Determination of formation constant of Er (III) systems with different N, S & O donor ligands

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Abstract

Formation Constant of Erbium (III) Systems with different N, S & O donor ligands have been measured with the help of Spectrophotometric method. Result indicates that solution of these complexes in solid state is difficult.

Keywords: Lanthanide (III), Complexes, Formation constant.

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Introduction

The term stability cannot be generalized for complexes, since a complex may be quite stable to one reagent and may decompose readily in presence of another reagent. [1-2]

The complexes of d-block transition metals and alkaline earth metals with various ligands have been studied extensively [3-16], where as in case of F-block lanthanide metal complexes, a limited study has so far been carried out, because of their poor tendency to form complexes. ⁽¹⁷⁻¹⁹⁾

Formation constant K shows the stability, although lanthanide complexes [20-24] are thermodynamically and kinetically less stable as compared to transition metal complexes and actinide complexes.

EXPERIMENTAL

Jobs method is a variation of spectrophotometric method [25-27] which is based on the fact that most of the complexes absorb light differently than the metal ions from which they are formed. The relationship between the absorbance or optical density at particular wavelength and concentration is expressed by Beer's law. [28-29]

In this method, standard grade chemicals which were used in the present study were of AR grade. $ErCl_{3.}6H_{2}O$ (M.W -273.61) was procured from Across India Ltd, USA and ligands with 99% purity were procured from HIMEDIA, India Ltd. The three N, S, and O donor ligands have been used for preparation of system with Er (III) and the solution spectra of these system have been recorded by using a standard spectrophotometer. Three representative ligands (L_1, L_3, L_4) have been used for this study.

Table 1. Observed values of absorbance at various concentrations of Er (III) with ligands containing N, S& O donor ligands

Metal ion concentration (in %) with ligands	Values of absorbance in mixed solutions with ligands			
	With	With	With	
	L_1	L_3	L_4	
10	0.0085	0.0089	0.0097	
20	0.0090	0.0096	0.0104	
30	0.0105	0.0112	0.0119	
40	0.0120	0.0128	0.0133	
50	0.0155	0.0169	0.0183	
60	0.0127	0.0141	0.0153	
70	0.0101	0.0123	0.0122	
80	0.0091	0.0098	0.0111	
90	0.0082	0.0088	0.0096	

		Absorbance in mixed solutions of Er (III) with			
Concentration	Concentration of				
of Er (III)	ligands	ligands			
		L1	L ₃	L_4	
M/40	M/40	0.021	0.020	0.024	
M/50	M/40	0.017	0.018	0.022	
M/60	M/40	0.016	0.017	0.019	
M/70	M/40	0.015	0.016	0.011	
M/80	M/40	0.012	0.010	0.010	

Table 2Values of absorbance in different
mixed solutions

Table 3Computed values of formationconstant from observed data

Li ga nd No	Initial conc. of Er (III) in mole/ lit A	Initial conc. of ligand in mole/ lit A	Equilib rium conc. of comple x in moles/l it x	Equilibrium conc. of Er (III) in moles/lit a-x	Equilib rium conc. of ligand in moles/l it a-x	x K = (a-x) (a- x)	log K
L1	M/50	M/50	M/65	M/50 - M/65	M/50 - M/65	692.6	2.8400
L ₃	M/50	M/50	M/70	M/50 - M/70	M/50 - M/70	422.11	2.6254
L_4	M/50	M/50	M/62	M/50 - M/62	M/50 - M/62	1058.514	3.0240

Results and discussion

Order of formation constant (k)

The computed values of the formation constants from the spectroscopic data have been tabulated in tables 3.

the following order-

 $\operatorname{Er}(\operatorname{III})$ - $\operatorname{L}_4 > \operatorname{Er}(\operatorname{III})$ - $\operatorname{L}_1 > \operatorname{Er}(\operatorname{III})$ - L_3

Comparative values of stability constants for various complexes have been shown in table-3

The formation constant data show that the stability of lanthanide complexes is similar to following type of complexes, which have been reported by earlier workers [27] at room temperature.

 $[Ag (NH_3)_2] + type of complex (logK = 3.24 - 3.81)$

1:1 Pr (III)-Oxy diacetic acid (logK=2.53)

1:1 Nd (III)-Oxy diacetic acid (logK=2.67)

1:1 Pr (III)-Sulphonanilide ligand (logK=2.9118-3.3551)

Low stability constant makes the isolation of these complexes in solid state difficult, so doped model technique has been taken as system in the electronic-spectral study.

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