



(43) International Publication Date
14 May 2015 (14.05.2015)

- | | |
|--|--|
| <p>(51) International Patent Classification:
 <i>C09K 8/72</i> (2006.01) <i>E21B 43/27</i> (2006.01)</p> | <p>(71) Applicant (<i>for GB, JP, NL only</i>): SCHLUMBERGER HOLDINGS LIMITED; P.O. Box 71, Craigmuir Chambers, Road Town, Tortola, Virgin Islands, British, 1110 (VG).</p> |
| <p>(21) International Application Number:
 PCT/US2014/061015</p> | <p>(71) Applicant (<i>for all designated States except AE, AO, BH, CA, CN, FR, GB, GH, IN, JP, LY, MA, MZ, NA, NG, NL, PH, SD, SY, US, VN</i>): SCHLUMBERGER TECHNOLOGY B.V. [NL/NL]; Parkstraat 83-89m, NL-2514 JG The Hague (NL).</p> |
| <p>(22) International Filing Date:
 17 October 2014 (17.10.2014)</p> | <p>(71) Applicant (<i>for AE, AO, BF, BH, BJ, CF, CG, CI, CM, CN, GA, GH, GN, GQ, GW, IN, KM, LY, MA, ML, MR, MZ, NA, NE, NG, PH, SD, SN, SY, TD, TG, VN only</i>): PRAD RESEARCH AND DEVELOPMENT LIMITED; P.O. Box 71, Craigmuir Chambers, Road Town, Tortola, Virgin Island, British, 1110 (VG).</p> |
| <p>(25) Filing Language:
 English</p> | <p>(71) Applicant (<i>for US only</i>): SCHLUMBERGER TECHNOLOGY CORPORATION [US/US]; 300 Schlumberger Drive, Sugar Land, Texas 77478 (US).</p> |
| <p>(26) Publication Language:
 English</p> | |
| <p>(30) Priority Data:
 14/072,437 5 November 2013 (05.11.2013) US</p> | |
| <p>(71) Applicant (<i>for CA only</i>): SCHLUMBERGER CANADA LIMITED [CA/CA]; 125 - 9 Avenue SE, Calgary, Alberta T2G 0P6 (CA).</p> | |
| <p>(71) Applicant (<i>for FR only</i>): SERVICES PETROLIERS SCHLUMBERGER [FR/FR]; 42 rue Saint Dominique, F-75007 Paris (FR).</p> | |

[Continued on next page]

(54) Title: AQUEOUS SOLUTION AND METHOD FOR USE THEREOF

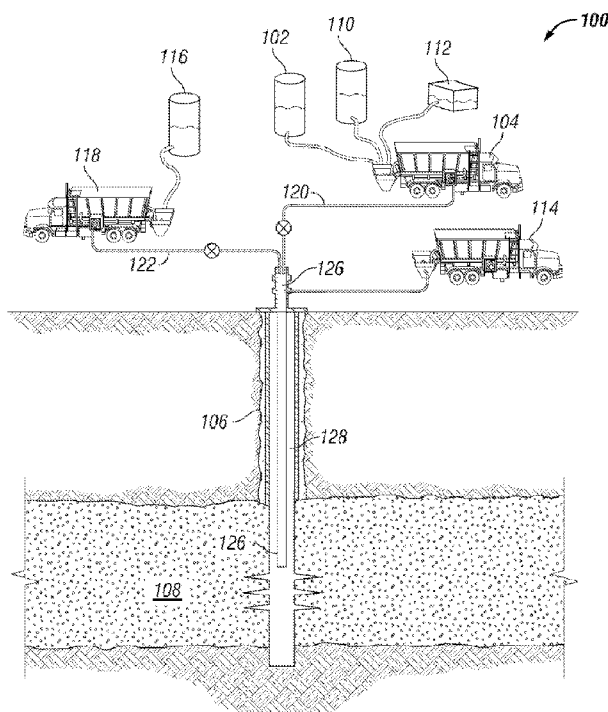


Figure 1

(57) Abstract: Oilfield treatment compositions contain water, hydrochloric acid at a concentration between 15 wt% and 45.7 wt% and a first fixing agent. The first fixing agent comprises urea, a urea derivative or both. The molar ratio between the first fixing agent and water may be higher than or equal to 0.5, and urea is the dominant solvent species in the compositions. In these compositions the first fixing agent provides corrosion inhibition when exposed to steel. The compositions may also contain a second fixing agent and an inhibitor aid.



(72) **Inventors:** **JIANG, Li**; 26014 Summer Savory Lane, Katy, Texas 77494 (US). **LECERF, Bruno**; 4601 Waycross Street, Houston, Texas 77035 (US). **JONES, Timothy G.J.**; 16 Denmark Road, Cottenham Cambridgeshire CB24 8QS (GB). **ZIAUDDIN, Murtaza**; 5215 Red Oak Valley Dr, Katy, Texas 77494 (US). **HUTCHINS, Richard**; 17507 Sunny Trail Lane, Sugar Land, Texas 77479 (US). **HE, Jian**; 5814 Bayberry Way, Sugar Land, Texas 77479 (US). **LI, Jack**; 1906 Cambria Lane, Sugar Land, Texas 77479 (US). **KRAEMER, Chad**; 26315 Cottage Springs Ct., Katy, Texas 77494 (US).

(74) **Agents:** **VANDERMOLEN, Mathieu** et al.; 10001 Richmond Avenue, IP Administration Center of Excellence, Room 4720, Houston, Texas 77042 (US).

(81) **Designated States** (*unless otherwise indicated, for every kind of national protection available*): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM,

GT, HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) **Designated States** (*unless otherwise indicated, for every kind of regional protection available*): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

Published:

— with international search report (Art. 21(3))

AQUEOUS SOLUTION AND METHOD FOR USE THEREOF**RELATED APPLICATION DATA**

[0001] None.

BACKGROUND

[0002] The statements in this section merely provide background information related to the present disclosure and may not constitute prior art.

[0003] The technical field generally, but not exclusively, relates to high-concentration hydrochloric acid (HCl) solutions with urea, and uses thereof. Previously known solutions of HCl with urea, for example as described in U.S. Patent 4,466,893, utilize urea with low HCl concentrations (at or below 15%) and in the presence of various plant-based polysaccharide gums. HCl above 15% was determined to be deleterious to the properties of previously available solutions.

SUMMARY

[0004] In an aspect, embodiments relate to compositions comprising water at a concentration lower than or equal to 20 wt%, hydrochloric acid at a concentration between 15 wt% and 45.7 wt%, and a first fixing agent comprising urea, a urea derivative or both. The first fixing agent:hydrochloric acid molar ratio is between 0.5 and 3.0 inclusive and the urea:water molar ratio or the urea derivative:water molar ratio is higher than or equal to 0.5. In these compositions, urea is the dominant solvent species of the fluid.

[0005] In a further aspect, embodiments relate to methods for inhibiting the corrosion of steel exposed to an acidic composition. A composition is prepared that comprises water at a concentration lower than or equal to 20 wt%, hydrochloric acid at a concentration between 15 wt% and 45.7 wt%, and a first fixing agent comprising urea, a urea derivative or both. The first fixing agent:hydrochloric acid molar ratio is between 0.5 and 3.0 inclusive and the urea:water molar ratio or the urea derivative:water molar ratio is higher than or equal to 0.5. In these compositions, urea is the dominant solvent species of the fluid.

[0006] In yet a further aspect, embodiments relate to methods for treating a subterranean well having a wellbore, at least one steel casing and a formation. A composition is prepared that comprises water at a concentration lower than or equal to 20 wt%, hydrochloric acid at a concentration between 15 wt% and 45.7 wt%, and a first fixing agent comprising urea, a urea derivative or both. The first fixing agent:hydrochloric acid molar ratio is between 0.5 and 3.0 inclusive and the urea:water molar ratio or the urea derivative:water molar ratio is higher than or equal to 0.5. In these compositions, urea is the dominant solvent species of the fluid. An oilfield treatment fluid that includes the composition is provided to a high-pressure pump. The high-pressure pump is operated to place the composition in the well such that the composition contacts the steel casing.

[0007] This summary is provided to introduce a selection of concepts that are further described below in the illustrative embodiments. This summary is not intended to identify key or essential features of the claimed subject matter, nor is it intended to be used as an aid in limiting the scope of the claimed subject matter. Further embodiments, forms, objects, features, advantages, aspects, and benefits shall become apparent from the following description and drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

[0008] Fig. 1 depicts example equipment to treat a wellbore and/or a formation fluidly coupled to the wellbore.

DETAILED DESCRIPTION OF SOME ILLUSTRATIVE EMBODIMENTS

[0009] For the purposes of promoting an understanding of the principles of the disclosure, reference will now be made to the embodiments illustrated in the drawings and specific language will be used to describe the same. It will nevertheless be understood that no limitation of the scope of the claimed subject matter is thereby intended, any alterations and further modifications in the illustrated embodiments, and any further applications of the principles of the application as illustrated therein as would normally occur to one skilled in the art to which the disclosure relates are contemplated herein.

[0010] At the outset, it should be noted that in the development of any such actual embodiment, numerous implementation—specific decisions must be made to achieve

the developer's specific goals, such as compliance with system related and business related constraints, which will vary from one implementation to another. Moreover, it will be appreciated that such a development effort might be complex and time consuming but would nevertheless be a routine undertaking for those of ordinary skill in the art having the benefit of this disclosure. In addition, the compositions used/disclosed herein can also comprise some components other than those cited. In the summary and this detailed description, each numerical value should be read once as modified by the term "about" (unless already expressly so modified), and then read again as not so modified unless otherwise indicated in context. Also, in the summary and this detailed description, it should be understood that a concentration range listed or described as being useful, suitable, or the like, is intended that any and every concentration within the range, including the end points, is to be considered as having been stated. For example, "a range of from 1 to 10" is to be read as indicating each and every possible number along the continuum between about 1 and about 10. Thus, even if specific data points within the range, or even no data points within the range, are explicitly identified or refer to only a few specific, it is to be understood that the Applicant appreciates and understands that any and all data points within the range are to be considered to have been specified, and that the Applicant possessed knowledge of the entire range and all points within the range.

[0011] The term "substantially no polysaccharides" as utilized herein should be understood broadly. An example solution having substantially no polysaccharides includes a solution without any polysaccharides intentionally present in the solution. Another example solution having substantially no polysaccharides includes a fluid having polysaccharides only incidentally, for example as part of an additive, and not in an amount sufficient to support development of higher viscosity in the fluid. Example amounts of polysaccharides present in a solution include less than 0.24 g/L (2 lbm/1000 gal), less than 0.12 g/L (1 lbm/1000 gal), less than 0.06 g/L (0.5 lbm/gal), less than 0.012 g/L (0.1 lbm/1000 gal) and a solution having no polysaccharides. Yet another example solution having substantially no polysaccharides includes a fluid having no detectable polysaccharides, where the detection is performed through rheological testing. Yet another example solution having substantially no polysaccharides contemplates that polysaccharides include materials such as: galactomannans such as guar gum, gum karaya, gum tragacanth, gum ghatti, gum acacia, gum konjak, shariz, locus, psyllium,

tamarind, gum tara, carrageenan, gum kauri, and modified guar such as hydroxy-propyl guar, hydroxy-ethyl guar, carboxy-methyl hydroxy-ethyl guar, and carboxy-methyl hydroxy-propyl guar.

[0012] The term "formation" as utilized herein should be understood broadly. A formation includes any underground fluidly porous formation, and can include without limitation any oil, gas, condensate, mixed hydrocarbons, paraffin, kerogen, water, and/or CO₂ accepting or providing formations. A formation can be fluidly coupled to a wellbore, which may be an injector well, a producer well, a monitoring well and/or a fluid storage well. The wellbore may penetrate the formation vertically, horizontally, in a deviated orientation, or combinations of these. The formation may include any geology, including at least a sandstone, limestone, dolomite, shale, tar sand, and/or unconsolidated formation. The wellbore may be an individual wellbore and/or a part of a set of wellbores directionally deviated from a number of close proximity surface wellbores (e.g. off a pad or rig) or single initiating wellbore that divides into multiple wellbores below the surface.

[0013] The term "oilfield treatment fluid" as utilized herein should be understood broadly. In certain embodiments, an oilfield treatment fluid includes any fluid having utility in an oilfield type application, including a gas, oil, geothermal, or injector well. In certain embodiments, an oilfield treatment fluid includes any fluid having utility in any formation or wellbore described herein. In certain embodiments, an oilfield treatment fluid includes a matrix acidizing fluid, a wellbore cleanup fluid, a pickling fluid, a near wellbore damage cleanup fluid, a surfactant treatment fluid, an unviscosified fracture fluid (e.g. slick water fracture fluid), and/or any other fluid consistent with the fluids otherwise described herein. An oilfield treatment fluid may include any type of additive known in the art, which are not listed herein for purposes of clarity of the present description, but which may include at least friction reducers, inhibitors, surfactants and/or wetting agents, fluid diverting agents, particulates, acid retarders (except where otherwise provided herein), organic acids, chelating agents, energizing agents (e.g. CO₂ or N₂), gas generating agents, solvents, emulsifying agents, flowback control agents, resins, breakers, and/or non-polysaccharide based viscosifying agents.

[0014] The term "high pressure pump" as utilized herein should be understood broadly. In certain embodiments, a high pressure pump includes a positive displacement pump that provides an oilfield relevant pumping rate – for example at least 80 L/min (0.5

bbl/min or bpm), although the specific example is not limiting. A high pressure pump includes a pump capable of pumping fluids at an oilfield relevant pressure, including at least 3.5 MPa (500 psi), at least 6.9 MPa (1,000 psi), at least 13.8 MPa (2,000 psi), at least 34.5 MPa (5,000 psi), at least 68.9 MPa (10,000 psi), up to 103.4 MPa (15,000 psi), and/or at even greater pressures. Pumps suitable for oilfield cementing, matrix acidizing, and/or hydraulic fracturing treatments are available as high pressure pumps, although other pumps may be utilized.

[0015] The term “treatment concentration” as utilized herein should be understood broadly. A treatment concentration in the context of an HCl concentration is a final concentration of the fluid before the fluid is placed in the wellbore and/or the formation for the treatment. The treatment concentration may be the mix concentration available from the HCl containing fluid at the wellsite or other location where the fluid is provided from. The treatment concentration may be modified by dilution before the treating and/or during the treating. Additionally, the treatment concentration may be modified by the provision of additives to the fluid. Example and non-limiting treatment concentrations include 7.5%, 15%, 20%, 28%, 36%, and/or up to 45.7% HCl concentration in the fluid. In certain embodiments, a treatment concentration is determined upstream of additives deliver (e.g. at a blender, hopper, or mixing tub) and the concentration change from the addition of the additives is ignored. In certain embodiments, the treatment concentration is a liquid phase or acid phase concentration of a portion of the final fluid – for example when the fluid is an energized or emulsified fluid. In certain embodiments the treatment concentration exceeds 15%. In certain embodiments, the fluid concentration exceeds 36% or exceeds 37%.

[0016] The term “urea derivative” as used herein should be understood broadly. An example urea derivative includes any urea compound having at least one of the four nitrogen bonded hydrogens substituted. The substitution products may be anything, but include at least any hydrocarbon group, and may include substitutions on one or both of the urea nitrogens. Additionally or alternatively, substitutions may include cyclic groups (e.g. ethylene urea), aromatic groups, and/or nitrogen containing hydrocarbon groups. The inclusion of a urea derivative in the present disclosure should not be read as limiting to other urea derivatives which may be used as an alternative or addition.

[0017] Applicant has determined that a first fixing agent (FA1) has utility in inhibiting corrosion of steel exposed to hydrochloric acid solutions. In an aspect, embodiments relate to compositions that comprise water at a concentration lower than or equal to 20 wt%, hydrochloric acid and a first fixing agent. The water concentration may be lower than 10 wt%. The hydrochloric acid concentration may be between 15 wt% and 45.7 wt%, or between 15 wt% and 40 wt% or between 15 wt% and 37 wt%. The FA1 comprises urea, a urea derivative or both. The molar ratio FA1:HCl may be between 0.5 and 3.0 inclusive, or between 0.75 and 2.4 or between 1.0 and 2.4. The molar ratio urea:water or urea derivative:water may be higher than or equal to 0.5 (corresponding to a mass ratio of $30/18 = 1.67$), or higher than or equal to 1.0 or higher than or equal to 1.5. Urea may therefore be considered as the dominant solvent species in these compositions. In embodiments, urea is the continuous phase.

[0018] In a further aspect, embodiments relate to methods for inhibiting the corrosion of steel exposed to an acidic composition. A composition is prepared that comprises water at a concentration lower than or equal to 20 wt%, hydrochloric acid and a first fixing agent. The water concentration may be lower than 10 wt%. The hydrochloric acid concentration may be between 15 wt% and 45.7 wt%, or between 15 wt% and 40 wt% or between 15 wt% and 37 wt%. The FA1 comprises urea, a urea derivative or both. The molar ratio FA1:HCl may be between 0.5 and 3.0 inclusive, or between 0.75 and 2.4 or between 1.0 and 2.4. The molar ratio urea:water or urea derivative:water may be higher than or equal to 0.5, or higher than or equal to 1.0 or higher than or equal to 1.5. Urea is the dominant solvent species in these compositions. The steel is then exposed to the composition. Without being held to any particular theory, Applicant believes the corrosion inhibition effect may originate from the fact that water is no longer the dominant solvent species of the treatment fluid. Further improvements are realized in that urea and urea derivatives as fixing agents are less costly and more environmentally friendly than other corrosion inhibitors known in the art.

[0019] In yet a further aspect, embodiments relate to methods for treating a subterranean well having a steel casing. A composition is prepared that comprises water at a concentration lower than or equal to 20 wt%, hydrochloric acid and a first fixing agent. The water concentration may be lower than 10 wt%. The hydrochloric acid concentration may be between 15 wt% and 45.7 wt%, or between 15 wt% and 40 wt% or between 15 wt% and 37 wt%. The FA1 comprises urea, a urea derivative or both. The

molar ratio FA1:HCl may be between 0.5 and 3.0 inclusive, or between 0.75 and 2.4 or between 1.0 and 2.4. The molar ratio urea:water or urea derivative:water may be higher than or equal to 0.5, or higher than or equal to 1.0 or higher than or equal to 1.5. Urea is the dominant solvent species in these compositions. An oilfield treatment fluid that includes the composition is provided to a high-pressure pump. The high-pressure pump is operated to place the composition in the well such that the composition contacts the steel casing.

[0020] The hydrochloric acid may be transported to a wellsite, the acid having a concentration between 28 wt% and 45.7 wt%. The acid may then be diluted to a treatment concentration before providing the oilfield treatment fluid to the high-pressure pump.

[0021] The operation of the pump may comprise at least one of (i) injecting the treatment fluid into the formation at matrix rates; (ii) injecting the treatment fluid into the formation at a pressure equal to that necessary to fracture the formation; and (iii) contacting at least one of the wellbore and the formation with the oilfield treatment fluid.

[0022] For all aspects, the urea derivatives may comprise 1,1-dimethylurea, 1,3-dimethylurea, 1,1-diethylurea, 1,3-diethylurea, 1,1-diallylurea, 1,3-diallylurea, 1,1-dipropylurea, 1,3-dipropylurea, 1,1-dibutylurea, 1,3-dibutylurea, 1,1,3,3-tetramethylurea, 1,1,3,3-tetraethylurea, 1,1,3,3-tetrapropylurea, 1,1,3,3-tetrabutylurea, ethyleneurea, propyleneurea, 1,3-dimethylpropyleneurea or 1,3-dimethylethyleneurea, or combinations thereof.

[0023] For all aspects, the compositions may further comprise a second fixing agent (FA2) that comprises a mixture of amines and alcohols. The FA2 concentration may be between 0.1 wt% and 0.5 wt% inclusive, or between 0.2 wt% and 0.5 wt% inclusive.

[0024] For all aspects, the compositions may further comprise an inhibitor aid (IA) that comprises a mixture of phenyl ketones and quaternary amines. The IA concentration may be between 0.4 wt% and 0.8 wt% inclusive, or between 0.5 wt% and 0.8 wt% inclusive.

[0025] For all aspects, the compositions may further comprise hydrofluoric acid (HF) at a concentration higher than or equal to 0.25 wt%. The HF may be present at concentrations up to 2%, up to 6%, up to 10%, up to 15%, or greater amounts. The HF

may be present in addition to the amount of HCl, and/or as a substitution for an amount of the HCl.

[0026] Referencing Fig. 1, a system **100** is depicted having example equipment to treat a wellbore **106** and/or a formation **108** fluidly coupled to the wellbore **106**. The formation **108** may be any type of formation with a bottomhole temperature up to at least 177°C (350°F). The wellbore **106** is depicted as a vertical, cased and cemented wellbore **106**, having perforations providing fluid communication between the formation **108** and the interior of the wellbore **106**. However, none of the particular features of the wellbore **106** are limiting, and the example is provided only to provide an example context **100** for a procedure.

[0027] The system **100** includes a high-pressure pump **104** having a source of an aqueous solution **102**. In a first example, the aqueous solution **102** includes a FA1 and HCl, the HCl in an amount between 5% and 45.7% inclusive, and the FA1 present in a molar ratio between 0.5 and 3.0 inclusive. The aqueous solution **102** further includes water in an amount sufficient to dissolve the HCl and the FA1, the aqueous solution **102** includes substantially no polysaccharides and water is not the dominant solvent species. The high pressure pump **104** is fluidly coupled to the wellbore **106**, through high pressure lines **120** in the example. The example system **100** includes a tubing **126** in the wellbore **106**. The tubing **126** is optional and non-limiting. In certain examples, the tubing **106** may be omitted, a coiled tubing unit (not shown) may be present, and/or the high pressure pump **104** may be fluidly coupled to the casing or annulus **128**. The tubing or casing may be made of steel.

[0028] Certain additives (not shown) may be added to the aqueous solution **102** to provide an oilfield treatment fluid. Additives may be added at a blender (not shown), at a mixing tub of the high pressure pump **104**, and/or by any other method. In certain embodiments, a second fluid **110** may be a diluting fluid, and the aqueous solution **102** combined with some amount of the second fluid **110** may make up the oilfield treatment fluid. The diluting fluid may contain no HCl, and/or HCl at a lower concentration than the aqueous solution **102**. The second fluid **110** may additionally or alternatively include any other materials to be added to the oilfield treatment fluid, including additional amounts of an FA1, or of FA2 or IA or both. In certain embodiments, an additional FA1 solution **112** is present and may be added to the aqueous solution **102** during a portion or all of the

times when the aqueous solution **102** is being utilized. The additional FA1 solution **112** may include the same or a different FA1 from the aqueous solution **102**, may include all of the FA1 for the oilfield treatment fluid, and/or may include FA1 at a distinct concentration from the aqueous solution.

[0029] The high-pressure pump **104** can treat the wellbore **106** and/or the formation **108**, for example by positioning fluid therein, by injecting the fluid into the wellbore **106**, and/or by injecting the fluid into the formation **108**. Example and non-limiting operations include any oilfield treatment without limitation. Potential fluid flows include flowing from the high-pressure pump **104** into the tubing **126**, into the formation **108**, and/or into the annulus **128**. The fluid may be recirculated out of the well before entering the formation **108**, for example utilizing a back side pump **114**. In the example, the annulus **128** is shown in fluid communication with the tubing **126**, although in certain embodiments the annulus **128** and the tubing **126** may be isolated (e.g. with a packer). Another example fluid flow includes flowing the oilfield treatment fluid into the formation at a matrix rate (e.g. a rate at which the formation is able to accept fluid flow through normal porous flow), and/or at a rate that produces a pressure exceeding a hydraulic fracturing pressure. The fluid flow into the formation may be either flowed back out of the formation, and/or flushed away from the near wellbore area with a follow up fluid. Fluid flowed to the formation may be flowed to a pit or containment (not shown), back into a fluid tank, prepared for treatment, and/or managed in any other manner known in the art. Acid remaining in the returning fluid may be recovered or neutralized.

[0030] Another example fluid flow includes the aqueous solution **102** including HCl, with FA1 being optional and in certain embodiments not present in the aqueous solution **102**. The example fluid flow includes a second aqueous solution **116** including FA1 (urea or a urea derivative). The fluid flow includes, sequentially, a first high pressure pump **104** and a second high pressure pump **118** treating the formation **108**. The second high-pressure pump **118** in the example is fluidly coupled to the tubing **126** through a second high pressure line **122**. The fluid delivery arrangement is optional and non-limiting. In certain embodiments, a single pump may deliver both the aqueous solution **102** and the second aqueous solution **116**. In the example, either the first aqueous solution **102** or the second aqueous solution **116** may be delivered first, and one or more of the solutions **102**, **116** may be delivered in multiple stages, including potentially some stages where the solutions **102**, **116** are mixed.

[0031] The schematic flow descriptions which follow provide illustrative embodiments of performing procedures for treating formations and/or wellbores. Operations illustrated are understood to be examples only, and operations may be combined or divided, and added or removed, as well as re-ordered in whole or part, unless stated explicitly to the contrary herein. Certain operations illustrated may be implemented by a computer executing a computer program product on a computer readable medium, where the computer program product comprises instructions causing the computer to execute one or more of the operations, or to issue commands to other devices to execute one or more of the operations.

[0032] Without limitation, it is contemplated the procedure includes any one of a number of specific embodiments. An example includes treating with the first oilfield treatment fluid and then the second oilfield treatment fluid, or treating with the second oilfield treatment fluid then the first oilfield treatment fluid. An example includes the first oilfield treatment fluid including no FA1, including FA1 in an amount distinct from the amount of FA1 in the second oilfield treatment fluid, and/or including FA1 in an amount that is the same or similar to the amount of FA1 in the second oilfield treatment fluid. An example includes the second oilfield treatment fluid including no HCl, including HCl in an amount distinct from the amount of HCl in the first oilfield treatment fluid, and/or including FA1 in an amount that is the same or similar to the amount of FA1 in the first oilfield treatment fluid. The first and second oilfield treatment fluids do not include both the HCl amount and the FA1 amount present in identical amounts with each other, although either one of the HCl amount or the FA1 amount may be present in identical amounts with each other. Additionally, it is contemplated that multiple stages of the first oilfield treatment fluid and/or the second oilfield treatment fluid may be performed, which stages may be equal or unequal in size or number, and/or which may include spacer fluids or not between any one or more of the stages.

[0033] As is evident from the figures and text presented above, a variety of embodiments according to the present disclosure are contemplated.

[0034] The present disclosure may be further illustrated by the following examples. These examples do not limit the scope of the disclosure.

EXAMPLES

[0035] The following examples disclose the results of corrosion tests performed with N80 steel coupons. The tests conformed to standard procedures published by the American Society for Testing and Materials (ASTM). The pitting index is a qualitative visual evaluation of the number of pits that have developed on the coupon surface. The index scale is between 0 and 5, and skilled practitioners endeavor to achieve a pitting index of at most 2. The corrosion rate is expressed in lb/ft², an oilfield unit that has no SI equivalent. In the oilfield, most practitioners limit the corrosion rate to at most 0.05 lb/ft².

[0036] Chemicals used during the tests were hydrochloric acid (HCl), and urea or ethyleneurea FA1.

EXAMPLE 1

[0037] A 21 wt% solution of HCl was prepared by passing HCl gas through to binary mixtures of predetermined urea/H₂O ratios. Corrosion tests were performed during which the HCl solution was tested alone and with urea at various urea:HCl molar ratios. The solutions were heated to 93°C (200°F) during which a steel coupon was immersed in each solution for four hours. The results, presented in Table 1, show that the presence of urea reduced the corrosion rate to less than 7% of that observed with the control solution containing acid alone. In particular, at the tested urea/H₂O molar ratios, urea was the dominant solvent species and the corrosion rate was satisfactory.

Acid Formulation			Corrosion Results	
Fluid	Urea/H ₂ O (molar ratio)	Urea/HCl (molar ratio)	Pitting Index	Corrosion Rate (lb/ft ²)
21% HCl	N/A	N/A	2	0.4873
21% HCl + Urea	1.5	1.9	0	0.0431
21% HCl + Urea	2.4	2.0	0	0.0375
21% HCl + Urea	3.0	2.1	0	0.0333

Table 1. Corrosion test results: N80 steel; 93°C; 4 h

EXAMPLE 2

[0038] A 21 wt% solution of HCl was prepared by passing HCl gas through to binary mixtures of predetermined urea (or urea derivative)/H₂O ratios. Corrosion tests were

performed with solutions containing urea or ethyleneurea (EU) at various concentrations. The solutions were heated to 135°C (275°F) during a steel coupon was immersed in the solutions for four hours.

[0039] The results, shown in Table 2, demonstrated that the presence of urea, in particular when the molar ratios of urea/H₂O were greater than unity, significantly improved corrosion inhibition. In addition the urea derivative, EU, exhibited significant corrosion inhibition abilities. Urea or EU was the dominant solvent species when present in the following solutions.

Acid Formulation			Corrosion Results	
Fluid	Urea/H ₂ O (molar ratio)	Urea/HCl (molar ratio)	Pitting Index	Corrosion Rate (lb/ft ²)
21% HCl	N/A	N/A	4	0.4530
21% HCl + Urea	1.5	1.9	1	0.0326
21% HCl + Urea	2.4	2.0	1	0.0213
21% HCl + EU	0.6	1.2	1	0.0406
21% HCl + EU	1.0	1.3	1	0.0312
21% HCl + EU	1.7	1.4	1	0.0292

Table 2. Corrosion test results: N80 steel; 135°C; 4 hr

[0040] The examples provided herein are illustrative only and do not limit the scope of the disclosure.

[0041] While the disclosure has provided specific and detailed descriptions to various embodiments, the same is to be considered as illustrative and not restrictive in character. Although only a few example embodiments have been described in detail above, those skilled in the art will readily appreciate that many modifications are possible in the example embodiments without materially departing from this invention. Accordingly, all such modifications are intended to be included within the scope of this disclosure as defined in the following claims. In the claims, means-plus-function clauses are intended to cover the structures described herein as performing the recited function and not only structural equivalents, but also equivalent structures. Thus, although a nail and a screw may not be structural equivalents in that a nail employs a cylindrical surface to secure wooden parts together, whereas a screw employs a helical surface, in the environment of fastening wooden parts, a nail and a screw may be equivalent structures.

[0042] Moreover, in reading the claims, it is intended that when words such as “a,” “an,” “at least one,” or “at least one portion” are used there is no intention to limit the claim to only one item unless specifically stated to the contrary in the claim. When the language “at least a portion” and/or “a portion” is used the item can include a portion and/or the entire item unless specifically stated to the contrary. It is the express intention of the applicant not to invoke 35 U.S.C. § 112, paragraph 6 for any limitations of any of the claims herein, except for those in which the claim expressly uses the words ‘means for’ together with an associated function.

We claim:

1. A composition, comprising:
 - (i) water at a concentration lower than or equal to 20 wt%;
 - (ii) hydrochloric acid at a concentration between 15 wt% and 45.7 wt%;and
 - (iii) a first fixing agent comprising urea, a urea derivative or both;wherein the first fixing agent:hydrochloric acid molar ratio is between 0.5 and 3.0 inclusive, the urea:water molar ratio or the urea derivative:water molar ratio is higher than or equal to 0.5, and urea is the dominant solvent species.
2. The composition of claim 1, wherein the urea derivative comprises 1,1-dimethylurea, 1,3-dimethylurea, 1,1-diethylurea, 1,3-diethylurea, 1,1-diallylurea, 1,3-diallylurea, 1,1-dipropylurea, 1,3-dipropylurea, 1,1-dibutylurea, 1,3-dibutylurea, 1,1,3,3-tetramethylurea, 1,1,3,3-tetraethylurea, 1,1,3,3-tetrapropylurea, 1,1,3,3-tetrabutylurea, ethyleneurea, propyleneurea, 1,3-dimethylpropyleneurea or 1,3-dimethylethyleneurea, or combinations thereof.
3. The composition of claim 1, further comprising a second fixing agent comprising a mixture of amines and alcohols.
4. The composition of claim 3, wherein the concentration of the second fixing agent is between 0.1 wt% and 0.5 wt% inclusive.
5. The composition of claim 3, further comprising an inhibitor aid comprising a mixture of phenyl ketones and quarternary amines.
6. The composition of claim 5, wherein the concentration of the inhibitor aid is between 0.4 wt% and 0.8 wt% inclusive.

7. The composition of claim 1, further comprising hydrofluoric acid at a concentration higher than or equal to 0.25 wt%.
8. A method for inhibiting corrosion of a steel exposed to an acidic composition, comprising:
 - (i) preparing a composition comprising water at a concentration lower than or equal to 20 wt%, hydrochloric acid at a concentration between 15 wt% and 45.7 wt% and a first fixing agent comprising urea, a urea derivative or both, wherein the first fixing agent:hydrochloric acid molar ratio is between 0.5 and 3.0 inclusive, the urea:water molar ratio or the urea derivative:water molar ratio is higher than or equal to 0.5, and urea is the dominant solvent species; and
 - (ii) exposing the steel to the composition.
9. The method of claim 8, wherein the urea derivative comprises 1,1-dimethylurea, 1,3-dimethylurea, 1,1-diethylurea, 1,3-diethylurea, 1,1-diallylurea, 1,3-diallylurea, 1,1-dipropylurea, 1,3-dipropylurea, 1,1-dibutylurea, 1,3-dibutylurea, 1,1,3,3-tetramethylurea, 1,1,3,3-tetraethylurea, 1,1,3,3-tetrapropylurea, 1,1,3,3-tetrabutylurea, ethyleneurea, propyleneurea, 1,3-dimethylpropyleneurea or 1,3-dimethylethyleneurea, or combinations thereof.
10. The method of claim 8, further comprising a second fixing agent comprising a mixture of amines and alcohols.
11. The method of claim 10, wherein the concentration of the second fixing agent is between 0.1 wt% and 0.5 wt% inclusive.

12. The method of claim 8, further comprising an inhibitor aid comprising a mixture of phenyl ketones and quaternary amines.
13. The method of claim 12, wherein the concentration of the inhibitor aid is between 0.4 wt% and 0.8 wt% inclusive.
14. The method of claim 8, further comprising hydrofluoric acid at a concentration higher than or equal to 0.25 wt%.
15. A method for treating a subterranean well having a wellbore, at least one steel casing and a formation, comprising:
 - (i) preparing a composition comprising water at a concentration lower than or equal to 20 wt%, hydrochloric acid at a concentration between 15 wt% and 45.7 wt% and a first fixing agent comprising urea, a urea derivative or both, wherein the first fixing agent:hydrochloric acid molar ratio is between 0.5 and 3.0 inclusive, the urea:water molar ratio or the urea derivative:water molar ratio is higher than or equal to 0.5, and urea is the dominant solvent species;
 - (ii) providing an oilfield treatment fluid including the composition to a high pressure pump; and
 - (iii) operating the high pressure pump to place the composition in the well such that the composition contacts the steel casing.
16. The method of claim 15, wherein the composition further comprises a second fixing agent comprising a mixture of amines and alcohols, wherein the concentration of the second fixing agent is between 0.1 wt% and 0.5 wt% inclusive.
17. The method of claim 15, wherein the composition further comprises an inhibitor aid comprising a mixture of phenyl ketones and quaternary

- amines, wherein the concentration of the inhibitor aid is between 0.4 wt% and 0.8 wt% inclusive.
18. The method of claim 15, wherein the composition further comprises hydrofluoric acid at a concentration higher than or equal to 0.25 wt%.
19. The method of claim 15, wherein the hydrochloric acid is transported to a wellsite, the acid having a concentration between 28 wt% and 45.7 wt%, and the treatment further comprises diluting the hydrochloric acid to a treatment concentration before providing the oilfield treatment fluid to the high pressure pump.
20. The method of claim 15, wherein operating the pump comprises at least one of:
- (i) injecting the treatment fluid into the formation at matrix rates;
 - (ii) injecting the treatment fluid into the formation at a pressure equal to or greater than that necessary to fracture the formation; and
 - (iii) contacting at least one of the wellbore and the formation with the oilfield treatment fluid.

1/1

DRAWINGS

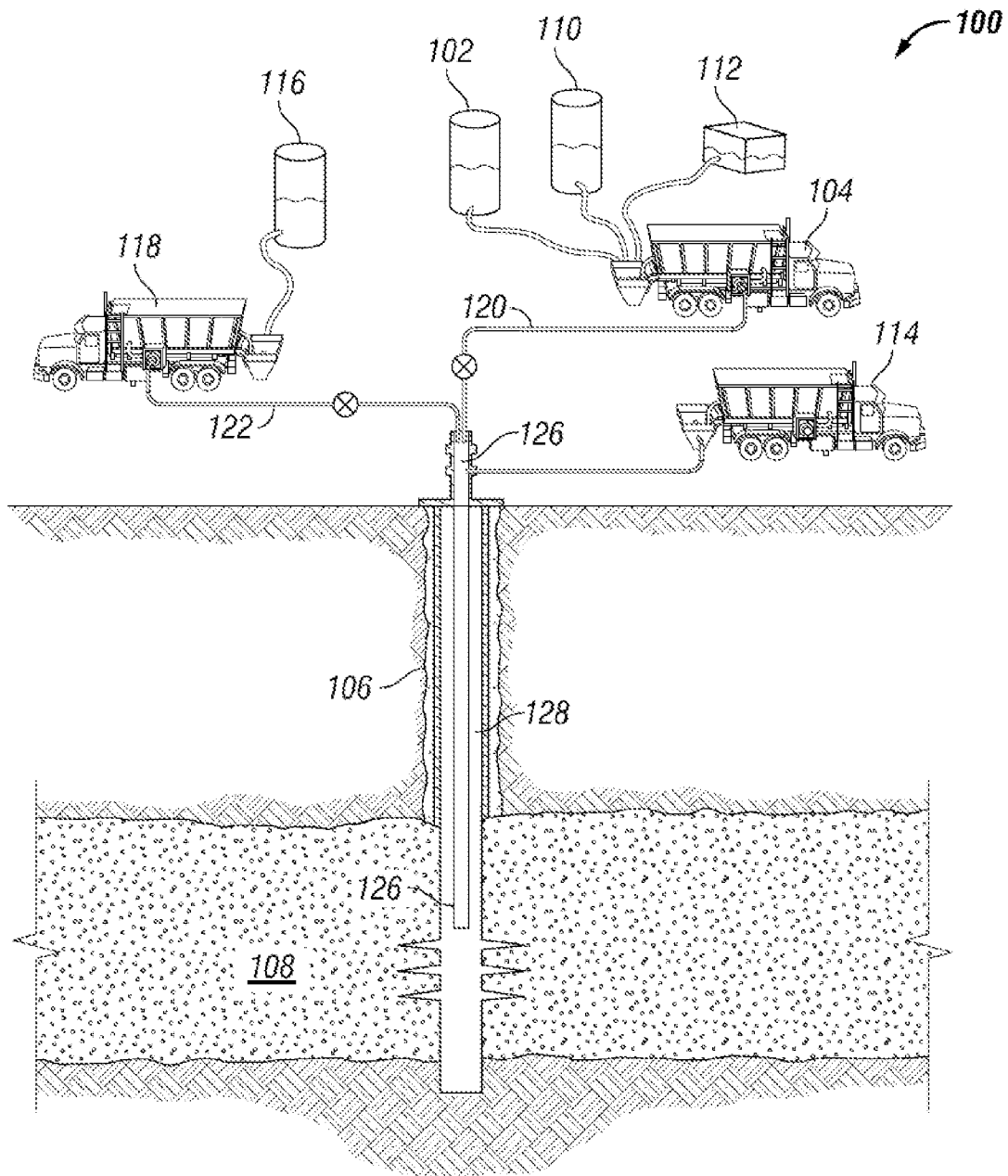


Figure 1

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2014/061015**A. CLASSIFICATION OF SUBJECT MATTER****C09K 8/72(2006.01)i, E21B 43/27(2006.01)i**

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

C09K 8/72; C23F 11/14; C11D 1/94; C11D 3/20; C23F 11/04; D06M 13/435; E21B 43/27; D06P 1/00

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Korean utility models and applications for utility models

Japanese utility models and applications for utility models

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

eKOMPASS(KIPO internal) & Keywords: water, hydrochloric acid, urea, amines, alcohols, subterranean, steel casing, corrosion inhibitor

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4466893 A (DILL, WALTER R.) 21 August 1984 See abstract; column 1; and claims 1-5 and 13-18.	1-20
A	US 5366643 A (WALKER, MICHAEL L.) 22 November 1994 See abstract; columns 3 and 4; and claims 1-12.	1-20
A	US 6340660 B1 (GASTGABER, CHARLES) 22 January 2002 See abstract; columns 2 and 3; and claims 1 and 14.	1-20
A	US 5616151 A (SARGENT, R. RICHARD et al.) 01 April 1997 See abstract; columns 2-4; and claim 1.	1-20
A	US 4631138 A (JOHNS, MARSHALL E. et al.) 23 December 1986 See abstract; column 1; and claim 1.	1-20



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

19 January 2015 (19.01.2015)

Date of mailing of the international search report

19 January 2015 (19.01.2015)

Name and mailing address of the ISA/KR

International Application Division
Korean Intellectual Property Office
189 Cheongsu-ro, Seo-gu, Daejeon Metropolitan City, 302-701,
Republic of Korea

Facsimile No. ++82 42 472 3473

Authorized officer

HEO, Joo Hyung

Telephone No. +82-42-481-8150



INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/US2014/061015

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4466893 A	21/08/1984	None	
US 5366643 A	22/11/1994	EP 489498 A1 EP 489498 B1 US 5411670 A	10/06/1992 11/05/1994 02/05/1995
US 6340660 B1	22/01/2002	None	
US 5616151 A	01/04/1997	US 2001-0045393 A1 US 2003-0038085 A1 US 5234466 A US 5672279 A US 5733463 A US 5919375 A US 7029553 B1 WO 94-02549 A1	29/11/2001 27/02/2003 10/08/1993 30/09/1997 31/03/1998 06/07/1999 18/04/2006 03/02/1994
US 4631138 A	23/12/1986	None	