earth and rare earth

metals have been synthesized by fusion techniques by Shobhna Sharma⁹ and characterized by various physico- chemical techniques in these laboratories.

In the present communication, the polymeric nature of ternary complex polymetaphosphate derivatives of the composition: $[Na_xM^{III}_{1-x/2a}M'^{III}_{1-x/2a}PO_3]_n$ (where $M^{III} \& M'^{III} = Fe(III)$, La(III), Sm(III), Gd(III), x = 2/3, 3/4, a = Valency of metal ion) have been synthesized by fusion technique. Metachromatic studies of ternary polymetaphosphate derivatives are carried out with some thiazene dyes like methylene blue and toludine blue. These metachromatic studies of ternary polymetaphosphate derivatives confirmed their long chain polymeric character.

MATERIAL AND METHODS:

All the reagents such as NaH_2PO_4 , $NH_4H_2PO_4$ and metal oxides used were of analytical grade. Double distilled water used for preparing the solutions. An Ostwald viscometer having efflux time between 80-120 sec. for water was used for viscosity measurements and a Pycknometer of volume about 10 ml was used for density measurements. A digital pH- meter having glass and a calomel electrode was used for pH measurements. Viscosity measurements were performed at $30^{\circ}\pm0.1^{\circ}C$. All the measurements were carried out after 24 hours of dissolution so as to attain equilibrium.

PREPARATION:

Preparation of complex polymetaphosphate derivatives of the composition $[Na_{x}M^{III}_{1-x/2a}M'^{III}_{1-x/2a}PO_{3}]_{n}$ (where $M^{III} \& M'^{III} = Fe(III)$, La(III), Sm(III), Gd(III), x = 2/3, 3/4, a = valency of metal ion)

When x=2/3

	$900 \pm 25^{\circ}\mathrm{C}$		
n 2/3 NaH ₂ PO ₄ + n 1/18 M ^{II} O + n 1/18 M' ^{II} O + 1/3 n NH ₄ H ₂ PO ₄			
$[Na_{2/3} \ M^{III}_{1/18} \ M'^{III}_{1/18} \ PO_3]_n + 1/3 \ n \ NH_3 + 7/8 \ n \ H_2O$	Δ	(1)	
<u>When x=3/4</u>			
	$900\pm25^{o}C$		
n 3/4 NaH ₂ PO ₄ + n 1/24 M ^{III} O + n 1/24 M ^{III} O + 1/4 n NH ₄ H ₂ PO ₄			
$[Na_{3/4} M^{III}_{1/24} M'^{III}_{1/24} PO_3]_n + 1/4 n NH_3 + 9/8 n H_2O$	Δ	((2)

Complex polymetaphosphate derivatives of the composition $[Na_xM^{III}_{1-x/2a}M'^{III}_{1-x/2a}PO_3]_n$ (where M^{III} & $M'^{III} = Fe(III)$, La(III), Sm(III), Gd(III), x = 2/3, 3/4, a = valency of metal ion) were prepared by fusion technique in accordance with the equation (1) & (2).

The reactant were taken in a platinum crucible, heated over a flame for 40-45 minutes and then put in a muffle furnace at $800 \pm 25^{\circ}$ C for one hour. A clear melt was obtained in almost all the cases which was then chilled between cold stainless steel plates to give a clear glass. The colour of the glass was green in the case of copper derivatives whereas in all the other cases it was colourless. The yield was almost theoretical and the composition was confirmed by estimating phosphorous and metal contents of the compounds.

Phosphorous was estimated volumetrically as ammonium molybdate and metals by standard methods as described by Vogel¹⁰ given in table (1).

Sr.	Complex	%	b P	% of	f M ^I	% of	[°] M ^{III}	%	of M' ^{III}
No	Polymetaphosphates								
		Exp	Cal.	Exp	Cal.	Exp	Cal.	Exp	Cal.
1	$[Na_{2/3} Fe_{1/36}La_{1/12}PO_3]_n$	28.73	28.85	14.20	14.26	1.38	1.44	10.65	10.77
2	$[Na_{3/4} Fe_{1/24}La_{1/24}PO_3]_n$	29.63	29.70	16.48	16.52	2.18	2.22	5.50	5.54
3	$[Na_{3/4} Sm_{1/24}Fe_{1/24}PO_3]_n$	29.44	29.56	16.42	16.44	5.90	5.97	2.16	2.21
4	$[Na_{3/4} Gd_{1/24}Fe_{1/24}PO_3]_n$	29.30	29.49	16.35	16.40	6.20	6.23	2.18	2.21
5	$[Na_{3/4} La_{1/24}Sm_{1/24}PO_3]_n$	28.47	28.62	15.86	15.92	5.30	5.34	5.72	5.78
6	$[Na_{3/4} La_{1/24}Gd_{1/24}PO_3]_n$	28.46	28.55	15.81	15.87	5.28	5.33	6.00	6.03

Table: 1 Analysis of complex polymetaphosphate derivatives:

The complex polymetaphosphates synthesized as above were analyzed for their constituents i.e. Phosphorus, Alkali Metals, Trivalent Metals for the confirmation of their compositions.

METACHROMATIC STUDIES

INTRODUCTION: Metachromatic reactions of ternary (bi-bivalent) complex polymetaphosphate derivatives with methylene blue (MB) and toludine blue (TB) have been studied. It has been found that the absorption maximum observed at 630 nm and 665 nm for TB and MB respectively are shifted to 530 nm and 580 nm in presence of complex polymetaphosphate derivatives. These shifts are similar to those reported in case of these dye solution in presence of Graham's salt as well as complex sodium and potassium polymetaphosphate derivatives. The metachromatic behaviour of complex polymetaphosphate derivatives is independent of their molecular weight and is exhibitive of their polyelectrolytic character.

Metachromacy is the variability of the colour of the dyes, produced by the addition of certain high molecular weight polyelectrolyte. The colour of the dyes produced in presence of polyelectrolyte is known as metachromatic colour. Such dyes are also known to disobey Beer's law deviation from which have been attributed to reversible polymerizations of dye molecules i.e. the polymer exhibits a colour different from the monomer.

Michaelis and Granick¹¹ have studied metachromacy of some thiazine dyes spectrophotometrically. The absorption maxima of the dye, according to these workers are shifted to lower wavelengths.Wiam¹² studied the metachromatic behaviour of sodium hexametaphosphate with toludine blue. It was observed that absorption maxima of aqueous solution of the dye are shifted to lower value i.e. 530 nm in presence of Graham's salt. Phosphates such as ortho-, pyro -, and tripoly-, other than polymetaphosphate derivatives do not produce metachromatic effect. Anne-Levine and Schubert¹³ have also studied the metachromatic reaction of TB in presence of polyanion and suggested that the loose binding of the former with later is cause of metachromacy. Mckay and Hillson¹⁴ studied the effect of different electrolytes in varying concentrations on the metachromatic behaviour of methylene blue. Mehrotra and Gupta¹⁵ and later Mehrotra, Vyas and Ojha¹⁶ have studied the metachromatic effect of complex polymetaphosphate derivatives of sodium, potassium, lithium and cesium on TB, MB and thionine dyes. Metachromacy produced by these derivatives was found to be similar to that of Graham's salt and indicative of their polyelectrolytic character.

These studies were used by these workers for elucidating the polymeric nature of the corresponding complex polymetaphosphates. The polymeric nature of $(BHPO_3)_n$ and complex polyphosphate of the composition $[BH_x M^{II}_{1-x/a}PO_3]$ was established by $Vyas^{17}$ and Mamta¹⁸ by studying the metachromatic reactions of these derivatives towards cationic dyes. The above study was also considered useful for elucidating the long chain polymeric character of these complex polymetaphosphate derivatives of the composition:

Experimental:

Materials: Dyes: Methylene blue and Toludine blue have been employed for studying the metachromatic reactions. Both the dyes used in the study were of E. Merck grade. The dye samples were recrystallized from alcohol before use.

Polyelectrolyte: Polymetaphosphate of the composition $[Na_xM^{III}_{1-x/2a}M'^{III}_{1-x/2a}PO_3]_n$ (where $M^{III} \& M'^{III} = Fe(III)$, La(III), Sm(III), Gd(III), x = 2/3, 3/4, a = valency of metal ion) was used for these studies.

Solvent: Water was used as a solvent for the preparation of dye and polyelectrolyte solutions.

Method: The stock solutions were prepared by dissolving the weighed samples in double distilled water. The concentration of polyphosphate derivatives was taken in the range $0.2-1.0X10^{-3}$ M. The stock solutions of dyes were also prepared in similar manner. The final concentration of the dye in the dye – polymer solution was maintained at a fixed value which is 1.6×10^{-4} M for TB and 3.2×10^{-5} M for MB. A lower concentration of MB was preferred because at higher concentration the particles of MB settled down on keeping the solution for some time after the addition of complex polymetaphosphate derivatives.

Measurements: The optical densities of the dyes (MB or TB) alone and in presence of polyphosphate were measured with the help of **UV- Visible Spectrophotometer (Model - 301E)**. The conductometric titrations of dyes were carried out by employing digital conductivity bridge (Model – CC 601) and a conductivity cell having cell factor 0.90. The titrations were performed by drop wise addition of 2 X10⁻² M solution of polymer to the 25 ml solution of 1.0 X 10⁻³ M dye in a 50 ml beaker. The solution mixtures were stirred vigorously and allowed to stand for some time to attain equilibrium before measuring the conductance.

Result and Discussions

The absorption spectra of the aqueous solution of MB and TB were recorded in the absence and presence of complex polymetaphosphate derivative of the composition $[Na_x M^{III}_{1-x/2a} M'^{III}_{1-x/2a} PO_3]_n$ (where $M^{III} \& M'^{III} = Fe(III)$, La(III), Sm(III), Gd(III), x = 2/3, 3/4, a = valency of metal ion). There representative spectra are given in figure 1 to 4. The position of the main band (α) and metachromatic band (β) observed in the case of dye solution alone and in the presence of polyphosphate derivatives recorded in table 1 to 4.

Sr. No.	Wave Length (λ) (nm)	Absorbance
1	400	0.03
2	420	0.04
3	440	0.05
4	460	0.07
5	480	0.08
6	500	0.10
7	520	0.11
8	540	0.14
9	560	0.25
10	580	0.44
11	600	0.68
12	620	1.0
13	640	1.26
14	660	1.53
15	665	1.61
16	670	1.51
17	680	1.09
18	700	0.36
19	720	0.21
20	740	0.16
21	760	0.14
22	780	0.12
23	800	0.10

Table 1: Absorbance of the aqueous solution of Methylene Blue $(3.2 \times 10^{-5} \text{ M})$

Sr. No.	Wave Length (λ) (nm)	Absorbance
1	400	0.06
2	420	0.07
3	440	0.09
4	460	0.11
5	480	0.13
6	500	0.17
7	520	0.21
8	540	0.33
9	560	0.54
10	580	0.77
11	600	1.11
12	620	1.51
13	630	1.65
14	640	1.46
15	660	1.16
16	680	0.92
17	700	0.71
18	720	0.53
19	740	0.42
20	760	0.31
21	780	0.26
22	800	0.22

Table 2: Absorbance of the aqueous solution of Toludine Blue (1.6 X 10^{-4} M)

S.	COMPLEX		CONCENTRATION								
No.	POLYMETAPHOSPHATE	0.2 X 1	$0.2 \times 10^{-3} M$ $0.4 \times 10^{-3} M$		0.6 X 10 ⁻³ M		0.8 X 10 ⁻³ M		1.0 X 10 ⁻³ M		
		А	В	А	В	α	В	А	В	А	В
1	$[Na_{2/3} Fe_{1/36}La_{1/12}PO_3]_n$	0.270	0.665	0.276	0.658	0.220	0.634	0.265	0.642	0.258	0.652
2	$[Na_{3/4} Fe_{1/24}La_{1/24}PO_3]_n$	0.302	0.643	0.298	0.656	0.266	0.609	0.275	0.634	0.292	0.657
3	$[Na_{3/4} \ Sm_{1/24} Fe_{1/24} PO_3]_n$	0.294	0.610	0.276	0.625	0.285	0.636	0.280	0.644	0.302	0.650
4	$[Na_{3/4}\ Gd_{1/24}Fe_{1/24}PO_3]_n$	0.298	0.610	0.282	0.625	0.276	0.632	0.294	0.642	0.302	0.653
5	$[Na_{3/4}\ La_{1/24}Sm_{1/24}PO_3]_n$	0.302	0.612	0.282	0.629	0.274	0.634	0.285	0.642	0.296	0.649
6	$[Na_{3/4} \ La_{1/24} Gd_{1/24} PO_3]_n$	0.295	0.678	0.302	0.688	0.278	0.649	0.310	0.683	0.316	0.693

Table 3: Absorbance of main band (α) and metachromatic band (β) of the aqueous solution of methylene blue (3.2 X 10⁻⁵ M) in the presence of complex polymetaphosphatederivatives

Table 4: Absorbance of main band (α) and metachromatic band (β) of the aqueous solution of toludine blue (1.6 X 10⁻⁵ M) in the presence of complex polymetaphosphatederivatives

	COMPLEX		CONCENTRATION								
S.	POLYMETAPHOSPHATE	0.2 X 1	0 ⁻³ M	0.4 X 1	0 ⁻³ M	0.6 X	10 ⁻³ M	0.8 X 1	0 ⁻³ M	1.0 X	10 ⁻³ M
No.		А	B	А	В	α	В	А	B	Α	В
1	$[Na_{2/3} Fe_{1/36}La_{1/12}PO_3]_n$	0.648	1.184	0.655	1.168	0.624	1.156	0.682	1.174	0.676	1.145
2	$[Na_{3/4} Fe_{1/24}La_{1/24}PO_3]_n$	0.630	1.128	0.654	1.122	0.618	1.166	0.675	1.175	0.662	1.148
3	$[Na_{3/4} Sm_{1/24}Fe_{1/24}PO_3]_n$	0.624	1.125	0.635	1.114	0.650	1.119	0.672	1.167	0.646	1.178
4	$[Na_{3/4} Gd_{1/24}Fe_{1/24}PO_3]_n$	0.628	1.124	0.634	1.127	0.615	1.112	0.657	1.146	0.662	1.151
5	[Na _{3/4} La _{1/24} Sm _{1/24} PO ₃] _n	0.616	1.121	0.632	1.123	0.638	1.119	0.665	1.148	0.646	1.151
6	$[Na_{3/4} La_{1/24} Gd_{1/24} PO_3]_n$	0.725	1.114	0.710	1.131	0.672	1.106	0.646	1.098	0.654	1.143

It can be seen from the figures and data's that the main band (α) observed at 630 & 665nm for TB and MB respectively are shifted to 530 and 580nm in presence of complex polymetaphosphate derivatives. These shifts are similar to those reported in case of these dye solution in presence of Graham's salt as well as for complex sodium and potassium phosphate derivatives. It confirms that complex polymetaphosphate derivatives possess polymeric character similar to Graham's salt. It is also evident

from these studies that polyanion induced metachromacy of these dyes is very much dependent on the concentration of polyphosphate.



Figure 1: Absorption spectra of aqueous solution of Methylene blue (3.2×10^{-5}) aloneand in presence of the varying concentration of complex polymetaphosphate $[Na_{2/3} Fe_{1/36}La_{1/12}PO_3]_n$



Figure 2: Absorption spectra of aqueous solution of Toludine Blue (1.6X10⁻⁴ M) aloneand in presence of the varying concentration of complex polymetaphosphate [Na_{2/3} Fe_{1/36}La_{1/12}PO₃]_n

The term metachromacy (R) can be defined as follows:

R = Absorbance at metachromatic band (β) /Absorbance at main band (α)

The metachromatic values for these dyes at varying concentration of polymetaphosphates have been recorded in table 5 to 6

Sr.	Complex	Metachromacy					
No.	Polymetaphosphate	P/D Ratio					
		6.25	12.50	18.75	25.00	31.25	
1	$[Na_{2/3} Fe_{1/36} La_{1/12} PO_3]_n$	2.46	2.38	2.88	2.42	2.52	
2	$[Na_{3/4} Fe_{1/24}La_{1/24}PO_3]_n$	2.12	2.20	2.28	2.30	2.25	
3	$[Na_{3/4} Sm_{1/24}Fe_{1/24}PO_3]_n$	2.07	2.26	2.23	2.30	2.15	
4	$[Na_{3/4} Gd_{1/24}Fe_{1/24}PO_3]_n$	2.04	2.21	2.28	2.18	2.16	
5	$[Na_{3/4} La_{1/24} Sm_{1/24} PO_3]_n$	2.02	2.23	2.31	2.25	2.19	
6	$[Na_{3/4} La_{1/24}Gd_{1/24}PO_3]_n$	2.29	2.27	2.33	2.20	2.19	

Table 5: Metachromacy solution of of aqueous Methylene blue (3.2X10-5 M) at different P/Dratio.

Sr.	Complex	Metachromacy					
	Porymetaphosphate	P/D Ratio					
		6.25	12.50	18.75	25.00	31.25	
1	$[Na_{2/3} Fe_{1/36} La_{1/12} PO_3]_n$	1.82	1.78	1.85	1.72	1.69	
2	$[Na_{3/4} Fe_{1/24} La_{1/24} PO_3]_n$	1.79	1.71	1.88	1.74	1.73	
3	$[Na_{3/4} Sm_{1/24}Fe_{1/24}PO_3]_n$	1.80	1.75	1.72	1.73	1.82	
4	$[Na_{3/4} Gd_{1/24}Fe_{1/24}PO_3]_n$	1.78	1.77	1.80	1.74	1.73	
5	$[Na_{3/4} La_{1/24}Sm_{1/24}PO_3]_n$	1.81	1.77	1.75	1.72	1.78	
6	$[Na_{3/4} La_{1/24}Gd_{1/24}PO_3]_n$	1.53	1.59	1.64	1.69	1.74	

Table 6: Metachromacy solution of of aqueous Toludine blue (1.6X10⁻⁴ M) at different P/D ratio.

From these data following conclusion can be drawn:

The values of metachromacy (R) for these dyes are independent of the molecular weight or chain length of the complex polymetaphosphates derivatives. Maximum metachromatic values (R) are obtained at P/D ratio equal to 12.5 and 25.00 for MB and TB respectively.

Krishnan and Coworkers¹⁹ have also found that metachromacy of TB is independent of the molecular weight of sodium polymetaphosphates. This type of behavior was further confirmed by Mehrotra and Gupta²⁰. A P/D ratio equal to 5 is observed for TB in presence of polyphosphates. Although Wiame has determined a P/D ratio equal to 8 for solution of TB in presence of sodium hexametaphosphate. This difference occurs because of difference in experimental conditions. For MB a P/D ratio equal to 12.5 is observed in presence of polyphosphates. However a critical range of P/D ratio, necessary for inducing metachromacy in MB has not been reported so far.

Stiochiometry of dye – polymer complex:

The dye ion association with polyanion was investigated conductometrically¹²⁻¹⁶. Figure 3 to 4 show the conductometric titration curves drawn on the basis of specific conductance data given in the conductometric titration curves at 1:1 molar ratio of dye and polymer in solution given in table 7 and table 8. This indicate that MB and TB can site-bound to a complex polyphosphate at 1:1 molar ratio irrespective of the chain length with phosphate residue of the polymer. Similar results were obtained by Yamaoka et al²¹ for a reaction between crystal violet and Graham's salt on the basis of conductometric studies. Further Pal et a²²⁻²³ established that polyanions such as ATP, HgCl₂, KI etc. form a complex with methylene blue.

Volume of	Specific conductance (k X10 ⁻⁴ M) in Ω^{-1} cm ⁻¹								
polymer	Ι	II	III	IV	V	VI			
solution									
(ml)									
0.00	.129	.138	.162	.158	.173	.145			
0.20	.131	.140	.165	.162	.176	.150			
0.40	.133	.142	.168	.164	.178	.154			
0.60	.135	.144	.170	.166	.182	.156			
0.80	.137	.146	.176	.170	.186	.158			
1.00	.140	.150	.182	.174	.190	.162			
1.20	.146	.154	.187	.177	.197	.167			
1.40	.153	.162	.190	.185	.204	.172			
1.60	.166	.175	.196	.192	.208	.184			
1.80	.187	.192	.224	.212	.232	.200			
2.00	.206	.215	.247	.236	.257	.226			
2.20	.233	.240	.272	.262	.284	.252			
2.40	.258	.266	.290	.287	.303	.276			
2.60	.276	.288	.318	.306	.326	.297			
2.80	.302	.310	.342	.331	.352	.322			
3.00	.333	.332	.364	.354	.376	.343			

	Table 7: Spec	ific conducta	10-4) nce (k X 10-4)	of 25 ml aque	eous soluti	on of	
methyleneblu	e (1.0 X 10 ⁻⁴ M)) in presence (of varying co	ncentration of	f complex]	polymetaj	phosphate

 $I = [Na_{2/3} Fe_{1/36}La_{1/12}PO_3]_n$ II= $[Na_{3/4} Fe_{1/24}La_{1/24}PO_3]_n$
$$\begin{split} &IV = [Na_{3/4} \ Gd_{1/24}Fe_{1/24}PO_3]_n \ V = [Na_{3/4} \ La_{1/24}Sm_{1/24}PO_3]_n \ VI = [Na_{3/4} \ La_{1/24}Gd_{1/24}PO_3]_n \\ &C = Concentration of polyphosphate solution (2 X 10^{-3} F) \end{split}$$

III= $[Na_{3/4} Sm_{1/24}Fe_{1/24}PO_3]_n$

Volume of	Specific conductance (k X10 ⁻⁴ M) in Ω^{-1} cm ⁻¹								
solution(ml)	Ι	II	III	IV	V	VI			
0.00	.410	.392	.380	.431	.361	.450			
0.20	.413	.394	.382	.433	.363	.452			
0.40	.417	.396	.384	.435	.365	.454			
0.60	.419	.398	.386	.438	.367	.456			
0.80	.421	.403	.389	.443	.369	.458			
1.00	.423	.405	.393	.445	.372	.463			
1.20	.425	.407	.395	.447	.374	.467			
1.40	.428	.409	.397	.454	.376	.475			
1.60	.436	.415	.404	.458	.379	.486			
1.80	.444	.427	.406	.466	.383	.493			
2.00	.462	.443	.425	.485	.395	.505			
2.20	.507	.486	.448	.526	.416	.548			
2.40	.544	.522	.483	.567	.448	.586			
2.60	.573	.564	.537	.595	.505	.628			
2.80	.615	.596	.578	.638	.549	.654			
3.00	.658	.634	.615	.676	.586	.692			

Table 8: Specific conductance (k X 10-4) of 25 ml aqueous solution of toludine blue (1.0 X 10-4M)in presence of varying concentration of complex polymetaphosphate

 $\begin{array}{ll} I = [Na_{2/3} \ Fe_{1/36} La_{1/12} PO_3]_n & II = [Na_{3/4} \ Fe_{1/24} La_{1/24} PO_3]_n & III = [Na_{3/4} \ Sm_{1/24} Fe_{1/24} PO_3]_n \\ IV = [Na_{3/4} \ Gd_{1/24} Fe_{1/24} PO_3]_n & V = [Na_{3/4} \ La_{1/24} Sm_{1/24} PO_3]_n & VI = [Na_{3/4} \ La_{1/24} Gd_{1/24} PO_3]_n \\ C = Concentration of polyphosphate solution (2 \ X \ 10^{-3} \ F) & VI = [Na_{3/4} \ La_{1/24} Gd_{1/24} PO_3]_n \\ \end{array}$







Figure 4: Specific Conductance (k X 10⁻⁴) of 25ml of aqueous solution of TB (1 X 10⁻⁴) in presence of varying concentration of complex polymetaphosphates

The observation of inflection in the conductometric titration curves of dye andpolymer at 1:1 molar ratio irrespective of chain length of polyphosphates, indicate that metachromacy of dye polymer is enhanced due to their site binding with polymer chains of polyelectrolytes like complex polyphosphate derivatives. Scheibe²⁴ and Robinowitch²⁵ attributed deviations (even at very low concentration) from Beer's law in the Spectrophotometric studies of aqueous solution of MB and TO to the formation of dimeric cations (MB)⁺² and (TO)⁺² respectively.

The metachromatic behaviour of thiazine dyes in presence of agar – agar has been attributed to the increased polymerization of dye molecules by Michaelis and Granick¹¹. According to Walton and Rickett²⁶ in the case of sulfonated dextranes, the reactive groups on dextrane are responsible for the metachromatic behaviour, which can be attributed to polymerization of the dye molecules. However Levine and Schubert have explained the metachromatic behaviour of dye molecule is due to loose site bonding with the polyanions.

In view of these observations the metachromatic behaviour of MB and TB in presence of polyphosphate derivatives can be attributed to increased polymerizations and counter ion associations with dye cations. The association of dye ions with polyphosphate chain can be explained by considering an ion – exchange reaction stated below:

$$Y(D)_{2}^{++} + [Na_{x} M^{I}_{1-x/2a} M^{II}_{1-x/2a} PO_{3}]_{n} \rightarrow [Na_{x-2y} D_{2y} M^{II}_{1-x/2a} M^{II}_{1-x/2a} PO_{3}]_{n} + n2YNa^{+}$$

where D = MB or TB, (where $M^{II} \& M'^{II} = Mg$ (II), Ba (II), Ca (II), Ni (II), Cu (II), Sr (II), Zn (II), x = 2/3, 3/4, a = valency of metal ion).

The dye ions are highly polarisable in comparison to cations and can be exchanged with the later on the polyanion. The most plausible explanation of the metachromatic effectof polyelctrolytes seems to be that the dye cations assume a more symmetrical conformation in aqueous solution while associating with the polymer chain and the system absorb at lower wave length.

Conclusion:

The important conclusion of the above study to us is that these metachromatic reactions are the best evidence of chain like character of the complex plymetaphosphate derivatives and a similar behaviour to that of Graham's salt.

REFERENCES:

- 1. T. Graham; Ann. Pharm. Liebigs, 12, 1, 1834, 29, 1, 1839.
- 2. T. Fleitmann, and W. Henneberg; Liebigs Ann. Chem., 65, 304-334, 387-390, 1849.
- **3.** G.Tammann; J. Prakt. Chem., 45, 417-474,1892.
- 4. R.E. Hall; U.S. Patent, 1, 956, 515, 1934.
- 5. F.T. Wall and R.H. Doremus, J. Amer. Chem. Soc., 76, 868, 1954.
- 6. Mehrotra, R.C. and Gupta, V.S.; J. Polymer Sci., 54, 613, 1961.
- 7. R.C. Mehrotra and C.K. Oza; Ind. J. Chem., 6, 455, 1968.
- 8. M. Oza, Ph.D. Thesis, Univ. of Rajasthan, Jaipur, 1987.
- 9. Sobhana Sharma, Ph.D. Thesis, Univ. of Rajasthan, Jaipur, 1996.
- **10.** A.I. Vogel, "A Text book of quantative inorganic analysis" Longmans Green and Co. London 3rd Edn., 1961.
- **11.** Michaelis, L. and Granick, S.; J.Amer.Chem. Soc., 67, 1212, 1945.2. Wiame, J.W.; J. Amer.Chem. Soc., 69, 3146, 1947.
- 12. Wiem, J.W.:J.Amer.Chem.Soc,69,3146, 1947
- 13. Anne, L., Levine, A. and Schubert, M; Ibid, 74, 91, 1952.
- 14. Mckay, R.B. and Hillson, P.J.; Trans Faraday Soc., 61, 1800, 1965.
- **15.** Mehrotra, R.C. and Gupta, V.S.; J. Polymer Sci., 42, 3963, 1964.
- 16. Mehrotra, R.C., Vyas, P.C. and Ojha, C.K.; J. Polym. Sci., A3, 2535, 1965.
- 17. Vyas, P.C.; Ph.D. Thesis, Univ. of Rajasthan, Jaipur, 1967.
- 18. Oza, Mamta; Ph.D. Thesis, Univ. of Rajasthan, Jaipur, 1987.
- 19. Krishnan, P.S. and Damle, S.P.; Arch Biochem. Biophys., 49, 58, 1954.
- 20. Mehrotra, R.C. and Gupta, V.S.; J. Polymer Sci., 55,81, 1961.
- **21.** Yamaoka,K., Suenaga,T., Fuyita,A. and Miura,M.; J. Sc. Hiroshima Univ. Ser. A II 34, 1, 1970
- 22. Pal,M.K. and Ash,S.K.; Histochem., 27, 36, 1971.
- 23. Pal, M.K. and Manju, B.; Histochem., 27, 36, 1971.
- **24.** Scheibe Guenter; Pallette 35, 28, 1970.
- 25. Robinowitch, E. and Epstein, L.F.; J.Amer. Chem. Soc., 63, 69, 1941.
- 26. Walton, K.W. and Rickett, C.R.; Brit.J.Exptl. Pathol, 35, 277 1954.