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RELATIONSHIP BETWEEN THE RHEOLOGICAL PARAMETERS AND ZETA POTENTIAL OF BENTONITES****

1. INTRODUCTION

One of the most important elements of each drilling application is drilling fluid. Drilling mud fulfils a number of roles. First of all, it helps remove cuttings from borehole, maintains hydrostatic pressure against the borehole wall and seals it with a thin "wall cake". Drilling mud also lubricates and cools the drill bit. Of course, there is a difference between drilling fluids for HDD and vertical drilling. It is connected with significant differences arising from flow characteristics and physical dissimilarity borehole.

Bentonite is used as a base or starting point for drilling fluids. To control soil conditions additives and polymers are used. The main component of bentonite is montmorillonite. Different contributions of the mineral makes that not all bentonite clays may be used for drilling muds. Therefore, it is important to estimate the utility of bentonites in drilling fluids technology [1].

2. ZETA POTENTIAL AND ELECTRIC DOUBLE LAYER

Electrokinetic potential, also known as the zeta potential is a term connected with chemistry and physics. Defining the zeta potential requires knowledge of colloid chemistry. Zeta potential is a value describing the surface properties of solids dispersed in liquid. Zeta potential characterizes the interaction of colloid particles, determines the amount of electric charge in the immediate vicinity of the surface of a solid. The determination of the potential characterizes the behavior of bentonite particles in suspension, their interaction and stability of the suspension [2].

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The development of a nett charge at the particle surface affects the distribution of ions in the surrounding interfacial region, resulting in an increased concentration of counter ions (ions of opposite charge to that of the particle) close to the surface. Thus, an electrical double layer exists around each particle. The liquid layer surrounding the particle exists as two parts; an inner region, called the Stern layer, where the ions are strongly bound and an outer, diffuse, region where they are less firmly attached. Within the diffuse layer there is a notional boundary inside which the ions and particles form a stable entity. When a particle moves ions within the boundary move with it, but any ions beyond the boundary do not travel with the particle. This boundary is called the surface of hydrodynamic shear or slipping planet. The potential that exists at this boundary is known as the Zeta potential (Fig. 1) [3].

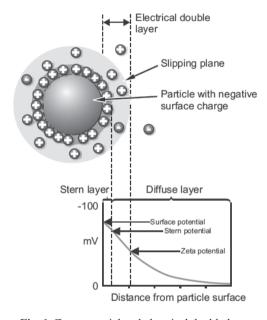


Fig. 1. Zeta potential and electrical double layer

For the purposes of the paper, Zeta potential was calculated by determining the Electrophoretic Mobility and then applying the Henry equation. The electrophoretic mobility is obtained by performing an electrophoresis experiment on the sample and measuring the velocity of the particles using Laser Doppler Velocimetry (LDV) [4].

The Henry equation is:

$$U_E = \frac{2\varepsilon \zeta f(ka)}{3\eta},$$

where:

 ζ – Zeta potential,

 $U_{\scriptscriptstyle E}$ – Electrophoretic mobility,

 ε – Dielectric constant,

η – Viscosity,

f(ka) – Henry's function.

If all the particles in suspension have a large negative or positive zeta potential, they will tend to repel each other and there is no tendency to aggregate. If the particles have low zeta potential values then there is no force to prevent the particles coming together and aggregating [5].

3. LABORATORY RESEARCH

For the purposes of the laboratory research, bentonite materials from CETCO-POLAND company were used. Rheological parameters tests have been done on the basis of API standards spec. 13A for the 3% bentonite suspensions. Studies have been completed with a full rotation speed range with 0.1 rpm and 0.036 rpm (Low Shear Rate Viscosity). Furthermore, gel strengths and LSYP (Low Shear Yield Point) were designated (Tab. 1, Fig. 2–7).

 Table 1

 Research results of rheological parameters and zeta potential values

	BCPD1	BCPD2	BCPG1	BCPG2	BCPY2	CPW14
600	2.8	19.6	3	9.2	26	27
300	1.7	17.5	1.7	5.3	22	19
200	1.4	17.7	1.2	3.8	19.9	16
100	0.8	14.8	0.6	2.3	17	11.7
60	0.5	13.8	0.4	1.9	15.9	10
30	0.4	12.9	0.3	1.3	14.8	8.7
20	0.3	12.7	0.3	1.2	14.6	8.4
10	0.2	12.5	0.2	0.9	14.3	8.1
6	0.2	12.6	0.2	0.8	14.2	8
3	0.2	12.5	0.2	0.7	14.1	7.9
2	0.2	12.6	0.2	0.7	14.1	7.9
1	0.2	12.7	0.2	0.6	14.2	8
0.1 [mPa·s]	510.9	37,200	515.5	1,580	41,700.8	24,740
0.036 (LSRV) [mPa·s]	1,318.8	107,780.4	1,394.1	6,691	121,750	75,460
0.1 [mPa·s]	0.213	8.094	0.320	0.320	12.993	8.201
0.036 (LSRV) [mPa·s]	0.320	33.015	0.320	7.242	31.950	24.495
PV [mPa·s]	1.1	2.1	1.3	3.9	4	8
AV [mPa·s]	1.4	9.8	1.5	4.6	13	13.5
YP [lb/100 sq ft]	0.639	16.401	0.426	1.491	19.17	11.715
LSYP [lb/100 sq ft]	0.213	13.206	0.213	0.639	14.91	8.307
LSYP/YP	0.333	0.805	0.500	0.429	0.778	0.709
3/600	0.071	0.638	0.067	0.076	0.542	0.293
Filtrate Volume [ml]	42.34	11.21	42.51	16.51	11.06	10.91
Moisture [%]	12.25	12.12	13.32	12.84	12.89	9.37
pH	8.13	9.99	8.28	9.63	9.26	8.86
Zeta potential [mV]	-14.5	-46.9	-23.6	-32.9	-36.8	-38.6

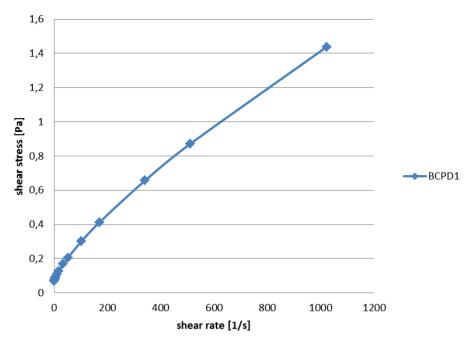


Fig. 2. The flow curve of the suspension prepared from a sample of BCPD1 based on Table 1 approximated by Herschel–Bulkley model

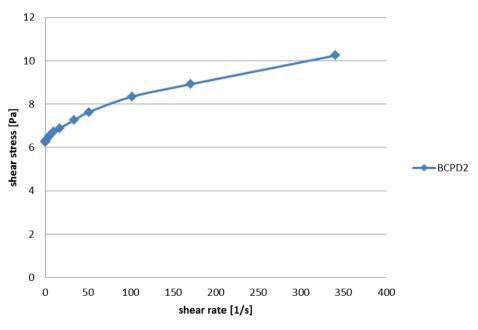


Fig. 3. The flow curve of the suspension prepared from a sample of BCPD2 based on Table 1 approximated by Herschel–Bulkley model

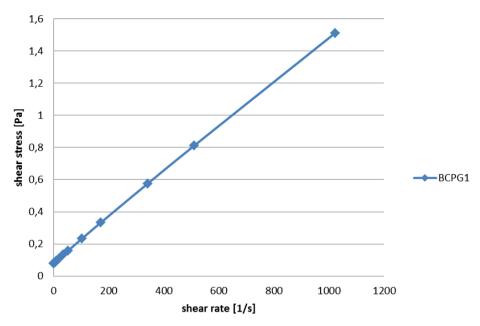


Fig. 4. The flow curve of the suspension prepared from a sample of BCPG1 based on Table 1 approximated by Herschel–Bulkley model

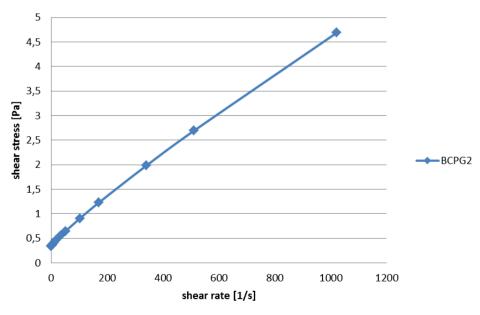


Fig. 5. The flow curve of the suspension prepared from a sample of BCPG2 based on Table 1 approximated by Herschel–Bulkley model

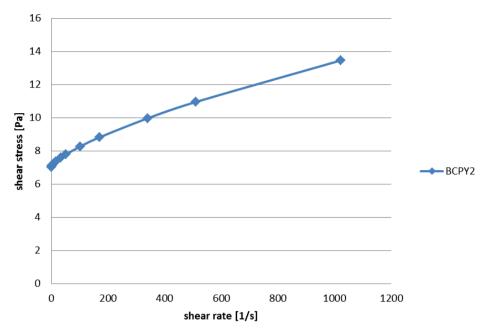


Fig. 6. The flow curve of the suspension prepared from a sample of BCPY2 based on Table 1 approximated by the Herschel–Bulkley model

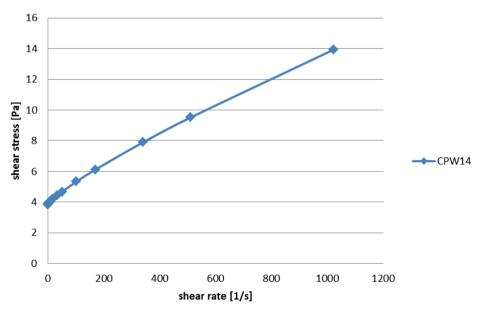


Fig. 7. The flow curve of the suspension prepared from a sample of CPW14 based on Table 1 approximated by the Herschel–Bulkley model

4. RESEARCH ANALYSIS AND CONCLUSION

The tested clay materials are characterized by great diversity. The most flat flow curves describe BCPD2 (Fig. 3) and BCPY2 (Fig. 6). For these materials shear stresses change at least relative to shear rate. The 3/600 parameter, describing the change in stress relative to shear rate, well characterizes changes in viscosity of the drilling fluid. This parameter for BCPD2 and BCPY2 is the highest and oscillates around 0.6. Both of these samples have a high viscosity at a low shear rate. Also the LSYP value is the highest and the ratio LSYP/YP exceeds 0.8. The highest structural strength is BCPY2, and the structural strength after the 10 s. BCPD2 sample is relatively low in comparison with the gelling after 10 minutes. BCPD1, BCPG1 and BCPG2 are characterized by the lowest values of the aforementioned parameters, and their flow curves are almost straight lines. When it comes to viscosity, both the plastic viscosity and the apparent, highest is the material CPW14, while the lowest values have BCPD1 and BCPG1. Other parameters, such as filtration are the most propitious for the suspension CPW14, BCPY2 and BCPD2. The filtrate volume for these samples was the lowest. In turn, filtrate volume suspensions BCPD1 and BCPG1 exceed 40 ml. Moistures of the samples are between 12–13% by weight, only for CPW14 is below 10%.

The study shows that the tested samples of the clays have a negative electrokinetic potentials. This indicates that the surface of the clay particles have a negative charge. The test suspensions have weakly alkaline and alkaline pH. The greatest absolute value of the zeta potential is characterized by BCPD2 (the most negative potential). BCPD2 suspension also exhibits the highest pH among all the tested materials. In turn, the least negative potential value has BCPD1 suspension, the pH of which is the lowest.

To sum up, electrokinetic potential analysis allows for the evaluation of the usefulness of bentonite. For high shear stresses, the relationship between viscosity and zeta potential is imperceptible but for low shear stresses the relationship is significant. The reason is, for high shear stresses friction forces play a key role and when they cease to act, when shear stress is very low, particle electrostatic attraction appears.

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