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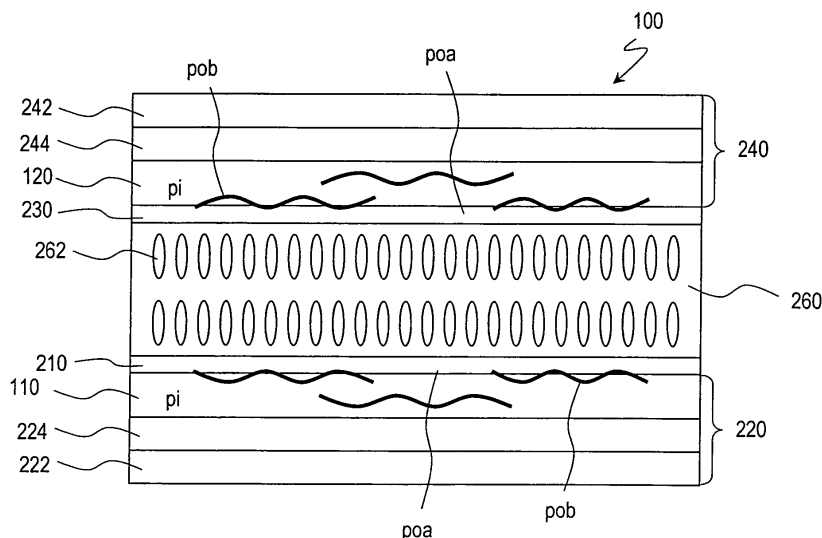
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(54) **LIQUID CRYSTAL DISPLAY AND METHOD FOR PRODUCING THE SAME**

(57) A liquid crystal display device (100) according to the present invention includes: an active matrix substrate (220) having a pixel electrode (224); a counter substrate (240) having a counter electrode (244); a vertical-alignment type liquid crystal layer (260); and alignment sustaining layers (210, 230) provided between the active matrix substrate (220) and the liquid crystal layer (260) and between the counter substrate (240) and the liquid

crystal layer (260). The alignment sustaining layers (210, 230) contain a polymerization product (poa) resulting from polymerization of a polyfunctional monomer. At least one of the active matrix substrate and the counter substrate further includes an alignment film (110, 120) located on the liquid crystal layer side, the alignment film (110, 120) containing polyimide (pi) and a polymerization product (pob) resulting from polymerization of a polyfunctional monomer.

FIG. 1



Description**TECHNICAL FIELD**

[0001] The present invention relates to a liquid crystal display device and a method of producing the same.

BACKGROUND ART

[0002] Liquid crystal display devices are used not only as small-sized display devices, e.g., the display sections of mobile phones, but also as large-sized television sets. Liquid crystal display devices of the TN (Twisted Nematic) mode, which have often been used conventionally, have relatively narrow viewing angles. In recent years, however, liquid crystal display devices with wide viewing angles have been produced, e.g., the IPS (In-Plane Switching) mode and the VA (Vertical Alignment) mode. Among such modes with wide viewing angles, the VA mode is adopted in a large number of liquid crystal display devices because of an ability to realize a high contrast ratio.

[0003] As one kind of VA mode, the MVA (Multi-domain Vertical Alignment) mode is known, under which a plurality of liquid crystal domains are created in one pixel region. An MVA-mode liquid crystal display device includes alignment regulating structures provided on the liquid-crystal-layer side of at least one of a pair of opposing substrates, between which a vertical-alignment type liquid crystal layer is interposed. The alignment regulating structures may be linear slits (apertures) or ribs (protruding structures) that are provided on electrodes, for example. The alignment regulating structures provide alignment regulating forces from one side or both sides of the liquid crystal layer, thus creating a plurality of liquid crystal domains (typically four liquid crystal domains) with different alignment directions, whereby the viewing angle characteristics are improved.

[0004] Also known as one kind of VA mode is the CPA (Continuous Pinwheel Alignment) mode. In a generic liquid crystal display device of the CPA mode, pixel electrodes of a highly symmetrical shape are provided, and on a counter electrode, protrusions are provided corresponding to the centers of liquid crystal domains. These protrusions are also referred to as rivets. When a voltage is applied, in accordance with an oblique electric field which is created with the counter electrode and a highly symmetrical pixel electrode, liquid crystal molecules take an inclined alignment of a radial shape. Moreover, the inclined alignment of the liquid crystal molecules are stabilized due to the alignment regulating forces of side slopes of the rivets. Thus, the liquid crystal molecules in one pixel are aligned in a radial shape, thereby improving the viewing angle characteristics.

[0005] Unlike in TN-mode liquid crystal display devices in which the pretilt direction of liquid crystal molecules is regulated by an alignment film, alignment regulating forces in an MVA-mode liquid crystal display device are applied to the liquid crystal molecules by linear slits or ribs. Therefore, depending on distances from the slits and ribs, the alignment regulating forces for the liquid crystal molecules within a pixel region will differ, thus resulting in differing response speeds of the liquid crystal molecules within the pixel. Similarly, also in the CPA mode, the response speeds of the liquid crystal molecules will differ within the pixel, and the differences in response speed will become more outstanding as the pixel electrodes increase in size. Furthermore, in a VA-mode liquid crystal display device, the light transmittance in the regions in which slits, ribs, or rivets are provided is low, thus making it difficult to realize a high luminance.

[0006] In order to avoid the above problems, use of an alignment film for applying alignment regulating forces to liquid crystal molecules in a VA-mode liquid crystal display device is also known, such that the liquid crystal molecules will tilt from the normal direction of a principal face of the alignment film in the absence of an applied voltage (see, for example, Patent Document 1). The alignment film regulates the liquid crystal molecules so that the liquid crystal molecules are inclined from the normal direction of a principal face thereof even in the absence of an applied voltage, whereby an improvement in response speed is realized. Furthermore, since the alignment film regulates the pretilt azimuth of liquid crystal molecules so that the liquid crystal molecules within one pixel will be symmetrically aligned, the viewing angle characteristics are improved. In a liquid crystal display device disclosed in Patent Document 1, four liquid crystal domains are formed in a liquid crystal layer in accordance with a combination of two alignment regions of a first alignment film and two alignment regions of a second alignment film, whereby a wide viewing angle is realized.

[0007] However, a liquid crystal display device that has a pretilt conferred thereto may not have adequate long-term reliability. Therefore, use of a Polymer Sustained Alignment Technology (hereinafter referred to as the "PSA technique") to improve the long-term reliability is known (see Patent Document 2, for example). The PSA technique is a technique where a small amount of polymerizable compound (e.g., a polymerizable monomer) is mixed in a liquid crystal material, and after the liquid crystal material is introduced between an active matrix substrate and a counter substrate, the polymerizable compound is irradiated with ultraviolet light to form a polymerization product while applying a voltage between the pixel electrodes and the counter electrode. By using the PSA technique, the alignment state of the liquid crystal molecules when generating the polymerization product is maintained (stored), thus making it possible to control the pretilt direction of the liquid crystal molecules.

CITATION LIST**PATENT LITERATURE**5 **[0008]**

[Patent Document 1] Japanese Laid-Open Patent Publication No. 11-352486

[Patent Document 2] Japanese Laid-Open Patent Publication No. 2008-076950

SUMMARY OF INVENTION**TECHNICAL PROBLEM**

15 **[0009]** Generally speaking, when a liquid crystal display device keeps displaying the same pattern for a long time, the previous pattern may remain even after the displayed image is changed. Such a phenomenon is also called image sticking. For example, after displaying white in a partial region of the screen and black in another region for a long time, if the entire liquid crystal panel is caused to display the same intermediate gray scale level, the region previously displaying white may appear slightly brighter than the region previously displaying black.

20 **[0010]** One cause of such image sticking is charge accumulation. The amount of charge accumulated in the region which was displaying black is different from the amount of charge accumulated in the region which was displaying white, and an electric field occurs because impurity ions in the liquid crystal accumulate at the interface between the alignment film and the liquid crystal layer. Therefore, when entirely switched to the same gray scale level, different voltages are applied to the layers of liquid crystal in the respective regions which were displaying white and black, thus being perceived as image sticking.

25 **[0011]** Image sticking caused by such charge accumulation can be somewhat suppressed by applying voltages of inverted polarities to the respective pixels. An image sticking caused by charge accumulation is also called DC image sticking, and the driving which involves applying voltages of inverted polarities for the sake of suppressing DC image sticking is also called polarity inversion driving. Strictly speaking, even with polarity inversion driving, it is difficult to apply voltages of completely symmetric polarities and thus a resultant image sticking may be perceived as flickering.

30 **[0012]** Moreover, image sticking will also occur when minute changes in the pretilt angle occur. When the pretilt angle changes, the V-T characteristics are affected, and thus the transmittance will vary even if the same voltage is applied. Since the applied voltage when displaying white is different from the applied voltage when displaying black, the amount of change in the pretilt angle will vary depending on the applied voltage. When later entirely switched to the same gray scale level, image sticking may be perceived due to changes in the pretilt angle. This is considered to be because the tilt of the liquid crystal molecules increases (i.e., the pretilt angle decreases) as the applied voltage increases. Such image sticking cannot be suppressed even by performing polarity inversion driving, and is also called AC image sticking. As is also disclosed in Patent Document 2, by using the aforementioned PSA technique, it is possible to suppress AC image sticking to a certain extent. However, some image sticking may still persist even if a polymerization product is formed from a monomer which is mixed in the liquid crystal material.

40 **[0013]** The present invention has been made in view of the above problems, and an objective thereof is to provide a liquid crystal display device whose image sticking is suppressed and a method of producing the same.

SOLUTION TO PROBLEM

45 **[0014]** A liquid crystal display device according to the present invention is a liquid crystal display device comprising: an active matrix substrate having a pixel electrode; a counter substrate having a counter electrode; a vertical-alignment type liquid crystal layer provided between the active matrix substrate and the counter substrate; and alignment sustaining layers provided between the active matrix substrate and the liquid crystal layer and between the counter substrate and the liquid crystal layer, the alignment sustaining layers containing a polymerization product resulting from polymerization of a polyfunctional monomer, wherein, at least one of the active matrix substrate and the counter substrate further includes an alignment film; and the alignment film contains polyimide and a polymerization product resulting from polymerization of a polyfunctional monomer.

50 **[0015]** In one embodiment, the polyfunctional monomer of the polymerization product contained in the alignment film has a plurality of vinyl groups.

55 **[0016]** In one embodiment, the polyfunctional monomer of the polymerization product contained in the alignment film is represented by general formula (1) $\text{Pb1-Ab1-(Zb1-Ab2)}_n\text{-Pb2}$ (in general formula (1), Pb1 and Pb2 are, independently, acrylate, methacrylate, acrylamide, or methacrylamide; Ab1 and Ab2 represent, independently, 1,4-phenylene, 1,4-

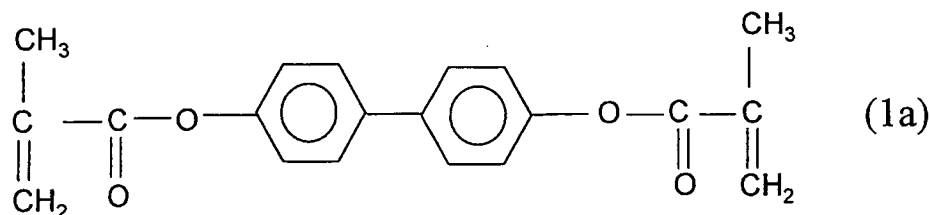
cyclohexane or 2, 5-thiophene, or naphthalene-(2,6)-diyl or anthracene-(2,7)-diyl; and Zb1 is a -COO- group, a -OCO- group, a -O- group, a -CONH- group or a single bond, where n is 0 or 1).

[0017] In one embodiment, at least one of Ab1 and Ab2 is substituted by at least one fluorine group.

[0018] In one embodiment, the polyfunctional monomer of the polymerization product contained in the alignment film includes a dimethacrylate monomer.

[0019] In one embodiment, the dimethacrylate monomer is represented by structural formula (1a).

[Formula 1]



[0020] In one embodiment, the polyfunctional monomer of the polymerization product contained in the alignment sustaining layers has a plurality of vinyl groups.

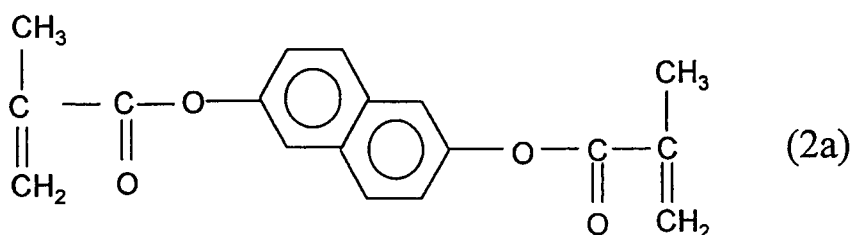
[0021] In one embodiment, the polyfunctional monomer of the polymerization product contained in the alignment sustaining layers is represented by general formula (2) Pa1-Aa1-(Za1-Aa2)_n-Pa2 (in general formula (2), Pa1 and Pa2 are, independently, acrylate, methacrylate, acrylamide, or methacrylamide; Aa1 and Aa2 represent, independently, 1,4-phenylene, 1,4-cyclohexane or 2,5-thiophene, or naphthalene-(2,6)-diyl or anthracene-(2,7)-diyl; and Za1 is a -COO- group, a -OCO- group, a -O- group, a -CONH- group or a single bond, where n is 0 or 1).

[0022] In one embodiment, at least one of Aa1 and Aa2 is substituted by at least one fluorine group.

[0023] In one embodiment, the polyfunctional monomer of the polymerization product contained in the alignment sustaining layers includes a dimethacrylate monomer.

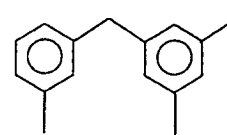
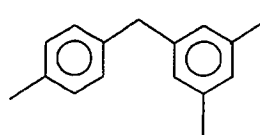
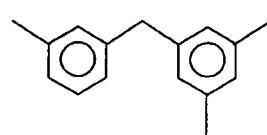
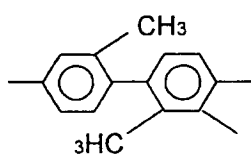
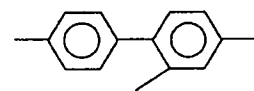
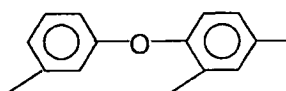
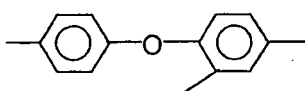
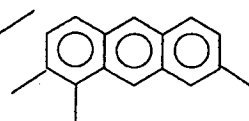
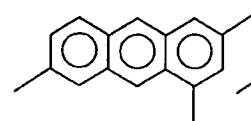
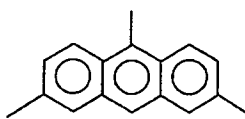
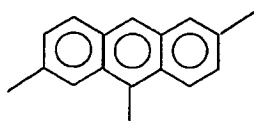
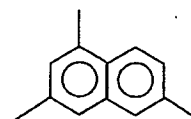
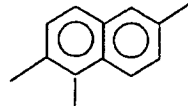
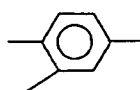
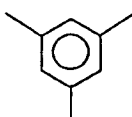
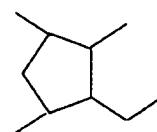
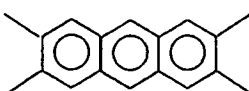
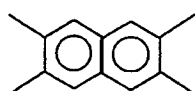
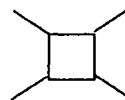
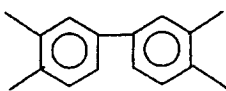
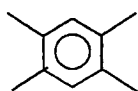
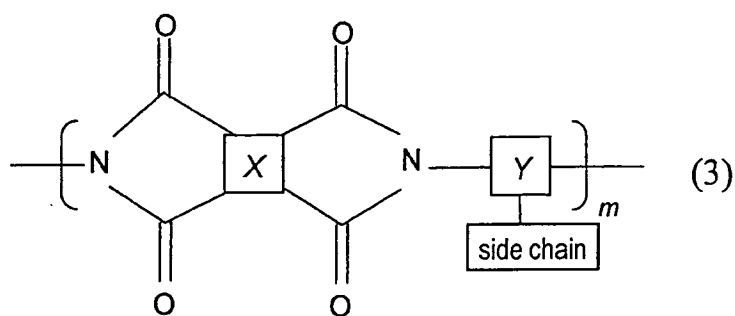
[0024] In one embodiment, the dimethacrylate monomer is represented by structural formula (2a).

[Formula 2]



[0025] In one embodiment, the polyimide is a polyimide including a structure represented by general formula (3).

[Formula 3]

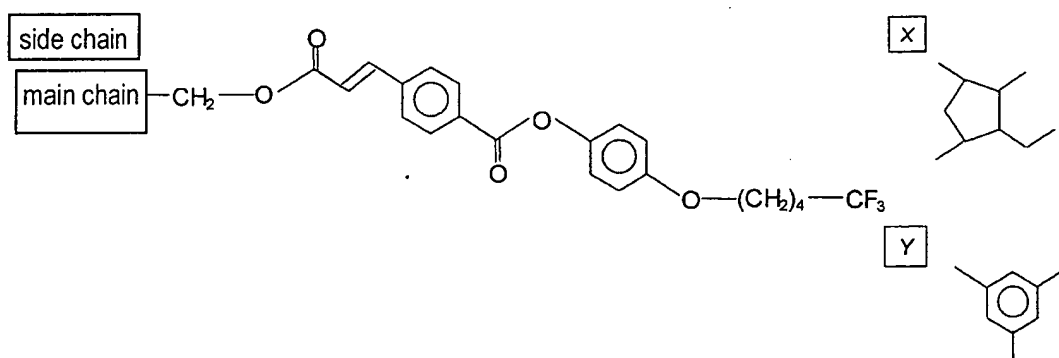
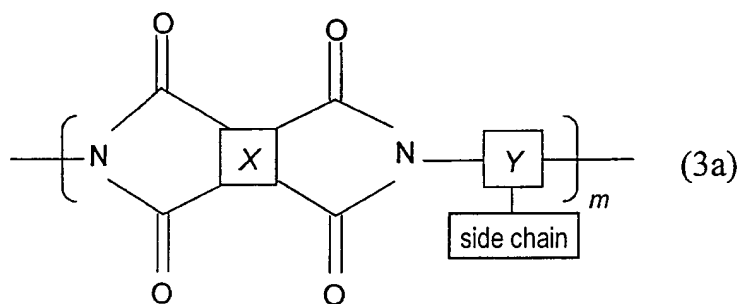


[0026] In one embodiment, the polyimide has a photoreactive functional group.

[0027] In one embodiment, the photoreactive functional group is one selected from the group consisting of a cinnamate group, a chalcone group, a tolan group, a coumarin group, and an azobenzene group.

[0028] In one embodiment, the polyimide is a polyimide including a structure represented by structural formula (3a).

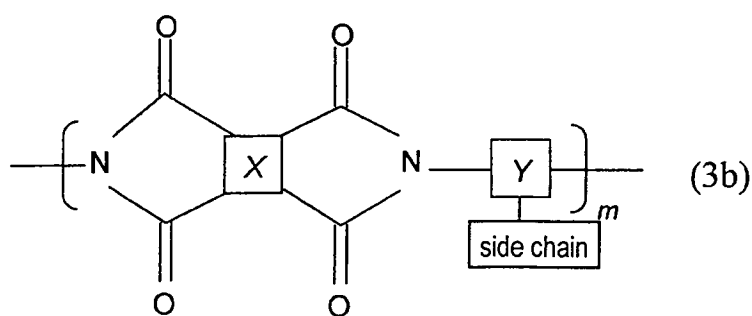
[Formula 4]



[0029] In one embodiment, the polyimide has a vertical alignment group in a side chain thereof.

[0030] In one embodiment, the polyimide is represented by structural formula (3b).

[Formula 5]



[0031] In one embodiment, the alignment film regulates liquid crystal molecules in the liquid crystal layer so that the liquid crystal molecules are inclined with respect to a normal direction of a principal face of the alignment film in the absence of an applied voltage.

[0032] In one embodiment, the liquid crystal display device has a plurality of pixels; and in each of the plurality of pixels, the liquid crystal layer has a plurality of liquid crystal domains having respectively different reference alignment

azimuths.

[0033] In one embodiment, the plurality of liquid crystal domains are four liquid crystal domains.

[0034] A method of producing a liquid crystal display device according to the present invention is a method of producing a liquid crystal display device, the method comprising the steps of: forming an active matrix substrate and a counter substrate; and forming a liquid crystal layer between the active matrix substrate and the counter substrate, and forming alignment sustaining layers between the active matrix substrate and the liquid crystal layer and between the counter substrate and the liquid crystal layer, the alignment sustaining layers containing a polymerization product resulting from polymerization of a polyfunctional monomer, wherein the step of forming the active matrix substrate and the counter substrate comprises a step of providing a first insulative substrate having a pixel electrode provided thereon and a second insulative substrate having a counter electrode provided thereon, and a step of forming an alignment film on at least one of the pixel electrode and the counter electrode, the alignment film containing polyimide and a polymerization product resulting from polymerization of a polyfunctional monomer.

[0035] In one embodiment, the step of forming the alignment film comprises: a step of providing an alignment film material containing a precursor of the polyimide and the polyfunctional monomer; and a step of applying the alignment film material and performing a heat treatment, thereby imidizing the precursor of the polyimide to form the polyimide and forming a polymerization product resulting from polymerization of the polyfunctional monomer.

[0036] In one embodiment, the step of performing a heat treatment comprises a step of conducting a pre-bake and thereafter conducting a post-bake at a higher temperature than in the pre-bake.

[0037] In one embodiment, the step of forming the alignment sustaining layers comprises a step of performing light irradiation.

ADVANTAGEOUS EFFECTS OF INVENTION

[0038] According to the present invention, a liquid crystal display device whose image sticking is suppressed can be provided.

BRIEF DESCRIPTION OF DRAWINGS

[0039]

[FIG. 1] A schematic diagram of an embodiment of a liquid crystal display device according to the present invention.

[FIG. 2] (a) to (c) are schematic diagrams each illustrating a method of producing a liquid crystal display device of the present embodiment.

[FIG. 3] (a) is a schematic diagram showing a side chain of polyimide; (b) is a schematic diagram showing emergence of an ionized site; (c) is a schematic diagram showing the ionized site dissolving into a liquid crystal layer; and (d) is a schematic diagram showing how dissolution of the ionized site into the liquid crystal layer is suppressed in the liquid crystal display device of the present embodiment.

[FIG. 4] (a) is a schematic diagram showing dissolution of unreacted cross-linker into a liquid crystal layer; and (b) is a schematic diagram showing how the dissolution of unreacted cross-linker into the liquid crystal layer is suppressed in the liquid crystal display device of the present embodiment.

[FIG. 5] (a) is a schematic diagram of an alignment film of the liquid crystal display device of the present embodiment; (b) is a schematic diagram of the alignment film; and (c) is a schematic diagram showing alignment directions of liquid crystal molecules in the centers of liquid crystal domains.

[FIG. 6] A schematic diagram showing alignment directions of liquid crystal molecules in one liquid crystal domain of the liquid crystal display device of the present embodiment.

[FIG. 7] A schematic diagram showing alignment treatment directions for first and second alignment films as viewed from the viewer's side.

[FIG. 8] A schematic diagram showing alignment treatment directions for the first and second alignment films as viewed from the viewer's side.

[FIG. 9] A graph showing amounts of change in the pretilt angle against powering time.

[FIG. 10] A graph showing amounts of change in the pretilt angle against powering time.

DESCRIPTION OF EMBODIMENTS

[0040] Hereinafter, with reference to the drawings, an embodiment of a liquid crystal display device according to the present invention and a method of producing the same will be described.

[0041] FIG. 1 shows a schematic diagram of a liquid crystal display device **100** of the present embodiment. The liquid crystal display device **100** includes an active matrix substrate **220** having a first alignment film **110**, a counter substrate **240** having a second alignment film **120**, and a liquid crystal layer **260** provided between the active matrix substrate **220** and the counter substrate **240**. The liquid crystal display device **100** further includes a first alignment sustaining layer **210** located between the liquid crystal layer **260** and the active matrix substrate **220** and a second alignment sustaining layer **230** located between the liquid crystal layer **260** and the counter substrate **240**.

[0042] The active matrix substrate **220** further includes a first insulative substrate **222** and pixel electrodes **224**, such that the first alignment film **110** covers the pixel electrodes **224**. Moreover, the counter substrate **240** further includes a second insulative substrate **242** and a counter electrode **244**, such that the second alignment film **120** covers the counter electrode **244**. For example, the first and second insulative substrates **222** and **242** are transparent glass substrates.

[0043] The liquid crystal display device **100** includes pixels composing a matrix of a plurality of rows and a plurality of columns. On the active matrix substrate **220**, at least one switching element (e.g., thin film transistor (Thin Film Transistor: TFT))(not shown in the figure) is provided for each pixel, and the active matrix substrate **220** is also referred to a TFT substrate. In the present specification, a "pixel" refers to the smallest unit that expresses a specific gray scale level in displaying; in the case of multicolor displaying, a "pixel" corresponds to a unit that expresses a gray scale level of each of R, G, and B, for example, and is also referred to as a dot. A combination of an R pixel, a G pixel, and a B pixel composes a single color displaying pixel. A "pixel region" refers to a region of the liquid crystal device **100** that corresponds to a "pixel" in displaying.

[0044] Although not shown, a polarizer is provided on each of the active matrix substrate **220** and the counter substrate **240**. Therefore, the two polarizers are disposed so as to oppose each other with the liquid crystal layer **260** interposed therebetween. The transmission axes (polarization axes) of the two polarizers are positioned so as to be orthogonal to each other, such that one of them extends along the horizontal direction (row direction), whereas the other extends along the vertical direction (column direction).

[0045] Although not shown in the figure, the liquid crystal display device **100** includes a liquid crystal panel, a driving circuit for driving the liquid crystal panel, and a control circuit for controlling the driving circuit. As necessary, the liquid crystal display device **100** may include a backlight.

[0046] The first alignment sustaining layer **210** is located between the first alignment film **110** and the liquid crystal layer **260**. The second alignment sustaining layer **230** is located between the second alignment film **120** and the liquid crystal layer **260**. Each of the first and second alignment sustaining layers **210** and **230** contains a polymerization product **poa**. The polymerization product **poa** is formed through polymerization of a precursor, which is a polyfunctional monomer. Specifically, after a liquid crystal material having the polyfunctional monomer mixed thereto is introduced between the active matrix substrate **220** and the counter substrate **240**, the precursor is polymerized to form the polymerization product **poa**, and the polymerization product **poa** becomes phase-separated from the liquid crystal layer **260**, whereby the first and second alignment sustaining layers **210** and **230** are formed.

[0047] The first alignment film **110** contains polyimide **pi** and a polymerization product **pob**. Similarly, the second alignment film **120** contains polyimide **pi** and the polymerization product **pob**. Polyimide **pi** is formed by imidizing a precursor thereof. The polymerization product **pob** is formed through polymerization of a polyfunctional monomer (precursor). The polymerization is performed by applying heat or light to the polyfunctional monomer.

[0048] The first alignment film **110** is formed from an alignment film material. The alignment film material is obtained by allowing the precursor of polyimide **pi** and the polyfunctional monomer to be dissolved in a solvent. For example, after applying the alignment film material, a heat treatment is performed to evaporate the solvent and effect imidization and polymerization, whereby the first alignment film **110** containing the polyimide **pi** and the polymerization product **pob** is formed. The heat treatment is performed twice at different temperatures, for example. Similarly, after applying the alignment film material, a heat treatment is performed to evaporate the solvent and effect imidization and polymerization, whereby the second alignment film **120** containing the polyimide **pi** and the polymerization product **pob** is formed.

[0049] As from the above, in the liquid crystal display device **100** of the present embodiment, the polymerization product **poa** within the first alignment sustaining layer **210** and the polymerization product **pob** within the first alignment film **110** are present in the neighborhood of the boundary between the active matrix substrate **220** and the liquid crystal layer **260**. Similarly, the polymerization product **poa** within the second alignment sustaining layer **230** and the polymerization product **pob** within the second alignment film **120** are present in the neighborhood of the boundary between the counter substrate **240** and the liquid crystal layer **260**. In the following description of the present specification, the polymerization product **poa** facing the liquid crystal layer **260** may be referred to as an upper polymerization product

poa, whereas the polymerization product **pob** facing the insulative substrates **222** and **242** may be referred to as a lower polymerization product **pob**. Also, in the following description, a polyfunctional monomer that is the precursor of the upper polymerization product **poa** may be represented as a polyfunctional monomer **ma**, whereas a polyfunctional monomer that is the precursor of the lower polymerization product **pob** may be represented as a polyfunctional monomer **mb**.

[0050] The liquid crystal layer **260** contains a nematic liquid crystal material (liquid crystal molecules **262**) having negative dielectric anisotropy. The first alignment film **110** and the second alignment film **120** are each treated so that the pretilt angle of the liquid crystal molecules **262** is less than 90° with respect to the surface of the vertical alignment film. The pretilt angle of the liquid crystal molecules **262** is an angle between principal faces of the first alignment film **110** and the second alignment film **120** and the major axis of each liquid crystal molecule **262** that is regulated in a pretilt direction.

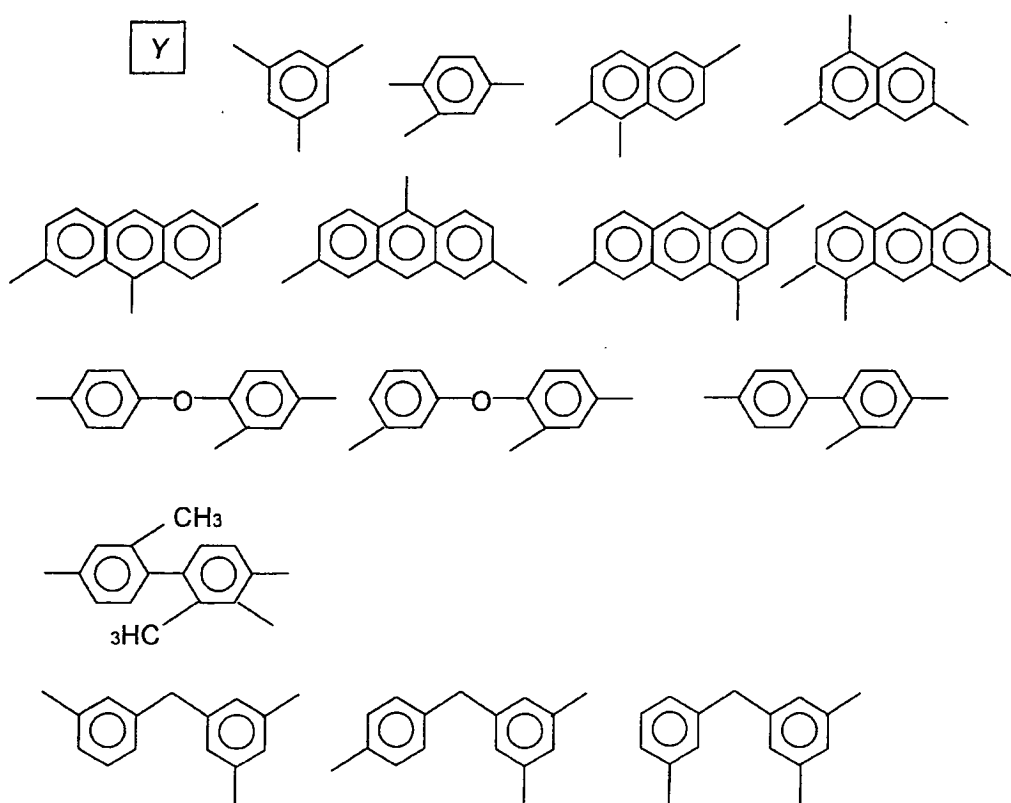
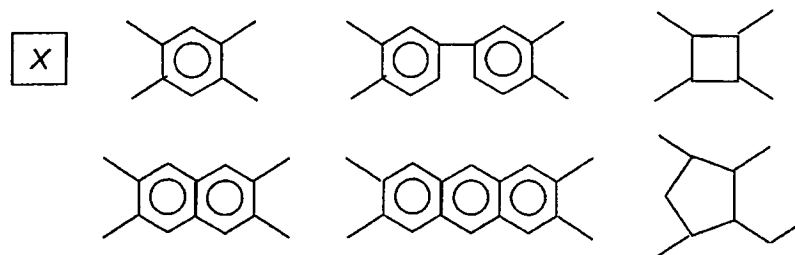
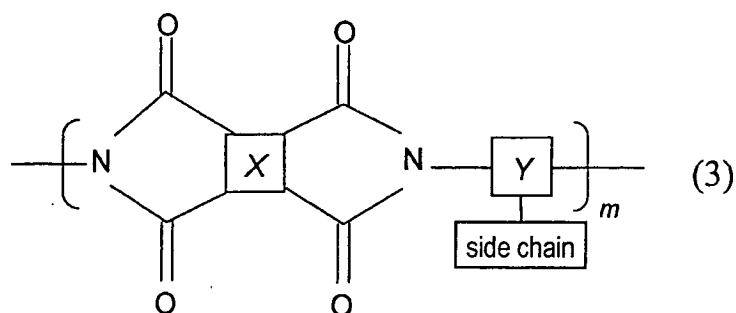
[0051] Although the liquid crystal layer **260** is of a vertical-alignment type, due to the first alignment film **110** and the second alignment film **120**, the liquid crystal molecules **262** in its neighborhood are slightly inclined from the normal directions of the principal faces of the first and second alignment films **110** and **120**. The pretilt angle is within a range from 85° to 89.7°, for example. The pretilt angle is measured by crystal rotation technique, for example. The side chain of the polyimide **pi** regulates the pretilt direction of the liquid crystal molecules **262**. In the following description, this component may also be referred to as a pretilt-angle-exhibiting component.

[0052] Note that the pretilt azimuth of the liquid crystal molecules **262** introduced by the first alignment film **110** is different from the pretilt azimuth of the liquid crystal molecules **262** introduced by the second alignment film **120**. For example, the pretilt azimuth of the liquid crystal molecules **262** introduced by the first alignment film **110** intersects, at 90°, the pretilt azimuth of the liquid crystal molecules **262** introduced by the second alignment film **120**. Herein, no chiral agent is added to the liquid crystal layer **260**, and when a voltage is applied across the liquid crystal layer **260**, the liquid crystal molecules **262** in the liquid crystal layer **260** take a twist alignment in accordance with the alignment regulating forces from the first and second alignment films **110** and **120**. However, a chiral agent may be added to the liquid crystal layer **260** as necessary. In combination with polarizers which are placed in crossed Nicols, the liquid crystal layer **260** performs displaying in a normally black mode.

[0053] Moreover, each of the first and second alignment films **110** and **120** may have a plurality of alignment regions for each pixel. For example, a portion of the first alignment film **110** may be masked, and after a predetermined region of the first alignment film **110** is irradiated with light from a certain direction, another region which was not irradiated with light may be irradiated with light from a different direction. Furthermore, the second alignment film **120** is also formed similarly. In this manner, regions that confer different alignment regulating forces can be formed in each of the first and second alignment films **110** and **120**.

[0054] The polyimide **pi** may include a photoreactive functional group in its main chain or side chain. The polyimide **pi** includes a structure represented by general formula (3), for example.

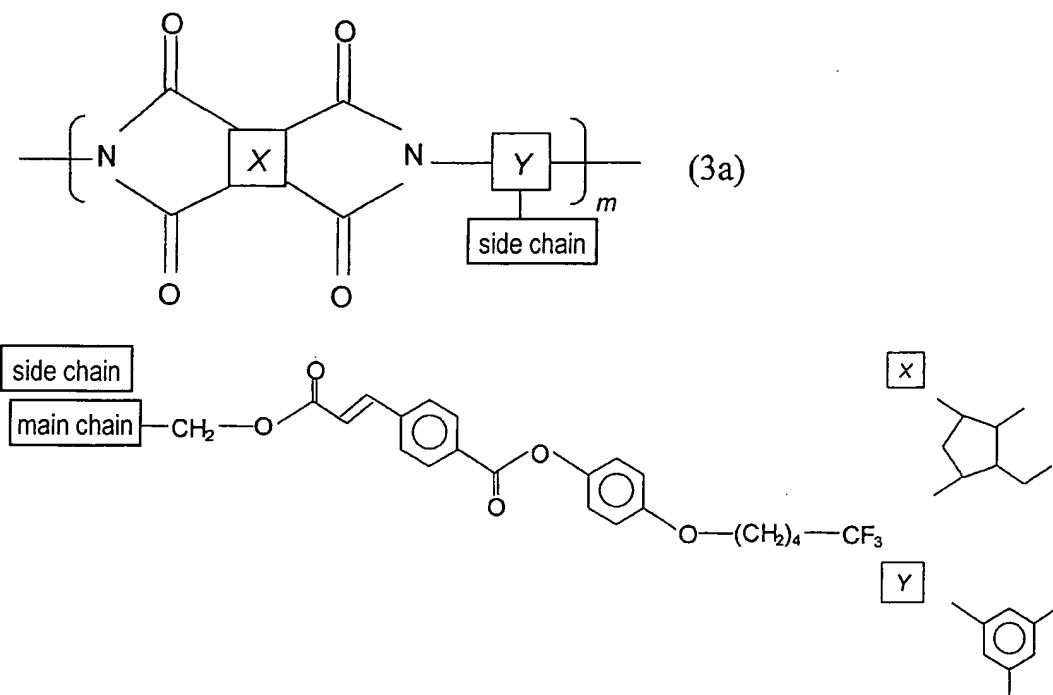
[Formula 6]



[0055] Moreover, the side chain of the polyimide **pi** may include a photoreactive functional group. In this case, a dimerization site is formed in the side chain through light irradiation. An alignment film containing such polyimide **pi** is also referred to as a photo-alignment film. The photoreactive functional group is a cinnamate group, a chalcone group, a tolan group, a coumarin group, or an azobenzene group, for example. In this case, by irradiating the first or second alignment film **110** or **120** with light from a direction which is oblique with respect to the normal direction of its principal face, an alignment regulating force is applied to the polyimide **pi** such that the liquid crystal molecules **262** are aligned

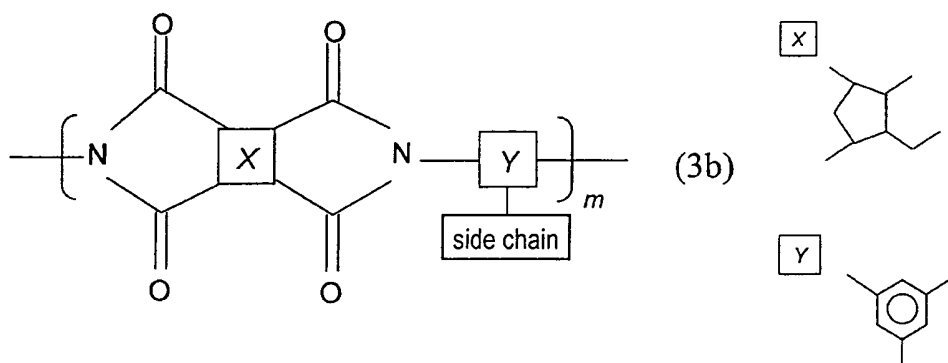
so as to be inclined from the normal directions of the principal faces of the first and second alignment films **110** and **120** in the absence of an applied voltage. Such polyimide **pi** is also referred to as a photo-alignable polyimide, and such a treatment is also referred to as a photo-alignment treatment. Since a photo-alignment treatment is performed without involving any contact, static electricity will not occur due to friction as in a rubbing treatment, and thus the production yield can be improved. Moreover, the side chain of the polyimide **pi** may contain a fluorine atom. When the side chain contains a fluorine atom, the aforementioned image sticking is suppressed to a certain extent. For example, the polyimide **pi** includes a structure represented by structural formula (3a).

[Formula 7]



[0056] Alternatively, the side chain of the polyimide **pi** may include a vertical alignment group. A polyimide **pi** having a vertical alignment group as a side chain is also referred to as a vertical-alignment type polyimide. After formation of the first and second alignment films **110** and **120**, the first and second alignment films **110** and **120** may be subjected to a rubbing treatment or an ion beam irradiation, whereby a pretilt can be conferred to the liquid crystal molecules **262**. For example, AL60101 manufactured by JSR Corporation is used as the precursor of the polyimide **pi**. In this case, the polyimide **pi** is represented by structural formula (3b).

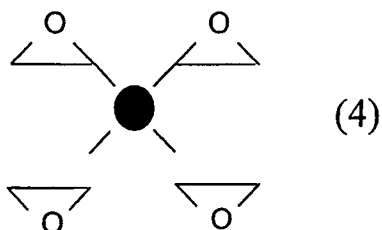
[Formula 8]



Herein, the side chain of the polyimide is a saturated aliphatic or unsaturated aliphatic compound.

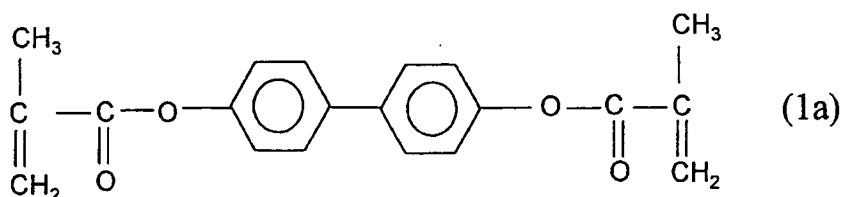
[0057] Moreover, a cross-linker may be mixed in the alignment film material. For example, the cross-linker is an epoxy-type cross-linker, e.g., a compound represented by structural formula (4). By using such a cross-linker, DC image sticking can be suppressed.

[Formula 9]



[0058] The polymerization product **pob** is obtained by polymerizing the polyfunctional monomer **mb**. For example, the polyfunctional monomer **mb** has a plurality of vinyl groups. For example, the polyfunctional monomer **mb** is represented by general formula (1) $\text{Pb1-Ab1-(Zb1-Ab2)}_n\text{-Pb2}$. In general formula (1), Pb1 and Pb2 are, independently, acrylate, methacrylate, acrylamide, or methacrylamide; Ab1 and Ab2 represent, independently, 1,4-phenylene, 1,4-cyclohexane or 2,5-thiophene, or naphthalene-(2,6)-diyl or anthracene-(2,7)-diyl; and Zb1 is a -COO- group, a -OCO- group, a -O- group, a -CONH- group or a single bond, where n is 0 or 1. Specifically, the polyfunctional monomer **mb** is biphenyldimethacrylate. The polyfunctional monomer **mb** is represented by structural formula (1a), for example.

[Formula 10]



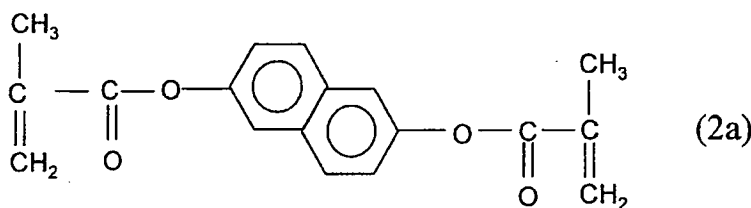
[0059] As for the polyfunctional monomer mb, at least one of Ab1 and Ab2 may be substituted by at least one fluorine group in the aforementioned general formula (1); for example, a part of the biphenyl group in biphenyldimethacrylate may be substituted by a fluorine group. When the polyfunctional monomer mb has any fluorine groups, more polyfunctional monomer mb and more polymerization product thereof will be present on the surface of the first and second alignment films **110** and **120**, thereby enhancing the effect of tilt angle stabilization. Moreover, it is preferable that the polyfunctional monomer mb has a symmetric structure; for example, it is preferable that an even number of hydrogen groups are substituted by fluorine groups.

[0060] The first and second alignment films **110** and **120** contain not only the polymerization product **pob** but also the polyimide **pi**, and thus, thermal resistance, solvent resistance, hygroscopicity, and other characteristics of the first and second alignment films **110** and **120** are substantially non-inferior to those of a commonly-used alignment film formed only of polyimide. Moreover, the first and second alignment films **110** and **120** contain the polymerization product **pob**, whereby the first and second alignment films **110** and **120** are structurally stabilized.

[0061] As described above, the alignment sustaining layers **210** and **230** contain the polymerization product **poa**. The polymerization product **poa** is obtained by polymerizing the polyfunctional monomer ma, and the polyfunctional monomer ma has a plurality of vinyl groups. For example, the polyfunctional monomer ma is represented by general formula (2) Pa1-Aal-(Za1-Aa2)*n*-Pa2. In general formula (2), Pa1 and Pa2 are, independently, acrylate, methacrylate, acrylamide, or methacrylamide; Aa1 and Aa2 represent, independently, 1,4-phenylene, 1,4-cyclohexane or 2,5-thiophene, or naphthalene-(2,6)-diyl or anthracene-(2,7)-diyl; and Za1 is a -COO-group, a -OCO- group, a -O- group, a -CONH- group or a single bond, where *n* is 0 or 1.

[0062] Specifically, the polyfunctional monomer ma is represented by structural formula (2a).

[Formula 11]



[0063] Moreover, at least one of Aa1 and Aa2 may be substituted by at least one fluorine group. In the liquid crystal display device **100**, the polyimide **pi** and the polymerization product **pob** are present in the first alignment film **110**; however, in the first alignment sustaining layer **210**, the polymerization product **poa** is present, but not the polyimide **pi**. Similarly, both of the polyimide **pi** and the polymerization product **pob** are present in the second alignment film **120**; however, in the second alignment sustaining layer **230**, the polymerization product **poa** is present, but not the polyimide **pi**.

[0064] In the liquid crystal display device **100** of the present embodiment, not only the upper polymerization product **poa** but also the lower polymerization product **pob** exists at the interface of the liquid crystal layer **260**. Therefore, changes in the alignment function are sufficiently suppressed, and the pretilt angle of the liquid crystal molecules **262** in the liquid crystal layer **260** is maintained. When the monomer is a monofunctional monomer, the resultant polymerization product, i.e., a long linear polymer, is liable to deformation, and therefore changes in the alignment function cannot be sufficiently suppressed. On the other hand, when the monomer is a polyfunctional monomer, its polymerization product can sufficiently suppress changes in the alignment function.

[0065] The respective concentrations of the upper polymerization product **poa** and the lower polymerization product **pob** can be measured through a chemical analysis, such as nuclear magnetic resonance (NMR), Fourier transform infrared spectrometry (FT-IR), or mass spectrometry (MS). For example, the respective concentrations of the upper polymerization product **poa** and the lower polymerization product **pob** are measured by time of flight-secondary ion mass spectrometry (Time Of Flight-Secondary Ion Mass Spectrometry: TOF-SIMS). Alternatively, the concentrations of the upper polymerization product **poa** and the lower polymerization product **pob** are also measurable by X-ray photoelectron spectroscopy (X-ray Photoelectron Spectroscopy: XPS). In the case of XPS, for example, an apparatus manufactured by ULVAC-PHI, INCORPORATED may be used to analyze the atoms in the depth direction while etching with C60.

[0066] In the liquid crystal display device **100** of the present embodiment, as described above, the polymerization products **poa** and **pob** are present at the interface of the liquid crystal layer **260**. As a result of this, the pretilt direction of the liquid crystal molecules **262** is fixed. This is presumably because the polymerization products **poa** and **pob** suppress deformation of the pretilt-angle-exhibiting component, thus allowing the alignment direction of the liquid crystal molecules **262** to be maintained.

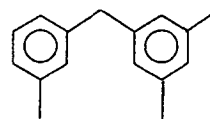
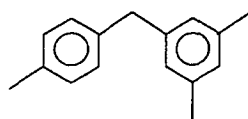
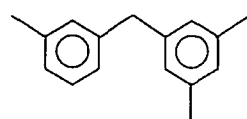
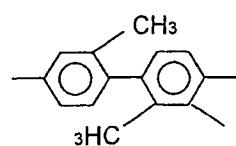
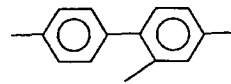
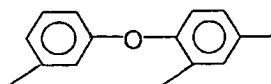
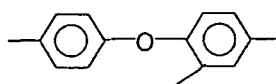
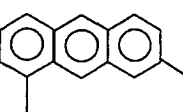
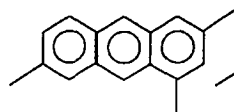
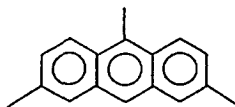
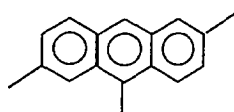
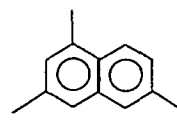
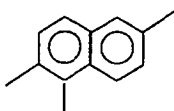
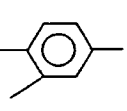
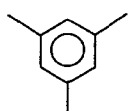
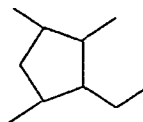
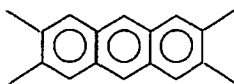
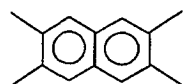
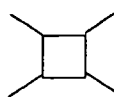
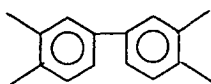
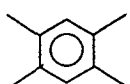
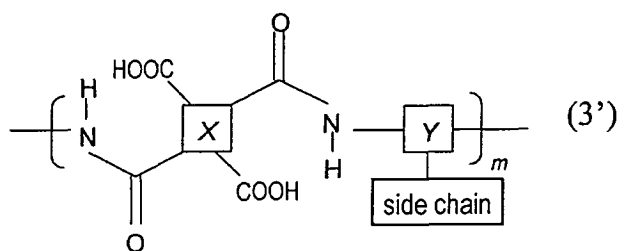
[0067] Moreover, since the first and second alignment films **110** and **120** contain the polymerization product **pob**, impurities and the like which occurred due to damage during the alignment treatment are stabilized, whereby occurrence of impurity ions is suppressed. As a result of this, the residual DC voltage and voltage holding ratio are improved, as a result of which the occurrence of image sticking is suppressed.

[0068] Hereinafter, with reference to FIG. 2, a method of producing the liquid crystal display device **100** will be described.

[0069] First, as shown in FIG. 2(a), the pixel electrodes **224** are formed on the first insulative substrate **222**. Although not shown in FIG. 2(a), TFTs and wiring lines and the like that are connected thereto are provided between the first insulative substrate **222** and the pixel electrodes **224**. Next, the first alignment film **110** covering the pixel electrodes **224** is formed.

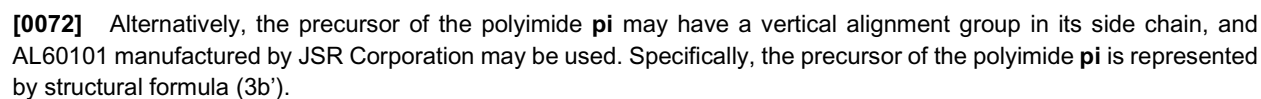
[0070] Formation of the first alignment film **110** is performed as follows. First, the alignment film material is obtained (as a mixture) by allowing the precursor of the polyimide **pi** and the polyfunctional monomer **mb** to be dissolved in a solvent. For example, the precursor (polyamic acid) of the polyimide **pi** includes a structure represented by general formula (3').

[Formula 12]

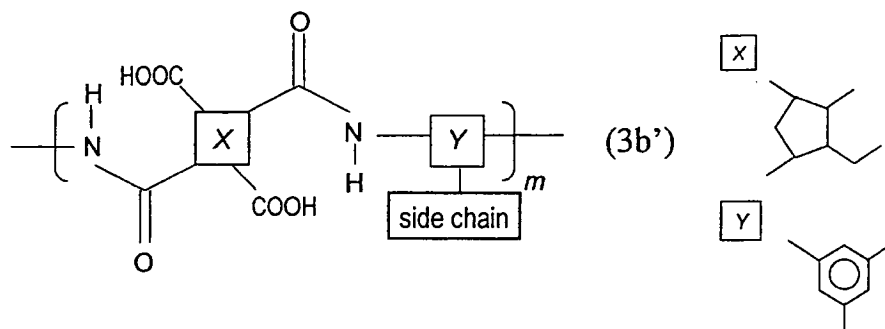


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[0071] The precursor of the polyimide **pi** may have a cinnamate group as the photoreactive functional group. Moreover, this side chain may contain a fluorine atom. When the side chain contains a fluorine atom, the aforementioned image sticking is suppressed to a certain extent. Specifically, the precursor of the polyimide **pi** includes a structure represented by structural formula (3a').



[Formula 14]



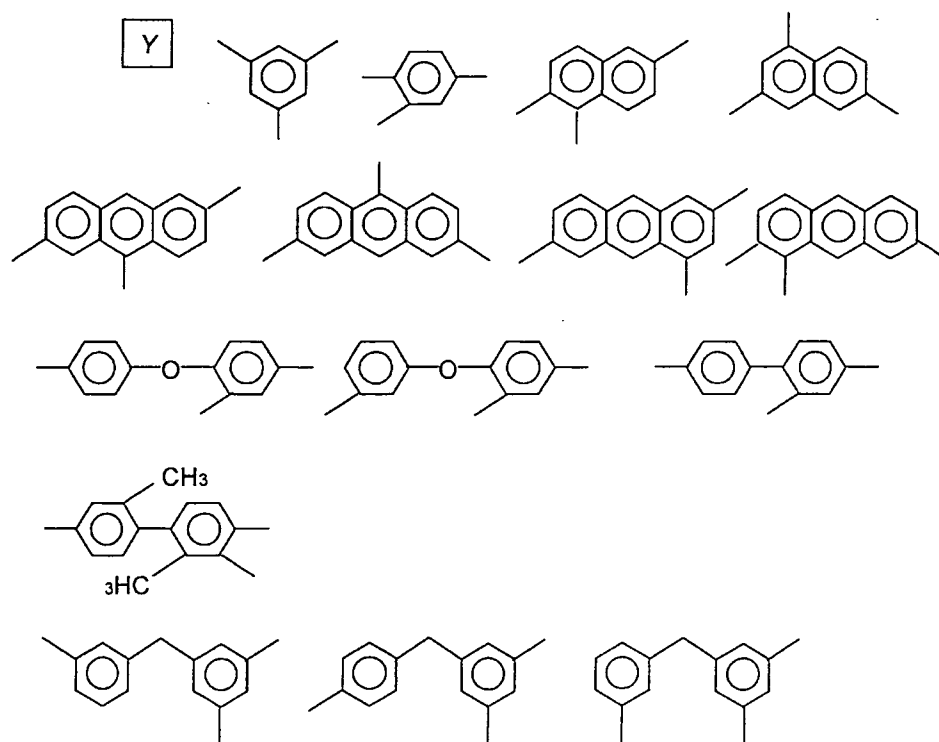
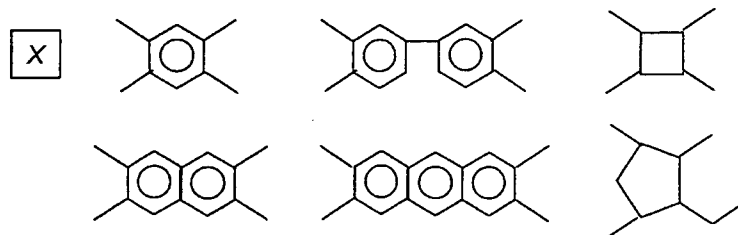
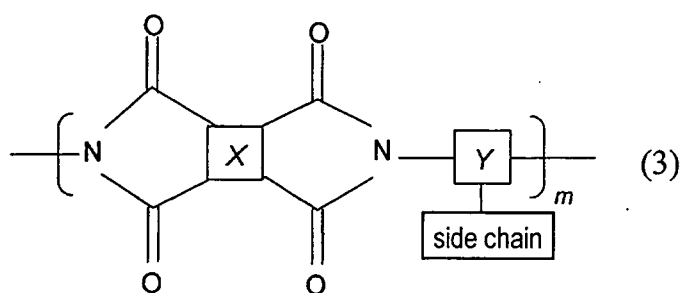
[0073] In the present embodiment, as described earlier, the alignment film material contains the polyfunctional monomer mb. The polyfunctional monomer mb may have two or more directly-bonded ring structures or one or more condensed ring structures, for example. For example, a methacrylate-type monomer or an acrylate-type monomer is used as the polyfunctional monomer mb. Specifically, the polyfunctional monomer mb is represented by structural formula (1a).

CC(=C)C(=O)Oc1ccc(cc1)-c2ccc(cc2)Oc3C(=O)C(=C)C (1a)

[0075] Next, the alignment film material is applied on the pixel electrodes **224**. Application of the alignment film material is performed by a printing technique, an ink jet technique, or a spin coating technique.

[0076] Next, a heat treatment is conducted. As the heat treatment, for example, two heat treatments may be performed at different temperatures. Specifically, after performing a first heat treatment, a second heat treatment is performed at a higher temperature than that of the first heat treatment. The first heat treatment removes most of the solvent. In the following description, what remains after the solvent has been substantially removed will be referred to as an alignment film. Through the subsequent second heat treatment, imidization progresses, whereby the alignment film is stabilized. The first heat treatment is also referred to as a preliminary bake or a pre-bake, and the second heat treatment is also referred to as a full bake or a post-bake. Through the heat treatment, the polyamic acid is imidized, whereby the polyimide **pi** is formed. The polyimide **pi** includes a structure represented by general formula (3), for example.

[Formula 16]



[0077] Note that the polyamic acid does not need to be completely imidized, and a portion thereof may still be polyamic acid. Moreover, through the heat treatment, the polyfunctional monomer **mb** is polymerized to form the polymerization product **pub**. The polyfunctional monomer **mb** polymerizes at a temperature which is equal to or less than that of the post-bake. The polymerization product **pub** exists also on the surface of the first alignment film **110**. In this manner, the first alignment film **110** is formed.

[0078] Next, the first alignment film **110** is subjected to an alignment treatment. The alignment treatment may be performed after the first heat treatment (pre-bake), or after the second heat treatment (post-bake), in the formation of the first alignment film **110**. For example, the alignment treatment is performed by irradiating the first alignment film **110** with light. For example, light of wavelengths in the range of no less than 250 nm and no more than 400 nm is radiated

onto the first alignment film **110** at an irradiation dose of no less than 20 mJ/cm² and no more than 200 mJ/cm², from a direction which is inclined from the normal direction of the principal face of the first alignment film **110**. If the irradiation dose increases from 200 mJ/cm², the alignment film may deteriorate so that the voltage holding ratio and the like may be lowered. Moreover, the irradiation angle of light may be in the range of no less than 5° and no more than 85°, and preferably no less than 40° and no more than 60°, from the normal direction of the principal face of the first alignment film **110**. Note that, when the irradiation angle is small, it becomes difficult to confer a pretilt angle; when the irradiation angle is large, it takes more time to confer the same pretilt angle. Moreover, light may be unpolarized light, linearly polarized light, elliptically polarized light, or circularly polarized light. However, linearly polarized light is to be used in the case where a cinnamate group is used as the photoreactive functional group. Alternatively, as the alignment treatment, the first alignment film **110** may be subjected to a rubbing treatment or an ion beam irradiation.

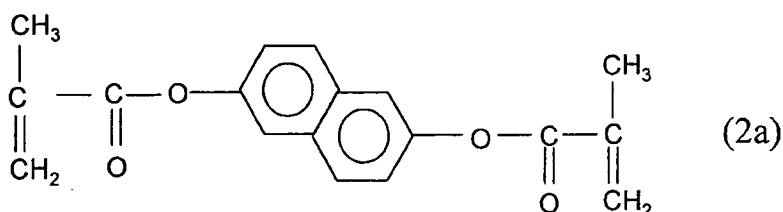
[0079] As shown in FIG. 2(b), the counter electrode **244** is formed on the second insulative substrate **242**. Next, the second alignment film **120** is formed on the counter electrode **244**.

[0080] Formation of the second alignment film **120** is performed as follows. First, an alignment film material is provided. This alignment film material may be similar to that of the first alignment film **110**. The alignment film material is applied, and a heat treatment is performed. The heat treatment removes most of the solvent. Through the subsequent second heat treatment, imidization progresses, whereby the alignment film is stabilized. The polyamic acid is imidized to form the polyimide **pi**. Moreover, through the heat treatment, the polyfunctional monomer **mb** is polymerized to form the polymerization product **pob**, such that the polymerization product **pob** exists also on the surface of the second alignment film **120**. Thereafter, the second alignment film **120** is subjected to an alignment treatment. The alignment treatment is performed in a similar manner to the first alignment film **110**.

[0081] Next, as shown in FIG. 2(c), the active matrix substrate **220** and the counter substrate **240** are attached together so that the first alignment film **110** and the second alignment film **120** oppose each other. In the present specification, the active matrix substrate and the counter substrate being attached together before formation of the liquid crystal layer may also be referred to as a "vacant panel".

[0082] Next, a liquid crystal material is provided. The polyfunctional monomer **ma** is mixed in the liquid crystal material. Specifically, the polyfunctional monomer **ma** is represented by structural formula (2a), for example.

[Formula 17]



[0083] The liquid crystal material is introduced between the first alignment film **110** and the second alignment film **120** of the vacant panel. As described above, the first and second alignment films **110** and **120** have been subjected to an alignment treatment, and thus the liquid crystal molecules **262** are aligned so as to be inclined from the normal directions of the principal faces of the first and second alignment films **110** and **120** even in the absence of an applied voltage. Moreover, the polymerization product **pob** has been formed, and even this alone can suppress image sticking caused by changes in the pretilt angle to a certain extent. In order to stabilize the alignment of the liquid crystal molecules **262**, after injecting the liquid crystal material, the liquid crystal panel may be heated for a certain time and thereafter quenched.

[0084] Thereafter, the liquid crystal panel is irradiated with light. The light irradiation may be performed while applying a voltage between the pixel electrodes **224** and the counter electrode **244**, or without applying a voltage. In the case where a voltage is applied when performing light irradiation, a large change in the pretilt angle of the liquid crystal molecules **262** can be attained.

[0085] In the light irradiation, a black light or fluorescent lamp is used as a light source, for example. Through the light irradiation, the polyfunctional monomer **ma** mixed in the liquid crystal material is polymerized to form the polymerization product **poa**. The polymerization product **poa** becomes phase-separated from the liquid crystal layer **260**, whereby the first and second alignment sustaining layers **210** and **230** containing the polymerization product **poa** are formed.

[0086] If any unreacted monomer remains in the liquid crystal material, the voltage holding ratio (Voltage Holding Ratio: VHR) will be lowered. Therefore, in order to reduce the residual monomer, ultraviolet light may be radiated for a

long time. For example, black light irradiation is continued until the peak that is attributable to the polyfunctional monomer **ma** is no longer observable by gas chromatography.

[0087] In this manner, a liquid crystal panel is formed. Thereafter, a driving circuit and a control circuit not shown are mounted on the liquid crystal panel, whereby the liquid crystal display device **100** is fabricated.

[0088] As the polyfunctional monomer **mb** in the alignment film material increases, the exposed amount of carboxylic acid in the polyamic acid at the surface of the first and second alignment films **110** and **120** decreases, and the concentration of the lower polymerization product **pob** increases. Since such a lower polymerization product **pob** undergoes a radical reaction with the polyfunctional monomer **ma** in the liquid crystal material as well as the upper polymerization product **poa** before phase separation, the polyfunctional monomer **ma** and the low-molecular weight upper polymerization product **poa** existing in the liquid crystal material will rapidly decrease, and the phase separation of the first and second alignment sustaining layers **210** and **230** containing the upper polymerization product **poa** from the liquid crystal layer **260** will be promoted.

[0089] In the liquid crystal display device **100** of the present embodiment, the residual DC voltage and voltage holding ratio are improved because the first and second alignment films **110** and **120** contain the polymerization product **pob**. The reason thereof will be described below.

[0090] In the case where the first and second alignment films of the liquid crystal display device do not contain the polymerization product, if light is radiated for polymerization of the polyfunctional monomer **ma**, a portion of the alignment film may dissolve into the liquid crystal layer, thus deteriorating the residual DC voltage and voltage holding ratio. For example, if the side chain of the polyimide **pi** represented by structural formula (3a) is irradiated with light as shown in FIG. 3(a), a portion thereof is separated to become an ionized site as shown in FIG. 3(b), and dissolves into the liquid crystal layer as shown in FIG. 3(c). On the other hand, in the liquid crystal display device **100** of the present embodiment, as shown in FIG. 3(d), the dissolution of the ionized site into the liquid crystal layer **260** is suppressed because the polymerization product **pob** exists in the first and second alignment films **110** and **120**.

[0091] In the case where the alignment film material contains a cross-linker, if light is radiated for polymerization of the polyfunctional monomer **ma**, as shown in FIG. 4(a), the unreacted cross-linker will dissolve into the liquid crystal layer. On the other hand, in the liquid crystal display device **100** of the present embodiment, as shown in FIG. 4(b), the dissolution of the cross-linker into the liquid crystal layer **260** is suppressed because the polymerization product **pob** exists in the first and second alignment films **110** and **120**.

[0092] In the aforementioned PSA technique, a polymerization product is formed under an applied voltage. In the case where ultraviolet light for the purpose of polymerization product formation is radiated while thus applying a voltage, a complex fabrication apparatus is required in which a device for applying a voltage across the liquid crystal panel and a device for radiating ultraviolet light are integrated. Moreover, since ultraviolet light irradiation is performed after a voltage is applied across the liquid crystal panel for a long time for obtaining a predetermined alignment, this fabrication apparatus needs to be used for a long time. Moreover, when forming the liquid crystal layer of a liquid crystal panel through dropwise application of a liquid crystal material, generally speaking, a plurality of liquid crystal panels are simultaneously produced by using a large-sized mother glass substrate, and thereafter each liquid crystal panel is cut out from the large-sized mother glass substrate. In this case of simultaneously producing a plurality of liquid crystal panels, a design must be adopted such that special wiring lines are formed on the mother glass substrate for allowing a voltage to be simultaneously applied to the plurality of liquid crystal panels.

[0093] Moreover, in the case where a liquid crystal panel of a particularly large size is to be produced, it is difficult to uniformly apply a voltage across the liquid crystal layer in the respective pixels. If ultraviolet light irradiation is performed with non-uniform voltages being applied, there will be fluctuations in the pretilt angle. Moreover, in the case of applying a voltage during polymerization product formation, ribs, slits, or rivets need to be provided on the pixel electrode and the counter electrode for improved viewing angle characteristics. This will result in an increased number of steps and a decrease in the effective aperture ratio.

[0094] On the other hand, according to the production method of the present embodiment, it is not necessary to apply a voltage when forming the polymerization products **poa** and **pob**. Therefore, the liquid crystal display device **100** can be easily produced without using a complex fabrication apparatus. Moreover, a liquid crystal panel can be easily produced even when producing the liquid crystal layer **260** through dropwise application of a liquid crystal material. Moreover, since it is not necessary to apply a voltage across the liquid crystal layer **260** of all pixels when forming the polymerization products **poa** and **pob**, fluctuations in the pretilt angle among liquid crystal molecules **262** can be suppressed. Furthermore, the transmittance, contrast ratio, and viewing angle can be improved without providing ribs, slits, or rivets on the pixel electrodes **224** and the counter electrode **244**.

[0095] Note that slits, ribs, and/or rivets may be provided on the pixel electrodes **224** and the counter electrode **244**. Alternatively, slits, ribs, and/or rivets may not be provided on the pixel electrodes **224** and the counter electrode **244**, and the liquid crystal molecules **262** may be aligned in accordance with an oblique electric field which is created by a highly-symmetrical pixel electrode **226** and the counter electrode **246**. As a result, the alignment regulating force of the liquid crystal molecules **262** under an applied voltage can be further increased.

[0096] Moreover, in order to increase the tilt of the liquid crystal molecules **262** from the normal directions of the principal faces of the first and second alignment films **110** and **120** in the absence of an applied voltage, a voltage may be applied between the pixel electrodes **224** and the counter electrode **244** during the light irradiation for forming the first and second alignment sustaining layers **210** and **230**.

[0097] Although the above description illustrates that the first and second alignment films **110** and **120** are made of the same alignment film material, and the characteristics of the second alignment film **120** are similar to those of the first alignment film **110**, the present invention is not limited thereto. The characteristics of the second alignment film **120** may be different from those of the first alignment film **110**, and the first and second alignment films **110** and **120** may be made of different alignment film materials. For example, at least one of the polyimide **pi** and the polymerization product **pob** of the first alignment film **110** may be different from at least one of the polyimide **pi** and the polymerization product **pob** of the second alignment film **120**.

[0098] Although the above description illustrates that the first and second alignment films **110** and **120** each contain the polymerization product **pob**, the present invention is not limited thereto. Only one of the first and second alignment films **110** and **120** may contain the polymerization product **pob**. However, if the second alignment film **120** differs in characteristics from the first alignment film **110**, the impurities remaining in the first and second alignment films **110** and **120** will tend to be asymmetric. Therefore, it is preferable that the characteristics of the second alignment film **120** are similar to those of the first alignment film **110**.

[0099] Although the above description illustrates that the active matrix substrate **220** and the counter substrate **240** respectively include the first and second alignment films **110** and **120**, the present invention is not limited thereto. Only one of the active matrix substrate **220** and the counter substrate **240** may include the first or second alignment film **110** or **120**.

[0100] Although the above description illustrates that polymerization product **pob** is formed through a heat treatment, the present invention is not limited thereto. The polymerization product **pob** may be formed through light irradiation. For example, in such light irradiation, a light source which mainly emits ultraviolet light (i-line) with a wavelength of 365 nm is suitably used. The irradiation time is about 500 seconds, for example, and the irradiation intensity of light is about 20 mW/cm². In the case where polymerization is effected through light irradiation, the polyfunctional monomer **mb** will sufficiently polymerize even if the irradiation intensity of light is 10 mW/cm² or less. The wavelength of light is preferably in the range of no less than 250 nm and no more than 400 nm, and more preferably in the range of no less than 300 nm and no more than 400 nm. However, polymerization will sufficiently occur with light of a wavelength greater than 400 nm. Although polymerization can occur with light of a wavelength of 300 nm or less, the irradiation dose should preferably be as small as possible because decomposition of organic matter will occur with irradiation of deep-ultraviolet with wavelengths near 200 nm.

[0101] Moreover, the liquid crystal display device **100** may be of the 4D-RTN (4 Domain-Reverse Twisted Nematic) mode in which four-split alignment is realized. Hereinafter, a liquid crystal display device of the 4D-RTN mode will be described with reference to FIG. 5 and FIG. 6.

[0102] FIG. 5(a) shows pretilt directions **PA1** and **PA2** of liquid crystal molecules as regulated on the first alignment film **110** of the active matrix substrate **220**, and FIG. 5(b) shows pretilt directions **PB1** and **PB2** of liquid crystal molecules as regulated on the second alignment film **120** of the counter substrate **240**. FIG. 5(c) shows alignment directions of liquid crystal molecules in the centers of liquid crystal domains **A** to **D** under an applied voltage, and regions (domain lines) **DL1** to **DL4** appearing dark due to alignment disorder. Note that the domain lines **DL1** to **DL4** are not so-called disclination lines.

[0103] FIG. 5(a) to FIG. 5(c) schematically show alignment directions of liquid crystal molecules as seen from the viewer side. FIG. 5(a) to FIG. 5(c) indicate that the end portions (essentially circular portions) of the cylindrical liquid crystal molecules are tilted toward the viewer.

[0104] As shown in FIG. 5(a), the first alignment film **110** includes a first alignment region **OR1** and a second alignment region **OR2**. The liquid crystal molecules regulated by the first alignment region **OR1** are tilted in the -y direction from the normal direction of the principal face of the first alignment film **110**, whereas the liquid crystal molecules regulated by the second alignment region **OR2** of the first alignment film **110** are tilted in the +y direction from the normal direction of the principal face of the first alignment film **110**. Moreover, the boundary between the first alignment region **OR1** and the second alignment region **OR2** extends in the column direction (y direction), and located in the substantial center along the row direction (x direction) of pixels. Thus, first and second alignment regions **OR1** and **OR2** of different pretilt azimuths are provided on the first alignment film **110**.

[0105] Moreover, as shown in FIG. 5(b), the second alignment film **120** includes a third alignment region **OR3** and a fourth alignment region **OR4**. The liquid crystal molecules regulated by the third alignment region **OR3** are tilted in the +x direction from the normal direction of the principal face of the second alignment film **120**, such that the -x direction end portions of these liquid crystal molecules are pointed toward the front face. The liquid crystal molecules regulated by the fourth alignment region **OR4** of the second alignment film **120** are tilted in the -x direction from the normal direction of the principal face of the second alignment film **120**, such that the +x direction end portions of these liquid crystal

molecules are pointed toward the front face. Thus, the second alignment film **120** includes third and fourth alignment regions **OR3** and **OR4** with different pretilt azimuths.

[0106] An alignment treatment direction corresponds to an azimuth angle component obtained by projecting a direction, which extends toward an alignment region along the major axes of the liquid crystal molecules, onto that alignment region. The alignment treatment directions of the first, second, third, and fourth alignment regions are also referred to as first, second, third, and fourth alignment treatment directions.

[0107] The first alignment region **OR1** of the first alignment film **110** has been subjected to an alignment treatment along a first alignment treatment direction **PD1**, whereas the second alignment region **OR2** has been subjected to an alignment treatment along a second alignment treatment direction **PD2** which is different from the first alignment treatment direction **PD1**. The first alignment treatment direction **PD1** is essentially antiparallel to the second alignment treatment direction **PD2**. Moreover, the third alignment region **OR3** of the second alignment film **120** has been subjected to an alignment treatment along a third alignment treatment direction **PD3**, whereas the fourth alignment region **OR4** has been subjected to an alignment treatment along a fourth alignment treatment direction **PD4** which is different from the third alignment treatment direction **PD3**. The third alignment treatment direction **PD3** is essentially antiparallel to the fourth alignment treatment direction **PD4**.

[0108] As shown in FIG. 5(c), four liquid crystal domains **A**, **B**, **C**, and **D** are formed in the liquid crystal layer of a pixel. In the liquid crystal layer **260**, a portion interposed between the first alignment region **OR1** of the first alignment film **110** and the third alignment region **OR3** of the second alignment film **120** defines the liquid crystal domain **A**; a portion interposed between the first alignment region **OR1** of the first alignment film **110** and the fourth alignment region **OR4** of the second alignment film **120** defines a liquid crystal domain **B**; a portion interposed between the second alignment region **OR2** of the first alignment film **110** and the fourth alignment region **OR4** of the second alignment film **120** defines a liquid crystal domain **C**; and a portion interposed between the second alignment region **OR2** of the first alignment film **110** and the third alignment region **OR3** of the second alignment film **120** defines a liquid crystal domain **D**. Note that the angle constituted by the first or second alignment treatment direction **PD1** or **PD2** and the third or fourth alignment treatment direction **PD3** or **PD4** is essentially 90°, and the twist angle in each of the liquid crystal domains **A**, **B**, **C**, and **D** is essentially 90°.

[0109] FIG. 6 shows the alignment directions of the liquid crystal molecules **262** in the liquid crystal domain **A**. The angle between the alignment treatment direction **PD1** of the first alignment film **110** and the alignment treatment direction **PD3** of the second alignment film **120** is 90°, and the liquid crystal molecules **262** have a twist angle of 90°. Although not shown in the figure, the liquid crystal molecules **262** in the other liquid crystal domains **B**, **C**, and **D** are also similarly twisted.

[0110] FIG. 5(c) is referred to again. The alignment direction of a liquid crystal molecule at the center of a liquid crystal domain **A** to **D** is an intermediate direction between the pretilt direction for liquid crystal molecules introduced by the first alignment film **110** and the pretilt direction for liquid crystal molecules introduced by the second alignment film **120**. In the present specification, the alignment direction of a liquid crystal molecule in the center of a liquid crystal domain is referred to as a reference alignment direction; and within the reference alignment direction, an azimuth angle component in a direction from the rear face toward the front face and along the major axis of the liquid crystal molecule (i.e., an azimuth angle component obtained by projecting the reference alignment direction onto the principal face of the first alignment film **110** or the second alignment film **120**) is referred to as a reference alignment azimuth. The reference alignment azimuth characterizes its corresponding liquid crystal domain, and exerts a predominant influence on the viewing angle characteristics of that liquid crystal domain. Now, by relying on the horizontal direction (right-left direction) of the display screen (plane of the figure) as a reference for the azimuthal direction, and defining the left turn as positive (i.e., if the display surface is compared to the face of a clock, counterclockwise is positive, the 3 o'clock direction being an azimuth angle of 0°), the reference alignment directions of the four liquid crystal domains **A** to **D** are set to be four directions such that the difference between any two directions is substantially equal to an integer multiple of 90°. Specifically, the reference alignment azimuths of the liquid crystal domains **A**, **B**, **C**, and **D** are, respectively, 225°, 315°, 45°, and 135°.

[0111] As shown in FIG. 5(c), the domain lines **DL1** to **DL4** are respectively formed in the liquid crystal domains **A**, **B**, **C**, and **D**. The domain line **DL1** is formed in parallel to a portion of an edge **EG1** of the pixel electrode **224**, whereas the domain line **DL2** is formed in parallel to a portion of an edge **EG2**. Moreover, the domain line **DL3** is formed in parallel to a portion of an edge **EG3** of the pixel electrode **224**, whereas the domain line **DL4** is formed in parallel to a portion of an edge **EG4**. Moreover, a disclination line **CL** indicated by a broken line is observed in a border region where each of the liquid crystal domains **A** to **D** adjoins another liquid crystal domain. The disclination lines **CL** are dark lines in the aforementioned central portion. The disclination lines **CL** and the domain lines **DL1** to **DL4** are continuous, thus resulting

in dark lines of a reverse  shape. Although the dark lines herein are in a reverse  shape, the dark lines may be in an 8 shape.

[0112] Although the above-described liquid crystal display device realizes four-split alignment, the present invention is not limited thereto. Two-split alignment may be realized in the liquid crystal display device **100**. FIG. 7 shows alignment treatment directions for the first and second alignment films **110** and **120**. As described above, the active matrix substrate **220** and the counter substrate **240** are attached together so that the angles between the alignment treatment directions **PD1** and **PD2** of the first alignment film **110** and the alignment treatment direction **PD3** of the second alignment film **120** are 90°, and the liquid crystal molecules **262** have a twist angle of 90°.

[0113] Alternatively, the liquid crystal display device **100** does not need to be split in terms of alignment. FIG. 8 shows alignment treatment directions for the first and second alignment films **110** and **120**. As described above, the active matrix substrate **220** and the counter substrate **240** are attached together so that the angle between the alignment treatment direction **PD1** of the first alignment film **110** and the alignment treatment direction **PD3** of the second alignment film **120** is 90°, and the liquid crystal molecules **262** have a twist angle of 90°.

[0114] Although the above-described liquid crystal display device is of the RTN mode, the present invention is not limited thereto. The liquid crystal display device may be of the CPA mode.

[0115] Hereinafter, liquid crystal display devices of the present Examples will be described.

[Example 1]

[0116] Hereinafter, with reference to FIG. 1, FIG. 6, and FIG. 8, a liquid crystal display device of Example 1 will be described. The liquid crystal display device of Example 1 also operates in the RTN mode.

[0117] First, on a principal face of the first insulative substrate **222**, although not shown in the figures, TFTs and wiring lines connected to the TFTs, and an insulating layer and the like were formed, upon which the pixel electrodes **224** were formed. Similarly, on a principal face of the second insulative substrate **242**, although not shown in the figures, a colored layer having color filters, and an insulating layer and the like were formed, upon which the counter electrode **244** was formed.

[0118] Next, an alignment film material was prepared. The alignment film material was obtained by allowing the precursor (polyamic acid) of the polyimide **pi** represented by structural formula (3a') to be dissolved in a solvent, and thereafter allowing the polyfunctional monomer **mb** to be further dissolved therein. The precursor of the polyimide **pi** had a cinnamate group in its side chain. The polyfunctional monomer **mb** was biphenyldimethacrylate as represented by structural formula (1a). The concentration of the polyfunctional monomer **mb** on the basis of the alignment film material was 10wt%.

[0119] First, the alignment film material was applied on the pixel electrodes **224**. The application of the alignment film material was performed by a printing technique, an ink jet technique, or a spin coating technique. As a first heat treatment (pre-bake), one minute of heating was conducted at 90°C to remove the solvent to a certain extent; and furthermore, 40 minutes of heating was conducted at 200°C as a second heat treatment (post-bake). Through such heat treatments, the first alignment film **110** containing the polyimide **pi**, into which the polyamic acid had imidized, and the polymerization product **pob**, into which the polyfunctional monomer **mb** had polymerized, was formed. The fraction of imide units of the polyimide **pi** was about 50%. Thereafter, obliquely from a 40° direction with respect to the normal direction of a principal face of the first alignment film **110**, P-polarized light with a peak wavelength of 330 nm was radiated at 50 mJ/cm², thus performing a photo-alignment treatment. Similarly, the aforementioned alignment film material was applied to form the second alignment film **120** on the counter electrode **244**, and a photo-alignment treatment was performed.

[0120] Next, the active matrix substrate **220** and the counter substrate **240** were attached together so that the first alignment film **110** and the second alignment film **120** opposed each other and that the angle between the alignment treatment direction of the first alignment film **110** and the alignment treatment direction of the second alignment film was 90°, and fixed so that the interspace between the active matrix substrate **220** and the counter substrate **240** was about 4 μm. Specifically, after applying a seal onto the active matrix substrate **220** and scattering beads over the counter substrate **240**, the active matrix substrate **220** and the counter substrate **240** were attached together.

[0121] Next, the polyfunctional monomer **ma** represented by structural formula (2a) was mixed in a nematic liquid crystal material having negative dielectric anisotropy. The concentration of the polyfunctional monomer **ma** on the basis of the liquid crystal material was 0.6wt%. Then, the liquid crystal material was injected between the active matrix substrate **220** and the counter substrate **240**, and a heating-quenching was performed at 130°C.

[0122] Next, black light irradiation was performed to form a polymerization product of the polyfunctional monomer **ma**. If any polyfunctional monomer **ma** remains in the liquid crystal material, the voltage holding ratio and residual DC voltage will be deteriorated. Therefore, the black light irradiation was continued until the peak attributable to the polyfunctional monomer **ma** was no longer observable by gas chromatography.

[0123] FIG. 6 shows an alignment state of the liquid crystal molecules **262** in the liquid crystal display device of Example 1. As shown in FIG. 8, the angle between the active matrix substrate **220** and the counter substrate **240** were attached together so that the alignment treatment direction **PD1** of the first alignment film **110** and the alignment treatment direction **PD3** of the second alignment film **120** was 90°, and the liquid crystal molecules **262** had a twist angle of 90°. Herein,

the polarization axis of the polarizer on the active matrix substrate **220** was parallel to the alignment treatment direction **PD1** of the first alignment film **110**, and the polarization axis of the polarizer on the counter substrate **240** was parallel to the alignment treatment direction **PD3** of the second alignment film **120**. In this manner, the liquid crystal panel of Example 1 was produced.

[0124] Moreover, three liquid crystal panels were produced for the sake of comparison. In producing a liquid crystal panel of Comparative Example 1-1, the polyfunctional monomer **ma** was not mixed in the liquid crystal material, and the polyfunctional monomer **mb** was not mixed in the alignment film material. As a result, the liquid crystal panel of Comparative Example 1-1 did not have any alignment sustaining layers, and its alignment films did not contain the polymerization product. In the liquid crystal panel of Comparative Example 1-1, since the monomer within the liquid crystal material was not subjected to polymerization, black light irradiation was not performed.

[0125] In producing a liquid crystal panel of Comparative Example 1-2, the polyfunctional monomer **ma** represented by structural formula (2a) was mixed in the liquid crystal material and a photopolymerization was carried out, but the polyfunctional monomer **mb** was not mixed in the alignment film material. As a result, the alignment films of the liquid crystal panel of Comparative Example 1-2 did not contain the polymerization product. Since polymerization of the monomer within the liquid crystal material was effected in the liquid crystal panel of Comparative Example 1-2, black light irradiation was performed. When black light irradiation was continued until the peak attributable to the polyfunctional monomer **ma** was no longer observable as described above, the irradiation time of black light had come to 30 minutes for the liquid crystal panel of Example 1, whereas the irradiation time was 2 hours for the liquid crystal panel of Comparative Example 1-2.

[0126] In producing the liquid crystal panel of Comparative Example 1-3, the polyfunctional monomer **ma** was not mixed in the liquid crystal material, but the polyfunctional monomer **mb** represented by structural formula (1a) was mixed in the alignment film material, and a thermal polymerization was carried out. As a result, the liquid crystal panel of Comparative Example 1-3 did not have any alignment sustaining layers, but the alignment films contained the polymerization product.

[0127] Next, a voltage of 1 V was applied to each of the liquid crystal panels of Comparative Examples 1-1, 1-2, 1-3, and Example 1 at 70°C, and their initial VHRs were measured. The initial VHR was 99% for the liquid crystal panel of Comparative Examples 1-1, 1-3, and Example 1, whereas the initial VHR was 98% for the liquid crystal panel of Comparative Example 1-2. Since it is considered that an initial VHR of 99% or more is necessary for ensuring long-term reliability, the liquid crystal panel of Comparative Example 1-2 did not provide a satisfactory value.

[0128] Moreover, the residual DC voltage was measured by flicker minimizing method. After applying a DC offset voltage of 2 V at 40°C, a residual DC voltage measurement was taken. The liquid crystal panel of Comparative Example 1-1 had a residual DC voltage of +150 mV; the liquid crystal panel of Comparative Example 1-2 had a residual DC voltage of +90 mV; and the liquid crystal panels of Comparative Example 1-3 and Example 1 had a residual DC voltage of -20 mV. Although the liquid crystal panel of Comparative Example 1-1 had a relatively good initial VHR, its residual DC voltage was very high. In the liquid crystal panel of Comparative Example 1-2, it is presumable from its initial VHR and residual DC voltage results that, since the alignment films did not contain the polymerization product, a portion of the alignment films dissolved into the liquid crystal layer as impurities during light irradiation.

[0129] Moreover, the resultant liquid crystal panels of Comparative Examples 1-1, 1-2, 1-3, and Example 1 were subjected to a power-on test of continuously applying a voltage of ± 10 V with a frequency of 60 Hz at 40°C, and thereafter an amount of change in the tilt angle was measured.

[0130] FIG. 9 is a graph showing amounts of change in the tilt angle against powering time, with respect to the liquid crystal panels of Comparative Examples 1-1, 1-2, 1-3, and Example 1. Herein, a comparison between the results of the liquid crystal panels of Comparative Examples 1-1, 1-2, and 1-3 indicates that: in the liquid crystal panel of Comparative Example 1-1, the amount of change in the pretilt angle increased as the powering time increased; however, in the liquid crystal panels of Comparative Examples 1-2 and 1-3, the increase in the amount of change in the pretilt angle was suppressed even when the powering time increased. This is presumably because the pretilt angle of the liquid crystal molecules was maintained due to the upper polymerization product **poa**, into which the polyfunctional monomer **ma** mixed in the liquid crystal material had polymerized, or the lower polymerization product **pob**, into which the polyfunctional monomer **mb** mixed in the alignment film material had polymerized. Note that the effect provided by the upper polymerization product **poa** was greater than the effect provided by the lower polymerization product **pob**.

[0131] Furthermore, a comparison between the results of the liquid crystal panels of Comparative Examples 1-2, 1-3, and Example 1 indicates that the amount of change in the pretilt angle of the liquid crystal panel of Example 1, which contained both the upper polymerization product **poa** and the lower polymerization product **pob**, was slightly smaller than those of the liquid crystal panels of Comparative Examples 1-2 and 1-3. Thus, the amount of change in the pretilt angle was improved in the liquid crystal panel of Example 1.

[Example 2]

[0132] Hereinafter, with reference to FIG. 1, FIG. 6, and FIG. 8, a liquid crystal display device of Example 2 will be described. The liquid crystal display device of Example 2 also operates in the RTN mode.

[0133] First, on a principal face of the first insulative substrate **222**, although not shown in the figures, TFTs and wiring lines connected to the TFTs, and an insulating layer and the like were formed, upon which the pixel electrodes **224** were formed. Similarly, on a principal face of the second insulative substrate **242**, although not shown in the figures, a colored layer having color filters, and an insulating layer and the like were formed, upon which the counter electrode **244** was formed.

[0134] Next, an alignment film material was prepared. The alignment film material was obtained by allowing a precursor (polyamic acid) of the polyimide **pi** to be dissolved in a solvent, and thereafter allowing the polyfunctional monomer **mb** and a cross-linker to be further dissolved therein. The precursor of the polyimide **pi** was the polyamic acid (AL60101 manufactured by JSR Corporation) represented by structural formula (3b'). The polyfunctional monomer **mb** was biphenyldimethacrylate as represented by structural formula (1a). The cross-linker was an epoxy-type cross-linker represented by structural formula (4).

[0135] First, the alignment film material was applied on the pixel electrodes **224**. The application of the alignment film material was performed by a printing technique, ink jet technique, or spin coating technique. As a first heat treatment (pre-bake), one minute of heating was conducted at 90°C to remove the solvent to a certain extent; and furthermore, 40 minutes of heating was conducted at 200°C as a second heat treatment (post-bake). Through such heat treatments, the first alignment film **110** containing the polyimide **pi**, into which the polyamic acid had imidized, and the polymerization product **pob**, into which the polyfunctional monomer **mb** had polymerized, was formed. The fraction of imide units of the polyimide **pi** was about 50%. Thereafter, a principal face of the first alignment film **110** was subjected to a rubbing treatment. Similarly, the aforementioned alignment film material was applied to form the second alignment film **120** on the counter electrode **244**, and a rubbing alignment treatment was performed.

[0136] Next, the active matrix substrate **220** and the counter substrate **240** were attached together so that the first alignment film **110** and the second alignment film **120** opposed each other and that the angle between the alignment treatment direction **PD1** of the first alignment film and the alignment treatment direction **PD3** of the second alignment film was 90°, and fixed so that the interspace between the active matrix substrate **220** and the counter substrate **240** was about 4 μm. Specifically, after applying a seal onto the active matrix substrate **220** and scattering beads over the counter substrate **240**, the active matrix substrate **220** and the counter substrate **240** were attached together.

[0137] Next, the polyfunctional monomer **ma** represented by structural formula (2a) was mixed in a nematic liquid crystal material having negative dielectric anisotropy. The concentration of the polyfunctional monomer **ma** on the basis of the liquid crystal material was 0.6wt%. Then, the liquid crystal material was injected between the active matrix substrate **220** and the counter substrate **240**, and a heating-quenching was performed at 130°C.

[0138] Thereafter, black light irradiation was performed to form a polymerization product of the polyfunctional monomer **ma**. If any polyfunctional monomer **ma** remains in the liquid crystal material, the voltage holding ratio and residual DC voltage will be deteriorated. Therefore, the black light irradiation was continued until the peak attributable to the polyfunctional monomer **ma** was no longer observable by gas chromatography, thus reducing the polyfunctional monomer **ma** mixed in the liquid crystal material. In this manner, the liquid crystal panel of Example 2 was produced.

[0139] Moreover, three liquid crystal panels were produced for the sake of comparison. In producing a liquid crystal panel of Comparative Example 2-1, the polyfunctional monomer **ma** was not mixed in the liquid crystal material, and the polyfunctional monomer **mb** was not mixed in the alignment film material although a cross-linker was added thereto. As a result, the liquid crystal panel of Comparative Example 2-1 did not have any alignment sustaining layers, and its alignment films did not contain the polymerization product. In the liquid crystal panel of Comparative Example 2-1, since the monomer within the liquid crystal material was not subjected to polymerization, black light irradiation was not performed.

[0140] In producing a liquid crystal panel of Comparative Example 2-2, the polyfunctional monomer **ma** represented by structural formula (2a) was mixed in the liquid crystal material and a photopolymerization was carried out, but the polyfunctional monomer **mb** was not mixed in the alignment film material although a cross-linker was added thereto. As a result, the alignment films of the liquid crystal panel of Comparative Example 2-2 did not contain the polymerization product. Since polymerization of the monomer within the liquid crystal material was effected in the liquid crystal panel of Comparative Example 2-2, black light irradiation was performed. When black light irradiation was continued until the peak attributable to the polyfunctional monomer **ma** was no longer observable as described above, the irradiation time of black light had come to 30 minutes for the liquid crystal panel of Example 2, whereas the irradiation time was 2 hours for the liquid crystal panel of Comparative Example 2-2.

[0141] In producing the liquid crystal panel of Comparative Example 2-3, the polyfunctional monomer **ma** was not mixed in the liquid crystal material, but the polyfunctional monomer **mb** represented by structural formula (1a) and a cross-linker were mixed in the alignment film material, and a thermal polymerization was carried out. As a result, the

liquid crystal panel of Comparative Example 2-3 did not have any alignment sustaining layers, but the alignment films contained the polymerization product.

[0142] Next, a voltage of 1 V was applied to each of the liquid crystal panels of Comparative Examples 2-1, 2-2, 2-3, and Example 2 at 70°C, and their initial VHRs were measured. The initial VHR was 99% for the liquid crystal panels of Comparative Examples 2-1, 2-3, and Example 2, whereas the initial VHR was 97% for the liquid crystal panel of Comparative Example 2-2. Thus, the liquid crystal panel of Comparative Example 2-2 did not provide a satisfactory value.

[0143] Moreover, the residual DC voltage was measured by flicker minimizing method. After applying a DC offset voltage of 2 V at 40°C, a residual DC voltage measurement was taken. The liquid crystal panels of Comparative Examples 2-1, 2-3, and Example 2 had a residual DC voltage of -10 mV, whereas the liquid crystal panel of Comparative Example 2-2 had a residual DC voltage of a +60 mV. It is presumable in the liquid crystal panel of Comparative Example 2-1 that the residual DC voltage was relatively low because of the cross-linker within the alignment films. In the liquid crystal panel of Comparative Example 2-2, it is presumable from its initial VHR and residual DC voltage results that, since the alignment films did not contain the polymerization product, a portion of the alignment films dissolved into the liquid crystal layer as impurities during light irradiation.

[0144] Moreover, the resultant liquid crystal panels of Comparative Examples 2-1, 2-2, 2-3, and Example 2 were subjected to a power-on test of continuously applying a voltage of ± 10 V with a frequency of 60 Hz at 40°C, and thereafter an amount of change in the tilt angle was measured.

[0145] FIG. 10 is a graph showing amounts of change in the tilt angle against powering time, with respect to the liquid crystal panels of Comparative Examples 2-1, 2-2, 2-3, and Example 2. Herein, a comparison between the results of the liquid crystal panels of Comparative Examples 2-1, 2-2, and 2-3 indicates that: in the liquid crystal panel of Comparative Example 2-1, the amount of change in the pretilt angle increased as the powering time increased; however, in the liquid crystal panels of Comparative Examples 2-2 and 2-3, the increase in the amount of change in the pretilt angle was suppressed even when the powering time increased. This is presumably because the pretilt angle of the liquid crystal molecules was maintained due to the upper polymerization product **poa**, into which the polyfunctional monomer **ma** mixed in the liquid crystal material had polymerized, or the lower polymerization product **pob**, into which the polyfunctional monomer **mb** mixed in the alignment film material had polymerized. Note that the effect provided by the upper polymerization product **poa** was greater than the effect provided by the lower polymerization product **pob**.

[0146] Furthermore, a comparison between the results of the liquid crystal panels of Comparative Examples 2-2, 2-3, and Example 2 indicates that the amount of change in the pretilt angle of the liquid crystal panel of Example 2, which contained both the upper polymerization product **poa** and the lower polymerization product **pob**, was smaller than those of the liquid crystal panels of Comparative Examples 2-2 and 2-3. Thus, the amount of change in the pretilt angle was improved in the liquid crystal panel of Example 2.

[0147] For reference sake, the entire disclosure of Japanese Patent Application No. 2008-303176, on which the present application claims priority, is incorporated herein by reference.

INDUSTRIAL APPLICABILITY

[0148] A liquid crystal display device according to the present invention is able to suppress image sticking caused by changes in the pretilt angle. Moreover, a liquid crystal display device according to the present invention is suitably used not only for liquid crystal television sets, but also for DIDs (Digital Information Displays).

REFERENCE SIGNS LIST

[0149]

100 liquid crystal display device

110 first alignment film

pi polyimide

pob lower polymerization product

120 second alignment film

210 first alignment sustaining layer

poa upper polymerization product

220 active matrix substrate

222 first insulative substrate

5 224 pixel electrode

230 second alignment sustaining layer

10 240 counter substrate

242 second insulative substrate

244 counter electrode

15 260 liquid crystal layer

262 liquid crystal molecules

20 Claims

1. A liquid crystal display device comprising:

25 an active matrix substrate having a pixel electrode;
a counter substrate having a counter electrode;
a vertical-alignment type liquid crystal layer provided between the active matrix substrate and the counter substrate; and
alignment sustaining layers provided between the active matrix substrate and the liquid crystal layer and between
30 the counter substrate and the liquid crystal layer, the alignment sustaining layers containing a polymerization product resulting from polymerization of a polyfunctional monomer, wherein,
at least one of the active matrix substrate and the counter substrate further includes an alignment film; and
the alignment film contains polyimide and a polymerization product resulting from polymerization of a polyfunctional monomer.

35 2. The liquid crystal display device of claim 1, wherein the polyfunctional monomer of the polymerization product contained in the alignment film has a plurality of vinyl groups.

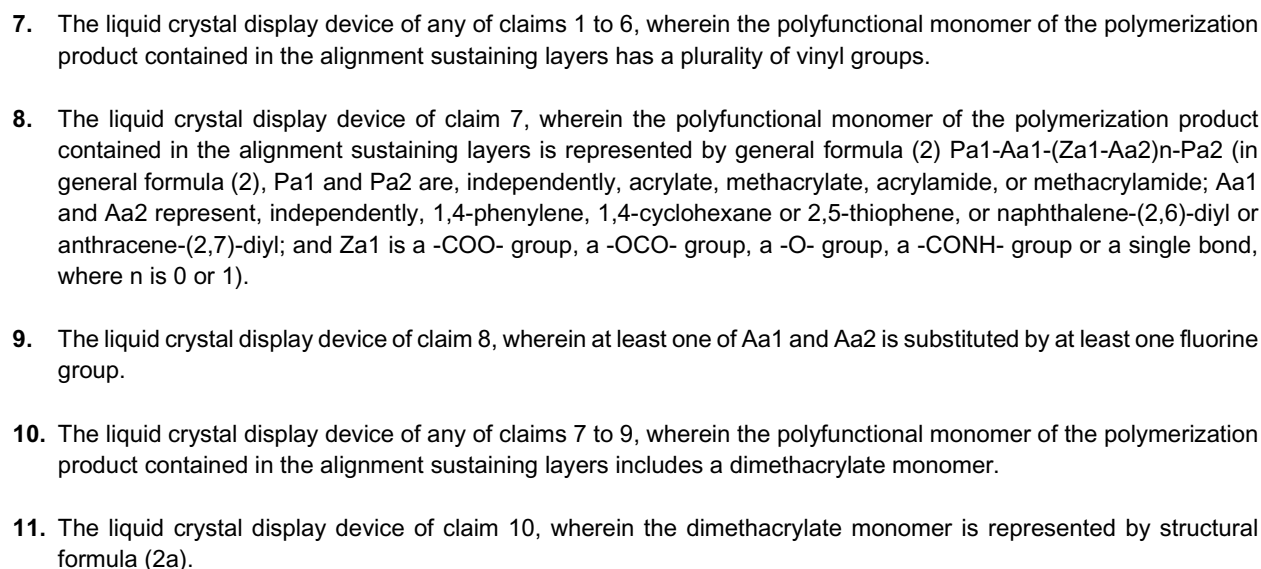
3. The liquid crystal display device of claim 1 or 2, wherein the polyfunctional monomer of the polymerization product contained in the alignment film is represented by general formula (1) $Pb1-Ab1-(Zb1-Ab2)_n-Pb2$ (in general formula
40 (1), Pb1 and Pb2 are, independently, acrylate, methacrylate, acrylamide, or methacrylamide; Ab1 and Ab2 represent, independently, 1,4-phenylene, 1,4-cyclohexane or 2,5-thiophene, or naphthalene-(2,6)-diyl or anthracene-(2,7)-diyl; and Zb1 is a -COO- group, a -OCO- group, a -O- group, a -CONH- group or a single bond, where n is 0 or 1).

4. The liquid crystal display device of claim 3, wherein at least one of Ab1 and Ab2 is substituted by at least one fluorine group.
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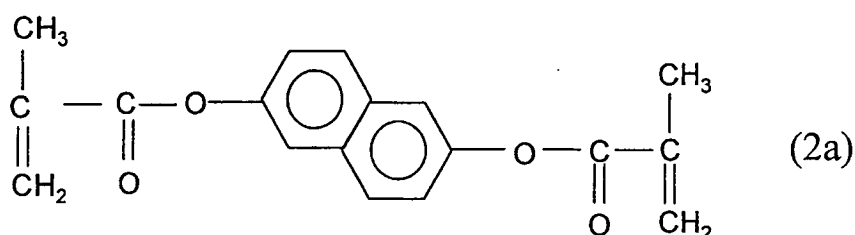
5. The liquid crystal display device of any of claims 1 to 4, wherein the polyfunctional monomer of the polymerization product contained in the alignment film includes a dimethacrylate monomer.

50 6. The liquid crystal display device of claim 5, wherein the dimethacrylate monomer is represented by structural formula (1a).

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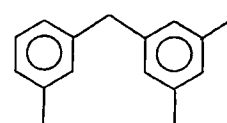
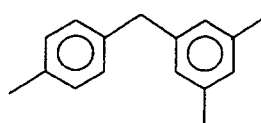
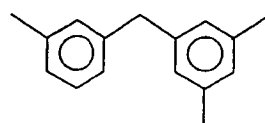
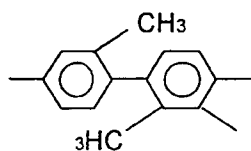
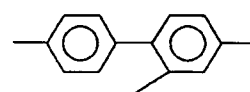
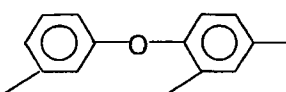
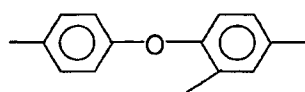
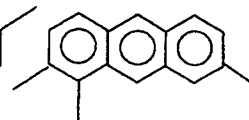
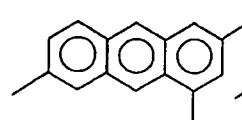
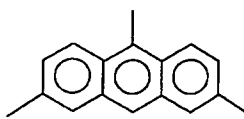
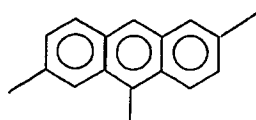
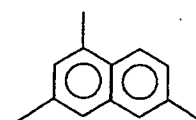
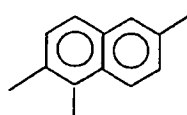
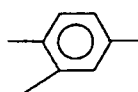
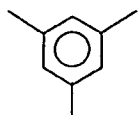
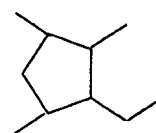
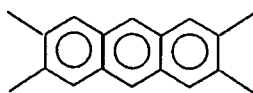
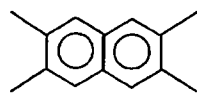
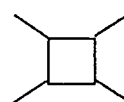
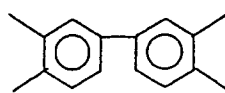
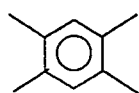
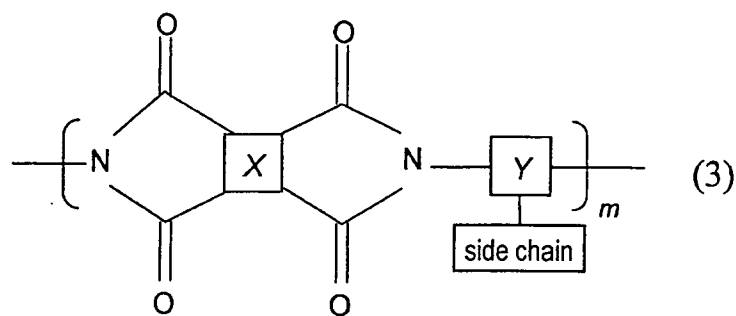


[Formula 2]



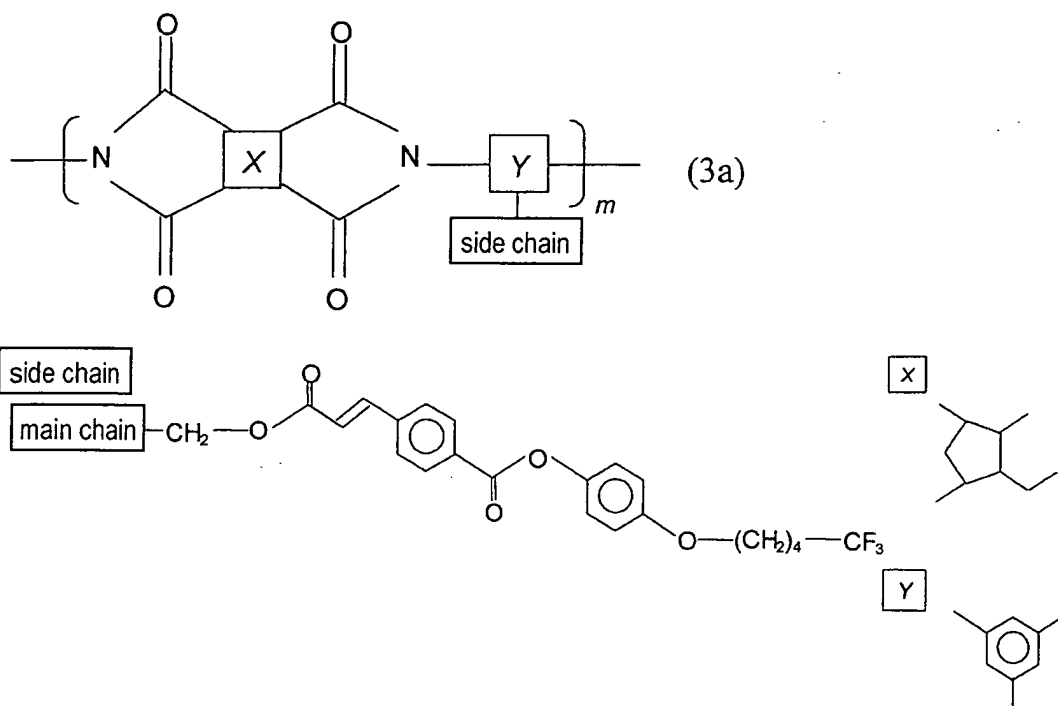
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[Formula 3]



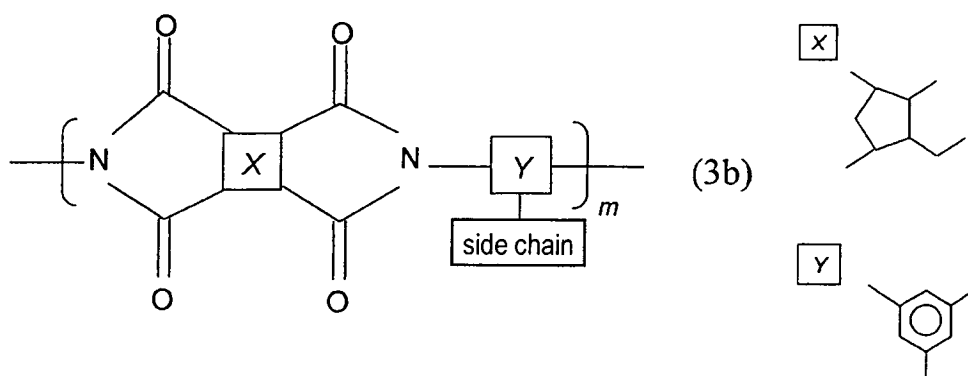
13. The liquid crystal display device of claim 12, wherein the polyimide has a photoreactive functional group.
14. The liquid crystal display device of claim 13, wherein the photoreactive functional group is one selected from the group consisting of a cinnamate group, a chalcone group, a tolan group, a coumarin group, and an azobenzene group.
15. The liquid crystal display device of claim 13 or 14, wherein the polyimide is a polyimide including a structure represented by structural formula (3a).

[Formula 4]



16. The liquid crystal display device of claim 12, wherein the polyimide has a vertical alignment group in a side chain thereof.
17. The liquid crystal display device of claim 16, wherein the polyimide is represented by structural formula (3b).

[Formula 5]



20 **18.** The liquid crystal display device of any of claims 1 to 17, wherein the alignment film regulates liquid crystal molecules in the liquid crystal layer so that the liquid crystal molecules are inclined with respect to a normal direction of a principal face of the alignment film in the absence of an applied voltage.

25 **19.** The liquid crystal display device of any of claims 1 to 18, wherein, the liquid crystal display device has a plurality of pixels; and in each of the plurality of pixels, the liquid crystal layer has a plurality of liquid crystal domains having respectively different reference alignment azimuths.

30 **20.** The liquid crystal display device of claim 19, wherein the plurality of liquid crystal domains are four liquid crystal domains.

21. A method of producing a liquid crystal display device, the method comprising the steps of:

35 forming an active matrix substrate and a counter substrate; and forming a liquid crystal layer between the active matrix substrate and the counter substrate, and forming alignment sustaining layers between the active matrix substrate and the liquid crystal layer and between the counter substrate and the liquid crystal layer, the alignment sustaining layers containing a polymerization product resulting from polymerization of a polyfunctional monomer, wherein the step of forming the active matrix substrate and the counter substrate comprises

40 a step of providing a first insulative substrate having a pixel electrode provided thereon and a second insulative substrate having a counter electrode provided thereon, and a step of forming an alignment film on at least one of the pixel electrode and the counter electrode, the alignment film containing polyimide and a polymerization product resulting from polymerization of a polyfunctional monomer.

45 **22.** The method of producing a liquid crystal display device of claim 21, wherein the step of forming the alignment film comprises:

50 a step of providing an alignment film material containing a precursor of the polyimide and the polyfunctional monomer; and a step of applying the alignment film material and performing a heat treatment, thereby imidizing the precursor of the polyimide to form the polyimide and forming a polymerization product resulting from polymerization of the polyfunctional monomer.

55 **23.** The method of producing a liquid crystal display device of claim 22, wherein the step of performing a heat treatment comprises a step of conducting a pre-bake and thereafter conducting a post-bake at a higher temperature than in the pre-bake.

- 24.** The method of producing a liquid crystal display device of any of claims 21 to 23, wherein the step of forming the alignment sustaining layers comprises a step of performing light irradiation.

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FIG. 1

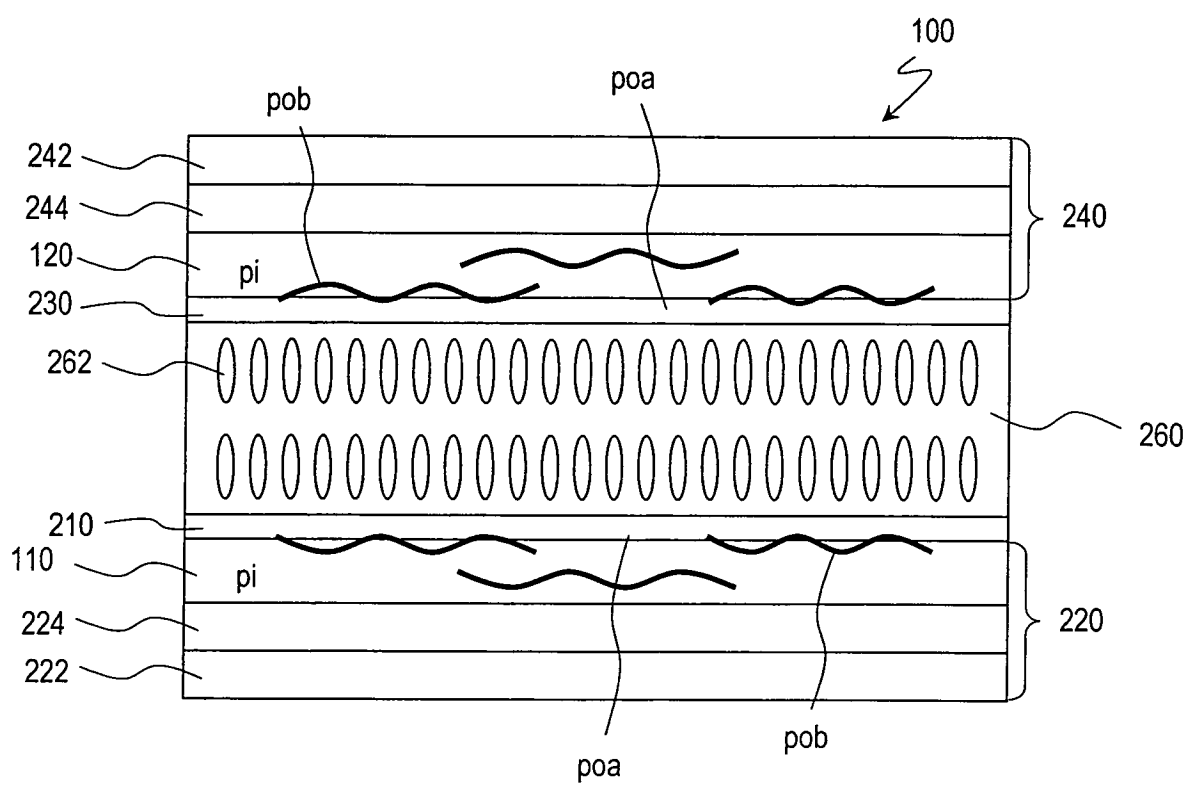


FIG. 2

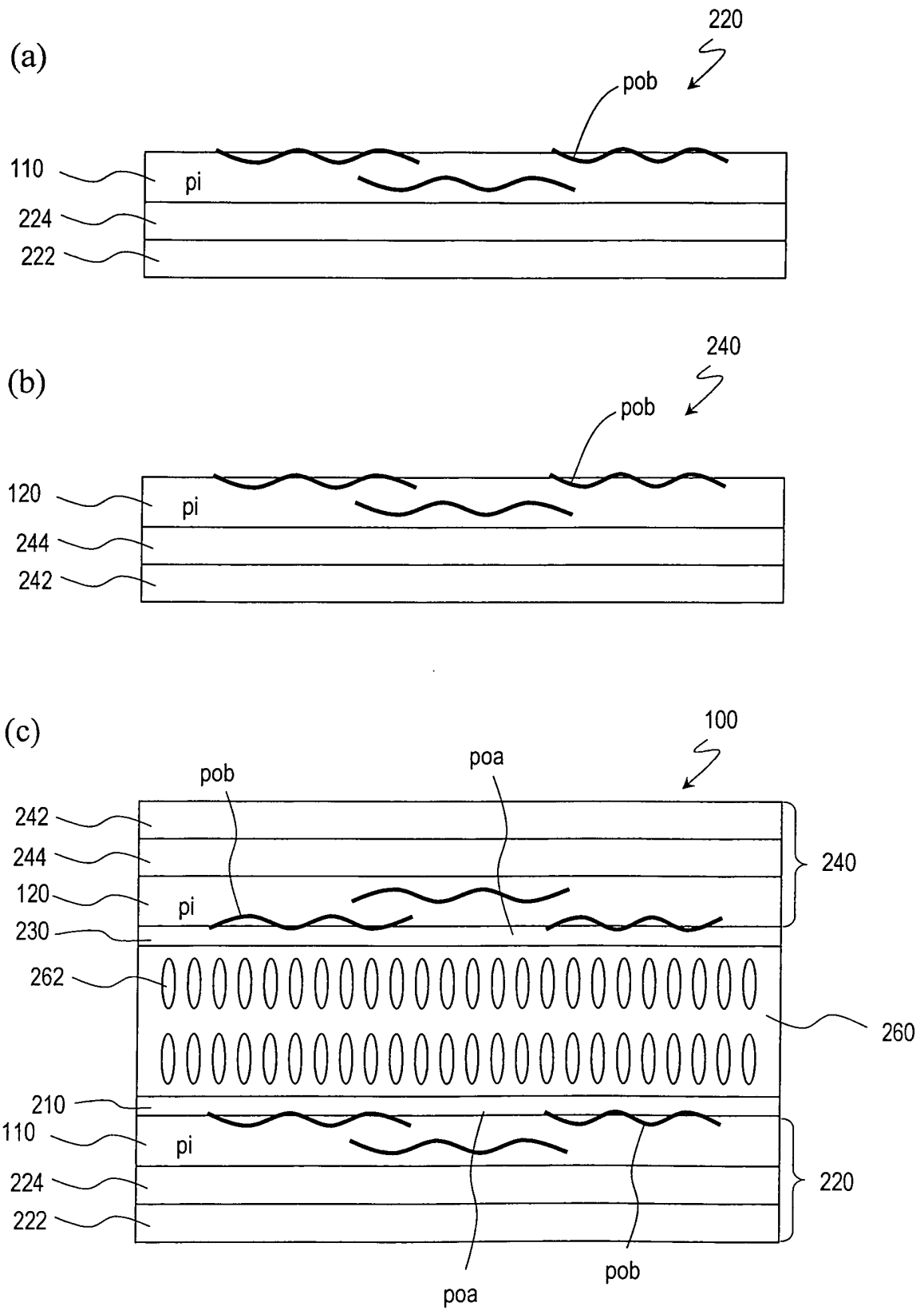


FIG. 3

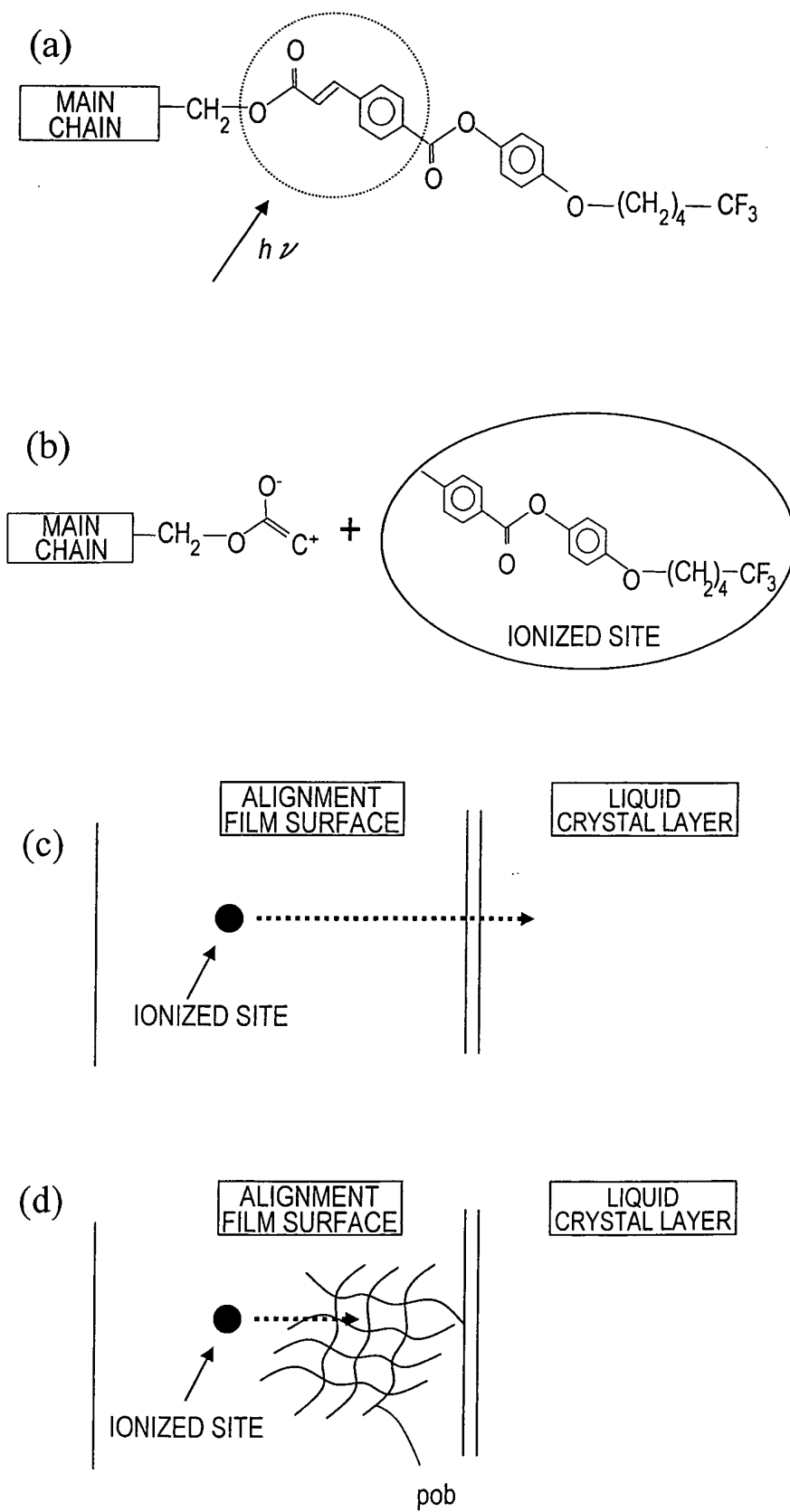


FIG. 4

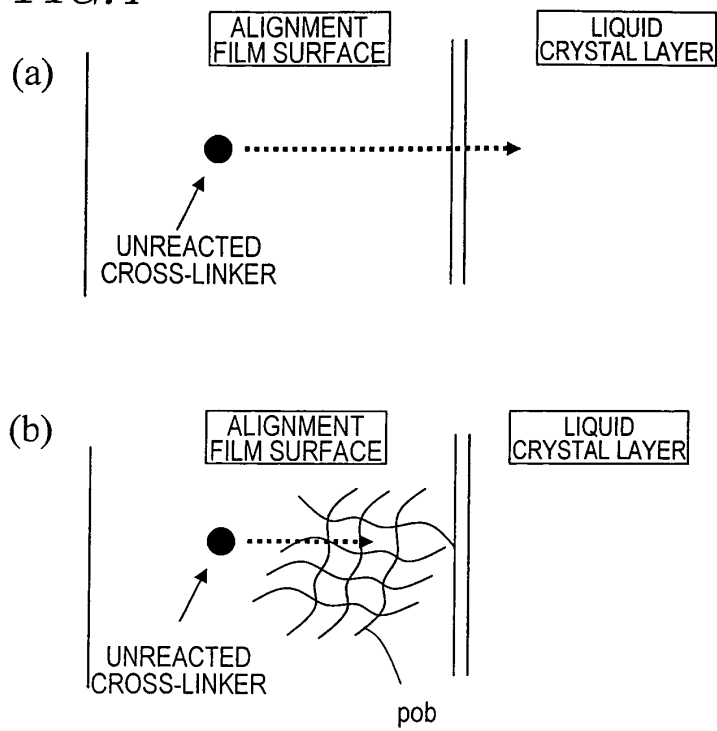


FIG. 5

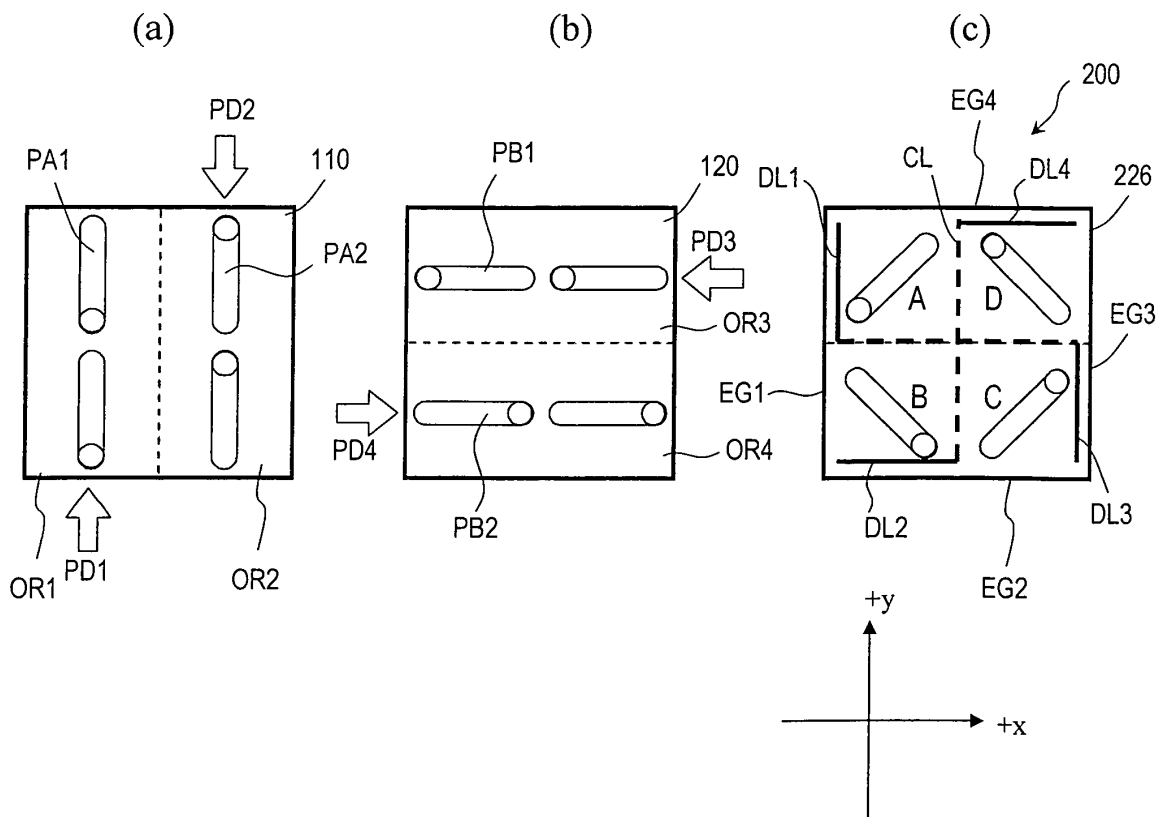


FIG. 6

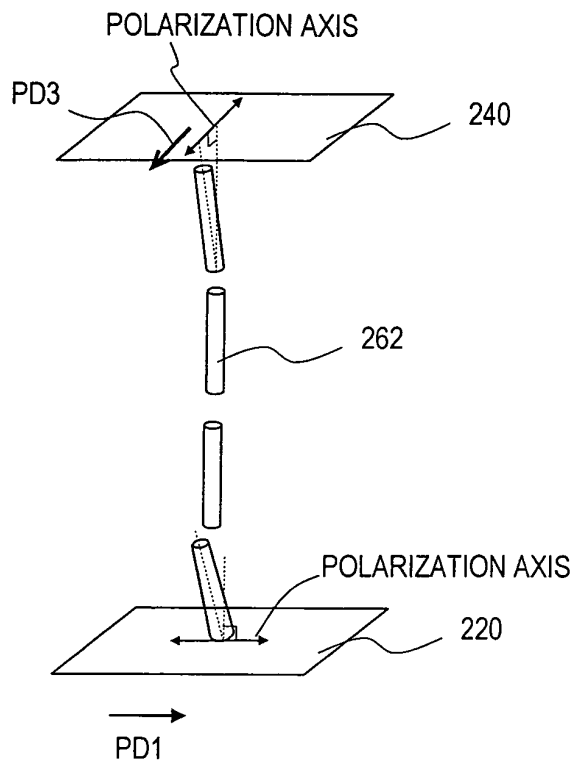


FIG. 7

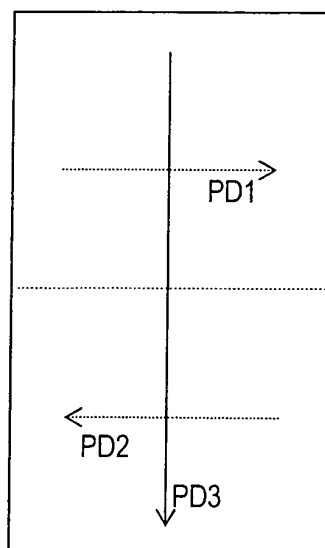


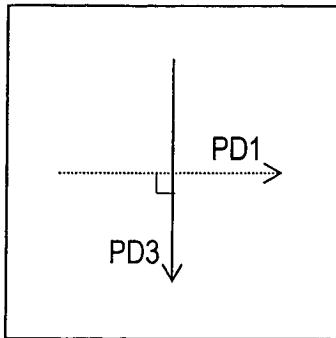
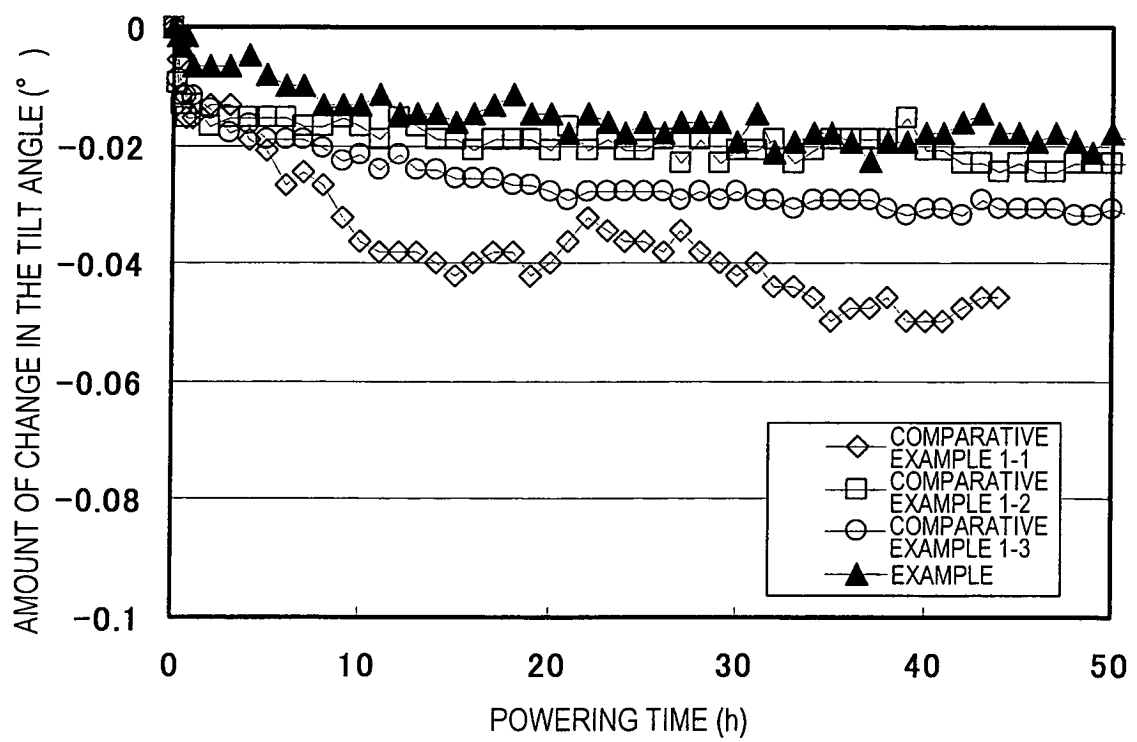
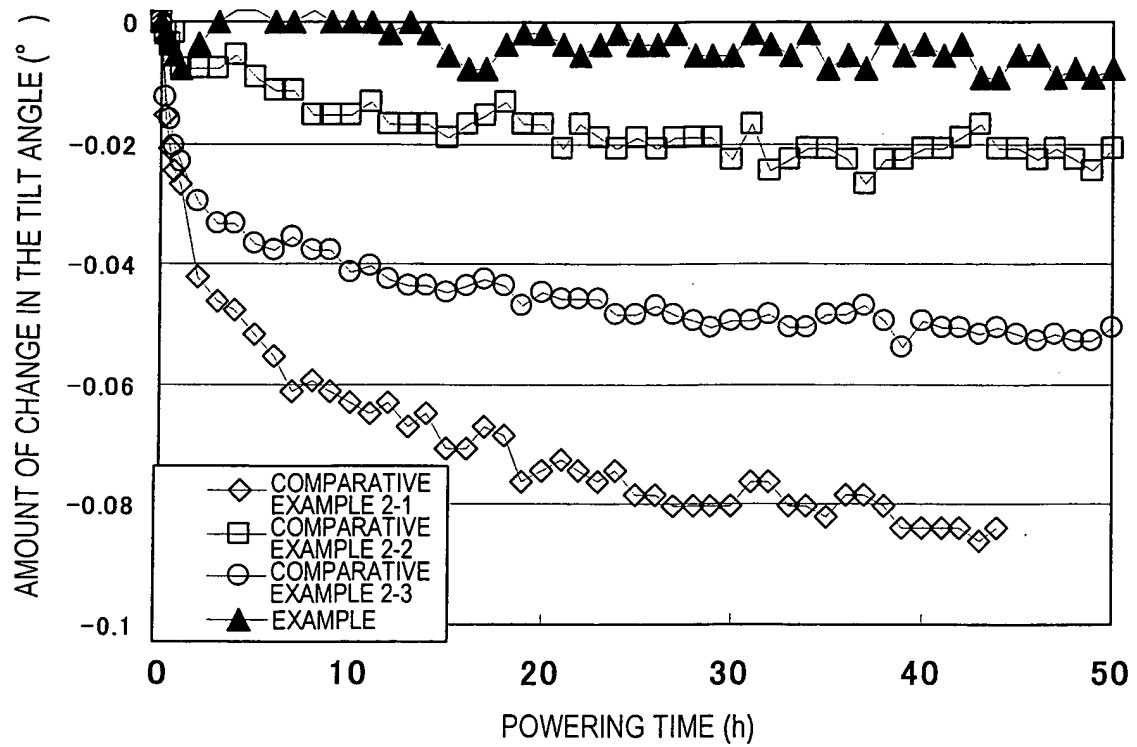
FIG. 8*FIG. 9*

FIG. 10



INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP2009/002931

A. CLASSIFICATION OF SUBJECT MATTER G02F1/1337(2006.01)i, C08F20/00(2006.01)i, C08F290/14(2006.01)i, C08L33/04(2006.01)i, C08L79/08(2006.01)i According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) G02F1/1337, C08F20/00, C08F290/14, C08L33/04, C08L79/08 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Jitsuyo Shinan Koho 1922-1996 Jitsuyo Shinan Toroku Koho 1996-2009 Kokai Jitsuyo Shinan Koho 1971-2009 Toroku Jitsuyo Shinan Koho 1994-2009 Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X A	JP 2001-4986 A (Optrex Corp.), 12 January, 2001 (12.01.01), Par. Nos. [0024], [0037] to [0041] (Family: none)	21-24 1-20
A	JP 2006-215184 A (JSR Corp.), 17 August, 2006 (17.08.06), Claims 1 to 4 (Family: none)	1-20
☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.		
* Special categories of cited documents: “A” document defining the general state of the art which is not considered to be of particular relevance “E” earlier application or patent but published on or after the international filing date “L” document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) “O” document referring to an oral disclosure, use, exhibition or other means “P” document published prior to the international filing date but later than the priority date claimed “T” later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention “X” document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone “Y” document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art “&” document member of the same patent family		
Date of the actual completion of the international search 14 September, 2009 (14.09.09)		Date of mailing of the international search report 29 September, 2009 (29.09.09)
Name and mailing address of the ISA/ Japanese Patent Office		Authorized officer
Facsimile No.		Telephone No.

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REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

- JP 11352486 A [0008]
- JP 2008076950 A [0008]
- JP 2008303176 A [0147]

专利名称(译)	液晶显示器及其制造方法		
公开(公告)号	EP2372440A1	公开(公告)日	2011-10-05
申请号	EP2009828740	申请日	2009-06-25
[标]申请(专利权)人(译)	夏普株式会社		
申请(专利权)人(译)	夏普株式会社		
当前申请(专利权)人(译)	夏普株式会社		
[标]发明人	MIZUSAKI MASANOBU NAKANISHI YOHEI		
发明人	MIZUSAKI, MASANOBU NAKANISHI, YOHEI		
IPC分类号	G02F1/1337 C08F20/00 C08F290/14 C08L33/04 C08L79/08 C08G73/10 C08G73/12 G02F1/1333		
CPC分类号	C08G73/1078 C08G73/12 C08L79/08 C08L79/085 G02F1/133723 G02F1/133753 G02F1/133788 G02F2001/133397 G02F2001/133715 G02F2001/133742 Y10T428/1005 Y10T428/1023		
优先权	2008303176 2008-11-27 JP		
其他公开文献	EP2372440A4 EP2372440B1		
外部链接	Espacenet		

摘要(译)

根据本发明的液晶显示装置 (100) 包括 : 具有像素电极 (224) 的有源矩阵基板 (220) ; 对向基板 (240) , 具有对电极 (244) ; 垂直取向型液晶层 (260) ; 和对准维持层 (210,230) 设置在有源矩阵基板 (220) 和液晶层 (260) 之间以及对置基板 (240) 和液晶层 (260) 之间。取向维持层 (210,230) 含有由多官能单体聚合得到的聚合产物 (poa) 。有源矩阵基板和対置基板中的至少一个还包括位于液晶层侧的取向膜 (110,120) , 包含聚酰亚胺 (pi) 的取向膜 (110,120) 和聚合产物 (pob) 由多官能单体聚合得到的。

FIG.1

