

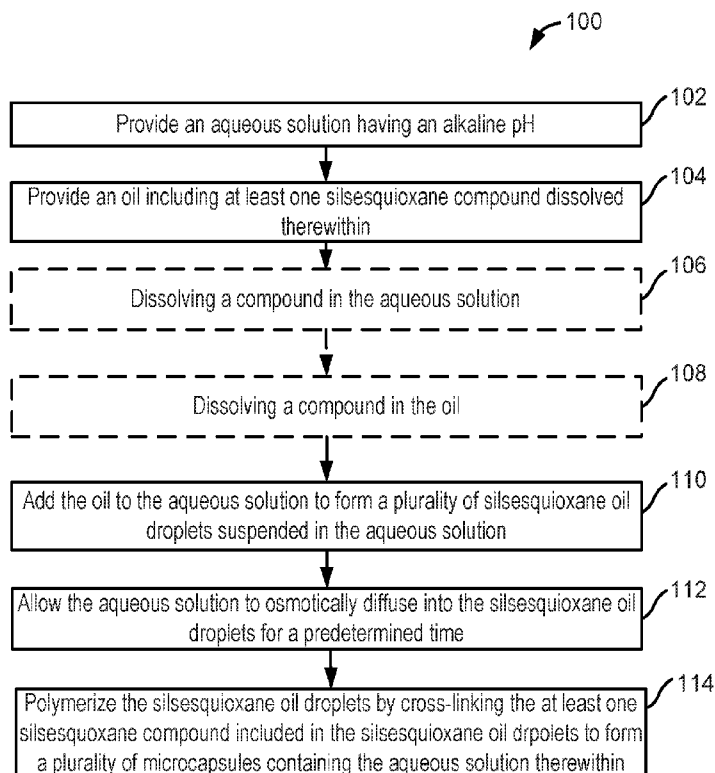


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[Continued on next page]

(54) **Title:** SELF-INFLATING MICROCAPSULES

FIG. 1



(57) **Abstract:** A method comprises providing an aqueous solution having an alkaline pH and providing an oil including at least one silsesquioxane compound. The oil is added to the aqueous solution. The oil forms a plurality of silsesquioxane oil droplets suspended in the aqueous solution such that an internal osmotic pressure is generated inside the oil droplets via a chemical reaction. The aqueous solution is allowed to osmotically diffuse into the plurality of oil droplets for a predetermined time. The silsesquioxane oil droplets are polymerized by cross-linking the at least one silsesquioxane compound included in the silsesquioxane oil droplets to form a plurality of solidified microcapsules containing the aqueous solution therein.



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Declarations under Rule 4.17:

- *as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii))*

SELF-INFLATING MICROCAPSULES

Cross-Reference to Related Patent Applications

[0001] This application claims priority from U.S. Provisional Application No. 62/196,218, filed July 23, 2015, incorporated herein by reference in its entirety.

Technical Field

[0002] The present disclosure relates generally to methods of preparing polymerized oil microcapsules.

Background

[0003] Microcapsules are colloidal containers that can house or contain drugs, enzymes, food ingredients or other materials on a micrometer scale or nanometer scale. Of particular relevance are microcapsule type micro and nanoparticles capable of encapsulating a payload and releasing the payload on demand. Such microcapsules find uses in various fields including targeted drug delivery, food science, cosmetics, biotechnology, paint industry, agriculture, environmental engineering, etc. Conventional manufacturing processes used to prepare microcapsules, microparticles or nanoparticles, are slow and cumbersome. For example, templating is often used for manufacturing such load carrying microcapsules or particles and consists of depositing material onto a sacrificial template via methods such as interfacial polymerization and layer by layer self-assembly. However, need of a template adds complexity to the synthetic process.

Summary

[0004] Embodiments described herein relate generally to methods of preparing microcapsules and in particular, to methods of preparing oil based microcapsules that can contain a solid, liquid or gas therewithin.

[0005] In some embodiments, a method comprises providing an aqueous solution having an alkaline pH and providing an oil including at least one silsesquioxane compound. The oil is then added to the aqueous solution such that the oil forms a plurality of silsesquioxane oil droplets suspended in the aqueous solution. Adding the oil to the aqueous solution generates an internal osmotic pressure inside the oil droplets by means of a chemical reaction. The aqueous solution is allowed to osmotically diffuse into the plurality of oil droplets for a

predetermined time. The oil droplets are polymerized by cross-linking at least one silsesquioxane oligomer included in the oil to form a plurality of solidified microcapsules containing the aqueous solution therewithin.

[0006] In some embodiments, a method comprises providing an aqueous solution having an alkaline pH. A solid compound is mixed with the aqueous solution to form a colloidal suspension. An oil is added to the colloidal suspension. The oil includes at least one silsesquioxane compound. The oil is allowed to undergo a hydrolysis-and-condensation reaction such that a plurality of silsesquioxane oil droplets form via a heterogeneous nucleation due to the presence of the solid compound. The colloidal suspension is allowed to diffuse into the plurality of oil droplets for a predetermined time. The oil droplets are polymerized by cross-linking the at least one silsesquioxane oligomer included in the silsesquioxane oil droplets to form a plurality of solidified microcapsules such that each of the plurality of solidified microcapsules contain at least one particle of the solid compound embedded therewithin.

[0007] In some embodiments, a method comprises providing an aqueous solution having an alkaline pH. An oil is added to the aqueous solution. The oil includes at least one silsesquioxane compound. The oil forms a plurality of silsesquioxane oil droplets suspended in the aqueous solution. Furthermore, an internal osmotic pressure is generated inside the oil droplets by means of a chemical reaction. The aqueous solution is allowed to osmotically diffuse into the plurality of silsesquioxane oil droplets for a predetermined time. The silsesquioxane oil droplets are polymerized by cross-linking the at least one silsesquioxane compound included in the silsesquioxane oil droplets to form a plurality of solidified microcapsules containing the aqueous solution therewithin. The plurality of solidified microcapsules are dried. The plurality of solidified microcapsules are exposed to a gas to infuse the gas into the plurality of solidified microcapsules. The plurality of solidified microcapsules are exposed to a hydrophobic volatile silane which seals the plurality of solidified microcapsules to trap the gas therewithin.

[0008] It should be appreciated that all combinations of the foregoing concepts and additional concepts discussed in greater detail below (provided such concepts are not mutually inconsistent) are contemplated as being part of the inventive subject matter disclosed herein. In particular, all combinations of claimed subject matter appearing at the

end of this disclosure are contemplated as being part of the inventive subject matter disclosed herein.

Brief Description of Drawings

[0009] The foregoing and other features of the present disclosure will become more fully apparent from the following description and appended claims, taken in conjunction with the accompanying drawings. Understanding that these drawings depict only several implementations in accordance with the disclosure and are therefore, not to be considered limiting of its scope, the disclosure will be described with additional specificity and detail through use of the accompanying drawings.

[0010] FIG. 1 is a schematic flow diagram of a method of preparing oil based microcapsules containing an aqueous solution therewithin, according to an embodiment.

[0011] FIG. 2 is a schematic flow diagram of another embodiment of a method of forming oil based microcapsules containing solid particles therewithin.

[0012] FIG. 3 is a schematic flow diagram of yet another embodiment of a method of forming oil based microcapsules containing a gas therewithin.

[0013] Reference is made to the accompanying drawings throughout the following detailed description. In the drawings, similar symbols typically identify similar components, unless context dictates otherwise. The illustrative implementations described in the detailed description, drawings, and claims are not meant to be limiting. Other implementations may be utilized, and other changes may be made, without departing from the spirit or scope of the subject matter presented here. It will be readily understood that the aspects of the present disclosure, as generally described herein, and illustrated in the figures, can be arranged, substituted, combined, and designed in a wide variety of different configurations, all of which are explicitly contemplated and made part of this disclosure.

Detailed Description of Various Embodiments

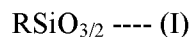
[0014] Embodiments described herein relate generally to methods of preparing microcapsules and in particular, to methods of preparing oil based microcapsules that can contain a solid, liquid, or gas therewithin.

[0015] Embodiments of the self-inflating microcapsules described herein may provide numerous benefits including, for example: (1) providing oil based droplets that can self-inflate due to osmotic diffusion by an aqueous solution into the oil to form microcapsules thus allowing rapid loading and fabrication of microcapsules; (2) allowing loading of the microcapsules with an aqueous solution which can contain dissolved compounds, solid particles or a gas; (3) protecting the load contained within the microcapsules; (4) allowing targeted release of the load contained within the microcapsules at specific pH conditions, temperature, or ionic strength; and (5) providing utility in numerous fields including targeted drug delivery, ultrasound imaging, self-healing materials, fragrance release, nutrient preservation, herbicides, insecticides, fungicides, in paints and coatings (e.g., controlled release of biocides), adhesives and sealants (e.g., encapsulation of catalysts, initiators and monomers), personal and household care products (e.g., fragrance delivery), food, nutraceuticals and pharmaceuticals (e.g., taste modifiers, nutrients, additives), etc.

[0016] FIG. 1 is a schematic flow diagram of a method 100 for preparing oil microcapsules containing or filled with an aqueous solution according to an embodiment. The method 100 includes providing an aqueous solution having an alkaline pH at 102. The aqueous solution can include, for example water (e.g., deionized water, distilled water or tap water) or a buffer solution, for example, ammonia, HEPES-buffered saline, tris-buffered saline, phosphate buffered saline (PBS), MES, PIPES, Lactated Ringer's solution, Dulbecco's minimum essential media (DMEM), Ham's F-10 media, Ham's F-12 media, minimum essential media (MEM), a cell culture medium, or any other aqueous solution. The pH of the aqueous solution is greater than the neutral pH of 7 (e.g., 8, 9, 10, 11, 12, 13 or 14).

[0017] An oil including at least one silsesquioxane compound is provided at 104. The silsesquioxane compound can include a hydrolysable silane monomer and/or a low molecular weight silsesquioxane oligomer. In various embodiments, the oil includes organofunctional alkoxysilane molecules including, but not limited to 3-(Trimethoxysilyl)propyl methacrylate, mercaptopropyl trimethoxysilane or aminopropyltrimethoxysilane

[0018] Furthermore, at least one silsesquioxane compound can include a compound of formula I:



where:

Si is silicon;

O is oxygen; and

R is hydrogen, alkyl, alkene, aryl or arylene.

[0019] Examples of silsesquioxanes include but are not limited to polyhedral oligomeric silsesquioxane (POSS) reagents such as POSS-methacrylates, POSS-Epoxyes, POSS-isocyanates, and POSS-Amines. The silsesquioxane oligomers can be synthesized using a hydrolysis and condensation reaction. The silsesquioxane oligomer can be formulated to polymerize under suitable conditions. For example, the silsesquioxane polymer can be formulated to self-polymerize, polymerize when exposed to ultraviolet (UV) light, thermoset (i.e., polymerize at increased temperature), polymerize by a condensation and/or hydrolysis reaction, or polymerize due to changes in pH.

[0020] In some embodiments, a compound can also be dissolved in the aqueous solution at 106. In various embodiments, the compound can include a fluorescent dye such as, for example ALEXAFLUOR®, 7-aminoactinomycin D, 8-anilidonaphthalene-1-sulfonic acid, ATTO dyes, auramine-rhodamine stain, benzanthrone, biman, 9,10-bis(phenylethynyl)anthracene, 5,12-bis(phenylethynyl)naphthacene, bisbenzimidazole, blacklight paint, brainbow, calcein, carboxyfluorescein, carboxyfluorescein diacetate succinimidyl ester, carboxyfluorescein succinimidyl ester, 1-chloro-9,10-bis(phenylethynyl)anthracene, 2-chloro-9,10-bis(phenylethynyl)anthracene, 2-chloro-9,10-diphenylanthracene, coumarin, DAPI, dark quencher, DiOC₆, dylight fluor, epicocconone, flash-EDT2, fluo-3, fluo-4, fluoProbes, fluoro-jade stain, FMN-binding fluorescent proteins, fura-2, fura-2-acetoxymethyl ester, green fluorescent protein, heptamethine dyes, hoechst stain, iminocoumarin indian yellow, indo-1, laurdan, lucifer yellow, luciferin, MCherry, merocyanine, Nile red, optical brightener, perylene, phloxine, phycobillin, phycoerythrin, phycoerythrobilin, pyranine, rhodamine, rhodamine 123, rhodamine 6G, ribogreen, RoGFP, rubrene, (E)-stilbene, (Z)-stilbene, sulforhodamine 101, sulforhodamine B, SYBR green I, SYBR safe, synapto-pHluorin, tetraphenyl butadiene, tetrasodium tris(bathophenanthroline

disulfonate)ruthenium(II), texas red, titan yellow, TSQ, umbelliferone, violanthrone, yellow fluorescent protein, YOYO-1 or any other suitable fluorescent dye.

[0021] In other embodiments, the compound can include a pharmaceutical or a nutraceutical. In various embodiment, the compound can include any representative classes of antibiotics, e.g., 1) aminoglycosides, such as gentamycin, kanamycin, neomycin, streptomycin or tobramycin; 2) cephalosporins, such as cefaclor, cefadroxil or cefotaxime; 3) macrolides, such as azithromycin, clarithromycin, or erythromycin; 4) penicillins, such as amoxicillin, carbenicillin or penicillin; 5) peptides, such as bacitracin, polymyxin B or vancomycin; 6) quinolones, such as ciprofloxacin, levofloxacin, or enoxacin; 7) sulfonamides, such as sulfamethazole, sulfacetimide; or sulfamethoxazole; 8) tetracyclines, such as doxycycline, minocycline or tetracycline; 8) other antibiotics with diverse mechanisms of action such as rifampin, chloramphenicol, or nitrofurantoin. Other antimicrobial agents, e.g., antifungal agents and antiviral agents can also be used.

[0022] In some embodiments, the pharmaceutical or nutraceutical can include a growth factor such as, for example, vascular endothelial cell growth factors (VEGF) (e.g., VEGF A, B, C, D, and E), platelet-derived growth factor (PDGF), insulin-like growth factor (IGF) I and IGF-II, interferons (IFN) (e.g., IFN - α , β or γ), fibroblast growth factors (FGF) (e.g., FGF 1, FGF-2, FGF-3, FGF-4-FGF-10), epidermal growth factor, keratinocyte growth factor, transforming growth factors (TGF) (e.g., TGF α or β), tumor necrosis factor- α , an interleukin (IL) (e.g., IL-1, IL-2, IL-17 -IL-18), Osterix, Hedgehogs (e.g., sonic or desert), SOX9, bone morphogenetic proteins (BMP's), in particular, BMP 2, 4, 6, and (BMP-7 is also called OP-1), parathyroid hormone, calcitonin prostaglandins, or ascorbic acid.

[0023] In other embodiments, the compound can include anti-inflammatory agents. The anti-inflammatory agents can include, for example non-steroidal anti-inflammatory drugs (NSAIDs) such as, for example, aspirin, choline and magnesium salicylates, celecoxib, diclofenac potassium, diclofenac sodium, diclofenac sodium with misoprostol, diflunisal, etodolac, fenoprofen, flurbiprofen, ibuprofen, indomethacin, ketoprofen, meclofenamate sodium, mefenamic acid, meloxicam, nabumetone, naproxen, naproxen sodium, oxaprozin, piroxicam, rofecoxib, salsalate, sodium salicylate, sulindac, tolmetin sodium, valdecoxib, any other suitable NSAID or a combination thereof. In some embodiments, the anti-inflammatory agents can include corticosteroids such as, for example, hydrocortisone, hydrocortisone acetate, cortisone acetate, tixocortol pivalate, prednisolone,

methylprednisolone, methylprednisolone aceponate, prednisone, triamcinolone acetonide, triamcinolone alcohol, mometasone, mometasone furoate, amcinonide, budesonide, desonide, fluocinonide, fluocinolone acetonide, halcinonide, betamethasone, betamethasone sodium phosphate, dexamethasone, dexamethasone sodium phosphate, fluocortolone, hydrocortisone-17-valerate, halometasone, alclometasone dipropionate, betamethasone valerate, betamethasone dipropionate, prednicarbate, clobetasone-17-butyrate, clobetasol-17-propionate, fluocortolone caproate, fluocortolone pivalate, fluprednidene acetate, hydrocortisone-17-butyrate, hydrocortisone-17-aceponate, hydrocortisone-17-buteprate, hydrocortisone valerate, flurandrenolide, triamcinolone acetonide, ciclesonide, halobetasol, diflorasone diacetate, fluocinonide, halicinonide, amcinonide, desoximetasone, fluticasone propionate, betamethasone dipropionate, desonide, alclometasone dipropionate, clobetasol propionate, prednicarbate, any other suitable corticosteroid or a combination thereof.

[0024] In various embodiments, the compound can also include a polymerizable entity such as acrylic monomers, alkene monomers, vinyl chloride, urethane, or any other polymerizable entity. Any of the compounds can diffuse into oil microcapsules such that the oil microcapsules include the aqueous solution with the compound contained therewithin.

[0025] In some embodiments, the method includes dissolving the compound in the oil at 108. In still other embodiments, a plurality of nanoparticles can be suspended in the oil. Such nanoparticles can include magnetic nanoparticles (e.g., magnetite nanoparticles, iron oxide nanoparticles), photocatalytic nanoparticles (e.g., titanium oxide nanoparticles), quantum dots, gold nanoparticles, silver nanoparticles, porous silica nanoparticles, poly(*n*-isopropylacrylamide) nanoparticles (pNIPAM) nanoparticles, etc. The nanoparticles can get incorporated in the walls of the microcapsules, as described herein. For example, the nanoparticles can include magnetic nanoparticles which are incorporated in the walls of the microcapsules allowing magnetic manipulation of the microcapsules.

[0026] The oil is added to the aqueous solution to form a plurality of silsesquioxane oil droplets suspended in the aqueous solution at 110. The hydrophobicity of the oil can urge the oil to form droplets suspended in the aqueous solution. In other words, an oil-aqueous solution microemulsion is formed. In various embodiments, the oil-aqueous solution mixture can be subjected to shaking, vibration, stirring or sonication (e.g., ultrasonication) to form the oil droplets suspended in the aqueous solution.

[0027] Adding the oil to the aqueous solution generates an internal osmotic pressure inside the oil droplets by means of a chemical reaction, for example a base catalyzed hydrolysis reaction. Suitable bases can include sodium hydroxide (NaOH), potassium hydroxide (KOH), tetramethylammonium hydroxide (TMAH) or any other suitable base. In one embodiment, the base is already present in the aqueous solution, for example is used to adjust the pH of the aqueous solution to the alkaline pH. In other embodiments, the base can be separately added to the oil aqueous solution once the oil has been added to the aqueous solution. In one embodiment, the base also includes Azobisisobutyronitrile (AIBN). AIBN can be added to base in any suitable ratio, for example base:AIBN by volume of 1:1, 1:2, 1:3, 1:4, 1:5 or 1:6.

[0028] The aqueous solution is allowed to osmotically diffuse into the plurality of silsesquioxane oil droplets for a predetermined time at 112. The osmotic pressure difference between the oil droplets and surrounding aqueous solution allows the aqueous solution to diffuse into the silsesquioxane oil droplets. This inflates the silsesquioxane oil droplets such that the silsesquioxane oil droplets are filled with a certain amount of the aqueous solution. In this manner, a solution-oil-solution (e.g., water-oil-water) double emulsion is formed.

[0029] In various implementations, the predetermined time is sufficient to allow the aqueous solution to diffuse into the silsesquioxane oil droplets until the osmotic pressure inside the silsesquioxane oil droplets is substantially equal to the osmotic pressure of the aqueous solution on the exterior of the oil droplets. In other implementations, the predetermined time to fill the silsesquioxane oil droplets can depend upon the porosity of the oil droplets. Thus, the predetermined time or filling time can be varied to control the amount of aqueous solution that diffuses into the plurality of silsesquioxane oil droplets. The time needed to inflate the droplets (loading time) and the final size of the silsesquioxane oil droplets (or the microcapsules) are controlled by changing temperature and/or the concentration of the catalyst that generates the internal osmotic pressure (for example, the base such as NaOH).

[0030] The silsesquioxane oil droplets are polymerized by cross-linking at least one silsesquioxane compound included in the silsesquioxane oil droplets to form a plurality of solidified microcapsules containing the aqueous solution therewithin at 114. The silsesquioxane polymers can be polymerized, for example by a radical polymerization, a condensation and polymerization reaction, exposure to UV light, increasing a temperature of

the aqueous solution in which the silsesquioxane oil droplets are suspended, or adjusting a pH of the aqueous solution. In various embodiments, the solidified microcapsules have a diameter in the range of 100 nm to 10 microns. The polymerization cross-links and hardens the silsesquioxane compound included in the oil vesicles yielding the solidified microcapsules which have a solid shell.

[0031] In various embodiments, the concentration and/or type of the silsesquioxane compounds (e.g., the silsesquioxane oligomers) can be chosen to provide different physical properties to the shell of the microcapsules. For example, the shell of the microcapsules can be formulated to be soft and deformable, or rigid and brittle. Furthermore, size, polydispersity, wall thickness, surface charge and/or mechanical strength of the microcapsules can be controlled by varying one or more of the polymerization rate, silsesquioxane oligomer concentration, type of silsesquioxane oligomer used, type of oil used and/or type of compound dissolved in the aqueous solution.

[0032] In some embodiments in which the aqueous solution includes the compound dissolved therein, the osmotic diffusion also allows the compound to diffuse into the oil droplets so that the silsesquioxane oil droplets contain the aqueous solution with the compound dissolved therein. In particular embodiments, the compound can also bind with the silsesquioxane compound (e.g., a silsesquioxane oligomer) included in the oil, or bind with the oil such that the compound is drawn into the oil droplets. The concentration or amount of the water or aqueous solution soluble compound builds up inside the droplets due to the lower diffusivity of the compound in the oil, thus also driving the aqueous solution (e.g., water) into the oil droplets. This can cause phase separation yielding the solution-oil-solution double emulsion microcapsules.

[0033] In some embodiments, in which the compound is included in the oil droplets, a combination of the diffusing and polymerization separates the compound from the oil such that the microcapsules contain the compound with the aqueous solution therewithin. For example, the self-inflation of the oil droplets and subsequent polymerization urges the compound to phase separate from polymerizing oil droplets and be trapped into the microcapsules. In still other embodiments in which nanoparticles are included in the oil, the polymerization traps the nanoparticles into the shell of the microcapsules.

[0034] In particular embodiments, the microcapsules can be configured to experience a change in diameters and/or porosity when exposed to a variation in pH or temperature. For example, a change in pH of a solution in which the microcapsules are suspended, from acidic to basic can urge the microcapsules to expand thereby causing an increase in the diameter of the microcapsules. The increase in diameter can increase the porosity of the shell of the microcapsules allowing the aqueous solution and, thereby the compound contained therewithin to leach out of the microcapsules. This can be used to deliver the compound to targeted location. For example, the microcapsule can be used for targeted drug delivery in cells. The microcapsules can have a first diameter within a blood stream of a patient. Once the microcapsules are phagocytized into cells, the higher pH of the cell cytoplasm can cause an increase in the diameter and porosity of the microcapsules thereby allowing the compound contained therewithin to diffuse into the cytoplasm. In other embodiments the microcapsules can also be used as contrast agent in ultrasound imaging.

[0035] In particular embodiments, the microcapsules can be removed from the aqueous solutions and heated, for example to a temperature of up to about 100 degrees Celsius to promote cross-linking of the silsesquioxane compound. This can dry the microcapsules by evaporating the aqueous solution therewithin. In such embodiments, the microcapsule can be refilled with the aqueous solution by reimmersing the microcapsules in the aqueous solution for a second predetermined time to at least partially fill the microcapsules with the aqueous solution.

[0036] In some embodiments, solid particles can also be included into the microcapsules. For example, FIG. 2 is a schematic flow diagram of an exemplary method 200 of preparing microcapsules containing solid particles therewithin. The method includes providing an aqueous solution having an alkaline pH at 202. The aqueous solution can include any of the aqueous solutions described before herein.

[0037] A solid compound is mixed with the aqueous solution to form a colloidal suspension at 204. For example, the solid can be crushed into a powder and mixed, stirred, shaken or subjected to vibration or ultrasonication to form a colloidal suspension of the solid particles suspended in the aqueous solution. The solid compound can include any compound which is insoluble in the aqueous solution. Such solid compounds can include, for example a pharmaceutical, a nutraceutical, a pigment, a dye, a pesticide, a catalyst or any other suitable solid compound.

[0038] In various embodiments, a surfactant is added to the colloidal suspension at 206. The surfactant can include, for example sodium dodecyl sulfate (SDS), cetrimonium bromide (CTAB), TWEEN® surfactants or any other surfactant. The presence of the surfactant can urge the solid compound to dewet the oil during the microcapsule formation, as described herein. The dewetting can be used to either trap particles of the solid compound within the microcapsules or expel the solid particles from the microcapsules.

[0039] An oil including at least one silsesquioxane compound is added to the colloidal suspension at 208. Any suitable silsesquioxane compound (e.g., a hydrolysable silane monomer or a low molecular weight silsesquioxane oligomer) can be used as described before herein, and mixed with any oil as described before herein. The oil is allowed to undergo a hydrolysis-and-condensation reaction so that a plurality of silsesquioxane oil droplets form in the oil via a heterogeneous nucleation reaction due to the presence of the solid compound at 210.

[0040] The colloidal suspension is allowed to diffuse into the oil for a predetermined time at 212. Diffusion of the colloidal suspension into the oil droplets also delivers particles of the solid compounds into the oil droplets. The oil droplets therefore self-inflate and are filled with particles of the colloidal suspension.

[0041] The oil droplets are polymerized by cross-linking at least one silsesquioxane oligomer included in the oil to form a plurality of solidified microcapsules containing at least one particle of the solid compound therewithin at 214. Any suitable polymerization mechanism can be used. For example, the silsesquioxane oil added to colloidal suspension forms a silsesquioxane oil emulsion which is polymerized, cross-linked or nucleated via a two-step hydrolysis and condensation reaction to form the microcapsule. Each of the plurality of microcapsules can have one or more particles of the solid compound contained therewithin.

[0042] In some embodiments, the microcapsules are prepared to contain a gas therewithin. For example, FIG. 3 is a schematic flow diagram of an exemplary method 300 for preparing microcapsules containing a gas or infused with a gas. The method includes providing an aqueous solution having an alkaline pH at 302. The aqueous solution can include any aqueous solution described before herein.

[0043] An oil including at least one silsesquioxane compound is added to the aqueous solution to form a plurality of silsesquioxane oil droplets suspended in the aqueous solution at 304. The oil and the at least one silsesquioxane compound can include any of the silsesquioxane compound and oils described before herein. The aqueous solution is allowed to osmotically diffuse into the plurality of silsesquioxane oil droplets for a predetermined time at 306. As described before herein, this urges the oil droplets to self-inflate and produce oil droplets containing the aqueous solution therewithin.

[0044] The silsesquioxane oil droplets are polymerized by cross-linking the at least one silsesquioxane compound included in the silsesquioxane oil droplets to form a plurality of solidified microcapsules containing the aqueous solution therewithin at 308. The oil droplets can be polymerized using any suitable polymerization mechanisms, for example exposure to UV light, self-polymerize, variations in temperature or a hydrolysis and condensation reaction.

[0045] The plurality of solidified microcapsules are dried at 310. For example, the plurality of solidified microcapsules can be removed from the aqueous solution using centrifugation, filtering, or any other suitable technique. The plurality of solidified microcapsules can then be dried by heating the plurality of solidified microcapsules or by vacuum drying the plurality of solidified microcapsules. The dried solidified microcapsules are exposed to a gas to infuse the gas into the microcapsules at 312. For example, the microcapsules can be infused with air, oxygen, nitrogen, carbon dioxide, an inert gas, a noble gas, a reactive gas, or any other suitable gas.

[0046] Once the plurality of microcapsules are infused and filled with the gas, the microcapsules are exposed to a hydrophobic volatile silane at 314. The silane can include, for example a fluorinated chlorosilane, or any other suitable silane which is evaporated such that the silane deposits into the pores of the microcapsules and seals the pores. Suitable examples of silanes which can be used include but are not limited to n-Octadecyldimethylchlorosilane, tridecafluorooctyl triethoxysilane, hexamethyl-disilazane or any other suitable silane. In other words, the silane acts as a chemical sealant effectively sealing a shell of the microcapsules to trap the gas therewithin. In some embodiments, the gas filled microcapsules are suspended in an aqueous solution which includes a surfactant at 316. For example, the microcapsules can be suspended in water which includes the surfactant such as, for example, PLURONICS®.

EXPERIMENTAL EXAMPLES

Preparation of the oil droplets (pre-inflation)

[0047] First, 60 μL NH_3 is added to 80 mL deionized water, and 300 μL trimethoxysilyl propyl methacrylate (TPM) oil is added to this solution under mild stirring. After one hour an emulsion consisting of monodisperse 700 nm TPM oil droplets form in the water (hereinafter “the seed emulsion”). The final size of the droplets can be finely adjusted via seeded growth. The seeded growth consist of feeding the seed emulsion with a growing solution prepared as follows: 2.5 ml TPM is added to 50 ml deionized water under mild stirring. After approximately two hours, the solution turns clear, denoting the hydrolysis, and subsequent solubilization of TPM oil.

Seeded Growth (to tune the droplets size)

[0048] The growing solution is added to the seed emulsion at a rate of one drop per second. The final size of the oil droplets is typically 1.8 μm . The emulsion is used immediately for the next step. As a grown emulsion ages, it becomes less susceptible to inflation which is attributed to additional siloxane cross-linking.

Inflating the capsules.

[0049] The grown emulsion is rapidly added in a ratio of 1:4 (v:v) to a solution of 100 mM NaOH which contains 1g/L Azobisisobutyronitrile (AIBN). The suspension is capped and shaken vigorously to distribute AIBN as much as possible.

Polymerizing the Capsules

[0050] The inflated capsules are put immediately into an oven at 100 degrees Celsius. The polymerization process is over within two hours.

[0051] As used herein, the singular forms “a”, “an” and “the” include plural referents unless the context clearly dictates otherwise.

[0052] It should be noted that the term “exemplary” as used herein to describe various embodiments is intended to indicate that such embodiments are possible examples, representations, and/or illustrations of possible embodiments (and such term is not intended to connote that such embodiments are necessarily extraordinary or superlative examples).

[0053] It is important to note that the construction and arrangement of the various exemplary embodiments are illustrative only. Although only a few embodiments have been described in detail in this disclosure, those skilled in the art who review this disclosure will readily appreciate that many modifications are possible (e.g., variations in sizes, dimensions, structures, shapes and proportions of the various elements, values of parameters, mounting arrangements, use of materials, colors, orientations, etc.) without materially departing from the novel teachings and advantages of the subject matter described herein. Other substitutions, modifications, changes and omissions may also be made in the design, operating conditions and arrangement of the various exemplary embodiments without departing from the scope of the present invention.

[0054] Reference throughout this specification to "one embodiment," "an embodiment," or similar language means that a particular feature, structure, or characteristic described in connection with the embodiment is included in at least one embodiment of the present disclosure. Appearances of the phrases "in one embodiment," "in an embodiment," and similar language throughout this specification may, but do not necessarily, all refer to the same embodiment. Similarly, the use of the term "implementation" means an implementation having a particular feature, structure, or characteristic described in connection with one or more embodiments of the present disclosure, however, absent an express correlation to indicate otherwise, an implementation may be associated with one or more embodiments.

[0055] While this specification contains many specific implementation details, these should not be construed as limitations on the scope of any inventions or of what may be claimed, but rather as descriptions of features specific to particular implementations of particular inventions. Certain features described in this specification in the context of separate implementations can also be implemented in combination in a single implementation. Conversely, various features described in the context of a single implementation can also be implemented in multiple implementations separately or in any suitable subcombination. Moreover, although features may be described above as acting in certain combinations and even initially claimed as such, one or more features from a claimed combination can in some cases be excised from the combination, and the claimed combination may be directed to a subcombination or variation of a subcombination.

[0056] Thus, particular implementations of the invention have been described. Other implementations are within the scope of the following claims. In some cases, the actions recited in the claims can be performed in a different order and still achieve desirable results. In addition, the processes depicted in the accompanying figures do not necessarily require the particular order shown, or sequential order, to achieve desirable results. In certain implementations, multitasking and parallel processing may be advantageous.

WHAT IS CLAIMED IS:

1. A method, comprising:
 - providing an aqueous solution having an alkaline pH;
 - providing an oil including at least one silsesquioxane compound;
 - adding the oil to the aqueous solution, the oil forming a plurality of silsesquioxane oil droplets suspended in the aqueous solution, the adding generating an internal osmotic pressure inside the oil droplets by means of a chemical reaction;
 - allowing the aqueous solution to osmotically diffuse into the plurality of silsesquioxane oil droplets for a predetermined time; and
 - polymerizing the silsesquioxane oil droplets by cross-linking the at least one silsesquioxane compound included in the silsesquioxane oil droplets to form a plurality of solidified microcapsules containing the aqueous solution therewithin.
2. The method of claim 1, wherein the silsesquioxane compound includes at least one of a hydrolysable silane monomer and a low molecular weight silsesquioxane oligomer.
3. The method of claim 2, wherein the at least one silsesquioxane compound includes a compound of formula I:

$$\text{RSiO}_{3/2} \text{ ---- (I)}$$
 where:
 - Si is silicon;
 - O is oxygen; and
 - R is hydrogen, alkyl, alkene, aryl or arylene.
4. The method of claim 1, wherein allowing the aqueous solution to osmotically diffuse into the oil droplets inflates the oil droplets.
5. The method of claim 1, wherein the chemical reaction includes a base catalyzed hydrolysis.
6. The method of claim 1, further comprising:
 - dissolving a compound in the aqueous solution,
 - wherein the diffusing urges the compound to also diffuse into the oil droplets such that the plurality of microcapsules contain the compound therewithin with the aqueous solution.

7. The method of claim 6, wherein the compound includes at least one of a fluorescent dye, a pharmaceutical, a nutraceutical, and a polymerizable moiety.
8. The method of claim 1, wherein the microcapsules have a diameter in the range of 100 nm to 10 microns.
9. The method of claim 1, further comprising:
dissolving a compound into the oil,
wherein a combination of the diffusing and polymerization separates the compound from the oil such that the microcapsules contain the compound with the aqueous solution contained therewithin.
10. The method of claim 1, further comprising:
suspending nanoparticles in the oil,
wherein the polymerization traps the nanoparticles into a shell of the plurality of microcapsules.
11. A method, comprising:
providing an aqueous solution having an alkaline pH;
mixing a solid compound with the aqueous solution to form a colloidal suspension;
adding an oil to the colloidal suspension, the oil including at least one silsesquioxane compound;
allowing the oil to undergo a hydrolysis-and-condensation reaction, a plurality of silsesquioxane oil droplets forming in the oil via a heterogeneous nucleation reaction due to the presence of the solid compound;
allowing the colloidal suspension to diffuse into the plurality of silsesquioxane oil droplets for a predetermined time; and
polymerizing the oil droplets by cross-linking the at least one silsesquioxane compound included in the silsesquioxane oil droplets to form a plurality of solidified microcapsules,
wherein the each of the plurality of solidified microcapsules contain at least one particle of the solid compound therewithin.
12. The method of claim 11, wherein the silsesquioxane compound includes at least one of a hydrolysable silane monomer and a low molecular weight silsesquioxane oligomer.

13. The method of claim 11, further comprising:
adding a surfactant to the colloidal suspension,
wherein the solid compound dewets the oil droplets during the polymerization.
14. The method of claim 9, wherein the solid compound includes at least one of
quantum nanodots, photocatalytic nanoparticles, magnetic nanoparticles, porous silica
nanoparticles and poly(n-isopropylacrylamide) nanoparticles.
15. A method, comprising:
providing an aqueous solution having an alkaline pH;
adding an oil to the aqueous solution, the oil including at least one silsesquioxane
compound, the oil forming a plurality of silsesquioxane oil droplets suspended in the
aqueous solution, the adding generating an internal osmotic pressure inside the oil droplets
by means of a chemical reaction;
allowing the aqueous solution to osmotically diffuse into the plurality of
silsesquioxane oil droplets for a predetermined time;
polymerizing the silsesquioxane oil droplets by cross-linking the at least one
silsesquioxane compound included in the silsesquioxane oil droplets to form a plurality of
solidified microcapsules containing the aqueous solution therewithin;
drying the plurality of solidified microcapsules;
exposing the plurality of solidified microcapsules to a gas to infuse the gas into the
plurality of solidified microcapsules; and
exposing the plurality of solidified microcapsules to a hydrophobic volatile silane,
wherein the silane seals the plurality of solidified microcapsules to trap the gas
therewithin.
16. The method of claim 11, further comprising:
suspending the plurality of solidified microcapsules in an aqueous solution including
a surfactant.

FIG. 1

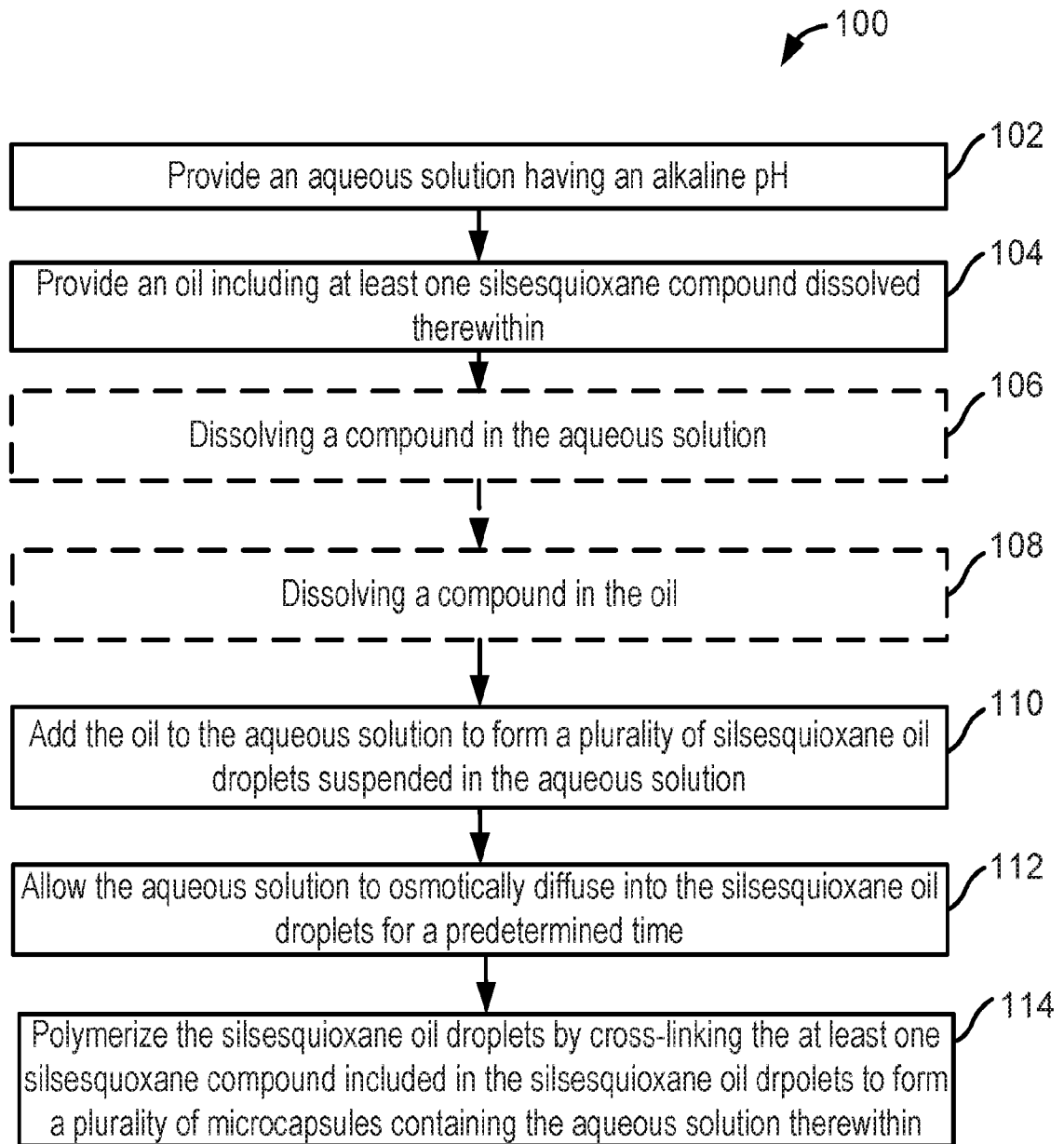


FIG. 2

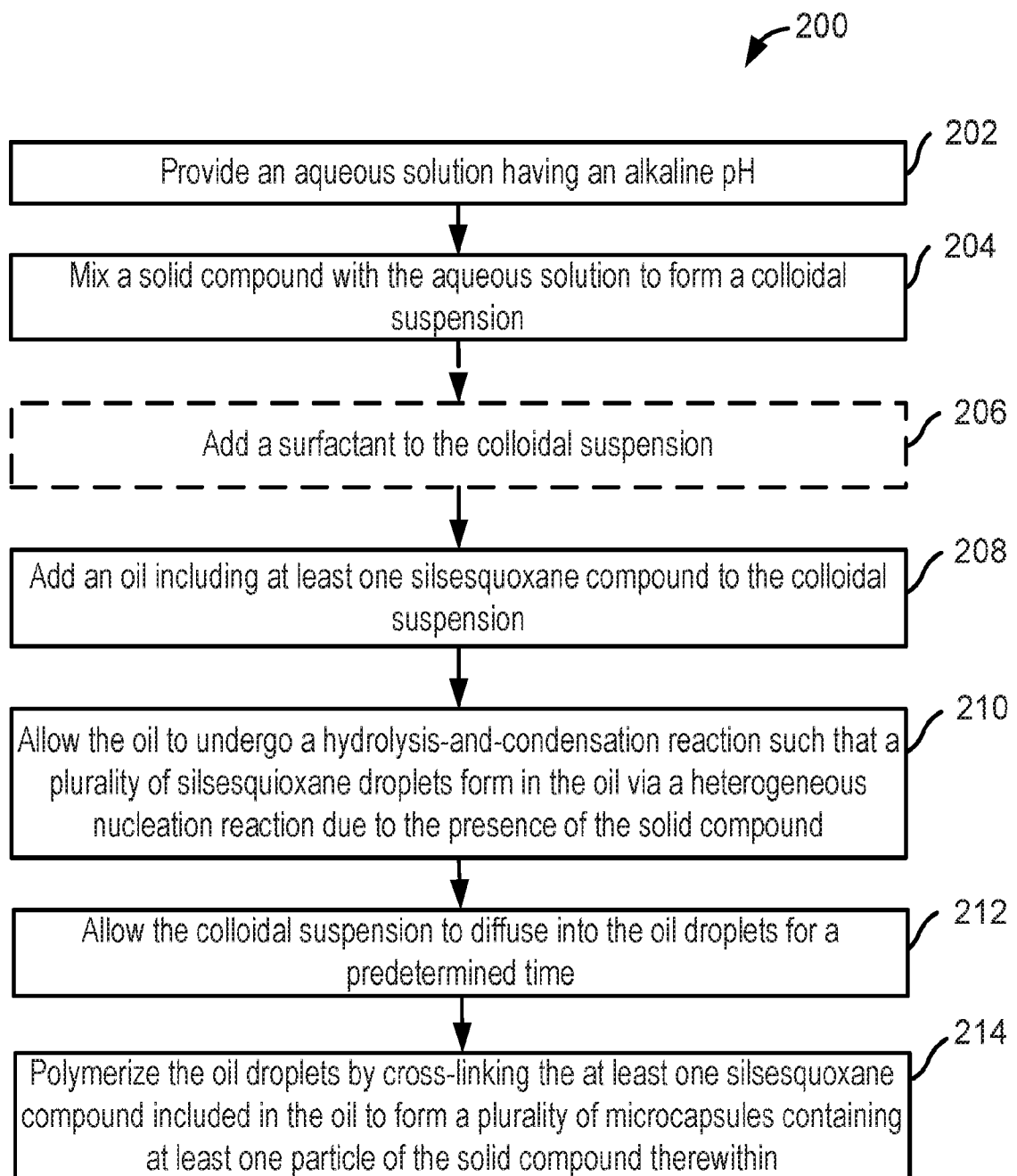
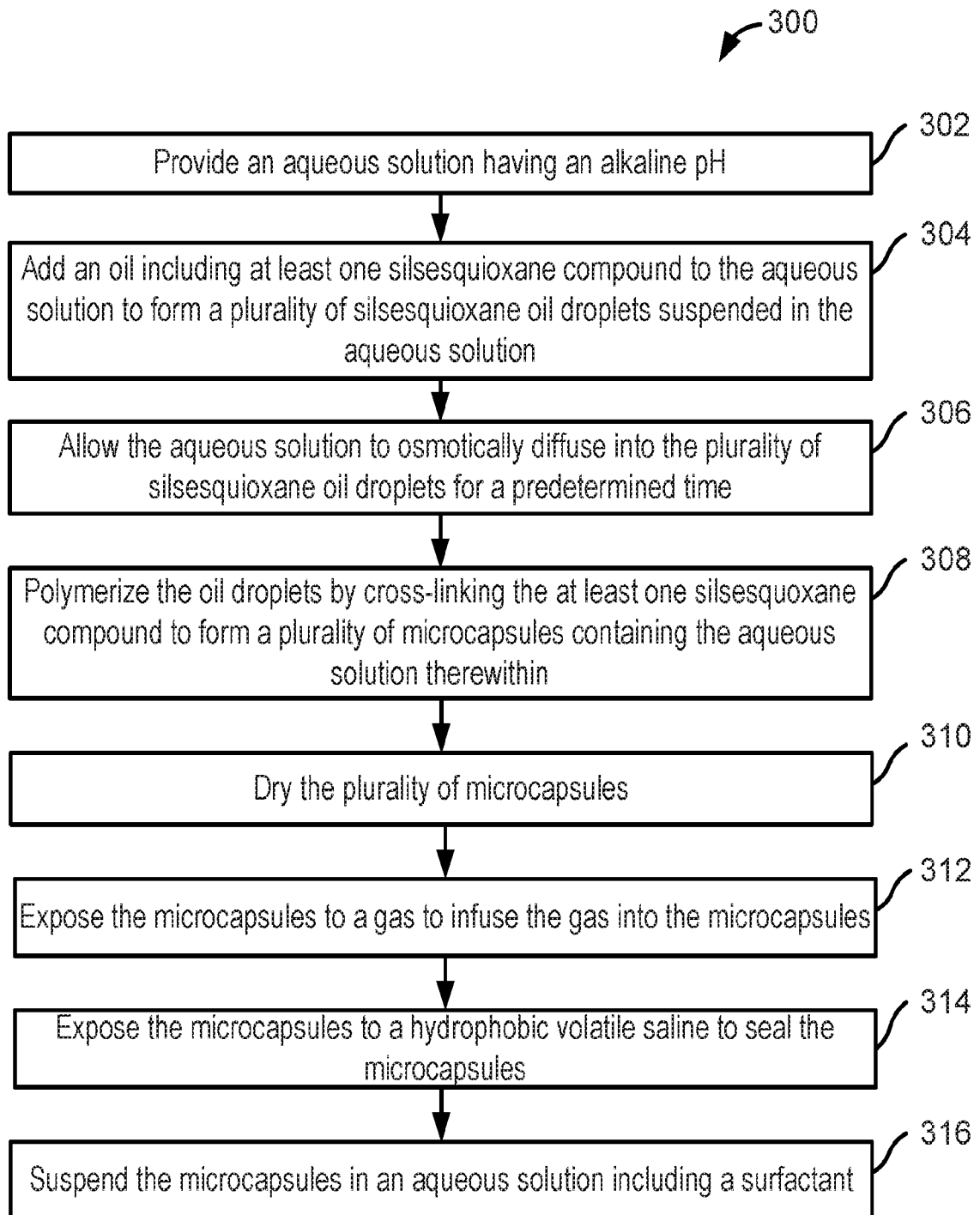


FIG. 3



INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 16/43699

A. CLASSIFICATION OF SUBJECT MATTER

IPC(8) - A61K 8/11; A61Q 19/00; B01J 13/18 (2016.01)

CPC - A61Q19/00; A61Q19/08; A61K2800/95; A61K8/11; A61Q17/04; A61K8/89; B01J13/18; B01J13/16

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)

IPC(8): A61K 8/11; A61Q 19/00; B01J 13/18 (2016.01)

CPC: A61Q19/00; A61Q19/08; A61K2800/95; A61K8/11; A61Q17/04; A61K8/89; B01J13/18; B01J13/16

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
USPC: 424/401; 512/4 (key word limited; see search terms below)

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

PatBase, Google Patents, Google Scholar

Search terms used: polysilsesquioxane microcapsules, encapsulated aqueous phase, in situ polymerization, silsesquioxane oil droplets, alkaline pH

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 2014/110148 A1 (NEW YORK UNIVERSITY) 17 July 2014 (17.07.2014); para [0007], [0011], [0024], [0036], [0041], [0043], [0062]; Fig. 7A	1-16
Y	US 5,611,971 A (MAEDERA et al.) 18 March 1997 (18.03.1997); col 1 ln 66-67; col 2 ln 10-12; col 6 ln 62-64; col 7 ln 62-63; col 8 ln 43-45; col 9 ln 56-58	1-10, 14-15
Y	US 6,337,089 B1 (YOSHIOKA et al.) 08 January 2002 (08.01.2002); col 7 ln 44-58; col 10 ln 52-54	5, 11-13, 16
Y	US 2011/0200654 A1 (HABAR) 18 August 2011 (18.08.2011); para [0001], [0019], [0030]-[0034]	2-3, 12
Y	US 2002/0082699 A1 (WARD et al.) 27 June 2002 (27.06.2002); para [0002], [0044]	4
Y	US 2008/0171077 A1 (GRAY) 17 July 2008 (17.07.2008); para [0015], [0018], [0031], [0076]-[0078]	9, 14
Y	US 2006/0071357 A1 (PILON et al.) 06 April 2006 (06.04.2006); para [0041]-[0042], [0057], [0066]	10
Y	US 2013/0064862 A1 (WEITZ et al.) 14 March 2013 (14.03.2013); para [0044], [0049], [0051], [0056], [0071], [0086]	13
A	US 2008/0175918 A1 (LAULICHT) 24 July 2008 (24.07.2008); entire document	1-16

☐ Further documents are listed in the continuation of Box C.


* Special categories of cited documents:

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

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"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

18 September 2016 (18.09.2016)

Date of mailing of the international search report

06 OCT 2016

Name and mailing address of the ISA/US

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