Sol Gel transition of PVC plastisols

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DETERMINING THE SOL-GEI TRANSITION OF PVC GELS

A polymeric gel is a three dimensional network forming from flexible chains through chemical cross-linking or physical interaction. The transition from the liquid state (sol) to the solid state (gel) is called the sol-gel transition and the critical point for the transition is defined as the gel point. To study a gelling material, it is important to have a method to determine the gel point in an accurate and reproducible manner. Rheologically, the independence of the loss tangent of frequency at the gel point has been widely used to measure the gel point for chemical and physical gels.

\[ \frac{G''(\omega)}{G'(\omega)} = \tan \delta = \tan \left( n \frac{\pi}{2} \right) \]  

(1)

\( n \) is the critical exponent characterizing the viscoelastic nature of a gelling material at the gel point. If the gel strength (elastic modulus) at the gel point is weak, an accurate measurement of the gel point requires high torque sensitivity of the rheometer.

Poly vinyl chloride (PVC) is known to form physical gels in numerous solvents (plasticizers). Bi(2-ethyhexyl) phthalate (DOP) is one of the most used plasticizers for PVC. A PVC/DOP system was prepared at room temperature as a function of polymer concentration. PVC has a weight-average molecular weight \( M_w \) of 250,000 and a polydispersity \( M_w/M_n \) of 2.2.
The dynamic measurements were carried out at 40 °C on an ARES rheometer equipped with a force transducer using cone/plate test fixtures. The elastic modulus $G'$, loss modulus $G''$, and loss tangent $\tan \delta$ were measured on each sample in a frequency run from 0.1 to 100 rad/s. The linear response of dynamic viscoelasticity was ensured by applying the suitable amplitude of shear.

The shear storage modulus $G'$ and the loss modulus $G''$ are shown in Figures 1 and 2 as a function of the angular frequency for the PVC/DOP samples. The polymer concentration ranges from 2.94 to 24.7 g/l. The PVC/DOP samples follow a liquid-like terminal behavior at very low concentrations while the deviation from the above relations becomes more pronounced for the higher concentrations of PVC.

$$G'(\omega) \propto \omega^{-2}, \quad G''(\omega) \propto \omega \quad \text{for} \quad \omega \to 0 \quad (2)$$

The behavior, $G'(\omega) = \text{constant at} \ \omega \to 0$ indicates the formation of a rubbery plateau. The slope of the $G'(\omega)$ versus $\omega$ (at $\omega \to 0$) in a log-log plot undergoes a change from 2 to 0, while that of the $G''(\omega)$ versus $\omega$ (at $\omega \to 0$) varies from 1 to 0. Therefore, it is reasonable to consider that the liquid-solid transition should take place at some value of the slope between 0 and 2 for the $G'(\omega)$ curves or at a slope between 0 and 1 for the $G''(\omega)$ curves. Furthermore, it is obvious that at the gel point both slopes are the same. This consideration is an approach to the onset of the rheological definition of the gel point.

Based on equation (1) the gel point can be easily determined. The method is known as frequency-independence of $\tan \delta$. It is convenient to create a multi-frequency plot of $\tan \delta$ versus gelation time, temperature, or concentration, depending on which parameter governs the gelation process. In the present case, $\tan \delta$ versus polymer concentration at different frequencies provides the desired information on the gel point, as shown in Figure 3. All curves in the figure cross at one common point on the polymer concentration axis (12.5 g/l). This point is defined as the gel point $c_{gel}$. The critical exponent $n$, directly calculated from the gel point using (1), was found to be 0.75. With a high sensitivity transducer, accurate and easy measurement of the gel point for PVC plastisols using the method based on the frequency independence of $\tan \delta$ could be performed.

![Figure 3: The cross over point of tan d as a function of concentration defines the gel point](image)