

Description

The invention relates to prepolymers especially useful as monomers that can be formed into biomedical devices.

Heretofore, biomedical materials especially useful commercially as contact lenses have been based upon polymers and/or copolymers of a select set of chemistries. Methylmethacrylic acid type chemistries form the oldest type, the polymers from which poly(methylmethacrylates) (PMMA) have been surpassed by hydrogel chemistries based upon poly(hydroxyethyl methacrylate) (pHEMA) or polyvinyl pyrrolidinone (pNVP), and copolymers of HEMA and NVP. These materials form the basis for most soft contact lenses.

Silicone chemistries have played a lesser role in the marketplace, but have offered higher oxygen permeabilities than their hydrogel counterparts. They have, however, presented certain performance characteristics which limit their application - specifically surface wettability problems.

Copolymers employing PMMA types of chemistry have been employed in conjunction with silicone chemistry and hydrogel chemistry to produce a wide assortment of materials which have the common characteristics of high oxygen permeability and high modulus (rigidity). These materials have been characterized as hard gas permeable or rigid gas permeable materials.

Urethane chemistries have not been employed commercially in the contact lens market despite significant work in the area such as US-A-3,786,034 which relates to hard, hydrophilic polyurethane material formed from reacting a specific polyol with a polyfunctional isocyanate. US-A-3,821,186 teaches similar materials as US-A-3,786,034.

US-A-4,136,250 teaches a polymer formed by reacting a high molecular weight polydimethyl siloxane diol with 2 mole equivalents of isophorone diisocyanate and then reacting with excess hydroxy-containing monomers. Essentially, this is a soft segment prepolymer endcapped with ethylenically reactive endcaps. These materials are relatively weak and in their hydrated form show low degrees of elongation.

US-A-4,309,526 teaches adhesive compositions which employ low molecular weight polyols reacted with diisocyanates and short chain cycloaliphatic or aromatic diols, endcapped with hydrophilic endcaps. Various characteristics such as oxygen permeability are not mentioned.

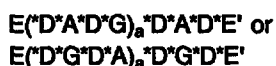
US-A-4,359,553 teaches polyurethane diacrylate compositions useful as biomedical materials which are formed by reacting a diol of MW 200 to 20,000 with 2 mole equivalents diisocyanate which is then reacted with diethyleneglycol diacrylate to form the water soluble polyurethane. Specific applications taught in the patent are as controlled release matrices.

US-A-4,740,533 teaches materials which are block copolymers of polyoxyalkylenes and polysiloxanes which have no hard segments.

US-A-4,780,488 teaches a prepolymer with only a central soft segment endcapped with hydroxyethyl methacrylate endcaps or the like.

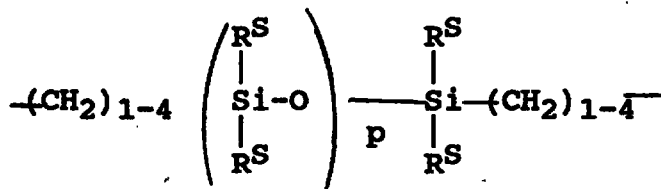
The art does not disclose polyurethane prepolymers useful as biomedical materials with the hard-soft-hard or the soft-hard-soft configurations disclosed herein which are oxygen permeable and still exhibit exemplary physical strength characteristics. Nor does the art teach that these prepolymers are useful as biomedical materials.

The present invention provides a prepolymer material consisting of prepolymers represented by the general formulae :



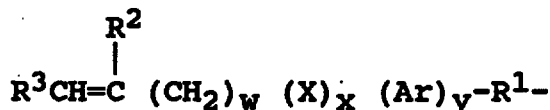
where

A denotes a divalent polymeric radical chosen from the group of



wherein

- 15 R^P denotes a straight or branched alkylene group with 1 to 6 carbon atoms and n, n' and n'' are the same or different integers such that the sum of n, n' and n'' provides for a moiety weight of said divalent polymer radical between 2,000 and 10,000;
- 20 R^F denotes a fluorinated straight or branched alkylene radical with 1 to 6 carbon atoms and m, m' and m'' are the same or different integers such that the sum of m, m' and m'' provides for a moiety weight of said divalent polymeric radical of between 400 and 10,000;
- 25 R^S denotes an alkyl radical or a short chain fluorinated alkyl radical with 1 to 3 carbon atoms; and
- P provides a moiety weight of 400 to 10,000;
- 30 D denotes an alkylene diradical, an alkyl cycloalkylene diradical, a cycloalkylene diradical, an alkylarylene diradical or an arylene diradical, with 6 to 30 carbon atoms;
- G denotes an alkylene diradical, a cycloalkylene diradical, an alkyl cycloalkylene diradical, an aromatic diradical or an alkylaromatic diradical with 1 to 40 carbon atoms which may have ether, thio, or amine linkages in the main chain;
- $*$ denotes a urethane or ureido linkage; and
- E and E' denote polymerizable unsaturated organic radicals represented by the general chemical formula

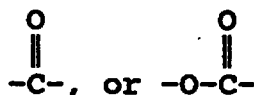


wherein

- 40 R^1 denotes a divalent alkylene radical with 1 to 10 carbon atoms;
- R^2 denotes a -H or -CH₃ radical;
- R^3 denotes a -H radical or an alkyl radical with 1 to 6 carbon atoms or a



- 50 radical where
- Y is -O-, -S- or -NH- and R^4 denotes an alkyl radical with 1 to 12 carbon atoms;
- X denotes

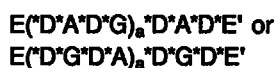


Ar denotes an aromatic radical with 6 to 30 carbon atoms;
 a is at least 1;
 w is 0 to 6;
 x is 0 or 1; and
 y is 0 or 1.

These prepolymers are especially useful in forming copolymerizates with ethylenically unsaturated monomers which are known and used in the biomedical materials field. The resultant copolymers have a combination of oxygen permeability, surface wettability and physical strength in dry and/or hydrated forms otherwise unavailable.

The prepolymers of the present invention are especially useful in making biomedical materials due to this combination of physical strength and high oxygen permeability when copolymerized with state of the art ethylenically unsaturated biomedical monomers. The combination of advantageous properties is achieved due to the specific chemistry inherent in the claimed prepolymer.

The prepolymers of the invention can be represented by the general formulae:



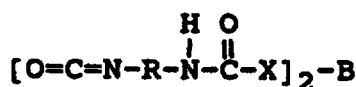
wherein A, D, E, E', G, * and a are as defined hereinabove.

The prepolymers upon polymerization form two domains which can be characterized as hard and soft domains, respectively. The soft domain generally have glass transition temperatures (Tg s) lower than room temperature whereas the hard domains have Tg s higher than room temperature. Upon polymerization, the hard segments of the prepolymer associate with one another and the soft segments form the soft domain which account for the oxygen permeability of the polymeric mass. The combination of hard and soft segments provides the polymer with superior mechanical properties otherwise unavailable.

The hard segments of the prepolymer are formed by the reaction of the short chain diol with the diisocyanate. Thus, in the formulae *supra*, the "hard segments" are represented by the "D*G*D*" portions of the formulae. Thus termed, formula (i) represents a hard-soft-hard prepolymer and formula (ii) represents a soft-hard-soft prepolymer.

The isocyanates which can be used in preparation of the invention include, toluene diisocyanate, 4,4'-diphenyl diisocyanate, 4,4'-diphenylene methane diisocyanate, p-phenylene diisocyanate, dianisidine diisocyanate, 1,5 naphthalene diisocyanate, 4,4'-diphenyl ether diisocyanate, 4,4'-(dicyclohexyl)methane diisocyanate, 1,3-bis-(isocyanato methyl)cyclohexane, cyclohexane diisocyanate, tetrachlorophenylene diisocyanate, isophorone diisocyanate, and 3,5-diethyl-4,4'-diisocyanato diphenyl methane.

Other diisocyanates which may be used are higher molecular weight diisocyanate formed by reacting polyamines which are terminally capped with primary or secondary amines, or polyhydric alcohols with excess of any of the above described diisocyanates. In general, these high molecular weight diisocyanates will have the general formula



wherein R is a divalent organic radical with 2 to about 20 carbon atoms, X is -O-, or -NR'-, where R is -H or a lower alkyl, and B is a divalent organic radical.

The diisocyanate is reacted with low molecular weight diols or glycols such as 2,2-(4,4' dihydroxydiphenyl)propane (bisphenol-A), 4,4'-iso-propylidene dicyclohexanol (hydrogenated biphenol-A), ethoxylated bisphenol-A, propoxylated bisphenol-A, 2,2-(4,4'-dihydroxydiphenyl)pentane, α,α' -(4,4'-dihydroxydiphenyl)-p-diisopropyl benzene, 1,3 cyclohexane diol, 1,4-cyclohexane diol-1,4-cyclohexane dimethanol, bicyclic and tricyclic diols such as 4,8-bis-(hydroxymethyl)-tricyclo [5.2.1.0^{2,6}] decane, neopentyl glycol, 1, 4 butanediol, 1,3-propanediol, 1,5-pentanediol, diethylene glycol, triethylene glycol and the like.

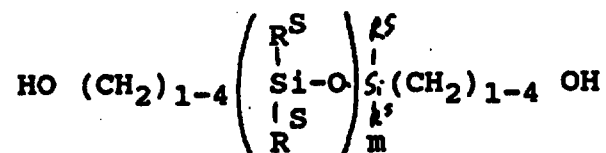
These hard segments as mentioned before form hard domains in the final polymer or copolymer by association via hydrogen bonding with other rigid segments. The degree of association within the hard domain can be modified by controlling the amount of hydrogen bonding between the segments by either 1) decreasing the overall weight content of the hard segment in the prepolymer by increasing the molecular weight of the soft segment or 2) by decreasing the amount of hydrogen bonding density in the hard segment by either using relatively soft, longer chained diols, or by using primary amines or secondary

amines capped low molecular weight compounds in conjunction with the diisocyanates rather than the diols.

The hard segments are then reacted with a relatively high molecular weight polymer which is α,ω -endcapped with two active hydrogens, usually hydroxyl groups. These segments form the so-called soft segment of the prepolymer. Various types of high molecular weight polymers can be used including in

general polymers of the following formulae

- a) $\text{HO}(\text{CH}_2)(\text{R}^{\text{P}}\text{O})_n(\text{CH}_2)\text{OH}$;
- b) $\text{HO}(\text{CH}_2)(\text{R}^{\text{P}}\text{O})_m(\text{CH}_2)\text{OH}$; and
- c)

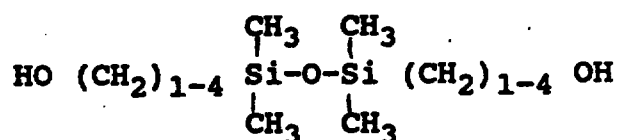


Formulae a) represents polyoxyalkyleneglycols which are generally commercially available in the molecular weight range called for in the present invention. These diols include polymers prepared from the epoxides: ethylene oxide, 1,2-propylene oxide, 1,2-butylene oxide, 2,2-epoxydecane, 1,2-epoxyoctane, 2,3-epoxy norborane, 1,2-epoxy-3-ethoxy propane, 2,2-epoxy-3-phenoxypropane, 2,3-epoxypropyl-4-methoxy phenyl ether, tetrahydrofuran, 1,2-epoxy-3-cyclohexyloxy propane, oxetane, 1,2-epoxy-5-hexene, 1,2-epoxyethylbenzene, 1,2-epoxy-1-methoxy-2-methylpropane, benzyloxy propylene oxide and combinations thereof.

The preferred polymers of this class are polypropylene glycols of molecular weights, 2000, 3000 and 4000 and more, and polyoxyethylene polyoxypropylene block copolymers with molecular weight greater than 2000.

Formulae b) represents polyfluoroethers with α,ω -active hydrogens. This class of polymers can be synthesized per the heading of US-A-3,810,874. Generally, these polymers should have molecular weights between 400 and 10,000 to be useful in the present invention.

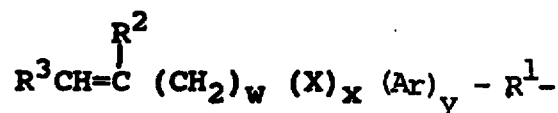
Formulae c) represents α,ω -dihydroxyl alkyl endblocked polysiloxane which for the purpose of the present invention should have a molecular weight in the range of 400 to 10,000. These polysiloxanes can be synthesized by reacting a disiloxane of the general formula



with cyclopolydimethyl siloxane under acidic conditions.

Alternately, the disiloxane can be replaced with dimethoxydimethylsilane or diethoxy dimethyl silane to produce the α,ω -dihydroxy endcapped polysiloxanes.

The endcapping monomers used in the prepolymer are generally represented by the formula



as defined *supra*. The Stage B reaction product is reacted with an excess of suitable acrylate or methacrylate esters containing a hydroxy or amine group on the non-acrylate or non-methacrylate portion of the monomer to form the endcaps of the above formula. Suitable endcap monomers include hydroxyethyl acrylate, hydroxyethyl methacrylate, aminoethyl methacrylate, 3 hydroxypropyl methacrylate, amino propyl methacrylate, hydroxyhexylacrylate, t-butylaminoethyl methacrylate, monoacrylate or monomethacrylate esters of bisphenol-A and/or bisphenol-B.

The prepolymers of the present invention are formed by two general synthetic approaches. One approach produces the hard-soft-hard prepolymer while the second approach produces the soft-hard-soft prepolymer. Variations of each scheme will be found to be necessary for specific rigid/soft segment combinations, as illustrated in the examples.

HARD-SOFT-HARD PREPOLYMER SYNTHETIC SCHEMES

The scheme used to produce this type of prepolymer employs three stages to produce the final prepolymer. The first stage (STAGE A) involves reacting 2 mole equivalents of diisocyanate with about 1 mole equivalent low molecular weight diols described herein. If these diols are represented by the symbol $\Delta G \Delta$, where Δ denotes a hydroxyl radical and G represents the rest of the diol compound, and the diisocyanate functional compound is represented by $\bullet D \bullet$ where $\bullet$ represents an isocyanate radical, the STAGE A reaction can be schematically represented as follows:



where * denotes a urethane or a ureido linkage. STAGE A produces a so-called "Hard" segment. As is known to those skilled in polymer chemistry, the product $\bullet D^* G^* D \bullet$ is the mathematical average of all reaction product molecules. The reaction product of the actual reaction will contain $\bullet O \bullet$ and $\bullet D^* (G^* D)_c^* G^* D$ with $c \geq 2$. Again, the formulae are numerical averages.

STAGE B involves reacting about one half mole equivalent of a α, ω -diol endcapped long chain polymer with the reaction product of STAGE A. If $\Delta A \Delta$ represents the long chain diol the STAGE B Reaction is



In STAGE C, the reaction product from STAGE B is reacted with a molar excess of an endcapping monomer which has: 1) hydroxyl or amine functionality; and 2) some polymerizable unsaturation. If the endcapper is represented by the symbol $E \Delta$, where Δ is $-OH$ or $-NH_2$ or $-NH-$, the reaction proceeds generally as



Optionally, STAGE B can be run with molar excess of A to produce multiblock polymers of the general formula $\bullet (D^* G^* D^* A)_a^* D^* G^* D \bullet$ where a is at least 1. This reaction product would be endcapped in STAGE C above.

SOFT-HARD-SOFT PREPOLYMER SYNTHETIC SCHEME

The second general synthetic scheme using the same nomenclature described is represented by the following general formulae:

STAGE A



STAGE B



STAGE C



In general, each of the reaction stages is run until the reactive step is complete. Reaction progress in STAGES A and B reactants may be monitored by acid base titration. The isocyanate content is calculated by the difference of acid equivalents between a stock solution dibutylamine and its reaction product with the diisocyanate reaction intermediate. The reaction can also be monitored by ATR-IR for the appearance/disappearance of peaks at 1700 cm^{-1} , which indicates the presence of



5 and 2250 cm^{-1} which indicates consumption of $-\text{N}=\text{C}=\text{O}$.

It is found that the synthesis of the prepolymer can be run neat or in solution. A wide range of aprotic solvents can be used to synthesize the prepolymers of the present invention. Solvents useful in the synthesis include toluene, methylene chloride, benzene, cyclohexane, hexane, heptane and the like. Preferred solvents are toluene, methylene chloride and mixtures thereof.

10 Reaction of the prepolymer precursors may be accomplished in the presence or absence of catalysts for urethane reactions, such catalysts being well known in the art. The first step of prepolymer synthesis where diisocyanate is first reacted with a short carbon chain (2 to 30 carbon atoms) diol, particularly where an aromatic diisocyanate is used, proceeds very rapidly, even in the absence of any catalyst. In fact, during the step of reacting diisocyanate and short chain diol, temperature control may be required in order to
15 avoid/minimize side reactions.

Preferably, the first step of prepolymer synthesis in accordance with the present invention is carried out below about 100°C, most suitably within the range of from about 60°C to about 90°C. Thereafter, the second step of the reaction is carried out at comparable temperatures, preferably within the range of from about 40°C to 70°C. The final step of prepolymer formation suitably is effected at temperatures of from
20 about room temperature to about 100°C, with a narrow range of from about 40°C to about 50°C being most preferred. As will be apparent to those skilled in the art, optimal reaction conditions, such as temperatures and duration, are selected for each individual reaction system to achieve conditions that produce a favorable rate of reaction without fostering undesirable side reactions.

Among the suitable catalysts for use in prepolymer formation are tin salts and organic tin esters, such
25 as dibutyl tin dilaurate, tertiary amines, such as triethyl diamine and other recognized catalysts, such as 1,4-diaza (2.2.2)-bicyclooctane (DABCO).

The prepolymers of the present invention are particularly useful as comonomers with ethylenically reactive monomers useful in the field of biomedical materials. In general, these monomers are the hydroxyalkyl acrylates and diacrylates such as hydroxyethyl acrylate, hydroxypropyl acrylate, and the
30 corresponding methacrylate compounds, including cyclohexyl methacrylate, methyl methacrylate, isobornyl methacrylate, lauryl methacrylate, triethylene glycol dimethacrylate, isobutyl methacrylate and tetrahydrofurfuryl methacrylate. Other useful unsaturated reactive monomers are acrylamides, methacrylamides, pyrrolidinones, styrene and acrylonitrile and other monomers known in the art including fluorinated analogs of all of the previously mentioned monomers and the organo silicone comonomers known in the art. Specific
35 fluorocomonomers include:

(2,2,2-trifluoroethyl) itaconate
(hexafluoroisopropyl) itaconate
(1H, 1H-perfluorooctyl) itaconate
(1H, 1H, 11H-perfluoroundecyl) itaconate
40 (perfluoro-t butyl) itaconate
(pentafluorophenyl) itaconate
(2H, 2H-perfluorobenzyl) itaconate
(pentafluorophenylmethyl) itaconate
(decafluorocyclohexyl) itaconate
45 (1H-perfluorocyclohexyl) methyl itaconate
(1,1,1-trifluoroisopropyl) itaconate
1-methyl-4-(hexafluoroisopropyl) monoitaconate
4-(hexafluoroisopropyl) monoitaconate
1-(1H, 1H-perfluorooctyl)-4-hexafluoroisopropyl itaconate

50 and methacrylate analogs thereof.

Specific organosilicon comonomers include:
tris(2-acetoxyethyl)dimethylsiloxy)silylpropyl acrylate and methacrylate
tris(2-carboxyethyl)dimethylsiloxy)silylpropyl acrylate and methacrylate
tris(3-hydroxypropyl)dimethylsiloxy)silylpropyl acrylate and methacrylate
55 acrylate and methacrylate functional, fluorosubstituted alkyl/aryl siloxanes such as:
tris(3,3,3 trifluoropropyl dimethylsiloxy) silyl propyl acrylate and methacrylate
tris[3-heptafluoroisopropoxy propyl] dimethylsiloxy silylpropyl acrylate and methacrylate
tris(pentafluorophenyl dimethylsiloxy)silyl propyl acrylate and methacrylate

Other potentially useful organosilicon comonomers include:

p-(pentamethyldisiloxanyl) styrene

bis(trimethylsiloxy)

pyrrolidinonyldimethyl

5 siloxy-silylpropyl acrylate and methacrylate.

When used as comonomers these materials can be used from 5 to 85 weight percent of the final copolymer weight with the balance comprising the prepolymer portion.

Other di-ethylenically reactive monomers can also be used to effect the mechanical and surface properties. Such crosslinks are generally employed in the 0.1 to 5 wt % range.

10 The polymers and copolymers are formed by a free radical mechanism using a wide variety of known free radical catalysts such as the diacyl peroxides such as benzoyl peroxide; dialkyl peroxides such as di-tert, -butyl peroxide; ketone peroxides such as methylethyl ketone peroxide; and peresters which readily hydrolyze, e.g. tert-butyl peracetate, tert-butyl perbenzoate, di-tert-butyl diperphthalate, etc. A particularly useful class of peroxy initiators are the organic hydroperoxides such as cumene hydroperoxide, methylethyl 15 ketone hydroperoxide, tert-butyl hydroperoxide, etc. The initiators should be used at a concentration of about 0.01 percent to about 10 percent by weight of the total formulation, preferably about 0.1 percent to about 5 percent by weight. Another useful class of initiators comprises carbonyl-containing ultraviolet-activated free radical generators, such as acetophenone, benzophenone, and the benzoin ethers. Other suitable UV initiators are known in the art. Initiator mixtures may also be used.

20 Solvents can be used in the final copolymerization and/or polymerization process. Solvent choice will depend upon the solubility parameters of the prepolymer and of the comonomers used, if any, and should be chosen to allow full solubilization of all polymerizable components.

In certain instances, the copolymerization process should be carried out without solvent. For instance, when 2-hydroxyethyl methacrylate (HEMA) is copolymerized with one of the prepolymers formed with 25 polyethylene glycol, use of toluene causes the HEMA to form heterogenous domains which are not stable under aggressive hydrolytic conditions.

Some of the preferred copolymers are polymerized from the following comonomer mixtures:

<u>Copolymer A</u>	<u>Wt. %</u>	<u>Component</u>
	80	prepolymer of the hard-soft-hard configuration made from 2-hydroxyethyl methacrylate, isophorone diisocyanate, neopentyl glycate and disilanol endcapped polydimethylsiloxane of molecular weight equal to 3000 (INS3H)
	20	2-hydroxyethylmethacrylate

Copolymer B

30-45 INS3H
 40-55 tris(trimethylsiloxy)silylpropyl
 15-25 methacrylate
 N-N dimethylacrylamide

Copolymer C

30-90 prepolymer of the hard-soft-hard
 configuration made from hydroxy-
 ethylmethacrylate, isophorone
 diisocyanate 1,4-butanediol and
 disilanol endcapped polydimethyl-
 siloxane of average molecular weight
 of about 3000 (IBS3H) or similar
 polysiloxane of about 4000 molecular
 weight (IBS4H)
 10-40 dimethylacrylamide

Copolymer D

30-90 IBS4H or IBS3H
 20-40 dimethylacrylamide
 5-30 tris(trimethylsiloxy)silyl propyl
 methacrylate

Copolymer E

30-90 prepolymer of the hard-soft-hard
 configuration made from 2-hydroxy-
 ethylmethacrylate, isophorone
 diisocyanate, diethylene glycol, and
 the disilanol endcapped polydimethyl-
 siloxane of molecular weight 3000 or
 4000 (IDS3H and IDS4H, respectively)
 20-40 dimethylacrylamide
 1-30 tris(trimethylsiloxy)silyl propyl
 methacrylate

Various homopolymers and copolymers films can be formed and characterized by standard testing procedures such as:

1. Tensile strength (g/mm²) and modulus of elasticity can be measured per ASTM test method D1708.
2. Elongation can be measured per ASTM 1708.
3. Initial tear strength and propagation tear strength can be measured per ASTM 1438.
4. Oxygen permeabilities can be measured by the method reported by Relojo, M. et al in Contact and Intraocular Lens Medical Journal, Vol. 3, issued p. 27 (1977) and edge effects can be accounted for per the methods described by Fatt, et al. in International Contact Lens Clinic, V. 14, p. 389 (1987).
5. Water content can be measured per a gravimetric method.
6. Refractive index can be measured per typical methods on hydrated samples using a refractometer.

As mentioned, the prepolymers of the present invention are particularly useful in forming shaped articles used in biomedical applications. These polymers and copolymers can be used to make biomedical devices i.e. shaped articles, such as dialyzer diaphragms, to prepare artificial kidneys and other biomedical implants, such as disclosed in US-A-2,976,576 and US-A-3,220,960. The instant polymers and copolymers can be used in preparing therapeutic bandages as disclosed in US-A-3,428,043. The instant polymers and copolymers can also be used in preparing medical surgical devices e.g. heart valves, vessel substitutes, intrauterine devices, membranes and other films, dialyzer diaphragms, catheters, mouth guards, denture liners and other such devices as disclosed in Shephard US-A-3,618,231. The instant polymers and copolymers can be used to modify collagen to make blood vessels, urinary bladders and other such devices as disclosed in US-A-3,563,925. The instant polymers and copolymers can be used to make catheters as disclosed in US-A-3,566,874. The instant polymers and copolymers can be used as semipermeable sheets for dialysis, artificial dentures and all of such disclosures as set forth in US-A-3,607,848. The instant polymers and copolymers can be used in ophthalmic prostheses and all other uses disclosed in US-A-3,679,504. They may also be used as a polymeric matrix for controlled release of active pharmaceutical agents.

In the following examples which illustrate the invention, the properties of such films derived from the claimed prepolymer and combinations of prepolymers with various comonomers are described. For certain comonomers which employed "hydrophilic" comonomers, the films were hydrated to "hydrogel" states- their physical properties were measured as hydrated.

EXAMPLE 1

SYNTHESIS OF HARD-SOFT-HARD PREPOLYMER BASED ON TOLUENE-2,4-DIISOCYANATE

2 mole equivalents of toluene-2,4-diisocyanate (TDI) were dissolved in toluene with one mole equivalent hydrogenated bisphenol A (HBPA) in a resin kettle with constant stirring under nitrogen. This STAGE A reaction was run at 60-90 °C for 2-3 hours in the presence of 0.5 wt% "DABCO" catalyst.

One mole equivalent of polypropylene glycol with a molecular weight of about 1000 was then slowly added to the reaction mixture and reacted for 2-3 hours at 60-80 °C under nitrogen.

Excess 2-hydroxyethyl methacrylate (HEMA) was added to the isocyanate terminated STAGE B reaction mixture and then reacted to completion.

Variations of the above synthesis were run by substituting neopentylglycol (NPG) for the hydrogenated Bisphenol A, and by substituting various molecular weight polypropylene glycols for the PPG-1000.

The resulting prepolymers were homopolymerized between treated glass plates in the presence of a free radical catalyst, benzoin methylether. The polymerization was UV initiated in toluene. The solvent was removed by vacuum and the resultant films were characterized. The results obtained are reported in TABLE

1.

The oxygen permeability of the homopolymers increase as the molecular weight of the polypropylene glycol increased.

EXAMPLE 2

HARD-SOFT-HARD PREPOLYMER USING ISOPHORONE DIISOCYANATE (IPDI)

The same basic synthetic procedure as described in the previous example was employed except that, due to the decreased reactivity of isophorone diisocyanate relative to toluene diisocyanate, higher levels of reaction catalyst were used. Reaction conditions and summary of the cured homopolymer films is reported in TABLE 2.

Once again, the oxygen permeability of the cured prepolymer increased with increasing molecular weight of the polypropylene glycol polymer. These prepolymers did not produce yellowed films as the TDI

materials did upon hydrolytic testing.

TABLE 1

<u>Composition</u>	<u>R_x Temp</u>	<u>Solvent</u>	<u>Catalyst</u> (Dabco)	<u>O₂ Permeability (DK)</u>
TD1-NPG-PPG-1000-HEMA	45	methylene chloride	A=0 B,C=0.05 wt%	2
TD1-NPG-PPG-2000-HEMA	45	methylene chloride/ 2 toluene	A=0 B,C=0.05	9
TD1-NPG-PPG-3000-HEMA	45	toluene	A=0 B,C=0.05	20
TD1-NPG-PPG-4000-HEMA	45	2 toluene/ 1 methylene chloride	A=0 B,C=0.05	30
TD1-HBPA-PPG-1000-HEMA	A ₁ B=75 C=60	toluene/ methylene chloride	A=0 B,C=0.05	2
TD1-HBPA-PPG-2000-HEMA	A ₁ B=75 C=60	toluene/ methylene chloride	A=0 B,C=0.05	5
TD1-HBPA-PPG-3000-HEMA	A ₁ B=75 C=60	toluene/ methylene chloride	A=0 B,C=0.05	15
TD1-HBPA-PPG-3000-HEMA	A ₁ B=75 C=60	toluene/ methylene chloride	A=0 B,C=0.05	25

TABLE 2 - IPDI BASED HARD-SOFT-HARD FILMS

Composition	Solvent	Rx Temp			Catalyst *			Oxygen Permeability (DK)	Contact Angle (°)
		A	B	C	A	B	C		
IPDI-NPG-PPG-1000-HEMA	Toluene	80	80	50	0.25	0.25	0.25	3.4	53
" -2000-HEMA	Toluene	80	80	60	0.5	0.5	0.5	13.8	35
" -3000-HEMA	Toluene	80	80	60	0.5	0.5	0.5	24.3	42
" -4000-HEMA	Toluene	75	85	50	0.25	0.25	0.25	33.2	41
IPDI-CHDM-PPG-1000-HEMA	Toluene	85	85	50	0.25	0.25	0.25	2.0	59
" -2000-HEMA	Toluene	85	85	45	0.5	0.25	0.25	10.9	43
" -3000-HEMA	Toluene	80	80	60	0.5	0.25	0.25	21.0	42
" -4000-HEMA	Toluene	85	85	50	0.25	0.25	0.25	30.1	41
IPDI-HBPA-PPG-1000-HEMA	Toluene	85	85	50	0.25	0.25	0.25	1.5	62
" -2000-HEMA	Toluene	85	85	45	0.25	0.5	0.5	8.0	41
" -3000-HEMA	Toluene	85	85	45	0.3	0.3	0.3	17.4	44
" -4000-HEMA	Toluene	80	80	50	0.25	0.25	0.25	24.1	44

EXAMPLE 3**4,4-DI-CYCLOHEXYL METHANE DIISOCYANATE BASED HARD-SOFT-HARD PREPOLYMERS**

5 Hard-soft-hard prepolymers were synthesized per the examples supra except that H₁₂-MDI was used. Films were cured from 30 • % prepolymer solution in toluene as reported in TABLE 3.

In general, the H₁₂-MDI prepolymers produced films with greater tear strengths than the TDI or IPDI diisocyanate based systems and were slightly less oxygen permeable.

10 **EXAMPLE 4**

COPOLYMERS OF IPDI BASED HARD-SOFT-HARD PREPOLYMERS WITH HYDROPHILIC MONOMERS

15 A hard-soft-hard prepolymer produced with isophorone diisocyanate (IPDI), neopentyl glycol (NPG), polypropylene glycol with a molecular weight of 4000 and 2-hydroxethyl methacrylate was copolymerized with a hydrophilic comonomer. Various copolymers of this prepolymer (INP4H) with various comonomers were made and tested.

The general scheme for making the copolymer involved mixing the prepolymer with the comonomer in a 30-70 wt% solution in toluene in the presence of a known free radical catalyst. The solutions were placed
20 between treated glass plates and polymerized. The films were then physically characterized.

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TABLE 3 - H₁₂-MDI HARD-SOFT-HARD PREPOLYMERS

Composition	Oxy (DKl)	Contact Angle (°)	Tear (P) (g/mm)	Tensile Strength (g/mm ²)	Modulus (g/mm ²)	Elongation (%)
NPG PPG-1000	2.3	41	*	132.9	828	170.1
PPG-2000	10.7	39	187	743.2	931.2	172.5
PPG-3000	20.4	33	91.8	514.5	386.5	190.8
PPG-4000	29.3	39	108	201.5	185.1	218.2
CHDM PPG-1000	1.8	44	*	1282	16000	173.8
PPG-2000	8.4	41	437	612.7	1676	282.1
PPG-3000	18.6	38	290	399.9	373.2	348.7
PPG-4000	28.0	41	170	169.9	170	301.9
HBPA PPG-1000	1.2	62	*	2271	32610	84.6
PPG-2000	6.4	43	256	1160	3590	221.7
PPG-3000	14.2	41	197	672.2	1607	217.5
PPG-4000	23.2	44	130	433.9	615.9	198.6

* too stiff to measure

Specific hydrophilic comonomers used were 2-hydroxyethyl methacrylate and N-2-vinylpyrrolidinone. The characteristics of these copolymers are disclosed in TABLE 4.

Several interesting features are displayed by the resultant films. In general, the contact angle of the material was lowered as the content of hydrophilic comonomers increased, as did water content. However, oxygen permeability decreased as the water content of the hydrated material increased. Normally, water

content and O₂ permeability of hydrogel materials is directly related rather than inversely related.

EXAMPLE 5

5 POLYMER ALLOYS OF POLYMERIZED HARD-SOFT-HARD MATERIALS

A homopolymer of INP4H was prepared and then soaked in a solution of N-vinyl pyrrolidinone containing a free radical catalyst. The films were then subjected to UV radiation in order to polymerize the N-vinylpyrrolidone. Based upon analysis of the extracted parties of the polymerized polymer alloy, wt% of
10 polymerized N-vinyl pyrrolidinone could be calculated. The characteristics of these polymer alloys are reported in TABLE 5.

It was observed that these alloy materials felt much more slippery than the copolymers made from the same monomer combinations.

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TABLE 4 - IPDI BASED HARD-SOFT-HARD/HYDROPHILIC COMONOMERS

Copolymer Composition		Water Content (%)	O ₂ (DK)	Contact Angle (°)	Tear (p) Strength (g/mm)	Tensile Strength (g/mm ²)	Tensile Modulus (g/mm ²)	Elong. %
INP4H	: HEMA							
100	: 0	--	33.2	41	74.2	219.3	117.8	294
80	: 20	8.3	19.7	36	55.2	65.4	162	285
60	: 40	19.4	13.3	38	31.0	97.4	91.1	258
40	: 60	26.5	10.4	37	13.0	58.8	80.0	155
0	: 100	38.6	9	--	4.2	56	57	225
INP4H	: NVP							
90	: 10	9.0	15.9	45	--	--	--	--
80	: 20	17.7	24.4	37	15.1	55.8	71.4	243
60	: 40	43.7	25.7	35	5.3	77.7	117.9	174
40	: 60	58.5	31.6	37	2.5	17.4	50.8	72.4

TABLE 5 - POLYMER ALLOYS

	Alloy Composition	Water Content (%)	O ₂ (DK)	Contact Angle (°)
	INP4H : NVP (IPN)			
	96.6 : 3.4	7.9	26.2	55
	94.8 : 5.2	12.6	25.7	43
	89.5 : 10.9	15.8	25.5	46
	81.3 : 18.7	27.7	17.9	40
	78.2 : 21.8	31.3	20.6	39
	74.8 : 25.2	38.7	22.9	38

EXAMPLE 6

COPOLYMERS OF HARD-SOFT-HARD PREPOLYMER (INP4H) WITH FLUOROMONOMERS

In the interest of producing copolymers with less lipid uptake values certain fluorinated comonomers, octafluoropentyl methacrylate (OFPMA) and hexafluoro isopropyl methacrylates, (HFIPMA) were copolymerized with INP4H prepolymers in various ratios. The cured copolymer films are characterized in TABLE 6.

As the fluoro content of the copolymer was increased the oleic acid pickup decreased with a concomitant decrease in oxygen permeability. The samples with 80 wt% fluoromonomer content were too brittle to characterize.

EXAMPLE 7

COPOLYMERS OF INP4H WITH SILICONE MONOMERS

Various copolymers were produced from combinations of INP4H with tris(trimethylsiloxy)silylpropyl methacrylate (referred to as tris) and 2-hydroxyethyl methacrylate and characterized in TABLE 7.

These copolymers exhibit very good oxygen permeabilities and oleic acid uptake characteristics as well as vastly superior tear strengths and tensile moduli with respect to pHEMA.

TABLE 6 - COPOLYMERS OF INP4H WITH FLUORINATED COMONOMERS

Composition	O ₂ (DK)	Contact Angle (°)	Tear Strength g/mm ²	Tensile Strength g/mm ²	Tensile Modulus g/mm ²	Elongation (%)
INP4H : OFPMA						
80 : 20	23.9	40	107	238.1	254.6	225
60 : 40	20.8	45	155	412.8	365.4	320
40 : 60	17.6	50	447	569.7	3020	318
20 : 80	--	--	--	--	--	--
INP4H : HGIPMA						
80 : 20	28.0	40	269	405.9	615.2	263
60 : 40	24.8	42	235	1043	2408	383
40 : 60	21.3	44	--	--	--	--
20 : 80	--	--	--	--	--	--

TABLE 7 - INP4H-TRIS-HEMA COPOLYMERS

Copolymer Composition		Water Content (%)	Appear.	O ₂ (BK)	Contact Angle (°)	Protein ug/mg	Oleic (dry film) %	Tear (P) (g/mm)	Tensile Modulus (g/mm ²)	Elong (%)
INP4H : HEMA : Tris										
100 :	0 : 0	33.2	Clear	33.2	41		348	74.2	117.8	254
80 :	20 : 0	8.3	Clear	13.3	36	0	179	55.2	162	285
60 :	40 : 0	19.4	Clear	19.7	38	0	113	31.0	91.1	258
40 :	60 : 0	26.5	Hazy	10.4	37	0	43.6	13.0	80.0	155
0 :	100 : 0	38.6	Clear	9.0	--	0-1	29.0	4.2	57.0	225
60 :	20 : 20	6.4	Clear	31.7	33	0	216	50.0	55.4	345
50 :	20 : 30	7.8	Clear	42.4	40	0	192	42.6	56.3	399
40 :	20 : 40	9.8	Clear	52.9	42	0	192.0	82.2	75.4	460
40 :	50 : 10	17.5	Hazy	15.9	32	0	62.5			
40 :	40 : 20	13.4	Clear	22.9	30	0	79.6			
40 :	30 : 30	10.5	Clear	26.7	33	0	112			
40 :	20 : 40	5.7	Clear	43.5	34	0	172			

EXAMPLE 8**HYDROLYTIC STABILITY OF INP4H/HEMA COPOLYMERS**

Hydrolytic stability testings of INPD4H/HEMA copolymers formed in toluene were conducted by subjecting the copolymeric samples to hostile aqueous conditions (either acidic or basic H₂O at elevated temperatures) for extended periods of up to 14 days. These copolymers showed consistent and significant loss of mass during the hydrolytic testing which indicated that the copolymerization process was not efficient.

Chemical analysis of the extracted portion revealed that the portion being extracted was largely an oligomeric product of the 2-hydroxyethyl methacrylate.

Subsequently, INP4H/HEMA/TRIS copolymers were produced without aid of a solvent. These copolymers proved to be hydrolytically stable under the same conditions as the toluene solution produced polymers had been treated under.

EXAMPLE 9**SYNTHESIS OF HARD-SOFT-HARD PREPOLYMERS USING POLYSILOXANE DIOLS AND COPOLYMERS THEREOF**

α,ω -bis(hydroxybutyldimethylsilyl) polysiloxane was prepared by reacting dimethoxydimethylsilane with 1,3-bis(4-hydroxybutyl)tetramethyldisiloxane in water under acidic conditions. The average molecular weight of the polymer was about 3100. Using the general synthetic scheme for hard-soft-hard prepolymers with the above long chain diol, isophorone diisocyanate (IPDI), neopentylglycol (NPG) and 2-hydroxyethyl methacrylate end blocker (HEMA), a prepolymer was formed (INS3H).

This prepolymer was copolymerized with 2-hydroxyethyl methacrylate in various ratios. The characteristics of these copolymers is reported in TABLE 8.

These films were all hydrolytically stable.

EXAMPLE 10**TERPOLYMERS OF INP4H PREPOLYMERS**

Terpolymers of INP4H/TRIS/glycerol methacrylate (GM) and INP4H/TRIS/N-N-dimethylacrylamide (DMM) were prepared and evaluated. The characteristics of these films is summarized in TABLES 9A and 9B.

These terpolymers provide soft hydrogel type materials with excellent oxygen permeability and physical strength characteristics.

TABLE 9B

INP4H: TRIS: DMA	O ₂ DK
30 : 50 : 20	80
25 : 50 : 25	75

TABLE 8 - INS3H/HEMA COPOLYMERS

Composition	O ₂ (BK)	% H ₂ O	Modulus	Elongation	Tear
INS3H : HEMA					
80 : 20	262	5	6600	140	10
70 : 30	189	9	4500	170	12
60 : 40	112	13	1800	160	80

TABLE 9A. - INP4H:TRIS:GM TERPOLYMERS

<u>Formulation</u>	<u>O₂</u> <u>(DK)</u>	<u>Contact</u> <u>Angle (°)</u>	<u>% H₂O</u>	<u>Tensile</u>	<u>Modulus</u>	<u>Strength</u>	<u>Tear</u>
INP4H:TRIS:GM							
30 : 50 : 20	78	48	14	62	94	380	83
25 : 50 : 25	86	--	11	82	370	300	90
20 : 50 : 30	62	--	17	96	250	170	45

EXAMPLE 11**HARD-SOFT-HARD URETHANE SILICONE PREPOLYMERS WITH "SOFTER" HARD PORTIONS**

5 Siloxane containing prepolymers were produced as before except 1,4 butanediol or diethyleneglycol was used in place of neopentyl glycol. In general, the homopolymers and copolymers of these materials had lower modulus and tear strengths than their neopentyl glycol based counterparts.

IBS4H Prepolymer is formed using the general scheme for hard-soft-hard prepolymers using isophorone diisocyanate, 1,4-butanediol, the α,ω -hydroxyalkyl endcapped polydimethylsiloxane described
10 earlier with a molecular weight of about 4000, and 2-hydroxyethyl methacrylate.

IBS3H Prepolymer is the same prepolymer as IBS4H except with a lower molecular weight polydimethylsiloxane. IDS4H and IDS3H are equivalent to IBS4H and IBS3H, respectively, except diethyleneglycol was used in place of 1,4 butanediol.

The physical characteristics of copolymers formed from these prepolymers with dimethylacrylamide
15 (DMA) and/or tris (trimethylsiloxy)silyl propyl methacrylate are reported in TABLE 10. All of the prepolymers were hydrolytically stable.

EXAMPLE 12**20 SOFT-HARD-SOFT PREPOLYMERS**

According to the general scheme for producing soft-hard-soft prepolymers were formed using 2-hydroxyethyl-

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TABLE 10 - HARD-SOFT-HARD SILICONE PREPOLYMERS

Formulation	O ₂ (DK)	Water%	Contact Angle (°)	Tensile (g/mm ²)	Modulus (g/mm ²)	Elongation %	Tear (g/mm ²)
IBS3H : DMA							
100 : 0	399.3	0	--	347.4	1242	70.4	28.2
70 : 30	110.4		55	124.0	531.0	70.5	13.2
60 : 40	49.2		59	75.6	239.2	62.1	6.4
IBS4H : DMA							
100 : 0	146.8			193.7	574.9	82.7	11.1
70 : 30	126.5			57.2	186.5	53.3	5.0
60 : 40	71.7	38.2		49.1	155.6	52.3	3.4
IDS3H : DMA : Tris							
100 : 0 : 0	478.8	--		480.3	2861	56.1	36.9
70 : 30 : 0	67.0	26.4		65.2	220.5	45.0	3.8
60 : 30 : 10	60.1	31.1		55.7	169.2	53.5	4.9
50 : 30 : 20	45.6	40.6		48.9	140.5	56.9	2.4
40 : 30 : 30	113.1	22.4		59.5	79.8	133.8	15.1
60 : 40	36.4	38.9		58.0	223.4	37.4	2.4
IDS4H : DMA : Tris							
100 : 0 : 0	74.2			602.7	1141	41.4	--
70 : 30 : 0	120.4			52.1	175.6	46.5	2.4
60 : 30 : 0	76.4	29.2		42.7	116.5	59.0	2.7
50 : 30 : 20	113.4	28.2		34.9	68.5	88.2	4.8
60 : 40	67.0			43.6	173.8	34.8	2.5

methacrylate (HEMA), isophorone diisocyanate (IPDI), neopentyl glycol (NPG) and a disilanol endcapped polydimethylsiloxane with an average molecular weight of about 1500 (Si-1500). The general chemical structure of the prepolymer was

HEMA-IPDI-Si-1500-IPDI-NPG-IPDI-Si-1500-HEMA

Copolymers of the above prepolymer were formed with N-N-Dimethylacrylamide and tris-(trimethylsiloxy)silylpropylmethacrylate as reported in TABLE 11.

5 COMPARATIVE EXAMPLE

SOFT SEGMENT PREPOLYMER

10 To illustrate the beneficial properties of the invention a comparative prepolymer was formed from 2-hydroxyethyl methacrylate (HEMA), isophorone diisocyanate (IPDI) and an α,ω -dihydroxyalkyl endcapped polydimethyl siloxane with an average molecular weight of about 3000 (SI-3000). The structure of the prepolymer was

HEMA-IPDI-SI-3000-IPDI-HEMA

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Copolymers of this prepolymer with N-N-dimethyl acrylamide and tris(trimethylsiloxy)silylpropylmethacrylate are characterized in TABLE 12.

These copolymers displayed much lower tear, modulus and tensile strength than the copolymers of the present invention.

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TABLE 11 - SOFT-HARD-SOFT PREPOLYMERS

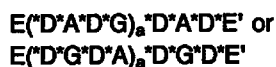
Formulation	(DK)	% Water	Contact Angle (°)	Tensile $\frac{g}{mm^2}$	Modulus $\frac{g}{mm^2}$	Elongation %	Tear $\frac{g}{mm}$
Prepolymer/DMA/Tris							
100/ 0/ 0	283	--	--	60	4500	164	70
70/30/ 0	114	25.7	51	111	350	70	35
35/30/35	114	23.8	52	84	140	150	35
40/30/30	99	24.3	--	63	100	130	35

TABLE 12 - COMPARATIVE SOFT SEGMENT PREPOLYMER

Formulation	(DK)	% Water	Contact Angle (°)	Tensile $\frac{g}{mm^2}$	Modulus $\frac{g}{mm^2}$	Elongation %	Tear $\frac{g}{mm}$
Prepolymer/DMA							
100 : 0 :	624	0	--	195	620	53	9
70 : 30 :	158	25	--	56	250	30	3

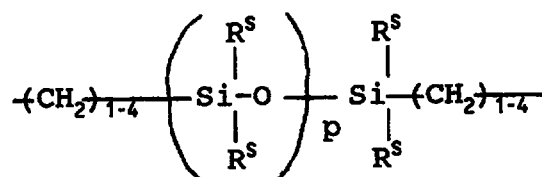
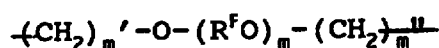
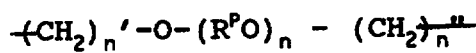
**Claims****Claims for the following Contracting States : DE, FR, GB, IT, SE**

1. A prepolymer material consisting of prepolymers represented by the general chemical formulae



where

A denotes a divalent polymeric radical chosen from the group of



wherein

R^P denotes a straight or branched alkylene group with 1 to 6 carbon atoms, and n , n' and n'' are the same or different integers such that the sum of n , n' and n'' provides for a moiety weight of said divalent polymeric radical of between 2,000 and 10,000;

R^F denotes a fluorinated straight or branched alkylene radical with 1 to 6 carbon atoms and m , m' and m'' are the same or different integers such that the sum of m , m' and m'' provides for a moiety weight of said divalent polymeric radical of between 400 and 10,000;

R^S denotes an alkyl radical or a short chain fluorinated alkyl radical with 1 to 3 carbon atoms; and

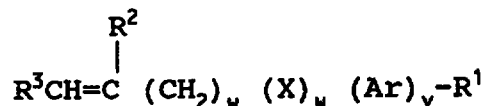
p is an integer such that the total moiety weight of said divalent polymeric radical is from 400 to 10,000;

D denotes an alkylene diradical, an alkyl cycloalkylene diradical, a cycloalkylene diradical, an alkylarylene diradical or an aryene diradical, with 6 to 30 carbon atoms;

G denotes an alkylene diradical, a cycloalkylene diradical, an alkyl cycloalkylene diradical, an aromatic diradical or an alkylaromatic diradical with 1 to 40 carbon atoms which may have ether, thio or amine linkages in the main chain;

$*$ denotes a urethane or ureido linkage; and

E and E' denote polymerizable unsaturated organic radicals represented by the general chemical formula:

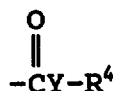


wherein

R^1 denotes a divalent alkylene radical with 1 to 10 carbon atoms;

R^2 denotes a $-H$ or $-CH_3$ radical;

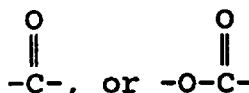
R^3 denotes a $-H$ radical or an alkyl radical with 1 to 6 carbon atoms or a



radical where

Y is -O-, -S- or -NH- and R⁴ denotes an alkyl radical with 1 to 12 carbon atoms;

X denotes



Ar denotes an aromatic radical with 6 to 30 carbon atoms;

a is at least 1;

w is 0 to 6;

x is 0 or 1; and

y is 0 or 1.

2. The prepolymer of Claim 1 wherein the polymeric radical, A, is selected from (1) a polyethylene glycol diradical with a molecular weight of at least about 2000; (2) a dialkyl endcapped polydimethylsiloxane radical with a molecular weight of at least 400; (3) dialkyl endcapped polydifluoroalkylsiloxane polymer with a molecular weight of at least 400; and (4) a fluoroalkyl ether with a molecular weight of at least about 400.
3. A prepolymer according to Claim 2, wherein the D radical is formed from the group of diisocyanates consisting of isophorone diisocyanate, toluene diisocyanate and 4,4'-dicyclohexyl methane diisocyanate.
4. A prepolymer according to Claim 2 or Claim 3, wherein the G radical is formed from the group of diols consisting of neopentyl glycol, 1,4-butanediol, diethylene glycol and triethylene glycol.
5. A copolymer material formed by polymerizing a comonomer mixture comprised of:
 - (I) the prepolymer described by the general chemical formulae:

$$\begin{array}{l} \text{E}(\text{D}^*\text{A}^*\text{D}^*\text{G})_a\text{D}^*\text{A}^*\text{D}^*\text{E}' \text{ or } \\ \text{E}(\text{D}^*\text{G}^*\text{D}^*\text{A})_a\text{D}^*\text{G}^*\text{D}^*\text{E}' \end{array}$$
 wherein,
 - A, D, E, E', G, * and a are as defined in Claim 1, and
 - (II) an ethylenically unsaturated comonomer.
6. The copolymer of Claim 5 wherein the polymeric radical of component (I), A, is polyethylene radical with a molecular weight of at least about 2,000.
7. The copolymer of Claim 6 wherein component (I) comprises 15 to 95 weight percent of the comonomer mixture and said ethylenically unsaturated comonomer comprises 5 to 85 weight percent of said comonomer mixture.
8. The copolymer of Claim 7 which further comprises a diethylenically reactive crosslinker.
9. The copolymer of Claim 8 wherein said ethylenically unsaturated comonomer is chosen from 2-hydroxyethyl methacrylate, N-vinyl-2-pyrrolidinone, methacrylic acid, dimethyl acrylamide and glycerol methacrylate.
10. The copolymer of Claim 5 wherein the polymeric radical of component (I), A, is a dialkyl endcapped polydimethylsiloxane radical with a molecular weight of at least 400.



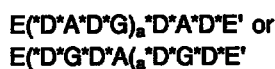
11. The copolymer of Claim 10 wherein component (I) comprises 15 to 95 weight percent of the comonomer mixture and said ethylenically unsaturated comonomer comprises 5 to 85 weight percent of said comonomer mixture.

12. The copolymer of Claim 11 wherein said ethylenically unsaturated comonomer is chosen from 2-hydroxyethyl methacrylate, N-vinyl-2-pyrrolidinone, methacrylic acid, dimethyl acrylamide and glycerol methacrylate.

13. The copolymer of Claim 12 which further comprises 5 to 40 weight percent organosilicon comonomer.

14. The copolymer of Claim 5 wherein the polymeric radical of component (I), A, is a poly(fluoroalkyl)ether with a molecular weight of at least 400.

15. A copolymer formed by polymerizing a comonomer mixture comprised of:
(I) the prepolymer described by the general chemical formulae:



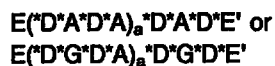
wherein A, D, E, E', G, * and a are as defined in Claim 1, and
(II) an organosilicon comonomer.

16. The copolymer of Claim 15 wherein said organosilicon monomer is tris(trimethylsiloxy)silylpropyl methacrylate.

17. The copolymer of Claim 16 which comprises 20-50 wt % prepolymer (I), and 20-40 weight percent tris(trimethylsiloxy)silyl propyl methacrylate.

18. The copolymer of Claim 17 which further comprises 10-35 weight percent of one of the following hydrophilic comonomers: dimethacrylamide; glycerol methacrylate; 2-hydroxyethyl methacrylate; and N-vinyl-2-pyrrolidinone.

19. A copolymer formed by polymerizing a comonomer mixture comprised of:
(I) the prepolymer described by the general chemical formulae:



wherein A, D, E, E', G, * and a are as defined in Claim 1, and
(II) an ethylenically unsaturated fluoroorgano comonomer.

20. A biomedical device made from a polymer comprising the prepolymer of any preceding claim.

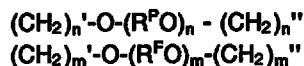
Claims for the following Contracting State : ES

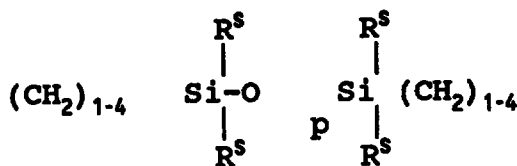
1. A process for preparing a prepolymer material consisting of prepolymers represented by the general chemical formula:



where

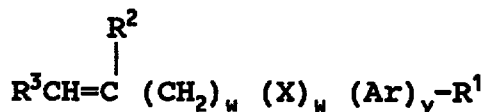
A denotes a divalent polymeric radical chosen from the group of





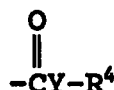
wherein

- R^p denotes a straight or branched alkylene group with 1 to 6 carbon atoms, and n, n' and n'' are the same or different integers such that the sum of n, n' and n'' provides for a moiety weight of said divalent polymeric radical of between 2,000 and 10,000;
- R^f denotes a fluorinated straight or branched alkylene radical with 1 to 6 carbon atoms and m, m' and m'' are the same or different integers such that the sum of m, m' and m'' provides for a moiety weight of said divalent polymeric radical of between 400 and 10,000;
- R^s denotes an alkyl radical or a short chain fluorinated alkyl radical with 1 to 3 carbon atoms; and
- p is an integer such that the total moiety weight of said divalent polymeric radical is from 400 to 10,000;
- D denotes an alkylene diradical, an alkyl cycloalkylene diradical, a cycloalkylene diradical, an alkylarylene diradical or an arylylene diradical, with 6 to 30 carbon atoms;
- G denotes an alkylene diradical, a cycloalkylene diradical, an alkyl cycloalkylene diradical, an aromatic diradical or an alkylaromatic diradical with 1 to 40 carbon atoms which may have ether, thio or amine linkages in the main chain;
- * denotes a urethane or ureido linkage; and
- E denotes a polymerizable unsaturated organic radical represented by the general chemical formula:



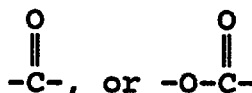
wherein

- R^1 denotes a divalent alkylene radical with 1 to 10 carbon atoms;
- R^2 denotes a -H or -CH₃ radical;
- R^3 denotes a -H radical or an alkyl radical with 1 to 6 carbon atoms or a



radical where

- Y is -O-, -S- or -NH- and R^4 denotes an alkyl radical with 1 to 12 carbon atoms;
- X denotes



- Ar denotes an aromatic radical with 6 to 30 carbon atoms;
- a is at least 1;
- w is 0 to 6;
- x is 0 or 1; and
- y is 0 or 1;

which process comprises:

- (a) reacting 2 mole equivalents of a diisocyanate of the formula:



wherein D is as defined above, and

$\bullet$ represents an isocyanate radical, with about 1 mole equivalent of a diol of the formula:



wherein G is as defined above, and

Δ represents a hydroxy radical, to form a product of the formula:



wherein * is as defined above,

(b) reacting the product of step (a) with a molar excess of a α, ω -diol endcapped long chain polymer of the formula:



wherein A and Δ are as defined above, to form a product of the formula:



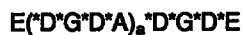
wherein a is as defined above; and

(c) reacting the product of step (b) with a molar excess of an endcapper of the formula:



wherein E and Δ are as defined above, to form the desired prepolymer.

2. A process for preparing a prepolymer material consisting of prepolymers represented by the general chemical formula:



where A, D, E, G and * are as defined in Claim 1, which process comprises:

(a) reacting 2 mole equivalents of a diisocyanate of the formula:



wherein D is as defined above and

$\bullet$ represents an isocyanate radical, with about 1 mole equivalent of a diol of the formula:



wherein A is as defined above and

Δ represents a hydroxy radical, to form a product of the formula:



wherein * is as defined above;

(b) reacting the product of step (a) with a molar excess of a diol of the formula:



wherein G and Δ are as defined above, to form a product of the formula:





wherein a is as defined above; and

(c) reacting the product of step (b) with a molar excess of an endcapper of the formula:

5 E Δ

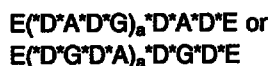
wherein E and Δ are as defined above, to form the desired prepolymer.

10 3. A process according to Claim 1 or Claim 2, wherein the polymeric radical, A, is selected from (1) a polyethylene glycol diradical with a molecular weight of at least about 2000; (2) a dialkyl endcapped polydimethylsiloxane radical with a molecular weight of at least 400; (3) dialkyl endcapped polydifluoroalkylsiloxane polymer with a molecular weight of at least 400; and (4) a fluoroalkyl ether with a molecular weight of at least about 400.

15 4. A process according to Claim 3, wherein said diisocyanate of the formula $\bullet D \bullet$ is selected from isophorone diisocyanate, toluene diisocyanate and 4,4'-dicyclohexyl methane diisocyanate.

20 5. A process according to Claim 3 or Claim 4, wherein said diol of the formula $\Delta G \Delta$ is selected from neopentyl glycol, 1,4-butanediol, diethylene glycol and triethylene glycol.

25 6. A process for making a copolymer, comprising preparing a prepolymer of the general chemical formulae:



wherein A, D, E, G, * and a are as defined in Claim 1, by a process according to any preceding claim, and thereafter polymerizing a mixture of said prepolymer with an ethylenically unsaturated comonomer.

30 7. A process according to Claim 6 wherein said polymeric radical, A, of said prepolymer is polyethylene radical with a molecular weight of at least about 2,000.

35 8. A process according to Claim 7, wherein said prepolymer comprises 15 to 95 weight percent of the comonomer mixture and said ethylenically unsaturated comonomer comprises 5 to 85 weight percent of said comonomer mixture.

9. A process according to Claim 8, wherein said polymerizable mixture further comprises a diethylenically reactive crosslinker.

40 10. A process according to Claim 9, wherein said ethylenically unsaturated comonomer is chosen from 2-hydroxyethyl methacrylate, N-vinyl-2-pyrrolidinone, methacrylic acid, dimethyl acrylamide and glycerol methacrylate.

45 11. A process according to Claim 6, wherein said polymeric radical, A, of said prepolymer is a dialkyl endcapped polydimethylsiloxane radical with a molecular weight of at least 400.

50 12. A process according to Claim 11, wherein said prepolymer comprises 15 to 95 weight percent of the comonomer mixture and said ethylenically unsaturated comonomer comprises 5 to 85 weight percent of said comonomer mixture.

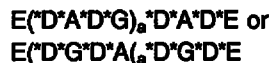
13. A process according to Claim 12, wherein said ethylenically unsaturated comonomer is chosen from 2-hydroxyethyl methacrylate, N-vinyl-2-pyrrolidinone, methacrylic acid, dimethyl acrylamide and glycerol methacrylate.

55 14. A process according to Claim 13, wherein said polymerizable mixture further comprises 5 to 40 weight percent organosilicon comonomer.



15. A process according to Claim 6, wherein said polymeric radical of said prepolymer is a poly-(fluoroalkyl)ether with a molecular weight of at least 400.

16. A process for making a copolymer, comprising preparing a prepolymer of the general chemical formulae:



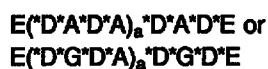
wherein A, D, E, G, * and a are as defined in Claim 1, by a process according to any one of Claims 1-5, and thereafter polymerizing a mixture of said prepolymer and an organosilicon comonomer.

17. A process according to Claim 16, wherein said organosilicon monomer is tris(trimethylsiloxy)silylpropyl methacrylate.

18. A process according to Claim 17, wherein there is made a copolymer which comprises 20-50 wt % of said prepolymer and 20-40 weight percent of tris(trimethylsiloxy)silyl propyl methacrylate.

19. A process according to Claim 18, wherein there is made a copolymer which further comprises 10-35 weight percent of one of the following hydrophilic comonomers: dimethacrylamide; glycerol methacrylate; 2-hydroxyethyl methacrylate; and N-vinyl-2-pyrrolidinone.

20. A process for making a copolymer, comprising preparing a prepolymer of the general chemical formulae:



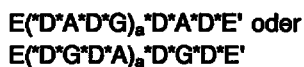
wherein A, D, E, G, * and a are as defined in Claim 1, by a process according to any one of Claims 1-5, and thereafter polymerizing an ethylenically unsaturated fluoroorgano comonomer.

21. A method of making a biomedical device, comprising fabricating said device from a polymer comprising a prepolymer prepared by a process according to any one of Claims 1-5.

Patentansprüche

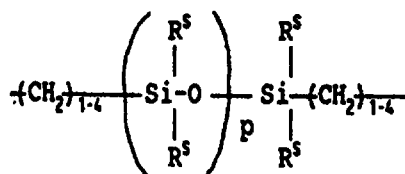
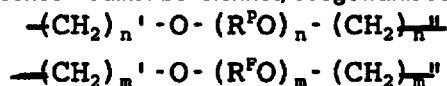
Patentansprüche für folgende Vertragsstaaten : DE, FR, GB, IT, SE

1. Ein Prepolymer-Material, bestehend aus Prepolymeren dargestellt durch die allgemeinen chemischen Formeln:



in der

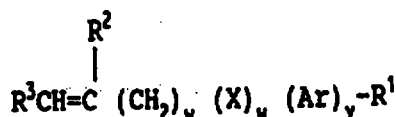
A ein bivalentes polymerisches Radikal bezeichnet, ausgewählt aus der Gruppe von



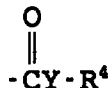
in der



- R^P** eine gerade oder verzweigte Alkylen-Gruppe mit 1 bis 6 Kohlenstoffatomen bezeichnet und n, n' und n'' dieselben oder verschiedene ganze Zahlen sind, so daß die Summe von n, n' und n'' für ein Mittelgewicht des besagten bivalenten polymerischen Radikals zwischen 2000 und 10000 sorgt;
- R^F** ein mit Fluor angereichertes gerades oder verzweigtes Alkylen-Radikal mit 1 bis 6 Kohlenstoffatomen bezeichnet und m, m' und m'' dieselben oder verschiedene ganze Zahlen sind, so daß die Summe von m, m' und m'' für ein Mittelgewicht des besagten bivalenten polymerischen Radikals zwischen 400 und 10000 sorgt;
- R^S** ein Alkyl-Radikal oder ein kurzkettiges, mit Fluor angereichertes Alkyl-Radikal mit 1 bis 3 Kohlenstoffatomen bezeichnet; und
- p** eine ganze Zahl ist, so daß das gesamte Mittelgewicht des besagten bivalenten polymerischen Radikals von 400 bis 10000 ist;
- D** ein Alkylen-Diradikal, ein Alkyl-Cycloalkylen-Diradikal, ein Cycloalkylen-Diradikal, ein Alkyl-Arylen-Diradikal oder ein Arylen-Diradikal mit 6 bis 30 Kohlenstoffatomen bezeichnet;
- G** ein Alkylen-Diradikal, ein Cycloalkylen-Diradikal, ein Alkyl-Cycloalkylen-Diradikal, ein aromatisches Diradikal oder ein alkylaromatisches Diradikal mit 1 bis 40 Kohlenstoffatomen bezeichnet, welche Äther-, Thio- oder Aminbindungen in der Hauptkette haben können;
- *** eine Urethan- oder Ureid-Bindung bezeichnet; und
- E und E'** polymerisierbare ungesättigte organische Radikale bezeichnen, dargestellt durch die allgemeine chemische Formel:



- in der
- R¹** ein bivalentes Alkylen-Radikal mit 1 bis 10 Kohlenstoffatomen bezeichnet;
- R²** ein -H oder -CH₃ Radikal bezeichnet;
- R³** ein -H Radikal oder ein Alkyl-Radikal mit 1 bis 6 Kohlenstoffatomen bezeichnet oder ein



- Radikal, in der
- X** Y -O-, -S- oder -NH- ist und R⁴ ein Alkyl-Radikal mit 1 bis 12 Kohlenstoffatomen bezeichnet;



- bezeichnet;
- Ar** ein aromatisches Radikal mit 6 bis 30 Kohlenstoffatomen bezeichnet;
- a** mindestens 1 ist,
- w** 0 bis 6 ist,
- x** 0 oder 1 ist, und
- y** 0 oder 1 ist.

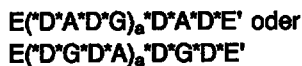
2. Das Prepolymer nach Anspruch 1, in dem das polymerische Radikal A ausgewählt wird aus (1) einem Polyäthylen-Glykol-Diradikal mit einem Molekulargewicht von mindestens ungefähr 2000; (2) einem Dialkyl-abgeschlossenen Polydimethylsiloxan-Radikal mit einem Molekulargewicht von mindestens 400; (3) Dialkyl-abgeschlossenen Polydifluoralkylsiloxan-Polymer mit einem Molekulargewicht von mindestens 400; und (4) einem Fluoralkyl-Äther mit einem Molekulargewicht von mindestens ungefähr 400.



3. Ein Prepolymer nach Anspruch 2, in dem das D-Radikal aus der Diisocyanat-Gruppe gebildet wird, die aus Isophoron-Diisocyanat, Toluol-Diisocyanat und 4,4'-Dicyclohexyl-Methan-Diisocyanat besteht.

4. Ein Prepolymer nach Anspruch 2 oder Anspruch 3, in dem das G-Radikal aus der Diol-Gruppe gebildet wird, die aus Neopentyl-Glykol, 1,4-Butandiol, Diäthylen-Glykol und Triäthylen-Glykol besteht.

5. Ein Copolymer-Material, gewonnen durch Polymerisation einer Comonomer-Mischung, bestehend aus:
(I) dem Prepolymer, beschrieben durch die allgemeinen chemischen Formeln:



worin A, D, E, E', G, * und a wie in Anspruch 1 definiert sind, und
(II) einem Äthylen-ungesättigten Comonomer.

6. Das Copolymer nach Anspruch 5, in dem das polymerische Radikal A der Komponente (I) ein Polyäthylen-Radikal mit einem Molekulargewicht von mindestens ungefähr 2000 ist.

7. Das Copolymer nach Anspruch 6, in dem die Komponente (I) 15 bis 95 Gewichtsprozent der Comonomer-Mischung enthält und das besagte Äthylen-ungesättigte Comonomer 5 bis 85 Gewichtsprozent der besagten Comonomer-Mischung enthält.

8. Das Copolymer nach Anspruch 7, das außerdem einen Diäthylen-reaktiven Vernetzer enthält.

9. Das Copolymer nach Anspruch 8, in dem das besagte Äthylen-ungesättigte Comonomer ausgewählt ist aus 2-Hydroxyäthyl-Methacrylat, N-vinyl-2-pyrrolidinon, Methacrylsäure, Dimethyl-Acrylamid und Glycerol-Methacrylat.

10. Das Copolymer nach Anspruch 5, in dem das polymerische Radikal A der Komponente (I) ein Dialkylabgeschlossenes Polydimethylsiloxan-Radikal mit einem Molekulargewicht von mindestens 400 ist.

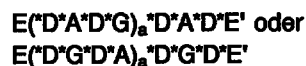
11. Das Copolymer nach Anspruch 10, in dem die Komponente (I) 15 bis 95 Gewichtsprozent der Comonomer-Mischung enthält und das besagte Äthylen-ungesättigte Comonomer 5 bis 85 Gewichtsprozent der besagten Comonomer-Mischung enthält.

12. Das Copolymer nach Anspruch 11, in dem das besagte Äthylen-ungesättigte Comonomer ausgewählt ist aus 2-Hydroxyäthyl-Methacrylat, N-vinyl-2-pyrrolidinon, Methacrylsäure, Dimethyl-Acrylamid und Glycerol-Methacrylat.

13. Das Copolymer nach Anspruch 12, das außerdem 5 bis 40 Gewichtsprozent eines Organosilikon-Comonomers enthält.

14. Das Copolymer nach Anspruch 5, in dem das polymerische Radikal A der Komponente (I) ein Poly-(Fluoralkyl)-Äther mit einem Molekulargewicht von mindestens 400 ist.

15. Ein Copolymer, gewonnen durch Polymerisation einer Comonomer-Mischung, bestehend aus:
(I) dem Prepolymer, beschrieben durch die allgemeinen chemischen Formeln:



worin A, D, E, E', G, * und a wie in Anspruch 1 definiert sind, und
(II) einem Organosilikon-Comonomer.

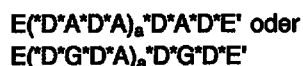
16. Das Copolymer nach Anspruch 15, in dem das besagte Organosilikon-Monomer ein Tris-(Trimethylsiloxy)-Silylpropyl-Methacrylat ist.



17. Das Copolymer nach Anspruch 16, das 20-50 Gewichtsprozent des Prepolymers (I) und 20-40 Gewichtsprozent des Tris(trimethylsiloxy)-silylpropyl-Methacrylats enthält.

18. Das Copolymer nach Anspruch 17, das außerdem 10-35 Gewichtsprozent eines der folgenden hydrophilen Comonomere enthält: Dimethacrylamid; Glycerol-Methacrylat; 2-Hydroxyäthyl-Methacrylat und N-vinyl-2-pyrrolidinon.

19. Ein Copolymer, gewonnen durch Polymerisation einer Comonomer-Mischung, bestehend aus:
(I) dem Prepolymer, beschrieben durch die allgemeinen chemischen Formeln:



worin A, D, E, E', G, * und a wie in Anspruch 1 definiert sind, und
(II) einem Äthylen-ungesättigten Fluororgano-Comonomers.

20. Eine biomedizinische Vorrichtung, die aus einem Polymer hergestellt wird, das das Prepolymer nach irgendeinem der vorhergehenden Ansprüche enthält.

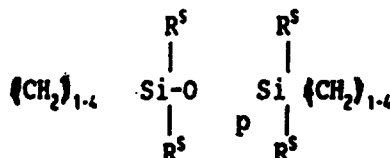
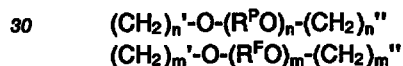
20 Patentansprüche für folgenden Vertragsstaat : ES

1. Ein Verfahren für die Zubereitung eines Prepolymer-Materials, bestehend aus Prepolymeren dargestellt durch die allgemeine chemische Formel:



in der

A ein bivalentes polymerisches Radikal bedeutet, ausgewählt aus der Gruppe von



in der

R^P eine gerade oder verzweigte Alkylen-Gruppe mit 1 bis 6 Kohlenstoffatomen bezeichnet und n, n' und n'' dieselben oder verschiedene ganze Zahlen sind, so daß die Summe von n, n' und n'' für ein Mittelgewicht des besagten bivalenten polymerischen Radikals zwischen 2000 und 10000 sorgt;

R^F ein mit Fluor angereichertes gerades oder verzweigtes Alkylen-Radikal mit 1 bis 6 Kohlenstoffatomen bezeichnet und m, m' und m'' dieselben oder verschiedene ganze Zahlen sind, so daß die Summe von m, m' und m'' für ein Mittelgewicht des besagten bivalenten polymerischen Radikals zwischen 400 und 10000 sorgt;

R^S ein Alkyl-Radikal oder ein kurzkettiges, mit Fluor angereichertes Alkyl-Radikal mit 1 bis 3 Kohlenstoffatomen bezeichnet; und

p eine ganze Zahl ist, so daß das gesamte Mittelgewicht des besagten bivalenten polymerischen Radikals von 400 bis 10000 ist;

D ein Alkylen-Diradikal, ein Alkyl-Cycloalkylen-Diradikal, ein Cycloalkylen-Diradikal, ein Alkyl-Arylen-Diradikal oder ein Arylen-Diradikal mit 6 bis 30 Kohlenstoffatomen bezeichnet;

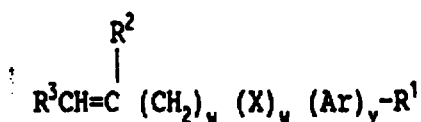
G ein Alkylen-Diradikal, ein Cycloalkylen-Diradikal, ein Alkyl-Cycloalkylen-Diradikal, ein aromatisches Diradikal oder ein alkylaromatisches Diradikal mit 1 bis 40 Kohlenstoffatomen bezeichnet, welche Äther-, Thio- oder Aminbindungen in der Hauptkette haben können;

* eine Urethan- oder Ureid-Bindung bezeichnet; und

E ein polymerisierbares ungesättigtes organisches Radikal bezeichnet, dargestellt durch die



allgemeine chemische Formel:

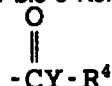


in der

R^1 ein bivalentes Alkyl-Radikal mit 1 bis 10 Kohlenstoffatomen bezeichnet;

R^2 ein -H oder -CH₃ Radikal bezeichnet;

R^3 ein -H Radikal oder ein Alkyl-Radikal mit 1 bis 6 Kohlenstoffatomen bezeichnet oder ein



Radikal, in der

Y -O-, -S- oder -NH- ist und R^4 ein Alkyl-Radikal mit 1 bis 12 Kohlenstoffatomen bezeichnet;

X



bezeichnet;

Ar ein aromatisches Radikal mit 6 bis 30 Kohlenstoffatomen bezeichnet;

a mindestens 1 ist,

w 0 bis 6 ist,

x 0 oder 1 ist, und

y 0 oder 1 ist,

und dieses Verfahren Folgendes umfaßt:

(a) Umsetzung von 2 Mol Äquivalenten eines Diisocyanats der Formel

●D●, in der D wie oben definiert ist und

● ein Isocyanat-Radikal darstellt mit ungefähr 1 Mol Äquivalent eines Diols der Formel:

▲G▲

in der G wie oben definiert ist und

▲ ein Hydroxy-Radikal darstellt, zur Bildung eines Produkts der Formel:

●D*G*D●

in der * wie oben definiert ist,

(b) Umsetzung vom Produkt gemäß Schritt (a) mit einem molaren Überschuß eines langkettigen Polymers mit α,ω -Diol-Endabschluß gemäß der Formel:

▲A▲

in der A und ▲ wie oben definiert sind, zur Bildung eines Produkts der Formel:

●(D*G*D*A)_a*D*G*D●

in der a wie oben definiert ist; und

(c) Umsetzung vom Produkt gemäß Schritt (b) mit einem molaren Überschuß eines Abschlusses gemäß der Formel:

E▲

in der E und ▲ wie oben definiert sind, um das gewünschte Prepolymer zu bilden.



2. Ein Verfahren für die Zubereitung eines Prepolymer-Materials, bestehend aus Prepolymeren dargestellt durch die allgemeine chemische Formel:



in der A, D, E, G und * wie in Anspruch 1 definiert sind; welches Verfahren Folgendes umfaßt:

(a) Umsetzung von 2 Mol Äquivalenten eines Diisocyanats der Formel

●D●, in der D wie oben definiert ist und

● ein Isocyanat-Radikal darstellt mit ungefähr 1 Mol Äquivalent eines Diols der Formel:



in der A wie oben definiert ist und

Δ ein Hydroxy-Radikal darstellt,

zur Bildung eines Produkts der Formel:



in der * wie oben definiert ist,

(b) Umsetzung vom Produkt gemäß Schritt (a) mit einem molaren Überschuß eines Diols der Formel



in der G und Δ wie oben definiert sind, zur Bildung eines Produkts der Formel:



in der a wie oben definiert ist; und

(c) Umsetzung vom Produkt gemäß Schritt (b) mit einem molaren Überschuß eines Abschlusses gemäß der Formel:



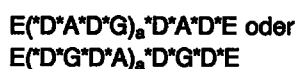
in der E und Δ wie oben definiert sind, um das gewünschte Prepolymer zu bilden.

3. Ein Verfahren nach Anspruch 1 oder Anspruch 2, in dem das polymerische Radikal A ausgewählt wird aus (1) einem Polyäthylen-Glykol-Diradikal mit einem Molekulargewicht von mindestens ungefähr 2000; (2) einem Dialkyl-abgeschlossenen Polydimethylsiloxan-Radikal mit einem Molekulargewicht von mindestens 400; (3) Dialkylabgeschlossenen Polydifluoralkylsiloxan-Polymer mit einem Molekulargewicht von mindestens 400; und (4) einem Fluoralkyl-Äther mit einem Molekulargewicht von mindestens ungefähr 400.

4. Ein Verfahren nach Anspruch 3, in dem das besagte Diisocyanat der Formel ●D● ausgewählt ist aus Isophoron-Diisocyanat, Toluol-Diisocyanat und 4,4'-Dicyclohexyl-Methan-Diisocyanat.

5. Ein Verfahren nach Anspruch 3 oder Anspruch 4, in dem das besagte Diol der Formel ΔGΔ ausgewählt ist aus Neopentyl-Glykol, 1,4-Butandiol, Diäthylen-Glykol und Triäthylen-Glykol.

6. Ein Verfahren zur Herstellung eines Copolymers, umfassend die Zubereitung eines Prepolymers der allgemeinen chemischen Formeln:



worin A, D, E, G, * und a wie in Anspruch 1 definiert sind, durch ein Verfahren entsprechend irgendeinem der vorhergehenden Ansprüche und nachfolgender Polymerisation einer Mischung des besagten Prepolymers mit einem Äthylenungesättigten Comonomer.



7. Ein Verfahren nach Anspruch 6, in dem das besagte polymerische Radikal A des besagten Prepolymers ein Polyäthylen-Radikal mit einem Molekulargewicht von mindestens ungefähr 2000 ist.
8. Ein Verfahren nach Anspruch 7, in dem das besagte Prepolymer 15 bis 95 Gewichtsprozent der Comonomer-Mischung enthält und das besagte Äthylen-ungesättigte Comonomer 5 bis 85 Gewichtsprozent der besagten Comonomer-Mischung enthält.
9. Ein Verfahren nach Anspruch 8, in dem die besagte polymerisierbare Mischung außerdem einen Diäthylenreaktiven Vernetzer enthält.
10. Ein Verfahren nach Anspruch 9, in dem das besagte Äthylen-ungesättigte Comonomer ausgewählt ist aus 2-Hydroxyäthyl-Methacrylat, N-vinyl-2-pyrrolidinon, Methacrylsäure, Dimethyl-Acrylamid und Glycerol-Methacrylat.
11. Ein Verfahren nach Anspruch 6, in dem das besagte polymerische Radikal A des besagten Prepolymers ein Dialkyl-abgeschlossenen Polydimethylsiloxan-Radikal mit einem Molekulargewicht von mindestens 400 ist.
12. Ein Verfahren nach Anspruch 11, in dem das besagte Prepolymer 15 bis 95 Gewichtsprozent der Comonomer-Mischung enthält und das besagte Äthylen-ungesättigte Comonomer 5 bis 85 Gewichtsprozent der besagten Comonomer-Mischung enthält.
13. Ein Verfahren nach Anspruch 12, in dem das besagte Äthylen-ungesättigte Comonomer ausgewählt ist aus 2-Hydroxyäthyl-Methacrylat, N-vinyl-2-pyrrolidinon, Methacrylsäure, Dimethyl-Acrylamid und Glycerol-Methacrylat.
14. Ein Verfahren nach Anspruch 13, in dem die besagte polymerisierbare Mischung außerdem 5 bis 40 Gewichtsprozent eines Organosilikon-Comonomers enthält.
15. Ein Verfahren nach Anspruch 6, in dem das besagte polymerische Radikal des besagten Prepolymers ein Poly(Fluoralkyl)-Äther mit einem Molekulargewicht von mindestens 400 ist.
16. Ein Verfahren zur Herstellung eines Copolymers, umfassend die Zubereitung eines Prepolymers der allgemeinen chemischen Formeln:

$$E(D^*A^*D^*G)_a^*D^*A^*D^*E \text{ oder } E(D^*G^*D^*A)_a^*D^*G^*D^*E$$

worin A, D, E, G, * und a wie in Anspruch 1 definiert sind, durch ein Verfahren entsprechend irgendeinem der vorhergehenden Ansprüche 1-5 und nachfolgender Polymerisation einer Mischung des besagten Prepolymers und eines Organosilikon-Comonomers.
17. Ein Verfahren nach Anspruch 16, in dem das besagte Organosilikon-Monomer ein Tris(Trimethylsiloxy)-Silylpropyl-Methacrylat ist.
18. Ein Verfahren nach Anspruch 17, in dem ein Copolymer hergestellt wird, das 20-50 Gewichtsprozent des besagten Prepolymers und 20-40 Gewichtsprozent des Tris(Trimethylsiloxy)-Silylpropyl-Methacrylats enthält.
19. Ein Verfahren nach Anspruch 18, in dem ein Copolymer hergestellt wird, das außerdem 10-35 Gewichtsprozent eines der folgenden hydrophilen Comonomere enthält: Dimethacrylamid; Glycerol-Methacrylat; 2-Hydroxyäthyl-Methacrylat und N-vinyl-2-pyrrolidinon.
20. Ein Verfahren zur Herstellung eines Copolymers, umfassend die Zubereitung eines Prepolymers der allgemeinen chemischen Formeln:

$$E(D^*A^*D^*A)_a^*D^*A^*D^*E \text{ oder } E(D^*G^*D^*A)_a^*D^*G^*D^*E$$



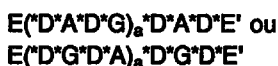
worin A, D, E, G, * und a wie in Anspruch 1 definiert sind, durch ein Verfahren entsprechend irgendeinem der Ansprüche 1-5 und nachfolgender Polymerisation eines Äthylen-ungesättigten Fluororgan-Comonomers.

21. Eine Methode zur Herstellung einer biomedizinischen Vorrichtung, die die Fabrikation der besagten Vorrichtung aus einem Polymer umfaßt, das ein Prepolymer, welches durch ein Verfahren nach irgendeinem der Ansprüche 1-5 zubereitet wurde, enthält.

Revendications

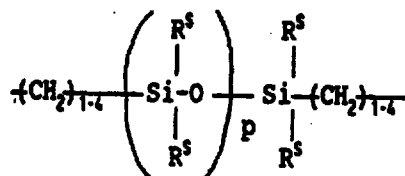
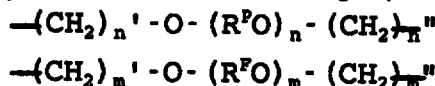
Revendications pour les Etats contractants suivants : DE, FR, GB, IT, SE

1. Un matériel prépolymère composé de prépolymères représentés par les formules chimiques générales suivantes



où

A représente un radical polymère bivalent choisi dans le groupe



où

R^p représente un groupe alkylène linéaire ou ramifié avec 1 à 6 atomes de carbone, et n , n' et n'' sont des nombres entiers identiques ou différents tels que de la somme de n , n' et n'' résulte une moitié en poids dudit radical polymère bivalent laquelle est comprise entre 2.000 et 10.000;

R^f représente un radical alkylène fluoré linéaire ou ramifié avec 1 à 6 atomes de carbone, et m , m' et m'' sont des nombres entiers identiques ou différents tels que de la somme de m , m' et m'' résulte une moitié en poids dudit radical polymère bivalent laquelle est comprise entre 400 et 10.000;

R^s représente un radical alkyl ou un radical alkyl fluoré de courte chaîne avec 1 à 3 atomes de carbone; et

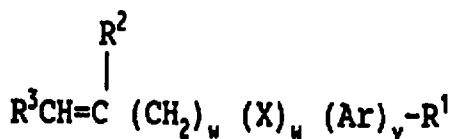
p est un nombre entier tel que la moitié en poids total dudit radical polymère bivalent est de 400 à 10.000;

D représente un biradical alkylène, un biradical alkyl cycloalkylène, un biradical cycloalkylène, un biradical alkylarylène ou un biradical arylène avec 6 à 30 atomes de carbone;

G représente un biradical alkylène, un biradical cycloalkylène, un biradical alkyl cycloalkylène, un biradical aromatique ou un biradical alkylaromatique avec 1 à 40 atomes de carbone qui peuvent avoir des liaisons éther, thio ou amine dans la chaîne principale;

* représente une liaison uréthane ou ureido; et

E et E' représentent des radicaux organiques insaturés polymérisables représentés par la formule chimique générale:



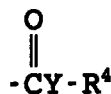


où

R¹ représente un radical alkylène bivalent avec 1 à 10 atomes de carbone;

R² représente un radical -H ou -CH₃;

R³ représente un radical -H ou un radical alkyl avec 1 à 6 atomes de carbone ou un



radical où Y est -O-, -S- ou -NH- et R⁴ représente un radical alkyl avec 1 à 12 atomes de carbone;

X représente



Ar représente un radical aromatique avec 6 à 30 atomes de carbone;

a est au moins égal à 1;

w est de 0 à 6;

x est égal à 0 ou 1; et

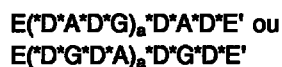
y est égal à 0 ou 1.

2. Le prépolymère de la revendication 1 dans lequel le radical polymère, A, est choisi entre (1) un biradical polyéthylène glycol d'un poids moléculaire d'au moins égal à environ 2.000; (2) un radical polydiméthylsiloxane à terminaison dialkyl d'un poids moléculaire d'au moins 400; (3) polymère polydifluoroalkylsiloxane à terminaison dialkyl d'un poids moléculaire d'au moins 400; et (4) un éther fluoroalkyl d'un poids moléculaire minimal d'environ 400.

3. Un prépolymère selon la revendication 2 où le radical D est formé à partir du groupe diisocyanates comprenant le diisocyanate isophorone, le diisocyanate toluène et le diisocyanate 4,4'-dicyclohexyl méthane.

4. Un prépolymère selon la revendication 2 ou la revendication 3, où le radical G est formé à partir du groupe des diols comprenant le néopentyl glycol, le 1,4-butanediol, le diéthylène glycol et le triéthylène glycol.

5. Un matériel copolymère formé par polymérisation d'un mélange de comonomères composé de:
(I) le prépolymère décrit dans la formule chimique générale:



où A, D, E, E', G, * et a sont tels que définis dans la revendication 1 et

(II) un comonomère éthyléniquement insaturé.

6. Le copolymère selon la revendication 5 où le radical polymère du composant (I), A, est un radical polyéthylène d'un poids moléculaire au moins égal à environ 2.000.

7. Le copolymère de la revendication 6 où le composant (I) comprend 15 à 95 pour cent en poids du mélange de comonomères et où le comonomère éthyléniquement insaturé comprend 5 à 85 pour cent en poids dudit mélange de comonomères.

8. Le copolymère de la revendication 7 comprenant aussi un agent de réticulation diéthyléniquement réactif.

9. Le copolymère de la revendication 8 où ledit comonomère éthyléniquement insaturé est choisi entre le 2-hydroxyéthyl méthacrylate, le N-vinyle-2-pyrrolidinone, l'acide méthacrylique, le diméthyl acrylamide



et le glycérol méthacrylate.

- 5
10. Le copolymère de la revendication 5 où le radical polymère du composant (I), A, est un radical polydiméthylsiloxane à terminaison dialkyl d'un poids moléculaire d'au moins 400.
11. Le copolymère de la revendication 10 où le composant (I) comprend 15 à 95 pour cent en poids du mélange de comonomères et où le comonomère éthyléniquement insaturé comprend 5 à 85 pour cent en poids dudit mélange de comonomères.
- 10 12. Le copolymère de la revendication 11 où ledit comonomère éthyléniquement insaturé est choisi entre le 2-hydroxyéthyl méthacrylate, le N-vinyle-2-pyrrolidinone, l'acide méthacrylique, le diméthyl acrylamide et le glycérol méthacrylate.
13. Le copolymère de la revendication 12 comprenant aussi 5 à 40 pour cent en poids de comonomère organosilicone.
- 15 14. Le copolymère de la revendication 5 où le radical polymère du composant (I), A, est un poly-(fluoroalkyl)éther d'un poids moléculaire d'au moins 400.
- 20 15. Un copolymère formé par polymérisation d'un mélange de comonomères comprenant:
(I) le prépolymère décrit par la formule chimique générale :
- $E(D^*A^*D^*G)_aD^*A^*D^*E'$ ou
 $E(D^*G^*D^*A)_aD^*G^*D^*E'$
- 25 où A, D, E, E', G, * et a sont tels que définis dans la revendication 1 et
(II) un comonomère organosilicone.
16. Le copolymère de la revendication 15' où le monomère organosilicone est un tris(triméthylsiloxyl)silylpropyl méthacrylate.
- 30 17. Le copolymère de la revendication 16 comprenant 20 à 50 pour cent en poids de prépolymère (I) et 20 à 40 pour cent en poids de tris(triméthylsiloxyl)silylpropyl méthacrylate.
- 35 18. Le copolymère de la revendication 17 comprenant aussi 10 à 35 pour cent en poids de l'un des comonomères hydrophiles suivants : diméthacrylamide; glycérol méthacrylate; 2-hydroxyéthyl méthacrylate; et N-vinyle-2-pyrrolidinone.
19. Un copolymère formé par polymérisation d'un mélange de comonomères comprenant:
(I) le prépolymère décrit par la formule chimique générale :
- $E(D^*A^*D^*A)_aD^*A^*D^*E'$ ou
 $E(D^*G^*D^*A)_aD^*G^*D^*E'$
- 45 où A, D, E, E', G, * et a sont tels que définis dans la revendication 1 et
(II) un comonomère fluororgano éthyléniquement insaturé.
20. Un dispositif biomédical fait d'un polymère comprenant le prépolymère de l'une des revendications précédentes.
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Revendications pour l'Etat contractant suivant : ES

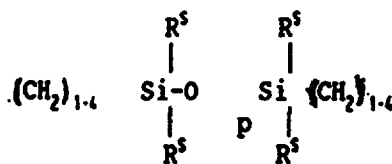
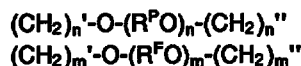
1. Un procédé pour la préparation d'un matériel prépolymère composé de prépolymères représentés par la formule chimique générale suivante



où



A représente un radical polymère bivalent choisi dans le groupe



où

R^{P} représente un groupe alkylène linéaire ou ramifié avec 1 à 6 atomes de carbone, et n , n' et n'' sont des nombres entiers identiques ou différents tels que de la somme de n , n' et n'' résulte une moitié en poids dudit radical polymère bivalent laquelle est comprise entre 2.000 et 10.000;

R^{F} représente un radical alkylène fluoré linéaire ou ramifié avec 1 à 6 atomes de carbone, et m , m' et m'' sont des nombres entiers identiques ou différents tels que de la somme de m , m' et m'' résulte une moitié en poids dudit radical polymère bivalent laquelle est comprise entre 400 et 10.000;

R^{S} représente un radical alkyl ou un radical alkyl fluoré de courte chaîne avec 1 à 3 atomes de carbone; et

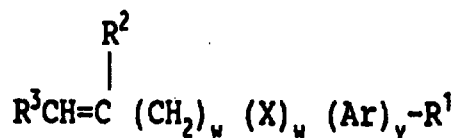
p est un nombre entier tel que la moitié en poids total dudit radical polymère bivalent est de 400 à 10.000;

D représente un biradical alkylène, un biradical alkyl cycloalkylène, un biradical cycloalkylène, un biradical alkylarylène ou un biradical arylène, avec 6 à 30 atomes de carbone;

G représente un biradical alkylène, un biradical cycloalkylène, un biradical alkyl cycloalkylène, un biradical aromatique ou un biradical alkylaromatique avec 1 à 40 atomes de carbone qui peuvent avoir des liaisons éther, thio ou amine dans la chaîne principale;

$*$ représente une liaison uréthane ou ureido; et

E représente un radical organique insaturé polymérisable représenté par la formule chimique générale:

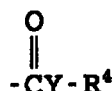


où

R^1 représente un radical alkylène bivalent avec 1 à 10 atomes de carbone;

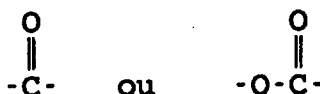
R^2 représente un radical -H ou -CH₃;

R^3 représente un radical -H ou un radical alkyl avec 1 à 6 atomes de carbone ou un



radical où Y est -O-, -S- ou -NH- et R^4 représente un radical alkyl avec 1 à 12 atomes de carbone;

X représente



Ar représente un radical aromatique avec 6 à 30 atomes de carbone;



a est au moins égal à 1;

w est de 0 à 6;

x est égal à 0 ou 1; et

y est égal à 0 ou 1;

5 lequel procédé comprend:

(a) la réaction de 2 équivalents de mole d'un diisocyanate de formule :



10 où D est tel que défini ci-dessus, et

● représente un radical isocyanate avec environ un équivalent de mole d'un diol de formule :



15 où G est tel que défini ci-dessus, et

▲ représente un radical hydroxy, pour former un produit de formule :



20 où * est tel que défini ci-dessus

(b) la réaction du produit de l'étape (a) avec un polymère de chaîne longue et avec terminaison α, ω -diol en excédant molaire de formule:



25

où A et Δ sont tels que définis ci-dessus, pour former un produit de formule :



30 Où a est tel que défini ci-dessus; et

(c) la réaction du produit de l'étape (b) avec un excès moléculaire d'un terminateur de la formule:



35

où E et Δ sont tels que définis ci-dessus, pour former le prépolymère désiré.

2. Un procédé pour la préparation d'un matériel prépolymère composé de prépolymères représentés par la formule chimique générale suivante

40



où A, D, E, G et * sont tels que définis dans la revendication 1, lequel procédé comprend:

(a) la réaction de 2 équivalents de mole d'un diisocyanate de formule :

45



où D est tel que défini ci-dessus, et

● représente un radical isocyanate avec environ 1 équivalent de mole d'un diol de formule :

50



où A est tel que défini ci-dessus et

▲ représente un radical hydroxy
pour former un produit de formule :

55



où * est tel que défini ci-dessus



(b) la réaction du produit de l'étape (a) avec un diol en excédant molaire de formule:



où G et Δ sont tels que définis ci-dessus, pour former un produit de formule :



Où a est tel que défini ci-dessus; et

(c) la réaction du produit de l'étape (b) avec une terminaison en excédant moléculaire de formule :



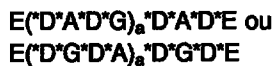
où E et Δ sont tels que définis ci-dessus, pour former le prépolymère désiré.

3. Un procédé selon la revendication 1 ou la revendication 2 dans lequel le radical polymère, A, est choisi entre (1) un biradical polyéthylène glycol d'un poids moléculaire d'au moins environ 2.000; (2) un radical polydiméthylsiloxane à terminaison dialkyl d'un poids moléculaire d'au moins 400; (3) polymère polydifluoroalkylsiloxane à terminaison dialkyl d'un poids moléculaire d'au moins 400; et (4) un éther fluoroalkyl d'un poids moléculaire minimal d'environ 400.

4. Un procédé selon la revendication 3 où le diisocyanate de la formule $\bullet D \bullet$ est choisi entre le diisocyanate isophorone, le diisocyanate toluène et le diisocyanate 4,4'-dicyclohexyl méthane.

5. Un procédé selon la revendication 3 ou la revendication 4, où le diol de la formule $\Delta G \Delta$ est choisi entre le néopentyl glycol, le 1,4-butanediol, le diéthylène glycol et le triéthylène glycol.

6. Un procédé pour la préparation d'un copolymère comprenant la préparation d'un prépolymère des formules chimiques générales:



où A, D, E, G, * et a sont tels que définis dans la revendication 1 selon un procédé conforme à l'une des revendications précédentes et polymérisation ultérieure du mélange dudit prépolymère avec un comonomère éthyléniquement insaturé.

7. Un procédé selon la revendication 6 où le radical polymère, A, dudit prépolymère est un radical polyéthylène d'un poids moléculaire au moins égal à environ 2.000.

8. Un procédé selon la revendication 7 où ledit prépolymère comprend 15 à 95 pour cent en poids du mélange de comonomères et où le comonomère éthyléniquement insaturé comprend 5 à 85 pour cent en poids dudit mélange de comonomères.

9. Un procédé selon la revendication 8 dans lequel le mélange polymérisable aussi-même comprend un agent de réticulation diéthyléniquement réactif.

10. Un procédé selon la revendication 9 où ledit comonomère éthyléniquement insaturé est choisi entre le 2-hydroxyéthyl méthacrylate, le N-vinyle-2-pyrrolidinone, l'acide méthacrylique, le diméthyl acrylamide et le glycérol méthacrylate.

11. Un procédé selon la revendication 6 où le radical polymère, A, dudit prépolymère est un radical polydiméthylsiloxane à terminaison dialkyl d'un poids moléculaire d'au moins 400.

12. Un procédé selon la revendication 11 où ledit prépolymère comprend 15 à 95 pour cent en poids du mélange de comonomères et où le comonomère éthyléniquement insaturé comprend 5 à 85 pour cent en poids dudit mélange de comonomères.



13. Un procédé selon la revendication 12 où ledit comonomère éthyléniquement insaturé est choisi entre le 2-hydroxyéthyl méthacrylate, le N-vinyle-2-pyrrolidinone, l'acide méthacrylique, le diméthyl acrylamide et le glycérol méthacrylate.
- 5 14. Un procédé selon la revendication 13 dans lequel le mélange polymérisable comprend aussi 5 à 40 pour cent en poids de comonomère organosilicone.
15. Un procédé selon la revendication 6 où le radical polymère dudit prépolymère est un poly(fluoroalkyl)-éther d'un poids moléculaire d'au moins 400.
- 10 16. Un procédé pour la préparation d'un copolymère comprenant la préparation d'un prépolymère des formules chimiques générales:
- E(*D*A*D*G)_a*D*A*D*E ou
E(*D*G*D*A)_a*D*G*D*E
- 15 où A, D, E, G, * et a sont tels que définis dans la revendication 1, selon un procédé conforme à l'une des revendications 1 à 5 et polymérisation ultérieure d'un mélange dudit prépolymère avec un comonomère organosilicone.
- 20 17. Un procédé selon la revendication 16 où le monomère organosilicone est un tris(triméthylsiloxy)-silylpropyl méthacrylate.
- 25 18. Un procédé selon la revendication 17 où est produit un copolymère comprenant 20 à 50 pour cent en poids dudit prépolymère et 20 à 40 pour cent en poids de tris(triméthylsiloxy)silylpropyl méthacrylate.
- 30 19. Un procédé selon la revendication 18 où est produit un copolymère comprenant aussi 10 à 35 pour cent en poids de l'un des comonomères hydrophiles suivants : diméthacrylamide; glycérol méthacrylate; 2-hydroxyéthyl méthacrylate; et N-vinyle-2-pyrrolidinone.
- 35 20. Un procédé pour la préparation d'un copolymère comprenant la préparation d'un prépolymère des formules chimiques générales:
- E(*D*A*D*A)_a*D*A*D*E ou
E(*D*G*D*A)_a*D*G*D*E
- 40 où A, D, E, G, * et a sont tels que définis dans la revendication 1, selon un procédé conforme à l'une des revendications 1 à 5 et polymérisation ultérieure d'un comonomère fluoroorgano éthyléniquement insaturé.
- 45 21. Une méthode de fabrication d'un dispositif biomédical comprenant la fabrication dudit dispositif à partir d'un polymère comprenant un prépolymère préparé selon un procédé conforme à l'une des revendications 1 à 5.
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- 55