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Molecular interactions in silicone rubber- nano hydroxyl apatite system in solution phase probed by ultrasonic technique

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Abstract

Ultrasonic investigations have been carried on the solutions of silicone rubber (SR) - nano hydroxyl apatite (n-HA) systems at an operating frequency of 2 MHz. Ultrasonic velocity (v), density (ρ) and refractive index (n) have been examined for the solutions of SR (average molecular weight = 1, 28,000 g/mol) and those of SR- n HA in toluene and xylene. The ' v ' value has been found to be increased with increase in concentration of SR in the solvents. For a given SR content, the ' v ' value has been observed to be higher for toluene than xylene. The results have been interpreted in terms of closer solubility parameter values. With the addition of n- HA, the ' v ' increases for a given SR content and solvent. This has been accounted in terms of better molecular interaction being developed in the medium with the incorporation of n- HA. Various acoustical parameters such as adiabatic compressibility, acoustic impedance, Rao's constant, van der Waals constant and space filling factor have been computed to complement the observations. The observations on SR/ n-HA systems is proposed for utilization in the encapsulation of drugs, for controlled release, under ultrasound stimulation.

Key words: Ultrasonic velocity; Silicone rubber; nano- Hydroxyl apatite; Refractive index; Density; Adiabatic compressibility

1. Introduction

Ultrasonic studies on polymer solutions have, by now, gained a lot of attention [1-4]. The sound velocity measurements in polymer solutions furnish impressive knowledge about polymer- polymer and polymer- solvent interactions, which are of greatest importance for processes involving polymer product development and their industrial applications [5-8]. Adiabatic compressibility and internal pressure, the functions of ultrasonic velocity, are prominent factors indicating molecular interactions in solutions, including hydrogen bonding [9-13].

Many interesting examinations related to ultrasonic studies on polymer solutions are available in literature. Typically, Bhatt et al [14] highlighted that the evaluation of propagation of ultrasonic

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