ISSN (Online): 2319-7064, Impact Factor (2013): 4.438

www.ijsr.net

Measurement of Acoustic and Thermodynamic Parameters of Tannic Acid and Ethanol at Various Temprature

G.M.Jamankar¹, M.S.Deshapande²

^{1, 2}Department of Physics, Ananad Niketan College, Ananadwan, Warora Dist. Chandrapur(M.S.),India

Abstract: Ultrasonic velocity, density and viscosity have been measured experimentally in binary mixture of ethanol and tannic acid at different temperature 298,303,308,313k for the frequency 2MHz. The thermodynamic and acoustical studies clarify the nature of interaction between binary solutions. acoustical parameter such as characteristics of acoustic impedance(z), adiabatic compressibility(β a), relaxation time(τ) are calculated for these solution at different concentration to study solute solvent interaction. The variation in ultrasonic velocity and other parameters play a significant role in understanding solute-solvent interaction between the constituent molecules. The variation of ultrasonic velocity, density, viscosity, relaxation time and acoustic impedance indicate the presence of strong intermolecular interaction between the components of constituents' molecules. Increase in velocity and decrease in adiabatic compressibility shows strong association and dipole-dipole interaction in the constituents' molecules.

Keywords: adiabatic compressibility, viscosity, relaxation time, tannic acid

1. Introduction

The study of intermolecular interaction plays an important role in the development of molecular sciences. A large number of studies have been made on the molecular interaction in liquid systems by various physical methods like Infrared [1,2], Raman effect [3,4], Nuclear Magnetic resonance, Dielectric constant[5], ultra violet[6] and ultrasonic method [7,8]. In recent years ultrasonic technique has become a powerful tool in providing information regarding the molecular behavior of liquids and solids owing to its ability of Characterizing physiochemical behavior of the medium. Scientific and practical interest has been stimulated by investigation of organic liquids by ultrasonic measurements. Studies involving density &viscosity measurements are important for revelation of ion-solvent, ion-ion & solvent-solvent interaction in mixed solvent systems .The ultrasonic studies in binary liquids are essential for utilizing them in medical technology. Therefore it was though worthwhile to understand ultrasonic study on some binary liquids & their mixtures [9-10]. Ultrasonic parameters are extensively being used to study molecular interaction in pure liquids, binary mixtures & ionic interaction [14-15] in single & mixed solution of tannins. Tannic acid is not a single constituent but a type of hydrolysable tannin that contain several units of Gallic or ellagic acid esterifies with glucoses -OH to produce complex tannin compounds [16].

Here the attempts have been made for experimental investigations of derived parameters such as the ultrasonic velocity, adiabatic compressibility (βa), density, and viscosity of pure binary liquid tannic acid in ethanol at various molar concentrations in the range of 0.1mol to 0.09mol. The ultrasound frequency used was 2MHz at 298K, 303Kand 308K.

Adiabatic compressibility has been calculated from the ultrasonic velocity (u) and density (ρ) of the medium using the equation as

$$\beta_a = 1 / u^2 \rho \text{ kg}^{-1} \text{ms}^2$$
(1)

It is a measure of solvent-solute chemical interaction and the point of reference for the molecular orientation. We have also calculated intermolecular free length as

$$L_f = K_T(\beta_a)^{1/2} m$$
(2)

Where K_T is Jacobson's constant.

Beyer and Letcher considered the case of plane harmonic wave & obtained a relation for acoustic impedance (Z) as

$$Z = \rho u \qquad \dots (3)$$

2. Materials and Methods

The liquid mixture of various concentrations in mole fraction was prepared by taking AR grade chemicals. The study was carried out for the temperatures 298K, 303K and 308K at fixed frequency 2MHz. The temperature of the liquid mixture was kept constant within an accuracy of $\pm 0.1^{\circ}$ C by using thermostat U-10. The experimental temperature was maintained constant by circulating water with the help of thermostatic water bath .Viscosity measurements were taken using Ostwald's viscometer with an accuracy of $\pm 0.1 \text{Kg/m}^3$. The flow of time was measured by a digital stop watch capable of registering time accurate to ±0.1sec. An average of three sets of flow of time for each solution was taken for the purpose of calculation of viscosity. The density of the solution was determined accurately using 10 ml specific gravity bottle and electronic balance and accuracy in the density measurement is $\pm 1 \times 10^{-1}$ ⁵gm/cm³. An average of triple measurements was taken into account.

Ultrasonic velocity has been measurement by Multi frequency interferometer (Mittal enterprises, M-83, New Delhi) operating at a frequency range 1MHz to 10MHz with an accuracy of ± 0.01 m/s. The source of ultrasonic waves was a quartz crystal excited by a radio frequency oscillator. The cell was filled with desired solution and water at constant temperature was circulated in the outer jacket.

ISSN (Online): 2319-7064, Impact Factor (2013): 4.438

www.ijsr.net

3. Results and Discussion

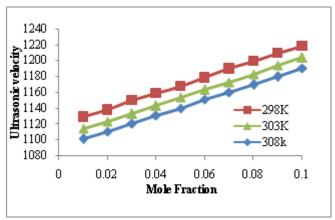


Figure 1: Variation of Ultrasonic Velocity versus molar concentration at different temperature

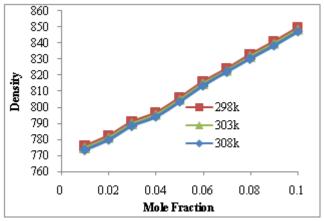


Figure 2: Variation of Density versus molar concentration at different temperature

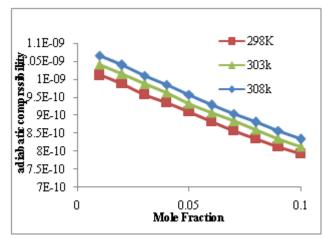


Figure 3: Variation of Adiabatic Compressibility versus molar concentration at different temperature

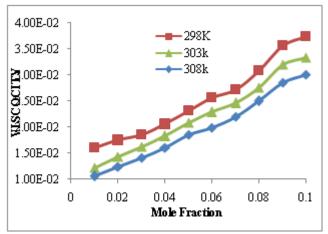


Figure 4: Variation of Viscosity versus molar concentration at different temperature

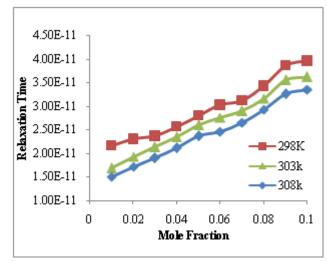


Figure 5: Variation of Relaxation Time versus molar concentration at different temperature

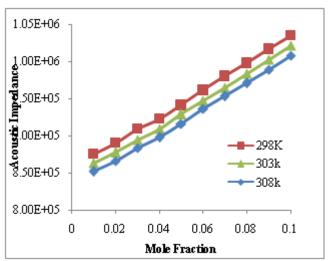


Figure 6: Variation of Acoustic Impedance versus molar concentration at different temperatures

ISSN (Online): 2319-7064, Impact Factor (2013): 4.438

www.ijsr.net

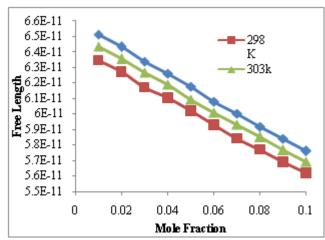


Figure 7: Variation of Free Length versus molar concentration at different temperatures

Figures (1-7) represents the variation of ultrasonic velocity, density, viscosity, adiabatic compressibility, acoustic impedance and relaxation time versus molar concentration of binary liquid mixtures of tannic acid with ethanol at temperatures 298 K, 303K and 308K.

It is observed that in fig.(1)higher mol concentration shows upward trend in ultrasonic velocity and trend is reversed for the rise in temperature With rise in mol concentration the number of particles in a given region increases and this leads to quick transfer of ultrasonic energy increasing ultrasound velocity .When temperature is increased the density of solution is decreased so that number of particles in a given region decreases hence there is no quick transfer of sound energy and thus velocity also decreases [10-13].

It is evident from fig. (2)that the density of solution increases continuously with the increase in mol concentration and decrease in temperature. Variation in viscosity indicates the presence of intermolecular interactions between the Tannic acid and ethanol.

It is observed from fig. (3)that adiabatic compressibility decreases with decreasing temperature and increase of solute concentration for the systems. The decrease of adiabatic compressibility with increasing concentration clearly indicates the presence of solute-solvent interactions due to aggregation of solvent molecules around solute molecules. A strong intermolecular H-bonding interaction exists between Tannic acid and ethanol.

It is observed that in fig. (4) viscosity of solution increases with the increase in molar concentration and decrease in temperature. The increasing concentration of solution supports non splitting of molecules hence there is increase in viscosity but when there is an increase in temperature of solution of molecules get apart hence viscosity decreases [14-15].

It is observed that in fig. (5) the relaxation time increases with increase in concentration and decrease in temperature of solution. The variation of relaxation time is a cumulative effect of the density ,viscosity and ultrasonic velocity.

It is evident from fig.(6) that the acoustic impedance increases with decrease in temperature and increase in concentration of solution. The increase of acoustic impendence with concentration is due to the effective solute-solvent interactions.

It is observed that in fig.(7) that free length decreases with increase of mole fraction of the solute in system. This may lead to the presence of specific molecular interaction between the molecules of the

Liquid mixture. Intermolecular free length which is the distance between the surfaces of the neighboring molecules. Generally, when the ultrasonic velocity increases, the value of free length decreases. The decrease in intermolecular free length indicates the interaction between the solute and solvent molecules due to which the structural arrangement in the neighborhood of constituent ions or molecules gets affected considerably Decrease in intermolecular free length in system leads to positive deviation in sound velocity and negative deviation in compressibility. This indicates that the molecules are nearer in the system

4. Conclusions

- 1)Increase in ultrasonic velocity with increase in molar concentration in this binary liquid system is due to molecular aggrigation between the constituents molecules.
- 2)Nonlinear variation of thermo acoustics parameters with molar concentration shows the strong intermolecular interactions between the constituent molecules. Decrease in adiabatic compressibility and increase in acoustic impedance with increase in molar concentration is due to association in the constituent molecules.
- 3) The nonlinear variation of thermo acoustic parameters with molar concentration provides useful information about nature of intermolecular forces existing in binary liquid mixtures.

5. Acknowledgement

M.S.Deshpande is grateful to University grant commission, New Delhi, for providing financial support to this work through Minor research project.

References

- [1] Eyring, H. andkincaid, J.F, J. Chem, Phys, 6,620.(1938).
- [2] Singh, S.,Singh,R.,Prasad, N.and prakash.S., Ind.J.Pure and Appl.Phy.,3,156 (1977).
- [3] Ramamurthy, M.and sastry, O.S., Indian. J.Pure and Appl. Phys., 21, 579, (1983).
- [4] Flethcer, A., J.Phys. Chem., 73, 2217, (1969).
- [5] Hammker, R., Clegg, R., Patterson, P., Riddick, P.and Rock, P., and Rock, S., J.Phys. Chem., 72, 1837, (1968).
 (i) Ramasamy.K.and Ranganathan.V., Indian J.Pure and Appl. Phys., 8,144, (1970).73
 - (ii) Venkateswaran.K.,Krishnapillai.M.G. and Ramasmy.K.,proc Indian Aca,Sci., 53,195, (1961).
- [6] Hobbs, M.E. and Bates, W.W., J.Am. Chem. Soc., 74,746, (1952).
- [7] Negakuva., J.Am. Chem. Soc., 76,3070, (1954).
- [8] Freedman, E., J.Chem. Phys., 21, 1784, (1955).

ISSN (Online): 2319-7064, Impact Factor (2013): 4.438

www.ijsr.net

- [9] Kannappan, A.N. and Rajendran, V., Indian J.Pure and Appl. Phys., 30,176, (1992).
- [10] Hyderkhan, V.and Subramanyam, S.V., Tras. Parad Soc. (GB)
- [11] AlkaTadkalkar, PravinaPawar and Govind K. Bichile J. Chem. Pharm. Res., 2011, 3(3):165-168
- [12] Sunanda S. Aswale, Shashikant R. Aswale, Rajesh S. Hajare Journal of Chemical and Pharmaceutical Research, 2012, 4(5):2671-2677
- [13] H.P.S.Makkar&K.Becker ,Vanillin-HCL method for condensed tannins;Journal of chemical ecology .vol.19 no.4,1993Karamali khanbabaee&Teunis van ree Nat.prod.Rep.2001,18,641-64
- [14] Rice, M.E., Ritchard, N.T., Ann E. Hagerman 2002 Food chem. 1998, 46 1887-1892.
- [15] Krzysztof Bebek Molecular & quantum acoustics vol.26(2005)
- [16] Bee Hive Digital concepts Cochin for Mahatma Gandhi University Kottyam.
- [17] A.Ali&A.K.Nainpramana-Journal of Physics vol.58, No.4 April2002 pp 695-701.
- [18] T.Sumathi&M. VaralakshmiRasayan J.Chem.vol.3 No.3(2010),550 -556.
- [19] V.S.Soilkar, S.N.Jajoo 1984 Acoustics let.79(2)1991-1996.
- [20] Pankaj, C Sharma Ultrasonic 1991,29, 344-347.
- [21] M.V.Kaulgud, K.J.Patil 1975, Indian J pure applPhys 13,238-242.
- [22] M,G.SheshagiriRao 1971. Indian J pure appl. Phys.9,169-172.
- [23] AlkaTadkalkar, K. J. Deshpande and Govind K. Bichile Archives of Physics Research, Journal of chemical and Pharmaceutical Research, 2012, 4(5):2671-2677