

Block Diagram of Time Domain Reflectometry

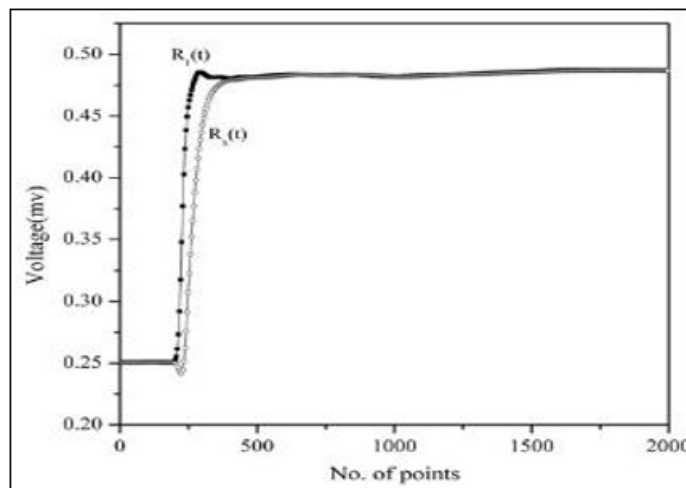


Figure 2: Reflected pulses without sample $R_1(t)$ and with sample $R_x(t)$

3. Result and Discussion

The recorded pulses are added [$q(t) = R_1(t) + R_x(t)$] and subtracted [$p(t) = R_1(t) - R_x(t)$]. Further the Fourier transformation of $p(t)$ and $q(t)$ was obtained by Summation and Samulon method for the frequency range 10 MHz to 30 GHz. The complex reflection spectra were determined as follows.

$$\rho^*(\omega) = \frac{c}{j\omega d} \frac{p(\omega)}{q(\omega)}$$

Where $p(\omega)$ and $q(\omega)$ are Fourier transformation of $p(t)$ and $q(t)$ respectively, c is speed of light, ω is the angular frequency, d is effective length and $j = \sqrt{-1}$

The complex permittivity spectrum of the Niacin (Nicotinic acid) mixtures solution is an asymmetric shape and it is determined by the Havriliak – Negami (HN) equation [11]. The complex dielectric permittivity data were fitted to HN model using non – linear least square fit method in order to extract dielectric relaxation time with the following expression [6, 11].

$$\varepsilon^*(\omega) = \varepsilon_\infty + \frac{(\varepsilon_0 - \varepsilon_\infty)}{[1 + (j\omega\tau)^{1-\alpha}]^\beta}$$

Where ε_0 is the static dielectric constant, ε_∞ is the dielectric constant at high frequency. τ is relaxation time, the α & β are symmetric and asymmetric distribution of relaxation time respectively. The Havriliak– Negami equation includes the relaxation model as a limiting form.

- 1) If $\alpha = 0$ and $\beta = 1$ then single Debye equation.
- 2) $0 \leq \alpha \leq 1$ then it would be Cole – Cole model of symmetric distribution of relaxation times.
- 3) $\alpha = 0$ and β varied such that $0 \leq \beta \leq 1$ this behavior is identified as Cole Davidson (CD) asymmetric distribution of relaxation time.

The value of static dielectric constant (ε_0), dielectric relaxation time (τ) and dielectric constant at high frequency (ε_∞) are reported in table – 1 for Nicotinic acid - water mixture. The relaxation time observed to decrease

systematically as increase in concentration as shown in table 1.

Table 1: Dielectric parameter of Niacin - water mixture.

Conc.	ε_0	τ (PS)	ε_∞	β
0.01	81.42	8.69	2.00	0.946
0.02	81.07	8.41	2.00	0.934
0.03	80.28	8.34	2.00	0.922

With the increase in concentration of solution the decrease in dielectric constant and systematic change in the dielectric parameters of the solution can be explained on the basis of molecular interaction. The decrease in τ values with increase in concentration is indicating that the number of dipoles decreases in solution. The intermediate structure formed rotates fast there by giving the decrease value of τ in solution.

Conclusion

The dielectric properties of Niacin - water mixture have been studied using time domain reflectometry technique in the frequency range 10 MHz to 30 GHz at 25°C temperature and various concentrations. The deviation in dielectric constant and relaxation times from ideality may be due to interaction in Niacin - water mixture dielectric constant and relaxation time goes on decreasing as concentration increases.

From the observed result we can conclude that dielectric constant, relaxation time of Niacin - water mixture are depending on concentration.

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