

THE RHEOLOGY OF SWELLING CLAY DISPERSIONS

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INTRODUCTION

Nearly all muds, in drilling applications, contain smectite clay minerals, such as montmorillonite or hectorite, which have the remarkable property of swelling in the presence of water. Such systems are known to exhibit long-term gelation and thixotropic behavior, of great importance in applications, depending on clay volume fraction, particle properties (origin of the clay, surface charge, etc.) and solvent conditions (pH, ionic strength, etc.) [1-6]. Mainly two mechanisms for gel formation of smectite clays have been proposed up to now: electrostatic attraction between the positively charged edges and negatively charged faces of the platelets, resulting in a card-house structure, [7] and long-range electrostatic repulsion between interacting double layers of clay particles [8]. According to recent works, the latter mechanism has appeared to be the most probable one.

The rheological results presented in the paper have been obtained with a synthetic hectorite-type clay, namely Laponite, often considered as a model reference smectite clay. The data will be compared to those of the literature.

1 EXPERIMENT

Laponite is a smectite clay of high purity composed of particles similar to circular discs of diameter 300 Å and thickness 10 Å.

For all Laponite dispersions, the electrolyte (NaCl) concentration is set to 10^{-3} mole dm^{-3} , corresponding to a Debye length of about 100 Å. All samples have been tested at 25°C using a Low-Shear 30 viscometer, at low volume fractions, and a Carri-Med CSL 50 constant stress rheometer at higher volume fractions. The clay concentration range studied lies from 0.5% w/w to 6% w/w.

All suspensions have been set at rest for about 45 d, the longest time needed for all samples to attain their equilibrium state.

2 RESULTS

The experimental results clearly show the existence of a Laponite concentration threshold separating two types of mechanical response. This critical concentration has been proved to be 1.5% w/w, in agreement with results from the literature [6]; this critical concentration corresponds to a mean interparticle distance of about 200 Å.

2.1 Viscous fluid behavior

Up to a concentration of about 1.5% w/w, all Laponite suspensions exhibit a Newtonian behavior at low shear rates. The zero-shear viscosity increases strongly with concentration from about 1% w/w, as shown on Figure 1.

2.2 Gel-like behavior

Beyond 1.5% w/w, creep tests clearly show the existence of a yield stress, that is a stress marking the transition from solid-like behavior to a liquid-like behavior, as illustrated in Figure 2.

The yield stress thus determined, attributed to the presence of a structure, has been proved to depend on concentration according to a power law with exponent 3.3 over the whole concentration range.

In order to characterize the viscoelastic behavior of the structure below the yield stress, oscillatory shear tests in the linear regime have been carried out. The results clearly show that the level of elastic modulus G' is nearly independent

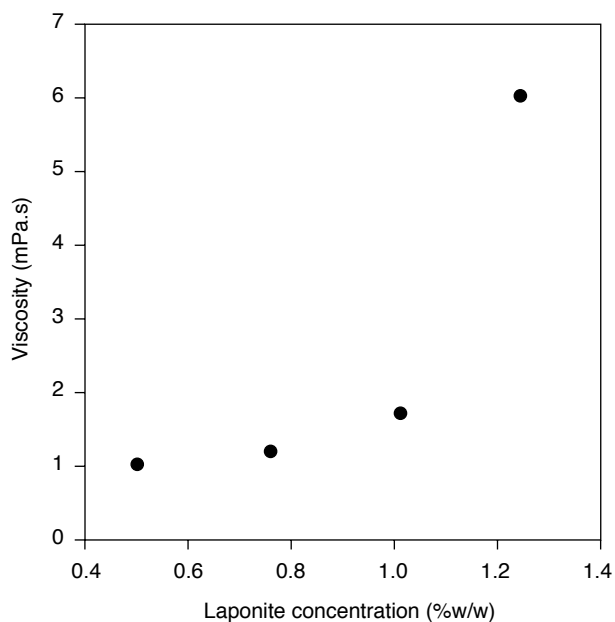


Figure 1

Zero-shear viscosity as a function of Laponite concentration.

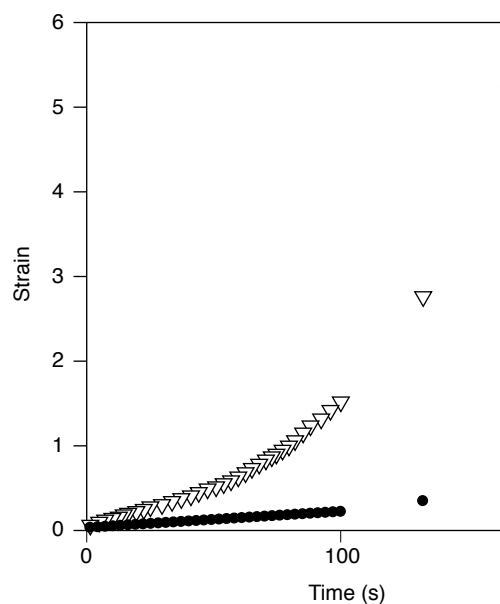


Figure 2

Illustration of the mechanical transition and yield determination: creep behavior of a 2% w/w suspension at 6 Pa and 7 Pa imposed shear stress.

The cohesive energy, that is the work required to disrupt the elastic structure, is then a power law of the concentration with exponent 3.

These results are in very good agreement with Sohm and Tadros' data [4] obtained with commercial sodium montmorillonite (Gelwhite H) suspensions.

CONCLUSION

The study of the rheological properties of Laponite dispersions at low ionic strength is essential to understand the mechanism of gelation and to characterize the dynamics and related structure of the gel inside swelling clay suspensions. Still, scattering investigations of the microscopic structures on different length scales are definitely required to precise and complete the rheological characterization. Future work should study in detail the influence of anisometry and dimensions of the clay particles on gel formation and structure.

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