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(54) **RESILIENT WELL CEMENT
COMPOSITIONS AND METHODS**

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(57) **ABSTRACT**

The present invention provides improved compositions and methods for sealing pipe in a well bore. The compositions which harden into highly resilient solid masses having high strengths are basically comprised of a hydraulic cement, an aqueous rubber latex, an aqueous rubber latex stabilizing surfactant and silica hydrophobicized with silicon oil.

RESILIENT WELL CEMENT COMPOSITIONS AND METHODS

[0001] This is a continuation of application Ser. No. 09/255,301 filed on Feb. 22, 1999

BACKGROUND OF THE INVENTION

[0002] 1. Field of the Invention

[0003] The present invention relates generally to cementing subterranean wells, and more particularly, to cement compositions which set into resilient solid masses having high strength.

[0004] 2. Description of the Prior Art

[0005] Hydraulic cement compositions are commonly utilized in primary cementing operations whereby pipe strings such as casings and liners are cemented in well bores. In performing primary cementing, a hydraulic cement composition is pumped into the annular space between the walls of the well bore and the exterior surfaces of the pipe string disposed therein. The cement composition is permitted to set in the annular space thereby forming an annular sheath of hardened substantially impermeable cement therein. The cement sheath physically supports and positions the pipe string in the well bore and bonds the exterior surfaces of the pipe string to the walls of the well bore whereby the undesirable migration of fluids between zones or formations penetrated by the well bore is prevented.

[0006] The development of wells including one or more laterals to increase production has recently taken place. Such multi-lateral wells include vertical or deviated (including horizontal) principal well bores having one or more ancillary laterally extending well bores connected thereto. Drilling and completion equipment has been developed which allows multi-laterals to be drilled from a principal cased and cemented well bore. Each of the lateral well bores can include a liner cemented therein which is tied into the principal well bore. The lateral well bores can be vertical or deviated and can be drilled into predetermined producing formations or zones at any time in the productive life cycle of the well.

[0007] In both conventional single bore wells and multi-lateral wells having several bores, the cement composition utilized for cementing casing or liners in the well bores must develop high strength after setting and also have sufficient resiliency, i.e., elasticity and ductility, to resist the loss of the bonds between the pipe and formation and the cement composition. Also, the cement composition must be able to resist cracking and/or shattering as a result of pipe movements, impacts and shocks subsequently generated by drilling and other well operations. The bond loss, cracking or shattering of the set cement allows leakage of formation fluids through at least portions of the well bore or bores which can be highly detrimental.

[0008] The cement sheath in the annulus between a pipe string and the walls of a well bore often fail due to pipe movements which cause shear and compressional stresses to be exerted on the set cement. Such stress conditions are commonly the result of relatively high fluid pressures and/or temperatures inside the cemented pipe string during testing,

perforating, fluid injection or fluid production. The high internal pipe pressure and/or temperature results in the expansion of the pipe string, both radially and longitudinally, which places stresses on the cement sheath causing it to crack or causing the cement bonds between the exterior surfaces of the pipe or the well bore walls, or both, to fail which allows leakage of formation fluids, etc.

[0009] Stress conditions also result from exceedingly high pressures which occur inside the cement sheath due to the thermal expansion of fluids trapped within the cement sheath. This condition often occurs as a result of high temperature differentials created during the injection or production of high temperature fluids through the well bore, e.g., wells subjected to steam recovery or the production of hot formation fluids from high temperature formations. Typically, the pressure of the trapped fluids exceeds the collapse pressure of the cement and pipe causing leaks and bond failure. Other compressional stress conditions occur as a result of outside forces exerted on the cement sheath due to formation shifting, overburden pressures, subsidence and/or tectonic creep.

[0010] Thus, there are needs for improved well cement compositions and methods whereby after setting, the cement compositions form highly resilient solid masses which have high compressive, tensile and bond strengths sufficient to withstand the above described stresses without failure.

SUMMARY OF THE INVENTION

[0011] The present invention provides improved cement compositions and methods for sealing pipe in well bores which meet the needs described above and overcome the deficiencies of the prior art. The improved compositions of the invention are basically comprised of a hydraulic cement, an aqueous rubber latex present in an amount in the range of from about 40% to about 55% by weight of hydraulic cement in the composition, an effective amount of an aqueous rubber latex stabilizing surfactant, and silica hydrophobicized with silicon oil present in an amount in the range of from about 0.5% to about 2% by weight of the composition.

[0012] The improved methods of this invention for cementing pipe in a well bore are comprised of the following steps. A cement composition of the invention is prepared which hardens into a highly resilient solid mass having high compressive, tensile and bond strengths. The cement composition is placed in the annulus between the pipe and the walls of the well bore and then allowed to harden therein.

[0013] It is, therefore, a general object of the present invention to provide improved cement compositions which harden into resilient solid masses having high strength and methods of using such cement compositions for sealing pipe in well bores.

[0014] Other and further objects, features and advantages of the present invention will be readily apparent to those skilled in the art upon a reading of the description of preferred embodiments which follows.

DESCRIPTION OF PREFERRED EMBODIMENTS

[0015] As mentioned, the present invention provides improved compositions and methods for cementing pipe in well bores whereby the hardened cement composition is a highly resilient solid mass having high compressive, tensile and bond strengths and which effectively withstands pipe movements due to expansion, contraction, impacts, shocks or the like. The compositions of this invention are basically comprised of a hydraulic cement, an aqueous rubber latex, an aqueous rubber latex stabilizing surfactant and silica which has been hydrophobicized with silicon oil.

[0016] A more preferred composition of this invention is comprised of a hydraulic cement, an aqueous rubber latex, an aqueous rubber latex stabilizing surfactant, an epoxy resin, an epoxy resin hardening agent and porous precipitated silica which has been hydrophobicized with silicon oil.

[0017] A variety of hydraulic cements can be utilized in accordance with the present invention including those comprised of calcium, aluminum, silicon, oxygen and/or sulfur which set and harden by reaction with water. Such hydraulic cements include Portland cements, pozzolana cements, gypsum cements, high aluminum content cements, silica cements and high alkalinity cements. Portland cements or their equivalents are generally preferred for use in accordance with the present invention. Portland cements of the types defined and described in *API Specification For Materials And Testing For Well Cements*, API Specification 10, 5th Edition, dated Jul. 1, 1990 of the American Petroleum Institute are particularly suitable. Preferred API Portland cements include classes A, B, C, G and H, with API classes G and H being more preferred and class G being the most preferred.

[0018] A variety of well known rubber materials which are commercially available in aqueous latex form, i.e., aqueous dispersions or emulsions, can be utilized in accordance with the present invention. For example, natural rubber (cis-1,4-polyisoprene) and most of its modified types can be utilized. Synthetic polymers of various types can also be used including nitrile rubber, ethylene-propylene rubbers (EPM and EPDM), styrene-butadiene rubber (SBR), nitrile-butadiene rubber (NBR), butyl rubber, neoprene rubber, cis-1,4-polybutadiene rubber and blends thereof with natural rubber or styrene-butadiene rubber, high styrene resin, silicone rubber, chlorosulfonated polyethylene rubber, crosslinked polyethylene rubber, epichlorohydrin rubber, fluorocarbon rubber, fluorosilicone rubber, polyurethane rubber, polyacrylic rubber and polysulfide rubber. The aqueous latex forms of one or more of the above rubbers can be utilized with the other components of the sealing composition being added directly to the latex.

[0019] Of the various aqueous rubber latexes which can be utilized, those formed of cis-polyisoprene rubber, nitrile rubber, ethylene-propylene rubber, styrene-butadiene rubber, nitrile-butadiene rubber, butyl rubber and neoprene rubber are generally preferred.

[0020] The most preferred aqueous rubber latex for use in accordance with this invention is a styrene-butadiene copolymer latex emulsion prepared by emulsion polymerization. The aqueous phase of the emulsion is an aqueous colloidal dispersion of the styrene-butadiene copolymer. The

latex dispersion usually includes water in an amount in the range of from about 40% to about 70% by weight of the latex, and in addition to the dispersed styrene-butadiene particles, the latex often includes small quantities of an emulsifier, polymerization catalysts, chain modifying agents and the like. The weight ratio of styrene to butadiene in the latex can range from about 10%:90% to about 90%:10%.

[0021] Styrene-butadiene latexes are often commercially produced as terpolymer latexes which include up to about 3% by weight of a third monomer to assist in stabilizing the latex emulsions. The third monomer, when present, generally is anionic in character and includes a carboxylate, sulfate or sulfonate group. Other groups that may be present on the third monomer include phosphates, phosphonates or phenolics. Non-ionic groups which exhibit steric effects and which contain long ethoxylate or hydrocarbon tails can also be present.

[0022] A particularly suitable and preferred styrene-butadiene aqueous latex contains water in an amount of about 50% by weight of the latex, and the weight ratio of styrene to butadiene in the latex is about 25%:75%. A latex of this type is available from Halliburton Energy Services of Duncan, Oklahoma, under the trade designation "LATEX 2000™."

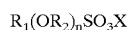
[0023] The aqueous rubber latex utilized is generally included in the cement compositions of this invention in an amount in the range of from about 40% to about 55% by weight of hydraulic cement in the compositions.

[0024] In order to prevent the aqueous latex from prematurely coagulating and increasing the viscosity of the sealing compositions, an effective amount of a rubber latex stabilizing surfactant can be included in the compositions. A suitable such surfactant has the formula



[0025] wherein R is an alkyl group having from about 5 to about 30 carbon atoms, Ph is phenyl and m is an integer in the range of from about 5 to about 50. A preferred surfactant in this group is ethoxylated nonylphenol containing in the range of from about 20 to about 30 moles of ethylene oxide.

[0026] Another latex stabilizing surfactant which can be used has the general formula



[0027] wherein R_1 is selected from the group consisting of alkyl groups having from 1 to about 30 carbon atoms, cycloalkyl groups having 5 or 6 carbon atoms, C_1 - C_4 alkyl substituted cycloalkyl groups, phenyl, alkyl substituted phenol of the general formula



[0028] wherein Ph is phenyl, R_3 is an alkyl group having from 1 to about 18 carbon atoms and a is an integer of from 1 to 3, and phenyl-alkyl groups wherein the alkyl groups have from 1 to about 18 carbon atoms and the phenyl-alkyl groups have a total of from about 8 to about 28 carbon atoms; R_2 is a substituted ethylene group of the formula



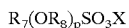
[0029] wherein R_4 is selected from hydrogen, methyl, ethyl or mixtures thereof; n is a number from 0 to about 40 provided that when R_1 is phenyl or alkyl substituted phenyl, n is at least 1; and X is any compatible cation.

[0030] Another latex stabilizing surfactant which can be utilized is a sodium salt having the general formula



[0031] wherein R_5 is an alkyl radical having in the range of from 1 to about 9 carbon atoms, R_6 is the group $-\text{CH}_2\text{CH}_2-$, o is an integer from about 10 to about 20 and X is a compatible cation.

[0032] Another surfactant which can be utilized is a sodium salt having the formula



[0033] wherein R_7 is an alkyl group having in the range of from about 5 to about 20 carbon atoms, R_8 is the group $-\text{CH}_2\text{CH}_2-$, p is an integer in the range of from about 10 to about 40 and X is a compatible cation. A preferred surfactant of this type is the sodium salt of a sulfonated compound derived by reacting a C_{12} - C_{15} alcohol with about 40 moles of ethylene oxide (hereinafter referred to as an "ethoxylated alcohol sulfonate") which is commercially available under the name "AVANEL S400™" from PPG Mazer, a division of PPG Industries, Inc. of Gurnee, Ill.

[0034] While different rubber latex stabilizers and amounts can be included in the cement compositions of this invention depending on the particular aqueous rubber latex used and other factors, the latex stabilizer is usually included in the cement compositions in an amount in the range of from about 10% to about 20% by weight of the aqueous rubber latex in the compositions.

[0035] A variety of hardenable epoxy resins can be utilized in the cement compositions of this invention. Preferred epoxy resins are those selected from the condensation products of epichlorohydrin and bisphenol A. A particularly suitable such resin is commercially available from the Shell Chemical Company under the trade designation "EPON®RESIN 828." This epoxy resin has a molecular weight of about 340 and a one gram equivalent of epoxide per about 180 to about 195 grams of resin. Another suitable epoxy resin is an epoxidized bisphenol A novolac resin which has a one gram equivalent of epoxide per about 205 grams of resin.

[0036] For ease of mixing, the epoxy resin utilized is preferably pre-dispersed in a non-ionic aqueous fluid. A non-ionic aqueous dispersion of the above described condensation product of epichlorohydrin and bisphenol A is commercially available from the Shell Chemical Company under the trade designation "EPI-REZ®-3510-W-60." Another non-ionic aqueous dispersion of an epoxy resin comprised of a condensation product of epichlorohydrin and bisphenol A having a higher molecular weight than the above described resin is also commercially available from the Shell Chemical Company under the trade designation "EPI-REZ®-3522-W-60." The above mentioned epoxidized bisphenol A novolac resin is commercially available in a non-ionic aqueous dispersion from the Shell Chemical Company under the trade designation "EPI-REZ®-5003-W-55." Of the foregoing non-ionic aqueous dispersions of epoxy resins, the aqueous dispersion of the condensation product of epichlorohydrin and bisphenol A having a molecular weight of about 340 and a one gram equivalent of epoxide per about 180 to about 195 grams of resin is the most preferred.

[0037] The epoxy resin utilized is preferably included in the cement compositions of this invention in an amount in the range of from about 5% to about 15% by weight of hydraulic cement in the compositions.

[0038] A variety of hardening agents, including, but not limited to, aliphatic amines, aliphatic tertiary amines, aromatic amines, cycloaliphatic amines, heterocyclic amines, amidoamines, polyamides, polyethyleneamines and carboxylic acid anhydrides can be utilized in the compositions of this invention containing the above described epoxy resins. Of these, aliphatic amines, aromatic amines and carboxylic acid anhydrides are the most suitable.

[0039] Examples of aliphatic and aromatic amine hardening agents are triethylenetetraamine, ethylenediamine, N-cocaoalkyltri-methylenediamine, isophoronediamine, diethyltoluenediamine, and tris(dimethylaminomethylphenol). Examples of suitable carboxylic acid anhydrides are methyltetrahydrophthalic anhydride, hexahydrophthalic anhydride, maleic anhydride, polyazelaic polyanhydride and phthalic anhydride. Of these, triethylenetetraamine, ethylenediamine, N-cocaoalkyltri-methylenediamine, isophoronediamine, diethyltoluenediamine and tris(dimethylaminomethylphenol) are preferred, with isophoronediamine, diethyltoluenediamine and tris(dimethylaminomethylphenol) being the most preferred.

[0040] The hardening agent or agents utilized are preferably included in the cement compositions of this invention in an amount in the range of from about 10% to about 30% by weight of epoxy resin in the compositions (from about 1% to about 3% by weight of hydraulic cement in the compositions).

[0041] It has been discovered that the addition of particulate silica hydrophobicized with silicon oil, i.e., polydialkylsiloxanes, to the cement compositions of this invention significantly improves the strengths of the hardened cement compositions, i.e., the compressive, tensile and shear bond strengths of the compositions. The particulate silica can be hydrophobicized by spraying it with a uniform coating of silicon oil followed by heating the sprayed silica to a temperature in the range of from about 300° F. to about 570° F. for a time period in the range of from about 1 hour to about 20 hours. Suitable commercially available silicon oils which can be utilized include the silicon oil which is commercially available under the trade designation "SWS 101™" from the Dow Corning Company or the silicon oil commercially available under the trade designation "L-45™" from the Union Carbide Corporation.

[0042] While various forms of silica can be utilized, porous precipitated silica is preferred. Porous precipitated silica can be prepared by adding sulfuric acid and a sodium silicate solution to water in a reaction vessel with high agitation. The mixture of acid, sodium silicate and water must be mixed at a high rate to prevent the formation of low pH areas where gelation will occur. Since silica dissolves to form silicate at a pH value above about 9, smaller particles are continuously dissolved during the precipitation process and therefore, uniform particle sizes are obtained. As the silica precipitation progresses, the small particles aggregate through siloxane bridges to form three dimensional networks that resist the high capillary pressure that develops during drying. After drying, the precipitated porous silica is sprayed with silicon oil as described above. The hydropho-

bicized silica is included in the cement compositions of this invention in an amount in the range of from about 0.5% to about 2% by weight of the hydraulic cement in the compositions.

[0043] A preferred composition of the present invention is comprised of a hydraulic cement, an aqueous rubber latex present in an amount in the range of from about 40% to about 55% by weight of hydraulic cement in the composition, an effective amount of an aqueous rubber latex stabilizing surfactant, and silica hydrophobicized with silicon oil present in an amount in the range of from about 0.5% to about 2% by weight of hydraulic cement in the composition.

[0044] A more preferred composition of this invention is comprised of a hydraulic cement, preferably Portland cement or the equivalent thereof; an aqueous styrene-butadiene latex which contains water in an amount of about 50% by weight of the latex and has a weight ratio of styrene to butadiene in the latex of about 25%:75%, the latex being present in an amount in the range of from about 44% to about 53% by weight of hydraulic cement in the composition; an aqueous rubber latex stabilizing surfactant comprised of an ethoxylated alcohol sulfonate present in an amount in the range of from about 10% to about 15% by weight of the aqueous rubber latex in the composition; an epoxy resin comprised of the condensation product of epichlorohydrin and bisphenol A present in an amount in the range of from about 10% to about 12% by weight of hydraulic cement in the composition; an epoxy resin hardening agent comprised of diethyltoluenediamine present in an amount in the range of from about 10% to about 20% by weight of epoxy resin in the composition (from about 1% to about 2% by weight of hydraulic cement in the composition); and porous precipitated silica hydrophobicized with silicon oil present in an amount in the range of from about 0.5% to about 1% by weight of hydraulic cement in the composition.

[0045] The improved methods of the invention for cementing pipe in a well bore are comprised of the steps of preparing a cement composition of this invention which hardens into a resilient solid mass having high strength as described above, placing the cement composition in the annulus between a pipe and the walls of a well bore and allowing the cement composition to harden therein.

[0046] In order to further illustrate the compositions and methods of the present invention, the following examples are given.

EXAMPLE

[0047] A first cement composition was prepared by combining 424.4 grams of an aqueous styrene-butadiene latex which contained 50% by weight water and had a weight ratio of styrene to butadiene of about 25%:75% with 45.6 grams of an aqueous rubber latex stabilizing surfactant comprised of an ethoxylated alcohol sulfonate. 800 grams of Premium Class G cement were added to the mixture of latex and stabilizer, and the resulting cement composition was vigorously mixed for 35 seconds after which it was cured at 140° F. for 72 hours. A second cement composition was prepared which was identical to the first composition described above except that 8 grams of porous precipitated silica hydrophobicized with silicon oil were combined with the composition. The second cement composition was also mixed for 35

seconds and cured at 140° F. for 72 hours. A third composition identical to the first composition described above was prepared except that 4 grams of the hydrophobicized silica were added to the composition. The third composition was also mixed and cured at 140° F. for 72 hours.

[0048] A fourth cement composition was prepared by combining 353.7 grams of the aqueous styrene-butadiene latex described above with 38 grams of the latex stabilizer described above. To that mixture, 800 grams of Premium Class G cement, 78.1 grams of an epoxy resin comprised of the condensation product of epichlorohydrin and bisphenol A and 10.9 grams of an epoxy resin hardening agent comprised of diethyltoluenediamine were added. The fourth cement composition was vigorously mixed for 35 seconds and then cured at 140° F. for 72 hours. A fifth cement composition identical to the above described fourth composition was prepared except that 8 grams of hydrophobicized silica was added to the composition prior to when it was mixed and cured at 140° F. for 72 hours. A sixth composition identical to the fourth composition was prepared except that 4 grams of hydrophobicized silica were added to the composition prior to when it was mixed and cured at 140° F. for 72 hours. A seventh cement composition identical to the fourth cement composition described above was prepared except that a bisphenol A novolac epoxy resin was substituted for the condensation product of epichlorohydrin and bisphenol A and 9.1 grams of diethyltoluenediamine hardening agent were included in the composition. The seventh composition was also mixed and cured at 140° F. for 72 hours. An eighth cement composition was prepared which was identical to the seventh composition described above except that 8 grams of hydrophobicized silica were added to the composition prior to when it was mixed and cured at 140° F. for 72 hours. A ninth cement composition which was also identical to the seventh composition was prepared except that 4 grams of hydrophobicized silica were added to the composition prior to when it was mixed and cured at 140° F. for 72 hours.

[0049] Cured samples of the nine cement compositions described above were used to measure the mechanical properties of the compositions. That is, the confined and unconfined compressive strengths of samples were determined in accordance with the procedure set forth in the *API Specification For Materials And Testing for Well Cements*, API Specification 10, 5th Edition, dated Jul. 1, 1990 of the American Petroleum Institute.

[0050] In addition, samples of the nine compositions were cured in the annuluses of pipe assemblies, i.e., small pipes centered inside larger pipes. The samples were cured in the pipe assemblies at 140° F. for 72 hours. After curing, the shear bond strength of each composition was determined by supporting the larger pipe and applying force to the smaller inner pipe. The shear bond strength is the total force applied divided by the bonded surface area which breaks. Additional samples of the nine cured compositions were tested for Brazilian tensile strength.

[0051] The results of these tests are given in the Table below.

TABLE

Cement Composition Strength Tests								
Composition No.	Cement ¹ ,		Epoxy	Shear	Compressive Strength, psi			
	Latex ² and Stabilizer ³	Hydrophobicized Silica	Resin and Hardening Agent ⁶	Bond Strength, psi	Unconfined	250 psi Confining Pressure	500 psi Confining Pressure	Tensile Strength, psi
1	Yes	No	No	107	587	1043	1420	100
2	Yes	Yes	No	187	653	1119	1358	104
3	Yes	Yes	No	145	645	1085	1335	112
4	Yes	No	Yes ⁴	160	878	1333	1731	107
5	Yes	Yes	Yes ⁴	125	1033	1511	1913	172
6	Yes	Yes	Yes ⁴	143	940	1461	1788	152
7	Yes	No	Yes ⁵	160	1045	1496	1804	116
8	Yes	Yes	Yes ⁵	186	894	1365	1733	165
9	Yes	Yes	Yes ⁵	186	886	1341	1678	137

¹Premium cement
²Styrene-butadiene latex (Latex 2000™)
³Ethoxylated alcohol sulfonate
⁴Condensation product of epichlorohydrin and bisphenol A
⁵Epoxidized bisphenol A novolac resin
⁶Diethyltoluenediamine

[0052] From the above Table, it can be seen that the test compositions containing hydrophobicized silica had improved compressive strengths, shear bond strengths and tensile strengths. Also, it can be seen that the compositions including both hydrophobicized silica and hardened epoxy resin had significantly higher compressive strengths, shear bond strengths and tensile strengths.

[0053] Thus, the present invention is well adapted to carry out the objects and advantages mentioned as well as those which are inherent therein. While numerous changes may be made by those skilled in the art, such changes are encompassed within the spirit of this invention as defined by the appended claims.

What is claimed is:

1. An improved method of cementing pipe in a well bore comprising the steps of:
 - (a) preparing a cement composition which hardens into a resilient solid mass having high strength comprised of a hydraulic cement, an aqueous rubber latex, an aqueous rubber latex stabilizing surfactant and silica hydrophobicized with silicon oil;
 - (b) placing said cement composition in the annulus between said pipe and the walls of said well bore; and
 - (c) allowing said cement composition to harden.
2. The method of claim 1 wherein said hydraulic cement is selected from the group consisting of Portland cements, pozzolana cements, gypsum cements, high aluminum content cements, silica cements and high alkalinity cements.
3. The method of claim 1 wherein said hydraulic cement is a Portland or equivalent cement.
4. The method of claim 1 wherein said hydraulic cement is selected from the group consisting of API Portland Classes A, B, C, G and H cements.

5. The method of claim 1 wherein said aqueous rubber latex in said cement composition is selected from the group of cis-polyisoprene rubber latex, nitrile rubber latex, ethylene-propylene rubber latex, styrene-butadiene rubber latex, nitrile-butadiene rubber latex, butyl rubber latex and neoprene rubber latex.

6. The method of claim 1 wherein said aqueous rubber latex in said cement composition is an aqueous styrene-butadiene latex.

7. The method of claim 6 wherein said aqueous styrene-butadiene latex contains water in the amount of about 50% by weight of said latex and the weight ratio of styrene to butadiene in said latex is about 25%:75%.

8. The method of claim 1 wherein said aqueous rubber latex is present in said cement composition in an amount in the range of from about 40% to about 55% by weight of said hydraulic cement in said composition.

9. The method of claim 1 wherein said aqueous rubber latex stabilizing surfactant is selected from the group consisting of ethoxylated nonylphenyl containing in the range of from about 20 to about 30 moles of ethylene oxide and an ethoxylated alcohol sulfonate.

10. The method of claim 9 wherein said aqueous rubber latex stabilizing agent is present in said composition in an amount in the range of from about 10% to about 20% by weight of said aqueous rubber latex in said composition.

11. The method of claim 1 wherein said silica hydrophobicized with silicon oil is porous precipitated silica.

12. The method of claim 12 wherein said silica hydrophobicized with silicon oil is present in said composition in an amount in the range of from about 0.5% to about 2% by weight of said hydraulic cement in said composition.

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