Characterizing the microstructure of 'worm-like micelles' using rheology



Introduction

The properties of worm-like micelles (WLMs) represent a key research area in both academia and industry. This is primarily due to the fact that they have widespread applications across a range of industries ranging from Personal Care to Oil Recovery. They offer a simple, cost effective way to generate remarkable viscosity and viscoelasticity. They can be made into 'smart' or stimuli responsive structures that can undergo transitions into another phase with strikingly different rheology. Such a response is of high interest for biomedical and drug delivery applications and also for separations using microfluidic devices.

Worm-like Micelles can be formed from a wide range of different surfactant systems (anionic, cationic and zwitterionic) and also from various block copolymers. The key interesting factor is that although they can be formed from such a wide variety of chemical species, their rheological response is strikingly similar and they have a distinct rheological signature. The theoretical developments, which are now well established and widely accepted, allow not only the detection of the structure (as revealed through the distinct rheological signature), but also allows the extraction of important structural parameters.



Table 1: Impact of Packing Parameter on the formed surfactant microstructure

Geometry	Packing Parameter	Microstructure
Spherical	P<1/3	
Cylindrical (worm-like)	1/2 <p<1 3<="" td=""><td></td></p<1>	
Lamellar	P>1/2	

This allows researchers to gain insight into how various formulation conditions such as electrolyte level, pH or surfactant composition impact on the microstructure of the formed worm-like micelle. Worm-like micelles are in a vast majority of cases formed from surfactants, which are amphiphilic molecules. Depending upon the surfactant packing parameter, surfactants can assemble into a wide variety of microstructures (see Table 1).

When the packing parameter is between 1/2 and 1/3, then the surfactant molecules can arrange into a rod like micellar arrangement. Based on their thermodynamics, these rod-like micelles can continue to grow with increasing concentration or on addition of an electrolyte or co-surfactant into worm like micelles and then onto nematic liquid crystals (Figure 1)

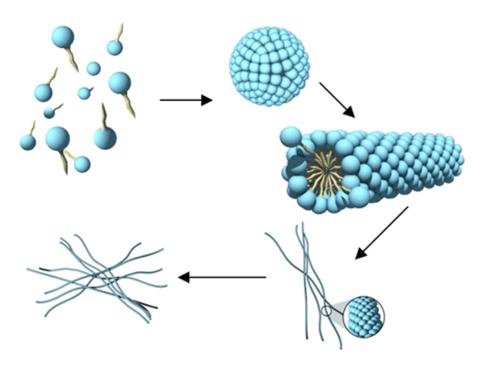


Figure 1: Phase behavior within the worm-like micelle microstructure

Each of the different phases illustrated in Figure 1 exhibits distinct rheological characteristics. The most pronounced and clear rheological signature is that of semi-dilute and concentrated worm-like micelle. Transitions from the dilute to the semi-dilute and from the concentrated to the nematic phases can also be followed through rheology.

Since they are the primary rheology building structures in a wide range of different applications, understanding their rheological signature and the changes in their structure and corresponding rheology on addition/changes in formulation is a key insight desired by both academic and industrial scientists. Rheology can provide specific insights into micellar growth, entanglement, branching, and shear induced transitions.

Theory

Worm like micelles are similar to polymers, they are long and flexible, and their spectacular viscosity and viscoelasticity is driven by entanglement of the worm-like micelles. Two key structural features which controls their rheological response is the contour length L (a measure of the end to end distance) and the persistence length I_p (a measure of the flexibility of the micelle). The elasticity of the system is impacted by the hydrodynamic correlation length ξ_H of the worm-like micelle.

Stress relaxation in a worm-like micelle, similar to polymers, can take place by reptation, (stress relaxation through snake-like motion of a polymer through a tube formed by its neighbours, until it exits the tube, at which point the stress is completely relaxed) and also by breaking and re-forming.

The reptation time is dependent on the volume fraction ϕ and is given by: $\tau_{rep} \sim L^3 \phi^{3/4}$

The breaking/forming time is given by: $\tau_{break} \sim 1/L$

When $\tau_{break} > \tau_{rep}$, the micelles behave very much like unbreakable polymers, with exponential polydispersity and the stress relaxation takes the form:

$$\sigma(t) \sim exp \left[\left(-\left(t/ au_{rep}
ight)
ight]^{0.25}$$
 Equation

If $\tau_{break} < \tau_{rep}$, the relaxation time is given by $\tau = (\tau_{break} \, \tau_{rep})^{1/2}$. Under these conditions, the fluid behaves as a Maxwell fluid for which

$$G' = \frac{G_P \omega^2 \tau^2}{1 + \omega^2 \tau^2}$$
 Equation 2

$$G^{"} = \frac{G_P \omega \tau}{1 + \omega^2 \tau^2}$$
 Equation 3

The zero shear viscosity η_0 can be linked to the plateau modulus G_p by

$$\eta = G_P \tau$$
 Equation 4

where τ is obtained from the inverse of the angular frequency ω_c at the cross-over of G' and G".

Hydrodynamic Correlation Length (ξH)

The hydrodynamic correlation length, ξ_H , can be extracted from the plateau modulus:

$$\xi_H \approx {k_B T \choose G_B}$$
 Equation 5

Where k_{B} is Boltzmann Constant and T is temperature in Kelvins. The hydrodynamic correlation length is in nanometers.

Entanglement Length (le)

If the persistence length is estimated (can easily be done) or extracted (from high frequency rheology through Microrheology or Small Angle Neutron Scattering), then one can calculate the entanglement length through

$$l_e \approx \frac{\xi_H^2}{l_p}^2 \hspace{1.5cm} \begin{array}{c} \text{Hydrodynamic Correlation length} \\ \text{Equation 6} \end{array}$$

Experimental

- In this experiment a worm-like micelle structured bodywash was evaluated in order to determine its relaxation time and hydrodynamic correlation length.
- Rotational rheometer measurements were made using a Kinexus rheometer with a Peltier plate cartridge and a cone and plate measuring system¹, utilizing standard pre-configured sequences in the rSpace software.
- A standard loading sequence was used to ensure that both samples were subject to a consistent and controllable loading protocol.
- · All rheology measurements were performed at 25°C.
- A frequency sweep test was performed between 0.2 and 40 rad/s using a strain value within the LVER.
- A Cole-Cole plot (plot of G" vs G') was produced automatically from the frequency sweep to establish whether or not the characteristic semi-circular shape (Maxwell response) of the worm-like micelle was obtained.
- Values for G_p and τ were extracted from the frequency sweep data and ξ_H calculated from the former.

Results and Discussion

The frequency response of G', G" for the bodywash product is shown in Figure 2(a) and the corresponding Cole-Cole plot is shown in Figure 2(b)

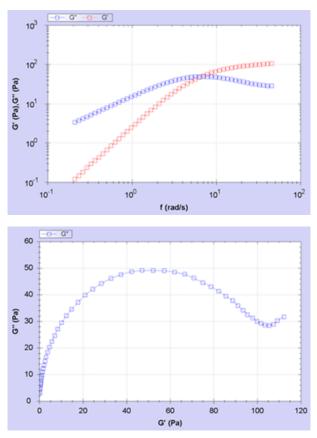


Figure 2: (a) G', G" plotted as a function of angular frequency and (b) plotted against each other (Cole-Cole plot)

The data shown in Figure 2(a) are similar to those expected for a single relaxation time Maxwell model with the onset of a plateau in G' at high frequencies (G_p) and a cross-over in G'/G" at ω_c = 1/ τ . The semicircular shape of the Cole-Cole plot confirms Maxwell behaviour. Most simple bodywash or transparent shampoo products generally conform to this behavior, the worm-like micelle structure resulting from the combination of anionic and zwitterionic surfactants in the presence of salt. In more complex formulations the presence of other additives such as perfume and pearlescent agents can cause a deviation from a purely entangled worm-like micelle system. If this deviation persists in the absence of any additives, it can then be attributed to changes in the microstructure and the structuring efficiency of the surfactant system. The ability to achieve a fully entangled worm-like micelle system at low surfactant and low salt levels is highly desirable as it implies a highly efficient structuring system.

The corresponding structural parameters extracted using theory are shown for this system in Table 2.

Table 2: Structural parameters extracted from measurement data using theory

Parameter	Hydrodynamic Correlation Length ξ _H (nm)	Relaxation Time τ (s)
Value	33.13	0.15

Conclusion

The properties of worm like micelles (WLMs) represent a key research area in both academia and industry, since they are employed in a wide range of products and applications, many of which are critically dependent on their underlying microstructure. By combining rheological measurements with theoretical understanding it has been shown that it is possible to extract key microstructural parameters including the relaxation time and hydrodynamic correlation length that are both characteristic and descriptive of the of the material and its rheological behavior.

Notes and References

Note that a parallel plate geometry or a cylindrical geometry can also be used. The
use of a solvent trap is also recommended for these tests since evaporation of
solvent (e.g. water) around the edges of the measuring system can invalidate the
test, particularly when working at higher temperatures.

Structure & Rheology of Complex Fluids-R.G.Larson

Giant Micelles-Properties & Applications-R.Zana & E.W.Kaler



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