

METHOD FOR MAKING SILICONE HYDROGEL CONTACT LENSES

The present invention is related to a method for making silicone hydrogel (SiHy) contact lenses.

BACKGROUND

Soft contact lenses are widely used for correcting many different types of vision deficiencies. Most of soft contact lenses in the market are manufactured by cast-molding. In the cast-molding method, a polymerizable composition (or lens formulation) in a mold is polymerized to obtain a molded polymer article which is in turn extracted and hydrated to form the final hydrated soft contact lens. The molded polymer article has a shape and size similar to the shape of size of the final hydrated soft contact lenses, except that allowance must be made for swelling of the soft contact lens during hydration of the molded polymer article. In order to achieve the prescribed optical correction, soft contact lenses must be designed and then produced to meet demanding dimensional requirements. The dimensions and properties can be varied by changes in polymerization rate or conditions, in the concentrations of polymerizable components in the lens formulation, and in processing conditions. Manufacturers have to establish rigorous quality control programs for lens design process, the quality of raw materials, processing conditions, etc., and to continuously evaluate and revise their operating procedures to minimize the incidences of defective lenses.

Accordingly, there is still a need for a lens manufacturing process according to which soft contact lenses can be produced to have a consistent, targeted lens dimension.

SUMMARY OF THE INVENTION

In accomplishing the foregoing, the present invention provides a method for producing hydrated soft contact lenses. The method comprises the steps of: (1) obtaining a fluid polymerizable composition, wherein the composition comprises polymerizable materials and an organic solvent, wherein the polymerizable materials comprises at least one silicone-containing vinylic monomer or crosslinker and at least one hydrophilic vinylic monomer, wherein the organic solvent has a boiling point of at least 95°C or higher and is present in the polymerizable composition in an amount of from about 4.0% to about 15% by weight relative to the total amount of the fluid polymerizable composition and is capable of functioning as a processing-aid tool for controlling lens diameter of a hydrated soft contact lens made from the polymerizable composition; (2) introducing the fluid polymerizable composition into a cavity formed by a mold, wherein the mold has a first mold half with a first molding surface defining the anterior surface of a contact lens and a

second mold half with a second molding surface defining the posterior surface of the contact lens, wherein said first and second mold halves are configured to receive each other such that a cavity is formed between said first and second molding surfaces; (3) curing thermally or actinically the fluid polymerizable composition in the mold to form a dry contact lens; (4) extracting the dry contact lens with a liquid extraction medium to remove the organic solvent, unpolymerized polymerizable components and oligomers from the dry lens to obtain an extracted contact lens; (5) hydrating the extracted contact lens with water or an aqueous solution to obtain the hydrated soft contact lens, wherein the amount of the organic solvent is adjusted to achieve a targeted lens diameter of the hydrated soft contact lens without causing a change of about 1.7% or less in equilibrium water content.

BRIEF DESCRIPTION OF DRAWINGS

Figure 1 illustrates the general dimensions of a contact lens.

Figure 2 shows the changes of lens diameter and equilibrium water content as function of solvent concentration (weight percent).

DETAILED DESCRIPTION OF EMBODIMENTS OF THE INVENTION

Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this invention belongs. Generally, the nomenclature used herein and the laboratory procedures are well known and commonly employed in the art. Conventional methods are used for these procedures, such as those provided in the art and various general references. Where a term is provided in the singular, the inventors also contemplate the plural of that term. The nomenclature used herein and the laboratory procedures described below are those well-known and commonly employed in the art. Also, as used in the specification including the appended claims, reference to singular forms such as "a," "an," and "the" include the plural, and reference to a particular numerical value includes at least that particular value, unless the context clearly dictates otherwise. "About" as used herein means that a number referred to as "about" comprises the recited number plus or minus 1-10% of that recited number.

"Contact Lens" refers to a structure that can be placed on or within a wearer's eye. A contact lens can correct, improve, or alter a user's eyesight, but that need not be the case. A contact lens can be of any appropriate material known in the art or later developed, and can be a soft lens, a hard lens, or a hybrid lens. A "silicone hydrogel contact lens" or "SiHy contact lens" refers to a contact lens comprising a silicone hydrogel bulk (core) material.

A “soft contact lens” refers to a contact lens which has an elastic modulus (i.e., Young’s modulus) of less than 2.5 MPa.

A “hydrogel” or “hydrogel material” refers to a crosslinked polymeric material which has three-dimensional polymer networks (i.e., polymer matrix), is insoluble in water, but can hold at least 10 percent by weight of water in its polymer matrix when it is fully hydrated.

A “silicone hydrogel” or “SiHy” refers to a silicone-containing hydrogel obtained by copolymerization of a polymerizable composition comprising at least one silicone-containing monomer or at least one silicone-containing macromer or at least one crosslinkable silicone-containing prepolymer.

As used in this application, the term “non-silicone hydrogel” refers to a hydrogel that is theoretically free of silicon.

“Hydrophilic,” as used herein, describes a material or portion thereof that will more readily associate with water than with lipids.

A “vinyl monomer” refers to a compound that has one sole ethylenically unsaturated group, is soluble in a solvent, and can be polymerized actinically or thermally.

The term “room temperature” refers to a temperature of about 21°C to about 27°C.

The term “soluble”, in reference to a compound or material in a solvent, means that the compound or material can be dissolved in the solvent to give a solution with a concentration of at least about 0.02% by weight at room temperature.

The term “insoluble”, in reference to a compound or material in a solvent, means that the compound or material can be dissolved in the solvent to give a solution with a concentration of less than 0.005% by weight at room temperature.

As used in this application, the term “ethylenically unsaturated group” is employed herein in a broad sense and is intended to encompass any groups containing at least one >C=C< group. Exemplary ethylenically unsaturated groups include without limitation

(meth)acryloyl ($\text{—}\overset{\text{O}}{\underset{\text{||}}{\text{C}}}\text{—}\overset{\text{CH}_3}{\underset{\text{|}}{\text{C}}}=\text{CH}_2$ and/or $\text{—}\overset{\text{O}}{\underset{\text{||}}{\text{C}}}\text{—CH=CH}_2$), allyl, vinyl, styrenyl, or other C=C containing groups.

The term “terminal (meth)acryloyl group” refers to one (meth)acryloyl group at one of the two ends of the main chain (or backbone) of an organic compound as known to a person skilled in the art.

The term “(meth)acrylamide” refers to methacrylamide and/or acrylamide.

The term “(meth)acrylate” refers to methacrylate and/or acrylate.

As used herein, “actinically” in reference to curing, crosslinking or polymerizing of a polymerizable composition, a prepolymer or a material means that the curing (e.g., crosslinked and/or polymerized) is performed by actinic irradiation, such as, for example,

UV/visible irradiation, ionizing radiation (e.g. gamma ray or X-ray irradiation), microwave irradiation, and the like. Thermal curing or actinic curing methods are well-known to a person skilled in the art.

A “hydrophilic vinylic monomer”, as used herein, refers to a vinylic monomer which as a homopolymer typically yields a polymer that is water-soluble or can absorb at least 10 percent by weight of water.

A “hydrophobic vinylic monomer”, as used herein, refers to a vinylic monomer which as a homopolymer typically yields a polymer that is insoluble in water and can absorb less than 10 percent by weight of water.

A “primary-amino-containing vinylic monomer” refers to a vinylic monomer having an amino group of —NH_2 .

A “blending vinylic monomer” refers to a vinylic monomer capable of dissolving both hydrophilic and hydrophobic components of a polymerizable composition to form a solution.

An “acrylic monomer” means a vinylic monomer having one sole (meth)acryloyl group.

An “N-vinyl amide monomer” refers to an amide compound having a vinyl group (—CH=CH_2) that is directly attached to the nitrogen atom of the amide group.

A “macromer” or “prepolymer” refers to a compound or polymer that is soluble in a solvent, contains ethylenically unsaturated groups, and has a number average molecular weight of greater than 700 Daltons.

As used in this application, the term “vinylic crosslinker” refers to an organic compound having at least two ethylenically unsaturated groups. A “vinylic crosslinking agent” refers to a vinylic crosslinker having a molecular weight of 700 Daltons or less.

As used in this application, the term “polymer” means a material formed by polymerizing one or more monomers or macromers or prepolymers or combinations thereof.

As used in this application, the term “molecular weight” of a polymeric material (including monomeric or macromeric materials) refers to the number average molecular weight unless otherwise specifically noted or unless testing conditions indicate otherwise.

A “polysiloxane segment” refers to a polymer chain consisting of at least three consecutively- and directly-linked siloxane units (divalent radical) each independent of one



another having a formula of —Si—O— in which R_1' and R_2' are two substituents independently selected from the group consisting of $\text{C}_1\text{—C}_{10}$ alkyl, $\text{C}_1\text{—C}_4$ alkyl- or $\text{C}_1\text{—C}_4$ -alkoxy-substituted phenyl, $\text{C}_1\text{—C}_{10}$ fluoroalkyl, $\text{C}_1\text{—C}_{10}$ fluoroether, $\text{C}_6\text{—C}_{18}$ aryl radical, $\text{—alk—(OC}_2\text{H}_4)_y\text{—OR}^0$ (in

which alk is C₁-C₆ alkyl diradical, R^o is H or C₁-C₄ alkyl and γ₁ is an integer from 1 to 10), a C₂-C₄₀ organic radical having at least one functional group selected from the group consisting of hydroxyl group (-OH), carboxyl group (-COOH), -NR₃'R₄', amino linkages of -NR₃'-, amide linkages of -CONR₃'-, amide of -CONR₃'R₄', urethane linkages of -OCONH-, and C₁-C₄ alkoxy group, or a linear hydrophilic polymer chain, in which R₃' and R₄' independent of each other are hydrogen or a C₁-C₁₅ alkyl.

A "polysiloxane vinylic crosslinker" refers to a compound comprising at least one polysiloxane segment and at least two ethylenically-unsaturated groups.

A "linear polysiloxane vinylic crosslinker" refers to a compound comprising a main chain which includes at least one polysiloxane segment and is terminated with one ethylenically-unsaturated group at each of the two ends of the main chain.

A "chain-extended polysiloxane vinylic crosslinker" refers to a compound comprising at least two ethylenically-unsaturated groups and at least two polysiloxane segments each pair of which is linked by one divalent radical.

A "polycarbosiloxane" refers to a compound containing at least one polycarbosiloxane segment which is a polymer chain consisting of at least three consecutively- and directly-linked siloxane units (divalent radical) each independent of one

another having a formula of
$$\left[\text{O}-\underset{\text{R}_2''}{\overset{\text{R}_1''}{\text{Si}}}(\text{CH}_2)_{n1}-\underset{\text{R}_4''}{\overset{\text{R}_3''}{\text{Si}}} \right]$$
 in which n₁ is an integer of 2 or 3, R₁'', R₂'', R₃'', and R₄'' independent of one another are a C₁-C₆ alkyl radical (preferably methyl).

A "polycarbosiloxane vinylic monomer" refers to a compound comprising at least one polycarbosiloxane segment and one sole ethylenically-unsaturated group.

A "polycarbosiloxane vinylic crosslinker" refers to a compound comprising at least one polycarbosiloxane segment and at least two ethylenically-unsaturated groups.

The term "fluid" means that a material is capable of flowing like a liquid.

As used in this application, the term "clear" in reference to a polymerizable composition means that the polymerizable composition is a transparent solution or liquid mixture (i.e., having a light transmissibility of 85% or greater, preferably 90% or greater in the range between 400 to 700 nm).

The term "alkyl" refers to a monovalent radical obtained by removing a hydrogen atom from a linear or branched alkane compound. An alkyl group (radical) forms one bond with one other group in an organic compound.

The term "alkylene divalent group" or "alkylene diradical" or "alkyl diradical" interchangeably refers to a divalent radical obtained by removing one hydrogen atom from an alkyl. An alkylene divalent group forms two bonds with other groups in an organic compound.

The term “alkoxy” or “alkoxyl” refers to a monovalent radical obtained by removing the hydrogen atom from the hydroxyl group of a linear or branched alkyl alcohol. An alkoxy group (radical) forms one bond with one other group in an organic compound.

In this application, the term “substituted” in reference to an alkyl diradical or an alkyl radical means that the alkyl diradical or the alkyl radical comprises at least one substituent which replaces one hydrogen atom of the alkyl diradical or the alkyl radical and is selected from the group consisting of hydroxyl (-OH), carboxyl (-COOH), -NH₂, sulfhydryl (-SH), C₁-C₄ alkyl, C₁-C₄ alkoxy, C₁-C₄ alkylthio (alkyl sulfide), C₁-C₄ acylamino, C₁-C₄ alkylamino, di-C₁-C₄ alkylamino, and combinations thereof.

A free radical initiator can be either a photoinitiator or a thermal initiator. A “photoinitiator” refers to a chemical that initiates free radical crosslinking/polymerizing reaction by the use of light. A “thermal initiator” refers to a chemical that initiates radical crosslinking/polymerizing reaction by the use of heat energy.

As used in this application, the term “phosphorylcholine” refers to a zwitterionic

group of
$$\begin{array}{c} \text{O} \\ \parallel \\ -\text{O}-\text{P}-\text{O}-(\text{CH}_2)_n-\text{N}^+-\text{R}_2''' \\ \mid \qquad \qquad \qquad \mid \\ \text{O}^- \qquad \qquad \qquad \text{R}_1''' \\ \qquad \qquad \qquad \qquad \qquad \mid \\ \qquad \qquad \qquad \qquad \qquad \text{R}_3''' \end{array}$$
 in which n is an integer of 1 to 5 and R₁′′′, R₂′′′ and R₃′′′ independently of each other are C₁-C₈ alkyl or C₁-C₈ hydroxyalkyl.

In general, the invention is directed to a method for producing SiHy contact lenses. A method of the invention is characterized by using an organic solvent as a processing-aid tool for controlling lens diameter without significantly affecting lens equilibrium water content during cast-molding of soft contact lenses from a polymerizable composition. The invention is partly based on the discovery that when an organic solvent having a boiling point of at least 90°C or higher (e.g., a C₃-C₈ alkyl alcohol) is used as a diluent in a polymerizable composition for cast molding SiHy contact lenses, the lens diameter of the fully-hydrated resultant contact lenses decreases, whereas their equilibrium water content remains substantially constant, as the weight percentage of that organic solvent increases within a range of from about 4.0% to about 15% by weight relative to the total weight of the polymerizable composition. By adjusting the weight percentage of the organic solvent in a polymerizable composition for making SiHy contact lenses, one can adjust lens diameter of produced SiHy lenses to meet target without affecting adversely the lens properties, e.g., equilibrium water content. The advantages of the invention are to easily control lens diameter in mass production of SiHy contact lenses, to enhance raw material specification or storage tolerance (e.g., vinylic monomers, vinylic crosslinkers, etc.), and thereby to lower the product cost.

The present invention provides a method for producing contact lenses. The method comprises the steps of: (1) obtaining a fluid polymerizable composition, wherein the

composition comprises an organic solvent and dissolved therein polymerizable materials, wherein the polymerizable materials comprises at least one silicone-containing vinylic monomer or crosslinker and at least one hydrophilic vinylic monomer, wherein the organic solvent has a boiling point of at least 95°C or higher (preferably at least 100°C or higher, more preferably at least 105°C or higher, even more preferably at least 110°C or higher) and is present in the polymerizable composition in an amount of from about 4% to about 15% (preferably from about 5.0% to about 12.5%, more preferably from about 5.5% to about 11%, even more preferably from about 5.5% to about 9.5%) by weight relative to the total amount of the fluid polymerizable composition and is capable of functioning as a processing-aid tool for controlling lens diameter of a hydrated soft contact lens made from the polymerizable composition; (2) introducing the fluid polymerizable composition into a cavity formed by a mold, wherein the mold has a first mold half with a first molding surface defining the anterior surface of a contact lens and a second mold half with a second molding surface defining the posterior surface of the contact lens, wherein said first and second mold halves are configured to receive each other such that a cavity is formed between said first and second molding surfaces; (3) curing thermally or actinically (preferably, curing thermally) the fluid polymerizable composition in the mold to form a dry contact lens; (4) extracting the dry contact lens with a liquid extraction medium to remove the organic solvent, unpolymerized polymerizable components and oligomers from the dry lens to obtain an extracted contact lens; and (5) hydrating the extracted contact lens with water or an aqueous solution to obtain the hydrated soft contact lens, wherein the amount of the organic solvent in the polymerizable composition is adjusted to achieve a targeted lens diameter of the hydrated soft contact lens without affecting adversely the equilibrium water content of the hydrated soft contact lens.

The term "dry contact lens" refers to a molded contact lens which has not been subjected to extraction and hydration processes.

The term "organic solvent" refers to an organic compound that cannot participate in free-radical polymerization reaction.

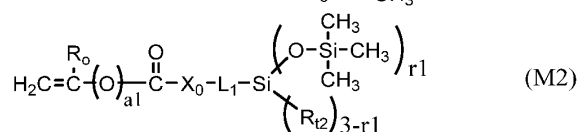
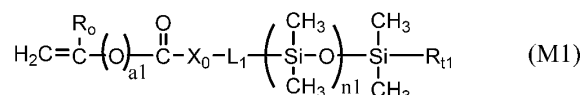
In accordance with a preferred embodiment of the invention, the organic solvent is a C₃-C₈ alkyl alcohol, preferably a C₄-C₈ alkyl alcohol, more preferably a C₄-C₆ alkyl alcohol.

Polymerizable materials (or lens-forming materials) for making contact lenses are well known to a person skilled in the art and are disclosed in numerous patents.

In accordance with the invention, a silicone-containing vinylic monomer can be any silicone-containing vinylic monomer known to a person skilled in the art. Examples of preferred silicone-containing vinylic monomers include without limitation vinylic monomers each having a bis(trialkylsilyloxy)alkylsilyl group or a tris(trialkylsilyloxy)silyl group,

polysiloxane vinylic monomers, polycarbosiloxane vinylic monomer, 3-methacryloxy propylpentamethyldisiloxane, t-butyl dimethyl-siloxyethyl vinyl carbonate, trimethylsilylethyl vinyl carbonate, and trimethylsilylmethyl vinyl carbonate, and combinations thereof.

In a preferred embodiment, the silicone-containing vinylic monomer is a siloxane-containing vinylic monomer of formula (M1) or (M2)



in which: a_1 is zero or 1; R_0 is H or methyl; X_0 is O or NR_1 ; L_1 is a C_2 - C_8 alkylene divalent radical or a divalent radical of $-\text{L}_1'-\text{X}_1-\text{L}_1''-$, $-(\text{C}_2\text{H}_4\text{O})_{q1}-\text{L}_1''-$, $-(\text{C}_2\text{H}_4\text{O})_{q1}-\text{CONH}-\text{L}_1''-$, $-\text{L}_1'-\text{NHCOO}-(\text{C}_2\text{H}_4\text{O})_{q1}-\text{L}_1''-$, $-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{X}_1'-(\text{C}_2\text{H}_4\text{O})_{q2}-\text{L}_1''-$, $-\text{L}_1'-\text{X}_1'-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O}-\text{L}_1''-$, or $-(\text{C}_2\text{H}_4\text{O})_{q1}-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O}-\text{L}_1''-$; L_1' is a C_2 - C_8 alkylene divalent radical which has zero or one hydroxyl group; L_1'' is C_3 - C_8 alkylene divalent radical which has zero or one hydroxyl group; X_1 is O, NR_1 , NHCOO , OCONH , CONR_1 , or NR_1CO ; R_1 is H or a C_1 - C_4 alkyl having 0 to 2 hydroxyl group; R_{t1} and R_{t2} independent of each other are a C_1 - C_4 alkyl; X_1' is O or NR_1 ; q_1 is an integer of 1 to 20; q_2 is an integer of 0 to 20; n_1 is an integer of 3 to 25; and r_1 is an integer of 2 or 3.

Examples of preferred siloxane-containing vinylic monomers of formula (M1) include without limitation α -(meth)acryloxypropyl terminated ω - C_1 - C_4 -alkyl terminated polydimethylsiloxane, α -(meth)acryloxy-2-hydroxypropyloxypropyl terminated ω - C_1 - C_4 -alkyl terminated polydimethylsiloxane, α -(2-hydroxyl-methacryloxypropyloxypropyl)- ω - C_1 - C_4 -alkyl-decamethylpentasiloxane, α -[3-(meth)acryloxyethoxy-2-hydroxypropyloxypropyl]-terminated ω - C_1 - C_4 -alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxy-propyloxy-2-hydroxypropyloxypropyl]-terminated ω - C_1 - C_4 -alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxyisopropyloxy-2-hydroxypropyloxypropyl]-terminated ω - C_1 - C_4 -alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxybutyloxy-2-hydroxypropyloxypropyl]-terminated ω - C_1 - C_4 -alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxyethylamino-2-hydroxypropyloxypropyl]-terminated ω - C_1 - C_4 -alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxypropylamino-2-hydroxypropyloxypropyl]-terminated ω - C_1 - C_4 -alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxy-butylamino-2-hydroxypropyloxypropyl]-terminated ω - C_1 - C_4 -alkyl terminated polydimethylsiloxane, α -(meth)acryloxy(polyethylenoxy)-2-hydroxypropyloxypropyl]-terminated ω - C_1 - C_4 -alkyl

terminated polydimethylsiloxane, α -[(meth)acryloxy-2-hydroxypropyloxy-ethoxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[(meth)acryloxy-2-hydroxypropyl-N-ethylaminopropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[(meth)acryloxy-2-hydroxypropyl-aminopropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[(meth)acryloxy-2-hydroxypropyloxy-(polyethylenoxy)propyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -(meth)acryloylamidopropoxypropyl terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -N-methyl-(meth)acryloylamidopropoxypropyl terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acrylamidoethoxy-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl polydimethylsiloxane, α -[3-(meth)acrylamidopropoxy-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acrylamidoisopropoxy-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acrylamidobutyloxy-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloylamido-2-hydroxypropyloxypropyl] terminated ω -C₁-C₄-alkyl polydimethylsiloxane, α -[3-[N-methyl-(meth)acryloylamido]-2-hydroxypropyloxypropyl] terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, N-methyl-N'-(propyltetra(dimethylsiloxy)dimethylbutylsilane) (meth)acrylamide, N-(2,3-dihydroxypropane)-N'-(propyltetra(dimethylsiloxy)dimethylbutylsilane) (meth)acrylamide, (meth)acryloylamidopropyltetra(dimethylsiloxy)dimethylbutylsilane, α -vinyl carbonate-terminated ω -C₁-C₄-alkyl-terminated polydimethylsiloxanes, α -vinyl carbamate-terminated ω -C₁-C₄-alkyl-terminated polydimethylsiloxane, hydrophilized siloxane-containing (meth)acrylamides disclosed in US 9,097,840, hydrophilized siloxane-containing acrylic monomers disclosed in US 9,103,965, and mixtures thereof. The above preferred polysiloxanes vinylic monomers of formula (M1) can be obtained from commercial suppliers (e.g., Shin-Etsu, Gelest, etc.) or prepared according to procedures described in patents, e.g., U.S. Pat. Nos. 5070215, 6166236, 6867245, 8415405, 8475529, 8614261, 9097840, 9103965, and 9217813, or by reacting a hydroxyalkyl (meth)acrylate or (meth)acrylamide or a (meth)acryloxypolyethylene glycol with a mono-epoxypropyloxypropyl-terminated polydimethylsiloxane, by reacting glycidyl (meth)acrylate with a mono-carbinol-terminated polydimethylsiloxane, a mono-aminopropyl-terminated polydimethylsiloxane, or a mono-ethylaminopropyl-terminated polydimethylsiloxane, or by reacting isocyanatoethyl (meth)acrylate with a mono-carbinol-terminated polydimethylsiloxane according to coupling reactions well known to a person skilled in the art.

Examples of preferred siloxane-containing vinylic monomers of formula (M2) include without limitation tris(trimethylsilyloxy)silylpropyl (meth)acrylate, [3-(meth)acryloxy-2-hydroxypropyloxy]propylbis(trimethylsiloxy)methylsilane, [3-(meth)acryloxy-2-hydroxypropyloxy]propylbis(trimethylsiloxy)butylsilane, 3-(meth)acryloxy-2-(2-hydroxyethoxy)-propyloxypropylbis(trimethylsiloxy)methylsilane, 3-(meth)acryloxy-2-hydroxypropyloxy-propyltris(trimethylsiloxy)silane, N-[tris(trimethylsiloxy)silylpropyl]-(meth)acrylamide, N-(2-hydroxy-3-(3-(bis(trimethylsilyloxy)methylsilyl)propyloxy)propyl)-2-methyl (meth)acrylamide, N-(2-hydroxy-3-(3-(bis(trimethylsilyloxy)methylsilyl)propyloxy)propyl) (meth)acrylamide, N-(2-hydroxy-3-(3-(tris(trimethylsilyloxy)silyl)propyloxy)propyl)-2-methyl acrylamide, N-(2-hydroxy-3-(3-(tris(trimethylsilyloxy)silyl)propyloxy)propyl) (meth)acrylamide, N-[tris(dimethylpropylsiloxy)silylpropyl]-(meth)acrylamide, N-[tris(dimethylphenylsiloxy)silylpropyl] (meth)acrylamide, N-[tris(dimethylethylsiloxy)silylpropyl] (meth)acrylamide, N,N-bis[2-hydroxy-3-(3-(bis(trimethylsilyloxy)methylsilyl)propyloxy)propyl]-2-methyl (meth)acrylamide, N,N-bis[2-hydroxy-3-(3-(bis(trimethylsilyloxy)methylsilyl)propyloxy)propyl] (meth)acrylamide, N,N-bis[2-hydroxy-3-(3-(tris(trimethylsilyloxy)silyl)propyloxy)propyl]-2-methyl (meth)acrylamide, N,N-bis[2-hydroxy-3-(3-(tris(trimethylsilyloxy)silyl)propyloxy)propyl] (meth)acrylamide, N-[2-hydroxy-3-(3-(t-butylidimethylsilyl)propyloxy)propyl]-2-methyl (meth)acrylamide, N-[2-hydroxy-3-(3-(t-butylidimethylsilyl)propyloxy)propyl] (meth)acrylamide, N,N-bis[2-hydroxy-3-(3-(t-butylidimethylsilyl)propyloxy)propyl]-2-methyl (meth)acrylamide, N-2-(meth)acryloxyethyl-O-(methyl-bis-trimethylsiloxy-3-propyl)silyl carbamate, 3-(trimethylsilyl)propylvinyl carbonate, 3-(vinylloxycarbonylthio)propyl-tris(trimethylsiloxy)silane, 3-[tris(trimethylsiloxy)silyl]propylvinyl carbamate, 3-[tris(trimethylsiloxy)silyl]propyl allyl carbamate, 3-[tris(trimethylsiloxy)silyl]propyl vinyl carbonate, those disclosed in U.S. Pat. Nos. 9097840, 9103965 and 9475827, and mixtures thereof. The above preferred silicone-containing vinylic monomers can be obtained from commercial suppliers or can be prepared according to procedures described in U.S. Pat. Nos. 5070215, 6166236, 7214809, 8475529, 8658748, 9097840, 9103965, and 9475827.

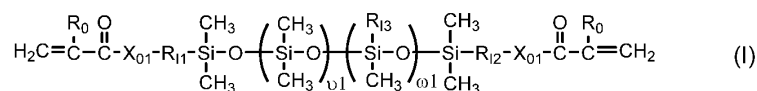
Any polycarbosiloxane vinylic monomers can be used in the invention. Examples of preferred polycarbosiloxane vinylic monomers include without limitation those disclosed in U.S. Pat. Nos. 7915323 and 8420711 and in U.S. Pat. Appl. Pub. Nos. 2012/244088A1 and 2012/245249A1.

Any silicone-containing vinylic crosslinkers can be used in the invention. Examples of preferred silicone-containing vinylic crosslinkers include without limitation polysiloxane vinylic crosslinkers, polycarbosiloxane vinylic crosslinkers, and combinations thereof.

Any polysiloxane vinylic crosslinkers can be used in the invention. Examples of preferred polysiloxane vinylic crosslinkers are di-(meth)acryloyl-terminated

polydimethylsiloxanes; di-vinyl carbonate-terminated polydimethylsiloxanes; di-vinyl carbamate-terminated polydimethylsiloxane; N,N,N',N'-tetrakis(3-methacryloxy-2-hydroxypropyl)-alpha,omega-bis-3-aminopropyl-polydimethylsiloxane; polysiloxane-containing macromer selected from the group consisting of Macromer A, Macromer B, Macromer C, and Macromer D described in US 5,760,100; polysiloxane-containing macromers disclosed in U.S. Pat. Nos. 4,136,250, 4,153,641, 4,182,822, 4,189,546, 4,343,927, 4,254,248, 4,355,147, 4,276,402, 4,327,203, 4,341,889, 4,486,577, 4,543,398, 4,605,712, 4,661,575, 4,684,538, 4,703,097, 4,833,218, 4,837,289, 4,954,586, 4,954,587, 5,010,141, 5,034,461, 5,070,170, 5,079,319, 5,039,761, 5,346,946, 5,358,995, 5,387,632, 5,416,132, 5,451,617, 5,486,579, 5,962,548, 5,981,675, 6,039,913, and 6,762,264; polysiloxane-containing macromers disclosed in U.S. Pat. Nos. 4,259,467, 4,260,725, and 4,261,875.

One class of preferred polysiloxane vinylic crosslinkers are di-(meth)acryloyloxy-terminated polysiloxane vinylic crosslinkers each having dimethylsiloxane units and hydrophilized siloxane units each having one methyl substituent and one monovalent C₄–C₄₀ organic radical substituent having 2 to 6 hydroxyl groups as disclosed in U.S. Pat. App. Pub. No. 2017/0166673 A1, more preferably a polysiloxane vinylic crosslinker of formula (I)



in which:

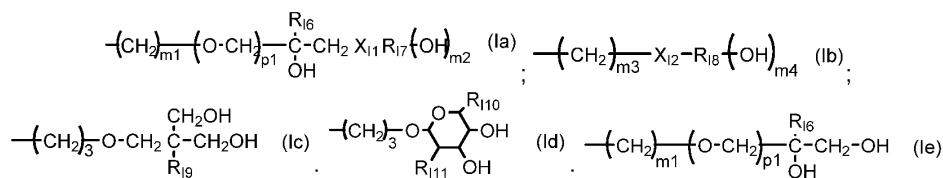
$\omega 1$ is an integer of from 30 to 500 and $\omega 1$ is an integer of from 1 to 75, provided that $\omega 1/\omega 1$ is from about 0.035 to about 0.15 (preferably from about 0.040 to about 0.12, even more preferably from about 0.045 to about 0.10);

X₀₁ is O or NR_N in which R_N is hydrogen or C₁–C₁₀-alkyl;

R₀ is hydrogen or methyl;

R₁₁ and R₁₂ independently of each other are a substituted or unsubstituted C₁–C₁₀ alkylene divalent radical or a divalent radical of –R₁₄–O–R₁₅– in which R₁₄ and R₁₅ independently of each other are a substituted or unsubstituted C₁–C₁₀ alkylene divalent radical;

R₁₃ is a monovalent radical of any one of formula (Ia) to (Ie)

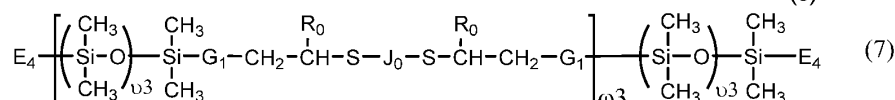
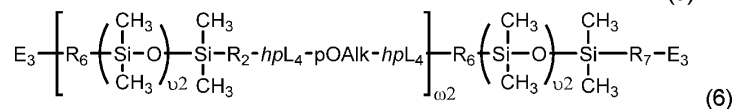
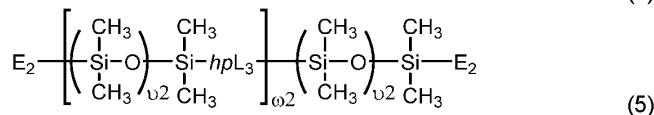
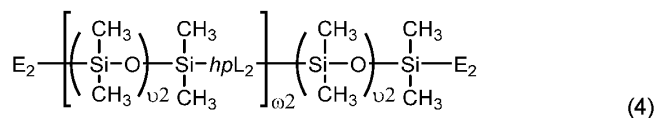
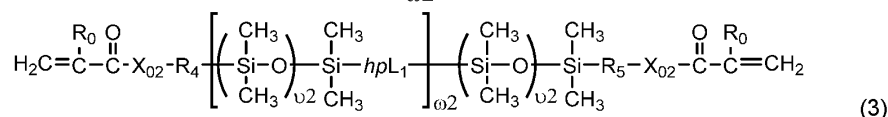
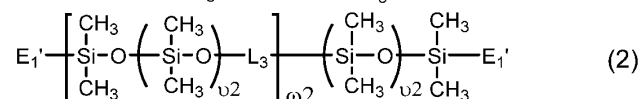
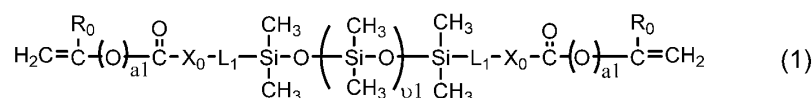


in which: p₁ is zero or 1; m₁ is an integer of 2 to 4; m₂ is an integer of 1 to 5; m₃ is an integer of 3 to 6; m₄ is an integer of 2 to 5; R₁₆ is hydrogen or methyl; R₁₇ is a C₂–C₆

hydrocarbon radical having (m2+1) valencies; R₁₈ is a C₂-C₆ hydrocarbon radical having (m4+1) valencies; R₁₉ is ethyl or hydroxymethyl; R₁₁₀ is methyl or hydromethyl; R₁₁₁ is hydroxyl or methoxy; X₁₁ is a sulfur linkage of -S- or a tertiary amino linkage of -NR₁₁₂- in which R₁₁₂ is C₁-C₁ alkyl, hydroxyethyl, hydroxypropyl, or 2,3-dihydroxypropyl; and X₁₂ is an amide linkage of -NR₁₁₃-C(=O)- or -C(=O)-NR₁₁₃- in which R₁₁₃ is hydrogen or C₁-C₁₀ alkyl.

A polysiloxane vinylic crosslinker of formula (I) can be prepared according to procedures described in U.S. Pat. Appl. Pub. No. 20120088843A1.

Other classes of preferred polysiloxane vinylic crosslinkers are vinylic crosslinkers of any one of formula (1) to (7)



in which: v₁ is an integer of from 30 to 500; v₂ is an integer of from 5 to 50; v₃ is an integer of from 5 to 100; ω₂ and ω₃ independent of each other are an integer of from 1 to 15; a₁ and g₁ independent of each other is zero or 1; h₁ is an integer of 1 to 20 and h₂ is an integer of 0 to 20; m₁ and m₃ independent of each other is 0 or 1, m₂ is an integer of 1 to 6, m₄ is an integer of 1 to 5, m₅ is 2 or 3; q₁ is an integer of 1 to 20, q₂ is an integer of 0 to 20, q₃ is an integer of 0 to 2, q₄ is an integer of 2 to 50, q₅ and q₆ independent of each other are a number of 0 to 35, provided that (q₄+q₅+q₆) is an integer of 2 to 50; x+y is an integer of from 10 to 30; e₁ is an integer of 5 to 100, p₁ and b₁ independent of each other

are an integer of 0 to 50, provided that $(e1+p1+b1) \geq 10$ and $\frac{e1}{(p1+b1)} \geq 2$ (preferably from about 2:1 to about 10:1, more preferably from about 3:1 to about 6:1) when $(p1+b1) \geq 1$; R_0 is H or methyl; $R_1, R_{1n}, R_{2n}, R_{3n}$, and R_{4n} independent of one another are H or a C₁-C₄ alkyl having 0 to 2 hydroxyl group; R_{n5} is H or a C₁-C₁₀ alkyl; R_2 is a C₄-C₁₄ hydrocarbon divalent radical; R_3 is a C₂-C₆ alkylene divalent radical; R_4 and R_5 independent of each other are a C₁-C₆ alkylene divalent radical or a C₁-C₆ alkylene-oxy-C₁-C₆ alkylene divalent radical; R_6 and R_7 independent of each other are a C₁-C₆ alkylene divalent radical or a C₁-C₆ alkoxy-C₁-C₆ alkylene divalent radical; R_8 and R_9 independent of each other are a substituted or unsubstituted C₁-C₁₂ alkylene divalent radical; X_0, X_1, X_{01}, X_{02} , and X_{03} independent of one another are O or NR₁; X_1 is O, NR₁, NHCOO, CONH, CONR₁, or NR₁CO; X_{04} is -COO- or -CONR_{n5}-; X_{05} and X_{07} independent of each other are a direct bond, -COO- or -CONR_{n5}-; X_{06} is a direct bond, a C₁-C₆ alkylene divalent radical, a C₁-C₆ alkyleneoxy divalent radical, -COO-, or -CONR_{n5}-; X_{08} is a direct bond or -COO-; X_{09} is O or NR_{n5}; X_{10} is a direct bond, a C₁-C₆ alkylene divalent radical, -COO-, or -CONR_{n5}-; E_1 is a

monovalent radical of $H_2C=\overset{\overset{R_0}{\parallel}}{\underset{\underset{O}{\parallel}}{C}}-X_{01}-L_4-$; E_2 is a monovalent radical of

$H_2C=\overset{\overset{R_0}{\parallel}}{\underset{\underset{O}{\parallel}}{C}}-Z_7-$; E_3 is a monovalent radical of $H_2C=\overset{\overset{R_0}{\parallel}}{\underset{\underset{O}{\parallel}}{C}}-X_{03}-$; E_4 is a monovalent radical

of $H_2C=\overset{\overset{R_0}{\parallel}}{\underset{\underset{O}{\parallel}}{C}}-X_{11}-G_2-$ or $H_2C=\overset{\overset{R_0}{\parallel}}{\underset{\underset{O}{\parallel}}{C}}-G_3-$; L_1 is a C₂-C₈ alkylene divalent radical or a

divalent radical of $-L_1'-X_1-L_1''-$, $-(C_2H_4O)_{q1}-L_1''-$, $-(C_2H_4O)_{q1}-CONH-L_1''-$,

$-L_1'-NHCOO-(C_2H_4O)_{q1}-L_1''-$, $-CH_2-CH(OH)-CH_2-X_1'-(C_2H_4O)_{q2}-L_1''-$,

$-L_1'-X_1'-CH_2-CH(OH)-CH_2-O-L_1''-$, or $-(C_2H_4O)_{q1}-CH_2-CH(OH)-CH_2-O-L_1''-$; L_1' is a C₂-C₈

alkylene divalent radical which has zero or one hydroxyl group; L_1'' is C₃-C₈ alkylene

divalent radical which has zero or one hydroxyl group; L_3 is a divalent radical of

$-L_3'-O-(C_2H_4O)_{q2}-CONH-R_2-(NHCO-PE-CONH-R_2)_{g1}-NHCO-(OC_2H_4)_{q2}-O-L_3'-$ in which PE is a

divalent radical of $-(CH_2CH_2O)_{q3}-Z_0-CF_2-(OCF_2)_x-(OCF_2CF_2)_y-OCF_2-Z_0-(OCH_2CH_2)_{q3}-$ or

$-Z_0-(CH_2CH_2O)_{q4}-\left(CH_2-\overset{\overset{CH_3}{\mid}}{CH}-O\right)_{q5}-\left(CH_2-\overset{\overset{C_2H_5}{\mid}}{CH}-O\right)_{q6}-Z_0-$; L_3' is C₃-C₈ alkylene divalent radical;

L_4 is a divalent radical of $-C_2H_4-NHCO-O-(C_2H_4O)_{q2}-L_3'-$,

$-(C_2H_4O)_{q1}-CONH-R_2-NHCO-O-(C_2H_4O)_{q2}-L_3'-$, $-R_3-O-CONH-R_2-NHCO-O-(C_2H_4O)_{q2}-L_3'-$,

$-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O}-\left(\text{C}_2\text{H}_4\text{O}\right)_{q2}-\text{L}_3'-$, or $-\left(\text{C}_2\text{H}_4\text{O}\right)_{q2}-\text{L}_3'-$; hpl_1 is a divalent radical of

$-\text{R}_4-\overset{\text{R}_0}{\underset{|}{\text{CH}}}-\text{CH}_2-\text{S}-\text{C}_2\text{H}_4-\overset{\text{H}_3\text{C}-\overset{\text{O}}{\parallel}{\text{C}}-\text{NH}}{\underset{\text{O}}{\parallel}{\text{C}}}-\text{NR}_{n2}-\text{Y}_1-\text{NR}_{n2}-\overset{\text{O}}{\parallel}{\text{C}}-\overset{\text{H}_3\text{C}-\overset{\text{O}}{\parallel}{\text{C}}-\text{NH}}{\underset{\text{O}}{\parallel}{\text{C}}}-\text{CH}_2-\text{CH}-\overset{\text{R}_0}{\underset{|}{\text{R}}}_5-$; hpl_2 is a

divalent radical of $-\text{R}_4-\text{NR}_{n2}-\overset{\text{O}}{\parallel}{\text{C}}-\overset{\text{H}_3\text{C}-\overset{\text{O}}{\parallel}{\text{C}}-\text{NH}}{\underset{\text{O}}{\parallel}{\text{C}}}-\text{CH}-\text{C}_2\text{H}_4-\text{S}-\text{Y}_2-\text{S}-\text{C}_2\text{H}_4-\overset{\text{H}_3\text{C}-\overset{\text{O}}{\parallel}{\text{C}}-\text{NH}}{\underset{\text{O}}{\parallel}{\text{C}}}-\text{CH}-\text{NR}_{n2}-\text{R}_5-$; hpl_3 is a

divalent radical of $-\text{R}_4-\text{NR}_{n2}-\overset{\text{O}}{\parallel}{\text{C}}-\overset{\text{H}_3\text{C}-\overset{\text{O}}{\parallel}{\text{C}}-\text{NH}}{\underset{\text{O}}{\parallel}{\text{C}}}-\text{CH}-\text{C}_2\text{H}_4-\text{S}-\text{Y}_3-\text{S}-\text{C}_2\text{H}_4-\overset{\text{H}_3\text{C}-\overset{\text{O}}{\parallel}{\text{C}}-\text{NH}}{\underset{\text{O}}{\parallel}{\text{C}}}-\text{CH}-\text{NR}_{n2}-\text{R}_5-$; hpl_4 is a

divalent radical of $-\text{X}_{03}-\overset{\text{O}}{\parallel}{\text{C}}-\overset{\text{R}_0}{\underset{|}{\text{CH}}}-\text{CH}_2-\text{S}-\text{C}_2\text{H}_4-\overset{\text{H}_3\text{C}-\overset{\text{O}}{\parallel}{\text{C}}-\text{NH}}{\underset{\text{O}}{\parallel}{\text{C}}}-\text{CH}-\text{NR}_{n4}-\text{R}_8-$ or

$-\text{R}_9-\text{NR}_{n4}-\overset{\text{O}}{\parallel}{\text{C}}-\overset{\text{H}_3\text{C}-\overset{\text{O}}{\parallel}{\text{C}}-\text{NH}}{\underset{\text{O}}{\parallel}{\text{C}}}-\text{CH}-\text{C}_2\text{H}_4-\text{S}-\text{CH}_2-\overset{\text{R}_0}{\underset{|}{\text{CH}}}-\overset{\text{O}}{\parallel}{\text{C}}-\text{X}_{03}-$; $pOAlk$ is a divalent radical of

$-(\text{EO})_e(\text{PO})_p(\text{BO})_b-$ in which EO is an oxyethylene unit ($-\text{CH}_2\text{CH}_2\text{O}-$), PO is an

oxypropylene unit ($-\text{CH}_2-\overset{\text{CH}_3}{\underset{|}{\text{CH}}}-\text{O}-$), and BO is an oxybutylene unit ($-\text{CH}_2-\overset{\text{CH}_2\text{CH}_3}{\underset{|}{\text{CH}}}-\text{O}-$); M_0 is

C_3-C_8 alkylene divalent radical; M_1 is a C_4-C_{14} hydrocarbon divalent radical; M_2 and M_3 independent of each other are a C_1-C_6 alkylene divalent radical; J_0 is a C_1-C_{12} hydrocarbon radical having 0 to 2 hydroxyl or carboxyl groups; G_1 is a direct bond, a C_1-C_4 alkylene

divalent radical, or a bivalent radical of $-\text{X}_{04}-\left(\text{C}_2\text{H}_4\text{O}\right)_{h1}-\text{CONH}-M_1-\text{NHCO}-\text{O}-\left(\text{C}_2\text{H}_4\text{O}\right)_{h2}-M_0-$,

$-\text{X}_{05}-M_2-\text{O}-\text{CONH}-M_1-\text{NHCO}-\text{O}-\left(\text{C}_2\text{H}_4\text{O}\right)_{h2}-M_0-$, $-\text{X}_{06}-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O}-\left(\text{C}_2\text{H}_4\text{O}\right)_{h2}-M_0-$,

$-\text{X}_{07}-\left(\text{C}_2\text{H}_4\text{O}\right)_{h2}-M_0-$, $-\text{X}_{08}-M_3-\text{NHCOO}-\left(\text{C}_2\text{H}_4\text{O}\right)_{h1}-M_0-$,

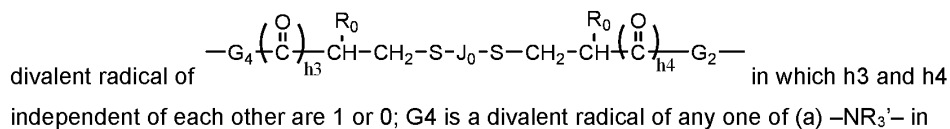
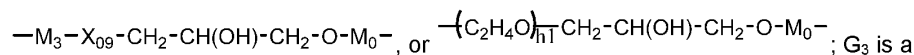
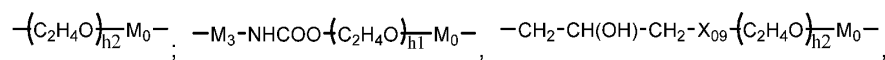
$-\text{X}_{10}-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{X}_{09}-\left(\text{C}_2\text{H}_4\text{O}\right)_{h2}-M_0-$, $-\text{X}_{07}-M_3-\text{X}_{09}-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O}-M_0-$, or

$-\text{X}_{08}-\left(\text{C}_2\text{H}_4\text{O}\right)_{h1}-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O}-M_0-$ in which M_0 is linked to Si atom while X_{04} to X_{10}

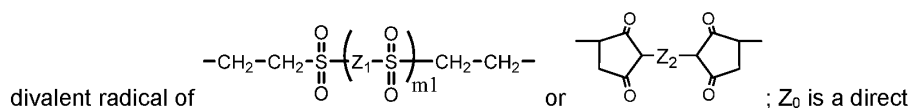
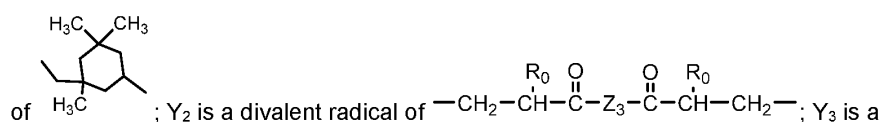
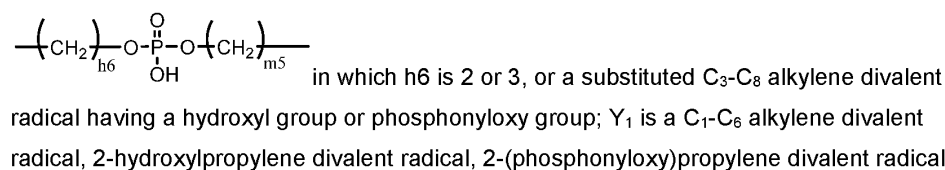
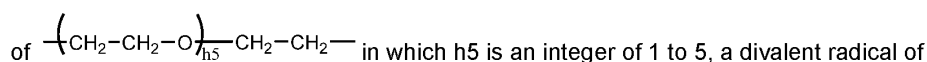
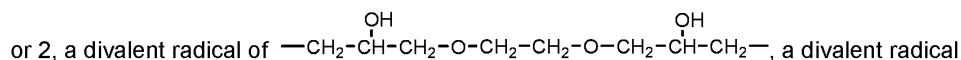
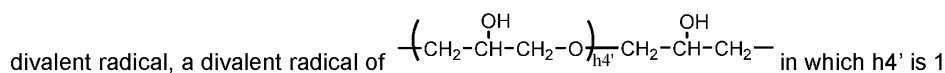
are linked to the group of $-\text{CH}_2-$ in formula (7) and at least one of J_0 and G_1 in formula (7) comprises at least one moieties selected from the group consisting of hydroxyl groups, urethane linkage of $-\text{OCONH}-$, amino groups of $-\text{NHR}^0$, amino linkages of $-\text{NH}-$, amide linkages of $-\text{CONH}-$, carboxyl groups, and combinations thereof; G_2 is a C_1-C_4 alkylene

divalent radical or a bivalent radical of $-\left(\text{C}_2\text{H}_4\text{O}\right)_{h1}-\text{CONH}-M_1-\text{NHCO}-\text{O}-\left(\text{C}_2\text{H}_4\text{O}\right)_{h2}-M_0-$,

$-\text{M}_2-\text{O}-\text{CONH}-M_1-\text{NHCO}-\text{O}-\left(\text{C}_2\text{H}_4\text{O}\right)_{h2}-M_0-$, $-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O}-\left(\text{C}_2\text{H}_4\text{O}\right)_{h2}-M_0-$,



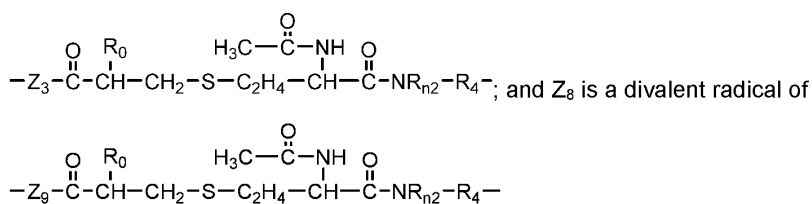
which R_3' is hydrogen or C_1 - C_3 alkyl, (b) $-\text{N} \begin{array}{c} \diagup \quad \diagdown \\ | \quad | \\ \text{---} \end{array} -$, (c) $-NR_0-G_5-NR_0-$ in which G_5 is a C_1 - C_6 alkylene divalent radical, 2-hydroxylpropylene divalent radical, 2-(phosphonyloxy)propylene divalent radical, 1,2-dihydroxyethylene divalent radical, 2,3-dihydroxybutylene divalent radical, and (d) $-O-G_6-O-$ in which G_6 is a C_1 - C_6 alkylene



bond or a C_1 - C_{12} alkylene divalent radical; Z_1 is a C_1 - C_6 alkylene divalent radical, a hydroxyl- or methoxy-substituted C_1 - C_6 alkylene divalent radical, or a substituted or unsubstituted phenylene divalent radical, Z_2 is a C_1 - C_6 alkylene divalent radical, a hydroxyl- or methoxy-substituted C_1 - C_6 alkylene divalent radical, a dihydroxyl- or dimethoxy-substituted C_2 - C_6 alkylene divalent radical, a divalent radical of $-C_2H_4-(O-C_2H_4)_{m2}-$, a divalent radical of $-Z_4-S-S-Z_4-$, a hydroxyl- or methoxy-substituted C_1 - C_6 alkylene divalent radical, or a substituted or unsubstituted phenylene divalent radical, Z_3 is a divalent radical

of any one of (a) $-\text{NR}_{n3}-$, (b) $-\text{N} \begin{array}{c} \diagup \diagdown \\ \diagdown \diagup \end{array} \text{N}-$, (c) $-\text{NR}_6-\text{Z}_5-\text{NR}_6-$, and (d) $-\text{O}-\text{Z}_6-\text{O}-$, Z_4 is a C_1 - C_8 alkylene divalent radical, Z_5 is a C_1 - C_6 alkylene divalent radical, 2-hydroxypropylene divalent radical, 2-(phosphonyloxy)propylene divalent radical, 1,2-dihydroxyethylene divalent radical, 2,3-dihydroxybutylene divalent radical, Z_6 is (a) a C_1 - C_6 alkylene divalent

radical, (b) a divalent radical of $-(\text{CH}_2-\overset{\text{OH}}{\underset{|}{\text{CH}}}-\text{CH}_2-\text{O})_{m3}-\text{CH}_2-\overset{\text{OH}}{\underset{|}{\text{CH}}}-\text{CH}_2-$, $-\text{CH}_2-\overset{\text{OH}}{\underset{|}{\text{CH}}}-\text{CH}_2-\text{O}-\text{CH}_2-\text{CH}_2-\text{O}-\text{CH}_2-\overset{\text{OH}}{\underset{|}{\text{CH}}}-\text{CH}_2-$, $-(\text{CH}_2-\text{CH}_2-\text{O})_{m4}-\text{CH}_2-\text{CH}_2-$, $-(\text{CH}_2)_{m5}-\text{O}-\overset{\text{O}}{\underset{\text{OH}}{\text{P}}}(\text{CH}_2)_{m5}-$, or (c) a substituted C_3 - C_8 alkylene divalent radical having a hydroxyl group or phosphonyloxy group, Z_7 is a divalent radical of



Polysiloxane vinylic crosslinkers of formula (1) can be obtained from commercial suppliers, prepared according to procedures described in US , or prepared by reacting glycidyl (meth)acrylate (meth)acryloyl chloride with a di-amino-terminated polydimethylsiloxane or a di-hydroxyl-terminated polydimethylsiloxane, reacting isocyanatoethyl (meth)acrylate with di-hydroxyl-terminated polydimethylsiloxanes, reacting an amino-containing acrylic monomer with di-carboxyl-terminated polydimethylsiloxane in the presence of a coupling agent (a carbodiimide); reacting a carboxyl-containing acrylic monomer with di-amino-terminated polydimethylsiloxane in the presence of a coupling agent (a carbodiimide). Or reacting a hydroxyl-containing acrylic monomer with a di-hydroxy-terminated polydisiloxane.

Preferred examples of polysiloxane vinylic crosslinkers of formula (1) include without limitation α,ω -bis[3-(meth)acrylamidopropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxyethoxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxypropyloxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxy-isopropyloxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxybutyloxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamidoethoxy-2-

hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamidopropoxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamidoisopropoxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamidobutyloxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxyethylamino-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxypropylamino-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxybutylamino-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acrylamidoethylamino-2-hydroxypropyloxy-propyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamidopropylamino-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamide-butylamino-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acryloxy-2-hydroxypropyloxy-ethoxypropyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acryloxy-2-hydroxypropyl-N-ethylaminopropyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acryloxy-2-hydroxypropyl-aminopropyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acryloxy-2-hydroxypropyloxy-(polyethylenoxy)propyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acryloxyethylamino-carbonyloxy-ethoxypropyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acryloxyethylamino-carbonyloxy-(polyethylenoxy)propyl]-terminated polydimethylsiloxane, combinations thereof.

Chain-extended polysiloxane vinylic crosslinkers of formula (2) can be prepared according to the procedures described in U.S. Pat. Nos. 5034461, 5416132, 5449729, 5760100, 7423074, and 8529057. Chain-extended polysiloxane vinylic crosslinkers of formula (3), (4) or (5) can be prepared according to the procedures described in detail in U.S. Pat. App. Ser. No. 62/406465 filed 11 Oct. 2016. Chain-extended polysiloxane vinylic crosslinkers of formula (6) can be prepared according to the procedures described in detail in U.S. Pat. App. Ser. No. 62/406467 filed 11 Oct. 2016. Chain-extended polysiloxane vinylic crosslinkers of formula (6) can be prepared according to the procedures described in US8993651.

Another class of preferred chain-extended polysiloxane vinylic crosslinkers are those which each comprise at least two polysiloxane segments and dangling hydrophilic polymer chains each covalently attached to a divalent radical separating each pair of adjacent polysiloxane segments and having at least two pendant hydrophilic groups and/or chains as disclosed in US20120088843A1; those which each comprise at least two polysiloxane segments and dangling hydrophilic polymer chains each covalently attached to a divalent radical separating each pair of adjacent polysiloxane segments as disclosed in US20120088844A1.

Any polycarbosiloxane vinylic crosslinkers can be used in the invention. Examples of preferred polycarbosiloxane vinylic crosslinkers include without limitation those disclosed in U.S. Pat. Nos. 7915323 and 8420711 and in U.S. Pat. Appl. Pub. Nos. 2012/0244088 and 2012/0245249.

Any hydrophilic vinylic monomers can be used in the invention. Examples of preferred hydrophilic vinylic monomers are alkyl (meth)acrylamides (as described below), hydroxyl-containing acrylic monomers (as described below), carboxyl-containing acrylic monomers (as described below), amino-containing acrylic monomers (as described below), N-vinyl amide monomers (as described below), methylene-containing pyrrolidone monomers (i.e., pyrrolidone derivatives each having a methylene group connected to the pyrrolidone ring at 3- or 5- position) (as described below), acrylic monomers having a C₁-C₄ alkoxyethoxy group (as described below), vinyl ether monomers (as described below), allyl ether monomers (as described below), phosphorylcholine-containing vinylic monomers (as described below), allyl alcohol, N-2-hydroxyethyl vinyl carbamate, N-carboxyvinyl- β -alanine (VINAL), N-carboxyvinyl- α -alanine, and combinations thereof.

Examples of alkyl (meth)acrylamides includes without limitation (meth)acrylamide, N,N-dimethyl (meth)acrylamide, N-ethyl (meth)acrylamide, N,N-diethyl (meth)acrylamide, N-propyl (meth)acrylamide, N-isopropyl (meth)acrylamide, N-3-methoxypropyl (meth)acrylamide, and combinations thereof.

Examples of hydroxyl-containing acrylic monomers include without limitation N-2-hydroxyethyl (meth)acrylamide, N,N-bis(hydroxyethyl) (meth)acrylamide, N-3-hydroxypropyl (meth)acrylamide, N-2-hydroxypropyl (meth)acrylamide, N-2,3-dihydroxypropyl (meth)acrylamide, *N*-tris(hydroxymethyl)methyl (meth)acrylamide, 2-hydroxyethyl (meth)acrylate, 3-hydroxypropyl (meth)acrylate, 2-hydroxypropyl (meth)acrylate, glycerol methacrylate (GMA), di(ethylene glycol) (meth)acrylate, tri(ethylene glycol) (meth)acrylate, tetra(ethylene glycol) (meth)acrylate, poly(ethylene glycol) (meth)acrylate having a number average molecular weight of up to 1500, poly(ethylene glycol)ethyl (meth)acrylamide having a number average molecular weight of up to 1500, and combinations thereof.

Examples of amino-containing acrylic monomers include without limitation N-2-aminoethyl (meth)acrylamide, N-2-methylaminoethyl (meth)acrylamide, N-2-ethylaminoethyl (meth)acrylamide, N-2-dimethylaminoethyl (meth)acrylamide, N-3-aminopropyl (meth)acrylamide, N-3-methylaminopropyl (meth)acrylamide, N-3-dimethylaminopropyl (meth)acrylamide, 2-aminoethyl (meth)acrylate, 2-methylaminoethyl (meth)acrylate, 2-ethylaminoethyl (meth)acrylate, 3-aminopropyl (meth)acrylate, 3-methylaminopropyl (meth)acrylate, 3-ethylaminopropyl (meth)acrylate, 3-amino-2-hydroxypropyl (meth)acrylate, trimethylammonium 2-hydroxy propyl (meth)acrylate

hydrochloride, dimethylaminoethyl (meth)acrylate, and combinations thereof.

Examples of carboxyl-containing acrylic monomers include without limitation 2-(meth)acrylamidoglycolic acid, (meth)acrylic acid, ethylacrylic acid, and combinations thereof.

Examples of preferred N-vinyl amide monomers include without limitation N-vinylpyrrolidone (*aka*, N-vinyl-2-pyrrolidone), N-vinyl-3-methyl-2-pyrrolidone, N-vinyl-4-methyl-2-pyrrolidone, N-vinyl-5-methyl-2-pyrrolidone, N-vinyl-6-methyl-2-pyrrolidone, N-vinyl-3-ethyl-2-pyrrolidone, N-vinyl-4,5-dimethyl-2-pyrrolidone, N-vinyl-5,5-dimethyl-2-pyrrolidone, N-vinyl-3,3,5-trimethyl-2-pyrrolidone, N-vinyl piperidone (*aka*, N-vinyl-2-piperidone), N-vinyl-3-methyl-2-piperidone, N-vinyl-4-methyl-2-piperidone, N-vinyl-5-methyl-2-piperidone, N-vinyl-6-methyl-2-piperidone, N-vinyl-6-ethyl-2-piperidone, N-vinyl-3,5-dimethyl-2-piperidone, N-vinyl-4,4-dimethyl-2-piperidone, N-vinyl caprolactam (*aka*, N-vinyl-2-caprolactam), N-vinyl-3-methyl-2-caprolactam, N-vinyl-4-methyl-2-caprolactam, N-vinyl-7-methyl-2-caprolactam, N-vinyl-7-ethyl-2-caprolactam, N-vinyl-3,5-dimethyl-2-caprolactam, N-vinyl-4,6-dimethyl-2-caprolactam, N-vinyl-3,5,7-trimethyl-2-caprolactam, N-vinyl-N-methyl acetamide, N-vinyl formamide, N-vinyl acetamide, N-vinyl isopropylamide, N-vinyl-N-ethyl acetamide, N-vinyl-N-ethyl formamide, and mixtures thereof. Preferably, the N-vinyl amide monomer is N-vinylpyrrolidone, N-vinyl-N-methyl acetamide, or combinations thereof.

Examples of preferred methylene-containing ($=CH_2$) pyrrolidone monomers include without limitations 1-methyl-3-methylene-2-pyrrolidone, 1-ethyl-3-methylene-2-pyrrolidone, 1-methyl-5-methylene-2-pyrrolidone, 1-ethyl-5-methylene-2-pyrrolidone, 5-methyl-3-methylene-2-pyrrolidone, 5-ethyl-3-methylene-2-pyrrolidone, 1-n-propyl-3-methylene-2-pyrrolidone, 1-n-propyl-5-methylene-2-pyrrolidone, 1-isopropyl-3-methylene-2-pyrrolidone, 1-isopropyl-5-methylene-2-pyrrolidone, 1-n-butyl-3-methylene-2-pyrrolidone, 1-tert-butyl-3-methylene-2-pyrrolidone, and combinations thereof.

Examples of preferred acrylic monomers having a C_1 - C_4 alkoxyethoxy group include without limitation ethylene glycol methyl ether (meth)acrylate, di(ethylene glycol) methyl ether (meth)acrylate, tri(ethylene glycol) methyl ether (meth)acrylate, tetra(ethylene glycol) methyl ether (meth)acrylate, C_1 - C_4 -alkoxy poly(ethylene glycol) (meth)acrylate having a weight average molecular weight of up to 1500, methoxy-poly(ethylene glycol)ethyl (meth)acrylamide having a number average molecular weight of up to 1500, and combinations thereof.

Examples of preferred vinyl ether monomers include without limitation ethylene glycol monovinyl ether, di(ethylene glycol) monovinyl ether, tri(ethylene glycol) monovinyl ether, tetra(ethylene glycol) monovinyl ether, poly(ethylene glycol) monovinyl ether, ethylene glycol methyl vinyl ether, di(ethylene glycol) methyl vinyl ether, tri(ethylene glycol)

methyl vinyl ether, tetra(ethylene glycol) methyl vinyl ether, poly(ethylene glycol) methyl vinyl ether, and combinations thereof.

Examples of preferred allyl ether monomers include without limitation ethylene glycol monoallyl ether, di(ethylene glycol) monoallyl ether, tri(ethylene glycol) monoallyl ether, tetra(ethylene glycol) monoallyl ether, poly(ethylene glycol) monoallyl ether, ethylene glycol methyl allyl ether, di(ethylene glycol) methyl allyl ether, tri(ethylene glycol) methyl allyl ether, tetra(ethylene glycol) methyl allyl ether, poly(ethylene glycol) methyl allyl ether, and combinations thereof.

Examples of preferred phosphorylcholine-containing vinylic monomers include without limitation (meth)acryloyloxyethyl phosphorylcholine (*aka*, MPC, or 2-((meth)acryloyloxy)ethyl-2'-(trimethylammonio)ethylphosphate), (meth)acryloyloxypropyl phosphorylcholine (*aka*, 3-((meth)acryloyloxy)propyl-2'-(trimethylammonio)ethylphosphate), 4-((meth)acryloyloxy)butyl-2'-(trimethylammonio)ethylphosphate, 2-[(meth)acryloylamino]ethyl-2'-(trimethylammonio)-ethylphosphate, 3-[(meth)acryloylamino]propyl-2'-(trimethylammonio)ethylphosphate, 4-[(meth)acryloylamino]butyl-2'-(trimethylammonio)ethylphosphate, 5-((meth)acryloyloxy)pentyl-2'-(trimethylammonio)ethyl phosphate, 6-((meth)acryloyloxy)hexyl-2'-(trimethylammonio)-ethylphosphate, 2-((meth)acryloyloxy)ethyl-2'-(triethylammonio)ethylphosphate, 2-((meth)acryloyloxy)ethyl-2'-(tripropylammonio)ethylphosphate, 2-((meth)acryloyloxy)ethyl-2'-(tributylammonio)ethyl phosphate, 2-((meth)acryloyloxy)propyl-2'-(trimethylammonio)-ethylphosphate, 2-((meth)acryloyloxy)butyl-2'-(trimethylammonio)ethylphosphate, 2-((meth)acryloyloxy)pentyl-2'-(trimethylammonio)ethylphosphate, 2-((meth)acryloyloxy)hexyl-2'-(trimethylammonio)ethyl phosphate, 2-(vinylloxy)ethyl-2'-(trimethylammonio)ethylphosphate, 2-(allyloxy)ethyl-2'-(trimethylammonio)ethylphosphate, 2-(vinylloxycarbonyl)ethyl-2'-(trimethylammonio)ethyl phosphate, 2-(allyloxycarbonyl)ethyl-2'-(trimethylammonio)-ethylphosphate, 2-(vinylcarbonylamino)ethyl-2'-(trimethylammonio)ethylphosphate, 2-(allyloxycarbonylamino)ethyl-2'-(trimethylammonio)ethyl phosphate, 2-(butenoyloxy)ethyl-2'-(trimethylammonio)ethylphosphate, and combinations thereof.

In accordance with the invention, any non-silicone vinylic crosslinkers can be used in this invention. Examples of preferred non-silicone vinylic cross-linking agents include without limitation ethyleneglycol di-(meth)acrylate, diethyleneglycol di-(meth)acrylate, triethyleneglycol di-(meth)acrylate, tetraethyleneglycol di-(meth)acrylate, glycerol di-(meth)acrylate, 1,3-propanediol di-(meth)acrylate, 1,3-butanediol di-(meth)acrylate, 1,4-butanediol di-(meth)acrylate, glycerol 1,3-diglycerolate di-(meth)acrylate, ethylenebis[oxy(2-hydroxypropane-1,3-diyl)] di-(meth)acrylate, bis[2-(meth)acryloxyethyl]

phosphate, trimethylolpropane di-(meth)acrylate, and 3,4-bis[(meth)acryloyl]tetrahydrofuran, diacrylamide, dimethacrylamide, N,N-di(meth)acryloyl-N-methylamine, N,N-di(meth)acryloyl-N-ethylamine, N,N'-methylene bis(meth)acrylamide, N,N'-ethylene bis(meth)acrylamide, N,N'-dihydroxyethylene bis(meth)acrylamide, N,N'-propylene bis(meth)acrylamide, N,N'-2-hydroxypropylene bis(meth)acrylamide, N,N'-2,3-dihydroxybutylene bis(meth)acrylamide, 1,3-bis(meth)acrylamidepropane-2-yl dihydrogen phosphate, piperazine diacrylamide, tetraethyleneglycol divinyl ether, triethyleneglycol divinyl ether, diethyleneglycol divinyl ether, ethyleneglycol divinyl ether, triallyl isocyanurate, triallyl cyanurate, trimethylolpropane trimethacrylate, pentaerythritol tetramethacrylate, bisphenol A dimethacrylate, allylmethacrylate, allylacrylate, N-allylmethacrylamide, N-allylacrylamide, and combinations thereof. A preferred non-silicone vinylic cross-linking agent is tetra(ethyleneglycol) di-(meth)acrylate, tri(ethyleneglycol) di-(meth)acrylate, ethyleneglycol di-(meth)acrylate, di(ethyleneglycol) di-(meth)acrylate, tetraethyleneglycol divinyl ether, triethyleneglycol divinyl ether, diethyleneglycol divinyl ether, ethyleneglycol divinyl ether, triallyl isocyanurate, triallyl cyanurate, and combinations thereof.

Any thermal polymerization initiators can be used in the invention. Suitable thermal polymerization initiators are known to the skilled artisan and comprise, for example peroxides, hydroperoxides, azo-bis(alkyl- or cycloalkylnitriles), persulfates, percarbonates, or mixtures thereof. Examples of preferred thermal polymerization initiators include without limitation benzoyl peroxide, t-butyl peroxide, t-amyl peroxybenzoate, 2,2-bis(tert-butylperoxy)butane, 1,1-bis(tert-butylperoxy)cyclohexane, 2,5-Bis(tert-butylperoxy)-2,5-dimethylhexane, 2,5-bis(tert-butylperoxy)-2,5-dimethyl-3-hexyne, bis(1-(tert-butylperoxy)-1-methylethyl)benzene, 1,1-bis(tert-butylperoxy)-3,3,5-trimethylcyclohexane, di-t-butyl-diperoxyphthalate, t-butyl hydroperoxide, t-butyl peracetate, t-butyl peroxybenzoate, t-butylperoxy isopropyl carbonate, acetyl peroxide, lauroyl peroxide, decanoyl peroxide, dicetyl peroxydicarbonate, di(4-t-butylcyclohexyl)peroxy dicarbonate (Perkadox 16S), di(2-ethylhexyl)peroxy dicarbonate, t-butylperoxy pivalate (Lupersol 11); t-butylperoxy-2-ethylhexanoate (Trigonox 21-C50), 2,4-pentanedione peroxide, dicumyl peroxide, peracetic acid, potassium persulfate, sodium persulfate, ammonium persulfate, 2,2'-azobis(4-methoxy-2,4-dimethylvaleronitrile) (VAZO 33), 2,2'-Azobis[2-(2-imidazolin-2-yl)propane]dihydrochloride (VAZO 44), 2,2'-azobis(2-amidinopropane) dihydrochloride (VAZO 50), 2,2'-azobis(2,4-dimethylvaleronitrile) (VAZO 52), 2,2'-azobis(isobutyronitrile) (VAZO 64 or AIBN), 2,2'-azobis-2-methylbutyronitrile (VAZO 67), 1,1-azobis(1-cyclohexanecarbonitrile) (VAZO 88); 2,2'-azobis(2-cyclopropylpropionitrile), 2,2'-azobis(methylisobutyrate), 4,4'-Azobis(4-cyanovaleric acid), and combinations thereof. Preferably, the thermal initiator is 2,2'-azobis(isobutyronitrile) (AIBN or VAZO 64).

Suitable photoinitiators are benzoin methyl ether, diethoxyacetophenone, a benzoylphosphine oxide, 1-hydroxycyclohexyl phenyl ketone and Darocur and Irgacur types, preferably Darocur 1173® and Darocur 2959®, Germanium-based Norrish Type I photoinitiators (e.g., those described in US 7,605,190). Examples of benzoylphosphine initiators include 2,4,6-trimethylbenzoyldiphenylphosphine oxide; bis-(2,6-dichlorobenzoyl)-4-N-propylphenylphosphine oxide; and bis-(2,6-dichlorobenzoyl)-4-N-butylphenylphosphine oxide. Reactive photoinitiators which can be incorporated, for example, into a macromer or can be used as a special monomer are also suitable. Examples of reactive photoinitiators are those disclosed in EP 632 329, herein incorporated by reference in its entirety.

Suitable photoinitiators are benzoin methyl ether, diethoxyacetophenone, a benzoylphosphine oxide, 1-hydroxycyclohexyl phenyl ketone and Darocur and Irgacur types, preferably Darocur 1173® and Darocur 2959®, Germane-based Norrish Type I photoinitiators. Examples of benzoylphosphine initiators include 2,4,6-trimethylbenzoyldiphenylphosphine oxide; bis-(2,6-dichlorobenzoyl)-4-N-propylphenylphosphine oxide; and bis-(2,6-dichlorobenzoyl)-4-N-butylphenylphosphine oxide. Reactive photoinitiators, which can be incorporated, for example, into a macromer or can be used as a special monomer, are also suitable. Examples of reactive photoinitiators are those disclosed in EP 632 329.

Where a vinylic monomer capable of absorbing ultra-violet radiation and high energy violet light (HEVL) is used in the invention, a Germane-based Norrish Type I photoinitiator and a light source including a light in the region of about 400 to about 550 nm are preferably used to initiate a free-radical polymerization. Any Germane-based Norrish Type I photoinitiators can be used in this invention, so long as they are capable of initiating a free-radical polymerization under irradiation with a light source including a light in the region of about 400 to about 550 nm. Examples of Germane-based Norrish Type I photoinitiators are acylgermanium compounds described in US 7,605,190.

In accordance with a preferred embodiment of the invention, a fluid polymerizable composition of the invention preferably can further comprise at least one blending vinylic monomer. Examples of preferred blending vinylic monomers include C₁-C₁₀ alkyl (meth)acrylate (e.g., methyl (meth)acrylate, ethyl (meth)acrylate, propyl (meth)acrylate, isopropyl (meth)acrylate, 2-ethylhexyl (meth)acrylate, pentyl (meth)acrylate, hexyl (meth)acrylate, etc.), cyclopentylacrylate, cyclohexylmethacrylate, cyclohexylacrylate, isobornyl (meth)acrylate, styrene, 4,6-trimethylstyrene, t-butyl styrene, trifluoroethyl (meth)acrylate, hexafluoro-isopropyl (meth)acrylate, hexafluorobutyl (meth)acrylate, or combinations thereof. Methyl methacrylate is the most preferred blending vinylic monomer.

In accordance with a preferred embodiment of the invention, a fluid polymerizable composition of the invention can further comprise at least one UV-absorbing vinylic monomer and optionally (but preferably) at least one UV/HEVL-absorbing vinylic monomer. The term "UV/HEVL-absorbing vinylic monomer" refers to a vinylic monomer that can absorb UV light and high-energy-violet-light (i.e., light having wavelength between 380 nm and 440 nm).

Any suitable UV-absorbing vinylic monomers and UV/HEVL-absorbing vinylic monomers can be used in a fluid polymerizable composition for preparing a polymer of the invention. Examples of preferred UV-absorbing and UV/HEVL-absorbing vinylic monomers include without limitation: 2-(2-hydroxy-5-vinylphenyl)-2H-benzotriazole, 2-(2-hydroxy-5-acryloyloxyphenyl)-2H-benzotriazole, 2-(2-hydroxy-3-methacrylamido methyl-5-tert octylphenyl) benzotriazole, 2-(2'-hydroxy-5'-methacrylamidophenyl)-5-chlorobenzotriazole, 2-(2'-hydroxy-5'-methacrylamidophenyl)-5-methoxybenzotriazole, 2-(2'-hydroxy-5'-methacryloxypropyl-3'-t-butyl-phenyl)-5-chlorobenzotriazole, 2-(2'-hydroxy-5'-methacryloxypropylphenyl) benzotriazole, 2-hydroxy-5-methoxy-3-(5-(trifluoromethyl)-2H-benzo[d][1,2,3]triazol-2-yl)benzyl methacrylate (WL-1), 2-hydroxy-5-methoxy-3-(5-methoxy-2H-benzo[d][1,2,3]triazol-2-yl)benzyl methacrylate (WL-5), 3-(5-fluoro-2H-benzo[d][1,2,3]triazol-2-yl)-2-hydroxy-5-methoxybenzyl methacrylate (WL-2), 3-(2H-benzo[d][1,2,3]triazol-2-yl)-2-hydroxy-5-methoxybenzyl methacrylate (WL-3), 3-(5-chloro-2H-benzo[d][1,2,3]triazol-2-yl)-2-hydroxy-5-methoxybenzyl methacrylate (WL-4), 2-hydroxy-5-methoxy-3-(5-methyl-2H-benzo[d][1,2,3]triazol-2-yl)benzyl methacrylate (WL-6), 2-hydroxy-5-methyl-3-(5-(trifluoromethyl)-2H-benzo[d][1,2,3]triazol-2-yl)benzyl methacrylate (WL-7), 4-allyl-2-(5-chloro-2H-benzo[d][1,2,3]triazol-2-yl)-6-methoxyphenol (WL-8), 2-{2'-Hydroxy-3'-tert-5'[3''-(4''-vinylbenzyloxy)propoxy]phenyl}-5-methoxy-2H-benzotriazole, phenol, 2-(5-chloro-2H-benzotriazol-2-yl)-6-(1,1-dimethylethyl)-4-ethenyl- (UVAM), 2-[2'-hydroxy-5'-(2-methacryloxyethyl)phenyl]-2H-benzotriazole (2-Propenoic acid, 2-methyl-, 2-[3-(2H-benzotriazol-2-yl)-4-hydroxyphenyl]ethyl ester, Norbloc), 2-{2'-Hydroxy-3'-tert-butyl-5'-[3'-methacryloyloxypropoxy]phenyl}-2H-benzotriazole, 2-{2'-Hydroxy-3'-tert-butyl-5'-[3'-methacryloyloxypropoxy]phenyl}-5-methoxy-2H-benzotriazole (UV13), 2-{2'-Hydroxy-3'-tert-butyl-5'-[3'-methacryloyloxypropoxy]phenyl}-5-chloro-2H-benzotriazole (UV28), 2-[2'-Hydroxy-3'-tert-butyl-5'-(3'-acryloyloxypropoxy)phenyl]-5-trifluoromethyl-2H-benzotriazole (UV23), 2-(2'-hydroxy-5-methacrylamidophenyl)-5-methoxybenzotriazole (UV6), 2-(3-allyl-2-hydroxy-5-methylphenyl)-2H-benzotriazole (UV9), 2-(2-Hydroxy-3-methyl-5-methylphenyl)-2H-benzotriazole (UV12), 2-3'-t-butyl-2'-hydroxy-5'-(3''-dimethylvinylsilylpropoxy)-2'-hydroxy-phenyl)-5-methoxybenzotriazole (UV15), 2-(2'-hydroxy-5'-methacryloylpropyl-3'-tert-butyl-phenyl)-5-methoxy-2H-benzotriazole (UV16), 2-(2'-hydroxy-5'-acryloylpropyl-3'-tert-butyl-phenyl)-5-methoxy-2H-benzotriazole (UV16A), 2-

Methylacrylic acid 3-[3-*tert*-butyl-5-(5-chlorobenzotriazol-2-yl)-4-hydroxyphenyl]-propyl ester (16-100, CAS#96478-15-8), 2-(3-(*tert*-butyl)-4-hydroxy-5-(5-methoxy-2H-benzo[d][1,2,3]triazol-2-yl)phenoxy)ethyl methacrylate (16-102); Phenol, 2-(5-chloro-2H-benzotriazol-2-yl)-6-methoxy-4-(2-propen-1-yl) (CAS#1260141-20-5); 2-[2-Hydroxy-5-[3-(methacryloyloxy)propyl]-3-*tert*-butylphenyl]-5-chloro-2H-benzotriazole; Phenol, 2-(5-ethenyl-2H-benzotriazol-2-yl)-4-methyl-, homopolymer (9Cl) (CAS#83063-87-0).

In a preferred embodiment, a fluid polymerizable composition of the invention comprises a UV-absorbing vinylic monomer and a UV/HEVL absorbing vinylic monomer. More preferably, the UV-absorbing vinylic monomer is 2-[2'-hydroxy-5'-(2-methacryloxyethyl)phenyl]-2H-benzotriazole (Norbloc), and the UV/HEVL absorbing vinylic monomer is 2-[2'-Hydroxy-3'-*tert*-butyl-5'-[3'-methacryloyloxypropoxy]phenyl]-2H-benzotriazole, 2-[2'-Hydroxy-3'-*tert*-butyl-5'-[3'-methacryloyloxypropoxy]phenyl]-5-methoxy-2H-benzotriazole (UV13), 2-[2'-Hydroxy-3'-*tert*-butyl-5'-[3'-methacryloyloxypropoxy]phenyl]-5-chloro-2H-benzotriazole (UV28), 2-[2'-Hydroxy-3'-*tert*-butyl-5'-(3'-acryloyloxypropoxy)phenyl]-5-trifluoromethyl-2H-benzotriazole (UV23), or combinations thereof. The resultant SiHy contact lens is characterized by having the UVB transmittance of about 10% or less (preferably about 5% or less, more preferably about 2.5% or less, even more preferably about 1% or less) between 280 and 315 nanometers and a UVA transmittance of about 30% or less (preferably about 20% or less, more preferably about 10% or less, even more preferably about 5% or less) between 315 and 380 nanometers and a Violet transmittance of about 70% or less, preferably about 60% or less, more preferably about 50% or less, even more preferably about 40% or less) between 380 nm and 440 nm.

In accordance with the invention, a fluid polymerizable composition of the invention comprises: (a) from about 20% to about 79% (preferably from about 20% to about 75%, more preferably from about 25% to about 70%, even more preferably from about 30% to about 65%) by weight of the silicone-containing vinylic monomer and/or the silicone-containing vinylic crosslinker; (b) 20% to about 79% (preferably from about 20% to about 75%, more preferably from about 25% to about 70%, even more preferably from about 30% to about 65%) by weight of the hydrophilic vinylic monomer; (c) from 0 to about 2.5% (preferably from 0 to about 2.0%, more preferably from about 0.01% to about 1.5%, even more preferably from about 0.05% to about 1.0%) by weight of the non-silicone vinylic crosslinker; (d) from about 0.05% to about 2.0% (preferably from about 0.1% to about 2.0%, more preferably from about 0.2% to about 1.5%, even more preferably from about 0.3% to about 1.2%) by weight of the free-radical initiator; (e) from 0 to about 25% (preferably from 0 to about 20%, more preferably from about 2.5% to about 17.5%, even more preferably from about 5% to about 15%) by weight of the blending vinylic monomer;

and (f) from 0 to about 3.0%, preferably about 0.1% to about 2.5%, more preferably about 0.2% to about 2.0%, by weight of the UV-absorbing vinylic monomer and/or the UV/HEVL-absorbing vinylic monomer, relative to the total amount of all polymerizable materials in the fluid polymerizable composition, provided that the sum of the amounts of polymerizable materials (a) to (f) and other not-listed polymerizable materials is 100%. Preferably, the sum of the amounts of polymerizable materials (a) and (b) is at least 70% (preferably at least 75%, more preferably at least 80, even more preferably at least 85%) by weight relative to the total amount of all polymerizable materials in the fluid polymerizable composition.

A fluid polymerizable composition of the invention can also comprise other necessary components known to a person skilled in the art, such as, for example, a visibility tinting agent (e.g., one or more polymerizable dyes, pigments, or mixtures thereof), antimicrobial agents (e.g., preferably silver nanoparticles), a bioactive agent, leachable lubricants, leachable tear-stabilizing agents, and mixtures thereof, as known to a person skilled in the art.

A fluid polymerizable composition (SiHy lens formulation) of the invention is a solution prepared by dissolving all of the desirable components in the organic solvent.

Lens molds for making contact lenses are well known to a person skilled in the art and, for example, are employed in cast molding or spin casting. For example, a mold (for cast molding) generally comprises at least two mold sections (or portions) or mold halves, i.e. first and second mold halves. The first mold half defines a first molding (or optical) surface and the second mold half defines a second molding (or optical) surface. The first and second mold halves are configured to receive each other such that a lens forming cavity is formed between the first molding surface and the second molding surface. The molding surface of a mold half is the cavity-forming surface of the mold and in direct contact with lens-forming material.

Methods of manufacturing mold sections for cast-molding a contact lens are generally well known to those of ordinary skill in the art. The process of the present invention is not limited to any particular method of forming a mold. In fact, any method of forming a mold can be used in the present invention. The first and second mold halves can be formed through various techniques, such as injection molding or lathing. Examples of suitable processes for forming the mold halves are disclosed in U.S. Pat. Nos. 4,444,711; 4,460,534; 5,843,346; and 5,894,002.

Virtually all materials known in the art for making molds can be used to make molds for making contact lenses. For example, polymeric materials, such as polyethylene, polypropylene, polystyrene, PMMA, Topas® COC grade 8007-S10 (clear amorphous copolymer of ethylene and norbornene, from Ticona GmbH of Frankfurt, Germany and

Summit, New Jersey), or the like can be used. Other materials that allow UV light transmission could be used, such as quartz glass and sapphire.

In accordance with the invention, a mold used in the invention is preferably made of a hydrophobic polymeric material (i.e., having a water contact angle by static sessile drop method of about 100 degrees or larger), more preferably made of a polypropylene.

In accordance with the invention, the fluid polymerizable composition can be introduced (dispensed) into a cavity formed by a mold according to any known methods.

After the fluid polymerizable composition is dispensed into the mold, it is polymerized to produce a SiHy contact lens precursor. Crosslinking may be initiated thermally or actinically (but preferably is initiated thermally) to crosslink the polymerizable materials in the fluid polymerizable composition.

The thermal polymerization is carried out conveniently in an above-mentioned solvent at elevated temperature, for example at a temperature of from 25 to 100°C and preferably 40 to 100°C. The reaction time may vary within wide limits, but is conveniently, for example, from 1 to 24 hours or preferably from 2 to 12 hours. It is advantageous to previously degas the components and solvents used in the polymerization reaction and to carry out copolymerization reaction under an inert atmosphere, for example under a nitrogen or argon atmosphere.

The actinic polymerization can then be triggered off by actinic radiation, for example light, in particular UV light or visible light of a suitable wavelength. The spectral requirements can be controlled accordingly, if appropriate, by addition of suitable photosensitizers.

Opening of the mold so that the molded article (i.e., the SiHy contact lens precursor) can be removed from the mold may take place in a manner known *per se*.

In accordance with the invention, the extraction liquid medium is any solvent capable of dissolving the organic solvent, unpolymerized polymerizable materials, and oligomers in the dry contact lens. Water, any organic solvents known to a person skilled in the art, or a mixture thereof can be used in the invention. Preferably, the organic solvents used extraction liquid medium are water, a buffered saline, a C₁-C₃ alkyl alcohol, 1,2-propylene glycol, a polyethyleneglycol having a number average molecular weight of about 400 Daltons or less, a C₁-C₆ alkylalcohol, or combinations thereof.

The extracted contact lens can then be hydrated according to any method known to a person skilled in the art.

The hydrated soft contact lens can further subject to further processes, such as, for example, surface treatment, packaging in lens packages with a packaging solution which is well known to a person skilled in the art; sterilization such as autoclave at from 118 to 124°C for at least about 30 minutes; and the like.

A hydrated soft contact lens of the invention has: an oxygen permeability of at least about 40 barrers (preferably at least about 60 barrers, more preferably at least about 80 barrers); an elastic modulus of from about 0.2 MPa to about 2.0 MPa (preferably from about 0.3 MPa to about 1.5 MPa or less, more preferably from about 0.4 MPa to about 1.2 or less); an equilibrium water content of from about 15% to about 75% (preferably from about 20% to about 70%, more preferably from about 30% to about 65%) by weight when fully hydrated.

Although various embodiments of the invention have been described using specific terms, devices, and methods, such description is for illustrative purposes only. The words used are words of description rather than of limitation. As would be obvious to one skilled in the art, many variations and modifications of the invention may be made by those skilled in the art without departing from the spirit and scope of the novel concepts of the disclosure. In addition, it should be understood that aspects of the various embodiments of the invention may be interchanged either in whole or in part or can be combined in any manner and/or used together, as illustrated below:

1. A method for making contact lenses, comprising the steps of:
 - (1) obtaining a fluid polymerizable composition, wherein the composition comprises an organic solvent and dissolved therein polymerizable materials, wherein the polymerizable materials comprises (a) from about 20% to about 79% by weight of at least one silicone-containing vinylic monomer and/or at least one silicone-containing vinylic crosslinker; (b) from 20% to about 79% by weight of at least one hydrophilic vinylic monomer; (c) from 0 to about 2.5% by weight of at least one non-silicone vinylic crosslinker; (d) from about 0.05% to about 2.0% by weight of at least one free-radical initiator; (e) from 0 to about 25% by weight of a blending vinylic monomer; and (f) from 0 to about 3.0% by weight of at least one UV-absorbing vinylic monomer and/or at least one UV/HEVL-absorbing vinylic monomer, relative to the total amount of all polymerizable materials in the fluid polymerizable composition, provided that the sum of the amounts of polymerizable materials (a) to (f) and other not-listed polymerizable materials is 100%, wherein the organic solvent has a boiling point of at least 95°C or higher and is present in the polymerizable composition in an amount of from about 4% to about 15% by weight relative to the total amount of the fluid polymerizable composition and is capable of functioning as a processing-aid tool for controlling lens diameter of a hydrated soft contact lens made from the polymerizable composition;
 - (2) introducing the fluid polymerizable composition into a cavity formed by a mold,

wherein the mold has a first mold half with a first molding surface defining the anterior surface of a contact lens and a second mold half with a second molding surface defining the posterior surface of the contact lens, wherein said first and second mold halves are configured to receive each other such that a cavity is formed between said first and second molding surfaces;

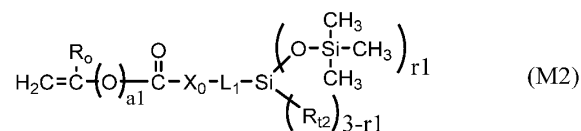
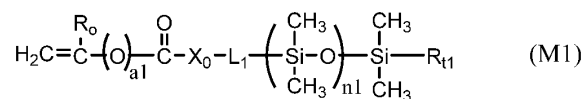
- (3) curing thermally or actinically the fluid polymerizable composition in the mold to form a dry contact lens;
- (4) extracting the dry contact lens with a liquid extraction medium to remove the organic solvent, unpolymerized polymerizable components and oligomers from the dry lens to obtain an extracted contact lens; and
- (5) hydrating the extracted contact lens with water or an aqueous solution to obtain the hydrated soft contact lens,

wherein the amount of the organic solvent in the polymerizable composition is adjusted to achieve a targeted lens diameter of the hydrated soft contact lens without affecting adversely the equilibrium water content of the hydrated soft contact lens.

- 2. The method of embodiment 1, wherein the step of curing is carried out thermally.
- 3. The method of embodiment 1 or 2, wherein the organic solvent has a boiling point of at least 100°C or higher.
- 4. The method of embodiment 1 or 2, wherein the organic solvent has a boiling point of at least 105°C or higher (preferably at least 105°C or higher).
- 5. The method of embodiment 1 or 2, wherein the organic solvent has a boiling point of at least 110°C or higher.
- 6. The method of any one of embodiments 1 to 5, wherein the organic solvent is present in the polymerizable composition in an amount of from about 5.0% to about 12.5% by weight relative to the total amount of the fluid polymerizable composition.
- 7. The method of any one of embodiments 1 to 5, wherein the organic solvent is present in the polymerizable composition in an amount of from about 5.5% to about 11% by weight relative to the total amount of the fluid polymerizable composition.
- 8. The method of any one of embodiments 1 to 5, wherein the organic solvent is present in the polymerizable composition in an amount of from about 5.5% to about 9.5% by weight relative to the total amount of the fluid polymerizable composition.
- 9. The method of any one of embodiments 1 to 8, wherein the organic solvent is a C₃-C₈ alkyl alcohol.
- 10. The method of any one of embodiments 1 to 8, wherein the organic solvent is a C₄-C₈ alkyl alcohol.
- 11. The method of any one of embodiments 1 to 8, wherein the organic solvent is a C₄-C₆ alkyl alcohol.

12. The method of any one of embodiments 1 to 11, wherein the polymerizable materials comprises from about 20% to about 75% by weight of said at least one silicone-containing vinylic monomer and/or said at least one silicone-containing vinylic crosslinker.
13. The method of any one of embodiments 1 to 11, wherein the polymerizable materials comprises from about 25% to about 70% by weight of said at least one silicone-containing vinylic monomer and/or said at least one silicone-containing vinylic crosslinker.
14. The method of any one of embodiments 1 to 11, wherein the polymerizable materials comprises from about 30% to about 65% by weight of said at least one silicone-containing vinylic monomer and/or said at least one silicone-containing vinylic crosslinker.
15. The method of any one of embodiments 1 to 14, wherein the polymerizable materials comprises from about 20% to about 75% by weight of said at least one hydrophilic vinylic monomer.
16. The method of any one of embodiments 1 to 14, wherein the polymerizable materials comprises from about 25% to about 70% by weight of said at least one hydrophilic vinylic monomer.
17. The method of any one of embodiments 1 to 14, wherein the polymerizable materials comprises from about 30% to about 65% by weight of said at least one hydrophilic vinylic monomer.
18. The method of any one of embodiments 1 to 17, wherein the polymerizable materials comprises from 0 to about 2.0% by weight of said at least one non-silicone vinylic crosslinker.
19. The method of any one of embodiments 1 to 17, wherein the polymerizable materials comprises from about 0.01% to about 1.5% by weight of said at least one non-silicone vinylic crosslinker.
20. The method of any one of embodiments 1 to 17, wherein the polymerizable materials comprises from about 0.05% to about 1.0% by weight of said at least one non-silicone vinylic crosslinker.
21. The method of any one of embodiments 1 to 20, wherein the polymerizable materials comprises from about 0.1% to about 2.0% by weight of said at least one free-radical initiator.
22. The method of any one of embodiments 1 to 20, wherein the polymerizable materials comprises from about 0.2% to about 1.5% by weight of said at least one free-radical initiator.

23. The method of any one of embodiments 1 to 20, wherein the polymerizable materials comprises from about 0.3% to about 1.2% by weight of said at least one free-radical initiator.
24. The method of any one of embodiments 1 to 23, wherein the polymerizable materials comprises from 0 to about 20% by weight of the blending vinylic monomer.
25. The method of any one of embodiments 1 to 23, wherein the polymerizable materials comprises from about 2.5% to about 17.5% by weight of the blending vinylic monomer.
26. The method of any one of embodiments 1 to 23, wherein the polymerizable materials comprises from about 5% to about 15% by weight of the blending vinylic monomer.
27. The method of any one of embodiments 1 to 26, wherein the polymerizable materials comprises from about 0.1% to about 2.5% by weight of said at least one UV-absorbing vinylic monomer and/or said at least one UV/HEVL-absorbing vinylic monomer.
28. The method of any one of embodiments 1 to 26, wherein the polymerizable materials comprises from about 0.2% to about 2.0% by weight of said at least one UV-absorbing vinylic monomer and/or said at least one UV/HEVL-absorbing vinylic monomer.
29. The method of any one of embodiments 1 to 28, wherein the polymerizable materials comprise at least one silicone containing vinylic monomer which is a vinylic monomer having a bis(trialkylsilyloxy)alkylsilyl group or a tris(trialkylsilyloxy)silyl group, a polysiloxane vinylic monomer, a polycarbosiloxane vinylic monomer, 3-methacryloxy propylpentamethyldisiloxane, t-butyldimethyl-siloxyethyl vinyl carbonate, trimethylsilylethyl vinyl carbonate, and trimethylsilylmethyl vinyl carbonate, and combinations thereof.
30. The method of any one of embodiments 1 to 28, wherein the polymerizable materials comprise at least one silicone containing vinylic monomer which is a vinylic monomer of formula (M1) or (M2)



in which: a1 is zero or 1; R₀ is H or methyl; X₀ is O or NR₁; L₁ is a C₂-C₈ alkylene

divalent radical or a divalent radical of $-\text{L}_1'-\text{X}_1-\text{L}_1''-$, $-\text{C}_2\text{H}_4\text{O}-\text{L}_1''-$,

$\text{-(C}_2\text{H}_4\text{O)}_{q1}\text{-CONH-L}_1\text{''-}$, $\text{-L}_1\text{'-NHCOO-(C}_2\text{H}_4\text{O)}_{q1}\text{-L}_1\text{''-}$,
 $\text{-CH}_2\text{-CH(OH)-CH}_2\text{-X}_1\text{'-(C}_2\text{H}_4\text{O)}_{q2}\text{-L}_1\text{''-}$, $\text{-L}_1\text{'-X}_1\text{'-CH}_2\text{-CH(OH)-CH}_2\text{-O-L}_1\text{''-}$, or
 $\text{-(C}_2\text{H}_4\text{O)}_{q1}\text{-CH}_2\text{-CH(OH)-CH}_2\text{-O-L}_1\text{''-}$; $\text{L}_1\text{'}$ is a $\text{C}_2\text{-C}_8$ alkylene divalent radical which
 has zero or one hydroxyl group; $\text{L}_1\text{'}$ is $\text{C}_3\text{-C}_8$ alkylene divalent radical which has zero
 or one hydroxyl group; X_1 is O, NR_1 , NHCOO , OCONH , CONR_1 , or NR_1CO ; R_1 is H or
 a $\text{C}_1\text{-C}_4$ alkyl having 0 to 2 hydroxyl group; R_{11} and R_{12} independent of each other are
 a $\text{C}_1\text{-C}_4$ alkyl; $\text{X}_1\text{'}$ is O or NR_1 ; q_1 is an integer of 1 to 20; q_2 is an integer of 0 to 20;
 n_1 is an integer of 3 to 25; and r_1 is an integer of 2 or 3.

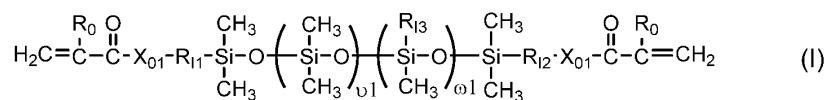
31. The method of any one of embodiments 1 to 28, wherein the polymerizable materials comprise at least one silicone containing vinylic monomer which is
- tris(trimethylsilyloxy)silylpropyl (meth)acrylate, [3-(meth)acryloxy-2-hydroxypropyloxy]propylbis(trimethylsiloxy) methylsilane, [3-(meth)acryloxy-2-hydroxypropyloxy]propylbis(trimethylsiloxy)butylsilane, 3-(meth)acryloxy-2-(2-hydroxyethoxy)-propyloxy]propylbis(trimethylsiloxy)methylsilane, 3-(meth)acryloxy-2-hydroxypropyloxy]propyltris(trimethylsiloxy)silane, N-[tris(trimethylsiloxy)silylpropyl]-(meth)acrylamide, N-(2-hydroxy-3-(3-(bis(trimethylsilyloxy)methylsilyl)propyloxy)propyl)-2-methyl (meth)acrylamide, N-(2-hydroxy-3-(3-(bis(trimethylsilyloxy)methylsilyl)propyloxy)propyl) (meth)acrylamide, N-(2-hydroxy-3-(3-(tris(trimethylsilyloxy)silyl)propyloxy)propyl)-2-methyl acrylamide, N-(2-hydroxy-3-(3-(tris(trimethylsilyloxy)silyl)propyloxy)propyl) (meth)acrylamide, N-[tris(dimethylpropylsiloxy)silylpropyl]-(meth)acrylamide, N-[tris(dimethylphenylsiloxy)silylpropyl] (meth)acrylamide, N-[tris(dimethylethylsiloxy)silylpropyl] (meth)acrylamide, N,N-bis[2-hydroxy-3-(3-(bis(trimethylsilyloxy)methylsilyl)propyloxy)propyl]-2-methyl (meth)acrylamide, N,N-bis[2-hydroxy-3-(3-(bis(trimethylsilyloxy)methylsilyl)propyloxy)-propyl] (meth)acrylamide, N,N-bis[2-hydroxy-3-(3-(tris(trimethylsilyloxy)silyl)propyloxy)-propyl]-2-methyl (meth)acrylamide, N,N-bis[2-hydroxy-3-(3-(tris(trimethylsilyloxy)silyl)propyloxy)propyl] (meth)acrylamide, N-[2-hydroxy-3-(3-(t-butyl dimethylsilyl)-propyloxy)propyl]-2-methyl (meth)acrylamide, N-[2-hydroxy-3-(3-(t-butyl dimethylsilyl)propyloxy)propyl] (meth)acrylamide, N,N-bis[2-hydroxy-3-(3-(t-butyl dimethylsilyl)propyloxy)propyl]-2-methyl (meth)acrylamide, N-2-(meth)acryloxyethyl-O-(methyl-bis-trimethylsiloxy-3-propyl)silyl carbamate, 3-(trimethylsilyl)propylvinyl carbonate, 3-(vinyl oxycarbonylthio)propyl-tris(trimethylsiloxy)silane, 3-[tris(trimethylsiloxy)silyl]propylvinyl carbamate, 3-[tris(trimethylsiloxy)silyl] propyl allyl carbamate, 3-[tris(trimethylsiloxy)silyl]propyl vinyl

carbonate, or a combination thereof.

32. The method of any one of embodiments 1 to 28, wherein the polymerizable materials comprise at least one silicone containing vinylic monomer which is α -(meth)acryloxypropyl terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -(meth)acryloxy-2-hydroxypropyloxypropyl terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -(2-hydroxyl-methacryloxypropyloxypropyl)- ω -C₁-C₄-alkyl-decamethylpentasiloxane, α -[3-(meth)acryloxyethoxy-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxy-propyloxy-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxyisopropyloxy-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxybutyloxy-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxyethylamino-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxypropylamino-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloxy-butylamino-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -(meth)acryloxy(polyethylenoxy)-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[(meth)acryloxy-2-hydroxypropyloxy-ethoxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[(meth)acryloxy-2-hydroxypropyl-N-ethylaminopropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[(meth)acryloxy-2-hydroxypropyl-aminopropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[(meth)acryloxy-2-hydroxypropyloxy-(polyethylenoxy)propyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -(meth)acryloylamidopropyloxypropyl terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -N-methyl-(meth)acryloylamidopropyloxypropyl terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acrylamidoethoxy-2-hydroxypropyloxy-propyl]-terminated ω -C₁-C₄-alkyl polydimethylsiloxane, α -[3-(meth)acrylamidopropyloxy-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acrylamidoisopropyloxy-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acrylamidobutyloxy-2-hydroxypropyloxypropyl]-terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, α -[3-(meth)acryloylamido-2-hydroxypropyloxypropyl] terminated ω -C₁-C₄-alkyl polydimethylsiloxane, α -[3-[N-methyl-(meth)acryloylamido]-2-hydroxypropyloxypropyl] terminated ω -C₁-C₄-alkyl terminated polydimethylsiloxane, N-methyl-N'-(propyltetra(dimethylsiloxy)dimethylbutylsilane) (meth)acrylamide, N-(2,3-

dihydroxypropane)-N'-(propyltetra(dimethylsiloxy)dimethylbutylsilane)
(meth)acrylamide, (meth)acryloylamidopropyltetra(dimethylsiloxy)dimethylbutylsilane,
 α -vinyl carbonate-terminated ω -C₁-C₄-alkyl-terminated polydimethylsiloxanes, α -vinyl
carbamate-terminated ω -C₁-C₄-alkyl-terminated polydimethylsiloxane, or a mixture
thereof.

33. The method of any one of embodiments 1 to 32, wherein the polymerizable materials comprise at least one silicone-containing vinylic crosslinker which is a polysiloxane vinylic crosslinker, a polycarbosiloxane vinylic crosslinker, or a combination thereof.
34. The method of any one of embodiments 1 to 32, wherein the polymerizable materials comprise at least one silicone-containing vinylic crosslinker which is a di-(meth)acryloyl-terminated polydimethylsiloxane, a di-vinyl carbonate-terminated polydimethylsiloxane; a di-vinyl carbamate-terminated polydimethylsiloxane; N,N,N',N'-tetrakis(3-methacryloxy-2-hydroxypropyl)- α,ω -bis-3-aminopropyl-polydimethylsiloxane, or a combination thereof.
35. The method of any one of embodiments 1 to 32, wherein the polymerizable materials comprise at least one silicone-containing vinylic crosslinker which is a vinylic crosslinker of formula (I)



in which:

$\omega 1$ is an integer of from 30 to 500 and $\omega 1$ is an integer of from 1 to 75, provided that

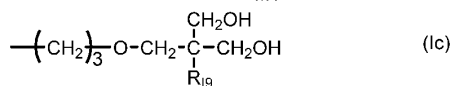
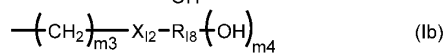
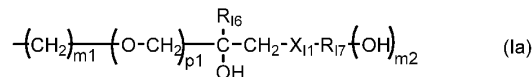
$\omega 1/\omega 1$ is from about 0.035 to about 0.15;

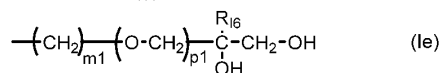
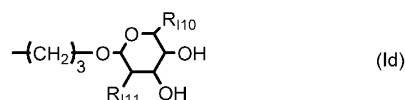
X_{01} is O or NR_N in which R_N is hydrogen or C₁-C₁₀-alkyl;

R_0 is hydrogen or methyl;

R_{11} and R_{12} independently of each other are a substituted or unsubstituted C₁-C₁₀ alkylene divalent radical or a divalent radical of $-\text{R}_{14}-\text{O}-\text{R}_{15}-$ in which R_{14} and R_{15} independently of each other are a substituted or unsubstituted C₁-C₁₀ alkylene divalent radical;

R_{13} is a monovalent radical of any one of formula (Ia) to (Ie)





p1 is zero or 1; m1 is an integer of 2 to 4; m2 is an integer of 1 to 5; m3 is an integer of 3 to 6; m4 is an integer of 2 to 5;

R₁₆ is hydrogen or methyl;

R₁₇ is a C₂-C₆ hydrocarbon radical having (m2+1) valencies;

R₁₈ is a C₂-C₆ hydrocarbon radical having (m4+1) valencies;

R₁₉ is ethyl or hydroxymethyl;

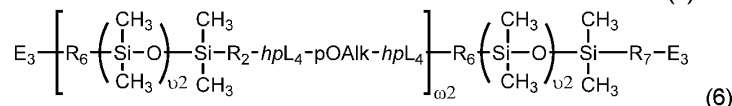
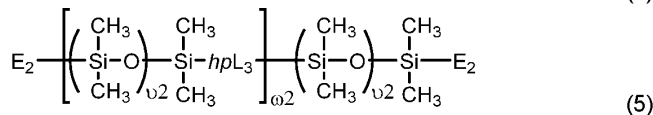
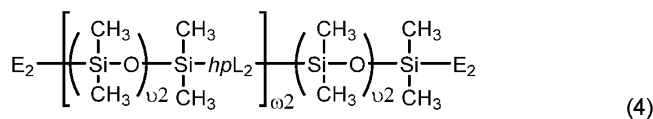
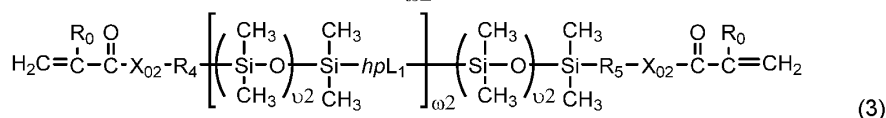
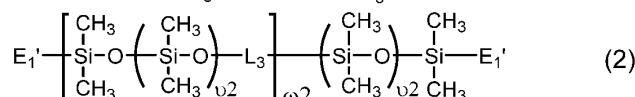
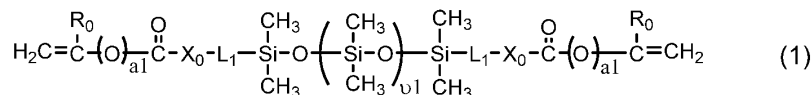
R₁₁₀ is methyl or hydromethyl;

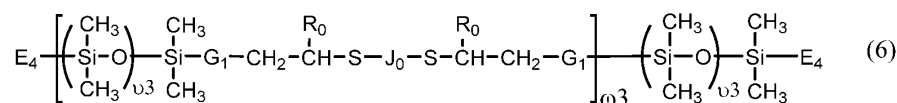
R₁₁₁ is hydroxyl or methoxy;

X₁₁ is a sulfur linkage of ---S--- or a tertiary amino linkage of $\text{---NR}_{112}\text{---}$ in which R₁₁₂ is C₁-C₁ alkyl, hydroxyethyl, hydroxypropyl, or 2,3-dihydroxypropyl; and

X₁₂ is an amide linkage of $\text{---NR}_{113}\text{---}\overset{\text{O}}{\parallel}\text{C---}$ or $\text{---}\overset{\text{O}}{\parallel}\text{C---NR}_{113}\text{---}$ in which R₁₁₃ is hydrogen or C₁-C₁₀ alkyl.

36. The method of any one of embodiments 1 to 32, wherein the polymerizable materials comprise at least one silicone-containing vinylic crosslinker which is a vinylic crosslinker of any one of formula (1) to (7)





in which:

$\nu 1$ is an integer of from 30 to 500;

$\nu 2$ is an integer of from 5 to 50;

$\nu 3$ is an integer of from 5 to 100;

$\omega 2$ and $\omega 3$ independent of each other are an integer of from 1 to 15;

a_1 and g_1 independent of each other is zero or 1;

h_1 is an integer of 1 to 20 and h_2 is an integer of 0 to 20;

m_1 and m_3 independent of each other is 0 or 1, m_2 is an integer of 1 to 6, m_4 is an integer of 1 to 5, m_5 is 2 or 3;

q_1 is an integer of 1 to 20, q_2 is an integer of 0 to 20, q_3 is an integer of 0 to 2, q_4 is an integer of 2 to 50, q_5 and q_6 independent of each other are a number of 0 to 35, provided that $(q_4+q_5+q_6)$ is an integer of 2 to 50;

$x+y$ is an integer of from 10 to 30;

e_1 is an integer of 5 to 100, p_1 and b_1 independent of each other are an integer of 0 to 50, provided that $(e_1+p_1+b_1) \geq 10$ and $\frac{e_1}{(p_1+b_1)} \geq 2$ when $(p_1+b_1) \geq 1$;

R_0 is H or methyl;

R_1 , R_{1n} , R_{2n} , R_{3n} , and R_{4n} independent of one another are H or a C_1 - C_4 alkyl having 0 to 2 hydroxyl group;

R_{n5} is H or a C_1 - C_{10} alkyl;

R_2 is a C_4 - C_{14} hydrocarbon divalent radical;

R_3 is a C_2 - C_6 alkylene divalent radical;

R_4 and R_5 independent of each other are a C_1 - C_6 alkylene divalent radical or a C_1 - C_6 alkylene-oxy- C_1 - C_6 alkylene divalent radical;

R_6 and R_7 independent of each other are a C_1 - C_6 alkylene divalent radical or a C_1 - C_6 alkoxy- C_1 - C_6 alkylene divalent radical;

R_8 and R_9 independent of each other are a substituted or unsubstituted C_1 - C_{12} alkylene divalent radical;

X_0 , X_1 , X_{01} , X_{02} , and X_{03} independent of one another are O or NR_1 ;

X_1 is O, NR_1 , $NHCOO$, $OCONH$, $CONR_1$, or NR_1CO ;

X_{04} is $-COO-$ or $-CONR_{n5}-$;

X_{05} and X_{07} independent of each other are a direct bond, $-COO-$ or $-CONR_{n5}-$;

X_{06} is a direct bond, a C_1 - C_6 alkylene divalent radical, a C_1 - C_6 alkyleneoxy divalent radical, $-COO-$, or $-CONR_{n5}-$;

X_{08} is a direct bond or $-\text{COO}-$;

X_{09} is O or NR_{n5} ;

X_{10} is a direct bond, a C_1 - C_6 alkylene divalent radical, $-\text{COO}-$, or $-\text{CONR}_{n5}-$;

E_1' is a monovalent radical of $\text{H}_2\text{C}=\overset{\text{R}_0}{\underset{\cdot}{\text{C}}}-\overset{\text{O}}{\parallel}{\text{C}}-\text{X}_{01}-\text{L}_4-$;

E_2 is a monovalent radical of $\text{H}_2\text{C}=\overset{\text{R}_0}{\underset{\cdot}{\text{C}}}-\overset{\text{O}}{\parallel}{\text{C}}-\text{Z}_7-$;

E_3 is a monovalent radical of $\text{H}_2\text{C}=\overset{\text{R}_0}{\underset{\cdot}{\text{C}}}-\overset{\text{O}}{\parallel}{\text{C}}-\text{X}_{03}-$;

E_4 is a monovalent radical of $\text{H}_2\text{C}=\overset{\text{R}_0}{\underset{\cdot}{\text{C}}}-\overset{\text{O}}{\parallel}{\text{C}}-\text{X}_{11}-\text{G}_2-$ or $\text{H}_2\text{C}=\overset{\text{R}_0}{\underset{\cdot}{\text{C}}}-\overset{\text{O}}{\parallel}{\text{C}}-\text{G}_3-$;

L_1 is a C_2 - C_8 alkylene divalent radical or a divalent radical of $-\text{L}_1'-\text{X}_1-\text{L}_1''-$,
 $-(\text{C}_2\text{H}_4\text{O})_{q1}-\text{L}_1''-$, $-(\text{C}_2\text{H}_4\text{O})_{q1}-\text{CONH}-\text{L}_1''-$, $-\text{L}_1'-\text{NHCOO}-(\text{C}_2\text{H}_4\text{O})_{q1}-\text{L}_1''-$,
 $-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{X}_1'-(\text{C}_2\text{H}_4\text{O})_{q2}-\text{L}_1''-$, $-\text{L}_1'-\text{X}_1'-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O}-\text{L}_1''-$, or
 $-(\text{C}_2\text{H}_4\text{O})_{q1}-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O}-\text{L}_1''-$;

L_1' is a C_2 - C_8 alkylene divalent radical which has zero or one hydroxyl group;

L_1'' is C_3 - C_8 alkylene divalent radical which has zero or one hydroxyl group;

L_3 is a divalent radical of

$-\text{L}_3'-\text{O}-(\text{C}_2\text{H}_4\text{O})_{q2}-\text{CONH}-\text{R}_2-(\text{NHCO}-\text{PE}-\text{CONH}-\text{R}_2)_{g1}-\text{NHCO}-(\text{OC}_2\text{H}_4)_{q2}-\text{O}-\text{L}_3'-$ in which

PE is a divalent radical of

$-(\text{CH}_2\text{CH}_2\text{O})_{q3}-\text{Z}_0-\text{CF}_2-(\text{OCF}_2)_x-(\text{OCF}_2\text{CF}_2)_y-\text{OCF}_2-\text{Z}_0-(\text{OCH}_2\text{CH}_2)_{q3}-$ or

$-\text{Z}_0-(\text{CH}_2\text{CH}_2\text{O})_{q4}-\left(\text{CH}_2-\overset{\text{CH}_3}{\underset{\cdot}{\text{CH}}}-\text{O}\right)_{q5}-\left(\text{CH}_2-\overset{\text{C}_2\text{H}_5}{\underset{\cdot}{\text{CH}}}-\text{O}\right)_{q6}-\text{Z}_0-$;

L_3' is C_3 - C_8 alkylene divalent radical;

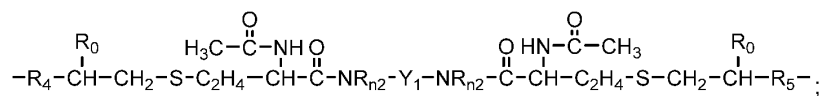
L_4 is a divalent radical of $-\text{C}_2\text{H}_4-\text{NHCO}-\text{O}-(\text{C}_2\text{H}_4\text{O})_{q2}-\text{L}_3'-$,

$-(\text{C}_2\text{H}_4\text{O})_{q1}-\text{CONH}-\text{R}_2-\text{NHCO}-\text{O}-(\text{C}_2\text{H}_4\text{O})_{q2}-\text{L}_3'-$,

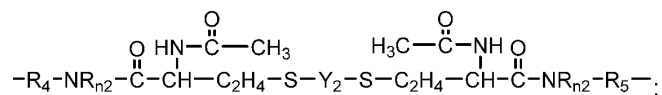
$-\text{R}_3-\text{O}-\text{CONH}-\text{R}_2-\text{NHCO}-\text{O}-(\text{C}_2\text{H}_4\text{O})_{q2}-\text{L}_3'-$, $-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O}-(\text{C}_2\text{H}_4\text{O})_{q2}-\text{L}_3'-$, or

$-(\text{C}_2\text{H}_4\text{O})_{q2}-\text{L}_3'-$;

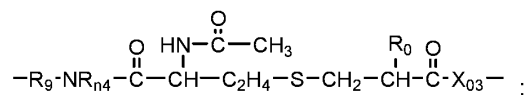
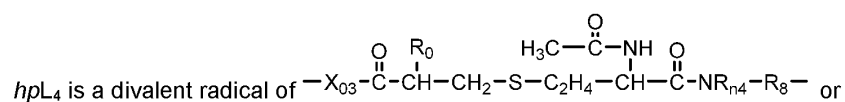
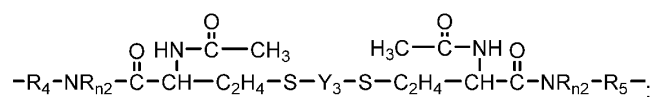
hpL_1 is a divalent radical of



hpL₂ is a divalent radical of



hpL₃ is a divalent radical of



pOAlk is a divalent radical of $-(\text{EO})_{e1}(\text{PO})_{p1}(\text{BO})_{b1}-$ in which EO is an oxyethylene

unit ($-\text{CH}_2\text{CH}_2\text{O}-$), PO is an oxypropylene unit ($-\text{CH}_2-\overset{\overset{\text{CH}-\text{O}-}{|}}{\text{CH}_3}-$), and BO is an

oxybutylene unit ($-\text{CH}_2-\overset{\overset{\text{CH}-\text{O}-}{|}}{\text{C}_2\text{H}_5}-$);

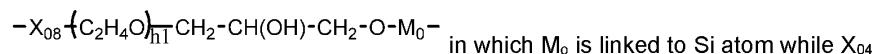
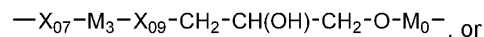
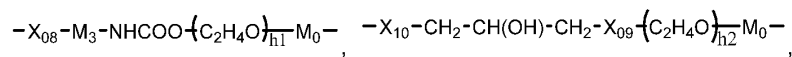
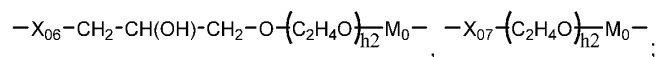
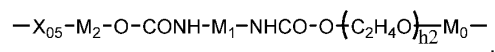
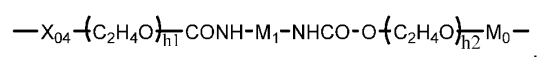
M₀ is C₃-C₈ alkylene divalent radical;

M₁ is a C₄-C₁₄ hydrocarbon divalent radical;

M₂ and *M₃* independent of each other are a C₁-C₆ alkylene divalent radical;

J₀ is a C₁-C₁₂ hydrocarbon radical having 0 to 2 hydroxyl or carboxyl groups;

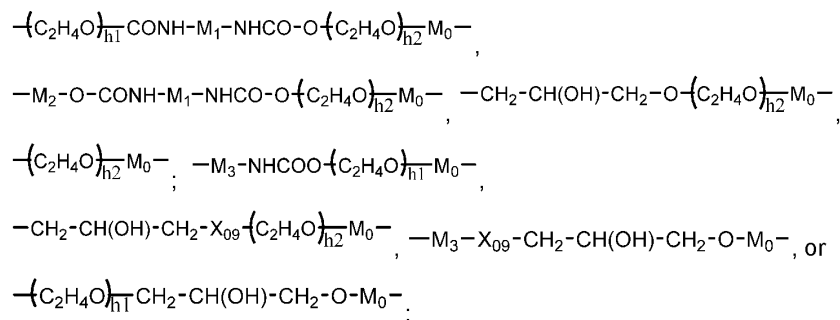
G1 is a direct bond, a C₁-C₄ alkylene divalent radical, or a divalent radical of



in which *M₀* is linked to Si atom while *X₀₄* to *X₁₀* are linked to the group of $-\text{CH}_2-$ in formula (7) and at least one of *J₀* and *G1* in formula (7) comprises at least one moieties selected from the group consisting of

hydroxyl groups, urethane linkage of $-\text{OCONH}-$, amino groups of $-\text{NHR}^\circ$, amino linkages of $-\text{NH}-$, amide linkages of $-\text{CONH}-$, carboxyl groups, and combinations thereof;

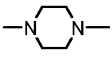
G_2 is a C_1 - C_4 alkylene divalent radical or a bivalent radical of



G_3 is a divalent radical of $-\text{G}_4-\left(\text{C}(\text{O})\right)_{h3}-\text{CH}-\text{CH}_2-\text{S}-\text{J}_0-\text{S}-\text{CH}_2-\text{CH}-\left(\text{C}(\text{O})\right)_{h4}-\text{G}_2-$ in

which h_3 and h_4 independent of each other are 1 or 0;

G_4 is a divalent radical of any one of (a) $-\text{NR}_3'-$ in which R_3' is hydrogen or C_1 - C_3

alkyl, (b) , (c) $-\text{NR}_0-\text{G}_5-\text{NR}_0-$ in which G_5 is a C_1 - C_6 alkylene divalent radical, 2-hydroxypropylene divalent radical, 2-(phosphonyloxy)propylene divalent radical, 1,2-dihydroxyethylene divalent radical, 2,3-dihydroxybutylene divalent radical, and (d) $-\text{O}-\text{G}_6-\text{O}-$ in which G_6 is a C_1 - C_6 alkylene divalent radical, a

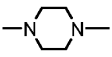
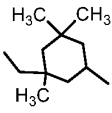
divalent radical of $-(\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O})_{h4'}-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-$ in which h_4' is 1 or 2, a

divalent radical of $-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{O}-\text{CH}_2-\text{CH}_2-\text{O}-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-$, a divalent radical

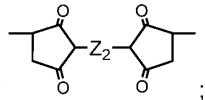
of $-(\text{CH}_2-\text{CH}_2-\text{O})_{h5}-\text{CH}_2-\text{CH}_2-$ in which h_5 is an integer of 1 to 5, a divalent

radical of $-(\text{CH}_2)_{h6}-\text{O}-\text{P}(\text{OH})(\text{O})-\text{O}-(\text{CH}_2)_{m5}-$ in which h_6 is 2 or 3, or a substituted C_3 - C_8 alkylene divalent radical having a hydroxyl group or phosphonyloxy group;

Y_1 is a C_1 - C_6 alkylene divalent radical, 2-hydroxypropylene divalent radical, 2-(phosphonyloxy)propylene divalent radical, 1,2-dihydroxyethylene divalent radical, a

divalent radical of , or a divalent radical of ;

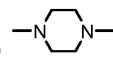
Y_2 is a divalent radical of $-\text{CH}_2-\overset{\text{R}_0}{\underset{\text{O}}{\overset{\text{O}}{\text{C}}}}-\text{CH}-\text{Z}_3-\overset{\text{R}_0}{\underset{\text{O}}{\overset{\text{O}}{\text{C}}}}-\text{CH}-\text{CH}_2-$;

Y_3 is a divalent radical of $-\text{CH}_2-\text{CH}_2-\overset{\text{O}}{\underset{\text{O}}{\overset{\text{O}}{\text{S}}}}-\left(\overset{\text{O}}{\underset{\text{O}}{\overset{\text{O}}{\text{S}}}}-\text{Z}_1-\overset{\text{O}}{\underset{\text{O}}{\overset{\text{O}}{\text{S}}}}\right)_{m1}-\text{CH}_2-\text{CH}_2-$ or ;

Z_0 is a direct bond or a C_1 - C_{12} alkylene divalent radical;

Z_1 is a C_1 - C_6 alkylene divalent radical, a hydroxyl- or methoxy-substituted C_1 - C_6 alkylene divalent radical, or a substituted or unsubstituted phenylene divalent radical,

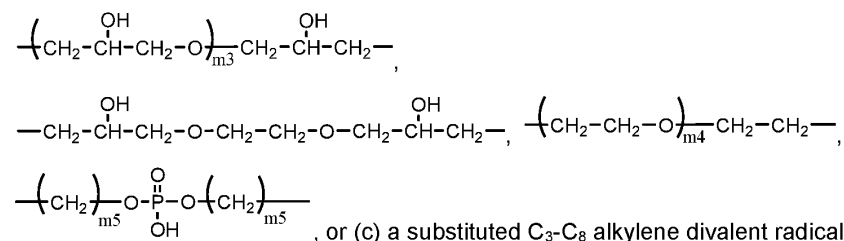
Z_2 is a C_1 - C_6 alkylene divalent radical, a hydroxyl- or methoxy-substituted C_1 - C_6 alkylene divalent radical, a dihydroxyl- or dimethoxy-substituted C_2 - C_6 alkylene divalent radical, a divalent radical of $-\text{C}_2\text{H}_4-(\text{O}-\text{C}_2\text{H}_4)_{m2}-$, a divalent radical of $-\text{Z}_4-\text{S}-\text{S}-\text{Z}_4-$, a hydroxyl- or methoxy-substituted C_1 - C_6 alkylene divalent radical, or a substituted or unsubstituted phenylene divalent radical,

Z_3 is a divalent radical of any one of (a) $-\text{NR}_{n3}-$, (b) , (c) $-\text{NR}_0-\text{Z}_5-\text{NR}_0-$, and (d) $-\text{O}-\text{Z}_6-\text{O}-$,

Z_4 is a C_1 - C_6 alkylene divalent radical,

Z_5 is a C_1 - C_6 alkylene divalent radical, 2-hydroxypropylene divalent radical, 2-(phosphonyloxy)propylene divalent radical, 1,2-dihydroxyethylene divalent radical, 2,3-dihydroxybutylene divalent radical,

Z_6 is (a) a C_1 - C_6 alkylene divalent radical, (b) a divalent radical of



having a hydroxyl group or phosphonyloxy group,

Z_7 is a divalent radical of $-\text{Z}_3-\overset{\text{O}}{\underset{\text{O}}{\overset{\text{O}}{\text{C}}}}-\overset{\text{R}_0}{\underset{\text{O}}{\overset{\text{O}}{\text{C}}}}-\text{CH}-\text{CH}_2-\text{S}-\text{C}_2\text{H}_4-\text{CH}-\overset{\text{O}}{\underset{\text{O}}{\overset{\text{O}}{\text{C}}}}-\text{NR}_{n2}-\text{R}_4-$; and

Z_8 is a divalent radical of $-\text{Z}_9-\overset{\text{O}}{\underset{\text{O}}{\overset{\text{O}}{\text{C}}}}-\overset{\text{R}_0}{\underset{\text{O}}{\overset{\text{O}}{\text{C}}}}-\text{CH}-\text{CH}_2-\text{S}-\text{C}_2\text{H}_4-\text{CH}-\overset{\text{O}}{\underset{\text{O}}{\overset{\text{O}}{\text{C}}}}-\text{NR}_{n2}-\text{R}_4-$.

37. The method of any one of embodiments 1 to 32, wherein the polymerizable materials comprise at least one silicone-containing vinylic crosslinker which is α,ω -bis[3-

(meth)acrylamidopropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxyethoxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxy-isopropoxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxybutyloxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamidoethoxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamidopropoxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamidoisopropoxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamidobutyloxy-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxyethylamino-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxypropylamino-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acryloxybutylamino-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acrylamidoethylamino-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamidopropylamino-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[3-(meth)acrylamide-butylamino-2-hydroxypropyloxypropyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acryloxy-2-hydroxypropyloxy-ethoxypropyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acryloxy-2-hydroxypropyl-N-ethylaminopropyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acryloxy-2-hydroxypropyl-aminopropyl]-polydimethylsiloxane, α,ω -bis[(meth)acryloxy-2-hydroxypropyloxy-(polyethylenoxy)propyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acryloxyethylamino-carbonyloxy-ethoxypropyl]-terminated polydimethylsiloxane, α,ω -bis[(meth)acryloxyethylamino-carbonyloxy-(polyethylenoxy)propyl]-terminated polydimethylsiloxane, or a combination thereof.

38. The method of any one of embodiments 1 to 37, wherein said at least one hydrophilic vinylic monomer comprises: (1) an alkyl (meth)acrylamide selected from the group consisting of (meth)acrylamide, N,N-dimethyl (meth)acrylamide, N-ethyl (meth)acrylamide, N,N-diethyl (meth)acrylamide, N-propyl (meth)acrylamide, N-isopropyl (meth)acrylamide, N-3-methoxypropyl (meth)acrylamide, and combinations thereof; (2) a hydroxyl-containing acrylic monomer selected from the group consisting of N-2-hydroxyethyl (meth)acrylamide, N,N-bis(hydroxyethyl) (meth)acrylamide, N-3-

hydroxypropyl (meth)acrylamide, N-2-hydroxypropyl (meth)acrylamide, N-2,3-dihydroxypropyl (meth)acrylamide, *N*-tris(hydroxymethyl)methyl (meth)acrylamide, 2-hydroxyethyl (meth)acrylate, 3-hydroxypropyl (meth)acrylate, 2-hydroxypropyl (meth)acrylate, glycerol methacrylate (GMA), di(ethylene glycol) (meth)acrylate, tri(ethylene glycol) (meth)acrylate, tetra(ethylene glycol) (meth)acrylate, poly(ethylene glycol) (meth)acrylate having a number average molecular weight of up to 1500, poly(ethylene glycol)ethyl (meth)acrylamide having a number average molecular weight of up to 1500, and combinations thereof; (3) a carboxyl-containing acrylic monomer selected from the group consisting of 2-(meth)acrylamidoglycolic acid, (meth)acrylic acid, ethylacrylic acid, and combinations thereof; (4) an amino-containing acrylic monomer selected from the group consisting of N-2-aminoethyl (meth)acrylamide, N-2-methylaminoethyl (meth)acrylamide, N-2-ethylaminoethyl (meth)acrylamide, N-2-dimethylaminoethyl (meth)acrylamide, N-3-aminopropyl (meth)acrylamide, N-3-methylaminopropyl (meth)acrylamide, N-3-dimethylaminopropyl (meth)acrylamide, 2-aminoethyl (meth)acrylate, 2-methylaminoethyl (meth)acrylate, 2-ethylaminoethyl (meth)acrylate, 3-aminopropyl (meth)acrylate, 3-methylaminopropyl (meth)acrylate, 3-ethylaminopropyl (meth)acrylate, 3-amino-2-hydroxypropyl (meth)acrylate, trimethylammonium 2-hydroxy propyl (meth)acrylate hydrochloride, dimethylaminoethyl (meth)acrylate, and combinations thereof; (5) an N-vinyl amide monomer selected from the group consisting of N-vinylpyrrolidone (*aka*, N-vinyl-2-pyrrolidone), N-vinyl-3-methyl-2-pyrrolidone, N-vinyl-4-methyl-2-pyrrolidone, N-vinyl-5-methyl-2-pyrrolidone, N-vinyl-6-methyl-2-pyrrolidone, N-vinyl-3-ethyl-2-pyrrolidone, N-vinyl-4,5-dimethyl-2-pyrrolidone, N-vinyl-5,5-dimethyl-2-pyrrolidone, N-vinyl-3,3,5-trimethyl-2-pyrrolidone, N-vinyl piperidone (*aka*, N-vinyl-2-piperidone), N-vinyl-3-methyl-2-piperidone, N-vinyl-4-methyl-2-piperidone, N-vinyl-5-methyl-2-piperidone, N-vinyl-6-methyl-2-piperidone, N-vinyl-6-ethyl-2-piperidone, N-vinyl-3,5-dimethyl-2-piperidone, N-vinyl-4,4-dimethyl-2-piperidone, N-vinyl caprolactam (*aka*, N-vinyl-2-caprolactam), N-vinyl-3-methyl-2-caprolactam, N-vinyl-4-methyl-2-caprolactam, N-vinyl-7-methyl-2-caprolactam, N-vinyl-7-ethyl-2-caprolactam, N-vinyl-3,5-dimethyl-2-caprolactam, N-vinyl-4,6-dimethyl-2-caprolactam, N-vinyl-3,5,7-trimethyl-2-caprolactam, N-vinyl-N-methyl acetamide, N-vinyl formamide, N-vinyl acetamide, N-vinyl isopropylamide, N-vinyl-N-ethyl acetamide, N-vinyl-N-ethyl formamide, and mixtures thereof; (6) a methylene-containing pyrrolidone monomer selected from the group consisting of 1-methyl-3-methylene-2-pyrrolidone, 1-ethyl-3-methylene-2-pyrrolidone, 1-methyl-5-methylene-2-pyrrolidone, 1-ethyl-5-methylene-2-pyrrolidone, 5-methyl-3-methylene-2-pyrrolidone, 5-ethyl-3-methylene-2-pyrrolidone, 1-n-propyl-3-methylene-2-pyrrolidone, 1-n-propyl-

5-methylene-2-pyrrolidone, 1-isopropyl-3-methylene-2-pyrrolidone, 1-isopropyl-5-methylene-2-pyrrolidone, 1-n-butyl-3-methylene-2-pyrrolidone, 1-tert-butyl-3-methylene-2-pyrrolidone, and combinations thereof; (7) acrylic monomers having a C₁-C₄ alkoxyethoxy group selected from the group consisting of ethylene glycol methyl ether (meth)acrylate, di(ethylene glycol) methyl ether (meth)acrylate, tri(ethylene glycol) methyl ether (meth)acrylate, tetra(ethylene glycol) methyl ether (meth)acrylate, C₁-C₄-alkoxy poly(ethylene glycol) (meth)acrylate having a weight average molecular weight of up to 1500, methoxy-poly(ethylene glycol)ethyl (meth)acrylamide having a number average molecular weight of up to 1500, and combinations thereof; (8) a vinyl ether monomer selected from the group consisting of ethylene glycol monovinyl ether, di(ethylene glycol) monovinyl ether, tri(ethylene glycol) monovinyl ether, tetra(ethylene glycol) monovinyl ether, poly(ethylene glycol) monovinyl ether, ethylene glycol methyl vinyl ether, di(ethylene glycol) methyl vinyl ether, tri(ethylene glycol) methyl vinyl ether, tetra(ethylene glycol) methyl vinyl ether, poly(ethylene glycol) methyl vinyl ether, and combinations thereof; (9) an allyl ether monomer selected from the group consisting of ethylene glycol monoallyl ether, di(ethylene glycol) monoallyl ether, tri(ethylene glycol) monoallyl ether, tetra(ethylene glycol) monoallyl ether, poly(ethylene glycol) monoallyl ether, ethylene glycol methyl allyl ether, di(ethylene glycol) methyl allyl ether, tri(ethylene glycol) methyl allyl ether, tetra(ethylene glycol) methyl allyl ether, poly(ethylene glycol) methyl allyl ether, and combinations thereof; (10) a phosphorylcholine-containing vinylic monomer selected from the group consisting of (meth)acryloyloxyethyl phosphorylcholine, (meth)acryloyloxypropyl phosphorylcholine, 4-((meth)acryloyloxy)butyl-2'-(trimethylammonio)ethylphosphate, 2-[(meth)acryloylamino]ethyl-2'-(trimethylammonio)-ethylphosphate, 3-[(meth)acryloylamino]propyl-2'-(trimethylammonio)ethylphosphate, 4-[(meth)acryloylamino]butyl-2'-(trimethylammonio)ethylphosphate, 5-((meth)acryloyloxy)pentyl-2'-(trimethylammonio)ethyl phosphate, 6-((meth)acryloyloxy)hexyl-2'-(trimethylammonio)-ethylphosphate, 2-((meth)acryloyloxy)ethyl-2'-(triethylammonio)ethylphosphate, 2-((meth)acryloyloxy)ethyl-2'-(tripropylammonio)ethylphosphate, 2-((meth)acryloyloxy)ethyl-2'-(tributylammonio)ethyl phosphate, 2-((meth)acryloyloxy)propyl-2'-(trimethylammonio)-ethylphosphate, 2-((meth)acryloyloxy)butyl-2'-(trimethylammonio)ethylphosphate, 2-((meth)acryloyloxy)pentyl-2'-(trimethylammonio)ethylphosphate, 2-((meth)acryloyloxy)hexyl-2'-(trimethylammonio)ethyl phosphate, 2-(vinylloxy)ethyl-2'-(trimethylammonio)ethylphosphate, 2-(allyloxy)ethyl-2'-(trimethylammonio)ethylphosphate, 2-(vinylloxycarbonyl)ethyl-2'-(trimethylammonio)ethylphosphate, 2-(vinylloxycarbonyl)ethyl-2'-(trimethylammonio)ethylphosphate, and combinations thereof.

(trimethylammonio)ethyl phosphate, 2-(allyloxycarbonyl)ethyl-2'-(trimethylammonio)-ethylphosphate, 2-(vinylcarbonylamino)ethyl-2'-(trimethylammonio)ethylphosphate, 2-(allyloxycarbonylamino)ethyl-2'-(trimethylammonio)ethyl phosphate, 2-(butenoyloxy)ethyl-2'-(trimethylammonio)ethylphosphate, and combinations thereof; (11) allyl alcohol; (12) N-2-hydroxyethyl vinyl carbamate; (13) N-carboxyvinyl- β -alanine (VINAL); (14) N-carboxyvinyl- α -alanine; (15) or combinations thereof.

39. The method of embodiment 38, wherein said at least one hydrophilic vinylic monomer comprises N-vinylpyrrolidone, N-vinyl-N-methyl acetamide, or combinations thereof.
40. The method of embodiment 38 or 39, wherein said at least one hydrophilic vinylic monomer comprises N,N-dimethyl (meth)acrylamide.
41. The method of any one of embodiments 38 to 40, wherein said at least one hydrophilic vinylic monomer comprises N-2-hydroxyethyl (meth)acrylamide, N,N-bis(hydroxyethyl) (meth)acrylamide, N-3-hydroxypropyl (meth)acrylamide, N-2-hydroxypropyl (meth)acrylamide, N-2,3-dihydroxypropyl (meth)acrylamide, *N*-tris(hydroxymethyl)methyl (meth)acrylamide, 2-hydroxyethyl (meth)acrylate, 3-hydroxypropyl (meth)acrylate, 2-hydroxypropyl (meth)acrylate, glycerol methacrylate (GMA), di(ethylene glycol) (meth)acrylate, tri(ethylene glycol) (meth)acrylate, tetra(ethylene glycol) (meth)acrylate, poly(ethylene glycol) (meth)acrylate having a number average molecular weight of up to 1500, poly(ethylene glycol)ethyl (meth)acrylamide having a number average molecular weight of up to 1500, or combinations thereof.
42. The method of any one of embodiments 1 to 41, wherein said at least one non-silicone vinylic cross-linking agents comprises ethyleneglycol di-(meth)acrylate, diethyleneglycol di-(meth)acrylate, triethyleneglycol di-(meth)acrylate, tetraethyleneglycol di-(meth)acrylate, glycerol di-(meth)acrylate, 1,3-propanediol di-(meth)acrylate, 1,3-butanediol di-(meth)acrylate, 1,4-butanediol di-(meth)acrylate, glycerol 1,3-diglycerolate di-(meth)acrylate, ethylenebis[oxy(2-hydroxypropane-1,3-diyl)] di-(meth)acrylate, bis[2-(meth)acryloxyethyl] phosphate, trimethylolpropane di-(meth)acrylate, and 3,4-bis[(meth)acryloyl]tetrahydrofuran, diacrylamide, dimethacrylamide, N,N-di(meth)acryloyl-N-methylamine, N,N-di(meth)acryloyl-N-ethylamine, N,N'-methylene bis(meth)acrylamide, N,N'-ethylene bis(meth)acrylamide, N,N'-dihydroxyethylene bis(meth)acrylamide, N,N'-propylene bis(meth)acrylamide, N,N'-2-hydroxypropylene bis(meth)acrylamide, N,N'-2,3-dihydroxybutylene bis(meth)acrylamide, 1,3-bis(meth)acrylamidepropane-2-yl dihydrogen phosphate, piperazine diacrylamide, tetraethyleneglycol divinyl ether, triethyleneglycol divinyl ether, diethyleneglycol divinyl ether, ethyleneglycol divinyl ether, triallyl isocyanurate, triallyl cyanurate, trimethylopropane trimethacrylate, pentaerythritol tetramethacrylate,

bisphenol A dimethacrylate, allylmethacrylate, allylacrylate, N-allyl-methacrylamide, N-allyl-acrylamide, or combinations thereof.

43. The method of any one of embodiments 1 to 42, wherein said at least one blending vinylic monomer comprises methyl methacrylate.
44. The method of any one of embodiments 1 to 43, wherein the polymerizable materials comprise 2-[2'-hydroxy-5'-(2-methacryloxyethyl)phenyl]-2H-benzotriazole and a UV/HEVL absorbing vinylic monomer selected from the group consisting of 2-{2'-Hydroxy-3'-*tert*-butyl-5'-[3'-methacryloyloxypropoxy]phenyl}-2H-benzotriazole, 2-{2'-Hydroxy-3'-*tert*-butyl-5'-[3'-methacryloyloxypropoxy]phenyl}-5-methoxy-2H-benzotriazole, 2-{2'-Hydroxy-3'-*tert*-butyl-5'-[3'-methacryloyloxypropoxy]phenyl}-5-chloro-2H-benzotriazole, 2-[2'-Hydroxy-3'-*tert*-butyl-5'-(3'-acryloyloxypropoxy)phenyl]-5-trifluoromethyl-2H-benzotriazole, and combinations thereof.
45. The method of any one of embodiments 1 to 44, wherein the liquid extraction medium is water or an aqueous solution.
46. The method of any one of embodiments 1 to 44, wherein the liquid extraction medium is a mixture of water with one or more organic solvents selected from the group consisting of 1,2-propylene glycol, a polyethyleneglycol having a number average molecular weight of about 400 Daltons or less, a C₁-C₃ alkyl alcohol, and combination thereof.
47. The method of any one of embodiments 1 to 44, wherein the liquid extraction medium is one or more organic solvents selected from the group consisting of 1,2-propylene glycol, a polyethyleneglycol having a number average molecular weight of about 400 Daltons or less, a C₁-C₃ alkylalcohol, and combination thereof.
48. A hydrated silicone hydrogel contact lens obtained according to the method of any one of claims 1 to 47.
49. The silicone hydrogel contact lens of embodiment 87, wherein the silicone hydrogel contact lens has an oxygen permeability of at least 50 barrers, an elastic modulus of from about 0.2 MPa to about 1.5 MPa, and an equilibrium water content of from about 15% to about 75% by weight.

The previous disclosure will enable one having ordinary skill in the art to practice the invention. Various modifications, variations, and combinations can be made to the various embodiment described herein. In order to better enable the reader to understand specific embodiments and the advantages thereof, reference to the following examples is suggested. It is intended that the specification and examples be considered as exemplary.

Example 1

Oxygen Permeability Measurements

Unless specified, the apparent oxygen permeability (Dk_{app}), the apparent oxygen transmissibility (Dk/t), the intrinsic (or edge-corrected) oxygen permeability (Dk_i or Dk_e) of a lens and a lens material are determined according to procedures described in Example 1 of U.S. Pat. Appl. Pub. No. 2012/0026457 A1.

Equilibrium Water Content

The equilibrium water content (EWC) of contact lenses is determined as follows.

Amount of water (expressed as percent by weight) present in a hydrated hydrogel contact lens, which is fully equilibrated in saline solution, is determined at room temperature. Quickly stack the lenses, and transfer the lens stack to the aluminum pan on the analytical balance after blotting lens in a cloth. The number of lenses for each sample pan is typically five (5) and three sample pans are used. Record the pan plus hydrated weight of the lenses. Cover the pan with aluminum foil. Place pans in a laboratory oven at 100 ± 2 °C to dry for 16-18 hours. Remove pan plus lenses from the oven and cool in a desiccator for at least 30 minutes. Remove a single pan from the desiccator, and discard the aluminum foil. Weigh the pan plus dried lens sample on an analytical balance. Repeat for all pans. The wet and dry weight of the lens samples can be calculated by subtracting the weight of the empty weigh pan and then being divided by the number of lenses for each sample pan. The measurements of three sample pans from one single batch of contact lenses are averaged to obtain the averaged equilibrium water content for that batch of contact lenses.

Determination of Diameter of Hydrated Contact Lens

Figure 1 shows schematically the general lens dimensions which are typically determined by a contact lens manufacturer. The general dimensions include center thickness (CT) (110), anterior sagittal height (ASag) (120), posterior sagittal height (PSag) (140), base curve equivalent (BCE) (150), edge thickness (ET) (160), and diameter (180). Measurements of the general lens dimensions can be carried out on fully hydrated contact lenses in a wetcell by using a low coherence interferometer similar to that described by Heidemana and Greivenkamp in their paper (Optical Engineering 55(3), 034106 (March 2016)).

For measurements, a contact lens is seated on the flat bottom surface of the wetcell which is filled with a phosphate buffered saline and a low coherence interferometer is placed at the geometric center of the lens using a motion controller. The interferometer measures thicknesses of material based off of reflections between different material surfaces. The center of the lens is determined by the measurement by the camera.

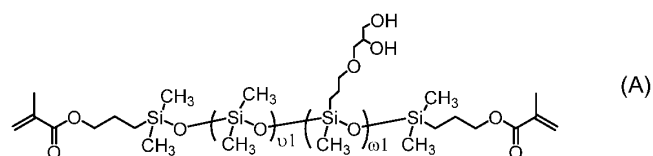
The Diameter is defined as the outermost edge of the lens viewed from above the lens. The edge points are fit to an ellipse and the diameter is calculated as the average of the major and minor ellipse diameters. Typically, contact lenses have highly circular diameters and either a circular or elliptical fitting will result in similar values. However, if a lens is slightly out of round, an ellipse more accurately describes the shape of the contact lens diameter than a circle. The lens diameters of 3 to 10 contact lenses from one single batch of contact lenses are measured and averaged to obtain the averaged lens diameter for that batch of contact lenses.

Elastic Modulus

The elastic modulus of a contact lens is determined using a MTS insight instrument. The contact lens is first cut into a 3.12 mm wide strip using Precision Concept two stage cutter. Five thickness values are measured within 6.5mm gauge length. The strip is mounted on the instrument grips and submerged in PBS (phosphate buffered saline) with the temperature controlled at 21 ± 2 °C. Typically 5N Load cell is used for the test. Constant force and speed is applied to the sample until the sample breaks. Force and displacement data are collected by the TestWorks software. The elastic modulus value is calculated by the TestWorks software which is the slope or tangent of the stress vs. strain curve near zero elongation, in the elastic deformation region.

Chemicals

The following abbreviations are used in the following examples: NVP represents N-vinylpyrrolidone; DMA represents N,N-dimethylacrylamide; MMA represents methyl methacrylate; TEGDMA represent triethyleneglycol dimethacrylate; VAZO 64 represents 2,2'-dimethyl-2,2'-azodipropiononitrile; Nobloc is 2-[3-(2H-Benzotriazol-2-yl)-4-hydroxyphenyl]ethyl methacrylate from Aldrich; TAA represents tert-amyl alcohol; ProOH represents 1-propanol; IPA represents isopropanol; DC 1173 represents Darocur 1173® photoinitiator; PBS represents a phosphate-buffered saline which has a pH of 7.2 ± 0.2 at 25°C and contains about 0.044 wt.% $\text{NaH}_2\text{PO}_4 \cdot \text{H}_2\text{O}$, about 0.388 wt.% $\text{Na}_2\text{HPO}_4 \cdot 2\text{H}_2\text{O}$, and about 0.79 wt.% NaCl and; wt.% represents weight percent; D9 represents monobutyl-terminated monomethacryloxypropyl-terminated polydimethylsiloxane ($M_w \sim 984$ g/mol from Shin-Etsu); CEPDMS represents a di-methacrylate-terminated chain-extended polydimethylsiloxane ($M_n \sim 9000$ g/mol), which has three polydimethylsiloxane (PDMS) segments linked via diurethane linkages between two PDMS segments and two urethane linkages each located between one terminal methacrylate group and one PDMS segment, is prepared according to method similar to what described in Example 2 of US8,529,057; "G4" macromer represents a di-methacryloxypropyl-terminated polysiloxane ($M_n \sim 13.5\text{K}$ g/mol, OH content ~ 1.8 meq/g) of formula (A).



Example 2

A lens formulation is purged with nitrogen at room temperature for 30 to 35 minutes. The N₂-purged lens formulation is introduced into polypropylene molds and thermally cured in an oven under the following curing conditions: ramping from room temperature to a first temperature and then holding at the first temperature for a first curing time period; ramping from the first temperature to a second temperature and holding at the second temperature for a second curing time period; optionally ramping from the second temperature to a third temperature and holding at the third temperature for a third curing time period; and optionally ramping from the third temperature to a fourth temperature and holding at the fourth temperature for a fourth curing time period.

Lens molds are opened by using a demolding machine with a push pin. Lenses are pushed onto base curve molds with a push pin and then molds are separated into base curve mold halves and front curve mold halves. The base curve mold halves with a lens thereon are placed in an ultrasonic device (e.g., Dukane's single horn ultrasonic device). With a certain energy force, a dry state lens is released from mold. The dry state lens is loaded in a designed extraction tray. Alternatively, lenses can be removed from the base curve mold halves by floating off (i.e., soaking in an organic solvent, e.g., IPA, without ultrasonic). The lenses removed from the molds are subjected to an extraction process using water or an organic solvent or a mixture of solvents for at least 30 minutes. For example, extracted in 50% IPA for 30 min, or in 100% IPA for 15 min then back to 50% IPA for 30 min, DI water for 30min and finally in PBS saline overnight. Inspected lens is packaged in lens packages containing a phosphate buffered saline (pH ~ 7.2) and autoclaved at 121°C for about 30-45 minutes.

Example 3

A lens formulation is purged with nitrogen at room temperature for 30 to 35 minutes. The N₂-purged lens formulation is introduced into polypropylene molds and cured by UV/visible light (Hamamatsu lamp) for a curing time period. The post cast molding procedures described in Example 2 are used in this process for producing SiHy contact lenses.

Examples 4-8

Lens formulations (fluid polymerizable compositions) are prepared to have compositions (in weight part units). All the prepared formulations consists two parts: 102.06 weight part units of the polymerizable materials (0.65 weight part units of **TEGDMA**, 10 weight part units of **G4**, 33 weight part units of **D9**, 46 weight part units of **NVP**, 10 weight part units of **MMA**, 1.5 weight part units of **Norbloc**, 0.40 weight part units of **UV28**, 0.01 weight part units of **RB247**, 0.5 weight part units of **V64**) and various weight part units of an alcohol as shown in Tables 1 to 5. The formulations are prepared by adding listed components in their targeted amounts into a clean bottle, with a stir bar to mix at 600 rpm for 30 minutes at room temperature. After all the solid is dissolved, a filtration of the formulation is carried out by using 2.7µm glass-microfiber-filter.

Table 1

	Composition (Weigh Part Units)				
	4-1	4-2	4-3	4-4	4-5
Polymerizable materials	102.06	102.06	102.06	102.06	102.06
Ethanol	6	8	10	12	14

Table 2

	Composition (Weigh Part Units)				
	5-1	5-2	5-3	5-4	5-5
Polymerizable materials	102.06	102.06	102.06	102.06	102.06
1-Propanol	6	8	10	12	14

Table 3

	Composition (Weigh Part Units)				
	6-1	6-2	6-3	6-4	6-5
Polymerizable materials	102.06	102.06	102.06	102.06	102.06
1-Butanol	6	8	10	12	14

Table 4

	Composition (Weigh Part Units)				
	7-1	7-2	7-3	7-4	7-5
Polymerizable materials	102.06	102.06	102.06	102.06	102.06
TAA	6	8	10	12	14

Table 4

	Composition (Weigh Part Units)				
	8-1	8-2	8-3	8-4	8-5
Polymerizable materials	102.06	102.06	102.06	102.06	102.06
1-Octanol	6	8	10	12	14

SiHy contact lenses are prepared from those fluid polymerizable compositions according to curing processes described in Example 2, with the following curing profile: 55°C/80°C/100°C for 40 minutes at each temperature.

Example 9

In examples 4-8, five organic solvents: ethanol, 1-propanol, tert-amyl alcohol, 1-butanol and 1-octanol, are used in the lens formulations to produce hydrated contact lenses. They have a boiling point of 78.5°C, 97°C, 102°C, 117.7°C, and 194.4°C respectively. The amounts of each of the five organic solvents are varied from 5.6wt% to 12.1wt% in order to examine the effects of the concentration of each organic solvent upon the diameter and equilibrium water content of hydrated contact lenses. The diameters and equilibrium water contents of hydrated contact lenses prepared in Examples 4 to 8 are determined according to the procedures described in Example 1. The changes of lens diameter and equilibrium water content (EWC) as function of the concentration (weight percentage) of the solvent present in the lens formulation, and the averaged results are reported in Table 5.

Table 5

[Alcohol] Wt%	Ethanol		1-Propanol		Tert-Amyl Alcohol	
	D (mm)	EWC (wt%)	D (mm)	EWC (wt%)	D (mm)	EWC (wt%)
5.6	14.01±0.14	53.79±0.21	14.25±0.13	53.84±0.07	14.32±0.02	53.56±0.28
7.3	13.88±0.04	53.46±0.49	14.15±0.04	54.05±0.18	14.23±0.02	53.91±0.26
8.9	13.71±0.04	52.96±0.22	13.97±0.02	53.61±0.05	14.15±0.02	54.47±0.16
10.5	13.51±0.02	52.35±0.26	13.81±0.02	53.34±0.29	14.04±0.01	54.29±0.07
12.1	13.37±0.01	52.13±0.30	13.66±0.01	53.31±0.04	13.95±0.01	54.57±0.16

Table 5 (continue)

[Alcohol] Wt%	1-Butanol		1-Octanol	
	D (mm)	EWC (wt%)	D (mm)	EWC (wt%)
5.6	14.19±0.01	53.85±0.24	14.22±0.03	53.54±0.29
7.3	14.05±0.02	53.86±0.18	14.09±0.04	54.00±0.22
8.9	13.91±0.02	53.91±0.09	13.98±0.02	54.31±0.35
10.5	13.74±0.01	53.43±0.22	13.84±0.02	54.05±0.18
12.1	13.59±0.02	52.94±0.31	13.76±0.11	54.01±0.15

All the publications and patents which have been cited herein above are hereby incorporated by reference in their entireties.