

DESCRIPTION

PATTERN FORMING METHOD, AND RESIST COMPOSITION, DEVELOPER AND RINSING SOLUTION USED IN THE PATTERN FORMING METHOD

TECHNICAL FIELD

The present invention relates to a pattern forming method for use in the process of producing a semiconductor such as IC, in the production of a circuit board for liquid crystal, thermal head and the like, and in the lithography process of other photo-applications; a resist composition for use in the pattern forming method; a developer for negative tone development used in the pattern forming method; and a rinsing solution for negative tone development used in the pattern forming method. More specifically, the present invention relates to a pattern forming method suitable for exposure with an ArF exposure apparatus using a light source that emits far ultraviolet light at a wavelength of 300 nm or less or with an immersion-type projection exposure apparatus; a resist composition for use in the pattern forming method; a developer for negative tone development used in the pattern forming method; and a rinsing solution for negative tone development used in the pattern forming method.

BACKGROUND ART

Since the advent of a resist for KrF excimer laser (248 nm), an image forming method called chemical amplification is used as an image forming method for a resist so as to compensate for sensitivity reduction caused by light absorption. For example, the image forming method by positive tone chemical amplification is an image forming method of decomposing an acid generator in the exposed area upon exposure to produce an acid, converting an alkali-insoluble group into an alkali-soluble group by using the generated acid as a reaction catalyst in the baking after exposure (PEB: Post Exposure Bake), and removing the exposed area by alkali development.

Along with miniaturization of a semiconductor device, there is becoming shorter the wavelength of the exposure light source and higher the numerical aperture (higher NA) of the projection lens, and an exposure machine using an ArF excimer laser having a wavelength of 193 nm as a light source has been so far developed. As commonly well known, these features can be expressed by the following formulae:

$$(\text{Resolution}) = k_1 \bullet (\lambda / \text{NA})$$

$$(\text{Depth of focus}) = \pm k_2 \bullet \lambda / \text{NA}^2$$

wherein λ is the wavelength of the exposure light source, NA is the numerical aperture of the

projection lens, and k_1 and k_2 are coefficients related to the process.

A so-called immersion method of filling a high refractive-index liquid (hereinafter sometimes referred to as an "immersion liquid") between the projection lens and the sample has been conventionally advocated as a technique for raising the resolution.

As for the "effect of immersion", assuming that $NA_0 = \sin\theta$, the above-described resolution and depth of focus in the immersion can be expressed by the following formulae:

$$(\text{Resolution}) = k_1 \bullet (\lambda_0/n)/NA_0$$

$$(\text{Depth of focus}) = \pm k_2 \bullet (\lambda_0/n)/NA_0^2$$

wherein λ_0 is the wavelength of exposure light in air, n is the refractive index of the immersion liquid based on air, and θ is the convergence half-angle of beam.

That is, the effect of immersion is equal to use of an exposure wavelength of $1/n$. In other words, in the case of a projection optical system with the same NA, the depth of focus can be made n times larger by the immersion. This is effective for all pattern profiles and can be combined with the super-resolution technology under study at present, such as phase-shift method and modified illumination method.

A double exposure technology or a double patterning technology is being advocated as a technique for more enhancing the resolution. This is to make small k_1 in the above-described formula of resolution and is positioned as a resolution-increasing technology.

In conventional pattern formation of an electronic device such as semiconductor device, a mask or reticle pattern in a size of 4 to 5 times larger than the pattern intended to form is reduced and transferred on an exposure target such as wafer by using a reduction projection exposure apparatus.

However, the dimensional miniaturization brings about a problem that in the conventional exposure system, lights irradiated on adjacent patterns interfere with each other to decrease the optical contrast. Therefore, in such technology, it is devised to divide the exposure mask design into two or more designs and synthesize an image by independently exposing these masks. In this double exposure system where the exposure mask design is divided, the image of the design must be again synthesized on an exposure target (wafer) and therefore, the division of the mask design must be devised so that the pattern on the reticle can be faithfully reproduced on the exposure target.

Studies of applying the effect of these double exposure systems to the transfer of a fine image pattern of a semiconductor device are introduced, for example, in Patent Document 1 (JP-A-2006-156422 (the term "JP-A" as used herein means an "unexamined published Japanese patent application")).

Also, the recent progress of double exposure technology is reported, for example, in Non-Patent Document 1 (SPIE Proc 5754, 1508 (2005)), Non-Patent Document 2 (SPIE Proc

5377, 1315 (2004)), and Non-Patent Document 3 (SPIE Proc 61531K-1 (2006)).

However, in these double exposure systems, the pattern formation needs to be performed in the vicinity of resolution limit of the resist and this incurs a problem that sufficient exposure margin or depth of focus cannot be obtained.

In other words, if a pattern forming process described, for example, in Patent Document 2 (JP-A-2001-109154) where a resist composition containing a resin capable of increasing the polarity upon exposure is coated on a substrate and the resist film is exposed and developed to dissolve the exposed area with an alkali developer, or a pattern forming process described, for example, in Patent Document 3 (JP-A-2003-76019) where a resist composition containing a resin capable of increasing the molecular weight upon exposure is coated on a substrate and the resist film is exposed and developed to dissolve the unexposed area with an alkali developer, is applied to a double exposure process, a sufficiently high resolving performance cannot be obtained.

With respect to the developer for g-line, i-line, KrF, ArF, EB or EUV lithography, an aqueous alkali developer of 2.38 mass% TMAH (tetramethylammonium hydroxide) is being used at present as a general-purpose developer.

Other than the above-described developer, for example, Patent Document 4 (JP-A-2001-215731) describes a developer for developing a resist material containing a copolymer of a styrene-based monomer and an acryl-based monomer and dissolving the exposed portion, where the developer contains an aliphatic linear ether-based solvent or aromatic ether-based solvent and a ketone-based solvent having a carbon number of 5 or more; Patent Document 5 (JP-A-2006-227174) describes a developer for developing a resist material capable of decreasing the molecular weight as a result of breakage of the polymer chain upon irradiation with radiation, thereby dissolving the exposed portion, where the developer has at least two or more acetic acid groups, ketone groups, ether groups or phenyl groups and has a molecular weight of 150 or more; and Patent Document 6 (JP-A-6-194847) describes a developer for developing a resist material mainly composed of a photosensitive polyhydroxy ether resin obtained by the reaction of a polyhydroxy ether resin with a diglycidyl (meth)acrylate, where the developer is an aromatic compound having a carbon number of 6 to 12 or a mixed solvent containing 50 mass% or more of an aromatic compound having a carbon number of 6 to 12.

However, these combinations of the resist composition and the developer mentioned above merely provide a system of forming a pattern by combining a specific resist composition with a high-polarity alkali developer or a developer containing a low-polarity organic solvent.

That is, as shown in Fig. 1, in the case of a positive tone system (a combination of a resist composition and a positive tone developer), a material system of performing the pattern

formation by selectively dissolving and removing a region having strong light irradiation intensity out of an optical aerial image (light intensity distribution) is merely provided. On the other hand, as for the combination of a negative tone system (a resist composition and a negative tone developer), a material system of performing the pattern formation by selectively dissolving and removing a region having a weak light irradiation intensity is merely provided.

The term "positive tone developer" as used herein indicates a developer that selectively dissolves and removes the exposed area not lower than a predetermined threshold value shown by a solid line in Fig. 1, and the "negative tone developer" indicates a developer that selectively dissolves and removes the exposed area not higher than the predetermined threshold value. A development step using a positive tone developer is called a positive tone development (sometimes referred to as a positive tone development step), and a development step using a negative tone developer is called a negative tone development (sometimes referred to as a negative tone development step).

Patent Document 7 (JP-A-2000-199953) describes a double development technology as the double patterning technology for improving the resolution. In this case, an image forming method by chemical amplification in general is utilized, and by making use of a property that the polarity of a resin in a resist composition when exposed becomes a high polarity in a region irradiated with a high light intensity and becomes a low polarity in a region irradiated with a low light intensity, positive tone development is performed by dissolving a high exposure region of the resist film with a high-polarity developer, while negative tone development is performed by dissolving a low exposure region with a low-polarity developer. More specifically, the region not lower than an exposure dose of E2 in Fig. 2 is dissolved using an aqueous alkali solution as the positive tone developer, the region not higher than an exposure dose of E1 is dissolved using a specific organic solvent as the negative tone developer, and, as shown in Fig. 2, the region with a medium exposure dose (E2 to E1) is allowed to remain without being developed, whereby an L/S pattern 3 with a pitch half the pitch of the exposure mask 2 pattern is formed on a wafer 4.

Patent Document 1: JP-A-2006-156422

Patent Document 2: JP-A-2001-109154

Patent Document 3: JP-A-2003-76019

Patent Document 4: JP-A-2001-215731

Patent Document 5: JP-A-2006-227174

Patent Document 6: JP-A-6-194847

Patent Document 7: JP-A-2000-199953

Non-Patent Document 1: SPIE Proc 5754, 1508 (2005)

Non-Patent Document 2: SPIE Proc 5377, 1315 (2004)

DISCLOSURE OF THE INVENTION

PROBLEMS THAT THE INVENTION IS TO SOLVE

In the conventional double development technology, a tert-butyl group is employed as an acid-decomposable group in the resin contained in the resist composition and therefore, the resin is insufficient to bring out a polarity change large enough to cause a difference in the dissolution characteristics by a chemical amplification reaction associated with the exposure.

Furthermore, since a resin having a styrene skeleton is used as the resin in the resist composition, the polarity in the low exposure region of the resist film becomes high and this causes a problem that the development rate at the development using a negative tone developer is low and the developability when using a negative tone developer is bad.

An object of the present invention is to solve these problems and provide a pattern forming method for stably forming a high-precision fine pattern particularly by reducing the line edge roughness and raising the dimensional uniformity of the pattern so as to produce a highly integrated electronic device with high precision.

MEANS FOR SOLVING THE PROBLEMS

The present invention has the following constructions, and the object of the present invention can be attained by these constructions.

(1) A pattern forming method, comprising:

(i) a step of applying a resist composition whose solubility in a positive tone developer increases and solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation and which contains a resin having an alicyclic hydrocarbon structure and a dispersity of 1.7 or less and being capable of increasing a polarity of the resin by an action of an acid;

(ii) an exposure step; and

(iv) a development step using a negative tone developer.

(2) A pattern forming method, comprising:

(i) a step of applying a resist composition whose solubility in a positive tone developer increases and solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation and which contains a resin having an alicyclic hydrocarbon structure and a weight average molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;

(ii) an exposure step; and

(iv) a development step using a negative tone developer.

- (3) A pattern forming method, comprising:
- (i) a step of applying a resist composition whose solubility in a positive tone developer increases and solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation and which contains a resin having an alicyclic hydrocarbon structure, a dispersity of 1.7 or less and a weight average molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;
 - (ii) an exposure step; and
 - (iv) a development step using a negative tone developer.
- (4) The pattern forming method according to any one of (1) to (3) above, wherein (iv) the development step using a negative tone developer is a step performed using a developer containing at least one kind of a solvent selected from organic solvents and having a vapor pressure of 5 kPa or less at 20°C.
- (5) The pattern forming method as described in any one of (1) to (4) above, which further comprises:
- (vi) a washing step using a rinsing solution containing an organic solvent.
- (6) The pattern forming method as described in (5) above, wherein the rinsing solution containing an organic solvent is a rinsing solution having a vapor pressure of 0.1 kPa or more at 20°C.
- (7) The pattern forming method as described in any one of (1) to (6), which further comprises:
- (iii) a development step using a positive tone developer.
- (8) A resist composition for negative tone development, comprising:
- (a1) a resin having an alicyclic hydrocarbon structure and a dispersity of 1.7 or less and being capable of increasing a polarity of the resin by an action of an acid;
 - (B) a photo-acid generator; and
 - (C) a solvent.
- (9) A resist composition for negative tone development, comprising:
- (a2) a resin having an alicyclic hydrocarbon structure and a weight average molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;
 - (B) a photo-acid generator; and
 - (C) a solvent.
- (10) A resist composition for negative tone development, comprising:
- (a3) a resin having an alicyclic hydrocarbon structure, a dispersity of 1.7 or less and a weight average molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;

- (B) a photo-acid generator; and
- (C) a solvent.
- (11) A resist composition for multiple development, comprising:
 - (a1) a resin having an alicyclic hydrocarbon structure and a dispersity of 1.7 or less and being capable of increasing a polarity of the resin by an action of an acid;
 - (B) a photo-acid generator; and
 - (C) a solvent.
- (12) A resist composition for multiple development, comprising:
 - (a2) a resin having an alicyclic hydrocarbon structure and a weight average molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;
 - (B) a photo-acid generator; and
 - (C) a solvent.
- (13) A resist composition for multiple development, comprising:
 - (a3) a resin having an alicyclic hydrocarbon structure, a dispersity of 1.7 or less and a weight average molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;
 - (B) a photo-acid generator; and
 - (C) a solvent.
- (14) A developer for negative tone development, which is used in the resist composition described in any one of (8) to (13) above, the developer comprising an organic solvent and having a vapor pressure of 5 kPa or less at 20°C.
- (15) A rinsing solution for negative tone development, which is used in the pattern forming method described in any one of (8) to (13) above, the rinsing solution comprising an organic solvent and having a vapor pressure of 0.1 kPa or more at 20°C.

ADVANTAGE OF THE INVENTION

According to the present invention, a method of stably forming a high-precision fine pattern with reduced line edge roughness and high dimensional uniformity, a resist composition for negative tone development or multiple development used in the method, a developer for negative tone development used in the method, and a rinsing solution for negative tone development used in the method can be provided.

BRIEF DESCRIPTION OF THE DRAWING

Fig. 1 is a schematic view showing the relationship among positive tone development, negative tone development and exposure dose.

Fig. 2 is a schematic view showing the pattern forming method using positive tone development and negative tone development in combination.

Fig. 3 is a schematic view showing the relationship among positive tone development, negative tone development and exposure dose.

Fig. 4 is a graph showing the relationship between exposure dose and residual film curve when a positive tone developer or a negative tone developer is used.

Fig. 5 is a schematic view showing the relationship among positive tone development, negative tone development and exposure dose.

Fig. 6 is a schematic view showing the relationship among positive tone development, negative tone development and exposure dose in the method of the present invention.

Fig. 7 is a schematic view showing the relationship among positive tone development, negative tone development and exposure dose.

Fig. 8 is a view showing the spatial intensity distribution of an optical image.

Fig. 9 is a schematic view showing the relationship among positive tone development, threshold value (a) and light intensity.

Fig. 10 is a schematic view showing the spatial intensity distribution of an optical image.

Fig. 11 is a schematic view showing the relationship among negative tone development, threshold value (b) and light intensity.

BEST MODE FOR CARRYING OUT THE INVENTION

The best mode for carrying out the present invention is described below.

Incidentally, in the present invention, when a group (atomic group) is denoted without specifying whether substituted or unsubstituted, the group includes both a group having no substituent and a group having a substituent. For example, an "alkyl group" includes not only an alkyl group having no substituent (unsubstituted alkyl group) but also an alkyl group having a substituent (substituted alkyl group).

First, the terms used in the context of the present invention are described. The pattern forming system includes a positive tone system and a negative tone system and in either system, a change in the solubility of a resist film in a developer due to a chemical reaction triggered by light irradiation is utilized. A system where the light-irradiated part dissolves in a developer is called a positive tone system, and a system where the light-unirradiated part dissolves in a developer is called a negative tone system. The developer used here includes two types of developers, that is, a positive tone developer and a negative tone developer. The positive tone developer is a developer that selectively dissolves and removes the exposed area not lower than a predetermined threshold value shown in Fig. 1,

and the negative tone developer is a developer that selectively dissolves and removes an exposed area not higher than the above-described threshold value. A development step using a positive tone developer is called a positive tone development (sometimes referred to as a positive tone development step), and a development step using a negative tone developer is called a negative tone development (sometimes referred to as a negative tone development step).

The present invention provides, as a technique for raising the resolution, a new pattern forming method by a combination of a developer (negative tone developer) for selectively dissolving and removing the exposed area not higher than a predetermined threshold value (b) as shown in Fig. 3, with a resist composition for negative tone development, which contains a resin that increases the polarity by the action of an acid and forms a film whose solubility in a positive tone developer (preferably an alkali developer) increases and solubility in a negative tone developer (preferably an organic developer) decreases upon irradiation with an actinic ray or radiation.

The present invention provides, as a technique for raising the resolution, a new pattern forming method preferably by a combination of a developer (positive tone developer) for selectively dissolving and removing the exposed area not lower than a predetermined threshold value (a) and a developer (negative tone developer) for selectively dissolving and removing the exposed area not higher than a predetermined threshold value (b), with a resist composition for multiple development, which forms a film whose solubility in a positive tone developer (preferably an alkali developer) increases and solubility in a negative tone developer (preferably an organic developer) decreases upon irradiation with an actinic ray or radiation.

That is, as shown in Fig. 3, when a pattern element on an exposure mask is projected on a wafer by light irradiation, the region having a strong light irradiation intensity (the exposed area not lower than a predetermined threshold value (a)) is dissolved and removed using a positive tone developer and the region having a weak light irradiation intensity (the exposed area not higher than a predetermined threshold value (b)) is dissolved and removed using a negative tone developer, whereby a pattern with resolution as high as 2 times the frequency of the optical spatial image (light intensity distribution) can be obtained. Also, in the method of the present invention, the design of the exposure mask need not be divided.

As regards the resist composition for multiple development when performing those two or more development operations at the same time, a resist composition for negative tone development may be used as it is.

The pattern forming process necessary for practicing the present invention comprises the following steps:

- (i) a step of coating a substrate with a resist composition whose solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation,
- (ii) an exposure step, and
- (iv) a step of developing the resist film with a negative tone developer.

The pattern forming method of the present invention preferably further comprises (vi) a step of washing the resist film with a rinsing solution for negative tone development.

The pattern forming method of the present invention preferably further comprises (iii) a step of developing the resist film with a positive tone developer.

The pattern forming method of the present invention preferably comprises (v) a heating step after the exposure step (ii).

In the pattern forming method of the present invention, the exposure step (ii) may be performed a plurality of times.

In the pattern forming method of the present invention, the heating step (v) may be performed a plurality of times.

In practicing the present invention, (a) a resist composition whose solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation and which contains a resin having an alicyclic hydrocarbon structure and a dispersity (molecular weight distribution, Mw/Mn) of 1.7 or less and/or a weight average molecular weight of 6,000 or less and being capable of increasing the polarity by the action of an acid, and (b) a negative tone developer (preferably an organic developer) are necessary.

In practicing the present invention, (c) a rinsing solution for negative tone development is preferably further used.

In practicing the present invention, (d) a positive tone developer (preferably an alkali developer) is preferably further used.

In the present invention, a pattern forming process using two kinds of developers, that is, a positive tone developer and a negative tone developer, is preferred. In this case, the order of developments is not particularly limited, but it is preferred to perform development by using a positive tone developer or a negative tone developer after first exposure and then perform negative or positive tone development by using a developer different from that in the first development. After the negative tone development, the resist film is preferably washed with an organic solvent-containing rinsing solution for negative tone development.

The pattern forming system includes a positive tone system and a negative tone system and in either system, a change in the solubility of a resist film in a developer due to a chemical reaction triggered by light irradiation is utilized. In general, a system where the light-irradiated part dissolves in a developer is called a positive tone system, and a system where the light-unirradiated part dissolves in a developer is called a negative tone system.

The positive tone resist utilizes a chemical reaction such as polarity conversion for enhancing the solubility in a developer, and the negative tone resist utilizes bond formation between molecules, such as crosslinking reaction or polymerization reaction.

Since the advent of a resist for KrF excimer laser (248 nm), an image forming method called chemical amplification is used as an image forming method for a resist so as to compensate for sensitivity reduction caused by light absorption. For example, the image forming method by positive tone chemical amplification is an image forming method of decomposing an acid generator in the exposed area upon exposure to generate an acid, converting an alkali-insoluble group into an alkali-soluble group by using the generated acid as a reaction catalyst in the baking after exposure (PEB: Post Exposure Bake), and removing the exposed area by alkali development.

In the present invention, one positive tone resist composition (a) acts as a positive tone resist for a positive tone developer and at the same time, acts as a negative tone resist for a negative tone developer.

In the present invention, an alkali developer (aqueous) can be used as the positive tone developer, and an organic developer containing an organic solvent can be used as the negative tone developer.

Also, the resist composition (a) is a "resin composition that forms a film capable of increasing the polarity as a result of a chemical reaction triggered by exposure to irradiation".

In conventionally employed negative tone image-forming systems, a material system utilizing a mechanism of increasing the molecular weight exclusively by the bonding between molecules and decreasing the solubility in a developer has been proposed. However, it has been difficult for the image forming mechanism utilizing a change in the molecular weight to establish a system of allowing one resist material system to act as a positive tone resist for one developer and act as a negative tone resist for another developer.

In the present invention, the resist composition (a) not only causes a decrease of the solubility in a negative tone developer due to a polarity conversion reaction of the polymer side chain but also brings about both an increase of the solubility in an alkali developer and a decrease of the solubility in an organic developer particularly by virtue of a specific chemical reaction (the polarity conversion reaction of the polymer side chain) at the same time.

In the present invention, the matter of importance is to control the "threshold value" of exposure dose (the exposure dose for solubilizing or insolubilizing the film by a developer in the light-irradiated region). The "threshold value" is the minimum exposure dose for solubilizing the film by a positive tone developer and the maximum exposure dose for insolubilizing the film by a negative tone developer so as to obtain a desired line width at the time of performing pattern formation.

The "threshold value" can be determined as follows.

That is, the "threshold value" is the maximum exposure dose for solubilizing the film by a positive tone developer and the minimum exposure dose for insolubilizing the film by a negative tone developer so as to obtain a desired line width at the time of performing pattern formation.

More strictly, the threshold value is defined as follows.

The residual film ratio of the resist film to the exposure dose is measured and at this time, as shown in Fig. 4, the minimum exposure dose giving a residual film ratio of 0% for the positive tone developer is designated as a threshold value (a) and the minimum exposure dose giving a residual film ratio of 100% for the negative tone developer is designated as a threshold value (b).

For example, as shown in Fig. 5, the threshold value (a) of the exposure dose for solubilizing the film by a positive tone developer is set to be higher than the threshold value (b) for solubilizing the film by a negative tone developer, whereby pattern formation can be achieved by one exposure operation. That is, as shown in Fig. 6, after a resist is coated on a wafer and exposed, the portion not lower than the threshold value (a) of the exposure dose is dissolved with a positive tone developer and then, the region not higher than the threshold value (b) of the exposure dose is dissolved with a negative tone developer, whereby pattern formation can be performed by one exposure operation. In this case, as for the order of the development with a positive tone developer and the development with a negative tone developer, either development may be performed earlier. After the negative tone development, when the resist film is washed with a rinsing solution containing an organic solvent, more successful pattern formation can be achieved.

The method for controlling the threshold value includes a method of controlling the material-related parameters of the resist composition (a) and the developer or controlling the parameters related to the process.

As for the material-related parameter, control of various physical values related to solubility of the resist composition (a) in the developer and the organic solvent, such as SP value (solubility parameter) and LogP value, is effective. Specific examples of the parameter include, for the polymer contained in the resist composition (a), the average molecular weight, the molecular weight dispersity, the monomer compositional ratio, the polarity of monomer, the monomer sequence, the polymer blend and the addition of low molecular additive, and for the developer, include the concentration of developer, the addition of low molecular additive and the addition of surfactant.

Also, specific examples of the process-related parameter include the film formation temperature, the film formation time, the temperature and time of post-heating after exposure,

the temperature at development, the development time, the nozzle system (puddle method) of developing apparatus, and the rinsing method after development.

Accordingly, for obtaining a good pattern in the pattern forming method using negative tone development as well as in the pattern forming method by multiple development using negative tone development and positive tone development in combination, it is important to combine the above-described material-related parameters and process parameters while appropriately controlling these parameters.

The pattern forming process using two kinds of developers, namely, a positive tone developer and a negative tone developer, may be a process where the exposure is performed once as described above or where the exposure is performed two or more times by the following process. That is, development using a positive tone developer or a negative tone developer is performed after first exposure, and negative or positive tone development using a developer different from that in the first development is performed after second exposure.

A pattern forming method comprising, in order:

(i) a step of coating a substrate with a resist composition for multiple development, whose solubility in a positive tone developer increases and solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation,

(ii-1) a first exposure step,

(v-1) a first heating step,

(iii) a step of developing the resist film with a positive tone developer,

(ii-2) a second exposure step,

(v-2) a second heating step, and

(iv) a step of developing the resist film with a negative tone developer.

A pattern forming method comprising, in order:

(i) a step of coating a substrate with a resist composition for multiple development, whose solubility in a positive tone developer increases and solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation,

(ii-1) a first exposure step,

(v-1) a first heating step,

(iv) a step of developing the resist film with a negative tone developer,

(ii-2) a second exposure step,

(v-2) a second heating step, and

(iii) a step of developing the resist film with a positive tone developer.

As regards the resist composition for multiple development, the resist composition described later can be used.

When exposure is performed two or more times, this is advantageous in that the

latitude of control of the threshold value in the development after first exposure and control of the threshold value in the development after second exposure increases.

In the case of performing the exposure two or more times, the second exposure dose is preferably set to be larger than the first exposure dose. Because, as shown in Fig. 7, in the second development, the threshold value is determined based on the amount to which the history of first and second exposure doses are added, and when the second exposure dose is sufficiently larger than the first exposure dose, the effect of the first exposure dose is reduced and depending on the case, can be neglected.

The exposure dose ($Eo1$ [mJ/cm^2]) in the step of performing the first exposure is preferably 5 [mJ/cm^2] or more smaller than the exposure dose ($Eo2$ [mJ/cm^2]) in the step of performing the second exposure. In this case, the history of the first exposure can be made to less affect the process of performing pattern formation by the second exposure.

In the case of performing the exposure two times, the first development is not limited to positive tone development, and development using a negative tone developer may be performed first.

For changing the first exposure dose and the second exposure dose, a method of adjusting the above-described various parameters related to the material and process is effective. In particular, control of the temperature in the first heating step and the temperature in the second heating step is effective. That is, in order to make the first exposure dose to be smaller than the second exposure dose, it is effective to set the temperature in the first heating step to be lower than the temperature in the second heating step.

The threshold value (a) in the positive tone development is determined as follows in the actual lithography process.

A film composed of a resist composition whose solubility in a positive tone developer increases and solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation is formed on a substrate, and the resist film is exposed through a photomask having a desired pattern size under desired illumination conditions. At this time, the exposure is performed by varying the exposure focus in 0.05 [μm] steps and the exposure dose in 0.5 [mJ/cm^2] steps.

After the exposure, the resist film is heated at a desired temperature for a desired time and then developed with an alkali developer in a desired concentration for a desired time. After the development, the line width of the pattern is measured using CD-SEM, and the exposure dose A [mJ/cm^2] and focus position for forming a desired line width are determined. Subsequently, the intensity distribution of an optical image when the above-described photomask is irradiated with a specific exposure dose A [mJ/cm^2] and a specific focus

position is calculated. The calculation can be performed using a simulation software (Prolith, ver. 9.2.0.15, produced by KLA). Details of the calculation method are described in Chris. A. Mack, Inside PROLITH, Chapter 2, "Aerial Image Formation", FINLE Technologies, Inc.

As a result of calculation, for example, the spatial intensity distribution shown in Fig. 8 of an optical image is obtained.

Here, as shown in Fig. 9, the light intensity at a position after the spatial position is shifted by 1/2 of the obtained pattern line width from the minimum value in the spatial intensity distribution of an optical image becomes the threshold value (a).

The threshold value (b) in the negative tone development is determined as follows in the actual lithography process.

A film composed of a resist composition whose solubility in a positive tone developer increases and solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation is formed on a substrate, and the resist film is exposed through a photomask having a desired pattern size under desired illumination conditions. At this time, the exposure is performed by varying the exposure focus in 0.05 [μm] steps and the exposure dose in 0.5 [mJ/cm^2] steps.

After the exposure, the resist film is heated at a desired temperature for a desired time and then developed with an organic developer in a desired concentration for a desired time. After the development, the line width of the pattern is measured using CD-SEM, and the exposure dose A [mJ/cm^2] and focus position for forming a desired line width are determined. Subsequently, the intensity distribution of an optical image when the above-described photomask is irradiated with a specific exposure dose A [mJ/cm^2] and a specific focus position is calculated. The calculation is performed using a simulation software (Prolith, produced by KLA).

For example, a spatial intensity distribution shown in Fig. 10 of an optical image is obtained.

Here, as shown in Fig. 11, the light intensity at a position after the spatial position is shifted by 1/2 of the obtained pattern line width from the maximum value in the spatial intensity distribution of an optical image is defined as the threshold value (b).

The threshold value (a) is preferably from 0.1 to 100 [mJ/cm^2], more preferably from 0.5 to 50 [mJ/cm^2], still more preferably from 1 to 30 [mJ/cm^2]. The threshold value (b) is preferably from 0.1 to 100 [mJ/cm^2], more preferably from 0.5 to 50 [mJ/cm^2], still more preferably from 1 to 30 [mJ/cm^2].

The difference between threshold values (a) and (b) is preferably from 0.1 to 80 [mJ/cm^2], more preferably from 0.5 to 50 [mJ/cm^2], still more preferably from 1 to 30 [mJ/cm^2].

At the time of performing positive tone development, an alkali developer is preferably used.

The alkali developer which can be used in performing positive tone development is, for example, an alkaline aqueous solution of inorganic alkalis such as sodium hydroxide, potassium hydroxide, sodium carbonate, sodium silicate, sodium metasilicate and aqueous ammonia, primary amines such as ethylamine and n-propylamine, secondary amines such as diethylamine and di-n-butylamine, tertiary amines such as triethylamine and methyldiethylamine, alcohol amines such as dimethylethanolamine and triethanolamine, quaternary ammonium salts such as tetramethylammonium hydroxide and tetraethylammonium hydroxide, and cyclic amines such as pyrrole and piperidine.

Furthermore, this alkaline aqueous solution may be used after adding thereto alcohols and a surfactant each in an appropriate amount.

The alkali concentration of the alkali developer is usually from 0.1 to 20 mass%.

The pH of the alkali developer is usually from 10.0 to 15.0.

In particular, an aqueous 2.38% tetramethylammonium hydroxide solution is preferred.

As for the rinsing solution in the rinsing treatment performed after positive tone development, pure water is used, and the pure water may be used after adding thereto a surfactant in an appropriate amount.

At the time of performing negative tone development, an organic developer containing an organic solvent is preferably used.

As for the organic developer which can be used in performing negative tone development, a polar solvent such as ketone-based solvent, ester-based solvent, alcohol-based solvent, amide-based solvent and ether-based solvent, and a hydrocarbon-based solvent can be used. For example, there may be used a ketone-based solvent such as 1-octanone, 2-octanone, 1-nonanone, 2-nonanone, acetone, 4-heptanone, 2-hexanone, diisobutyl ketone, cyclohexanone, methylcyclohexanone, phenylacetone, methyl ethyl ketone and methyl isobutyl ketone; and an ester-based solvent such as methyl acetate, butyl acetate, ethyl acetate, isopropyl acetate, amyl acetate, propylene glycol monomethyl ether acetate, ethylene glycol monoethyl ether acetate, diethylene glycol monobutyl ether acetate, diethylene glycol monoethyl ether acetate, ethyl-3-ethoxypropionate, 3-methoxybutyl acetate, 3-methyl-3-methoxybutyl acetate, methyl formate, ethyl formate, butyl formate, propyl formate, ethyl lactate, butyl lactate and propyl lactate.

Examples of the alcohol-based solvent include an alcohol such as methyl alcohol, ethyl alcohol, n-propyl alcohol, isopropyl alcohol, n-butyl alcohol, sec-butyl alcohol, tert-butyl alcohol, isobutyl alcohol, n-hexyl alcohol, n-heptyl alcohol, n-octyl alcohol and n-

decanol; a glycol-based solvent such as ethylene glycol, diethylene glycol and triethylene glycol; and a glycol ether-based solvent such as ethylene glycol monomethyl ether, propylene glycol monomethyl ether, ethylene glycol, propylene glycol, diethylene glycol monomethyl ether, triethylene glycol monoethyl ether and methoxymethyl butanol.

Examples of the ether-based solvent include, in addition to the glycol ether-based solvents above, dioxane and tetrahydrofuran.

Examples of the amide-based solvent which can be used include N-methyl-2-pyrrolidone, N,N-dimethylacetamide and N,N-dimethylformamide.

Examples of the hydrocarbon-based solvent include an aromatic hydrocarbon-based solvent such as toluene and xylene, and an aliphatic hydrocarbon-based solvent such as pentane, hexane, octane and decane.

A plurality of these solvents may be mixed, or the solvent may be used by mixing it with a solvent other than those described above or water.

As regards the development system, for example, a method of raising the developer on a substrate surface by the effect of a surface tension and keeping it still for a fixed time, thereby performing the development (puddle method), a method of spraying the developer on a substrate surface (spray method), and a method of continuously ejecting the developer on a substrate rotating at a constant speed while scanning the developer ejecting nozzle at a constant rate (dynamic dispense method) may be applied. In using such a development method, if the vapor pressure of the negative tone developer is high, the substrate surface is cooled due to evaporation of the developer to reduce the temperature of the developer and the film of the resist composition formed on the substrate is not dissolved at a constant rate, giving rise to deterioration of the dimensional uniformity. For this reason, the vapor pressure at 20°C of the developer which can be used in performing negative tone development is preferably 5 kPa or less, more preferably 3 kPa or less, and most preferably 2 kPa or less.

Specific examples of the developer having a vapor pressure of 5 kPa or less at 20°C include a ketone-based solvent such as 1-octanone, 2-octanone, 1-nonanone, 2-nonanone, 4-heptanone, 2-hexanone, diisobutyl ketone, cyclohexanone, methylcyclohexanone, phenylacetone and methyl isobutyl ketone; an ester-based solvent such as butyl acetate, amyl acetate, propylene glycol monomethyl ether acetate, ethylene glycol monoethyl ether acetate, diethylene glycol monobutyl ether acetate, diethylene glycol monoethyl ether acetate, ethyl-3-ethoxypropionate, 3-methoxybutyl acetate, 3-methyl-3-methoxybutyl acetate, butyl formate, propyl formate, ethyl lactate, butyl lactate and propyl lactate; an alcohol-based solvent such as n-propyl alcohol, isopropyl alcohol, n-butyl alcohol, sec-butyl alcohol, tert-butyl alcohol, isobutyl alcohol, n-hexyl alcohol, n-heptyl alcohol, n-octyl alcohol and n-decanol; a glycol-based solvent such as ethylene glycol, diethylene glycol and triethylene glycol; a glycol ether-

based solvent such as propylene glycol monomethyl ether, ethylene glycol, propylene glycol, diethylene glycol monomethyl ether, triethylene glycol monoethyl ether and methoxymethylbutanol; an ether-based solvent such as tetrahydrofuran; an amide-based solvent such as N-methyl-2-pyrrolidone, N,N-dimethylacetamide and N,N-dimethylformamide; an aromatic hydrocarbon-based solvent such as toluene and xylene; and an aliphatic hydrocarbon-based solvent such as octane and decane.

Specific examples of the developer having a vapor pressure of 2 kPa or less at 20°C, which is a most preferred range, include a ketone-based solvent such as 1-octanone, 2-octanone, 1-nonanone, 2-nonanone, 4-heptanone, 2-hexanone, diisobutyl ketone, cyclohexanone, methylcyclohexanone and phenylacetone; an ester-based solvent such as butyl acetate, amyl acetate, propylene glycol monomethyl ether acetate, ethylene glycol monoethyl ether acetate, diethylene glycol monobutyl ether acetate, diethylene glycol monoethyl ether acetate, ethyl-3-ethoxypropionate, 3-methoxybutyl acetate, 3-methyl-3-methoxybutyl acetate, ethyl lactate, butyl lactate and propyl lactate; an alcohol-based solvent such as n-butyl alcohol, sec-butyl alcohol, tert-butyl alcohol, isobutyl alcohol, n-hexyl alcohol, n-heptyl alcohol, n-octyl alcohol and n-decanol; a glycol-based solvent such as ethylene glycol, diethylene glycol and triethylene glycol; a glycol ether-based solvent such as propylene glycol monomethyl ether, ethylene glycol, propylene glycol, diethylene glycol monomethyl ether, triethylene glycol monoethyl ether and methoxymethylbutanol; an amide-based solvent such as N-methyl-2-pyrrolidone, N,N-dimethylacetamide and N,N-dimethylformamide; an aromatic hydrocarbon-based solvent such as xylene; and an aliphatic hydrocarbon-based solvent such as octane and decane.

In the developer usable when performing negative tone development, a surfactant can be added in an appropriate amount, if desired.

The surfactant is not particularly limited but, for example, an ionic or nonionic fluorine-containing and/or silicon-containing surfactant can be used. Examples of such a fluorine-containing and/or silicon-containing surfactant include the surfactants described in JP-A-62-36663, JP-A-61-226746, JP-A-61-226745, JP-A-62-170950, JP-A-63-34540, JP-A-7-230165, JP-A-8-62834, JP-A-9-54432, JP-A-9-5988 and U.S. Patents 5,405,720, 5,360,692, 5,529,881, 5,296,330, 5,436,098, 5,576,143, 5,294,511 and 5,824,451. The surfactant is preferably a nonionic surfactant. The nonionic surfactant is not particularly limited, but a fluorine-containing surfactant or a silicon-containing surfactant is more preferred.

The amount of the surfactant used is usually from 0.001 to 5 mass%, preferably from 0.005 to 2 mass%, more preferably from 0.01 to 0.5 mass%, based on the entire amount of the developer.

After the step of performing negative tone development, a step of stopping the

development by the replacement with another solvent may be practiced.

A step of washing the resist film with a rinsing solution containing an organic solvent is preferably provided after the negative tone development.

A method where after washing with a rinsing solution, the rinsing solution is removed from the substrate surface by rotating the substrate at a rotation number of 2,000 rpm to 4,000 rpm, is preferred. In the case where the vapor pressure of the rinsing solution is low, the rinsing solution remains on the substrate even after removing the rinsing solution by rotating the substrate and penetrates into the resist pattern formed on the substrate to swell the resist pattern, as a result, the dimensional uniformity of the resist pattern is deteriorated. For this reason, the vapor pressure at 20°C of the rinsing solution is preferably 0.05 kPa or more, more preferably 0.1 kPa or more, and most preferably 0.12 kPa or more.

In the rinsing step after negative tone development, the washing is preferably performed using a rinsing solution containing at least one kind of an organic solvent selected from an alkane-based solvent, a ketone-based solvent, an ester-based solvent, an alcohol-based solvent, an amide-based solvent and an ether-based solvent. Preferably, a step of washing the resist film by using a rinsing solution containing at least one kind of an organic solvent selected from a ketone-based solvent, an ester-based solvent, an alcohol-based solvent and an amide-based solvent is performed after negative tone development; more preferably, a step of washing the resist film by using a rinsing solution containing an alcohol-based solvent or an ester-based solvent is performed after negative tone development; still more preferably, a step of washing the resist film by using a rinsing solution containing a monohydric alcohol having a carbon number of 6 to 8 is performed after negative tone development. The monohydric alcohol having a carbon number of 6 to 8, which is used in the rinsing step after negative tone development, includes a linear, branched or cyclic monohydric alcohol, and specific examples of the monohydric alcohol which can be used include 1-hexanol, 1-heptanol, 1-octanol, 2-hexanol, 2-heptanol, 2-octanol, 3-hexanol, 3-heptanol, 3-octanol, 4-octanol and benzyl alcohol, with 1-hexanol, 2-hexanol and 2-heptanol being preferred.

A plurality of these components may be mixed, or the solvent may be used by mixing it with an organic solvent other than those described above.

The water content in the rinsing solution is preferably 10 mass% or less, more preferably 5 mass% or less, still more preferably 3 mass% or less. By setting the water content to 10 mass% or less, good development characteristics can be obtained.

The rinsing solution may also be used after adding thereto a surfactant in an appropriate amount.

In the rinsing step, the wafer after negative tone development is washed using the above-described organic solvent-containing rinsing solution. The method of washing

treatment is not particularly limited but, for example, a method of continuously ejecting the rinsing solution on a substrate rotating at a constant speed (spin coating method), a method of dipping the substrate in a bath filled with the rinsing solution for a fixed time (dipping method), and a method of spraying the rinsing solution on the substrate surface (spraying method) may be applied. Above all, a method where the washing treatment is performed by the spin coating method and after the washing, the rinsing solution is removed from the substrate surface by rotating the substrate at a rotation number of 2,000 rpm to 4,000 rpm, is preferred.

As regards the development method, for example, a method of dipping the substrate in a bath filled with the developer for a fixed time (dipping method), a method of raising the developer on the substrate surface by the effect of a surface tension and keeping it still for a fixed time, thereby performing the development (puddle method), a method of spraying the developer on a substrate surface (spraying method), and a method of continuously ejecting the developer on the substrate rotating at a constant speed while scanning the developer ejecting nozzle at a constant rate (dynamic dispense method) may be applied.

In the pattern forming method of the present invention, the step of forming a film on a substrate by using a resist composition whose solubility in a positive tone developer increases and solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation, the step of exposing the film, the step of heating the film, and the step of applying positive tone development to the film may be performed by generally known methods.

The exposure apparatus for use in the present invention is not limited in the light source wavelength, but, for example, a KrF excimer laser wavelength (248 nm), an ArF excimer laser wavelength (193 nm) and an F₂ excimer laser wavelength (157 nm) can be applied.

In the step of performing exposure of the present invention, an immersion exposure method can be applied.

The immersion exposure method is a technique for raising the resolution, and this is a technique of performing the exposure by filling a high refractive-index liquid (hereinafter sometimes referred to as an "immersion liquid") between the projection lens and the sample.

As for the "effect of immersion", assuming that $NA_0 = \sin\theta$, the resolution and depth of focus when immersed can be expressed by the following formulae:

$$(\text{Resolution}) = k_1 \bullet (\lambda_0/n)/NA_0$$

$$(\text{Depth of focus}) = \pm k_2 \bullet (\lambda_0/n)/NA_0^2$$

wherein λ_0 is the wavelength of exposure light in air, n is the refractive index of the immersion liquid based on air, and θ is the convergence half-angle of beam.

That is, the effect of immersion is equal to use of an exposure wavelength of $1/n$. In other words, when the projection optical system has the same NA, the depth of focus can be made n times larger by the immersion. This is effective for all pattern profiles and can be combined with the super-resolution technology under study at present, such as phase-shift method and modified illumination method.

In the case of performing immersion exposure, a step of washing the film surface with an aqueous chemical solution may be performed (1) before the exposure step after forming the film on a substrate and/or (2) after the step of exposing the film through an immersion liquid but before the step of heating the film.

The immersion liquid is preferably a liquid being transparent to light at the exposure wavelength and having as small a temperature coefficient of refractive index as possible so as to minimize the distortion of an optical image projected on the film. Particularly, when the exposure light source is an ArF excimer laser (wavelength: 193 nm), water is preferably used in view of easy availability and easy handleability, in addition to the above-described aspects.

In the case of using water, an additive (liquid) capable of decreasing the surface tension of water and increasing the surface activity may be added in a small ratio. This additive is preferably a liquid that does not dissolve the resist layer on the wafer and at the same time, gives only a negligible effect on the optical coat at the undersurface of the lens element.

Such an additive is preferably, for example, an aliphatic alcohol having a refractive index nearly equal to that of water, and specific examples thereof include methyl alcohol, ethyl alcohol and isopropyl alcohol. By virtue of adding an alcohol having a refractive index nearly equal to that of water, even when the alcohol component in water is evaporated and its content concentration is changed, the change in the refractive index of the entire liquid can be advantageously made very small.

On the other hand, if a substance opaque to light at 193 nm or an impurity greatly differing in the refractive index from water is mingled, this incurs distortion of the optical image projected on the resist. Therefore, the water used is preferably distilled water. Pure water after further filtration through an ion exchange filter or the like may also be used.

In the present invention, the substrate on which the film is formed is not particularly limited, and an inorganic substrate such as silicon, SiN, SiO₂ and SiN, a coating-type inorganic substrate such as SOG, or a substrate generally used in the process of producing a semiconductor such as IC or producing a circuit board of liquid crystal, thermal head or the like or in the lithography process of other photo-applications can be used. If desired, an organic antireflection film may be formed between the film and the substrate.

In the present invention, the film formed on the substrate is a film composed of a

resist composition whose solubility in a positive tone developer increases and solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation.

The resist composition which can be used in the present invention is described below.

(A) Resin capable of increasing the polarity by the action of an acid

The resin capable of increasing the polarity by the action of an acid, which is used in the resist composition of the present invention, is a resin having a group that decomposes by the action of an acid to produce an alkali-soluble group (hereinafter sometimes referred to as an "acid-decomposable group"), on either one or both of the main chain and the side chain of the resin (sometimes referred to as an "acid-decomposable resin", an "acid-decomposable resin (A)" or a "resin (A)"). The resin is preferably a resin whose solubility in a positive tone developer increases and which has a monocyclic or polycyclic alicyclic hydrocarbon structure and can increase the porality, increase the solubility in an alkali developer and decrease the solubility in an organic solvent by the action of an acid (hereinafter sometimes referred to as an "alicyclic hydrocarbon-based acid-decomposable resin"). Because, the polarity of the resin is greatly changed between before and after irradiation of an actinic ray or radiation and when the resist film is developed using a positive tone developer (preferably an alkali developer) and a negative tone developer (preferably an organic solvent), the dissolution contrast rises. Furthermore, the resin having a monocyclic or polycyclic alicyclic hydrocarbon structure generally has high hydrophobicity and ensures a high development rate at the time of developing the resist film in the region of weak light irradiation intensity with a negative tone developer (preferably an organic developer), and the developability in using a negative tone developer is enhanced.

Examples of the alkali-soluble group include groups having a phenolic hydroxyl group, a carboxylic acid group, a fluorinated alcohol group, a sulfonic acid group, a sulfonamide group, a sulfonylimide group, an (alkylsulfonyl)(alkylcarbonyl)methylene group, an (alkylsulfonyl)(alkylcarbonyl)imide group, a bis(alkylcarbonyl)methylene group, a bis(alkylcarbonyl)imide group, a bis(alkylsulfonyl)methylene group, a bis(alkylsulfonyl)imide group, a tris(alkylcarbonyl)methylene group or a tris(alkylsulfonyl)methylene group.

The alkali-soluble group is preferably a carboxylic acid group, a fluorinated alcohol group (preferably hexafluoroisopropanol) or a sulfonic acid group.

The group capable of decomposing by the action of an acid (acid-decomposable groups) is preferably a group obtained by substituting the hydrogen atom of the above-described alkali-soluble group with a group capable of leaving by the action of an acid.

Examples of the group capable of leaving by the action of an acid include $-C(R_{36})(R_{37})(R_{38})$, $-C(R_{36})(R_{37})(OR_{39})$ and $-C(R_{01})(R_{02})(OR_{39})$.

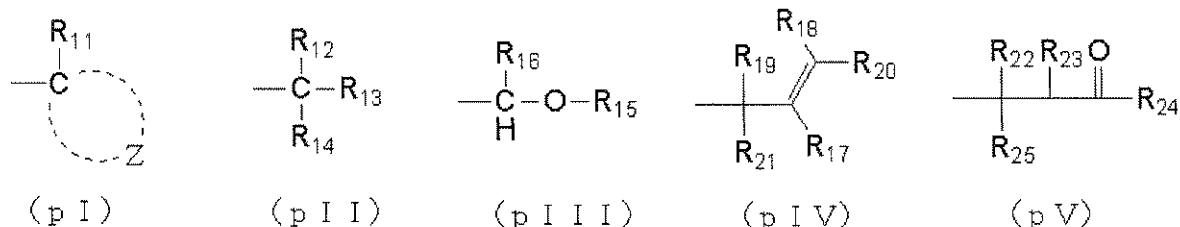
In the formulae, each of R_{36} to R_{39} independently represents an alkyl group, a cycloalkyl group, an aryl group, an aralkyl group or an alkenyl group. R_{36} and R_{37} may combine with each other to form a ring.

Each of R_{01} and R_{02} independently represents a hydrogen atom, an alkyl group, a cycloalkyl group, an aryl group, an aralkyl group or an alkenyl group.

The acid-decomposable group is preferably a cumyl ester group, an enol ester group, an acetal ester group, a tertiary alkyl ester group or the like, more preferably a tertiary alkyl ester group.

The resist composition of the present invention containing a resin (a1) having a monocyclic or polycyclic alicyclic hydrocarbon structure and being capable of increasing the polarity by the action of an acid can be suitably used when ArF excimer laser light is irradiated.

The resin having a monocyclic or polycyclic alicyclic hydrocarbon structure and being capable of increasing the polarity by the action of an acid (hereinafter sometimes referred to as an "alicyclic hydrocarbon-based acid-decomposable resin") is preferably a resin containing at least one member selected from the group consisting of a repeating unit having an alicyclic hydrocarbon-containing partial structure represented by any one of the following formulae (pI) to (pV) and a repeating unit represented by the following formula (II-AB).



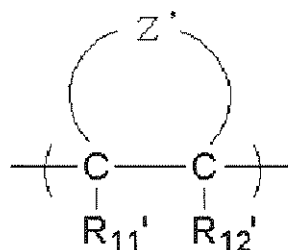
In formulae (pI) to (pV), R_{11} represents a methyl group, an ethyl group, an n-propyl group, an isopropyl group, an n-butyl group, an isobutyl group or a sec-butyl group.

Z represents an atomic group necessary for forming a cycloalkyl group together with the carbon atom.

Each of R_{12} to R_{16} independently represents a linear or branched alkyl group having a carbon number of 1 to 4 or a cycloalkyl group, provided that at least one of R_{12} to R_{14} or either one of R_{15} and R_{16} represents a cycloalkyl group.

Each of R_{17} to R_{21} independently represents a hydrogen atom, a linear or branched alkyl group having a carbon number of 1 to 4 or a cycloalkyl group, provided that at least one of R_{17} to R_{21} represents a cycloalkyl group and that either one of R_{19} and R_{21} represents a linear or branched alkyl group having a carbon number of 1 to 4 or a cycloalkyl group.

Each of R_{22} to R_{25} independently represents a hydrogen atom, a linear or branched alkyl group having a carbon number of 1 to 4 or a cycloalkyl group, provided that at least one of R_{22} to R_{25} represents a cycloalkyl group. R_{23} and R_{24} may combine with each other to form a ring.

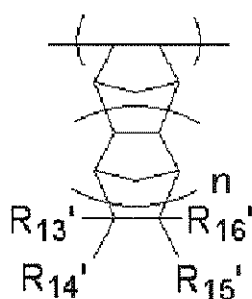


(I I - A B)

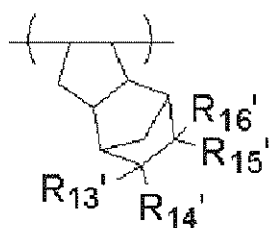
In formula (II-AB), each of R_{11}' and R_{12}' independently represents a hydrogen atom, a cyano group, a halogen atom or an alkyl group.

Z' represents an atomic group for forming an alicyclic structure containing two bonded carbon atoms (C-C).

Formula (II-AB) is preferably the following formula (II-AB1) or (II-AB2):



(I I - A B 1)



(I I - A B 2)

In formulae (II-AB1) and (II-AB2), each of R_{13}' to R_{16}' independently represents a hydrogen atom, a halogen atom, a cyano group, $-\text{COOH}$, $-\text{COOR}_5$, a group capable of decomposing by the action of an acid, $-\text{C}(=\text{O})-\text{X}-\text{A}'-\text{R}_{17}'$, an alkyl group or a cycloalkyl group, and at least two members out of R_{13}' to R_{16}' may combine to form a ring.

R_5 represents an alkyl group, a cycloalkyl group or a group having a lactone structure.

X represents an oxygen atom, a sulfur atom, $-\text{NH}-$, $-\text{NHSO}_2-$ or $-\text{NHSO}_2\text{NH}-$.

A' represents a single bond or a divalent linking group.

R_{17}' represents $-\text{COOH}$, $-\text{COOR}_5$, $-\text{CN}$, a hydroxyl group, an alkoxy group, $-\text{CO}-\text{NH}-\text{R}_6$, $-\text{CO}-\text{NH}-\text{SO}_2-\text{R}_6$ or a group having a lactone structure.

R_6 represents an alkyl group or a cycloalkyl group.

n represents 0 or 1.

In formulae (pI) to (pV), the alkyl group of R_{12} to R_{25} is a linear or branched alkyl

group having a carbon number of 1 to 4.

The cycloalkyl group of R_{11} to R_{25} and the cycloalkyl group formed by Z together with the carbon atom may be monocyclic or polycyclic. Specific examples thereof include a group having a carbon number of 5 or more and having a monocyclo, bicyclo, tricyclo or tetracyclo structure or the like. The carbon number thereof is preferably from 6 to 30, more preferably from 7 to 25. These cycloalkyl groups each may have a substituent.

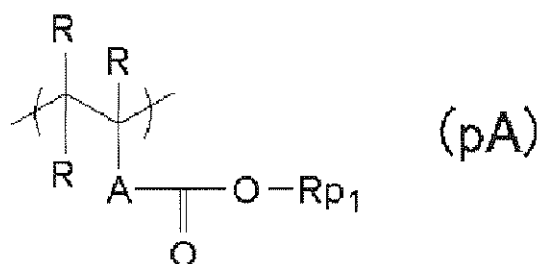
Preferred examples of the cycloalkyl group include an adamantyl group, a noradamantyl group, a decalin residue, a tricyclodecanyl group, a tetracyclododecanyl group, a norbornyl group, a cedrol group, a cyclopentyl group, a cyclohexyl group, a cycloheptyl group, a cyclooctyl group, a cyclodecanyl group and a cyclododecanyl group. Among these, more preferred are an adamantyl group, a norbornyl group, a cyclohexyl group, a cyclopentyl group, a tetracyclododecanyl group and a tricyclodecanyl group.

Examples of the substituent which the alkyl group and cycloalkyl group each may further have include an alkyl group (having a carbon number of 1 to 4), a halogen atom, a hydroxyl group, an alkoxy group (having a carbon number of 1 to 4), a carboxyl group and an alkoxycarbonyl group (having a carbon number of 2 to 6). Examples of the substituent which these alkyl group, alkoxy group, alkoxycarbonyl group and the like each may further have include a hydroxyl group, a halogen atom and an alkoxy group.

The structures represented by formulae (pI) to (pV) each can be used for the protection of an alkali-soluble group in the resin. Examples of the alkali-soluble group include various groups known in this technical field.

Specific examples thereof include a structure where the hydrogen atom of a carboxylic acid group, a sulfonic acid group, a phenol group or a thiol group is replaced by the structure represented by any one of formulae (pI) to (pV). Among these, preferred is a structure where the hydrogen atom of a carboxylic acid group or a sulfonic acid group is replaced by the structure represented by any one of formulae (pI) to (pV).

The repeating unit having an alkali-soluble group protected by the structure represented by any one of formulae (pI) to (pV) is preferably a repeating unit represented by the following formula (pA):



In the formula, R represents a hydrogen atom, a halogen atom or a linear or branched alkyl group having a carbon number of 1 to 4, and a plurality of R's may be the same or different from each other.

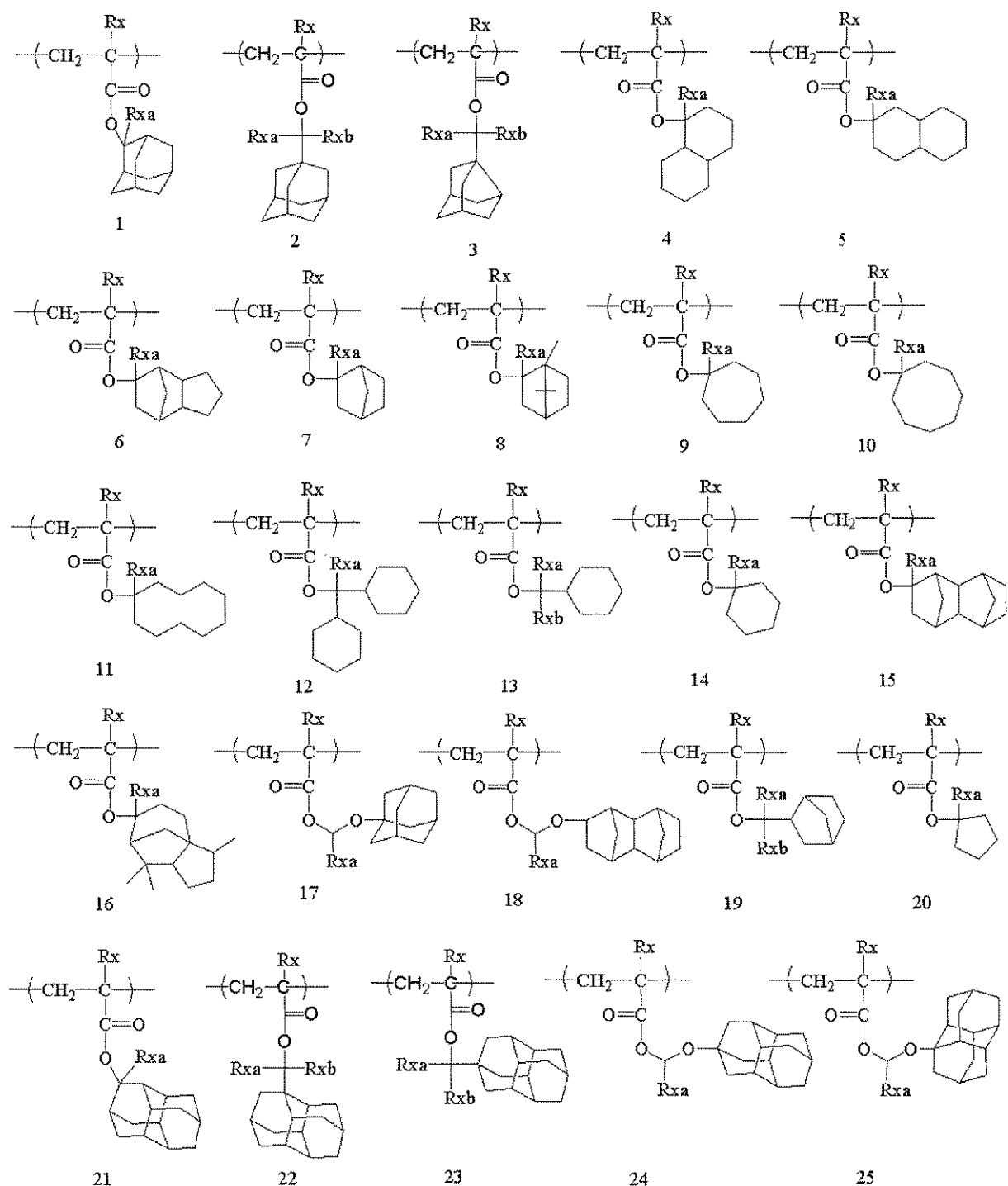
A represents a single bond, or a sole group or a combination of two or more groups selected from the group consisting of an alkylene group, an ether group, a thioether group, a carbonyl group, an ester group, an amido group, a sulfonamido group, a urethane group and a urea group, and is preferably a single bond.

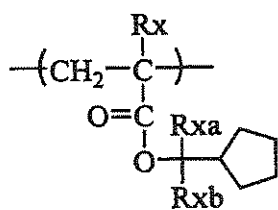
Rp₁ represents a group represented by any one of formulae (pI) to (pV).

The repeating unit represented by formula (pA) is more preferably a repeating unit composed of a 2-alkyl-2-adamantyl (meth)acrylate or a dialkyl(1-adamantyl)methyl (meth)acrylate.

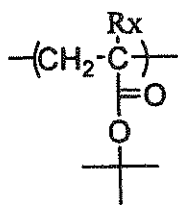
Specific examples of the repeating unit represented by formula (pA) are set forth below, but the present invention is not limited thereto.

(In the formulae, Rx represents H, CH₃ or CH₂OH, and each of Rxa and Rxb independently represents an alkyl group having a carbon number of 1 to 4.)

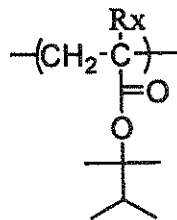




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Examples of the halogen atom of R_{11}' and R_{12}' in formula (II-AB) include a chlorine atom, a bromine atom, a fluorine atom and an iodine atom.

The alkyl group of R_{11}' and R_{12}' includes a linear or branched alkyl group having a carbon number of 1 to 10.

The atomic group of Z' for forming an alicyclic structure is an atomic group for forming, in the resin, a repeating unit composed of an alicyclic hydrocarbon which may have a substituent. Above all, an atomic group for forming a crosslinked alicyclic structure to form a crosslinked alicyclic hydrocarbon repeating unit is preferred.

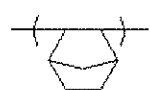
Examples of the skeleton of the alicyclic hydrocarbon formed are the same as those of the alicyclic hydrocarbon group of R_{12} to R_{25} in formulae (pI) to (pV).

The skeleton of the alicyclic hydrocarbon may have a substituent, and examples of the substituent include R_{13}' to R_{16}' in formulae (II-AB1) and (II-AB2).

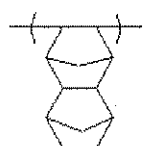
In the alicyclic hydrocarbon-based acid-decomposable resin for use in the present invention, the group capable of decomposing by the action of an acid may be contained in at least one repeating unit out of the repeating unit having an alicyclic hydrocarbon-containing partial structure represented by any one of formulae (pI) to (pV), the repeating unit represented by formula (II-AB), and the repeating unit composed of a copolymerization component described later. The group capable of decomposing by the action of an acid is preferably contained in the repeating unit having an alicyclic hydrocarbon-containing partial structure represented by any one of formulae (pI) to (pV).

Various substituents R_{13}' to R_{16}' in formulae (II-AB1) and (II-AB2) may become substituents of the atomic group for forming an alicyclic structure or the atomic group Z for forming a crosslinked alicyclic structure in formula (II-AB).

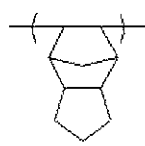
Specific examples of the repeating units represented by formulae (II-AB1) and (II-AB2) are set forth below, but the present invention is not limited to these specific examples.



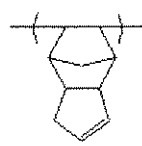
[II-1]



[II-2]



[II-3]



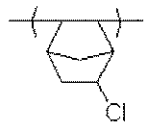
[II-4]



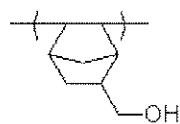
[II-5]



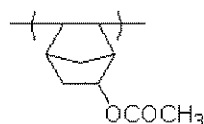
[II-6]



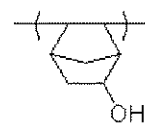
[II-7]



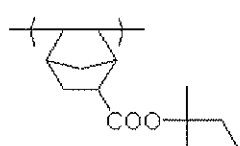
[II-8]



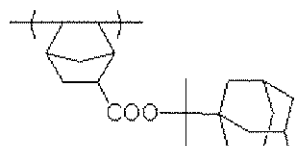
[II-9]



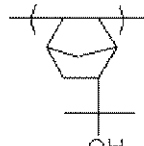
[II-10]



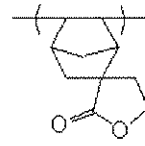
[II-11]



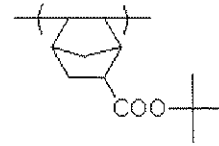
[II-12]



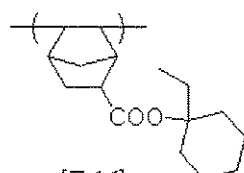
[II-13]



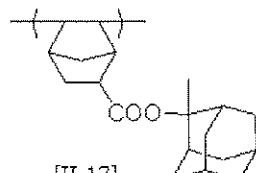
[II-14]



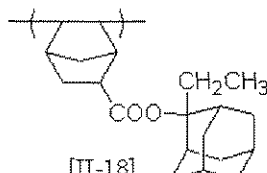
[II-15]



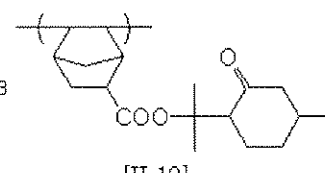
[II-16]



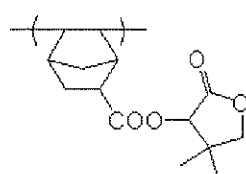
[II-17]



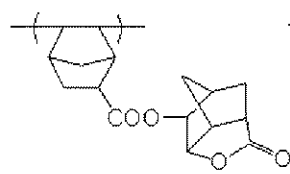
[II-18]



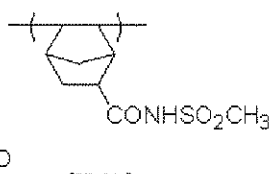
[II-19]



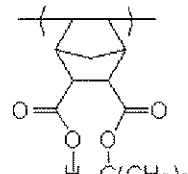
[II-20]



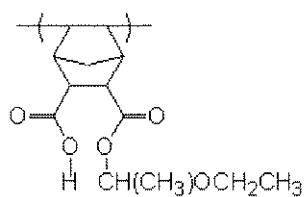
[II-21]



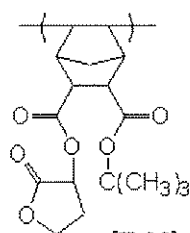
[II-22]



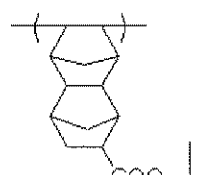
[II-23]



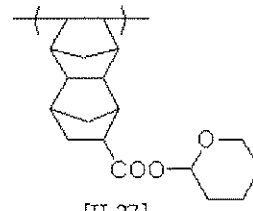
[II-24]



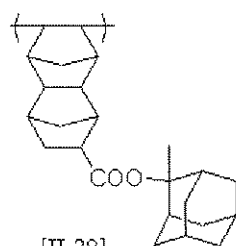
[II-25]



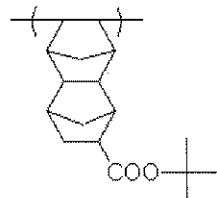
[II-26]



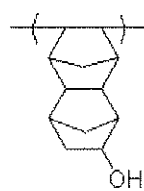
[II-27]



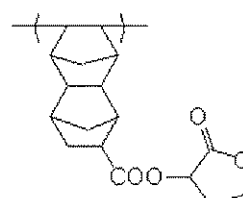
[II-28]



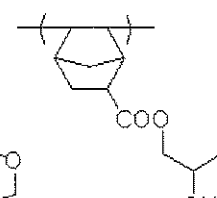
[II-29]



[II-30]

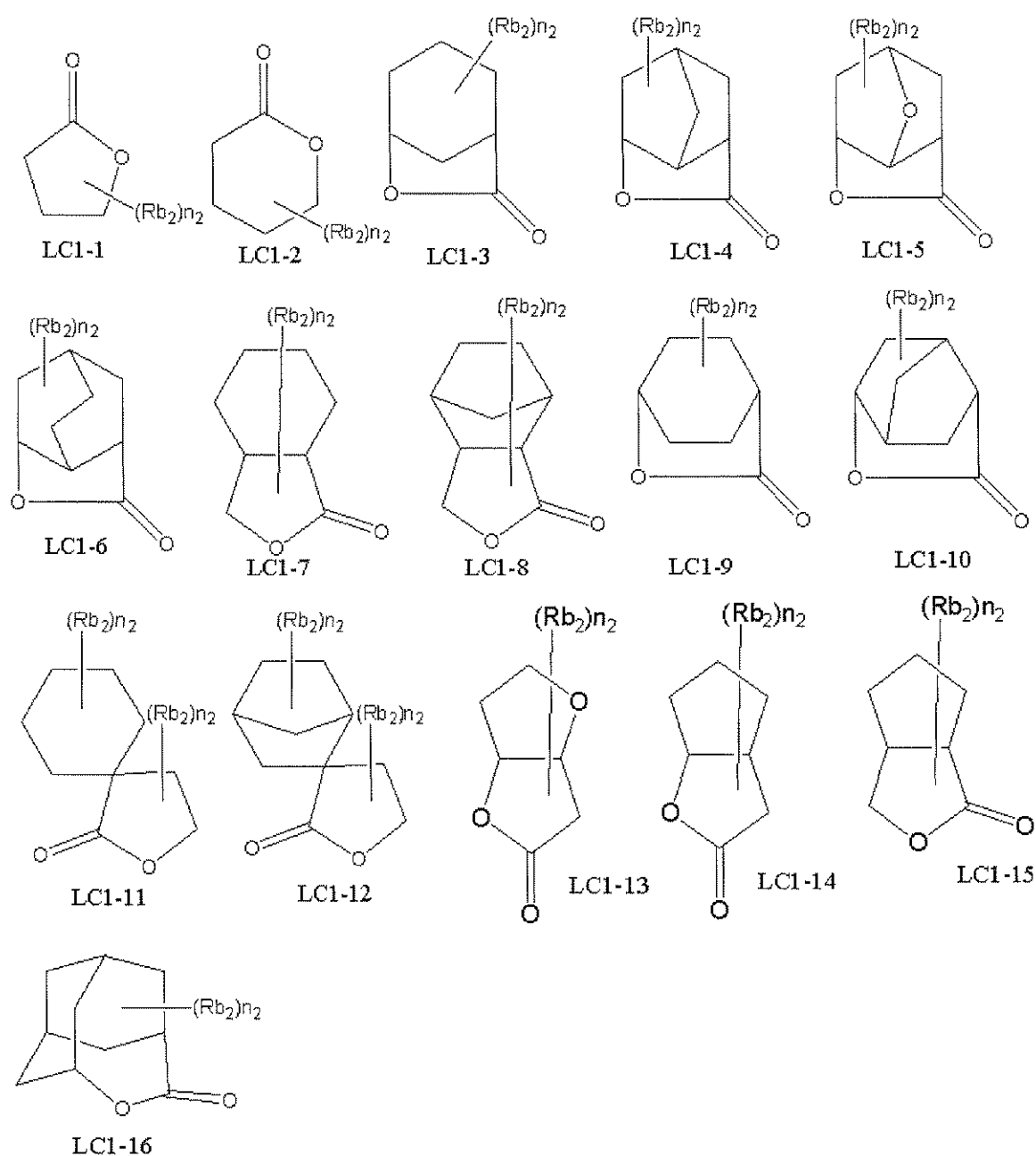


[II-31]



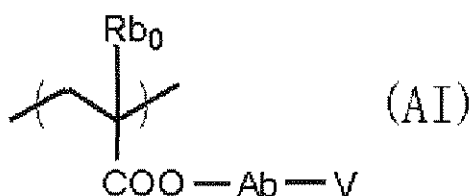
[II-32]

The alicyclic hydrocarbon-based acid-decomposable resin for use in the present invention preferably has a lactone group. As for the lactone group, any group may be used as long as it has a lactone structure, but a group having a 5- to 7-membered ring lactone structure is preferred. The 5- to 7-membered ring lactone structure is preferably condensed with another ring structure in the form of forming a bicyclo or spiro structure. The resin more preferably has a repeating unit containing a group having a lactone structure represented by any one of the following formulae (LC1-1) to (LC1-16). The group having a lactone structure may be bonded directly to the main chain. Among these lactone structures, preferred are groups represented by formulae (LC1-1), (LC1-4), (LC1-5), (LC1-6), (LC1-13) and (LC1-14). By virtue of using a specific lactone structure, the line edge roughness and development defect are improved.



The lactone structure moiety may or may not have a substituent (Rb_2). Preferred examples of the substituent (Rb_2) include an alkyl group having a carbon number of 1 to 8, a cycloalkyl group having a carbon number of 4 to 7, an alkoxy group having a carbon number of 1 to 8, an alkoxy carbonyl group having a carbon number of 1 to 8, a carboxyl group, a halogen atom, a hydroxyl group, a cyano group and an acid-decomposable group. n_2 represents an integer of 0 to 4. When n_2 is an integer of 2 or more, each Rb_2 may be the same as or different from every other Rb_2 and also, the plurality of substituents (Rb_2) may combine with each other to form a ring.

Examples of the repeating unit containing a group having a lactone structure represented by any one of formulae (LC1-1) to (LC1-16) include a repeating unit where at least one of R_{13}' to R_{16}' in formula (II-AB1) or (II-AB2) has a group represented by any one of formulae (LC1-1) to (LC1-16) (for example, R_5 of $-COOR_5$ is a group represented by any one of formulae (LC1-1) to (LC1-16)), and a repeating unit represented by the following formula (AI):



In formula (AI), Rb_0 represents a hydrogen atom, a halogen atom or an alkyl group having a carbon number of 1 to 4.

Preferred examples of the substituent which the alkyl group of Rb_0 may have include a hydroxyl group and a halogen atom.

The halogen atom of Rb_0 includes a fluorine atom, a chlorine atom, a bromine atom and an iodine atom.

Rb_0 is preferably a hydrogen atom or a methyl group.

Ab represents a single bond, an alkylene group, a divalent linking group having a monocyclic or polycyclic alicyclic hydrocarbon structure, an ether group, an ester group, a carbonyl group, or a divalent group formed by combining these groups and is preferably a single bond or a linking group represented by $-Ab_1-CO_2-$. Ab_1 represents a linear or branched alkylene group or a monocyclic or polycyclic cycloalkylene group and is preferably a methylene group, an ethylene group, a cyclohexylene group, an adamantylene group or a norbornylene group.

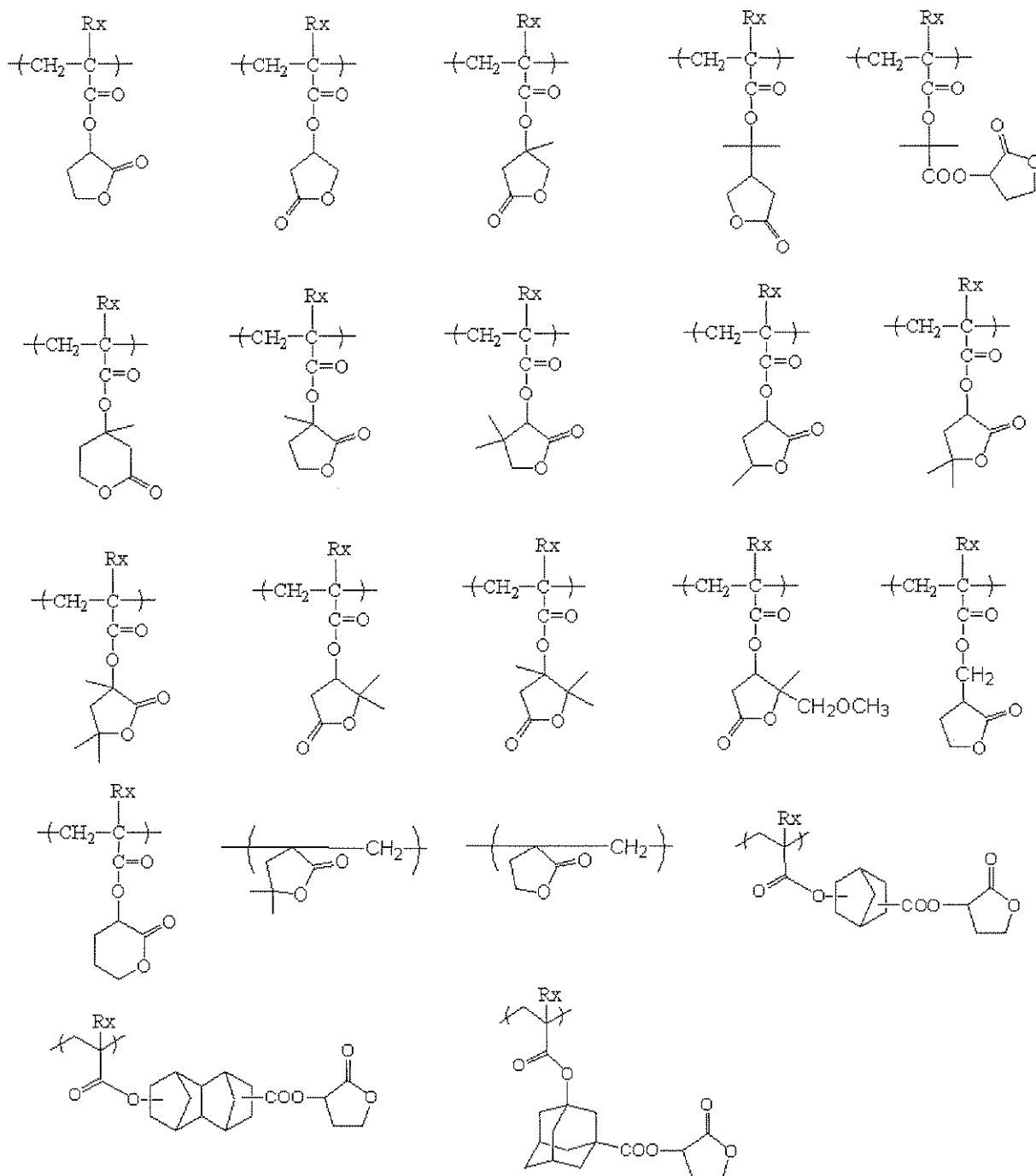
V represents a group represented by any one of formulae (LC1-1) to (LC1-16).

The repeating unit having a lactone structure usually has an optical isomer, but any optical isomer may be used. One optical isomer may be used alone or a mixture of a

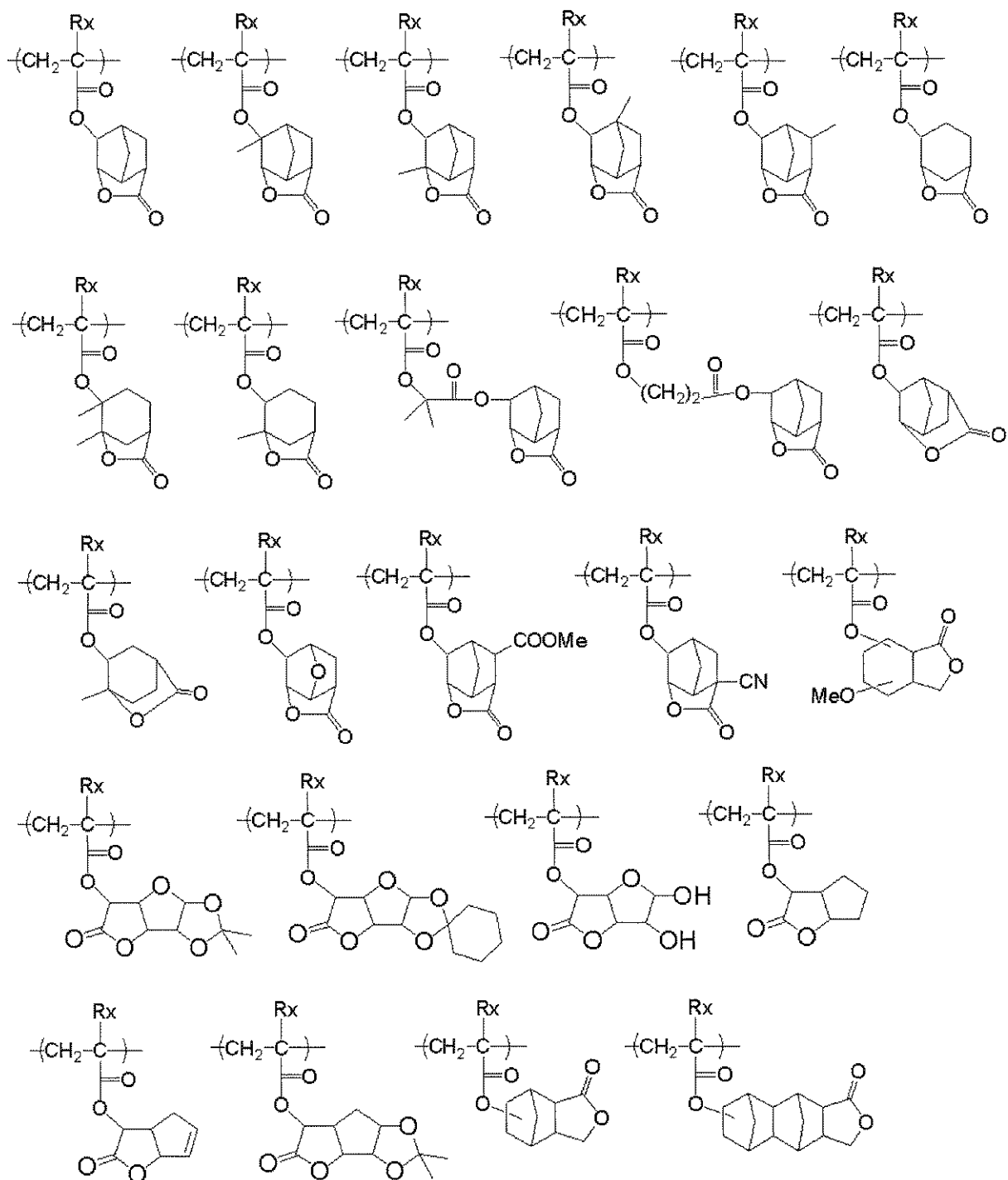
plurality of optical isomers may be used. In the case of mainly using one optical isomer, the optical purity (ee) thereof is preferably 90 or more, more preferably 95 or more.

Specific examples of the repeating unit having a lactone structure are set forth below, but the present invention is not limited thereto.

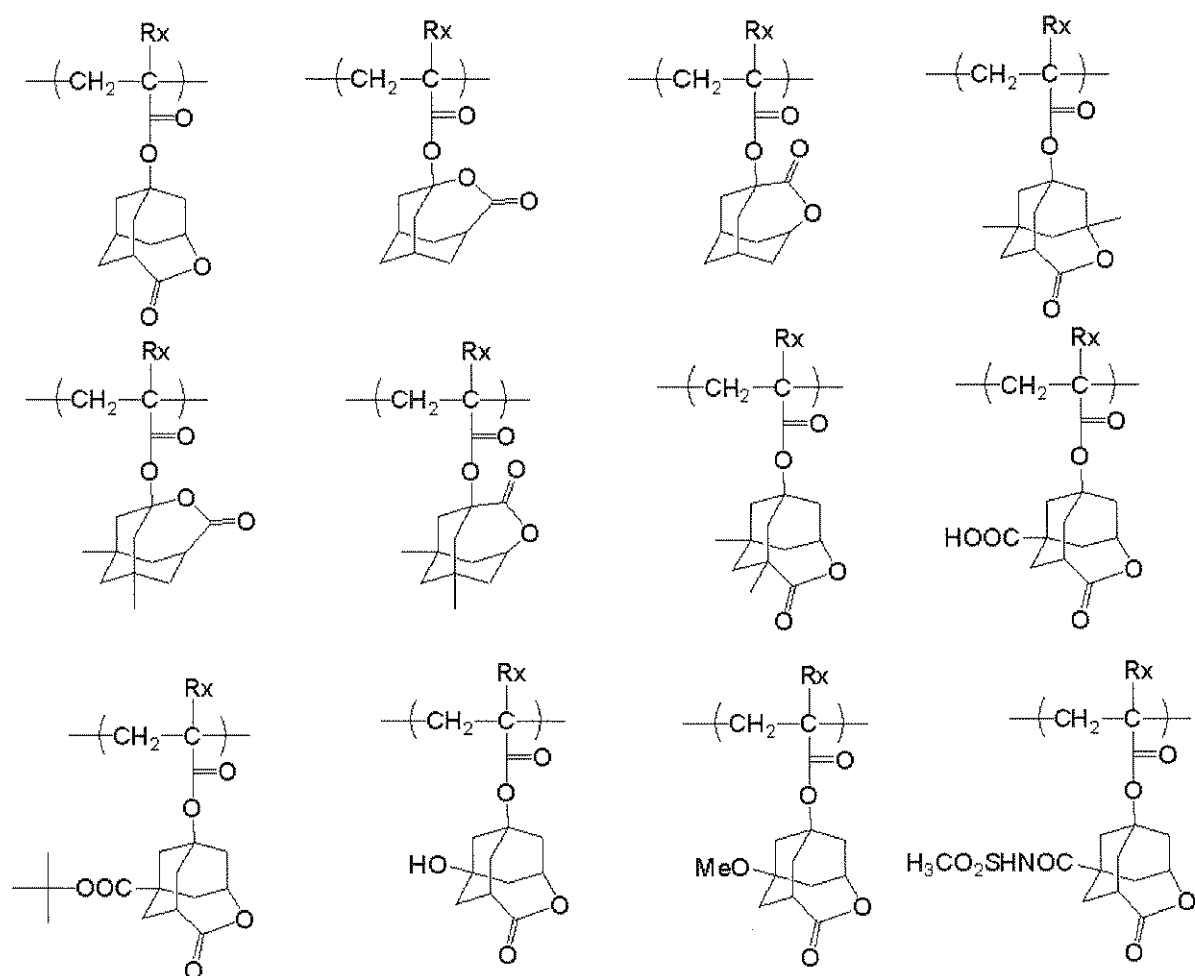
(In the formulae, Rx is H, CH₃, CH₂OH or CF₃.)



(In the formulae, Rx is H, CH₃, CH₂OH or CF₃.)

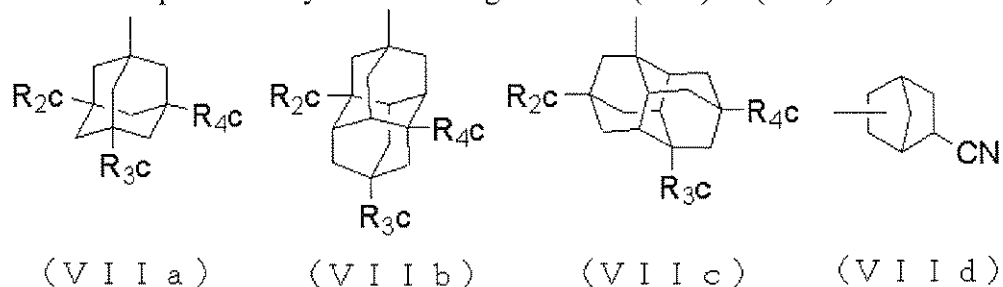


(In the formulae, Rx is H, CH₃, CH₂OH or CF₃.)



The alicyclic hydrocarbon-based acid-decomposable resin for use in the present invention preferably has a repeating unit containing an organic group having a polar group, more preferably a repeating unit having an alicyclic hydrocarbon structure substituted by a polar group. By virtue of this repeating unit, the adherence to substrate and the affinity for developer are enhanced. The alicyclic hydrocarbon structure of the polar group-substituted alicyclic hydrocarbon structure is preferably an adamantyl group, a diamantyl group or a norbornane group. The polar group is preferably a hydroxyl group or a cyano group.

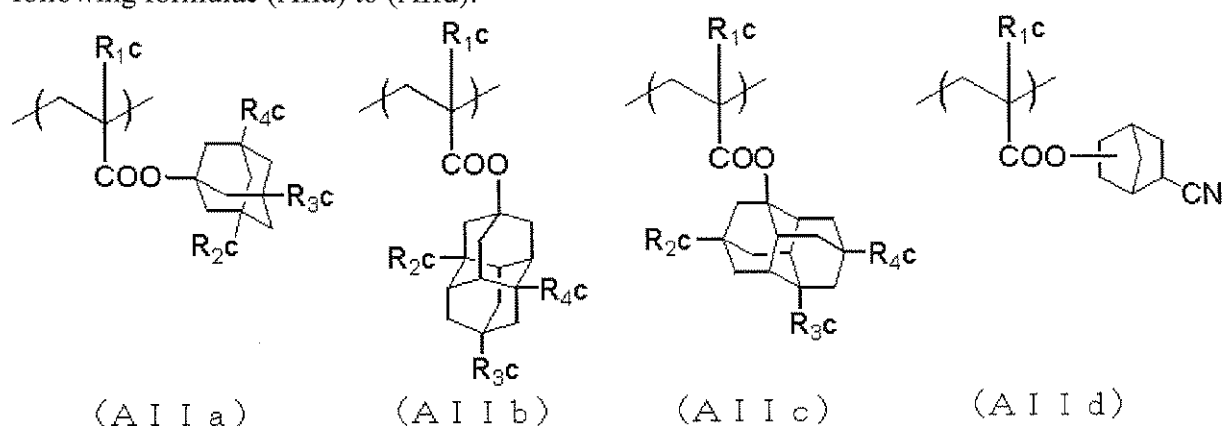
The polar group-substituted alicyclic hydrocarbon structure is preferably a partial structure represented by the following formulae (VIIa) to (VIId):



In formulae (VIIa) to (VIIc), each of R_{2c} to R_{4c} independently represents a hydrogen atom, a hydroxyl group or a cyano group, provided that at least one of R_{2c} to R_{4c} represents a hydroxyl group or a cyano group. A structure where one or two members out of R_{2c} to R_{4c} are a hydroxyl group with the remaining being a hydrogen atom is preferred.

In formula (VIIa), it is more preferred that two members out of R_{2c} to R_{4c} are a hydroxyl group and the remaining is a hydrogen atom.

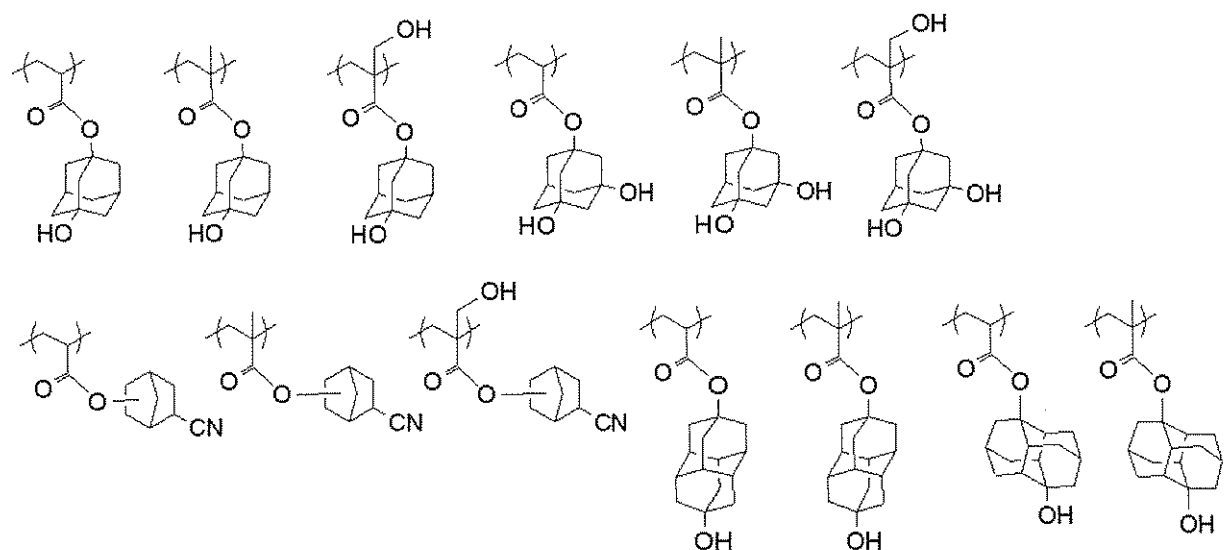
The repeating unit having a group represented by any one of formulae (VIIa) to (VIIId) includes a repeating unit where at least one of $R_{13'}$ to $R_{16'}$ in formula (II-AB1) or (II-AB2) has a group represented by formula (VII) (for example, R_5 of $-\text{COOR}_5$ is a group represented by any one of formulae (VIIa) to (VIIId)), and repeating units represented by the following formulae (AIIa) to (AIIId):



In formulae (AIIa) to (AIIId), R_{1c} represents a hydrogen atom, a methyl group, a trifluoromethyl group or a hydroxymethyl group.

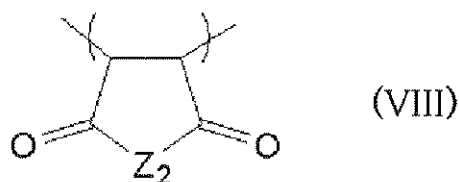
R_{2c} to R_{4c} have the same meanings as R_{2c} to R_{4c} in formulae (VIIa) to (VIIc).

Specific examples of the repeating unit having a structure represented by any one of formulae (AIIa) to (AIIId) are set forth below, but the present invention is not limited thereto.



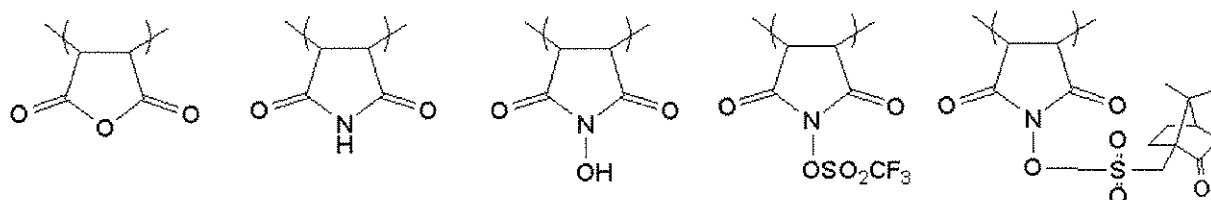
The alicyclic hydrocarbon-based acid-decomposable resin for use in the present

invention may contain a repeating unit represented by the following formula (VIII):



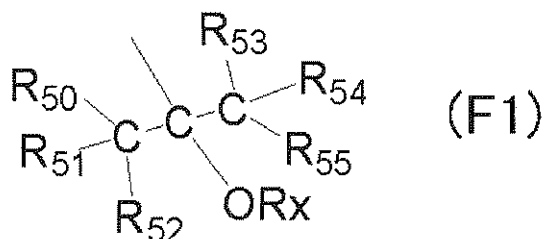
In formula (VIII), Z_2 represents -O- or -N(R₄₁)-. R₄₁ represents a hydrogen atom, a hydroxyl group, an alkyl group or -OSO₂-R₄₂. R₄₂ represents an alkyl group, a cycloalkyl group or a camphor residue. The alkyl group of R₄₁ and R₄₂ may be substituted by a halogen atom (preferably fluorine atom) or the like.

Specific examples of the repeating unit represented by formula (VIII) are set forth below, but the present invention is not limited thereto.



The alicyclic hydrocarbon-based acid-decomposable resin for use in the present invention preferably contains a repeating unit having an alkali-soluble group, more preferably a repeating unit having a carboxyl group. By virtue of containing this repeating unit, the resolution increases in the usage of forming contact holes. As for the repeating unit having a carboxyl group, a repeating unit where a carboxyl group is directly bonded to the resin main chain, such as repeating unit by an acrylic acid or a methacrylic acid, a repeating unit where a carboxyl group is bonded to the resin main chain through a linking group, and a repeating unit where a carboxyl group is introduced into the terminal of the polymer chain by using a polymerization initiator or chain transfer agent having an alkali-soluble group at the polymerization, all are preferred. The linking group may have a monocyclic or polycyclic hydrocarbon structure. A repeating unit by an acrylic acid or a methacrylic acid is more preferred.

The alicyclic hydrocarbon-based acid-decomposable resin for use in the present invention may further contain a repeating unit having from 1 to 3 groups represented by formula (F1). Thanks to this repeating unit, the performance in terms of line edge roughness is enhanced.



In formula (F1), each of R₅₀ to R₅₅ independently represents a hydrogen atom, a fluorine atom or an alkyl group, provided that at least one of R₅₀ to R₅₅ is a fluorine atom or an alkyl group with at least one hydrogen atom being substituted by a fluorine atom.

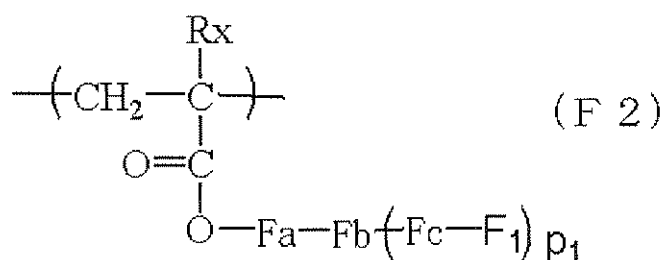
Rx represents a hydrogen atom or an organic group (preferably an acid-decomposable protective group, an alkyl group, a cycloalkyl group, an acyl group or an alkoxycarbonyl group).

The alkyl group of R₅₀ to R₅₅ may be substituted by a halogen atom (e.g., fluorine), a cyano group or the like, and the alkyl group is preferably an alkyl group having a carbon number of 1 to 3, such as methyl group and trifluoromethyl group.

It is preferred that R₅₀ to R₅₅ all are a fluorine atom.

The organic group represented by Rx is preferably an acid-decomposable protective group, an alkyl group, a cycloalkyl group, an acyl group, an alkylcarbonyl group, an alkoxycarbonyl group, an alkoxycarbonylmethyl group, an alkoxymethyl group or a 1-alkoxyethyl group, each of which may have a substituent.

The repeating unit having a group represented by formula (F1) is preferably a repeating unit represented by the following formula (F2):



In formula (F2), Rx represents a hydrogen atom, a halogen atom or an alkyl group having a carbon number of 1 to 4. Preferred examples of the substituent which the alkyl group of Rx may have include a hydroxyl group and a halogen atom.

Fa represents a single bond or a linear or branched alkylene group and is preferably a single bond.

Fb represents a monocyclic or polycyclic hydrocarbon group.

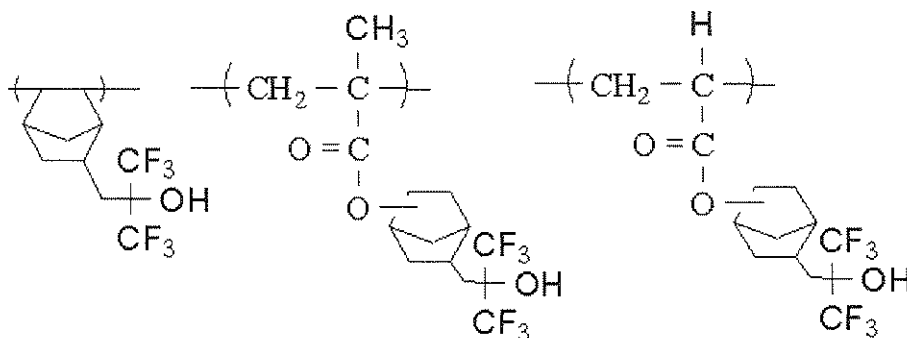
Fc represents a single bond or a linear or branched alkylene group and is preferably a single bond or a methylene group.

F₁ represents a group represented by formula (F1).

p_1 represents a number of 1 to 3.

The cyclic hydrocarbon group in Fb is preferably a cyclopentyl group, a cyclohexyl group or a norbornyl group.

Specific examples of the repeating unit having a group represented by formula (F1) are set forth below, but the present invention is not limited thereto.



The alicyclic hydrocarbon-based acid-decomposable resin for use in the present invention may further contain a repeating unit having an alicyclic hydrocarbon structure and not exhibiting acid decomposability. Thanks to this repeating unit, the dissolving out of low molecular components from the resist film to the immersion liquid at the immersion exposure can be reduced. Examples of this repeating unit include 1-adamantyl (meth)acrylate, tricyclodecanyl (meth)acrylate and cyclohexyl (meth)acrylate.

The alicyclic hydrocarbon-based acid-decomposable resin for use in the present invention may contain, in addition to the above-described repeating units, various repeating structural units for the purpose of controlling dry etching resistance, suitability for standard developer, adherence to substrate, resist profile and properties generally required of the resist, such as resolution, heat resistance and sensitivity.

Examples of such a repeating structural unit include, but are not limited to, repeating structural units corresponding to the monomers described below.

Thanks to such a repeating structural unit, the performance required of the alicyclic hydrocarbon-based acid-decomposable resin, particularly, (1) solubility in coating solvent, (2) film-forming property (glass transition point), (3) solubility in positive or negative tone developer, (4) film loss (selection of hydrophilic, hydrophobic and alkali-soluble group), (5) adherence of unexposed area to substrate, (6) dry etching resistance, and the like, can be subtly controlled.

Examples of the monomer include a compound having one addition-polymerizable unsaturated bond selected from acrylic acid esters, methacrylic acid esters, acrylamides, methacrylamides, allyl compounds, vinyl ethers and vinyl esters.

Other than these, an addition-polymerizable unsaturated compound copolymerizable with the monomers corresponding to the above-described various repeating structural units

may be copolymerized.

In the alicyclic hydrocarbon-based acid-decomposable resin, the molar ratio of respective repeating structural units contained is appropriately determined to control the dry etching resistance of resist, suitability for standard developer, adherence to substrate, resist profile and performances generally required of the resist, such as resolution, heat resistance and sensitivity.

The preferred embodiment of the alicyclic hydrocarbon-based acid-decomposable resin for use in the present invention includes the followings:

(1) a resin containing a repeating unit having an alicyclic hydrocarbon-containing partial structure represented by any one of formulae (pI) to (pV) (side chain type), preferably a resin containing a (meth)acrylate repeating unit having a structure represented by any one of formulae (pI) to (pV), and

(2) a resin containing a repeating unit represented by formula (II-AB) (main chain type).

The resin of (2) further includes:

(3) a resin having a repeating unit represented by formula (II-AB), a maleic anhydride derivative and a (meth)acrylate structure (hybrid type).

In the alicyclic hydrocarbon-based acid-decomposable resin, the content of the repeating unit having an acid-decomposable group is preferably from 10 to 60 mol%, more preferably from 20 to 50 mol%, still more preferably from 30 to 50 mol%, based on all repeating structural units.

In the alicyclic hydrocarbon-based acid-decomposable resin, the content of the repeating unit having an alicyclic hydrocarbon-containing partial structure represented by any one of formulae (pI) to (pV) is preferably from 20 to 70 mol%, more preferably from 20 to 50 mol%, still more preferably from 30 to 50 mol%, based on all repeating structural units.

In the alicyclic hydrocarbon-based acid-decomposable resin, the content of the repeating unit represented by formula (II-AB) is preferably from 10 to 60 mol%, more preferably from 15 to 55 mol%, still more preferably from 20 to 50 mol%, based on all repeating structural units.

In the acid-decomposable resin, the content of the repeating unit having a lactone ring is preferably from 10 to 70 mol%, more preferably from 20 to 60 mol%, still more preferably from 25 to 50 mol%, based on all repeating structural units.

In the acid-decomposable resin, the content of the repeating unit having a polar group-containing organic group is preferably from 1 to 40 mol%, more preferably from 5 to 30 mol%, still more preferably from 5 to 20 mol%, based on all repeating structural units.

The content of the repeating structural unit based on the monomer as the further

copolymerization component in the resin can also be appropriately selected according to the desired resist performance but in general, the content thereof is preferably 99 mol% or less, more preferably 90 mol% or less, still more preferably 80 mol% or less, based on the total molar number of the repeating structural unit having an alicyclic hydrocarbon-containing partial structure represented by any one of formulae (pI) to (pV) and the repeating unit represented by formula (II-AB).

In the case of using the resist composition of the present invention for ArF exposure, the resin preferably has no aromatic group in view of transparency to ArF light.

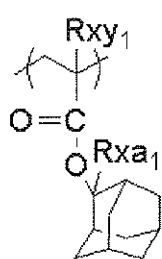
The alicyclic hydrocarbon-based acid-decomposable resin for use in the present invention is preferably a resin where all repeating units are composed of a (meth)acrylate-based repeating unit. In this case, the repeating units may be all a methacrylate-based repeating unit, all an acrylate-based repeating unit, or all a mixture of methacrylate-based repeating unit/acrylate-based repeating unit, but the acrylate-based repeating unit preferably accounts for 50 mol% or less of all repeating units.

The alicyclic hydrocarbon-based acid-decomposable resin is preferably a copolymer having at least three kinds of repeating units, that is, a (meth)acrylate-based repeating unit having a lactone ring, a (meth)acrylate-based repeating unit having an organic group substituted by at least either a hydroxyl group or a cyano group, and a (meth)acrylate-based repeating unit having an acid-decomposable group.

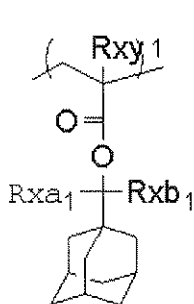
The copolymer is preferably a ternary copolymerization polymer containing from 20 to 50 mol% of the repeating unit having an alicyclic hydrocarbon-containing partial structure represented by any one of formulae (pI) to (pV), from 20 to 50 mol% of the repeating unit having a lactone structure and from 5 to 30 mol% of the repeating unit having a polar group-substituted alicyclic hydrocarbon structure, or a quaternary copolymerization polymer further containing from 0 to 20 mol% of other repeating units.

In particular, the resin is preferably a ternary copolymerization polymer containing from 20 to 50 mol% of an acid-decomposable group-containing repeating unit represented by any one of the following formulae (ARA-1) to (ARA-5), from 20 to 50 mol% of a lactone group-containing repeating unit represented by any one of the following formulae (ARL-1) to (ARL-6), and from 5 to 30 mol% of a repeating unit having a polar group-substituted alicyclic hydrocarbon structure represented by any one of the following formulae (ARH-1) to (ARH-3), or a quaternary copolymerization polymer further containing from 5 to 20 mol% of a repeating unit having a carboxyl group or a structure represented by formula (F1) and a repeating unit having an alicyclic hydrocarbon structure and not exhibiting acid decomposability.

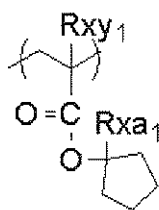
(In the formulae, Rxy₁ represents a hydrogen atom or a methyl group, and Rxa₁ and Rxb₁ each represents a methyl group or an ethyl group).



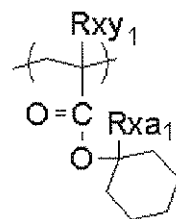
ARA-1



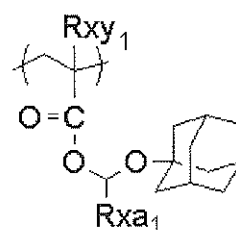
ARA-2



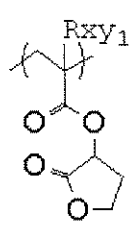
ARA-3



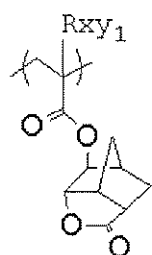
ARA-4



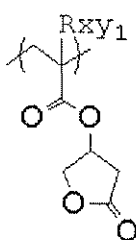
ARA-5



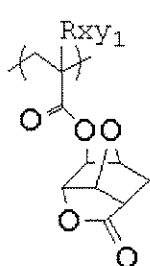
ARL-1



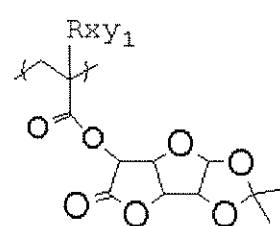
ARL-2



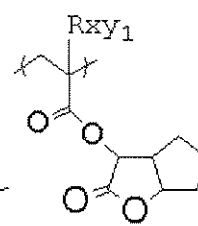
ARL-3



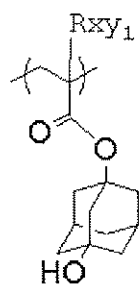
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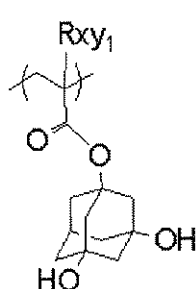
ARL-5



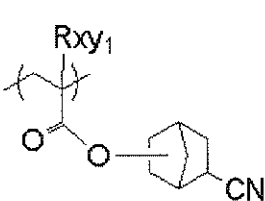
ARL-6



ARH-1



ARH-2



ARH-3

The alicyclic hydrocarbon-based acid-decomposable resin for use in the present invention can be synthesized by an ordinary method (for example, radical polymerization). Examples of the synthesis method in general include a batch polymerization method of dissolving the monomer species and an initiator in a solvent and heating the solution, thereby effecting the polymerization, and a dropping polymerization method of adding dropwise a solution containing monomer species and an initiator to a heated solvent over 1 to 10 hours. A dropping polymerization method is preferred. Examples of the reaction solvent include tetrahydrofuran, 1,4-dioxane, ethers such as diisopropyl ether, ketones such as methyl ethyl ketone and methyl isobutyl ketone, an ester solvent such as ethyl acetate, an amide solvent such as dimethylformamide and dimethylacetamide, and the later-described solvent capable of dissolving the composition of the present invention, such as propylene glycol monomethyl ether acetate, propylene glycol monomethyl ether and cyclohexanone. The polymerization

is more preferably performed using the same solvent as the solvent used in the resist composition of the present invention. By the use of this solvent, production of particles during storage can be suppressed.

The polymerization reaction is preferably performed in an inert gas atmosphere such as nitrogen and argon. As for the polymerization initiator, the polymerization is started using a commercially available radical initiator (e.g., azo-based initiator, peroxide). The radical initiator is preferably an azo-based initiator, and an azo-based initiator having an ester group, a cyano group or a carboxyl group is preferred. Preferred examples of the initiator include azobisisobutyronitrile, azobisdimethylvaleronitrile and dimethyl 2,2'-azobis(2-methylpropionate). The initiator is added additionally or in parts, if desired. After the completion of reaction, the reaction product is charged into a solvent, and the desired polymer is recovered by a method such as powder or solid recovery. The reaction concentration is from 5 to 50 mass%, preferably from 10 to 30 mass%.

The reaction temperature is usually from 10 to 150°C, preferably from 30 to 120°C, more preferably from 60 to 100°C.

The purification may be performed by the same method as that for the resin (C) described later, and a normal method, for example, a liquid-liquid extraction method of combining water washing with an appropriate solvent to remove residual monomers or oligomer components, a purification method in a solution state, such as ultrafiltration of extracting and removing only polymers having a molecular weight lower than a specific molecular weight, a reprecipitation method of adding dropwise the resin solution in a poor solvent to solidify the resin in the poor solvent and thereby remove residual monomers and the like, or a purification method in a solid state, such as washing of the resin slurry with a poor solvent after separation by filtration, may be employed.

The weight average molecular weight of the resin for use in the present invention is preferably 6,000 or less, more preferably from 1,000 to 5,000, still more preferably from 2,000 to 3,500, in terms of polystyrene by the GPC method. When the weight average molecular weight of the resin is 6,000 or less, the line edge roughness of the resist pattern is reduced.

The dispersity (molecular weight distribution, M_w/M_n) is preferably 1.7 or less, more preferably 1.5 or less, still more preferably 1.3 or less. As the dispersity is smaller, the line edge roughness of the resist pattern is more reduced.

To speak in more detail, the resin for use in the present invention is (a1) a resin having an alicyclic hydrocarbon structure and a dispersity of 1.7 or less and being capable of increasing the polarity by the action of an acid, or (a2) a resin having an alicyclic hydrocarbon structure and a weight average molecular weight of 6,000 or less and being capable of

increasing the polarity by the action of an acid, preferably (a3) a resin having an alicyclic hydrocarbon structure, a dispersity of 1.7 or less and a weight average molecular weight of 6,000 or less and being capable of increasing the polarity by the action of an acid.

The molecular weight can be controlled by adjusting the amount added of the radical initiator, and as the amount added of the initiator material is larger, the molecular weight becomes smaller. As for the control of dispersity, a living polymerization method or the like disclosed in JP-A-2004-220009 may be used, and in addition, the dispersity can also be reduced by raising the proportion of the good solvent for the resin during precipitation when synthesizing the resin.

In the resin for use in the present invention, by combining the weight average molecular weight and the dispersity in respective preferred ranges, the line edge roughness can be more reduced.

In the resist composition of the present invention, the amount of all resins for use in the present invention blended in the entire composition is preferably from 50 to 99.9 mass%, more preferably from 60 to 99.0 mass%, based on the entire solid content.

In the present invention, one kind of a resin may be used or a plurality of kinds of resins may be used in combination.

The alicyclic hydrocarbon-based acid-decomposable resin for use in the present invention preferably contains no fluorine or silicon atom in view of compatibility with the resin (D).

(B) Compound capable of generating an acid upon irradiation with an actinic ray or radiation

The resist composition of the present invention contains a compound capable of generating an acid upon irradiation with an actinic ray or radiation (sometimes referred to as a "photo-acid generator" or "component (B)").

The photo-acid generator which can be used may be appropriately selected from a photo-initiator for cationic photopolymerization, a photo-initiator for radical photopolymerization, a photodecoloring agent for dyes, a photodiscoloring agent, a known compound used for microresist or the like and capable of generating an acid upon irradiation with an actinic ray or radiation, and a mixture thereof.

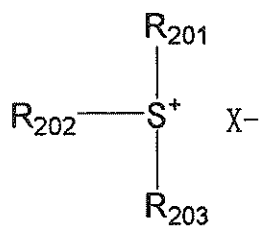
Examples thereof include a diazonium salt, a phosphonium salt, a sulfonium salt, an iodonium salt, an imidosulfonate, an oxime sulfonate, a diazodisulfone, a disulfone and an o-nitrobenzyl sulfonate.

Also, a compound where such a group or compound capable of generating an acid upon irradiation with an actinic ray or radiation is introduced into the main or side chain of the polymer, for example, compounds described in U.S. Patent 3,849,137, German Patent 3,914,407, JP-A-63-26653, JP-A-55-164824, JP-A-62-69263, JP-A-63-146038, JP-A-63-

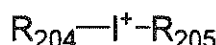
163452, JP-A-62-153853 and JP-A-63-146029, may be used.

Furthermore, compounds capable of generating an acid by the effect of light described, for example, in U.S. Patent 3,779,778 and European Patent 126,712 may also be used.

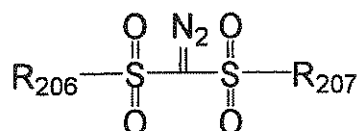
Out of the compounds capable of generating an acid upon irradiation with an actinic ray or radiation, preferred are the compounds represented by the following formulae (ZI), (ZII) and (ZIII):



ZI



ZII

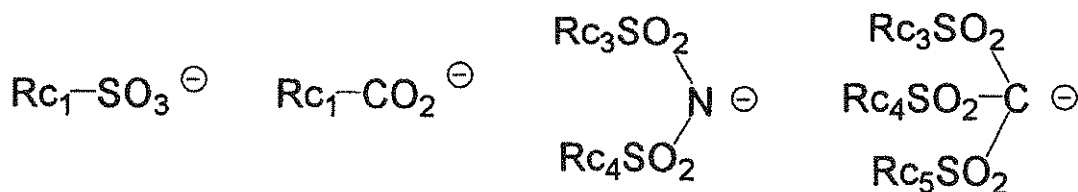


ZIII

In formula (ZI), each of R_{201} , R_{202} and R_{203} independently represents an organic group.

X^- represents a non-nucleophilic anion, and preferred examples thereof include sulfonate anion, carboxylate anion, bis(alkylsulfonyl)amide anion, tris(alkylsulfonyl)methide anion, BF_4^- , PF_6^- and SbF_6^- . The anion is preferably an organic anion containing a carbon atom.

The preferred organic anion includes organic anions represented by the following formulae:



In the formulae, Rc_1 represents an organic group.

The organic group of Rc_1 includes an organic group having a carbon number of 1 to 30, and preferred examples thereof include an alkyl group which may be substituted, an aryl group which may be substituted, and a group where a plurality of these groups are connected through a single bond or a linking group such as $-\text{O}-$, $-\text{CO}_2-$, $-\text{S}-$, $-\text{SO}_3-$ and $-\text{SO}_2\text{N}(\text{Rd}_1)-$. Rd_1 represents a hydrogen atom or an alkyl group.

Each of Rc_3 , Rc_4 and Rc_5 independently represents an organic group. Preferred organic groups of Rc_3 , Rc_4 and Rc_5 are the same as preferred organic groups in Rc_1 . The organic group is most preferably a perfluoroalkyl group having a carbon number of 1 to 4.

Rc₃ and Rc₄ may combine to form a ring. The group formed by combining Rc₃ and Rc₄ includes an alkylene group and an arylene group, and a perfluoroalkylene group having a carbon number of 2 to 4 is preferred.

The organic group of Rc₁ and Rc₃ to Rc₅ is particularly preferably an alkyl group with the 1-position being substituted by a fluorine atom or a fluoroalkyl group, or a phenyl group substituted by a fluorine atom or a fluoroalkyl group. By virtue of having a fluorine atom or a fluoroalkyl group, the acidity of the acid generated upon irradiation with light is raised and the sensitivity is enhanced. Also, when Rc₃ and Rc₄ are combined to form a ring, the acidity of the acid generated upon irradiation with light is raised and the sensitivity is enhanced.

The carbon number of the organic group as R₂₀₁, R₂₀₂ and R₂₀₃ is generally from 1 to 30, preferably from 1 to 20.

Two members out of R₂₀₁ to R₂₀₃ may combine to form a ring structure, and the ring may contain an oxygen atom, a sulfur atom, an ester bond, an amide bond or a carbonyl group. Examples of the group formed by combining two members out of R₂₀₁ to R₂₀₃ include an alkylene group (e.g., butylene, pentylene).

Specific examples of the organic group as R₂₀₁, R₂₀₂ and R₂₀₃ include corresponding groups in the compounds (ZI-1), (ZI-2) and (ZI-3) which are described later.

The compound may be a compound having a plurality of structures represented by formula (ZI). For example, the compound may be a compound having a structure where at least one of R₂₀₁ to R₂₀₃ in the compound represented by formula (ZI) is bonded to at least one of R₂₀₁ to R₂₀₃ in another compound represented by formula (ZI).

The component (ZI) is more preferably a compound (ZI-1), (ZI-2) or (ZI-3) described below.

The compound (ZI-1) is an arylsulfonium compound where at least one of R₂₀₁ to R₂₀₃ in formula (ZI) is an aryl group, that is, a compound having an arylsulfonium as the cation.

In the arylsulfonium compound, R₂₀₁ to R₂₀₃ all may be an aryl group or a part of R₂₀₁ to R₂₀₃ may be an aryl group with the remaining being an alkyl group or a cycloalkyl group.

Examples of the arylsulfonium compound include a triarylsulfonium compound, a diarylalkylsulfonium compound, an aryldialkylsulfonium compound, a diarylcycloalkylsulfonium compound and an aryldicycloalkylsulfonium compound.

The aryl group in the arylsulfonium compound is preferably an aryl group such as phenyl group and naphthyl group, or a heteroaryl group such as indole residue and pyrrole residue, more preferably a phenyl group or an indole residue. In the case where the

arylsulfonium compound has two or more aryl groups, these two or more aryl groups may be the same or different.

The alkyl group which is present, if desired, in the arylsulfonium compound is preferably a linear or branched alkyl group having a carbon number of 1 to 15, and examples thereof include a methyl group, an ethyl group, a propyl group, an n-butyl group, a sec-butyl group and a tert-butyl group.

The cycloalkyl group which is present, if desired, in the arylsulfonium compound is preferably a cycloalkyl group having a carbon number of 3 to 15, and examples thereof include a cyclopropyl group, a cyclobutyl group and a cyclohexyl group.

The aryl group, alkyl group and cycloalkyl group of R_{201} to R_{203} each may have, as the substituent, an alkyl group (for example, an alkyl group having a carbon number of 1 to 15), a cycloalkyl group (for example, a cycloalkyl group having a carbon number of 3 to 15), an aryl group (for example, an aryl group having a carbon number of 6 to 14), an alkoxy group (for example, an alkoxy group having a carbon number of 1 to 15), a halogen atom, a hydroxyl group or a phenylthio group. The substituent is preferably a linear or branched alkyl group having a carbon number of 1 to 12, a cycloalkyl group having a carbon number of 3 to 12, or a linear, branched or cyclic alkoxy group having a carbon number of 1 to 12, more preferably an alkyl group having a carbon number of 1 to 4 or an alkoxy group having a carbon number of 1 to 4. The substituent may be substituted to any one of three members R_{201} to R_{203} or may be substituted to all of these three members. In the case where R_{201} to R_{203} are an aryl group, the substituent is preferably substituted at the p-position of the aryl group.

The compound (ZI-2) is described below. The compound (ZI-2) is a compound where each of R_{201} to R_{203} in formula (ZI) independently represents an aromatic ring-free organic group. The aromatic ring as used herein includes an aromatic ring containing a heteroatom.

The aromatic ring-free organic group as R_{201} to R_{203} generally has a carbon number of 1 to 30, preferably from 1 to 20.

Each of R_{201} to R_{203} is independently preferably an alkyl group, a cycloalkyl group, an allyl group or a vinyl group, more preferably a linear, branched or cyclic 2-oxoalkyl group or an alkoxycarbonylmethyl group, still more preferably a linear or branched 2-oxoalkyl group.

The alkyl group as R_{201} to R_{203} may be either linear or branched and includes a linear or branched alkyl group preferably having a carbon number of 1 to 10 (e.g., methyl, ethyl, propyl, butyl, pentyl). The alkyl group as R_{201} to R_{203} is preferably a linear or branched 2-oxoalkyl group or an alkoxycarbonylmethyl group.

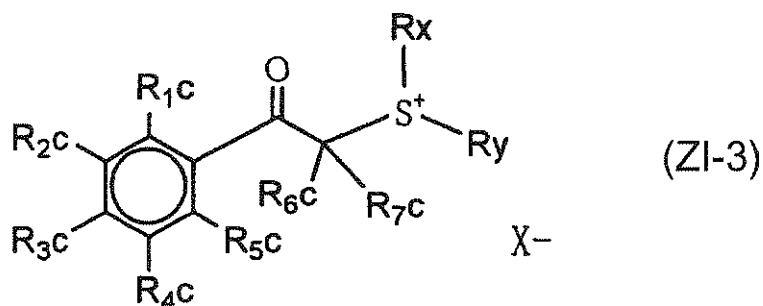
The cycloalkyl group as R_{201} to R_{203} includes a cycloalkyl group preferably having a carbon number of 3 to 10 (e.g., cyclopentyl, cyclohexyl, norbornyl). The cycloalkyl group as R_{201} to R_{203} is preferably a cyclic 2-oxoalkyl group.

The linear, branched or cyclic 2-oxoalkyl group as R_{201} to R_{203} is preferably a group having $>C=O$ at the 2-position of the above-described alkyl or cycloalkyl group.

The alkoxy group in the alkoxycarbonylmethyl group as R_{201} to R_{203} includes an alkoxy group preferably having a carbon number of 1 to 5 (e.g., methoxy, ethoxy, propoxy, butoxy, pentoxy).

R_{201} to R_{203} each may be further substituted by a halogen atom, an alkoxy group (for example, an alkoxy group having a carbon number of 1 to 5), a hydroxyl group, a cyano group or a nitro group.

The compound (ZI-3) is a compound represented by the following formula (ZI-3), and this is a compound having a phenacysulfonium salt structure.



In formula (ZI-3), each of R_{1c} to R_{5c} independently represents a hydrogen atom, an alkyl group, a cycloalkyl group, an alkoxy group or a halogen atom.

Each of R_{6c} and R_{7c} independently represents a hydrogen atom, an alkyl group or a cycloalkyl group.

Each of R_x and R_y independently represents an alkyl group, a cycloalkyl group, an allyl group or a vinyl group.

Any two or more members out of R_{1c} to R_{7c} , and a pair of R_x and R_y , may combine with each other to form a ring structure, respectively, and the ring structure may contain an oxygen atom, a sulfur atom, an ester bond or an amide bond. Examples of the group formed by combining any two or more members out of R_{1c} to R_{7c} or a pair of R_x and R_y include a butylene group and a pentylene group.

X^- represents a non-nucleophilic anion, and examples thereof are the same as those of the non-nucleophilic anion of X^- in formula (ZI).

The alkyl group as R_{1c} to R_{7c} may be linear or branched and includes, for example, a linear or branched alkyl group having a carbon number of 1 to 20, preferably a linear or branched alkyl group having a carbon number of 1 to 12 (such as methyl group, ethyl group,

linear or branched propyl group, linear or branched butyl group and linear or branched pentyl group).

The cycloalkyl group as R_{1c} to R_{7c} includes a cycloalkyl group preferably having a carbon number of 3 to 8 (e.g., cyclopentyl, cyclohexyl).

The alkoxy group as R_{1c} to R_{5c} may be linear, branched or cyclic and includes, for example, an alkoxy group having a carbon number of 1 to 10, preferably a linear or branched alkoxy group having a carbon number of 1 to 5 (such as methoxy group, ethoxy group, linear or branched propoxy group, linear or branched butoxy group and linear or branched pentoxy group), and a cyclic alkoxy group having a carbon number of 3 to 8 (e.g., cyclopentyloxy, cyclohexyloxy).

A compound where any one of R_{1c} to R_{5c} is a linear or branched alkyl group, a cycloalkyl group or a linear, branched or cyclic alkoxy group is preferred, and a compound where the sum of carbon numbers of R_{1c} to R_{5c} is from 2 to 15 is more preferred. Thanks to this construction, the solubility in a solvent is more enhanced and generation of particles during storage is suppressed.

The alkyl group as R_x and R_y is the same as the alkyl group of R_{1c} to R_{7c} . The alkyl group as R_x and R_y is preferably a linear or branched 2-oxoalkyl group or an alkoxycarbonylmethyl group.

The cycloalkyl group as R_x and R_y is the same as the cycloalkyl group of R_{1c} to R_{7c} . The cycloalkyl group as R_x and R_y is preferably a cyclic 2-oxoalkyl group.

The linear, branched or cyclic 2-oxoalkyl group includes a group having $>C=O$ at the 2-position of the alkyl group or cycloalkyl group of R_{1c} to R_{7c} .

The alkoxy group in the alkoxycarbonylmethyl group is the same as the alkoxy group of R_{1c} to R_{5c} .

R_x and R_y each is preferably an alkyl group having a carbon number of 4 or more, more preferably 6 or more, still more preferably 8 or more.

In formulae (ZII) and (ZIII), each of R_{204} to R_{207} independently represents an aryl group, an alkyl group or a cycloalkyl group.

The aryl group of R_{204} to R_{207} is preferably a phenyl group or a naphthyl group, more preferably a phenyl group.

The alkyl group as R_{204} to R_{207} may be linear or branched and includes a linear or branched alkyl group preferably having a carbon number of 1 to 10 (e.g., methyl, ethyl, propyl, butyl, pentyl).

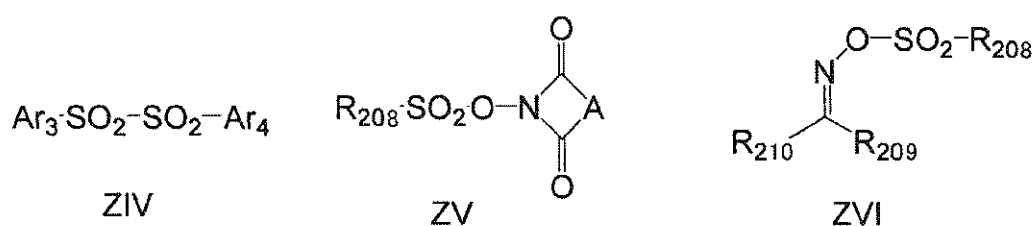
The cycloalkyl group as R_{204} to R_{207} includes a cycloalkyl group preferably having a carbon number of 3 to 10 (e.g., cyclopentyl, cyclohexyl, norbornyl).

R_{204} to R_{207} each may have a substituent. Examples of the substituent which R_{204} to

R₂₀₇ each may have include an alkyl group (for example, an alkyl group having a carbon number of 1 to 15), a cycloalkyl group (for example, a cycloalkyl group having a carbon number of 3 to 15), an aryl group (for example, an aryl group having a carbon number of 6 to 15), an alkoxy group (for example, an alkoxy group having a carbon number of 1 to 15), a halogen atom, a hydroxyl group and a phenylthio group.

X⁻ represents a non-nucleophilic anion and is the same as the non-nucleophilic anion of X⁻ in formula (ZI).

Out of the compounds capable of generating an acid upon irradiation with an actinic ray or radiation, preferred compounds further include the compounds represented by the following formulae (ZIV), (ZV) and (ZVI):



In formulae (ZIV) to (ZVI), each of Ar₃ and Ar₄ independently represents an aryl group.

R₂₀₈ represents an alkyl group or an aryl group.

Each of R₂₀₉ and R₂₁₀ independently represents an alkyl group, an aryl group or an electron-withdrawing group.

R₂₀₈ is preferably an aryl group.

R₂₀₉ is preferably an electron-withdrawing group, more preferably a cyano group or a fluoroalkyl group.

A represents an alkylene group, an alkenylene group or an arylene group.

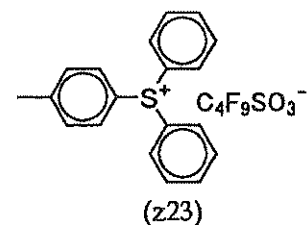
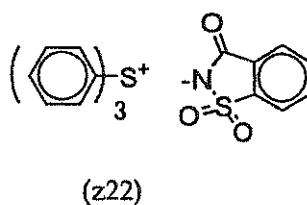
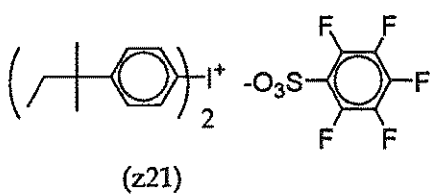
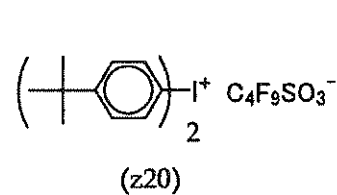
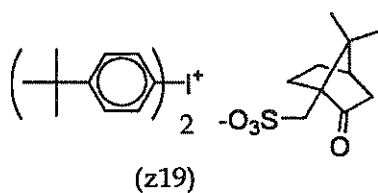
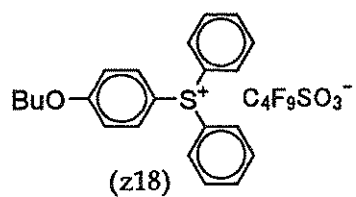
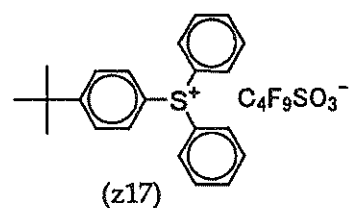
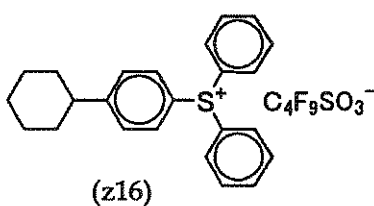
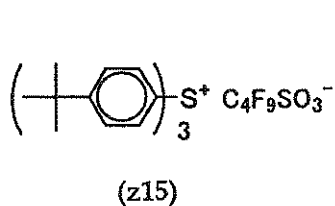
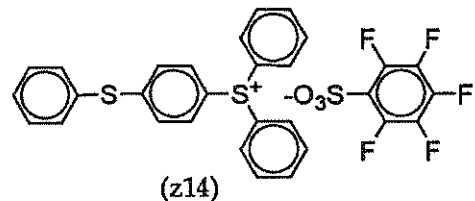
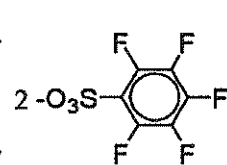
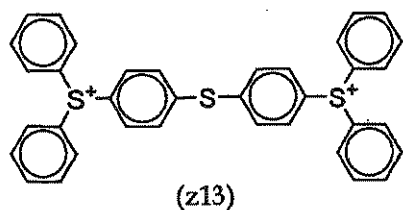
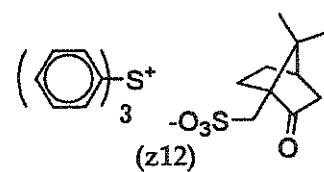
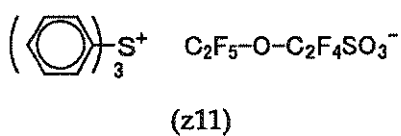
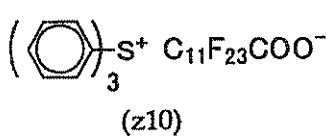
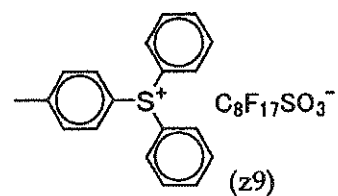
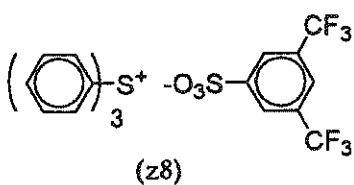
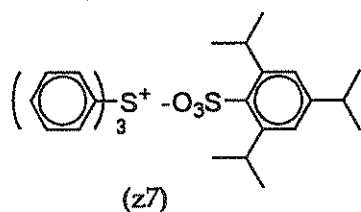
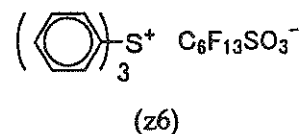
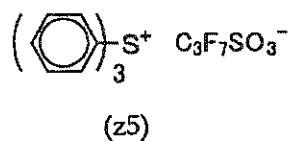
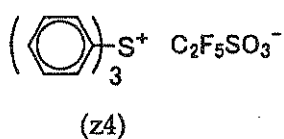
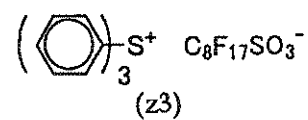
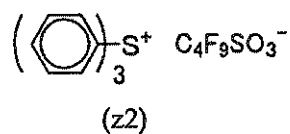
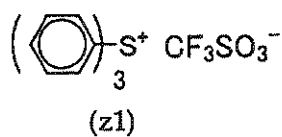
The compound capable of generating an acid upon irradiation with an actinic ray or radiation is preferably a compound represented by any one of formulae (ZI) to (ZIII).

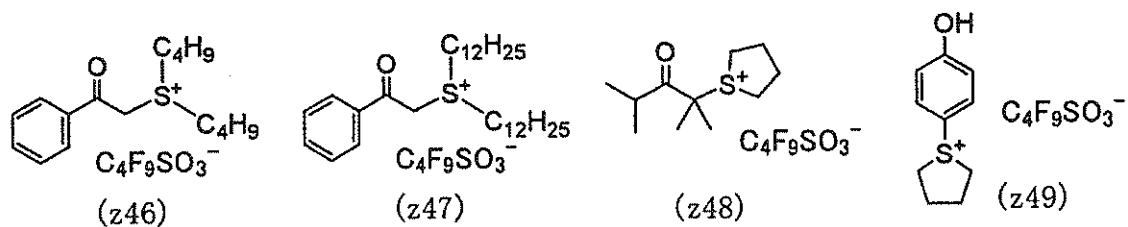
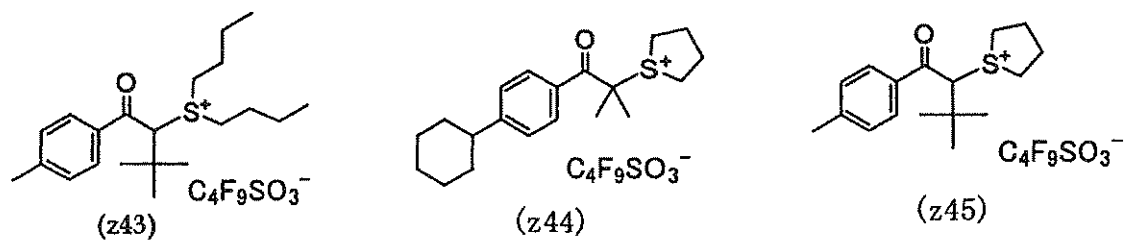
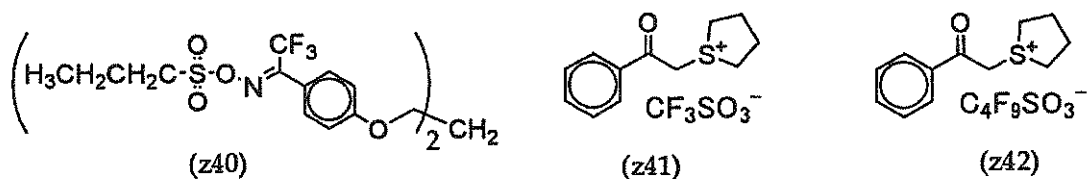
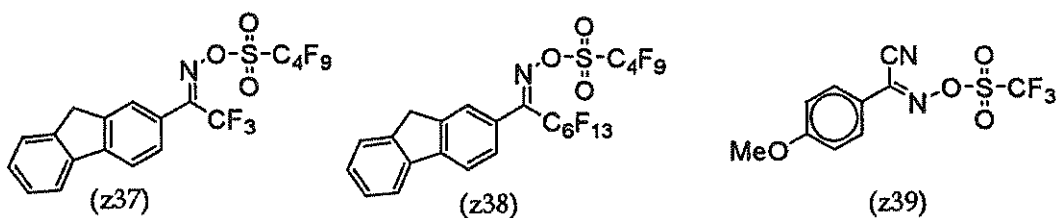
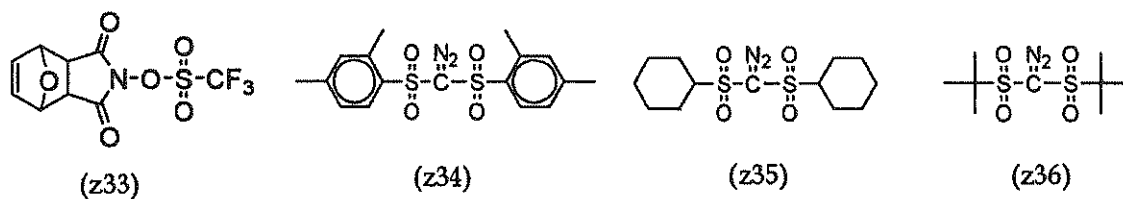
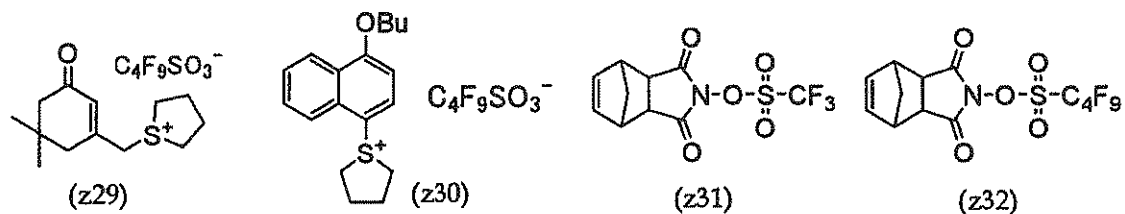
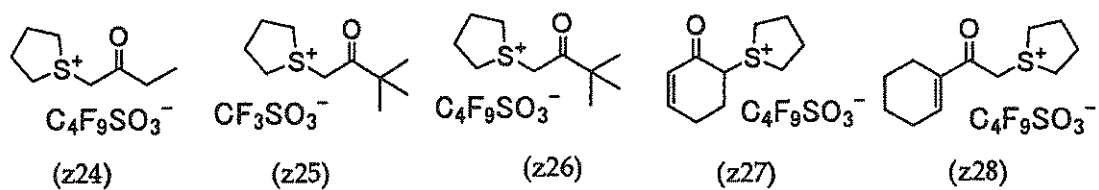
The compound (B) is preferably a compound capable of generating a fluorine atom-containing aliphatic sulfonic acid or a fluorine atom-containing benzenesulfonic acid upon irradiation with an actinic ray or radiation.

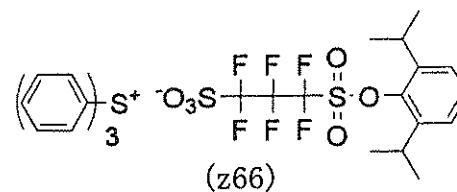
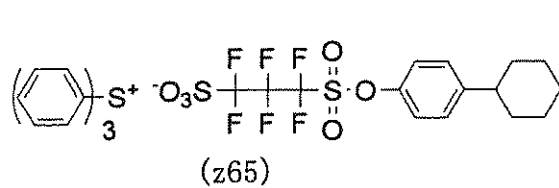
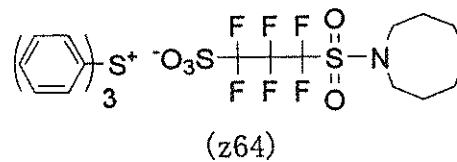
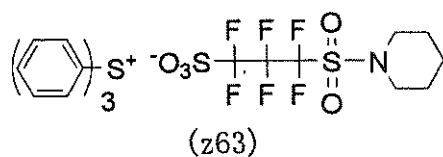
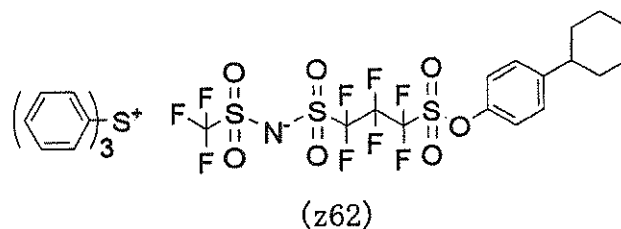
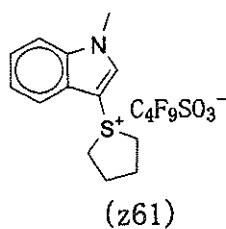
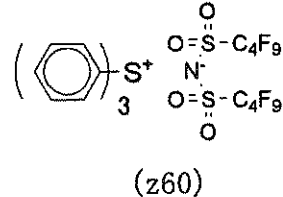
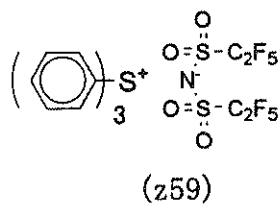
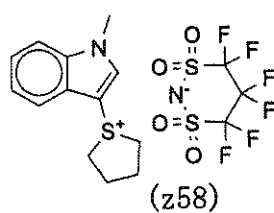
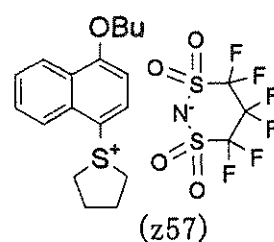
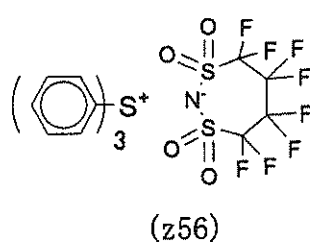
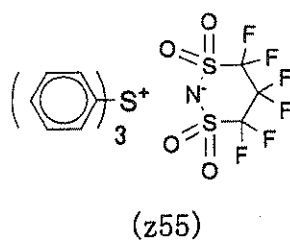
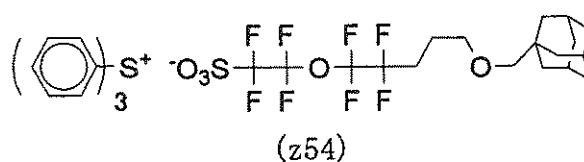
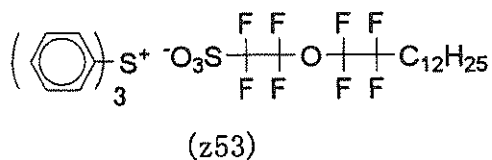
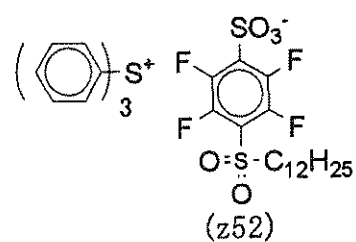
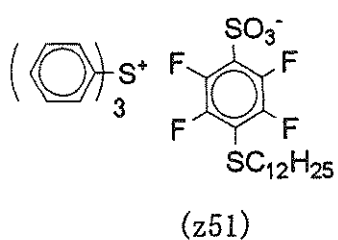
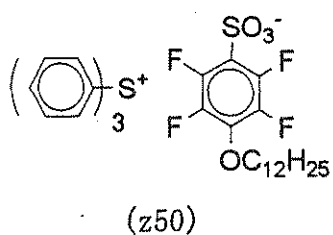
The compound (B) preferably has a triphenylsulfonium structure.

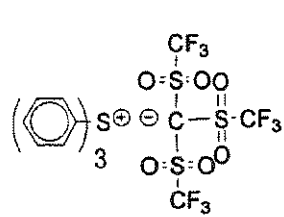
The compound (B) is preferably a triphenylsulfonium salt compound having a fluorine-unsubstituted alkyl or cycloalkyl group in the cation moiety.

Particularly preferred examples out of the compounds capable of generating an acid upon irradiation with an actinic ray or radiation are set forth below.

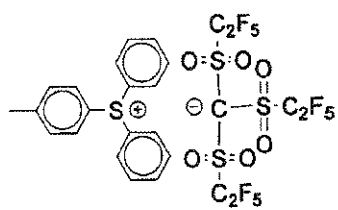




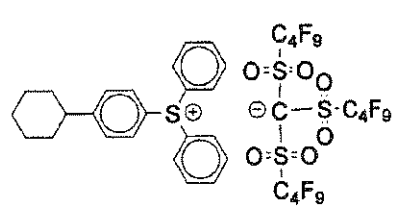




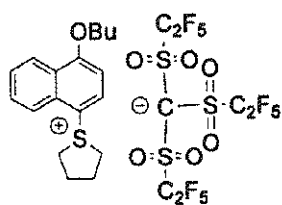
(z67)



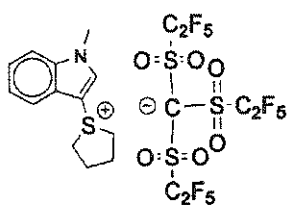
(z68)



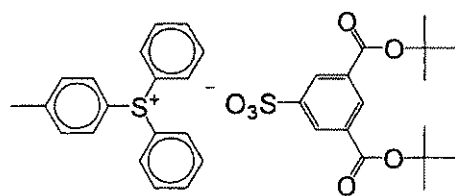
(z69)



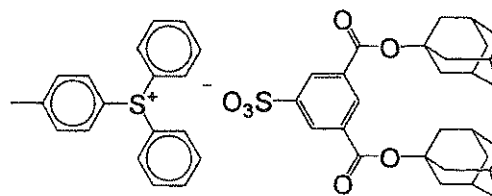
(z70)



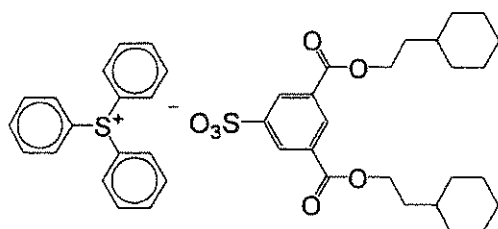
(z71)



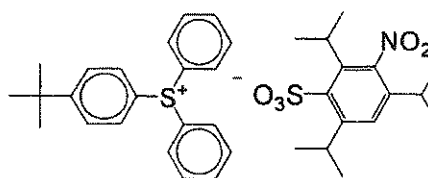
(z72)



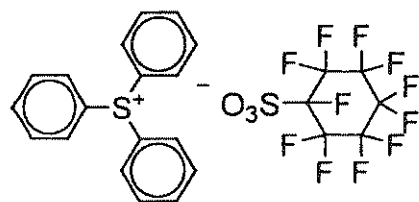
(z73)



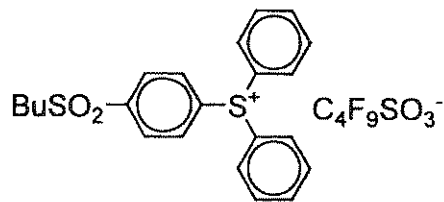
(z74)



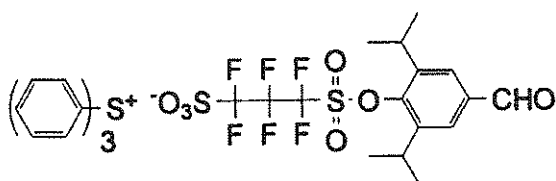
(z75)



(z76)



(z77)



(z78)

One of these photo-acid generators may be used alone, or two or more kinds thereof may be used in combination. In the case of using two or more kinds of photo-acid generators in combination, compounds capable of generating two kinds of organic acids differing in the total atom number except for hydrogen atom by 2 or more are preferably combined.

The content of the photo-acid generator is preferably from 0.1 to 20 mass%, more preferably from 0.5 to 10 mass%, still more preferably from 1 to 7 mass%, based on the entire solid content of the resist composition.

(C) Solvent

Examples of the solvent which can be used for dissolving respective components described above to prepare a positive tone resist composition include an organic solvent such as alkylene glycol monoalkyl ether carboxylate, alkylene glycol monoalkyl ether, alkyl lactate, alkyl alkoxypropionate, cyclic lactone having a carbon number of 4 to 10, monoketone compound having a carbon number of 4 to 10 which may contain a ring, alkylene carbonate, alkyl alkoxyacetate and alkyl pyruvate.

Preferred examples of the alkylene glycol monoalkyl ether carboxylate include propylene glycol monomethyl ether acetate, propylene glycol monoethyl ether acetate, propylene glycol monopropyl ether acetate, propylene glycol monobutyl ether acetate, propylene glycol monomethyl ether propionate, propylene glycol monoethyl ether propionate, ethylene glycol monomethyl ether acetate and ethylene glycol monoethyl ether acetate.

Preferred examples of the alkylene glycol monoalkyl ether include propylene glycol monomethyl ether, propylene glycol monoethyl ether, propylene glycol monopropyl ether, propylene glycol monobutyl ether, ethylene glycol monomethyl ether and ethylene glycol monoethyl ether.

Preferred examples of the alkyl lactate include methyl lactate, ethyl lactate, propyl lactate and butyl lactate.

Preferred examples of the alkyl alkoxypropionate include ethyl 3-ethoxypropionate, methyl 3-methoxypropionate, methyl 3-ethoxypropionate and ethyl 3-methoxypropionate.

Preferred examples of the cyclic lactone having a carbon number of 4 to 10 include β -propiolactone, β -butyrolactone, γ -butyrolactone, α -methyl- γ -butyrolactone, β -methyl- γ -butyrolactone, γ -valerolactone, γ -caprolactone, γ -octanoic lactone and α -hydroxy- γ -

butyrolactone.

Preferred examples of the monoketone compound having a carbon number of 4 to 10 which may contain a ring include 2-butanone, 3-methylbutanone, pinacolone, 2-pentanone, 3-pentanone, 3-methyl-2-pentanone, 4-methyl-2-pentanone, 2-methyl-3-pentanone, 4,4-dimethyl-2-pentanone, 2,4-dimethyl-3-pentanone, 2,2,4,4-tetramethyl-3-pentanone, 2-hexanone, 3-hexanone, 5-methyl-3-hexanone, 2-heptanone, 3-heptanone, 4-heptanone, 2-methyl-3-heptanone, 5-methyl-3-heptanone, 2,6-dimethyl-4-heptanone, 2-octanone, 3-octanone, 2-nonanone, 3-nonanone, 5-nonanone, 2-decanone, 3-decanone, 4-decanone, 5-hexen-2-one, 3-penten-2-one, cyclopentanone, 2-methylcyclopentanone, 3-methylcyclopentanone, 2,2-dimethylcyclopentanone, 2,4,4-trimethylcyclopentanone, cyclohexanone, 3-methylcyclohexanone, 4-methylcyclohexanone, 4-ethylcyclohexanone, 2,2-dimethylcyclohexanone, 2,6-dimethylcyclohexanone, 2,2,6-trimethylcyclohexanone, cycloheptanone, 2-methylcycloheptanone and 3-methylcycloheptanone.

Preferred examples of the alkylene carbonate include propylene carbonate, vinylene carbonate, ethylene carbonate and butylene carbonate.

Preferred examples of the alkyl alkoxyacetate include 2-methoxyethyl acetate, 2-ethoxyethyl acetate, 2-(2-ethoxyethoxy)ethyl acetate, 3-methoxy-3-methylbutyl acetate and 1-methoxy-2-propyl acetate.

Preferred examples of the alkyl pyruvate include methyl pyruvate, ethyl pyruvate and propyl pyruvate.

The solvent which can be preferably used is a solvent having a boiling point of 130°C or more at ordinary temperature under atmospheric pressure, and specific examples thereof include cyclopentanone, γ -butyrolactone, cyclohexanone, ethyl lactate, ethylene glycol monoethyl ether acetate, propylene glycol monomethyl ether acetate, ethyl 3-ethoxypropionate, ethyl pyruvate, 2-ethoxyethyl acetate, 2-(2-ethoxyethoxy)ethyl acetate and propylene carbonate.

In the present invention, one of these solvents may be used alone, or two or more kinds thereof may be used in combination.

In the present invention, a mixed solvent prepared by mixing a solvent containing a hydroxyl group in the structure and a solvent not containing a hydroxyl group may be used as the organic solvent.

Examples of the solvent containing a hydroxyl group include ethylene glycol, ethylene glycol monomethyl ether, ethylene glycol monoethyl ether, propylene glycol, propylene glycol monomethyl ether, propylene glycol monoethyl ether and ethyl lactate. Among these, propylene glycol monomethyl ether and ethyl lactate are preferred.

Examples of the solvent not containing a hydroxyl group include propylene glycol

monomethyl ether acetate, ethyl ethoxypropionate, 2-heptanone, γ -butyrolactone, cyclohexanone, butyl acetate, N-methylpyrrolidone, N,N-dimethylacetamide and dimethylsulfoxide. Among these, propylene glycol monomethyl ether acetate, ethyl ethoxypropionate, 2-heptanone, γ -butyrolactone, cyclohexanone and butyl acetate are preferred, and propylene glycol monomethyl ether acetate, ethyl ethoxypropionate and 2-heptanone are most preferred.

The mixing ratio (by mass) between the solvent containing a hydroxyl group and the solvent not containing a hydroxyl group is from 1/99 to 99/1, preferably from 10/90 to 90/10, more preferably from 20/80 to 60/40. A mixed solvent in which the solvent not containing a hydroxyl group is contained in an amount of 50 mass% or more is preferred in view of coating uniformity.

The solvent is preferably a mixed solvent of two or more kinds of solvents including propylene glycol monomethyl ether acetate.

The entire solid content concentration in the resist composition is generally from 1 to 10 mass%, preferably from 1 to 8.0 mass%, more preferably from 1.0 to 6.0 mass%.

(D) Resin having at least either a fluorine atom or a silicon atom

The resist composition of the present invention preferably contains (D) a resin having at least either a fluorine atom or a silicon atom.

In the resin (D), the fluorine atom or silicon atom may be present in the main chain of the resin or may be substituted to the side chain.

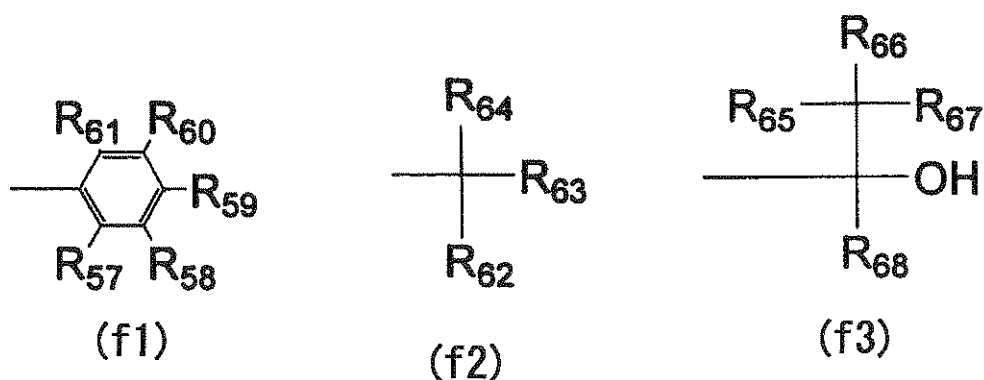
The resin (D) is preferably a resin having a fluorine atom-containing alkyl group, a fluorine atom-containing cycloalkyl group or a fluorine atom-containing aryl group, as a fluorine atom-containing partial structure.

The fluorine atom-containing alkyl group (preferably having a carbon number of 1 to 10, more preferably from 1 to 4) is a linear or branched alkyl group with at least one hydrogen atom being substituted by a fluorine atom and may further have another substituent.

The fluorine atom-containing cycloalkyl group is a monocyclic or polycyclic cycloalkyl group with at least one hydrogen atom being substituted by a fluorine atom and may further have another substituent.

The fluorine atom-containing aryl group is an aryl group (e.g., phenyl, naphthyl) with at least one hydrogen atom being substituted by a fluorine atom and may further have another substituent.

Specific examples of the fluorine atom-containing alkyl group, fluorine atom-containing cycloalkyl group and fluorine atom-containing aryl group are set forth below, but the present invention is not limited thereto.



In formulae (f1) to (f3), each of R_{57} to R_{68} independently represents a hydrogen atom, a fluorine atom or an alkyl group, provided that at least one of R_{57} to R_{61} , at least one of R_{62} to R_{64} and at least one of R_{65} to R_{68} are a fluorine atom or an alkyl group (preferably having a carbon number of 1 to 4) with at least one hydrogen atom being substituted by a fluorine atom. It is preferred that R_{57} to R_{61} and R_{65} to R_{67} all are a fluorine atom. Each of R_{62} , R_{63} and R_{68} is preferably an alkyl group (preferably having a carbon number of 1 to 4) with at least one hydrogen atom being substituted by a fluorine atom, more preferably a perfluoroalkyl group having a carbon number of 1 to 4. R_{62} and R_{63} may combine with each other to form a ring.

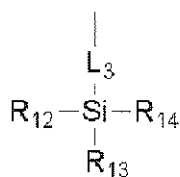
Specific examples of the group represented by formula (f1) include p-fluorophenyl group, pentafluorophenyl group and 3,5-di(trifluoromethyl)phenyl group.

Specific examples of the group represented by formula (f2) include trifluoroethyl group, pentafluoropropyl group, pentafluoroethyl group, heptafluorobutyl group, hexafluoroisopropyl group, heptafluoroisopropyl group, hexafluoro(2-methyl)isopropyl group, nonafluorobutyl group, octafluoroisobutyl group, nonafluorohexyl group, nonafluoro-tert-butyl group, perfluoroisopentyl group, perfluorooctyl group, perfluoro(trimethyl)hexyl group, 2,2,3,3-tetrafluorocyclobutyl group and perfluorocyclohexyl group. Among these, hexafluoroisopropyl group, heptafluoroisopropyl group, hexafluoro(2-methyl)isopropyl group, octafluoroisobutyl group, nonafluoro-tert-butyl group and perfluoroisopentyl group are preferred, and hexafluoroisopropyl group and heptafluoroisopropyl group are more preferred.

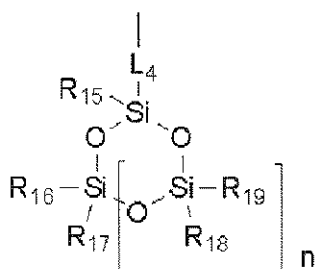
Specific examples of the group represented by formula (f3) include $-C(CF_3)_2OH$, $-C(C_2F_5)_2OH$, $-C(CF_3)(CH_3)OH$ and $-CH(CF_3)OH$, with $-C(CF_3)_2OH$ being preferred.

The resin (D) is preferably a resin having an alkylsilyl structure (preferably a trialkylsilyl group) or a cyclic siloxane structure, as a silicon atom-containing partial structure.

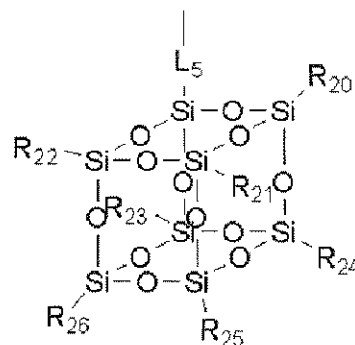
Specific examples of the alkylsilyl structure and cyclic siloxane structure include the groups represented by the following formulae (CS-1) to (CS-3):



(CS-1)



(CS-2)



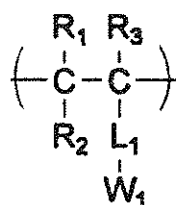
(CS-3)

In formulae (CS-1) to (CS-3), each of R_{12} to R_{26} independently represents a linear or branched alkyl group (preferably having a carbon number of 1 to 20) or a cycloalkyl group (preferably having a carbon number of 3 to 20).

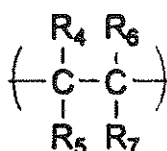
Each of L_3 to L_5 represents a single bond or a divalent linking group. The divalent linking group is a sole group or a combination of two or more groups selected from the group consisting of an alkylene group, a phenyl group, an ether group, a thioether group, a carbonyl group, an ester group, an amide group, a urethane group and a urea group.

n represents an integer of 1 to 5.

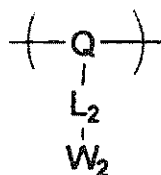
The resin (D) includes a resin containing at least one member selected from the group consisting of repeating units represented by the following formulae (C-I) to (C-V):



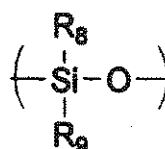
(C-I)



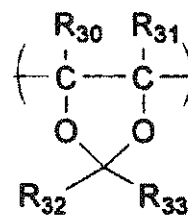
(C-II)



(C-III)



(C-IV)



(C-V)

In formulae (C-I) to (C-V), each of R_1 to R_3 independently represents a hydrogen atom, a fluorine atom, a linear or branched alkyl group having a carbon number of 1 to 4, or a linear or branched fluorinated alkyl group having a carbon number of 1 to 4.

Each of W_1 and W_2 represents an organic group having at least either a fluorine atom or a silicon atom.

Each R_4 to R_7 independently represents a hydrogen atom, a fluorine atom, a linear or branched alkyl group having a carbon number of 1 to 4, or a linear or branched fluorinated alkyl group having a carbon number of 1 to 4, provided that at least one of R_4 to R_7 represents

a fluorine atom. R_4 and R_5 , or R_6 and R_7 may form a ring.

R_8 represents a hydrogen atom or a linear or branched alkyl group having a carbon number of 1 to 4.

R_9 represents a linear or branched alkyl group having a carbon number of 1 to 4, or a linear or branched fluorinated alkyl group having a carbon number of 1 to 4.

Each of L_1 and L_2 represents a single bond or a divalent linking group and is the same as L_3 to L_5 above.

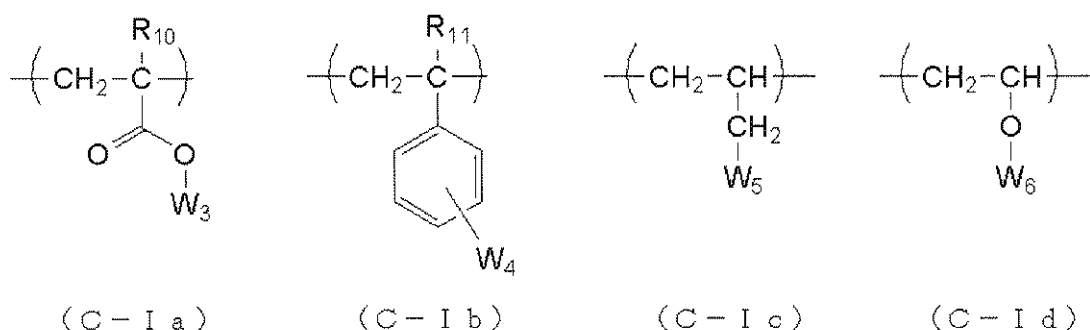
Q represents a monocyclic or polycyclic aliphatic group, that is, an atomic group for forming an alicyclic structure, including the two bonded carbon atoms (C-C).

Each of R_{30} and R_{31} independently represents a hydrogen or fluorine atom.

Each of R_{32} and R_{33} independently represents an alkyl group, a cycloalkyl group, a fluorinated alkyl group or a fluorinated cycloalkyl group.

Here, the repeating unit represented by formula (C-V) has at least one fluorine atom in at least one member out of R_{30} , R_{31} , R_{32} and R_{33} .

The resin (D) preferably has a repeating unit represented by formula (C-I), more preferably a repeating unit represented by any one of the following formulae (C-Ia) to (C-Id):



In formulae (C-Ia) to (C-Id), each of R_{10} and R_{11} represents a hydrogen atom, a fluorine atom, a linear or branched alkyl group having a carbon number of 1 to 4, or a linear or branched fluorinated alkyl group having a carbon number of 1 to 4.

Each of W_3 to W_6 represents an organic group having one or more atoms of at least either a fluorine atom or a silicon atom.

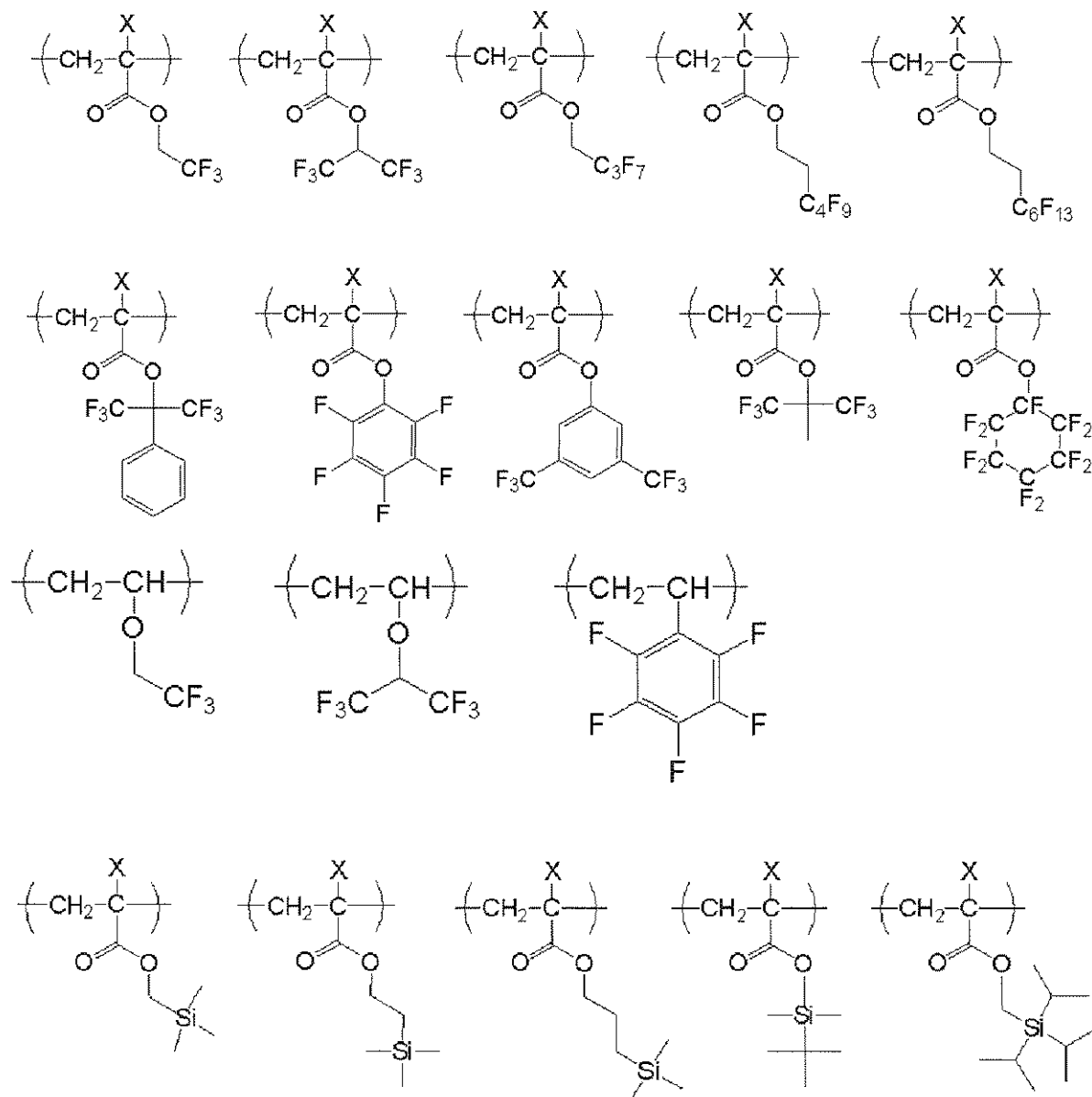
When W_1 to W_6 are an organic group having a fluorine atom, the organic group is preferably a fluorinated linear or branched alkyl group or cycloalkyl group having a carbon number of 1 to 20, or a fluorinated linear, branched or cyclic alkyl ether group having a carbon number of 1 to 20.

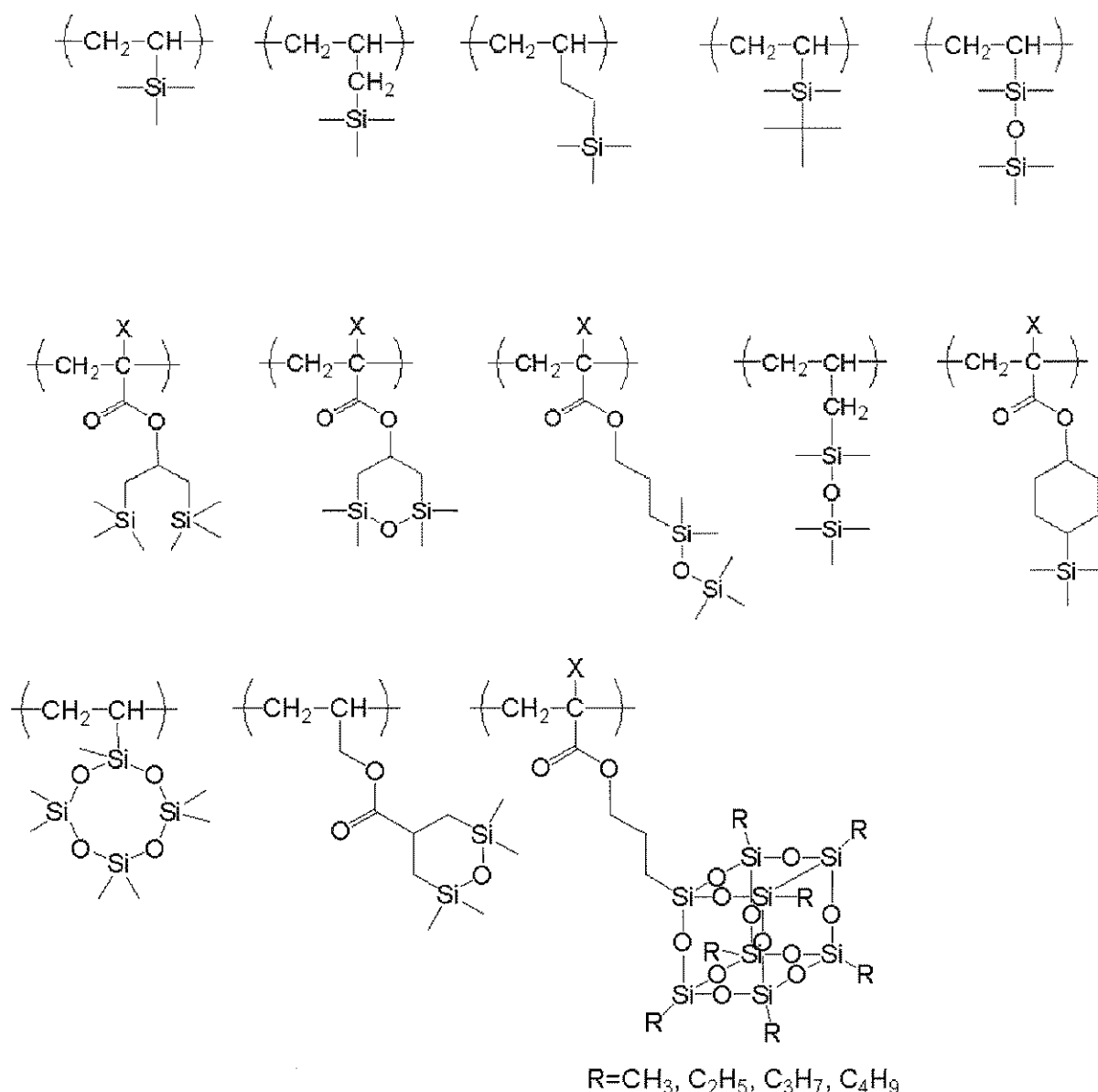
Examples of the fluorinated alkyl group of W_1 to W_6 include a trifluoroethyl group, a pentafluoropropyl group, a hexafluoroisopropyl group, a hexafluoro(2-methyl)isopropyl group, a heptafluorobutyl group, a heptafluoroisopropyl group, an octafluoroisobutyl group, a

nonafluorohexyl group, a nonafluoro-tert-butyl group, a perfluoroisopentyl group, a perfluorooctyl group and a perfluoro(trimethyl)hexyl group.

When W_1 to W_6 are an organic group having a silicon atom, the organic group preferably has an alkylsilyl structure or a cyclic siloxane structure. Specific examples thereof include the groups represented by formulae (CS-1) to (CS-3).

Specific examples of the repeating unit represented by formula (C-I) are set forth below. X represents a hydrogen atom, $-\text{CH}_3$, $-\text{F}$ or $-\text{CF}_3$.





The resin (D) is preferably any one resin selected from the following (D-1) to (D-6):

(D-1) a resin containing (a) a repeating unit having a fluoroalkyl group (preferably having a carbon number of 1 to 4), more preferably a resin containing only the repeating unit (a),

(D-2) a resin containing (b) a repeating unit having a trialkylsilyl group or a cyclic siloxane structure, more preferably a resin containing only the repeating unit (b),

(D-3) a resin containing (a) a repeating unit having a fluoroalkyl group (preferably having a carbon number of 1 to 4) and (c) a repeating unit having a branched alkyl group (preferably having a carbon number of 4 to 20), a cycloalkyl group (preferably having a carbon number of 4 to 20), a branched alkenyl group (preferably having a carbon number of 4 to 20), a cycloalkenyl group (preferably having a carbon number of 4 to 20) or an aryl group

(preferably having a carbon number of 4 to 20), more preferably a copolymerization resin of the repeating unit (a) and the repeating unit (c),

(D-4) a resin containing (b) a repeating unit having a trialkylsilyl group or a cyclic siloxane structure and (c) a repeating unit having a branched alkyl group (preferably having a carbon number of 4 to 20), a cycloalkyl group (preferably having a carbon number of 4 to 20), a branched alkenyl group (preferably having a carbon number of 4 to 20), a cycloalkenyl group (preferably having a carbon number of 4 to 20) or an aryl group (preferably having a carbon number of 4 to 20), more preferably a copolymerization resin of the repeating unit (b) and the repeating unit (c),

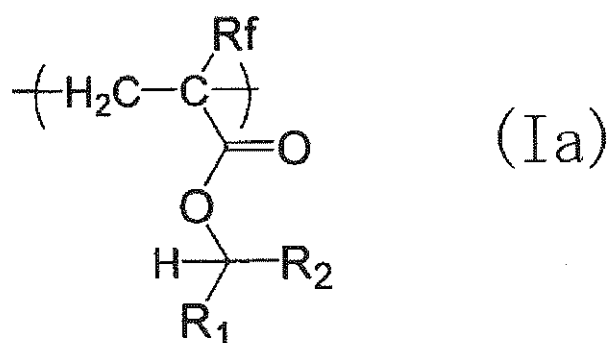
(D-5) a resin containing (a) a repeating unit having a fluoroalkyl group (preferably having a carbon number of 1 to 4) and (b) a repeating unit having a trialkylsilyl group or a cyclic siloxane structure, more preferably a copolymerization resin of the repeating unit (a) and the repeating unit (b), and

(D-6) a resin containing (a) a repeating unit having a fluoroalkyl group (preferably having a carbon number of 1 to 4), (b) a repeating unit having a trialkylsilyl group or a cyclic siloxane structure, and (c) a repeating unit having a branched alkyl group (preferably having a carbon number of 4 to 20), a cycloalkyl group (preferably having a carbon number of 4 to 20), a branched alkenyl group (preferably having a carbon number of 4 to 20), a cycloalkenyl group (preferably having a carbon number of 4 to 20) or an aryl group (preferably having a carbon number of 4 to 20), more preferably a copolymerization resin of the repeating unit (a), the repeating unit (b) and the repeating unit (c).

As for the repeating unit (c) having a branched alkyl group, a cycloalkyl group, a branched alkenyl group, a cycloalkenyl group or an aryl group in the resins (D-3), (D-4) and (D-6), an appropriate functional group can be introduced considering the hydrophilicity/hydrophobicity, interaction and the like, but in view of followability of immersion liquid or receding contact angle, a functional group having no polar group is preferred.

In the resins (D-3), (D-4) and (D-6), the content of the repeating unit (a) having a fluoroalkyl group and/or the repeating unit (b) having a trialkylsilyl group or a cyclic siloxane structure is preferably from 20 to 99 mol%.

The resin (D) is preferably a resin having a repeating unit represented by the following formula (Ia):



In formula (Ia), Rf represents a fluorine atom or an alkyl group with at least one hydrogen atom being substituted by a fluorine atom.

R₁ represents an alkyl group.

R₂ represents a hydrogen atom or an alkyl group.

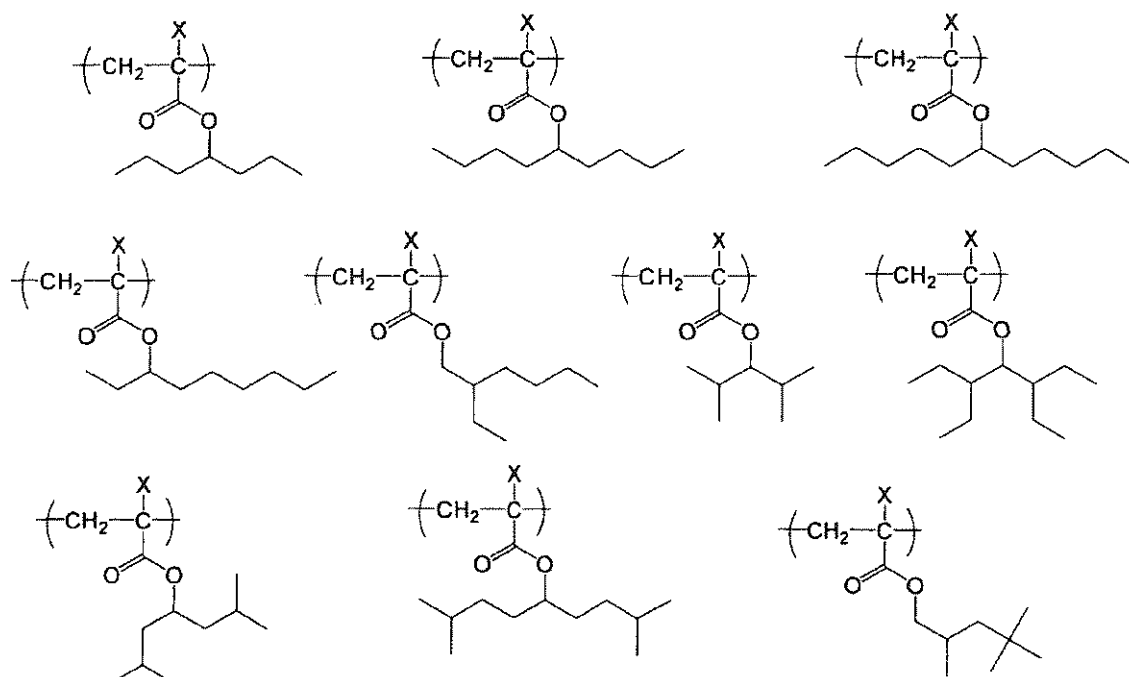
In formula (Ia), the alkyl group with at least one hydrogen atom being substituted by a fluorine atom of Rf is preferably an alkyl group having a carbon number of 1 to 3, more preferably a trifluoromethyl group.

The alkyl group of R₁ is preferably a linear or branched alkyl group having a carbon number of 3 to 10, more preferably a branched alkyl group having a carbon number of 3 to 10.

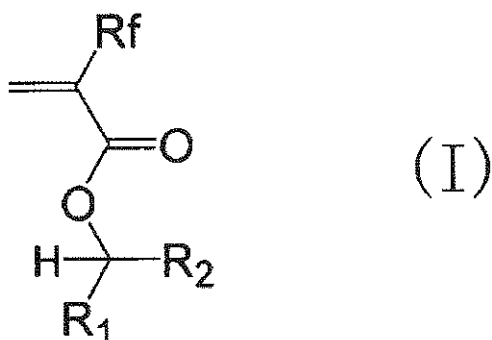
R₂ is preferably a linear or branched alkyl group having a carbon number of 1 to 10, more preferably a linear or branched alkyl group having a carbon number of 3 to 10.

Specific examples of the repeating unit represented by formula (Ia) are set forth below, but the present invention is not limited thereto.

X=F or CF₃



The repeating unit represented by formula (Ia) can be formed by polymerizing a compound represented by the following formula (I):



In formula (I), Rf represents a fluorine atom or an alkyl group with at least one hydrogen atom being substituted by a fluorine atom.

R₁ represents an alkyl group.

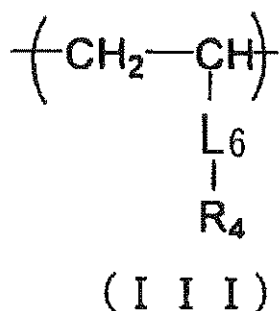
R₂ represents a hydrogen atom or an alkyl group.

Rf, R₁ and R₂ in formula (I) have the same meanings as Rf, R₁ and R₂ in formula (Ia).

The compound represented by formula (I) is a novel compound.

As for the compound represented by formula (I), a commercially available product or a compound synthesized may be used. In the case of synthesizing the compound, this can be attained by converting a 2-trifluoromethyl methacrylic acid into an acid chloride and then esterifying the acid chloride.

The resin (D) containing a repeating unit represented by formula (Ia) preferably further contains a repeating unit represented by the following formula (III):



In formula (III), R₄ represents an alkyl group, a cycloalkyl group, an alkenyl group, a cycloalkenyl group, a trialkylsilyl group or a group having a cyclic siloxane structure.

L₆ represents a single bond or a divalent linking group.

In formula (III), the alkyl group of R₄ is preferably a linear or branched alkyl group having a carbon number of 3 to 20.

The cycloalkyl group is preferably a cycloalkyl group having a carbon number of 3 to 20.

The alkenyl group is preferably an alkenyl group having a carbon number of 3 to 20.

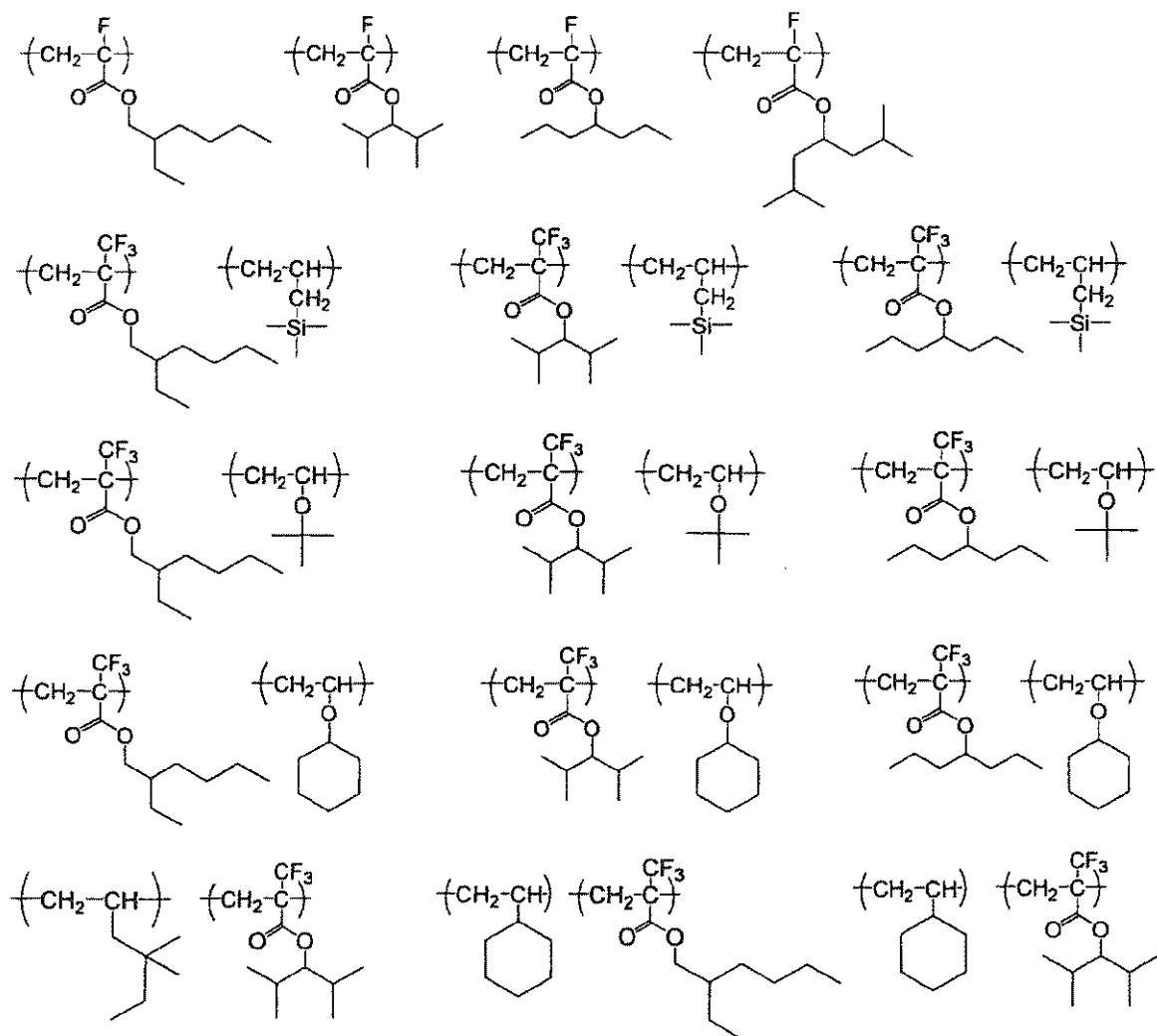
The cycloalkenyl group is preferably a cycloalkenyl group having a carbon number of 3 to 20.

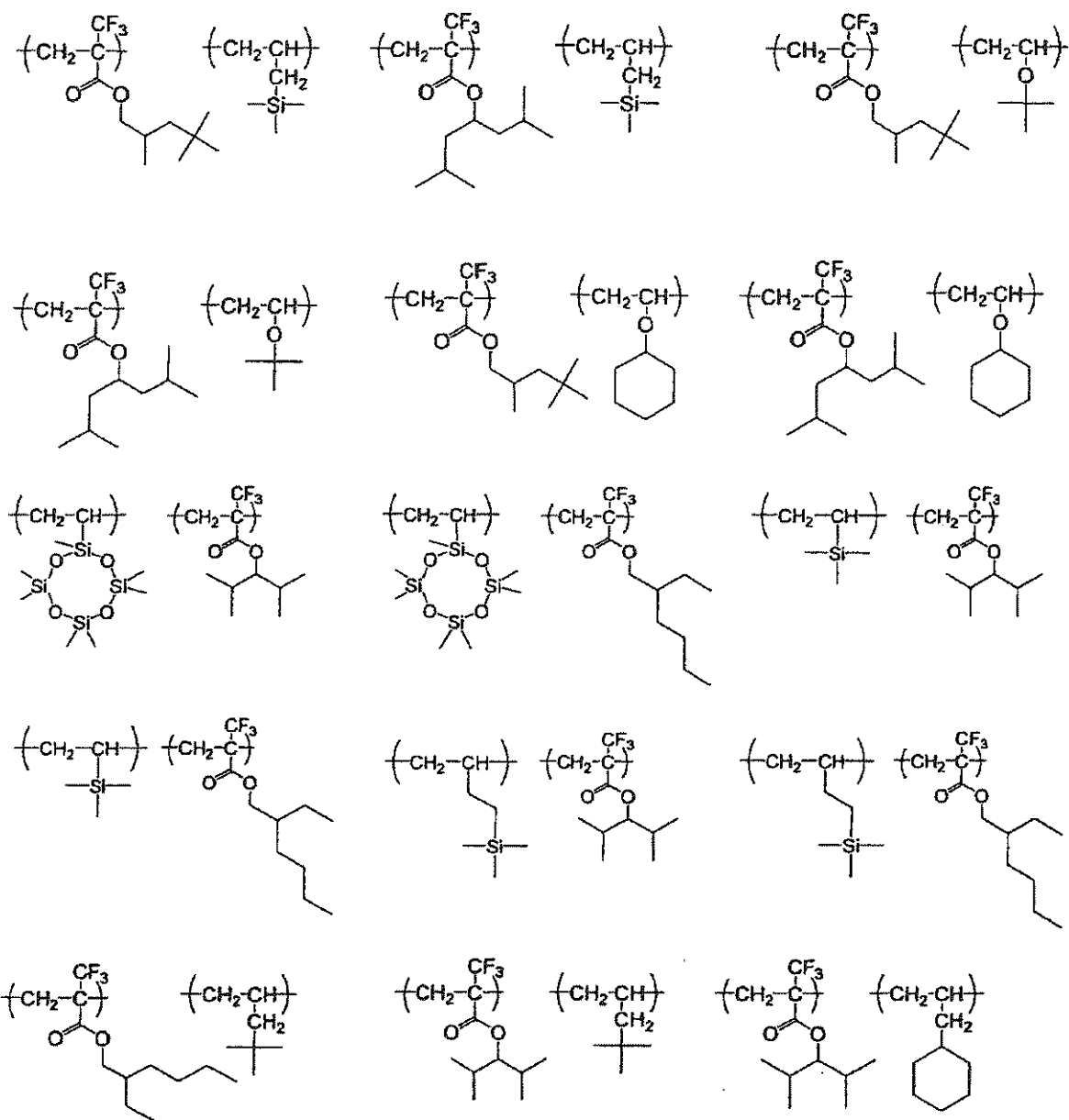
The trialkylsilyl group is preferably a trialkylsilyl group having a carbon number of 3 to 20.

The group having a cyclic siloxane structure is preferably a group containing a cyclic siloxane structure having a carbon number of 3 to 20.

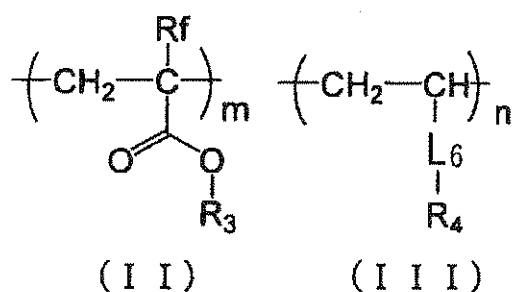
The divalent linking group of L₆ is preferably an alkylene group (preferably having a carbon number of 1 to 5) or an oxy group.

Specific examples of the resin (D) having a repeating unit represented by formula (Ia) are set forth below, but the present invention is not limited thereto.





The resin (D) is preferably a resin containing a repeating unit represented by the following formula (II) and a repeating unit represented by the following formula (III):



In formulae (II) and (III), Rf represents a fluorine atom or an alkyl group with at least one hydrogen atom being substituted by a fluorine atom.

R₃ represents an alkyl group, a cycloalkyl group, an alkenyl group, a cycloalkenyl

group, or a group formed by combining two or more members thereof.

R_4 represents an alkyl group, a cycloalkyl group, an alkenyl group, a cycloalkenyl group, a trialkylsilyl group, a group having a cyclic siloxane structure, or a group formed by combining two or more members thereof.

In the alkyl group, cycloalkyl group, alkenyl group, cycloalkenyl group and trialkylsilyl group of R_3 and R_4 , a functional group can be introduced but in view of followability of immersion liquid, a functional group having no polar group is preferred, and an unsubstituted functional group is more preferred.

L_6 represents a single bond or a divalent linking group.

$0 < m < 100$.

$0 < n < 100$.

In formula (II), R_f has the same meaning as R_f in formula (Ia).

The alkyl group of R_3 is preferably a linear or branched alkyl group having a carbon number of 3 to 20.

The cycloalkyl group is preferably a cycloalkyl group having a carbon number of 3 to 20.

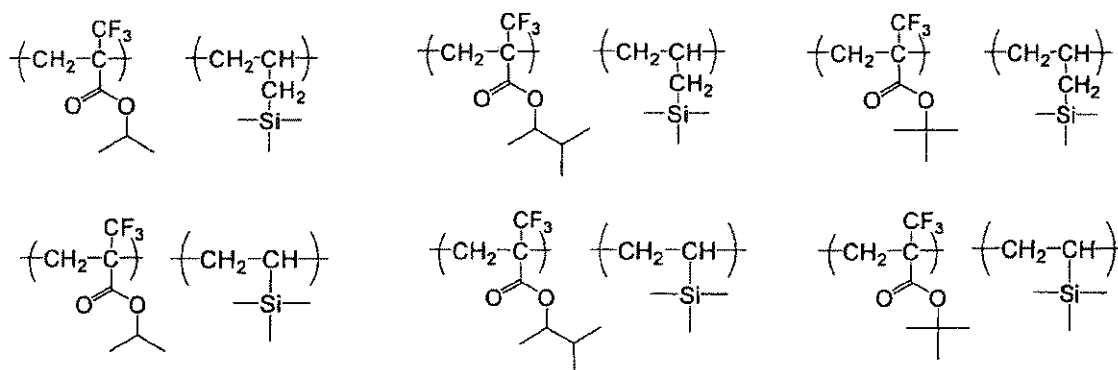
The alkenyl group is preferably an alkenyl group having a carbon number of 3 to 20.

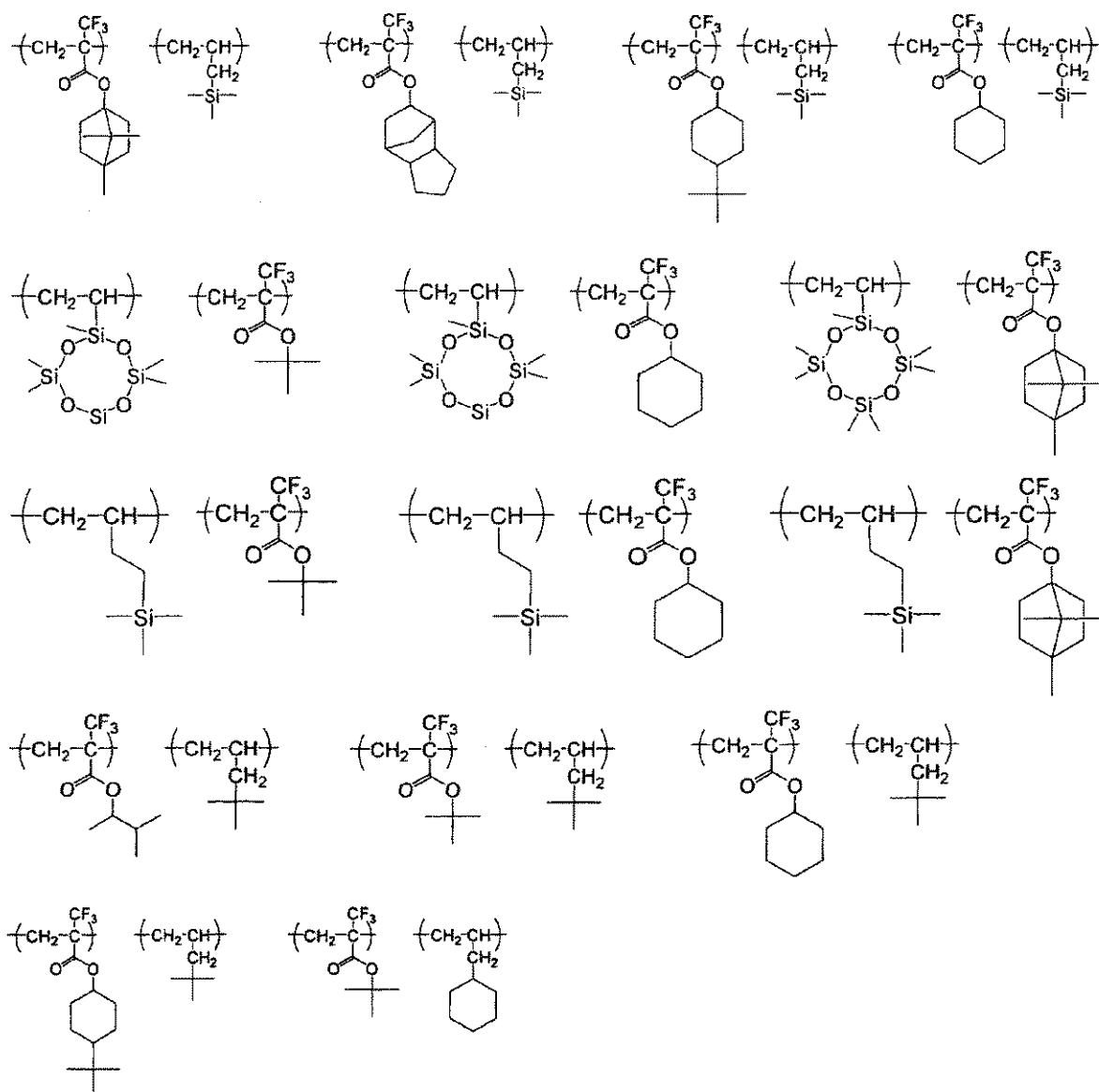
The cycloalkenyl group is preferably a cycloalkenyl group having a carbon number of 3 to 20.

L_6 is preferably a single bond, a methylene group, an ethylene group or an ether group.

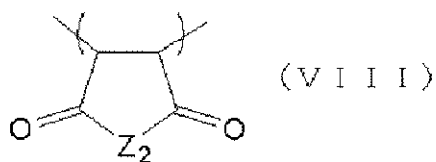
m and n are preferably from 30 to 70 and from 30 to 70, respectively, more preferably from 40 to 60 and from 40 to 60, respectively.

Specific examples of the resin (D) containing a repeating unit represented by formula (II) and a repeating unit represented by formula (III) are set forth below, but the present invention is not limited thereto.





The resin (D) may contain a repeating unit represented by the following formula (VIII):



In formula (VIII), Z_2 represents $-\text{O}-$ or $-\text{N}(\text{R}_{41})-$. R_{41} represents a hydrogen atom, an alkyl group or $-\text{OSO}_2-\text{R}_{42}$. R_{42} represents an alkyl group, a cycloalkyl group or a camphor residue. The alkyl group of R_{41} and R_{42} may be substituted by a halogen atom (preferably fluorine atom) or the like.

The resin (D) is preferably solid at ordinary temperature (25°C). Furthermore, the

glass transition temperature (T_g) is preferably from 50 to 200°C, more preferably from 80 to 160°C.

When the resin is solid at 25°C, this means that the melting point is 25°C or more.

The glass transition temperature (T_g) can be measured by a scanning calorimeter (Differential Scanning Calorimeter). For example, the glass transition temperature can be measured by once heating and then cooling the sample, again raising the temperature at 5°C/min, and analyzing the value when the specific volume is changed.

The resin (D) is preferably stable to an acid and insoluble in an alkali developer.

In view of followability of immersion liquid, the resin (D) preferably contains none of (x) an alkali-soluble group, (y) a group capable of decomposing by the action of an alkali (alkali developer) to increase the solubility in an alkali developer, and (z) a group capable of decomposing by the action of an acid to increase the solubility in a developer.

In the resin (D), the total amount of repeating units having an alkali-soluble group or a group whose solubility in a developer increases by the action of an acid or alkali is preferably 20 mol% or less, more preferably from 0 to 10 mol%, still more preferably from 0 to 5 mol%, based on all repeating units constituting the resin (D).

Also, unlike a surfactant generally used for resists, the resin (D) does not have an ionic bond or a hydrophilic group such as (poly(oxyalkylene)) group. If the resin (D) contains a hydrophilic polar group, the followability of immersion water tends to decrease. Therefore, it is more preferred not to contain a polar group selected from a hydroxyl group, alkylene glycols and a sulfone group. Furthermore, an ether group bonded to the carbon atom of the main chain through a linking group is preferably not contained because the hydrophilicity increases and the followability of immersion liquid deteriorates. On the other hand, an ether group bonded directly to the carbon atom of the main chain as in formula (III) can sometimes express activity as a hydrophobic group and is preferred.

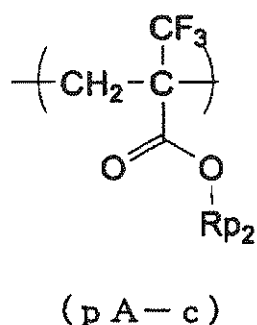
Examples of the alkali-soluble group (x) include groups having a phenolic hydroxyl group, a carboxylic acid group, a fluorinated alcohol group, a sulfonic acid group, a sulfonamide group, a sulfonylimide group, an (alkylsulfonyl)(alkylcarbonyl)methylene group, an (alkylsulfonyl)(alkylcarbonyl)imide group, a bis(alkylcarbonyl)methylene group, a bis(alkylcarbonyl)imide group, a bis(alkylsulfonyl)methylene group, a bis(alkylsulfonyl)imide group, a tris(alkylcarbonyl)methylene group or a tris(alkylsulfonyl)methylene group.

Examples of the group (y) capable of decomposing by the action of an alkali (alkali developer) to increase the solubility in an alkali developer include a lactone group, an ester group, a sulfonamide group, an acid anhydride and an acid imide group.

Examples of the group (z) capable of decomposing by the action of an acid to

increase the solubility in a developer include the same groups as those of the acid-decomposable group in the acid-decomposable resin (A).

However, the repeating unit represented by the following formula (pA-c) is not or scarcely decomposed by the action of an acid as compared with the acid-decomposable group of the resin (A) and is regarded as substantially non-acid-decomposable.



In formula (pA-c), Rp₂ represents a hydrocarbon group having a tertiary carbon atom bonded to the oxygen atom in the formula.

In the case where the resin (D) contains a silicon atom, the silicon atom content is preferably from 2 to 50 mass%, more preferably from 2 to 30 mass%, based on the molecular weight of the resin (D). Also, the silicon atom-containing repeating unit preferably occupies from 10 to 100 mass%, more preferably from 20 to 100 mass%, in the resin (D).

In the case where the resin (D) contains a fluorine atom, the fluorine atom content is preferably from 5 to 80 mass%, more preferably from 10 to 80 mass%, based on the molecular weight of the resin (D). Also, the fluorine atom-containing repeating unit preferably occupies from 10 to 100 mass%, more preferably from 30 to 100 mass%, in the resin (D).

The standard polystyrene-reduced weight average molecular of the resin (D) is preferably from 1,000 to 100,000, more preferably from 1,000 to 50,000, still more preferably from 2,000 to 15,000, yet still more preferably from 3,000 to 15,000.

The residual monomer amount in the resin (D) is preferably from 0 to 10 mass%, more preferably from 0 to 5 mass%, still more preferably from 0 to 1 mass%. Also, in view of the resolution, resist profile, and side wall, roughness or the like of the resist pattern, the molecular weight distribution (Mw/Mn, also called dispersity) is preferably from 1 to 5, more preferably from 1 to 3, still more preferably from 1 to 1.5.

The amount added of the resin (D) in the resist composition is preferably from 0.1 to 20 mass%, more preferably from 0.1 to 10 mass%, still more preferably from 0.1 to 5 mass%, even still more preferably from 0.2 to 3.0 mass%, yet even still more preferably from 0.3 to 2.0 mass%, based on the entire solid content of the resist composition.

In the resin (D), similarly to the acid-decomposable resin (A), impurities such as metal are of course little contained and as well, the content of residual monomers or oligomer components is preferably not more than a specific value, for example, 0.1 mass% by HPLC. By satisfying these conditions, not only the resist can be improved in the sensitivity, resolution, process stability, pattern profile and the like but also a resist free from extraneous substances in liquid or change with aging in the sensitivity and the like can be obtained.

As for the resin (D), various commercially available products may be used or the resin may be synthesized by an ordinary method (for example, radical polymerization)). Examples of the synthesis method in general include a batch polymerization method of dissolving monomer species and an initiator in a solvent and heating the solution, thereby effecting the polymerization, and a dropping polymerization method of adding dropwise a solution containing monomer species and an initiator to a heated solvent over 1 to 10 hours. A dropping polymerization method is preferred. Examples of the reaction solvent include tetrahydrofuran, 1,4-dioxane, ethers such as diisopropyl ether, ketones such as methyl ethyl ketone and methyl isobutyl ketone, an ester solvent such as ethyl acetate, an amide solvent such as dimethylformamide and dimethylacetamide, and the later-described solvent capable of dissolving the composition of the present invention, such as propylene glycol monomethyl ether acetate, propylene glycol monomethyl ether and cyclohexanone. The polymerization is more preferably performed using the same solvent as the solvent used in the resist composition of the present invention. By the use of this solvent, generation of particles during storage can be suppressed.

The polymerization reaction is preferably performed in an inert gas atmosphere such as nitrogen and argon. As for the polymerization initiator, the polymerization is started using a commercially available radical initiator (e.g., azo-based initiator, peroxide). The radical initiator is preferably an azo-based initiator, and an azo-based initiator having an ester group, a cyano group or a carboxyl group is preferred. Preferred examples of the initiator include azobisisobutyronitrile, azobisdimethylvaleronitrile and dimethyl 2,2'-azobis(2-methylpropionate). A chain transfer agent may also be used, if desired. The reaction concentration is usually from 5 to 50 mass%, preferably from 20 to 50 mass%, more preferably from 30 to 50 mass%, and the reaction temperature is usually from 10 to 150°C, preferably from 30 to 120°C, more preferably from 60 to 100°C.

After the completion of reaction, the reaction product is allowed to cool to room temperature and purified. The purification may be performed by a normal method, for example, a liquid-liquid extraction method of combining water washing with an appropriate solvent to remove residual monomers or oligomer components; a purification method in a solution state, such as ultrafiltration of extracting and removing only polymers having a

molecular weight lower than a specific molecular weight; a reprecipitation method of adding dropwise the resin solution in a poor solvent to solidify the resin in the poor solvent and thereby remove residual monomers and the like; or a purification method in a solid state, such as washing of the resin slurry with a poor solvent after separation by filtration. For example, the resin is precipitated as a solid through contact with a solvent in which the resin is sparingly soluble or insoluble (poor solvent) and which is in a volume amount of 10 times or less, preferably from 10 to 5 times, the reaction solution.

The solvent used at the operation of precipitation or reprecipitation from the polymer solution (precipitation or reprecipitation solvent) may be sufficient if it is a poor solvent for the polymer, and the solvent used may be appropriately selected according to the kind of the polymer from, for example, a hydrocarbon (an aliphatic hydrocarbon such as pentane, hexane, heptane and octane; an alicyclic hydrocarbon such as cyclohexane and methylcyclohexane; and an aromatic hydrocarbon such as benzene, toluene and xylene), a halogenated hydrocarbon (a halogenated aliphatic hydrocarbon such as methylene chloride, chloroform and carbon tetrachloride; and a halogenated aromatic hydrocarbon such as chlorobenzene and dichlorobenzene), a nitro compound (e.g., nitromethane, nitroethane), a nitrile (e.g., acetonitrile, benzonitrile), an ether (a chain ether such as diethyl ether, diisopropyl ether and dimethoxyethane; and a cyclic ether such as tetrahydrofuran and dioxane), a ketone (e.g., acetone, methyl ethyl ketone, diisobutyl ketone), an ester (e.g., ethyl acetate, butyl acetate), a carbonate (e.g., dimethyl carbonate, diethyl carbonate, ethylene carbonate, propylene carbonate), an alcohol (e.g., methanol, ethanol, propanol, isopropyl alcohol, butanol), a carboxylic acid (e.g., acetic acid), water, and a mixed solvent containing such a solvent.

Among these, the precipitation or reprecipitation solvent is preferably a solvent containing at least an alcohol (particularly methanol or the like) or water. In such a solvent containing at least a hydrocarbon, the ratio of the alcohol (particularly methanol or the like) to other solvents (for example, an ester such as ethyl acetate, and ethers such as tetrahydrofuran) is approximately, for example, the former/the latter (volume ratio, at 25°C) = from 10/90 to 99/1, preferably the former/the latter (volume ratio, at 25°C) = from 30/70 to 98/2, more preferably the former/the latter (volume ratio, at 25°C) = from 50/50 to 97/3.

The amount of the precipitation or reprecipitation solvent used may be appropriately selected by taking into account the efficiency, yield and the like, but in general, the amount used is from 100 to 10,000 parts by mass, preferably from 200 to 2,000 parts by mass, more preferably from 300 to 1,000 parts by mass, per 100 parts by mass of the polymer solution.

The nozzle bore diameter at the time of feeding the polymer solution into a precipitation or reprecipitation solvent (poor solvent) is preferably 4 mmφ or less (for example, from 0.2 to 4 mmφ), and the feeding rate (dropping rate) of the polymer solution

into the poor solvent is, for example, in terms of the linear velocity, from 0.1 to 10 m/sec, preferably on the order of 0.3 to 5 m/sec.

The precipitation or reprecipitation operation is preferably performed under stirring. Examples of the stirring blade which can be used for stirring include a disc turbine, a fan turbine (including paddle), a curved vane turbine, a feathering turbine, a Pfaudler type, a bull margin type, an angled vane fan turbine, a propeller, a multistage type, an anchor type (or horseshoe type), a gate type, a double ribbon and a screw. The stirring is preferably further performed for 10 minutes or more, more preferably 20 minutes or more, after the completion of feeding of the polymer solution. If the stirring time is short, the monomer content in the polymer particle may not be sufficiently reduced. The mixing and stirring of the polymer solution and the poor solvent may also be performed using a line mixer instead of the stirring blade.

The temperature at the precipitation or reprecipitation may be appropriately selected by taking into account the efficiency or operability, but the temperature is usually on the order of 0 to 50°C, preferably in the vicinity of room temperature (for example, approximately from 20 to 35°C). The precipitation or reprecipitation operation may be performed using a commonly employed mixing vessel such as stirring tank according to a known method such as batch system and continuous system.

The precipitated or reprecipitated particulate polymer is usually subjected to commonly employed solid-liquid separation such as filtration and centrifugation, then dried and used. The filtration is performed using a solvent-resistant filter element preferably under pressure. The drying is performed under atmospheric pressure or reduced pressure (preferably under reduced pressure) at a temperature of approximately from 30 to 100°C, preferably on the order of 30 to 50°C.

Incidentally, after the resin is once precipitated and separated, the resin may be again dissolved in a solvent and then put into contact with a solvent in which the resin is sparingly soluble or insoluble.

More specifically, there may be used a method comprising, after the completion of radical polymerization reaction, precipitating a resin by bringing the polymer into contact with a solvent in which the polymer is sparingly soluble or insoluble (step a), separating the resin from the solution (step b), anew dissolving the resin in a solvent to prepare a resin solution A (step c), precipitating a resin solid by bringing the resin solution A into contact with a solvent in which the resin is sparingly soluble or insoluble and which is in a volume amount of less than 10 times (preferably a volume amount of 5 times or less) the resin solution A (step d), and separating the precipitated resin (step e).

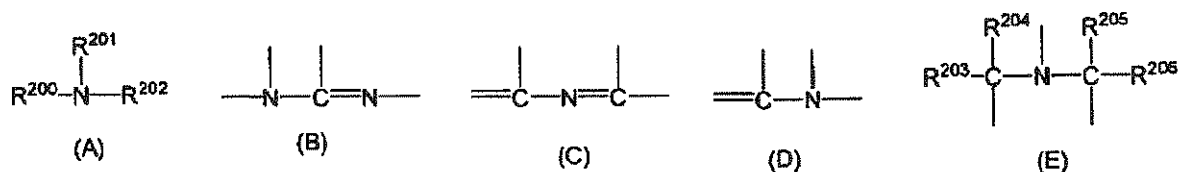
As for the solvent used at the preparation of the resin solution A, a solvent similar to

the solvent used for dissolving the monomer at the polymerization reaction may be used, and the solvent may be the same as or different from the solvent used at the polymerization reaction.

(E) Basic compound

The resist composition of the present invention preferably contains (E) a basic compound for reducing the change of performance with aging from exposure to heating.

Preferred examples of the basic compound include compounds having a structure represented by any one of the following formulae (A) to (E):



In formulae (A) to (E), each of R^{200} , R^{201} and R^{202} which may be the same or different represents a hydrogen atom, an alkyl group (preferably having a carbon number of 1 to 20), a cycloalkyl group (preferably having a carbon number of 3 to 20) or an aryl group (having a carbon number of 6 to 20), and R^{201} and R^{202} may combine with each other to form a ring.

As for the alkyl group, the alkyl group having a substituent is preferably an aminoalkyl group having a carbon number of 1 to 20, a hydroxyalkyl group having a carbon number of 1 to 20, or a cyanoalkyl group having a carbon number of 1 to 20.

Each of R^{203} , R^{204} , R^{205} and R^{206} which may be the same or different represents an alkyl group having a carbon number of 1 to 20.

The alkyl group in these formulae (A) to (E) is more preferably unsubstituted.

Preferred examples of the compound include guanidine, aminopyrrolidine, pyrazole, pyrazoline, piperazine, aminomorpholine, aminoalkylmorpholine and piperidine. More preferred examples of the compound include a compound having an imidazole structure, a diazabicyclo structure, an onium hydroxide structure, an onium carboxylate structure, a trialkylamine structure, an aniline structure or a pyridine structure; an alkylamine derivative having a hydroxyl group and/or an ether bond; and an aniline derivative having a hydroxyl group and/or an ether bond.

Examples of the compound having an imidazole structure include imidazole, 2,4,5-triphenylimidazole and benzimidazole. Examples of the compound having a diazabicyclo structure include 1,4-diazabicyclo[2,2,2]octane, 1,5-diazabicyclo[4,3,0]non-5-ene and 1,8-diazabicyclo[5,4,0]undec-7-ene. Examples of the compound having an onium hydroxide structure include triarylsulfonium hydroxide, phenacylsulfonium hydroxide and sulfonium hydroxide having a 2-oxoalkyl group, specifically, triphenylsulfonium hydroxide, tris(tert-

butylphenyl)sulfonium hydroxide, bis(tert-butylphenyl)iodonium hydroxide, phenacylthiophenium hydroxide and 2-oxopropylthiophenium hydroxide. Examples of the compound having an onium carboxylate structure include a compound where the anion moiety of the compound having an onium hydroxide structure is converted into a carboxylate, such as acetate, adamantane-1-carboxylate and perfluoroalkyl carboxylate. Examples of the compound having a trialkylamine structure include tri(n-butyl)amine and tri(n-octyl)amine. Examples of the aniline compound include 2,6-diisopropylaniline, N,N-dimethylaniline, N,N-dibutylaniline and N,N-dihexylaniline. Examples of the alkylamine derivative having a hydroxyl group and/or an ether bond include ethanolamine, diethanolamine, triethanolamine and tris(methoxyethoxyethyl)amine. Examples of the aniline derivative having a hydroxyl group and/or an ether bond include N,N-bis(hydroxyethyl)aniline.

One of these basic compounds is used alone, or two or more kinds thereof are used in combination.

The amount of the basic compound used is usually from 0.001 to 10 mass%, preferably from 0.01 to 5 mass%, based on the solid content of the resist composition.

The ratio between the acid generator and the basic compound used in the composition is preferably acid generator/basic compound (by mol) = from 2.5 to 300. That is, the molar ratio is preferably 2.5 or more in view of sensitivity and resolution and preferably 300 or less from the standpoint of suppressing the reduction in resolution due to thickening of the resist pattern with aging after exposure until heat treatment. The acid generator/basic compound (by mol) is more preferably from 5.0 to 200, still more preferably from 7.0 to 150.

(F) Surfactant

The resist composition of the present invention preferably further contains (F) a surfactant, more preferably any one fluorine-containing and/or silicon-containing surfactant (a fluorine-containing surfactant, a silicon-containing surfactant or a surfactant containing both a fluorine atom and a silicon atom) or two or more kinds thereof.

When the resist composition of the present invention contains the surfactant (F), a resist pattern with good sensitivity, resolution and adherence as well as less development defects can be obtained in using an exposure light source of 250 nm or less, particularly 220 nm or less.

Examples of the fluorine-containing and/or silicon-containing surfactant include surfactants described in JP-A-62-36663, JP-A-61-226746, JP-A-61-226745, JP-A-62-170950, JP-A-63-34540, JP-A-7-230165, JP-A-8-62834, JP-A-9-54432, JP-A-9-5988, JP-A-2002-277862 and U.S. Patents 5,405,720, 5,360,692, 5,529,881, 5,296,330, 5,436,098, 5,576,143, 5,294,511 and 5,824,451. The following commercially available surfactants each may also

be used as it is.

Examples of the commercially available surfactant which can be used include a fluorine-containing or silicon-containing surfactant such as Eftop EF301 and EF303 (produced by Shin-Akita Kasei K.K.); Florad FC430, 431 and 4430 (produced by Sumitomo 3M Inc.); Megaface F171, F173, F176, F189, F113, F110, F177, F120 and R08 (produced by Dainippon Ink & Chemicals, Inc.); Surflon S-382, SC101, 102, 103, 104, 105 and 106 (produced by Asahi Glass Co., Ltd.); Troysol S-366 (produced by Troy Chemical); GF-300 and GF-150 (produced by Toagosei Chemical Industry Co., Ltd.); Surflon S-393 (produced by Seimi Chemical Co., Ltd.); Eftop EF121, EF122A, EF122B, RF122C, EF125M, EF135M, EF351, 352, EF801, EF802 and EF601 (produced by JEMCO Inc.); PF636, PF656, PF6320 and PF6520 (produced by OMNOVA); and FTX-204D, 208G, 218G, 230G, 204D, 208D, 212D, 218 and 222D (produced by NEOS Co., Ltd.). In addition, polysiloxane polymer KP-341 (produced by Shin-Etsu Chemical Co., Ltd.) may also be used as the silicon-containing surfactant.

Other than those known surfactants, a surfactant using a polymer having a fluoro-aliphatic group derived from a fluoro-aliphatic compound which is produced by a telomerization process (also called a telomer process) or an oligomerization process (also called an oligomer process), may be used. The fluoro-aliphatic compound can be synthesized by the method described in JP-A-2002-90991.

The polymer having a fluoro-aliphatic group is preferably a copolymer of a fluoro-aliphatic group-containing monomer with a (poly(oxyalkylene)) acrylate and/or a (poly(oxyalkylene)) methacrylate, and the copolymer may have an irregular distribution or may be a block copolymer. Examples of the poly(oxyalkylene) group include a poly(oxyethylene) group, a poly(oxypropylene) group and a poly(oxybutylene) group. This group may also be a unit having alkylenes differing in the chain length within the same chain, such as block-linked poly(oxyethylene, oxypropylene and oxyethylene) and block-linked poly(oxyethylene and oxypropylene). Furthermore, the copolymer of a fluoro-aliphatic group-containing monomer and a (poly(oxyalkylene)) acrylate (or methacrylate) is not limited only to a binary copolymer but may also be a ternary or greater copolymer obtained by simultaneously copolymerizing two or more different fluoro-aliphatic group-containing monomers or two or more different (poly(oxyalkylene)) acrylates (or methacrylates).

Examples thereof include, as the commercially available surfactant, Megaface F178, F-470, F-473, F-475, F-476 and F-472 (produced by Dainippon Ink & Chemicals, Inc.) and further include a copolymer of a C_6F_{13} group-containing acrylate (or methacrylate) with a (poly(oxyalkylene)) acrylate (or methacrylate), and a copolymer of a C_3F_7 group-containing acrylate (or methacrylate) with a (poly(oxyethylene)) acrylate (or methacrylate) and a

(poly(oxypropylene)) acrylate (or methacrylate).

In the present invention, a surfactant other than the fluorine-containing and/or silicon-containing surfactant may also be used. Specific examples thereof include a nonionic surfactant such as polyoxyethylene alkyl ethers (e.g., polyoxyethylene lauryl ether, polyoxyethylene stearyl ether, polyoxyethylene cetyl ether, polyoxyethylene oleyl ether), polyoxyethylene alkylallyl ethers (e.g., polyoxyethylene octylphenol ether, polyoxyethylene nonylphenol ether), polyoxyethylene-polyoxypropylene block copolymers, sorbitan fatty acid esters (e.g., sorbitan monolaurate, sorbitan monopalmitate, sorbitan monostearate, sorbitan monooleate, sorbitan trioleate, sorbitan tristearate) and polyoxyethylene sorbitan fatty acid esters (e.g., polyoxyethylene sorbitan monolaurate, polyoxyethylene sorbitan monopalmitate, polyoxyethylene sorbitan monostearate, polyoxyethylene sorbitan trioleate, polyoxyethylene sorbitan tristearate).

One of these surfactants may be used alone, or several surfactants may be used in combination.

The amount of the surfactant (F) used is preferably from 0.01 to 10 mass%, more preferably from 0.1 to 5 mass%, based on the entire amount of the resist composition (excluding the solvent).

(G) Onium carboxylate

The resist composition of the present invention may contain (G) an onium carboxylate. Examples of the onium carboxylate include sulfonium carboxylate, iodonium carboxylate and ammonium carboxylate. In particular, the onium carboxylate (G) is preferably an iodonium salt or a sulfonium salt. Furthermore, the carboxylate residue of the onium carboxylate (H) for use in the present invention preferably contains no aromatic group and no carbon-carbon double bond. The anion moiety is preferably a linear, branched, monocyclic or polycyclic alkylcarboxylate anion having a carbon number of 1 to 30, more preferably an anion of the carboxylic acid with the alkyl group being partially or entirely fluorine-substituted. The alkyl chain may contain an oxygen atom. Thanks to such a construction, the transparency to light of 220 nm or less is ensured, the sensitivity and resolution are enhanced, and the iso/dense bias and exposure margin are improved.

Examples of the anion of the fluorine-substituted carboxylic acid include anions of fluoroacetic acid, difluoroacetic acid, trifluoroacetic acid, pentafluoropropionic acid, heptafluorobutyric acid, nonafluoropentanoic acid, perfluorododecanoic acid, perfluorotridecanoic acid, perfluorocyclohexanecarboxylic acid and 2,2-bistrifluoromethylpropionic acid.

These onium carboxylates (G) can be synthesized by reacting a sulfonium, iodonium or ammonium hydroxide and a carboxylic acid with silver oxide in an appropriate solvent.

The content of the onium carboxylate (G) in the composition is generally from 0.1 to 20 mass%, preferably from 0.5 to 10 mass%, more preferably from 1 to 7 mass%, based on the entire solid content of the composition.

(H) Other additives

The resist composition of the present invention may further contain, for example, a dye, a plasticizer, a photosensitizer, a light absorber, an alkali-soluble resin, a dissolution inhibitor and a compound for accelerating dissolution in a developer (for example, a phenol compound having a molecular weight of 1,000 or less, or a carboxyl group-containing alicyclic or aliphatic compound), if desired.

The phenol compound having a molecular weight of 1,000 or less can be easily synthesized by one skilled in the art with reference to the methods described, for example, in JP-A-4-122938, JP-A-2-28531, U.S. Patent 4,916,210 and European Patent 219294.

Specific examples of the carboxyl group-containing alicyclic or aliphatic compound include, but are not limited to, a carboxylic acid derivative having a steroid structure, such as cholic acid, deoxycholic acid and lithocholic acid, an adamantanecarboxylic acid derivative, an adamantanedicarboxylic acid, a cyclohexanecarboxylic acid and a cyclohexanedicarboxylic acid.

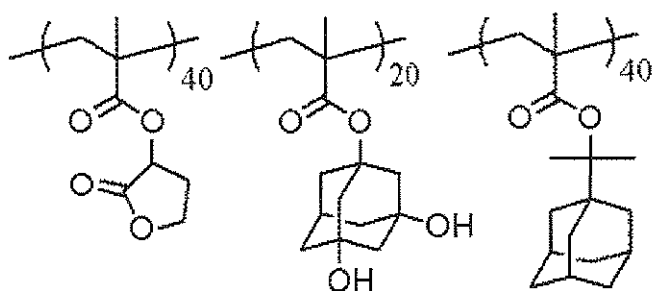
EXAMPLES

The present invention is described below by referring to Examples, but the present invention should not be construed as being limited thereto.

Synthesis Example 1 (Synthesis of Resin (A1)):

Under a nitrogen stream, 8.4 g of methyl isobutyl ketone was charged into a three-neck flask and heated at 80°C. Thereto, a solution obtained by dissolving 9.4 g of 2-(1-adamantyl)propan-2-yl methacrylate, 4.5 g of 3,5-dihydroxy-1-adamantyl methacrylate, 6.1 g of β -methacryloyloxy- γ -butyrolactone and azobisisobutyronitrile corresponding to 6 mol% based on the entire monomer amount, in 75.3 g of methyl isobutyl ketone was added dropwise over 6 hours. After the completion of dropwise addition, the reaction was further allowed to proceed at 80°C for 2 hours. The resulting reaction solution was allowed to cool and then poured in 720 ml of heptane/80 ml of ethyl acetate, and the powder precipitated was collected by filtration and dried, as a result, 18.3 g of Resin (A1) was obtained. The weight average molecular weight of the obtained resin was 7,000 and the dispersity (Mw/Mn) was 1.80.

Resin (A1):



Weight average molecular weight: 7,000

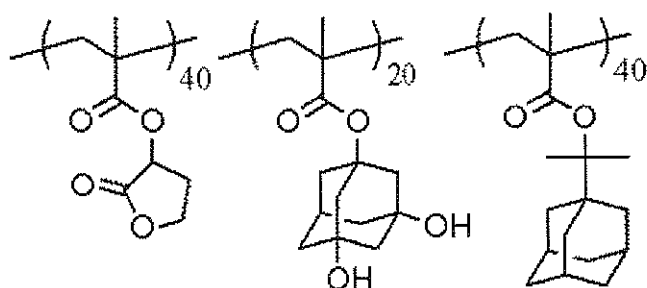
Dispersity: 1.80

Molar compositional ratio: 40/20/40

Synthesis Example 2 (Synthesis of Resin (A2)):

Under a nitrogen stream, 8.4 g of methyl isobutyl ketone was charged into a three-neck flask and heated at 80°C. Thereto, a solution obtained by dissolving 9.4 g of 2-(1-adamantyl)propan-2-yl methacrylate, 4.5 g of 3,5-dihydroxy-1-adamantyl methacrylate, 6.1 g of β -methacryloyloxy- γ -butyrolactone and azobisisobutyronitrile corresponding to 7 mol% based on the entire monomer amount, in 75.3 g of methyl isobutyl ketone was added dropwise over 6 hours. After the completion of dropwise addition, the reaction was further allowed to proceed at 80°C for 2 hours. The resulting reaction solution was allowed to cool and then poured in 680 ml of heptane/120 ml of ethyl acetate, and the powder precipitated was collected by filtration and dried, as a result, 17.5 g of Resin (A2) was obtained. The weight average molecular weight of the obtained resin was 5,800 and the dispersity (M_w/M_n) was 1.53.

Resin (A2):



Weight average molecular weight: 5,800

Dispersity: 1.53

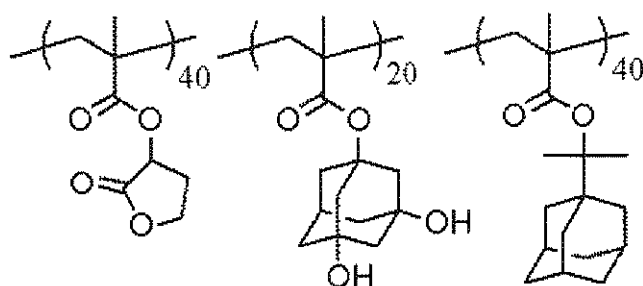
Molar compositional ratio: 40/20/40

Synthesis Example 3 (Synthesis of Resin (A3)):

Under a nitrogen stream, 19.4 g of methyl isobutyl ketone was charged into a three-neck flask and heated at 80°C. Thereto, a solution obtained by dissolving 9.4 g of 2-(1-

adamantyl)propan-2-yl methacrylate, 4.5 g of 3,5-dihydroxy-1-adamantyl methacrylate, 6.1 g of β -methacryloyloxy- γ -butyrolactone and azobisisobutyronitrile corresponding to 12 mol% based on the entire monomer amount, in 65.3 g of methyl isobutyl ketone was added dropwise over 8 hours. After the completion of dropwise addition, the reaction was further allowed to proceed at 80°C for 2 hours. The resulting reaction solution was allowed to cool and then poured in 720 ml of heptane/80 ml of ethyl acetate, and the powder precipitated was collected by filtration and dried, as a result, 15.5 g of Resin (A3) was obtained. The weight average molecular weight of the obtained resin was 2,800 and the dispersity (Mw/Mn) was 1.26.

Resin (A3):



Weight average molecular weight: 2,800

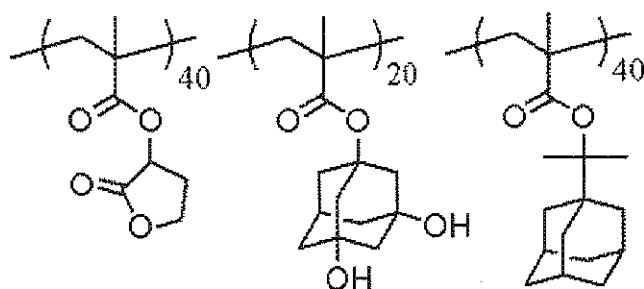
Dispersity: 1.26

Molar compositional ratio: 40/20/40

Synthesis Example 4 (Synthesis of Resin (A4)):

Under a nitrogen stream, 8.4 g of methyl isobutyl ketone was charged into a three-neck flask and heated at 80°C. Thereafter, a solution obtained by dissolving 9.4 g of 2-(1-adamantyl)propan-2-yl methacrylate, 4.5 g of 3,5-dihydroxy-1-adamantyl methacrylate, 6.1 g of β -methacryloyloxy- γ -butyrolactone and azobisisobutyronitrile corresponding to 6 mol% based on the entire monomer amount, in 75.3 g of methyl isobutyl ketone was added dropwise over 6 hours. After the completion of dropwise addition, the reaction was further allowed to proceed at 80°C for 2 hours. The resulting reaction solution was allowed to cool and then poured in 680 ml of heptane/120 ml of ethyl acetate, and the powder precipitated was collected by filtration and dried, as a result, 17.6 g of Resin (A4) was obtained. The weight average molecular weight of the obtained resin was 7,000 and the dispersity (Mw/Mn) was 1.55.

Resin (A4):



Weight average molecular weight: 7,000

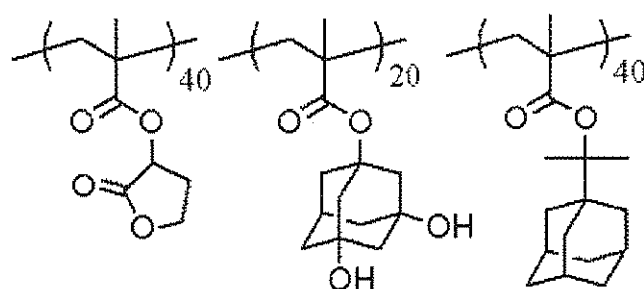
Dispersity: 1.55

Molar compositional ratio: 40/20/40

Synthesis Example 5 (Synthesis of Resin (A5)):

Under a nitrogen stream, 8.4 g of methyl isobutyl ketone was charged into a three-neck flask and heated at 80°C. Thereto, a solution obtained by dissolving 9.4 g of 2-(1-adamantyl)propan-2-yl methacrylate, 4.5 g of 3,5-dihydroxy-1-adamantyl methacrylate, 6.1 g of β -methacryloyloxy- γ -butyrolactone and azobisisobutyronitrile corresponding to 7 mol% based on the entire monomer amount, in 75.3 g of methyl isobutyl ketone was added dropwise over 6 hours. After the completion of dropwise addition, the reaction was further allowed to proceed at 80°C for 2 hours. The resulting reaction solution was allowed to cool and then poured in 720 ml of heptane/80 ml of ethyl acetate, and the powder precipitated was collected by filtration and dried, as a result, 18.5 g of Resin (A5) was obtained. The weight average molecular weight of the obtained resin was 6,000 and the dispersity (Mw/Mn) was 1.75.

Resin (A5):



Weight average molecular weight: 6,000

Dispersity: 1.75

Molar compositional ratio: 40/20/40

Resist Composition (A1):

A solution having a solid content concentration of 5.8 mass% obtained by dissolving the components shown below in a mixed solvent of polyethylene glycol monomethyl ether acetate/polyethylene glycol monomethyl ether (60:40) was filtered through a polyethylene

filter having a pore size of 0.1 μm to prepare Resist Composition (A1).

Resin (A1): 1.83 g, triphenylsulfonium nonaflate: 69.6 mg, diphenylaniline: 8.7 mg, and PF6320 (fluorine-containing surfactant produced by OMNOVA): 1.7 mg.

Resist Compositions (A2) to (A5):

Resist Compositions (A2) to (A5) were prepared in the same manner except for using Resins (A2) to (A5) in place of Resin (A1).

Comparative Example 1:

An organic antireflection film, ARC29A (produced by Nissan Chemical Industries, Ltd.), was coated on a 8-inch silicon wafer and baked at 205°C for 60 seconds to form a 78-nm antireflection film, and Resist Composition (A1) prepared above was coated thereon and baked at 120°C for 60 seconds to form a 150-nm resist film. The obtained wafer was subjected to pattern exposure using an ArF excimer laser scanner (NA: 0.75). Thereafter, the resist film was heated at 120°C for 60 seconds, developed with an aqueous tetramethylammonium hydroxide solution (2.38 mass%) (positive tone developer) for 30 seconds (positive tone development), and rinsed with pure water to obtain a pattern having a pitch of 600 nm and a line width of 450 nm. Furthermore, the resist film was developed with butyl acetate (negative tone developer) for 30 seconds (negative tone development) and rinsed with 1-hexanol (rinsing solution for negative tone development) for 30 seconds, and thereafter, the wafer was rotated at a rotation number of 3,000 rpm for 30 seconds to obtain a 150-nm (1:1) line-and-space resist pattern.

Examples 1 to 4:

In the same manner as in Comparative Example 1, 150-nm (1:1) resist patterns were obtained using Resist Compositions (A2) to (A5), respectively.

Evaluation of Line Edge Roughness (LER):

The 150-nm (1:1) line-and-space resist patterns obtained in Comparative Example 1 and Examples 1 to 4 were observed by a scanning microscope (S9260, manufactured by Hitachi Ltd.). With respect to the range of 2 μm edge in the longitudinal direction of the 150-nm line pattern, the distance from the reference line where the edge should be present was measured at 50 points and after determining the standard deviation, 3σ (unit: nm) was calculated. A smaller value indicates better performance.

The results are shown in Table 1.

Table 1

	Resist Composition	Resin	Molecular Weight	Dispersity	LER (nm)
Comparative Example 1	A1	A1	7000	1.80	9.5
Example 1	A2	A2	5800	1.53	6.8
Example 2	A3	A3	2800	1.26	5.2
Example 3	A4	A4	7000	1.55	7.0
Example 4	A5	A5	6000	1.75	7.1

Example 5:

An organic antireflection film, ARC29A (produced by Nissan Chemical Industries, Ltd.), was coated on a 8-inch silicon wafer and baked at 205°C for 60 seconds to form a 78-nm antireflection film, and Resist Composition (A2) prepared above was coated thereon and baked at 120°C for 60 seconds to form a 150-nm resist film. The obtained wafer was subjected to pattern exposure using an ArF excimer laser scanner (NA: 0.75). Thereafter, the resist film was heated at 120°C for 60 seconds, developed with an aqueous tetramethylammonium hydroxide solution (2.38 mass%) (positive tone developer) for 30 seconds (positive tone development), and rinsed with pure water to obtain a pattern having a pitch of 600 nm and a line width of 450 nm. Furthermore, the resist film was developed with ethyl acetate (negative tone developer) for 30 seconds (negative tone development) and rinsed with 1-hexanol (rinsing solution for negative tone development) for 30 seconds, and thereafter, the wafer was rotated at a rotation number of 3,000 rpm for 30 seconds to obtain a 150-nm (1:1) line-and-space resist pattern.

Examples 6 to 18:

In the same manner as in Example 5, 150-nm (1:1) resist patterns were obtained by using Resist Composition (A2) and combining the negative tone developer and the rinsing solution for negative tone development as shown in Table 2.

Evaluation of Dimensional Uniformity:

The 150-nm (1:1) line-and-space resist patterns obtained in Examples 1 and 5 to 18 each was measured for the dimension at 50 portions at intervals of 2 nm by using a scanning microscope (S9260, manufactured by Hitachi Ltd.). The standard deviation of 50 portions was determined, and 3σ (unit: nm) was calculated. A smaller value indicates better performance. The results are shown in Table 2.

Table 2

	Negative Tone Developer	Rinsing Solution for Negative Tone Development	Dimensional Uniformity (nm)
Example 1	butyl acetate	1-hexanol	3.1
Example 5	ethyl acetate	1-hexanol	8.4
Example 6	isoamyl acetate	1-hexanol	3.1
Example 7	methyl isobutyl ketone	1-hexanol	3.5
Example 8	2-hexanone	1-hexanol	2.7
Example 9	methyl ethyl ketone	1-hexanol	9.6
Example 10	dipropyl ether	1-hexanol	11.6
Example 11	dibutyl ether	1-hexanol	4.2
Example 12	butyl acetate/2-hexanone (80/20)	1-hexanol	3.5
Example 13	isoamyl acetate/ dibutyl ether (70/30)	1-hexanol	3.8
Example 14	isoamyl acetate	1-heptanol	8.8
Example 15	isoamyl acetate	2-heptanol	3.3
Example 16	isoamyl acetate	decane	4.3
Example 17	isoamyl acetate	dodecane	9.2
Example 18	isoamyl acetate	1-heptanol/decane (50/50)	3.7

The vapor pressure and boiling point of each of the solvent for negative tone developer and the solvent for rinsing solution for negative tone development used in Examples 1 and 5 to 18 are shown in Table 3.

Table 3

Solvent Name	Vapor Pressure (20°C)	Boiling Point
butyl acetate	1.2 kPa	126°C
ethyl acetate	10 kPa	77°C
isoamyl acetate	0.53 kPa	142°C
methyl isobutyl ketone	2.1 kPa	117-118°C
2-hexanone	0.36 kPa	126-128°C
methyl ethyl ketone	10.5 kPa	80°C
dipropyl ether	8.3 kPa	88-90°C
dibutyl ether	0.64 kPa	142°C
1-hexanol	0.13 kPa	157°C
1-heptanol	0.015 kPa	175°C
2-heptanol	0.13 kPa	150-160°C
decane	0.17 kPa	174.2°C
dodecane	0.040 kPa	216.2°C

Example 19:

An organic antireflection film, ARC29A (produced by Nissan Chemical Industries, Ltd.), was coated on a 8-inch silicon wafer and baked at 205°C for 60 seconds to form a 78-nm antireflection film, and Resist Composition (A2) prepared above was coated thereon and baked at 120°C for 60 seconds to form a 150-nm resist film. The obtained wafer was subjected to pattern exposure using an ArF excimer laser scanner (NA: 0.75). Subsequently, the resist film was developed with butyl acetate (negative tone developer) for 30 seconds (negative tone development) and rinsed with 1-hexanol (rinsing solution for negative tone development) for 30 seconds, and thereafter, the wafer was rotated at a rotation number of 3,000 rpm for 30 seconds to obtain a pattern having a pitch of 400 nm and a line width of 200 nm.

Examples 20 to 29:

In the same manner as in Example 19, patterns having a pitch 400 nm and a line width of 200 nm were obtained using Resist Compositions (A2) to (A5) and (B1) to (B6), respectively. The formulations of Resist Compositions (A2) to (A5) are as described above, and the formulations of Resist Compositions (B1) to (B6) are shown in Table 5 below.

Evaluation of Line Edge Roughness (LER):

The patterns having a pitch of 400 nm and a line width of 200 nm obtained in Examples 19 to 29 were observed by a scanning microscope (S9260, manufactured by Hitachi

Ltd.). With respect to the range of 2 μm edge in the longitudinal direction of the 200-nm line pattern, the distance from the reference line where the edge should be present was measured at 50 points and after determining the standard deviation, 3σ (unit: nm) was calculated. A smaller value indicates better performance. The results are shown in Tables 4 and 5.

Table 4

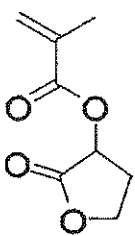
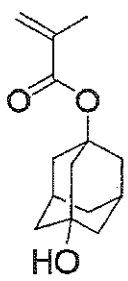
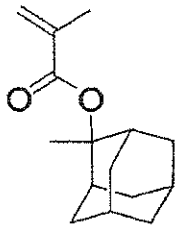
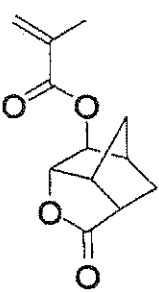
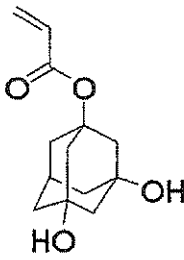
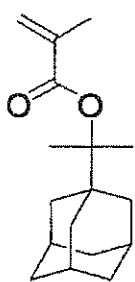
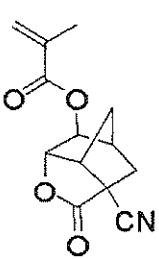
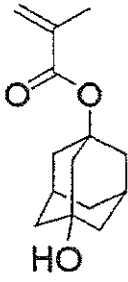
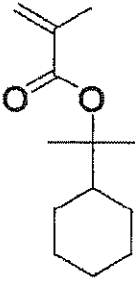
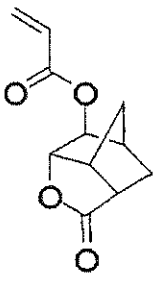
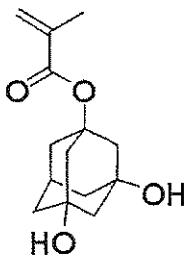
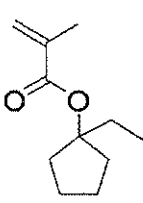
Example	Resist Composition	Resin	Molecular Weight	Dispersity	LER (nm)
19	A1	A1	7000	1.80	8.9
20	A2	A2	5800	1.53	6.3
21	A3	A3	2800	1.26	5.3
22	A4	A4	7000	1.55	7.2
23	A5	A5	6000	1.75	7.4

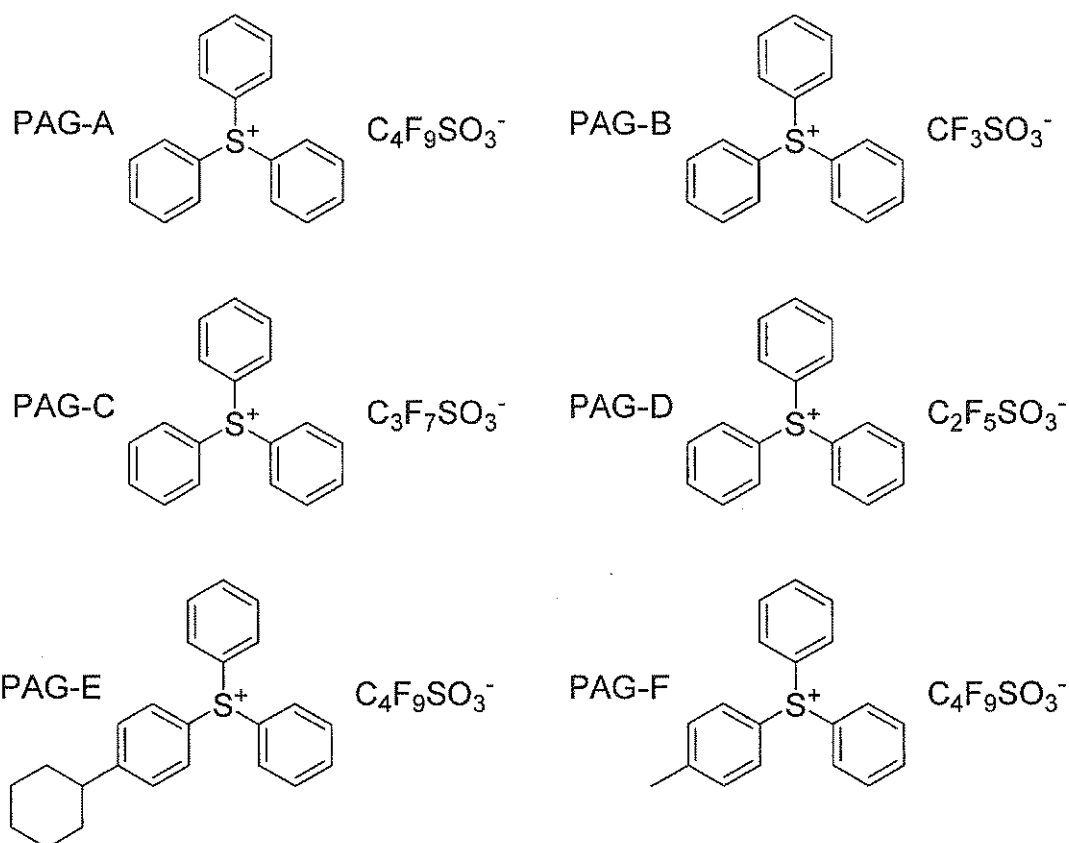
Table 5

Example	Resist Composition	Resin (10 g)	Acid Generator (g)	Basic Compound (g)	Surfactant (0.02 g)	Solvent (mass ratio)	LER (nm)
24	B1	A6	PAG-A (0.50)	PEA (0.05)	W-5	A1/B1 (60/40)	6.0
25	B2	A7	PAG-E (0.40)	DIA (0.05)	W-2	A1/B2 (40/60)	5.8
26	B3	A8	PAG-B (0.80)	TPA (0.06)	W-6	A2/B3 (95/5)	6.3
27	B4	A9	PAG-C (0.60)	PBI (0.07)	W-1	A2/B1 (50/50)	6.1
28	B5	A7	PAG-F (0.20)	DIA (0.10)	W-4	A2/B2 (40/60)	6.5
29	B6	A9	PAG-D (0.90)	TOA (0.08)	W-3	A1/B3 (90/10)	6.2

The abbreviations used in Examples indicate those set forth as specific examples above or the followings.

Table 6

No.	Monomer (1)	Monomer (2)	Monomer (3)	Compositional Ratio (by mol)	Mw	Mw/Mn
A6				30/30/40	5000	1.4
A7				50/20/30	5500	1.5
A8				50/10/40	4800	1.5
A9				40/20/40	5300	1.6



DIA: 2,6-diisopropylaniline

TPA: triphenylamine

PEA: N-phenyldiethanolamine

TOA: trioctylamine

PBI: 2-phenylbenzimidazole

W-1: Megaface F176 (produced by Dainippon Ink & Chemicals, Inc.) (fluorine-containing)

W-2: Megaface R08 (produced by Dainippon Ink & Chemicals, Inc.) (fluorine and silicon-containing)

W-3: Polysiloxane Polymer KP-341 (produced by Shin-Etsu Chemical Co., Ltd.) (silicon-containing)

W-4: Troysol S-366 (produced by Troy Chemical)

W-5: PF656 (produced by OMNOVA, fluorine-containing)

W-6: PF6320 (produced by OMNOVA, fluorine-containing)

A1: propylene glycol monomethyl ether acetate

A2: cyclohexanone

B1: propylene glycol monomethyl ether

B2: ethyl lactate

B3: γ -butyrolactone

As apparent from these Examples, thanks to the combination of resist composition, negative tone developer and rinsing solution for negative tone development of the present invention, the line edge roughness is reduced and furthermore, the dimensional uniformity is high.

INDUSTRIAL APPLICABILITY

By the pattern forming method and the resist composition of the present invention, a pattern with reduced line edge roughness and high dimensional uniformity is obtained. In particular, a highly practical negative tone development technique and a double development technique using the same are provided. These techniques enables finer patterning under the same light source as that in conventional technology. The pattern forming method of the present invention is suitably used in the process of producing a semiconductor such as IC, in the production of a circuit board for liquid crystal, thermal head and the like, and in the lithography process of other photo-applications.

While the invention has been described in detail and with reference to specific embodiments thereof, it will be apparent to one skilled in the art that various changes and modifications can be made therein without departing from the spirit and scope thereof.

This application is based on Japanese Patent Application (Japanese Patent Application No. 2007-106175) filed on April 13, 2007 and Japanese Patent Application (Japanese Patent Application No. 2007-198054) filed on July 30, 2007, and their contents are incorporated herein by way of reference.

CLAIMS

1. A pattern forming method, comprising:
 - (i) a step of applying a resist composition whose solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation and which contains a resin having an alicyclic hydrocarbon structure and a dispersity of 1.7 or less and being capable of increasing a polarity of the resin by an action of an acid;
 - (ii) an exposure step; and
 - (iv) a development step using a negative tone developer.
2. A pattern forming method, comprising:
 - (i) a step of applying a resist composition whose solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation and which contains a resin having an alicyclic hydrocarbon structure and a weight average molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;
 - (ii) an exposure step; and
 - (iv) a development step using a negative tone developer.
3. A pattern forming method, comprising:
 - (i) a step of applying a resist composition whose solubility in a negative tone developer decreases upon irradiation with an actinic ray or radiation and which contains a resin having an alicyclic hydrocarbon structure, a dispersity of 1.7 or less and a weight average molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;
 - (ii) an exposure step; and
 - (iv) a development step using a negative tone developer.
4. The pattern forming method according to any one of claims 1 to 3,
wherein (iv) the development step using a negative tone developer is a step performed using a developer containing at least one kind of a solvent selected from organic solvents and having a vapor pressure of 5.0 kPa or less at 20°C.
5. The pattern forming method according to any one of claims 1 to 4, which further comprises:
 - (vi) a washing step using a rinsing solution containing an organic solvent.

6. The pattern forming method according to claim 5,
wherein the rinsing solution containing an organic solvent is a rinsing solution having a vapor pressure of 0.1 kPa or more at 20°C.

7. The pattern forming method according to any one of claims 1 to 6, which further comprises:

(iii) a development step using a positive tone developer.

8. A resist composition for negative tone development, comprising:

(a1) a resin having an alicyclic hydrocarbon structure and a dispersity of 1.7 or less and being capable of increasing a polarity of the resin by an action of an acid;

(B) a photo-acid generator; and

(C) a solvent.

9. A resist composition for negative tone development, comprising:

(a2) a resin having an alicyclic hydrocarbon structure and a weight average molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;

(B) a photo-acid generator; and

(C) a solvent.

10. A resist composition for negative tone development, comprising:

(a3) a resin having an alicyclic hydrocarbon structure, a dispersity of 1.7 or less and a weight average molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;

(B) a photo-acid generator; and

(C) a solvent.

11. A resist composition for multiple development, comprising:

(a1) a resin having an alicyclic hydrocarbon structure and a dispersity of 1.7 or less and being capable of increasing a polarity of the resin by an action of an acid;

(B) a photo-acid generator; and

(C) a solvent.

12. A resist composition for multiple development, comprising:

(a2) a resin having an alicyclic hydrocarbon structure and a weight average

molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;

(B) a photo-acid generator; and

(C) a solvent.

13. A resist composition for multiple development, comprising:

(a3) a resin having an alicyclic hydrocarbon structure, a dispersity of 1.7 or less and a weight average molecular weight of 6,000 or less and being capable of increasing a polarity of the resin by an action of an acid;

(B) a photo-acid generator; and

(C) a solvent.

14. A developer for negative tone development, which is used in the resist composition claimed in any one of claims 8 to 13, the developer comprising an organic solvent and having a vapor pressure of 5 kPa or less at 20°C.

15. A rinsing solution for negative tone development, which is used in the pattern forming method claimed in any one of claims 8 to 13, the rinsing solution comprising an organic solvent and having a vapor pressure of 0.1 kPa or more at 20°C.