

ORIENTAL JOURNAL OF CHEMISTRY

An International Open Free Access, Peer Reviewed Research Journal

www.orientjchem.org

ISSN: 0970-020 X CODEN: OJCHEG 2013, Vol. 29, No. (2): Pg. 735-739

Viscosity of Thorium Soaps

RAMAKANT SHARMA and MEERA SHARMA*

Department of Chemistry, Agra College, Agra - 282 002, India

DOI: http://dx.doi.org/10.13005/ojc/290251

(Received: March 25, 2013; Accepted: April 30, 2013)

ABSTRACT

The density and viscosity results of thorium soaps in benzene methanol mixture have been explained satisfactorily in terms of the equations proposed by Einstein, Vand and Jones-Dole. The values of the CMC and molar volume of thorium soaps calculated from these equations are in close agreement.

Key words: Thorium (II), Soaps, Viscosity.

INTRODUCTION

Metallic soaps are widely used in industries and allied sciences as catalysts, cosmetics, lubricants, greases, medicines, softeners, flatteners, stabilizers, plasticizers, emulsifiers, surface active agents waterproofing agents. Therefore, a detailed study of these soaps is required for their great importance in industrial and academic fields. Extensive work has been done on the alkali, alkaline and transition metal soaps but comparatively less work has been done on lanthanide and actinide soaps1-4. Lanthanum soaps were prepared by Skrylev et af by the reaction of lanthanum chloride and corresponding salt of fatty acid. Mehrotra et al6 prepared the soaps of lanthanum, cerium, praseodymium and neodymium by double decomposition method. Skellon and Andrews7 studied the rate of oxidation of fatty acids in the presence of thorium soaps. Volatility and analysis of cerium soaps by treating with oxalic acid was studied by Marwedel⁸. Catalytic activity of cerium, thorium and urany¹ soaps was studied by Skellon and Spence⁹ Physico-chemical studies, IR, X-ray and TGA, of thorium and lanthanum soaps in solid state was studied by Mehrotra *et al*¹⁰ Ultrasonic and conductivity studies of thorium soap solutions have beendone by Mehrotra *et al*¹¹

The present work solutions have been of density and viscosity of thorium soap solutions in benzene-methanol mixture at different temperatures in order to examine their Micellar behavior and to check the validity of various known equations.

EXPERIMENTAL

All the chemicals used for the preparation of thorium soaps were of AR grade and were purified by standard methods. Thorium soaps were prepared by direct metathesis of the corresponding sodium soaps with the required amount of aqueous solution of thorium nitrate with constant stirring. The

precipitated soap was filtered and washed first with distilled water and finally with alcohol. The metal soaps were first dried in an air oven and finally under reduced pressure and further purified by recrystallization.

The viscosity and density of the solutions of thorium soaps were measured by Ostwald's Viscometer and Pyknometer at different temperatures.

RESULTS AND DISCUSSION

Density

The density (p) of the solutions of thorium soaps Lourate and Myristate is determined at different temperatures (40, 50 and 60°C). The density increases first rapidly and then slowly with the increase in the soap concentration and temperature. The plots of density vs. soap concentration (g mol dm⁻³) are characterized by an intersection of two straight lines at a definite soap concentration which corresponds to the critical micelle concentration (CMC) of the soaps, indicating a marked change in the aggregation of the soap molecules at the CMC. The values of the CMC decrease with the chain length of soaps but increase with temperature. The plots of density vs. soap concentration below the CMC have been extrapolated to zero soap concentration and the extrapolated values of the density, ρ_0 are in agreement with the experimental values of the density of the solvent. The density results have been explained in terms of Root's equations.

$$\rho = \rho_0 + AC - BC^{3/2}$$

Where C is the concentration of the soaps (g mol dm⁻³), r and r₀ are the densities of the soap solution and solvent, respectively and the constants A and B refer to the solute-solvent and solute-solute interactions, respectively. The values of the constants A and B have been obtained from the intercept and slope of the plots of $(p-p_0)/C$ vs. $C^{1/2}$ below the CMC. The results confirm that the soapsolvent interaction is larger than the solute-solute interaction in dilute soap solution. It is, therefore, concluded that there is a marked increase in the aggregation of the soap molecules and the soap molecules do not show appreciable aggregation below the CMC.

Viscosity

The viscosity (η) of the solutions of thorium soaps (Laurate and Myristate) in benzene-methanol mixture increases with the increase in the soap concentration and chain length of soap and decreases with temperature and *vice versa*. The plots of the viscosity η *vs.* the soap concentration vare characterized by the intersection of two straight lines at a definite soap concentration which corresponds to the critical micelle concentration. The values of the CMC are affected by the chain length of the anion in the soap. The decrease in the CMC with the length of hydrocarbon chain and increase with temperature may be due to the increase in the stability of the micelles as well as due to the increase in the tendency of aggregation. The plots

Table 1: values of molar volume, $\overline{\ }_V$ and constants a and b obtained from different-equations at different temperatures

Soap	Einstein's	Vand's	Jones-Dole's eqn.		Root's eqn.	
	eqn. x 10 ⁻² dr	eqn. n³ (mol-1)	A	B x 10 ⁻²	Α	B x 10 ⁻²
Temperature 40°C						
Laurate	3.13	3.07	1.00	7.50	23.00	3.00
Myristate	3.20	3.29	1.25	7.69	23.50	3.13
Temperature 50°						
Laurate	2.75	2.76	0.75	6.67	29.75	2.50
Myristate	2.91	2.93	1.00	6.78	30.25	2.73
Temperature 60°						
Laurate	3.00	3.15	0.25	6.00	40.25	2.50
Myristate	3.20	3.22	0.75	6.11	35.25	2.85

of the viscosity vs. soap concentration below the CMC have been extrapolated to zero soap concentration and it is observed that the extrapolated values of the viscosity for zero soap

concentration are in agreement with the viscosity of pure solvents. This again confirms that the soap molecules do not aggregate to an appreciable extent below the CMC. The viscosity results are

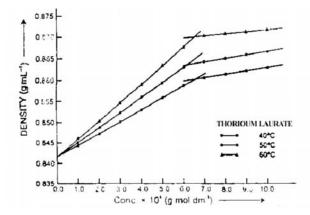


Fig. 1: Density vs Concentration (Solvent : Benzene : Methanol (1:1) Mixture)

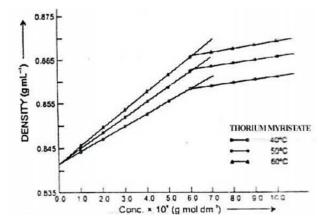


Fig. 2: Density vs Concentration (Solvent : Benzene : Methanol (1:1) Mixture)

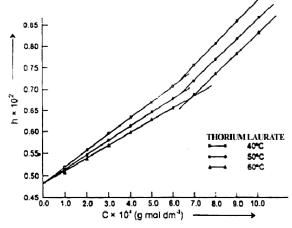


Fig. 3: Viscosity vs Concentration (Solvent : Benzene : Methanol (1:1) Mixture)

Table 2: Values of the CMC (g mol dm⁻³), C x 10⁴

Soap	Temperatures (°C)				
	40	50	60		
Thorium Laurate	6.10	6.30	6.35		
Thorium Myristate	6.00	6.20	6.30		
Conc. of Soap,	Vicosity at different temperatures (°C)				
CX10⁴(g mol dm ⁻³)	40	50	60		
Thorium laurate					
1.0	5.22	5.16	5.12		
2.0	5.58	5.47	5.40		
3.0	5.96	5.82	5.64		
4.0	6.34	6.13	5.99		
5.0	6.74	6.39	6.20		
6.0	7.12	6.77	6.52		
7.0	7.60	7.21	6.89		
8.0	8.11	7.73	7.40		
9.0	8.62	8.67	7.85		
10.0	9.00	8.62	8.34		
Thorium myristate					
1.0	5.21	5.14	5.09		
2.0	5.60	5.47	5.34		
3.0	5.97	5.84	5.62		
4.0	6.40	6.21	5.95		
5.0	6.73	6.50	6.21		
6.0	7.12	6.88	6.48		
7.0	7.64	7.35	6.90		
8.0	8.11	7.89	7.44		
9.0	8.64	8.34	7.98		
10.0	9.10	9.01	8.47		

satisfactorily explained on the basis of the following equations.

Einstein¹²:
$$\eta_{sp} = 2.5\overline{V}C$$

$$\mathsf{Vand^{13}}: \ \frac{1}{c} = \left[\frac{0.921}{V}\right]^{-1} \frac{1}{Log\left(\frac{\eta}{\eta_0}\right)} + \varnothing \overline{V}$$

$$\mathsf{Jones\text{-}Dole^{14}:} \quad \frac{\mathsf{\eta}\, sp}{C^{1/2}} = A + B \ C^{1/2}$$

Where, \overline{V} , C, ϕ and η_{sp} Are molar volume of the soap (dm³ mol¹), concentration of the soap (g mol dm³), interaction coefficient and specific viscosity of the solution, respectively.

The values of the molar volume of the soap have been calculated from the slope of the plots of $\eta_{\mbox{\tiny sp}}$ vs. C (Einstein's equation) and from the plots of 1/C vs. 1/[log (η/η_0)] (Vand's equation) and it is observed that the values obtained from both the equations are in agreement indicating that these equations are applicable to these soap solutions. The applicability of Jones-Dole's equation was checked by the plots of h_{so}/C^{12} vs. $C^{1/2}$ which are linear below the CMC. The values of coefficients A and B are calculated from the intercept and slope of the plots below the CMC. The values of B are larger than the values of coefficient A, which confirms that the molecules of the soap do not aggregate appreciably below the CMC and there is sulden change in the aggregation above the CMC.

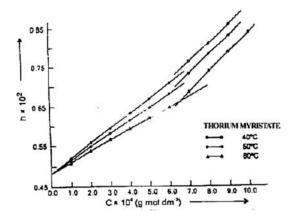


Fig. 4: Viscosity vs Concentration (Solvent : Benzene : Methanol (1:1) Mixture)

It is, therefore, concluded that the equations of Einstein, Vand and Jones-Dole are applicable to dilute solutions of thorium soaps in

benzene-methanol mixture. The values of the CMC and molar volume of thorium soaps calculate from these equations are in close agreement.

REFERENCES

- K.N. Mehrotra, A.S. Gahlaut and M. Sharma,
 J. Am. Oil Chem. Soc., 63: 157 (1986).
- 2. R.P. Varma and R. Jindal, *Tenside Detergents*, **20:** 193 (1983).
- 3. W. Brzyska and W. Hubicki, *Ann. Univ. Maide Curie Sklodowksa Soc.*,(*A/P*), 24/25, 69 (1970).
- 4. A.M. Bhandari, S. Dubey and R.N. Kapoor, *J. Am. Oil Chem. Soc.*, **47**: 47 (1990).
- 5. L.D. Skrylev, V.F. Sazonoma, M.E. Kornelli and N.A. Shukiline, *Khim. Khim. I khnol.*, **21:** 491 (1978).
- 6. R.C. Mehrotra, *Wizz. Z. Friedrich-Schillan Univ. Jena, Math. Naturewise Ket.*, **14:** 171 (1965).
- 7. J.H. Skellon and K.E. Andrews, *J. Appl. Chem.*

- (London), 5: 245 (1955).
- 8. G. Marwedel, *Farbe U. Lack*, **60:** 530 (1954); **62:** 92 (1956).
- 9. J.H. Skellon and J.W. Spence, *J. Appl. Chem.* (London), **3:** 10 (1953).
- 10. K.N. Mehrotra, A.S. Gahlaut and M. Sharma, *J. Indian Chem. Soc.*, **65:** 397 (1988).
- J. Indian Chem. Soc., 64, 729 (1987);
 Acoustics Lett. 12: 107 (1988); Recl. Trav.
 Pays Bas., 107: 310 (1988); J. Phy. Chem.
 Liq., 20: 147 (1988).
- 12. A. Einstein, Ann. Phys., 19, 289 (1906).
- 13. V. Vand. *J. Phys. Colloid Chem.*, **52**: 277 (1948).
- 14. G. Jones and M. Dole, *J. Am. Chem. Soc.*, **51**: 2950 (1929).