High Pressure and High Temperature Wells Cementation using Calcium Oxide Additives: Review

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Abstract: The challenges created by HPHT conditions make drilling and cementing difficult. The overwhelming challenge stands for materials used for wells operated at higher temperature and pressures in the face of producing downhole. There is need for better choice of cement for cementing operations during development and production of wells. The review method is adopted in this article considering Calcium Oxide which poses as good cement expander. In this paper, a broad overview of the fundamental understanding and use of calcium oxide as additive is considered for cementation of HPHT production wells will be discussed.

Keywords: High, Pressure, Temperature, Cement, Calcium, Oxide, Production.

1. INTRODUCTION

A. High Pressure High Temperature Wells

High pressure/high temperature (HP/HT) wells are those that are at greater temperature or pressure than most other oil wells having downhole (Shado and Ogbonna, 2011) temperature and pressure that ranges between 150 – 200°C and 70 – 140MPa, with maximum projected pore pressure of porous formation which exceeds about 0.8 psi/ft of hydrostatic gradient, or has rated working pressure in excess of about 10000 psi at their farthest depth (Arpit et al., 2019). Wells in this categories and conditions display challenges. HPHT are pushing workable limits for engineers working bottomhole and for manufacturers in fluid control components to ensure products continue to meet these challenges (Woha, Ogbonna and Boniface, 2011).

Other challenges produced by HPHT situations make drilling/completion and cementing operations difficult which should be addressed for execution of successful job and delivery of perfect job (Zhaoguang, 2012). Temperature speeds up the process of hydration in cement, hence limiting the time it stays alterable slurry in the well. This aim refers to its ability to flow inside a defined operational window. The hydration process can chemically be delayed by the inclusion of retarders. The primary objective of cementing is maximizing cement coverage. The secondary purpose are casing protection, zonal isolation which can achieved in the annular gap if the presence of cement is optimised and maximized (Boma, Babs and Islam, 2016). During cementing, cement coverage is possible only if mud is absolutely displaced from the hole. Furthermore, one of the most challenges in HPHT is the rheology of cement slurry. It is common in HPHT wells to have a narrow margin (Michael et al, 2017) between fracture and pore pressure limiting the allowable displacement rates and ECD, not to surpass the fracture gradient. The exact implications for cement slurry and its placement is the lower pumping rates due to limited Equivalent Circulating Density (ECD) which prevent achieving proper density but having demands on chemicals concentrations, especially retarders, to cause longer thickening times.

Features of High-Density Cement Slurry:

1. They are hard to mix at surface because they are dependent strictly on temperature and because of their high solid content which supports settling and/or gelling.
2. It demands a careful and conscious design taking cognisance of variations in temperatures, density and concentration of additives.
3. It must be stable and able to pump at elevated circulating temperature and at surface.
4. It must be retarded more than enough and yet be able to set at lower temperatures at the top of the cement.

Cement Slurry Design Criteria for HPHT
1. Surface conditions of 300 rpm reading and yield point lower than 35 lbs/100 ft²;
2. Condition HPHT consistometer at 185 °F and yield point higher than 10 lbs/100ft²;
3. Monitor the temperature stability of the drilling fluid using HPHT rheometer or viscometer to evaluate its rheological properties at up to 260 °C and 20,000 psi.
4. Ramp-up and down readings at different speed. Settling tendency and gelling tendencies will be indicated.
5. Condition HPHT consistometer to less than 30 Be
6. Shut-off motor and check for a consistency spike in the HPHT consistometer. Static gel strength development is indicated.
7. Free fluid to zero with a test if well is deviated
8. Static Sedimentation test, less than 5 % deviation from theoretical density;
9. Use stirred fluid loss cell and API fluid Loss that is less than 50 ml per every 30 minutes.
11. Run sensitivity tests using retarder concentration of about 0.02 gallon per second and density of about 0.3 part per gallon.
*Avoid excessive time from reaching 50 psi compressive strength to reduce a longer wait-on-cement.

Well Construction in HPHT
The HPHT condition implicates several challenges, with wells often deep and having narrow pressure operational window, undesirable long non-productive times and higher-than-planned costs. Proper cementing can provide tremendous benefits to keep positive well economics.
B. Calcium Oxide

Calcium oxide, also known as quicklime, is a chemical compound and alkaline substance with the formula CaO. Calcium oxide molecules contain one calcium cation (which holds a charge of +2) and one oxygen anion (which holds a charge of -2), i.e. Ca$^{2+}$O$^{2-}$. It is the key ingredient for the process of making cement. The tests on CaO-type expansive cements to isolate gas entry, for some years, has produced great interest in oil well cementing.
Figure 8: Chemical structure of Calcium Oxide (cnx.org)

Calcium Oxide has the following properties:
(a) It is an amorphous white solid.
(b) It is an ionic compound featuring an ionic bond between calcium and oxygen.
(c) It is soluble in water and glycerol.
(d) Its molar mass is 56.0774 g/mol.
(e) Its density is 3.34 g/cm³.
(f) It is basic in nature and forms salts when it comes in contact with an acid.
(g) It forms slaked lime in the presence of water, i.e. CaO(s) + H₂O(l) → Ca(OH)₂(aq)
(h) Its melting point is 2572°C.
(i) Its boiling point is 2850°C.
(j) It is a very stable compound and withstands high temperatures.
(k) Its standard molar entropy corresponds to 40 joules/mole kelvin.
(l) It crystallizes in a cubic crystal
(m) It emits an intense glow when heated to temperatures above 2400°C
(n) It reacts with water to release enough heat which can ignite combustible materials.
(o) It is favourable heterogeneous base catalysts
(p) It is nontoxic
(q) It has low solubility in methanol
(r) It has a medium viscosity
(s) It has a high surface tension
(t) It has a high to intermediate expansion and contraction rate
(u) It is not volatile at ceramic temperatures.
(v) It has a moderate effect on colour,
(w) It has a bleaching effect on iron oxide.
(x) It exists in the colour of kaki/tomato reds.
(y) It causes severe irritation the nose, skin and eyes.

How Calcium Oxide is Prepared
Chemical Reaction: \( \text{CaCO}_3(s) \rightarrow \text{CaO}(s) + \text{CO}_2(g) \)

Figure 9: How Calcium Oxide is prepared (youtube.com)

Calcium oxide is produced by decomposition through heat in a rotary lime kiln at high temperatures ranging between 1070°C and 1270°C (below melting point) on substances that contain calcium carbonate (CaCO₃) which forms burnt lime and carbon dioxide. This process is called calcination.

II. LITERATURE REVIEW

The term high pressure / high temperature (HPHT) was first mentioned around 1990 (Drake and Callaway, 2014; Wilson, 2006). The calcium oxide has a white color, a shape that is spherical, narrow particles size distribution with high porosity (Tangboriboon, Kunanurukaspong and Sirivat, 2012). Cement pastes which contain ceramic waste produced hydrated products of low Ca/Si ratios when combined with silica flour that sustains the compressive strength levels satisfactorily (Souza et al, 2012). A more reliable of fluid loss additive is seen in fresh water cement slurry than in salty one (Qiang, Pinhua and Wenfa , 2014).

Volumetric shrinkage of cement slurry during oilwell cementing has been a problem that is of priority (Beirute and Tragesser, 1973; Mehta, 1978; Chenevert and Shrestha, 1987). The estimated apparent activation energy of the different cements at 2 MPa varies from 38.7 kJ/mol to 41.4 kJ/mol for the temperature range of 25 °C to 40 °C, which reduces a little with increasing curing temperature and pressure (Xueyu et al, 2013). CaO plays an essential role in biologic response during the mineralization process (Estrela et al, 2012). The application of CaO-based expansive agent permits obtaining low porous cement matrix (Valeria, Jacopo and Alessandro, 2015).
III. MATERIAL AND METHODS

This paper discusses HPHT and the Calcium Oxide as the right additive for cementation in production HPHT wells. The method adopted is review of calcium oxide as a better additive candidate for cement slurry production utilized for HPHT wells.

IV. DATA AND METHODOLOGY

Considering Appah D. and Reichetseder P. (2001), using the:

Table 1: Permeabilities of 3 different CaO-cement paste which depends on temperature.

<table>
<thead>
<tr>
<th>Temperature °C</th>
<th>CaO-expanding cements 1</th>
<th>CaO-expanding cements 2</th>
<th>CaO-expanding cements 3</th>
</tr>
</thead>
<tbody>
<tr>
<td>50</td>
<td>0.08</td>
<td>0.03</td>
<td>0.05</td>
</tr>
<tr>
<td>60</td>
<td>0.2</td>
<td>0.57</td>
<td>0.17</td>
</tr>
<tr>
<td>65</td>
<td>0.23</td>
<td>0.65</td>
<td>0.18</td>
</tr>
<tr>
<td>70</td>
<td>0.2</td>
<td>0.6</td>
<td>0.19</td>
</tr>
<tr>
<td>78</td>
<td>0.17</td>
<td>0.36</td>
<td>0.19</td>
</tr>
<tr>
<td>80</td>
<td>0.1</td>
<td>0.29</td>
<td>0.2</td>
</tr>
<tr>
<td>87</td>
<td>0.18</td>
<td>0.2</td>
<td>0.23</td>
</tr>
<tr>
<td>90</td>
<td>0.2</td>
<td>0.18</td>
<td>0.3</td>
</tr>
<tr>
<td>95</td>
<td>0.78</td>
<td>0.23</td>
<td>0.4</td>
</tr>
<tr>
<td>110</td>
<td>0.78</td>
<td>1</td>
<td>1.14</td>
</tr>
</tbody>
</table>

Regression (R²)

- Permeability of 3 different CaO-cement paste which depends on temperature.
- Consistency curves of 4 different CaO-expanding cements at a temperature of 70 °C and pressure of 43.5 MPa.
- Compressive strength (MPa) curves for 3 different CaO-cement paste.

Equation

- $y = 0.0049x + 0.0282$
- $y = 0.0106x - 0.5406$
- $y = 0.0137x - 0.7716$

Figure 10: Permeability versus Temperature
Table 2: Consistency curves of 4 different CaO-expanding cements at a temperature of 70 °C and pressure of 43.5 MPa.

<table>
<thead>
<tr>
<th>Consistency (Bc)</th>
<th>Time (Hour)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CaO-expanding cements 1</td>
<td>CaO-expanding cements 2</td>
</tr>
<tr>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td>14</td>
<td>11</td>
</tr>
<tr>
<td>15</td>
<td>13</td>
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<td>16</td>
<td>16</td>
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<td>18</td>
<td>18</td>
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<tr>
<td>40</td>
<td>37</td>
</tr>
<tr>
<td>100</td>
<td>100</td>
</tr>
<tr>
<td>100</td>
<td>100</td>
</tr>
</tbody>
</table>

Regression (R²) 0.654 0.6818 0.8818 0.6773
Equation y = 106.32x - 11.347 y = 108.76x - 11.993 y = 31.688x + 6.0443 y = 109.72x - 13.446

Figure 11: Consistency versus Time

Table 3: Compressive strength (MPa) curves for 3 different CaO-cement paste.

<table>
<thead>
<tr>
<th>Compressive Strength (Mpa)</th>
<th>Temperature °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>CaO-expanding cements 1</td>
<td>40.8 45.1</td>
</tr>
<tr>
<td>CaO-expanding cements 2</td>
<td>42 43.8 41.9</td>
</tr>
<tr>
<td>CaO-expanding cements 3</td>
<td>42.8 43.8 41</td>
</tr>
<tr>
<td></td>
<td>43.3 44 40.9</td>
</tr>
<tr>
<td></td>
<td>43.9 45.2 40.7</td>
</tr>
<tr>
<td></td>
<td>45 47.4 40.3</td>
</tr>
<tr>
<td></td>
<td>45.8 48.3 40.5</td>
</tr>
<tr>
<td></td>
<td>44.5 45.5 43.1</td>
</tr>
<tr>
<td></td>
<td>43.8 43.8 44.1</td>
</tr>
<tr>
<td></td>
<td>43.7 43 44.3 98</td>
</tr>
<tr>
<td></td>
<td>43.5 42.8 44.1</td>
</tr>
<tr>
<td></td>
<td>43.4 41.4 43</td>
</tr>
</tbody>
</table>

Regression (R²) 0.4611 0.1713 0.2291
Equation y = 0.0554x + 37.727 y = -0.0433x + 47.969 y = 0.0339x + 40.83
In figure 10, the permeability of three cement pastes was considered against different temperature. The three samples selected had similar trends beginning from 50°C. Cement 1 rose to its first permeability peak of 0.22 microDarcy at 70°C which fell to 0.05 microDarcy at 80°C but later peaked straight to 0.8 microDarcy at 90°C that eventually plunged to 0.9 microDarcy at 110°C. Cement 2 rose to its first permeability peak of 0.65 microDarcy at 65°C which fell to 0.08 microDarcy at 90°C but later skyrocketed straight to 1.0 microDarcy at 110°C. Cement 3 rose to its first permeability peak of 0.60 microDarcy at 70°C which fell to 0.28 microDarcy at 90°C but later skyrocketed straight to 1.7 microDarcy at 110°C.

In figure 11, the consistency of four cement pastes were considered against time. The four samples chosen had similar trends beginning from 0Bc. Cement 1 rose proportionately up to 40Bc for 0.67 hours which further curved away till 100Bc for another 0.13 hours. Cement 2 went up as well proportionately up to 37Bc for the same length of time till 100Bc. Cement 3 followed suit up to 35Bc for same length of time till 100Bc. Cement 4 rose up to 33Bc for same length of time, but decreased to 32Bc for another 0.13 hours.

In figure 12, the compressive strength of three cement pastes were considered against temperature. The three samples used had similar trends beginning from 50°C. Cement 1 initially possessed the compressive strength of 42.5 MPa at this temperature which peaked at 43.3 MPa at 110°C. Cement 2 initially had the compressive strength of 45 MPa at this temperature which peaked at 48.3 MPa at 110°C. Cement 3 initially had the compressive strength of 42 MPa at this temperature which peaked at 43.3 MPa at 110°C. Cement 2 and 3 went below average in their cement compressive strength, but cement 1 is the best.

V. RESULTS AND DISCUSSION

The compressive strength of three cement pastes was considered against temperature. The three samples used had similar trends beginning from 50°C. Cement 1 initially possessed the compressive strength of 42.5 MPa at this temperature, which peaked at 43.3 MPa at 110°C. Cement 2 initially had the compressive strength of 45 MPa at this temperature which peaked at 48.3 MPa at 110°C. Cement 3 initially had the compressive strength of 42 MPa at this temperature, which peaked at 43.3 MPa at 110°C.

Considering correlation regression in table 4, cement 1 performed poorly in its cement permeability; cement 2 was on the average and cement 3 went above average posing as the best. Cement 1, 2 and 3 went above average in their cement consistency, but cement 3 is the best. Cement 1, 2 and 3 all went below average in their cement compressive strength, but cement 1 is the best.

Table 4: Summary of Results

<table>
<thead>
<tr>
<th>S/No.</th>
<th>Permeability Versus Temperature</th>
<th>Consistency Versus Time</th>
<th>Compressive Strength Versus Temperature</th>
</tr>
</thead>
<tbody>
<tr>
<td>Regression (R²)</td>
<td>0.0903</td>
<td>0.527</td>
<td>0.6387</td>
</tr>
<tr>
<td>Equation</td>
<td>( y = 0.0049x + 0.0106x )</td>
<td>( y = 0.0137x )</td>
<td>( y = 106.32x )</td>
</tr>
</tbody>
</table>
VI. CONCLUSION AND RECOMMENDATION

For Calcium Oxide to perform its functions as good cement expanders, the permeability should be increased to about 63%; its consistency to 88% and its compressive strength to above 46%. This will be able to meet up the challenge of high pressure and high temperature downhole cement operations.

REFERENCES

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