

## 特許協力条約に基づく国際出願願書

紙面による写し(注意 電子データが原本となります)

|        |                                            |                                                                  |
|--------|--------------------------------------------|------------------------------------------------------------------|
| 0      | 受理官庁記入欄                                    |                                                                  |
| 0-1    | 国際出願番号                                     |                                                                  |
| 0-2    | 国際出願日                                      |                                                                  |
| 0-3    | (受付印)                                      |                                                                  |
| 0-4    | 様式 PCT/RO/101<br>この特許協力条約に基づく国際出願願書は、      |                                                                  |
| 0-4-1  | 右記によって作成された。                               | JPO-PAS<br>0333                                                  |
| 0-5    | 申立て<br>出願人は、この国際出願が特許協力条約に従って処理されることを請求する。 |                                                                  |
| 0-6    | 出願人によって指定された受理官庁                           | 日本国特許庁 (RO/JP)                                                   |
| 0-7    | 出願人又は代理人の書類記号                              | A65124H                                                          |
| I      | 発明の名称                                      | コンタクトレンズ材料の製造方法およびソフトコンタクトレンズの製造方法                               |
| II     | 出願人                                        |                                                                  |
| II-1   | この欄に記載した者は                                 | 出願人である (applicant only)                                          |
| II-2   | 右の指定国についての出願人である。                          | 米国を除く全ての指定国 (all designated States except US)                    |
| II-4ja | 名称                                         | HOYA 株式会社                                                        |
| II-4en | Name:                                      | HOYA CORPORATION                                                 |
| II-5ja | あて名                                        | 1618525<br>日本国<br>東京都新宿区中落合2丁目7番5号                               |
| II-5en | Address:                                   | 7-5, Naka-Ochiai 2-chome, Shinjuku-ku, Tokyo<br>1618525<br>Japan |
| II-6   | 国籍(国名)                                     | 日本国 JP                                                           |
| II-7   | 住所(国名)                                     | 日本国 JP                                                           |
| II-8   | 電話番号                                       | 042-596-1154                                                     |
| II-9   | ファクシミリ番号                                   | 042-596-6466                                                     |
| II-11  | 出願人登録番号                                    | 000113263                                                        |

## 特許協力条約に基づく国際出願願書

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|           |                                                                                                                                                                                      |                                                                                               |
|-----------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------|
| III-1     | その他の出願人又は発明者                                                                                                                                                                         |                                                                                               |
| III-1-1   | この欄に記載した者は                                                                                                                                                                           | 出願人及び発明者である (applicant and inventor)                                                          |
| III-1-2   | 右の指定国についての出願人である。                                                                                                                                                                    | 米国のみ (US only)                                                                                |
| III-1-4ja | 氏名(姓名)                                                                                                                                                                               | 今福 元                                                                                          |
| III-1-4en | Name (LAST, First):                                                                                                                                                                  | IMAFUKU, Suguru                                                                               |
| III-1-5ja | あて名                                                                                                                                                                                  | 1618525<br>日本国<br>東京都新宿区中落合2丁目7番5号 HOYA株式会社内                                                  |
| III-1-5en | Address:                                                                                                                                                                             | c/o Hoya Corporation, 7-5, Naka-Ochiai 2-chome, Shinjuku-ku, Tokyo<br>1618525<br>Japan        |
| III-1-6   | 国籍(国名)                                                                                                                                                                               | 日本国 JP                                                                                        |
| III-1-7   | 住所(国名)                                                                                                                                                                               | 日本国 JP                                                                                        |
| IV-1      | 代理人又は共通の代表者、通知のあて名<br>下記の者は国際機関において右記のごとく出願人のために行動する。                                                                                                                                | 代理人 (agent)                                                                                   |
| IV-1-1ja  | 名称                                                                                                                                                                                   | 特許業務法人特許事務所サイクス                                                                               |
| IV-1-1en  | Name:                                                                                                                                                                                | SIKs & Co.                                                                                    |
| IV-1-2ja  | あて名                                                                                                                                                                                  | 1040031<br>日本国<br>東京都中央区京橋一丁目8番7号 京橋日殖ビル8階                                                    |
| IV-1-2en  | Address:                                                                                                                                                                             | 8th Floor, Kyobashi-Nisshoku Bldg., 8-7, Kyobashi 1-chome, Chuo-ku, Tokyo<br>1040031<br>Japan |
| IV-1-3    | 電話番号                                                                                                                                                                                 | 03-3538-5680                                                                                  |
| IV-1-4    | ファクシミリ番号                                                                                                                                                                             | 03-3538-5686                                                                                  |
| IV-1-5    | 電子メール                                                                                                                                                                                | mail@siks.jp                                                                                  |
| IV-1-6    | 代理人登録番号                                                                                                                                                                              | 110000109                                                                                     |
| V         | 国の指定                                                                                                                                                                                 |                                                                                               |
| V-1       | この願書を用いてされた国際出願は、規則4.9(a)に基づき、国際出願の時点で拘束される全てのPCT締約国を指定し、取得しうるあらゆる種類の保護を求め、及び該当する場合には広域と国内特許の両方を求める国際出願となる。                                                                          |                                                                                               |
| V-2       | V-2欄に示される国は、それらの国々の国内法令に基づき、国際出願が主張する優先権主張の基礎となる先の国内出願の効果が消滅することを避けることを目的に、当該国の指定を除外される。しかし、いったん除外した指定は、それを変更することはできない。(これらの国及びそのような制度を有する国が持つ国内法令手続の結果に関しては、PCT-SAFEヘルプの国の指定の項を参照。) | JP                                                                                            |

## 特許協力条約に基づく国際出願願書

紙面による写し (注意 電子データが原本となります)

|        |                                                                                    |                              |            |
|--------|------------------------------------------------------------------------------------|------------------------------|------------|
| VI-1   | 先の国内出願に基づく優先権主張                                                                    |                              |            |
| VI-1-1 | 出願日                                                                                | 2005年 09月 05日 (05. 09. 2005) |            |
| VI-1-2 | 出願番号                                                                               | 2005-256647                  |            |
| VI-1-3 | 国名                                                                                 | 日本国 JP                       |            |
| VI-2   | 優先権証明書送付の請求<br>上記の先の出願のうち、右記の番号のものについては、出願書類の認証謄本を作成し国際事務局へ送付することを、受理官庁に対して請求している。 | VI-1                         |            |
| VII-1  | 特定された国際調査機関(ISA)                                                                   | 日本国特許庁 (ISA/JP)              |            |
| VIII   | 申立て                                                                                | 申立て数                         |            |
| VIII-1 | 発明者の特定に関する申立て                                                                      | —                            |            |
| VIII-2 | 出願し及び特許を与えられる国際出願日における出願人の資格に関する申立て                                                | —                            |            |
| VIII-3 | 先の出願の優先権を主張する国際出願日における出願人の資格に関する申立て                                                | —                            |            |
| VIII-4 | 発明者である旨の申立て(米国を指定国とする場合)                                                           | —                            |            |
| VIII-5 | 不利にならない開示又は新規性喪失の例外に関する申立て                                                         | —                            |            |
| IX     | 照合欄                                                                                | 用紙の枚数                        | 添付された電子データ |
| IX-1   | 願書(申立てを含む)                                                                         | 4                            | ✓          |
| IX-2   | 明細書                                                                                | 18                           | ✓          |
| IX-3   | 請求の範囲                                                                              | 1                            | ✓          |
| IX-4   | 要約                                                                                 | 1                            | ✓          |
| IX-5   | 図面                                                                                 | 3                            | ✓          |
| IX-7   | 合計                                                                                 | 27                           |            |
| IX-8   | 添付書類                                                                               | 添付                           | 添付された電子データ |
|        | 手数料計算用紙                                                                            | —                            | ✓          |
| IX-17  | PCT-SAFE 電子出願                                                                      | —                            | —          |
| IX-19  | 要約書とともに提示する図の番号                                                                    |                              |            |
| IX-20  | 国際出願の使用言語名                                                                         | 日本語                          |            |
| X-1    | 出願人、代理人又は代表者の記名押印                                                                  | /110000109/                  |            |
| X-1-1  | 氏名(姓名)                                                                             | 特許業務法人特許事務所サイクス              |            |
| X-1-2  | 署名者の氏名                                                                             |                              |            |
| X-1-3  | 権限                                                                                 |                              |            |

## 特許協力条約に基づく国際出願願書

紙面による写し(注意 電子データが原本となります)

## 受理官庁記入欄

|        |                                                                      |        |
|--------|----------------------------------------------------------------------|--------|
| 10-1   | 国際出願として提出された書類<br>の実際の受理の日                                           |        |
| 10-2   | 図面                                                                   |        |
| 10-2-1 | 受理された                                                                |        |
| 10-2-2 | 不足図面がある                                                              |        |
| 10-3   | 国際出願として提出された書類<br>を補完する書類又は図面であつ<br>てその後期間内に提出されたも<br>のの実際の受理の日(訂正日) |        |
| 10-4   | 特許協力条約第11条(2)に基づ<br>く必要な補完の期間内の受理の日                                  |        |
| 10-5   | 出願人により特定された国際調査機関                                                    | ISA/JP |
| 10-6   | 調査手数料未払いにつき、国際<br>調査機関に調査用写しを送付していない                                 |        |

## 国際事務局記入欄

|      |           |  |
|------|-----------|--|
| 11-1 | 記録原本の受理の日 |  |
|------|-----------|--|

## PCT手数料計算用紙(願書付属書)

紙面による写し(注意 電子データが原本となります)

[この用紙は、国際出願の一部を構成せず、国際出願の用紙の枚数に算入しない]

|         |                                                            |                                                                  |          |  |
|---------|------------------------------------------------------------|------------------------------------------------------------------|----------|--|
| 0       | 受理官庁記入欄                                                    |                                                                  |          |  |
| 0-1     | 国際出願番号                                                     |                                                                  |          |  |
| 0-2     | 受理官庁の日付印                                                   |                                                                  |          |  |
| 0-4     | 様式 PCT/RO/101(付属書)<br>このPCT手数料計算用紙は、<br>0-4-1 右記によって作成された。 | JP0-PAS<br>0333                                                  |          |  |
| 0-9     | 出願人又は代理人の書類記号                                              | A65124H                                                          |          |  |
| 2       | 出願人                                                        | HOYA株式会社                                                         |          |  |
| 12      | 所定の手数料の計算                                                  | 金額/係数                                                            | 小計 (JPY) |  |
| 12-1    | 送付手数料 T                                                    | ⇒                                                                | 13000    |  |
| 12-2    | 調査手数料 S                                                    | ⇒                                                                | 97000    |  |
| 12-3    | 国際出願手数料<br>(最初の30枚まで) i1                                   | 121800                                                           |          |  |
| 12-4    | 30枚を越える用紙の枚数                                               | 0                                                                |          |  |
| 12-5    | 用紙1枚の手数料 (X)                                               | 0                                                                |          |  |
| 12-6    | 合計の手数料 i2                                                  | 0                                                                |          |  |
| 12-7    | i1 + i2 = i                                                | 121800                                                           |          |  |
| 12-12   | fully electronic filing fee reduction R                    | -26100                                                           |          |  |
| 12-13   | 国際出願手数料の合計 (i-R) I                                         | ⇒                                                                | 95700    |  |
| 12-17   | 納付すべき手数料の合計 (T+S+I+P)                                      | ⇒                                                                | 205700   |  |
| 12-19   | 支払方法                                                       | 送付手数料: 予納口座引き落としの承認<br>調査手数料: 予納口座引き落としの承認<br>国際出願手数料: 銀行口座への振込み |          |  |
| 12-20   | 予納口座<br>受理官庁                                               | 日本国特許庁 (RO/JP)                                                   |          |  |
| 12-20-1 | 上記手数料合計額の請求に対する承認                                          | ✓                                                                |          |  |
| 12-21   | 予納口座番号                                                     | 170347                                                           |          |  |
| 12-22   | 日付                                                         | 2006年 08月 30日 (30.08.2006)                                       |          |  |
| 12-23   | 記名押印                                                       |                                                                  |          |  |

## SPECIFICATION

Method of manufacturing contact lens material and method of manufacturing soft contact lens

### TECHNICAL FIELD

[0001] The present invention relates to a method of manufacturing a contact lens material having a smooth surface with good water wettability and durability, and to a method of manufacturing a soft contact lens employing the contact lens material obtained by the above method.

### TECHNICAL BACKGROUND

[0002] Clinical results have indicated that the supply of oxygen from the atmosphere diminishes when wearing contact lenses, resulting in the inhibition of corneal epithelial cell mitosis and thickening of the cornea. Accordingly, attempts have been made to improve the oxygen permeability of materials to provide safer contact lenses.

[0003] Water-containing soft contact lenses are known to produce a pleasant wear comfort due to the flexibility of the material, but the oxygen permeability they afford is lower than that of hard contact lenses due to the moisture content of the lenses. For example, in the case of water-containing soft contact lenses, the oxygen permeation coefficient of a material with an 80 percent water content is about  $40 \times 10^{-11}$  (cm<sup>2</sup>/s) · (mLO<sub>2</sub>/mL x mmHg); an adequate amount of oxygen does not reach the cornea. For such reasons, soft contact lenses having lens components in the form of silicone-containing monomers and siloxane macromers to increase oxygen permeability and contact lenses having lens components in the form of fluorine-containing monomers to prevent contamination have been proposed. For example, Japanese Unexamined Patent Publication (KOKAI) No. 2001-311917 discloses a soft contact lens material having good flexibility and oxygen permeability employing a macromer having a dimethylsiloxane structure in a side chain. However, employing silicon components and fluorine components in lens materials generally tends to decrease the water wettability of the lens surface. As a result, contaminants

in lacrimal fluid components tend to adhere, sometimes compromising wear comfort.

[0004] Under such a situation, various surface treatment methods have been proposed for contact lenses to enhance the water wettability of the surface. Japanese Examined Patent Publication (KOKOKU) Showa No. 63-040293 discloses a treatment method in which coating of silicone or polyurethane lenses is conducted under a hydrocarbon atmosphere essentially not comprising nitrogen or oxygen, after which the lenses are exposed to oxygen, thereby increasing the hydrophilic property of the lens surface. In such a coating method, a silicone hydrogel lens is treated with plasma while not containing water. However, to obtain the final lens product, after treating the lens with plasma, an organic solvent such as alcohol is employed to remove unpolymerized monomers, oligomers, and the like the retention of which in the lens is biologically undesirable. A physiological saline, soft contact lens storage solution (packing solution), or the like is then substituted, followed by high-pressure steam sterilization processing. In such steps of extraction with an organic solvent such as alcohol or the like and water introduction using physiological saline or a soft contact lens storage solution, the volume of the lens swells by about 10 to 50 percent. Thus, since the coating film is a rigid film with an extremely high crosslinking density that is formed in a hydrocarbon atmosphere, there is a problem in that the polymer film separates, forming cracks in the lens surface. The defect level of separation and cracking of the film caused by the extraction and swelling steps is exacerbated by the subsequent high-pressure steam sterilization treatment, rendering the lens unsuitable for use as a contact lens worn in direct contact with the eye.

[0005] Japanese Unexamined Patent Publication (KOKAI) Showa No. 62-031803 discloses a plasma treatment method with a mixed gas of a saturated hydrocarbon and oxygen. However, in plasma polymerization with a mixed gas of saturated hydrocarbon and oxygen, the polymer film that is formed by the plasma is etched away more rapidly than the film formed on the lens due to etching effect attributed by oxygen plasma. Thus, considerable time is required to achieve a desired film thickness, which is problematic for the commercialization of this method.

[0006] "Application of plasma polymerization on silicone rubber contact lens surfaces and its effects" in *Collected Articles on Polymers* (Vol. 42, No. 11, pp. 841-847, 1985) describes a polymer film with good water wettability and durability employing N-vinyl-2-pyrrolidone ("NVP" hereinafter) monomer. This method permits the formation of a polymer film with good water wettability and durability. However, the NVP employed in the coating is a known carcinogen when in a vapor state. When employing organic monomers such as NVP as gases in plasma processing, great care must be exercised for the safety of workers and at worksites. Thus, this method also presents problems for commercialization.

#### DISCLOSURE OF THE INVENTION

[0007] Accordingly, the object of the present invention is to provide a contact lens material having a smooth surface with good water wettability and durability such that, after coating the lens surface while the lens contains no water, an extraction step with an organic solvent such as alcohol, a water introduction step employing physiological saline or a soft contact lens storage solution, and a high-pressure steam sterilization step, the film that is formed on the lens surface is able to withstand volumetric swelling of the lens and the surface of the lens does not have deficiencies that become problems when the lens is worn, as well as having surface water wettability and durability adequate for wearing.

[0008] The present invention relates to a method of manufacturing a contact lens material comprising a contact lens base material and a coating formed on at least a portion of a surface of said base material, wherein

said contact lens base material is processed by plasma polymerization in a mixed gas atmosphere of methane and moist air, and then treated with plasma in a nonpolymerizable gas atmosphere to form said coating.

Furthermore, the present invention relates to a method of manufacturing a soft contact lens, wherein

water is introduced into the contact lens material manufactured by the above method to obtain a soft contact lens.

[0009] The present invention can provide a soft contact lens having a smooth surface possessing good water wettability and durability, and more particularly, a soft contact lens comprised of hydrogel that contains silicone as a lens component.

#### BEST MODE FOR CARRYING OUT THE INVENTION

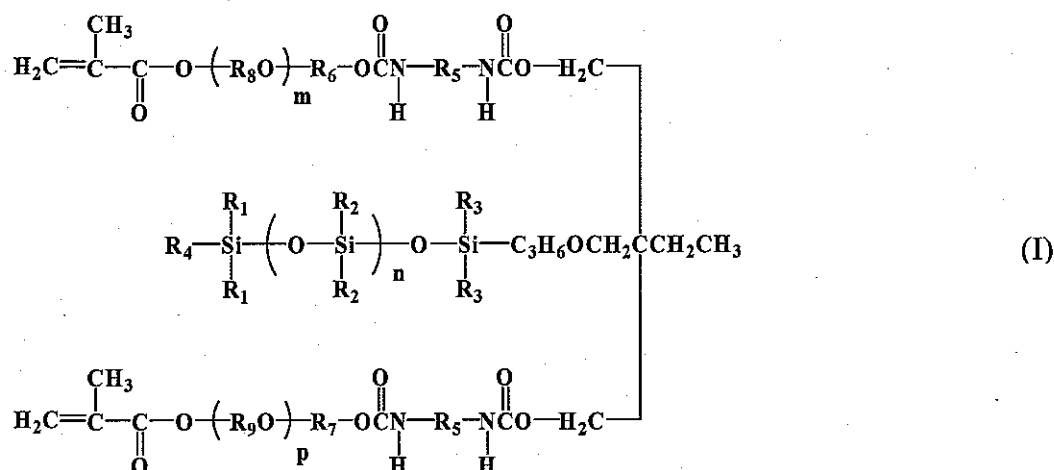
[0010] The present invention will be described in greater detail below.

[Method of manufacturing contact lens material]

The present invention relates to a method of manufacturing a contact lens material comprising a contact lens base material and a coating formed on at least a portion of a surface of said base material. In the method of manufacturing a contact lens material of the present invention, said contact lens base material is processed by plasma polymerization in a mixed gas atmosphere of methane and moist air, and then treated with plasma in a nonpolymerizable gas atmosphere to form said coating.

[0011] The aforementioned contact lens base material can be a polymer capable of becoming a hydrogel, preferably a silicone-containing copolymer capable of becoming a hydrogel, having the shape of a contact lens. Base materials that are commonly known as those for soft contact lens may be employed. Specific examples of such contact lens base materials are base materials comprised of copolymers obtained by polymerizing a mixture of at least one silicone-containing monomer, or silicone-containing macromonomer, and at least one hydrophilic monomer. An example of an applicable silicone monomer is tris(trimethylsiloxy)-*gamma*-methacryloxypropylsilane. One example of a silicone-containing macromonomer is a siloxane macromonomer having a number average molecular weight of about 1,000 to 10,000 with a polysiloxane structure in a side chain, such as that shown in General Formula (I) below.

[Chemical formula 1]



[In the formula,  $\text{R}_1$ ,  $\text{R}_2$  and  $\text{R}_3$  are each independently selected from C1 to C4 alkyl groups,  $\text{R}_4$  is selected from C1 to C6 alkyl groups,  $\text{R}_5$  is a residue in which NCO is removed from an aliphatic, alicyclic or aromatic diisocyanate,  $\text{R}_6$ ,  $\text{R}_7$ ,  $\text{R}_8$  and  $\text{R}_9$  are each independently selected from C1 to C3 alkylene,  $n$  is an integer of 4 to 80, and  $m$  and  $p$  are each independently an integer of 3 to 40.]

Examples of hydrophilic monomers are 2-hydroxyethyl methacrylate, N,N-dimethylacrylamide, N-vinyl-2-pyrrolidone, and methacrylic acid. Japanese Unexamined Patent Publication (KOKAI) No. 2001-311917, Published Japanese Translation of a PCT International Application (TOKUHYO) Heisei No. 10-509763, Japanese Unexamined Patent Publication (KOKAI) No. 2003-228029, Published Japanese Translation of a PCT International Application (TOKUHYO) Heisei No. 11-502894, and the like can be referred to for details regarding contact lens base materials that are suitable for use in the present invention.

[0012] The contact lens base material can be manufactured by various conventional techniques (such as lathe cutting, spin casting, and cast molding manufacturing methods), in the cast molding manufacturing method, the lens is removed from the mold, and then processed to form a coating (processed by plasma polymerization under a mixed gas atmosphere and treated with plasma under a nonpolymerizable gas atmosphere as set forth above).

[0013] In the present invention, the coating is formed on at least a portion of the surface of the contact lens base material. The coating is formed by processing the

contact lens base material by plasma polymerization in a mixed gas atmosphere of methane and moist air (referred to as "first step", hereinafter), followed by plasma treatment in a nonpolymerizable gas atmosphere (referred to as "second step", hereinafter). In the present invention, the term "processing by plasma polymerization" refers to generating a plasma state by an electrical discharge in a suitable degree of vacuum in a plasma polymerization device and forming a thin film (polymer film) formed with polymerizable gas on the surface of the base material, and "treatment with plasma" refers to modification of the outermost layer of the surface of the base material with a nonpolymerizable gas.

[0014]     FIRST STEP

In the first step, the contact lens base material is processed by plasma polymerization in a mixed gas atmosphere of methane and moist air. Specifically, the contact lens base material is introduced into a plasma polymerization device and a vacuum is generated to bring the arrival pressure within the device to below a certain range. When the contact lens base material is conveyed into a vacuum device and a vacuum is drawn, in addition to gas adsorbing to the surface of the device, occluded gas on the interior, and gas that is discharged through seal materials, gas, moisture and the like that are adsorbed to the contact lens base material being processed is discharged. Thus, keeping the arrival pressure in the device constant prior to processing by plasma polymerization also contributes to reducing the variation in quality between processing lots and within lots, and is desirable from both practical and commercial perspectives. The arrival pressure when drawing a vacuum is preferably equal to or less than 1.35 Pa, more preferably equal to or less than 1.30 Pa. Equal to or less than 1.35 Pa is preferable because the variation in quality (the variation in coating thickness) between processing lots and within lots can be reduced due to the effects of gas adsorbed on the surface of the device and gas adsorbed to the contact lens base material, as mentioned above. To discharge gas within the device to within a certain range in this manner, it suffices to employ a vacuum pump having the ability to reach the targeted degree of vacuum; a hydraulic rotation pump, dry pump, or

any commonly known pump may be employed. Any measuring apparatus capable of measuring the pressure within the prescribed range may be employed to measure the degree of vacuum within the device; examples are diaphragm vacuum gages and Pirani vacuum gages. Further, in the present invention, to uniformly and efficiently process the surfaces (front curved surface and base curved surface) of the contact lens base material being processed, the contact lens base material is preferably placed on a tray for support. Since the interior of the device is in a state of near vacuum, it is desirable to support the contact lens base material through linear contact, such as contact of equal to or greater than 1 percent, with the surface of the contact lens base material to increase the stability of the contact lens base material during processing. Any material that is commonly employed in vacuum devices, such as stainless steel, may be employed as the tray material.

[0015] Once the degree of vacuum within the device has reached the prescribed pressure range, a mixed gas of methane and moist air is introduced into the device, and a base for a coating that is flexible, able to withstand swelling of the lens, and possesses good water wettability and durability is formed on the surface of the lens. The use of moist air in the first step results in a coating film having flexibility on the surface of the lens, and permits the formation of a smooth surface such that cracks and the like do not form in the coating film on the lens surface even after extraction with an organic solvent such as alcohol (referred to as "alcohol extraction", hereinafter) or a water introduction step employing physiological saline or the like. In the present invention, the term "moist air" refers to air containing equal to or greater than 100 ppm of moisture. Moist air in the form of dry air containing equal to or greater than 150 ppm of moisture is preferable, 150 to 1,000 ppm is more preferable, and 150 to 400 ppm is of even greater preference. In this process, moisture content of less than 150 ppm in dry air is undesirable because the crosslinking density in the film formed on the surface of the lens increases, tending to form a rigid film, and cracks and the like tend to develop in swelling steps such as alcohol extraction. Examples of moist air preparation methods are: mixing moisture as an impurity during dry air

manufacturing; mixing moisture by causing dry air to pass through distilled water when introducing dry air into the device; and boiling distilled water in a glass container such as a flask with a rounded bottom, for example, mixing the steam that is obtained with dry air, and then introducing the mixture into the device.

[0016] The mixing ratio of methane to moist air (methane:moist air) employed in the first step is preferably from 50:50 to 70:30 based on volume. A proportion of moist air exceeding this ratio is undesirable because the rate of film formation on the surface of the lens tends to decrease (the processing time increases) and the water wettability following high-pressure steam sterilization that is carried out prior to obtaining the final product sometimes decreases. A proportion of methane exceeding this ratio is undesirable as a film formed on a flexible soft contact lens because the film that is formed on the surface of the lens tends to become rigid, and separation, cracking, and the like of the rigid polymer film occur due to size change caused by swelling. The mixing ratio is more preferably from 55:45 to 65:35.

[0017] In the first step, a mixed gas of methane and moist air can be introduced into the device, or the methane and moist air can be separately introduced to form a mixed gas within the device. In the first step, it is desirable for the gas to be supplied without interruption to the interior of the device, with processing by plasma polymerization being conducted while using a vacuum pump to maintain a constant pressure within the device. The flow rate of the mixed gas of methane and moist air that is introduced into the device is preferably 1.5 to 20 sccm, more preferably 2 to 10 sccm, for a device with an internal volume of about 150 to 700 L, for example.

[0018] The processing by plasma polymerization in the first step is desirably conducted after the gas has been introduced into the device and the pressure within the device has been stabilized. The processing conditions during electrical discharge are suitably selected; for example, a pressure within the device of 4 to 10 Pa, a discharge output of 10 to 80 W, and a power source with low frequency of about 6 to 15 kHz during plasma generation are desirable. Internal electrode-type and external electrode-type devices may be employed, for example. In all cases, known devices

may be employed to be carried out. The duration of the processing by plasma polymerization in the first step can be set taking into account the desired thickness of the film, and may be, for example, 3 to 20 minutes, preferably 4 to 10 minutes.

[0019]     SECOND STEP

In the second step, to the contact lens base material following the aforementioned first step, plasma treatment is conducted in a nonpolymerizable gas atmosphere. In the present invention, the hydrophilic property of the coating can be enhanced by conducting the second step after forming a coating base on the surface of the contact lens base material in the first step. The nonpolymerizable gas employed in the second step has an etching effect, functioning to increase the hydrophilic property of the coating by etching the coating that has been formed on the lens base material in the first step. The second step can be conducted with the same plasma device as the first step, or with a different plasma device. From the perspective of workability and the like, it is desirable to successively conduct the second step following the first step in the same plasma device. In that case, once the processing gas in the device has been discharged following the first step, the nonpolymerizable gas employed in the second step is introduced. In the first step, in the case of a silicone hydrogel, it is possible to increase the water wettability, for example, to the extent that the contact angle of 105 degree in unprocessed state becomes about 50 to 60 degree, in the contact angle measurement by a liquid-drop method employing distilled water. The second step can impart a greater hydrophilic property to the lens surface, and it is possible to increase the contact angle to about 40 degree, in the aforementioned case.

[0020]     The "nonpolymerizable gas" employed in the second step refers to a gas not having the property of depositing on the surface by plasma processing. Such gases can be broadly divided into those that do not contribute to chemical reactions, such as inert gases, and those that contribute to chemical reactions but do not have the property of depositing on the surface. Specific examples are helium, argon, H<sub>2</sub>, O<sub>2</sub>, N<sub>2</sub>, H<sub>2</sub>O, NH<sub>3</sub>, and air. Of these, oxygen, argon, and air are preferred, and oxygen is

of even greater preference. The "air" referred to here can be moist air. The moist air described in the first step above can be employed.

[0021] In the second step as well, it is desirable to conduct the plasma treatment while introducing the gas into the device without interruption and using a vacuum pump to maintain a constant pressure within the device. A pressure within the plasma device of 4 to 10 Pa, a discharge output of 10 to 80 W, and a power source with low frequency of about 6 to 15 kHz during plasma generation are desirable. When the internal volume of the plasma device is about 150 to 700 L, the flow rate of the nonpolymerizable gas is preferably 1.5 to 20 sccm, more preferably 2 to 10 sccm. The duration of the plasma processing in the second step can be set taking into account the etching rate and the like of the processing gas employed in the second step; for example, it can be set 30 seconds to 5 minutes, and preferably 1 to 3 minutes.

[0022] A coating can be formed on the surface of the contact lens base material by the above steps. It suffices for the coating to be formed on at least a portion of the surface of the base material, but it is desirably formed over the entire surface of the base material. The thickness of the coating is important with regard to the water wettability and durability of the lens surface. The thickness of the coating can be measured with an automatic ellipsometer. Instead of directly measuring the thickness of the coating formed on the contact lens base material with an automatic ellipsometer, it is also possible to place both the lens and a silicon wafer on the conveyor tray, measure the thickness of the film that is formed on the silicon wafer, and adopt this film thickness as the thickness of the coating that has been formed on the contact lens base material. The thickness is preferably 90 to 250 Angstroms, more preferably 100 to 200 Angstroms. A coating thickness of equal to or greater than 90 Angstroms is preferable because it affords high coating heat resistance and does not result in a decrease in water wettability following high-pressure steam sterilization. A coating thickness of equal to or less than 250 Angstroms yields high oxygen permeability.

[0023] [Method of manufacturing soft contact lens]

The method of manufacturing a soft contact lens of the present invention is

the method wherein water is introduced into the contact lens material manufactured by the aforementioned method to obtain a soft contact lens.

The water introduction can be conducted by a known method. Specifically, the contact lens material obtained by the aforementioned method can be immersed in physiological saline or a soft contact lens storage solution to introduce water.

[0024] Following processing to introduce water, the soft contact lens can be sterilized by processing with high-pressure steam, for example. The contact lens material can also be subjected to the extraction treatment with an organic solvent prior to the water introduction treatment. This extraction treatment can be conducted by known methods; such treatment can remove unpolymerized monomers, oligomers, and the like remaining in the contact lens material that are biologically undesirable. Since there is a highly flexible coating on the surface of the contact lens material that is obtained by the method of the present invention, it is possible to reduce or prevent separation, cracking and the like of the coating during the above-described extraction treatment and water introduction treatment. Thus, according to the present invention, a soft contact lens of good water wettability and durability in addition to high surface smoothness can be obtained.

## EXAMPLES

[0025] The present invention will be further described below based on examples. However, the present invention is not limited to the following examples.

[0026] 1. Manufacturing contact lens base material

Manufacturing Example A: Synthesis of macromer

To a three-necked flask were charged 8.88 g of isophorone diisocyanate, 0.025 g of a catalyst in the form of dibutyltin dilaurate, and 45 mL of methylene chloride and the mixture was stirred under a nitrogen gas flow. Next, 20 g of *alpha*-butyl-*omega*-[3-(2,2-(dihydroxymethyl)butoxy)propyl]polydimethylsiloxane was precisely weighed out and added dropwise to the flask over about three hours, and the mixture was reacted. When the reaction had progressed for 48 hours at room temperature, another 0.025 g of dibutyltin dilaurate and 23.3 g of polyethylene glycol

monomethacrylate (PE-350) were precisely weighed out and added dropwise to the flask over about 30 minutes. The mixture was covered with aluminum foil and stirred until the absorption band ( $2,260\text{ cm}^{-1}$ ) corresponding to the isocyanate in IR (infrared radiation absorption spectra) analysis disappeared (a reaction of about 48 hours at room temperature). Methylene chloride was further added to the solution, after which the mixture was washed with a large quantity of water, dehydrated, and filtered. The solvent was then distilled off, yielding a macromer with a number average molecular weight of 2,000 (based on polystyrene conversion).

[0027] Manufacturing Example A-1: Manufacturing contact lens base material

About 3.9 g of the siloxane macromonomer obtained in Manufacturing Example A, about 15 g of tris(trimethylsiloxy)-*gamma*-methacryloxypropylsilane, about 11.1 g of N,N-dimethylacrylamide, about 0.003 g of a coloring material in the form of 1-anilino-4-(4-vinylbenzyl)aminoanthraquinone, and about 0.18 g of a polymerization initiator in the form of phenylbis(2,4,6-trimethylbenzoyl)phosphineoxide were mixed for about 20 hours at room temperature to prepare a mixed monomer solution. The mixed monomer solution was then placed in a contact lens-shaped casting mold comprised of polypropylene, the upper and lower molds were combined, and polymerization was completed by irradiation for 20 minutes with radiation in the UV to visible light (380 to 450 nm) at about  $35\text{ mW/cm}^2$ . When polymerization had ended, the polymerization product was removed from the mold, yielding a contact lens base material.

[0028] Manufacturing Example A-2: Manufacturing contact lens

About 4.5 g of the siloxane macromonomer obtained in Manufacturing Example A, about 15 g of tris(trimethylsiloxy)-*gamma*-methacryloxypropylsilane, about 10.5 g of N,N-dimethylacrylamide, and about 0.18 g of phenylbis(2,4,6-trimethylbenzoyl)phosphineoxide were mixed for about 20 hours at room temperature to prepare a mixed monomer solution. The mixed monomer solution was then placed in a contact lens-shaped casting mold comprised of

polypropylene, the upper and lower molds were combined, and polymerization was completed by irradiation for 20 minutes with radiation in the UV to visible light (380 to 450 nm) at about 35 mW/cm<sup>2</sup>. When polymerization had ended, the polymerization product was removed from the mold, yielding a contact lens base material.

[0029] Example 1

The contact lens obtained in Manufacturing Example A-1 was placed in a plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of 1.24 Pa. Next, a mixed gas of methane and moist air (moisture concentration 150 ppm) with a mixing ratio (methane:moist air) of 60:40 (gas flow rate: methane = 1.8 (sccm), moist air = 1.2 (sccm)) was introduced into the device and processing by plasma polymerization was conducted for 6 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was introduced into the device and plasma treatment was conducted for 2 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. The thickness of a film formed on a silicon wafer processed together with the lens was measured at about 200 Angstroms with an automatic ellipsometer.

[0030] Following processing, the lens was immersed for about 5 hours in methanol, replaced with a soft contact lens filler solution, packed into a lens shipping case, and subjected to high-pressure steam sterilization processing at 121°C for 30 minutes. The following methods were employed to observe the surface of the lens obtained and evaluate it for water wettability (contact angle measurement) and the durability (scrubbing durability) of the hydrophilic film on the surface. The results are given in Table 1. As shown in Table 1, the lens obtained in the present example had a smooth, soft surface (Fig. 1), good water wettability and durability, and adequate physical properties as a soft contact lens.

[0031] (a) Inspection of external appearance

After wiping moisture off the surface of the lens, a VH-8000 digital HF

microscope made by Keyence Corp. was used to examine the surface of the lens for scratches and cracks.

[0032] (b) Evaluation of surface water wettability (contact angle measurement)

After wiping moisture off the surface of the lens, the lens was attached on a support and the contact angle was measured by the liquid-drop method with distilled water.

[0033] (c) Scrubbing durability test

The lens was placed in the palm of the hand, a soft contact lens cleaning solution was employed to scrub the front and back sides of the lens, and the lens was thoroughly rinsed with distilled water. This cleaning cycle was repeated, and the method described in (b) was then used to measure the contact angle after 10, 20, and 30 cleaning cycles to evaluate the durability of the hydrophilic film on the surface.

[0035] Example 2

The contact lens obtained in Manufacturing Example A-1 was placed in a plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of 1.24 Pa. Next, a mixed gas of methane and moist air (moisture concentration 150 ppm) with a mixing ratio (methane:moist air) of 60:40 (gas flow rate: methane = 1.8 (sccm), moist air = 1.2 (sccm)) was introduced into the device and processing by plasma polymerization was conducted for 5 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was introduced into the device and plasma treatment was conducted for 2 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. The thickness of a film formed on a silicon wafer processed together with the lens was measured at about 170 Angstroms with an automatic ellipsometer. Subsequent processing and evaluation of the lens after processing were conducted in the same manner as in Example 1. The evaluation results for the lens are given in Table 1. The lens had a smooth, soft surface (Fig. 2), good water wettability and durability, and adequate physical properties as a soft contact lens.

[0036] Example 3

The contact lens obtained in Manufacturing Example A-1 was placed in a plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of 1.28 Pa. Next, a mixed gas of methane and moist air (moisture concentration 150 ppm) with a mixing ratio (methane:moist air) of 60:40 (gas flow rate: methane = 1.8 (sccm), moist air = 1.2 (sccm)) was introduced into the device and processing by plasma polymerization was conducted for 4.5 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was introduced into the device and plasma treatment was conducted for 2 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. The thickness of a film formed on a silicon wafer processed together with the lens was measured at about 150 Angstroms with an automatic ellipsometer. Subsequent processing and evaluation of the lens after processing were conducted in the same manner as in Example 1. The evaluation results for the lens are given in Table 1. The lens had a smooth, soft surface (Fig. 3), good water wettability and durability, and adequate physical properties as a soft contact lens.

[0036] Example 4

The contact lens obtained in Manufacturing Example A-2 was placed in a plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of about 1.25 Pa. Next, a mixed gas of methane and moist air (moisture concentration 250 ppm) with a mixing ratio (methane:moist air) of 67:33 (gas flow rate: methane = 2 (sccm), moist air = 1 (sccm)) was introduced into the device and processing by plasma polymerization was conducted for 6 minutes at a pressure of 4 Pa, a frequency of 15 kHz, and a discharge power of 44 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was introduced into the device and plasma treatment was conducted for 1 minute at a pressure of 4 Pa, a frequency of 15 kHz, and a discharge power of 44 W. The thickness of a film formed on a silicon wafer processed together with the lens was

measured at about 220 Angstroms with an automatic ellipsometer. Subsequent processing and evaluation of the lens after processing were conducted in the same manner as in Example 1. The evaluation results for the lens are given in Table 1. The lens had a smooth, soft surface, good water wettability and durability, and thus it was adequately sufficient as a soft contact lens.

[0037] Example 5

The contact lens obtained in Manufacturing Example A-1 was placed in a plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of about 1.24 Pa. Next, a mixed gas of methane and moist air (moisture concentration 300 ppm) with a mixing ratio (methane:moist air) of 60:40 (gas flow rate: methane = 1.8 (sccm), moist air = 1.2 (sccm)) was introduced into the device and processing by plasma polymerization was conducted for 6 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was introduced into the device and plasma treatment was conducted for 2 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. The thickness of a film formed on a silicon wafer processed together with the lens was measured at about 190 Angstroms with an automatic ellipsometer. Subsequent processing and evaluation of the lens after processing were conducted in the same manner as in Example 1. The evaluation results for the lens are given in Table 1. The lens had a smooth, soft surface (Fig. 4), good water wettability and durability, and adequate physical properties as a soft contact lens.

[0038] Example 6

The contact lens obtained in Manufacturing Example A-1 was placed in a plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of about 1.24 Pa. Next, a mixed gas of methane and moist air (moisture concentration 300 ppm) with a mixing ratio (methane:moist air) of 60:40 (gas flow rate: methane = 1.8 (sccm), moist air = 1.2 (sccm)) was introduced into the device and processing by plasma polymerization was conducted for 4 minutes

at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was introduced into the device and plasma treatment was conducted for 2 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. The thickness of a film formed on a silicon wafer processed together with the lens was measured at about 120 Angstroms with an automatic ellipsometer. Subsequent processing and evaluation of the lens after processing were conducted in the same manner as in Example 1. The evaluation results for the lens are given in Table 1. The lens had a smooth, soft surface, good water wettability and durability, and adequate physical properties as a soft contact lens.

[0039] Example 7

The contact lens obtained in Manufacturing Example A-1 was placed in a plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of about 1.24 Pa. Next, a mixed gas of methane and moist air (moisture concentration 300 ppm) with a mixing ratio (methane:moist air) of 53:47 (gas flow rate: methane = 1.6 (sccm), moist air = 1.4 (sccm)) was introduced into the device and processing by plasma polymerization was conducted for 4.5 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was introduced into the device and plasma treatment was conducted for 2 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. The thickness of a film formed on a silicon wafer processed together with the lens was measured at about 100 Angstroms with an automatic ellipsometer. Subsequent processing and evaluation of the lens after processing were conducted in the same manner as in Example 1. The evaluation results for the lens are given in Table 1. The lens had a smooth, soft surface, good water wettability and durability, and adequate physical properties as a soft contact lens.

[0040] Comparative Example 1 (Processing with hydrocarbon alone)

The contact lens obtained in Manufacturing Example A-2 was placed in a

plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of about 1.23 Pa. Next, 3 (sccm) of methane was introduced into the device and processing by plasma polymerization was conducted for 6 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 45 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was introduced into the device and plasma treatment was conducted for 2 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 45 W. The thickness of a film formed on a silicon wafer processed together with the lens was measured at about 200 Angstroms with an automatic ellipsometer. Subsequent processing and evaluation of the lens after processing were conducted in the same manner as in Example 1. The evaluation results for the lens are given in Table 1. Observation of the lens surface revealed cracks, rendering the lens unsuitable as a contact lens (Fig. 5). It was thought that the film formed with methane was formed as a rigid coating on the lens, and the coating was unable to withstand the degree of swelling of the lens in the alcohol extraction and water introduction steps, resulting in cracks.

[0041] Comparative Example 2 (Processing with a mixed gas of saturated hydrocarbon and oxygen)

The contact lens obtained in Manufacturing Example A-1 was placed in a plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of 1.27 Pa. Next, a mixed gas was introduced into the device at flow rates of 1 (sccm) of methane and 2 (sccm) of oxygen and processing by plasma polymerization was conducted for 7 minutes at a pressure of 4 Pa, a frequency of 15 kHz, and a discharge power of 35 W. The thickness of a film formed on a silicon wafer processed together with the lens was measured at about 70 Angstroms with an automatic ellipsometer. Subsequent processing and evaluation of the lens after processing were conducted in the same manner as in Example 1. The evaluation results for the lens are given in Table 1. Since the film formed on the lens surface was thin, water wettability following high-pressure steam sterilization was

poor, and the lens was unsuitable for use as a contact lens.

[0042] Comparative Example 3 (Processing with a mixed gas of methane and dry air)

The contact lens obtained in Manufacturing Example A-1 was placed in a plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of about 1.23 Pa. Next, a mixed gas of methane and air (moisture concentration 0.53 ppm or less) with a mixing ratio (methane:air) of 60:40 (gas flow rate: methane = 1.8 (sccm), air = 1.2 (sccm)) was introduced into the device and processing by plasma polymerization was conducted for 7 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 35 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was introduced into the device and plasma treatment was conducted for 2 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was introduced into the device and plasma treatment was conducted for 2 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. The thickness of a film formed on a silicon wafer processed together with the lens was measured at about 220 Angstroms with an automatic ellipsometer. Subsequent processing and evaluation of the lens after processing were conducted in the same manner as in Example 1. The evaluation results for the lens are given in Table 1. Observation of the lens surface revealed cracks, rendering the lens unsuitable as a contact lens (Fig. 6). It was thought that the film formed with the mixed gas of methane and dry air was formed as a rigid coating on the lens, and the coating was unable to withstand the degree of swelling of the lens in the alcohol extraction and water introduction steps, resulting in cracks. A comparison with Example 1 shows that the processing gas employed in the first step differed, being moist air in the one case and dry air in the other, revealing that the admixing of moisture produced a major difference in the results.

[0043] Comparative Example 4 (Processing with a mixed gas of methane and dry

air)

The contact lens obtained in Manufacturing Example A-1 was placed in a plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of about 1.25 Pa. Next, a mixed gas of methane and air (moisture concentration of equal to or less than 0.53 ppm) with a mixing ratio (methane:air) of 67:33 (gas flow rate: methane = 2 (sccm), air = 1 (sccm)) was introduced into the device and processing by plasma polymerization was conducted for 6 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 35 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was introduced into the device and plasma treatment was conducted for 2 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. The thickness of a film formed on a silicon wafer processed together with the lens was measured at about 150 Angstroms with an automatic ellipsometer. Subsequent processing method of the lens after processing was conducted in the same manner as in Example 1. The evaluation results for the lens are given in Table 1. Observation of the lens surface revealed cracks, rendering the lens unsuitable as a contact lens (Fig. 7). A comparison with Example 1 shows that the processing gas employed in the first step differed, being moist air in the one case and dry air in the other, revealing that admixing of moisture produced in a major difference in the results.

[0044] Comparative Example 5 (Processing with a mixed gas of methane and moist air (moisture concentration 50 ppm))

The contact lens obtained in Manufacturing Example A-1 was placed in a plasma polymerization device (with an internal volume of about 170 L) and a vacuum was drawn to an arrival pressure of about 1.23 Pa. Next, a mixed gas of methane and air (moisture concentration 50 ppm) with a mixing ratio (methane:air) of 60:40 (gas flow rate: methane = 1.8 (sccm), moist air = 1.2 (sccm)) was introduced into the device and processing by plasma polymerization was conducted for 7 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 35 W. Next, once the processing gas had been discharged from the device, 3 (sccm) of oxygen was

introduced into the device and plasma treatment was conducted for 2 minutes at a pressure of 4 Pa, a frequency of 10 kHz, and a discharge power of 40 W. The thickness of a film formed on a silicon wafer processed together with the lens was measured at about 195 Angstroms with an automatic ellipsometer. Subsequent processing and evaluation of the lens after processing were conducted in the same manner as in Example 1. The evaluation results for the lens are given in Table 1. Observation of the lens surface revealed cracks, rendering the lens unsuitable as a contact lens (Fig. 8). A comparison of Comparative Examples 3 and 4, in which processing was conducted with a mixed gas of methane and dry air, revealed substantial improvement in the degree of cracking, but a moisture concentration of 50 ppm was considered inadequate.

[0045] As shown in Table 1 and Figs. 5 to 8, when the coating was formed with a hydrocarbon (such as methane) essentially containing no nitrogen or oxygen; when the coating was formed with a mixed gas of methane and dry air; and when the coating was formed with a mixed gas of methane and air containing a moisture content of about 50 ppm, as in Comparative Examples 1, 3 to 5, cracks appeared in the lens surface, rendering the lens unsuitable as a contact lens. When the coating was formed using a mixed gas of a saturated hydrocarbon and oxygen, as in Comparative Example 2, the film formed by the saturated hydrocarbon was etched away by the oxygen plasma, precluding the formation of a film of adequate thickness on the lens surface and imparting insufficient water wettability.

By contrast, as shown in the Examples, lenses that were coated with a mixed gas of methane and moist air had surfaces of good smoothness, water wettability, and durability, and were satisfactory as contact lenses.

[0046] [Table 1]

| Example  | External appearance of lens | Film thickness (Angstroms) | Water wettability (Contact angle (Degree)) | Scrubbing durability (Contact angle (degree)) |          |          |
|----------|-----------------------------|----------------------------|--------------------------------------------|-----------------------------------------------|----------|----------|
|          |                             |                            |                                            | 10 times                                      | 20 times | 30 times |
| 1        | Good                        | 200                        | 44                                         | 45                                            | 44       | 46       |
| 2        | Good                        | 170                        | 43                                         | 40                                            | 38       | 40       |
| 3        | Good                        | 150                        | 45                                         | 45                                            | 45       | 45       |
| 4        | Good                        | 220                        | 43                                         | 45                                            | 42       | 40       |
| 5        | Good                        | 190                        | 41                                         | 44                                            | 44       | 45       |
| 6        | Good                        | 120                        | 44                                         | 42                                            | 44       | 45       |
| 7        | Good                        | 100                        | 48                                         | 47                                            | 48       | 49       |
| Comp.Ex. |                             |                            |                                            |                                               |          |          |
| 1        | Cracks                      | 200                        | —                                          | —                                             | —        | —        |
| 2        | Good                        | 70                         | 89                                         | —                                             | —        | —        |
| 3        | Cracks                      | 220                        | —                                          | —                                             | —        | —        |
| 4        | Cracks                      | 150                        | —                                          | —                                             | —        | —        |
| 5        | Cracks                      | 195                        | —                                          | —                                             | —        | —        |

## INDUSTRIAL APPLICABILITY

[0047] The present invention can provide a soft contact lens, and more particularly, a soft contact lens comprised of a hydrogel that contains silicone as a lens component, having a surface that is smooth and has good water wettability and durability.

## BRIEF DESCRIPTION OF THE DRAWINGS

[0048]

[Fig. 1] Fig. 1 shows a microscope photograph (175X) of the surface of the lens obtained in Example 1.

[Fig. 2] Fig. 2 shows a microscope photograph (175X) of the surface of the lens obtained in Example 2.

[Fig. 3] Fig. 3 shows a microscope photograph (175X) of the surface of the lens obtained in Example 3.

[Fig. 4] Fig. 4 shows a microscope photograph (175X) of the surface of the lens obtained in Example 5.

[Fig. 5] Fig. 5 shows a microscope photograph (175X) of the surface of the lens obtained in Comparative Example 1.

[Fig. 6] Fig. 6 shows a microscope photograph (175X) of the surface of the lens obtained in Comparative Example 3.

[Fig. 7] Fig. 7 shows a microscope photograph (175X) of the surface of the lens obtained in Comparative Example 4.

[Fig. 8] Fig. 8 shows a microscope photograph (175X) of the surface of the lens obtained in Comparative Example 5.

## CLAIMS

[1] A method of manufacturing a contact lens material comprising a contact lens base material and a coating formed on at least a portion of a surface of said base material, wherein

said contact lens base material is processed by plasma polymerization in a mixed gas atmosphere of methane and moist air, and then treated with plasma in a nonpolymerizable gas atmosphere to form said coating.

[2] The method of manufacturing according to claim 1, wherein said contact lens base material comprises silicone.

[3] The method of manufacturing according to claim 1 or 2, wherein said moist air comprises equal to or greater than 150 ppm of moisture.

[4] The method of manufacturing according to any of claims 1 to 3, wherein the mixing rate of methane and moisture air, methane: moist air, in said mixed gas ranges from 50:50 to 70:30.

[5] The method of manufacturing according to any of claims 1 to 4, wherein said nonpolymerizable gas is at least one selected from the group consisting of oxygen, argon, and air.

[6] A method of manufacturing a soft contact lens, wherein

water is introduced into the contact lens material manufactured by the method according to any of claims 1 to 5 to obtain a soft contact lens.

[7] The method of manufacturing according to claim 6, which comprises subjecting the contact lens material prior to said water introduction to extraction treatment with an organic solvent.