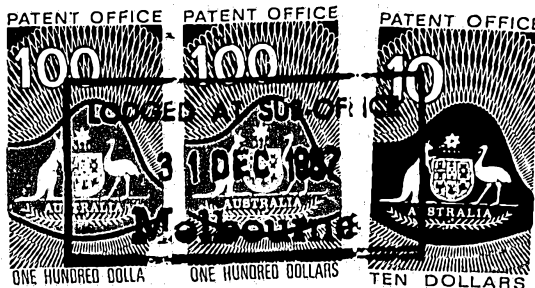


(CONVENTION. By one or more persons and

COMMONWEALTH

Patents



CONVENTION APPLICATION FOR A PATENT

605533

(1) Here
insert (in
full) Name
or Names of
Applicant or
Applicants,
followed by
Address (es).

K (1) DOW CORNING CORPORATION

We of Midland,

State of Michigan,

United States of America

(2) Here
insert Title
of Invention.

hereby apply for the grant of a Patent for an invention entitled: (2) HAIR CARE

COMPOSITION CONTAINING MODIFIED AMINOALKYL

SUBSTITUTED POLYORGANOSILOXANE

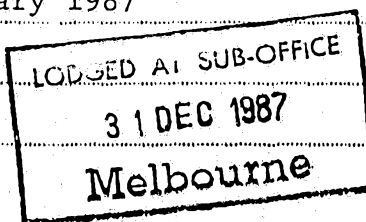
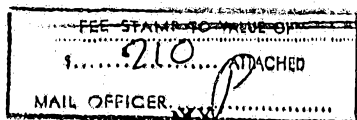
(3) Here insert
number(s)
of basic
application(s)

which is described in the accompanying complete specification. This application is a
Convention application and is based on the application numbered (3)

294

(4) Here insert
Name of basic
Country or
Countries, and
basic date or
dates

for a patent or similar protection made in (4) UNITED STATES OF AMERICA
on the 2nd day of January 1987



Our address for service is Messrs. Edwd. Waters & Sons, Patent Attorneys,

50 Queen Street, Melbourne, Victoria, Australia.

APPLICATION ACCEPTED AND AMENDMENTS

ALLOWED 17.10.90 DATED this 30th day of December 1987

(5) Signa-
ture (s) of
Applicant (s)
or
Seal of
Company and
Signatures of
its Officers as
prescribed by
its Articles of
Association.

(5)

DOW CORNING CORPORATION

IAN A. SCOTT
Registered Patent Attorney

To:

(CONVENTION. Company.)

Form 8

COMMONWEALTH OF AUSTRALIA

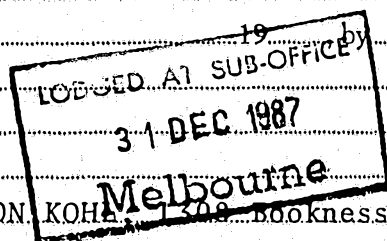
Patents Act 1952-1969

DECLARATION IN SUPPORT OF A CONVENTION
APPLICATION FOR A PATENT OR PATENT OF ADDITION(1) Here
insert (in
full) Name of
Company.In support of the Convention Application made by⁽¹⁾
DOW CORNING CORPORATION of Midland, State of Michigan,
United States of America

(hereinafter referred to as the applicant) for a Patent

(2) Here
insert title
of Invention.for an invention entitled:⁽²⁾
HAIR CARE COMPOSITION CONTAINING MODIFIED AMINOALKYL
SUBSTITUTED POLYDIORGANOSILOXANE(3) Here
insert full Name
and Address,
of Company
official
authorized
to make
declaration.I, ⁽³⁾ NORMAN EDWARD LEWIS, General Patent Counsel & Assistant
of DOW CORNING CORPORATION of Midland, Secretary
State of Michigan, United States of America

do solemnly and sincerely declare as follows:

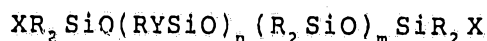
1. I am authorised by the applicant for the patent
to make this declaration on its behalf.(4) Here
insert basic
Country or
Countries
followed by
date or dates
and basic
Applicant or
Applicants.2. The basic application as defined by Section 141 of the Act was
made in ⁽⁴⁾ U.S.A., Serial No. 294
on the 2nd day of January 19 87, by
GRETCHEN SUTTON KOHL and THOMAS HOWARD LANE
on the day of 19 87 by(5) Here
insert (in
full) Name
and Address
of Actual
Inventor or
Inventors.3. ⁽⁵⁾ GRETCHEN SUTTON KOHL, 1308 Brookness, Midland,
Michigan, U.S.A. and THOMAS HOWARD LANE, 107 Varner Court,
Midland, Michigan, U.S.A.is/are the actual inventors of the invention and the facts upon which the applicant
is entitled to make the application are as follow:The applicant is the assignee of
GRETCHEN SUTTON KOHL and THOMAS HOWARD LANE4. The basic application referred to in paragraph 2 of this Declaration
was the first application made in a Convention country in
respect of the invention the subject of the application.DECLARED at Midland, Michigan, U.S.A.
this 9th day of December 19 87

(12) PATENT ABRIDGMENT (11) Document No. AU-B-83181/87
(19) AUSTRALIAN PATENT OFFICE (10) Acceptance No. 605533

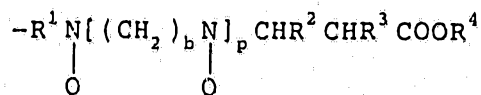
- (54) Title
HAIR CARE COMPOSITION MODIFIED AMINOALKYL SUBSTITUTED
POLYDIORGANOSILOXANE
- International Patent Classification(s)
(51)⁴ A61K 007/08 A61K 007/075
- (21) Application No. : 83181/87 (22) Application Date : 31.12.87
- (30) Priority Data
- (31) Number (32) Date (33) Country
000294 02.01.87 US UNITED STATES OF AMERICA
- (43) Publication Date : 07.07.88
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- (71) Applicant(s)
DOW CORNING CORPORATION
- (72) Inventor(s)
GRETCHEN SUTTON KOHL; THOMAS HOWARD LANE
- (74) Attorney or Agent
WATERMARK PATENT & TRADEMARK ATTORNEYS, Locked Bag 5, HAWTHORN VIC
3122
- (56) Prior Art Documents
FR 2548019
EP 113992
EP 174092

(57) Claim

1. The method of treating or conditioning hair which comprises contacting hair with a polydiorganosiloxane represented by the formula



wherein R denotes a monovalent hydrocarbon radical having 1 to 6 carbon atoms; X denotes -OH, -OR, Y, or R; Y denotes a substituent of the formula



wherein R¹ denotes an alkylene radical having 3 to 6 carbon atoms, R² denotes a hydrogen atom, phenyl, or an alkyl radical of 1 to 6 carbon atoms, R³ denotes a hydrogen atom or methyl radical and R⁴ denotes an alkyl radical having 1 to 6 carbon atoms, b has a value of 2, 3, or 4, p has a value of 0, 1, or 2, and Q denotes a hydrogen atom or a radical of the formula -CHR²CHR³COOR⁴ wherein R², R³ and R⁴ have the same meanings as defined above; m has an average value from 10 to 600; n has an average value of 0 to 100 with the proviso that when n is 0, X denotes Y.

605533
Form 10

COMMONWEALTH OF AUSTRALIA

PATENTS ACT 1952-69

COMPLETE SPECIFICATION

(ORIGINAL)

Class

Int. Class

Application Number:

Lodged:

Complete Specification Lodged:

Accepted:

Published:

Priority:

Related Art:

This document contains the
amendments made under
Section 49 and is correct for
printing.

Name of Applicant: DOW CORNING CORPORATION

Address of Applicant: Midland,
State of Michigan,
United States of America

Actual Inventor: Gretchen Sutton Kohl and Thomas Howard Lane

Address for Service: EDWD. WATERS & SONS,
50 QUEEN STREET, MELBOURNE, AUSTRALIA, 3000.

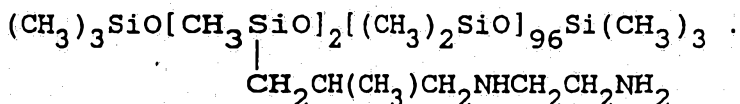
Complete Specification for the invention entitled: HAIR CARE COMPOSITION MODIFIED AMINOALKYL
SUBSTITUTED POLYDIORGANOSILOXANE

The following statement is a full description of this invention, including the best method of performing it known to : US

HAIR CARE COMPOSITION CONTAINING MODIFIED AMINOALKYL
SUBSTITUTED POLYDIORGANOSILOXANE

This invention relates to the use of modified amino-alkyl substituted polydiorganosiloxane in hair care compositions such as conditioners and shampoos. More specifically, the invention relates to compositions and methods for treating hair with polydiorganosiloxane containing amino-alkyl groups modified by alkoxycarbonylalkyl substituents.

Aminoalkyl substituted polydiorganosiloxanes are well known copolymers which are typically composed of dimethylsiloxane units and aminoalkylmethylsiloxane units. For example, Starch, U.S. Patent No. 4,563,347, January 7, 1986, describes a copolymer represented by the formula



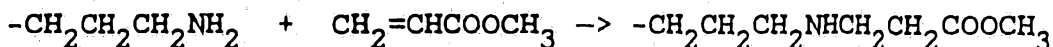
Such copolymers are further described in the CTFA Cosmetic Ingredient Dictionary, 3rd edition, 1982, published by The Cosmetic, Toiletry and Fragrance Association, Inc., 1133 Fifteenth Street NW Washington D.C. (USA) under the nomenclature amodimethicone and trimethylsilylamodimethicone.

The use of aminoalkyl substituted polydiorganosiloxane in hair care compositions is also well known. For instance, Starch, U.S. Patent No. 4,563,347, teaches using these copolymers in hair conditioning compositions to improve combing and feel characteristics. Similarly, Grollier et al., UK Patent Application Nos. 2,157,168 and 2,143,434, October 23, 1985 and February 13, 1985 respectively, teach using the aminoalkyl substituted polydiorganosiloxanes with other components in hair care compositions which may be formulated as a shampoo, a rinsed or non-rinsed lotion, a restructuring

composition, a composition for blow drying, or a composition for permanent waving.

The aminoalkyl substituents are credited with providing the copolymers with cationic sites that make the polymer more substantive to hair than nonsubstituted polydimethylsiloxane. However, the chemical reactivity of the aminoalkyl groups may also present a problem in regard to compatibility with other common components of hair care formulations. Compatibility problems may be observed especially with anionic surfactants. Tangney et al., U.S. Patent No. 4,507,455, March 26, 1985, describe a method of modifying the chemical reactivity of aminoalkyl substituents of polydiorganosiloxane by reacting the polymer with a monocarboxylic acid anhydride to acylate the amine groups and thus convert them to amides. While acylation was effective to improve compatibility, it has also been found to reduce the substantivity of the polymer to hair so that the utility of the polymers in hair care compositions is limited. It is a purpose of the present invention to provide hair care compositions containing a conditioning silicone polymer which has the advantage of being highly substantive to hair but at the same time has a more moderate chemical reactivity with correspondingly reduced compatibility problems.

Aminoalkyl substituted silanes and polydiorganosiloxanes have been reacted with acrylate esters to provide silanes and polydiorganosiloxanes in which the amine is modified by addition of an alkoxycarbonylalkyl group to the nitrogen atom. For example, the reaction is illustrated as follows for methylacrylate addition to an aminopropyl substituent of a polydiorganosiloxane.



More specifically, Pike et al., U.S. Patent

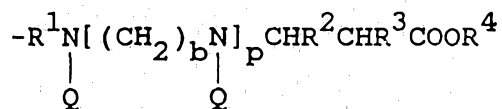
No. 3,033,815, May 8, 1962, describe the reaction of methylacrylate with a silicone copolymer of 90 percent dimethylsiloxane units and 10 percent delta-aminobutylmethylsiloxane units. The resulting silicone is described as containing delta-(N-2-carbomethoxyethyl)aminobutylmethylsiloxane units. Pike et al. teach that these compounds find use as sizes for fibrous materials, particularly fibrous glass materials employed in combination with thermosetting resins and that the monomeric and polymeric compounds can also be employed as adhesives or as flocculation agents.

Similarly, Ryan, U.S. Patent No. 3,317,577, May 2, 1967, describes the addition of methylacrylate to silanes and organosiloxanes having diaminoalkyl groups such as N-(2-aminoethyl)-3-aminopropyl groups ($-\text{CH}_2\text{CH}_2\text{CH}_2\text{NHCH}_2\text{CH}_2\text{NH}_2$). Ryan teaches that the materials are useful as binding agents for binding pigments to glass fabric and other siliceous surfaces, that they are useful as priming agents on metallic surfaces for securing adhesion of subsequently applied resin coatings, that they are useful as binding agents in the preparation of glass fiber polyester laminates, and that fluids prepared from the polymers and copolymers are also of interest as heat exchange agents and for further reaction to modify organic materials.

Neither Pike et al. nor Ryan suggest that their modified aminoalkyl substituted polydiorganosiloxanes are useful in hair care compositions. The use of these materials in hair treating compositions is believed to be novel.

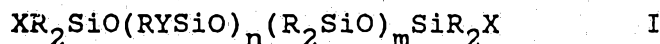
The present invention relates to a hair treating composition comprising (a) from 0.01 to 10 parts by weight of an organopolysiloxane represented by the formula

$$\text{XR}_2\text{SiO}(\text{RYSiO})_n(\text{R}_2\text{SiO})_m\text{SiR}_2\text{X}$$
wherein R denotes a monovalent hydrocarbon radical having 1 to 6 carbon atoms; X denotes -OH, -OR, Y, or R; Y denotes a substituent of the formula



wherein R^1 denotes an alkylene radical having 3 to 6 carbon atoms, R^2 denotes a hydrogen atom, phenyl, or an alkyl radical of 1 to 6 carbon atoms, R^3 denotes a hydrogen atom or methyl radical, and R^4 denotes an alkyl radical having 1 to 6 carbon atoms, b has a value of 2, 3, or 4, p has a value of 0, 1, or 2, and Q denotes a hydrogen atom or a radical of the formula $-CHR^2CHR^3COOR^4$ wherein R^2 , R^3 , and R^4 have the same meanings as defined above; m has an average value from 10 to 600; n has an average value of 0 to 100 with the proviso that when n is 0, X denotes Y and (b) from 50 to 100 parts by weight of a physiologically acceptable carrier. The invention also relates to the method of conditioning hair which comprises contacting hair with the treating composition defined above.

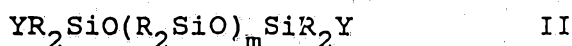
In the present invention, certain modified aminoalkyl substituted polydiorganosiloxanes are used in hair treating compositions such as conditioners, rinses, and shampoos. The useful polydiorganosiloxanes are represented by the formula



wherein R denotes a monovalent hydrocarbon radical having 1 to 6 carbon atoms. The organic groups denoted by R in formula I may be alkyl groups such as methyl, ethyl, or hexyl; alkenyl groups such as vinyl or allyl; or an aryl group such as phenyl. The diorganosiloxane polymers most useful in this invention are predominantly methyl substituted polymers wherein at least 90 percent of R radicals in the formula are methyl groups. It is most preferred to use polydimethylsiloxanes wherein essentially all R radicals are methyl groups.

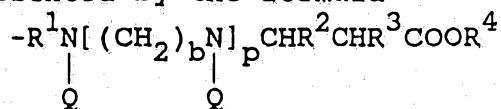
In formula I, X denotes $-OH$, $-OR$, Y , or R where R has the same meaning described above and Y denotes a modified aminoalkyl substituent. Thus, the polymer may be terminated by a hydroxyl group, a hydrocarbonoxy group, or a

triorganosiloxane unit. When n is not zero so that the polymer has pendant modified aminoalkyl substituents, it is preferred that the polymer be terminated by triorganosiloxane units such as dimethylvinylsiloxane units, dimethylphenylsiloxane units or trimethylsiloxane units, with trimethylsiloxane units being most preferred. When n is zero, X denotes a modified aminoalkyl substituent so that useful polymers have the formula



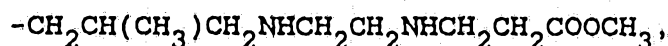
wherein R has the same meaning as described above.

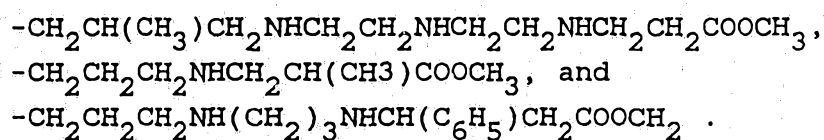
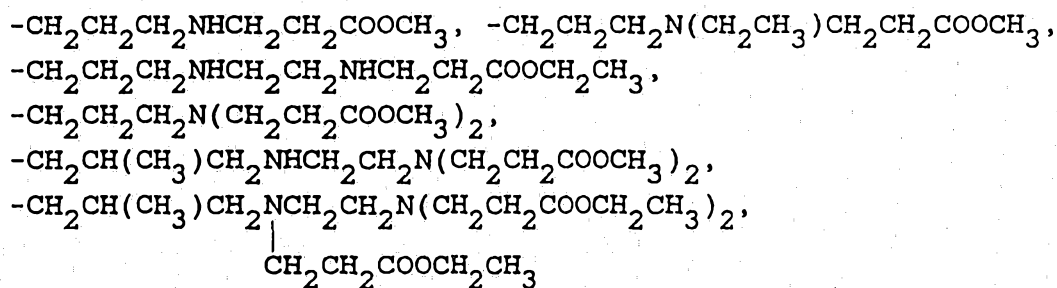
In formula I and II, Y denotes a modified aminoalkyl substituent represented by the formula



wherein R^1 denotes an alkylene radical having 3 to 6 carbon atoms. Alkylene radical, R^1 , may be a straight chain radical such as trimethylene, tetramethylene, or hexamethylene or a branched radical such as 2-methylpropylene, $-CH_2CH(CH_3)CH_2-$. R^2 denotes a hydrogen atom, phenyl, or an alkyl radical of 1 to 6 carbon atoms such as methyl, isopropyl, butyl, or hexyl. R^3 denotes a hydrogen atom of a methyl radical. R^4 denotes an alkyl radical having 1 to 6 carbon atoms, for example, methyl, ethyl, propyl, pentyl, or isobutyl. In the modified aminoalkyl substituent, b may have a value of 2, 3, or 4 and p may have a value of 0, 1, or 2. Q denotes a hydrogen atom, an alkyl radical having 1 to 4 carbon atoms, or a radical of the formula $-CHR^2CHR^3COOR^4$ wherein R^2 , R^3 , and R^4 have the same meanings as defined previously. Alkyl radicals having 1 to 4 carbon atoms include methyl, ethyl, isopropyl and butyl. Generally, silicones are preferred wherein Q denotes a hydrogen atom or a radical of the formula $-CHR^2CHR^3COOR^4$.

According to the present invention, useful modified aminoalkyl substituents include, for example,





The modified aminoalkyl substituted polydiorgano-siloxane may vary widely in viscosity and degree of polymerization. In formula I, for example, m may vary from an average value of 10 to 600 and n may vary from an average value of 0 to 100. However, polymers wherein m has an average value from 40 to 400 and n has an average value from 1 to 60 with the ratio of n to m being less than or equal to 0.15 are generally preferred for use in the hair treating compositions of this invention. Such polymers are preferred because they have viscosities which allow more easy formulation with other components typically used in hair treating compositions and because they have sufficient amino functionality to provide substantivity to hair.

Methods of preparing the modified aminoalkyl substituted polydiorganosiloxanes that are employed in the hair treating compositions according to this invention are known in the art. For example, U.S. Patent Nos. 3,033,815 and 3,317,577 teach methods whereby aminoalkyl substituted polydimethylsiloxanes are reacted with acrylate esters to prepare polymers useful in the present invention.

The modified aminoalkyl substituted polydiorgano-
siloxane ^{are preferably} ~~must be~~ dissolved or dispersed in a carrier to be
useful for treating hair. Typically, useful hair treating
compositions will contain from 0.01 to 10 parts of silicone

polymer diluted in 50 to 100 parts of a carrier liquid. Since the carrier serves only to dilute the silicone polymer to allow uniform application of appropriately small quantities, any carrier that is physiologically acceptable for use on the human body may be used. For example, the silicone polymer can be dissolved in organic solvents including chlorinated alkanes such as 1,1,1-trichloroethane and methylene dichloride, alcohols such as ethanol and isopropanol, and polyols such as ethylene glycol and propylene glycol.

The silicone polymer can also be used to treat hair in an aqueous dispersion or emulsion. Treating hair with an emulsion of silicone in water is an especially preferred embodiment of the present invention. Aqueous emulsions of silicone polymer may be prepared by high shear mixing of the polymer in water using a suitable emulsifying surfactant as is well known in the art. Microemulsions of silicone polymers in water are also highly preferred for use in the present invention. Microemulsions of modified aminoalkyl substituted polydiorganosiloxane can be prepared by the method described in U.S. Patent No. 4,620,878 which describes generally the preparation of emulsions of silicones containing polar substituents.

Depending on the specific hair treating application, the compositions of this invention may be thickened or unthickened creams, gels, aerosol foams, or sprays. Suitable thickeners include, among others, sodium alginate, gum arabic, polyoxyethylene, cellulose derivatives such as methylcellulose, hydroxymethylcellulose, hydroxyethylcellulose, hydroxypropylcellulose, hydroxypropylmethylcellulose, guar gum or its derivatives, starch and starch derivatives such as hydroxyethylamylose and starch amylose, locust bean gum, electrolytes such as NaCl, saccharides such as fructose and glucose, and derivatives of saccharides such as PEG-120 methyl glucose

dioleate. The hair treating compositions may be applied in the form of shampoo; rinsing products to be applied after shampooing, before or after tinting or bleaching, and before or after permanent waving or straightening; products for setting or brushing; conditioning compositions, restoring compositions; and compositions for permanent-waved hair.

The types of hair treating compositions encompassed by the present invention is further illustrated, but not limited, by specific reference to the following preferred embodiments. Hair treating compositions of this invention may form a shampoo, in which case, the compositions contain a cleansing surfactant in addition to the silicone polymer and carrier. The concentration of cleansing surfactant can range from about 2 to 40 parts by weight per 100 parts of total composition. Cleansing surfactants selected from the group consisting of anionic surfactants, nonionic surfactants, and amphoteric surfactants are well known for use in shampoo formulations. For example, the anionic, nonionic and amphoteric surfactants useful as cleansing agents in shampoos are further described in U.S. Patent No. 4,559,227.

Typical cleansing surfactants include the anionics such as the sodium, ammonium, or triethanolamine salts of lauryl sulfate and lauryl ether sulfate; the nonionics such as fatty acid alkanolamides like lauric acid diethanolamide; and the amphoterics such as N-cocamidopropyl dimethyl glycine. Generally the anionic surfactants, especially the sodium, ammonium, and triethanolamine salts of lauryl sulfate, are preferred since they provide richer, denser foams than other types of cleansing surfactants at comparable concentrations.

In another embodiment of the present invention, the hair treating composition may form a conditioning product for application to hair after shampooing. The hair is typically rinsed in running water after treating with the conditioning

product. Conditioners facilitate combing out hair and impart softness and suppleness to the hair. Conditioning products may also contain other components such as thickeners and auxiliary conditioning compounds. Auxiliary conditioning agents may be used to provide further improved conditioning benefits such as antistatic characteristics. Auxiliary conditioning agents useful in the compositions of this invention include the organic cationic compounds and polymers such as stearyldimethylbenzylammonium chloride, quaternary nitrogen derivatives of cellulose ethers, and homopolymers and copolymers of dimethyldiallylammonium chloride, and other quaternary ammonium compounds which are known for use in hair conditioning formulations.

In another embodiment of the present invention, the hair treating composition may be in the form of a product for permanent waving hair. The conventional technique for these permanent waves consists in applying, in a first stage, a composition containing a reducing agent and then, after having rinsed the hair if appropriate, applying a composition containing an oxidizing agent. According to the present invention, at least one of the two compositions is a hair treating composition containing modified aminoalkyl substituted polydiorganosiloxane and carrier in addition to the reducing or oxidizing agent.

One advantage of the hair treating compositions of this invention is the increased amount of silicone polymer deposited on the hair in comparison to the amount of silicone deposited when hair is treated by prior art silicone materials. Also, the hair treating compositions of the invention are more easily formulated with other hair care components because of the reduced reactivity of the modified aminoalkyl substituted polydiorganosiloxane with other components.

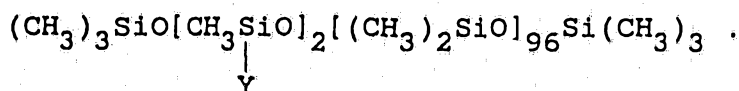
The following examples are presented to illustrate the invention to those skilled in the art and should not be construed as limiting the invention, which is properly delineated in the appended claims. All proportions by parts or percents are by weight unless otherwise stated.

Example 1

This example shows that aminoalkyl substituted polydiorganosiloxane which has been modified by reaction with methylacrylate is useful to condition hair. Also, the combing improvement provided by the methylacrylate modified silicone is compared to the improvement provided by the unmodified silicone and to silicone modified by reaction with acetic anhydride.

Hair treating compositions suitable for use as hair conditioners were prepared by making an emulsion of the various silicone polymers in water. Stock emulsions were prepared by mixing 4.4 g of isolaureth-6 ($C_{12}H_{25}(OCH_2CH_2)_6OH$ sold under the trademark TERGITOL TMN-6 by Union Carbide Corp., Danbury, CT), 12.1 g of octoxynol-40 ($C_8H_{17}C_6H_4(OCH_2CH_2)_{40}OH$ sold under the trademark TRITON X-405 by Rohm and Haas Company, Inc., Philadelphia, PA), and 76.1 g of water and then adding 50 g of silicone polymer while homogenizing the mixture on a colloid mill. The stock emulsions contained 35% silicone, 11.6% emulsifying surfactants, and 53.4% water. Portions of each stock emulsion were diluted to 0.25% silicone with additional water to provide baths for treating hair in the following tests.

The silicone polymers tested generally conformed to the average formula



The specific structure of the Y substituents for the polymers in each emulsion tested is shown in Table 1. Silicone Emulsions A and B illustrate the present invention while

Silicone Emulsions C, D, and E represent the prior art and are presented for comparison purposes.

TABLE 1

<u>Silicone Emulsion</u>	<u>Y</u>
A	$-\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{NHCH}_2\text{CH}_2\text{NHCH}_2\text{CH}_2\text{COOCH}_3$
B*	$ \begin{array}{c} -\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{NCH}_2\text{CH}_2\text{NCH}_2\text{CH}_2\text{COOCH}_3 \\ \quad \quad \quad \quad \quad \quad \\ \quad \quad \quad \text{O} \quad \quad \quad \text{O} \end{array} $
C	$-\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{NHCH}_2\text{CH}_2\text{NH}_2$
D	$ \begin{array}{c} \text{O} \\ \\ -\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{NHCH}_2\text{CH}_2\text{NHCCH}_3 \end{array} $
E	$ \begin{array}{c} \text{O} \\ \\ -\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{NCH}_2\text{CH}_2\text{NHCCH}_3 \\ \quad \quad \quad \\ \quad \quad \quad \text{O}=\text{CCH}_3 \end{array} $

* O groups are half hydrogen and half methoxycarbonylethyl groups of the formula $-\text{CH}_2\text{CH}_2\text{COOCH}_3$

The conditioning efficacy of each silicone emulsion was evaluated by measuring the force required to comb both wet and dry hair after it had been treated with the compositions. Hair tresses were made by assembling 12 g bundles of 30 cm natural brown, virgin European hair and binding the root ends in wax. All reported combing results are averages for three tresses treated with each composition.

Combing forces were measured on an INSTRON strain gauge adapted with a hard rubber comb. The combing force was recorded as the comb was moved down through the tress which was attached to the end of a stationary load cell. The average combing load (ACL) was determined by integrating the total combing force over the entire tress length and is reported in grams. Four combing strokes were conducted for each treatment

condition of a tress and averaged to obtain more representative values.

Tresses were washed with a conventional shampoo and dried at least 18 hr under ambient conditions prior to initial combing force measurement in the dry/untreated state. Tresses were then soaked in water for 12-30 minutes and combing forces measured again in the wet/untreated state. Between each of the four wet combing strokes, the hair was dipped three times in water to uniformly retangle the hair. Tresses were then treated by dipping for 30 sec. in a 200 g bath of silicone emulsion (0.25% silicone solids). The tresses were rinsed for 30 sec. in 40°C. running tap water and the combing forces measured in the wet/treated state. Combing forces were measured again in the dry/treated state after the hair had dried for at least 18 hr under ambient conditions. Combs were changed between treated and untreated tresses, and a new comb is used for each treatment material.

The percent change in ACL was calculated as

$$\% \text{ Change} = \frac{ACL_T - ACL_U}{ACL_U} \times 100$$

where ACL_U

is the average combing load for the untreated tress, and ACL_T is the average combing load for the treated tress. Percent change values are shown in table 2 for both the dry and the wet state.

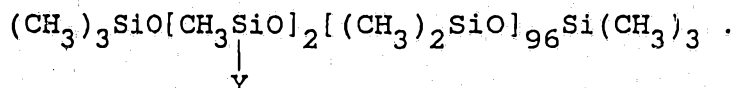
TABLE 2

Silicone Emulsion Treatment	Percent Change in ACL	
	Dry	Wet
A	-37	-69
B	-35	-53
C	-65	-66
D	-35	-63
E	-10	+1

Example 2

This example shows an additional type of methylacrylate modified aminoalkyl substituted polydiorganosiloxane that is useful to condition hair. Also, the combing improvement provided by the methylacrylate modified silicone is compared to the improvement provided by the unmodified silicone.

Hair treating compositions were prepared by making an emulsion of the various silicone polymers in water according to the procedure described in Example 1. The silicone polymers tested generally conformed to the average formula



The specific structure of the Y substituents for the polymers in each emulsion tested is shown in Table 3. Silicone Emulsion F illustrates the present invention while Silicone Emulsions G and H represent the prior art and are presented for comparison purposes.

TABLE 3

<u>Silicone Emulsion</u>	<u>Y</u>
F	$-\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{N}(\text{CH}_3)\text{CH}_2\text{CH}_2\text{COOCH}_3$
G	$-\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{NHCH}_3$
H	$-\text{CH}_2\text{CH}(\text{CH}_3)\text{CH}_2\text{NHCH}_2\text{CH}_2\text{NH}_2$

The conditioning efficacy of each silicone emulsion was evaluated by measuring the force required to comb dry hair after it had been treated with the compositions. The hair tresses were treated and combing forces measured by the same procedures described in Example 1. The percent change in average combing load between treated and untreated tresses is shown in Table 4.

TABLE 4

Silicone Emulsion	Percent Change in ACL
Treatment	Dry
F	-15
G	-52
H	-12

Example 3

This example demonstrates the substantivity of the silicones of this invention on hair and compares their substantivity to that of prior silicones used in hair care.

Clippings of the treated hair tresses from Example 1 were analyzed to determine the amount of silicone bound or adsorbed on the hair. In addition, after initial clippings were removed, the tresses were washed with 0.6 g of conventional shampoo for 60 sec., blow dried, and additional clippings taken for analysis. Additional washing steps were performed with further clippings taken for analysis after the 3rd and 5th wash.

Hair clippings were analyzed by atomic absorption determination of silicon after the hair had been digested with papain enzyme. The digested hair samples were extracted in acidified methyl isobutyl ketone for analysis. The results are presented as parts per million (ppm) of total silicon in Table 5.

The data in Table 5 demonstrates that the acrylate modified silicones surprisingly deposit more effectively on hair than the other materials, yet they are effectively removed after 1 to 3 washing so that undesirable build up on the hair does not occur.

Table 5

Silicone Treatment	Silicon Detected (ppm)			
	Number of Washes After Treatment			
	0	1	3	5
A	1460	430	260	270
B	1060	180	160	180
C*	580	350	270	270
D*	550	190	130	140
E*	180	100	130	100
Untreated Hair	180	110	60	100

*Comparison examples, not encompassed by the present invention

Example 4

This example illustrates the preparation of a shampoo formulation within the hair treating compositions of the present invention. The shampoo employs ammonium lauryl sulfate as an anionic cleansing surfactant.

A shampoo was prepared by first heating a solution of 9 g of ammonium lauryl sulfate in 61 g of water to 80°C. A separate mixture of 6 g of lauric acid diethanolamide and 2 g of the silicone polymer from Silicone Emulsion A of Example 1 (a methylacrylate modified aminoalkyl substituted polydimethylsiloxane) was heated to 80°C. and then added with stirring to the aqueous surfactant solution. A clear solution resulted which was thickened by addition of 2.5 g of PEG-120 methyl glucose dioleate (a methyl glucoside derivative with two polyoxyethylene substituent groups containing a total of 120 oxyethylene units). The thickener was melted and then mixed into the shampoo composition. After cooling, the pH of the

composition was adjusted to 6.8 with 25% aqueous citric acid solution and sufficient additional water added to make 100 g of shampoo composition.

Example 5

This example illustrates the preparation of a shampoo formulation within the hair treating compositions of the present invention. The shampoo employs a mixture of ammonium lauryl sulfate (anionic) and N-cocamidopropyl dimethyl glycine (amphoteric) as cleansing surfactant.

A mixture of 5.25 g of N-cocamidopropyl dimethyl glycine, 4 g of lauric acid diethanolamide, 2 g of the silicone polymer from Silicone Emulsion A of Example 1, and 12.25 g of water was heated to 80°C. and added to a similarly heated solution of 5.25 g of ammonium lauryl sulfate in 24.25 g of water. After cooling, an additional 43 g of water was added. The pH was adjusted to 5.5-6.0 with 25% aqueous citric acid. The solution was thickened by adding NaCl and sufficient additional water added to make 100 g of shampoo composition.

Example 6

This example illustrates the improved skin toxicological characteristics of the silicones used in the hair treating compositions of the present invention. The silicone polymers described in Example 1 were tested for acute eye and dermal irritation in rabbits. The results are presented in

Table 6.

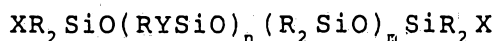
Table 6

Silicone Polymer	Acute Eye Irritation	Acute Dermal Irritation
A	slight irritant	slight irritant
B	slight redness	non-irritating
C*	moderate irritant	moderate to severe irritation
D*	moderate irritant	moderate irritation
E*	slight redness	non-irritating

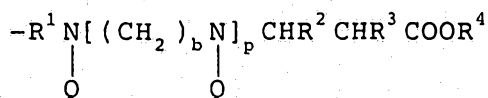
*Comparison examples, not encompassed by the present invention

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. The method of treating or conditioning hair which comprises contacting hair with a polydiorganosiloxane represented by the formula



wherein R denotes a monovalent hydrocarbon radical having 1 to 6 carbon atoms; X denotes -OH, -OR, Y, or R; Y denotes a substituent of the formula



wherein R¹ denotes an alkylene radical having 3 to 6 carbon atoms, R² denotes a hydrogen atom, phenyl, or an alkyl radical of 1 to 6 carbon atoms, R³ denotes a hydrogen atom or methyl radical and R⁴ denotes an alkyl radical having 1 to 6 carbon atoms, b has a value of 2, 3, or 4, p has a value of 0, 1, or 2, and Q denotes a hydrogen atom or a radical of the formula -CHR²CHR³COOR⁴ wherein R², R³ and R⁴ have the same meanings as defined above; m has an average value from 10 to 600; n has an average value of 0 to 100 with the proviso that when n is 0, X denotes Y.

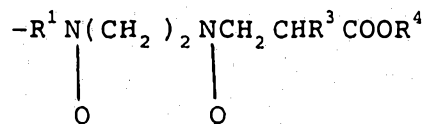
2. A method as claimed in claim 1 in which the polydiorganosiloxane is applied together with a physiologically acceptable carrier.

3. A method as claimed in claim 1 or 2 in which the polydiorganosiloxane is applied together with from 2 to 40 parts by weight of a cleansing surfactant selected from the group consisting of anionic surfactants, nonionic surfactants and amphoteric surfactants.

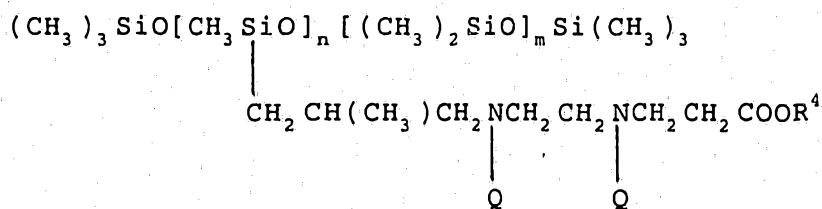
4. A method as claimed in any one of claims 1 to 3 wherein R denotes methyl and p has a value of 1.



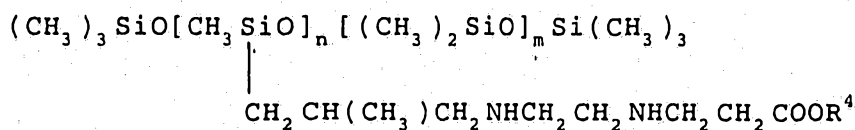
5. A method as claimed in any one of claims 1 to 4 wherein X denotes methyl, n has an average value of 1 to 100 and Y denotes a substituent of the formula



6. A method as claimed in any one of claims ^{1 to 5} ~~12 to 16~~ wherein the polydiorganosiloxane is represented by the formula



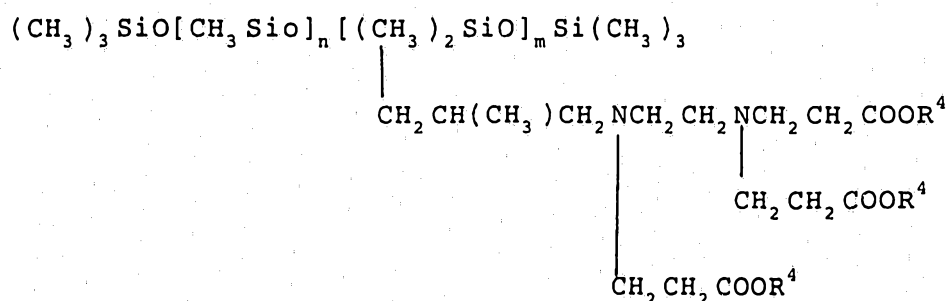
7. A method as claimed in any one of claims 1 to 6 wherein the polydiorganosiloxane is represented by the formula



8. A method as claimed in any one of claims 1 to 7 wherein about 50 percent of the substituents denoted by Q are $-CH_2CH_2COOR^4$ and the remainder are hydrogen.



9. A method as claimed in any one of claims 1 to 8 wherein the polydiorganosiloxane is represented by the formula



10. A method as claimed in any one of claims 1 to 9 wherein m has an average value from 40 to 400, n has an average value from 1 to 60, and ratio of n to m is less than or equal to 0.15.

11. A method as claimed in any one of claims 1 to 10 wherein the carrier is water.

DATED this 13th day of February, 1990

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