Development and research of cutting fluids for stimulation of oil production on the basis of water-salt and acid solutions containing surfactant compounds and water-soluble polymers

Reorientation in the issue of safety in oil and gas production is a recent feature of oil and gas industry in Russia and CIS countries. It remains at constant level not so much on the account of commissioning more new wells but due to improvement in the operation quality in every specific well to ensure its production capability closest to its potential [3].

The focus of attention of oil and gas industry specialists on the issues of applying modified cutting fluids in the system of oil and gas production results from their key role in production stimulation.

Cutting fluids ensure successful performance of different manufacturing operations which affect the quality of construction, productivity and duration of the well maintenance as a capital facility. Because of this, cutting fluids have to possess the range of specific properties that ensure successful implementation of various technological operations and borehole repair zone permeability [7].

The methods of oil production stimulation in domestic practice are the actual methods of oil field development, but they require further evolution of technologies in the direction of cutting fluids improvement. Such cutting fluids may contain surfactants with predetermined and regulated viscoelastic and thermochemical properties. Also, these cutting fluids must be easily destructible in the borehole zone and must not cause pollution of the reservoir.

The oil and gas industry viscoelastic compositions based on polyacrylamide, polysaccharides and other polymers have been used in a wide range of applications.

Polyacrylamide is applied for various purposes, e.g. in drilling as a stabilizer and regulator of mud rheological properties, as rock tunneling accelerator and as soil structure generator for wellbore strengthening. Recently polyacrylamide application has been reduced in connection with difficulties in its removal from the formation as its presence deteriorates the reservoir permeability [1, 2].

The main advantage of the natural vegetable polysaccharides is their low cost. But their technological indices are rather low, which narrows the field of their application. Because of this, chemically modified cellulose derivatives, i.e. starch and guar are used, as they possess a set of technological features [2].

For further effective viscosity increase, polymeric structures are often cross-linked. After pumping polymeric fluids into the formation there is no need to ensure viscosity of the composition. Therefore, the destruction of molecular skeleton of polymeric structures occurs under the influence of a destructing factor. The efficiency of the destructing factor reaches no more than 80% nowadays. Incomplete gel destruction after hydraulic fracturing leads to fracture permeability decrease.

The application of viscoelastic compositions based on polymeric substances has the following disadvantages: the necessity of cross-linking; the necessity of using a destructing factor; high friction pressure loss during pumping; incomplete recovery of un-decomposed gel after destruction.

The application of viscoelastic compositions based on surfactants helps avoid the disadvantages.

Specified surface-active substances (SAS) are able to form viscoelastic fluids in aqueous solutions. Such surfac-
Surface-active substance concentration in aqueous viscoelastic fluid reaches from 1.0 to 10.0 wt. %, preferably from 1.5 to 5.0 wt. % [6].

Compositions based on viscoelastic surfactants have a set of advantages over viscoelastic compositions based on polymeric substances. These advantages are as follows: absence of cross-linking agents, ability of application in high-temperature and high-permeable formations, low rate of leakages, structure destruction at high fluid distribution rate and its repairing at low one, structure destruction after interaction with hydrocarbon phase and complete removal of composition components from a formation. Interaction with hydrocarbons cause micelle transition from cylindrical to spherical form that results in destruction of micellar structure (Figure 2).

In the Center of Field Chemistry in Gubkin State University compositions of viscoelastic cutting fluids based on well-known industrial surfactants were developed and studied [6]. Every composition was created on the basis of principle of interaction between cationic and anionic surfactants. As a result, a surfactant molecular complex is formed. Such complexes generate micellar structures in aqueous medium.

Obtained effective viscosity values of developed compositions are not high enough with regard to hydraulic fracturing technology. So polymeric substances (0.1 wt. %) were admixed for their rheological properties improvement. Polymers integrate into micellar structure of viscoelastic surfactants and favour its strengthening (Figure 3). The following polysaccharide reagents were used for research: carboxymethyl cellulose (CMC-600), Hydroxyethyl cellulose, Hydroxypropyl guar and xanthan-type biopolymers of brands Xanthan DRILLUX 1405 and Xanthan Gum E415.

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The choice of xanthan brands is not accidental. It was established that xanthan-type exopolysaccharides of chosen brands are the best viscosifiers by a set of physical and chemical properties for water-salt and acid compositions in hydraulic fracturing technology for low-temperature carbonate reservoirs.

Rheological characteristics of complexes containing mixture of viscoelastic surfactants composition (9.9 wt.%) and polymer substance (0.1 wt.%) in aqueous solutions (10 wt.%) of sodium, potassium, calcium and magnesium chlorides, inhibited HCl (12 wt.%) solution and in soft water were studied by the method of rotary viscosimetry using apparatus Rheotest-2. For investigated complexes effective viscosity in dependence from fluid distribution rate is presented in table 1 (for the mixture of Composition 1 and polymer substance used as an example).

To conclude, addition of a polymer substance to Composition 1 improves rheological properties of studied complexes. Effective viscosity reaches maximum in NaCl (10 wt.%) water-solution medium. Also, it can be observed that fluids under consideration are non-Newtonian fluids because their effective viscosities decrease with fluid distribution rate increase. The best viscosifiers for developed visco-elastic surfactants are Xanthan Gum E415 and Hydroxyethyl cellulose in a medium of 10 wt. % NaCl solution.

Further destruction of developed complexes with polymer addition was investigated at interaction with a hydrocarbon phase. This research was conducted in the same way as study of destruction of developed complexes (without any polymeric substances) at interaction with hydrocarbon phase. Xanthan Gum E415 was used as hydrocarbon phase. Kerosene was admixed to prepared complexes with ratio 75/25, 50/50, 25/75 and rigorously mixed. Obtained mixtures were left for 1 hour.

After mixing with kerosene at different ratios an aqueous part was taken from the mixture of Composition 1 (9.9 wt. %) and Xanthan Gum E415 (0.1 wt. %) in a medium of water.

<table>
<thead>
<tr>
<th>Medium</th>
<th>Solution with polymer addition in a medium of 10 wt. % NaCl solution</th>
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<tbody>
<tr>
<td>9.9 wt. % Composition 1 solution without polymer</td>
<td></td>
</tr>
<tr>
<td>Water</td>
<td></td>
</tr>
<tr>
<td>NaCl 10 wt. %</td>
<td></td>
</tr>
<tr>
<td>KCl 10 wt. %</td>
<td></td>
</tr>
<tr>
<td>CaCl₂ 10 wt. %</td>
<td></td>
</tr>
<tr>
<td>MgCl₂ 10 wt. %</td>
<td></td>
</tr>
<tr>
<td>Inhib. HCl 12 wt. %</td>
<td></td>
</tr>
</tbody>
</table>

Table 1. Sensitivity analysis of polymer effect on properties of developed complexes.
of sodium chloride (10 wt. %) solution. Rheological properties were investigated using the case of a study of obtained aqueous part by observation effective viscosity response to fluid distribution rate increase. Results are shown in Figure 4.

As it can be seen from Figure 4, after Composition 1 (9.9 wt. %) and Xanthan Gum E415 (0.1 wt. %) mixture interacts with hydrocarbon phase in the medium of sodium chloride solutions (10 wt. %), its effective viscosity decreases significantly. In this case we can conclude that micellar structure of developed complexes was destroyed.

The hydrocarbon part, obtained after the mixture of Composition 1 (9.9 wt. %) and Xanthan Gum E415 (0.1 wt. %) interacted with kerosene in a medium of sodium chloride (10 wt. %) solution, was examined for its interfacial tension on the boundary with distilled water. The results of these measurements show (Table 2) that after the surfactant composition 1 interacts with kerosene in sodium chloride solution (10 wt. %), the medium interfacial tension is about 10 times less than for kerosene TS-1. Addition of the polymer substance to surfactant enables the decrease in interfacial tension of studied complex hydrocarbon part by 35 times. Such intense decrease in interfacial tension occurs as the result of synergistic effect of interaction between polymeric substance with micellar structures. It allows to produce the flow stream in the best possible way and to the clean out the formation pay zone to the fullest extent when covered complexes are employed as components of cutting fluids.

![Fig. 4. Dependence of effective viscosity from fluid distribution rate for mixture of Composition 1 (9.9 wt. %) and Xanthan Gum E415 (0.1 wt. %) in sodium chloride solution medium and its aqueous parts after mixing with kerosene](image)

**Table 2. Interfacial tension on the boundary hydrocarbon part – distilled water**

<table>
<thead>
<tr>
<th>Examined composition</th>
<th>Interfacial tension [mH/m]</th>
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<tr>
<td>Kerosene TS-1</td>
<td>47.50</td>
</tr>
<tr>
<td>Hydrocarbon part obtained after Xanthan Gum E415 (0.1 wt. %) interaction with kerosene in sodium chloride solution (10 wt. %) medium</td>
<td>46.70</td>
</tr>
<tr>
<td>Hydrocarbon part obtained after surfactant composition 1 (9.9 wt. %) interaction with kerosene in sodium chloride solution (10 wt. %) medium</td>
<td>4.24</td>
</tr>
<tr>
<td>Hydrocarbon part obtained after surfactant composition 1 (9.9 wt. %) and Xanthan Gum E415 (0.1 wt. %) mixture interaction with kerosene in sodium chloride solution (10 wt. %) medium</td>
<td>1.38</td>
</tr>
<tr>
<td>Hydrocarbon part obtained after Xanthan Gum E415 (0.1 wt. %) interaction with kerosene in potassium chloride solution (10 wt. %) medium</td>
<td>49.80</td>
</tr>
<tr>
<td>Hydrocarbon part obtained after surfactant composition 1 (9.9 wt. %) interaction with kerosene in potassium chloride solution (10 wt. %) medium</td>
<td>4.24</td>
</tr>
<tr>
<td>Hydrocarbon part obtained after surfactant composition 1 (9.9 wt. %) and Xanthan Gum E415 (0.1 wt. %) mixture interaction with kerosene in potassium chloride solution (10 wt. %) medium</td>
<td>1.82</td>
</tr>
</tbody>
</table>

**Conclusions**

1. The effect of water-soluble polymeric substances addition (0.1 wt. %) to the developed viscoelastic surfactant compounds on their rheological and surfactant properties was studied.
2. It was found that xanthan-type exopolysaccharide and hydroxyethyl cellulose additives favoured effective viscosity increase in developed compounds and decrease in interfacial tension of the boundary hydrocarbon phase – distilled water.

3. Destruction of developed viscoelastic surfactant compounds containing polymer substance was studied in interaction with hydrocarbon phase.

4. Addition of a polymer substance enhances viscosity of developed surfactant compounds significantly and does not influence the phenomena of viscosity decrease in interaction with hydrocarbons for such compounds.

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References


