

EXCESS PARAMETERS OF ACOUSTICAL AND THERMODYNAMIC STUDIES IN BINARY LIQUID MIXTURES OF TETRACHLOROETHYLENE WITH NITROBENZENE AT FOUR (303.15, 308.15, 313.15 AND 318.15) K DIFFERENT TEMPERATURES

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Abstract: The density, viscosity, the speed of sound at Temperatures (303.15, 308.15, 313.15 and 318.15) K, in binary mixtures of Tetrachloroethylene with Nitrobenzene, were measured over the entire mole fraction range of the binary mixtures. Using these data, The Excess Acoustic and Thermodynamic parameters, deviations in Experimental and Theoretical values of Ultrasonic velocity were calculated. The computed quantities were fitted to the Redlich-Kister equation to derive the coefficients and estimate the standard error values. The results are discussed and interpreted in terms of structural and specific interaction predominated by hydrogen bonding.

IndexTerms - Ultrasonic velocity, density, binary liquid mixture, excess parameters, tetrachloroethylene, Nitrobenzene.

I. INTRODUCTION

Thermodynamic properties of non-electrolyte solutions have proved to be a useful tool in elucidating the structural interactions among component molecules [1]. For example, excess volume and density data can be used to study solvent-solvent specific interactions as a function of temperature, while the composition dependence provides valuable un-substitutable information about the presence and the stoichiometry of complex adducts. The intermolecular interactions influence the structural arrangement along with the shape of the molecules. The sign and magnitude of these properties guide us to understand possible interactions between the component molecules. The knowledge of physicochemical properties of non-aqueous binary liquid mixtures has relevance in theoretical and applied areas of research, and such results are frequently used in the design process (flow, mass transfer or heat transfer calculations) in many chemical and industrial processes. The excess properties derived from these physical property data reflect the physicochemical behaviour of the liquid mixtures with respect to the solution structure and intermolecular interactions between the component molecules of the mixture [2, 3]. In continuation of our studies of acoustic and volumetric properties of non-electrolyte liquid mixtures [4], the present study reports densities and viscosity at, 303.15, 308.15, 313.15, 318.15 K and speed of sound data at 303.15 and 308.15 313.15 and 318.15 K for binary mixtures of Tetrachloroethylene with Nitrobenzene, over the entire composition range.

From these data, excess molar volumes (V^E), excess speed of sound (u^E) and excess isentropic compressibility (i^E) were calculated. Further, Excess Acoustic and Thermodynamic parameters of Ultrasonic Velocity (U^E), Adiabatic Compressibility (βad^E), Intermolecular Free Length (L_f^E), Acoustical Impedance (Z^E), Molar Volume (V_m^E), Available Volume (Va^E), Viscosity ($\Delta\eta$), Free Volume (V_f^E), Internal Pressure (π^E), Enthalpy (H^E), Gibb's Free Energy (ΔG^E) at temperatures (303.15, 308.15, 313.15, 318.15) K for mole fraction Tetrachloroethylene with Nitrobenzene. The experimental speed of sound data were compared with Schaaff's collision factor theory (CFT) and Jacobson's free length theory (FLT). Liquids that were chosen in the present investigation are of much interest due to their various industrial and consumer applications. The applications of tetrachloroethylene are in dry cleaning, textile processing, degreasing metals, insulating fluid, and cooling gas in electrical transformers. The application of nitrobenzene are used to mask unpleasant odors in shoe and floor polishes, leather dressings, paint solvents, and other materials. Redistilled, as oil of mirbane, nitrobenzene had been used as an inexpensive perfume for soaps. It has been replaced by less toxic chemicals for this purpose. To the best of our knowledge, acoustic and volumetric properties of binary mixtures of Tetrachloroethylene with the nitrobenzene considered in this work are reported in literature.

II. EXPERIMENTAL SECTION

2.1 Chemicals Used:

All the chemicals used in the present work were of analytical reagent grade procured from Merck India and their purities were as follows: Tetrachloroethylene (99.5 %), Nitrobenzene (99.0 %), prior to experimental measurements, all the liquids were purified as described in the literature [5-7].

2.2 Analysis of Water Content in Chemicals:

The water content of solvents used in this work was measured using an Analab (Micro Aqua Cal 100) Karl Fischer Titrator and Karl Fisher reagent from Merck. It can detect water content from less than 10 ppm to 100 % by conduct metric titration with dual platinum electrodes.

The water contents are given in Table 1 along with their CAS number, supplier and manufacturer's stated purities. The purities of chemicals, after distillation, were checked by comparing the measured densities and speeds of sound, which are in good agreement with literature values [8 - 14] and these are given in Table 2

Table 1: List of chemicals with details of supplier, CAS number and purity

Chemical	Supplier	CAS number	Purity in mole fraction(as received from supplier)	Purity in mole fraction(after purification)	Water content
Tetrachloroethylene	Merck	127-18-4	0.990	0.995	0.012
Nitrobenzene	Merck	98-95-3	0.989	0.990	0.046

Table 2: Comparison of experimental Ultrasonic Velocity (U), Density (ρ), Viscosity (η), of pure liquid with literature value at 303.15K

Components	Velocity (U) m.s ⁻¹		Density (ρ) Kg.m ⁻³		Viscosity (η) Ns.m ⁻²	
	Exp	Lit	Exp	Lit	Exp	Lit
Tetrachloroethylene	1028.70	1028.0(8)	1606.36	1606.34(9)	0.797	0.798(10,11)
Nitrobenzene	1413.97	1432(12)	1193.3	1193.4(13)	1.6592	1.6595(14)

2.3 Measurements:

All binary liquid mixtures were prepared by weighing appropriate amounts of pure liquids on an electronic balance (Afoset, ER-120A, and India) with a precision of ± 0.1 mg, by syringing each component into airtight stopper bottles to minimize evaporation losses. The uncertainty of the mole fraction was $\pm 1.9 \times 10^{-4}$. After mixing, a bubble free homogenous sample was transferred into the U-tube of the densimeter through a syringe. The density measurements were performed with a Rudolph Research Analytical digital densimeter (Model DDM-2911), equipped with a built-in solid state thermostat and a resident program, with an accuracy of temperature of ± 0.03 K. The uncertainty in the density measurements is $\pm 2.9 \times 10^{-5}$ g.cm⁻³. Calibration of the densimeter, at each temperature, was with doubly distilled, deionized water and with air as standards. The ultrasonic speeds in the pure liquids and in their mixtures were measured by using a multi frequency ultrasonic interferometer (M-82 Model, Mittal Enterprise, New Delhi, India) single-crystal variable-path at (303.15, 308.15, 313.15 and 318.15) K. The uncertainty in the measurement of ultrasonic sound velocity is ± 0.3 %. The temperature stability is maintained within ± 0.01 K by a circulating thermostatic water bath around the cell with a circulating pump. The present investigation has been devoted to the study of densities, speed of sounds of binary liquid mixtures at different temperatures at a pressure of 0.1 MPa.

2.4 Calculations

Using the measured data, the acoustical parameters have been calculated

- ❖ Excess Ultrasonic Velocity

$$U^E = U_{(mix)} - [(1-x)U_1 + xU_2] \quad (1)$$

- ❖ Excess Viscosity

$$\eta^E = \eta_{(mix)} - [(1-x)\eta_1 + x\eta_2] \quad (2)$$

- ❖ Excess Adiabatic Compressibility

$$\beta_{ad}^E = \beta_{ad(mix)} - [(1-x)\beta_{ad1} + x\beta_{ad2}] \quad (3)$$

- ❖ Excess Intermolecular Free Length

$$L_f^E = L_{f(mix)} - [(1-x)L_{f1} + xL_{f2}] \quad (4)$$

❖ Excess Acoustic Impedance

$$Z^E = Z_{(mix)} - [(1-x)Z_1 + xZ_2] \tag{5}$$

❖ Excess Molar Volume

$$V_m^E = V_{m(mix)} - [(1-x)V_{m1} + xV_{m2}] \tag{6}$$

❖ Excess Free Volume

$$V_f^E = V_{f(mix)} - [(1-x)V_{f1} + xV_{f2}] \tag{7}$$

❖ Excess Internal Pressure

$$\pi_i^E = \pi_{i(mix)} - [(1-x)\pi_{i1} + x\pi_{i2}] \tag{8}$$

Where, x represents mole fraction of the component and subscript 1 and 2 stands for components 1 & 2.

The excess properties were fitted to a Redlich, Kister-type polynomial equation

$$Y^E = x_1 x_2 \sum_{i=0}^j A_i (1 - 2x_1)^i \tag{9}$$

The optimum number of coefficients, j, was ascertained from an examination of the variation of the standard deviation σ. The values of coefficients, Ai, were evaluated by using the method of least squares, with all points weighted equally.

$$\sigma = \left[\left(\sum \{ Y_{exp} - Y_{cal} \}^2 \right) / (N - n) \right]^{1/2} \tag{10}$$

Table 3: Values of Ultrasonic Velocity (U), Density (ρ), Adiabatic Compressibility (βad), Intermolecular Free Length (Lf), Acoustical Impedance (Z), Molar Volume (Vm), Rao’s Constant (R), Wada’s constant (W), Available Volume (Va), Degree of Molecular Interaction (χu) at temperatures (303.15,308.15,313.15,318.15)K for molefraction Tetrachloroethylene with Nitrobenzene.

X1	U m.s-1	ρ Kg.m-3	βad. 10 ⁻¹⁰ m2.N-1	Lf x 10 ⁻¹⁰ m	Z x10 ⁶ Kg.m ² . s ⁻¹	R ×10 ⁻³ m10/3 s-1/3 mol-1	W ×10 ⁻¹ (m3/mol- 1) (N/m2)1/7	Vm ×10 ⁻⁶ m3/mol-1	Va ×10 ⁻⁶ m3/mol- 1	χu ×10 ⁻²
303.15 K										
0.0000	1028.70	1606.34	5.8828	0.5032	1.6524	3.5910	5.9499	76.6400	2.7365	0.0000
0.0993	1110.65	1501.84	5.3978	0.4820	1.6680	4.0761	6.6647	84.7985	2.5935	0.0799
0.2161	1202.64	1389.3	4.9765	0.4628	1.6708	4.7020	7.5743	95.2593	2.3657	0.1615
0.3554	1247.14	1337.67	4.8064	0.4549	1.6682	5.1653	8.2614	103.3852	2.2800	0.1322
0.5244	1318.79	1296.8	4.4337	0.4369	1.7102	5.7116	9.0707	112.2116	1.9721	0.1307
0.7338	1374.39	1247.64	4.2431	0.4274	1.7147	6.3889	10.0706	123.8022	1.7456	0.0765
1.0000	1431.97	1193.3	4.0867	0.4194	1.7087	7.2703	11.3650	138.9675	1.4594	0.0000
308.15 K										
0.0000	1007.80	1597.72	6.1624	0.5200	1.6101	3.5857	5.9425	77.0535	2.8519	0.0000
0.0993	1095.04	1494.42	5.5804	0.4949	1.6364	4.0771	6.6662	85.2208	2.6895	0.0900
0.2161	1186.20	1382.01	5.1424	0.4750	1.6393	4.7053	7.5789	95.7644	2.4767	0.1689
0.3554	1257.23	1331.62	4.7510	0.4566	1.6741	5.2029	8.3130	103.8585	2.2249	0.1855
0.5244	1301.73	1291.07	4.5709	0.4479	1.6806	5.7123	9.0717	112.71380	2.1011	0.1300
0.7338	1357.08	1242.28	4.37088	0.4379	1.6858	6.3896	10.0716	124.3398	1.8877	0.0715
1.0000	1420.89	1187.49	4.1710	0.4278	1.6872	7.2870	11.3874	139.6474	1.5632	0.0000

313.15 K										
0.0000	993.00	1589.35	6.3809	5.3425	1.5782	3.5869	5.9441	77.4593	2.9386	0.0000
0.0993	1079.51	1486.96	5.7709	5.0808	1.6051	4.0782	6.6678	85.6511	2.7862	0.0903
0.2161	1169.71	1374.49	5.3174	4.8770	1.6077	4.7093	7.5845	96.2939	2.5896	0.1691
0.3554	1240.46	1325.57	4.9026	4.6830	1.6443	5.2037	8.3141	104.3404	2.3446	0.1861
0.5244	1284.69	1285.33	4.7139	4.5920	1.6512	5.7131	9.0729	113.2260	2.2313	0.1303
0.7338	1339.79	1236.91	4.5038	4.4885	1.6571	6.3904	10.0727	124.8868	2.0310	0.0717
1.0000	1403.24	1182.53	4.2946	4.3830	1.6593	7.2872	11.3876	140.2332	1.7245	0.0000
318.15 K										
0.0000	978.26	1580.9	6.6097	5.4889	1.5465	3.5881	5.9459	77.8733	3.0260	0.0000
0.0993	1064.17	1479.34	5.9691	5.2161	1.5742	4.0798	6.6701	86.0952	2.8832	0.0909
0.2161	1153.71	1365.96	5.5000	5.0070	1.5759	4.7173	7.5955	96.9011	2.7028	0.1701
0.3554	1223.74	1319.49	5.0607	4.8029	1.6147	5.2045	8.3153	104.8294	2.4651	0.1867
0.5244	1267.73	1279.59	4.8626	4.7079	1.6221	5.7139	9.0740	113.7430	2.3620	0.1306
0.7338	1322.51	1231.54	4.6425	4.6001	1.6287	6.3909	10.0735	125.4388	2.1755	0.0717
1.0000	1385.72	1177.54	4.4225	4.4898	1.6317	7.2875	11.3880	140.8274	1.8860	0.0000

Table 4: Values of Viscosity (η), Free Volume (V_f), Relaxation Time (τ), Internal Pressure (π), Gibb's Free Energy (ΔG), Enthalpy (H), Classical Absorption coefficient (α/f^2), Ultrasonic Attenuation (α) at temperatures(303.15,308.15,313.15,318.15)K for molefraction Tetrachloroethylene with Nitrobenzene.

X	$\eta \times 10^{-3}$ NSm ⁻²	$V_f \times 10^{-7}$ m ³ mol ⁻¹	$\tau \times 10^{-12}$ sec	$\pi \times 10^6$ Nm ⁻²	$\Delta G \times 10^{-20}$ K J mol ⁻¹	$H \times 10^{-3}$ J mol ⁻¹	$\alpha/f^2 \times 10^{14}$ m ⁻¹ s ²	α Neper/m
303.15 K								
0.0000	0.7972	2.2613	0.6251	458.5863	0.1077	35.1460	3.6709	48.0211
0.0993	0.9981	1.9048	0.7182	453.9051	0.1082	38.4905	5.4088	51.1038
0.2161	1.1911	1.7441	0.7902	432.5652	0.1085	41.2058	7.5811	51.9262
0.3554	1.3540	1.6230	0.8677	419.5296	0.1089	43.3731	9.2540	54.9796
0.5244	1.4511	1.7171	0.8577	389.8390	0.1088	43.7444	11.3680	51.3979
0.7338	1.5863	1.7483	0.8972	362.9293	0.1090	44.9314	13.5313	51.5895
1.0000	1.6194	2.0054	0.8822	320.9983	0.1090	44.6083	14.9422	48.6825
308.15 K								
0.0000	0.7562	2.3732	0.6211	457.0427	0.1095	35.2167	3.2565	48.7052
0.0993	0.9433	2.0304	0.7016	450.1863	0.1099	38.3652	4.8740	50.6322
0.2161	1.1165	1.8836	0.7652	427.0553	0.1103	40.8967	6.7805	50.9750
0.3554	1.2590	1.8323	0.7975	408.3106	0.1104	42.4065	8.7754	50.1278
0.5244	1.3484	1.8806	0.8215	383.2902	0.1105	43.2021	10.1116	49.8718
0.7338	1.4682	1.9263	0.8555	356.1477	0.1107	44.2833	12.0054	49.8159
1.0000	1.6010	2.0157	0.8903	324.6783	0.1108	45.3405	14.3654	49.5174

313.15 K								
0.0000	0.732	2.4365	0.6227	458.8106	0.1113	35.5391	3.0005	49.5589
0.0993	0.894	2.1531	0.6878	447.12810	0.1117	38.2970	4.4048	50.3542
0.2161	1.05	2.0212	0.7444	422.3451	0.1120	40.6692	6.0839	50.2910
0.3554	1.178	1.9843	0.7700	402.8097	0.1122	42.0293	7.8508	49.0537
0.5244	1.257	2.0479	0.7900	377.4612	0.1122	42.7384	9.0232	48.5963
0.7338	1.365	2.1077	0.8197	350.2068	0.1124	43.7362	10.6954	48.3462
1.0000	1.461	2.2693	0.8365	316.2826	0.1125	44.3533	12.5740	47.11076
318.15 K								
0.0000	0.711	2.4888	0.6266	461.2074	0.1132	35.9157	2.7712	50.6151
0.0993	0.85	2.2732	0.6764	444.5859	0.1135	38.2767	3.9915	50.2337
0.2161	0.99	2.1628	0.7260	417.7609	0.1138	40.4815	5.4699	49.7261
0.3554	1.104	2.1433	0.7449	397.6194	0.1139	41.6822	7.0317	48.1031
0.5244	1.176	2.2187	0.7624	372.2519	0.1140	42.3410	8.0756	47.5263
0.7338	1.273	2.2954	0.7879	344.8109	0.1141	43.2526	9.5519	47.0826
1.0000	1.359	2.4823	0.8013	310.9878	0.1142	43.7956	11.2160	45.6978

Table 5: Excess Acoustic and Thermodynamic parameters of Ultrasonic Velocity (U^E), Adiabatic Compressibility (β_{ad}^E), Intermolecular Free Length (L_f^E), Acoustical Impedance (Z^E), Molar Volume (V_m^E), Available Volume (V_a^E), Viscosity ($\Delta\eta$), Free Volume (V_f^E), Internal Pressure (π^E), Enthalpy (H^E), Gibb's Free Energy (ΔG^E) at temperatures (303.15, 308.15, 313.15, 318.15)K for mole fraction Tetrachloroethylene with Nitrobenzene.

X	U^E m/sec	β_{ad}^E $X10^{-10}$ $N^{-1}ms^{-2}$	L_f^E $X10^{-10}$ m	Z^E $X10^6$ Kgm^2 s^{-1}	V_m^E $X10^{-6}$ m^3/mol	V_a^E $X10^{-6}$ m^3/mol	$\Delta\eta$ $X 10^{-3}$ NSm^{-2}	V_f^E $X10^{-7}$ m^3/mol	π^E $X10^6$ N/m^2	H^E $X10^{-3}$ J/mol	ΔG^E $X10^{-20}$ J/mol
303.15K											
0.0000	0.0000	6.1136	0.000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000
0.0993	46.2365	5.9497	-0.1512	0.0199	1.9667	-0.0161	0.1234	-0.3529	9.3217	2.4044	0.0049
0.2161	89.4578	5.8439	-0.2414	0.0332	5.1473	-0.0947	0.2343	-0.4759	2.9834	4.0145	0.0085
0.3554	101.0125	5.7667	-0.2944	0.0382	4.5901	-0.0025	0.2758	-0.4632	2.6243	4.8636	0.0094
0.5244	83.9636	5.6954	-0.2240	0.0328	2.8816	-0.0945	0.2254	-0.3623	-1.2321	3.9624	0.0082
0.7338	45.7954	5.6448	-0.1435	0.0209	1.4222	-0.0536	0.1175	-0.1945	-0.1245	2.1123	0.0049
1.0000	0.0000	5.5913	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000
308.15K											
-0.0000	0.0000	6.1979	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000
0.0993	46.1843	6.0365	-0.1600	0.0223	1.9463	-0.0342	0.1104	-0.3075	6.2988	2.1423	0.0048
0.2161	89.0771	5.9125	-0.2504	0.0366	5.1761	-0.0965	0.2087	-0.4346	-1.3660	3.6124	0.0080
0.3554	102.5453	5.8376	-0.3063	0.0408	4.5481	-0.1687	0.2514	-0.4325	-1.6665	4.2369	0.0088
0.5244	84.5245	5.7701	-0.2379	0.0354	2.8227	-0.0746	0.1975	-0.3321	-4.3125	3.2874	0.0076
0.7338	46.0877	5.7113	-0.1440	0.0235	1.3446	-0.0183	0.1045	-0.1845	-2.2432	1.7245	0.0044
1.0000	0.0000	5.6535	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000
313.15K											
0.0000	0.0000	6.2597	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000
0.0993	45.6984	6.1097	-0.1663	0.0256	1.9469	-0.0315	0.0894	-0.2668	2.4964	1.8810	0.0044
0.2161	87.9292	5.9852	-0.2578	0.0405	5.2496	-0.0862	0.1602	-0.3914	-5.6207	3.2226	0.0078
0.3554	101.4884	5.8982	-0.3181	0.0452	4.5449	-0.1619	0.1904	-0.3927	-5.2865	3.6589	0.0082
0.5244	83.5254	5.8314	-0.2468	0.0386	2.8180	-0.0700	0.1457	-0.3001	-6.5396	2.8297	0.0069
0.7338	45.6040	5.7698	-0.1495	0.0263	1.3407	-0.0162	0.0804	-0.1755	-3.9639	1.3654	0.0039
1.0000	0.0000	5.7188	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000

318.15K											
0.0000	0.0000	6.3463	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000
0.0993	45.3347	6.1874	-0.1733	0.0296	1.9434	-0.0292	0.0744	-0.2149	-1.6625	1.5762	0.0041
0.2161	87.1947	6.0664	-0.2655	0.0453	5.2345	-0.0762	0.1386	-0.3458	-10.9091	2.8589	0.0072
0.3554	100.3946	5.9659	-0.3303	0.0493	4.5332	-0.1549	0.1692	-0.3431	-10.0989	3.1254	0.0078
0.5244	83.8987	5.8973	-0.2563	0.0423	2.8645	-0.0652	0.1313	-0.2666	-10.0673	2.3123	0.0062
0.7338	45.0174	5.8359	-0.1550	0.0288	1.3123	-0.0133	0.0713	-0.1725	-6.0774	1.1124	0.0033
1.0000	0.0000	5.7841	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000	0.0000

Table 6: Experimental and Theoretical values of Ultrasonic velocity at temperatures (303.15, 308.15, 313.15, 318.15) K for molefraction Tetrachloroethylene with Nitrobenzene

Mole Fraction X	U_{EXP} ms^{-1}	U_J ms^{-1}	U_{NOM} ms^{-1}	U_{FLT} ms^{-1}	U_{IDR} ms^{-1}	U_{IMR} ms^{-1}	U_{RAO} ms^{-1}
303.15 K							
0.0000	1028.70	1028.70	1.0287	102.87	1028.70	1028.70	1028.70
0.0993	1110.65	1079.29	1.0898	100.03	1059.24	1043.90	973.15
0.2161	1202.64	1135.14	1.1533	93.81	1097.26	1065.75	907.07
0.3554	1247.14	1197.12	1.2192	98.11	1145.89	1098.45	973.09
0.5244	1318.79	1266.30	1.2876	107.91	1210.31	1150.42	1092.61
0.7338	1374.39	1344.01	1.3585	121.03	1299.68	1241.40	1237.10
1.0000	1431.97	1431.97	1.4319	143.19	1431.97	1431.97	1431.97
308.15 K							
0.0000	1007.80	1007.80	1.0078	100.78	1007.8	1007.8	1007.8
0.0993	1095.04	1058.99	1.07018	98.45	1039.11	1023.22	956.20
0.2161	1186.2	1115.70	1.13507	92.76	1078.09	1045.38	892.00
0.3554	1257.23	1178.90	1.2025	97.40	1127.93	1078.54	960.78
0.5244	1301.73	1249.75	1.2726	107.29	1193.94	1131.34	1081.21
0.7338	1357.08	1329.77	1.3454	120.60	1285.47	1224.18	1227.10
1.0000	1420.89	1420.89	1.4208	142.08	1420.89	1420.89	1420.89
313.15 K							
0.0000	993.00	993.00	0.9930	99.30	993.00	993.00	993.00
0.0993	1079.51	1043.72	1.0549	97.30	1024.15	1008.30	942.86
0.2161	1169.71	1099.94	1.1193	91.99	1062.92	1030.28	878.35
0.3554	1240.46	1162.64	1.1863	96.97	1112.49	1063.18	948.57
0.5244	1284.69	1233.01	1.2559	106.84	1178.07	1115.56	1067.62
0.7338	1339.79	1312.56	1.3282	119.94	1268.94	1207.72	1211.82
1.0000	1403.24	1403.24	1.4032	140.32	1403.24	1403.24	1403.24
318.15K							
0.0000	978.26	978.26	0.9782	97.82	978.26	978.26	978.26
0.0993	1064.17	1028.51	1.0285	96.13	1009.27	993.45	929.42
0.2161	1153.71	1084.26	1.0842	91.02	1047.84	1015.25	862.98
0.3554	1223.74	1146.47	1.1464	96.47	1097.12	1047.89	936.46
0.5244	1267.73	1216.35	1.2163	106.30	1162.30	1099.87	1054.22
0.7338	1322.51	1295.44	1.2954	119.18	1252.53	1191.35	1196.75
1.0000	1385.72	1385.72	1.3857	138.572	1385.72	1385.72	1385.72

Table 7: Percentage deviation between Experimental and Theoretical values of Ultrasonic velocity at temperatures (303.15, 308.15, 313.15, 318.15) K for mole fraction Tetrachloroethylene with Nitrobenzene

Mole Fraction X	% U_{NOM}	% U_J	% U_{IDR}	% U_{FLT}	% U_{IMR}	% U_{RAO}	U^2_{exp}/U^2_{imx}
303.15 K							
0.0000	0.0000	-2.2103	0.0000	-90.0000	0.0000	-2.2103	1.0000
0.0993	0.2046	-2.8231	-4.6287	-90.9934	-6.0094	-12.3798	1.1319
0.2161	0.3180	-5.6120	-8.7621	-92.1988	-11.3818	-24.5767	1.2733
0.3554	0.3557	-4.0102	-8.1177	-92.1327	-11.9219	-21.9740	1.2890
0.5244	0.3047	-3.9798	-8.2251	-91.8170	-12.7663	-17.1500	1.3141
0.7338	0.1823	-2.2097	-5.4356	-91.1933	-9.6756	-9.9886	1.2257
1.0000	0.0000	-1.5878	0.0000	-90.0000	0.0000	3.1756	1.0000
308.15 K							
0.0000	1.0078	0.0000	1.1280	-90.0000	0.0000	4.5122	1.0000
0.0993	1.0701	-3.2918	-5.1072	-91.0093	-6.5579	-12.6789	1.1452
0.2161	1.1350	-5.9425	-9.1139	-92.1794	-11.8714	-24.8013	1.2875
0.3554	1.2025	-6.2302	-10.2838	-92.2522	-14.2128	-23.5792	1.3587
0.5244	1.2726	-3.9925	-8.2803	-91.7573	-13.0893	-16.9402	1.3238
0.7338	1.3454	-2.0117	-5.2765	-91.1126	-9.7925	-9.5775	1.2288
1.0000	1.4208	0.0000	0.0000	-90.0000	0.0000	-8.0014	1.0000
313.15K							
0.0000	0.993	-2.2897	0.0000	-90.0000	-1.1448	-1.1448	5.3425
0.0993	1.0549	-3.3153	-5.1274	-90.9868	-6.5956	-12.6577	5.6365
0.2161	1.1193	-5.9638	-9.1287	-92.1354	-11.9194	-24.9083	6.2013
0.3554	1.1863	-6.2727	-10.3162	-92.1822	-14.2911	-23.5305	5.9902
0.5244	1.2559	-4.0225	-8.2989	-91.6831	-13.1643	-16.8965	5.5213
0.7338	1.3282	-2.0323	-5.2874	-91.0471	-9.8570	-9.5513	5.0136
1.0000	1.4032	1.6203	0.0000	-90.0000	1.6203	-1.6203	4.3830
318.15K							
0.0000	0.0785	1.1621	-1.1621	-90.0000	-1.1621	-8.1349	1.0000
0.0993	1.0014	-3.3508	-5.1588	-90.9666	-6.6453	-12.6622	1.1474
0.2161	1.1500	-6.0195	-9.1762	-92.1103	-12.0006	-25.1945	1.2913
0.3554	1.2000	-6.3138	-10.3464	-92.1167	-14.3694	-23.4749	1.3637
0.5244	1.2226	-4.0521	-8.3162	-91.6144	-13.2408	-16.8418	1.3285
0.7338	1.2745	-2.0462	-5.2900	-90.9879	-9.9172	-9.5091	1.2323
1.0000	1.3208	0.0000	0.0000	-90.0000	0.0000	-3.2816	1.0000

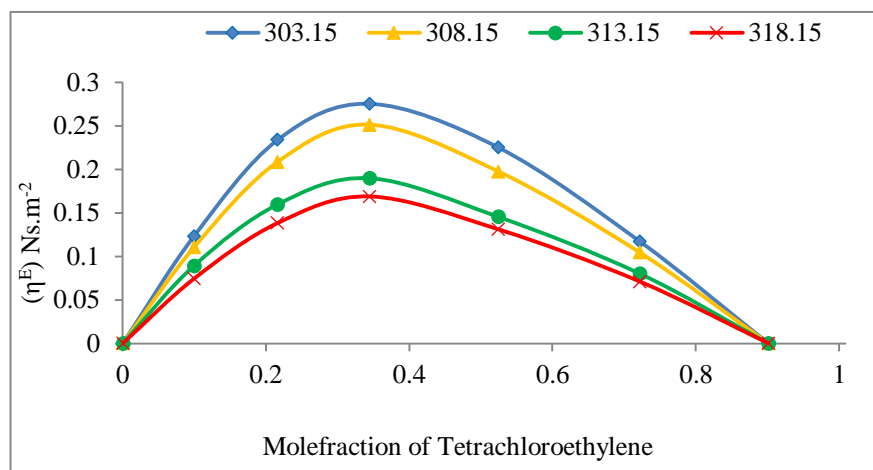


Fig.1. Variation of Excess Viscosity with molefraction of Tetrachloroethylene + Nitrobenzene.

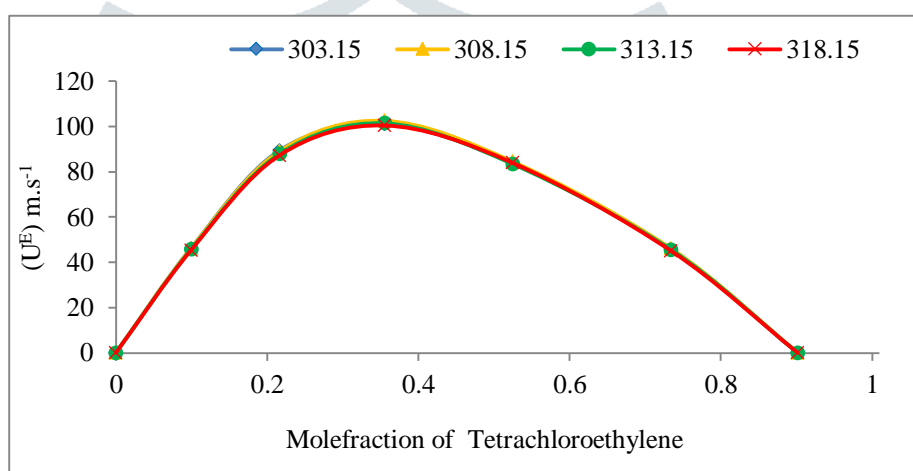


Fig.2. Variation of Excess Velocity with molefraction of Tetrachloroethylene + Nitrobenzene.

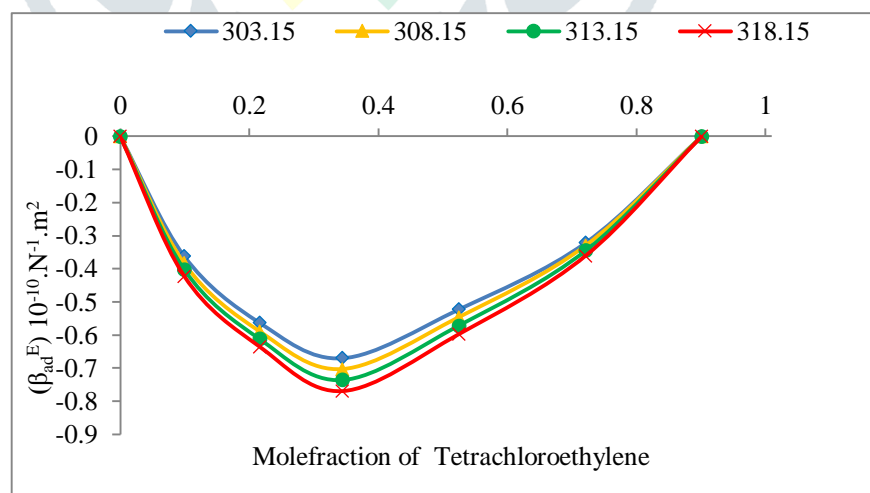


Fig.3. Variation of Excess Adiabatic compressibility with molefraction of Tetrachloroethylene + Nitrobenzene

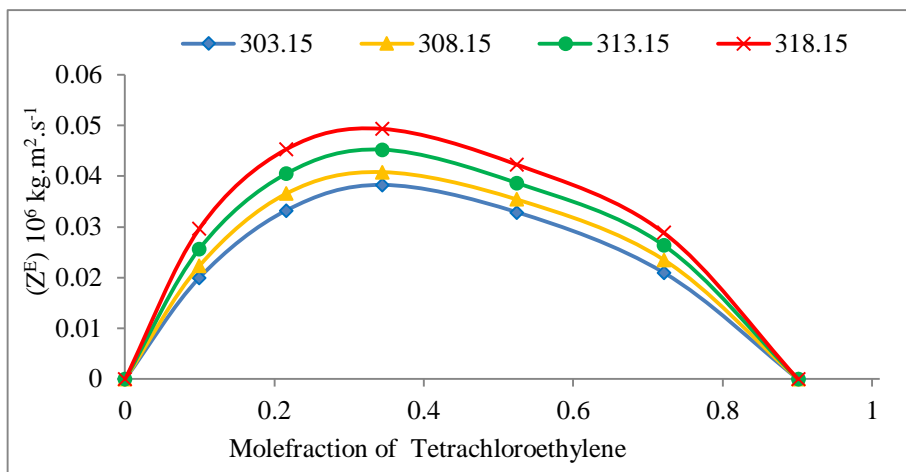


Fig.4. Variation of Excess Acoustic impedance with molefraction of Tetrachloroethylene + Nitrobenzene

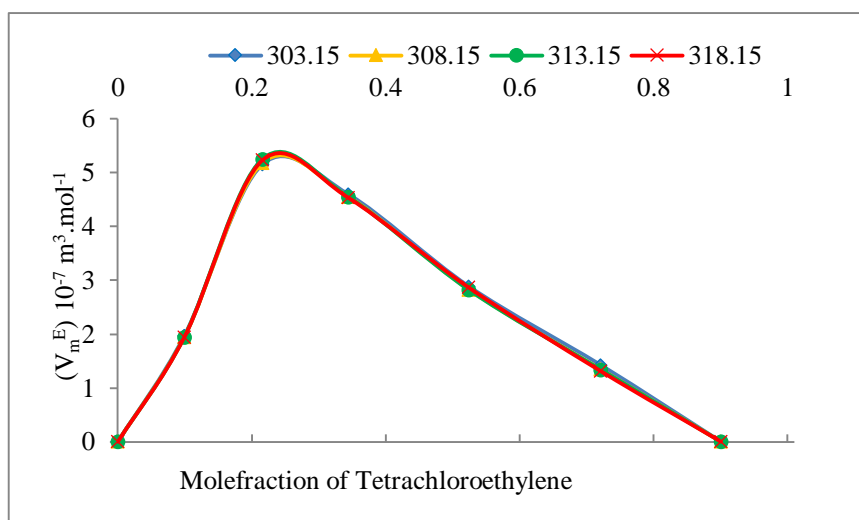


Fig.5. Variation of Excess molar volume with molefraction Tetrachloroethylene +Nitrobenzene

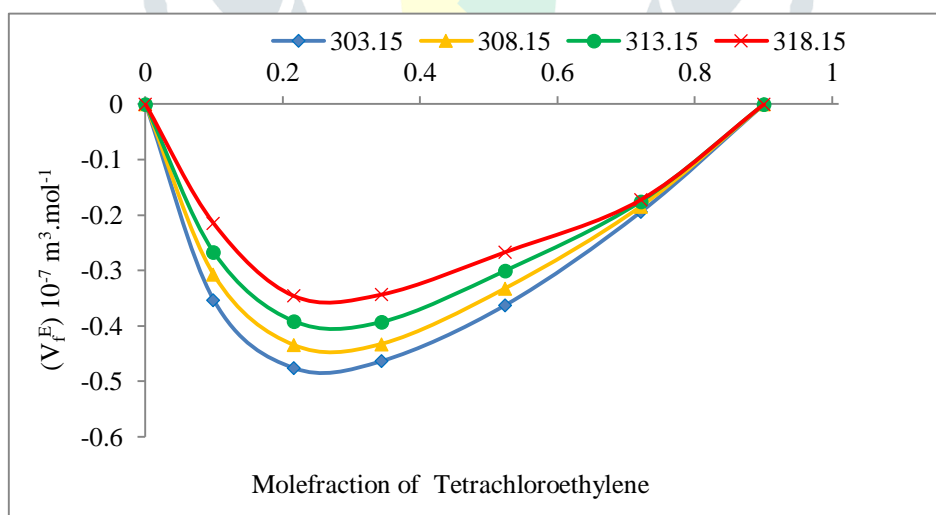


Fig.6. Variation of Excess Acoustic impedance with molefraction of Tetrachloroethylene +Nitrobenzene

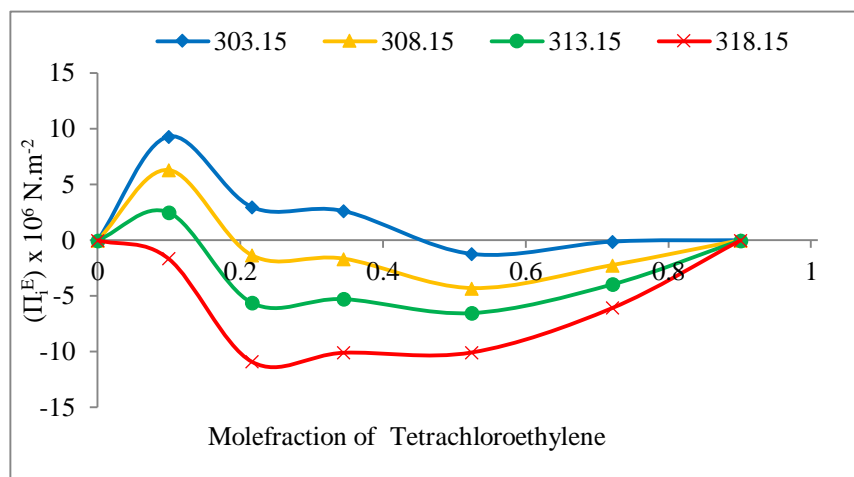


Fig.7. Variation of Excess Internal pressure with molefraction of Tetrachloroethylene +Nitrobenzene

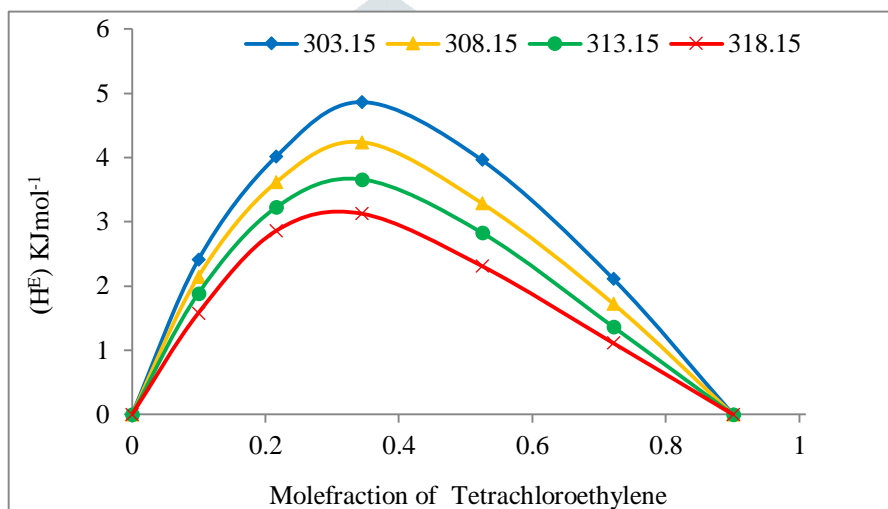


Fig.8. Variation of Excess Enthalpy with molefraction of Tetrachloroethylene +Nitrobenzene.

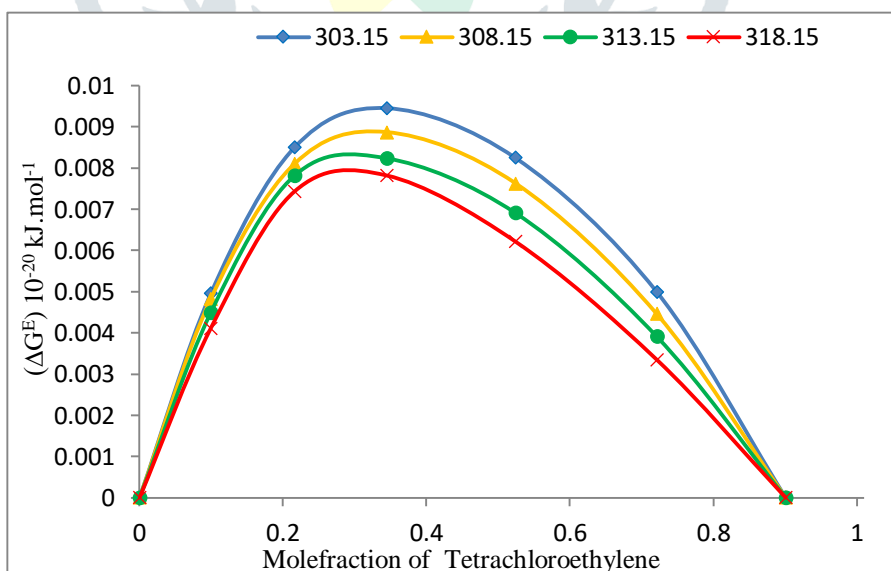


Fig.9. Variation of Excess Gibb's Free Energy with molefraction of Tetrachloroethylene +Nitrobenzene

III. Result and discussions:

The experimentally measured values of Density (ρ), Ultrasonic velocity (U), Viscosity (η) and thermodynamic parameters like adiabatic compressibility (β_{ad}), Intermolecular free length (L_f), Acoustic impedance (Z), Molar volume (V_m), Rao's Constant (R), Wada's constant (W), Internal pressure (π) and Free volume (V_f) of with nitrobenzene binary liquid system at different temperatures over the whole concentration of Tetrachloroethylene with Nitrobenzene are presented in table-2 shows us ultrasonic velocity decreases with increase in the concentration of Tetrachloroethylene and decreases with increase in

temperature¹⁵⁻¹⁸. The decrease in velocity is perhaps due to structural changes occurring in the mixtures resulting in weakening of intermolecular forces. Further the ultrasonic velocity decreases with increase in temperature at any concentrations as rise in temperature leads to less disordered structure and more spacing between the molecules. Also Density increases with increasing the concentration tetrachloroethylene of and it decreases with increasing the temperature. It suggests that a solute-solvent interaction exist between Tetrachloroethylene and Nitrobenzene system. In other words the decrease in density may be interpreted to the structure maker of the solvent due to H-bonding.

The induced dipole moment that creates induced dipole–induced dipole force of attraction between pair of atoms. This type of interaction is weaker which is given by the least ultrasonic velocity¹⁹. From the Table-2, the adiabatic compressibility and free length decreases with increasing mole fraction of the Tetrachloroethylene and increases with increasing temperature. Which suggest that making and breaking of H- bonding between molecules of the system²⁰⁻²². The intermolecular free length depends upon the intermolecular attractive and repulsive forces. Eyring and Kincaid have proposed that L_f is a predominating factor in determining the variation of ultrasonic velocity of solution. Hence it can be concluded that there is significant interaction between solute and solvent molecules due to which the structural arrangement is also affected. From the above parameters it is clear that there is a strong association between and Nitrobenzene system.

The acoustic impedance (Z) (which is the product of ultrasonic velocity and density of the solution) increases with increase in concentration of Tetrachloroethylene. It represents that there is strong interaction between the Tetrachloroethylene and Nitrobenzene system. In this system, viscosity increases with increasing molefraction of Tetrachloroethylene and decreases with increasing temperature. The decrease in density and viscosity with temperature indicates that decrease in intermolecular forces due to increase in thermal energy of the system, which cause increase in volume expansion and hence increase in free length.

The table-3 parameters investigation of Viscosity increases with concentration of Tetrachloroethylene confirms that increase of cohesive forces because of strong interaction²³⁻²⁵. The internal pressure decreases with increasing mole fraction of Tetrachloroethylene. The reduction in internal pressure may be due to the loosening of cohesive forces and adhesive force leading to breaking the structure of the solution. This gives the information regarding the nature and strength of forces existing between the molecules. The free volume decreases with increase molefraction of Tetrachloroethylene. The free volume is the space available for the molecules to move in an imaginary unit cell²⁶⁻²⁹. It clearly indicates the existence of intermolecular interaction, due to which the structural arrangement is considerably affected. International Letters of Chemistry, Physics and Astronomy Vol. 52-55. These binary systems exhibit non-linear increase/decrease in U , V_f , Z and π_i values with composition of T. This indicates the presence of intermolecular interactions between the component molecules of the mixture³⁰⁻³². In order to substantiate the presence of interactions (either adhesive or cohesive forces) between the molecules.

Table-4 the plot of excess molar volume (V^E) versus molefraction (x_1) of Tetrachloroethylene with Nitrobenzene at $T = (303.15, 308.15, 313.15, 318.15)$ K is represented in Fig. 5, and it is clear that the curves of excess molar volume (V^E) for the binary system (Tetrachloroethylene + Nitrobenzene) are positive. The observed positive values of V^E over entire range of mole fraction can be attributed to the presence of specific interactions between similar molecules of the mixtures [33]. Dipole–dipole interactions and molecular complex formation between unlike molecules may be responsible for the specific interactions [34].

Tables (5, 6) shows us the positive values of deviation in ultrasonic speed (from Fig. 1), generally, indicate dispersion forces due to strong interactions [35–37]. The observed negative values of excess acoustic impedance (Fig. 4) and positive excess free length (Fig. 3) values over the entire composition range of the liquid mixtures support the variation of deviation in ultrasonic velocity as well as deviation in isentropic compressibility[38,39]. The excess enthalpies for tetrachloroethylene with nitrobenzene are positive over the entire range of mole fractions. The curves of excess molar enthalpies vs. composition vary almost symmetrically and maximum positive values are excess enthalpy decreases as the size of the tetrachloroethylene increases. The same trend is found in the results of Kiyohara et al. [40]. In general, negative $\Delta\alpha$ values indicate the presence of strong interactions, whereas, positive $\Delta\alpha$ values are attributed to the weak interactions between the components in the mixtures [41].

CONCLUSION

From the data of ultrasonic speed, density and viscosity, various acoustical parameters and their excess values for the binary liquid mixture of Tetrachloroethylene with Nitrobenzene was measured at (303.15, 308.15, 313.15 and 318.15) K, it is obvious that there exist strong molecular interactions between Tetrachloroethylene and Nitrobenzene. The existence of type of molecular interactions in solute-solvent is favoured in the system, confirmed from the U , ρ , β_{ad} , L_f , Z , V_m , R , W , η , π and V_f data. Weak dispersive type intermolecular interactions are confirmed in the systems investigated. The observed negative and positive values of deviation/excess properties are attributed to the dispersion forces between the unlike molecules of the mixtures. Further theoretical values of sound velocity in the mixtures have been evaluated using various theories and have been compared with experimental sound velocities to verify the applicability of such theories to the systems studied. All the experimental determinations of acoustic parameters are weakly correlated between Tetrachloroethylene with Nitrobenzene.

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