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Determination of the Critical Micelle Concentration of Some Surfactants in the Presence of 2-Hydroxy Propyl Methacrylate

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ABSTRACT

In this study, the critical micelle concentration (CMC) of different surfactants in the presence of 2-hydroxy propyl methacrylate (2-HPMA) as monomer has been investigated by the using of conductometry and viscometry methods. For this aim, sodium dodecyl sulfate (SDS) as anionic, dodecyl trimethyl ammonium bromide (DTAB) and cetyl trimethyl ammonium bromide (CTAB) as cationic and lecithin as nonionic surfactants have been used. The main aim of this research was to introduce of a new compatible co-surfactant for different applications in industry. Finding of this research showed that, by increasing of the monomer, the CMC values showed decreasing for lecithin and SDS. The results of viscometry confirmed that this technique can be use as a drastic and simple method for determination of the CMC.

Keywords: Critical Micelle Concentration, Conductometry, Viscometry

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INTRODUCTION

Surfactants or surface active agents or detergents have applications in various fields like biology, pharmacy and etc. Their structures are amphiphilic which consist both apolar long chain hydrocarbon and polar groups. In polar solvent like water, this compartment causes to self-associated or micellization. Depending on the surfactants structures, their micelles can be anionic, cationic, nonionic and amphotitic. CMC is the most important physical change in a solution including surfactant, when its concentration increases. Below the CMC, they are monomeric and above the CMC they are self-associated and observe as disperse form to lower their free energy. Thus, the CMC, for a surfactant in a definite solvent, is a elementary means of surfactant characterization. Various methods were used to determine the CMC in aqueous system. The most important techniques are based on conductometry, voltammetry, calorimetry, scattering methods, surface tension, UV-Vis and fluorescence spectroscopy. [1-14] Recently, interactions of the polymers and surfactants have been attended by many researchers [15-17]. In this study, the micellization of different surfactants such as SDS, DTAB, CTAB and lecithin in the presence of 2-HPMA in aqueous medium was investigated by conductometry and viscometry.

MATERIALS AND METHODS

2-HPMA as monomer was purchased from Merck.co (Germany) and after distillation under reduced pressure (23mmHg, $T_{bp}=372K$) was used. Both of cationic surfactants (DTAB, CTAB) were purchased from Merck.co and were recrystallized from 50/50 (v%)=acetone/ethanol mixture and egg lecithin was provided from Aldrich.co(England) and prepared as 40% ethanolic solution and after evaporation of all alcohol was used. SDS was supplied by Merck.co and was used as received. Crison GLP 32 conductometer and Ostwald viscometer were used to determine the CMC values at 298K. For this aim, the electric conductivity and relative viscosity versus surfactants concentration were plotted and the break points of the curves were recorded the CMC. [18]

$$\eta_r = \eta / \eta_0 = t \cdot \rho / t_0 \cdot \rho_0$$

In this equation, η and η_0 are the solvent and the solution viscosity, respectively. t is the flow time of the solution, t_0 is the flow time of the solvent and ρ and ρ_0 are the density of the solution and the solvent, respectively. [18]

RESULTS AND DISCUSSION

As shown in Figures 1-24, in all cases the electrical conductivity and the relative viscosity in the presence of monomer were increased. However, the effect of monomer in the presence of SDS and lecithin were tangible, there were not any changing in the CMC for cationic surfactants. The CMC values for SDS and lecithin decreased from 0.008M to 0.005M and 0.01M to 0.004M, respectively. In the presence of cationic surfactants (DTAB and CTAB) the CMC values were standstill and were recorded (0.006M and 0.003M), respectively. At the CMC point, the relative viscosity showed a maximum value and after complexation of monomer-surfactant due to the formation of stable drops of micelles, the Gibbs free energy decreased. At higher concentrations (post-micellar region) due to increasing of the number of micelles in system, the electrical conductivity and the viscosity was increased. The finding research, identify, the presence of 2-HPMA had positive effect to the formation of stable micelles in 2-HPMA/SDS and 2-HPMA/lecithin systems and the compatibility of monomer and SDS or lecithin is favorable. [15-17]

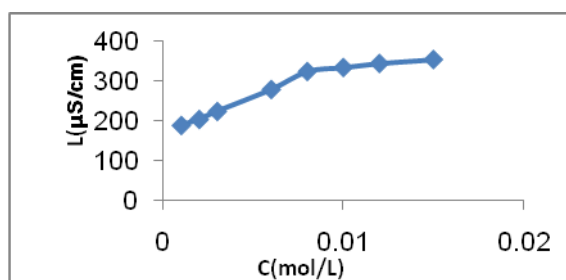


Figure 1: SDS electrical conductivity in the absence of monomer at 298K

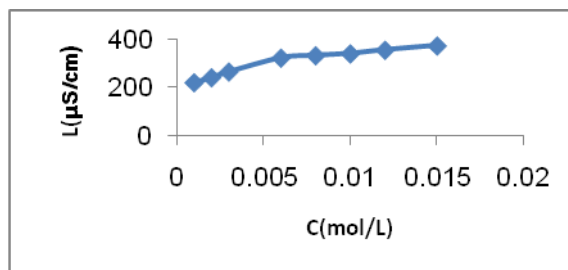


Figure 2: SDS electrical conductivity in the presence of 0.5mL monomer at 298K

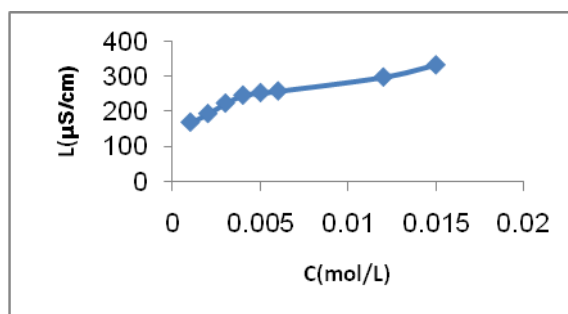


Figure 3: SDS electrical conductivity in the presence of 1.0mL monomer at 298K

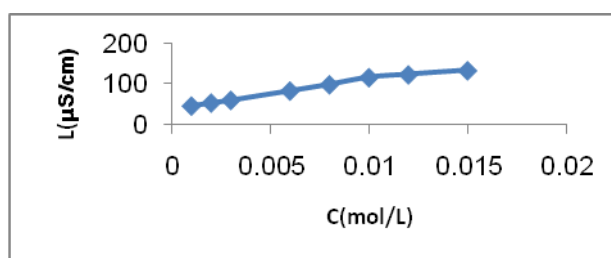


Figure 4: lecithin electrical conductivity in the absence of monomer at 298K

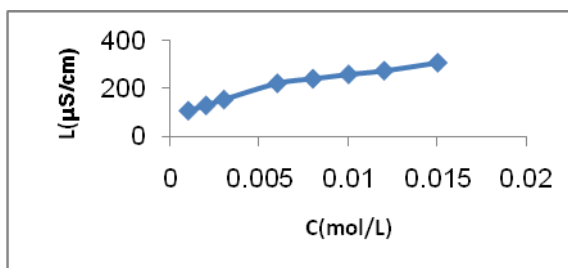


Figure 5: lecithin electrical conductivity in the presence of 0.5mL monomer at 298K

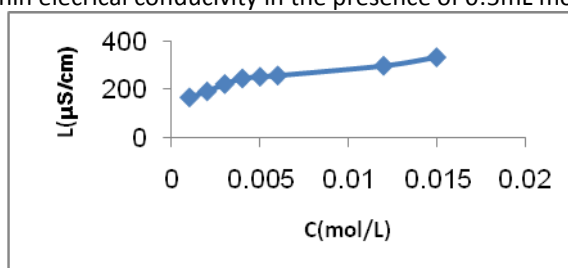


Figure 6: lecithin electrical conductivity in the presence of 1.0mL monomer at 298K

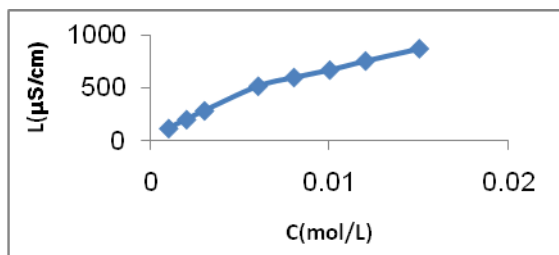


Figure 7: DTAB electrical conductivity in the absence of monomer at 298K

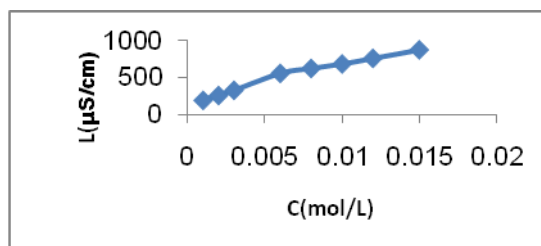


Figure 8: DTAB electrical conductivity in the presence of 0.5mL monomer at 298K

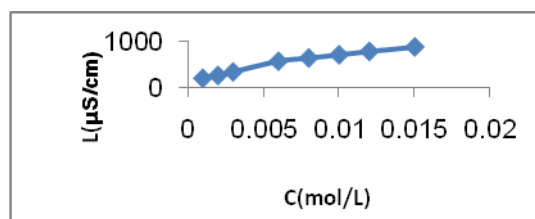


Figure 9: DTAB electrical conductivity in the presence of 1.0mL monomer at 298K

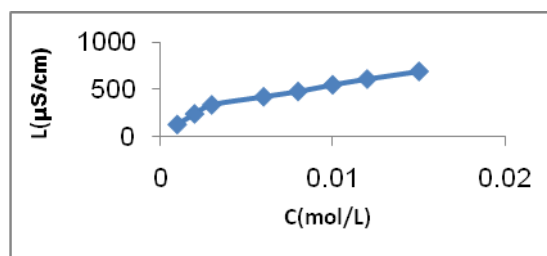


Figure 10: CTAB electrical conductivity in the absence of monomer at 298K

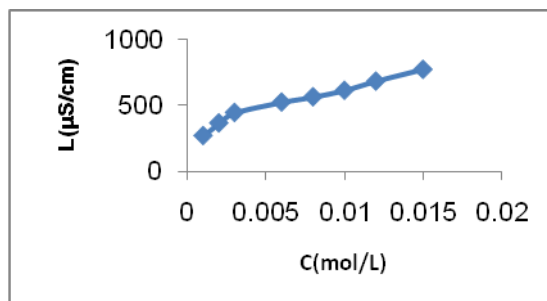


Figure 11: CTAB electrical conductivity in the presence of 0.5mL monomer at 298K

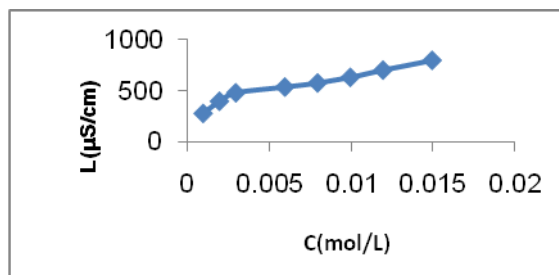


Figure 12: CTAB electrical conductivity in the presence of 1:0mL monomer at 298K

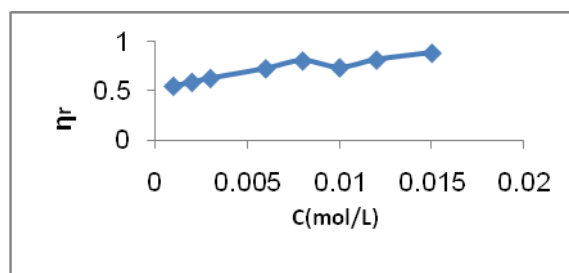


Figure 13: SDS relative viscosity in the absence of monomer at 298K

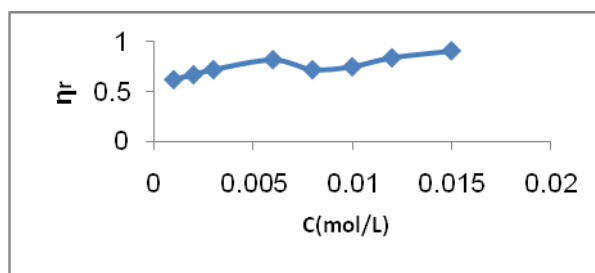


Figure 14: SDS relative viscosity in the presence of 0:5mL monomer at 298K

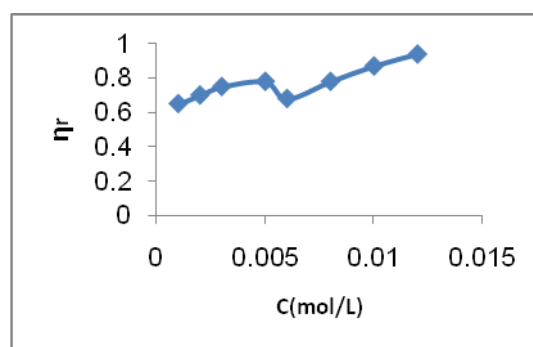


Figure 15: SDS relative viscosity in the presence of 1:0mL monomer at 298K

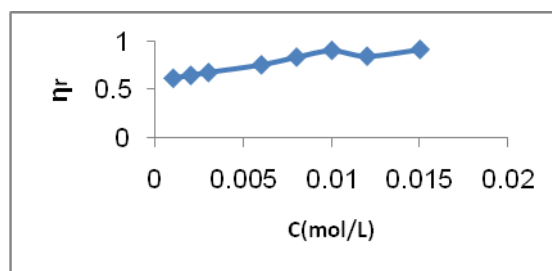


Figure 16: lecithin relative viscosity in the absence of monomer at 298K

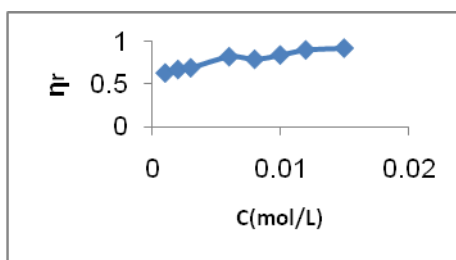


Figure 17: lecithin relative viscosity in the presence of 0:5mL monomer at 298K

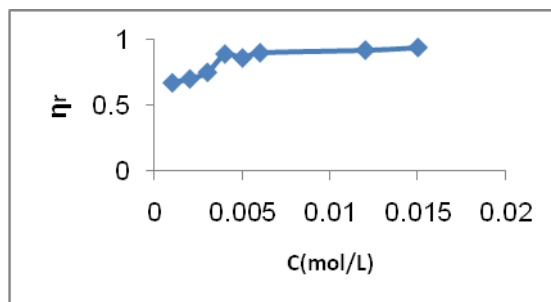


Figure 18: lecithin relative viscosity in the presence of 1:0mL monomer at 298K

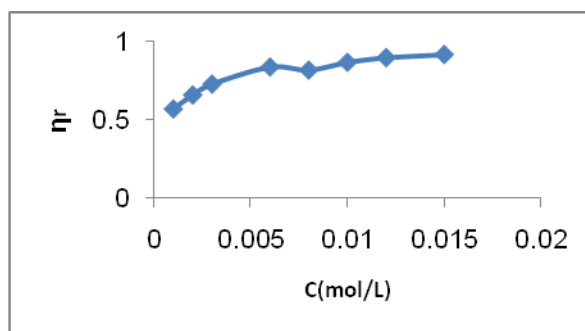


Figure 19: DTAB relative viscosity in the absence of monomer at 298K

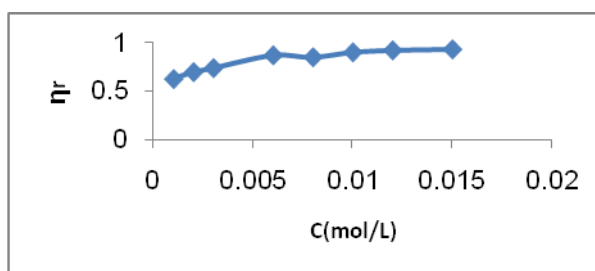


Figure 20: DTAB relative viscosity in the presence of 0:5mL monomer at 298K

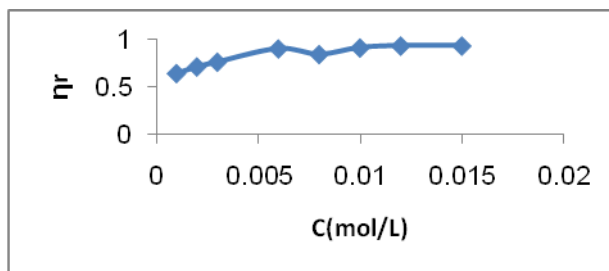


Figure 21: DTAB relative viscosity in the presence of 1:0mL monomer at 298K

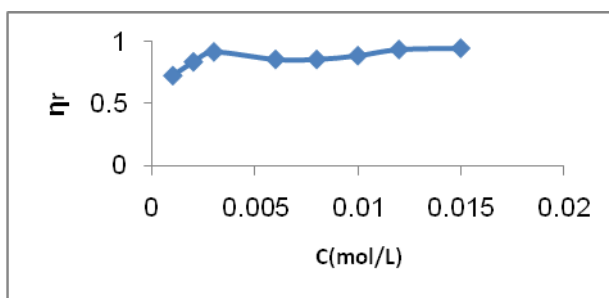


Figure 22: CTAB relative viscosity in the absence of monomer at 298K

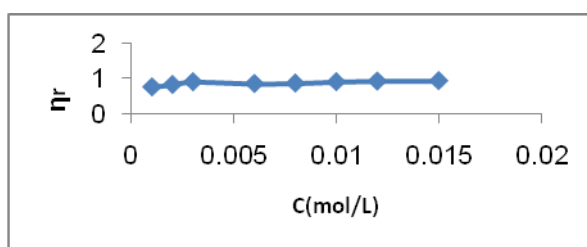


Figure 23: CTAB relative viscosity in the presence of 0:5mL monomer at 298K

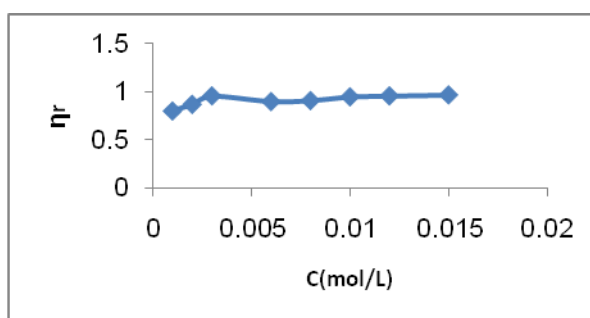


Figure 24: CTAB relative viscosity in the presence of 1:0mL monomer at 298K

CONCLUSION

A comparison of the effect of 2-HPMA in the presence of lecithin and SDS in aqueous system was recorded a synergistic effect to decreasing of the CMC. Besides of other known methods, the viscometry was introduced as an optimum and simple method to determination of the CMC.

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