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Abstract

The influence of concentration (0.125-1% w/v) on viscosity and surface tension of tamarind gum has been studied. This research establishes a direct relationship between concentration and viscosity. The intrinsic viscosity was found out to be 0.696 dl/g. The value of the Huggins constant was about 0.32 indicating the presence of molecular association. Huggins parameter ‘b’ (5.042) indicates that tamarind gum has random coil confirmation. A Kraemer constant of 0.180 gives information about the molecular association between gum molecules. Power law model has been used for predicting the molecular confirmation. A coil overlaps parameter of 0.4484 is the indication of presence of molecular entanglement. Their Values indicates polymer chain structure which are coiled & interpenetrating & overlapping with one another. The present investigation shows that surface tension decreases with an increase in the concentration of gum in solution. From these findings, it can be concluded that the gum molecules have a large interaction with more liquid than present in the air.

Introduction

Viscosity is a quantitative property of a fluid. Resistance to the flow of a fluid is termed as viscosity. The SI unit of viscosity is the poise (P); dyne·sec/cm²or g·cm/s [1]. The tendency of fluid to acquire the least surface area is termed as surface tension. The SI unit of surface tension is force/length or energy/area. Importance of surface tension of liquids is to tell about the requirement of free energy per unit area for forming interface of liquid–air at constant composition, pressure and temperature. On the other hand, viscosity determination of liquids mainly characterizes its resistance to flow. In the present work, natural polysaccharides have been used for developing controlled as well as sustained release formulations. The naturally obtained polymers are in the form of macromolecules. The natural gum has a great utilization in pharmaceutical industry [7].

Material and Methods

Tamarind seed was procured from a local market in New Delhi. Distilled water has been taken as a reference to compare the effect of mixing in the estimation of viscosity. It is given that at 25°C, 0.809 centipoise is the viscosity of water [1]. Solution of TSP prepared by dissolving required amount of TSP powder in a specified volume of distilled water.

Experimentation

Seeds of tamarind were taken. The outer coverings of seed were peeled off in such a way that the only white portion of the tamarind seed was obtained from the seed of Tamarind indica [4]. Xyloglucan is the major structure found in tamarind seeds. Broad pH tolerance, gelling capacity, high viscosity and great adhesion are all properties possessed by tamarind gum [5]. These properties help in understanding solution preparations, fundamental and structural organization. These properties have key role in predicting the heat transfer phenomenon in fluid products. These products are easily getting affected by the dispersion state, its concentration, and temperature [6]. Information regarding processing effects, effects of aging dealing with stability and changes in formulation can be predicted by measuring rheological parameters. For tamarind gum, the rheological properties serve as applicability factor in the formulation of various dosage forms in the pharmaceutical industry [7].
tamarind seeds observed. These should be crushed properly in order to get a fine powder. Crushed seeds were placed in water (24 hr) for soaking purposes. Soaked seeds were filtered through a muslin cloth to get gum. Marc was discarded and subject for multiple extraction with increasing volume of water. The formed gum was washed with ethyl alcohol (in equal amount) to precipitate the same. Further filtration was performed with ethyl alcohol. Extraction was continued in repeated manner until the material gets free from gum. The obtained gum was subjected to drying process. Drying was done at temperature 40°C. After drying, the gum was powdered and passed through sieve. Dried gum was stored at room temperature in air tight containers [8].

**Influence of concentration on viscosity**

The intrinsic viscosity is a measure of the hydrodynamic volume which is occupied by the macromolecule related closely to the confirmation and size of chains of macromolecule in its specific solvent system. It facilitates measurement of size as well as confirmation of chains of macromolecules [9]. When c approaches 0, reduced viscosity \( \eta_{rel} \) becomes \( \frac{\eta_\text{sp}}{c} \). Then expression can be represented as Huggins equation [10].

\[
\eta_{rel} = \eta_\text{sp} + kH [n]c
\]

Where, \( \eta_{rel} \) is the reduced viscosity; \([n]\) is the intrinsic viscosity and \(C\) is the intercept and \(kH\) is the intrinsic viscosity.

Huggins equation can be determined through following equation:

\[
\eta_{sp} = [n] + kH [n]^2 c
\]

Where, \( \eta_{sp} \) is the reduced viscosity; \([n]\) is the intrinsic viscosity and \(C\) is the intercept and \(kH\) is the intrinsic viscosity.

Graphical relationship between \( \ln \) relative viscosity and concentration estimated on extrapolation of line to zero concentration [11]. It gives \( \ln \) intrinsic viscosity, determined through the following equation:

\[
\ln \eta_{rel}/c = [n] + kH [n]^2 c
\]

Where, \( \ln \eta_{rel} \) is the log value of relative viscosity; \(kH\) is the Kraemer constant; \([n]\) is the intrinsic viscosity; \(C\) is the intercept and \(c\) is the concentration.

Other viscosities can also be estimated such as the relative viscosity; the specific viscosity; the inherent viscosity; and the reduced viscosity

a. The relative viscosity is the ratio of the viscosity of a solution to the viscosity of solvent used . It can be determined by using the eqn 4. Figure 2a shows the

\[
\eta_{rel}/\eta_\text{solvent} = \eta_{solvent}
\]

b. Specific viscosity can be expressed as the ratio of absolute viscosity of fluid that of a reference. It can also be determined by:

\[
\eta_s = \eta_{rel} - 1
\]

c. The inherent viscosity is the ratio of the natural logarithm of the relative viscosity to the mass concentration of gum.

\[
\eta_{inh} = \frac{[\eta]}{c} \quad \text{.....Equation 6}
\]

\[
\eta_{inh} = \frac{[\eta]}{c} \quad \text{.....Equation 6}
\]

d. The reduced viscosity is equal to the ratio of relative viscosity increment to the mass concentration of gum.

\[
\eta_{red} = \eta_{rel}/c \quad \text{.....Equation 7}
\]

A power law equation can be used to explain the polymer interactions as shown by equation 8

\[
\eta_p = aC^b \quad \text{.....Equation 8}
\]

This equation can be used to estimate the ‘b’ value from the slope of a double logarithm plot of \( \eta_p \) against concentrations. It provides information regarding polysaccharide confirmation [10]. The Huggins equation expresses \( \eta_p \) as a function of concentration of polymer in the following equation:

\[
\eta_p = (c_a + k_c [n]^2 c)
\]

\[
\eta_p = (c_a + k_c [n]^2 c)
\]

Where, \( b \) and \( k \) are Huggins parameters and \([n]\) is intrinsic viscosity respectively.

The molecular conformation and the coil overlap can be estimated via the power law model equation shown in the following equation has greater utility in studying molecular interactions are in the solution of polymer and its confirmation

\[
\ln \eta_{rel}/c = [n] + kH [n]^2 c
\]

On simplification, equation 4, can be represented as

\[
\eta_p = aC^b
\]

Where ‘a’ and ‘b’ are the power law constants; \(kH\) is the Kraemer constant; \(\ln \eta_{rel} \) is the log value of relative viscosity and \(c\) is the concentration of sample respectively.

\[
\ln \eta_p = \ln a + b\ln c
\]

The surface tension is another important parameter to be considered during formulation of better products. The chemistry which deals with surface properties, interface mechanism between two phases. The chemistry deals with chemical processes at the interface between two phases [12]. The surface tension of gum solution determined using a Stalagmometer and formula:

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determine viscosity. Viscosity is determined through Ostwald viscometer. It has been shown in following figure 1a that increase in relative viscosity of gum is observed, as we increase the concentration of the gum. It is due to increment at internal resistance flow of gum and an exertion of intermolecular friction between fluids layers which are slide over one another. Observed values gives direct relationship between viscosity with concentration while inverse relationship between concentration and the surface tension. The intrinsic viscosity can be estimated from intercept Figure 1a and was found to be 0.696dl/g. The value of Huggins constant is 0.32. Kraemer constant estimated from equation-3 shown in figure 1b. The value observed is about 0.180 and gives information about molecular association between gum molecules. In a theta solvent, the value of 'k' is between 0.5 to 0.7 ranges [13]. The intrinsic viscosity can also be determined from the Kraemer equation, gives the line between ln relative viscosity and concentration and fitting to kraemer equation. This equation provides the information regarding molecular association between gum molecules [14]. A Figure 1a shown below for gum reveals the extent degree of linearity in graph (R²=0.989). The intrinsic viscosity can also be estimated from intercept. The intercept from figure 1a is found to be 0.696dl/g. The Kₚ value can be calculated from equation 10. The value of Kₚ observed is 0.32 in a theta solvent, Kₚ value should lies in between the range of 0.3-0.8 [15]. On putting values in above equation we get Kₚ=0.17. On calculating, the Kraemer constant is found to be 0.180. The sum of Huggins and Kraemer constant should be below 0.5%+10% and signifies an absence of molecular association. The Kraemer equation gives the information regarding molecular association between gums molecules. Hence tamarind gum has molecular association between gum molecules.

The relative viscosity; specific viscosity; inherent viscosity; and reduced viscosity were determined by using equation 4,5,6,7 and their plots are shown via figure 2a-d. Intrinsic viscosity also involves Huggins parameters such as 'b' value which is found to be 5.042. This value indicates a random coil confirmation for tamarind gum. Calculated value of critical overlap concentration is found to be 0.4484 and represents the presence of molecular entalgment in gum.

After getting the equation 13, a graph has been plotted in between ln(ɳₛₚ) vs concentration which gives intercept equals to ln(a) as well as slope b. Equation for above graph is given by equation 13 and shown in figure 3. A good correlation is exhibited i.e.R²=0.766 and the value of 'b' is found to be 5.042. The value 'b' designate index for polysaccharides conformation [16]. Tamarind gum contains polymers which get coiled & started interpenetrating and overlapping to one another. This phenomenon is observed when we increase the concentration [17]. This confirms that tamarind gum contains molecular confirmation. The coil overlap property is also exhibited by the gum. If 'b' >1= Random coil confirmation; b' <1 is in rod like confirmation. The observed value of 'b' is found to be greater than unity, i.e.1 hence, the gum have random coil confirmation.
to change in confirmation of polymer termed as critical concentration \( C^* \). Critical concentration for coil can be determined through interpretation from figure 3 [18]. At zero shear rate, the concentration dependence of specific viscosity on parameter \( C^* \). For tamarind, \( C^* \) is of 0.4484 indicating the presence of molecular entanglement. From this value, it can be predicted that gum has polymer which is coiled and started interpenetrating & overlapping to one another. Hence, it confirms that gum contains molecular confirmation.

**Influence of concentration on surface tension**

Stalagmometer filled with distilled water up to the mark. The number of bubble drops counted flowing between marked volumes of stalagmometer. Three readings were taken. The average is also calculated to minimize error. Stalagmometer washed with sample gum solution. Same procedure has been repeated for all sample concentration. Using equation –14, the surface tension can be estimated. Density was determined by R.D bottle and is expressed in g/ml unit. It was found that as we increase the concentration of gum solution from 0.125, 0.25%, 0.5%, 0.75% to 1%, the surface tension decreases from 55.91, 53.22, 50.75, 47.47, 46.62. The surface tension (\( \gamma \)) of 1% solution is comparatively lower than 0.75%, 0.5%, 0.25% and 0.125%.

The surface tension can be determined from equation 14. The graph is plotted between surface tension (\( \gamma \)) and concentration (\( \text{conc.} \)) as shown in figure 4. The observed values shows that surface tension is inversely proportional to the concentration as shown in figure 4. Hence, study shows that surface tension decreases as we increase the concentration of gum solution. This occurs due to larger interaction between molecules of liquid than molecules present in air or in non-polar solvents. The cohesion forces of liquid molecule are greater than adhesion force of air molecules and leads to inward force at the surface. Tension exerted on the surface due to presence of the imbalanced force. Hence surface tension phenomena occurs [19].

**Conclusions**

The present investigation has shown that tamarind gum is precipitated in ethyl alcohol and extracted with water efficiently. Rheological properties have been determined by using Huggins model, Arrhenius model, Kraemer model and Power law models. The result obtained from the present study lead one to the conclusion that the gum has random coil confirmation and molecular entanglement. Values computed from these studies predict the influence of concentration establishes a direct relationship with viscosity. The observed value shows that surface tension gets decreases with increase in concentration of gum solution.

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**References**


